ORDIN Nr. 1301/500 din 11 iulie 2008 - Partea a II-a

pentru aprobarea protocoalelor terapeutice privind prescrierea medicamentelor aferente denumirilor comune internaţionale prevăzute în Lista cuprinzând denumirile comune internaţionale corespunzătoare medicamentelor de care beneficiază asiguraţii, cu sau fără contribuţie personală, pe bază de prescripţie medicală, în sistemul de asigurări sociale de sănătate, aprobată prin Hotărârea Guvernului nr. 720/2008

*Text în vigoare începând cu data de 21 august 2013*

*REALIZATOR: COMPANIA DE INFORMATICĂ NEAMŢ*

*Text actualizat prin produsul informatic legislativ LEX EXPERT în baza actelor normative modificatoare, publicate în Monitorul Oficial al României, Partea I, până la 21 august 2013.*

***Act de bază***

**#B**: *Ordinul ministrului sănătăţii publice şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 1301/500/2008*

***Acte modificatoare***

**#M1**: *Ordinul ministrului sănătăţii publice şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 1745/780/2008*

**#M2**: *Ordinul ministrului sănătăţii publice şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 1941/872/2008*

**#M3**: *Ordinul ministrului sănătăţii şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 461/477/2010*

**#M4**: *Ordinul ministrului sănătăţii şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 423/118/2012*

**#M5**: *Ordinul ministrului sănătăţii şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 961/536/2013*

*Modificările şi completările efectuate prin actele normative enumerate mai sus sunt scrise cu font italic. În faţa fiecărei modificări sau completări este indicat actul normativ care a efectuat modificarea sau completarea respectivă, în forma* ***#M1****,* ***#M2*** *etc.*

**#B**

Văzând Referatul de aprobare al Direcţiei generale politici, strategii şi managementul calităţii în sănătate din cadrul Ministerului Sănătăţii Publice nr. E.N. 7.547 din 11 iulie 2008 şi al directorului general al Casei Naţionale de Asigurări de Sănătate nr. D.G. 2.004 din 11 iulie 2008,

având în vedere prevederile:

- art. 406 alin. (1) lit. g) şi art. 243 din Legea nr. 95/2006 privind reforma în domeniul sănătăţii, cu modificările şi completările ulterioare;

- art. 4 din Hotărârea Guvernului nr. 720/2008 pentru aprobarea Listei cuprinzând denumirile comune internaţionale corespunzătoare medicamentelor de care beneficiază asiguraţii, cu sau fără contribuţie personală, pe bază de prescripţie medicală, în sistemul de asigurări sociale de sănătate;

în temeiul art. 281 alin. (2) din Legea nr. 95/2006, cu modificările şi completările ulterioare, al art. 7 alin. (4) din Hotărârea Guvernului nr. 862/2006\*) privind organizarea şi funcţionarea Ministerului Sănătăţii Publice, cu modificările şi completările ulterioare, şi al art. 17 alin. (5) din Statutul Casei Naţionale de Asigurări de Sănătate, aprobat prin Hotărârea Guvernului nr. 972/2006, cu modificările şi completările ulterioare,

ministrul sănătăţii publice şi preşedintele Casei Naţionale de Asigurări de Sănătate emit următorul ordin:

**#CIN**

***\*)*** *Hotărârea Guvernului nr. 862/2006 a fost abrogată. A se vedea Hotărârea Guvernului nr. 144/2010.*

**#B**

ART. 1

Se aprobă protocoalele terapeutice privind prescrierea medicamentelor aferente denumirilor comune internaţionale prevăzute în Lista cuprinzând denumirile comune internaţionale corespunzătoare medicamentelor de care beneficiază asiguraţii, cu sau fără contribuţie personală, pe bază de prescripţie medicală, în sistemul de asigurări sociale de sănătate, aprobată prin Hotărârea Guvernului nr. 720/2008, denumite în continuare protocoale terapeutice, prevăzute în anexele nr. 1 şi 2, care fac parte integrantă din prezentul ordin.

ART. 2

(1) În înţelesul prezentului ordin, termenii şi noţiunile folosite au următoarele semnificaţii:

a) prescriere limitată - prescrierea medicamentelor în cadrul sistemului de asigurări sociale de sănătate este limitată la indicaţia/indicaţiile medicală/medicale prevăzută/prevăzute în protocoalele terapeutice;

b) cod de restricţie - cod unic atribuit unei prescrieri limitate. Modalitatea de implementare a codurilor de restricţie se va stabili prin ordin al ministrului sănătăţii şi al preşedintelui Casei Naţionale de Asigurări de Sănătate.

(2) Condiţiile privind prescrierile limitate ale medicamentelor aferente denumirilor comune internaţionale prevăzute în Lista cuprinzând denumirile comune internaţionale corespunzătoare medicamentelor de care beneficiază asiguraţii, cu sau fără contribuţie personală, pe bază de prescripţie medicală, în sistemul de asigurări sociale de sănătate, aprobată prin Hotărârea Guvernului nr. 720/2008, şi codurile de restricţie ale acestora sunt prevăzute în anexa nr. 2.

ART. 3

(1) Protocoalele terapeutice constituie baza de prescriere şi monitorizare a medicamentelor care se acordă asiguraţilor pe bază de prescripţie medicală eliberată de medicii care sunt în relaţie contractuală cu casele de asigurări de sănătate.

(2) Respectarea schemelor terapeutice stabilite conform protocoalelor terapeutice prevăzute în anexele nr. 1 şi 2 este obligatorie pentru medicii aflaţi în relaţie contractuală cu casele de asigurări de sănătate.

**#M2**

*(3) Până la data de 31 decembrie 2008, medicii aflaţi în relaţie contractuală cu casele de asigurări de sănătate au obligaţia de a proceda la evaluarea bolnavilor pe care îi au în evidenţă, în vederea adaptării schemelor terapeutice în conformitate cu prevederile prezentului ordin.*

**#B**

ART. 4

Iniţierea şi continuarea tratamentului specific unei afecţiuni de către medicii aflaţi în relaţie contractuală cu casele de asigurări de sănătate se realizează cu respectarea prevederilor fiecărui protocol terapeutic.

ART. 5

Prescrierea, eliberarea şi decontarea medicamentelor corespunzătoare denumirilor comune internaţionale prevăzute în Lista cuprinzând denumirile comune internaţionale corespunzătoare medicamentelor de care beneficiază asiguraţii, cu sau fără contribuţie personală, pe bază de prescripţie medicală, în sistemul de asigurări sociale de sănătate, aprobată prin Hotărârea Guvernului nr. 720/2008, în baza protocoalelor terapeutice, se realizează după cum urmează:

- în conformitate cu prevederile Contractului-cadru privind condiţiile acordării asistenţei medicale în cadrul sistemului de asigurări sociale de sănătate pentru anul 2008, aprobat prin Hotărârea Guvernului nr. 324/2008, cu modificările şi completările ulterioare, şi ale Normelor metodologice de aplicare a Contractului-cadru privind condiţiile acordării asistenţei medicale în cadrul sistemului de asigurări sociale de sănătate pentru anul 2008, aprobate prin Ordinul ministrului sănătăţii publice şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 522/236/2008, pentru denumirile comune internaţionale cuprinse în lista menţionată mai sus, notate cu (\*\*) şi (\*\*\*) în sublista A, (\*\*), (\*\*\*) şi (\*\*\*\*) în sublista B, (\*\*), (\*\*\*) şi (\*\*\*\*) în secţiunea C1 a sublistei C şi (\*\*) în secţiunea C3 a sublistei C;

- în conformitate cu prevederile Hotărârii Guvernului nr. 357/2008 pentru aprobarea programelor naţionale de sănătate în anul 2008, cu modificările şi completările ulterioare, şi ale Normelor tehnice de realizare a programelor naţionale de sănătate în anul 2008, aprobate prin Ordinul ministrului sănătăţii publice şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 574/269/2008, pentru denumirile comune internaţionale cuprinse în lista menţionată mai sus, notate cu (\*\*), (\*\*\*) şi (\*\*\*\*) în secţiunea C2 a sublistei C.

ART. 6

Protocoalele terapeutice vor fi revizuite periodic.

ART. 7

Direcţiile de specialitate ale Ministerului Sănătăţii Publice, Casa Naţională de Asigurări de Sănătate, autorităţile de sănătate publică, casele de asigurări de sănătate şi furnizorii de servicii medicale vor duce la îndeplinire prevederile prezentului ordin.

ART. 8

Prezentul ordin se publică în Monitorul Oficial al României, Partea I.

ANEXA 1

Anexa nr. 1 se găseşte în Ordinul ministrului sănătăţii publice şi al preşedintelui Casei Naţionale de Asigurări de Sănătate nr. 1301/500/2008, partea I.

ANEXA 2

SUBLISTA A - MEDICAMENTE CU NIVEL DE COMPENSARE 90% DIN PREŢUL DE REFERINŢĂ

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| 1 |A02BA02| RANITIDINUM | |

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NOTĂ:

Terapia de eradicare a infecţiei cu Helicobacter pylori trebuie avută în vedere înaintea iniţierii tratamentului ulcerului peptic cu acest medicament. Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BA02 RANITIDINUM CAPS. 150 mg

ULCORAN 150 mg EUROPHARM SA

A02BA02 RANITIDINUM COMPR. 150 mg

RANITIDIN 150 mg ARENA GROUP SA

RANITIDINA 150 mg 150 mg MAGISTRA C&C

A02BA02 RANITIDINUM COMPR. EFF. 150 mg

DIGENEFF 150 mg 150 mg OZONE LABORATORIES LTD.

A02BA02 RANITIDINUM COMPR. FILM. 150 mg

GERTOCALM 150 mg FARAN LABORATORIES SA.

RANITIDIN 150 mg 150 mg AC HELCOR SRL

RANITIDINA 150 mg 150 mg FILDAS TRADING SRL

RANITIDINA ANTIBIOTICE 150 mg 150 mg ANTIBIOTICE SA

RANITIDINA LPH 150 mg 150 mg LABORMED PHARMA SA

ZANTAC 150 mg 150 mg GLAXO WELLCOME UK LTD.

A02BA02 RANITIDINUM SOL. INJ. 25 mg/ml

ARNETIN 25 mg/ml MEDOCHEMIE LTD.

ZANTAC SOLUŢIE INJECTABILĂ 25 mg/ml GLAXO WELLCOME UK LTD.

A02BA02 RANITIDINUM COMPR. FILM. 300 mg

RANITIDIN 300 mg 300 mg AC HELCOR SRL

RANITIDINA 300 mg 300 mg FILDAS TRADING SRL

RANITIDINA LPH 300 mg 300 mg LABORMED PHARMA SA

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| 2 |A02BA03| FAMOTIDINUM | |

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NOTĂ:

Terapia de eradicare a infecţiei cu Helicobacter pylori trebuie avută în vedere înaintea iniţierii tratamentului ulcerului peptic cu acest medicament. Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BA03 FAMOTIDINUM COMPR. FILM. 10 mg

FAMODAR ABR 10 mg DAR AL DAWA PHARMA S.R.L.

A02BA03 FAMOTIDINUM COMPR. FILM. 20 mg

FAMODAR 20 20 mg DAR AL DAWA PHARMA S.R.L.

FAMODIN 20 20 mg AC HELCOR PHARMA SRL

FAMOTAK 20 mg SEDICO IMPEX S.R.L.

QUAMATEL(R) 20 mg 20 mg GEDEON RICHTER PLC

A02BA03 FAMOTIDINUM COMPR. FILM. 20 mg

FAMODAR 20 20 mg DAR AL DAWA PHARMA S.R.L

FAMODIN 20 20 mg AC HELCOR PHARMA SRL

FAMOTAK 20 mg SEDICO IMPEX S.R.L.

QUAMATEL(R) 20 mg 20 mg GEDEON RICHTER PLC

A02BA03 FAMOTIDINUM LIOF. + SOLV. PT.

SOL. INJ. 20 mg

QUAMATEL(R) 20 mg 20 mg GEDEON RICHTER PLC

A02BA03 FAMOTIDINUM COMPR. FILM. 40 mg

FAMODAR 40 40 mg DAR AL DAWA PHARMA SRL

FAMODIN 40 40 mg AC HELCOR PHARMA SRL

FAMOTAK 40 mg SEDICO IMPEX S.R.L.

FAMOTIDINA ZENTIVA 40 mg 40 mg ZENTIVA S.A.

GASTROSIDIN 40 mg ECZACIBASI

PHARMACEUTICALS S.R.L.

QUAMATEL(R) 40 mg 40 mg GEDEON RICHTER PLC

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| 3 |A02BA04| NIZATIDINUM | |

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NOTĂ:

Terapia de eradicare a infecţiei cu Helicobacter pylori trebuie avută în vedere înaintea iniţierii tratamentului ulcerului peptic cu acest medicament. Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BA04 NIZATIDINUM CAPS. 150 mg

NIZATIDIN 150 mg 150 mg VIM SPECTRUM SRL

NIZATIDIN ALAROPHARM 150 mg 150 mg LAROPHARM S.R.L.

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| 4 |A02BC01| OMEPRAZOLUM | |

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NOTĂ:

Terapia de eradicare a infecţiei cu Helicobacter pylori trebuie avută în vedere înaintea iniţierii tratamentului ulcerului peptic cu acest medicament. Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BC01 OMEPRAZOLUM CAPS. GASTROREZ. 10 mg

ORTANOL(R) S 10 mg LEK PHARMACEUTICALS D.D.

Prescriere limitată: Boala de reflux gastro-esofagian.

Se aplică pentru toate denumirile comerciale şi formele farmaceutice ale concentraţiei de 10 mg.

A02BC01 OMEPRAZOLUM COMPR. FILM. GASTROREZ. 10 mg

LOSEC MUPS 10 mg 10 mg ASTRAZENECA AB

A02BC01 OMEPRAZOLUM CAPS. GASTROREZ. 20 mg

HELICID 20 20 mg ZENTIVA AS

OMEPRAZOL 20 mg 20 mg GEDEON RICHTER ROMANIA SA

OMEPRAZOL BIOFARM 20 mg 20 mg BIOFARM S.A.

OMEPRAZOL LPH 20 mg 20 mg LABORMED PHARMA SA

OMEPRAZOL TERAPIA 20 mg 20 mg TERAPIA S.A.

OMERAN 20 20 mg EUROPHARM SA

OMEZ 20 mg DR. REDDY'S LABORATORIES

ORTANOL(R) 20 mg 20 mg LEK PHARMACEUTICALS D.D.

ULTOP 20 mg KRKA D.D.

Prescriere limitată: Boala de reflux gastro-esofagian

Prescriere limitată: Sclerodermia esofagului.

Prescriere limitată: Sindromul Zollinger - Ellison

Prescriere limitată: Tratamentul ulcerului gastric şi duodenal.

Prescriere limitată: Profilaxia ulcerului gastro - duodenal la pacienţii în tratament cu AINS pe termen lung şi care prezintă factori de risc gastrointestinal.

NOTĂ:

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiilor de 20 mg.

A02BC01 OMEPRAZOLUM CAPS. ENTER. 20 mg

RISEK 20 mg GULF PHARMACEUTICAL

INDUSTRIES S.R.L.

A02BC01 OMEPRAZOLUM COMPR. FILM. GASTROREZ. 20 mg

LOSEC MUPS 20 mg 20 mg ASTRAZENECA AB

OMEDAR(R) 20 mg DAR AL DAWA PHARMA S.R.L.

A02BC01 OMEPRAZOLUM CAPS. GASTROREZ. 40 mg

OMERAN 40 40 mg EUROPHARM SA

ORTANOL(R) 40 mg LEK PHARMACEUTICALS D.D.

ULTOP 40 mg KRKA D.D. NOVO MESTO

Prescriere limitată: Boala de reflux gastro-esofagian

Prescriere limitată: Sclerodermia esofagului.

Prescriere limitată: Sindromul Zollinger-Ellison

Prescriere limitată: Tratamentul ulcerului gastric şi duodenal.

NOTĂ:

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiilor de 40 mg.

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| 5 |A02BC03| LANSOPRAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BC03 LANSOPRAZOLUM CAPS. GASTROREZ. 15 mg

LEVANT 15 mg 15 mg RANBAXY UK LTD.

Prescriere limitată: Boala de reflux gastro-esofagian

A02BC03 LANSOPRAZOLUM CAPS. GASTROREZ. 30 mg

LANZAP 30 mg 30 mg DR. REDDY'S LABORATORIES

LANZUL 30 mg KRKA D.D. NOVO MESTO

LEVANT 30 mg 30 mg RANBAXY UK LTD.

Prescriere limitată: Boala de reflux gastro-esofagian

Prescriere limitată: Sindromul Zollinger-Ellison.

Prescriere limitată: Tratamentul ulcerului gastric şi duodenal.

Prescriere limitată: Profilaxia ulcerului gastro-duodenal la pacienţii în tratament cu AINS pe termen lung şi care prezintă factori de risc gastrointestinal.

NOTĂ:

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiei de 30 mg.

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| 6 |A03AA04| MEBEVERINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A03AA04 MEBEVERINUM CAPS. ELIB. PREL. 200 mg

DUSPATALIN(R) 200 mg 200 mg SOLVAY PHARMACEUTICALS BV

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| 7 |A03AA05| TRIMEBUTINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A03AA05 TRIMEBUTINUM COMPR. 100 mg

COLOBUTINE(R) 100 mg LAB. FOURNIER SA

COLPERIN 100 mg 100 mg GEDEON RICHTER

ROMANIA S.A.

PROMEBUTIN(R) 100 mg 100 mg TERAPIA SA

TRIMEBUTIN 100 100 mg MAGISTRA C & C

A03AA05 TRIMEBUTINUM COMPR. FILM. 100 mg

DEBRIDAT 100 mg 100 mg PFIZER EUROPE MA EEIG

A03AA05 TRIMEBUTINUM GRAN. PT. SUSP. ORALĂ 24 mg/5 ml

DEBRIDAT 24 mg/5 ml PFIZER EUROPE MA EEIG

A03AA05 TRIMEBUTINUM COMPR. FILM. ELIB. 300 mg

PREL.

IBUTIN(R) 300 mg 300 mg ZENTIVA SA

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| 8 |A03AD01| PAPAVERINI HYDROCHLORIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A03AD01 PAPAVERINI COMPR. 100 mg

HYDROCHLORIDUM

PAPAVERINA 100 mg 100 mg SINTOFARM SA

A03AD01 PAPAVERINI COMPR. 200 mg

HYDROCHLORIDUM

PAPAVERINA 200 mg FARMACOM SA

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| 9 |A03FA01| METOCLOPRAMIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A03FA01 METOCLOPRAMIDUM COMPR. 10 mg

METOCLOPRAMID 10 mg SLAVIA PHARM SRL

METOCLOPRAMID 10 mg 10 mg TERAPIA SA

N-METOCLOPRAMID 10 mg MEDUMAN SA

A03FA01 METOCLOPRAMIDUM SIROP 1 mg/5 ml

METOCLOPRAMID BIOFARM 1 mg/5 ml BIOFARM S.A.

1 mg/5 ml

A03FA01 METOCLOPRAMIDUM SOL. INJ. 5 mg/ml

METOCLOPRAMID 10 mg 5 mg/ml TERAPIA SA

A03FA01 METOCLOPRAMIDUM PIC. ORALE - SOL. 7 mg/ml

METOCLOPRAMID 7 mg/ml BIOFARM SA

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| 10 |A03FA03| DOMPERIDONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A03FA03 DOMPERIDONUM COMPR. FILM. 10 mg

MOTILIUM 10 mg JANSSEN PHARMACEUTICA NV

MOTILIUM 10 mg 10 mg TERAPIA SA

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| 11 |A05AA02| ACIDUM URSODEOXYCHOLICUM\* | |

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NOTĂ:

Nu se va prescrie în regim compensat pentru tratamentul litiazei biliare.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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A05AA02 ACIDUM CAPS. 250 mg

URSODEOXYCHOLICUM

URSOFALK(R) 250 mg DR. FALK PHARMA GMBH

URSOSAN 250 mg PRO. MED. CS PRAHA AS

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| 12 |A06AD11| LACTULOSUM | |

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Prescriere limitată: Tratamentul encefalopatiei cronice porto-sistemice

Tratamentul constipaţiei la pacienţii neoplazici.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A06AD11 LACTULOSUM LICHID ORAL 66.7%

DUPHALAC(R) 66.7% SOLVAY PHARMACEUTICALS BV

A06AD11 LACTULOSUM SIROP 66.7%

LACTULOSE AL SIROP 66.7% ALIUD PHARMA GMBH & CO.KG

A06AD11 LACTULOSUM SOL. ORALĂ 670 mg/ml

LAEVOLAC 670 mg/ml 670 mg/ml FRESENIUS KABI AUSTRIA

GMBH

A06AD11 LACTULOSUM SIROP 66.7%

LACTULOSE AL SIROP 66.7% ALIUD PHARMA GMBH & CO.KG

A06AD11 LACTULOSUM SOL. ORALĂ 670 mg/ml

LAEVOLAC 670 mg/ml 670 mg/ml FRESENIUS KABI AUSTRIA

GMBH

A06AD11 LACTULOSUM SIROP 66.7%

LACTULOSE AL SIROP 66.7% ALIUD PHARMA GMBH & CO.KG

A06AD11 LACTULOSUM SOL. ORALĂ 670 mg/ml

LAEVOLAC 670 mg/ml 670 mg/ml FRESENIUS KABI AUSTRIA

GMBH

A06AD11 LACTULOSUM LICHID ORAL 66.7%

DUPHALAC(R) 66.7% SOLVAY PHARMACEUTICALS BV

A06AD11 LACTULOSUM SIROP 65%

LACTULOSE 65% E.I.P.I.CO. MED S.R.L.

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| 13 |A07AA02| NYSTATINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A07AA02 NYSTATINUM COMPR. FILM. 500.000 ui

NISTATINA 500.000 ui ANTIBIOTICE SA

A07AA02 NYSTATINUM COMPR. FILM. 500.000 ui

STAMICIN(R) 500.000 U.I. 500.000 ui ZENTIVA SA

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| 14 |A07EC01| SULFASALAZINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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A07EC01 SULFASALAZINUM COMPR. FILM. 500 mg

SALAZIDIN 500 mg AC HELCOR SRL

A07EC01 SULFASALAZINUM COMPR. FILM. GASTROREZ. 500 mg

SULFASALAZIN EN 500 mg KRKA D.D. NOVO MESTO

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| 15 |A07EC02| MESALAZINUM\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A07EC02 MESALAZINUM SUPOZ. 1 g

PENTASA 1 g FERRING A/S

Prescriere limitată: Episod acut de proctită ulcerativă forma moderată.

NOTĂ:

A nu se administra în tratamentul bolii Crohn.

A07EC02 MESALAZINUM COMPR. GASTROREZ. 250 mg

SALOFALK 250 mg COMPRIMATE 250 mg DR. FALK PHARMA GMBH

GASTROREZISTENTE

Cod restricţie 1708: Colită ulcerativă asociată cu hipersensibilitate la sulfonamide

Cod restricţie 1709: Colită ulcerativă asociată cu intoleranţă la sulfasalazinum

Cod restricţie 2268: Boala Crohn în cazul în care există hipersensibilitate la sulfonamide

Cod restricţie 2269: Boala Crohn în cazul în care există intoleranţă la sulfasalazinum

A07EC02 MESALAZINUM SUPOZ. 250 mg

SALOFALK(R) 250 mg 250 mg DR. FALK PHARMA GMBH

Prescriere limitată: Episod acut de proctită ulcerativă forma moderată

A07EC02 MESALAZINUM SUSP. RECTALĂ 4 g/60 ml

SALOFALK 4 g/60 ml 4 g/60 ml DR. FALK PHARMA GMBH

Cod restricţie 1707: Episod acut de colită ulcerativă forma moderată

A07EC02 MESALAZINUM COMPR. ELIB. PREL. 500 mg

PENTASA 500 mg FERRING A/S

Cod restricţie 1708: Colită ulcerativă asociată cu hipersensibilitate la sulfonamide

Cod restricţie 1709: Colită ulcerativă asociată cu intoleranţă la sulfasalazinum

Cod restricţie 2268: Boala Crohn în cazul în care există hipersensibilitate la sulfonamide

Cod restricţie 2269: Boala Crohn în cazul în care există intoleranţă la sulfasalazinum

A07EC02 MESALAZINUM COMPR. GASTROREZ. 500 mg

SALOFALK 500 mg COMPRIMATE 500 mg DR. FALK PHARMA GMBH

GASTROREZISTENTE

Cod restricţie 1708: Colită ulcerativă asociată cu hipersensibilitate la sulfonamide

Cod restricţie 1709: Colită ulcerativă asociată cu intoleranţă la sulfasalazinum

Cod restricţie 2268: Boala Crohn în cazul în care există hipersensibilitate la sulfonamide

Cod restricţie 2269: Boala Crohn în cazul în care există intoleranţă la sulfasalazinum

A07EC02 MESALAZINUM SUPOZ. 500 mg

SALOFALK(R) 500 mg 500 mg DR. FALK PHARMA GMBH

Prescriere limitată: Episod acut de proctită ulcerativă forma moderată

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| 16 |B01AA07| ACENOCUMAROLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AA07 ACENOCUMAROLUM COMPR. 2 mg

TROMBOSTOP 2 mg 2 mg TERAPIA SA

B01AA07 ACENOCUMAROLUM COMPR. 4 mg

SINTROM(R) 4 mg NOVARTIS PHARMA GMBH

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| 17 |B01AC05| TICLOPIDINUM (1) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AC05 TICLOPIDINUM COMPR. FILM. 250 mg

IPATON 250 mg EGIS PHARMACEUTICALS PLC

TICLID(R) 250 mg 250 mg SANOFI-SYNTHELABO FRANCE

TICLODIN 250 mg AC HELCOR SRL

TICLOPIDIN SANDOZ 250 mg HEXAL AG

Cod restricţie 1719: Prevenţia recurentei accidentului vascular ischiemic sau a accidentului ischemic tranzitor la pacienţii cu istoric de episoade ischiemice cerebrovasculare în timpul terapiei cu doze reduse de aspirină

Cod restricţie 1720: Prevenţia recurenţei accidentului vascular ischiemic sau a accidentului ischemic tranzitor la pacienţii la care terapia cu doze reduse de aspirină prezintă un risc major (inacceptabil) de sângerare gastrointestinală;

Cod restricţie 1721: Prevenţia recurenţei accidentului vascular ischiemic sau a accidentului ischemic tranzitor la pacienţii cu istoric de reacţie anafilactică, urticarie sau astm bronşic în decurs de 4 ore de la administrarea de aspirină, alţi salicilaţi sau AINS

Neutropenia severă este un efect advers comun în primele luni de terapie. Monitorizarea hematologică se impune la începutul tratamentului şi apoi la fiecare două săptămâni în primele patru luni de tratament.

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| 18 |B02BA01| PHYTOMENADIONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B02BA01 PHYTOMENADIONUM SOL. INJ. 10 mg/ml

FITOMENADION 10 mg/ml 10 mg/ml TERAPIA SA

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| 19 |B03AA07| FERROSI SULFAS | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B03AA07 FERROSI SULFAS COMPR. ELIB. PREL. 105 mg

FERROGRADUMET(R) 105 mg TEOFARMA SRL

B03AA07 FERROSI SULFAS DRAJ. ELIB. PREL. 80 mg

TARDYFERON 80 mg 80 mg LAB. PIERRE FABRE

B03AA07 FERROSI SULFAS COMPR. FILM. ELIB.

PREL.

FERRO-GRADUMET POLIPHARMA

INDUSTRIES S.R.L.

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| 20 |B03AB05| COMPLEX DE HIDROXID DE FER (III) | |

| | | POLIMALTOZAT | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B03AB05 COMPLEX DE HIDROXID COMPR. MAST. 100 mg

DE FER (III)

POLIMALTOZAT

MALTOFER 100 mg VIFOR FRANCE SA

B03AB05 COMPLEX DE HIDROXID SIROP 10 mg/ml

DE FER (III)

POLIMALTOZAT

FERRUM HAUSMANN(R) 10 mg/ml VIFOR FRANCE SA

B03AB05 COMPLEX DE HIDROXID SIROP 50 mg/5 ml

DE FER (III)

POLIMALTOZAT

FERGLUROM 50 mg/5 ml 50 mg/5 ml BIOFARM S.A.

B03AB05 COMPLEX DE HIDROXID PICĂTURI ORALE - SOL. 5%

DE FER (III)

POLIMALTOZAT

PHARMA-FERRUM(R) 5% TERAPIA SA

B03AB05 COMPLEX DE HIDROXID PICĂTURI ORALE - SOL. 50 mg/ml

DE FER (III)

POLIMALTOZAT

FERRUM HAUSMANN(R) 50 mg/ml VIFOR FRANCE SA

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| 21 |B03AEN1| COMBINAŢII (FERROSI SULFAS + ACIDUM | |

| | | ASCORBICUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B03AEN1 COMBINAŢII (FERROSI COMPR. FILM.

SULFAS + ACIDUM

ASCORBICUM)

SORBIFER DURULES EGIS PHARMACEUTICALS LTD.

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| 22 |B03BB01| ACIDUM FOLICUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B03BB01 ACIDUM FOLICUM COMPR. FILM. 5 mg

ACIFOL 5 mg 5 mg ZENTIVA SA

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| 23 |C01AA05| DIGOXINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01AA05 DIGOXINUM SOL. ORALĂ 0.05 mg/ml

LANOXIN SOLUŢIE ORALĂ 0.05 mg/ml THE WELLCOME

FOUNDATION LTD.

C01AA05 DIGOXINUM COMPR. 0.25 mg

DIGOXIN 0,25 mg 0,25 mg ZENTIVA SA

C01AA05 DIGOXINUM SOL. INJ. 0.5 mg/ml

DIGOXIN 0,5 mg/2 ml 0,5 mg/ml ZENTIVA SA

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| 24 |C01BA01| CHINIDINI SULFAS | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01BA01 CHINIDINI SULFAS COMPR. 200 mg

CHINIDINA LAROPHARM 200 mg 200 mg LAROPHARM SRL

CHINIDINA SULFAT 200 mg 200 mg ARENA GROUP SA

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| 25 |C01BC03| PROPAFENONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01BC03 PROPAFENONUM COMPR. 150 mg

PROPAFENONA 150 mg 150 mg ARENA GROUP SA

C01BC03 PROPAFENONUM COMPR. FILM. 150 mg

PROPAFENON AL 150 150 mg ALIUD(R) PHARMA GMBH &

CO.KG

PROPAFENON SANDOZ 150 mg 150 mg HEXAL AG

RYTMONORM(R) 150 mg ABBOTT GMBH & CO.KG

C01BC03 PROPAFENONUM SOL. INJ. 70 mg/20 ml

RYTMONORM 70 mg/20 ml ABBOTT GMBH & CO.KG

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| 26 |C01BD01| AMIODARONUM | |

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Prescriere limitată: Tratamentul tulburărilor de ritm severe care nu răspund

la alte terapii sau când alte antiaritmice nu pot fi

folosite: a) tahiaritmii asociate sindromului Wolff -

Parkinson - White; b) flutter/fibrilaţie atrială,

atunci când alte antiaritmice nu pot fi folosite; c)

toate tahiaritmiile paroxistice, incluzând tahicardii

supraventriculare, tahicardii ventriculare şi nodale,

fibrilaţie ventriculară, atunci când alte antiaritmice

nu pot fi folosite.

Există dovezi că amiodarona poate produce toxicitate frecventă şi potenţial severă. Se recomandă monitorizarea periodică a funcţiilor hepatice şi tiroidiene.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01BD01 AMIODARONUM COMPR. 200 mg

AMIODARONA 200 mg 200 mg ARENA GROUP SA

AMIODARONA LPH 200 mg 200 mg LABORMED PHARMA SA

DARITMIN(R) 200 mg 200 mg GEDEON RICHTER ROMANIA SA

SEDACORON(R) 200 mg EBEWE PHARMA GMBH NFG. KG

C01BD01 AMIODARONUM COMPR. DIVIZ. 200 mg

CORDARONE 200 mg 200 mg SANOFI-AVENTIS FRANCE

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| 27 |C01DA02| NITROGLYCERINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01DA02 NITROGLYCERINUM SPRAY SUBLINGUAL 0.4 mg/doza

NITROMINT 0.4 mg/doza EGIS PHARMACEUTICALS

P.L.C.

NOTĂ:

Sprayul nu trebuie inhalat.

C01DA02 NITROGLYCERINUM COMPR. SUBLING. 0.5 mg

NITROGLICERINA 0,5 mg 0.5 mg ZENTIVA SA

C01DA02 NITROGLYCERINUM COMPR. ELIB. PREL. 2.6 mg

NITROMINT 2,6 mg 2.6 mg EGIS PHARMACEUTICALS LTD.

C01DA02 NITROGLYCERINUM SIST. TERAP. TRANSDERM. 25 mg

NITRODERM(R) TTS 5 25 mg NOVARTIS PHARMA GMBH

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| 28 |C01DA08| ISOSORBIDI DINITRAS | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01DA08 ISOSORBIDI DINITRAS COMPR. 10 mg

ISOSORBIDE DINITRATE 10 mg 10 mg E.I.P.I.CO. MED S.R.L.

C01DA08 ISOSORBIDI DINITRAS CAPS. ELIB. PREL. 20 mg

DINITER SR 20 mg 20 mg TERAPIA SA

C01DA08 ISOSORBIDI DINITRAS COMPR. ELIB. PREL. 20 mg

ISODINIT(R) RETARD 20 mg BALKAN PHARMA DUPNITZA AD

C01DA08 ISOSORBIDI DINITRAS CAPS. ELIB. PREL. 40 mg

DINITER SR 40 mg 40 mg TERAPIA SA

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| 29 |C01DA14| ISOSORBIDI MONONITRAS | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01DA14 ISOSORBIDI MONONITRAS CAPS. ELIB. PREL. 40 mg

OLICARD 40 mg RETARD 40 mg SOLVAY PHARMACEUTICALS

GMBH

C01DA14 ISOSORBIDI MONONITRAS CAPS. ELIB. PREL. 60 mg

OLICARD 60 mg RETARD 60 mg SOLVAY PHARMACEUTICALS

GMBH

C01DA14 ISOSORBIDI MONONITRAS COMPR. FILM. ELIB. 60 mg

PREL.

MONONITRON(R) EP 60 mg 60 mg ZENTIVA S.A.

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| 30 |C01EA01| ALPROSTADILUM\*\*\* | Protocol: C002I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01EA01 ALPROSTADILUM CONC. PT. SOL. PERF. 20 micrograme

ALPROSTADIL "PINT" 20 micrograme PINT-PHARMA GMBH

20 micrograme

C01EA01 ALPROSTADILUM LIOF. PT. SOL. PERF. 20 micrograme

VASAPROSTAN 20 20 micrograme SCHWARZ PHARMA AG

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| 31 |C01EB15| TRIMETAZIDINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01EB15 TRIMETAZIDINUM CAPS. 20 mg

TRIMETAZIDIN 20 mg 20 mg VIM SPECTRUM SRL

C01EB15 TRIMETAZIDINUM COMPR. FILM. 20 mg

DILATAN 20 mg 20 mg TERAPIA SA

MODUXIN(R) 20 mg 20 mg GEDEON RICHTER ROMÂNIA SA

PREDOZONE 20 mg 20 mg OZONE LABORATORIES LTD.

PREDUCTAL(R) 20 mg LES LAB. SERVIER IND.

TRIMETAZIDINA LPH(R) 20 mg 20 mg LABORMED PHARMA SA

TRIVEDON 20 20 mg CIPLA (UK) LIMITED

C01EB15 TRIMETAZIDINUM DRAJ. 20 mg

TRIMETAZIDINA 20 mg 20 mg TERAPIA SA

C01EB15 TRIMETAZIDINUM COMPR. FILM. ELIB. 35 mg

MODIF.

DILATAN MR 35 mg 35 mg TERAPIA SA

PREDUCTAL MR 35 mg 35 mg LES LAB. SERVIER IND.

TRIMETAZIDINA LPH(R) 35 mg 35 mg LABORMED PHARMA SA

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| 32 |C02AB01| METHYLDOPUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C02AB01 METHYLDOPUM COMPR. 250 mg

DOPEGYT 250 mg EGIS PHARMACEUTICALS PLC

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| 33 |C02AC01| CLONIDINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C02AC01 CLONIDINUM COMPR. 0.15 mg

CLONIDINA 0,15 mg 0.15 mg ARENA GROUP SA

CLONIDINA SINTOFARM 0,15 mg 0.15 mg SINTOFARM SA

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| 34 |C02CA04| DOXAZOSINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C02CA04 DOXAZOSINUM COMPR. 1 mg

DOXAZOSIN 1 MEDOCHEMIE 1 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 1 1 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 1 mg KRKA D.D. NOVO MESTO

MAGUROL 1 mg MEDOCHEMIE LTD.

C02CA04 DOXAZOSINUM COMPR. 2 mg

DOXAZOSIN 2 MEDOCHEMIE 2 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 2 2 mg ALIUD(R) PHARMA GMBH &

CO.KG

DOXAZOSIN SANDOZ 2 mg 2 mg HEXAL AG

KAMIREN 2 mg KRKA, D.D. NOVO MESTO

MAGUROL 2 mg MEDOCHEMIE LTD.

C02CA04 DOXAZOSINUM COMPR. 4 mg

DOXAZOSIN 4 MEDOCHEMIE 4 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 4 4 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 4 mg KRKA D.D. NOVO MESTO

MAGUROL 4 mg MEDOCHEMIE LTD.

C02CA04 DOXAZOSINUM COMPR. ELIB. MODIF. 4 mg

CARDURA XL 4 mg 4 mg PFIZER H.C.P. CORPORATION

C02CA04 DOXAZOSINUM COMPR. FILM. ELIB. 4 mg

PREL.

KAMIREN XL 4 mg 4 mg KRKA D.D. NOVO MESTO

C02CA04 DOXAZOSINUM COMPR. 1 mg

DOXAZOSIN 1 MEDOCHEMIE 1 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 1 1 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 1 mg KRKA D.D. NOVO MESTO

MAGUROL 1 mg MEDOCHEMIE LTD.

C02CA04 DOXAZOSINUM COMPR. 2 mg

DOXAZOSIN 2 MEDOCHEMIE 2 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 2 2 mg ALIUD(R) PHARMA GMBH &

CO.KG

DOXAZOSIN SANDOZ 2 mg 2 mg HEXAL AG

KAMIREN 2 mg KRKA D.D. NOVO MESTO

MAGUROL 2 mg MEDOCHEMIE LTD.

C02CA04 DOXAZOSINUM COMPR. 4 mg

DOXAZOSIN 4 MEDOCHEMIE 4 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 4 4 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 4 mg KRKA D.D. NOVO MESTO

MAGUROL 4 mg MEDOCHEMIE LTD.

C02CA04 DOXAZOSINUM COMPR. ELIB. MODIF. 4 mg

CARDURA XL 4 mg 4 mg PFIZER H.C.P. CORPORATION

C02CA04 DOXAZOSINUM COMPR. FILM. ELIB. 4 mg

PREL.

KAMIREN XL 4 mg 4 mg KRKA D.D. NOVO MESTO

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| 35 |C03AA03| HYDROCHLOROTHIAZIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03AA03 HYDROCHLOROTHIAZIDUM COMPR. 25 mg

NEFRIX 25 mg 25 mg ZENTIVA SA

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| 36 |C03BA11| INDAPAMIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03BA11 INDAPAMIDUM COMPR. FILM. ELIB. 1,5 mg

PREL.

IMPAMID SR 1,5 mg 1,5 mg GEDEON RICHTER

ROMANIA S.A.

INDATER SR 1,5 mg 1,5 mg TERAPIA SA

C03BA11 INDAPAMIDUM COMPR. FILM. ELIB. 1.5 mg

PREL.

INDAPAMID SR 1,5 mg LAROPHARM 1.5 mg LAROPHARM SRL

C03BA11 INDAPAMIDUM COMPR. ELIB. PREL. 1.5 mg

INDAPAMID MCC 1,5 mg 1.5 mg MAGISTRA C & C SRL

C03BA11 INDAPAMIDUM COMPR. FILM. ELIB. 1.5 mg

PREL.

INDAPAMID LPH(R) 1,5 mg 1.5 mg LABORMED PHARMA SA

RAWEL SR 1,5 mg 1.5 mg KRKA D.D. NOVO MESTO

TERTENSIF(R) SR 1.5 mg LES LAB. SERVIER IND.

C03BA11 INDAPAMIDUM COMPR. 2.5 mg

IMPAMID(R) 2,5 mg 2.5 mg GEDEON RICHTER ROMANIA SA

C03BA11 INDAPAMIDUM COMPR. FILM. 2.5 mg

INDAPAMID LPH (R) 2,5 mg 2.5 mg LABORMED PHARMA SA

INDAPAMIDE 2,5 mg 2.5 mg HEMOFARM S.R.L.

