

MRF 15	MRF 15
Registration number:	Registration number:
41/2.5/0711	41/2.5/0712
Name of medicine:	Name of medicine:
EPITOZ 25	EPITOZ 50
Dosage form:	Dosage form:
TABLET	TABLET
Active ingredients:	Active ingredients:
EACH TABLET CONTAINS: TOPIRAMATE 25,0 mg	EACH TABLET CONTAINS: TOPIRAMATE 50,0 mg
Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7, 8	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	Applicant:
ZYDUS HEALTHCARE S.A. (PTY) LTD	ZYDUS HEALTHCARE S.A. (PTY) LTD
Manufacturer:	Manufacturer:
ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA	ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA
Packer:	Packer:
ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA	ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA
Laboratory:	Laboratory:
FPRC: ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA	FPRC: ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
FPRC:	FPRC:
ZYDUS HEALTHCARE S.A., VAN DER HOFF PARK, POTCHEFSTROOM	ZYDUS HEALTHCARE S.A., VAN DER HOFF PARK, POTCHEFSTROOM
Shelf-life:	Shelf-life:
24 months (Provisional)	24 months (Provisional)
Date of registration:	Date of registration:
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/2.5/0713	Registration number: 41/20.2.3/0723
Name of medicine: EPI TOZ 100	Name of medicine: RITIB
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: TOPIRAMATE 100,0 mg	Active ingredients: EACH TABLET CONTAINS: RIFAMPICIN 150,0 mg ISONIAZID 75,0 mg ETHAMBUTOL HYDROCHLORIDE 275,0 mg PYRAZINAMIDE 400,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: ZYDUS HEALTHCARE S.A. (PTY) LTD	Applicant: SVIZERA SA (PTY) LTD
Manufacturer: ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA	Manufacturer: SVIZERA LABS PVT LTD, TURBHE, NEW MUMBAI, INDIA
Packer: ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA	Packer: SVIZERA LABS PVT LTD, TURBHE, NEW MUMBAI, INDIA
Laboratory: ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM	Laboratory: SVIZERA LABS PVT LTD, TURBHE, NEW MUMBAI, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRC:	FPRC:
FPRR: ZYDUS HEALTHCARE S.A., VAN DER HOFF PARK, POTCHEFSTROOM	FPRR: SVIZERA SA, LENASIA, JOHANNESBURG
Shelf-life: 24 months (Provisional)	Shelf-life: 36 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15

Registration number: 41/2.5/0724
 Name of medicine: NEUSEIZE 100
 Dosage form: CAPSULE
 Active ingredients: EACH CAPSULE CONTAINS:
 GABAPENTIN 100,0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: RANBAXY (S.A.) (PTY) LTD
 Manufacturer: RANBAXY LABORATORIES LTD, PAONTA SAHIB,
 HIMACHAL PRADESH, INDIA
 Packer: RANBAXY LABORATORIES LTD, PAONTA SAHIB,
 HIMACHAL PRADESH, INDIA
 Laboratory: FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB,
 HIMACHAL PRADESH, INDIA
 CONSULTING CHEMICAL LABORATORIES,
 ATLASVILLE, BOKSBURG
 Shelf-life: 36 months
 Date of registration: 18 APRIL 2008
 FPRC: RANBAXY S.A., CENTURION, RSA

MRF 15

Registration number: 41/2.5/0725
 Name of medicine: NEUSEIZE 300
 Dosage form: CAPSULE
 Active ingredients: EACH CAPSULE CONTAINS:
 GABAPENTIN 300,0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: RANBAXY (S.A.) (PTY) LTD
 Manufacturer: RANBAXY LABORATORIES LTD, PAONTA
 SAHIB, HIMACHAL PRADESH, INDIA
 Packer: RANBAXY LABORATORIES LTD, PAONTA
 SAHIB, HIMACHAL PRADESH, INDIA
 Laboratory: FPRC: RANBAXY LABORATORIES LTD, PAONTA
 SAHIB, HIMACHAL PRADESH, INDIA
 CONSULTING CHEMICAL LABORATORIES,
 ATLASVILLE, BOKSBURG
 Shelf-life: 36 months
 Date of registration: 18 APRIL 2008
 FPRC: RANBAXY S.A., CENTURION, RSA

