
GENERAL NOTICE ALGEMENE KENNISGEWING

NOTICE 905 OF 2008

CONDITIONS OF REGISTRATION OF A MEDICINE IN TERMS OF THE PROVISIONS OF SECTION 15(7) OF THE MEDICINES AND RELATED SUBSTANCES ACT, 1965 (ACT 101 OF 1965)

1. The applicant shall ensure that the medicine is manufactured and controlled in terms of the current Good Manufacturing Practices as determined by Council
2. The manufacture of this medicine is subject to regular investigation and inspections by the inspectors appointed in terms of Section 26 of the Act, to assess compliance with current Good Manufacturing Practices.
3. The information in the package insert shall be updated on a regular basis to conform to the package insert recently approved by Council.
4. The applicant must comply with all the legal requirements of the Medicines and Related Substances Act, 1965 (Act 101 of 1965).
5. The registration of this medicine shall be subject to review at intervals as determined by Council regarding its quality, safety and efficacy, and the registration of this medicine may be varied subject to issues Council may deem fit.
6. The first two production batches must be fully validated in terms of the detailed process validation protocol submitted at the time of application for registration, and the validation report must be submitted within a month after completion of the validation.
7. The product may be advertised to the professions only.
8. The provisional shelf-life allocated must be confirmed with stability data over the full shelf-life period on the first two production batches (well-known API) or first three production batches (NCE) in accordance with the Stability Guideline. Stability testing submitted on any pilot batches must also be completed and reported on. The stability report must be submitted within six months after completion of the stability trial. However, Council has to be informed immediately if any parameter shows a significant change or out-of-specification result during the stability trial.
9. A post-registration inspection must be conducted on the first production batch manufactured by each local manufacturer.
10. A post-registration inspection must be conducted on the first production batch of the imported product.
11. Marketing of the product may only commence following a satisfactory post-registration inspection report.
12. One sample of every batch, together with four copies of the protocol for testing of the bulk lot and filling lot, and six copies of the certificate of release issued by a competent authority in the country in which the product was manufactured, must be submitted to the Council for lot release purposes.
13. The expiry date allocated shall be modified by adding a statement that the virus strains are currently recommended for South African usage in the specific year.
14. The strains of the master seed viruses must be approved by the Department of Health for each year.

MRF 15

Registration number: 3677.5/0281

Name of medicine: BEZACHOLE SR

Dosage form: TABLET

Active ingredients: EACH TABLET CONTAINS: 400,0 mg
BEZAFIBRATE

Conditions of registration: 1, 2, 3, 4, 5, 6, 7

Applicant: PHARMACARE LIMITED

Manufacturer: VALPHARMA INTERNATIONAL S.p.A.,
PENNABILLI, PERSARO-URBINO, ITALY

Packer: ASPEN PHARMACARE EAST LONDON, WILSONIA,
EAST LONDON
PHARMACARE LTD, KORSTEN, PORT ELIZABETH

Laboratory: FPRC:
VALPHARMA INTERNATIONAL S.p.A.,
PENNABILLI, PERSARO-URBINO, ITALY
SEDEK AGRIKEM, KAMEELDRIEF EAST, PRETORIA
INSTITUTE FOR PHARMACEUTICAL SERVICES,
SILVERTONDALE, PRETORIA
SOUTH AFRICAN BUREAU OF STANDARDS,
GROENKLOOF, PRETORIA
M&L LABORATORY SERVICES, ORMONDE,
JOHANNESBURG
RESEARCH INSTITUTE FOR INDUSTRIAL
PHARMACY, NORTH-WEST UNIVERSITY,
POTCHEFSTROOM
ASPEN PHARMACARE EAST LONDON, WILSONIA,
EAST LONDON

FPRC/FPRR: PHARMACARE LTD, KORSTEN, PORT ELIZABETH

Shelf-life: 24 months

Date of registration: 18 APRIL 2008

MRF 15

Registration number: 37/26/0483

Name of medicine: NEOTALEM

Dosage form: INJECTION

Active ingredients: EACH 10,0 ml VIAL CONTAINS:
MITOXANTRONE HYDROCHLORIDE 20,0 mg
EQUIVALENT TO MITOXANTRONE

Conditions of registration: 1, 2, 3, 4, 5, 6, 7

Applicant: KEY ONCOLOGICS (PTY) LTD

Manufacturer: LEMERY S.A. de C.V, HUICHAPAN, XOCH,
MEXICO

Packer: LEMERY S.A. de C.V, HUICHAPAN, XOCH,
MEXICO

Laboratory: FPRC:
LEMERY S.A. de C.V, HUICHAPAN, XOCH,
MEXICO
INSTITUTE FOR PHARMACEUTICAL SERVICES,
SILVERTONDALE, PRETORIA

FPRR: KEY ONCOLOGICS, SANDTON, JOHANNESBURG

Shelf-life: 24 months

Date of registration: 18 APRIL 2008

MRF 15	MRF 15		
Registration number:	38/7.1.3/0032	Registration number:	38/7.1.3/0033
Name of medicine:	AUSTELL-LISINAPRIL 10 mg	Name of medicine:	AUSTELL-LISINAPRIL 5 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 10,0 mg	Active ingredients:	EACH TABLET CONTAINS: LISINOPRIL DIHYDRATE EQUIVALENT TO LISINOPRIL 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AUSTELL LABORATORIES (PTY) LTD	Applicant:	AUSTELL LABORATORIES (PTY) LTD
Manufacturer:	IPCA LABORATORIES LTD, SILVASSA, DADRA & NAGAR HAVELI, INDIA	Manufacturer:	IPCA LABORATORIES LTD, SILVASSA, DADRA & NAGAR HAVELI, INDIA
Packer:	IPCA LABORATORIES LTD, SILVASSA, DADRA & NAGAR HAVELI, INDIA	Packer:	IPCA LABORATORIES LTD, SILVASSA, DADRA & NAGAR HAVELI, INDIA
Laboratory:	FPRC: IPCA LABORATORIES LTD, SILVASSA, DADRA & NAGAR HAVELI, INDIA SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA	Laboratory:	FPRC: IPCA LABORATORIES LTD, SILVASSA, DADRA & NAGAR HAVELI, INDIA 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
FPRR:	AUSTELL LABORATORIES, SPRINGFIELD, JOHANNESBURG	FPRR:	AUSTELL LABORATORIES, SPRINGFIELD, JOHANNESBURG

