
GENERAL NOTICE ALGEMENE KENNISGEWING

NOTICE 915 OF 2012

MEDICINES CONTROL COUNCIL

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	MRF 15	MRF 15
Registration number: 41/10.2.1/0765	Registration number: 42/21.12/0461	Registration number: 44/10.2.2/0203
Name of medicine: CO-IPRASAL SOLUTION	Name of medicine: FINIDE 5 TABLETS	Name of medicine: SINTRINE 10
Dosage form: SOLUTION FOR INHALATION	Dosage form: TABLET	Dosage form: TABLET
Active ingredients: EACH 2.5 ml VIAL CONTAINS: IPRATROPIUM BROMIDE 0.5 mg SALBUTAMOL SULPHATE 3.0 mg	Active ingredients: EACH TABLET CONTAINS: FINASTERIDE 5.0 mg	Active ingredients: EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10.0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: ARROW PHARMA (PTY) LTD	Applicant: ZYDUS HEALTHCARE (PTY) LTD	Applicant: SANDOZ SA (PTY) LTD
Manufacturer: LABORATOIRE UNITHER, AMIENS, FRANCE	Manufacturer: ZYDUS CADILA HEALTHCARE LIMITED, SANAND, AHMEDABAD, INDIA	Manufacturer: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY
Packer: LABORATOIRE UNITHER, AMIENS, FRANCE DIVPHARM MANUFACTURING AND PACKAGING (PTY) LTD, LONGDALE, INDUSTRIAL, RSA TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA	Packer: ZYDUS CADILA HEALTHCARE LIMITED, SANAND, AHMEDABAD, INDIA	Packer: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Laboratory: FPRC: LABORATOIRE UNITHER, AMIENS, FRANCE TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory: FPRC: ZYDUS CADILA HEALTHCARE LIMITED, SANAND, AHMEDABAD, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory: FPRC: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
FPRR: ARROW PHARMA (PTY) LTD, WOODMEAD, SANDTON, RSA	FPRR: ZYDUS HEALTHCARE (PTY) LTD, POTCHEFSTROOM, RSA	FPRR: SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Shelf-life: 36 months	Shelf-life: 24 months	Shelf-life: 24 months (Provisional)
Date of registration: 18 MAY 2012	Date of registration: 18 MAY 2012	Date of registration: 18 MAY 2012

MRF 15

Registration number:	44/10.2.2/0204
Name of medicine:	SANDOZ MONTELUKAST 10
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	SANDOZ SA (PTY) LTD
Manufacturer:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY
Packer:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Laboratory: FPRC:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
FPRR:	SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	18 MAY 2012

MRF15

Registration number:	44/10.2.2/0485
Name of medicine:	SINTRINE 4
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	SANDOZ SA (PTY) LTD
Manufacturer:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY
Packer:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Laboratory: FPRC:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
FPRR:	SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	18 MAY 2012

MRF 15

Registration number:	44/10.2.2/0486
Name of medicine:	SINTRINE 5
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANDOZ SA (PTY) LTD
Manufacturer:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY
Packer:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Laboratory: FPRC:	SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
FPRR:	SANDOZ SA, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months
Date of registration:	18 MAY 2012

MRF 15	MRF 15	MRF 15
Registration number: 44/10.2.2/0487 Name of medicine: SANDOZ MONTELUKAST 4 Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: SANDOZ SA (PTY) LTD Manufacturer: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY Packer: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA Laboratory: FPRC: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA FPRR: SANDOZ SA, SPARTAN, KEMPTON PARK, RSA Shelf-life: 24 months Date of registration: 18 MAY 2012	Registration number: 44/10.2.2/0488 Name of medicine: SANDOZ MONTELUKAST 5 Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: SANDOZ SA (PTY) LTD Manufacturer: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY Packer: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA Laboratory: FPRC: SANDOZ ILAC SANAYI VE TICARET A.S., GEBZE-KOCAELI, TURKEY CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA SANDOZ SA, SPARTAN, KEMPTON PARK, RSA FPRR: SANDOZ SA, SPARTAN, KEMPTON PARK, RSA Shelf-life: 24 months Date of registration: 18 MAY 2012	Registration number: 38/21.12/0173 Name of medicine: ASPEN FLUTAMIDE 250 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: FLUTAMIDE 250,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: PHARMACARE LIMITED Manufacturer: GENERICS (UK) LIMITED, POTTERS BAR, HERTFORDSHIRE, UNITED KINGDOM GERARD LABORATORIES, GRANGE ROAD, DUBLIN, IRELAND PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA Packer: GENERICS (UK) LIMITED, POTTERS BAR, HERTFORDSHIRE, UNITED KINGDOM GERARD LABORATORIES, GRANGE ROAD, DUBLIN, IRELAND PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA Laboratory: FPRC: GENERICS (UK) LIMITED, POTTERS BAR, HERTFORDSHIRE, UNITED KINGDOM GERARD LABORATORIES, GRANGE ROAD, DUBLIN, IRELAND SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA FPRC/FPRR: PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA FPRR: PHARMACARE LTD, WOODMEAD, SANDTON, RSA Shelf-life: 24 months Date of registration: 07 JUNE 2012

MRF 15		MRF 15		MRF 15	
Registration number:	A38/7.5/0729	Registration number:	A40/15.1/0048	Registration number:	A40/2.9/0355
Name of medicine:	OMACOR	Name of medicine:	CEPROLEN EYE DROPS	Name of medicine:	SABAX TRAMADOL 100 mg/2 ml
Dosage form:	CAPSULE	Dosage form:	EYE DROP SOLUTION	Dosage form:	INJECTION
Active ingredients:	EACH CAPSULE CONTAINS: Eicosapent (EPA) ethyl esters 46,0 % Docosahexa (DHA) ethyl esters 38,0 %	Active ingredients:	EACH 100,0 ml SOLUTION CONTAINS: Ciprofloxacin Hydrochloride equivalent to Ciprofloxacin 0,3 g	Active ingredients:	EACH 2,0 ml CONTAINS: TRAMADOL HYDROCHLORIDE 100,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ABBOTT LABORATORIES SA (PTY) LTD	Applicant:	MEDIVISION (PTY) LTD	Applicant:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD
Manufacturer:	CARDINAL HEALTH UK, SWINDON, WILTSHIRE, UNITED KINGDOM	Manufacturer:	INDOCO REMEDIES, VERNA INDUSTRIAL ESTATE, GOA, INDIA	Manufacturer:	PHARMA-Q (PTY) LTD, INDUSTRIAL, JOHANNESBURG, RSA
Packer:	CARDINAL HEALTH UK, SWINDON, WILTSHIRE, UNITED KINGDOM GMPACK, HADSUND, DENMARK	Packer:	INDOCO REMEDIES, VERNA INDUSTRIAL ESTATE, GOA, INDIA	Packer:	PHARMA-Q (PTY) LTD, INDUSTRIAL, JOHANNESBURG, RSA
Laboratory:	FPRC: CARDINAL HEALTH UK, SWINDON, WILTSHIRE, UNITED KINGDOM PRONOVA BIO CARE A.S, LYSAKER, NORWAY SOLVAY PHARMACEUTICALS GmbH, NEUSTADT, GERMANY	Laboratory:	FPRC: INDOCO REMEDIES, VERNA INDUSTRIAL ESTATE, GOA, INDIA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory:	FPRC: PHARMA-Q (PTY) LTD, INDUSTRIAL, JOHANNESBURG, RSA
FPRC/FPRR:	ABBOTT LABORATORIES (PTY) LTD, CONSTANTIA KLOOF, RSA	FPRR:	MEDIVISION (PTY) LTD, KRAMERVILLE, SANDTON, RSA	FPRR:	PHARMA-Q (PTY) LTD, INDUSTRIAL, JOHANNESBURG, RSA ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA
Shelf-life:	36 months	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15

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MRF 15

Registration number: A40/21.5.1/0748	Registration number: 42/2.2/0045	Registration number: 42/2.2/0046
Name of medicine: FRESENIUS DEXAMETHASONE 4 mg/1 ml	Name of medicine: MIDAZOLAM SAFELINE 15 mg	Name of medicine: MIDAZOLAM SAFELINE 50 mg
Dosage form: INJECTION	Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH 1,0 ml CONTAINS: Dexamethasone sodium phosphate equivalent to Dexamethasone phosphate 4,0 mg	Active ingredients: EACH 3,0 ml AMPOULE CONTAINS: MIDAZOLAM HYDROCHLORIDE 15,0 mg	Active ingredients: EACH 3,0 ml AMPOULE CONTAINS: MIDAZOLAM HYDROCHLORIDE 50,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: BODENE (PTY) LIMITED TRADING AS INTRAMED	Applicant: SAFELINE PHARMACEUTICALS (PTY) LTD	Applicant: SAFELINE PHARMACEUTICALS (PTY) LTD
Manufacturer: BODENE (PTY) LIMITED TRADING AS INTRAMED, KORSTEN, PORT ELIZABETH, RSA	Manufacturer: DEMO S.A., PHARMACEUTICAL INDUSTRY, ATHENS, GREECE	Manufacturer: DEMO S.A., PHARMACEUTICAL INDUSTRY, ATHENS, GREECE
Packer: BODENE (PTY) LIMITED TRADING AS INTRAMED, KORSTEN, PORT ELIZABETH, RSA	Packer: DEMO S.A., PHARMACEUTICAL INDUSTRY, ATHENS, GREECE	Packer: DEMO S.A., PHARMACEUTICAL INDUSTRY, ATHENS, GREECE
Laboratory: FPRC: BODENE (PTY) LIMITED TRADING AS INTRAMED, KORSTEN, PORT ELIZABETH, RSA	Laboratory: FPRC: DEMO S.A., PHARMACEUTICAL INDUSTRY, ATHENS, GREECE INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	Laboratory: FPRC: DEMO S.A., PHARMACEUTICAL INDUSTRY, ATHENS, GREECE INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA
FPRR: BODENE (PTY) LIMITED TRADING AS INTRAMED, KORSTEN, PORT ELIZABETH, RSA	FPRR: SAFELINE PHARMACEUTICALS (PTY) LTD, FLORIDA, RSA	FPRR: SAFELINE PHARMACEUTICALS (PTY) LTD, FLORIDA, RSA
Shelf-life: 24 months (Provisional)	Shelf-life: 36 months	Shelf-life: 36 months
Date of registration: 07 JUNE 2012	Date of registration: 07 JUNE 2012	Date of registration: 07 JUNE 2012

MRF-15	F 15									
	Registration number:	Name of medicine:	Dosage form:	Active ingredients:	Conditions of registration:	Applicant:	Manufacturer:	Packer:	Laboratory:	FPRC:
MRF-15	42/20.1/1/0511	ASPEN TEICOPLANIN 200	POWDER FOR SOLUTION FOR INJECTION	EACH 3.2 ml CONTAINS: TEICOPLANIN 200.0 mg	1, 2, 3, 4, 5, 6, 7, 8	PHARMACARE LIMITED	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA M & L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	FPRC/FPRR: FPRR: Shelf-life: Date of registration:
MRF-15	42/20.1/1/0512	ASPEN TEICOPLANIN 400	POWDER FOR SOLUTION FOR INJECTION	EACH 3.2 ml CONTAINS: TEICOPLANIN 400.0 mg	1, 2, 3, 4, 5, 6, 7, 8	PHARMACARE LIMITED	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA M & L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	FPRC/FPRR: FPRR: Shelf-life: Date of registration:
MRF-15	42/34/0513	ASPEN DILUENT FOR TEICOPLANIN	SOLUTION FOR INJECTION	EACH 3.2 ml CONTAINS: WATER FOR INJECTION 3.2 ml	1, 2, 3, 4, 5, 6, 7, 8	PHARMACARE LIMITED	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	LABIANA PHARMACEUTICALS, S.L.U, BARBERA DEL VALLES, BARCELONA, SPAIN PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH- WEST UNIVERSITY, POTCHEFSTROOM, RSA M & L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	FPRC/FPRR: FPRR: Shelf-life: Date of registration:

MRF 15	MRF 15	MRF 15
Registration number: 42/24/0518	Registration number: 42/20.1.1/0617	Registration number: 42/20.1.1/0618
Name of medicine: RINGERFUNDIN	Name of medicine: AURO CEFUROXIME 250 mg	Name of medicine: AURO CEFUROXIME 750 mg
Dosage form: SOLUTION FOR INFUSION	Dosage form: INJECTION	Dosage form: INJECTION
Active ingredients: EACH PLASTIC INFUSION BAG CONTAINS: Calcium Chloride Dihydrate 0,37 g Magnesium Chloride 0,20 g Malic Acid 0,67 g Potassium Chloride 0,30 g Sodium Acetate 3,27 g Sodium Chloride 6,80 g	Active ingredients: EACH VIAL CONTAINS: CEFUROXIME SODIUM EQUIVALENT TO CEFUROXIME 250,0 mg	Active ingredients: EACH VIAL CONTAINS: CEFUROXIME SODIUM EQUIVALENT TO CEFUROXIME 750,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8
Applicant: B BRAUN MEDICAL (PTY) LTD	Applicant: AUROBINDO PHARMA (PTY) LTD	Applicant: AUROBINDO PHARMA (PTY) LTD
Manufacturer: B BRAUN MELSUNGEN AG, GERMANY PRODUCTION PHARMA PIEFFEWIESEN, GERMANY B BRAUN MEDICAL S.A., BARCELONA, SPAIN B BRAUN MEDICAL S.A., SWITZERLAND	Manufacturer: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA	Manufacturer: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA
Packer: B BRAUN MELSUNGEN AG, GERMANY PRODUCTION PHARMA PIEFFEWIESEN, GERMANY B BRAUN MEDICAL S.A., BARCELONA, SPAIN B BRAUN MEDICAL S.A., SWITZERLAND	Packer: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA	Packer: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA
Laboratory: FPRC: B BRAUN MELSUNGEN AG, GERMANY M & L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA	Laboratory: FPRC: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA
FPRR: B BRAUN MEDICAL (PTY) LTD, FOURWAYS, JOHANNESBURG, RSA	FPRR: AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRR: AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA
Shelf-life: 24 months (Provisional)	Shelf-life: 24 months (Provisional)	Shelf-life: 24 months (Provisional)
Date of registration: 07 JUNE 2012	Date of registration: 07 JUNE 2012	Date of registration: 07 JUNE 2012

MRF 15	MRF 15	MRF 15
Registration number: 42/20.1.1/0619 Name of medicine: AURO CEFUROXIME 1500 mg Dosage form: INJECTION Active ingredients: EACH VIAL CONTAINS: CEFUROXIME SODIUM EQUIVALENT TO CEFUROXIME 1500,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: AUROBINDO PHARMA (PTY) LTD Manufacturer: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA Packer: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA Laboratory: FPRC: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA FPRR: AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012	Registration number: 42/26/0739 Name of medicine: HYCAMTIN 0,25 mg Dosage form: CAPSULE Active ingredients: EACH CAPSULE CONTAINS: Topotecan Hydrochloride equivalent to Topotecan 0,25 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: GLAXOSMITHKLINE SA (PTY) LTD Manufacturer: GLAXOSMITHKLINE MANUFACTURING S.p.A, PARMA, ITALY Packer: GLAXOSMITHKLINE MANUFACTURING S.p.A, PARMA, ITALY Laboratory: FPRC: GLAXOSMITHKLINE MANUFACTURING S.p.A, PARMA, ITALY FPRR: GLAXOSMITHKLINE SA (PTY) LTD, EPPING, CAPE TOWN, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012	Registration number: 42/26/0740 Name of medicine: HYCAMTIN 1,0 mg Dosage form: CAPSULE Active ingredients: EACH CAPSULE CONTAINS: Topotecan Hydrochloride equivalent to Topotecan 1,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: GLAXOSMITHKLINE SA (PTY) LTD Manufacturer: GLAXOSMITHKLINE MANUFACTURING S.p.A, PARMA, ITALY Packer: GLAXOSMITHKLINE MANUFACTURING S.p.A, PARMA, ITALY Laboratory: FPRC: GLAXOSMITHKLINE MANUFACTURING S.p.A, PARMA, ITALY FPRR: GLAXOSMITHKLINE SA (PTY) LTD, EPPING, CAPE TOWN, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012