MRF 15	MRF 15
Registration number:	41/20.1.1/0751
Name of medicine:	ZINOXIME 125 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 125,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
FPRR:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG
Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	41/2.5/0726
Name of medicine:	NEUSEIZE 400
Dosage form:	CAPSULE
Active ingredients:	EACH CAPSULE CONTAINS: GABAPENTIN 400,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	RANBAXY (S.A.) (PTY) LTD
Manufacturer:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Packer:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Laboratory:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR:	RANBAXY S.A., CENTURION, RSA
Shelf-life:	36 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/20.1.1/0752	Registration number: 41/20.1.1/0753
Name of medicine: ZINOXIME 250 mg	Name of medicine: ZINOXIME 500 mg
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 250,0 mg	Active ingredients: EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 500,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: AUROBINDO PHARMA (PTY) LTD	Applicant: AUROBINDO PHARMA (PTY) LTD
Manufacturer: AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer: AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Packer: AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Packer: AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Laboratory: FPRC: AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Laboratory: FPRC: AUROBINDO PHARMA LTD, UNIT VI, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
FPRC: AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG	FPRC: AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG
Shelf-life: 24 months (Provisional)	Shelf-life: 24 months (Provisional)
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/7. 1/0754
Name of medicine:	AURO-AMLODIPINE TABLETS 5 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: AMLODIPINE BESILATE EQUIVALENT TO AMLODIPINE 5.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	41/7. 1/0755
Name of medicine:	AURO-AMLODIPINE TABLETS 10 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: AMLODIPINE BESILATE EQUIVALENT TO AMLODIPINE 10.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/17.3/0788	Registration number: 41/20.2.8/0806
Name of medicine: MIGRESS 50	Name of medicine: LAZIVIR TABLETS
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: SUMATRIPTAN SUCCINATE EQUIVALENT TO SUMATRIPTAN 50.0 mg	Active ingredients: EACH TABLET CONTAINS: LAMIVUDINE 150,0 mg ZIDOVUDINE 300,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: ARROW PHARMA SOUTH AFRICA (PTY) LTD	Applicant: AUROBINDO PHARMA (PTY) LTD
Manufacturer: ARROW PHARMACEUTICALS INC, MISSISSAUGA, ONTARIO, CANADA	Manufacturer: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer: ARROW PHARMACEUTICALS INC, MISSISSAUGA, ONTARIO, CANADA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA	Packer: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory: FPRC: ARROW PHARMACEUTICALS INC, MISSISSAUGA, ONTARIO, CANADA SELAMINE LTD 1/2 ARROW GENERICS LTD, DUBLIN, IRELAND M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA	Laboratory: FPRC: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life: 24 months (Provisional)	Shelf-life: 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	Registration number:
411/10.2.1/0849	4117.3/0861
Name of medicine:	Name of medicine:
SPIRIVA RESPIMAT	MIGRESS 100
Dosage form:	Dosage form:
SOLUTION	TABLET
Active ingredients:	Active ingredients:
EACH DOSE CONTAINS: TIOTROPIUM 5,0 ug	EACH TABLET CONTAINS: SUMATRIPTAN SUCCINATE EQUIVALENT TO SUMATRIPTAN 100,0 mg
Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	Applicant:
INGELHEIM PHARMACEUTICALS (PTY) LTD	ARROW PHARMA SOUTH AFRICA (PTY) LTD
Manufacturer:	Manufacturer:
BOEHRINGER INGELHEIM PHARMA GmbH & CO, INGELHEIM AM RHEIN, GERMANY	ARROW PHARMACEUTICALS INC, MISSISSAUGA, ONTARIO, CANADA
Packer:	Packer:
BOEHRINGER INGELHEIM PHARMA GmbH & CO, INGELHEIM AM RHEIN, GERMANY	ARROW PHARMACEUTICALS INC, MISSISSAUGA, ONTARIO, CANADA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA
Laboratory:	Laboratory:
FPRC:	FPRC:
BOEHRINGER INGELHEIM PHARMA GmbH & CO, INGELHEIM AM RHEIN, GERMANY WINTHROP PHARMACEUTICALS, WALTLOO, PRETORIA	ARROW PHARMACEUTICALS INC, MISSISSAUGA, ONTARIO, CANADA SELAMINE LTD t/a ARROW GENERICS LTD, DUBLIN, IRELAND M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA
FPRR:	FPRR:
INGELHEIM PHARMACEUTICALS, RANDSBURG, JOHANNESBURG	ARROW PHARMA SA, WOODMEAD, RSA
Shelf-life:	Shelf-life:
24 months	24 months (Provisional)
Date of registration:	Date of registration:
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/20.1.1/0866	Registration number: 41/20.1.1/0867
Name of medicine: SANDOZ CEFTRIAXONE 0,5 g	Name of medicine: SANDOZ CEFTRIAXONE 1,0 g
Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 0,5 g	Active ingredients: EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 1,0 g
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: SANDOZ S.A. (PTY) LTD	Applicant: SANDOZ S.A. (PTY) LTD
Manufacturer: SANDOZ GmbH, KUNDL, AUSTRIA	Manufacturer: SANDOZ GmbH, KUNDL, AUSTRIA
Packer: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK	Packer: NOVARTIS SA, SPARTAN, KEMPTON PARK
Laboratory: FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA	Laboratory: FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
Shelf-life: 36 months	Shelf-life: 36 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008
FPRC: SANDOZ S.A., SPARTAN, KEMPTON PARK	FPRC: SANDOZ S.A., SPARTAN, KEMPTON PARK