MRF 15	MRF 15
Registration number: 38/26/0057	Registration number: 38/20.1.1/0204
Name of medicine: LITAK 10	Name of medicine: CEFUROXIME-SAFELINE 1,5 g
Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH 5,0 ml VIAL CONTAINS: CLADRIBINE 10,0 mg	Active ingredients: EACH VIAL CONTAINS: CEFUROXIME SODIUM EQUIVALENT TO CEFUROXIME 1,5 g
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: KEY ONCOLOGICS (PTY) LTD	Applicant: SAFELINE PHARMACEUTICALS (PTY) LTD
Manufacturer: HAUPT PHARMA GmbH, WOLFRATSHAUSEN, GERMANY	Manufacturer: DEMO S.A. PHARMACEUTICAL, KRYONERI, ATHENS, GREECE
Packer: HAUPT PHARMA GmbH, WOLFRATSHAUSEN, GERMANY	Packer: DEMO S.A. PHARMACEUTICAL, KRYONERI, ATHENS, GREECE
Laboratory: FPRC: LIPOMED AG, ARLESHEIM, SWITZERLAND INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA	Laboratory: FPRC: DEMO S.A. PHARMACEUTICAL, KRYONERI, ATHENS, GREECE INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA
FPRC: KEY ONCOLOGICS, SANDTON, JOHANNESBURG	FPRC: SAFELINE PHARMACEUTICALS, FLORIDA, JOHANNESBURG
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: 38/20.1.1/0205	Registration number: A04/3.1.5/08
Name of medicine: CEFUROXIME-SAFELINE 750 mg	Name of medicine: TILDREN
Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH VIAL CONTAINS: CEFUROXIME SODIUM EQUIVALENT TO CEFUROXIME 750 mg	Active ingredients: EACH VIAL CONTAINS: TILDURONIC ACID 50,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: SAFELINE PHARMACEUTICALS (PTY) LTD	Applicant: CEVA ANCHORPHARM ANIMAL HEALTH S.A (PTY) LTD
Manufacturer: DEMO S.A. PHARMACEUTICAL, KRYONERI, ATHENS, GREECE	Manufacturer: CEVA SANTE ANIMALE SA, CEDEX, FRANCE
Packer: DEMO S.A. PHARMACEUTICAL, KRYONERI, ATHENS, GREECE	Packer: CEVA SANTE ANIMALE SA, CEDEX, FRANCE
Laboratory: FPRC: DEMO S.A. PHARMACEUTICAL, KRYONERI, ATHENS, GREECE INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA	Laboratory: FPRC: CEVA SANTE ANIMALE SA, CEDEX, FRANCE CEVA LABORATOIRES STERILYO, SANT AMAND LES EAUX, FRANCE CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRC: SAFELINE PHARMACEUTICALS, FLORIDA, JOHANNESBURG	FPRC: CEVA ANCHORPHARM ANIMAL HEALTH, BRAMLEY, JOHANNESBURG
Shelf-life: 24 months	Shelf-life: 36 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	A05/22.6.2/10
Name of medicine:	ALIZINE
Dosage form:	INJECTION
Active ingredients:	EACH 1,0 ml SOLUTION CONTAINS: AGLEPRISTONE 30,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	VIRBAC RSA (PTY) LTD
Manufacturer:	VIRBAC SA, CEDEX, FRANCE
Packer:	VIRBAC SA, CEDEX, FRANCE VIRBAC RSA, CENTURION, RSA
Laboratory:	FPRC: M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
	FPRR
Shelf-life:	36 months
Date of registration:	18 APRIL 2008
Registration number:	A38/3.2/0381
Name of medicine:	BONDRONAT 6 mg/6 ml
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: IBANDRONIC ACID 6,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ROCHE PRODUCTS (PTY) LTD
Manufacturer:	ROCHE DIAGNOSTIC GmbH, MANNHEIM, GERMANY
Packer:	ROCHE DIAGNOSTIC GmbH, MANNHEIM, GERMANY ROCHE PRODUCTS, ISANDO, RSA
Laboratory:	FPRC: ROCHE DIAGNOSTIC GmbH, MANNHEIM, GERMANY
	FPRC/FPRR:
Shelf-life:	60 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: A39/21.12/0064	Registration number: A39/21.12/0065
Name of medicine: CETROTIDE 0,25 mg	Name of medicine: CETROTIDE 3 mg
Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH VIAL CONTAINS: CETRORELIX ACETATE EQUIVALENT TO CETRORELIX 0,25 mg	Active ingredients: EACH VIAL CONTAINS: CETRORELIX ACETATE EQUIVALENT TO CETRORELIX 3,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: MERCK (PTY) LTD	Applicant: MERCK (PTY) LTD
Manufacturer: BAXTER ONCOLOGY GmbH, HALLE, GERMANY	Manufacturer: BAXTER ONCOLOGY GmbH, HALLE, GERMANY
Packer: BAXTER ONCOLOGY GmbH, HALLE, GERMANY SOLVAY PHARMACEUTICALS BV, OLST, THE NETHERLANDS	Packer: BAXTER ONCOLOGY GmbH, HALLE, GERMANY SOLVAY PHARMACEUTICALS BV, OLST, THE NETHERLANDS
Laboratory: FPRC: BAXTER ONCOLOGY GmbH, HALLE, GERMANY SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA	Laboratory: FPRC: BAXTER ONCOLOGY GmbH, HALLE, GERMANY SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRC	FPRC
Shelf-life: 24 months	Shelf-life: 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008
MERCK, MODDERFONTEIN, RSA	MERCK, MODDERFONTEIN, RSA

MRF 15	MRF 15
Registration number:	A39/2.1/0077
Name of medicine:	B. BRAUN PROPOFOL 1 % (10 mg/ml)
Dosage form:	EMULSION
Active ingredients:	EACH 1,0 ml EMULSION CONTAINS: PROPOFOL 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	B BRAUN MEDICAL (PTY) LTD
Manufacturer:	B BRAUN MELSUNGEN AG, BERLIN, GERMANY
Packer:	B BRAUN MELSUNGEN AG, BERLIN, GERMANY
Laboratory:	FPRC: B BRAUN MELSUNGEN AG, BERLIN, GERMANY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
Shelf-life:	24 months
Date of registration:	18 APRIL 2008
Registration number:	A39/30.4/0417
Name of medicine:	CEPROTIN 500 iu
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: PROTEIN C 500,0 iu
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD
Manufacturer:	BAXTER AG, INDUSTRIESTRASSE, VIENNA, AUSTRIA BAXTER AG, BENATZKYGASSE, VIENNA, AUSTRIA BAXTER S.p.A., RUFINA, ITALY
Packer:	BAXTER AG, INDUSTRIESTRASSE, VIENNA, AUSTRIA ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG
Laboratory:	FPRC: BAXTER AG, SMOLAGASSE, VIENNA, AUSTRIA
Shelf-life:	24 months
Date of registration:	18 APRIL 2008
FPRC/FPRR:	FPRC/FPRR:
ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG	ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG

MRF 15	MRF 15
Registration number: A39/30.4/0418	Registration number: A39/1.2/0500
Name of medicine: CEPROTIN 1000 iu	Name of medicine: TAZERON 15 mg
Dosage form: INJECTION	Dosage form: TABLET
Active ingredients: EACH VIAL CONTAINS: PROTEIN C 1000,0 iu	Active ingredients: EACH TABLET CONTAINS: MIRTAZAPINE 15,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: ADCOCK INGRAM CRITICAL CARE (PTY) LTD	Applicant: PHARMAPLAN (PTY) LTD
Manufacturer: BAXTER AG, INDUSTRIESTRASSE, VIENNA, AUSTRIA BAXTER AG, BENATZKYGASSE, VIENNA, AUSTRIA BAXTER S.p.A., RUFINA, ITALY	Manufacturer: KERN PHARMA S.L., TERRASSA, BARCELONA, SPAIN
Packer: BAXTER AG, INDUSTRIESTRASSE, VIENNA, AUSTRIA ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG	Packer: KERN PHARMA S.L., TERRASSA, BARCELONA, SPAIN
Laboratory: FPRC:	Laboratory: FPRC: COMBINO PHARM S.L., SANT JOAN DESPI, BACELONA, SPAIN CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRC/FPRR: ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG	FPRR: PHARMAPLAN, MIDRAND, RSA
Shelf-life: 24 months	Shelf-life: 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15

Registration number: A39/1.2/0501
 Name of medicine: TAZERON 30 mg
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 MIRTAZAPINE 30,0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: PHARMAPLAN (PTY) LTD
 Manufacturer: KERN PHARMA S.L., TERRASSA, BARCELONA,
 SPAIN
 Packer: KERN PHARMA S.L., TERRASSA, BARCELONA,
 SPAIN
 Laboratory: FPRC: COMBINO PHARM S.L., SANT JOAN DESPI,
 BACELONA, SPAIN
 CONSULTING CHEMICAL LABORATORIES,
 ATLASVILLE, BOKSBURG
 Shelf-life: 24 months
 Date of registration: 18 APRIL 2008

MRF 15

Registration number: A39/21.5.1/0584
 Name of medicine: SANDOZ BECLOMETHASONE
 DIPROPIONATE AQ 100
 Dosage form: SUSPENSION
 Active ingredients: EACH METERED NASAL SPRAY DOSE
 CONTAINS:
 BECLOMETHASONE DIPROPIONATE
 100,0 ug
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: SANDOZ S.A. (PTY) LTD
 Manufacturer: CHIESI FARMACEUTICI SpA, LEONARDO,
 PARMA, ITALY
 Packer: CHIESI FARMACEUTICI SpA, LEONARDO,
 PARMA, ITALY
 CHIESI FARMACEUTICI SpA, PALERMO,
 PARMA, ITALY
 NOVARTIS S.A., SPARTAN, KEMPTON
 PARK
 Laboratory: FPRC: CHIESI FARMACEUTICI SpA, LEONARDO,
 PARMA, ITALY
 SOUTH AFRICAN BUREAU OF STANDARDS,
 GROENKLOOF, PRETORIA
 NOVARTIS S.A., SPARTAN, KEMPTON
 PARK
 Shelf-life: 36 months
 Date of registration: 18 APRIL 2008