MRF 15		MRF 15		F 15	
Registration number:	42/2.6.5/0792	Registration number:	42/2.6.5/0793	Registration number:	42/2.6.5/0794
Name of medicine:	RISNIA 0,5 mg	Name of medicine:	RISNIA 1 mg	Name of medicine:	RISNIA 2 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 0,5 mg	Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 1,0 mg	Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 2,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	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:	CIPLA MEDPRO (PTY) LTD	Applicant:	CIPLA MEDPRO (PTY) LTD
Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA
Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA
Laboratory: FPRC	CIPLA LTD, KURKUMBH, PUNE, INDIA	Laboratory: FPRC:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Laboratory: FPRC:	CIPLA LTD, KURKUMBH, PUNE, INDIA
FPRR:	CIPLA MEDPRO (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA	FPRR:	CIPLA MEDPRO (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA	FPRR:	CIPLA MEDPRO (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15		MRF 15		MRF 15	
Registration number:	42/2.6.5/0795	Registration number:	42/2.6.5/0796	Registration number:	42/2.6.5/0797
Name of medicine:	RISNIA 3 mg	Name of medicine:	RISNIA 4 mg	Name of medicine:	RISNIA 6 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 3,0 mg	Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 4,0 mg	Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 6,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	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:	CIPLA MEDPRO (PTY) LTD	Applicant:	CIPLA MEDPRO (PTY) LTD
Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA
Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA
Laboratory:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Laboratory:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Laboratory:	CIPLA LTD, KURKUMBH, PUNE, INDIA
FPRC:		FPRC:		FPRC:	
FPRR:	CIPLA MEDPRO (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA	FPRR:	CIPLA MEDPRO (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA	FPRR:	CIPLA MEDPRO (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15		MRF 15		MRF 15	
Registration number:	42/2.6.5/0798	Registration number:	42/2.6.5/0799	Registration number:	42/2.6.5/0800
Name of medicine:	CIPLA RISPERIDONE 0,5 mg	Name of medicine:	CIPLA RISPERIDONE 1 mg	Name of medicine:	CIPLA RISPERIDONE 2 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 0,5 mg	Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 1,0 mg	Active ingredients:	EACH TABLET CONTAINS: RISPERIDONE 2,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	CIPLA LIFE SCIENCES (PTY) LTD	Applicant:	CIPLA LIFE SCIENCES (PTY) LTD	Applicant:	CIPLA LIFE SCIENCES (PTY) LTD
Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Manufacturer:	CIPLA LTD, KURKUMBH, PUNE, INDIA
Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA	Packer:	CIPLA LTD, KURKUMBH, PUNE, INDIA
Laboratory:	FPRC:	Laboratory:	FPRC:	Laboratory:	FPRC
	FPRR:		FPRR:		FPRR:
	CIPLA LIFE SCIENCES (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA		CIPLA LIFE SCIENCES (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA		CIPLA LIFE SCIENCES (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15	Registration number: 42/2.6.5/0801 Name of medicine: CIPLA RISPERIDONE 3 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: RISPERIDONE 3,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: CIPLA LIFE SCIENCES (PTY) LTD Manufacturer: CIPLA LTD, KURKUMBH, PUNE, INDIA Packer: CIPLA LTD, KURKUMBH, PUNE, INDIA Laboratory: FPRC: CIPLA LTD, KURKUMBH, PUNE, INDIA FPRR: CIPLA LIFE SCIENCES (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA Shelf-life: 24 months Date of registration: 07 JUNE 2012	MRF 15	Registration number: 42/2.6.5/0802 Name of medicine: CIPLA RISPERIDONE 4 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: RISPERIDONE 4,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: CIPLA LIFE SCIENCES (PTY) LTD Manufacturer: CIPLA LTD, KURKUMBH, PUNE, INDIA Packer: CIPLA LTD, KURKUMBH, PUNE, INDIA Laboratory: FPRC: CIPLA LTD, KURKUMBH, PUNE, INDIA FPRR: CIPLA LIFE SCIENCES (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA Shelf-life: 24 months Date of registration: 07 JUNE 2012	MRF 15	Registration number: 42/2.6.5/0803 Name of medicine: CIPLA RISPERIDONE 6 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: RISPERIDONE 6,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: CIPLA LIFE SCIENCES (PTY) LTD Manufacturer: CIPLA LTD, KURKUMBH, PUNE, INDIA Packer: CIPLA LTD, KURKUMBH, PUNE, INDIA Laboratory: FPRC: CIPLA LTD, KURKUMBH, PUNE, INDIA FPRR: CIPLA LIFE SCIENCES (PTY) LTD, ROSEN HEIGHTS, BELLVILLE, RSA Shelf-life: 24 months Date of registration: 07 JUNE 2012
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MRF 15		MRF 15		MRF 15	
Registration number:	42/20.1.1/0888	Registration number:	42/20.1.1/0889	Registration number:	42/34/1056
Name of medicine:	SPEC TEICOPLANIN 200	Name of medicine:	SPEC TEICOPLANIN 400	Name of medicine:	SPEC TEICOPLANIN SOLVENT
Dosage form:	POWDER FOR SOLUTION FOR INJECTION	Dosage form:	POWDER FOR SOLUTION FOR INJECTION	Dosage form:	LIQUID
Active ingredients:	EACH VIAL CONTAINS: TEICOPLANIN 200,0 mg	Active ingredients:	EACH VIAL CONTAINS: TEICOPLANIN 400,0 mg	Active ingredients:	EACH AMPOULE CONTAINS: WATER FOR INJECTION 3,0 ml
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SPECPHARM (PTY) LTD	Applicant:	SPECPHARM (PTY) LTD	Applicant:	SPECPHARM (PTY) LTD
Manufacturer:	SIRTON, VILLA GUARDIA, ITALY	Manufacturer:	SIRTON, VILLA GUARDIA, ITALY	Manufacturer:	SIRTON, VILLA GUARDIA, ITALY
Packer:	SIRTON, VILLA GUARDIA, ITALY	Packer:	SIRTON, VILLA GUARDIA, ITALY	Packer:	SIRTON, VILLA GUARDIA, ITALY
Laboratory:	FPRC: SIRTON, VILLA GUARDIA, ITALY CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory:	FPRC: SIRTON, VILLA GUARDIA, ITALY CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory:	FPRC: SIRTON, VILLA GUARDIA, ITALY CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	FPRR:	SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	FPRR:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Shelf-life:	24 months (Provisional)*	Shelf-life:	24 months (Provisional)*	Shelf-life:	24 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

*A provisional shelf-life of 24 months is approved for this product manufactured by Sirton with API manufactured by Zhejiang Medicine. A shelf-life of 24 hours at 2-8 °C is approved for the reconstituted and diluted product.

MRF 15		MRF 15		MRF 15	
Registration number:	43/5.3/0251	Registration number:	43/5.3/0252	Registration number:	43/5.3/0253
Name of medicine:	DONESPES 5	Name of medicine:	DONESPES 10	Name of medicine:	DONERIN 5
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: DONEPEZIL HCl 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: DONEPEZIL HCl 10,0 mg	Active ingredients:	EACH TABLET CONTAINS: DONEPEZIL HCl 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	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	LASARA TRADERS (PTY) LTD	Applicant:	LASARA TRADERS (PTY) LTD	Applicant:	LASARA TRADERS (PTY) LTD
Manufacturer:	SPECIFAR S.A, VARVARA, ATHENS, GREECE	Manufacturer:	SPECIFAR S.A, VARVARA, ATHENS, GREECE	Manufacturer:	SPECIFAR S.A, VARVARA, ATHENS, GREECE
Packer:	SPECIFAR S.A, VARVARA, ATHENS, GREECE SPECPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA	Packer:	SPECIFAR S.A, VARVARA, ATHENS, GREECE SPECPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA	Packer:	SPECIFAR S.A, VARVARA, ATHENS, GREECE SPECPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA
Laboratory: FPRC:	SPECIFAR S.A, VARVARA, ATHENS, GREECE SPECPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA	Laboratory: FPRC:	SPECIFAR S.A, VARVARA, ATHENS, GREECE SPECPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA	Laboratory: FPRC:	SPECIFAR S.A, VARVARA, ATHENS, GREECE SPECPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA
FPRR:	LASARA TRADERS (PTY) LTD, MORELETA PARK, PRETORIA, RSA	FPRR:	LASARA TRADERS (PTY) LTD, MORELETA PARK, PRETORIA, RSA	FPRR:	LASARA TRADERS (PTY) LTD, MORELETA PARK, PRETORIA, RSA
Shelf-life:	24 months (Provisional)*	Shelf-life:	24 months (Provisional)*	Shelf-life:	24 months (Provisional)*
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

*A provisional shelf-life of 24 months is approved for this product manufactured with API from Jubilant Oranosys and Ranbaxy Laboratories Limited

MRF 15		MRF 15		MRF 15	
Registration number:	43/5.3/0254	Registration number:	43/8.3/0675	Registration number:	43/7.1.3/0716
Name of medicine:	DONERIN 10	Name of medicine:	FERROUS GLUCONATE SYRUP BARRS	Name of medicine:	DYNA IRBESARTAN 75 mg
Dosage form:	TABLET	Dosage form:	SYRUP	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: DONEPEZIL HCl 10.0 mg	Active ingredients:	EACH 5.0 ml CONTAINS: FERROUS GLUCONATE 350.0 mg	Active ingredients:	EACH TABLET CONTAINS: IRBESARTAN 75.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	LASARA TRADERS (PTY) LTD	Applicant:	BARRS PHARMACEUTICAL INDUSTRIES cc	Applicant:	PHARMA DYNAMICS (PTY) LTD
Manufacturer:	SPECIFAR S.A. VARVARA, ATHENS, GREECE	Manufacturer:	BARRS PHARMACEUTICAL INDUSTRIES cc, NDABENI, CAPE TOWN, RSA	Manufacturer:	LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN
Packer:	SPECIFAR S.A. VARVARA, ATHENS, GREECE SPECIPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA	Packer:	BARRS PHARMACEUTICAL INDUSTRIES cc, NDABENI, CAPE TOWN, RSA	Packer:	LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES (PTY) LTD, N'DABENI, PINELANDS, RSA SPECIPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Laboratory: FPRC:	SPECIFAR S.A. VARVARA, ATHENS, GREECE SPECIPHARM HOLDINGS (PTY) LTD, MIDRAND, RSA	Laboratory: FPRC:	BARRS PHARMACEUTICAL INDUSTRIES cc, NDABENI, CAPE TOWN, RSA	Laboratory: FPRC:	LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN LABORATORIOS ECHEVARNE, BARCELONA, SPAIN CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	LASARA TRADERS (PTY) LTD, MORELETA PARK, PRETORIA, RSA	FPRR:	BARRS PHARMACEUTICAL INDUSTRIES cc, NDABENI, CAPE TOWN, RSA	FPRR:	PHARMA DYNAMICS (PTY) LTD, WESTLAKE, CAPE TOWN, RSA
Shelf-life:	24 months (Provisional)*	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

A provisional shelf-life of 24 months is approved for this product manufactured with API from Jubilant Oranosys and Ranbaxy Laboratories Limited

MRF 15	MRF 15	MRF 15
Registration number: 43/7.1.3/0717 Name of medicine: DYNA IRBESARTAN 150 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: IRBESARTAN 150,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: PHARMA DYNAMICS (PTY) LTD Manufacturer: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN Packer: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES (PTY) LTD, N'DABENI, PINELANDS, RSA SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA Laboratory: FPRC: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN LABORATORIOS ECHEVARNE, BARCELONA, SPAIN CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA FPRR: PHARMA DYNAMICS (PTY) LTD, WESTLAKE, CAPE TOWN, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012	Registration number: 43/7.1.3/0718 Name of medicine: DYNA IRBESARTAN 300 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: IRBESARTAN 300,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: PHARMA DYNAMICS (PTY) LTD Manufacturer: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN Packer: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES (PTY) LTD, N'DABENI, PINELANDS, RSA SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA Laboratory: FPRC: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN LABORATORIOS ECHEVARNE, BARCELONA, SPAIN CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA FPRR: PHARMA DYNAMICS (PTY) LTD, WESTLAKE, CAPE TOWN, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012	Registration number: 43/7.1.3/0719 Name of medicine: DYNARB 75 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: IRBESARTAN 75,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: PHARMA DYNAMICS (PTY) LTD Manufacturer: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN Packer: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES (PTY) LTD, N'DABENI, PINELANDS, RSA SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA Laboratory: FPRC: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN LABORATORIOS ECHEVARNE, BARCELONA, SPAIN CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA FPRR: PHARMA DYNAMICS (PTY) LTD, WESTLAKE, CAPE TOWN, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012