INDATER(R) 2,5 mg 2.5 mg TERAPIA SA

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| 37 |C03CA01| FUROSEMIDUM | |

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Electroliţii serici trebuie să fie verificaţi periodic.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03CA01 FUROSEMIDUM SOL. INJ. 10 mg/ml

FUROSEMID 20 mg/2 ml 10 mg/ml ZENTIVA SA

C03CA01 FUROSEMIDUM COMPR. 40 mg

FUROSEMID ARENA 40 mg 40 mg ARENA GROUP S.A.

FUROSEMID EEL 40 mg BIO EEL SRL

FUROSEMID LPH 40 mg 40 mg LABORMED PHARMA SA

FUROSEMID MCC 40 mg 40 mg MAGISTRA C & C SRL

FUROSEMID SLAVIA 40 mg SLAVIA PHARM SRL

FUROSEMID ZENTIVA 40 mg ZENTIVA SA

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| 38 |C03DA01| SPIRONOLACTONUM | |

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Electroliţii serici trebuie să fie verificaţi periodic.

Femeile la vârsta fertilă la care s-a iniţiat tratament cu spironolactona trebuie să ia măsuri adecvate de contracepţie.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03DA01 SPIRONOLACTONUM CAPS. 100 mg

VEROSPIRON 100 mg GEDEON RICHTER LTD.

C03DA01 SPIRONOLACTONUM COMPR. 25 mg

SPIRONOLACTONA 25 mg 25 mg BIO EEL SRL

C03DA01 SPIRONOLACTONUM COMPR. FILM. 25 mg

ALSPIRON 25 mg 25 mg AC HELCOR PHARMA SRL

SPIRONOLACTONA 25 mg 25 mg TERAPIA SA

C03DA01 SPIRONOLACTONUM CAPS. 50 mg

VEROSPIRON 50 mg GEDEON RICHTER LTD.

C03DA01 SPIRONOLACTONUM COMPR. FILM. 50 mg

ALSPIRON 50 mg 50 mg AC HELCOR PHARMA SRL

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| 39 |C03EB01| COMBINAŢII (SPIRONOLACTONUM + FUROSEMIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03EB01 COMBINAŢII CAPS.

(SPIRONOLACTONUM +

FUROSEMIDUM)

DIUREX 50 TERAPIA SA

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| 40 |C04AD03| PENTOXIFYLLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C04AD03 PENTOXIFYLLINUM COMPR. FILM. ELIB. 400 mg

PREL.

ANGIOPENT 400 mg 400 mg AC HELCOR PHARMA SRL

PENTOXI RETARD 400 mg 400 mg TERAPIA SA

C04AD03 PENTOXIFYLLINUM DRAJ. ELIB. PREL. 400 mg

TRENTAL(R) 400 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

C04AD03 PENTOXIFYLLINUM COMPR. FILM. ELIB. 600 mg

PREL.

ANGIOPENT 600 mg 600 mg AC HELCOR PHARMA SRL

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| 41 |C07AA05| PROPRANOLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AA05 PROPRANOLOLUM COMPR. 10 mg

N-PROPRANOLOL 10 mg 10 mg SC MEDUMAN SA

PROPRANOLOL 10 mg BIO EEL SRL

PROPRANOLOL 10 mg 10 mg SINTOFARM SA

C07AA05 PROPRANOLOLUM COMPR. 40 mg

PROPRANOLOL 40 mg 40 mg SINTOFARM SA

PROPRANOLOL EEL 40 mg 40 mg BIO EEL SRL

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| 42 |C07AA07| SOTALOLUM | |

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Prescriere limitată: Tratamentul aritmiilor ventriculare severe.

Tratamentul aritmiilor supraventriculare.

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare denumirii comune internaţionale.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AA07 SOTALOLUM COMPR. 160 mg

ALS-SOTALOL 160 mg 160 mg ALSIFCOM INTERMED SRL

DAROB(R) 160 mg 160 mg ABBOTT GMBH & CO.KG

SOTAGAMMA 160 160 mg WORWAG PHARMA GMBH & CO.KG

SOTALOL AL 160 160 mg ALIUD PHARMA GMBH & CO.KG

C07AA07 SOTALOLUM COMPR. 80 mg

ALS-SOTALOL 80 mg 80 mg ALSIFCOM INTERMED SRL

DAROB(R) 80 mg 80 mg ABBOTT GMBH & CO.KG

SOTAGAMMA 80 80 mg WORWAG PHARMA GMBH & CO.KG

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| 43 |C07AB02| METOPROLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB02 METOPROLOLUM COMPR. 100 mg

BETAPROL 100 mg 100 mg AC HELCOR PHARMA SRL

BLOXAN 100 mg KRKA D.D. NOVO MESTO

EGILOK 100 mg 100 mg EGIS PHARMACEUTICALS

P.L.C.

METOPRO TAD 100 100 mg TAD PHARMA GMBH

METOPROLOL 100 mg 100 mg MAGISTRA C & C SRL

METOPROLOL AL 100 100 mg ALIUD(R) PHARMA GMBH &

CO.KG

METOPROLOL LPH 100 mg 100 mg LABORMED PHARMA SA

METOPROLOL MEDICO UNO 100 mg 100 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

METOPROLOL TERAPIA 100 mg 100 mg TERAPIA SA

VASOCARDIN(R) 100 100 mg SLOVAKOFARMA

C07AB02 METOPROLOLUM COMPR. ELIB. PREL. 100 mg

METOPROLOL RETARD 100 mg 100 mg TERAPIA S.A.

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. 100 mg

PREL.

BETALOCR ZOC 100 mg 100 mg ASTRAZENECA AB

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. MOD. 190 mg

METOSUCCINAT SANDOZ 190 mg 190 mg HEXAL AG

C07AB02 METOPROLOLUM COMPR. ELIB. PREL. 200 mg

VASOCARDIN(R) SR 200 200 mg ZENTIVA AS

C07AB02 METOPROLOLUM COMPR. 25 mg

EGILOK 25 mg 25 mg EGIS PHARMACEUTICALS

P.L.C.

METOPROLOL 25 mg 25 mg ARENA GROUP SA

METOPROLOL LPH 25 mg 25 mg LABORMED PHARMA SA

METOPROLOL MEDICO UNO 25 mg 25 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. MOD. 47.5 mg

METOSUCCINAT SANDOZ 47,5 mg 47.5 mg HEXAL AG

C07AB02 METOPROLOLUM COMPR. 50 mg

BETAPROL 50 mg 50 mg AC HELCOR PHARMA SRL

EGILOK 50 mg 50 mg EGIS PHARMACEUTICALS

P.L.C.

METOPRO TAD 50 50 mg TAD PHARMA GMBH

METOPROLOL 50 mg 50 mg MAGISTRA C & C SRL

METOPROLOL AL 50 50 mg ALIUD PHARMA GMBH & CO.KG

METOPROLOL LPH 50 mg 50 mg LABORMED PHARMA SA

METOPROLOL MEDICO UNO 50 mg 50 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

METOPROLOL TERAPIA 50 mg 50 mg TERAPIA SA

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. 50 mg

PREL.

BETALOCR ZOC 50 mg 50 mg ASTRAZENECA AB

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. MOD. 95 mg

METOSUCCINAT SANDOZ 95 mg 95 mg HEXAL AG

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| 44 |C07AB03| ATENOLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA FIRMA

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C07AB03 ATENOLOLUM COMPR. 100 mg

ATECOR 100 mg WIN - MEDICARE LTD.

ATENOCOR 100 mg 100 mg AC HELCOR SRL

ATENOLOL 100 mg 100 mg ARENA GROUP SA

ATENOLOL LPH 100 mg 100 mg LABORMED PHARMA SA

C07AB03 ATENOLOLUM COMPR. FILM. 100 mg

ATENOLOL 100 mg 100 mg TERAPIA SA

ATENOLOL 100 mg MEDO 100 mg MEDOCHEMIE ROMANIA SRL

VASCOTEN 100 mg MEDOCHEMIE LTD.

C07AB03 ATENOLOLUM COMPR. 50 mg

ATENOCOR 50 mg 50 mg AC HELCOR SRL

ATENOLOL 50 mg 50 mg SLAVIA PHARM SRL

ATENOLOL LPH 50 mg 50 mg LABORMED PHARMA SA

C07AB03 ATENOLOLUM COMPR. FILM. 50 mg

ATENOLOL 50 mg 50 mg TERAPIA SA

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| 45 |C07AB05| BETAXOLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB05 BETAXOLOLUM COMPR. FILM. 20 mg

BETAC 20 mg MEDOCHEMIE LTD.

LOKREN 20 mg 20 mg SANOFI-AVENTIS FRANCE

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| 46 |C07AB07| BISOPROLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB07 BISOPROLOLUM COMPR. 10 mg

BISOBLOCK 10 mg 10 mg KERI PHARMA GENERICS LTD.

C07AB07 BISOPROLOLUM COMPR. FILM. 10 mg

BISOGAMMA(R) 10 10 mg WORWAG PHARMA GMBH & CO.KG

BISOTENS 10 mg 10 mg ANTIBIOTICE SA

CONCOR 10 mg 10 mg MERCK KGAA

C07AB07 BISOPROLOLUM COMPR. FILM. 2.5 mg

CONCOR COR 2,5 mg 2.5 mg MERCK KGAA

C07AB07 BISOPROLOLUM COMPR. 5 mg

BISOBLOCK 5 mg 5 mg KERI PHARMA GENERICS LTD.

C07AB07 BISOPROLOLUM COMPR. FILM. 5 mg

BISOGAMMA(R) 5 5 mg WORWAG PHARMA GMBH & CO.KG

BISOTENS 5 mg 5 mg ANTIBIOTICE SA

CONCOR 5 mg 5 mg MERCK KGAA

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| 47 |C07AG02| CARVEDILOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AG02 CARVEDILOLUM COMPR. 12,5 mg

CARVEDILOL SANDOZ 12,5 mg HEXAL AG

C07AG02 CARVEDILOLUM COMPR. 12.5 mg

ATRAM 12.5 12.5 mg ZENTIVA AS

CARVEDILOL 12,5 mg 12.5 mg VIM SPECTRUM SRL

CARVEDILOL HELCOR 12,5 mg 12.5 mg AC HELCOR PHARMA SRL

CARVEDILOL LPH 12,5 mg 12.5 mg LABORMED PHARMA SA

CARVEDILOL TEVA 12,5 mg 12.5 mg TEVA PHARMACEUTICAL S.R.L.

CORYOL(R) 12.5 mg KRKA D.D.

DILATREND(R) 12,5 mg 12.5 mg ROCHE ROMANIA S.R.L.

TALLITON(R) 12,5 mg 12.5 mg EGIS PHARMACEUTICALS LTD.

C07AG02 CARVEDILOLUM COMPR. FILM. 12.5 mg

CARVEDIGAMMA 12,5 mg 12.5 mg WORWAG PHARMA GMBH & CO.KG

C07AG02 CARVEDILOLUM COMPR. 25 mg

ATRAM 25 25 mg ZENTIVA AS

CARVEDILOL HELCOR 25 mg 25 mg AC HELCOR PHARMA SRL

CARVEDILOL LPH 25 mg 25 mg LABORMED PHARMA SA

CARVEDILOL SANDOZ 25 mg HEXAL AG

CARVEDILOL TEVA 25 mg 25 mg TEVA PHARMACEUTICAL S.R.L.

CORYOL(R) 25 mg KRKA D.D.

DILATREND(R) 25 mg 25 mg ROCHE ROMANIA S.R.L.

TALLITON(R) 25 mg 25 mg EGIS PHARMACEUTICALS LTD.

C07AG02 CARVEDILOLUM COMPR. FILM. 25 mg

CARVEDIGAMMA 25 mg 25 mg WORWAG PHARMA GMBH & CO.KG

C07AG02 CARVEDILOLUM COMPR. 3.125 mg

CORYOL(R) 3,125 mg 3.125 mg KRKA D.D.

C07AG02 CARVEDILOLUM COMPR. 6.25 mg

ATRAM 6.25 6.25 mg ZENTIVA AS

CARVEDILOL 6,25 mg 6.25 mg VIM SPECTRUM SRL

CARVEDILOL LPH 6,25 mg 6.25 mg LABORMED PHARMA SA

CARVEDILOL SANDOZ 6.25 mg HEXAL AG

CARVEDILOL TEVA 6,25 mg 6.25 mg TEVA PHARMACEUTICAL S.R.L.

CORYOL(R) 6.25 mg KRKA D.D.

DILATREND(R) 6,25 mg 6.25 mg ROCHE ROMANIA S.R.L.

TALLITON 6,25 mg 6.25 mg EGIS PHARMACEUTICALS LTD.

C07AG02 CARVEDILOLUM COMPR. FILM. 6.25 mg

CARVEDIGAMMA 6,25 mg 6.25 mg WORWAG PHARMA GMBH & CO.KG

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| 48 |C08CA01| AMLODIPINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C08CA01 AMLODIPINUM COMPR. 10 mg

ALOZUR 10 mg 10 mg OZONE LABORATORIES BV

AMLO TAD 10 mg 10 mg TAD PHARMA GMBH

AMLODIPIN 10 mg 10 mg VIM SPECTRUM SRL

AMLODIPINE-TEVA 10 mg 10 mg TEVA PHARMACEUTICAL S.R.L.

AMLOHEXAL 10 mg 10 mg HEXAL AG

NORVASC 10 mg 10 mg PFIZER EUROPE MA EEIG

STAMLO M 10 mg 10 mg DR. REDDY'S LABORATORIES

TENOX(R) 10 10 mg KRKA D.D.

VASOREX 10 mg 10 mg LABORMED PHARMA S.A.

C08CA01 AMLODIPINUM COMPR. 5 mg

ALOZUR 5 mg 5 mg OZONE LABORATORIES BV

AMLO TAD 5 mg 5 mg TAD PHARMA GMBH

AMLODIPIN 5 mg 5 mg VIM SPECTRUM SRL

AMLODIPINE-TEVA 5 mg 5 mg TEVA PHARMACEUTICAL S.R.L.

AMLOHEXAL 5 mg 5 mg HEXAL AG

NORVASC 5 mg 5 mg PFIZER EUROPE MA EEIG

STAMLO M 5 mg 5 mg DR. REDDY'S LABORATORIES

TENOX(R) 5 5 mg KRKA D.D.

VASOREX 5 mg 5 mg LABORMED PHARMA S.A.

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| 49 |C08CA02| FELODIPINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C08CA02 FELODIPINUM COMPR. ELIB. MODIF. 10 mg

FELODIPIN AL 10 RETARD 10 mg ALIUD(R) PHARMA GMBH &

CO.KG

C08CA02 FELODIPINUM COMPR. ELIB. PREL. 10 mg

PLENDIL 10 mg ASTRAZENECA AB

C08CA02 FELODIPINUM COMPR. FILM. ELIB. 10 mg

MODIF.

SISTAR 10 mg 10 mg GEDEON RICHTER ROMANIA SA

C08CA02 FELODIPINUM COMPR. FILM. ELIB. 10 mg

PREL.

AURONAL 10 mg 10 mg EGIS PHARMACEUTICALS PLC

MIVARA 10 mg 10 mg STADA ARZNEIMITTEL AG

PRESID(R) 10 mg 10 mg IVAX-PHARMACEUTICALS

S.R.O.

C08CA02 FELODIPINUM COMPR. ELIB. MODIF. 2.5 mg

FELODIPIN AL 2,5 RETARD 2.5 mg ALIUD(R) PHARMA GMBH &

CO.KG

C08CA02 FELODIPINUM COMPR. ELIB. PREL. 2.5 mg

PLENDIL 2.5 mg ASTRAZENECA AB

C08CA02 FELODIPINUM COMPR. FILM. ELIB. 2.5 mg

MODIF.

SISTAR 2,5 mg 2.5 mg GEDEON RICHTER ROMANIA SA

C08CA02 FELODIPINUM COMPR. FILM. ELIB. 2.5 mg

PREL.

AURONAL 2,5 mg 2.5 mg EGIS PHARMACEUTICALS PLC

PRESID(R) 2,5 mg 2.5 mg IVAX-PHARMACEUTICALS

S.R.O.

C08CA02 FELODIPINUM COMPR. ELIB. MODIF. 5 mg

FELODIPIN AL 5 RETARD 5 mg ALIUD(R) PHARMA GMBH &

CO.KG

C08CA02 FELODIPINUM COMPR. ELIB. PREL. 5 mg

PLENDIL 5 mg ASTRAZENECA AB

C08CA02 FELODIPINUM COMPR. FILM. ELIB. 5 mg

MODIF.

SISTAR 5 mg 5 mg GEDEON RICHTER ROMANIA SA

C08CA02 FELODIPINUM COMPR. FILM. ELIB. 5 mg

PREL.

AURONAL 5 mg 5 mg EGIS PHARMACEUTICALS PLC

MIVARA 5 mg 5 mg STADA ARZNEIMITTEL AG

PRESID(R) 5 mg 5 mg IVAX-PHARMACEUTICALS

S.R.O.

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| 50 |C08CA05| NIFEDIPINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C08CA05 NIFEDIPINUM COMPR. ELIB. MODIF. 20 mg

ADALAT CR 20 20 mg BAYER HEALTHCARE AG

C08CA05 NIFEDIPINUM COMPR. FILM. ELIB. 20 mg

PREL.

ADALAT(R) RETARD 20 mg BAYER HEALTHCARE AG

NIFEDIPIN RETARD TERAPIA 20 mg TERAPIA SA

20 mg

C08CA05 NIFEDIPINUM DRAJ. 20 mg

EPILAT RETARD 20 mg E.I.P.I.CO. MED S.R.L.

C08CA05 NIFEDIPINUM COMPR. ELIB. MODIF. 30 mg

ADALAT CR 30 30 mg BAYER HEALTHCARE AG

C08CA05 NIFEDIPINUM COMPR. ELIB. MODIF. 20 mg

ADALAT CR 20 20 mg BAYER HEALTHCARE AG

C08CA05 NIFEDIPINUM COMPR. FILM. ELIB. 20 mg

PREL.

ADALAT(R) RETARD 20 mg BAYER HEALTHCARE AG

NIFEDIPIN RETARD TERAPIA 20 mg TERAPIA SA

20 mg

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| 51 |C08DA01| VERAPAMILUM | |

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Efectele de inhibare a funcţiei miocardice ale acestui medicament se cumulează cu cele ale beta-blocantelor.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C08DA01 VERAPAMILUM COMPR. FILM. ELIB. 240 mg

PREL.

ISOPTIN RR 240 mg ABBOTT GMBH & CO.KG

VEROGALID(R) ER 240 mg 240 mg IVAX-PHARMACEUTICALS

S.R.O.

C08DA01 VERAPAMILUM COMPR. FILM. 40 mg

ISOPTIN(R) 40 mg 40 mg ABBOTT GMBH & CO.KG

VERAPAMIL AL 40 40 mg ALIUD(R) PHARMA GMBH &

CO.KG

C08DA01 VERAPAMILUM COMPR. FILM. 80 mg

ISOPTIN(R) 80 mg 80 mg ABBOTT GMBH & CO.KG

VERAPAMIL AL 80 80 mg ALIUD(R) PHARMA GMBH &

CO.KG

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| 52 |C08DB01| DILTIAZEMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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C08DB01 DILTIAZEMUM COMPR. 60 mg

DILTIAZEM ALKALOID 60 mg 60 mg ALKALOID DOO

DILTIAZEM EIPICO 60 mg 60 mg E.I.P.I.CO. MED S.R.L.

DILTIAZEM LPH 60 mg 60 mg LABORMED PHARMA SA

DILZEM 60 mg 60 mg PFIZER EUROPE MA EEIG

C08DB01 DILTIAZEMUM COMPR. 90 mg

DILTIAZEM ALKALOID 90 mg 90 mg ALKALOID DOO

C08DB01 DILTIAZEMUM COMPR. FILM. ELIB. 90 mg

PREL.

DILZEM 90 mg RETARD 90 mg PFIZER EUROPE MA EEIG

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| 53 |C09AA01| CAPTOPRILUM | |

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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C09AA01 CAPTOPRILUM COMPR. 12,5 mg

CAPTOPRIL MCC 12,5 mg 12,5 mg MAGISTRA C & C

C09AA01 CAPTOPRILUM COMPR. 12.5 mg

CAPTOPRIL 12,5 mg 12.5 mg EGIS PHARMACEUTICALS LTD.

C09AA01 CAPTOPRILUM COMPR. 25 mg

CAPTOPRIL-AC 25 mg 25 mg AC HELCOR PHARMA SRL

CAPTOPRIL SINTOFARM 25 mg 25 mg SINTOFARM SA

CAPTOPRIL 25 EEL 25 mg BIO EEL SRL

CAPTOPRIL 25 mg 25 mg EGIS PHARMACEUTICALS LTD.

CAPTOPRIL LPH 25 mg 25 mg LABORMED PHARMA SA

CAPTOPRIL MCC 25 mg 25 mg MAGISTRA C & C

C09AA01 CAPTOPRILUM COMPR. 50 mg

CAPTOPRIL 50 mg 50 mg ARENA GROUP SA

C09AA01 CAPTOPRILUM COMPR. 50 mg

CAPTOPRIL-AC 50 mg 50 mg AC HELCOR PHARMA SRL

CAPTOPRIL 50 EEL 50 mg BIO EEL SRL

CAPTOPRIL 50 mg 50 mg EGIS PHARMACEUTICALS LTD.

CAPTOPRIL LPH 50 mg 50 mg LABORMED PHARMA SA

CAPTOPRIL MCC 50 mg 50 mg MAGISTRA C & C

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| 54 |C09AA02| ENALAPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA02 ENALAPRILUM SOL. INJ. 1.25 mg/ml

ENAP(R) 1.25 mg/ml KRKA D.D.

C09AA02 ENALAPRILUM COMPR. 10 mg

EDNYT(R) 10 mg 10 mg GEDEON RICHTER LTD.

ENAHEXAL(R) 10 mg 10 mg HEXAL AG

ENALA TAD 10 10 mg TAD PHARMA GMBH

ENALAP 10 mg E.I.P.I.CO. MED S.R.L.

ENALAPRIL 10 mg 10 mg MAGISTRA C & C

ENALAPRIL AL 10 10 mg ALIUD(R) PHARMA GMBH &

CO.KG

ENALAPRIL FABIOL 10 mg 10 mg FABIOL SA

ENALAPRIL LPH 10 mg 10 mg LABORMED PHARMA SA

ENALAPRIL SANDOZ 10 mg 10 mg SANDOZ SRL

ENALAPRIL TERAPIA 10 mg 10 mg TERAPIA SA

ENAM 10 mg 10 mg REPREZENTANTA DR. REDDY'S

LABORATORIES LTD.

ENAP 10 mg 10 mg KRKA D.D. NOVO MESTO

RENITEC 10 mg 10 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C09AA02 ENALAPRILUM COMPR. 2.5 mg

EDNYT(R) 2,5 mg 2.5 mg GEDEON RICHTER LTD.

C09AA02 ENALAPRILUM COMPR. 20 mg

EDNYT(R) 20 mg 20 mg GEDEON RICHTER LTD.

ENAHEXAL(R) 20 mg 20 mg HEXAL AG

ENALA TAD 20 20 mg TAD PHARMA GMBH

ENALAPRIL 20 mg OZONE LABORATORIES LTD.

ENALAPRIL 20 mg 20 mg MAGISTRA C & C

ENALAPRIL AL 20 20 mg ALIUD(R) PHARMA GMBH &

CO.KG

ENALAPRIL FABIOL 20 mg 20 mg FABIOL SA

ENALAPRIL LPH 20 mg 20 mg LABORMED PHARMA SA

ENALAPRIL SANDOZ 20 mg 20 mg SANDOZ SRL

ENALAPRIL TERAPIA 20 mg 20 mg TERAPIA SA

ENAP 20 mg 20 mg KRKA D.D. NOVO MESTO

RENITEC 20 mg 20 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C09AA02 ENALAPRILUM COMPR. 5 mg

EDNYT(R) 5 mg 5 mg GEDEON RICHTER LTD.

ENAHEXAL(R) 5 mg 5 mg HEXAL AG

ENALA TAD 5 5 mg TAD PHARMA GMBH

ENALAPRIL 5 mg 5 mg OZONE LABORATORIES LTD.

ENALAPRIL AL 5 5 mg ALIUD(R) PHARMA GMBH &

CO.KG

ENALAPRIL LPH(R) 5 mg 5 mg LABORMED PHARMA SA

ENALAPRIL SANDOZ 5 mg 5 mg SANDOZ SRL

ENAP 5 mg 5 mg KRKA D.D. NOVO MESTO

RENITEC 5 mg 5 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

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| 55 |C09AA03| LISINOPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA03 LISINOPRILUM COMPR. 10 mg

LISIGAMMA 10 mg 10 mg WORWAG PHARMA GMBH & CO.KG

LISINOPRIL 10 MEDO 10 mg MEDOCHEMIE ROMANIA SRL

LISINOPRIL ANTIBIOTICE 10 mg 10 mg ANTIBIOTICE SA

LISINOPRIL SANDOZ 10 mg 10 mg HEXAL AG

LISIREN 10 mg 10 mg AC HELCOR SRL

MEDAPRIL 10 10 mg MEDOCHEMIE LTD.

RANOLIP 10 mg RANBAXY U.K. LIMITED

SINOPRYL(R) 10 10 mg ECZACIBASI PHARMACEUTICALS

S.R.L.

TONOLYSIN 10 mg 10 mg GEDEON RICHTER LTD.

C09AA03 LISINOPRILUM COMPR. 2.5 mg

TONOLYSIN 2,5 mg 2.5 mg GEDEON RICHTER LTD.

C09AA03 LISINOPRILUM COMPR. 20 mg

USIGAMMA 20 mg 20 mg WORWAG PHARMA GMBH & CO.KG

LISINOPRIL 20 MEDO 20 mg MEDOCHEMIE ROMANIA SRL

LISINOPRIL ANTIBIOTICE 20 mg 20 mg ANTIBIOTICE SA

LISINOPRIL SANDOZ 20 mg 20 mg HEXAL AG

LISIREN 20 mg 20 mg AC HELCOR SRL

MEDAPRIL 20 20 mg MEDOCHEMIE LTD.

RANOLIP 20 mg RANBAXY U.K. LIMITED

TONOLYSIN 20 mg 20 mg GEDEON RICHTER LTD.

C09AA03 LISINOPRILUM COMPR. 40 mg

LISINOPRIL ANTIBIOTICE 40 mg 40 mg ANTIBIOTICE SA

C09AA03 LISINOPRILUM COMPR. 5 mg

LISIGAMMA 5 mg 5 mg WORWAG PHARMA GMBH & CO.KG

LISINOPRIL 5 MEDO 5 mg MEDOCHEMIE ROMANIA SRL

LISINOPRIL SANDOZ 5 mg 5 mg HEXAL AG

MEDAPRIL 5 5 mg MEDOCHEMIE LTD.

RANOLIP 5 mg RANBAXY U.K. LIMITED

TONOLYSIN 5 mg 5 mg GEDEON RICHTER LTD.

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| 56 |C09AA05| RAMIPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA05 RAMIPRILUM COMPR. FILM. 1.25 mg

RAMIRAN 1,25 mg 1.25 mg RANBAXY U.K. LIMITED

C09AA05 RAMIPRILUM COMPR. 10 mg

EMREN 10 mg 10 mg GEDEON RICHTER ROMANIA

S.A.

VIVACE 10 mg 10 mg ACTAVIS GROUP HF

C09AA05 RAMIPRILUM COMPR. 10 mg

AMPRIL 10 mg 10 mg KRKA D.D. NOVO MESTO

PIRAMIL 10 mg 10 mg SANDOZ SRL

RAMIGAMMA 10 mg 10 mg WORWAG PHARMA GMBH & CO.KG

RAMIPRIL-AC 10 mg 10 mg AC HELCOR PHARMA SRL

TRITACE 10 10 mg SANOFI-AVENTIS DEUTSCHLAND

GMBH

ZENRA 10 10 mg ZENTIVA S.A.

C09AA05 RAMIPRILUM COMPR. FILM. 10 mg

RAMIRAN 10 mg 10 mg RANBAXY U.K. LIMITED

C09AA05 RAMIPRILUM COMPR. 2,5 mg

EMREN 2,5 mg 2,5 mg GEDEON RICHTER ROMANIA

S.A.

VIVACE 2,5 mg 2,5 mg ACTAVIS GROUP HF

C09AA05 RAMIPRILUM COMPR. 2.5 mg

AMPRIL 2,5 mg 2.5 mg KRKA D.D. NOVO MESTO

PIRAMIL 2,5 mg 2.5 mg SANDOZ SRL

RAMIPRIL-AC 2,5 mg 2.5 mg AC HELCOR PHARMA SRL

TRITACE(R) 2,5 2.5 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

ZENRA 2,5 2.5 mg ZENTIVA S.A.

C09AA05 RAMIPRILUM COMPR. FILM. 2.5 mg

RAMIRAN 2,5 mg 2.5 mg RANBAXY U.K. LIMITED

C09AA05 RAMIPRILUM COMPR. 5 mg

EMREN 5 mg 5 mg GEDEON RICHTER ROMANIA

S.A.

VIVACE 5 mg 5 mg ACTAVIS GROUP HF

C09AA05 RAMIPRILUM COMPR. 5 mg

AMPRIL 5 mg 5 mg KRKA D.D. NOVO MESTO

PIRAMIL 5 mg 5 mg SANDOZ SRL

RAMIGAMMA 5 mg 5 mg WORWAG PHARMA GMBH & CO.KG

RAMIPRIL-AC 5 mg 5 mg AC HELCOR PHARMA SRL

TRITACE(R) 5 5 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

ZENRA 5 5 mg ZENTIVA S.A.

C09AA05 RAMIPRILUM COMPR. FILM. 5 mg

RAMIRAN 5 mg 5 mg RANBAXY U.K. LIMITED

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| 57 |C09AA06| QUINAPRILUM | |

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA06 QUINAPRILUM COMPR. FILM. 10 mg

QUINAPRIL SANDOZ 10 mg 10 mg HEXAL AG

C09AA06 QUINAPRILUM COMPR. FILM. 10 mg

ACCUPRO 10 mg 10 mg PFIZER EUROPE MA EEIG

AQUIRIL 10 mg 10 mg LABORMED PHARMA SA

QUINARAN 10 mg 10 mg RANBAXY UK LIMITED

C09AA06 QUINAPRILUM COMPR. FILM. 20 mg

QUINAPRIL SANDOZ 20 mg 20 mg HEXAL AG

C09AA06 QUINAPRILUM COMPR. FILM. 20 mg

ACCUPRO(R) 20 20 mg PFIZER EUROPE MA EEIG

QUINARAN 20 mg 20 mg RANBAXY UK LIMITED

C09AA06 QUINAPRILUM COMPR. FILM. 40 mg

QUINAPRIL SANDOZ 40 mg 40 mg HEXAL AG

C09AA06 QUINAPRILUM COMPR. FILM. 40 mg

QUINARAN 40 mg 40 mg RANBAXY UK LIMITED

C09AA06 QUINAPRILUM COMPR. FILM. 5 mg

QUINAPRIL SANDOZ 5 mg 5 mg HEXAL AG

C09AA06 QUINAPRILUM COMPR. FILM. 5 mg

ACCUPRO(R) 5 5 mg PFIZER EUROPE MA EEIG

QUINARAN 5 mg 5 mg RANBAXY UK LIMITED

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| 58 |C09AA09| FOSINOPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA09 FOSINOPRILUM COMPR. 10 mg

FOSINOPRIL TERAPIA 10 mg 10 mg TERAPIA SA

FOSYPRIL 10 mg 10 mg TERAPIA S.A.

MONOPRIL 10 mg 10 mg BRISTOL MYERS SQUIBB KFT

C09AA09 FOSINOPRILUM COMPR. 20 mg

FOSINOPRIL TERAPIA 20 mg 20 mg TERAPIA SA

FOSINOPRIL TEVA 20 mg 20 mg TEVA PHARMACEUTICAL S.R.L.

FOSYPRIL 20 mg 20 mg TERAPIA SA.

MONOPRIL 20 mg 20 mg BRISTOL-MYERS SQUIBB KFT

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| 59 |C09BA02| COMBINAŢII (ENALAPRILUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09BA02 COMBINAŢII COMPR. 10 mg/25 mg

(ENALAPRILUM + 10 mg/25 mg HEXAL AG

HYDROCHLOROTHIAZIDUM)

ENALAPRIL HCT SANDOZ

10 mg/25 mg

C09BA02 COMBINAŢII COMPR. 10 mg + 12.5 mg

(ENALAPRILUM +

HYDROCHLOROTHIAZIDUM)

ENAP HL 10 mg + 12.5 mg KRKA D.D. NOVO MESTO

C09BA02 COMBINAŢII COMPR. 10 mg + 25 mg

(ENALAPRILUM +

HYDROCHLOROTHIAZIDUM)

ENAP H 10 mg + 25 mg KRKA D.D. NOVO MESTO

C09BA02 COMBINAŢII COMPR. 20 mg/12.5 mg

(ENALAPRILUM +

HYDROCHLOROTHIAZIDUM)

ENAP(R) HL 20 mg/12,5 mg 20 mg/12.5 mg KRKA D.D. NOVO MESTO

C09BA02 COMBINAŢII COMPR. 20 mg/12.5 mg

(ENALAPRILUM + 20 mg/12.5 mg HEXAL AG

HYDROCHLOROTHIAZIDUM)

ENALAPRIL HCT SANDOZ

20 mg/12,5 mg

C09BA02 COMBINAŢII COMPR.

(ENALAPRILUM +

HYDROCHLOROTHIAZIDUM)

VIMAPRIL H VIM SPECTRUM SRL

VIMAPRIL HL VIM SPECTRUM SRL

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| 60 |C09BA05| COMBINAŢII (RAMIPRILUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09BA05 COMBINAŢII COMPR. 2.5 mg/12.5 mg

(RAMIPRILUM +

HYDROCHLOROTHIAZIDUM)

HARTIL HCT 2,5 mg/12,5 mg 2.5 mg/12.5 mg EGIS PHARMACEUTICALS PLC

PIRAMIL HCT 2,5 mg/12,5 mg 2.5 mg/12.5 mg HEXAL AG

RAMIPRIL HCT-MEDOCHEMIE 2.5 mg/12.5 mg MEDOCHEMIE LTD.

C09BA05 COMBINAŢII COMPR. 2.5 mg + 12.5 mg

(RAMIPRILUM +

HYDROCHLOROTHIAZIDUM)

AMPRIL HL 2.5 mg + 12.5 mg KRKA D.D. NOVO MESTO

C09BA05 COMBINAŢII COMPR. 5 mg/25 mg

(RAMIPRILUM +

HYDROCHLOROTHIAZIDUM)

HARTIL HCT 5 mg/25 mg 5 mg/25 mg EGIS PHARMACEUTICALS PLC

PIRAMIL HCT 5 mg/25 mg 5 mg/25 mg HEXAL AG

RAMIPRIL HCT-MEDOCHEMIE 5 mg/25 mg MEDOCHEMIE LTD.

C09BA05 COMBINAŢII COMPR. 5 mg + 25 mg

(RAMIPRILUM +

HYDROCHLOROTHIAZIDUM)

AMPRIL HD 5 mg + 25 mg KRKA D.D. NOVO MESTO

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| 61 |C09CA01| LOSARTANUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09CA01 LOSARTANUM COMPR. FILM. 50 mg

COZAAR 50 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

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| 62 |C10AA01| SIMVASTATINUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AA01 SIMVASTATINUM COMPR. FILM. 10 mg

SIMVASTATIN 10 mg 10 mg TERAPIA S.A.

C10AA01 SIMVASTATINUM COMPR. FILM. 10 mg

SIMGAL(R) 10 mg 10 mg IVAX-PHARMACEUTICALS

S.R.O.

SIMVA TAD 10 mg 10 mg TAD PHARMA GMBH

SIMVACARD(R) 10 10 mg ZENTIVA AS

SIMVAGAMMA 10 mg 10 mg WORWAG PHARMA GMBH & CO.KG

SIMVAHEXAL(R) 10 mg 10 mg HEXAL AG

SIMVASTATIN LPH 10 mg 10 mg LABORMED PHARMA SA

SIMVOR 10 mg 10 mg RANBAXY U.K. LIMITED

SINTENAL 10 mg 10 mg AC HELCOR PHARMA SRL

VABADIN 10 mg 10 mg MENARINI INTERNATIONAL

OPERATIONS LUXEMBURG S.A.

VASILIP 10 mg 10 mg KRKA D.D.

ZAREDIL 10 mg 10 mg OZONE LABORATORIES LTD.

ZEPLAN(R) 10 mg 10 mg GEDEON RICHTER ROMANIA SA

ZOCOR 10 mg 10 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C10AA01 SIMVASTATINUM COMPR. FILM. 20 mg

SIMVASTATIN 20 mg 20 mg TERAPIA S.A.

C10AA01 SIMVASTATINUM COMPR. FILM. 20 mg

SIMCOR 20 mg 20 mg ANTIBIOTICE SA

SIMGAL(R) 20 mg 20 mg IVAX-PHARMACEUTICALS

S.R.O.

SIMVA TAD 20 mg 20 mg TAD PHARMA GMBH

SIMVACARD(R) 20 20 mg ZENTIVA AS

SIMVAGAMMA 20 mg 20 mg WORWAG PHARMA GMBH & CO.KG

SIMVAHEXAL(R) 20 mg 20 mg HEXAL AG

SIMVASTATIN LPH 20 mg 20 mg LABORMED PHARMA SA

SIMVOR 20 mg 20 mg RANBAXY U.K. LIMITED

SINTENAL 20 mg 20 mg AC HELCOR PHARMA SRL

VABADIN 20 mg 20 mg MENARINI INTERNATIONAL

OPERATIONS LUXEMBURG S.A.

VASILIP 20 mg 20 mg KRKA D.D.

ZAREDIL 20 mg 20 mg OZONE LABORATORIES LTD.

ZEPLAN(R) 20 mg 20 mg GEDEON RICHTER ROMANIA SA

ZOCOR 20 mg 20 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C10AA01 SIMVASTATINUM COMPR. FILM. 30 mg

SIMVAHEXAL(R) 30 mg 30 mg HEXAL AG

C10AA01 SIMVASTATINUM COMPR. FILM. 40 mg

SIMVASTATIN 40 mg 40 mg TERAPIA S.A.

C10AA01 SIMVASTATINUM COMPR. FILM. 40 mg

SIMCOR 40 mg 40 mg ANTIBIOTICE S.A.

SIMGAL(R) 40 mg 40 mg IVAX-PHARMACEUTICALS

S.R.O.

SIMVA TAD 40 mg 40 mg TAD PHARMA GMBH

SIMVACARD(R) 40 40 mg ZENTIVA AS

SIMVAGAMMA 40 mg 40 mg WORWAG PHARMA GMBH & CO.KG

SIMVAHEXAL(R) 40 mg 40 mg HEXAL AG

SIMVASTATIN LPH 40 mg 40 mg LABORMED PHARMA S.A.

SIMVOR 40 mg 40 mg RANBAXY U.K. LIMITED

SINTENAL 40 mg 40 mg AC HELCOR PHARMA SRL

VABADIN 40 mg 40 mg MENARINI INTERNATIONAL

OPERATIONS LUXEMBURG S.A.

VASILIP(R) 40 mg 40 mg KRKA D.D.

ZAREDIL 40 mg 40 mg OZONE LABORATORIES LTD.

ZEPLAN(R) 40 mg 40 mg GEDEON RICHTER ROMANIA SA

ZOCOR FORTE 40 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C10AA01 SIMVASTATINUM COMPR. FILM. 5 mg

SIMVOR 5 mg 5 mg RANBAXY U.K. LIMITED

C10AA01 SIMVASTATINUM COMPR. FILM. 80 mg

SIMVASTATIN LPH 80 mg 80 mg LABORMED PHARMA S.A.

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| 63 |C10AA02| LOVASTATINUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AA02 LOVASTATINUM COMPR. 20 mg

MEDOSTATIN 20 mg MEDOCHEMIE LTD.

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| 64 |C10AA03| PRAVASTATINUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AA03 PRAVASTATINUM COMPR. 10 mg

PRALIP 10 mg 10 mg SANDOZ SRL

PRAVATOR 10 mg 10 mg RANBAXY U.K. LIMITED

C10AA03 PRAVASTATINUM COMPR. 20 mg

PRALIP 20 mg 20 mg SANDOZ SRL

PRAVATOR 20 mg 20 mg RANBAXY U.K. LIMITED

C10AA03 PRAVASTATINUM COMPR. 40 mg

PRALIP 40 mg 40 mg SANDOZ SRL

PRAVATOR 40 mg 40 mg RANBAXY U.K. LIMITED

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| 65 |C10AB05| FENOFIBRATUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

NOTĂ:

Riscul de toxicitate musculară severă creşte dacă fenofibratum este utilizat simultan cu inhibitori de HMG CoA reductaza sau alţi fibraţi. Terapia combinată trebuie folosită cu precauţie la pacienţii cu dislipidemie mixtă severă combinată cu risc cardiovascular înalt, în absenţa antecedentelor de afecţiune musculară. Pacienţii vor fi monitorizaţi regulat pentru semne cronice de toxicitate musculară.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AB05 FENOFIBRATUM CAPS. 100 mg

LIPANTHYL(R) 100 100 mg LAB. FOURNIER SA

C10AB05 FENOFIBRATUM CAPS. 160 mg

LIPOFIB 160 mg 160 mg TERAPIA S.A.