MRF 15	MRF 15		
Registration number:	A40/13.4.1/0035	Registration number:	A40/34/0045
Name of medicine:	CLOBEX SHAMPOO	Name of medicine:	CIPLA-TAMSULOSIN HYDROCHLORIDE
Dosage form:	SHAMPOO	Dosage form:	CAPSULE
Active ingredients:	EACH 1.0 g SHAMPOO CONTAINS: CLOBETASOL PROPIONATE	Active ingredients:	EACH CAPSULE CONTAINS: TAMSULOSIN HYDROCHLORIDE
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	GALDERMA LABORATORIES S.A. (PTY) LTD	Applicant:	CIPLA LIFE SCIENCES (PTY) LTD
Manufacturer:	LABORATOIRES GALDERMA, MONTDESIR, ALBY-SUR-CHERAN, FRANCE	Manufacturer:	CIPLA LTD, VIRGONAGAR, BANGALORE, INDIA
Packer:	LABORATOIRES GALDERMA, MONTDESIR, ALBY-SUR-CHERAN, FRANCE	Packer:	CIPLA LTD, VIRGONAGAR, BANGALORE, INDIA
Laboratory:	LABORATOIRES GALDERMA, MONTDESIR, ALBY-SUR-CHERAN, FRANCE	Laboratory:	CIPLA LTD, VIRGONAGAR, BANGALORE, INDIA
FPRC:	FPRC:	FPRC:	FPRC:
FPRC:	GALDERMA LABORATORIES S.A., BRYANSTON, JOHANNESBURG	FPRC:	CIPLA LIFE SCIENCES, ROSEN PARK, BELLVILLE
Shelf-life:	36 months	Shelf-life:	24 months
Date of registration:	18 APRIL 2008	Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/34/0046	Registration number: A40/20.1.1/0102
Name of medicine: UROMAX	Name of medicine: ZITHROMAX ONE
Dosage form: CAPSULE	Dosage form: GRANULES
Active ingredients: EACH CAPSULE CONTAINS: TAMSULOSIN HYDROCHLORIDE 400,0 ug	Active ingredients: EACH BOTTLE CONTAINS: AZITHROMYCIN DIHYDRATE EQUIVALENT TO AZITHROMYCIN 2,0 g
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: CIPLA MEDPRO (PTY) LTD	Applicant: PFIZER LABORATORIES (PTY) LTD
Manufacturer: CIPLA LTD, VIRGONAGAR, BANGALORE, INDIA	Manufacturer: PFIZER PHARMACEUTICALS LLC, VEGA BAJA, PUERTO RICO
Packer: CIPLA LTD, VIRGONAGAR, BANGALORE, INDIA	Packer: PFIZER PHARMACEUTICALS LLC, VEGA BAJA, PUERTO RICO PFIZER ITALIA S.r.l., LATINA, ITALY PFIZER GLOBAL MANUFACTURING, RETREAT, CAPE TOWN
Laboratory: FPRC:	Laboratory: FPRC: PFIZER PHARMACEUTICALS LLC, VEGA BAJA, PUERTO RICO PFIZER ITALIA S.r.l., LATINA, ITALY
FPRC: CIPLA MEDPRO, ROSEN PARK, BELLVILLE	FPRC/FPRR: PFIZER GLOBAL MANUFACTURING, RETREAT, CAPE TOWN
Shelf-life: 24 months	Shelf-life: FPRR PFIZER LABORATORIES, SANDTON, JOHANNESBURG 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	A40/21.2/0132
Name of medicine:	GLIMPID 1
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: GLIMEPIRIDE 1,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	RANBAXY (SA) (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, RSA
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	A40/21.2/0133
Name of medicine:	GLIMPID 2
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: GLIMEPIRIDE 2,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	RANBAXY (SA) (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, RSA
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/21.2/0134	Registration number: A40/2.5/0166
Name of medicine: GLIMPID 4	Name of medicine: DYNA-LAMOTRIGINE 100 mg
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: GLIMEPIRIDE 4,0 mg	Active ingredients: EACH TABLET CONTAINS: LAMOTRIGINE 100,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: RANBAXY (SA) (PTY) LTD	Applicant: PHARMA DYNAMICS (PTY) LTD
Manufacturer: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA	Manufacturer: MEDOCHEMIE LTD, LIMASSOL, CYPRUS
Packer: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA	Packer: MEDOCHEMIE LTD, LIMASSOL, CYPRUS DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES, N'DABENI, KZN IMPILO DRUGS, ISITHEBE, KZN
Laboratory: FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC: MEDOCHEMIE LTD, LIMASSOL, CYPRUS CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA TECHNIKON LABORATORIES, FLORIDA, RSA
FPRC: RANBAXY (SA), CENTURION, RSA	FPRC: PHARMA DYNAMICS, SILVERWOOD, WESTLAKE, RSA
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:	A40/2.5/0167
Name of medicine:	DYNA-LAMOTRIGINE 200 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LAMOTRIGINE 200,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	PHARMA DYNAMICS (PTY) LTD
Manufacturer:	MEDOCHEMIE LTD, LIMASSOL, CYPRUS
Packer:	MEDOCHEMIE LTD, LIMASSOL, CYPRUS DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES, N'DABENI, KZN
Laboratory:	MEDOCHEMIE LTD, LIMASSOL, CYPRUS CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA TECHNIKON LABORATORIES, FLORIDA, RSA
FPRC:	FPRC:
FPRR:	PHARMA DYNAMICS, SILVERWOOD, WESTLAKE, RSA
Shelf-life:	24 months
Date of registration:	18 APRIL 2008
Registration number:	A40/2.5/0169
Name of medicine:	DYNA-LAMOTRIGINE 50 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LAMOTRIGINE 50,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	PHARMA DYNAMICS (PTY) LTD
Manufacturer:	MEDOCHEMIE LTD, LIMASSOL, CYPRUS
Packer:	MEDOCHEMIE LTD, LIMASSOL, CYPRUS DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES, N'DABENI, KZN
Laboratory:	MEDOCHEMIE LTD, LIMASSOL, CYPRUS CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA TECHNIKON LABORATORIES, FLORIDA, RSA
FPRC:	FPRC:
FPRR:	PHARMA DYNAMICS, SILVERWOOD, WESTLAKE, RSA
Shelf-life:	24 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/2.5/0173	Registration number: A40/3.1/0240
Name of medicine: DYNA-LAMOTRIGINE 25 mg	Name of medicine: CATAFAST-D TABLETS
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: LAMOTRIGINE 25,0 mg	Active ingredients: EACH TABLET CONTAINS: DICLOFENAC SODIUM 50,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: PHARMA DYNAMICS (PTY) LTD	Applicant: NOVARTIS SOUTH AFRICA (PTY) LTD
Manufacturer: MEDOCHEMIE LTD, LIMASSOL, CYPRUS	Manufacturer: NOVARTIS PHARMA GmbH, WEHR, GERMANY
Packer: MEDOCHEMIE LTD, LIMASSOL, CYPRUS DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES, N'DABENI, KZN	Packer: NOVARTIS PHARMA GmbH, WEHR, GERMANY NOVARTIS S.A., SPARTAN, KEMPTON PARK
Laboratory: FPRC: MEDOCHEMIE LTD, LIMASSOL, CYPRUS CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA TECHNIKON LABORATORIES, FLORIDA, RSA	Laboratory: FPRC: NOVARTIS PHARMA GmbH, WEHR, GERMANY M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRR: PHARMA DYNAMICS, SILVERWOOD, WESTLAKE, RSA	FPRC/FPRR: NOVARTIS 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:	A40/7.1.3/0287
Name of medicine:	COIRBESARTAN WINTHROP 300/12,5
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: IRBESARTAN 300,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANOFI-SYNTHELABO (PTY) LTD
Manufacturer:	BRISTOL-MYERS SQUIBB CO., EVANSVILLE INDIANA, USA SANOFI WINTHROP INDUSTRIE, AMBARES, FRANCE
Packer:	SANOFI WINTHROP INDUSTRIE, AMBARES, FRANCE AVENTIS PHARMA, WALTLOO, PRETORIA PHARMACEUTICAL CONTRACTORS, ISANDO