MRF 15	MRF 15
Registration number:	Registration number:
Name of medicine:	Name of medicine:
Dosage form:	Dosage form:
Active ingredients:	Active ingredients:
Conditions of registration:	Conditions of registration:
Applicant:	Applicant:
Manufacturer:	Manufacturer:
Packer:	Packer:
Laboratory:	Laboratory:
FPRC:	FPRC:
Shelf-life:	Shelf-life:
Date of registration:	Date of registration:
41/20.1.1/0868	41/20.1.1/0869
SANDOZ CEFTRIAXONE 2,0g	ROKEF 0,5 g
INJECTION	INJECTION
EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 2,0 g	EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 0,5 g
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
SANDOZ S.A. (PTY) LTD	SANDOZ S.A. (PTY) LTD
SANDOZ GmbH, KUNDL, AUSTRIA	SANDOZ GmbH, KUNDL, AUSTRIA
SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK
SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRC:	FPRC:
36 months	36 months
18 APRIL 2008	18 APRIL 2008
SANDOZ S.A., SPARTAN, KEMPTON PARK	SANDOZ S.A., SPARTAN, KEMPTON PARK

MRF 15	MRF 15
Registration number:	41/20.1.1/0870
Name of medicine:	ROKEF 1,0 g
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 1,0 g
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANDOZ S.A. (PTY) LTD
Manufacturer:	SANDOZ GmbH, KUNDL, AUSTRIA
Packer:	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK
Laboratory:	FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRC:	SANDOZ S.A., SPARTAN, KEMPTON PARK
Shelf-life:	36 months
Date of registration:	18 APRIL 2008
Registration number:	41/20.1.1/0871
Name of medicine:	ROKEF 2,0 g
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 2,0 g
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANDOZ S.A. (PTY) LTD
Manufacturer:	SANDOZ GmbH, KUNDL, AUSTRIA
Packer:	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK
Laboratory:	FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRC:	SANDOZ S.A., SPARTAN, KEMPTON PARK
Shelf-life:	36 months
Date of registration:	18 APRIL 2008

MRF 15

Registration number: 41/20.1.1/0872
 Name of medicine: ORZID 1 g
 Dosage form: INJECTION
 Active ingredients: EACH VIAL CONTAINS:
 CEFTAZIDIME PENTAHYDRATE EQUIVALENT TO
 CEFTAZIDIME 1,0 g
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: ORCHID PHARMACEUTICALS SA (PTY) LTD
 Manufacturer: ORCHID HEALTHCARE; KANCHEEPUURAM
 DISTRICT, TAMIL NADU, INDIA
 Packer: ORCHID HEALTHCARE; KANCHEEPUURAM
 DISTRICT, TAMIL NADU, INDIA
 Laboratory: FPRC: ORCHID HEALTHCARE; KANCHEEPUURAM
 DISTRICT, TAMIL NADU, INDIA
 CONSULTING CHEMICAL LABORATORIES,
 ATLASVILLE, BOKSBURG
 RESEARCH INSTITUTE FOR INDUSTRIAL
 PHARMACY, NORTH-WEST UNIVERSITY,
 POTCHEFSTROOM
 INSTITUTE FOR PHARMACEUTICAL SERVICES,
 SILVERTONDALE, PRETORIA
 FPRR: ORCHID PHARMACEUTICALS SA,
 POTCHEFSTROOM
 Shelf-life: 24 months
 Date of registration: 18 APRIL 2008