C10AB05 FENOFIBRATUM COMPR. FILM. ELIB. 160 mg

MODIF.

LIPANTHYL(R) SUPRA 160 mg 160 mg LAB. FOURNIER SA

C10AB05 FENOFIBRATUM CAPS. 200 mg

LIPOFIB 200 mg 200 mg TERAPIA S.A.

C10AB05 FENOFIBRATUM CAPS. PULB. MICRONIZATĂ 200 mg

FENOFIBRAT LPH 200 mg 200 mg LABORMED PHARMA SA

LIPANTHYL 200 M 200 mg LAB. FOURNIER SA

LIPIVIM 200 mg VIM SPECTRUM SRL

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| 66 |D01AC02| MICONAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D01AC02 MICONAZOLUM CREMĂ 20 mg/g

MICONAL ECOBI 20 mg/g FARMACEUTICI ECOBI S.A.S.

D01AC02 MICONAZOLUM CREMĂ 2%

MEDACTER 2% FARAN LABORATORIES S.A.

MICONAZOL NITRAT 2% 2% SLAVIA PHARM SRL

D01AC02 MICONAZOLUM GEL 2%

DERMOZOL 2% PHARCOIMPEX 93 S.R.L.

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| 67 |D01AC10| BIFONAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D01AC10 BIFONAZOLUM SOL. CUT. 0,01 g/ml

MYCOSPOR 0,01 g/ml BAYER HEALTHCARE AG

D01AC10 BIFONAZOLUM CREMĂ 100 g

MYCO-FLUSEMIDON 100 g ANFARM HELLAS S.A.

PHARMACEUTICALS

D01AC10 BIFONAZOLUM CREMĂ 10 mg/g

BIAZOL 10 mg/g 10 mg/g GEDEON RICHTER ROMANIA SA

D01AC10 BIFONAZOLUM CREMĂ 1%

MYCOSPOR 1% BAYER HEALTHCARE AG

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| 68 |D06BB03| ACICLOVIRUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D06BB03 ACICLOVIRUM CREMĂ 50 mg/g

ACICLOVIR HYPERION 50 mg/g HYPERION S.A.

ACIKLOVIR CREMĂ 50 mg/g A & G MED TRADING S.R.L.

D06BB03 ACICLOVIRUM CREMĂ 5%

ACICLOVIR 5% OZONE LABORATORIES LTD.

ACICLOVIR 5% 5% GEDEON RICHTER ROMANIA SA

CLOVIRAL(R) 5% 5% ANTIBIOTICE SA

ZOVIRAX 5% THE WELLCOME FOUNDATION

LTD.

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| 69 |D07AA02| HYDROCORTISONUM | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AA02 HYDROCORTISONUM UNGUENT 1,00%

HIDROCORTIZON 1% 1% ANTIBIOTICE SA

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| 70 |D07AC01| BETAMETHASONUM | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AC01 BETAMETHASONUM CREMĂ 0,5 mg/g

BELODERM CREMĂ 0,5 mg/g A & G MED TRADING S.R.L.

D07AC01 BETAMETHASONUM UNGUENT 0,5 mg/g

BELODERM UNGUENT 0,5 mg/g A & G MED TRADING S.R.L.

D07AC01 BETAMETHASONUM CREMĂ 0.1%

BETADERM 0.1% E.I.P.I.CO. MED S.R.L.

D07AC01 BETAMETHASONUM UNGUENT 0.1%

BETADERM 0.1% E.I.P.I.CO. MED S.R.L.

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| 71 |G01AF04| MICONAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G01AF04 MICONAZOLUM CAPS. MOI VAG. 1.2 g

MICONAL ECOBI 1.2 g FARMACEUTICI ECOBI S.A.S

G01AF04 MICONAZOLUM CREMĂ VAG. 20 mg/g

MICONALECOBI 20 mg/g FARMACEUTICI ECOBI S.A.S

G01AF04 MICONAZOLUM SOL. VAGINALĂ 2 mg/ml

MICONAL ECOBI 2 mg/ml FARMACEUTICI ECOBI S.A.S

G01AF04 MICONAZOLUM OVULE 50 mg

MICONALECOBI 50 mg FARMACEUTICI ECOBI S.A.S

G01AF04 MICONAZOLUM CREMĂ VAG. 2%

MEDACTER 2% FARAN LABORATORIES S.A.

MYCOHEAL 2% DAR AL DAWA PHARMA S.R.L.

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| 72 |G02AB03| ERGOMETRINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G02AB03 ERGOMETRINUM SOL. INJ. 0.2 mg/ml

MALEAT DE ERGOMETRINA 0.2 mg/ml ZENTIVA S.A.

0,2 mg/ml

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| 73 |G03BA03| TESTOSTERONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03BA03 TESTOSTERONUM SOL. INJ. 1000 mg/4 ml

NEBIDO 1000 mg/4 ml 1000 mg/4 ml SCHERING AG

G03BA03 TESTOSTERONUM CAPS. MOI 40 mg

UNDESTO(R) TESTOCAPS(R) 40 mg ORGANON NV

Prescriere limitată: Deficit androgenic la bărbaţi de cauză primară testiculară sau secundară hipotalamo-hipofizară (confirmat prin nivel plasmatic al testosteronului la cel puţin 2 determinări în două dimineţi diferite mai mic sau egal cu limita inferioară a valorilor reactivului utilizat).

Prescriere limitată: Micropenis, inducerea pubertăţii sau întârzierea constituţională a creşterii sau a pubertăţii la băieţi sub 18 ani.

G03BA03 TESTOSTERONUM GEL 50 mg

ANDROGEL 50 mg 50 mg LAB. BESINS INTERNATIONAL

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| 74 |G03CA03| ESTRADIOLUM\* | |

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Prescriere limitată: Utilizat în simptomele caracteristice post-menopauzei, în cazul în care terapia estrogenică în doze reduse a demonstrat intoleranţa la administrarea orală cu estrogeni.

Această limitare este valabilă doar pentru formele farmaceutice Sistem terapeutic transdermic şi plasture transdermic.

NOTĂ:

Estradiol trebuie folosit împreună cu un progestativ oral la femeile nehisterectomizate.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03CA03 ESTRADIOLUM GEL 0.06%

OESTROGEL(R) 0.06% LAB. BESINS INTERNATIONAL

G03CA03 ESTRADIOLUM GEL 0.1%

ESTREVA(R) 0.1% LAB. THERAMEX

G03CA03 ESTRADIOLUM COMPR. FILM. 1 mg

ESTROFEM 1 mg 1 mg NOVO NORDISK A/S

G03CA03 ESTRADIOLUM COMPR. VAG. 25 micrograme

VAGIFEM 25 micrograme NOVO NORDISK A/S

G03CA03 ESTRADIOLUM IMPLANT 25 mg

RISELLE 25 mg 25 mg ORGANON NV

G03CA03 ESTRADIOLUM PLASTURE TRANSDERM. 50 micrograme/24 ore

CLIMARA 50 micrograme/24 ore SCHERING AG

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| 75 |G04CAN1| DOXAZOSINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04CAN1 DOXAZOSINUM COMPR. 1 mg

DOXAZOSIN 1 MEDOCHEMIE 1 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 1 1 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 1 mg KRKA D.D. NOVO MESTO

MAGUROL 1 mg MEDOCHEMIE LTD.

G04CAN1 DOXAZOSINUM COMPR. 2 mg

DOXAZOSIN 2 MEDOCHEMIE 2 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 2 2 mg ALIUD(R) PHARMA GMBH &

CO.KG

DOXAZOSIN SANDOZ 2 mg 2 mg HEXAL AG

KAMIREN 2 mg KRKA D.D. NOVO MESTO

MAGUROL 2 mg MEDOCHEMIE LTD.

G04CAN1 DOXAZOSINUM COMPR. 4 mg

DOXAZOSIN 4 MEDOCHEMIE 4 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 4 4 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 4 mg KRKA D.D. NOVO MESTO

MAGUROL 4 mg MEDOCHEMIE LTD.

G04CAN1 DOXAZOSINUM COMPR. ELIB. MODIF. 4 mg

CARDURA XL 4 mg 4 mg PFIZER H.C.P. CORPORATION

G04CAN1 DOXAZOSINUM COMPR. FILM. ELIB. 4 mg

PREL.

KAMIREN XL 4 mg 4 mg KRKA D.D. NOVO MESTO

G04CAN1 DOXAZOSINUM COMPR. 1 mg

DOXAZOSIN 1 MEDOCHEMIE 1 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 1 1 mg ALIUD(R) PHARMA GMBH &

CO.KG

KAMIREN 1 mg KRKA D.D. NOVO MESTO

MAGUROL 1 mg MEDOCHEMIE LTD.

G04CAN1 DOXAZOSINUM COMPR. 2 mg

DOXAZOSIN 2 MEDOCHEMIE 2 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 2 2 mg ALIUD(R) PHARMA GMBH &

CO.KG

DOXAZOSIN SANDOZ 2 mg 2 mg HEXAL AG

KAMIREN 2 mg KRKA D.D. NOVO MESTO

MAGUROL 2 mg MEDOCHEMIE LTD.

G04CAN1 DOXAZOSINUM COMPR. 4 mg

DOXAZOSIN 4 MEDOCHEMIE 4 mg MEDOCHEMIE ROMANIA SRL

DOXAZOSIN AL 4 4 mg ALIUD(R) PHARMA GMBH &

CO.KG

DOXAZOSIN SANDOZ 4 mg 4 mg HEXAL AG

KAMIREN 4 mg KRKA D.D. NOVO MESTO

MAGUROL 4 mg MEDOCHEMIE LTD.

G04CAN1 DOXAZOSINUM COMPR. ELIB. MODIF. 4 mg

CARDURA XL 4 mg 4 mg PFIZER H.C.P. CORPORATION

G04CAN1 DOXAZOSINUM COMPR. FILM. ELIB. 4 mg

PREL.

KAMIREN XL 4 mg 4 mg KRKA D.D. NOVO MESTO

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| 76 |H02AB04| METHYLPREDNISOLONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 125 mg

LEMOD SOLU 125 mg 125 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. ŞI SOLV. PT. SOL. 125 mg/2 ml

INJ.

SOLU-MEDROL ACT-O-VIAL 125 mg/2 ml PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 20 mg

LEMOD SOLU 20 mg 20 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. ŞI SOLV. PT. SOL. 250 mg/4 ml

INJ.

SOLU-MEDROL ACT-O-VIAL 250 mg/4 ml PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 40 mg

LEMOD SOLU 40 mg 40 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. ŞI SOLV. PT. SOL. 40 mg/1 ml

INJ.

SOLU-MEDROL ACT-O-VIAL 40 mg/1 ml PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 500 mg

LEMOD SOLU 500 mg 500 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. + SOLV. PT. SOL. 500 mg/7.8 ml

INJ.

SOLU-MEDROL 500 mg/7,8 ml 500 mg/7.8 ml PFIZER EUROPE MA EEIG

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| 77 |H02AB07| PREDNISONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AB07 PREDNISONUM COMPR. 5 mg

N-PREDNISON 5 mg MEDUMAN SA

PREDNISON 5 mg 5 mg SINTOFARM SA

PREDNISON ARENA 5 mg 5 mg ARENA GROUP SA

PREDNISON GEDEON RICHTER 5 mg 5 mg GEDEON RICHTER ROMÂNIA SA

PREDNISON MAGISTRA 5 mg 5 mg MAGISTRA C & C

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| 78 |H02AB09| HYDROCORTISONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AB09 HYDROCORTISONUM LIOF. + SOLV. PT. SOL. 100 mg

INJ.

HYDROCORTISONE 100 mg 100 mg HEMOFARM S.R.L.

HYDROCORTISONE SUCCINAT SODIC 100 mg E.I.P.I.CO. MED S.R.L.

H02AB09 HYDROCORTISONUM SOL. INJ. I.V. 25 mg/5 ml

HIDROCORTIZON HEMISUCCINAT 25 mg/5 ml ZENTIVA S.A.

H02AB09 HYDROCORTISONUM LIOF. PT. SOL. INJ. + 500 mg

SOLV.

HYDROCORTISONE 500 mg 500 mg HEMOFARM S.R.L.

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| 79 |J01AA02| DOXYCYCLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01AA02 DOXYCYCLINUM CAPS. 100 mg

DOXICICLINA 100 mg EUROPHARM SA

DOXICICLINA 100 mg 100 mg ANTIBIOTICE SA

DOXICICLINA SANDOZ 100 mg 100 mg SANDOZ SRL

J01AA02 DOXYCYCLINUM COMPR. DISP. 100 mg

UNIDOX SOLUTAB 100 mg ASTELLAS PHARMA EUROPE

B.V.

J01AA02 DOXYCYCLINUM CAPS. 100 mg

DOXICICLINA 100 mg ARENA GROUP SA

DOXICICLINA 100 mg 100 mg ANTIBIOTICE SA

DOXICICLINA SANDOZ 100 mg 100 mg SANDOZ SRL

J01AA02 DOXYCYCLINUM COMPR. DISP. 100 mg

UNIDOX SOLUTAB 100 mg ASTELLAS PHARMA EUROPE

B.V.

J01AA02 DOXYCYCLINUM CAPS. 100 mg

DOXICICLINA 100 mg ARENA GROUP SA

DOXICICLINA 100 mg 100 mg ANTIBIOTICE SA

DOXICICLINA SANDOZ 100 mg 100 mg SANDOZ SRL

J01AA02 DOXYCYCLINUM COMPR. DISP. 100 mg

UNIDOX SOLUTAB 100 mg ASTELLAS PHARMA EUROPE

B.V.

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| 80 |J01CA04| AMOXICILLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CA04 AMOXICILLINUM COMPR. FILM. 1000 mg

OSPAMOX 1000 mg 1000 mg SANDOZ GMBH

J01CA04 AMOXICILLINUM PULB. PT. SUSP. ORALĂ 125 mg/5 ml

AMOXICILINA 125 mg/5 ml 125 mg/5 ml OZONE LABORATORIES LTD.

AMOXICILINA SANDOZ 125 mg/5 ml SANDOZ S.R.L.

125 mg/5 ml PULBERE PENTRU

SUSPENSIE ORALĂ

JULPHAMOX 125 mg/5 ml GULF PHARMACEUTICAL

INDUSTRIES S.R.L.

OSPAMOX(R) 125 mg/5 ml 125 mg/5 ml SANDOZ GMBH

J01CA04 AMOXICILLINUM PULB. SUSP. 125 mg/5 ml

MOXILEN 125 mg/5 ml MEDOCHEMIE LTD.

J01CA04 AMOXICILLINUM CAPS. 250 mg

AMOXICILINA 250 mg EUROPHARM SA

AMOXICILINA SANDOZ 250 mg 250 mg SANDOZ SRL

CAPSULE

AMOXICILINA 250 mg 250 mg OZONE LABORATORIES LTD.

AMOXICILINA ANTIBIOTICE 250 mg ANTIBIOTICE SA

250 mg

AMOXICILINA ARENA 250 mg 250 mg ARENA GROUP SA

AMOXICILINA MEDICO UNO 250 mg 250 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

OSPAMOX 250 mg 250 mg SANDOZ GMBH

J01CA04 AMOXICILLINUM PULB. PT. SUSP. ORALĂ 250 mg/5 ml

AMOXICILINA SANDOZ 250 mg/5 ml SANDOZ S.R.L.

250 mg/5 ml PULBERE PENTRU

SUSPENSIE ORALĂ

AMOXICILINA 250 mg/5 ml 250 mg/5 ml OZONE LABORATORIES LTD.

E-MOX 250 mg/5 ml E.I.P.I.CO. MED S.R.L.

JULPHAMOX 250 mg/5 ml GULF PHARMACEUTICAL

INDUSTRIES S.R.L.

OSPAMOX 250 mg/5 ml 250 mg/5 ml SANDOZ GMBH

J01CA04 AMOXICILLINUM PULB. SUSP. 250 mg/5 ml

MOXILEN FORTE 250 mg/5 ml MEDOCHEMIE LTD.

J01CA04 AMOXICILLINUM CAPS. 500 mg

AMOXICILINA ANTIBIOTICE 500 mg ANTIBIOTICE SA

500 mg

AMOXICILINA ARENA 500 mg 500 mg ARENA GROUP SA

AMOXICILINA FORTE 500 mg 500 mg OZONE LABORATORIES LTD.

AMOXICILINA SANDOZ 500 mg 500 mg SANDOZ SRL

CAPSULE

E-MOX 500 mg E.I.P.I.CO. MED S.R.L.

EPHAMOX 500 mg EUROPHARM SA

MOXILEN 500 mg MEDOCHEMIE LTD.

J01CA04 AMOXICILLINUM COMPR. FILM. 500 mg

OSPAMOX 500 mg 500 mg SANDOZ GMBH

J01CA04 AMOXICILLINUM COMPR. PT. DISPERSIE 1000 mg

ORALĂ

DUOMOX 1000 mg 1000 mg ASTELLAS PHARMA EUROPE BV

J01CA04 AMOXICILLINUM COMPR. PT. DISPERSIE 250 mg

ORALĂ

DUOMOX 250 mg 250 mg ASTELLAS PHARMA EUROPE BV

J01CA04 AMOXICILLINUM COMPR. PT. DISPERSIE 500 mg

ORALĂ

DUOMOX 500 mg 500 mg ASTELLAS PHARMA EUROPE BV

J01CA04 AMOXICILLINUM COMPR. PT. DISPERSIE 750 mg

ORALĂ

DUOMOX 750 mg 750 mg ASTELLAS PHARMA EUROPE BV

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| 81 |J01CE02| PHENOXYMETHYLPENICILLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CE02 PHENOXY- COMPR. 1000000 ui

METHYLPENICILLINUM

PENICILINA V 1 000 000 U.I. 1000000 ui EUROPHARM SA

J01CE02 PHENOXY- COMPR. FILM. 1000000 ui

METHYLPENICILLINUM

OSPEN(R) 1000 1000000 ui SANDOZ GMBH

J01CE02 PHENOXY- COMPR. FILM. 1500000 ui

METHYLPENICILLINUM

OSPEN(R) 1500 1500000 ui SANDOZ GMBH

J01CE02 PHENOXY- SIROP 400000 ui/5 ml

METHYLPENICILLINUM

OSPEN 400 400000 ui/5 ml SANDOZ GMBH

J01CE02 PHENOXY- COMPR. FILM. 500000 ui

METHYLPENICILLINUM

OSPEN(R) 500 500000 ui SANDOZ GMBH

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| 82 |J01CE08| BENZATHINI BENZYLPENICILLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CE08 BENZATHINI PULB. PT. SUSP. INJ. 1200000 ui

BENZYLPENICILLINUM

MOLDAMIN(R) 1200000 ui ANTIBIOTICE SA

J01CE08 BENZATHINI PULB. PT. SUSP. INJ. 600000 ui

BENZYLPENICILLINUM

MOLDAMIN(R) 600000 ui ANTIBIOTICE SA

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| 83 |J01CF04| OXACILLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CF04 OXACILLINUM CAPS. 250 mg

OXACILINA 250 mg FARMACOM SA

OXACILINA 250 mg 250 mg ANTIBIOTICE SA

OXACILINA ARENA 250 mg 250 mg ARENA GROUP SA

OXACILINA FARMEX 250 mg 250 mg FARMEX COMPANY SRL

OXACILINA SANDOZ 250 mg 250 mg SANDOZ SRL

J01CF04 OXACILLINUM CAPS. 500 mg

OXACILINA 500 mg 500 mg ANTIBIOTICE SA

OXACILINA ARENA 500 mg 500 mg ARENA GROUP SA

OXACILINA FORTE 500 mg 500 mg OZONE LABORATORIES LTD.

OXACILINA SANDOZ 500 mg 500 mg SANDOZ SRL

OXALIN 500 mg 500 mg EUROPHARM SA

J01CF04 OXACILLINUM PULB. PT. SOL. INJ./ 1 g

PERF. I.M./I.V.

OXACILINA ANTIBIOTICE 1 g 1 g ANTIBIOTICE SA

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| 84 |J01CR02| AMOXICILLINUM + ACIDUM CLAVULANICUM | |

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Prescriere limitată: Tratamentul de scurtă durată a infecţiilor bacteriene

atunci când este suspectată rezistenţa la amoxicilină.

Infecţii la care este dovedită rezistenţa la

amoxicilină.

A fost raportată hepatotoxicitate la acest medicament.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CR02 AMOXICILLINUM + COMPR. FILM. 1000 mg

ACIDUM CLAVULANICUM

AMOKSIKLAV 2 x 1000 mg 1000 mg LEK PHARMACEUTICALS D.D.

MEDOCLAV 1000 mg 1000 mg MEDOCHEMIE LTD.

J01CR02 AMOXICILLINUM + COMPR. FILM. ELIB. 1062.5 mg

ACIDUM CLAVULANICUM PREL.

AUGMENTIN(TM) SR 1062.5 mg BEECHAM GROUP PLC

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 125 mg + 31.25/5 m

ACIDUM CLAVULANICUM

MEDOCLAV 156,25 mg/5 ml 125 mg + 31.25/5 ml MEDOCHEMIE LTD.

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 156.25 mg/5 ml

ACIDUM CLAVULANICUM

AMOKSIKLAV 156,25 mg/5 ml 156.25 mg/5 ml LEK PHARMACEUTICALS D.D.

BIOCLAVID 156.25 mg/5 ml 156.25 mg/5 ml SANDOZ GMBH

J01CR02 AMOXICILLINUM + COMPR. DISP. 250/62,5 mg

ACIDUM CLAVULANICUM

FORCID SOLUTAB 250/62,5 250/62,5 mg ASTELLAS PHARMA EUROPE BV

J01CR02 AMOXICILLINUM + COMPR. FILM. 250 mg + 125 mg

ACIDUM CLAVULANICUM

MEDOCLAV 375 mg 250 mg + 125 mg MEDOCHEMIE LTD.

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 250 mg +

ACIDUM CLAVULANICUM 62.5 mg/5 ml

MEDOCLAV FORTE 312,5 mg/5 ml 250 mg + 62.5 mg/5 ml MEDOCHEMIE LTD.

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 312.5 mg/5 ml

ACIDUM CLAVULANICUM

BIOCLAVID 312.5 mg/5 ml 312.5 mg/5 ml SANDOZ GMBH

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 312.5 mg/5 ml

ACIDUM CLAVULANICUM

AMOKSIKLAV 312.5 mg/5 ml 312.5 mg/5 ml LEK PHARMACEUTICALS D.D.

ENHANCIN 312,5 mg/5 ml 312.5 mg/5 ml RANBAXY UK LIMITED

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 400 mg + 57 mg/5 m

ACIDUM CLAVULANICUM

AUGMENTIN BIS 400 mg + 57 mg/5 ml SMITHKLINE BEECHAM PLC

J01CR02 AMOXICILLINUM + COMPR. DISP. 500/125 mg

ACIDUM CLAVULANICUM

FORCID SOLUTAB 500/125 500/125 mg ASTELLAS PHARMA EUROPE BV

J01CR02 AMOXICILLINUM + COMPR. FILM. 500 mg + 125 mg

ACIDUM CLAVULANICUM

AUGMENTIN 625 mg 500 mg + 125 mg BEECHAM GROUP PLC

MEDOCLAV 625 mg 500 mg + 125 mg MEDOCHEMIE LTD.

J01CR02 AMOXICILLINUM + COMPR. FILM. 625 mg

ACIDUM CLAVULANICUM

AMOKSIKLAV 2 x 625 mg 625 mg LEK PHARMACEUTICALS D.D.

BIOCLAVID 625 mg 625 mg SANDOZ GMBH

J01CR02 AMOXICILLINUM + PULB. PT. SUSP. ORALĂ 642.90 mg/5 ml

ACIDUM CLAVULANICUM

AUGMENTIN(R) ES 642.90 mg/5 ml SMITHKLINE BEECHAM PLC

J01CR02 AMOXICILLINUM + COMPR. DISP. 875 mg + 125 mg

ACIDUM CLAVULANICUM

FORCID SOLUTAB 875/125 875 mg + 125 mg ASTELLAS PHARMA EUROPE BV

J01CR02 AMOXICILLINUM + COMPR. FILM. 875 mg + 125 mg

ACIDUM CLAVULANICUM

AUGMENTIN 1 g 875 mg + 125 mg BEECHAM GROUP PLC

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| 85 |J01DB01| CEFALEXINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DB01 CEFALEXINUM COMPR. FILM. 1000 mg

OSPEXIN 1000 mg 1000 mg SANDOZ GMBH

J01DB01 CEFALEXINUM GRAN. PT. SUSP. ORALĂ 125 mg/5 ml

CEFALEXIN SANDOZ 125 mg/5 ml 125 mg/5 ml SANDOZ SRL

GRANULE PENTRU SUSPENSIE

ORALĂ

KEFLEX(R) 125 mg/5 ml 125 mg/5 ml ACTAVIS GROUP HF

OSPEXIN(R) 125 mg/5 ml 125 mg/5 ml SANDOZ GMBH

J01DB01 CEFALEXINUM PULB. PT. SUSP. ORALĂ 125 mg/5 ml

CEFALEXIN 125 mg/5 ml 125 mg/5 ml OZONE LABORATORIES LTD.

J01DB01 CEFALEXINUM CAPS. 250 mg

CEFALEXIN 250 mg 250 mg OZONE LABORATORIES LTD.

CEFALEXIN SANDOZ 250 mg 250 mg SANDOZ SRL

CAPSULE

CEFALEXINA 250 mg 250 mg ANTIBIOTICE SA

CEFALEXINA ARENA 250 mg 250 mg ARENA GROUP SA

CEFALEXINA MEDICO UNO 250 mg 250 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

OSPEXIN(R) 250 mg 250 mg SANDOZ GMBH

SPORIDEX 250 mg 250 mg TERAPIA SA

J01DB01 CEFALEXINUM GRAN. PT. SUSP. ORALĂ 250 mg/5 ml

CEFALEXIN SANDOZ 250 mg/5 ml 250 mg/5 ml SANDOZ SRL

GRANULE PENTRU SUSPENSIE

ORALĂ

KEFLEX(R) 250 mg/5 ml 250 mg/5 ml ACTAVIS GROUP HF

OSPEXIN(R) 250 mg/5 ml 250 mg/5 ml SANDOZ GMBH

J01DB01 CEFALEXINUM PULB. PT. SUSP. ORALĂ 250 mg/5 ml

CEFALEXIN 250 mg/5 ml 250 mg/5 ml OZONE LABORATORIES LTD.

J01DB01 CEFALEXINUM CAPS. 500 mg

CEFALEXIN 500 mg 500 mg OZONE LABORATORIES LTD.

CEFALEXINA 500 mg 500 mg ANTIBIOTICE SA

CEFALEXINA ARENA 500 mg 500 mg ARENA GROUP SA

SPORIDEX 500 mg 500 mg TERAPIA S.A.

J01DB01 CEFALEXINUM COMPR. FILM. 500 mg

CEFALEXIN SANDOZ 500 mg 500 mg SANDOZ SRL

COMPRIMATE FILMATE

OSPEXIN 500 mg 500 mg SANDOZ GMBH

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| 86 |J01DB05| CEFADROXILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DB05 CEFADROXILUM COMPR. FILM. 1000 mg

CEXYL 1000 mg 1000 mg SANDOZ S.R.L.

J01DB05 CEFADROXILUM GRAN. PT. SUSP. ORALĂ 125 mg/5 ml

CEXYL 125 mg/5 ml 125 mg/5 ml SANDOZ S.R.L.

J01DB05 CEFADROXILUM CAPS. 250 mg

CEXYL 250 mg 250 mg SANDOZ S.R.L.

J01DB05 CEFADROXILUM GRAN. SUSP. ORALĂ 250 mg/5 ml

CEXYL 250 mg/5 ml 250 mg/5 ml SANDOZ S.R.L.

J01DB05 CEFADROXILUM CAPS. 500 mg

CEFADROXIL 500 mg OZONE LABORATORIES LTD.

CEFADROXIL TERAPIA 500 mg TERAPIA S.A.

CEFORAN(R) 500 mg 500 mg ANTIBIOTICE SA

CEXYL 500 mg 500 mg SANDOZ S.R.L.

J01DB05 CEFADROXILUM GRAN. SUSP. ORALĂ 500 mg/5 ml

CEXYL 500 mg/5 ml 500 mg/5 ml SANDOZ S.R.L.

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| 87 |J01DC02| CEFUROXIMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DC02 CEFUROXIMUM COMPR. ACOPERITE 125 mg

AXYCEF(R) 125 125 mg SANDOZ SRL

J01DC02 CEFUROXIMUM COMPR. FILM. 125 mg

ZINNAT(R) 125 mg 125 mg GLAXO WELLCOME UK LTD.

J01DC02 CEFUROXIMUM GRAN. PT. SUSP. ORALĂ 125 mg/5 ml

CEROXIM 125 mg/5 ml 125 mg/5 ml RANBAXY UK LIMITED

ZINNAT 125 mg/5 ml GLAXO WELLCOME UK LTD.

J01DC02 CEFUROXIMUM PULB. PT. SUSP. ORALĂ 125 mg/5 ml

AXYCEF(R) 125 mg/5 ml 125 mg/5 ml SANDOZ SRL

J01DC02 CEFUROXIMUM COMPR. 250 mg

CEROXIM 250 mg 250 mg RANBAXY U.K. LIMITED

J01DC02 CEFUROXIMUM COMPR. ACOPERITE 250 mg

AXYCEF(R) 250 250 mg SANDOZ SRL

J01DC02 CEFUROXIMUM COMPR. FILM. 250 mg

ZINNAT(R) 250 mg 250 mg GLAXO WELLCOME UK LTD.

J01DC02 CEFUROXIMUM GRAN. PT. SUSP. ORALĂ 250 mg/5 ml

CEROXIM 250 mg/5 ml 250 mg/5 ml RANBAXY UK LIMITED

J01DC02 CEFUROXIMUM COMPR. 500 mg

CEROXIM 500 mg 500 mg RANBAXY U.K. LIMITED

J01DC02 CEFUROXIMUM COMPR. ACOPERITE 500 mg

AXYCEF(R) 500 500 mg SANDOZ SRL

J01DC02 CEFUROXIMUM COMPR. FILM. 500 mg

ZINNAT(R) 500 mg 500 mg GLAXO WELLCOME UK LTD.

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| 88 |J01DC04| CEFACLORUM | |

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Reacţii asemănătoare bolii serului au fost raportate la utilizarea acestui medicament, în special de către copii.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DC04 CEFACLORUM GRAN. PT. SUSP. ORALĂ 125 mg/5 ml

CECLODYNE 125 mg/5 ml 125 mg/5 ml SANDOZ SRL

CECLOR(R) 125 mg/5 ml 125 mg/5 ml ACTAVIS GROUP HF

CEFACLOR 125 mg/5 ml 125 mg/5 ml OZONE LABORATORIES LTD.

VERCEF 125 mg/5 ml 125 mg/5 ml TERAPIA S.A.

J01DC04 CEFACLORUM PULB. PT. SUSP. ORALĂ 125 mg/5 ml

CEFAKLOR 125 mg/5 ml 125 mg/5 ml HEMOFARM S.R.L.

CLORACEF(R) 125 125 mg/5 ml DAR AL DAWA PHARMA S.R.L

J01DC04 CEFACLORUM CAPS. 250 mg

CECLODYNE 250 mg 250 mg SANDOZ SRL

CEFACLOR 250 mg 250 mg OZONE LABORATORIES LTD.

CLORACEF(R) 250 250 mg DAR AL DAWA PHARMA S.R.L.

MEDOCLOR 250 250 mg MEDOCHEMIE LTD.

J01DC04 CEFACLORUM GRAN. PT. SUSP. ORALĂ 250 mg/5 ml

CECLODYNE 250 mg/5 ml 250 mg/5 ml SANDOZ SRL

CECLOR(R) 250 mg/5 ml 250 mg/5 ml ACTAVIS GROUP HF

CEFACLOR 250 mg/5 ml 250 mg/5 ml OZONE LABORATORIES LTD.

J01DC04 CEFACLORUM PULB. PT. SUSP. ORALĂ 250 mg/5 ml

CEC HEXAL FORTE 250 mg/5 ml 250 mg/5 ml HEXAL AG

CEFAKLOR 250 mg/5 ml 250 mg/5 ml HEMOFARM S.R.L.

CLORACEF(R) 250 250 mg/5 ml DAR AL DAWA PHARMA S.R.L.

J01DC04 CEFACLORUM COMPR. ELIB. PREL. 375 mg

CECLOR(R) MR 375 mg ACTAVIS GROUP HF

CECLOZONE MR 375 mg 375 mg OZONE LABORATORIES LTD.

J01DC04 CEFACLORUM COMPR. FILM. ELIB. 375 mg

MODIF.

CECLODYNE(R) MR 375 mg 375 mg SANDOZ SRL

J01DC04 CEFACLORUM CAPS. 500 mg

CECLODYNE FORTE 500 mg 500 mg SANDOZ SRL

CEFACLOR 500 mg 500 mg OZONE LABORATORIES LTD.

CLORACEF(R) FORTE 500 500 mg DAR AL DAWA PHARMA S.R.L.

MEDOCLOR 500 500 mg MEDOCHEMIE LTD.

J01DC04 CEFACLORUM COMPR. ELIB. PREL. 500 mg

CECLOR(R) MR 500 mg ACTAVIS GROUP HF

J01DC04 CEFACLORUM COMPR. FILM. ELIB. 500 mg

MODIF.

CECLODYNE(R) MR 500 mg 500 mg SANDOZ SRL

J01DC04 CEFACLORUM COMPR. ELIB. PREL. 750 mg

CECLOR(R) MR 750 mg ACTAVIS GROUP HF

CECLOZONE MR 750 mg 750 mg OZONE LABORATORIES LTD.

J01DC04 CEFACLORUM COMPR. FILM. ELIB. 750 mg

MODIF.

CECLODYNE(R) MR 750 mg 750 mg SANDOZ SRL

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| 89 |J01DD08| CEFIXIMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DD08 CEFIXIMUM GRAN. PT. SUSP. ORALĂ 100 mg/5 ml

SUPRAX 100 mg/5 ml GEDEON RICHTER LTD.

J01DD08 CEFIXIMUM CAPS. 200 mg

EFICEF(R) 200 mg 200 mg ANTIBIOTICE SA

J01DD08 CEFIXIMUM COMPR. FILM. 200 mg

SUPRAX 200 mg GEDEON RICHTER LTD.

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| 90 |J01EE01| SULFAMETHOXAZOLUM + TRIMETHOPRIMUM | |

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Există un risc crescut de reacţie adversă severă la administrarea acestui medicament la persoane în vârstă.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01EE01 SULFAMETHOXAZOLUM + SUSP. ORALĂ 200 mg/40 mg/5 ml

TRIMETHOPRIMUM

EPITRIM 200 mg/40 mg/5 ml E.I.P.I.CO. MED S.R.L.

J01EE01 SULFAMETHOXAZOLUM + SIROP 25 mg/5 mg/ml

TRIMETHOPRIMUM

SUMETROLIM 25 mg/5 mg/ml EGIS PHARMACEUTICALS

P.L.C.

J01EE01 SULFAMETHOXAZOLUM + COMPR. 400 mg/80 mg

TRIMETHOPRIMUM

BISEPTRIM 400 mg/80 mg EUROPHARM SA

CO-TRIM ELL 400 mg/80 mg ARENA GROUP S.A.

SUMETROLIM 400 mg/80 mg EGIS PHARMACEUTICALS

P.L.C.

TAGREMIN 400 mg/80 mg ZENTIVA S.A.

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| 91 |J01FA01| ERYTHROMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01FA01 ERYTHROMYCINUM COMPR. 200 mg

ERITROMAGIS 200 mg 200 mg ARENA GROUP SA

ERITROMICINA 200 mg 200 mg ANTIBIOTICE SA

ERITROMICINA EUROPHARM 200 mg 200 mg EUROPHARM SA

ERITROMICINA SANDOZ 200 mg 200 mg SANDOZ S.R.L.

COMPRIMATE

J01FA01 ERYTHROMYCINUM PULB. PT. SUSP. ORALĂ 200 mg/5 ml

ERITRO 200 200 mg/5 ml LEK PHARMATECH SRL

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| 92 |J01FA09| CLARITHROMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01FA09 CLARITHROMYCINUM GRAN. PT. SUSP. ORALĂ 125 mg/5 ml

FROMILID(R) 125 mg/5 ml 125 mg/5 ml KRKA D.D.

KLABAX 125 mg/5 ml 125 mg/5 ml TERAPIA S.A.

KLACID(R) 125 mg/5 ml ABBOTT SPA

LEKOKLAR 125 mg/5 ml 125 mg/5 ml SANDOZ S.R.L.

J01FA09 CLARITHROMYCINUM COMPR. FILM. 250 mg

CLAR 250 250 mg MEDICAROM GROUP S.R.L.

CLARITROMICINA 250 mg 250 mg OZONE LABORATORIES LTD.

FROMILID 250 250 mg KRKA D.D. NOVO MESTO

KLABAX 250 mg 250 mg TERAPIA S.A.

KLACID(R) 250 mg ABBOTT SPA

KLERIMED(R) 250 250 mg MEDOCHEMIE LTD.

LEKOKLAR(R) 250 mg 250 mg LEK PHARMACEUTICALS D.D.

J01FA09 CLARITHROMYCINUM GRAN. PT. SUSP. ORALĂ 250 mg/5 ml

KLABAX 250 mg/5 ml 250 mg/5 ml TERAPIA S.A.

LEKOKLAR 250 mg/5 ml 250 mg/5 ml SANDOZ SRL

J01FA09 CLARITHROMYCINUM COMPR. FILM. 500 mg

CLAR 500 500 mg MEDICAROM GROUP S.R.L.

CLARITROMICINA 500 mg 500 mg OZONE LABORATORIES LTD.

FROMILID 500 500 mg KRKA D.D. NOVO MESTO

KLABAX 500 mg 500 mg TERAPIA S.A.

KLERIMED(R) 500 500 mg MEDOCHEMIE LTD.

LEKOKLAR(R) 500 mg 500 mg LEK PHARMACEUTICALS D.D.

J01FA09 CLARITHROMYCINUM COMPR. FILM. ELIB. 500 mg

MODIF.

KLABAX MR 500 mg 500 mg TERAPIA S.A.

LEKOKLAR XL 500 mg 500 mg LEK PHARMACEUTICALS D.D.

J01FA09 CLARITHROMYCINUM COMPR. FILM. ELIB. 500 mg

PREL.

FROMILID(R) UNO 500 mg KRKA D.D.

KLACID SR 500 mg ABBOTT LABORATORIES LTD.

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| 93 |J01MA01| OFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MA01 OFLOXACINUM COMPR. FILM. 200 mg

OFLOXACIN 200 mg 200 mg ANTIBIOTICE SA

OFLOXIN 200 200 mg ZENTIVA AS

ZANOCIN 200 mg RANBAXY UK LIMITED

J01MA01 OFLOXACINUM COMPR. FILM. ELIB. 400 mg

PREL.

ZANOCIN(R) OD 400 400 mg RANBAXY U.K. LIMITED

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| 94 |J01MA02| CIPROFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MA02 CIPROFLOXACINUM CAPS. 250 mg

EUCIPRIN 250 mg EUROPHARM SA

J01MA02 CIPROFLOXACINUM COMPR. FILM. 250 mg

CIFRAN 250 mg RANBAXY UK LIMITED

CIPHIN 250 250 mg ZENTIVA A.S.

CIPRINOL 250 mg 250 mg KRKA D.D. NOVO MESTO

CIPROCIN 250 mg 250 mg E.I.P.I.CO. MED S.R.L.

CIPROFLOXACINA ALKALOID 250 mg ALKALOID D.O.O.

250 mg

CIPROLEN(R) 250 mg 250 mg AC HELCOR SRL

CIPROLET 250 mg 250 mg DR. REDDY'S LABORATORIES

CIPROZONE 250 mg 250 mg OZONE LABORATORIES LTD.

CUMINOL 250 mg 250 mg GEDEON RICHTER ROMÂNIA

J01MA02 CIPROFLOXACINUM COMPR. FILM. 500 mg

CIFRAN 500 mg RANBAXY UK LIMITED

CIPHIN 500 500 mg ZENTIVA A.S.

CIPRINOL 500 mg 500 mg KRKA D.D. NOVO MESTO

CIPRO QUIN(R) 500 mg ANTIBIOTICE SA

CIPROBAY(R) 500 500 mg BAYER HEALTHCARE AG

CIPROCIN 500 mg 500 mg E.I.P.I.CO MED S.R.L.