Laboratory:	SANOFI WINTHROP INDUSTRIE, AMBARES, FRANCE M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRC/FPRR:	AVENTIS PHARMA, WALTLOO, PRETORIA
FPRR:	SANOFI-SYNTHELABO, MIDRAND, RSA
Shelf-life:	36 months
Date of registration:	18 APRIL 2008
Registration number:	A40/7.1.3/0290
Name of medicine:	COIRBESARTAN WINTHROP 150/12,5
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: IRBESARTAN 150,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANOFI-SYNTHELABO (PTY) LTD
Manufacturer:	BRISTOL-MYERS SQUIBB CO., EVANSVILLE INDIANA, USA SANOFI WINTHROP INDUSTRIE, AMBARES, FRANCE
Packer:	SANOFI WINTHROP INDUSTRIE, AMBARES, FRANCE AVENTIS PHARMA, WALTLOO, PRETORIA PHARMACEUTICAL CONTRACTORS, ISANDO
Laboratory:	SANOFI WINTHROP INDUSTRIE, AMBARES, FRANCE M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRR:	AVENTIS PHARMA, WALTLOO, PRETORIA
Shelf-life:	SANOFI-SYNTHELABO, MIDRAND, RSA
Date of registration:	36 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/34/0299	Registration number: A40/20.1.1/0309
Name of medicine: CYCLEAN	Name of medicine: LEVOFLOXACIN-WINTHROP 250
Dosage form: SOLUTION	Dosage form: TABLET
Active ingredients: EACH 1,0 ml SOLUTION CONTAINS: POLYHEXANIDE 0,1 µg	Active ingredients: EACH TABLET CONTAINS: LEVOFLOXACIN HEMIHYDRATE EQUIVALENT TO LEVOFLOXACIN 250,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: REVISION S.A. (PTY) LTD	Applicant: AVENTIS PHARMA (PTY) LTD
Manufacturer: SAUFLOX PHARMACEUTICALS LTD, TWICKENHAM, MIDDLESEX, U.K.	Manufacturer: AVENTIS PHARMA SPECIALITES, COMPEIGNE, FRANCE
Packer: SAUFLOX PHARMACEUTICALS LTD, TWICKENHAM, MIDDLESEX, U.K.	Packer: AVENTIS PHARMA DEUTSCHLAND GmbH, FRANKFURT, GERMANY
Laboratory: FPRC: SAUFLOX PHARMACEUTICALS LTD, TWICKENHAM, MIDDLESEX, U.K. TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA	Laboratory: FPRC: AVENTIS PHARMA SPECIALITES, COMPEIGNE, FRANCE AVENTIS PHARMA DEUTSCHLAND GmbH, FRANKFURT, GERMANY
FPRC: REVISION S.A., BEDFORDVIEW, JOHANNESBURG, RSA	FPRC/FPRR: WINTHROP PHARMACEUTICALS, WALTLOO, PRETORIA
Shelf-life: 24 months (Provisional)	Shelf-life: FPRR: AVENTIS PHARMA, MIDRAND, RSA 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/20.1.1/0310	Registration number: A40/34/0314
Name of medicine: LEVOFLOXACIN-WINTHROP 500	Name of medicine: OCTREOTIDE HEXAL 0,05 mg
Dosage form: TABLET	Dosage form: INJECTION
Active ingredients: EACH TABLET CONTAINS: LEVOFLOXACIN HEMIHYDRATE EQUIVALENT TO LEVOFLOXACIN 500,0 mg	Active ingredients: EACH 1,0 ml SOLUTION CONTAINS: OCTREOTIDE 0,05 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: AVENTIS PHARMA (PTY) LTD	Applicant: HEXAL PHARMA (SA) (PTY)
Manufacturer: AVENTIS PHARMA SPECIALITES, COMPEIGNE, FRANCE	Manufacturer: WASSERBURGER ARZNEIMITTELWERK GmbH, WASSERBURG AM INN, GERMANY
Packer: AVENTIS PHARMA DEUTSCHLAND GmbH, FRANKFURT, GERMANY	Packer: WASSERBURGER ARZNEIMITTELWERK GmbH, WASSERBURG AM INN, GERMANY SALUTAS PHARMA GmbH, BARLEBEN, GERMANY DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA PHARMA-Q, INDUSTRIA, JOHANNESBURG
Laboratory: FPRC: AVENTIS PHARMA SPECIALITES, COMPEIGNE, FRANCE FPRR: AVENTIS PHARMA DEUTSCHLAND GmbH, FRANKFURT, GERMANY	Laboratory: FPRC: WASSERBURGER ARZNEIMITTELWERK GmbH, WASSERBURG AM INN, GERMANY FPRR: SALUTAS PHARMA GmbH, BARLEBEN, GERMANY LABOR L+S AG, BAD BOCKLET-GROSSENBACH, GERMANY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG ANALYTICON, TERENURE, KEMPTON PARK
FPRC/FPRR: WINTHROP PHARMACEUTICALS, WALTLOO, PRETORIA	FPRC/FPRR: HEXAL PHARMA, PINETOWN, KZN
Shelf-life: 60 months	Shelf-life: 24 months
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	Registration number:
A40/34/0315	A40/20.1.1/0522
Name of medicine:	Name of medicine:
OCTREOTIDE HEXAL 0,1 mg	TOBI 300 mg/5 ml
Dosage form:	Dosage form:
INJECTION	SOLUTION
Active ingredients:	Active ingredients:
EACH 1,0 ml SOLUTION CONTAINS: OCTREOTIDE 0,1 mg	EACH 5,0 ml SOLUTION CONTAINS: TOBRAMYCIN 300,0 mg
Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
Applicant:	Applicant:
HEXAL PHARMA (SA) (PTY)	NOVARTIS SA (PTY) LTD
Manufacturer:	Manufacturer:
WASSERBURGER ARZNEIMITTELWERK GmbH, WASSERBURG AM INN, GERMANY	CARDINAL HEALTH INC, WOODSTOCK, ILLINOIS, USA
Packer:	Packer:
WASSERBURGER ARZNEIMITTELWERK GmbH, WASSERBURG AM INN, GERMANY SALUTAS PHARMA GmbH, BARLEBEN, GERMANY DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA PHARMA-Q, INDUSTRIA, JOHANNESBURG	CARDINAL HEALTH INC, WOODSTOCK, ILLINOIS, USA CARDINAL HEALTH LTD, BOLTON, LANCASHIRE, UK
Laboratory:	Laboratory:
FPRC: WASSERBURGER ARZNEIMITTELWERK GmbH, WASSERBURG AM INN, GERMANY SALUTAS PHARMA GmbH, BARLEBEN, GERMANY LABOR L+S AG, BAD BOCKLET-GROSSENBACH, GERMANY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG ANALYTICON, TERENCE, KEMPTON PARK	FPRC: CARDINAL HEALTH LTD, BOLTON, LANCASHIRE, UK CHIRON CORPORATION, ANNANDALE, NEW JERSEY, USA TEPNEL LIFE SCIENCES, EDINBURGH, UK INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA
FPRR:	FPRR:
HEXAL PHARMA, PINETOWN, KZN	NOVARTIS SA, SPARTAN, KEMPTON PARK
Shelf-life:	Shelf-life:
24 months	36 months
Date of registration:	Date of registration:
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/21.5.1/0524	Registration number: A40/20.2.8/0619
Name of medicine: AP METHYLPRED INJECTION	Name of medicine: SONKE-LASTAD 30 TABLET
Dosage form: EACH 8.0 ml SOLUTION CONTAINS: METHYLPREDNISOLONE SODIUM SUCCINATE EQUIVALENT TO METHYLPREDNISOLONE 500,0 mg	Dosage form: EACH TABLET CONTAINS: LAMIVUDINE 150,0 mg STAVUDINE 30,0 mg
Active ingredients:	Active ingredients:
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: PHARMACARE LIMITED	Applicant: RANBAXY (S.A.) (PTY) LTD
Manufacturer: STRIDES ARCOLAB, ANEKAL TALUK, BANGALORE, INDIA	Manufacturer: RANBAXY LABORATORIES LTD, PAONTA SAHIB, DISTRICT SIRMOUR, HIMACHAL PRADESH, INDIA
Packer: STRIDES ARCOLAB, ANEKAL TALUK, BANGALORE, INDIA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON PHARMACARE LTD, KORSTEN, PORT ELIZABETH	Packer: RANBAXY LABORATORIES LTD, PAONTA SAHIB, DISTRICT SIRMOUR, HIMACHAL PRADESH, INDIA
Laboratory: FPRC: STRIDES ARCOLAB, ANEKAL TALUK, BANGALORE, INDIA M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM	Laboratory: FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB, DISTRICT SIRMOUR, HIMACHAL PRADESH, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRC/FPRR: ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON PHARMACARE LTD, KORSTEN, PORT ELIZABETH	FPRR: RANBAXY, CENTURION, RSA
Shelf-life: 24 months (Provisional)	Shelf-life: 24 months (provisional)
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15