MRF 15

MRF 15

MRF 15

Registration number:	43/7.1.3/0721	Registration number:	43/20.1.1/1026
Name of medicine:	DYNARB 300 mg	Name of medicine:	AURO CEFTAZIDIME 250 mg
Dosage form:	TABLET	Dosage form:	POWDER FOR INJECTION
Active ingredients:	EACH TABLET CONTAINS: IRBESARTAN 300,0 mg	Active ingredients:	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 250,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	PHARMA DYNAMICS (PTY) LTD	Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN	Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA
Packer:	LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA PHARMACEUTICAL ENTERPRISES (PTY) LTD, N'DABENI, PINELANDS, RSA SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	Packer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH INDIA
Laboratory:	FPRC: LABORATORIOS CINFA, S.A., HUARTE-PAMPLONA, SPAIN LABORATORIOS LICONSA, S.A., AZUQUECA DE HENARES (GUADALAJARA), SPAIN LABORATORIOS ECHEVARNE, BARCELONA, SPAIN CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory:	FPRC: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	36 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15	MRF 15	MRF 15	MRF 15
Registration number:	43/20.1.1/1027	43/20.1.1/1028	43/20.1.1/1029
Name of medicine:	AURO CEFTAZIDIME 500 mg	AURO CEFTAZIDIME 1,0 g	AURO CEFTAZIDIME 2,0 g
Dosage form:	POWDER FOR INJECTION	POWDER FOR INJECTION	POWDER FOR INJECTION
Active ingredients:	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 500,0 mg	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 1,0 g	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 2,0 g
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
Applicant:	AUROBINDO PHARMA (PTY) LTD	AUROBINDO PHARMA (PTY) LTD	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA	FPRC: AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA
FPRC:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRC: AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRC: AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA
Shelf-life:	36 months	36 months	36 months
Date of registration:	07 JUNE 2012	07 JUNE 2012	07 JUNE 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/20.1.1/1035	Registration number:	43/20.1.1/1036	Registration number:	43/20.1.1/1037
Name of medicine:	AURIZEF 250 mg	Name of medicine:	AURIZEF 500 mg	Name of medicine:	AURIZEF 1,0 g
Dosage form:	POWDER FOR INJECTION	Dosage form:	POWDER FOR INJECTION	Dosage form:	POWDER FOR INJECTION
Active ingredients:	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 250,0 mg	Active ingredients:	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 500,0 mg	Active ingredients:	EACH VIAL CONTAINS: Ceftazidime Pentahydrate equivalent to Ceftazidime 1,0 g
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	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	Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Packer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Packer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA
Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA
FPRR:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRR:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRR:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA
Shelf-life:	36 months	Shelf-life:	36 months	Shelf-life:	36 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/20.1.1/1038	Registration number:	44/30.1/0039	Registration number:	44/10.2.2/0059
Name of medicine:	AURIZEF 2.0 g	Name of medicine:	MENCEVAX ACW135Y MULTIDOSE	Name of medicine:	MONTE-AIR 4 mg
Dosage form:	POWDER FOR INJECTION	Dosage form:	INJECTION	Dosage form:	TABLET
Active ingredients:	EACH VIAL CONTAINS: Ceftriaxime Pentahydrate equivalent to Ceftriaxime 2.0 g	Active ingredients:	EACH 0.5 ml DOSE CONTAINS: NEISSERIA MENINGITIDIS GROUP A 50.0 µg NEISSERIA MENINGITIDIS GROUP C 50.0 µg NEISSERIA MENINGITIDIS GROUP Y 50.0 µg NEISSERIA MENINGITIDIS GROUP W135 50.0 µg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	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:	GLAXOSMITHKLINE SOUTH AFRICA (PTY) LTD	Applicant:	MSD (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	GLAXOSMITHKLINE BIOLOGICALS S.A., Rixensart, Belgium GLAXOSMITHKLINE BIOLOGICALS, Dresden, Germany	Manufacturer:	MERCK SHARP & DOHME, Crawlington, Northumberland, UK
Packer:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA	Packer:	GLAXOSMITHKLINE BIOLOGICALS S.A., Rixensart, Belgium GLAXOSMITHKLINE BIOLOGICALS S.A., Wavre, Belgium GLAXOSMITHKLINE BIOLOGICALS, Dresden, Germany GLAXOSMITHKLINE, EPPING, CAPE TOWN	Packer:	MERCK SHARP & DOHME, Crawlington, Northumberland, UK MMD HAARLEM, HAARLEM, THE NETHERLANDS MSD, HALFWAY HOUSE, MIDRAND, RSA
Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT VI, PATANCHERU MANDAL, MEDAK DISTRICT, ANDHRA PRADESH, INDIA M & L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA KHULULEKANI LABORATORY SERVICES cc, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	GLAXOSMITHKLINE BIOLOGICALS S.A., Rixensart, Belgium	Laboratory: FPRC:	MERCK SHARP & DOHME, Crawlington, Northumberland, UK MMD HAARLEM, HAARLEM, THE NETHERLANDS
FPRR:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRC/FPRR:	GLAXOSMITHKLINE, EPPING, CAPE TOWN	FPRC/FPRR:	MSD, HALFWAY HOUSE, MIDRAND, RSA
Shelf-life:	36 months	Shelf-life:	36 months at 2 – 8 °C (Lyophilised vaccine) 8 hours at 2 – 8 °C (Reconstituted vaccine) 60 months at 2 – 25 °C (Diluent)	Shelf-life:	24 months
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15	MRF 15	MRF 15	MRF 15
Registration number:	Registration number:	Registration number:	Registration number:
Name of medicine:	Name of medicine:	Name of medicine:	Name of medicine:
Dosage form:	Dosage form:	Dosage form:	Dosage form:
Active ingredients:	Active ingredients:	Active ingredients:	Active ingredients:
Conditions of registration:	Conditions of registration:	Conditions of registration:	Conditions of registration:
Applicant:	Applicant:	Applicant:	Applicant:
Manufacturer:	Manufacturer:	Manufacturer:	Manufacturer:
Packer:	Packer:	Packer:	Packer:
Laboratory: FPRC:	Laboratory: FPRC:	Laboratory: FPRC:	Laboratory: FPRC:
Shelf-life:	Shelf-life:	Shelf-life:	Shelf-life:
Date of registration:	Date of registration:	Date of registration:	Date of registration:
44/10.2.2/0060	44/10.2.2/0061	44/10.2.2/0062	44/10.2.2/0062
MONTE-AIR 5 mg	MONTE-AIR 10 mg	MONTE-AIR SPRINKLES	MONTE-AIR SPRINKLES
TABLET	TABLET	GRANULES	GRANULES
EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10,0 mg	EACH SACHET OF GRANULES CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg	EACH SACHET OF GRANULES CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
MSD (PTY) LTD	MSD (PTY) LTD	MSD (PTY) LTD	MSD (PTY) LTD
MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	MERCK MANUFACTURING DIVISION, WEST POINT, PENNSYLVANIA, USA	MERCK MANUFACTURING DIVISION, WEST POINT, PENNSYLVANIA, USA
MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	MERCK MANUFACTURING DIVISION, WILSON, NORTH CAROLINA, USA	MERCK MANUFACTURING DIVISION, WILSON, NORTH CAROLINA, USA
MMD HAARLEM, HAARLEM, THE NETHERLANDS MSD, HALFWAY HOUSE, MIDRAND, RSA	MMD HAARLEM, HAARLEM, THE NETHERLANDS MSD, HALFWAY HOUSE, MIDRAND, RSA	ANDERSON PACKAGING INC, ROCKFORD, ILLINOIS, USA MMD HAARLEM, HAARLEM, THE NETHERLANDS MSD, HALFWAY HOUSE, MIDRAND, RSA	ANDERSON PACKAGING INC, ROCKFORD, ILLINOIS, USA MMD HAARLEM, HAARLEM, THE NETHERLANDS MSD, HALFWAY HOUSE, MIDRAND, RSA
MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	MERCK MANUFACTURING DIVISION, WEST POINT, PENNSYLVANIA, USA	MERCK MANUFACTURING DIVISION, WEST POINT, PENNSYLVANIA, USA
MMD HAARLEM, HAARLEM, THE NETHERLANDS	MMD HAARLEM, HAARLEM, THE NETHERLANDS	MERCK MANUFACTURING DIVISION, WILSON, NORTH CAROLINA, USA	MERCK MANUFACTURING DIVISION, WILSON, NORTH CAROLINA, USA
MSD, HALFWAY HOUSE, MIDRAND, RSA	MSD, HALFWAY HOUSE, MIDRAND, RSA	MMD HAARLEM, HAARLEM, THE NETHERLANDS	MMD HAARLEM, HAARLEM, THE NETHERLANDS
FPRC:	FPRC:	FPRC/FPRR:	FPRC/FPRR:
MSD, HALFWAY HOUSE, MIDRAND, RSA	MSD, HALFWAY HOUSE, MIDRAND, RSA	MSD, HALFWAY HOUSE, MIDRAND, RSA	MSD, HALFWAY HOUSE, MIDRAND, RSA
24 months	24 months	24 months	24 months
07 JUNE 2012	07 JUNE 2012	07 JUNE 2012	07 JUNE 2012

MRF 15		MRF 15	
Registration number:	44/10.2.1/0544	Registration number:	44/10.2.2/0791
Name of medicine:	ONBREZ BREEZHALER 150 µg	Name of medicine:	MONTELUKAST ZYDUS 10 mg
Dosage form:	DRY POWDER INHALATION CAPSULES	Dosage form:	TABLET
Active ingredients:	EACH CAPSULE CONTAINS: INDACATEROL MALEATE EQUIVALENT TO INDACATEROL 150,0 µg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	NOVARTIS SA (PTY) LTD	Applicant:	ZYDUS HEALTHCARE (PTY) LTD
Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND	Manufacturer:	ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA
Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTEIN, SWITZERLAND NOVARTIS PHARMA GmbH, WEHRBADEN, GERMANY NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND	Packer:	ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA
Laboratory: FPRC:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory: FPRC:	ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA
FPR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	FPRC/FPRR:	ZYDUS HEALTHCARE (PTY) LTD, VAN DER HOFF PARK, POTCHEFSTROOM, RSA
Shelf-life:	24 months	Shelf-life:	24 months (Provisional)
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15	MRF 15	MRF 15
Registration number: 44/10.2.2/0821 Name of medicine: MONTELUKAST ZYDUS 4 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: ZYDUS HEALTHCARE (PTY) LTD Manufacturer: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA Packer: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA Laboratory: FPRC: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA FPRC/FPRR: ZYDUS HEALTHCARE (PTY) LTD, VAN DER HOFF PARK, POTCHEFSTROOM, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012	Registration number: 44/10.2.2/0822 Name of medicine: MONTELUKAST ZYDUS 5 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: ZYDUS HEALTHCARE (PTY) LTD Manufacturer: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA Packer: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA Laboratory: FPRC: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA FPRC/FPRR: ZYDUS HEALTHCARE (PTY) LTD, VAN DER HOFF PARK, POTCHEFSTROOM, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012	Registration number: 44/10.2.2/0823 Name of medicine: LUMONT 4 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: ZYDUS HEALTHCARE (PTY) LTD Manufacturer: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA Packer: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA Laboratory: FPRC: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA FPRC/FPRR: ZYDUS HEALTHCARE (PTY) LTD, VAN DER HOFF PARK, POTCHEFSTROOM, RSA Shelf-life: 24 months (Provisional) Date of registration: 07 JUNE 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/10.2/20824	Registration number:	44/7.1.3/0857	Registration number:	44/7.1.3/0858
Name of medicine:	LUMONT 5 mg	Name of medicine:	TWYNSTA 40/5 mg TABLET	Name of medicine:	TWYNSTA 40/10 mg TABLET
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: TELISARTAN 40,0 mg AMLODIPINE BESYLATE EQUIVALENT TO AMLODIPINE 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: TELISARTAN 40,0 mg AMLODIPINE BESYLATE EQUIVALENT TO AMLODIPINE 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	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ZYDUS HEALTHCARE (PTY) LTD	Applicant:	INGELHEIM PHARMACEUTICALS (PTY) LTD	Applicant:	INGELHEIM PHARMACEUTICALS (PTY) LTD
Manufacturer:	ZYDUS CADILA HEALTHCARE LTD, MORAIVA, SANAND, AHMEDABAD, INDIA	Manufacturer:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA	Manufacturer:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA
Packer:	ZYDUS CADILA HEALTHCARE LTD, MORAIVA, SANAND, AHMEDABAD, INDIA	Packer:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA	Packer:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA
Laboratory:	FPRC: ZYDUS CADILA HEALTHCARE LTD, MORAIVA, SANAND, AHMEDABAD, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	Laboratory:	FPRC: BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA PHAST GmbH, HOMBURG/SAAR, GERMANY A&M STABTEST, MAINZ, GERMANY LABOR L + S AG, BAD BOCKLET-GROSSENBRACH, GERMANY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory:	FPRC: BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA PHAST GmbH, HOMBURG/SAAR, GERMANY A&M STABTEST, MAINZ, GERMANY LABOR L + S AG, BAD BOCKLET-GROSSENBRACH, GERMANY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA
FPRC/FPRR:	ZYDUS HEALTHCARE (PTY) LTD, VAN DER HOFF PARK, POTCHEFSTROOM, RSA	FPRR:	INGELHEIM PHARMACEUTICALS (PTY) LTD, RANDBURG, RSA	FPRR:	INGELHEIM PHARMACEUTICALS (PTY) LTD, RANDBURG, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15		3F 15	
Registration number:	447/1.30859	Registration number:	45/20.2.8/0171
Name of medicine:	TWYNSTA 80/5 mg TABLET	Name of medicine:	EFLATEN
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TELISARTAN 80,0 mg AMLODIPINE BESYLATE EQUIVALENT TO AMLODIPINE 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: TENOFIVIR DISOPROXIL FUMARATE 300,0 mg LAMIVUDINE 300,0 mg EFAVIRENZ 600,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8*
Applicant:	INGELHEIM PHARMACEUTICALS (PTY) LTD	Applicant:	MYLAN (PTY) LTD
Manufacturer:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA	Manufacturer:	MATRIX LABORATORIES LIMITED, SINNAR, NASHIK, MAHARASHTRA, INDIA
Packer:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA	Packer:	MATRIX LABORATORIES LIMITED, SINNAR, NASHIK, MAHARASHTRA, INDIA
Laboratory: FPRC:	BOEHRINGER INGELHEIM PHARMA GmbH & CO. KG, INGELHEIM AM RHEIN, GERMANY CIPLA LIMITED (Unit IV), VERNA, GOA, INDIA PHAST GmbH, HOMBURG/SAAR, GERMANY A&M STABTEST, MAINZ, GERMANY LABOR L + S AG, BAD BOCKLET-GROSSENBRACH, GERMANY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory: FPRC:	MATRIX LABORATORIES LIMITED, SINNAR, NASHIK, MAHARASHTRA, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA ACORN PHARMACEUTICALS (PTY) LTD, NORTH RIDING, JOHANNESBURG, RSA
FPRR:	INGELHEIM PHARMACEUTICALS (PTY) LTD, RANDSBURG, RSA	FPRR:	MYLAN (PTY) LTD, MODDERFONTEIN, JOHANNESBURG, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	07 JUNE 2012	Date of registration:	07 JUNE 2012

MRF 15

Registration number:	45/20.2.8/0172
Name of medicine:	ATROIZA
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TENOFIVIR DISOPROXIL 300,0 mg FUMARATE 300,0 mg EFAVIRENZ 600,0 mg EMTRICITABINE 200,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8*
Applicant:	MYLAN (PTY) LTD
Manufacturer:	MATRIX LABORATORIES LIMITED, SINNAR, NASHIK, MAHARASHTRA, INDIA
Packer:	MATRIX LABORATORIES LIMITED, SINNAR, NASHIK, MAHARASHTRA, INDIA
Laboratory: FPRC:	MATRIX LABORATORIES LIMITED, SINNAR, NASHIK, MAHARASHTRA, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA ACORN PHARMACEUTICALS (PTY) LTD, NORTH RIDING, JOHANNESBURG, RSA
FPRC:	MYLAN (PTY) LTD, MODDERFONTEIN, JOHANNESBURG, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	07 JUNE 2012

MRF 15

Registration number:	07/3.1.2.2/23
Name of medicine:	ONSIOR INJECTION
Dosage form:	INJECTION
Active ingredients:	EACH 1,0 ml SOLUTION CONTAINS: ROBENACOXIB 20,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	NOVARTIS SA (PTY) LTD
Manufacturer:	NOVARTIS SANTE ANIMALE S.A.S, HUNINGUE, FRANCE
Packer:	NOVARTIS SANTE ANIMALE S.A.S, HUNINGUE, FRANCE
Laboratory: FPRC:	NOVARTIS SANTE ANIMALE S.A.S, HUNINGUE, FRANCE NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA
FPRR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months stored at 2-8 °C In-use shelf life of 28 days stored at 2-8 °C
Date of registration:	27 JULY 2012

MRF-15

Registration number:	37/15.4/0224
Name of medicine:	LIPOSIC EYE GEL
Dosage form:	EYE GEL
Active ingredients:	EACH 1,0 g GEL CONTAINS: CARBOMER 2,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SOFLENS (PTY) LTD
Manufacturer:	Dr MANN PHARMA, CHEM- PHARM FABRIK GmbH, BERLIN, GERMANY
Packer:	Dr MANN PHARMA, CHEM- PHARM FABRIK GmbH, BERLIN, GERMANY COLUMBIA PHARMACEUTICALS (PTY) LTD, MIDDEL RD, BARDENE, BOKSBURG, RSA
Laboratory: FPRC:	Dr MANN PHARMA, CHEM- PHARM FABRIK GmbH, BERLIN, GERMANY CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	SOFLENS (PTY) LTD, RIVONIA, SANDTON, RSA
Shelf-life:	24 months
Date of registration:	27 JULY 2012