CIPRODAR 500 mg DAR AL DAWA PHARMA S.R.L.

CIPROFLOXACINA ALKALOID 500 mg ALKALOID D.O.O.

500 mg

CIPROLEN(R) 500 mg 500 mg AC HELCOR SRL

CIPROLET 500 mg 500 mg DR. REDDY'S LABORATORIES

CIPROZONE FORTE 500 mg 500 mg OZONE LABORATORIES LTD.

CUMINOL 500 mg 500 mg GEDEON RICHTER ROMÂNIA

SIFLOKS(R) 500 mg ECZACIBASI ILAC SANAYI VE

TICARET AS

J01MA02 CIPROFLOXACINUM COMPR. FILM. 750 mg

CIPRINOL 750 mg 750 mg KRKA D.D. NOVO MESTO

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| 95 |J01MA03| PEFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MA03 PEFLOXACINUM COMPR. FILM. 400 mg

PEFLOXACIN LAROPHARM 400 mg LAROPHARM S.R.L.

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| 96 |J01MA06| NORFLOXACINUM | |

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Prescriere limitată: Enterocolită bacteriană.

Infecţii de tract urinar.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MA06 NORFLOXACINUM COMPR. FILM. 400 mg

EPINOR 400 mg E.I.P.I.CO. MED S.R.L.

H-NORFLOXACIN 400 mg 400 mg AC HELCOR SRL

NOLICIN 400 mg 400 mg KRKA D.D. NOVO MESTO

NORFLOX - 400 400 mg LEK PHARMATECH SRL

NORFLOXACIN 400 mg 400 mg OZONE LABORATORIES LTD.

NORFLOXACIN OZONE 400 mg 400 mg OZONE LABORATORIES LTD.

NORFLOXACINA LPH 400 mg 400 mg LABORMED PHARMA SA

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| 97 |J01MB02| ACIDUM NALIDIXICUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MB02 ACIDUM NALIDIXICUM CAPS. 500 mg

NALIXID 500 mg 500 mg ZENTIVA SA

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| 98 |J01XD01| METRONIDAZOLUM | |

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Prescriere limitată: Tratamentul infecţiilor cu anaerobi.

Tratamentul infecţiilor cu protozoare sensibile la

metronidazol.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01XD01 METRONIDAZOLUM COMPR. 250 mg

METRONIDAZOL 250 mg (J01XD01) 250 mg ARENA GROUP SA

J01XD01 METRONIDAZOLUM COMPR. FILM. 250 mg

FLAGYL 250 mg LABORATOIRE AVENTIS

J01XD01 METRONIDAZOLUM SUSP. ORALĂ 4%

FLAGYL 4% (J01XD01) 4% LABORATOIRE AVENTIS

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| 99 |J01XD02| TINIDAZOLUM | |

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Prescriere limitată: Tratamentul infecţiilor cu anaerobi.

Tratamentul infecţiilor cu protozoare sensibile la

tinidazol.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01XD02 TINIDAZOLUM COMPR. FILM. 500 mg

FASIGYN (J01XD02) 500 mg PFIZER EUROPE MA EEIG

TINIZOL(R) 500 mg (J01XD02) 500 mg ZENTIVA S.A.

TIPROGYN 500 500 mg AC HELCOR SRL

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| 100 |J02AB02| KETOCONAZOLUM | |

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Prescriere limitată: Candidoză genitală simptomatică recurentă după

tratamentul local a cel puţin două episoade.

Tratamentul candidozelor muco-cutanate cronice şi

candidozelor orofaringiene care nu pot fi tratate cu

antifungice locale, la pacienţii care prezintă

rezistenţă sau intoleranţă la fluconazol şi

intraconazol.

Micoze sistemice la care alte forme de terapie

antifungică nu au avut efect.

A fost raportată hepatotoxicitate la ketoconazol.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J02AB02 KETOCONAZOLUM COMPR. 200 mg

KEFUNGIN 200 mg ANTIBIOTICE SA

KETOCONAZOL 200 mg 200 mg MAGISTRA C & C

KETOSTIN 200 mg 200 mg AC HELCOR SRL

NIZORAL 200 mg JANSSEN PHARMACEUTICA NV

NIZORAL 200 mg 200 mg TERAPIA SA

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| 101 |J02AC01| FLUCONAZOLUM | |

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Prescriere limitată: Candidoze genitale.

Candidoze ale mucoaselor orofaringiene, esofagiene,

bronhopulmonare non-invazive.

Infecţii cu candida ale pielii.

Candidoze sistemice.

Profilaxia candidozelor la pacienţii cu risc aflaţi în

tratament cu antibiotic.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J02AC01 FLUCONAZOLUM CAPS. 100 mg

DIFLAZON 100 mg 100 mg KRKA D.D.

FLUCONAZOLE TEVA 100 mg 100 mg TEVA PHARMACEUTICALS SRL

FLUCORIC 100 mg 100 mg TERAPIA S.A.

FLUCOVIM 100 100 mg VIM SPECTRUM SRL

FUNGOLON 100 mg 100 mg BALKANPHARMA RAZGRAD AD

J02AC01 FLUCONAZOLUM CAPS. 150 mg

DIFLAZON 150 mg 150 mg KRKA D.D.

DIFLUCAN 150 mg 150 mg PFIZER EUROPE MA EEIG

FLUCONAZOL 150 mg 150 mg OZONE LABORATORIES LTD.

FLUCONAZOL MEDICO UNO 150 mg 150 mg MEDICO UNO PHARMACEUTICAL

SRL

FLUCONAZOL MEDOCHEMIE 150 mg 150 mg MEDOCHEMIE ROMANIA SRL

FLUCONAZOL SANDOZ(R) 150 150 mg SANDOZ SRL

FLUCONAZOL TERAPIA 150 mg 150 mg TERAPIA SA

FLUCONAZOLE TEVA 150 mg 150 mg TEVA PHARMACEUTICALS SRL

FLUCORIC 150 mg 150 mg TERAPIA S.A.

FLUCOVIM 150 150 mg VIM SPECTRUM SRL

MYCOMAX 150 150 mg ZENTIVA AS

MYCOSYSTA(R) 150 mg 150 mg GEDEON RICHTER PLC.

J02AC01 FLUCONAZOLUM CAPS. 200 mg

DIFLAZON 200 mg 200 mg KRKA D.D.

FLUCONAZOLE TEVA 200 mg 200 mg TEVA PHARMACEUTICALS SRL

J02AC01 FLUCONAZOLUM CAPS. 50 mg

DIFLAZON 50 mg 50 mg KRKA D.D.

DIFLUCAN 50 mg 50 mg PFIZER EUROPE MA EEIG

FLUCONAZOL 50 mg 50 mg SLAVIA PHARM SRL

FLUCONAZOL MEDOCHEMIE 50 mg 50 mg MEDOCHEMIE ROMANIA SRL

FLUCONAZOL SANDOZ(R) 50 50 mg SANDOZ SRL

FLUCONAZOL TERAPIA 50 mg 50 mg TERAPIA S.A.

FLUCONAZOLE TEVA 50 mg 50 mg TEVA PHARMACEUTICALS SRL

FLUCORIC 50 mg 50 mg TERAPIA S.A.

FLUCOVIM 50 50 mg VIM SPECTRUM SRL

FUNGOLON 50 mg 50 mg BALKANPHARMA RAZGRAD AD

J02AC01 FLUCONAZOLUM PULB. PT. SUSP. ORALĂ 50 mg/5 ml

DIFLUCAN(R) 50 mg/5 ml 50 mg/5 ml PFIZER EUROPE MA EEIG

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| 102 |J05AB01| ACICLOVIRUM | |

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Prescriere limitată: Herpes genital.

Tratamentul episodic sau supresiv al herpesului genital

recurent.

Herpes cutanat iniţial moderat/sever.

Tratamentul episodic sau supresiv al herpesului cutanat

recurent (forme moderate/severe).

Herpes zoster.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J05AB01 ACICLOVIRUM CAPS. 200 mg

ACICLOVIR 200 mg 200 mg SLAVIA PHARM SRL

EUVIROX 200 mg 200 mg EUROPHARM SA

J05AB01 ACICLOVIRUM COMPR. 200 mg

ACICLOVIR 200 mg 200 mg TERAPIA SA

CLOVIRAL 200 mg 200 mg ANTIBIOTICE SA

ZOVIRAX 200 mg GLAXO WELLCOME FOUNDATION

LTD.

J05AB01 ACICLOVIRUM COMPR. DISP. 200 mg

ACICLOVIR 200 mg 200 mg OZONE LABORATORIES LTD.

LOVIR 200 mg 200 mg RANBAXY UK LIMITED

J05AB01 ACICLOVIRUM PULB. PT. SOL. INJ./ 250 mg

PERF.

VIROLEX 250 mg KRKA D.D. NOVO MESTO

J05AB01 ACICLOVIRUM CAPS. 400 mg

ACICLOVIR 400 mg 400 mg ARENA GROUP S.A.

J05AB01 ACICLOVIRUM COMPR. 400 mg

ACICLOVIR 400 mg 400 mg EGIS PHARMACEUTICALS

P.L.C.

J05AB01 ACICLOVIRUM COMPR. FILM. 400 mg

ACIKLOVIR 400 mg A & G MED TRADING S.R.L.

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| 103 |L02BG01| AMINOGLUTETHIMIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L02BG01 AMINOGLUTETHIMIDUM COMPR. 250 mg

ROGLUTEN(R) 250 mg 250 mg ACTAVIS S.R.L.

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| 104 |M01AB01| INDOMETACINUM | |

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AB01 INDOMETACINUM SUPOZ. 50 mg

INDOMETACIN 50 mg 50 mg MAGISTRA C & C

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| 105 |M01AB05| DICLOFENACUM | |

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AB05 DICLOFENACUM COMPR. FILM. ELIB. 100 mg

PREL.

DICLOTARD(R) 100 mg 100 mg TERAPIA SA

REFEN(R) RETARD 100 mg HEMOFARM S.R.L.

VOLTAREN(R) RETARD 100 mg NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM SUPOZ. 100 mg

CLAFEN 100 mg 100 mg ANTIBIOTICE SA

DICLOFENAC 100 mg 100 mg SINTOFARM SA

DICLOFENAC SODIC 100 mg 100 mg MAGISTRA C & C

DICLOGESIC 100 SUPOZITOARE 100 mg DAR AL DAWA PHARMA SRL

EPIFENAC 100 mg E.I.P.I.CO. MED S.R.L.

VOLTAREN(R) 100 mg NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM SUPOZ. 120 mg

TRATUL 120 120 mg GEROT PHARMAZEUTIKA GMBH

M01AB05 DICLOFENACUM CAPS. ELIB. PREL. 150 mg

DICLOREUM 150 mg 150 mg ALFA WASSERMANN SPA

M01AB05 DICLOFENACUM COMPR. FILM. GASTROREZ. 25 mg

RHEUMAVEK 25 mg FARAN LABORATORIES S.A.

VOLTAREN(R) 25 25 mg NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM COMPR. GASTROREZ. 25 mg

EPIFENAC 25 mg E.I.P.I.CO. MED S.R.L.

M01AB05 DICLOFENACUM DRAJ. GASTROREZ. 25 mg

DICLOFENAC 25 mg 25 mg ARENA GROUP SA

M01AB05 DICLOFENACUM SOL. INJ. 25 mg/ml

RHEUMAVEK 25 mg/ml FARAN LABORATORIES S.A.

VOLTAREN(R) 25 mg/ml NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM SOL. INJ. 30 mg/ml

TRATUL(R) 30 mg/ml GEROT PHARMAZEUTIKA GMBH

M01AB05 DICLOFENACUM CAPS. GASTROREZ. 50 mg

TRATUL 50 50 mg GEROT PHARMAZEUTIKA GMBH

M01AB05 DICLOFENACUM COMPR. FILM. GASTROREZ. 50 mg

CLAFEN 50 mg 50 mg ANTIBIOTICE SA

DICLOFENAC 50 mg 50 mg AC HELCOR PHARMA SRL

VOLTAREN(R) 50 50 mg NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM DRAJ. 50 mg

VOLTAREN RAPID(R) 50 mg NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM DRAJ. GASTROREZ. 50 mg

DICLOFENAC 50 mg 50 mg ARENA GROUP SA

M01AB05 DICLOFENACUM SUPOZ. 50 mg

VOLTAREN(R) 50 mg NOVARTIS PHARMA GMBH

M01AB05 DICLOFENACUM SUPOZ. 60 mg

TRATUL 60 60 mg GEROT PHARMAZEUTIKA GMBH

M01AB05 DICLOFENACUM COMPR. ELIB. MODIF. 75 mg

DICLAC(R) 75 ID 75 mg HEXAL AG

M01AB05 DICLOFENACUM SOL. INJ. 75 mg

DICLAC 75 mg HEXAL AG

VURDON 75 mg HELP S.A. PHARMACEUTICALS

M01AB05 DICLOFENACUM SOL. INJ. 75 mg/3 ml

ALMIRAL 75 mg/3 ml MEDOCHEMIE LTD.

DICLOFENAC 75 mg 75 mg/3 ml TERAPIA SA

DICLOFENAC AL I.M. 75 mg/3 ml ALIUD PHARMA GMBH & CO.KG

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| 106 |M01AB15| KETOROLACUM TROMETHAMIN | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AB15 KETOROLACUM COMPR. FILM. 10 mg

TROMETHAMIN

KETANOV 10 mg 10 mg TERAPIA S.A.

KETOROL 10 mg DR. REDDY'S LABORATORIES

M01AB15 KETOROLACUM SOL. INJ. 30 mg/ml

TROMETHAMIN

KETANOV 30 mg/ml TERAPIA S.A.

KETOROL 30 mg/ml DR. REDDY'S LABORATORIES

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| 107 |M01AB16| ACECLOFENACUM | |

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Prescriere limitată: Tratamentul simptomatic al puseelor de poliartrită reumatoidă, osteoartrită, spondilită anchilozantă, artroză sau afecţiunilor musculo-scheletale acute.

A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AB16 ACECLOFENACUM COMPR. FILM. 100 mg

AFLAMIL 100 mg GEDEON RICHTER PLC.

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| 108 |M01AC02| TENOXICAMUM | |

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AC02 TENOXICAMUM CAPS. 20 mg

TENOXICAM LPH 20 mg 20 mg LABORMED PHARMA SA

M01AC02 TENOXICAMUM COMPR. 20 mg

TENOXICAM 20 mg 20 mg ARENA GROUP SA

M01AC02 TENOXICAMUM COMPR. FILM. 20 mg

NEO-ENDUSIX(R) 20 mg ANFARM HELLAS S.A.

PHARMACEUTICALS

TILCOTIL 20 mg ROCHE ROMANIA S.R.L.

M01AC02 TENOXICAMUM LIOF. + SOLV. PT. SOL. 20 mg

INJ.

NEO-ENDUSIX(R) 20 mg ANFARM HELLAS S.A.

PHARMACEUTICALS

M01AC02 TENOXICAMUM LIOF. ŞI SOLV. PT. SOL. 20 mg

INJ.

TILCOTIL 20 mg ROCHE ROMANIA S.R.L.

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| 109 |M01AC06| MELOXICAMUM | |

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AC06 MELOXICAMUM COMPR. 15 mg

CELOMIX 15 mg ACTAVIS GROUP HF.

MELARTRIN 15 mg 15 mg TERAPIA SA

MELOX 15 mg 15 mg MEDOCHEMIE LTD

MELOXICAM 15 mg 15 mg VIM SPECTRUM SRL

MELOXICAM LPH 15 mg 15 mg LABORMED PHARMA SA

MELOXICAM MCC 15 mg 15 mg MAGISTRA C & C SRL

MELOXICAM SANDOZ 15 mg 15 mg HEXAL AG

MOVALIS(R) 15 mg 15 mg BOEHRINGER INGELHEIM INT.

GMBH

RECOXA 15 15 mg ZENTIVA AS

M01AC06 MELOXICAMUM SUPOZ. 15 mg

MELOXICAM 15 mg 15 mg MAGISTRA C & C

MOVALIS(R) 15 mg 15 mg BOEHRINGER INGELHEIM INT.

GMBH

M01AC06 MELOXICAMUM SOL. INJ. 15 mg/1.5 ml

MOVALIS(R) 15 mg/1.5 ml BOEHRINGER INGELHEIM INT.

GMBH

M01AC06 MELOXICAMUM COMPR. 7.5 mg

MELOX 7.5 mg 7.5 mg MEDOCHEMIE LTD

M01AC06 MELOXICAMUM COMPR. 7.5 mg

MELOXICAM 7,5 mg 7.5 mg VIM SPECTRUM SRL

MELOXICAM LPH 7,5 mg 7.5 mg LABORMED PHARMA SA

MELOXICAM MCC 7,5 mg 7.5 mg MAGISTRA C & C SRL

MELOXICAM SANDOZ 7,5 mg 7.5 mg HEXAL AG

MOVALIS(R) 7,5 mg 7.5 mg BOEHRINGER INGELHEIM INT.

GMBH

M01AC06 MELOXICAMUM SUPOZ. 7.5 mg

MELOXICAM 7,5 mg 7.5 mg MAGISTRA C & C

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| 110 |M01AE03| KETOPROFENUM | |

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AE03 KETOPROFENUM COMPR. FILM. 100 mg

RUBIFEN 100 mg 100 mg ANTIBIOTICE SA

M01AE03 KETOPROFENUM CAPS. ELIB. PREL. 100 mg

KETOPROFEN SR 100 mg 100 mg TERAPIA SA

M01AE03 KETOPROFENUM COMPR. FILM. 100 mg

KETONAL FORTE 100 mg 100 mg LEK PHARMACEUTICALS D.D.

KETOPROXIN 100 mg 100 mg AC HELCOR PHARMA SRL

PROFENID(R) 100 mg 100 mg LAB. AVENTIS

M01AE03 KETOPROFENUM LIOF. + SOLV. PT. SOL. 100 mg

INJ.

PROFENID(R) 100 mg 100 mg LAB. AVENTIS

M01AE03 KETOPROFENUM SUPOZ. 100 mg

KETOMAG 100 mg MAGISTRA C & C

KETONAL 100 mg 100 mg LEK PHARMACEUTICALS D.D.

RUBIFEN(R) 100 mg 100 mg ANTIBIOTICE SA

M01AE03 KETOPROFENUM SOL. INJ. 100 mg/2 ml

KETONAL 100 mg/2 ml 100 mg/2 ml LEK PHARMACEUTICALS D.D.

M01AE03 KETOPROFENUM SOL. INJ./CONC. SOL. 100 mg/2 ml

PERF.

KETOPROFEN 100 mg/2 ml 100 mg/2 ml TERAPIA SA

M01AE03 KETOPROFENUM CAPS. ELIB. PREL. 150 mg

KETONAL(R) DUO 150 mg 150 mg SANDOZ S.R.L.

M01AE03 KETOPROFENUM COMPR. ELIB. PREL. 150 mg

KETONAL RETARD 150 mg LEK PHARMACEUTICALS D.D.

M01AE03 KETOPROFENUM CAPS. ELIB. PREL. 200 mg

KETALGON 200 mg 200 mg NOVIA FARM SOLUTIONS SRL

KETONAL UNO 200 mg SANDOZ S.R.L.

KETOPROFEN SR 200 mg 200 mg TERAPIA SA

M01AE03 KETOPROFENUM COMPR. FILM. ELIB. 200 mg

PREL.

PROFENID(R) LP 200 mg 200 mg LAB. AVENTIS

M01AE03 KETOPROFENUM COMPR. FILM. 50 mg

KETOPROXIN 50 mg 50 mg AC HELCOR PHARMA SRL

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| 111 |M03BX07| TETRAZEPAMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M03BX07 TETRAZEPAMUM COMPR. FILM. 50 mg

MYOLASTAN(R) 50 mg 50 mg SANOFI-SYNTHELABO FRANCE

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| 112 |M04AA01| ALLOPURINOLUM\* | |

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NOTĂ:

Doza trebuie ajustată în concordanţă cu funcţia renală.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M04AA01 ALLOPURINOLUM COMPR. 100 mg

MILURIT 100 mg 100 mg EGIS PHARMACEUTICALS LTD.

M04AA01 ALLOPURINOLUM COMPR. 300 mg

MILURIT 300 mg 300 mg EGIS PHARMACEUTICALS LTD.

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| 113 |M04AC01| COLCHICINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M04AC01 COLCHICINUM COMPR. 1 mg

COLCHICINA 1 mg BIOFARM SA

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| 114 |N02AD01| PENTAZOCINUM | |

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Prescriere limitată: Durere severă, care nu răspunde la analgezice non-opioide.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N02AD01 PENTAZOCINUM SOL. INJ. 30 mg/ml

FORTRAL 30 mg/ml KRKA D.D. NOVO MESTO

FORTWIN(R) 30 mg/ml TERAPIA S.A.

N02AD01 PENTAZOCINUM COMPR. 50 mg

FORTRAL 50 mg 50 mg KRKA D.D. NOVO MESTO

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| 115 |N02AX02| TRAMADOLUM | |

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Prescriere limitată: Durere severă, care nu răspunde la analgezice

non-opioide.

Pentru durere acută la care tratamentul cu aspirină

şi/sau paracetamol este contraindicat sau nu a dat

rezultate.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice (mai puţin formele injectabile) şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N02AX02 TRAMADOLUM COMPR. ELIB. MODIF. 100 mg

TRAMADOLOR(R) 100 ID 100 mg HEXAL AG

N02AX02 TRAMADOLUM COMPR. ELIB. PREL. 100 mg

TRALGIT SR 100 100 mg ZENTIVA AS

TRAMADOL(R) RETARD 100 mg KRKA D.D.

N02AX02 TRAMADOLUM COMPR. FILM. ELIB. 100 mg

PREL.

TRAMAL RETARD 100 mg 100 mg GRUNENTHAL GMBH

N02AX02 TRAMADOLUM SOL. INJ. 100 mg

MABRON 100 mg MEDOCHEMIE LTD.

TRADOLAN 100 mg LANNACHER HEILMITTEL GMBH

Prescriere limitată: Tratamentul de scurtă durată al durerii acute.

N02AX02 TRAMADOLUM SUPOZ. 100 mg

TRADOLAN 100 mg LANNACHER HEILMITTEL GMBH

TRAMADOL 100 mg KRKA D.D.

TRAMAG 100 100 mg MAGISTRA C & C

TRAMAL(R) 100 mg GRUNENTHAL GMBH

N02AX02 TRAMADOLUM SOL. INJ. 100 mg/2 ml

TRALGIT 100 100 mg/2 ml ZENTIVA A.S.

Prescriere limitată: Tratamentul de scurtă durată al durerii acute.

N02AX02 TRAMADOLUM PIC. ORALE, SOL. 100 mg/ml

TRALGIT 100 mg/ml ZENTIVA A.S.

N02AX02 TRAMADOLUM SOL. ORALA 100 mg/ml

TRADOLAN 100 mg/ml LANNACHER HEILMITTEL GMBH

N02AX02 TRAMADOLUM COMPR. ELIB. PREL. 150 mg

TRALGIT SR 150 150 mg ZENTIVA AS

N02AX02 TRAMADOLUM COMPR. FILM. ELIB. 150 mg

PREL.

TRAMADOL RETARD 150 mg 150 mg KRKA D.D. NOVO MESTO

TRAMAL RETARD 150 mg 150 mg GRUNENTHAL GMBH

N02AX02 TRAMADOLUM COMPR. ELIB. PREL. 200 mg

TRALGIT SR 200 200 mg ZENTIVA AS

N02AX02 TRAMADOLUM COMPR. FILM. ELIB. 200 mg

PREL.

TRAMADOL RETARD 200 mg 200 mg KRKA D.D. NOVO MESTO

TRAMAL RETARD 200 mg 200 mg GRUNENTHAL GMBH

N02AX02 TRAMADOLUM CAPS. 50 mg

K-ALMA(R) 50 mg ANTIBIOTICE SA

MABRON 50 mg 50 mg MEDOCHEMIE LTD.

TRALGIT 50 50 mg ZENTIVA A.S.

TRAMACALM 50 mg AC HELCOR SRL

TRAMADOL 50 mg KRKA D.D.

TRAMADOL LPH 50 mg 50 mg LABORMED PHARMA SA

TRAMADOL AL 50 50 mg ALIUD(R) PHARMA GMBH &

CO.KG

TRAMADOL ARENA 50 mg ARENA GROUP S.A.

TRAMAL(R) 50 mg GRUNENTHAL GMBH

URGENDOL 50 mg MEDICAROM GROUP SRL

N02AX02 TRAMADOLUM COMPR. 50 mg

TRAMADOL 50 mg 50 mg OZONE LABORATORIES LTD.

TRAMADOL EEL 50 mg BIO EEL SRL

TRAMAG 50 50 mg MAGISTRA C & C

N02AX02 TRAMADOLUM COMPR. FILM. 50 mg

TRADOLAN 50 mg 50 mg LANNACHER HEILMITTEL GMBH

N02AX02 TRAMADOLUM SOL. INJ. 50 mg

TRADOLAN 50 mg 50 mg LANNACHER HEILMITTEL GMBH

Prescriere limitată: Tratamentul de scurtă durată al durerii acute.

N02AX02 TRAMADOLUM SOL. INJ. 50 mg/ml

TRALGIT 50 50 mg/ml ZENTIVA A.S.

TRAMADOL 50 mg/ml KRKA D.D.

TRAMADOL(R) AL 100 Fiole 50 mg/ml ALIUD(R) PHARMA GMBH &

CO.KG

TRAMAL(R) 100 50 mg/ml GRUNENTHAL GMBH

TRAMAL(R) 50 50 mg/ml GRUNENTHAL GMBH

URGENDOL 50 mg/ml MEDICAROM GROUP SRL

Prescriere limitată: Tratamentul de scurtă durată al durerii acute.

N02AX02 TRAMADOLUM PIC. ORALE, SOL.

TRAMADOL AL PICATURI ALIUD PHARMA GMBH & CO.KG

N02AX02 TRAMADOLUM PICATURI ORALE - SOL. 100 mg/ml

TRAMAL(R) 100 mg/ml GRUNENTHAL GMBH

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| 116 |N02CC01| SUMATRIPTANUM\* | |

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Prescriere limitată: Atacurile de migrenă la pacienţii care primesc sau au primit medicaţie profilactică şi la care crizele migrenoase nu au răspuns la tratamentul oral cu ergotamină sau alte medicamente antimigrenoase, sau la care aceste medicamente sunt contraindicate.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N02CC01 SUMATRIPTANUM COMPR. 100 mg

SUMIGRA 100 mg 100 mg SANDOZ S.R.L.

N02CC01 SUMATRIPTANUM COMPR. FILM. 100 mg

IMIGRAN(R) 100 100 mg GLAXO WELLCOME UK LTD.

SUMACTA 100 mg 100 mg ACTAVIS GROUP PTC EHF.

SUMATRIPTAN 100 mg DR. REDDY'S LABORATORIES

N02CC01 SUMATRIPTANUM COMPR. 50 mg

SUMIGRA 50 mg 50 mg SANDOZ S.R.L.

N02CC01 SUMATRIPTANUM COMPR. FILM. 50 mg

IMIGRAN(R) 50 50 mg GLAXO WELLCOME UK LTD.

SUMACTA 50 mg 50 mg ACTAVIS GROUP PTC EHF.

SUMATRIPTAN 50 mg DR. REDDY'S LABORATORIES

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| 117 |N03AF01| CARBAMAZEPINUM | Protocol: N025G |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N03AF01 CARBAMAZEPINUM SUSP. ORALA 100 mg/5 ml

TIMONIL SIROP 100 mg/5 ml DESITIN ARZNEIMITTEL GMBH

N03AF01 CARBAMAZEPINUM COMPR. RET. 150 mg

TIMONIL 150 RETARD 150 mg DESITIN

N03AF01 CARBAMAZEPINUM COMPR. 200 mg

CARBAMAZEPIN 200 mg SLAVIA PHARM SRL

CARBAMAZEPIN EEL 200 mg BIO EEL SRL

CARBAMAZEPINA 200 mg OZONE LABORATORIES LTD.

CARBAMAZEPINA ARENA 200 mg 200 mg ARENA GROUP SA

CARBAMAZEPINA LPH 200 mg 200 mg LABORMED PHARMA SA

CARBAVIM 200 mg VIM SPECTRUM SRL

CARBEPSIL 200 200 mg AC HELCOR SRL

FINLEPSIN 200 mg AWD PHARMA GMBH & CO.KG

NEUROTOP 200 mg GEROT PHARMAZEUTIKA GMBH

TAVER 200 mg MEDOCHEMIE LTD.

TEGRETOL(R) 200 200 mg NOVARTIS PHARMA GMBH

N03AF01 CARBAMAZEPINUM COMPR. FILM. ELIB. 200 mg

PREL.

TEGRETOL CR 200 200 mg NOVARTIS PHARMA GMBH

N03AF01 CARBAMAZEPINUM COMPR. ELIB. PREL. 300 mg

NEUROTOP(R) RETARD 300 mg 300 mg GEROT PHARMAZEUTIKA GMBH

N03AF01 CARBAMAZEPINUM COMPR. RET. 300 mg

TIMONIL 300 RETARD 300 mg DESITIN

N03AF01 CARBAMAZEPINUM COMPR. 400 mg

CARBEPSIL 400 400 mg AC HELCOR SRL

TEGRETOL(R) 400 400 mg NOVARTIS PHARMA GMBH

N03AF01 CARBAMAZEPINUM COMPR. FILM. ELIB. 400 mg

PREL.

TEGRETOL CR 400 400 mg NOVARTIS PHARMA GMBH

N03AF01 CARBAMAZEPINUM COMPR. RET. 400 mg

FINLEPSIN 400 RETARD 400 mg AWD PHARMA GMBH & CO.KG

N03AF01 CARBAMAZEPINUM COMPR. ELIB. PREL. 600 mg

NEUROTOP(R) RETARD 600 mg 600 mg GEROT PHARMAZEUTIKA GMBH

N03AF01 CARBAMAZEPINUM COMPR. RET. 600 mg

TIMONIL 600 RETARD 600 mg DESITIN

N03AF01 CARBAMAZEPINUM COMPR. FILM. ELIB. 200 mg

PREL.

FINLEPSIN 200 RETARD 200 mg AWD PHARMA GMBH & CO.KG

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| 118 |N03AX12| GABAPENTINUM | Protocol: N025G |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N03AX12 GABAPENTINUM CAPS. 100 mg

GABAGAMMA 100 mg 100 mg WORWAG PHARMA GMBH & CO.KG

GABALEPT 100 mg 100 mg PLIVA LJUBLJANA D.O.O.

N03AX12 GABAPENTINUM CAPS. 300 mg

GABAGAMMA 300 mg 300 mg WORWAG PHARMA GMBH & CO.KG

GABALEPT 300 mg 300 mg PLIVA LJUBLJANA D.O.O.

GABARAN 300 mg 300 mg RANBAXY UK LTD.

N03AX12 GABAPENTINUM CAPS. 400 mg

GABAGAMMA 400 mg 400 mg WORWAG PHARMA GMBH & CO.KG

GABALEPT 400 mg 400 mg PLIVA LJUBLJANA D.O.O.

GABARAN 400 mg 400 mg RANBAXY UK LTD.

N03AX12 GABAPENTINUM COMPR. FILM. 600 mg

GABARAN 600 mg 600 mg RANBAXY UK LTD.

N03AX12 GABAPENTINUM COMPR. FILM. 800 mg

GABARAN 800 mg 800 mg RANBAXY UK LTD.

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| 119 |N05AD01| HALOPERIDOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05AD01 HALOPERIDOLUM SOL. INJ. 50 mg/ml

HALOPERIDOL DECANOAT 50 mg/ml GEDEON RICHTER LTD.

N05AD01 HALOPERIDOLUM COMPR. 5 mg

HALDOL 5 mg JANSSEN PHARMACEUTICA NV

N05AD01 HALOPERIDOLUM SOL. INJ. 5 mg/ml

HALOPERIDOL 5 mg/ml 5 mg/ml GEDEON RICHTER LTD.

N05AD01 HALOPERIDOLUM PIC. ORALE - SOL. 2 mg/ml

HALOPERIDOL 2 mg/ml 2 mg/ml GEDEON RICHTER ROMANIA SA

N05AD01 HALOPERIDOLUM PICATURI ORALE - SOL. 2 mg/ml

HALOPERIDOL 2 mg/ml TERAPIA SA

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| 120 |N05BA01| DIAZEPAMUM | |

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NOTĂ:

Diazepamum este utilizat ca agent anxiolitic, anticonvulsivant şi relaxant muscular de tip central. În tratamentul anxietăţii se utilizează ca sedativ în anxietatea acută de tip sever cât şi în tratamentul agitaţiei asociată cu delirium tremens.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05BA01 DIAZEPAMUM COMPR. 10 mg

DIAZEPAM 10 mg TERAPIA SA

DIAZEPAM 10 mg 10 mg GEDEON RICHTER ROMANIA SA

N05BA01 DIAZEPAMUM SOL. RECTALA 10 mg/2.5 ml

DIAZEPAM DESITIN(R) SOLUŢIE 10 mg/2.5 ml DESITIN

RECTALA 10 mg

N05BA01 DIAZEPAMUM SOL. INJ. 10 mg/2 ml

DIAZEPAM 10 mg 10 mg/2 ml TERAPIA SA

N05BA01 DIAZEPAMUM SOL. RECTALA 5 mg/2.5 ml

DIAZEPAM DESITIN(R) SOLUŢIE 5 mg/2.5 ml DESITIN

RECTALĂ 5 mg

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| 121 |N05BA03| MEDAZEPAMUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05BA03 MEDAZEPAMUM COMPR. 10 mg

MEDAZEPAM 10 mg ARENA GROUP SA

N05BA03 MEDAZEPAMUM CAPS. 10 mg

ANSILAN 10 mg 10 mg LEK PHARMACEUTICALS D.D.

N05BA03 MEDAZEPAMUM COMPR. 10 mg

MEDAZEPAM 10 mg 10 mg LABORMED PHARMA SA

RUDOTEL 10 mg AWD PHARMA GMBH & CO.KG

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| 122 |N05BA06| LORAZEPAMUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05BA06 LORAZEPAMUM COMPR. 1 mg

ANXIAR(R) 1 mg 1 mg GEDEON RICHTER ROMANIA SA

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| 123 |N05BA08| BROMAZEPAMUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05BA08 BROMAZEPAMUM COMPR. 1.5 mg

BROMAZEPAM LPH 1.5 mg 1.5 mg LABORMED PHARMA SA

CALMEPAM(R) 1.5 mg GLAXOSMITHKLINE (GSK) SRL

N05BA08 BROMAZEPAMUM COMPR. 3 mg

BROMAZEPAM LPH 3 mg 3 mg LABORMED PHARMA SA

CALMEPAM(R) 3 mg GLAXOSMITHKLINE (GSK) SRL

LEXOTAN 3 mg ROCHE ROMANIA S.R.L.

LEXOTANIL 3 mg TERAPIA S.A.

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| 124 |N05BA12| ALPRAZOLAMUM\* | |

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Prescriere limitată: Tulburări de panică, în cazul eşecului la tratamente similare.

Pentru tratamentul de scurtă durată al anxietăţii moderate sau severe şi al anxietăţii asociate cu depresia.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05BA12 ALPRAZOLAMUM COMPR. 0,25 mg

XANAX 0,25 mg 0,25 mg PFIZER EUROPE MA EEIG

N05BA12 ALPRAZOLAMUM COMPR. 0,5 mg

XANAX 0,5 mg 0,5 mg PFIZER EUROPE MA EEIG

N05BA12 ALPRAZOLAMUM COMPR. 0.25 mg

ALPRAZOLAMLPH(R) 0,25 mg 0.25 mg LABORMED PHARMA SA

FRONTIN 0,25 mg 0.25 mg EGIS PHARMACEUTICALS PLC

PRAZOLEX(R) 0,25 mg 0.25 mg GEDEON RICHTER ROMANIA SA

N05BA12 ALPRAZOLAMUM COMPR. 0.5 mg

ALPRAZOLAM LPH(R) 0,5 mg 0.5 mg LABORMED PHARMA SA

FRONTIN 0,5 mg 0.5 mg EGIS PHARMACEUTICALS PLC

PRAZOLEX(R) 0,5 mg 0.5 mg GEDEON RICHTER ROMANIA SA

N05BA12 ALPRAZOLAMUM COMPR. ELIB. PREL. 0.5 mg

NEUROL(R) SR 0.5 0.5 mg ZENTIVA AS

N05BA12 ALPRAZOLAMUM COMPR. 1 mg

ALPRAZOLAM LPH(R) 1 mg 1 mg LABORMED PHARMA SA

FRONTIN 1 mg EGIS PHARMACEUTICALS PLC

PRAZOLEX(R) 1 mg 1 mg GEDEON RICHTER ROMANIA SA

XANAX 1 mg 1 mg PFIZER EUROPE MA EEIG

N05BA12 ALPRAZOLAMUM COMPR. ELIB. PREL. 1 mg

NEUROL(R) SR 1 1 mg ZENTIVA AS

N05BA12 ALPRAZOLAMUM COMPR. 2 mg

XANAX 2 mg 2 mg PFIZER EUROPE MA EEIG

N05BA12 ALPRAZOLAMUM COMPR. ELIB. PREL. 2 mg

NEUROL(R) SR 2 2 mg ZENTIVA AS

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| 125 |N05CD02| NITRAZEPAMUM | |

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Cod restricţie 3007: Tratamentul de scurtă durată al insomniei.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05CD02 NITRAZEPAMUM COMPR. 2.5 mg

NITRAZEPAM LPH(R) 2,5 mg 2.5 mg LABORMED PHARMA SA

N05CD02 NITRAZEPAMUM COMPR. 5 mg

NITRAZEPAM 5 mg 5 mg GEDEON RICHTER ROMANIA SA

NITRAZEPAM LPH(R) 5 mg 5 mg LABORMED PHARMA SA

N05CD02 NITRAZEPAMUM COMPR. 5 mg

NITRAZEPAM 5 mg 5 mg GEDEON RICHTER ROMANIA SA

NITRAZEPAM LPH(R) 5 mg 5 mg LABORMED PHARMA SA

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| 126 |N05CF01| ZOPICLONUM\* | |

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Cod restricţie 3007: Pentru tratamentul de scurtă durată al insomniei.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05CF01 ZOPICLONUM COMPR. FILM. 7.5 mg

ALS-ZOPICLON 7.5 mg 7.5 mg ALSIFCOM INTERMED SRL

IMOVANE 7.5 mg 7.5 mg SANOFI AVENTIS

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| 127 |N05CF02| ZOLPIDEMUM\* | |

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Cod restricţie 3007: Pentru tratamentul de scurtă durată al insomniei.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05CF02 ZOLPIDEMUM COMPR. FILM. 10 mg

HYPNOGEN 10 mg 10 mg ZENTIVA AS

LADINOX 10 mg GEDEON RICHTER ROMANIA

S.A.

SANVAL(R) 10 mg 10 mg LEK PHARMACEUTICALS D.D.

ZOLPIDEM 10 mg 10 mg LABORMED PHARMA SA

ZOLPIDEM ANTIBIOTICE 10 mg 10 mg ANTIBIOTICE S.A.

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| 128 |N06AA04| CLOMIPRAMINUM\* | |

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Prescriere limitată: Tratamentul simptomatic al depresiei în special atunci

când este necesară sedarea.

Tratamentul stărilor fobice şi obsesiv-compulsive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AA04 CLOMIPRAMINUM DRAJ. 10 mg

ANAFRANIL(R) 10 mg 10 mg NOVARTIS PHARMA GMBH

N06AA04 CLOMIPRAMINUM DRAJ. 25 mg

ANAFRANIL(R) 25 mg 25 mg NOVARTIS PHARMA GMBH

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| 129 |N06AA09| AMITRIPTYLINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AA09 AMITRIPTYLINUM CAPS. RET. 25 mg

AMITRIPTILIN 25 R. DESITIN 25 mg DESITIN

N06AA09 AMITRIPTYLINUM COMPR. FILM. 25 mg

AMITRIPTILINA ARENA 25 mg 25 mg ARENA GROUP S.A.

N06AA09 AMITRIPTYLINUM CAPS. RET. 50 mg

AMITRIPTILIN 50 R. DESITIN 50 mg DESITIN

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| 130 |N06AA12| DOXEPINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AA12 DOXEPINUM DRAJ. 25 mg

DOXEPIN 25 mg 25 mg TERAPIA SA

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| 131 |N06AB03| FLUOXETINUM\* | |

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Prescriere limitată: Tratamentul depresiilor cu sau fără anxietate în

special când componenta de sedare nu este necesară.

Tratamentul bulimiei nervoase.

Tratamentul tulburărilor obsesiv-compulsive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AB03 FLUOXETINUM CAPS. 20 mg

FLUOXETINE 20 mg 20 mg DR. REDDY'S LABORATORIES

FLUOXIN 20 mg VIM SPECTRUM SRL

FLURAN 20 mg RANBAXY U.K. LIMITED

MAGRILAN 20 mg 20 mg MEDOCHEMIE LTD.