Registration number: A40/20.2.8/0620
 Name of medicine: SONKE-LASTAD 40
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 LAMIVUDINE 150,0 mg
 STAVUDINE 40,0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
 Applicant: RANBAXY (S.A.) (PTY) LTD
 Manufacturer: RANBAXY LABORATORIES LTD, PAONTA SAHIB,
 DISTRICT SIRMOUR, HIMACHAL PRADESH, INDIA
 Packer: RANBAXY LABORATORIES LTD, PAONTA SAHIB,
 DISTRICT SIRMOUR, HIMACHAL PRADESH, INDIA
 Laboratory: FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB,
 DISTRICT SIRMOUR, HIMACHAL PRADESH, INDIA
 CONSULTING CHEMICAL LABORATORIES,
 ATLASVILLE, BOSKBURG
 Shelf-life: 24 months (provisional)
 Date of registration: 18 APRIL 2008

MRF 15

Registration number: A40/3.1/0664
 Name of medicine: ARROW MELOXICAM 7,5
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 MELOXICAM 7,5 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: ARROW PHARMA SOUTH AFRICA (PTY)
 LTD
 Manufacturer: RX MANUFACTURING INC, MISSISSAUGA,
 ONTARIO, CANADA
 Packer: RX MANUFACTURING INC, MISSISSAUGA,
 ONTARIO, CANADA
 DIVPHARM MANUFACTURING &
 PACKAGING, LONGDALE, JOHANNESBURG
 Laboratory: FPRC: RX MANUFACTURING INC, MISSISSAUGA,
 ONTARIO, CANADA
 RESEARCH INSTITUTE FOR INDUSTRIAL
 PHARMACY, NORTH-WEST UNIVERSITY,
 POTCHEFSTROOM
 SEDEK AGRIKEM CC, KAMEELDRIEF,
 PRETORIA
 Shelf-life: 24 months
 Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number: A40/3.1/0665	41/7.1.3/0001
Name of medicine: ARROW MELOXICAM 15	ASPEN CARVEDILOL 25 mg
Dosage form: TABLET	TABLET
Active ingredients: EACH TABLET CONTAINS: MELOXICAM 15,0 mg	EACH TABLET CONTAINS: CARVEDILOL 25,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
Applicant: ARROW PHARMA SOUTH AFRICA (PTY) LTD	PHARMACARE LIMITED
Manufacturer: RX MANUFACTURING INC, MISSISSAUGA, ONTARIO, CANADA	PHARMASCIENCE INC, MONTREAL, CANADA PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
Packer: RX MANUFACTURING INC, MISSISSAUGA, ONTARIO, CANADA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG	PHARMASCIENCE INC, MONTREAL, CANADA PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
Laboratory: FPRC: RX MANUFACTURING INC, MISSISSAUGA, ONTARIO, CANADA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM SEDEK AGRIKEM CC, KAMEELDRIFT, PRETORIA	FPRC: PHARMASCIENCE INC, MONTREAL, CANADA SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRC: ARROW PHARMA SA, WOODMEAD, JOHANNESBURG	FPRC: PHARMACARE LTD, WOODMEAD, JOHANNESBURG
Shelf-life: 24 months	FPRC/FPRR: PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON 36 months
Date of registration: 18 APRIL 2008	Shelf-life: 36 months Date of registration: 18 APRIL 2008

MRF 15

Registration number: 417.1.3/0002
 Name of medicine: ASPEN CARVEDILOL 6,25 mg
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS: CARVEDILOL 6,25 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: PHARMACARE LIMITED
 Manufacturer: PHARMASCIENCE INC, MONTREAL, CANADA
 PHARMACARE LTD, KORSTEN, PORT ELIZABETH
 ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
 Packer: PHARMASCIENCE INC, MONTREAL, CANADA
 PHARMACARE LTD, KORSTEN, PORT ELIZABETH
 ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
 Laboratory: FPRC: PHARMASCIENCE INC, MONTREAL, CANADA
 SOUTH AFRICAN BUREAU OF STANDARDS,
 GROENKLOOF, PRETORIA
 RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY,
 NORTH-WEST UNIVERSITY, POTCHEFSTROOM
 M&L LABORATORY SERVICES, ORMONDE,
 JOHANNESBURG
 FPRR: PHARMACARE LTD, WOODMEAD, JOHANNESBURG
 FPRC/FPRR: PHARMACARE LTD, KORSTEN, PORT ELIZABETH
 ASPEN PHARMACARE EAST LONDON, WILSONIA,
 EAST LONDON
 Shelf-life: 36 months
 Date of registration: 18 APRIL 2008