MRF 15

Registration number:	37/15.2/0588
Name of medicine:	LOTEMAX OPHTHALMIC SUSPENSION
Dosage form:	EYE DROPS
Active ingredients:	EACH 1,0 ml SUSPENSION CONTAINS: LOTEPREDNOL ETABONATE 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SOFLENS (PTY) LTD
Manufacturer:	BAUSCH & LOMB INC, HIDDEN RIVER PARKWAY, TAMPA, FLORIDA, USA
Packer:	BAUSCH & LOMB INC, HIDDEN RIVER PARKWAY, TAMPA, FLORIDA, USA COLUMBIA PHARMACEUTICALS (PTY) LTD, MIDDEL RD, BARDENE, BOKSBURG, RSA
Laboratory: FPRC:	BAUSCH & LOMB INC, HIDDEN RIVER PARKWAY, TAMPA, FLORIDA, USA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	SOFLENS (PTY) LTD, RIVONIA, SANDTON, RSA
Shelf-life:	24 months
Date of registration:	27 JULY 2012

MRF 15	Registration number: A39/7.1/0419 Name of medicine: APEX TERAZOSIN 1 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 1,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: CAMOX PHARMACEUTICALS (PTY) LTD Manufacturer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA Packer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA Laboratory: FPRC: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA FPRR: CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA Shelf-life: 36 months Date of registration: 27 JULY 2012
MRF15	Registration number: A39/7.1/0420 Name of medicine: APEX TERAZOSIN 2 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 2,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: CAMOX PHARMACEUTICALS (PTY) LTD Manufacturer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA Packer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA Laboratory: FPRC: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA FPRR: CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA Shelf-life: 36 months Date of registration: 27 JULY 2012
MRF 15	Registration number: A39/7.1/0421 Name of medicine: APEX TERAZOSIN 5 mg Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 5,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7 Applicant: CAMOX PHARMACEUTICALS (PTY) LTD Manufacturer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA Packer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA Laboratory: FPRC: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA FPRR: CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA Shelf-life: 36 months Date of registration: 27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	Registration number:	Registration number:	Registration number:	Registration number:	Registration number:
A39/7.1/0422	A39/7.1/0423	A39/7.1/0424	A39/7.1/0424	A39/7.1/0424	A39/7.1/0424
Name of medicine:	Name of medicine:	Name of medicine:	Name of medicine:	Name of medicine:	Name of medicine:
APEX TERAZOSIN 10 mg	APEX TERAZOSIN BPH STARTER PACK	APEX TERAZOSIN BPH STARTER PACK	APEX TERAZOSIN BPH STARTER PACK	TERAPRESS BPH STARTER PACK	TERAPRESS BPH STARTER PACK
Dosage form:	Dosage form:	Dosage form:	Dosage form:	Dosage form:	Dosage form:
TABLET	TABLET	TABLET	TABLET	TABLET	TABLET
Active ingredients:	Active ingredients:	Active ingredients:	Active ingredients:	Active ingredients:	Active ingredients:
EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 10,0 mg	EACH PACK CONTAINS 3 x Apex Terazosin 1 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 1,0 mg 11 x Apex Terazosin 2 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 2,0 mg	EACH PACK CONTAINS 3 x Apex Terazosin 1 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 1,0 mg 11 x Apex Terazosin 2 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 2,0 mg	EACH PACK CONTAINS 3 x Apex Terazosin 1 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 1,0 mg 11 x Apex Terazosin 2 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 2,0 mg	EACH PACK CONTAINS 3 x Terapress 1 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 1,0 mg 11 x Terapress 2 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 2,0 mg	EACH PACK CONTAINS 3 x Terapress 1 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 1,0 mg 11 x Terapress 2 mg tablets containing Terazosin hydrochloride equivalent to Terazosin 2,0 mg
Conditions of registration:	Conditions of registration:	Conditions of registration:	Conditions of registration:	Conditions of registration:	Conditions of registration:
1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7
Applicant:	Applicant:	Applicant:	Applicant:	Applicant:	Applicant:
CAMOX PHARMACEUTICALS (PTY) LTD	CAMOX PHARMACEUTICALS (PTY) LTD	CAMOX PHARMACEUTICALS (PTY) LTD	CAMOX PHARMACEUTICALS (PTY) LTD	CAMOX PHARMACEUTICALS (PTY) LTD	CAMOX PHARMACEUTICALS (PTY) LTD
Manufacturer:	Manufacturer:	Manufacturer:	Manufacturer:	Manufacturer:	Manufacturer:
INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA
Packer:	Packer:	Packer:	Packer:	Packer:	Packer:
INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA
Laboratory:	Laboratory:	Laboratory:	Laboratory:	Laboratory:	Laboratory:
FPRC:	FPRC:	FPRC:	FPRC:	FPRC:	FPRC:
SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA	SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA	SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA	SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA	SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA	SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA
M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA	M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA	M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA	M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA	M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA	M&L LABORATORY SERVICES, LIMBRO BUSINESS PARK, SANDTON, RSA
INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA
FPRC:	FPRC:	FPRC:	FPRC:	FPRC:	FPRC:
CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA
Shelf-life:	Shelf-life:	Shelf-life:	Shelf-life:	Shelf-life:	Shelf-life:
36 months	36 months	36 months	36 months	36 months	36 months
Date of registration:	Date of registration:	Date of registration:	Date of registration:	Date of registration:	Date of registration:
27 JULY 2012	27 JULY 2012	27 JULY 2012	27 JULY 2012	27 JULY 2012	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	A39/7.1/0425	Registration number:	A39/7.1/0426	Registration number:	A39/7.1/0427
Name of medicine:	TERAPRESS 1 mg	Name of medicine:	TERAPRESS 2 mg	Name of medicine:	TERAPRESS 5 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 1,0 mg	Active ingredients:	EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 2,0 mg	Active ingredients:	EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	CAMOX PHARMACEUTICALS (PTY) LTD	Applicant:	CAMOX PHARMACEUTICALS (PTY) LTD	Applicant:	CAMOX PHARMACEUTICALS (PTY) LTD
Manufacturer:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	Manufacturer:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	Manufacturer:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA
Packer:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	Packer:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	Packer:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA
Laboratory: FPRC:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	Laboratory: FPRC:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	Laboratory: FPRC:	INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA
FPRR:	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	FPRR:	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	FPRR:	CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA
Shelf-life:	36 months	Shelf-life:	36 months	Shelf-life:	36 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

F 15

MRF 15

MRF 15

Registration number: A39/7.1/0428	Registration number: A40/4/0214	Registration number: 41/26/0034
Name of medicine: TERAPRESS 10 mg	Name of medicine: SABAX BUPIVACAINE 0,1 % VIAFLEX	Name of medicine: CIPLA DOXORUBICIN 10
Dosage form: TABLET	Dosage form: SOLUTION FOR INFUSION	Dosage form: INJECTION
Active ingredients: EACH TABLET CONTAINS: Terazosin hydrochloride equivalent to Terazosin 10,0 mg	Active ingredients: EACH 1 000,0 ml CONTAINS BUPIVACAINE HYDROCHLORIDE 1,0 g	Active ingredients: EACH VIAL CONTAINS: DOXORUBICIN HYDROCHLORIDE 10,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: CAMOX PHARMACEUTICALS (PTY) LTD	Applicant: ADCOCK INGRAM CRITICAL CARE (PTY) LTD	Applicant: CIPLA MEDPRO (PTY) LTD
Manufacturer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	Manufacturer: ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA	Manufacturer: CIPLA LTD, UNIT V, VERNA, GOA, INDIA
Packer: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA	Packer: ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA	Packer: CIPLA LTD, UNIT V, VERNA, GOA, INDIA
Laboratory, FPRC: INTAS PHARMACEUTICALS LTD, MATODA, GUJARAT, INDIA SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA	Laboratory, FPRC: ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA	Laboratory: FPRC CIPLA LTD, UNIT V, VERNA, GOA, INDIA
FPRR: CAMOX PHARMACEUTICALS (PTY) LTD, CROWN EXTENSION 3, JOHANNESBURG, RSA	FPRR ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA	FPRR: CIPLA MEDPRO (PTY) LTD, ROSENPARK, BELLVILLE, RSA
Shelf-life: 36 months	Shelf-life: 24 months	Shelf-life: 24 months
Date of registration: 27 JULY 2012	Date of registration: 27 JULY 2012	Date of registration: 27 JULY 2012

MRF 15		F 15	
Registration number:	41/26/0035	Registration number:	41/5.10/0241
Name of medicine:	CIPLA DOXORUBICIN 50	Name of medicine:	ONDANSETRON SAFELINE 4 mg INJECTION
Dosage form:	INJECTION	Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: DOXORUBICIN HYDROCHLORIDE 50.0 mg	Active ingredients:	EACH AMPOULE CONTAINS: Ondansetron hydrochloride dihydrate equivalent to Ondansetron 4.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	CIPLA MEDPRO (PTY) LTD	Applicant:	SAFELINE PHARMACEUTICALS (PTY) LTD
Manufacturer:	CIPLA LTD, (UNIT V), VERNA, GOA, INDIA	Manufacturer:	DEMO S.A., ATHENS-LAMIA, ATHENS, GREECE
Packer:	CIPLA LTD, (UNIT V), VERNA, GOA, INDIA	Packer:	DEMO S.A., ATHENS-LAMIA, ATHENS, GREECE
Laboratory: FPRC:	CIPLA LTD, (UNIT V), VERNA, GOA, INDIA	Laboratory: FPRC:	DEMO S.A., ATHENS-LAMIA, ATHENS, GREECE INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA
FPRC:	CIPLA MEDPRO (PTY) LTD, ROSENPARK, RSA	FPRC:	SAFELINE PHARMACEUTICALS (PTY) LTD, FLORIDA, RSA
Shelf-life:	24 months	Shelf-life:	24 months (Provisional) for product stored at or below 25 °C 24 hours for diluted solutions stored at 2 – 8 °C
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

Registration number:	41/5.10/0242
Name of medicine:	ONDANSETRON SAFELINE 8 mg INJECTION
Dosage form:	INJECTION
Active ingredients:	EACH AMPOULE CONTAINS: Ondansetron hydrochloride dihydrate equivalent to Ondansetron 8.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	SAFELINE PHARMACEUTICALS (PTY) LTD
Manufacturer:	DEMO S.A., ATHENS-LAMIA, ATHENS, GREECE
Packer:	DEMO S.A., ATHENS-LAMIA, ATHENS, GREECE
Laboratory: FPRC:	DEMO S.A., ATHENS-LAMIA, ATHENS, GREECE INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA
FPRC:	SAFELINE PHARMACEUTICALS (PTY) LTD, FLORIDA, RSA
Shelf-life:	24 months (Provisional) for product stored at or below 25 °C 24 hours for diluted solutions stored at 2 – 8 °C
Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	41/20.1.1/0947	Registration number:	41/20.1.1/0948	Registration number:	41/20.1.1/0949
Name of medicine:	TEICOPLANIN WINTHROP 200	Name of medicine:	TEICOPLANIN WINTHROP 400	Name of medicine:	TEICOPLANIN SANOFI-AVENTIS 200
Dosage form:	INJECTION	Dosage form:	INJECTION	Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: TEICOPLANIN 200,0 mg	Active ingredients:	EACH VIAL CONTAINS: TEICOPLANIN 400,0 mg	Active ingredients:	EACH VIAL CONTAINS: TEICOPLANIN 200,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD	Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD	Applicant:	SANOFI-AVENTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY	Manufacturer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY	Manufacturer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY
Packer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory: FPRC:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Laboratory: FPRC:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Laboratory: FPRC:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA	FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA	FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA
Shelf-life:	36 months	Shelf-life:	36 months	Shelf-life:	36 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	41/20.1.1/0950	Registration number:	41/34/1134	Registration number:	41/34/1135
Name of medicine:	TEICOPLANIN SANOFI-AVENTIS 400	Name of medicine:	TEICOPLANIN WINTHROP SOLVENT	Name of medicine:	TEICOPLANIN SANOFI-AVENTIS SOLVENT
Dosage form:	INJECTION	Dosage form:	INJECTION	Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: TEICOPLANIN 400,0 mg	Active ingredients:	EACH AMPOULE CONTAINS: WATER FOR INJECTION 3,0 ml	Active ingredients:	EACH AMPOULE CONTAINS: WATER FOR INJECTION 3,0 ml
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANOFI-AVENTIS SOUTH AFRICA (PTY) LTD	Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD	Applicant:	SANOFI-AVENTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY	Manufacturer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY	Manufacturer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY
Packer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory: FPRC:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Laboratory: FPRC:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Laboratory: FPRC:	GRUPPO LEPETIT Srl, ANAGNI, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA SANOFI-AVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA	FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA	FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA SANOFI-AVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA
Shelf-life:	36 months	Shelf-life:	36 months	Shelf-life:	36 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

F 15

MRF 15

MRF 15

Registration number: 42/28/0102	Registration number: 42/28/0103	Registration number: 42/28/0104
Name of medicine: OPTIMARK 10	Name of medicine: OPTIMARK 15	Name of medicine: OPTIMARK 20
Dosage form: SOLUTION FOR INJECTION	Dosage form: SOLUTION FOR INJECTION	Dosage form: SOLUTION FOR INJECTION
Active ingredients: EACH 10,0 ml CONTAINS: GADOVERSETAMIDE 3 309,0 mg	Active ingredients: EACH 15,0 ml CONTAINS: GADOVERSETAMIDE 4963,5 mg	Active ingredients: EACH 20,0 ml CONTAINS: GADOVERSETAMIDE 6618,0 mg
Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7	Conditions of registration: 1, 2, 3, 4, 5, 6, 7
Applicant: COVIDIEN (PTY) LTD	Applicant: COVIDIEN (PTY) LTD	Applicant: COVIDIEN (PTY) LTD
Manufacturer: TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA	Manufacturer: TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA	Manufacturer: TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA
Packer: TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA	Packer: TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA	Packer: TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA
Laboratory: FPRC	Laboratory: FPRC:	Laboratory: FPRC:
		TYCO HEALTHCARE INC, RALEIGH, NORTH CAROLINA, USA
		PETLABS PHARMACEUTICALS (PTY) LTD, GROENKLOOF, PRETORIA, RSA
		BIOCHEMICAL AND SCIENTIFIC cc, HILTON, KWA-ZULU NATAL, RSA
FPRC:	FPRC:	FPRC:
		COVIDIEN (PTY) LTD, RANDJESPAK, MIDRAND, RSA
Shelf-life: 24 months (Glass vials) 36 months (Pre-filled syringes)	Shelf-life: 24 months (Glass vials) 36 months (Pre-filled syringes)	Shelf-life: 24 months (Glass vials) 36 months (Pre-filled syringes)
Date of registration: 27 JULY 2012	Date of registration: 27 JULY 2012	Date of registration: 27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	42/28/0105	Registration number:	42/17.1.3/0262	Registration number:	42/20.1.1/0427
Name of medicine:	OPTIMARK 30	Name of medicine:	TRANDOPRESS 2 mg	Name of medicine:	LEVOFLOXACIN SPECPHARM 250 IV
Dosage form:	SOLUTION FOR INJECTION	Dosage form:	CAPSULE	Dosage form:	SOLUTION FOR IV INFUSION
Active ingredients:	EACH 30.0 ml CONTAINS: GADOVERSETAMIDE 9927.0 mg	Active ingredients:	EACH CAPSULE CONTAINS: TRANDOLAPRIL 2.0 mg	Active ingredients:	EACH 50.0 ml SOLUTION CONTAINS: LEVOFLOXACIN 250.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	COVIDIEN (PTY) LTD	Applicant:	PHARMACARE LIMITED	Applicant:	SPECPHARM (PTY) LTD
Manufacturer:	TYCO HEALTHCARE INC. RALEIGH, NORTH CAROLINA, USA	Manufacturer:	PHARMATHEN S.A, PALLINI, ATTIKIS, GREECE	Manufacturer:	BIOMENDI S.A, BERNEDO, ALVA, SPAIN
Packer:	TYCO HEALTHCARE INC. RALEIGH, NORTH CAROLINA, USA	Packer:	PHARMATEN S.A, PALLINI, ATTIKIS, GREECE PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	Packer:	BIOMENDI S.A, BERNEDO, ALVA, SPAIN
Laboratory:	FPRC: TYCO HEALTHCARE INC. RALEIGH, NORTH CAROLINA, USA PETLABS PHARMACEUTICALS (PTY) LTD, GROENKLOOF, PRETORIA, RSA BIOCHEMICAL AND SCIENTIFIC CC, HILTON, KWA-ZULU NATAL, RSA	Laboratory:	FPRC: PHARMATEN S.A, PALLINI, ATTIKIS, GREECE PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, POTCHEFSTROOM, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory:	FPRC: BIOMENDI S.A, BERNEDO, ALVA, SPAIN SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRC:	COVIDIEN (PTY) LTD, RANDJESPAK, MIDRAND, RSA	FPRC/FPRC:	PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	FPRC:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Shelf-life:	24 months (Glass vials) 36 months (Pre-filled syringes)	FPRC:	PHARMACARE LIMITED, WOODMEAD, SANDTON, RSA	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Shelf-life:	24 months (Provisional)	Date of registration:	27 JULY 2012
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012		