N06AB03 FLUOXETINUM COMPR. DISP. 20 mg

PROZAC 20 mg 20 mg ELI LILLY SA

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| 132 |N06AB04| CITALOPRAMUM\*\* | Protocol: N008F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AB04 CITALOPRAMUM COMPR. FILM. 10 mg

CITALORAN 10 mg 10 mg RANBAXY UK LIMITED

LINISAN 10 mg 10 mg GEDEON RICHTER ROMANIA

S.A.

N06AB04 CITALOPRAMUM COMPR. FILM. 20 mg

CITALORAN 20 mg 20 mg RANBAXY UK LIMITED

LINISAN 20 mg 20 mg GEDEON RICHTER ROMANIA

S.A.

N06AB04 CITALOPRAMUM COMPR. FILM. 40 mg

CITALORAN 40 mg 40 mg RANBAXY UK LIMITED

LINISAN 40 mg 40 mg GEDEON RICHTER ROMANIA

S.A.

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| 133 |N06AB05| PAROXETINUM\* | |

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Prescriere limitată: Tratamentul tulburărilor depresive majore.

Tratamentul tulburărilor obsesiv-compulsive.

Tratamentul atacurilor de panică.

Tratamentul anxietăţii.

Tratamentul fobiei sociale.

Tratamentul bolii de stres posttraumatic.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AB05 PAROXETINUM COMPR. 20 mg

ARKETIS 20 mg MEDOCHEMIE LTD.

N06AB05 PAROXETINUM COMPR. FILM. 20 mg

ALS-PAROXETIN 20 mg 20 mg ALSIFCOM INTERMED SRL

PALUXETIL 20 mg 20 mg HEXAL AG

PAXETEN 20 mg 20 mg ACTAVIS GROUP HF.

REXETIN 20 mg GEDEON RICHTER PLC.

SEROXAT 20 mg 20 mg SMITHKLINE BEECHAM PLC

N06AB05 PAROXETINUM COMPR. FILM. 40 mg

PALUXETIL 40 mg 40 mg HEXAL AG

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| 134 |N06AB06| SERTRALINUM\* | |

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Prescriere limitată: Tratamentul tulburărilor depresive inclusiv cu

componenta anxioasă.

Tratamentul tulburări obsesiv-compulsive.

Tratamentul atacurilor de panică.

Tratamentul fobiei sociale.

Tratamentul bolii de stres posttraumatic.

Eficienţa tratamentului în boala de stres posttraumatic a fost demonstrată doar pentru sexul feminin.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AB06 SERTRALINUM COMPR. FILM. 100 mg

ALS-SERTRALINA 100 mg 100 mg ALSIFCOM INTERMED SRL

ASENTRA(R) 100 mg KRKA D.D.

SERLIFT 100 mg 100 mg TERAPIA S.A.

SERTRALIN SANDOZ 100 mg 100 mg HEXAL AG

SERTRALINA 100 mg 100 mg ARENA GROUP S.A.

SERTRALINA DR. REDDY'S 100 mg 100 mg DR. REDDY'S LABORATORIES

STIMULOTON(R) 100 mg 100 mg EGIS PHARMACEUTICALS LTD.

N06AB06 SERTRALINUM SOL. ORALA 20 mg/ml

ZOLOFT(R) 20 mg/ml PFIZER EUROPE MA EEIG

N06AB06 SERTRALINUM COMPR. FILM. 50 mg

ALS-SERTRALINA 50 mg 50 mg ALSIFCOM INTERMED SRL

ASENTRA(R) 50 mg KRKA D.D.

SERLIFT 50 mg 50 mg TERAPIA S.A.

SERTRALIN 50 mg 50 mg OZONE LABORATORIES LTD.

SERTRALIN SANDOZ 50 mg 50 mg HEXAL AG

SERTRALINA 50 mg 50 mg ARENA GROUP SA

SERTRALINA DR. REDDY'S 50 mg 50 mg DR. REDDY'S LABORATORIES

STIMULOTON(R) 50 mg 50 mg EGIS PHARMACEUTICALS LTD.

ZOLOFT 50 mg 50 mg PFIZER EUROPE MA EEIG

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| 135 |N06AB08| FLUVOXAMINUM\* | |

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Prescriere limitată: Tratamentul tulburărilor depresive majore.

Tratamentul tulburărilor obsesiv-compulsive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AB08 FLUVOXAMINUM COMPR. FILM. 100 mg

FEVARIN(R) 100 100 mg SOLVAY PHARMACEUTICALS BV

FLUVOXAMINE TEVA 100 mg 100 mg TEVA PHARMACEUTICALS SRL

N06AB08 FLUVOXAMINUM COMPR. FILM. 50 mg

FEVARIN(R) 50 50 mg SOLVAY PHARMACEUTICALS BV

FLUVOXAMINE TEVA 50 mg 50 mg TEVA PHARMACEUTICALS SRL

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| 136 |N06AX03| MIANSERINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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N06AX03 MIANSERINUM COMPR. FILM. 10 mg

MIANSERIN 10 10 mg REMEDICA LTD.

N06AX03 MIANSERINUM DRAJ. 10 mg

MIANSERIN 10 mg 10 mg TERAPIA SA

N06AX03 MIANSERINUM COMPR. FILM. 30 mg

MIANSERIN 30 30 mg REMEDICA LTD.

N06AX03 MIANSERINUM DRAJ. 30 mg

MIANSERIN 30 mg 30 mg TERAPIA SA

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| 137 |N06AX11| MIRTAZAPINUM\* | |

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Prescriere limitată: Tratamentul episoadelor depresive majore.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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N06AX11 MIRTAZAPINUM COMPR. DISP. 15 mg

REMERON(R) SOLTAB 15 mg 15 mg ORGANON NV

N06AX11 MIRTAZAPINUM COMPR. FILM. 15 mg

ESPRITAL(R) 15 15 mg ZENTIVA AS

MIRZATEN(R) 15 mg 15 mg KRKA D.D.

N06AX11 MIRTAZAPINUM COMPR. DISP. 30 mg

REMERON(R) SOLTAB 30 mg 30 mg ORGANON NV

N06AX11 MIRTAZAPINUM COMPR. FILM. 30 mg

ESPRITAL(R) 30 30 mg ZENTIVA AS

MIRTAZAPINE-TEVA 30 mg 30 mg TEVA PHARMACEUTICALS SRL

MIRZATEN(R) 30 mg 30 mg KRKA D.D.

PHARMATAZ 30 mg 30 mg ACTAVIS GROUP HF.

N06AX11 MIRTAZAPINUM COMPR. DISP. 45 mg

REMERON(R) SOLTAB 45 mg 45 mg ORGANON NV

N06AX11 MIRTAZAPINUM COMPR. FILM. 45 mg

ESPRITAL(R) 45 45 mg ZENTIVA AS

MIRZATEN(R) 45 mg 45 mg KRKA D.D.

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| 138 |N06AX16| VENLAFAXINUM\*\* | Protocol: N013F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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N06AX16 VENLAFAXINUM CAPS. ELIB. PREL. 150 mg

EFECTIN ER 150 mg 150 mg WYETH LEDERLE PHARMA GMBH

VELAXIN 150 mg 150 mg EGIS PHARMACEUTICALS

P.L.C.

N06AX16 VENLAFAXINUM COMPR. 37.5 mg

VELAXIN 37,5 mg 37.5 mg EGIS PHARMACEUTICALS PLC

VENLAFAXINA LPH 37,5 mg 37.5 mg LABORMED PHARMA SA

N06AX16 VENLAFAXINUM CAPS. ELIB. PREL. 75 mg

EFECTIN ER 75 mg 75 mg WYETH LEDERLE PHARMA GMBH

VELAXIN 75 mg 75 mg EGIS PHARMACEUTICALS

P.L.C.

N06AX16 VENLAFAXINUM COMPR. 75 mg

VELAXIN 75 mg 75 mg EGIS PHARMACEUTICALS PLC

VENLAFAXINA LPH 75 mg 75 mg LABORMED PHARMA SA

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| 139 |N07AA01| NEOSTIGMINI BROMIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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N07AA01 NEOSTIGMINI BROMIDUM SOL. INJ. 0.5 mg/ml

MIOSTIN 0,5 mg/ml 0.5 mg/ml ZENTIVA S.A.

N07AA01 NEOSTIGMINI BROMIDUM COMPR. 15 mg

NEOSTIGMINA LPH 15 mg 15 mg LABORMED PHARMA SA

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| 140 |P01AB01| METRONIDAZOLUM | |

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Prescriere limitată: Tratamentul infecţiilor cu anaerobi.

Tratamentul infecţiilor cu protozoare sensibile la

metronidazol.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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P01AB01 METRONIDAZOLUM COMPR. 250 mg

METRONIDAZOL 250 mg (P01AB01) 250 mg ZENTIVA SA

METRONIDAZOL 250 mg (P01AB1) 250 mg ARENA GROUP SA

P01AB01 METRONIDAZOLUM COMPR. FILM. 250 mg

FLAGYL 250 mg LABORATOIRE AVENTIS

P01AB01 METRONIDAZOLUM SUSP. ORALA 4%

FLAGYL 4% (P01AB01) 4% LABORATOIRE AVENTIS

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| 141 |P01AB02| TINIDAZOLUM | |

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Prescriere limitată: Tratamentul infecţiilor cu anaerobi.

Tratamentul infecţiilor cu protozoare sensibile la

metronidazol.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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P01AB02 TINIDAZOLUM COMPR. FILM. 500 mg

FASIGYN (P01AB02) 500 mg PFIZER EUROPE MA EEIG

TINIZOL(R) 500 mg (P01AB02) 500 mg ZENTIVA S.A.

TIPROGYN 500 500 mg AC HELCOR SRL

P01AB02 TINIDAZOLUM COMPR. FILM. 500 mg

FASIGYN (P01AB02) 500 mg PFIZER EUROPE MA EEIG

TINIZOL(R) 500 mg (J01XD02) 500 mg ZENTIVA S.A.

TIPROGYN 500 500 mg AC HELCOR SRL

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| 142 |P02CA03| ALBENDAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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P02CA03 ALBENDAZOLUM SUSP. ORALA 0.4 g/10 ml

ZENTEL 0,4 g/10 ml 0.4 g/10 ml GLAXOSMITHKLINE EXPORT

LTD.

P02CA03 ALBENDAZOLUM COMPR. FILM. 200 mg

DUADOR 200 mg 200 mg GEDEON RICHTER ROMANIA SA

ZENTEL 200 mg 200 mg GLAXOSMITHKLINE EXPORT

LTD.

P02CA03 ALBENDAZOLUM COMPR. 400 mg

ESKAZOLE 400 mg GLAXO SMITHKLINE EXPORT

LTD.

P02CA03 ALBENDAZOLUM SUSP. ORALA 400 mg

ALBENDAZOL BIOFARM 400 mg 400 mg BIOFARM S.A.

P02CA03 ALBENDAZOLUM COMPR. FILM 200 mg

ALBENDAZOL 200 mg OZONE LABORATORIES LTD.

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| 143 |P02CE01| LEVAMISOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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P02CE01 LEVAMISOLUM COMPR. 150 mg

DECARIS 150 mg 150 mg GEDEON RICHTER ROMANIA SA

P02CE01 LEVAMISOLUM COMPR. 50 mg

DECARIS 50 mg 50 mg GEDEON RICHTER ROMANIA SA

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| 144 |R03AC02| SALBUTAMOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03AC02 SALBUTAMOLUM SUSP. INHAL PRESURIZATĂ 100 micrograme/doză

ASTHALIN INHALER 100 micrograme/doză CIPLA (UK) LIMITED

ECOSAL Easi-Breath 100 micrograme/doză IVAX-PHARMACEUTICALS

S.R.O.

VENTOLIN 100 INHALER CFC-FREE 100 micrograme/doză GLAXOWELLCOME UK LTD.

R03AC02 SALBUTAMOLUM SOL. INHAL. 5 mg/ml

VENTOLIN(R) 5 mg/ml GLAXOWELLCOME UK LTD.

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| 145 |R03BA01| BECLOMETASONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03BA01 BECLOMETASONUM AEROSOL SOL. INHAL. 100 micrograme/doză

ECOBEC 100 micrograme 100 micrograme/doză IVAX-PHARMACEUTICALS

CFC FREE S.R.O.

ECOBEC EASI-BREATHE 100 micrograme/doză IVAX-PHARMACEUTICALS

100 micrograme CFC FREE S.R.O.

R03BA01 BECLOMETASONUM SPRAY NAZ., SUSP. 100 micrograme/doză

RINOCLENIL 100 100 micrograme/doză CHIESI FARMACEUTICI S.P.A.

R03BA01 BECLOMETASONUM AEROSOL SOL. INHAL. 250 micrograme/doză

ECOBEC 250 micrograme/doză 250 micrograme/doză IVAX-PHARMACEUTICALS

CFC FREE S.R.O.

ECOBEC EASI-BREATHE 250 micrograme/doză IVAX-PHARMACEUTICALS

250 micrograme/doză CFC FREE S.R.O.

R03BA01 BECLOMETASONUM SOL. DE INHALAT 250 micrograme/doză

PRESURIZATĂ

BECLOFORTER(R) CFC-Free 250 micrograme/doză GLAXO WELLCOME UK LTD.

R03BA01 BECLOMETASONUM SOL. INHALAT 250 micrograme/doză

PRESURIZATĂ

CLENIL(R) JET 250 micrograme/doză CHIESI FARMACEUTICI SPA

250 micrograme/doză

R03BA01 BECLOMETASONUM SOL. DE INHALAT 50 micrograme/doză

PRESURIZATĂ

BECOTIDER(R) CFC-Free 50 micrograme/doză GLAXO WELLCOME UK LTD.

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| 146 |R03BA02| BUDESONIDUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03BA02 BUDESONIDUM PULB. INHAL. 200 micrograme/doză

FRENOLYN 200 200 micrograme/doză MEDOCHEMIE ROMÂNIA SRL

R03BA02 BUDESONIDUM PULB. INHAL. 200 micrograme/doză

PULMICORT TURBUHALER 200 micrograme/doză ASTRAZENECA AB

200 micrograme/doză

R03BA02 BUDESONIDUM PULB. INHAL. 400 micrograme/doză

FRENOLYN 400 400 micrograme/doză MEDOCHEMIE ROMÂNIA SRL

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| 147 |R03BB01| IPRATROPII BROMIDUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03BB01 IPRATROPII BROMIDUM AEROSOL 20 micrograme/doză

IPRAVENT 20 - INHALER 20 micrograme/doză CIPLA (UK) LIMITED

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| 148 |R03CC02| SALBUTAMOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03CC02 SALBUTAMOLUM SOL. ORALA 0.04%

SALBUTAMOL T 0.04% TIS FARMACEUTIC SA

R03CC02 SALBUTAMOLUM SOL. INJ. 0.5 mg/ml

VENTOLIN(R) 0.5 mg/ml GLAXO WELLCOME UK LTD.

R03CC02 SALBUTAMOLUM SIROP 2 mg/5 ml

SALBUTAMOL EIPICO 2 mg/5 ml 2 mg/5 ml E.I.P.I.CO. MED S.R.L.

VENTOLIN(R) 2 mg/5 ml GLAXOWELLCOME UK LTD.

R03CC02 SALBUTAMOLUM SOL. ORALA 2 mg/5 ml

SALBUTAMOL 2 mg/5 ml TERAPIA S.A.

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| 149 |R03DA04| THEOPHYLLINUM | |

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Datorită efectelor variabile ale alimentelor asupra absorbţiei teofilinei, pacienţii care folosesc un anumit preparat nu trebuie să îi schimbe cu un altul fără o monitorizare adecvată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03DA04 THEOPHYLLINUM CAPS. ELIB. PREL. 100 mg

TEOFIUNA SR 100 mg 100 mg TERAPIA SA

THEO SR 100 100 mg GLAXOSMITHKLINE (GSK) SRL

R03DA04 THEOPHYLLINUM CAPS. ELIB. PREL. 200 mg

TEOFIUNA SR 200 mg 200 mg TERAPIA SA

TEOTARD(R) 200 200 mg KRKA D.D.

THEO SR 200 200 mg GLAXOSMITHKLINE (GSK) SRL

R03DA04 THEOPHYLLINUM CAPS. ELIB. PREL. 300 mg

TEOFILINA SR 300 mg 300 mg TERAPIA SA

THEO SR 300 300 mg GLAXOSMITHKLINE (GSK) SRL

R03DA04 THEOPHYLLINUM CAPS. ELIB. PREL. 350 mg

TEOTARD(R) 350 350 mg KRKA D.D.

R03DA04 THEOPHYLLINUM CAPS. ELIB. PREL. 50 mg

TEOFILINA SR 50 mg 50 mg TERAPIA SA

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| 150 |R05DA04| CODEINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R05DA04 CODEINUM COMPR. 15 mg

CODEINA FOSFAT 15 mg SLAVIA PHARM SRL

CODEINA FOSFAT 15 mg 15 mg OZONE LABORATORIES LTD.

CODEINA FOSFAT LPH 15 mg 15 mg LABORMED PHARMA SA

CODEINA FOSFORICĂ 15 mg BIO EEL SRL

CODEINA FOSFORICĂ 15 mg 15 mg MAGISTRA C & C

FARMACOD 15 mg FARMACOM SA

FOSFAT DE CODEINĂ 15 mg 15 mg SINTOFARM SA

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| 151 |R06AB04| CHLORPHENAMINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R06AB04 CHLORPHENAMINUM COMPR. 4 mg

CLORFENIRAMIN 4 mg LABORMED PHARMA SA

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| 152 |R06AE07| CETIRIZINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R06AE07 CETIRIZINUM COMPR. FILM. 10 mg

CELERG 10 mg AC HELCOR PHARMA SRL

R06AE07 CETIRIZINUM COMPR. FILM. 10 mg

LETIZEN(R) 10 mg 10 mg KRKA D.D.

R06AE07 CETIRIZINUM SOL. ORALA 1 mg/ml

LETIZEN 1 mg/ml KRKA D.D.

R06AE07 CETIRIZINUM PICĂTURI ORALE - SOL. 10 mg/ml

ZYRTEC(R) 10 mg/ml U.C.B. GMBH

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| 153 |R06AX17| KETOTIFENUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R06AX17 KETOTIFENUM CAPS. 1 mg

KETOF 1 mg 1 mg HEXAL AG

R06AX17 KETOTIFENUM COMPR. 1 mg

H-KETOTIFEN 1 mg AC HELCOR SRL

KETOTIFEN 1 mg MAGISTRA C & C

KETOTIFEN LPH(R) 1 mg 1 mg LABORMED PHARMA SA

R06AX17 KETOTIFENUM SIROP 1 mg/5 ml

FRENASMA 1 mg/5 ml FARAN LABORATORIES S.A.

KETOF 1 mg/5 ml 1 mg/5 ml HEXAL AG

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| 154 |S01AX13| CIPROFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01AX13 CIPROFLOXACINUM UNG. OFT. 0.3%

CIPLOX 0.3% CIPLA (UK) LIMITED

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| 155 |S01BC03| DICLOFENACUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01BC03 DICLOFENACUM PICĂTURI OFT. - SOL. 0.1%

DICLOGESIC 0,1% PICĂTURI 0.1% DAR AL DAWA PHARMA S.R.L.

OFTALMICE, SOLUŢIE

UNICLOPHEN(R) 0,1% 0.1% UNIMED PHARMA LTD.

VOLTAREN OPHTHA CD 0.1% NOVARTIS PHARMA GMBH

S01BC03 DICLOFENACUM PIC. OFT. - SOL. 1 mg/ml

DICLOFENAC RPH 1 mg/ml ROMPHARM COMPANY SRL

S01BC03 DICLOFENACUM PIC. OFT. - SOL. 1 mg/ml

VURDON 1 mg/ml HELP S.A. PHARMACEUTICALS

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| 156 |S01CA05| COMBINAŢII (BETHAMETASONUM + | |

| | | ANTIINFECŢIOASE) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01CA05 COMBINAŢII UNG. OFT. 0,2 g + 0,5 g

(BETHAMETASONUM +

ANTIINFECŢIOASE)

BETABIOPTAL 0,2 g + 0,5 g FARMILA FARMACEUTICI

S01CA05 COMBINAŢII PICĂTURI OFT. SUSP. 0,2 g + 0,5 g

(BETHAMETASONUM +

ANTIINFECŢIOASE)

BETABIOPTAL 0,2 g + 0,5 g FARMILA FARMACEUTICI

MILANO SPA

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SUBLISTA B - MEDICAMENTE CU NIVEL DE COMPENSARE 50% DIN PREŢUL DE REFERINŢĂ

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| 157 |A02BC02| PANTOPRAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BC02 PANTOPRAZOLUM COMPR. FILM. GASTROREZ. 20 mg

CONTROLOC 20 mg 20 mg ALTANA PHARMA AG

Prescriere limitată: Menţinerea rezultatelor terapiei refluxului gastro-esofagian

Prescriere limitată: Boala de reflux gastro-esofagian

Prescriere limitată: Profilaxia ulcerului gastro-duodenal asociat tratamentului cu antiinflamatoare nesteroidiene pe termen lung la pacienţii cu factori de risc gastrointestinal.

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiei de 20 mg.

A02BC02 PANTOPRAZOLUM COMPR. FILM. GASTROREZ. 40 mg

CONTROLOC 40 mg 40 mg ALTANA PHARMA AG

Prescriere limitată: Boala de reflux gastro-esofagian.

Prescriere limitată: Sindromul Zollinger-Ellison

Prescriere limitată: Tratamentul ulcerului gastric şi duodenal.

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiei de 40 mg.

A02BC02 PANTOPRAZOLUM LIOF. PT. SOL. INJ. 40 mg

CONTROLOC 40 mg LIOFILIZAT 40 mg ALTANA PHARMA AG

PENTRU SOLUŢIE INJECTABILĂ

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| 158 |A02BC05| ESOMEPRAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BC05 ESOMEPRAZOLUM COMPR. FILM. GASTROREZ. 40 mg

NEXIUM 40 mg 40 mg ASTRAZENECA AB

Prescriere limitată: Terapia refluxului gastro-esofagian.

Prescriere limitată: Sindromul Zollinger-Ellison.

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiei de 40 mg.

A02BC05 ESOMEPRAZOLUM LIOF. PT. SOL. INJ./ 40 mg

PERF.

NEXIUM(R) 40 mg ASTRAZENECA AB

A02BC05 ESOMEPRAZOLUM COMPR. FILM. GASTROREZ. 20 mg

NEXIUM 20 mg 20 mg ASTRAZENECA AB

Prescriere limitată: Tratamentul ulcerului gastric şi duodenal.

Prescriere limitată: Menţinerea rezultatelor terapiei refluxului gastro-esofagian.

Prescriere limitată: Profilaxia ulcerului gastro-duodenal asociat tratamentului cu antiinflamatoare nesteroidiene pe termen lung la pacienţii cu factori de risc gastrointestinali.

Se aplică pentru toate denumirile comerciale şi formele farmaceutice corespunzătoare concentraţiei de 20 mg.

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| 159 |A02BX02| SUCRALFATUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A02BX02 SUCRALFATUM COMPR. 1 g

GASTROFAIT 1 g E.I.P.I.CO. MED S.R.L.

SUCRALAN(R) 1 g LANNACHER HEILMITTEL GMBH

VENTER 1 g KRKA D.D.

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| 160 |A02BX05| BISMUTHI SUBCITRAS | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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A02BX05 BISMUTHI SUBCITRAS COMPR. FILM. 120 mg

DE-NOL 120 mg ASTELLAS PHARMA EUROPE

B.V.

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| 161 |A03AB06| OTILONIUM BROMIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A03AB06 OTILONIUM BROMIDUM COMPR. FILM. 40 mg

SPASMOMEN(R) 40 mg A. MENARINI IND. FARM.

RIUNITE SRL

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| 162 |A06AD15| MACROGOLUM\* (2) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A06AD15 MACROGOLUM PULB. PT. SOL. ORALĂ

FORTRANS(R) BEAUFOUR IPSEN INT.

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| 163 |A07AA11| RIFAXIMINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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A07AA11 RIFAXIMINUM COMPR. FILM. 200 mg

NORMIX 200 mg 200 mg ALFA WASSERMANN SPA

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| 164 |A07EA06| BUDESONIDUM\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Inducerea remisiunii formelor uşoare/moderate de boala Crohn cu afectare ileală sau/şi de colon ascendent.

A07EA06 BUDESONIDUM CAPS. GASTROREZ. 3 mg

BUDENOFALK 3 mg DR. FALK PHARMA GMBH

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| 165 |A07XA04| RACECADOTRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A07XA04 RACECADOTRILUM CAPS. 100 mg

HIDRASEC 100 mg 100 mg LAB. FOURNIER SA

A07XA04 RACECADOTRILUM PULB. PT. SOL. ORALĂ 10 mg

HIDRASEC 10 mg 10 mg LAB. FOURNIER SA

A07XA04 RACECADOTRILUM PULB. PT. SOL. ORALĂ 30 mg

HIDRASEC 30 mg 30 mg LAB. FOURNIER SA

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| 166 |A08AA10| SIBUTRAMINUM\*\*\*\* | Protocol: A003E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A08AA10 SIBUTRAMINUM CAPS. 10 mg

LINDAXA 10 10 mg ZENTIVA A.S.

REDUCTIL 10 mg 10 mg ABBOTT GMBH & CO.KG

A08AA10 SIBUTRAMINUM CAPS. 15 mg

LINDAXA 15 15 mg ZENTIVA A.S.

REDUCTIL 15 mg 15 mg ABBOTT GMBH & CO.KG

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| 167 |A08AB01| ORLISTATUM\*\*\*\* | Protocol: A001E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A08AB01 ORLISTATUM CAPS. 120 mg

XENICAL 120 mg 120 mg ROCHE REGISTRATION LTD.

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| 168 |A08AX01| RIMONABANTUM\*\*\*\* | Protocol: A031E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A08AX01 RIMONABANTUM COMPR. FILM. 20 mg

ACOMPLIA 20 mg 20 mg SANOFI AVENTIS

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| 169 |A11CC03| ALFACALCIDOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A11CC03 ALFACALCIDOLUM CAPS. MOI 0.25 micrograme

ALPHA D3 0,25 micrograme 0.25 micrograme TEVA PHARMACEUTICALS SRL

A11CC03 ALFACALCIDOLUM CAPS. MOI 0.50 micrograme

ALPHA D3 0.50 micrograme 0.50 micrograme TEVA PHARMACEUTICALS

S.R.L.

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| 170 |A16AX01| ACIDUM TIOCTICUM (ALFA-LIPOICUM)\*\*# | Protocol: A021E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A16AX01 ACIDUM TIOCTICUM SOL. PERF. 12 mg/ml

(ALFA-LIPOICUM)

THIOGAMMA(R) TURBO-SE 12 mg/ml WORWAG PHARMA GMBH & CO.KG

THIOGAMMA(R) TURBO-SET 12 mg/ml WORWAG PHARMA GMBH & CO.KG

A16AX01 ACIDUM TIOCTICUM CONC. PT. SOL. PERF. 30 mg/ml

(ALFA-LIPOICUM)

THIOGAMMA(R) 600 INJEKT 30 mg/ml WORWAG PHARMA GMBH & CO.KG

A16AX01 ACIDUM TIOCTICUM COMPR. FILM. 600 mg

(ALFA-LIPOICUM)

THIOGAMMA(R) 600 oral 600 mg WORWAG PHARMA GMBH & CO.KG

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| 171 |B01AB04| DALTEPARINUM\*\*# | Protocol: B008D |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AB04 DALTEPARINUM SOL. INJ. 10000 ui/ml

FRAGMIN 10000 UI/ml 10000 ui/ml PFIZER EUROPE MA EEIG

B01AB04 DALTEPARINUM SOL. INJ. 15000 ui/0.6 ml

FRAGMIN 15000 UI/0,6 ml 15000 ui/0.6 ml PFIZER EUROPE MA EEIG

B01AB04 DALTEPARINUM SOL. INJ. 2500 ui/0.2 ml

FRAGMIN 2500 UI/0,2 ml 2500 ui/0.2 ml PFIZER EUROPE MA EEIG

B01AB04 DALTEPARINUM SOL. INJ. 5000 ui/0.2 ml

FRAGMIN 5000 UI/0,2 ml 5000 ui/0.2 ml PFIZER EUROPE MA EEIG

B01AB04 DALTEPARINUM SOL. INJ. 7500 ui/0.3 ml

FRAGMIN 7500 UI/0,3 ml 7500 ui/0.3 ml PFIZER EUROPE MA EEIG

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| 172 |B01AB05| ENOXAPARINUM\*\*# | Protocol: B008D |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AB05 ENOXAPARINUM SOL. INJ. 2000 ui

anti-Xa/0.2 ml

CLEXANE 2000 ui 2000 ui anti-Xa/0.2 ml LAB. AVENTIS

anti-Xa/0.2 ml

B01AB05 ENOXAPARINUM SOL. INJ. 4000 ui

anti-Xa/0,4 ml

CLEXANE 4000 ui 4000 ui anti-Xa/0,4 ml LAB. AVENTIS

anti-Xa/0.4 ml

B01AB05 ENOXAPARINUM SOL. INJ. 6000 ui

anti-Xa/0.6 ml

CLEXANE 6000 ui 6000 ui anti-Xa/0.6 ml LAB. AVENTIS

anti-Xa/0.6 ml

B01AB05 ENOXAPARINUM SOL. INJ. 8000 ui

anti-Xa/0.8 ml

CLEXANE 8000 ui 8000 ui anti-Xa/0.8 ml LAB. AVENTIS

anti-Xa/0.8 ml

B01AB05 ENOXAPARINUM SOL. INJ. 2000 ui

anti-Xa/0.2 ml

CLEXANE 2000 ui 2000 ui anti-Xa/0.2 ml LAB. AVENTIS

anti-Xa/0.2 ml

B01AB05 ENOXAPARINUM SOL. INJ. 4000 ui

anti-Xa/0,4 ml

CLEXANE 4000 ui 4000 ui LAB. AVENTIS

anti-Xa/0.4 ml anti-Xa/0,4 ml

B01AB05 ENOXAPARINUM SOL. INJ. 6000 ui

anti-Xa/0.6 ml

CLEXANE 6000 ui 6000 ui anti-Xa/0.6 ml LAB. AVENTIS

anti-Xa/0.6 ml

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| 173 |B01AB06| NADROPARINUM\*\*# | Protocol: B008D |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AB06 NADROPARINUM SOL. INJ. 11400 ui

AXa/0.6 ml

FRAXODI 11400 UI 11400 ui AXa/0.6 ml GLAXO GROUP LTD.

anti-factor Xa/0,6 ml

B01AB06 NADROPARINUM SOL. INJ. 15200 ui

AXa/0.8 ml

FRAXODI 15200 UI 15200 ui AXa/0.8 ml GLAXO GROUP LTD.

anti-factor Xa/0.8 ml

B01AB06 NADROPARINUM SOL. INJ. 2850 ui

AFXa/0.3 ml

FRAXIPARINE(R) 2850 UI 2850 ui AFXa/0.3 ml GLAXO GROUP LTD.

anti-factor Xa/0,3 ml

B01AB06 NADROPARINUM SOL. INJ. 3800 ui

AFXa/0.4 ml

FRAXIPARINE(R) 3800 UI 3800 ui AFXa/0.4 ml GLAXO GROUP LTD.

anti-factor Xa/0,4 ml

B01AB06 NADROPARINUM SOL. INJ. 5700 ui

AFXa/0.6 ml

FRAXIPARINE(R) 5700 UI 5700 ui AFXa/0.6 ml GLAXO GROUP LTD.

anti-factor Xa/0.6 ml

B01AB06 NADROPARINUM SOL. INJ. 7600 ui

AXa/0.8 ml

FRAXIPARINE(R) 7600 UI 7600 ui AXa/0.8 ml GLAXO GROUP LTD.

anti-factor Xa/0,8 ml

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| 174 |B01AB08| REVIPARINUM\*\*# | Protocol: B008D |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AB08 REVIPARINUM SOL. INJ. 1432 ui/0.25 ml

CLIVARIN(R) 1432 UI/0.25 ml 1432 ui/0.25 ml ABBOTT GMBH & CO.KG

B01AB08 REVIPARINUM SOL. INJ. 3436 ui/0.6 ml

CLIVARIN(R) 3436 UI/0,6 ml 3436 ui/0.6 ml ABBOTT GMBH & CO.KG

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| 175 |B01AB10| TINZAPARINUM\*\*# | Protocol: B008D |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AB10 TINZAPARINUM SOL. INJ. 10000 u ANTIF. Xa/ml

INNOHEP 10000 U ANTIF. Xa/ml LEO PHARMACEUTICAL

PRODUCTS

B01AB10 TINZAPARINUM SOL. INJ. 20000 u ANTIF. Xa/ml

INNOHEP 20000 U ANTIF. Xa/ml LEO PHARMACEUTICAL

PRODUCTS

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| 176 |B01AB11| SULODEXIDUM\*\* | Protocol: B014I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AB11 SULODEXIDUM CAPS. MOI 250 ULS

VESSEL DUE F 250 ULS ALFA WASSERMANN SPA

B01AB11 SULODEXIDUM SOL. INJ. 600 ULS/2 ml

VESSEL DUE F 600 ULS/2 ml ALFA WASSERMANN SPA

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| 177 |B01AC04| CLOPIDOGRELUM\*\* | Protocol: B009I;|

| | | | B010I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AC04 CLOPIDOGRELUM COMPR. FILM. 75 mg

PLAVIX 75 mg 75 mg SANOFI PHARMA - BRISTOL

MYERS SQUIBB

Cod restricţie 1719: Prevenţia recurenţei accidentului vascular ischemic sau a accidentului ischemic tranzitor la pacienţii cu istoric de episoade ischemice cerebrovasculare conform protocolului de prescriere.

Cod restricţie 1722: Prevenţia recurenţei infarctului de miocard sau a anginei instabile la pacienţii cu istoric de evenimente ischemice cardiace simptomatice conform protocolului de prescriere.

Cod restricţie 3008: Tratamentul antiagregant al ateromatozei extensive (carotidiene, coronariene şi periferice).

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| 178 |B01AC30| COMBINATII (DIPYRIDAMOLUM + ACIDUM | Protocol: B010I |

| | | ACETYLSALICYLICUM)\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AC30 COMBINATII CAPS. ELIB. MODIF.

(DIPYRIDAMOLUM +

ACIDUM

ACETYLSALICYLICUM)

AGGRENOX BOEHRINGER INGELHEIM

INTERNATIONAL GMBH

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| 179 |B01AX05| FONDAPARINUX SODIUM\*\*# | Protocol: B008D |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AX05 FONDAPARINUX SODIUM SOL. INJ. ÎN SERINGĂ 10 mg/0.8 ml

PREUMPLUTĂ

ARIXTRA 10 mg/0.8 ml 10 mg/0.8 ml GLAXO GROUP LTD.

B01AX05 FONDAPARINUX SODIUM SOL. INJ. ÎN SERINGĂ 2.5 mg/0.5 ml

PREUMPLUTĂ

ARIXTRA 2.5 mg/0.5 ml 2.5 mg/0.5 ml GLAXO GROUP LTD.

B01AX05 FONDAPARINUX SODIUM SOL. INJ. ÎN SERINGĂ 5 mg/0.4 ml

PREUMPLUTĂ

ARIXTRA 5 mg/0.4 ml 5 mg/0.4 ml GLAXO GROUP LTD.

B01AX05 FONDAPARINUX SODIUM SOL. INJ. ÎN SERINGĂ 7.5 mg/0.6 ml

PREUMPLUTĂ

ARIXTRA 7.5 mg/0.6 ml 7.5 mg/0.6 ml GLAXO GROUP LTD.

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| 180 |B03AC02| COMPLEX DE HIDROXID DE FER (III) SUCROZA | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B03AC02 COMPLEX DE HIDROXID SOL. INJ./PERF. 20 mg/ml

DE FER (III) SUCROZA

VENOFER(R) 20 mg/ml VIFOR FRANCE S.A.

Cod restricţie 2070: Anemie feriprivă, la pacienţii care au prezentat o reacţie de hipersensibilitate documentată la fier polimaltozat şi la care este indicată administrarea continuă intravenoasă.

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| 181 |B01AC18| TRIFLUSALUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AC18 TRIFLUSALUM CAPS. 300 mg

AFLEN(R) 300 mg ZENTIVA S.A.

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| 182 |C01BB02| MEXILETINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01BB02 MEXILETINUM CAPS. 200 mg

MEXITIL(R) 200 mg 200 mg BOEHRINGER INGELHEIM INT.

GMBH

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| 183 |C01EB17| IVABRADINUM\*\* | Protocol: C003I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01EB17 IVABRADINUM COMPR. FILM. 5 mg

CORLENTOR 5 mg 5 mg LES LAB. SERVIER

C01EB17 IVABRADINUM COMPR. FILM. 7.5 mg

CORLENTOR 7,5 mg 7.5 mg LES LAB. SERVIER

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| 184 |C02AC05| MOXONIDINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Hipertensiune arterială esenţială la pacienţii care primesc tratament antihipertensiv concomitent.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C02AC05 MOXONIDINUM COMPR. FILM. 0.2 mg

MOXOGAMMA 0,2 mg 0.2 mg WORWAG PHARMA GMBH

PHYSIOTENS 0,2 0.2 mg SOLVAY PHARMACEUTICALS

GMBH

C02AC05 MOXONIDINUM COMPR. FILM. 0.3 mg

MOXOGAMMA 0,3 mg 0.3 mg WORWAG PHARMA GMBH

C02AC05 MOXONIDINUM COMPR. FILM. 0.4 mg

MOXOGAMMA 0,4 mg 0.4 mg WORWAG PHARMA GMBH

PHYSIOTENS 0.4 0.4 mg SOLVAY PHARMACEUTICALS

GMBH

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| 185 |C02AC06| RILMENIDINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C02AC06 RILMENIDINUM COMPR. 1 mg

TENAXUM 1 mg LES LABORATOIRES SERVIER

INDUSTRIE

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| 186 |C02CA01| PRAZOSINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C02CA01 PRAZOSINUM COMPR. 1 mg

MINIPRESS(R) 1 mg 1 mg PFIZER EUROPE MA EEIG

C02CA01 PRAZOSINUM COMPR. 2 mg

MINIPRESS(R) 2 mg 2 mg PFIZER EUROPE MA EEIG

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| 187 |C03DA04| EPLERENONUM\*# | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Insuficienţă cardiacă cu o fracţie de ejecţie a ventriculului stâng de 40% sau mai puţin, apărută la 3 - 14 zile de la un infarct miocardic acut. Tratamentul cu eplerenone trebuie început la maximum 14 zile de la data apariţiei infarctului miocardic acut. Data infarctului miocardic acut şi data iniţierii tratamentului cu eplerenone trebuie documentate în fişa pacientului.

Electroliţii serici trebuie să fie verificaţi periodic.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C03DA04 EPLERENONUM COMPR. FILM. 25 mg

INSPRA 25 mg 25 mg PFIZER EUROPE MA EEIG

C03DA04 EPLERENONUM COMPR. FILM. 50 mg

INSPRA 50 mg 50 mg PFIZER EUROPE MA EEIG

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| 188 |C04AE02| NICERGOLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C04AE02 NICERGOLINUM COMPR. FILM. 10 mg

NICERGOLINA LPH 10 mg 10 mg LABORMED PHARMA SA

C04AE02 NICERGOLINUM DRAJ. 10 mg

NICERIUM(R) 10 10 mg HEXAL AG

SINERGOLIN 10 10 mg SINTOFARM SA

C04AE02 NICERGOLINUM CAPS. ELIB. MODIF. 15 mg

NICERIUM(R) 15 15 mg HEXAL AG

C04AE02 NICERGOLINUM CAPS. ELIB. MODIF. 30 mg

NICERIUM(R) 30 UNO 30 mg HEXAL AG

C04AE02 NICERGOLINUM COMPR. FILM. 30 mg

NICERGOLINA LPH 30 mg 30 mg LABORMED PHARMA SA

SERMION 30 mg PFIZER EUROPE MA EEIG

C04AE02 NICERGOLINUM DRAJ. 30 mg

SINERGOLIN 30 30 mg SINTOFARM SA

C04AE02 NICERGOLINUM COMPR. FILM. 5 mg

NICERGOLINA LPH 5 mg 5 mg LABORMED PHARMA SA

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| 189 |C04AX07| VINCAMINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C04AX07 VINCAMINUM DRAJ. 10 mg

VINCAMINA 10 mg BIOFARM SA

C04AX07 VINCAMINUM COMPR. 20 mg

VINCAMIL 20 mg FARMACEUTICI ECOBI S.A.S.