MRF 15

Registration number: 41/20.1.1/0873
 Name of medicine: ORZID 2 g
 Dosage form: INJECTION
 Active ingredients: EACH VIAL CONTAINS:
 CEFTAZIDIME PENTAHYDRATE
 EQUIVALENT TO
 CEFTAZIDIME 2,0 g
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: ORCHID PHARMACEUTICALS SA (PTY) LTD
 Manufacturer: ORCHID HEALTHCARE; KANCHEEPUURAM
 DISTRICT, TAMIL NADU, INDIA
 Packer: ORCHID HEALTHCARE; KANCHEEPUURAM
 DISTRICT, TAMIL NADU, INDIA
 Laboratory: FPRC: ORCHID HEALTHCARE; KANCHEEPUURAM
 DISTRICT, TAMIL NADU, INDIA
 CONSULTING CHEMICAL LABORATORIES,
 ATLASVILLE, BOKSBURG
 RESEARCH INSTITUTE FOR INDUSTRIAL
 PHARMACY, NORTH-WEST UNIVERSITY,
 POTCHEFSTROOM
 INSTITUTE FOR PHARMACEUTICAL
 SERVICES, SILVERTONDALE, PRETORIA
 FPRR: ORCHID PHARMACEUTICALS SA,
 POTCHEFSTROOM
 Shelf-life: 24 months
 Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	Registration number:
41/7.1.3/1050	41/7.1.3/1051
Name of medicine:	Name of medicine:
LISINOZIDE 10 mg	LISINOZIDE 20 mg
Dosage form:	Dosage form:
TABLET	TABLET
Active ingredients:	Active ingredients:
EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 10,0 mg HYDROCHLOROTHIAZIDE 12,5 mg	EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 20,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
Applicant:	Applicant:
AUROBINDO PHARMA (PTY) LTD	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	Manufacturer:
AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	Packer:
AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH INDIA	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH INDIA
Laboratory:	Laboratory:
FPRC:	FPRC:
AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
FPRR:	FPRR:
AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA
Shelf-life:	Shelf-life:
24 months	24 months
Date of registration:	Date of registration:
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15		
Registration number:	41/21.12/1054	Registration number:	41/21.12/1055
Name of medicine:	AURO-FINASTERIDE TABLETS 5 mg	Name of medicine:	PROFINA 5 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: FINASTERIDE 5.0 mg	Active ingredients:	EACH TABLET CONTAINS: FINASTERIDE 5.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD	Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Packer:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Laboratory:	FPRC: AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
FPRR:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG	FPRR:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008	Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/20.1.1/1063
Name of medicine:	SANDOZ CEFOTAXIME 0,5 g
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: CEFOTAXIME SODIUM EQUIVALENT TO CEFOTAXIME 0,5 g
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANDOZ S.A. (PTY) LTD
Manufacturer:	SANDOZ GmbH, KUNDL, AUSTRIA
Packer:	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK
Laboratory:	FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA ANALYTICON, TERENURE, KEMPTON PARK
Shelf-life:	FPRC: SANDOZ S.A., SPARTAN, KEMPTON PARK 24 months
Date of registration:	18 APRIL 2008
Registration number:	41/20.1.1/1064
Name of medicine:	SANDOZ CEFOTAXIME 1,0 g
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: CEFOTAXIME SODIUM EQUIVALENT TO CEFOTAXIME 1,0 g
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANDOZ S.A. (PTY) LTD
Manufacturer:	SANDOZ GmbH, KUNDL, AUSTRIA
Packer:	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK
Laboratory:	FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA ANALYTICON, TERENURE, KEMPTON PARK
Shelf-life:	FPRC: SANDOZ S.A., SPARTAN, KEMPTON PARK 24 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/20.1.1/1065	Registration number: 41/20.1.1/1066
Name of medicine: CLATAX 0,5 g	Name of medicine: CLATAX 1,0 g
Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH VIAL CONTAINS: CEFOTAXIME SODIUM EQUIVALENT TO CEFOTAXIME 0,5 g	Active ingredients: EACH VIAL CONTAINS: CEFOTAXIME SODIUM EQUIVALENT TO CEFOTAXIME 1,0 g