MRF 15	MRF 15	MRF 15	MRF 15
Registration number:	42/20.1.1/0428	42/20.1.1/0439	42/20.1.1/0440
Name of medicine:	SPEC LEVOFLOXACIN 500 IV	LEVOFLOXACIN SPECPHARM 500 IV	SPEC LEVOFLOXACIN 250 IV
Dosage form:	SOLUTION FOR IV INFUSION	SOLUTION FOR IV INFUSION	SOLUTION FOR IV INFUSION
Active ingredients:	EACH 100,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 500,0 mg	EACH 100,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 500,0 mg	EACH 50,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 250,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	1, 2, 3, 4, 5, 6, 7, 8	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	SPECPHARM (PTY) LTD	SPECPHARM (PTY) LTD	SPECPHARM (PTY) LTD
Manufacturer:	BIOMENDI S.A, BERNEDO, ALVA, SPAIN	BIOMENDI S.A, BERNEDO, ALVA, SPAIN	BIOMENDI S.A, BERNEDO, ALVA, SPAIN
Packer:	BIOMENDI S.A, BERNEDO, ALVA, SPAIN	BIOMENDI S.A, BERNEDO, ALVA, SPAIN	BIOMENDI S.A, BERNEDO, ALVA, SPAIN
Laboratory:	FPRC:	FPRC:	FPRC:
FPRC:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
FPRC:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Shelf-life:	24 months (Provisional)	24 months (Provisional)	24 months (Provisional)
Date of registration:	27 JULY 2012	27 JULY 2012	27 JULY 2012

MRF 15	Registration number: 42/20.1.1/0505 Name of medicine: ZITRO 500 TABLETS Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: Azithromycin dihydrate equivalent to Azithromycin 500,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: CIPLA MEDPRO (PTY) LTD Manufacturer: CIPLA LTD, UNIT VII, VERNA, GOA, INDIA Packer: CIPLA LTD, UNIT VII, VERNA, GOA, INDIA Laboratory: FPRC: CIPLA LTD, UNIT VII, VERNA, GOA, INDIA FPRR: CIPLA MEDPRO (PTY) LTD, ROSENPARK, SOUTH AFRICA Shelf-life: 24 months (Provisional) Date of registration: 27 JULY 2012
MRF 15	Registration number: 42/20.1.2/0771 Name of medicine: ADCO AMOXYCLAV BD Dosage form: TABLET Active ingredients: EACH TABLET CONTAINS: AMOXYCILLIN TRIHYDRATE EQUIVALENT TO AMOXYCILLIN 875,0 mg POTASSIUM CLAVULANATE EQUIVALENT TO CLAVULANIC ACID 125,0 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: ADCOCK INGRAM LIMITED Manufacturer: MEDREICH LTD, UNIT 1, VIRGONAGAR, BANGALORE, INDIA Packer: MEDREICH LTD, UNIT 1, VIRGONAGAR, BANGALORE, INDIA ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON, RSA Laboratory: FPRC: MEDREICH LTD, UNIT 1, VIRGONAGAR, BANGALORE, INDIA ADCOCK INGRAM LIMITED, RESEARCH AND DEVELOPMENT, AEROTON, JOHANNESBURG, RSA FPRC/FPRR: ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON, RSA FPRR: ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA Shelf-life: 24 months (Provisional) Date of registration: 27 JULY 2012
MRF 15	Registration number: 42/34/0786 Name of medicine: ADVAGRAF 0,5 mg Dosage form: CAPSULE Active ingredients: EACH CAPSULE CONTAINS: TACROLIMUS 0,5 mg Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8 Applicant: ASTELLAS PHARMA (PTY) LTD Manufacturer: ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND Packer: ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND Laboratory: FPRC: ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA FPRR: ASTELLAS PHARMA (PTY) LTD, BEDFORDVIEW, JOHANNESBURG, RSA Shelf-life: 24 months (Provisional) Date of registration: 27 JULY 2012

MRF 15		MRF 15	
Registration number:	42/34/0787	Registration number:	42/34/0788
Name of medicine:	ADVAGRAF 1 mg	Name of medicine:	ADVAGRAF 5 mg
Dosage form:	CAPSULE	Dosage form:	CAPSULE
Active ingredients:	EACH CAPSULE CONTAINS: TACROLIMUS 1,0 mg	Active ingredients:	EACH CAPSULE CONTAINS: TACROLIMUS 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:	ASTELLAS PHARMA (PTY) LTD	Applicant:	ASTELLAS PHARMA (PTY) LTD
Manufacturer:	ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND	Manufacturer:	ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND
Packer:	ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND	Packer:	ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND
Laboratory: FPRC:	ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	ASTELLAS IRELAND CO. LTD, KILLORGLIN, KERRY, IRELAND CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA
FPRR:	ASTELLAS PHARMA (PTY) LTD, BEDFORDVIEW, JOHANNESBURG, RSA	FPRR:	ASTELLAS PHARMA (PTY) LTD, BEDFORDVIEW, JOHANNESBURG, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012
		Registration number:	42/2.5/0810
		Name of medicine:	RESTAN TOPIRAMATE 25 mg
		Dosage form:	TABLET
		Active ingredients:	EACH TABLET CONTAINS: TOPIRAMATE 25,0 mg
		Conditions of registration:	1, 2, 3, 4, 5, 6, 7
		Applicant:	ADCOCK INGRAM LIMITED
		Manufacturer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
		Packer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
		Laboratory: FPRC:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
		FPRC/FPRR:	ADCOCK INGRAM HEALTHCARE (PTY) LTD, WADEVILLE, GERMISTON, RSA ADCOCK INGRAM LTD, RESEARCH AND DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
		Shelf-life:	36 months
		Date of registration:	27 JULY 2012

MRF 15

Registration number:	42/2.5/0811
Name of medicine:	RESTAN TOPIRAMATE 50 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TOPIRAMATE 50,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
Packer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
Laboratory: FPRC:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
FPRC/FPRR:	ADCOCK INGRAM HEALTHCARE (PTY) LTD, WADEVILLE, GERMISTON, RSA ADCOCK INGRAM LTD, RESEARCH AND DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPRR:	ADCOCK INGRAM LTD, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	36 months
Date of registration:	27 JULY 2012

MRF15

Registration number:	42/2.5/0812
Name of medicine:	RESTAN TOPIRAMATE 100 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TOPIRAMATE 100,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
Packer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
Laboratory: FPRC:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
FPRC/ FPRR:	ADCOCK INGRAM HEALTHCARE (PTY) LTD, WADEVILLE, GERMISTON, RSA ADCOCK INGRAM LTD, RESEARCH AND DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPRR:	ADCOCK INGRAM LTD, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	36 months
Date of registration:	27 JULY 2012

MRF 15

Registration number:	42/2.5/0813
Name of medicine:	RESTAN TOPIRAMATE 200 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TOPIRAMATE 200,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
Packer:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
Laboratory: FPRC:	ACTAVIS hf, HAFNARFJORDUR, ICELAND
FPRC/FPRR:	ADCOCK INGRAM HEALTHCARE (PTY) LTD, WADEVILLE, GERMISTON, RSA ADCOCK INGRAM LTD, RESEARCH AND DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPRR:	ADCOCK INGRAM LTD, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	36 months
Date of registration:	27 JULY 2012

MRF 15

Registration number:	42/21.12/0995
Name of medicine:	CASTRA 1
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: ANASTROZOLE 1,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ZYDUS HEALTHCARE SA (PTY) LTD
Manufacturer:	ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA
Packer:	ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA
Laboratory:	FPRC: ZYDUS CADILA HEALTHCARE LTD, MORAIYA, SANAND, AHMEDABAD, INDIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH- WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA PHARMASPEC LABORATORIES, FERNDALE, RANDBURG, RSA
FPRC:	ZYDUS HEALTHCARE SA (PTY) LTD, VAN DER HOFF PARK, POTCHEFSTROOM, RSA
Shelf-life:	24 months
Date of registration:	27 JULY 2012

MRF 15

Registration number:	42/21.12/1037
Name of medicine:	ASPEN ANASTROZOLE
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: ANASTROZOLE 1,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	PHARMACARE LIMITED
Manufacturer:	HAUPT PHARMA MUNSTER GmbH, MUNSTER, GERMANY PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA
Packer:	HAUPT PHARMA MUNSTER GmbH, MUNSTER, GERMANY PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA
Laboratory:	FPRC: SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA M & L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA
FPRC/FPFR:	PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA
FPFR:	PHARMACARE LIMITED, WOODMEAD, SANDTON, RSA
Shelf-life:	48 months
Date of registration:	27 JULY 2012

MRF 15

Registration number:	43/21.2/0002
Name of medicine:	GLAMARYL COMBI 1/250
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: GLIMEPIRIDE 1,0 mg METFORMIN HYDROCHLORIDE 250,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD
Manufacturer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA
Packer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory:	FPRC HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA
FPRC/FPFR:	WINTHROP PHARMACEUTICALS, (PTY) LTD WALTLOO, PRETORIA, RSA
FPFR:	WINTHROP PHARMACEUTICALS, (PTY) LTD MIDRAND, JOHANNESBURG, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/21.2/0003	Registration number:	43/21.2/0004	Registration number:	43/21.2/0005
Name of medicine:	GLAMARYL COMBI 2/500	Name of medicine:	AMARYL COMBI 1/250	Name of medicine:	AMARYL COMBI 2/500
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: GLIMEPIRIDE 2,0 mg METFORMIN HYDROCHLORIDE 500,0 mg	Active ingredients:	EACH TABLET CONTAINS: GLIMEPIRIDE 1,0 mg METFORMIN HYDROCHLORIDE 250,0 mg	Active ingredients:	EACH TABLET CONTAINS: GLIMEPIRIDE 2,0 mg METFORMIN HYDROCHLORIDE 500,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD	Applicant:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD	Applicant:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA	Manufacturer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA	Manufacturer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA
Packer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory: FPRC:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA	Laboratory: FPRC:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA	Laboratory: FPRC:	HANDOK PHARMACEUTICALS Co LTD, EUMSEONG-GUN, CHUNGCHONGBUK-DO, KOREA
FPRC/FPRR:	WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA	FPRC/FPRR:	WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA	FPRC/FPRR:	WINTHROP PHARMACEUTICALS, (PTY) LTD, WALTLOO, PRETORIA, RSA
FPRR:	WINTHROP PHARMACEUTICALS, (PTY) LTD, MIDRAND, JOHANNESBURG, RSA	FPRR:	SANOFLAVENTIS S.A. (PTY) LTD, MIDRAND, JOHANNESBURG, RSA	FPRR:	SANOFLAVENTIS S.A. (PTY) LTD, MIDRAND, JOHANNESBURG, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15	Registration number:	Name of medicine:	Dosage form:	Active ingredients:	Conditions of registration:	Applicant:	Manufacturer:	Packer:	Laboratory:	FPRC:	FPRR:	Shelf-life:	Date of registration:
MRF 15	43/26/0103	OXALIPLATIN PCH 100 CONCENTRATE FOR SOLUTION FOR INFUSION	EACH 20.0 ml VIAL CONTAINS: OXALIPLATIN 100.0 mg		1, 2, 3, 4, 5, 6, 7, 8	PHARMACHEMIE (PTY) LTD	PHARMACHEMIE B.V. HAARLEM, THE NETHERLANDS	PHARMACHEMIE B.V. HAARLEM, THE NETHERLANDS	PHARMACHEMIE B.V. HAARLEM, THE NETHERLANDS SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY NORTH WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD ATLASVILLE, BOKSBURG, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA		FPRR:	24 months (Provisional)	27 JULY 2012
MRF 15	43/30, 1/0127	PRIORIX TETRA POWDER AND DILUENT FOR SOLUTION FOR INJECTION	EACH 0.5 ml CONTAINS: Live attenuated measles virus (Schwarz strain) $\geq 10^{3.0}$ CCID ₅₀ Live attenuated mumps virus (RIT4385 strain) $\geq 10^{4.4}$ CCID ₅₀ Live attenuated rubella virus (Wistar RA 27/3 strain) $\geq 10^{3.0}$ CCID ₅₀ Live attenuated varicella virus (OKA strain) $\geq 10^{3.5}$ PFU		1, 2, 3, 4, 5, 6, 7	GLAXOSMITHKLINE SA (PTY) LTD	GLAXOSMITHKLINE BIOLOGICALS S.A. RIXENSART, BELGIUM SACHSISCHES SERUWERK DRESDEN, NI der SMITHKLINE BEECHAM PHARMA GmbH, DRESDEN, GERMANY	GLAXOSMITHKLINE BIOLOGICALS S.A. WAVRE, BELGIUM SACHSISCHES SERUWERK DRESDEN, NI der SMITHKLINE BEECHAM PHARMA GmbH, DRESDEN, GERMANY GLAXOSMITHKLINE SA (PTY) LTD, EPPING, CAPE TOWN, RSA	GLAXOSMITHKLINE BIOLOGICALS S.A. RIXENSART, BELGIUM GLAXOSMITHKLINE BIOLOGICALS S.A. WAVRE, BELGIUM		FPRC/FPRR:	60 months (Diluent stored at or below 25 °C) 24 months (Product stored at -20 °C) 18 months (Product stored at 2 – 8 °C) In-use shelf-life of 24 hours for product stored at 2 – 8 °C	27 JULY 2012
MRF 15	43/21, 2/0140	ACCORD PIOGLITAZONE 15 mg TABLET	EACH TABLET CONTAINS: PIOGLITAZONE HYDROCHLORIDE EQUIVALENT TO PIOGLITAZONE 15.0 mg		1, 2, 3, 4, 5, 6, 7, 8	ACCORD HEALTHCARE (PTY) LTD	INTAS PHARMACEUTICALS LIMITED, MATODA, TALUKA, AHMEDABAD, GUJARAT, INDIA	INTAS PHARMACEUTICALS LIMITED, MATODA, TALUKA, AHMEDABAD, GUJARAT, INDIA	INTAS PHARMACEUTICALS LIMITED, MATODA, TALUKA, AHMEDABAD, GUJARAT, INDIA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA		FPRR:	24 months (Provisional)	27 JULY 2012