C04AX07 VINCAMINUM CAPS. ELIB. PREL. 30 mg

OXYBRAL SR 30 mg 30 mg GLAXO SMITHKLINE (GSK) SRL

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| 190 |C04AXN1| GINKGO BILOBA\*\* | Protocol: C001I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C04AXN1 GINKGO BILOBA COMPR. FILM. 120 mg

GINGIUM 120 m 120 mg HEXAL AG

C04AXN1 GINKGO BILOBA COMPR. FILM. 40 mg

GINGIUM 40 mg 40 mg HEXAL AG

TEBOKAN 40 mg DR. WILLMAR SCHWABE GMBH &

CO KG

C04AXN1 GINKGO BILOBA CAPS. 80 mg

BILOBIL FORTE 80 mg KRKA D.D. NOVO MESTO

C04AXN1 GINKGO BILOBA COMPR. FILM. 80 mg

GINGIUM 80 mg 80 mg HEXAL AG

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| 191 |C05CA53| DIOSMINUM (COMBINATII)\*\* | Protocol: B016I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C05CA53 DIOSMINUM COMPR. FILM. 500 mg

(COMBINATII)

DETRALEX(R) 500 mg LES LAB. SERVIER IND.

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| 192 |C07AB12| NEBIVOLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB12 NEBIVOLOLUM COMPR. 5 mg

NEBILET(R) 5 mg BERLIN CHEMIE AG MENARINI

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| 193 |C07BB07| COMBINATII (BISOPROLOLUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07BB07 COMBINATII COMPR. FILM.

(BISOPROLOLUM +

HYDROCHLOROTHIAZIDUM)

LODOZ 10 mg MERCK KGAA

LODOZ 2,5 mg MERCK KGAA

LODOZ 5 mg MERCK KGAA

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| 194 |C07EBN1| COMBINATII (METOPROLOLUM + FELODIPINUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07EBN1 COMBINATII COMPR. ELIB. PREL. 50 mg + 5 mg

(METOPROLOLUM +

FELODIPINUM)

LOGIMAX(R) 50 mg + 5 mg ASTRAZENECA AB

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| 195 |C08CA09| LACIDIPINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C08CA09 LACIDIPINUM COMPR. FILM. 4 mg

LACIPIL 4 mg 4 mg GLAXO OPERATIONS UK LTD.

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| 196 |C08CA13| LERCANIDIPINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C08CA13 LERCANIDIPINUM COMPR. FILM. 10 mg

LERIDIP 10 mg BERLIN CHEMIE AG MENARINI

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C08CA13 LERCANIDIPINUM COMPR. FILM. 20 mg

LERIDIP 20 20 mg BERLIN CHEMIE AG MENARINI

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| 197 |C09AA04| PERINDOPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09AA04 PERINDOPRILUM COMPR. FILM. 10 mg

PRESTARIUM 10 mg 10 mg LES LAB. SERVIER IND.

C09AA04 PERINDOPRILUM COMPR. 4 mg

PRESTARIUM(R) 4 mg 4 mg LES LAB. SERVIER IND.

C09AA04 PERINDOPRILUM COMPR. FILM. 5 mg

PRESTARIUM 5 mg 5 mg LES LAB. SERVIER IND.

C09AA04 PERINDOPRILUM COMPR. 8 mg

PRESTARIUM(R) 8 mg 8 mg LES LAB. SERVIER IND.

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| 198 |C09AA07| BENAZEPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09AA07 BENAZEPRILUM COMPR. FILM. 10 mg

BENAZEPRIL STADA 10 mg 10 mg STADA ARZNEIMITTEL AG

CIBACEN(R) 10 mg 10 mg NOVARTIS PHARMA GMBH

C09AA07 BENAZEPRILUM COMPR. FILM. 20 mg

BENAZEPRIL STADA 20 mg 20 mg STADA ARZNEIMITTEL AG

CIBACEN(R) 20 mg 20 mg NOVARTIS PHARMA GMBH

C09AA07 BENAZEPRILUM COMPR. FILM. 5 mg

BENAZEPRIL STADA 5 mg 5 mg STADA ARZNEIMITTEL AG

CIBACEN(R) 5 mg 5 mg NOVARTIS PHARMA GMBH

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| 199 |C09AA10| TRANDOLAPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09AA10 TRANDOLAPRILUM CAPS. 0.5 mg

GOPTEN(R) 0,5 mg 0.5 mg ABBOTT GMBH & CO.KG

C09AA10 TRANDOLAPRILUM CAPS. 2 mg

GOPTEN(R) 2 mg 2 mg ABBOTT GMBH & CO.KG

C09AA10 TRANDOLAPRILUM CAPS. 4 mg

GOPTEN(R) 4 mg 4 mg ABBOTT GMBH & CO.KG

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| 200 |C09AA15| ZOFENOPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

C09AA15 ZOFENOPRILUM COMPR. FILM. 30 mg

ZOMEN(R) 30 mg 30 mg BERLIN CHEMIE AG

C09AA15 ZOFENOPRILUM COMPR. FILM. 7.5 mg

ZOMEN(R) 7,5 mg 7.5 mg BERLIN CHEMIE AG

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| 201 |C09BA04| COMBINATII (PERINDOPRILUM + INDAPAMIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09BA04 COMBINATII COMPR. 2 mg + 0.625 mg

(PERINDOPRILUM +

INDAPAMIDUM)

NOLIPREL(R) 2 mg + 0.625 mg LES LAB. SERVIER IND.

C09BA04 COMBINATII COMPR. 4 mg + 1.25 mg

(PERINDOPRILUM +

INDAPAMIDUM)

NOLIPREL FORTE(R) 4 mg + 1.25 mg LES LAB. SERVIER IND.

PRESTARIUM PLUS 4 mg + 1.25 mg LES LAB. SERVIER IND.

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| 202 |C09BA06| COMBINATII (QUINAPRILUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09BA06 COMBINATII COMPR. FILM. 10 mg/12.5 mg

(QUINAPRILUM +

HYDROCHLOROTHIAZIDUM)

ACCUZIDE 10 mg/12.5 mg PFIZER EUROPE MA EEIG

C09BA06 COMBINATII COMPR. FILM. 20 mg/12.5 mg

(QUINAPRILUM +

HYDROCHLOROTHIAZIDUM)

ACUZIDE FORTE 20 mg/12.5 mg PFIZER EUROPE MA EEIG

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| 203 |C09BA09| COMBINATII (FOSINOPRILUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09BA09 COMBINATII COMPR. 20 mg/12.5 mg

(FOSINOPRILUM +

HYDROCHLOROTHIAZIDUM)

FOSINOZIDE 20 mg/12,5 mg 20 mg/12.5 mg BRISTOL MYERS SQUIBB KFT

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| 204 |C09BB02| COMBINATII (NITRENDIPINUM + ENALAPRILUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09BB02 COMBINATII COMPR. 10 mg + 20 mg

(NITRENDIPINUM +

ENALAPRILUM)

ENEAS 10 mg + 20 mg VITA CIENTIFICA SL

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| 205 |C09BB10| COMBINATII (VERAPAMILUM + TRANDOLAPRILUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09BB10 COMBINATII COMPR. FILM. ELIB. 180 mg/2 mg

(VERAPAMILUM + MODIF.

TRANDOLAPRILUM)

TARKA 180 mg/2 mg ABBOTT GMBH & CO.KG

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| 206 |C09CA03| VALSARTANUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09CA03 VALSARTANUM COMPR. FILM. 160 mg

DIOVAN 160 mg 160 mg NOVARTIS PHARMA GMBH

C09CA03 VALSARTANUM COMPR. FILM. 80 mg

DIOVAN 80 mg 80 mg NOVARTIS PHARMA GMBH

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| 207 |C09CA04| IRBESARTANUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09CA04 IRBESARTANUM COMPR. FILM. 150 mg

APROVEL 150 mg 150 mg SANOFI PHARMA-BRISTOL

MYERS SQUIBB

C09CA04 IRBESARTANUM COMPR. FILM. 300 mg

APROVEL 300 mg 300 mg SANOFI PHARMA-BRISTOL

MYERS SQUIBB

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| 208 |C09CA06| CANDESARTANUM CILEXETIL | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09CA06 CANDESARTANUM COMPR. 16 mg

CILEXETIL

ATACAND 16 mg ASTRAZENECA AB

C09CA06 CANDESARTANUM COMPR. 8 mg

CILEXETIL

ATACAND 8 mg ASTRAZENECA AB

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| 209 |C09CA07| TELMISARTANUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09CA07 TELMISARTANUM COMPR. 40 mg

MICARDIS 40 mg 40 mg BOEHRINGER INGELHEIM INT.

GMBH

PRITOR 40 mg 40 mg BAYER HEALTHCARE AG

C09CA07 TELMISARTANUM COMPR. 80 mg

MICARDIS 80 mg 80 mg BOEHRINGER INGELHEIM INT.

GMBH

PRITOR 80 mg 80 mg BAYER HEALTHCARE AG

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| 210 |C09DA01| COMBINATII (LOSARTANUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09DA01 COMBINATII COMPR. FILM. 50 mg + 12.5 mg

(LOSARTANUM +

HYDROCHLOROTHIAZIDUM)

HYZAAR(R) 50 mg/12,5 mg 50 mg + 12.5 mg MERCK SHARP & DOHME

ROMANIA S.R.L

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| 211 |C09DA03| COMBINATII (VALSARTANUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09DA03 COMBINATII COMPR. FILM. 80 mg/12.5 mg

(VALSARTANUM +

HYDROCHLOROTHIAZIDUM)

CO-DIOVAN 80 mg/12,5 mg 80 mg/12.5 mg NOVARTIS PHARMA GMBH

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| 212 |C09DA04| COMBINATII (IRBERSARTANUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09DA04 COMBINATII COMPR. 15 u mg/12.5 mg

(IRBERSARTANUM +

HYDROCHLOROTHIAZIDUM)

COAPROVEL 150 mg/12.5 mg 150 mg/12.5 mg SANOFI PHARMA-BRISTOL

MYERS SQUIBB SNC

C09DA04 COMBINATII COMPR. 300 mg/12.5 mg

(IRBERSARTANUM +

HYDROCHLOROTHIAZIDUM)

COAPROVEL 300 mg/12.5 mg 300 mg/12.5 mg SANOFI PHARMA-BRISTOL

MYERS SQUIBB SNC

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| 213 |C09DA07| COMBINATII (TELMISARTANUM + | |

| | | HYDROCHLOROTHIAZIDUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09DA07 COMBINATII COMPR. 80/12.5 mg

(TELMISARTANUM +

HYDROCHLOROTHIAZIDUM)

MICARDISPLUS 80/12.5 mg 80/12.5 mg BOEHRINGER INGELHEIM INT.

GMBH

C09DA07 COMBINATII COMPR. 80 mg/12.5 mg

(TELMISARTANUM +

HYDROCHLOROTHIAZIDUM)

PRITOR PLUS 80 mg/12,5 mg 80 mg/12.5 mg BAYER HEALTHCARE AG

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| 214 |C09DB01| COMBINATII (VALSARTANUM + AMLODIPINUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul hipertensiunii arteriale la pacienţii cu

intoleranţă la inhibitorii enzimei de conversie ai

angiotensinei.

Tratamentul hipertensiunii arteriale la pacienţii ale

căror valori tensionale nu pot fi controlate adecvat cu

celelalte clase de medicamente antihipertensive.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

C09DB01 COMBINATII COMPR. FILM. 10 mg/160 mg

(VALSARTANUM +

AMLODIPINUM)

EXFORGE 10 mg/160 mg 10 mg/160 mg NOVARTIS EUROPHARM LTD.

C09DB01 COMBINATII COMPR. FILM. 5 mg/160 mg

(VALSARTANUM +

AMLODIPINUM)

EXFORGE 5 mg/160 mg 5 mg/160 mg NOVARTIS EUROPHARM LTD.

C09DB01 COMBINATII COMPR. FILM. 5 mg/80 mg

(VALSARTANUM +

AMLODIPINUM)

EXFORGE 5 mg/80 mg 5 mg/80 mg NOVARTIS EUROPHARM LTD.

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| 215 |C10AA04| FLUVASTATINUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AA04 FLUVASTATINUM CAPS. 20 mg

LESCOL(R) 20 20 mg NOVARTIS PHARMA GMBH

C10AA04 FLUVASTATINUM CAPS. 40 mg

LESCOL(R) 40 40 mg NOVARTIS PHARMA GMBH

C10AA04 FLUVASTATINUM COMPR. ELIB. PREL. 80 mg

LESCOL(R) XL 80 mg NOVARTIS PHARMA GMBH

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| 216 |C10AA05| ATORVASTATINUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AA05 ATORVASTATINUM COMPR. FILM. 10 mg

SORTIS 10 mg 10 mg PFIZER EUROPE MA EEIG

C10AA05 ATORVASTATINUM COMPR. FILM. 20 mg

SORTIS 20 mg 20 mg PFIZER EUROPE MA EEIG

C10AA05 ATORVASTATINUM COMPR. FILM. 40 mg

SORTIS 40 mg 40 mg PFIZER EUROPE MA EEIG

C10AA05 ATORVASTATINUM COMPR. FILM. 80 mg

SORTIS 80 mg 80 mg PFIZER EUROPE MA EEIG

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| 217 |C10AA07| ROSUVASTATINUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AA07 ROSUVASTATINUM COMPR. FILM. 10 mg

CRESTOR 10 mg 10 mg ASTRAZENECA UK LTD.

C10AA07 ROSUVASTATINUM COMPR. FILM. 20 mg

CRESTOR 20 mg 20 mg ASTRAZENECA UK LTD.

C10AA07 ROSUVASTATINUM COMPR. FILM. 40 mg

CRESTOR 40 mg 40 mg ASTRAZENECA UK LTD.

C10AA07 ROSUVASTATINUM COMPR. FILM. 5 mg

CRESTOR 5 mg 5 mg ASTRAZENECA UK LTD.

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| 218 |C10AB02| BEZAFIBRATUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AB02 BEZAFIBRATUM DRAJ. 200 mg

REGADRIN(R) B 200 mg BERLIN CHEMIE AG MENARINI

GROUP

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| 219 |C10AB08| CIPROFIBRATUM | Protocol: CE01E |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AB08 CIPROFIBRATUM CAPS. 100 mg

LIPANOR(R) 100 mg 100 mg SANOFI-AVENTIS FRANCE

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| 220 |C10AX06| ACID OMEGA-3-ESTERI ETILICI 90\*\* | Protocol: C004I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AX06 ACID OMEGA-3-ESTERI CAPS. MOI 1000 mg

ETILICI 90

OMACOR(R) 1000 mg PRONOVA BIOCARE AS

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| 221 |C10AX09| EZETIMIBUM | |

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Prescriere limitată: Se utilizează pentru pacienţii care îndeplinesc criteriile de eligibilitate stabilite prin Protocolul de prescriere a medicamentelor hipolipemiante.

Cod restricţie 2649: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (a) boală coronariană. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetinib.

Cod restricţie 2650: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (b) diabet zaharat. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2651: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (c) boala vasculară periferică. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doza zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2652: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (d) hipercolesterolemie familială heterozigotă. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doza zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2653: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (e) boala cerebrovasculară simptomatică. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doza zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2667: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (f) istoric familial de boala coronariană. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doza zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2668: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (g) HTA. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 1989: Pacienţi care îndeplinesc criteriile de eligibilitate pentru prescrierea de hipolipemiante (în concordanţă cu criteriile stabilite în Protocoalele de prescriere în vederea decontării) atunci când tratamentul cu un inhibitor de HMG CoA reductaza (statine) este contraindicat.

Cod restricţie 2669: Pacienţi care îndeplinesc criteriile de eligibilitate pentru prescrierea de hipolipemiante (în concordanţă cu criteriile stabilite în Protocoalele de prescriere în vederea decontării) atunci când tratamentul cu un inhibitor de HMG CoA reductaza (statin) trebuie întrerupt sau redus la mai puţin de 20 mg/zi datorită apariţiei efectelor secundare următoare: (i) Mialgie severă (simptome musculare fără creşterea CK) care survine pe perioada tratamentului cu statine; sau (ii) Miozita (creşterea importantă a CK, cu sau fără simptomatologie musculară) demonstrată prin valori de două ori mai mari decât limita superioară a normalului la o singură determinare sau o tendinţă de creştere la determinări consecutive, neexplicate de alte cauze; sau (iii) Persistenţa unor valori crescute ale transaminazelor, neexplicată de alte cauze (mai mult de trei ori decât valoarea limită maximă a normalului) în timpul tratamentului cu o statină.

Cod restricţie 1991: Homozygous sitosterolaemia;

Cod restricţie 2438: Pacienţi cu hipercolesterolemie familială (homozigotă) care sunt eligibili pentru tratamentul cu hipolipemiante (în concordanţă cu Protocolul de prescriere în vederea decontării), în combinaţie cu un inhibitor de HMG CoA reductase (statin).

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10AX09 EZETIMIBUM COMPR. 10 mg

EZETROL(R) 10 mg MERCK SHARP & DOHME

ROMANIA SRL

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| 222 |C10BA02| COMBINATII (EZETIMIBUM + SIMVASTATINUM) | |

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Cod restricţie 2654: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (a) boală coronariană. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2655: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (b) diabet zaharat. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dieta şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2656: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (c) boala vasculară periferică. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice.

Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2657: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (d) hipercolesterolemie familială heterozigotă. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2658: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (e) boală cerebrovasculară simptomatică. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dieta şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetimib. Valoarea colesterolului controlat inadecvat trebuie să fie mai recentă de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2678: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (f) istoric familial de boală coronariană. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SAU (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2679: Tratamentul, în combinaţie cu dietă şi exerciţiu fizic, ce presupune asocierea cu un inhibitor de HMG CoA reductaza (statină) la pacienţii al căror nivel de colesterol nu este controlat adecvat cu o statină şi care au: (g) HTA. Controlul inadecvat sub tratament cu o statină este definit astfel: (1) nivel al colesterolului peste pragul iniţial după cel puţin 3 luni de tratament cu o statină în doza zilnică optimă de statine, concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetinib; SA U (2) Controlul inadecvat al colesterolului cu nivele ale acestuia mai mari decât 155 mg/dl după cel puţin 3 luni de tratament cu o doză zilnică optimă de statine concomitent cu dietă şi exerciţii fizice. Doza, durata tratamentului cu statine şi nivelul colesterolului controlat inadecvat trebuie să fie înregistrate în dosarul medical al pacientului la momentul iniţierii tratamentului cu ezetinimb. Valoarea colesterolului controlat inadecvat trebuie să fie mai recent de două luni la momentul iniţierii tratamentului cu ezetimib.

Cod restricţie 2431: Pacienţii cu hipercolesterolemie familială (heterozigotă) care sunt eligibili pentru tratament cu hipo-lipemiante (în concordanţă cu criteriile stabilite prin Protocolul de prescriere în vederea decontării).

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10BA02 COMBINATII COMPR. 10 mg/10 mg

(EZETIMIBUM +

SIMVASTATINUM)

INEGY(R) 10 mg/10 mg 10 mg/10 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C10BA02 COMBINATII COMPR. 10 mg/20 mg

(EZETIMIBUM +

SIMVASTATINUM)

INEGY(R) 10 mg/20 mg 10 mg/20 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C10BA02 COMBINATII COMPR. 10 mg/40 mg

(EZETIMIBUM +

SIMVASTATINUM)

INEGY(R) 10 mg/40 mg 10 mg/40 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C10BA02 COMBINATII COMPR. 10 mg/80 mg

(EZETIMIBUM +

SIMVASTATINUM)

INEGY(R) 10 mg/80 mg 10 mg/80 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

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| 223 |C10BX03| COMBINATII (ATORVASTATINUM + AMLODIPINUM) | Protocol: CE01E |

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Prescriere limitată: Pentru utilizarea la pacienţii cu hipertensiune

arterială şi/sau angină pectorală care îndeplinesc

criteriile stabilite de Protocolul de prescriere a

medicamentelor hipolipemiante în vederea decontării,

şi: (a) care primesc în mod curent tratament cu un

blocant de canale de calciu.

Pentru utilizarea la pacienţii cu hipertensiune

arterială şi/sau angină pectorală care îndeplinesc

criteriile stabilite de Protocolul de prescriere a

medicamentelor hipolipemiante în vederea decontării,

şi: (b) al căror nivel al tensiunii arteriale şi/sau

angină pectorală sunt insuficient controlate cu alte

clase de antihipertensive şi/sau medicamente

anti-angină şi la care terapia adjuvantă cu blocanţi ai

canalelor de calciu ar fi adecvată;

Pentru utilizarea la pacienţii cu hipertensiune

arterială şi/sau angină pectorală care îndeplinesc

criteriile stabilite de Protocolul de prescriere a

medicamentelor hipolipemiante în vederea decontării, şi

(c) care nu tolerează efectele secundare ale altor

clase de antihipertensive şi/sau medicamente

anti-angină şi la care înlocuirea terapiei cu un

blocant de canale de calciu ar fi indicată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C10BX03 COMBINATII COMPR. FILM. 10 mg/10 mg

(ATORVASTATINUM +

AMLODIPINUM)

CADUET 10 mg/10 mg 10 mg/10 mg PFIZER EUROPE MA EEIG

C10BX03 COMBINATII COMPR. FILM. 5 mg/10 mg

(ATORVASTATINUM +

AMLODIPINUM)

CADUET 5 mg/10 mg 5 mg/10 mg PFIZER EUROPE MA EEIG

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| 224 |D01AA02| NATAMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D01AA02 NATAMYCINUM CREMA 20 mg/g

PIMAFUCIN 20 mg/g ASTELLAS EUROPE B.V.

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| 226 |D01BA02| TERBINAFINUM | |

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Prescriere limitată: Tratamentul de primă intenţie al onicomicozelor proximale sau extinse la care tratamentul topic a eşuat.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D01BA02 TERBINAFINUM COMPR. 250 mg

LAMISIL(R) 250 mg NOVARTIS PHARMA GMBH

TERBINARAN 250 mg RANBAXY UK LIMITED

TERBISIL(R) 250 mg 250 mg GEDEON RICHTER LTD.

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| 227 |D05AX02| CALCIPOTRIOLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D05AX02 CALCIPOTRIOLUM CREMA 50 micrograme/g

DAIVONEX 50 micrograme/g LEO PHARMACEUTICAL

PRODUCTS

D05AX02 CALCIPOTRIOLUM UNGUENT 50 micrograme/g

DAIVONEX UNGUENT 50 micrograme/g LEO PHARMACEUTICAL

PRODUCTS

D05AX02 CALCIPOTRIOLUM SOL. CUT. 50 micrograme/ml

DAIVONEX SOLUŢIE CUTANATĂ 50 micrograme/ml LEO PHARMACEUTICAL

PENTRU SCALP PRODUCTS

D05AX02 CALCIPOTRIOLUM UNGUENT 0.005%

SOREL(R) unguent 0,005% 0.005% LEK PHARMACEUTICALS D.D.

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| 228 |D05AX52| COMBINATII (CALCIPOTRIOLUM + | |

| | | BETAMETHASONUM)\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D05AX52 COMBINATII UNGUENT

(CALCIPOTRIOLUM +

BETAMETHASONUM)

DAIVOBET(R) UNGUENT LEO PHARMACEUTICAL

PRODUCTS

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| 230 |D06AX01| ACIDUM FUSIDICUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D06AX01 ACIDUM FUSIDICUM CREMA 2,00%

FUCIDIN(R) 2% LEO PHARMACEUTICAL

PRODUCTS

D06AX01 ACIDUM FUSIDICUM UNGUENT 2,00%

FUCIDIN(R) 2% LEO PHARMACEUTICAL

PRODUCTS

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| 231 |D06AXN1| COMBINATII (NEOMYCINUM + BACITRACINUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D06AXN1 COMBINATII PULB. CUT.

(NEOMYCINUM +

BACITRACINUM)

BANEOCIN(R) SANDOZ GMBH

D06AXN1 COMBINATII UNGUENT

(NEOMYCINUM +

BACITRACINUM)

BANEOCIN(R) SANDOZ GMBH

NEOBACIN DAR AL DAWA PHARMA SRL

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| 232 |D06BA01| SULFADIAZINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D06BA01 SULFADIAZINUM CREMA 1,00%

DERMAZIN(R) 1% 1% LEK PHARMACEUTICALS D.D.

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| 233 |D06BB04| PODOPHYLLOTOXINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D06BB04 PODOPHYLLOTOXINUM CREMA 1,5 mg/g

WARTEC 1,5 mg/g 1,5 mg/g STIEFEL LABORATORIES (UK)

LTD.

D06BB04 PODOPHYLLOTOXINUM SOL. CUT. 5 mg/ml

CONDYLINE 5 mg/ml ASTELLAS PHARMA EUROPE

B.V.

Prescriere limitată: Pentru tratamentul verucilor.

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| 235 |D07AB02| HYDROCORTISONUM BUTYRATUM | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AB02 HYDROCORTISONUM CREMA 0.1%

BUTYRATUM

LOCOID(R) cremă 0,1% 0.1% ASTELLAS PHARMA EUROPE BV

LOCOID(R)LIPOCREAM 0,1% 0.1% ASTELLAS PHARMA EUROPE

B.V.

D07AB02 HYDROCORTISONUM EMULSIE CUT. 0.1%

BUTYRATUM

LOCOID CRELO(R) 0,1% 0.1% ASTELLAS PHARMA EUROPE BV

D07AB02 HYDROCORTISONUM SOL. CUT. 0.1%

BUTYRATUM

LOCOID(R) 0.1% ASTELLAS PHARMA EUROPE

B.V.

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| 236 |D07AC13| MOMETASONUM | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AC13 MOMETASONUM CREMA 1 mg/g

ELOCOM 1 mg/g SCHERING PLOUGH EUROPE

D07AC13 MOMETASONUM SOL. CUT. 1 mg/g

ELOCOM 1 mg/g SCHERING PLOUGH EUROPE

D07AC13 MOMETASONUM UNGUENT 1 mg/g

ELOCOM 1 mg/g SCHERING PLOUGH EUROPE

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| 237 |D07AC14| METHYLPREDNISOLONUM ACEPONAT | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AC14 METHYLPREDNISOLONUM EMULSIE CUT. 0.1%

ACEPONAT

ADVANTAN(R) MILK 0.1% INTENDIS GMBH

D07AC14 METHYLPREDNISOLONUM CREMA 1 mg/g

ACEPONAT

ADVANTAN 1 mg/g INTENDIS GmbH

D07AC14 METHYLPREDNISOLONUM UNGUENT 1 mg/g

ACEPONAT

ADVANTAN 1 mg/g INTENDIS GmbH

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| 238 |D07AC17| FLUTICASONUM | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AC17 FLUTICASONUM CREMA 0.05%

CUTIVATE 0.05% GLAXO WELLCOME UK LIMITED

D07AC17 FLUTICASONUM UNGUENT 0.005%

CUTIVATE 0.005% GLAXO WELLCOME UK LIMITED

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| 239 |D07AD01| CLOBETASOLUM\* | Protocol: D001L |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07AD01 CLOBETASOLUM CREMA 0.05%

DERMOVATE 0.05% GLAXOWELLCOME UK LTD.

D07AD01 CLOBETASOLUM SOL. CUT. 0.05%

DERMOVATE(R) 0.05% GLAXO WELLCOME UK LTD.

D07AD01 CLOBETASOLUM UNGUENT 0.05%

DERMIONE 0.05% OZONE LABORATORIES LTD.

DERMOVATE 0.05% GLAXOWELLCOME UK LTD.

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| 240 |D07CA01| COMBINATII (HYDROCORTISONUM + | Protocol: D001L |

| | | ANTIINFECTIOASE) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07CA01 COMBINATII CREMA

(HYDROCORTISONUM +

ANTIINFECTIOASE)

FUCIDIN(R) H LEO PHARMACEUTICAL

PRODUCTS

PIMAFUCORT ASTELLAS PHARMA EUROPE

B.V.

D07CA01 COMBINATII UNGUENT

(HYDROCORTISONUM +

ANTIINFECTIOASE)

PIMAFUCORT ASTELLAS PHARMA EUROPE

B.V.

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| 241 |D07CC01| COMBINATII (BETAMETHASONUM + | Protocol: D001L |

| | | ANTIINFECTIOASE) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07CC01 COMBINATII CREMA

(BETAMETHASONUM +

ANTIINFECTIOASE)

FUCICORT(R) LEO PHARMACEUTICAL

PRODUCTS

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| 242 |D07XC03| COMBINATII (MOMETASONUM + | Protocol: D001L |

| | | AC. SALICILICUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D07XC03 COMBINATII UNGUENT

(MOMETASONUM + AC.

SALICILICUM)

ELOSALIC SCHERING PLOUGH EUROPE

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| 243 |D10AD54| COMBINATII (MOMETASONUM + ERITROMICINUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D10AD54 COMBINATII GEL 0,05%/0,2%

(ISOTRETINOINUM +

ERITROMICINUM)

ISOTREXIN GEL 0,05%/0,2% STIEFEL LABORATORIES (UK)

LTD.

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| 244 |D10AX03| ACIDUM AZELAICUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D10AX03 ACIDUM AZELAICUM CREMA 200 mg/g

SKINOREN 200 mg/g INTENDIS GmbH

D10AX03 ACIDUM AZELAICUM GEL 15%

SKINOREN(R) 15% 15% INTENDIS GMBH

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| 245 |D10BA01| ISOTRETINOINUM\*\* | |

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Cod restricţie 1354: Acnee chistică severă care nu răspunde la alte tipuri de tratamente.

Acest medicament poate cauza defecte la naştere. Isotretinoin a fost incriminat pentru cauzarea frecventă a altor efecte toxice cu potenţial sever.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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D10BA01 ISOTRETINOINUM CAPS. MOI 10 mg

ROACCUTANE 10 mg 10 mg ROCHE ROMANIA SRL

SOTRET 10 mg 10 mg RANBAXY U.K. LIMITED

D10BA01 ISOTRETINOINUM CAPS. MOI 20 mg

SOTRET 20 mg 20 mg RANBAXY U.K. LIMITED

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| 246 |G01AA02| NATAMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G01AA02 NATAMYCINUM OVULE 100 mg

PIMAFUCIN(R) 100 mg ASTELLAS PHARMA EUROPE

B.V.

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| 247 |G01AA51| COMBINATII | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G01AA51 COMBINATII CREMA VAG. 10 g/400000 UI

MACMIROR COMPLEX 10 g/400000 UI POLICHEM SA

G01AA51 COMBINATII CAPS. MOI VAG. 500 mg/200000 UI

MACMIROR COMPLEX 500 mg/200000 UI POLICHEM SA

G01AA51 COMBINATII CAPS. MOI VAG.

POLYGYNAX LAB. INNOTECH INT.

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| 248 |G01AF12| FENTICONAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G01AF12 FENTICONAZOLUM CAPS. MOI VAG. 600 mg

LOMEXIN 600 mg 600 mg RECORDATI SPA

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| 249 |G01AF15| BUTOCONAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G01AF15 BUTOCONAZOLUM CREMA VAG.

GYNOFORT 2% GEDEON RICHTER LTD.

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| 251 |G01AX05| NIFURATELUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G01AX05 NIFURATELUM DRAJ. 200 mg

MACMIROR 200 mg POLICHEM SA

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| 252 |G02CB03| CABERGOLINUM\* | Protocol: G001C |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G02CB03 CABERGOLINUM COMPR. 0.5 mg

DOSTINEX 0.5 mg PFIZER EUROPE MA EEG

Prescriere limitată: Prevenirea apariţiei lactaţiei în lăuzie.

Cod restricţie 2659: Hiperprolactinemia patologică pentru care nu este indicată intervenţia chirurgicală.

Cod restricţie 2660: Hiperprolactinemia patologică pentru care a fost utilizat tratament chirurgical dar cu rezultate incomplete.

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| 253 |G03AC02| LYNESTRENOLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03AC02 LYNESTRENOLUM COMPR. 0.5 mg

EXLUTON(R) 0.5 mg ORGANON NV

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| 254 |G03AC03| LEVONORGESTRELUM\*\*\*# | Protocol: G005N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03AC03 LEVONORGESTRELUM DISPOZITIV INTRAUTERIN 52 mg

MIRENA 20 micrograme/24 h 52 mg SCHERING OY (SCHERING AG)

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| 255 |G03CA04| ESTRIOLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03CA04 ESTRIOLUM CREMA VAG. 0.1%

OVESTIN 0.1% ORGANON NV

G03CA04 ESTRIOLUM OVULE 0.5 mg

OVESTIN 0.5 mg ORGANON NV

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| 256 |G03CA09| PROMESTRIENUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03CA09 PROMESTRIENUM CAPS. MOI VAG. 10 mg

COLPOTROPHINE(R) 10 mg LAB. THERAMEX

G03CA09 PROMESTRIENUM CREMA VAG. 1%

COLPOTROPHINE(R) 1% LAB. THERAMEX

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| 257 |G03DA04| PROGESTERONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03DA04 PROGESTERONUM CAPS. MOI 100 mg

UTROGESTAN(R) 100 mg 100 mg LAB. BESINS INTERNATIONAL

G03DA04 PROGESTERONUM GEL 1%

MASTOPROFEN 1% 1% ANTIBIOTICE SA

PROGESTOGEL 1% LAB. BESINS INTERNATIONAL

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| 258 |G03DB01| DYDROGESTERONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03DB01 DYDROGESTERONUM COMPR. FILM. 10 mg

DUPHASTON(R) 10 mg SOLVAY PHARMACEUTICALS BV

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| 259 |G03DC03| LYNESTRENOLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03DC03 LYNESTRENOLUM COMPR. 5 mg

ORGAMETRIL 5 mg ORGANON NV

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| 260 |G03DC05| TIBOLONUM\*\* | Protocol: G007N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03DC05 TIBOLONUM COMPR. 2,5 mg

LADYBON 2,5 mg ZENTIVA A.S.

G03DC05 TIBOLONUM COMPR. 2.5 mg

LIVIAL(R) 2,5 mg 2.5 mg ORGANON NV

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| 261 |G03FA01| COMBINATII\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FA01 COMBINATII COMPR. FILM. 1 mg/0,5 mg

ACTIVELLE 1 mg/0,5 mg 1 mg/0,5 mg NOVO NORDISK A/S

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| 262 |G03FA14| COMBINATII\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FA14 COMBINATII COMPR. FILM.

FEMOSTON(R) conti 1/5 SOLVAY PHARMACEUTICALS BV

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| 263 |G03FA15| ESTRADIOLUMVALERAT + DIENOGEST\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FA15 ESTRADIOLUMVALERAT + DRAJ.

DIENOGEST

KLIMODIEN SCHERING AG

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| 264 |G03FA17| COMBINATII\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FA17 COMBINATII COMPR. FILM.

ANGELIQ SCHERING AG

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| 265 |G03FB01| COMBINATII\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FB01 COMBINATII DRAJ.

CYCLO PROGINOVA(R) SCHERING AG

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| 266 |G03FB05| COMBINATII\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FB05 COMBINATII COMPR. FILM.

NOVOFEM NOVO NORDISK A/S

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| 267 |G03FB08| COMBINATII\*\* | Protocol: G002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03FB08 COMBINATII COMPR. FILM.

FEMOSTON 2/10 SOLVAY PHARMACEUTICALS BV

G03FB08 COMBINATII COMPR. FILM.

FEMOSTON 2/10 SOLVAY PHARMACEUTICALS BV

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| 268 |G03GA05| FOLITROPINUM ALFA\*\*\*\*# | Protocol: G003N |

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Prescriere limitată: Infertilitate anovulatorie.

NOTĂ:

Cu excepţia cazurilor de hipopituitarism sau amenoree primară, pacienta trebuie să fi fost tratată anterior cu citrat de clomifen şi/sau gonadorelin, iar tratamentul să fi rămas fără efect (sarcina nu a fost obţinută). Femeile care au urmat tratament pentru inducerea ovulaţiei cu alte clase de medicamente şi nu au obţinut o sarcină necesită evaluare laparoscopică pentru a exclude alte cauze care împiedică apariţia unei sarcini.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03GA05 FOLITROPINUM ALFA PULB. + SOLV. PT. SOL. 150 UI

INJ.

GONAL-f 150 UI 150 UI SERONO EUROPE LTD.

G03GA05 FOLITROPINUM ALFA PULB. + SOLV. PT. SOL. 75 UI

INJ.

GONAL-f 75 UI 75 UI SERONO EUROPE LTD.

Prescriere limitată: Infertilitate anovulatorie

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| 269 |G03GA06| FOLLITROPINUM BETA\*\*\*\*# | Protocol: G008N |

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Prescriere limitată: Infertilitate anovulatorie.

NOTĂ:

Cu excepţia cazurilor de hipopituitarism sau amenoree primară, pacienta trebuie să fi fost tratată anterior cu citrat de clomifen şi/sau gonadorelin, iar tratamentul să fi rămas fără efect (sarcina nu a fost obţinută). Femeile care au urmat tratament pentru inducerea ovulaţiei cu alte clase de medicamente şi nu au obţinut o sarcină necesită evaluare laparoscopică pentru a exclude alte cauze care împiedică apariţia unei sarcini.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03GA06 FOLLITROPINUM BETA SOL. INJ. 100 UI/0.5 ml

PUREGON 100 UI/0,5 ml 100 UI/0.5 ml N.V. ORGANON

G03GA06 FOLLITROPINUM BETA SOL. INJ. 300 UI/0.36 ml

PUREGON 300 UI/0,36 ml 300 UI/0.36 ml N.V. ORGANON

G03GA06 FOLLITROPINUM BETA SOL. INJ. 50 UI/0.5 ml

PUREGON 50 UI/0,5 ml 50 UI/0.5 ml N.V. ORGANON

G03GA06 FOLLITROPINUM BETA SOL. INJ. 600 UI/0.72 ml

PUREGON 600 UI/0.72 ml 600 UI/0.72 ml N.V. ORGANON

G03GA06 FOLLITROPINUM BETA PULB + SOLV. PT. SOL. 100 UI/ml

INJ.

PUREGON 100 UI 100 UI/ml ORGANON NV

G03GA06 FOLLITROPINUM BETA PULB + SOLV. PT. SOL. 50 UI/ml

INJ

PUREGON 50 UI 50 UI/ml ORGANON NV

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| 271 |G03GB02| CLOMIFENUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03GB02 CLOMIFENUM COMPR. 50 mg

CLOSTILBEGYT 50 mg EGIS PHARMACEUTICALS PLC

OVA-MIT 50 mg REMEDICA LTD.

Prescriere limitată: Infertilitate anovulatorie

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| 272 |G03XC01| RALOXIFENUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G03XC01 RALOXIFENUM COMPR. FILM. 60 mg

EVISTA 60 mg 60 mg ELI LILLY NEDERLAND BV

Cod restricţie 3006: Monoterapie cu medicamente antiresorbtive pentru tratamentul şi profilaxia osteoporozei sexoidoprive la femei până la vârsta de 60 ani.

NOTĂ:

Agenţii antiresorbtivi utilizaţi în tratamentul osteoporozei instalate sunt: ALENDRONATE SODIUM, RISEDRONAT SODIUM, RALOXIFEN HYDROCHLORIDE, IBANDRONATE, ZOLENDRONATE ŞI STRONTIUM RANELATE (agent antiresorbtiv şi formator osos).

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| 273 |G04BD04| OXYBUTYNINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04BD04 OXYBUTYNINUM COMPR. 5 mg

DRIPTANE(R) 5 mg 5 mg LAB. FOURNIER SA

Prescriere limitată: Hipereactivitate a detrusorului

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| 274 |G04BD07| TOLTERODINUM\*\*# | Protocol: G010N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04BD07 TOLTERODINUM CAPS. ELIB. PREL. 4 mg

DETRUSITOL SR 4 mg 4 mg PFIZER EUROPE MA EEIG

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| 275 |G04BD08| SOLIFENACINUM SUCCINATE\*\*# | Protocol: G009N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04BD08 SOLIFENACINUM COMPR. FILM. 10 mg

SUCCINATE

VESICARE 10 mg 10 mg ASTELLAS PHARMA EUROPE

B.V.

G04BD08 SOLIFENACINUM COMPR. FILM. 5 mg

SUCCINATE

VESICARE 5 mg 5 mg ASTELLAS PHARMA EUROPE

B.V.

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| 276 |G04BD09| TROSPIUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04BD09 TROSPIUM COMPR. FILM. 15 mg

INKONTAN 15 mg 15 mg PHARMAZEUTISCHE FABRIK

MONTAVIT GES. M.B.H.

G04BD09 TROSPIUM COMPR. FILM. 30 mg

INKONTAN 30 mg 30 mg PHARMAZEUTISCHE FABRIK

MONTAVIT GES. M.B.H.

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| 277 |G04CA01| ALFUZOSINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04CA01 ALFUZOSINUM COMPR. ELIB. PREL. 10 mg

ALFURAN MR 10 mg 10 mg TERAPIA S.A.

XATRAL SR 10 mg 10 mg SANOFI-SYNTHELABO FRANCE

G04CA01 ALFUZOSINUM COMPR. FILM. ELIB. 5 mg

PREL.