MRF 15

Registration number: 417.1.3/0003
 Name of medicine: ASPEN CARVEDILOL 12,5 mg
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS: CARVEDILOL 12,5 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: PHARMACARE LIMITED
 Manufacturer: PHARMASCIENCE INC, MONTREAL, CANADA
 PHARMACARE LTD, KORSTEN, PORT ELIZABETH
 ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
 Packer: PHARMASCIENCE INC, MONTREAL, CANADA
 PHARMACARE LTD, KORSTEN, PORT ELIZABETH
 ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
 Laboratory: FPRC: PHARMASCIENCE INC, MONTREAL, CANADA
 SOUTH AFRICAN BUREAU OF STANDARDS,
 GROENKLOOF, PRETORIA
 RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY,
 NORTH-WEST UNIVERSITY,
 POTCHEFSTROOM
 M&L LABORATORY SERVICES, ORMONDE,
 JOHANNESBURG
 FPRR: PHARMACARE LTD, WOODMEAD,
 JOHANNESBURG
 FPRC/FPRR: PHARMACARE LTD, KORSTEN, PORT ELIZABETH
 ASPEN PHARMACARE EAST LONDON, WILSONIA,
 EAST LONDON
 Shelf-life: 36 months
 Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/20.2.3/0079
Name of medicine:	RIFIZID 150/75
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RIFAMPICIN 150,0 mg ISONIAZID 75,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	DEZZO TRADING (392) (PTY) LTD t/a INDO PHARMA
Manufacturer:	SVIZERA LABS PRIVATE LTD, MUMBAI, INDIA
Packer:	SVIZERA LABS PRIVATE LTD, MUMBAI, INDIA
Laboratory:	SVIZERA LABS PRIVATE LTD, MUMBAI, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA
FPRG/FPRR:	FPRR:
Shelf-life:	24 months
Date of registration:	18 APRIL 2008
Registration number:	41/30.1/0025
Name of medicine:	VIVAXIM
Dosage form:	INJECTION
Active ingredients:	EACH 1,0 ml DOSE CONTAINS: SALMONELLA TYPHI VI POLYSACCHARIDE (TY2 STRAIN) 25,0 ug HEPATITIS A VIRUS 160,0 antigen units
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AVENTIS PHARMA (PTY) LTD
Manufacturer:	SANOI PASTEUR, MARCY, L'ETOILE, FRANCE SANOI PASTEUR, VAL DE REUIL, FRANCE
Packer:	SANOI PASTEUR, MARCY, L'ETOILE, FRANCE SANOI PASTEUR, VAL DE REUIL, FRANCE
Laboratory:	SANOI PASTEUR, MARCY, L'ETOILE, FRANCE SANOI PASTEUR, VAL DE REUIL, FRANCE
FPRG/FPRR:	FPRG/FPRR:
Shelf-life:	36 months
Date of registration:	18 APRIL 2008
Registration number:	41/30.1/0025
Name of medicine:	VIVAXIM
Dosage form:	INJECTION
Active ingredients:	EACH 1,0 ml DOSE CONTAINS: SALMONELLA TYPHI VI POLYSACCHARIDE (TY2 STRAIN) 25,0 ug HEPATITIS A VIRUS 160,0 antigen units
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AVENTIS PHARMA (PTY) LTD
Manufacturer:	SANOI PASTEUR, MARCY, L'ETOILE, FRANCE SANOI PASTEUR, VAL DE REUIL, FRANCE
Packer:	SANOI PASTEUR, MARCY, L'ETOILE, FRANCE SANOI PASTEUR, VAL DE REUIL, FRANCE
Laboratory:	SANOI PASTEUR, MARCY, L'ETOILE, FRANCE SANOI PASTEUR, VAL DE REUIL, FRANCE
FPRG/FPRR:	FPRG/FPRR:
Shelf-life:	36 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/5.7.1/0085
Name of medicine:	DYNA-CETIRIZINE SYRUP
Dosage form:	SYRUP
Active ingredients:	EACH 1,0 ml SYRUP CONTAINS: CETIRIZINE DIHYDROCHLORIDE 1,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	PHARMA DYNAMICS (PTY) LTD
Manufacturer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND ACTAVIS hf, KOPAVOGUR, ICELAND TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA
Packer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND ACTAVIS hf, KOPAVOGUR, ICELAND TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG PHARMACEUTICAL ENTERPRISES, N'DABENI, PINELANDS, KZN
Laboratory:	FPRC: ACTAVIS hf, HAFNARFJORDUR, ICELAND ACTAVIS hf, KOPAVOGUR, ICELAND TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
Shelf-life:	FPRR: 36 months
Date of registration:	18 APRIL 2008
Registration number:	41/7.5/0093
Name of medicine:	SIMZOR 10
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SIMVASTATIN 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
Shelf-life:	FPRR: 24 months (Provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/7.5/0094
Name of medicine:	SIMZOR 20
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SIMVASTATIN 20,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	41/7.5/0095
Name of medicine:	SIMZOR 40
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SIMVASTATIN 40,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/7.5/0096
Name of medicine:	SIMZOR 80
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SIMVASTATIN 80,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
Shelf-life:	FPRC: 24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	41/7.1.3/0110
Name of medicine:	AURO-LISINAPRIL CO 10 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 10,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life:	FPRC: 24 months (provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/7.1.3/0111
Name of medicine:	AURO-LISINAPRIL CO 20 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 20,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life:	FPRR: AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA 24 months (provisional)
Date of registration:	18 APRIL 2008
Registration number:	41/7.1.3/0125
Name of medicine:	VEDIBLOK 3,125 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: CARVEDILOL 3,125 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ARROW PHARMA SOUTH AFRICA (PTY) LTD
Manufacturer:	ELPEN PHARMACEUTICAL CO INC, PIKERM ATTIKIS, GREECE
Packer:	ELPEN PHARMACEUTICAL CO INC, PIKERM ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA
Laboratory:	FPRC: ELPEN PHARMACEUTICAL CO INC, PIKERM ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
Shelf-life:	FPRR: ARROW PHARMA S.A., WOODMEAD, RSA 24 months
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/7.1.3/0126	Registration number: 41/7.1.3/0127
Name of medicine: VEDIBLOK 6,25 mg	Name of medicine: VEDIBLOK 12,5 mg
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: CARVEDILOL 6,25 mg	Active ingredients: EACH TABLET CONTAINS: CARVEDILOL 12,5 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: ARROW PHARMA SOUTH AFRICA (PTY) LTD	Applicant: ARROW PHARMA SOUTH AFRICA (PTY) LTD
Manufacturer: ELPEN PHARMACEUTICAL CO INC, PIKERMII ATTIKIS, GREECE	Manufacturer: ELPEN PHARMACEUTICAL CO INC, PIKERMII ATTIKIS, GREECE
Packer: ELPEN PHARMACEUTICAL CO INC, PIKERMII ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA	Packer: ELPEN PHARMACEUTICAL CO INC, PIKERMII ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA
Laboratory: FPRC: ELPEN PHARMACEUTICAL CO INC, PIKERMII ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG	Laboratory: FPRC: ELPEN PHARMACEUTICAL CO INC, PIKERMII ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRC: ARROW PHARMA S.A., WOODMEAD, RSA	FPRC: ARROW PHARMA S.A., WOODMEAD, RSA
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: 417.1.3/0126	Registration number: 41/16/0190
Name of medicine: VEDIBLOK 6,25 mg	Name of medicine: ANDOSEPT-CO
Dosage form: TABLET	Dosage form: SOLUTION
Active ingredients: EACH TABLET CONTAINS: CARVEDILOL 6,25 mg	Active ingredients: EACH 15,0 ml SOLUTION CONTAINS: BENZYLAMINE HYDROCHLORIDE 22,5 mg CHLORHEXIDINE GLUCONATE 18,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 8
Applicant: ARROW PHARMA SOUTH AFRICA (PTY) LTD	Applicant: PHARMACARE LIMITED
Manufacturer: ELPEN PHARMACEUTICAL CO INC, PIKERM ATTIKIS, GREECE	Manufacturer: PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
Packer: ELPEN PHARMACEUTICAL CO INC, PIKERM ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA	Packer: PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
Laboratory: FPRC: ELPEN PHARMACEUTICAL CO INC, PIKERM ATTIKIS, GREECE TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG	Laboratory: FPRC: SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRC: ARROW PHARMA S.A., WOODMEAD, RSA	FPRC/FPRR: PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON
Shelf-life: 24 months	Shelf-life: 24 months (Provisional)
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number: 417.3/0245	Registration number: 41/26/0246
Name of medicine: TRIPTAM 100 mg	Name of medicine: FLORACOR 50 mg/ml INJECTION
Dosage form: TABLET	Dosage form: INJECTION
Active ingredients: EACH TABLET CONTAINS: SUMATRIPTAM 100,0 mg	Active ingredients: EACH 1,0 ml SOLUTION CONTAINS: FLUOROURACIL 50,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: PHARMACARE LIMITED	Applicant: ACCORD HEALTHCARE (PTY) LTD
Manufacturer: GENPHARM PHARMACEUTICALS INC, ETOBICOKE, ONTARIO, CANADA PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON	Manufacturer: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA
Packer: GENPHARM PHARMACEUTICALS INC, ETOBICOKE, ONTARIO, CANADA PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON GERARD LABORATORIES, DUBLIN, IRELAND	Packer: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA
Laboratory: FPRC: GENPHARM PHARMACEUTICALS INC, ETOBICOKE, ONTARIO, CANADA GERARD LABORATORIES, DUBLIN, IRELAND SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH WEST UNIVERSITY, POTCHEFSTROOM	Laboratory: FPRC: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG ANALYTICON, TERENURE, KEMPTON PARK
FPRC/FPRR: PHARMACARE LTD, KORSTEN, PORT ELIZABETH ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON	FPRR: ACCORD HEALTHCARE, RIVONIA, RSA
Shelf-life: 24 months	Shelf-life: 24 months (Provisional)
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/20.2.8/0254	Registration number: 41/21.12/0268
Name of medicine: SEBIVO	Name of medicine: FINIDE
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: TELBIVUDINE 600,0 mg	Active ingredients: EACH TABLET CONTAINS: FINASTERIDE 1,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: NOVARTIS SOUTH AFRICA (PTY) LTD	Applicant: DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND	Manufacturer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND IVERS-LEE AG, BURGDORF, SWITZERLAND NOVARTIS S.A., SPARTAN, KEMPTON PARK	Packer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA DRA PHARMACEUTICALS, IRENE, CENTURION
Laboratory: FPRC:	Laboratory: FPRC:
FPRC/FPRR: NOVARTIS S.A., SPARTAN, KEMPTON PARK	FPRC/FPRR: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA
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:	41/20.2.8/0254
Name of medicine:	SEBIVO
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TELIVUDINE 600,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	NOVARTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND
Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND IVERS-LEE AG, BURGDORF, SWITZERLAND NOVARTIS S.A., SPARTAN, KEMPTON PARK
Laboratory:	FPRC: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMALYTICA S.A., LOCARNO, SWITZERLAND M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG
FPRC/FPRR:	NOVARTIS S.A., SPARTAN, KEMPTON PARK
Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008
Registration number:	41/21.12/0268
Name of medicine:	FINIDE
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: FINASTERIDE 1,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA DRA PHARMACEUTICALS, IRENE, CENTURION
Laboratory:	FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA
FPRR:	DR REDDY'S LABORATORIES, ROSEBANK, JOHANNESBURG
Shelf-life:	24 months
Date of registration:	18 APRIL 2008