MRF 15		JRF 15	
Registration number:	43/21.2/0141	Registration number:	43/20.1.1/0310
Name of medicine:	ACCORD PIOGLITAZONE 30 mg	Name of medicine:	BACTRICEF 1 g
Dosage form:	TABLET	Dosage form:	POWDER FOR INJECTION
Active ingredients:	EACH TABLET CONTAINS: PIOGLITAZONE HYDROCHLORIDE EQUIVALENT TO PIOGLITAZONE 30,0 mg	Active ingredients:	EACH VIAL CONTAINS: CEFEPIME HYDROCHLORIDE EQUIVALENT TO CEFEPIME 1 000,0 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:	PHARMACARE LIMITED
Manufacturer:	INTAS PHARMACEUTICALS LIMITED, MATODA, TALUKA, AHMEDABAD, GUJARAT, INDIA	Manufacturer:	STRIDES ARCOLAB LIMITED, ANEKAL TALUK, BANGALORE, INDIA
Packer:	INTAS PHARMACEUTICALS LIMITED, MATODA, TALUKA, AHMEDABAD, GUJARAT, INDIA	Packer:	STRIDES ARCOLAB LIMITED, ANEKAL TALUK, BANGALORE, INDIA PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA
Laboratory: FPRC:	INTAS PHARMACEUTICALS LIMITED, MATODA, TALUKA, AHMEDABAD, GUJARAT, INDIA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	STRIDES ARCOLAB LIMITED, ANEKAL TALUK, BANGALORE, INDIA SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA
FPRR:	ACCORD HEALTHCARE (PTY) LTD, RIVONIA, GAUTENG, RSA	FPRC/FPRR:	PHARMACARE LTD, KORTSEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	PHARMACARE LTD, WOODMEAD, SANDTON, RSA
Date of registration:	27 JULY 2012	Date of registration:	24 months (Provisional) 27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/20.2.8/0433	Registration number:	43/5.10/0542	Registration number:	43/5.10/0543
Name of medicine:	VALCYTE 50 mg/ml	Name of medicine:	ONDANSETRON FRESENIUS 2 mg/ml (4 mg/2 ml)	Name of medicine:	ONDANSETRON FRESENIUS 2 mg/ml (8 mg/4 ml)
Dosage form:	POWDER FOR ORAL SOLUTION	Dosage form:	SOLUTION FOR INJECTION	Dosage form:	SOLUTION FOR INJECTION
Active ingredients:	EACH 1.0 ml SOLUTION CONTAINS: Valganciclovir hydrochloride equivalent to Valganciclovir 50.0 mg	Active ingredients:	EACH AMPOULE CONTAINS: ONDANSETRON HYDROCHLORIDE DIHYDRATE EQUIVALENT TO ONDANSETRON 4.0 mg	Active ingredients:	EACH AMPOULE CONTAINS: ONDANSETRON HYDROCHLORIDE DIHYDRATE EQUIVALENT TO ONDANSETRON 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	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ROCHE PRODUCTS (PTY) LTD	Applicant:	FRESENIUS KABI SOUTH AFRICA (PTY) LTD	Applicant:	FRESENIUS KABI SOUTH AFRICA (PTY) LTD
Manufacturer:	PANTHEON INC, MISSISSAUGA, ONTARIO, CANADA	Manufacturer:	LABESFAL, LABORATORIOS ALIRO S.A, FRESENIUS KABI GROUP, SANTIAGO DE BESTEIROS, PORTUGAL	Manufacturer:	LABESFAL, LABORATORIOS ALIRO S.A, FRESENIUS KABI GROUP, SANTIAGO DE BESTEIROS, PORTUGAL
Packer:	PATHEON INC, MISSISSAUGA, ONTARIO, CANADA F. HOFFMANN-LA ROCHE LTD KAISERAUGST SWITZERLAND AKACIA HEALTH CARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA	Packer:	LABESFAL, LABORATORIOS ALIRO S.A, FRESENIUS KABI GROUP, SANTIAGO DE BESTEIROS, PORTUGAL	Packer:	LABESFAL, LABORATORIOS ALIRO S.A, FRESENIUS KABI GROUP, SANTIAGO DE BESTEIROS, PORTUGAL
Laboratory: FPRC:	PATHEON INC, MISSISSAUGA, ONTARIO, CANADA ROCHE PHARMA AG, GRENZACH-WYHLEN, GERMANY AKACIA HEALTH CARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA	Laboratory: FPRC:	LABESFAL, LABORATORIOS ALIRO S.A, FRESENIUS KABI GROUP, SANTIAGO DE BESTEIROS, PORTUGAL SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA KHULULEKANI LABORATORY SERVICES, RANDJESPAK, MIDRAND, RSA	Laboratory: FPRC:	LABESFAL, LABORATORIOS ALIRO S.A, FRESENIUS KABI GROUP, SANTIAGO DE BESTEIROS, PORTUGAL SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA KHULULEKANI LABORATORY SERVICES, RANDJESPAK, MIDRAND, RSA
FPRR:	ROCHE PRODUCTS (PTY) LTD, ILLOVO, JOHANNESBURG, RSA	FPRR:	FRESENIUS KABI SOUTH AFRICA (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	FPRR:	FRESENIUS KABI SOUTH AFRICA (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MMRF 15	
Registration number:	43/5.4.1/0622	Registration number:	43/5.4.1/0624
Name of medicine:	PRAMEX 0,5	Name of medicine:	PRAMEX 0,125
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: PRAMIPEXOLE DIHYDROCHLORIDE 0,5 mg	Active ingredients:	EACH TABLET CONTAINS: PRAMIPEXOLE DIHYDROCHLORIDE 0,125 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AKACIA HEALTHCARE (PTY) LTD	Applicant:	AKACIA HEALTHCARE (PTY) LTD
Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE
Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE	Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE
Laboratory:	FPRC: PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH- WEST UNIVERSITY, POTCHEFSTROOM, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory:	FPRC PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA	FPRC/FPRR:	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA
Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/5.4.1/0625	Registration number:	43/5.4.1/0626	Registration number:	43/7.1.3/0652
Name of medicine:	PRAMEX 1.0	Name of medicine:	PRAMEX 1.5	Name of medicine:	PERIPREX 2/0.625
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: PRAMIPEXOLE DIHYDROCHLORIDE 1.0 mg	Active ingredients:	EACH TABLET CONTAINS: PRAMIPEXOLE DIHYDROCHLORIDE 1.5 mg	Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL TERT-BUTYLAMINE 2.0 mg INDAPAMIDE 0.625 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	AKACIA HEALTHCARE (PTY) LTD	Applicant:	AKACIA HEALTHCARE (PTY) LTD	Applicant:	ARROW PHARMA SOUTH AFRICA (PTY) LTD
Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA
Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE	Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE	Packer:	ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA DIVPHARM MANUFACTURING AND PACKAGING (PTY) LTD, LONGDALE, INDUSTRIAL, RSA TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA
Laboratory:	FPRC	Laboratory:	FPRC	Laboratory:	FPRC
	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA		PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA		ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA SEDEK AGRICHEM, SILVERTONDALE, PRETORIA, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRC		FPRC/FPFR		FPFR	
	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA		AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA		ARROW PHARMA SOUTH AFRICA (PTY) LTD, NORWICH CLOSE, SANDOWN, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/7.1.3/0653	Registration number:	43/7.1.3/0654	Registration number:	43/7.1.3/0655
Name of medicine:	PERIPREX 4/1,25	Name of medicine:	ARIPREL PLUS 2/0,625	Name of medicine:	ARIPREL PLUS 4/1,25
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL TERT-BUTYLAMINE 4,0 mg INDAPAMIDE 1,25 mg	Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL TERT-BUTYLAMINE 2,0 mg INDAPAMIDE 0,625 mg	Active ingredients:	EACH TABLET CONTAINS: PERINDOPRIL TERT-BUTYLAMINE 4,0 mg INDAPAMIDE 1,25 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ARROW PHARMA SOUTH AFRICA (PTY) LTD	Applicant:	ARROW PHARMA SOUTH AFRICA (PTY) LTD	Applicant:	ARROW PHARMA SOUTH AFRICA (PTY) LTD
Manufacturer:	ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA	Manufacturer:	ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA	Manufacturer:	ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA
Packer:	ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA	Packer:	ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA	Packer:	ARROW PHARMA (MALTA) LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA
Laboratory:	FPRR: ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA SEDEK AGRICHEM, SILVERTONDALE, PRETORIA, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory:	FPRR: ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA SEDEK AGRICHEM, SILVERTONDALE, PRETORIA, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory:	FPRC: ARROW LABORATORIES LIMITED, CROYDON SOUTH, VICTORIA, AUSTRALIA ARROW PHARMA (MALTA) LIMITED, HAL FAR INDUSTRIAL ESTATE, BIRZEBBUGIA, MALTA SEDEK AGRICHEM, SILVERTONDALE, PRETORIA, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/32.16/0681	Registration number:	43/30.1/0727	Registration number:	43/30.1/0728
Name of medicine:	SEQUENT PLEASE	Name of medicine:	STELARA 45 mg	Name of medicine:	STELARA 90 mg
Dosage form:	PTCA BALLOON CATHETER	Dosage form:	SOLUTION FOR INJECTION	Dosage form:	SOLUTION FOR INJECTION
Active ingredients:	EACH mm ² SURFACE CONTAINS: PACLITAXEL 3.0 µg	Active ingredients:	EACH VIAL CONTAINS: USTEKINUMAB 45.0 mg	Active ingredients:	EACH VIAL CONTAINS: USTEKINUMAB 90.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	B BRAUN MEDICAL (PTY) LTD	Applicant:	JANSSEN PHARMACEUTICA (PTY) LTD	Applicant:	JANSSEN PHARMACEUTICA (PTY) LTD
Manufacturer:	B. BRAUN MELSUNGEN AG – VASCULAR SYSTEMS, BERLIN, GERMANY B. BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY HEMOTEC GmbH, WURSELEN, GERMANY	Manufacturer:	CILAG AG, SCHAFFHAUSEN, SWITZERLAND BAXTER PHARMACEUTICAL SOLUTIONS, BLOOMINGTON, INDIANA, USA	Manufacturer:	CILAG AG, SCHAFFHAUSEN, SWITZERLAND BAXTER PHARMACEUTICAL SOLUTIONS, BLOOMINGTON, INDIANA, USA
Packer:	B. BRAUN MELSUNGEN AG – VASCULAR SYSTEMS, BERLIN, GERMANY B. BRAUN MELSUNGEN AG, MELSUNGEN, GERMANY HEMOTEC GmbH, WURSELEN, GERMANY	Packer:	CILAG AG, SCHAFFHAUSEN, SWITZERLAND BAXTER PHARMACEUTICAL SOLUTIONS, BLOOMINGTON, INDIANA, USA	Packer:	CILAG AG, SCHAFFHAUSEN, SWITZERLAND BAXTER PHARMACEUTICAL SOLUTIONS, BLOOMINGTON, INDIANA, USA
Laboratory: FPRC:	BIOPROOF AG, MUNCHEN, GERMANY MEDICAL DEVICE SERVICES – Dr ROSSBERGER GmbH, GILCHING, GERMANY M & L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	CILAG AG, SCHAFFHAUSEN, SWITZERLAND CENTOR B.V., LEIDEN, THE NETHERLANDS	Laboratory: FPRC:	CILAG AG, SCHAFFHAUSEN, SWITZERLAND CENTOR B.V., LEIDEN, THE NETHERLANDS
FPRR:	B BRAUN MEDICAL (PTY) LTD, FOURWAYS, RANDBURG, RSA	FPRR:	JANSSEN PHARMACEUTICA (PTY) LTD, WOODMEAD, RSA	FPRR:	JANSSEN PHARMACEUTICA (PTY) LTD, WOODMEAD, RSA
Shelf-life:	24 months	Shelf-life:	12 months (Vial) 18 months (Pre-filled syringe)	Shelf-life:	12 months (Vial) 18 months (Pre-filled syringe)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/20.1.1/0783	Registration number:	43/20.1.1/0784	Registration number:	43/5.7.1/0814
Name of medicine:	LEVOTAV IV 250	Name of medicine:	LEVOTAV IV 500	Name of medicine:	CETLEV 5
Dosage form:	INFUSION	Dosage form:	INFUSION	Dosage form:	TABLET
Active ingredients:	EACH 50.0 ml SOLUTION CONTAINS: LEVOFLOXACIN 250.0 mg	Active ingredients:	EACH 50.0 ml SOLUTION CONTAINS: LEVOFLOXACIN 500.0 mg	Active ingredients:	EACH TABLET CONTAINS: Levofloxacin dihydrochloride 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	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ADCOCK INGRAM LIMITED	Applicant:	ADCOCK INGRAM LIMITED	Applicant:	DR REDDY'S LABORATORIES (PTY) LTD
Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Manufacturer:	DR REDDY'S LABORATORIES LIMITED, QUTHUBULLAPUR MANDAL, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Packer:	DR REDDY'S LABORATORIES LIMITED, QUTHUBULLAPUR MANDAL, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA DRA PHARMACEUTICALS, IRENE, CENTURION, RSA DIPHARM MANUFACTURING AND PACKAGING (PTY) LTD, LONGDALE, INDUSTRIAL, RSA TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA
Laboratory, FPRC:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Laboratory, FPRC:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Laboratory, FPRC:	DR REDDY'S LABORATORIES LIMITED, QUTHUBULLAPUR MANDAL, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA
FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA	FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA	FPRR:	DR REDDY'S LABORATORIES (PTY) LTD, SOUTH WING, THE PLACE, SANDTON, RSA
Shelf-life:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA 24 months (Provisional)	Shelf-life:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA 24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/5.7.1/0815	Registration number:	43/2.5/0840	Registration number:	43/20.1.1/0859
Name of medicine:	DRL LEVOCETIRIZINE 5	Name of medicine:	GULF CARBAMAZEPINE 200	Name of medicine:	CLARITHEXAL 500 XL
Dosage form:	TABLET	Dosage form:	TABLETS	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: Levocetirizine dihydrochloride 5.0 mg	Active ingredients:	EACH TABLET CONTAINS: CARBAMAZEPINE 200.0 mg	Active ingredients:	EACH TABLET CONTAINS: CLARITHROMYCIN 500.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	DR REDDY'S LABORATORIES (PTY) LTD	Applicant:	GULF DRUG COMPANY (PTY) LTD	Applicant:	SANDOZ (PTY) LTD
Manufacturer:	DR REDDY'S LABORATORIES LIMITED, QUTHUBULLAPUR MANDAL, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	ALEMBIC LIMITED (FORMULATION DIVISION), TAL-HALOL, DISTRICT PANCHMAHALS, GUJARAT, INDIA	Manufacturer:	LEK PHARMACEUTICALS d.d, LJUBLJANA, SLOVENIA
Packer:	DR REDDY'S LABORATORIES LIMITED, QUTHUBULLAPUR MANDAL, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA DRA PHARMACEUTICALS, IRENE, CENTURION, RSA DIVPHARM MANUFACTURING AND PACKAGING (PTY) LTD, LONGDALE, INDUSTRIA, RSA TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA	Packer:	ALEMBIC LIMITED (FORMULATION DIVISION), TAL-HALOL, DISTRICT PANCHMAHALS, GUJARAT, INDIA	Packer:	LEK PHARMACEUTICALS d.d, LJUBLJANA, SLOVENIA TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA
laboratory: FPRC:	DR REDDY'S LABORATORIES LIMITED, QUTHUBULLAPUR MANDAL, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory: FPRC:	ALEMBIC LIMITED (FORMULATION DIVISION), TAL-HALOL, DISTRICT PANCHMAHALS, GUJARAT, INDIA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA SWIFT MICRO LABORATORIES (PTY) LTD, CONSTANTIA, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORY (PTY) LTD, MOREHILL, BENONI, RSA	Laboratory: FPRC	LEK PHARMACEUTICALS d.d, LJUBLJANA, SLOVENIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA
FPRR:	DR REDDY'S LABORATORIES (PTY) LTD, SOUTH WING, THE PLACE, SANDTON, RSA	FPRR:	GULF DRUG COMPANY (PTY) LTD, MOUNT EDGEcombe, RSA	FPRC/FPRR:	SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/20.1.1/0860	Registration number:	43/20.1.1/0955	Registration number:	43/20.1.1/0956
Name of medicine:	SANDOX CLARITHROMYCIN 500 XL	Name of medicine:	MYLAN CIPROFLOXACIN 200 mg/100 ml	Name of medicine:	MYLAN CIPROFLOXACIN 400 mg/200 ml
Dosage form:	TABLET	Dosage form:	INFUSION	Dosage form:	INFUSION
Active ingredients:	EACH TABLET CONTAINS: CLARITHROMYCIN 500,0 mg	Active ingredients:	EACH 100,0 ml SOLUTION CONTAINS: CIPROFLOXACIN 200,0 mg	Active ingredients:	EACH 200,0 ml SOLUTION CONTAINS: CIPROFLOXACIN 400,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANDOZ (PTY) LTD	Applicant:	SCP PHARMACEUTICALS (PTY) LTD	Applicant:	SCP PHARMACEUTICALS (PTY) LTD
Manufacturer:	LEK PHARMACEUTICALS d.d., LJUBLJANA, SLOVENIA	Manufacturer:	MACO PRODUCTIONS SAS, MOUVAUX, FRANCE	Manufacturer:	MACO PRODUCTIONS SAS, MOUVAUX, FRANCE
Packer:	LEK PHARMACEUTICALS d.d., LJUBLJANA, SLOVENIA TECHNIKON LABORATORIES (PTY) LTD, ROBERTVILLE, FLORIDA, RSA SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	Packer:	MACO PRODUCTIONS SAS, MOUVAUX, FRANCE	Packer:	MACO PRODUCTIONS SAS, MOUVAUX, FRANCE
Laboratory: FPRC:	LEK PHARMACEUTICALS d.d., LJUBLJANA, SLOVENIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA	Laboratory: FPRC:	MACO PRODUCTIONS SAS, MOUVAUX, FRANCE MERCK PHARMACEUTICALS MANUFACTURING (PTY) LTD, WADEVILLE, GERMISTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA ACORN PHARMACEUTICALS (PTY) LTD, NORTH RIDING, JOHANNESBURG, RSA	Laboratory: FPRC:	MACO PRODUCTIONS SAS, MOUVAUX, FRANCE MERCK PHARMACEUTICALS MANUFACTURING (PTY) LTD, WADEVILLE, GERMISTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA ACORN PHARMACEUTICALS (PTY) LTD, NORTH RIDING, JOHANNESBURG, RSA
FPRR:	SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	FPRR:	SCP PHARMACEUTICALS (PTY) LTD, MODDERFONTEIN, JOHANNESBURG, RSA	FPRR:	SCP PHARMACEUTICALS (PTY) LTD, MODDERFONTEIN, JOHANNESBURG, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15	
Registration number:	43/21.2/1055	Registration number:	43/21.2/1056
Name of medicine:	GALVUS MET 50 mg/850 mg	Name of medicine:	GALVUS MET 50 mg/1 000 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: VILDAGLIPTIN 50.0 mg METFORMIN HYDROCHLORIDE 850.0 mg	Active ingredients:	EACH TABLET CONTAINS: VILDAGLIPTIN 50.0 mg METFORMIN HYDROCHLORIDE 1 000.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	NOVARTIS SA (PTY) LTD	Applicant:	NOVARTIS SA (PTY) LTD
Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND	Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND
Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND	Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND
Laboratory: FPRC:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory: FPRC:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA
FPRR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	FPRR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