XATRAL(R) LP 5 mg 5 mg SANOFI-SYNTHELABO FRANCE

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| 278 |G04CA02| TAMSULOSINUM | |

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Prescriere limitată: Tratamentul hiperplaziei benigne de prostată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04CA02 TAMSULOSINUM CAPS. ELIB. MODIF. 0,4 mg

TAMSOL 0,4 mg 0,4 mg GEDEON RICHTER ROMANIA

TAMSULOSIN ACTAVIS 0,4 mg ACTAVIS GROUP HF.

G04CA02 TAMSULOSINUM CAPS. ELIB. MODIF. 0.4 mg

FOKUSIN 0.4 mg ZENTIVA AS

G04CA02 TAMSULOSINUM CAPS. ELIB. PREL. 0.4 mg

CONTIFLO MR 0.4 mg RANBAXY UK LTD.

TANYZ 0.4 mg KRKA D.D. NOVO MESTO

G04CA02 TAMSULOSINUM COMPR. FILM. ELIB. 0.4 mg

PREL.

OMNIC TOCAS(R) 0,4 0.4 mg ASTELLAS PHARMA EUROPE

B.V.

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| 279 |G04CB01| FINASTERIDUM\* | |

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Prescriere limitată: Tratamentul hiperplaziei benigne de prostată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04CB01 FINASTERIDUM COMPR. FILM. 5 mg

MOSTRAFIN 5 mg PLIVA LJUBLIJANA D.O.O.

PROSCAR 5 mg 5 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

TAREDOX 5 mg 5 mg DR. REDDY'S LABORATORIES

G04CB01 FINASTERIDUM COMPR. FILM. 5 mg

MOSTRAFIN 5 mg PLIVA LJUBLIJANA D.O.O.

PROSCAR 5 mg 5 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

TAREDOX 5 mg 5 mg DR. REDDY'S LABORATORIES

G04CB01 FINASTERIDUM COMPR. FILM. 5 mg

FINASTERID SANDOZ 5 mg 5 mg HEXAL AG

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| 280 |G04CB02| DUTASTERIDUM\* | |

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Prescriere limitată: Tratamentul hiperplaziei benigne de prostată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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G04CB02 DUTASTERIDUM CAPS. MOI 0.5 mg

AVODART(R) 0.5 mg GLAXO GROUP LTD.

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| 281 |H01AA02| TETRACOSACTIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H01AA02 TETRACOSACTIDUM SUSP. INJ. 1 mg/ml

SYNACHTEN DEPOT 1 mg/ml NOVARTIS PHARMA GMBH

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| 282 |H01AC01| SOMATROPINUM\*\*\*# | Protocol: H009E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H01AC01 SOMATROPINUM SOL. INJ. 10 mg/1.5 ml

NORDITROPIN 10 mg/1.5 ml NOVO NORD ISK A/S

SIMPLEX x 10 mg/1,5 ml

H01AC01 SOMATROPINUM SOL. INJ. 10 mg/2 ml

NUTROPINAq 10 mg/2 ml 10 mg/2 ml IPSEN LIMITED

H01AC01 SOMATROPINUM SOL. INJ. 3,3 mg/ml

OMNITROPE 3,3 mg/ml 3,3 mg/ml SANDOZ GMBH

H01AC01 SOMATROPINUM LIOF. + SOLV. PT. SOL. 4 mg (12 ui)

INJ.

ZOMACTON 4 mg (12 ui) FERING GMBH

H01AC01 SOMATROPINUM PULB. + SOLV. PT. SOL. 5.3 mg/ml (16 ui)

INJ.

GENOTROPIN(R) 16 ui (5,3 mg) 5.3 mg/ml (16 ui) PFIZER EUROPE MA EEIG

H01AC01 SOMATROPINUM SOL. INJ. 5 mg/1.5 ml

NORDITROPIN 5 mg/1.5 ml NOVO NORDISK A/S

SIMPLEX x 5 mg/1,5 ml

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| 283 |H01CC01| GANIRELIXUM\*\*\*# | Protocol: G004N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H01CC01 GANIRELIXUM SOL. INJ. 0.25 mg/0.5 ml

ORGALUTRAN 0.25 mg/0.5 ml 0.25 mg/0.5 ml N.V. ORGANON

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| 284 |H01CC02| CETRORELIXUM\*\*\*# | Protocol: H004E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H01CC02 CETRORELIXUM LIOF. + SOLV. PT. SOL. 0.25 mg

INJ.

CETROTIDE 0,25 mg 0.25 mg SERONO EUROPE LTD.

H01CC02 CETRORELIXUM LIOF. + SOLV. PT. SOL. 3 mg

INJ.

CETROTIDE 3 mg 3 mg SERONO EUROPE LTD.

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| 285 |H02AA02| FLUDROCORTISONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AA02 FLUDROCORTISONUM COMPR. 0.1 mg

ASTONIN H 0.1 mg MERCK KGAA

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| 286 |H02AB01| BETAMETHASONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AB01 BETAMETHASONUM SUSP. INJ. I.M. 7 mg/ml

DIPROPHOS(R) 7 mg/ml SCHERING PLOUGH EUROPE

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| 287 |H02AB06| PREDNISOLONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AB06 PREDNISOLONUM LIOF. + SOLV. PT. SOL. 250 mg

INJ.

SOLU-DECORTIN H 250 250 mg MERCK KGAA

H02AB06 PREDNISOLONUM LIOF. + SOLV. PT. SOL. 50 mg

INJ.

SOLU-DECORTIN H 50 50 mg MERCK KGAA

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| 288 |H03BB02| THIAMAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H03BB02 THIAMAZOLUM COMPR. FILM. 10 mg

THYROZOL(R) 10 mg 10 mg MERCK KGAA

H03BB02 THIAMAZOLUM COMPR. FILM. 20 mg

THYROZOL(R) 20 mg 20 mg MERCK KGAA

H03BB02 THIAMAZOLUM COMPR. FILM. 5 mg

THYROZOL(R) 5 mg 5 mg MERCK KGAA

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| 289 |H05BA01| CALCITONINUM (SOMON) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H05BA01 CALCITONINUM (SOMON) SOL. INJ. 100 UI

TONOCALCIN 100 UI 100 UI ALFA WASSERMANN SPA

H05BA01 CALCITONINUM (SOMON) SPRAY NAZ., SOL. 100 ui/doza

NYLEX 100 Ul/doza 100 UI/doza PHARMACEUTICAL IND. PROEL

EPAM. G. CORONIS SA

H05BA01 CALCITONINUM (SOMON) SOL. INJ. 100 UI/ml

NYLEX(R) 100 UI/ml PROEL E.P. CORONIS SA

H05BA01 CALCITONINUM (SOMON) SPRAY NAZAL-SOL. 200 UI/doza

MIACALCIC(R) NASAL 200 200 ui/doza NOVARTIS PHARMA GMBH

NYLEX(R) 200 UI/doza PROEL E.P. CORONIS SA

H05BA01 CALCITONINUM (SOMON) SOL. INJ. 50 ui/ml

MIACALCIC(R) 50 ui/ml NOVARTIS PHARMA GMBH

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| 290 |J01CR01| AMPICILLINUM + SULBACTAM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CR01 AMPICILLINUM + PULB. PT. SOL. INJ. 1 g + 500 mg

SULBACTAM

AMPIPLUS(R) 1,5 g 1 g + 500 mg ANTIBIOTICE SA

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| 291 |J01CR04| SULTAMICILLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01CR04 SULTAMICILLINUM PULB. PT. SUSP. ORALĂ 250 mg/5 ml

UNASYN 250 mg/5 ml PFIZER EUROPE MA EEIG

J01CR04 SULTAMICILLINUM COMPR. FILM. 375 mg

UNASYN 375 mg PFIZER EUROPE MA EEIG

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| 292 |J01DC10| CEFPROZILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DC10 CEFPROZILUM PULB. PT. SUSP. ORALĂ 125 mg/5 ml

CEFZIL 125 mg/5 ml 125 mg/5 ml BRISTOL-MYERS SQUIBB KFT.

J01DC10 CEFPROZILUM COMPR. FILM. 250 mg

CEFZIL 250 mg 250 mg BRISTOL-MYERS SQUIBB KFT.

J01DC10 CEFPROZILUM PULB. PT. SUSP. ORALĂ 250 mg/5 ml

CEFZIL 250 mg/5 ml 250 mg/5 ml BRISTOL-MYERS SQUIBB KFT.

J01DC10 CEFPROZILUM COMPR. FILM. 500 mg

CEFZIL 500 mg 500 mg BRISTOL-MYERS SQUIBB KFT.

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| 293 |J01DD14| CEFTIBUTENUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01DD14 CEFTIBUTENUM PULB. PT. SUSP. ORALĂ 36 mg/ml

CEDAX 36 mg/ml SCHERING PLOUGH EUROPE

J01DD14 CEFTIBUTENUM CAPS. 400 mg

CEDAX 400 mg SCHERING PLOUGH EUROPE

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| 294 |J01FA02| SPIRAMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01FA02 SPIRAMYCINUM COMPR. FILM. 1.5 M ui

ROVAMYCINE(R) 1,5 Mil. UI 1.5 M ui LAB. AVENTIS

J01FA02 SPIRAMYCINUM COMPR. FILM. 3 M ui

ROVAMYCINE(R) 3 Mil. UI 3 M ui LAB. AVENTIS

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| 295 |J01FA10| AZITHROMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01FA10 AZITHROMYCINUM PULB. PT. SUSP. ORALĂ 100 mg/5 ml

AZITROMICINA 100 mg/5 ml SANDOZ SRL

SANDOZ 100 mg/5 ml

J01FA10 AZITHROMYCINUM COMPR. FILM. 125 mg

SUMAMED 125 mg 125 mg PLIVA LJUBLJANA D.O.O.

J01FA10 AZITHROMYCINUM PULB. PT. SUSP. ORALĂ 200 mg/5 ml

AZITROMICINA 200 mg/5 ml SANDOZ SRL

SANDOZ 200 mg/5 ml

J01FA10 AZITHROMYCINUM PULB. + SOLV. SUSP. 200 mg/5 ml

ORALĂ

AZITROX 200 mg/5 ml 200 mg/5 ml ECZACIBASI PHARMACEUTICALS

S.R.L.

J01FA10 AZITHROMYCINUM PULB. PT. SUSP. ORALĂ 200 mg/5 ml

SUMAMED FORTE 200 mg/5 ml PLIVA LJUBLIJANA D.O.O.

J01FA10 AZITHROMYCINUM CAPS. 250 mg

AZATRIL 250 mg 250 mg BALKANPHARMA RAZGRAD AD

J01FA10 AZITHROMYCINUM COMPR. FILM. 250 mg

AZITROMICINA SANDOZ 250 mg 250 mg SANDOZ S.R.L.

COMPRIMATE FILMATE

AZITROX(R) 250 250 mg ZENTIVA AS

J01FA10 AZITHROMYCINUM COMPR. FILM. 500 mg

AZITROMICINA SANDOZ 500 mg 500 mg SANDOZ S.R.L.

COMPRIMATE FILMATE

AZITROX(R) 500 500 mg ZENTIVA AS

AZRO(R) 500 mg 500 mg ECZACIBASI PHARMACEUTICALS

S.R.L.

SUMAMED 500 mg 500 mg PLIVA LJUBLJANA D.O.O.

ZITROCIN 500 mg 500 mg OZONE LABORATORIES LTD.

J01FA10 AZITHROMYCINUM GRAN. ELIB. PREL. PT. 2 g

SUSP. ORALĂ

ZMAX 2 g 2 g PFIZER EUROPE MA EEIG

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| 296 |J01FF01| CLINDAMYCINUM | |

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Prescriere limitată: Infecţii cu coci Gram pozitivi care nu pot fi tratate

eficient cu peniciline.

Infecţii severe cu germeni anaerobi.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01FF01 CLINDAMYCINUM CAPS. 150 mg

DALACIN C 150 mg 150 mg PFIZER EUROPE MA EEIG

J01FF01 CLINDAMYCINUM CAPS. 300 mg

DALACIN C 300 mg 300 mg PFIZER EUROPE MA EEIG

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| 297 |J01MA12| LEVOFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MA12 LEVOFLOXACINUM COMPR. FILM. 500 mg

TAVANIC(R) 500 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

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| 298 |J01MA14| MOXIFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J01MA14 MOXIFLOXACINUM COMPR. FILM. 400 mg

AVELOX(R) 400 mg 400 mg BAYER HEALTHCARE AG

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| 299 |J02AC02| ITRACONAZOLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J02AC02 ITRACONAZOLUM CAPS. 100 mg

ITRACONAZOL 100 mg 100 mg TERAPIA S.A.

OMICRAL 100 mg 100 mg MEDICO UNO PHARMACEUTICAL

S.R.L

ORUNGAL 100 mg JANSSEN PHARMACEUTICA NV

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| 300 |J05AB11| VALACYCLOVIRUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J05AB11 VALACYCLOVIRUM COMPR. FILM. 500 mg

VALTREX 500 mg 500 mg THE WELLCOME FOUNDATION

LTD

Prescriere limitată: Tratamentul pacienţilor cu herpes zoster în decurs de 72 de ore de la debutul rash-ului

Prescriere limitată: Herpes zoster oftalmic.

Prescriere limitată: Herpes genital iniţial moderat/sever.

Prescriere limitată: Tratamentul episodic sau supresiv al herpesului genital recurent (forme moderate/severe).

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| 301 |J05ABN1| BRIVUDINUM\*# | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul pacienţilor cu herpes zoster în decurs de 72 de ore de la debutul rash-ului.

J05ABN1 BRIVUDINUM COMPR. 125 mg

BRIVAL(R) 125 mg BERLIN CHEMIE AG MENARINI

GROUP

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| 302 |L01AA01| CYCLOPHOSPHAMIDUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L01AA01 CYCLOPHOSPHAMIDUM DRAJ. 50 mg

ENDOXAN(R) 50 mg 50 mg BAXTER ONCOLOGY GMBH

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| 305 |L02AE04| TRIPTORELINUM\*\*\*# | Protocol: L013E |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul de scurtă durată (până la 6 luni) al

endometriozei confirmate histologic.

Tratamentul pubertăţii precoce.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

L02AE04 TRIPTORELINUM LIOF. + SOLV. PT. 0.1 mg

SOL. INJ.

DIPHERELINE 0,1 mg 0.1 mg BEAUFOUR IPSEN PHARMA

L02AE04 TRIPTORELINUM LIOF. + SOLV. PT. SUSP. 11.25 mg

INJ. I.M. ELIB. PREL

DIPHERELINE(R) 11,25 mg 11.25 mg BEAUFOUR IPSEN PHARMA

L02AE04 TRIPTORELINUM LIOF. +SOLV. PT. SUSP. 3.75 mg

INJ. I.M. ELIB. PREL.

DIPHERELINE(R) 3,75 mg 3.75 mg BEAUFOUR IPSEN PHARMA

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| 306 |L04AX01| AZATHIOPRINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L04AX01 AZATHIOPRINUM COMPR. FILM. 50 mg

IMURAN(R) 50 mg THE WELLCOME FOUNDATION

LTD.

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| 307 |M01AC01| PIROXICAMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M01AC01 PIROXICAMUM COMPR. 20 mg

FLAMEXIN 20 mg CHIESI FARMACEUTICI SPA

N-PIROXICAM MEDUMAN 20 mg 20 mg MEDUMAN S.A.

PIROXICAM 20 mg 20 mg ARENA GROUP SA

PIROXICAM LPH 20 mg 20 mg LABORMED PHARMA SA

PIROXSAL 20 mg SLAVIA PHARM SRL

M01AC01 PIROXICAMUM COMPR. EFF. 20 mg

FLAMEXIN 20 mg CHIESI FARMACEUTICI S.P.A.

M01AC01 PIROXICAMUM SUPOZ. 20 mg

PIROXICAM 20 mg 20 mg SINTOFARM SA

M01AC01 PIROXICAMUM SOL. INJ. 20 mg/ml

FELDENE(R) 20 mg/ml 20 mg/ml PFIZER EUROPE MA EEIG

HOTEMIN 20 mg/ml EGIS PHARMACEUTICALS PLC

M01AC01 PIROXICAMUM PULB. PT. SOL. ORALĂ 20 mg/plic

FLAMEXIN(R) 20 mg/plic CHIESI FARMACEUTICI SPA

M01AC01 PIROXICAMUM SUPOZ. 40 mg

PIROXICAM 40 mg 40 mg ANTIBIOTICE SA

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| 308 |M01AC05| LORNOXICAMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M01AC05 LORNOXICAMUM COMPR. FILM. 4 mg

XEFO(R) 4 mg 4 mg NYCOMED AUSTRIA GMBH

M01AC05 LORNOXICAMUM COMPR. FILM. 8 mg

XEFO(R) 8 mg 8 mg NYCOMED AUSTRIA GMBH

M01AC05 LORNOXICAMUM PULB. + SOLV. PT. 8 mg/2 ml

SOL. INJ.

XEFO 8 mg/2 ml 8 mg/2 ml NYCOMED AUSTRIA GMBH

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| 309 |M01AE17| DEXKETOPROFENUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M01AE17 DEXKETOPROFENUM COMPR. FILM. 25 mg

TADOR 25 mg BERLIN CHEMIE AG MENARINI

GROUP

M01AE17 DEXKETOPROFENUM SOL. INJ./CONC. 50 mg/2 ml

PT. SOL. PERF.

TADOR INJECT 50 mg/2 ml MENARINI INTERNATIONAL

OPERATIONS S.A.

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| 310 |M01AH01| CELECOXIBUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul simptomatic antiinflamator la pacienţii cu intoleranţă la AINS neselective.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M01AH01 CELECOXIBUM CAPS. 100 mg

CELEBREX 100 mg 100 mg PFIZER EUROPE MA EEIG

M01AH01 CELECOXIBUM CAPS. 200 mg

CELEBREX 200 mg 200 mg PFIZER EUROPE MA EEIG

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| 311 |M01AH05| ETORICOXIBUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul simptomatic antiinflamator la pacienţii cu intoleranţă la AINS neselective.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M01AH05 ETORICOXIBUM COMPR. FILM. 120 mg

ARCOXIA(R) 120 mg 120 mg MERCK SHARP & DOHME

ROMANIA S.R.L

M01AH05 ETORICOXIBUM COMPR. FILM. 60 mg

ARCOXIA(R) 60 mg 60 mg MERCK SHARP & DOHME

ROMANIA S.R.L

M01AH05 ETORICOXIBUM COMPR. FILM. 90 mg

ARCOXIA(R) 90 mg 90 mg MERCK SHARP & DOHME

ROMANIA S.R.L

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| 312 |M01AX05| GLUCOSAMINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M01AX05 GLUCOSAMINUM PULB. PT. SOL. ORALĂ 150 mg/plic

DONA(R) 150 mg/plic ROTTAPHARM SPA

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| 313 |M01AX17| NIMESULIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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A se administra cu precauţie la pacienţii cu istoric de factori de risc sau afecţiuni gastrointestinale.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M01AX17 NIMESULIDUM COMPR. 100 mg

APONIL 100 mg MEDOCHEMIE LTD

AULIN(R) 100 mg 100 mg CSC PHARMACEUTICALS

HANDELS GMBH

COXTRAL 100 mg ZENTIVA AS

LEMESIL 100 mg ANFARM HELLAS S.A.

PHARMACEUTICALS

NIMESULID LPH 100 mg 100 mg LABORMED PHARMA SA

NIMESULID 100 mg 100 mg MAGISTRA C & C

NIMESULID ARENA 100 mg 100 mg ARENA GROUP S.A.

NIMESULID SLAVIA 100 mg SLAVIA PHARM SRL

NISE 100 mg 100 mg DR. REDDY'S LABORATORIES

M01AX17 NIMESULIDUM PULB. PT. SUSP. ORALĂ 100 mg

SULIDAMOR 100 mg FARMACEUTICI DAMOR SPA

M01AX17 NIMESULIDUM GRAN. PT. SUSP. ORALĂ 100 mg/plic

AULIN(R) 100 mg/plic CSC PHARMACEUTICALS

HANDELS GMBH

NIMESIL 100 mg/plic LAB. GUIDOTTI SPA

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| 314 |M03BX01| BACLOFENUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M03BX01 BACLOFENUM COMPR. 10 mg

LIORESAL(R) 10 mg NOVARTIS PHARMA GMBH

M03BX01 BACLOFENUM COMPR. 25 mg

LIORESAL(R) 25 mg NOVARTIS PHARMA GMBH

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| 315 |M05BA04| ACIDUM ALENDRONICUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Cod restricţie 3001: Tratamentul osteoporozei la pacienţi cu scor T </= - 2,5 măsurat DEXA (coloană vertebrală sau şold).

Cod restricţie 3002: Tratamentul osteoporozei la pacienţi cu scor T </= - 2 măsurat DEXA (coloană vertebrală sau şold) şi istoric de fracturi pe structuri osoase fragile.

Cod restricţie 3003: Tratamentul osteoporozei la paciente în postmenopauză care au suferit fracturi vertebrale osteoporotice în antecedente.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M05BA04 ACIDUM ALENDRONICUM COMPR. 10 mg

FOSAMAX 10 mg 10 mg MERCK SHARP & DOHME S.R.L

M05BA04 ACIDUM ALENDRONICUM COMPR. 70 mg

FOSAMAX 70 mg 70 mg MERCK SHARP & DOHME S.R.L

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| 316 |M05BA06| ACIDUM IBANDRONICUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Cod restricţie 3001: Tratamentul osteoporozei la pacienţi cu scor T </= - 2,5 măsurat DEXA (coloană vertebrală sau şold).

Cod restricţie 3002: Tratamentul osteoporozei la pacienţi cu scor T </= - 2 măsurat DEXA (coloană vertebrală sau şold) şi istoric de fracturi pe structuri osoase fragile.

Cod restricţie 3003: Tratamentul osteoporozei la paciente în postmenopauză care au suferit fracturi vertebrale osteoporotice în antecedente.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M05BA06 ACIDUM IBANDRONICUM COMPR. FILM. 150 mg

BONVIVA 150 mg 150 mg ROCHE REGISTRATION LTD.

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| 317 |M05BA07| ACIDUM RISEDRONICUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Cod restricţie 3001: Tratamentul osteoporozei la pacienţi cu scor T </= - 2,5 măsurat DEXA (coloană vertebrală sau şold).

Cod restricţie 3002: Tratamentul osteoporozei la pacienţi cu scor T </= - 2 măsurat DEXA (coloană vertebrală sau şold) şi istoric de fracturi pe structuri osoase fragile.

Cod restricţie 3003: Tratamentul osteoporozei la paciente în postmenopauză care au suferit fracturi vertebrale osteoporotice în antecedente.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M05BA07 ACIDUM RISEDRONICUM COMPR. FILM. 35 mg

ACTONEL(R) SAPTAMANAL 35 mg AVENTIS PHARMA AB

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| 318 |M05BX03| STRONTIUM RANELATUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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M05BX03 STRONTIUM RANELATUM GRAN. PT. SUSP. ORALĂ 2 g

OSSEOR 2 g 2 g LES LAB. SERVIER

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| 319 |M05BB03| COMBINATII (ACIDUM ALENDRONICUM + | |

| | | COLECALCIFEROLUM)\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Cod restricţie 3001: Tratamentul osteoporozei la pacienţi cu scor T </= - 2,5 măsurat DEXA (coloană vertebrală sau şold).

Cod restricţie 3002: Tratamentul osteoporozei la pacienţi cu scor T </= - 2 măsurat DEXA (coloană vertebrală sau şold) şi istoric de fracturi pe structuri osoase fragile.

Cod restricţie 3003: Tratamentul osteoporozei la paciente în postmenopauză care au suferit fracturi vertebrale osteoporotice în antecedente.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

M05BB03 COMBINATII (ACIDUM COMPR. 70 mg/2800 UI

ALENDRONICUM +

COLECALCIFEROLUM)

FOSAVANCE 70 mg/2800 UI 70 mg/2800 UI MERCK SHARP & DOHME LTD.

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| 320 |N02AA05| OXYCODONUM\*# | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Durere severă, invalidantă, care nu răspunde la

analgezice non-opioide.

Durere cronică severă, invalidantă, care nu răspunde la

analgezice non-opioide, unde durata totală a

tratamentului opioid este mai scurtă de 12 luni.

Risc înalt de apariţie a dependenţei.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

N02AA05 OXYCODONUM COMPR. FILM. ELIB. 10 mg

MODIF.

OXYCONTIN(R) 10 mg 10 mg MUNDIPHARMA GMBH

N02AA05 OXYCODONUM COMPR. FILM. ELIB. 20 mg

MODIF.

OXYCONTIN(R) 20 mg 20 mg MUNDIPHARMA GMBH

N02AA05 OXYCODONUM COMPR. FILM. ELIB. 40 mg

MODIF.

OXYCONTIN(R) 40 mg 40 mg MUNDIPHARMA GMBH

N02AA05 OXYCODONUM COMPR. FILM. ELIB. 80 mg

MODIF.

OXYCONTIN(R) 80 mg 80 mg MUNDIPHARMA GMBH

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| 321 |N02AA08| DIHYDROCODEINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N02AA08 DIHYDROCODEINUM COMPR. ELIB. PREL. 120 mg

DHC CONTINUS 120 mg 120 mg MUNDIPHARMA GMBH

N02AA08 DIHYDROCODEINUM COMPR. ELIB. PREL. 60 mg

DHC CONTINUS 60 mg 60 mg MUNDIPHARMA GMBH

N02AA08 DIHYDROCODEINUM COMPR. ELIB. PREL. 90 mg

DHC CONTINUS 90 mg 90 mg MUNDIPHARMA GMBH

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| 322 |N02AX52| COMBINATII (TRAMADOLUM + PARACETAMOLUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Durere severă, care nu răspunde la analgezice

non-opioide.

Pentru durere acută la care tratamentul cu aspirină

şi/sau paracetamol este contraindicat sau nu a dat

rezultate.

N02AX52 COMBINATII COMPR. FILM. 37,5 mg + 325 mg

(TRAMADOLUM +

PARACETAMOLUM)

ZALDIAR(R) 37,5 mg + 325 mg GRUNENTHAL GMBH

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| 323 |N03AE01| CLONAZEPAMUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N03AE01 CLONAZEPAMUM COMPR. 0.5 mg

RIVOTRIL 0.5 mg ROCHE ROMANIA S.R.L

N03AE01 CLONAZEPAMUM COMPR. 2 mg

RIVOTRIL(R) 2 mg ROCHE ROMANIA S.R.L.

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| 324 |N03AX16| PREGABALINUM\*\*# | Protocol: N025G;|

| | | | N032G |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N03AX16 PREGABALINUM CAPS. 150 mg

LYRICA 150 mg 150 mg PFIZER LTD.

N03AX16 PREGABALINUM CAPS. 300 mg

LYRICA 300 mg 300 mg PFIZER LTD.

N03AX16 PREGABALINUM CAPS. 75 mg

LYRICA 75 mg 75 mg PFIZER LTD.

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| 326 |N04BC05| PRAMIPEXOLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul simptomatic al sindromului idiopatic al

picioarelor neliniştite, forme moderate/severe.

La acest medicament au fost raportate episoade de

instalare bruscă a somnului fără avertizare în timpul

activităţii.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

N04BC05 PRAMIPEXOLUM COMPR. 0.18 mg

MIRAPEXIN 0,18 mg 0.18 mg BOEHRINGER INGELHEIM

INTERNATIONAL GMBH

N04BC05 PRAMIPEXOLUM COMPR. 0.7 mg

MIRAPEXIN 0,7 mg 0.7 mg BOEHRINGER INGELHEIM

INTERNATIONAL GMBH

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| 327 |N04BXN1| PIRIBEDILUM\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N04BXN1 PIRIBEDILUM DRAJ. ELIB. PREL. 50 mg

PRONORAN(R) 50 mg LP 50 mg LES LAB. SERVIER IND.

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| 328 |N05AL03| TIAPRIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N05AL03 TIAPRIDUM COMPR. 100 mg

TIAPRIDAL(R) 100 mg 100 mg SANOFI-SYNTHELABO FRANCE

N05AL03 TIAPRIDUM SOL. INJ. 100 mg/2 ml

TIAPRIDAL 100 mg/2 ml 100 mg/2 ml SANOFI-AVENTIS FRANCE

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| 329 |N05BE01| BUSPIRONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Pentru tratamentul de scurta durata al anxietăţii.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

N05BE01 BUSPIRONUM COMPR. 10 mg

SPITOMIN 10 mg 10 mg EGIS PHARMACEUTICALS PLC

STRESSIGAL 10 mg ANFARM HELLAS S.A.

PHARMACEUTICALS

N05BE01 BUSPIRONUM COMPR. 5 mg

SPITOMIN 5 mg 5 mg EGIS PHARMACEUTICALS PLC

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| 330 |N06AA21| MAPROTILINUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AA21 MAPROTILINUM COMPR. FILM. 10 mg

LUDIOMIL(R) 10 mg 10 mg NOVARTIS PHARMA GMBH

N06AA21 MAPROTILINUM COMPR. FILM. 25 mg

LUDIOMIL(R) 25 mg 25 mg NOVARTIS PHARMA GMBH

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| 331 |N06AB10| ESCITALOPRAMUM\*\* | Protocol: N009F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AB10 ESCITALOPRAMUM COMPR. FILM. 10 mg

CIPRALEX 10 mg 10 mg H. LUNDBECK A/S

N06AB10 ESCITALOPRAMUM COMPR. FILM. 5 mg

CIPRALEX 5 mg 5 mg H. LUNDBECK A/S

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| 332 |N06AX05| TRAZODONUM\*\* | Protocol: N010F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AX05 TRAZODONUM COMPR. ELIB. PREL. 150 mg

TRITTICO AC 150 mg 150 mg ANGELINI FRANCESCO SPA

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| 333 |N06AX14| TIANEPTINUM\*\* | Protocol: N011F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AX14 TIANEPTINUM DRAJ. 12.5 mg

COAXIL(R) 12.5 mg LES LAB. SERVIER IND.

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| 334 |N06AX17| MILNACIPRANUM\*\* | Protocol: N002F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AX17 MILNACIPRANUM CAPS. 25 mg

IXEL 25 mg PIERRE FABRE MEDICAMENT

N06AX17 MILNACIPRANUM CAPS. 50 mg

IXEL 50 mg PIERRE FABRE MEDICAMENT

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| 335 |N06AX21| DULOXETINUM\*\* | Protocol: N014F |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AX21 DULOXETINUM CAPS. GASTROREZ. 30 mg

CYMBALTA 30 mg 30 mg ELI LILLY NEDERLAND BV

N06AX21 DULOXETINUM SOL. PERF. 100 mg/50 ml

CYMBALTA 60 mg 60 mg ELI LILLY NEDERLAND BV

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| 336 |N06AXN1| BUPROPIONUM\*\*# | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06AXN1 BUPROPIONUM COMPR. FILM. ELIB. PREL. 150 mg

WELLBUTRIN SR 150 mg 150 mg GLAXO WELLCOME UK LTD.

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| 337 |N06BX16| PRAMIRACETAMUM\*\*# | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06BX16 PRAMIRACETAMUM COMPR. FILM. 600 mg

PRAMISTAR 600 mg F.I.R.M.A. S.p.a.

(MENARINI GROUP)

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| 338 |N06BX18| VINPOCETINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N06BX18 VINPOCETINUM COMPR. 10 mg

CAVINTON(R) FORTE 10 mg GEDEON RICHTER LTD.

N06BX18 VINPOCETINUM CAPS. 5 mg

VIMPOCETIN 5 mg 5 mg VIM SPECTRUM SRL

N06BX18 VINPOCETINUM COMPR. 5 mg

CAVINTON(R) 5 mg GEDEON RICHTER LTD.

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| 339 |N07CA01| BETAHISTINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N07CA01 BETAHISTINUM COMPR. 16 mg

BETASERC(R) 16 mg 16 mg SOLVAY PHARMACEUTICALS BV

N07CA01 BETAHISTINUM COMPR. 24 mg

BETASERC(R) 24 mg 24 mg SOLVAY PHARMACEUTICALS BV

N07CA01 BETAHISTINUM COMPR. 80 mg

URUTAL 8 mg 80 mg A & G MED TRADING S.R.L.

N07CA01 BETAHISTINUM COMPR. 8 mg

BETASERC(R) 8 mg 8 mg SOLVAY PHARMACEUTICALS BV

MICROSER 8 mg PRODOTTI FORMENTI

VESTIBO 8 mg 8 mg ACTAVIS GROUP HF.

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| 340 |N07CA52| COMBINATII (CINNARIZINUM + DIMENHIDRATUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N07CA52 COMBINATII COMPR.

(CINNARIZINUM +

DIMENHIDRATUM)

ARLEVERT HENNIG ARZNEIMITTEL GMBH &

CO.KG

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| 341 |N07XN01| HIDROLIZAT DE PROTEINA DIN CREIER DE | Protocol: N026F |

| | | PORCINA\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N07XN01 HIDROLIZAT DE SOL. INJ./PERF. 215.2 mg/ml

PROTEINA DIN CREIER

DE PORCINA

CEREBROLYSIN(R) 215.2 mg/ml EBEWE PHARMA GMBH NFG.KG

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| 342 |P01BA02| HYDROXYCHLOROQUINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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P01BA02 HYDROXYCHLOROQUINUM COMPR. FILM. 200 mg

PLAQUENIL(R) 200 mg SANOFI-SYNTHELABO LTD.

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| 343 |R03AC04| FENOTEROLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03AC04 FENOTEROLUM SOL. DE INHALAT 100 micrograme/doză

PRESURIZATĂ

BEROTEC N-100 micrograme/doză 100 micrograme/doză BOEHRINGER INGELHEIM

INT. GMBH

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| 344 |R01AD05| BUDESONIDUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Profilaxia şi tratamentul rinitei alergice.

R01AD05 BUDESONIDUM SPRAY NAZAL SUSP. 32 micrograme/doză

RHINOCORT(R) AQUA 32 micrograme/doză ASTRAZENECA AB

R01AD05 BUDESONIDUM SPRAY NAZ. SUSP. 50 micrograme/doză

TAFEN NASAL 50 micrograme/doză LEK PHARMACEUTICALS D.D.

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| 345 |R01AD08| FLUTICASONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R01AD08 FLUTICASONUM SPRAY NAZAL SUSP. 50 micrograme/doză

FLIXONASE(R) 50 micrograme/doză GLAXOWELLCOME UK LTD.

Prescriere limitată: Profilaxia şi tratamentul rinitei alergice

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| 346 |R01AD09| MOMETASONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R01AD09 MOMETASONUM SPRAY NAZAL SUSP. 50 micrograme/doză

NASONEX 50 micrograme/doză SCHERING PLOUGH EUROPE

Prescriere limitată: Profilaxia şi tratamentul rinitei alergice.

Prescriere limitată: Tratamentul polipozelor nazale a adultului.

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| 347 |R03AK03| COMBINATII (FENOTEROLUM + IPRATROPIUM)\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03AK03 COMBINATII AEROSOL SOL. INHAL. 0,020 mg + 0,050 mg

(FENOTEROLUM +

IPRATROPIUM)

BERODUAL(R) N 0,020 mg + 0,050 mg BOEHRINGER INGELHEIM

INT. GMBH

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| 348 |R03AK06| COMBINATII (SALMETEROLUM + FLUTICASONUM)\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratament simptomatic al bolii pulmonare cronice

obstructive (BPOC) la pacienţii cu FEV1 mai mică decât

50% faţă de normal şi cu istoric de exacerbări repetate

şi simptome importante în timpul tratamentului

bronhodilatator cu agonişti ai receptorilor beta-2

adrenergici.

Tratamentul astmului bronşic care nu este controlat

adecvat cu corticosteroizi inhalatori şi

beta-2-agonişti inhalatori de scurtă durată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

R03AK06 COMBINATII AEROSOL SUSP. INHAL. 25 mg/125 mg

(SALMETEROLUM +

FLUTICASONUM)

SERETIDE(R) 25/125 micrograme 25 mg/125 mg GLAXOWELLCOME UK LTD.

INHALER CFC FREE

R03AK06 COMBINATII AEROSOL SUSP. INHAL. 25 mg/250 mg

(SALMETEROLUM +

FLUTICASONUM)

SERETIDE(R) 25/250 micrograme 25 mg/250 mg GLAXOWELLCOME UK LTD.

NHALER CFC FREE

R03AK06 COMBINATII AEROSOL SUSP. INHAL. 25 mg/50 mg

(SALMETEROLUM +

FLUTICASONUM)

SERETIDE(R) 25/50 micrograme 25 mg/50 mg GLAXOWELLCOME UK LTD.

INHALER CFC FREE

R03AK06 COMBINATII PULB. INHAL. 50 mg/100 mg

(SALMETEROLUM +

FLUTICASONUM)

SERETIDE DISKUS 50/100 50 mg/100 mg GLAXO WELLCOME UK LIMITED

R03AK06 COMBINATII PULB. INHAL. 50 mg/250 mg

(SALMETEROLUM +

FLUTICASONUM)

SERETIDE DISKUS 50/250 50 mg/250 mg GLAXOWELLCOME UK LTD.

R03AK06 COMBINATII PULB. INHAL. 50 mg/500 mg

(SALMETEROLUM +

FLUTICASONUM)

SERETIDE DISKUS 50/500 50 mg/500 mg GLAXOWELLCOME UK LTD.

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| 349 |R03AK07| COMBINATII (BUDESONIDUM + FORMOTEROLUM)\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratament simptomatic al bolii pulmonare cronice

obstructive (BPCO) la pacienţii cu FEV1 mai mică decât

50% faţă de normal şi cu istoric de exacerbări repetate

şi simptome importante în timpul tratamentului

bronhodilatator cu agonişti ai receptorilor beta-2

adrenergici.

Tratamentul astmului bronşic care nu este controlat

adecvat cu corticosteroizi inhalatori şi

beta-2-agonişti inhalatori de scurtă durată.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

R03AK07 COMBINATII PULB. INHAL. 160/4.5 micrograme

(BUDESONIDUM +

FORMOTEROLUM)

SYMBIOCORT(R) TURBUHALER(R) 160/4.5 micrograme ASTRAZENECA AB

160/4,5 micrograme

R03AK07 COMBINATII PULB. INHAL. 320/9 micrograme

(BUDESONIDUM +

FORMOTEROLUM)

SYMBIOCORT(R) TURBUHALER(R) 320/9 micrograme ASTRAZENECA AB

320/9 micrograme

R03AK07 COMBINATII PULB. INHAL. 80/4.5 micrograme

(BUDESONIDUM +

FORMOTEROLUM)

SYMBIOCORT(R) TURBUHALER(R) 80/4.5 micrograme ASTRAZENECA AB

80/4,5 micrograme

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| 350 |R03BA05| FLUTICASONUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul de control al astmului bronşic persistent.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

R03BA05 FLUTICASONUM SUSP. INHAL. 0.5 mg/2 ml

FLIXOTIDE(R) NEBULES(R) 0.5 mg/2 ml GLAXO WELLCOME UK LTD.

0.5 mg/2 ml

R03BA05 FLUTICASONUM SUSP. INHAL. 125 micrograme/doză

PRESURIZATĂ

FLIXOTIDE 125 INHALER 125 micrograme/doză GLAXO WELLCOME UK LTD.

CFC - Free

R03BA05 FLUTICASONUM SUSP. INHAL. 2 mg/2 ml

FLIXOTIDE(R) NEBULES(R) 2 mg/2 ml GLAXO WELLCOME UK LTD.

2 mg/2 ml

R03BA05 FLUTICASONUM SUSP. INHAL. 50 micrograme/doză

PRESURIZATĂ

FLIXOTIDE(R) 50 INHALER 50 micrograme/doză GLAXO WELLCOME UK LTD.

CFC - Free

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| 352 |R03BA08| CICLESONIDUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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Prescriere limitată: Tratamentul de fond al astmului bronşic persistent.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

R03BA08 CICLESONIDUM SOL. DE INHALAT 160 micrograme/doză

PRESURIZATĂ

ALVESCO 160 INHALER 160 micrograme/doză ALTANA PHARMA AG

R03BA08 CICLESONIDUM SOL. DE INHALAT 80 micrograme/doză

PRESURIZATĂ

ALVESCO 80 INHALER 80 micrograme/doză ALTANA PHARMA AG

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| 353 |R03BB04| TIOTROPIUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03BB04 TIOTROPIUM CAPS. CU PULB. INHAL. 18 micrograme

SPIRIVA(R) 18 micrograme 18 micrograme BOEHRINGER INGELHEIM

PHARMA GMBH & CO.KG

Prescriere limitată: Pentru tratamentul de întreţinere pe termen lung al bronhospasmului şi dispneei asociate bolii pulmonare obstructive cronice

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| 354 |R03DA05| AMINOPHYLLINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03DA05 AMINOPHYLLINUM CAPS. 100 mg

MIOFILIN 100 mg 100 mg ZENTIVA SA

R03DA05 AMINOPHYLLINUM COMPR. 100 mg

AMINOFILINA EEL 100 mg BIO EEL SRL

R03DA05 AMINOPHYLLINUM COMPR. 200 mg

AMINOFILINA 200 mg 200 mg ARENA GROUP SA

R03DA05 AMINOPHYLLINUM SOL. INJ. 24 mg/ml

MIOFILIN 24 mg/ml ZENTIVA S.A.