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:
Shelf-life:	Shelf-life:
Date of registration:	Date of registration:
41/26/0304	41/26/0305
METHACOR 5 mg/ml	METHACOR 25 mg/ml
INJECTION	INJECTION
EACH 1.0 ml SOLUTION CONTAINS: METHOTREXATE 5,0 mg	EACH 1.0 ml SOLUTION CONTAINS: METHOTREXATE 25 mg
1, 2, 3, 4, 5, 6, 7, 8	1, 2, 3, 4, 5, 6, 7, 8
ACCORD HEALTHCARE (PTY) LTD	ACCORD HEALTHCARE (PTY) LTD
INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA	INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA
INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA	INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA
FPRC: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG	FPRC: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR: ACCORD HEALTHCARE, RIVONIA, RSA	FPRR: ACCORD HEALTHCARE, RIVONIA, RSA
12 months (Provisional)	12 months (Provisional)
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/26/0306	Registration number: 41/20.2.8/0330
Name of medicine: METHACOR 100 mg/ml	Name of medicine: RETLAM SOLUTION
Dosage form: INJECTION	Dosage form: SOLUTION
Active ingredients: EACH 1,0 ml SOLUTION CONTAINS: METHOTREXATE 100,0 mg	Active ingredients: EACH 1,0 ml SOLUTION CONTAINS: LAMIVUDINE 10,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: ACCORD HEALTHCARE (PTY) LTD	Applicant: ADCOCK INGRAM LIMITED
Manufacturer: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA	Manufacturer: ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON
Packer: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA	Packer: ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON
Laboratory: FPRC: INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG ANALYTICON, TERENCE, KEMPTON PARK	Laboratory: FPRC: ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON
FPRR: ACCORD HEALTHCARE, RIVONIA, RSA	FPRC/FPRR: ADCOCK INGRAM LTD, BRYANSTON, JOHANNESBURG
Shelf-life: 12 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:	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: FPRC/FPRR:	Laboratory: FPRC:
Shelf-life:	Shelf-life:
Date of registration:	Date of registration:
41/1.2/0394	41/1.2/0395
LEXAMIL 5	LEXAMIL 10
TABLET	TABLET
EACH TABLET CONTAINS: ESCITALOPRAM OXALATE EQUIVALENT TO ESCITALOPRAM 5,0 mg	EACH TABLET CONTAINS: ESCITALOPRAM OXALATE EQUIVALENT TO ESCITALOPRAM 10,0 mg
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
CIPLA MEDPRO (PTY) LTD	CIPLA MEDPRO (PTY) LTD
CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA	CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA
CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA	CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA
CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA	CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA
FPRC:	CIPLA MEDPRO, ROSENPARK, BELLVILLE
24 months	24 months
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15
Registration number:	41/1.2/0396
Name of medicine:	LEXAMIL 20
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: ESCITALOPRAM OXALATE EQUIVALENT TO ESCITALOPRAM 20,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	CIPLA MEDPRO (PTY) LTD
Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA
Packer:	CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA
Laboratory:	CIPLA LTD, KURKUMBH, PUNE, MAHASHTRA, INDIA
FPRR:	CIPLA MEDPRO, ROSENPARK, BELLVILLE
Shelf-life:	24 months
Date of registration:	18 APRIL 2008
Registration number:	41/3.2/0432
Name of medicine:	BONIRAN 70
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SODIUM ALENDRONATE EQUIVALENT TO ALENDRONIC ACID 70,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	RANBAXY (SA) (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, BOKSBUURG
FPRR:	RANBAXY (SA), CENTURION, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/3.2/0433	Registration number: 41/3.2/0434
Name of medicine: RAN-ALENDRONATE 10	Name of medicine: RAN-ALENDRONATE 70
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: SODIUM ALENDRONATE EQUIVALENT TO ALENDRONIC ACID 10,0 mg	Active ingredients: EACH TABLET CONTAINS: SODIUM ALENDRONATE EQUIVALENT TO ALENDRONIC ACID 70,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: RANBAXY (SA) (PTY) LTD	Applicant: RANBAXY (SA) (PTY) LTD
Manufacturer: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA	Manufacturer: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA
Packer: 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	Laboratory: FPRC: RANBAXY LABORATORIES LTD, PAONTA SAHIB, HIMACHAL PRADESH, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
Shelf-life: 24 months (Provisional)	Shelf-life: 24 months (Provisional)
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008
FPRC: RANBAXY (SA), CENTURION, RSA	FPRC: RANBAXY (SA), CENTURION, RSA

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:
FPRR:	FPRR:
Shelf-life:	Shelf-life:
Date of registration:	Date of registration:
41/26/0435	41/7.5/0549
ACCORD-CARBOPLATIN 10 mg/ml INJECTION	REDICOR 10
INJECTION	TABLET
EACH 1,0 ml SOLUTION CONTAINS: CARBOPLATIN 10,0 mg	EACH TABLET CONTAINS: SIMVASTATIN 10,0 mg
1, 2, 3, 4, 5, 6, 7, 8	1, 2, 3, 4, 5, 6, 7, 8
ACCORD HEALTHCARE (PTY) LTD	DR REDDY'S LABORATORIES (PTY) LTD
INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
INTAS PHARMACEUTICALS LTD, SANAND, AHMEDABAD, INDIA	DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA
ANALYTICON, TERENURE, KEMPTON PARK	RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
ACCORD HEALTHCARE, RIVONIA, RSA	DR REDDY'S LABORATORIES, ROSEBANK, JOHANNESBURG
24 months (Provisional)	24 months (Provisional)
18 APRIL 2008	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/7.5/0550	Registration number: 41/7.5/0551
Name of medicine: REDICOR 20	Name of medicine: REDICOR 40
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: SIMVASTATIN 20,0 mg	Active ingredients: EACH TABLET CONTAINS: SIMVASTATIN 40,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: DR REDDY'S LABORATORIES (PTY) LTD	Applicant: DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Packer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory: FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA	Laboratory: FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA
Shelf-life: 24 months (Provisional)	Shelf-life: 24 months (Provisional)
Date of registration: 18 APRIL 2008	Date of registration: 18 APRIL 2008
FPRR: DR REDDY'S LABORATORIES, ROSEBANK, JOHANNESBURG	FPRR: DR REDDY'S LABORATORIES, ROSEBANK, JOHANNESBURG
POTCHEFSTROOM	POTCHEFSTROOM

MRF 15

Registration number: 41/7.5/0552
 Name of medicine: REDICOR 80
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 SIMVASTATIN 80,0 mg

Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
 Applicant: DR REDDY'S LABORATORIES (PTY) LTD
 Manufacturer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
 Packer: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
 Laboratory: FPRC: DR REDDY'S LABORATORIES LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
 FPRC: DR REDDY'S LABORATORIES, ROSEBANK, JOHANNESBURG
 Shelf-life: 24 months (Provisional)
 Date of registration: 18 APRIL 2008

MRF 15

Registration number: 41/1.2/0584
 Name of medicine: ZOLID 50 mg
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 SERTRALINE HYDROCHLORIDE EQUIVALENT TO SERTRALINE 50,0 mg

Conditions of registration: 1, 2, 3, 4, 5, 6, 7
 Applicant: AUROBINDO PHARMA (PTY) LTD
 Manufacturer: AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
 Packer: AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
 Laboratory: FPRC: AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
 FPRC: AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA
 Shelf-life: 24 months
 Date of registration: 18 APRIL 2008

MRF 15	MRF 15		
Registration number:	411/1.2/0585	Registration number:	411/1.1.3/0673
Name of medicine:	ZOLID 100 mg	Name of medicine:	CIPLA-PERINDOPRIL 2 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SERTRALINE HYDROCHLORIDE EQUIVALENT TO SERTRALINE 100.0 mg	Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL ERBUMINE 2.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AUROBINDO PHARMA (PTY) LTD	Applicant:	CIPLA LIFE SCIENCES (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA
Packer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Packer:	CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Laboratory:	FPRC: CIPLA LTD, (UNIT IV), SALCETTE, GOA
FPRR:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA	FPRR:	CIPLA LIFE SCIENCES, ROSEN PARK, BELLVILLE
Shelf-life:	24 months	Shelf-life:	24 months (Provisional)
Date of registration:	18 APRIL 2008	Date of registration:	18 APRIL 2008