Registration number:	43/7.1/1127
Name of medicine:	TRITACE PLUS 2,5/12,5 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RAMIPRIL 2,5 mg HYDROCHLOROTHIAZIDE 12,5 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	SANOFLAVENTIS DEUTSCHLAND GmbH, FRANKFURT, GERMANY SANOFLAVENTIS S.p.A., SCOPPIO, ITALY
Packer:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory: FPRC:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
FPRR:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Shelf-life:	36 months
Date of registration:	27 July 2012

MRF 15		MRF 15	
Registration number:	43/7.1/1128	Registration number:	43/7.1/1129
Name of medicine:	TRITACE PLUS 5/12,5 mg	Name of medicine:	TRITACE PLUS 10/12,5 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RAMIPRIL 5,0 mg HYDROCHLOROTHIAZIDE 12,5 mg	Active ingredients:	EACH TABLET CONTAINS: RAMIPRIL 10,0 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:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD	Applicant:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	SANOFLAVENTIS DEUTSCHLAND GmbH, FRANKFURT, GERMANY SANOFLAVENTIS S.p.A., SCOPPIO, ITALY	Manufacturer:	SANOFLAVENTIS DEUTSCHLAND GmbH, FRANKFURT, GERMANY SANOFLAVENTIS S.p.A., SCOPPIO, ITALY
Packer:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory: FPRC:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM	Laboratory: FPRC:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
FPRR:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	FPRR:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Shelf-life:	36 months	Shelf-life:	36 months
Date of registration:	27 July 2012	Date of registration:	27 July 2012

Registration number:	43/7.1/1130
Name of medicine:	TRITACE PLUS 5/25 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RAMIPRIL 5,0 mg HYDROCHLOROTHIAZIDE 25,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD
Manufacturer:	SANOFLAVENTIS DEUTSCHLAND GmbH, FRANKFURT, GERMANY SANOFLAVENTIS S.p.A., SCOPPIO, ITALY
Packer:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory: FPRC:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
FPRR:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Shelf-life:	36 months
Date of registration:	27 July 2012

MRF 15		MRF 15		MRF 15	
Registration number:	43/7.1/1131	Registration number:	44/7.1.3/0075	Registration number:	44/7.1.3/0076
Name of medicine:	TRITACE PLUS 10/25 mg	Name of medicine:	CO EXFORGE 5 mg/160 mg/12,5 mg	Name of medicine:	CO EXFORGE 5 mg/160 mg/25 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: RAMIPRIL 10,0 mg HYDROCHLOROTHIAZIDE 25,0 mg	Active ingredients:	EACH TABLET CONTAINS: Amlodipine besylate equivalent to Amlodipine 5,0 mg Valsartan 160,0 mg Hydrochlorothiazide 12,5 mg	Active ingredients:	EACH TABLET CONTAINS: Amlodipine besylate equivalent to Amlodipine 5,0 mg Valsartan 160,0 mg Hydrochlorothiazide 25,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD	Applicant:	NOVARTIS SA (PTY) LTD	Applicant:	NOVARTIS SA (PTY) LTD
Manufacturer:	SANOFLAVENTIS DEUTSCHLAND GmbH, FRANKFURT, GERMANY SANOFLAVENTIS S.p.A., SCOPPIO, ITALY	Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND	Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND
Packer:	SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND	Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND
Laboratory:	FPRC: SANOFLAVENTIS S.p.A., SCOPPIO, ITALY WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM	Laboratory:	FPRC: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory:	FPRC: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA
FPRC:	SANOFLAVENTIS SOUTH AFRICA (PTY) LTD, MIDRAND, RSA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	FPRC:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	FPRC:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	36 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 July 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/7.1.3/0077	Registration number:	44/7.1.3/0078	Registration number:	44/7.1.3/0079
Name of medicine:	CO EXFORGE 10 mg/160 mg/12,5 mg	Name of medicine:	CO EXFORGE 10 mg/160 mg/25 mg	Name of medicine:	CO EXFORGE 10 mg/320 mg/25 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: Amlodipine besylate equivalent to Amlodipine 10,0 mg Valsartan 160,0 mg Hydrochlorothiazide 12,5 mg	Active ingredients:	EACH TABLET CONTAINS: Amlodipine besylate equivalent to Amlodipine 10,0 mg Valsartan 160,0 mg Hydrochlorothiazide 25 mg	Active ingredients:	EACH TABLET CONTAINS: Amlodipine besylate equivalent to Amlodipine 10,0 mg Valsartan 320,0 mg Hydrochlorothiazide 25 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	NOVARTIS SA (PTY) LTD	Applicant:	NOVARTIS SA (PTY) LTD	Applicant:	NOVARTIS SA (PTY) LTD
Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND	Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND	Manufacturer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND
Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND	Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND	Packer:	NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND ALLPACK AG, REINACH, SWITZERLAND KONAPHARMA AG, PRATTELN, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA IVERS-LEE AG, BURGDORF, SWITZERLAND
Laboratory:	FPRC: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory:	FPRC: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA	Laboratory:	FPRC: NOVARTIS PHARMA STEIN AG, STEIN, SWITZERLAND NOVARTIS PHARMANALYTICA SA, LOCARNO, SWITZERLAND SANDOZ SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA
FPRR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	FPRR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA	FPRR:	NOVARTIS SA (PTY) LTD, SPARTAN, KEMPTON PARK, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15	
Registration number:	44/10.2.2/0200	Registration number:	44/20.1.1/0220
Name of medicine:	MONTEWIN 5 mg	Name of medicine:	PROPAN LEVOFLOXACIN IV 250
Dosage form:	CHEWABLE TABLETS	Dosage form:	INFUSION
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg	Active ingredients:	EACH 50.0 ml SOLUTION CONTAINS: LEVOFLOXACIN 250.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD	Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	NYCOMED, LYSZKOWICE, POLAND	Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Packer:	NYCOMED, LYSZKOWICE, POLAND WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Laboratory: FPRC:	NYCOMED, LYSZKOWICE, POLAND RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory: FPRC:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
FPRC/FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPRR:	WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA	FPRR:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15

Registration number:	44/20.1.1/0221
Name of medicine:	PROPAN LEVOFLOXACIN IV 500
Dosage form:	INFUSION
Active ingredients:	EACH 100,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 500,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Laboratory:	FPRC: ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
FPRC/FPFR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPFR:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF15

Registration number:	44/20.1.1/0222
Name of medicine:	RESTAN LEVOFLOXACIN IV 250
Dosage form:	INFUSION
Active ingredients:	EACH 50,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 250,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Laboratory:	FPRC: ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
FPRC/FPFR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPFR:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15

Registration number:	44/20.1.1/0223
Name of medicine:	RESTAN LEVOFLOXACIN IV 500
Dosage form:	INFUSION
Active ingredients:	EACH 100,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 500,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
Laboratory:	FPRC: ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND
FPFR/FPFR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA
FPFR:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/20.1.1/0224	Registration number:	44/20.1.1/0225	Registration number:	44/10.2.2/0257
Name of medicine:	COVAN LEVOFLOXACIN IV 250	Name of medicine:	COVAN LEVOFLOXACIN IV 500	Name of medicine:	MONTEFLO 4
Dosage form:	INFUSION	Dosage form:	INFUSION	Dosage form:	CHEWABLE TABLET
Active ingredients:	EACH 50,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 250,0 mg	Active ingredients:	EACH 100,0 ml SOLUTION CONTAINS: LEVOFLOXACIN 500,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ADCOCK INGRAM LIMITED	Applicant:	ADCOCK INGRAM LIMITED	Applicant:	TRINITY PHARMA SA (PTY) LTD
Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Manufacturer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Manufacturer:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Packer:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Packer:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
Laboratory: FPRC:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Laboratory: FPRC:	ACS DOBFAR INFO SA, CAMPASCIO, SWITZERLAND	Laboratory: FPRC:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA	FPRC/FPRR:	ADCOCK INGRAM CRITICAL CARE (PTY) LTD, AEROTON, JOHANNESBURG, RSA ADCOCK INGRAM LIMITED - RESEARCH & DEVELOPMENT, AEROTON, JOHANNESBURG, RSA	FPRR:	INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA
FPRR:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA	FPRR:	ADCOCK INGRAM LIMITED, ERAND GARDENS, MIDRAND, RSA	FPRR:	TRINITY PHARMA SA (PTY) LTD, MURRAYFIELD, PRETORIA, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15

Registration number:	44/10.2.2/0262
Name of medicine:	MONTEFLO 5
Dosage form:	CHEWABLE TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	TRINITY PHARMA SA (PTY) LTD
Manufacturer:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
Packer:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
Laboratory:	FPRC: TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA
FPRR:	TRINITY PHARMA SA (PTY) LTD, MURRAYFIELD, PRETORIA, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF15

Registration number:	44/10.2.2/0286
Name of medicine:	MONTEFLO 10
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	TRINITY PHARMA SA (PTY) LTD
Manufacturer:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
Packer:	TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA
Laboratory:	FPRC: TORRENT PHARMACEUTICALS LTD, MEHSANA, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA
FPRR:	TRINITY PHARMA SA (PTY) LTD, MURRAYFIELD, PRETORIA, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15

Registration number:	44/1.3.2/0339
Name of medicine:	ALTREX
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: NALTREXONE HYDROCHLORIDE 50,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	PHARMAPLAN (PTY) LTD
Manufacturer:	MALLINCKRODT INC, HOBART, NEW YORK, USA
Packer:	MALLINCKRODT INC, HOBART, NEW YORK, USA
Laboratory:	FPRC MALLINCKRODT INC, HOBART, NEW YORK, USA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	PHARMAPLAN (PTY) LTD, MIDRAND, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/10.2.2/0384	Registration number:	44/10.2.2/0431	Registration number:	44/10.2.2/0493
Name of medicine:	MONTEWIN 4 mg	Name of medicine:	LEUKOBLOC 10 mg	Name of medicine:	MINAIR 4
Dosage form:	CHEWABLE TABLETS	Dosage form:	TABLET	Dosage form:	CHEWABLE TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4.0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10.0 mg	Active ingredients:	EACH TABLET CONTAINS: Montelukast sodium equivalent to Montelukast 4.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD	Applicant:	PHARMACARE LIMITED	Applicant:	AKACIA HEALTHCARE (PTY) LTD
Manufacturer:	NYCOMED, LYSZKOWICE, POLAND	Manufacturer:	ABDI IBRAHIM ILAÇ SAN, VE TIC. A.Ş., HADIMKOY/İSTANBUL, TURKEY PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	Manufacturer:	PHARMATHEN S.A. PALLINI, ATTIKIS, GREECE
Packer:	NYCOMED, LYSZKOWICE, POLAND WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	Packer:	ABDI IBRAHIM ILAÇ SAN, VE TIC. A.Ş., HADIMKOY/İSTANBUL, TURKEY PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	Packer:	PHARMATHEN S.A. PALLINI, ATTIKIS, GREECE FAMAR S.A. ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A. ATHENS-JAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE
Laboratory:	FPRC: NYCOMED, LYSZKOWICE, POLAND RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory:	FPRC: SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA	Laboratory:	FPRC: PHARMATHEN S.A. PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORY (PTY) LTD, MOREHILL, BENONI, RSA PHARMASPEC CONSULTING LABORATORIES, FERNDALE, RANDBURG, RSA
FPRC:	WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA	FPRC/FPRC:	PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE EAST LONDON, WILSONIA, EAST LONDON, RSA	FPRC:	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA
Shelf-life:	WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA	FPRC:	PHARMACARE LIMITED, WOODMEAD, SANDTON, RSA	Shelf-life:	24 months
Date of registration:	24 months (Provisional) 27 JULY 2012	Shelf-life:	24 months (Provisional)	Date of registration:	27 JULY 2012