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: SANDOZ S.A. (PTY) LTD	Applicant: SANDOZ S.A. (PTY) LTD
Manufacturer: SANDOZ GmbH, KUNDL, AUSTRIA	Manufacturer: SANDOZ GmbH, KUNDL, AUSTRIA
Packer: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK	Packer: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK
Laboratory: FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA ANALYTICON, TERENURE, KEMPTON PARK	Laboratory: FPRC: SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA ANALYTICON, TERENURE, KEMPTON PARK
FPRC: SANDOZ S.A., SPARTAN, KEMPTON PARK	FPRC: SANDOZ S.A., SPARTAN, KEMPTON PARK
Shelf-life: 24 months	Shelf-life: 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	42/7.1.3/0106
Name of medicine:	PERIVAS 4
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL TERT-BUTYLAMINE 4,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	RANBAXY (S.A.) (PTY) LTD
Manufacturer:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Packer:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Laboratory:	FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR:	RANBAXY (SA), CENTURION, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	42/7.1.3/0107
Name of medicine:	RAN-PERINDOPRIL 4
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL TERT-BUTYLAMINE 4,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	RANBAXY (S.A.) (PTY) LTD
Manufacturer:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Packer:	RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Laboratory:	FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR:	RANBAXY (SA), CENTURION, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	A38/2.6.5/0622
Name of medicine:	MOXOTENS 0,2 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MOXONIDINE 0,2 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	HEXAL PHARMA (S.A.) (PTY) LTD
Manufacturer:	PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY
Packer:	PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG, RSA
Laboratory:	FPRC: PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA ANALYTICON, TERENURE, KEMPTON PARK, RSA
Shelf-life:	24 months
Date of registration:	18 APRIL 2008
Registration number:	A38/2.6.5/0623
Name of medicine:	MOXOTENS 0,3 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MOXONIDINE 0,3 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	HEXAL PHARMA (S.A.) (PTY) LTD
Manufacturer:	PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY
Packer:	PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG, RSA
Laboratory:	FPRC: PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA ANALYTICON, TERENURE, KEMPTON PARK, RSA
Shelf-life:	24 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	Registration number:
A38/2.6.5/0624	37/28/0521
Name of medicine:	Name of medicine:
MOXOTENS 0,4 mg	AXIM READICAT 2 %
Dosage form:	Dosage form:
TABLET	SUSPENSION
Active ingredients:	Active ingredients:
EACH TABLET CONTAINS: MOXONIDINE 0,4 mg	EACH 100,0 ml SUSPENSION CONTAINS: BARIUM SULPHATE 2,0940 g
Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6
Applicant:	Applicant:
HEXAL PHARMA (S.A.) (PTY) LTD	AXIM PHARMACEUTICALS (PTY) LTD
Manufacturer:	Manufacturer:
PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY	E-Z-EM CANADA INC, ANJOU, QUEBEC, CANADA
Packer:	Packer:
PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG, RSA	E-Z-EM CANADA INC, ANJOU, QUEBEC, CANADA
Laboratory:	Laboratory:
FPRC:	FPRC:
PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA ANALYTICON, TERENCE, KEMPTON PARK, RSA	E-Z-EM CANADA INC, ANJOU, QUEBEC, CANADA SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRR:	FPRR:
HEXAL PHARMA, PINETOWN, KZN, RSA	AXIM PHARMACEUTICALS, MIDRAND, JOHANNESBURG
Shelf-life:	Shelf-life:
24 months	36 months
Date of registration:	Date of registration:
18 APRIL 2008	13 JUNE 2008