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| 355 |R03DC03| MONTELUKASTUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03DC03 MONTELUKASTUM COMPR. FILM. 10 mg

SINGULAIR(R) 10 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

Prescriere limitată: Tratamentul combinat al astmului bronşic persistent la pacienţii cu simptomatologie necontrolată corespunzător cu corticosteroizi inhalatori şi beta-2-agonişti cu durată scurtă de acţiune.

R03DC03 MONTELUKASTUM COMPR. MAST. 4 mg

SINGULAIR 4 mg MERCK SHARP & DOHME S.R.L.

Prescriere limitată: Tratamentul combinat al astmului bronşic persistent la pacienţii cu simptomatologie necontrolată corespunzător cu corticosteroizi inhalatori şi beta-2-agonişti cu durată scurtă de acţiune.

Cod restricţie 2617: Tratamentul astmului bronşic persistent la copii cu vârste cuprinse între 2 şi 5 ani la care nu se poate administra terapie cu corticosteroizi inhalatori.

R03DC03 MONTELUKASTUM GRANULE 4 mg/plic

SINGULAIR(R) 4 mg/plic MERCK SHARP & DOHME

ROMANIA S.R.L.

Prescriere limitată: Tratamentul combinat al astmului bronşic persistent la pacienţii cu simptomatologie necontrolată corespunzător cu corticosteroizi inhalatori şi beta-2-agonişti cu durată scurtă de acţiune.

Cod restricţie 2617: Tratamentul astmului bronşic persistent la copii cu vârste cuprinse între 2 şi 5 ani la care nu se poate administra terapie cu corticosteroizi inhalatori.

R03DC03 MONTELUKASTUM COMPR. MAST. 5 mg

SINGULAIR(R) 5 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

Prescriere limitată: Tratamentul combinat al astmului bronşic persistent la pacienţii cu simptomatologie necontrolată corespunzător cu corticosteroizi inhalatori şi beta-2-agonişti cu durată scurtă de acţiune.

Cod restricţie 2617: Tratamentul astmului bronşic persistent la copii cu vârste cuprinse între 6 şi 14 ani la care nu se poate administra terapie cu corticosteroizi inhalatori.

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| 356 |R03DX03| FENSPIRIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R03DX03 FENSPIRIDUM SIROP 0.2%

EURESPAL(R) 0.2% LES LAB. SERVIER IND.

R03DX03 FENSPIRIDUM COMPR. FILM. 80 mg

EURESPAL 80 mg 80 mg LES LAB. SERVIER IND.

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| 357 |R05CB15| ERDOSTEINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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R05CB15 ERDOSTEINUM PULB. PT. SUSP. ORALĂ 175 mg/5 ml

ERDOMED(R) 175 175 mg/5 ml CSC PHARMACEUTICALS

HANDELS GMBH

R05CB15 ERDOSTEINUM CAPS. 300 mg

ERDOMED 300 mg MEDICOM INTERNATIONAL SRO

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| 358 |R05DA09| DEXTROMETHORPHANUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R05DA09 DEXTROMETHORPHANUM SIROP 0.1%

HUMEX 0.1% LAB. URGO SA

TUSSIN SIROP 0.1% GLOBAL PHARMACEUTICALS

S.R.L.

R05DA09 DEXTROMETHORPHANUM SIROP 0.13%

HUMEX 0.13% LAB. URGO SA

R05DA09 DEXTROMETHORPHANUM COMPR. 10 mg

TUSSIN 10 mg EUROPHARM SA

R05DA09 DEXTROMETHORPHANUM SIROP 15 mg/5 ml

ROFEDEX 15 mg/5 ml BIOFARM SA

R05DA09 DEXTROMETHORPHANUM COMPR. 20 mg

TUSSIN FORTE 20 mg EUROPHARM SA

R05DA09 DEXTROMETHORPHANUM SOL. ORALĂ 3.75 mg/5 ml

ROBITUSSIN JUNIOR 3.75 mg/5 ml WYETH WHITEHALL EXPORT

GMBH

R05DA09 DEXTROMETHORPHANUM SOL. ORALĂ 7.5 mg/5 ml

ROBITUSSIN ANTITUSSICUM 7.5 mg/5 ml WYETH WHITEHALL EXPORT

GMBH

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| 359 |R06AE09| LEVOCETIRIZINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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R06AE09 LEVOCETIRIZINUM COMPR. FILM. 5 mg

XYZAL(R) 5 mg U.C.B. GMBH

R06AE09 LEVOCETIRIZINUM PIC. ORALE, SOL. 5 mg/ml

XYZAL 5 mg/ml U.C.B. GMBH

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| 360 |R06AX26| FEXOFENADINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R06AX26 FEXOFENADINUM COMPR. FILM. 120 mg

ALTIVA 120 mg 120 mg RANBAXY UK LTD.

TELFAST(R) 120 mg 120 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

R06AX26 FEXOFENADINUM COMPR. FILM. 180 mg

ALTIVA 180 mg 180 mg RANBAXY UK LTD.

TELFAST(R) 180 mg 180 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

R06AX26 FEXOFENADINUM COMPR. FILM. 30 mg

TELFAST(R) 30 mg 30 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

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| 361 |R06AX27| DESLORATADINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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R06AX27 DESLORATADINUM SIROP 0.5 mg/ml

AERIUS 0.5 mg/ml 0.5 mg/ml SP EUROPE

R06AX27 DESLORATADINUM COMPR. FILM. 5 mg

AERIUS 5 mg 5 mg SP EUROPE

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| 362 |S01AA11| GENTAMICINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01AA11 GENTAMICINUM PICĂTURI OFT. - SOL. 0.3%

OPHTAGRAM(R) 0,3% 0.3% LAB. CHAUVIN

S01AA11 GENTAMICINUM SOL. OFT. 0.3%

GENTICOL 0.3% S.I.F.I. SPA

S01AA11 GENTAMICINUM UNG. OFT. 0.3%

GENTICOL 0.3% S.I.F.I. SPA

OPHTAGRAM(R) 0,3% 0.3% LAB. CHAUVIN

S01AA11 GENTAMICINUM PIC. OFT., SOL. 0.3%

GENTAMICIN SULPHATE 0.3% E.I.P.I.CO. MED S.R.L.

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| 363 |S01AA12| TOBRAMYCINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01AA12 TOBRAMYCINUM UNG. OFT. 0.3%

TOBREX(R) 0.3% ALCON COUVREUR NV

S01AA12 TOBRAMYCINUM PIC. OFT., SOL. 0.3%

TOBISOL 0.3% E.I.P.I.CO. MED S.R.L.

S01AA12 TOBRAMYCINUM PIC. OFT., SOL. 3 mg/ml

TOBREX 3 mg/ml ALCON COUVREUR NV

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| 364 |S01AA23| NETILMICINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01AA23 NETILMICINUM PICĂTURI OFT. - SOL. 0.3%

NETTACIN(R) 0.3% S.I.F.I. SPA

S01AA23 NETILMICINUM UNG. OFT. 3 mg/g

NETTAVISC 3 mg/g 3 mg/g S.I.F.I. SPA

S01AA23 NETILMICINUM PIC. OFT., SOL. 0.3%

NETTACIN(R) 0.3% S.I.F.I. SPA

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| 365 |S01AX11| OFLOXACINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01AX11 OFLOXACINUM PICĂTURI OFT. - SOL. 0.3%

FLOXAL(R) 0.3% DR. GERHARD MANN

CHEM-PHARM. FABRIK GMBH

S01AX11 OFLOXACINUM UNG. OFT. 0.3%

FLOXAL(R) 0.3% DR. GERHARD MANN

CHEM-PHARM. FABRIK GMBH

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| 366 |S01BA01| DEXAMETHASONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01BA01 DEXAMETHASONUM PICĂTURI OFT. - SUSP. 0.1%

MAXIDEX(R) 0.1% ALCON COUVREUR NV

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| 367 |S01BA06| BETAMETHASONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01BA06 BETAMETHASONUM SOL. OFT. 0.1%

OPHTAMESONE 0.1% DAR AL DAWA PHARMA S.R.L.

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| 368 |S01CA01| COMBINAŢII (NETILMICINUM + DEXAMETHASONUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01CA01 COMBINAŢII PIC. OFT., SOL.

(NETILMICINUM +

DEXAMETHASONUM)

NETILDEX S.I.F.I. SPA

S01CA01 COMBINAŢII PIC. OFT. SOL. UNIDOZĂ

(NETILMICINUM +

DEXAMETHASONUM)

NETILDEX S.I.F.I. SPA

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| 369 |S01CA01| COMBINAŢII (TOBRAMYCINUM + DEXAMETHASONUM) | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01CA01 COMBINAŢII UNG. OFT. 1 mg + 3 mg

(TOBRAMYCINUM +

DEXAMETHASONUM)

TOBRADEX(R) 1 mg + 3 mg ALCON COUVREUR NV

S01CA01 COMBINAŢII PICĂTURI OFT. - SUSP. 1 mg + 3 mg

(TOBRAMYCINUM +

DEXAMETHASONUM)

TOBRADEX(R) 1 mg + 3 mg ALCON COUVREUR NV

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| 370 |S01CA01| COMBINAŢII (CHLORAMPHENICOLUM + | |

| | | DEXAMETHASONUM)| | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01CA01 COMBINAŢII PICĂTURI OFT. - SOL. 5 mg/ml + 1 mg/ml

(CHLORAMPHENICOLUM +

DEXAMETHASONUM)

SPERSADEX COMP 5 mg/ml + 1 mg/ml NOVARTIS PHARMA GMBH

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| 371 |S01GX09| OLOPATADINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01GX09 OLOPATADINUM PIC. OFT., SOL. 1 mg/ml

OPATANOL 1 mg/ml 1 mg/ml ALCON LABORATORIES LTD.

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| 372 |S01XA02| RETINOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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S01XA02 RETINOLUM GEL OFT. 10 mg/g

OCULOTECT 10 mg/g NOVARTIS PHARMA GMBH

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| 1219 |H03BB01| CARBIMAZOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H03BB01 CARBIMAZOLUM COMPR. FILM. 5 mg

CARBIMAZOLE 5 5 mg REMEDICA LTD.

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SUBLISTA C1 - G1 INSUFICIENŢA CARDIACĂ CRONICĂ (CLASA III SAU IV NYHA).

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| 373 |B01AA07| ACENOCUMAROLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AA07 ACENOCUMAROLUM COMPR. 2 mg

TROMBOSTOP 2 mg 2 mg TERAPIA SA

B01AA07 ACENOCUMAROLUM COMPR. 4 mg

SINTROM(R) 4 mg NOVARTIS PHARMA GMBH

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| 374 |C01AA05| DIGOXINUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C01AA05 DIGOXINUM SOL. ORALĂ 0.05 mg/ml

LANOXIN SOLUŢIE ORALĂ 0.05 mg/ml THE WELLCOME FOUNDATION

LTD.

C01AA05 DIGOXINUM COMPR. 0.25 mg

DIGOXIN 0.25 mg 0.25 mg ZENTIVA S.A.

C01AA05 DIGOXINUM SOL. INJ. 0.5 mg/ml

DIGOXIN 0.5 mg/2 ml 0.5 mg/ml ZENTIVA S.A.

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| 375 |C03AA03| HYDROCHLOROTHIAZIDUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03AA03 HYDROCHLOROTHIAZIDUM COMPR. 25 mg

NEFRIX 25 mg 25 mg ZENTIVA SA

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| 376 |C03CA01| FUROSEMIDUM | |

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Electroliţii serici trebuie să fie verificaţi periodic.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03CA01 FUROSEMIDUM SOL. INJ. 10 mg/ml

FUROSEMID 20 mg/2 ml 10 mg/ml ZENTIVA SA

C03CA01 FUROSEMIDUM COMPR. 40 mg

FUROSEMID ARENA 40 mg 40 mg ARENA GROUP SA

FUROSEMID EEL 40 mg BIO EEL SRL

FUROSEMID LPH 40 mg 40 mg LABORMED PHARMA SA

FUROSEMID MCC 40 mg 40 mg MAGISTRA C & C SRL

FUROSEMID SLAVIA 40 mg SLAVIA PHARM SRL

FUROSEMID ZENTIVA 40 mg ZENTIVA SA

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| 377 |C03DA01| SPIRONOLACTONUM | |

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Electroliţii serici trebuie să fie verificaţi periodic.

Femeile la vârsta fertilă la care s-a iniţiat tratament cu spironolactonă trebuie să ia măsuri adecvate de contracepţie.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C03DA01 SPIRONOLACTONUM CAPS. 100 mg

VEROSPIRON 100 mg GEDEON RICHTER LTD.

C03DA01 SPIRONOLACTONUM COMPR. 25 mg

SPIRONOLACTONA 25 mg 25 mg BIO EEL SRL

C03DA01 SPIRONOLACTONUM COMPR. FILM. 25 mg

ALSPIRON 25 mg 25 mg AC HELCOR PHARMA SRL

SPIRONOLACTONA 25 mg 25 mg TERAPIA SA

C03DA01 SPIRONOLACTONUM CAPS. 50 mg

VEROSPIRON 50 mg GEDEON RICHTER LTD.

C03DA01 SPIRONOLACTONUM COMPR. FILM. 50 mg

ALSPIRON 50 mg 50 mg AC HELCOR PHARMA SRL

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| 378 |C07AB02| METOPROLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB02 METOPROLOLUM COMPR. 100 mg

BETAPROL 100 mg 100 mg AC HELCOR PHARMA SRL

BLOXAN 100 mg KRKA D.D. NOVO MESTO

EGILOK 100 mg 100 mg EGIS PHARMACEUTICALS

P.L.C.

METOPRO TAD 100 100 mg TAD PHARMA GMBH

METOPROLOL 100 mg 100 mg OZONE LABORATORIES LTD.

METOPROLOL AL 100 100 mg ALIUD(R) PHARMA GMBH &

CO.KG

METOPROLOL LPH 100 mg 100 mg LABORMED PHARMA SA

METOPROLOL MEDICO UNO 100 mg 100 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

METOPROLOL TERAPIA 100 mg 100 mg TERAPIA SA

VASOCARDIN(R) 100 100 mg SLOVAKOFARMA

C07AB02 METOPROLOLUM COMPR. ELIB. PREL. 100 mg

METOPROLOL RETARD 100 mg 100 mg TERAPIA S.A.

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. 100 mg

PREL.

BETALOC(R) ZOC 100 mg 100 mg ASTRAZENECA AB

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. 190 mg

MOD.

METOSUCCINAT SANDOZ 190 mg 190 mg HEXAL AG

C07AB02 METOPROLOLUM SOL. INJ. 1 mg/ml

BETALOC 1 mg/ml ASTRAZENECA AB

C07AB02 METOPROLOLUM COMPR. ELIB. PREL. 200 mg

VASOCARDIN(R) SR 200 200 mg ZENTIVA AS

C07AB02 METOPROLOLUM COMPR. 25 mg

EGILOK 25 mg 25 mg EGIS PHARMACEUTICALS

P.L.C.

METOPROLOL 25 mg 25 mg ARENA GROUP S.A.

METOPROLOL LPH 25 mg 25 mg LABORMED PHARMA SA

METOPROLOL MEDICO UNO 25 mg 25 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. 47.5 mg

MOD.

METOSUCCINAT SANDOZ 47,5 mg 47.5 mg HEXAL AG

C07AB02 METOPROLOLUM COMPR. 50 mg

BETAPROL 50 mg 50 mg AC HELCOR PHARMA SRL

EGILOK 50 mg 50 mg EGIS PHARMACEUTICALS

P.L.C.

METOPRO TAD 50 50 mg TAD PHARMA GMBH

METOPROLOL 50 mg 50 mg OZONE LABORATORIES LTD.

METOPROLOL AL 50 50 mg ALIUD PHARMA GMBH & CO.KG

METOPROLOL LPH 50 mg 50 mg LABORMED PHARMA SA

METOPROLOL MEDICO UNO 50 mg 50 mg MEDICO UNO PHARMACEUTICAL

S.R.L.

METOPROLOL TERAPIA 50 mg 50 mg TERAPIA SA

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. 50 mg

PREL.

BETALOC(R) ZOC 50 mg 50 mg ASTRAZENECA AB

C07AB02 METOPROLOLUM COMPR. FILM. ELIB. MOD. 95 mg

METOSUCCINAT SANDOZ 95 mg 95 mg HEXAL AG

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| 379 |C07AB07| BISOPROLOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB07 BISOPROLOLUM COMPR. FILM. 2.5 mg

CONCOR COR 2,5 mg 2.5 mg MERCK KGAA

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| 380 |C07AB12| NEBIVOLOLUM\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AB12 NEBIVOLOLUM COMPR. 5 mg

NEBILET(R) 5 mg BERLIN CHEMIE AG MENARINI

GROUP

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| 381 |C07AG02| CARVEDILOLUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C07AG02 CARVEDILOLUM COMPR. 12,5 mg

CARVEDILOL SANDOZ 12,5 mg HEXAL AG

C07AG02 CARVEDILOLUM COMPR. 12.5 mg

ATRAM 12,5 12.5 mg ZENTIVA AS

CARVEDILOL 12,5 mg 12.5 mg VIM SPECTRUM SRL

CARVEDILOL HELCOR 12,5 mg 12.5 mg AC HELCOR PHARMA SRL

CARVEDILOL LPH 12,5 mg 12.5 mg LABORMED PHARMA SA

CARVEDILOL TEVA 12,5 mg 12.5 mg TEVA PHARMACEUTICAL S.R.L.

CORYOL(R) 12.5 mg KRKA D.D.

DILATREND(R) 12,5 mg 12.5 mg ROCHE ROMANIA S.R.L.

TALLITON(R) 12,5 mg 12.5 mg EGIS PHARMACEUTICALS LTD.

C07AG02 CARVEDILOLUM COMPR. FILM. 12.5 mg

CARVEDIGAMMA 12,5 mg 12.5 mg WORWAG PHARMA GMBH & CO.KG

C07AG02 CARVEDILOLUM COMPR. 25 mg

ATRAM 25 25 mg ZENTIVA AS

CARVEDILOL HELCOR 25 mg 25 mg AC HELCOR PHARMA SRL

CARVEDILOL LPH 25 mg 25 mg LABORMED PHARMA SA

CARVEDILOL SANDOZ 25 mg HEXAL AG

CARVEDILOL TEVA 25 mg 25 mg TEVA PHARMACEUTICAL S.R.L.

CORYOL(R) 25 mg KRKA D.D.

DILATREND(R) 25 mg 25 mg ROCHE ROMANIA S.R.L.

TALLITON(R) 25 mg 25 mg EGIS PHARMACEUTICALS LTD.

C07AG02 CARVEDILOLUM COMPR. FILM. 25 mg

CARVEDIGAMMA 25 mg 25 mg WORWAG PHARMA GMBH & CO.KG

C07AG02 CARVEDILOLUM COMPR. 3.125 mg

CORYOL(R) 3,125 mg 3.125 mg KRKA D.D.

C07AG02 CARVEDILOLUM COMPR. 6.25 mg

ATRAM 6,25 6.25 mg ZENTIVA AS

CARVEDILOL 6,25 mg 6.25 mg VIM SPECTRUM SRL

CARVEDILOL LPH 6,25 mg 6.25 mg LABORMED PHARMA SA

CARVEDILOL SANDOZ 6.25 mg HEXAL AG

CARVEDILOL TEVA 6,25 mg 6.25 mg TEVA PHARMACEUTICAL S.R.L.

CORYOL(R) 6.25 mg KRKA D.D.

DILATREND(R) 6,25 mg 6.25 mg ROCHE ROMANIA S.R.L.

TALLITON 6,25 mg 6.25 mg EGIS PHARMACEUTICALS LTD.

C07AG02 CARVEDILOLUM COMPR. FILM. 6.25 mg

CARVEDIGAMMA 6,25 mg 6.25 mg WORWAG PHARMA GMBH & CO.KG

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| 382 |C09AA01| CAPTOPRILUM | |

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA01 CAPTOPRILUM COMPR. 12,5 mg

CAPTOPRIL MCC 12,5 mg 12,5 mg MAGISTRA C & C

C09AA01 CAPTOPRILUM COMPR. 12.5 mg

CAPTOPRIL 12,5 mg 12.5 mg EGIS PHARMACEUTICALS LTD.

C09AA01 CAPTOPRILUM COMPR. 25 mg

CAPTOPRIL - AC 25 mg 25 mg AC HELCOR PHARMA SRL

CAPTOPRIL SINTOFARM 25 mg 25 mg SINTOFARM SA

CAPTOPRIL 25 EEL 25 mg BIO EEL SRL

CAPTOPRIL 25 mg 25 mg EGIS PHARMACEUTICALS LTD.

CAPTOPRIL LPH 25 mg 25 mg LABORMED PHARMA SA

CAPTOPRIL MCC 25 mg 25 mg MAGISTRA C & C

C09AA01 CAPTOPRILUM COMPR. 50 mg

CAPTOPRIL 50 mg 50 mg ARENA GROUP SA

C09AA01 CAPTOPRILUM COMPR. 50 mg

CAPTOPRIL - AC 50 mg 50 mg AC HELCOR PHARMA SRL

CAPTOPRIL 50 EEL 50 mg BIO EEL SRL

CAPTOPRIL 50 mg 50 mg EGIS PHARMACEUTICALS LTD.

CAPTOPRIL LPH 50 mg 50 mg LABORMED PHARMA SA

CAPTOPRIL MCC 50 mg 50 mg MAGISTRA C & C

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| 383 |C09AA02| ENALAPRILUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA02 ENALAPRILUM SOL. INJ. 1.25 mg/ml

ENAP(R) 1.25 mg/ml KRKA D.D.

C09AA02 ENALAPRILUM COMPR. 10 mg

EDNYT(R) 10 mg 10 mg GEDEON RICHTER LTD.

ENAHEXAL(R) 10 mg 10 mg HEXAL AG

ENALA TAD 10 10 mg TAD PHARMA GMBH

ENALAP 10 mg E.I.P.I.CO. MED S.R.L.

ENALAPRIL 10 mg 10 mg MAGISTRA C & C

ENALAPRIL AL 10 10 mg ALIUD(R) PHARMA GMBH &

CO.KG

ENALAPRIL FABIOL 10 mg 10 mg FABIOL SA

ENALAPRIL LPH 10 mg 10 mg LABORMED PHARMA SA

ENALAPRIL SANDOZ 10 mg 10 mg SANDOZ SRL

ENALAPRIL TERAPIA 10 mg 10 mg TERAPIA SA

ENAM 10 mg 10 mg REPREZENTANTA DR. REDDY'S

LABORATORIES LTD.

ENAP 10 mg 10 mg KRKA D.D. NOVO MESTO

RENITEC 10 mg 10 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C09AA02 ENALAPRILUM COMPR. 2.5 mg

EDNYT(R) 2,5 mg 2.5 mg GEDEON RICHTER LTD.

C09AA02 ENALAPRILUM COMPR. 20 mg

EDNYT(R) 20 mg 20 mg GEDEON RICHTER LTD.

ENAHEXAL(R) 20 mg 20 mg HEXAL AG

ENALA TAD 20 20 mg TAD PHARMA GMBH

ENALAPRIL 20 mg OZONE LABORATORIES LTD.

ENALAPRIL 20 mg 20 mg MAGISTRA C & C

ENALAPRIL AL 20 20 mg ALIUD(R) PHARMA GMBH &

CO.KG

ENALAPRIL FABIOL 20 mg 20 mg FABIOL SA

ENALAPRIL LPH 20 mg 20 mg LABORMED PHARMA SA

ENALAPRIL SANDOZ 20 mg 20 mg SANDOZ SRL

ENALAPRIL TERAPIA 20 mg 20 mg TERAPIA SA

ENAP 20 mg 20 mg KRKA D.D. NOVO MESTO

RENITEC 20 mg 20 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

C09AA02 ENALAPRILUM COMPR. 5 mg

EDNYT(R) 5 mg 5 mg GEDEON RICHTER LTD.

ENAHEXAL(R) 5 mg 5 mg HEXAL AG

ENALA TAD 5 5 mg TAD PHARMA GMBH

ENALAPRIL 5 mg 5 mg OZONE LABORATORIES LTD.

ENALAPRIL AL 5 5 mg ALIUD(R) PHARMA

GMBH & CO.KG

ENALAPRIL LPH(R) 5 mg 5 mg LABORMED PHARMA SA

ENALAPRIL SANDOZ 5 mg 5 mg SANDOZ SRL

ENAP 5 mg 5 mg KRKA D.D. NOVO MESTO

RENITEC 5 mg 5 mg MERCK SHARP & DOHME

ROMANIA S.R.L.

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| 384 |C09AA03| LISINOPRILUM | |

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA03 LISINOPRILUM COMPR. 10 mg

LISIGAMMA 10 mg 10 mg WORWAG PHARMA GMBH & CO.KG

LISINOPRIL 10 MEDO 10 mg MEDOCHEMIE ROMANIA SRL

LISINOPRIL ANTIBIOTICE 10 mg 10 mg ANTIBIOTICE SA

LISINOPRIL SANDOZ 10 mg 10 mg HEXAL AG

LISIREN 10 mg 10 mg AC HELCOR SRL

MEDAPRIL 10 10 mg MEDOCHEMIE LTD.

RANOLIP 10 mg RANBAXY U.K. LIMITED

SINOPRYL(R) 10 10 mg ECZACIBASI PHARMACEUTICALS

S.R.L.

TONOLYSIN 10 mg 10 mg GEDEON RICHTER LTD.

C09AA03 LISINOPRILUM COMPR. 2.5 mg

TONOLYSIN 2,5 mg 2.5 mg GEDEON RICHTER LTD.

C09AA03 LISINOPRILUM COMPR. 20 mg

LISIGAMMA 20 mg 20 mg WORWAG PHARMA GMBH & CO.KG

LISINOPRIL 20 MEDO 20 mg MEDOCHEMIE ROMANIA SRL

LISINOPRIL ANTIBIOTICE 20 mg 20 mg ANTIBIOTICE SA

LISINOPRIL SANDOZ 20 mg 20 mg HEXAL AG

LISIREN 20 mg 20 mg AC HELCOR SRL

MEDAPRIL 20 20 mg MEDOCHEMIE LTD.

RANOLIP 20 mg RANBAXY U.K. LIMITED

TONOLYSIN 20 mg 20 mg GEDEON RICHTER LTD.

C09AA03 LISINOPRILUM COMPR. 40 mg

LISINOPRIL ANTIBIOTICE 40 mg 40 mg ANTIBIOTICE SA

C09AA03 LISINOPRILUM COMPR. 5 mg

LISIGAMMA 5 mg 5 mg WORWAG PHARMA GMBH & CO.KG

LISINOPRIL 5 MEDO 5 mg MEDOCHEMIE ROMANIA SRL

LISINOPRIL SANDOZ 5 mg 5 mg HEXAL AG

MEDAPRIL 5 5 mg MEDOCHEMIE LTD.

RANOLIP 5 mg RANBAXY U.K. LIMITED

TONOLYSIN 5 mg 5 mg GEDEON RICHTER LTD.

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| 385 |C09AA05| RAMIPRILUM | |

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Utilizarea inhibitorilor enzimelor de conversie a angiotensinei în timpul sarcinii este contraindicată deoarece medicamentele din această categorie au fost asociate cu moartea fătului in utero.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09AA05 RAMIPRILUM COMPR. FILM. 1.25 mg

RAMIRAN 1,25 mg 1.25 mg RANBAXY U.K. LIMITED

C09AA05 RAMIPRILUM COMPR. 10 mg

EMREN 10 mg 10 mg GEDEON RICHTER ROMANIA

S.A.

VIVACE 10 mg 10 mg ACTAVIS GROUP HF

C09AA05 RAMIPRILUM COMPR. 10 mg

AMPRIL 10 mg 10 mg KRKA D.D. NOVO MESTO

PIRAMIL 10 mg 10 mg SANDOZ SRL

RAMIGAMMA 10 mg 10 mg WORWAG PHARMA GMBH & CO.KG

RAMIPRIL-AC 10 mg 10 mg AC HELCOR PHARMA SRL

TRITACE 10 10 mg SANOFI-AVENTIS

DEUTSCHLAND GMBH

ZENRA 10 10 mg ZENTIVA S.A.

C09AA05 RAMIPRILUM COMPR. FILM. 10 mg

RAMIRAN 10 mg 10 mg RANBAXY U.K. LIMITED

C09AA05 RAMIPRILUM COMPR. 2,5 mg

EMREN 2,5 mg 2,5 mg GEDEON RICHTER ROMANIA SA

VIVACE 2,5 mg 2,5 mg ACTAVIS GROUP HF

C09AA05 RAMIPRILUM COMPR. 2.5 mg

AMPRIL 2,5 mg 2.5 mg KRKA D.D. NOVO MESTO

PIRAMIL 2,5 mg 2.5 mg SANDOZ SRL

RAMIPRIL-AC 2,5 mg 2.5 mg AC HELCOR PHARMA SRL

TRITACE(R) 2,5 2.5 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

ZENRA 2,5 2.5 mg ZENTIVA S.A.

C09AA05 RAMIPRILUM COMPR. FILM. 2.5 mg

RAMIRAN 2,5 mg 2.5 mg RANBAXY U.K. LIMITED

C09AA05 RAMIPRILUM COMPR. 5 mg

EMREN 5 mg 5 mg GEDEON RICHTER ROMANIA

S.A.

VIVACE 5 mg 5 mg ACTAVIS GROUP HF

C09AA05 RAMIPRILUM COMPR. 5 mg

AMPRIL 5 mg 5 mg KRKA D.D. NOVO MESTO

PIRAMIL 5 mg 5 mg SANDOZ SRL

RAMIGAMMA 5 mg 5 mg WORWAG PHARMA GMBH & CO.KG

RAMIPRIL-AC 5 mg 5 mg AC HELCOR PHARMA SRL

TRITACE(R) 5 5 mg AVENTIS PHARMA DEUTSCHLAND

GMBH

ZENRA 5 5 mg ZENTIVA S.A.

C09AA05 RAMIPRILUM COMPR. FILM. 5 mg

RAMIRAN 5 mg 5 mg RANBAXY U.K. LIMITED

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| 386 |C09CA03| VALSARTANUM\*\* | Protocol: C005I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09CA03 VALSARTANUM COMPR. FILM. 160 mg

DIOVAN 160 mg 160 mg NOVARTIS PHARMA GMBH

C09CA03 VALSARTANUM COMPR. FILM. 80 mg

DIOVAN 80 mg 80 mg NOVARTIS PHARMA GMBH

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| 387 |C09CA06| CANDESARTANUM CILEXETIL\*\* | Protocol: C005I |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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C09CA06 CANDESARTANUM COMPR. 16 mg

CILEXETIL

ATACAND 16 mg ASTRAZENECA AB

C09CA06 CANDESARTANUM COMPR. 8 mg

CILEXETIL

ATACAND 8 mg ASTRAZENECA AB

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SUBLISTA C1 - G2 BOLNAVI CU PROTEZE VALVULARE ŞI VASCULARE.

Protocol: BB01I

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| 388 |B01AA07| ACENOCUMAROLUM\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AA07 ACENOCUMAROLUM COMPR. 2 mg

TROMBOSTOP 2 mg 2 mg TERAPIA SA

B01AA07 ACENOCUMAROLUM COMPR. 4 mg

SINTROM(R) 4 mg NOVARTIS PHARMA GMBH

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SUBLISTA C1 - G3 BOLNAVI CU PROCEDURI INTERVENŢIONALE PERCUTANE, NUMAI DUPĂ IMPLANTAREA UNEI PROTEZE ENDOVASCULARE (STENT).

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| 389 |B01AC04| CLOPIDOGRELUM\*\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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B01AC04 CLOPIDOGRELUM COMPR. FILM. 75 mg

PLAVIX 75 mg 75 mg SANOFI PHARMA - BRISTOL

MYERS SQUIBB

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SUBLISTA C1 - G4 HEPATITELE CRONICE DE ETIOLOGIE VIRALĂ B, C ŞI D.

Protocol: LB01B; LB02E

**#M4**

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*| 390 | \*\*\* Abrogată |*

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*| 391 | \*\*\* Abrogată |*

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| 392 |J05AB04| RIBAVIRINUM\*\*\*\* | Protocol: J002N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J05AB04 RIBAVIRINUM CAPS. 200 mg

REBETOL 200 mg 200 mg SP EUROPE

J05AB04 RIBAVIRINUM COMPR. FILM. 200 mg

COPEGUS(R) 200 mg ROCHE ROMANIA SRL

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| 393 |J05AF05| LAMIVUDINUM\*\*\*\* | Protocol: J005N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J05AF05 LAMIVUDINUM COMPR. FILM. 100 mg

ZEFFIX 100 mg 100 mg GLAXO GROUP LTD.

J05AF05 LAMIVUDINUM SOL. ORALĂ 10 mg/ml

EPIVIR 10 mg/ml 10 mg/ml GLAXO GROUP LTD.

J05AF05 LAMIVUDINUM COMPR. FILM. 150 mg

EPIVIR 150 mg 150 mg GLAXO GROUP LTD.

J05AF05 LAMIVUDINUM SOL. ORALĂ 5 mg/ml

ZEFFIX 5 mg/ml 5 mg/ml GLAXO GROUP LTD.

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| 395 |J05AF10| ENTECAVIRUM\*\*\*\* | Protocol: J008N |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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J05AF10 ENTECAVIRUM COMPR. FILM. 0,5 mg

BARACLUDE 0,5 mg 0,5 mg BRISTOL-MYERS SQUIBB

PHARMA EEIG

J05AF10 ENTECAVIRUM COMPR. FILM. 1 mg

BARACLUDE 1 mg 1 mg BRISTOL-MYERS SQUIBB

PHARMA EEIG

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| 396 |L03AA02| FILGRASTIMUM (G-CSF)\*\* | Protocol: B013K |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L03AA02 FILGRASTIMUM (G-CSF) SOL. INJ. 30 MU/0.5 ml

NEUPOGEN(R) 30 MU/0.5 ml AMGEN EUROPE B.V.

L03AA02 FILGRASTIMUM (G-CSF) SOL. INJ. 48 MU/0.5 ml

NEUPOGEN(R) 48 MU/0.5 ml AMGEN EUROPE B.V.

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| 397 |L03AB04| INTERFERONUM ALFA 2a\*\*\*\* | Protocol: J007N |

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Tratamentul cu interferon alfa a fost asociat cu depresie şi suicid la unii pacienţi. Pacienţii cu istoric de ideaţie suicidară sau boală depresivă trebuie avertizaţi de riscuri. Trebuie monitorizat statusul psihiatric în timpul tratamentului.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 18 Mui/0.6 ml

ROFERON A 18 Mui/0.6 ml ROCHE ROMANIA S.R.L.

L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 3 Mui/0.5 ml

ROFERON A 3 Mui/0.5 ml ROCHE ROMANIA S.R.L.

L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 4.5 Mui/0.5 ml

ROFERON A 4.5 Mui/0.5 ml ROCHE ROMANIA S.R.L.

L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 9 Mui/0.5 ml

ROFERON A 9 Mui/0.5 ml ROCHE ROMANIA S.R.L.

L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 3 Mui/0.5 ml

ROFERON A 3 Mui/0.5 ml ROCHE ROMANIA S.R.L.

L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 4.5 Mui/0.5 ml

ROFERON A 4.5 Mui/0.5 ml ROCHE ROMANIA S.R.L.

L03AB04 INTERFERONUM ALFA 2a SOL. INJ. 9 Mui/0.5 ml

ROFERON A 9 Mui/0.5 ml ROCHE ROMANIA S.R.L.

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| 398 |L03AB05| INTERFERONUM ALFA 2b\*\*\*\* | Protocol: J006N |

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Tratamentul cu interferon alfa a fost asociat cu depresie şi suicid la unii pacienţi. Pacienţii cu istoric de ideaţie suicidară sau boală depresivă trebuie avertizaţi de riscuri. Trebuie monitorizat statusul psihiatric în timpul tratamentului.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L03AB05 INTERFERONUM ALFA 2b SOL. INJ. PEN MULTIDOZĂ 18 milioane U.I.

INTRON A 18 milioane U.I. 18 milioane U.I. SP EUROPE

L03AB05 INTERFERONUM ALFA 2b SOL. INJ. PEN MULTIDOZĂ 30 milioane U.I.

INTRON A 30 milioane U.I. 30 milioane U.I. SP EUROPE

INTRON A 60 milioane U.I. 60 milioane U.I. SP EUROPE

L03AB05 INTERFERONUM ALFA 2b SOL. INJ. PEN MULTIDOZĂ 60 milioane U.I.

INTRON A 30 milioane U.I. 30 milioane U.I. SP EUROPE

INTRON A 60 milioane U.I. 60 milioane U.I. SP EUROPE

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| 399 |L03AB10| PEGINTERFERON alfa-2b\*\*\*\* | Protocol: J003N |

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Tratamentul cu interferon alfa a fost asociat cu depresie şi suicid la unii pacienţi. Pacienţii cu istoric de ideaţie suicidară sau boală depresivă trebuie avertizaţi de riscuri. Trebuie monitorizat statusul psihiatric în timpul tratamentului.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L03AB10 PEGINTERFERON alfa-2b PULB. + SOLV. PT. SOL. 100 micrograme

INJ.

PEGINTRON 100 micrograme 100 micrograme SP EUROPE

L03AB10 PEGINTERFERON alfa-2b PULB. + SOLV. PT. SOL. 120 micrograme

INJ.

PEGINTRON 120 micrograme 120 micrograme SP EUROPE

L03AB10 PEGINTERFERON alfa-2b PULB. + SOLV. PT. SOL. 150 micrograme

INJ.

PEGINTRON 150 micrograme 150 micrograme SP EUROPE

L03AB10 PEGINTERFERON alfa-2b PULB. + SOLV. PT. SOL. 50 micrograme

INJ.

PEGINTRON 50 micrograme 50 micrograme SP EUROPE

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| 400 |L03AB11| PEGINTERFERON alfa-2a\*\*\*\* | Protocol: J004N |

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Tratamentul cu interferon alfa a fost asociat cu depresie şi suicid la unii pacienţi. Pacienţii cu istoric de ideaţie suicidară sau boală depresivă trebuie avertizaţi de riscuri. Trebuie monitorizat statusul psihiatric în timpul tratamentului.

Se aplică tuturor denumirilor comerciale, formelor farmaceutice şi concentraţiilor corespunzătoare DCI-ului.

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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L03AB11 PEGINTERFERON alfa-2a SOL. INJ. 135 micrograme/ml

PEGASYS 135 micrograme/ml 135 micrograme/ml ROCHE REGISTRATION LTD.

L03AB11 PEGINTERFERON alfa-2a SOL. INJ. ÎN SERINGĂ 180 micrograme/

PREUMPLUTĂ 0.5 ml

PEGASYS 180 micrograme/0,5 ml 180 micrograme/0.5 ml ROCHE REGISTRATION LTD.

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| 401 |N04BB01| AMANTADINUM\*\* | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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N04BB01 AMANTADINUM CAPS. 100 mg

VIREGYT(R)-K 100 mg EGIS PHARMACEUTICALS LTD.

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SUBLISTA C1 - G5 HEPATITA AUTOIMUNĂ.

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| 402 |H02AB04| METHYLPREDNISOLONUM | |

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DCI/DENUMIRE COMERCIALĂ FORMA FARM.

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CONCENTRAŢIE FIRMA

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H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 125 mg

LEMOD SOLU 125 mg 125 mg HEMOFARM S.R.L

H02AB04 METHYLPREDNISOLONUM LIOF. ŞI SOLV. PT. 125 mg/2 ml

SOL. INJ.

SOLU - MEDROL ACT-O-VIAL 125 mg/2 ml PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM COMPR. 16 mg

MEDROL A 16 16 mg PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 20 mg

LEMOD SOLU 20 mg 20 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. ŞI SOLV. PT. 250 mg/4 ml

SOL. INJ.

SOLU - MEDROL ACT-O-VIAL 250 mg/4 ml PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM COMPR. 32 mg

MEDROL 32 32 mg PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 40 mg

LEMOD SOLU 40 mg 40 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. ŞI SOLV. PT. 40 mg/1 ml

SOL. INJ.

SOLU - MEDROL ACT-O-VIAL 40 mg/1 ml PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM COMPR. 4 mg

MEDROL 4 mg 4 mg PFIZER EUROPE MA EEIG

H02AB04 METHYLPREDNISOLONUM LIOF. PT. SOL. INJ. 500 mg

LEMOD SOLU 500 mg 500 mg HEMOFARM S.R.L.

H02AB04 METHYLPREDNISOLONUM LIOF. + SOLV. PT. SOL. 500 mg/7.8 ml

INJ.

SOLU - MEDROL 500 mg/7,8 ml 500 mg/7.8 ml PFIZER EUROPE MA EEIG

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| 403 |H02AB06| PREDNISOLONUM