MRF 15	MRF 15
Registration number: 41/7.1.3/0674	Registration number: 41/7.1.3/0675
Name of medicine: CIPLA-PERINDOPRIL 4 mg	Name of medicine: CIPLA-PERINDOPRIL 8 mg
Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH TABLET CONTAINS: PERINDOPRIL ERBUMINE 4,0 mg	Active ingredients: EACH TABLET CONTAINS: PERINDOPRIL ERBUMINE 8,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: CIPLA LIFE SCIENCES (PTY) LTD	Applicant: CIPLA LIFE SCIENCES (PTY) LTD
Manufacturer: CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA	Manufacturer: CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA
Packer: CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA	Packer: CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA
Laboratory: FPRC: CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA	Laboratory: FPRC: CIPLA LTD, (UNIT IV), SALCETTE, GOA, INDIA
FPRR: CIPLA LIFE SCIENCES, ROSEN PARK, BELLVILLE	FPRR: CIPLA LIFE SCIENCES, ROSEN PARK, BELLVILLE
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:	41/20.1.2/0687
Name of medicine:	YOMAX 250 mg
Dosage form:	CAPSULE
Active ingredients:	EACH CAPSULE CONTAINS: AMOXICILLIN TRIHYDRATE EQUIVALENT TO AMOXICILLIN 250.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, UNIT XII, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, UNIT XII RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, UNIT XII, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life:	FPRC: AUROBINDO PHARMA LTD, ROSEBANK, JOHANNESBURG 24 months
Date of registration:	18 APRIL 2008
Registration number:	41/20.1.2/0688
Name of medicine:	YOMAX 500 mg
Dosage form:	CAPSULE
Active ingredients:	EACH CAPSULE CONTAINS: AMOXICILLIN TRIHYDRATE EQUIVALENT TO AMOXICILLIN 500.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, UNIT XII, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, UNIT XII, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	FPRC: AUROBINDO PHARMA LTD, UNIT XII, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf-life:	FPRC: AUROBINDO PHARMA LTD, ROSEBANK, JOHANNESBURG 24 months
Date of registration:	18 APRIL 2008

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
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

Registration number: 41/20.1.1/0752
 Name of medicine: ZINOXIME 250 mg
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 CEFUROXIME AXETIL EQUIVALENT TO
 CEFUROXIME 250,0 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
 FPRC:
 Shelf-life: 24 months (Provisional)
 Date of registration: 18 APRIL 2008

MRF 15

Registration number: 41/20.1.1/0753
 Name of medicine: ZINOXIME 500 mg
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 CEFUROXIME AXETIL EQUIVALENT TO
 CEFUROXIME 500,0 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
 FPRC:
 Shelf-life: 24 months (Provisional)
 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:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
FPRC:	FPRC:
FPRR:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG
Shelf-life:	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:	AUROBINDO PHARMA LTD, UNIT III, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
FPRC:	FPRC:
FPRR:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG
Shelf-life:	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/a 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
Name of medicine:	SANDOZ CEFTRIAXONE 0,5 g
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: CEFTRIAXONE SODIUM EQUIVALENT TO CEFTRIAXONE 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:	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRR:	SANDOZ S.A., SPARTAN, KEMPTON PARK
Shelf-life:	36 months
Date of registration:	18 APRIL 2008
Registration number:	41/20.1.1/0867
Name of medicine:	SANDOZ CEFTRIAXONE 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:	SANDOZ GmbH, KUNDL, AUSTRIA NOVARTIS SA, SPARTAN, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRR:	SANDOZ S.A., SPARTAN, KEMPTON PARK
Shelf-life:	36 months
Date of registration:	18 APRIL 2008

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

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
 FPRR: SANDOZ S.A., SPARTAN, KEMPTON PARK
 Shelf-life: 36 months
 Date of registration: 18 APRIL 2008

MRF 15

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
 FPRR: 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

FPRC: 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

FPRC: ORCHID PHARMACEUTICALS SA,
POTCHEFSTROOM

Shelf-life: 24 months

Date of registration: 18 APRIL 2008

MRF 15	MRF 15		
Registration number:	41/7.1.3/1050	Registration number:	41/7.1.3/1051
Name of medicine:	LISINOZIDE 10 mg	Name of medicine:	LISINOZIDE 20 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 10,0 mg HYDROCHLOROTHIAZIDE 12,5 mg	Active ingredients:	EACH TABLET CONTAINS: LISINAPRIL DIHYDRATE EQUIVALENT TO LISINAPRIL 20,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
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:	Laboratory:	FPRC:
FPRC:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA	FPRC:	AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG, RSA
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:	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:	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
FPRR:	SANDOZ S.A., SPARTAN, KEMPTON PARK	FPRR:	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: A38/2.6.5/0624	Registration number: 37/28/0521
Name of medicine: MOXOTENS 0,4 mg	Name of medicine: AXIM READICAT 2 %
Dosage form: TABLET	Dosage form: SUSPENSION
Active ingredients: EACH TABLET CONTAINS: MOXONIDINE 0,4 mg	Active ingredients: EACH 100,0 ml SUSPENSION CONTAINS: BARIUM SULPHATE 2,0940 g
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6
Applicant: HEXAL PHARMA (S.A.) (PTY) LTD	Applicant: AXIM PHARMACEUTICALS (PTY) LTD
Manufacturer: PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY	Manufacturer: E-Z-EM CANADA INC, ANJOU, QUEBEC, CANADA
Packer: PERRIGO ISRAEL PHARMACEUTICALS LTD, YERUHAM, ISRAEL CHANELLE PHARMACEUTICALS MANUFACTURING, GALWAY, IRELAND SALUTAS PHARMA GmbH, BARLEBEN, GERMANY DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, JOHANNESBURG, RSA	Packer: E-Z-EM CANADA INC, ANJOU, QUEBEC, CANADA
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, TERENCE, KEMPTON PARK, RSA	Laboratory: FPRC: E-Z-EM CANADA INC, ANJOU, QUEBEC, CANADA SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRR: HEXAL PHARMA, PINETOWN, KZN, RSA	FPRR: AXIM PHARMACEUTICALS, MIDRAND, JOHANNESBURG
Shelf-life: 24 months	Shelf-life: 36 months
Date of registration: 18 APRIL 2008	Date of registration: 13 JUNE 2008

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

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

MRF 15	MRF 15		
Registration number:	A40/5.10/0213	Registration number:	A40/21.5.1/0224
Name of medicine:	ADCO-NETRIN 2 mg/ml	Name of medicine:	SPEC-BUDESONIDE 100
Dosage form:	INJECTION	Dosage form:	NASAL SPRAY
Active ingredients:	EACH 2.0 ml SOLUTION CONTAINS: ONDANSETRON 4.0 mg	Active ingredients:	EACH METERED DOSE CONTAINS: BUDESONIDE 100.0 ug
Conditions of registration:	1, 2, 3, 4, 5, 6	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ADCOCK INGRAM LIMITED	Applicant:	SPECPHARM (PTY) LTD
Manufacturer:	PHARMAMED PARENTERALS LTD, ZEJTUN, MALTA FAMAR S.A., P. FALIRO, GREECE	Manufacturer:	MIPHARM S.p.A, MILAN, ITALY
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	Packer:	MIPHARM S.p.A, MILAN, ITALY
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	Laboratory:	FPRC: MIPHARM S.p.A, MILAN, ITALY ANALYTICON, TERENURE, KEMPTON PARK
FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE, AEROTON, JOHANNESBURG	FPRR:	SPECPHARM, HALFWAY HOUSE, MIDRAND
FPRR:	ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON ADCOCK INGRAM LTD, BRYANSTON, JOHANNESBURG	Shelf-life:	24 months
Shelf-life:	24 months	Date of registration:	13 JUNE 2008
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
Packer: B BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY B BRAUN MELSUNGEN PRODUCTION PHARMA PFIFFEWIESEN, MELSUNGEN, GERMANY	ASTRAZENECA UK LTD, MACCLESFIELD, CHESHIRE, UK ASTRAZENECA PHARMACEUTICALS, ALRODE, ALBERTON
Laboratory: FPRC: B BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY M&L LABORATORY SERVICES, ORMONDE, JOHANNESBURG COSI PHARMACEUTICALS, INDUSTRIA WEST, JOHANNESBURG CONSULTING CHEMICAL LABORATORIES, BOKSBURG	Laboratory: FPRC: ASTRAZENECA UK LTD, MACCLESFIELD, CHESHIRE, UK ASTRAZENECA GmbH, PLANKSTADT, GERMANY ANALYTICON, TERENURE, KEMPTON PARK CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR: 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