MRF 15	MRF 15
Registration number:	A39/5.4/0212
Name of medicine:	LYRINEL 5 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: OXYBUTYNIN HYDROCHLORIDE 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	JANSSEN PHARMACEUTICA (PTY) LTD
Manufacturer:	ALZA CORPORATION, MOUNTAIN VIEW, CALIFORNIA, USA ALZA CORPORATION, VACAVILLE, CALIFORNIA, USA
Packer:	ALZA CORPORATION, VACAVILLE, CALIFORNIA, USA JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY
Laboratory:	ALZA IRELAND LTD, TIPPERARY, IRELAND JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY JANSSEN PHARMACEUTICA NV, BEERSE, BELGIUM
FPRC:	FPRC:
FPRR:	JANSSEN PHARMACEUTICA, WOODMEAD, JOHANNESBURG
Shelf-life:	24 months
Date of registration:	13 JUNE 2008
Registration number:	A39/5.4/0224
Name of medicine:	LYRINEL 10 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: OXYBUTYNIN HYDROCHLORIDE 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	JANSSEN PHARMACEUTICA (PTY) LTD
Manufacturer:	ALZA CORPORATION, MOUNTAIN VIEW, CALIFORNIA, USA ALZA CORPORATION, VACAVILLE, CALIFORNIA, USA13 JUNE 2008
Packer:	ALZA CORPORATION, VACAVILLE, CALIFORNIA, USA JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY
Laboratory:	ALZA IRELAND LTD, TIPPERARY, IRELAND JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY JANSSEN PHARMACEUTICA NV, BEERSE, BELGIUM
FPRC:	FPRC:
FPRR:	ALZA IRELAND LTD, TIPPERARY, IRELAND JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY JANSSEN PHARMACEUTICA NV, BEERSE, BELGIUM
Shelf-life:	18 months
Date of registration:	13 JUNE 2008

MRF 15	MRF 15
Registration number:	Registration number:
A39/5.4/0226	A39/16.2/0544
Name of medicine:	Name of medicine:
LYRINEL 15 mg	CILODEX EAR DROPS
Dosage form:	Dosage form:
TABLET	SUSPENSION
Active ingredients:	Active ingredients:
EACH TABLET CONTAINS: OXYBUTYNIIN HYDROCHLORIDE 15,0 mg	EACH 1,0 ml SUSPENSION CONTAINS: CIPROFLOXACIN HYDROCHLORIDE EQUIVALENT TO CIPROFLOXACIN 3,0 mg DEXAMETHASONE 1,0 mg
Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
Applicant:	Applicant:
JANSSEN PHARMACEUTICA (PTY) LTD	ALCON LABORATORIES (S.A.) (PTY) LTD
Manufacturer:	Manufacturer:
ALZA CORPORATION, MOUNTAIN VIEW, CALIFORNIA, USA ALZA CORPORATION, VACAVILLE, CALIFORNIA, USA	S.A. ALCON-COUVREUR N.V., PUURS, BELGIUM
Packer:	Packer:
ALZA CORPORATION, VACAVILLE, CALIFORNIA, USA JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY	S.A. ALCON-COUVREUR N.V., PUURS, BELGIUM
Laboratory:	Laboratory:
FPRC: ALZA IRELAND LTD, TIPPERARY, IRELAND JANSSEN-CILAG SpA, BORGO S. MICHELLE, LATINA, ITALY JANSSEN PHARMACEUTICA NV, BEERSE, BELGIUM	FPRC: S.A. ALCON-COUVREUR N.V., PUURS, BELGIUM RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
Shelf-life:	Shelf-life:
18 months	24 months
Date of registration:	Date of registration:
13 JUNE 2008	13 JUNE 2008
FPRC: JANSSEN PHARMACEUTICA, WOODMEAD, JOHANNESBURG	FPRC: ALCON LABORATORIES, BRYANSTON, JOHANNESBURG

MRF 15	MRF 15
Registration number:	A40/5.10/0213
Name of medicine:	ADCO-NETRIN 2 mg/ml
Dosage form:	INJECTION
Active ingredients:	EACH 2.0 ml SOLUTION CONTAINS: ONDANSETRON 4.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	PHARMAMED PARENTERALS LTD, ZEJTUN, MALTA FAMAR S.A., P. FALIRO, GREECE
Packer:	PHARMAMED PARENTERALS LTD, ZEJTUN, MALTA FAMAR S.A., P. FALIRO, GREECE SYNTHON HISPANIA, BARCELONA, SPAIN NYCOMED AUSTRIA GmbH, LINZ, AUSTRIA ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG PHARMA-Q, INDUSTRIA, JOHANNESBURG
Laboratory:	FPRC: PHARMAMED PARENTERALS LTD, ZEJTUN, MALTA FAMAR S.A., P. FALIRO, GREECE SYNTHON HISPANIA, BARCELONA, SPAIN SYNTHON BV, NIJMEGEN, THE NETHERLANDS NYCOMED AUSTRIA GmbH, LINZ, AUSTRIA PHARMA-Q, INDUSTRIA, JOHANNESBURG
FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG
FPRR:	ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON ADCOCK INGRAM LTD, BRYANSTON, JOHANNESBURG
Shelf-life:	24 months
Date of registration:	13 JUNE 2008
Registration number:	A40/21.5.1/0224
Name of medicine:	SPEC-BUDESONIDE 100
Dosage form:	NASAL SPRAY
Active ingredients:	EACH METERED DOSE CONTAINS: BUDESONIDE 100.0 ug
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SPECPHARM (PTY) LTD
Manufacturer:	MIPHARM S.p.A, MILAN, ITALY
Packer:	MIPHARM S.p.A, MILAN, ITALY
Laboratory:	FPRC: MIPHARM S.p.A, MILAN, ITALY ANALYTICON, TERENURE, KEMPTON PARK
FPRR:	SPECPHARM, HALFWAY HOUSE, MIDRAND
Shelf-life:	24 months
Date of registration:	13 JUNE 2008

MRF 15	MRF 15
Registration number: 41/24/0136	Registration number: 41/7.5/0298
Name of medicine: 0.9 % SODIUM CHLORIDE INFUSION B. BRAUN	Name of medicine: CRESTOR 5
Dosage form: INFUSION	Dosage form: TABLET
Active ingredients: EACH 1000,0 ml SOLUTION CONTAINS: SODIUM CHLORIDE	Active ingredients: EACH TABLET CONTAINS: ROSUVASTATIN CALCIUM EQUIVALENT TO: ROSUVASTATIN 5,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: B BRAUN MEDICAL (PTY) LTD	Applicant: ASTRAZENECA PHARMACEUTICALS (PTY) LTD
Manufacturer: B BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY B BRAUN MELSUNGEN PRODUCTION PHARMA PFIFFEWIESEN, MELSUNGEN, GERMANY B BRAUN MEDICAL S.A, RUBI, BARCELONA, SPAIN	Packer: IPR PHARMACEUTICALS Inc, CAROLINA, PUERTO RICO IPR PHARMACEUTICALS Inc, CANOVANAS, PUERTO RICO ASTRAZENECA GmbH, PLANKSTADT, GERMANY ASTRAZENECA UK LTD, MACCLESFIELD, CHESHIRE, UK ASTRAZENECA PHARMACEUTICALS, ALRODE, ALBERTON
Packer: B BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY B BRAUN MELSUNGEN PRODUCTION PHARMA PFIFFEWIESEN, MELSUNGEN, GERMANY	Laboratory: ASTRAZENECA UK LTD, MACCLESFIELD, CHESHIRE, UK ASTRAZENECA GmbH, PLANKSTADT, GERMANY ANALYTICON, TERENURE, KEMPTON PARK CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
Laboratory: FPRC: B BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG COSI PHARMACEUTICALS, INDUSTRIA WEST, JOHANNESBURG CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG	Laboratory: FPRC: ASTRAZENECA UK LTD, MACCLESFIELD, CHESHIRE, UK ASTRAZENECA GmbH, PLANKSTADT, GERMANY ANALYTICON, TERENURE, KEMPTON PARK CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRC: B BRAUN MEDICAL, HONEYDEW, GAUTENG	FPRC/FPRR: ASTRAZENECA PHARMACEUTICALS, ALRODE, ALBERTON
Shelf-life: 36 months	Shelf-life: 36 months
Date of registration: 13 JUNE 2008	Date of registration: 13 JUNE 2008