MRF 15	MRF 15		MRF 15		MRF 15	
	Registration number:	44/10.2.2/0494	Registration number:	44/10.2.2/0495	Registration number:	44/20.1.1/0582
Name of medicine:	MINAIR 5	MINAIR 10	Name of medicine:	MINAIR 10	Name of medicine:	ZOCEF 250 mg
Dosage form:	CHEWABLE TABLET	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: Montelukast sodium equivalent to Montelukast: 5,0 mg	EACH TABLET CONTAINS: Montelukast sodium equivalent to Montelukast: 10,0 mg	Active ingredients:	EACH TABLET CONTAINS: Montelukast sodium equivalent to Montelukast: 10,0 mg	Active ingredients:	EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 250,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AKACIA HEALTHCARE (PTY) LTD	AKACIA HEALTHCARE (PTY) LTD	Applicant:	AKACIA HEALTHCARE (PTY) LTD	Applicant:	ALKEM LABORATORIES (PTY) LTD
Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA
Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE	Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., ATHENS-LAMIA HIGHWAY, SCHIMATRI, ATHENS, GREECE	Packer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA
Laboratory: FPRC:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORIES (PTY) LTD, MOREHILL, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORY (PTY) LTD, MOREHILL, BENONI, RSA PHARMASPEC CONSULTING LABORATORIES, FERNDALE, RANDBURG, RSA	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORY (PTY) LTD, MOREHILL, BENONI, RSA PHARMASPEC CONSULTING LABORATORIES, FERNDALE, RANDBURG, RSA	Laboratory: FPRC:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SABS COMMERCIAL (PTY) LTD, PHARMACEUTICAL CHEMISTRY DEPARTMENT, GROENKLOOF, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA M&L LABORATORY SERVICES, (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORY (PTY) LTD, MOREHILL, BENONI, RSA PHARMASPEC CONSULTING LABORATORIES, FERNDALE, RANDBURG, RSA	Laboratory: FPRC:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH- WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA
FPRC:	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA	FPRC:	AKACIA HEALTHCARE (PTY) LTD, ISANDO, JOHANNESBURG, RSA	FPRC:	ALKEM LABORATORIES (PTY) LTD, POTCHEFSTROOM, RSA
Shelf-life:	24 months	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 JULY 2012	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/20.1.1/0583	Registration number:	44/10.2.2/0666	Registration number:	44/10.2.2/0667
Name of medicine:	ZOCEF 500 mg	Name of medicine:	MSD MONTELUKAST 4 mg	Name of medicine:	MSD MONTELUKAST 5 mg
Dosage form:	TABLET	Dosage form:	CHEWABLE TABLET	Dosage form:	CHEWABLE TABLET
Active ingredients:	EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 500,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	ALKEM LABORATORIES (PTY) LTD	Applicant:	MSD (PTY) LTD	Applicant:	MSD (PTY) LTD
Manufacturer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA	Manufacturer:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	Manufacturer:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK
Packer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA	Packer:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK MMD HAARLEM, PC HAARLEM, THE NETHERLANDS MSD (PTY) LTD, HALFWAY HOUSE, RSA	Packer:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK MMD HAARLEM, PC HAARLEM, THE NETHERLANDS MSD (PTY) LTD, HALFWAY HOUSE, RSA
Laboratory: FPRC:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH- WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK MMD HAARLEM, PC HAARLEM, THE NETHERLANDS	Laboratory: FPRC:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK MMD HAARLEM, PC HAARLEM, THE NETHERLANDS
FPRR:	ALKEM LABORATORIES (PTY) LTD, POTCHEFSTROOM, RSA	FPRR:	MSD (PTY) LTD, HALFWAY HOUSE, RSA	FPRR:	MSD (PTY) LTD, HALFWAY HOUSE, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/10.2.2/0668	Registration number:	44/10.2.2/0669	Registration number:	44/10.2.2/0844
Name of medicine:	MSD MONTELUKAST 10 mg	Name of medicine:	MSD MONTELUKAST SPRINKLES	Name of medicine:	MONTELUKAST BRIMPHEM 4
Dosage form:	TABLET	Dosage form:	GRANULES	Dosage form:	CHEWABLE TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10.0 mg	Active ingredients:	EACH PACKET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4.0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	MSD (PTY) LTD	Applicant:	MSD (PTY) LTD	Applicant:	BRIMPHEM SA (PTY) LTD
Manufacturer:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK	Manufacturer:	MERCK MANUFACTURING DIVISION, WEST POINT, PENNSYLVANIA, USA	Manufacturer:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA
Packer:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK MMD HAARLEM, PC HAARLEM, THE NETHERLANDS MSD (PTY) LTD, HALFWAY HOUSE, RSA	Packer:	MERCK MANUFACTURING DIVISION, WILSON, NORTH CAROLINA, USA MMD HAARLEM, PC HAARLEM, THE NETHERLANDS MSD (PTY) LTD, HALFWAY HOUSE, RSA	Packer:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA, JOHANNESBURG, RSA
Laboratory: FPRC:	MERCK SHARP & DOHME, CRAMLINGTON, NORTHUMBERLAND, UK MMD HAARLEM, PC HAARLEM, THE NETHERLANDS	Laboratory: FPRC:	MERCK MANUFACTURING DIVISION, WEST POINT, PENNSYLVANIA, USA MMD HAARLEM, PC HAARLEM, THE NETHERLANDS	Laboratory: FPRC:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA CONFARMA FRANCE S.A.R.L., HOMBOURG, FRANCE CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA
FPRR:	MSD (PTY) LTD, HALFWAY HOUSE, RSA	FPRC/FPRR:	MSD (PTY) LTD, HALFWAY HOUSE, RSA	FPRR:	BRIMPHEM SA (PTY) LTD, CLAREMONT, CAPE TOWN, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15	
Registration number:	44/10.2.2/0845	Registration number:	44/10.2.2/0846
Name of medicine:	MONTELUKAST BRIMPARM 5	Name of medicine:	MONTELUKAST BRIMPARM 10
Dosage form:	CHEWABLE TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 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:	BRIMPARM SA (PTY) LTD	Applicant:	BRIMPARM SA (PTY) LTD
Manufacturer:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA	Manufacturer:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA
Packer:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA, JOHANNESBURG, RSA	Packer:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA DIVPHARM MANUFACTURING & PACKAGING, LONGDALE, INDUSTRIA, JOHANNESBURG, RSA
Laboratory: FPRC:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA CONFARMA FRANCE S.A.R.L., HOMBOURG, FRANCE CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	SIEGFRIED LTD, ZOFINGEN, SWITZERLAND SIEGFRIED GENERICS (MALTA) LTD, HAL FAR, MALTA CONFARMA FRANCE S.A.R.L., HOMBOURG, FRANCE CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA
FPRR:	BRIMPARM SA (PTY) LTD, CLAREMONT, CAPE TOWN, RSA	FPRR:	BRIMPARM SA (PTY) LTD, CLAREMONT, CAPE TOWN, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012
Registration number:	44/10.2.2/0889	Registration number:	44/10.2.2/0889
Name of medicine:	MONTELUKAST SPECPHARM 4	Name of medicine:	MONTELUKAST SPECPHARM 4
Dosage form:	CHEWABLE TABLET	Dosage form:	CHEWABLE TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	SPECPHARM (PTY) LTD	Applicant:	SPECPHARM (PTY) LTD
Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE
Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., LAMIA HIGHWAY, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., LAMIA HIGHWAY, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Laboratory: FPRC:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA
FPRR:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	FPRR:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	44/10.2.2/0890	Registration number:	44/10.2.2/0891	Registration number:	44/2.5/0924
Name of medicine:	MONTELUKAST SPECPHARM 5	Name of medicine:	MONTELUKAST SPECPHARM 10	Name of medicine:	GDC CARBAMAZEPINE 200
Dosage form:	CHEWABLE TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10,0 mg	Active ingredients:	EACH TABLET CONTAINS: CARBAMAZEPINE 200,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	SPECPHARM (PTY) LTD	Applicant:	SPECPHARM (PTY) LTD	Applicant:	GULF DRUG COMPANY (PTY) LTD
Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE	Manufacturer:	ALEMbic LIMITED (FORMULATION DIVISION), TAL-HALOL, DISTRICT PANCMAHALS, GUJARAT, INDIA
Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., LAMIA HIGHWAY, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	Packer:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE FAMAR S.A., ANTHOUSA, ATTIKIS, GREECE PHARMANEL PHARMACEUTICALS S.A., LAMIA HIGHWAY, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	Packer:	ALEMbic LIMITED (FORMULATION DIVISION), TAL-HALOL, DISTRICT PANCMAHALS, GUJARAT, INDIA
Laboratory: FPRC:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	PHARMATHEN S.A., PALLINI, ATTIKIS, GREECE SPECPHARM HOLDINGS (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	ALEMbic LIMITED (FORMULATION DIVISION), TAL-HALOL, DISTRICT PANCMAHALS, GUJARAT, INDIA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA SWIFT MICRO LABORATORIES (PTY) LTD, CONSTANTIA, MIDRAND, RSA CONSULTING CHEMICAL LABORATORIES (PTY) LTD, ATLASVILLE, BOKSBURG, RSA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA CONSULTING MICROBIOLOGICAL LABORATORY (PTY) LTD, MOREHILL, BENONI, RSA
FPRC:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	FPRC:	SPECPHARM (PTY) LTD, HALFWAY HOUSE, MIDRAND, RSA	FPRC:	GULF DRUG COMPANY (PTY) LTD, MOUNT EDGEcombe, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	45/10.2.2/0058	Registration number:	45/10.2.2/0059	Registration number:	45/10.2.2/0060
Name of medicine:	DILAIR 4	Name of medicine:	DILAIR 5	Name of medicine:	DILAIR 10
Dosage form:	CHEWABLE TABLET	Dosage form:	CHEWABLE TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 4,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: MONTELUKAST SODIUM EQUIVALENT TO MONTELUKAST 10,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	RANBAXY (S.A.) (PTY) LTD	Applicant:	RANBAXY (S.A.) (PTY) LTD	Applicant:	RANBAXY (S.A.) (PTY) LTD
Manufacturer:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA	Manufacturer:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA	Manufacturer:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA
Packer:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA	Packer:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA	Packer:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA
Laboratory: FPRC:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA KHULULEKANI LABORATORY SERVICES, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA KHULULEKANI LABORATORY SERVICES, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	TORRENT PHARMACEUTICALS LTD, INDRAD, MEHSANA, INDIA CONSULTING CHEMICAL LABORATORIES, (PTY) LTD, ATLASVILLE, BOKSBURG, RSA KHULULEKANI LABORATORY SERVICES, COVENTRY PARK, MIDRAND, RSA
FPRR:	RANBAXY (S.A.) (PTY) LTD, CENTURION, RSA	FPRR:	RANBAXY (S.A.) (PTY) LTD, CENTURION, RSA	FPRR:	RANBAXY (S.A.) (PTY) LTD, CENTURION, RSA
Shelf-life:	24 months	Shelf-life:	24 months	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	45/20.2.8/0082	Registration number:	45/20.2.8/0083	Registration number:	45/20.1.1/0216
Name of medicine:	COTIRAL 100/25 mg	Name of medicine:	COTIRAL 200/50 mg	Name of medicine:	CEFUROXIME ALKEM 250 mg
Dosage form:	TABLET	Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LOPINAVIR 100,0 mg RITONAVIR 25,0 mg	Active ingredients:	EACH TABLET CONTAINS: LOPINAVIR 200,0 mg RITONAVIR 50,0 mg	Active ingredients:	EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 250,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8	Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	AUROBINDO PHARMA (PTY) LTD	Applicant:	AUROBINDO PHARMA (PTY) LTD	Applicant:	ALKEM LABORATORIES (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT 111, QUTHUBULLAPUR MANDAL, RANGA RADDY DISTRICT, HYDERABAD, ANDHRA PRADESH, INDIA	Manufacturer:	AUROBINDO PHARMA LIMITED, UNIT 111, QUTHUBULLAPUR MANDAL, RANGA RADDY DISTRICT, HYDERABAD, ANDHRA PRADESH, INDIA	Manufacturer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA
Packer:	AUROBINDO PHARMA LIMITED, UNIT 111, QUTHUBULLAPUR MANDAL, RANGA RADDY DISTRICT, HYDERABAD, ANDHRA PRADESH, INDIA	Packer:	AUROBINDO PHARMA LIMITED, UNIT 111, QUTHUBULLAPUR MANDAL, RANGA RADDY DISTRICT, HYDERABAD, ANDHRA PRADESH, INDIA	Packer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA
Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT 111, QUTHUBULLAPUR MANDAL, RANGA RADDY DISTRICT, HYDERABAD, ANDHRA PRADESH, INDIA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA KHULLEKANI LABORATORY SERVICES, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	AUROBINDO PHARMA LIMITED, UNIT 111, QUTHUBULLAPUR MANDAL, RANGA RADDY DISTRICT, HYDERABAD, ANDHRA PRADESH, INDIA M&L LABORATORY SERVICES (PTY) LTD, LIMBRO BUSINESS PARK, SANDTON, RSA KHULLEKANI LABORATORY SERVICES, COVENTRY PARK, MIDRAND, RSA	Laboratory: FPRC:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA
FPRR:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRR:	AUROBINDO PHARMA (PTY) LTD, MEYERSDAL, JOHANNESBURG, RSA	FPRR:	ALKEM LABORATORIES (PTY) LTD, POTCHEFSTROOM, RSA
Shelf-life:	24 months (Provisional)	Shelf-life:	24 months (Provisional)	Shelf-life:	24 months
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15		MRF 15		MRF 15	
Registration number:	45/20 1.1/0217	Registration number:	45/20 2/0431	Registration number:	45/20 2.8/0537
Name of medicine:	CEFUROXIME ALKEM 500 mg	Name of medicine:	GSK PNEUMOCOCCAL VACCINE	Name of medicine:	MACLEODS EFAVIRENZ 600 mg
Dosage form:	TABLET	Dosage form:	SUSPENSION FOR INJECTION	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: CEFUROXIME AXETIL EQUIVALENT TO CEFUROXIME 500.0 mg	Active ingredients:	EACH 0.5 ml DOSE CONTAINS: Polysaccharide for Serotype 1 1.0 µg Polysaccharide for Serotype 5 1.0 µg Polysaccharide for Serotype 6B 1.0 µg Polysaccharide for Serotype 7F 1.0 µg Polysaccharide for Serotype 9V 1.0 µg Polysaccharide for Serotype 14 1.0 µg Polysaccharide for Serotype 23F 1.0 µg Polysaccharide for Serotype 4 3.0 µg Polysaccharide for Serotype 18C 3.0 µg Polysaccharide for Serotype 19F 3.0 µg	Active ingredients:	EACH TABLET CONTAINS: EFAVIRENZ 600.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7	Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	ALKEM LABORATORIES (PTY) LTD	Applicant:	GLAXOSMITHKLINE SOUTH AFRICA (PTY) LTD	Applicant:	MACLEODS PHARMACEUTICALS SA (PTY) LTD
Manufacturer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA	Manufacturer:	GLAXOSMITHKLINE BIOLOGICALS S.A, RIXENSART, BELGIUM GLAXOSMITHKLINE BIOLOGICALS S.A, WAVRE, BELGIUM	Manufacturer:	MACLEODS PHARMACEUTICALS LTD, KACHIGAM, DAMAN, INDIA
Packer:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA	Packer:	GLAXOSMITHKLINE BIOLOGICALS S.A, WAVRE, BELGIUM GLAXOSMITHKLINE SOUTH AFRICA (PTY) LTD, EPPING, CAPE TOWN, RSA	Packer:	MACLEODS PHARMACEUTICALS LTD, KACHIGAM, DAMAN, INDIA
Laboratory: FPRC:	ALKEM LABORATORIES LIMITED, AMALIYA, DAMAN, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM, RSA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA	Laboratory: FPRC:	GLAXOSMITHKLINE BIOLOGICALS S.A, RIXENSART, BELGIUM GLAXOSMITHKLINE BIOLOGICALS S.A, WAVRE, BELGIUM	Laboratory: FPRC:	MACLEODS PHARMACEUTICALS LTD, KACHIGAM, DAMAN, INDIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG, RSA SABS COMMERCIAL (PTY) LTD PHARMACEUTICAL CHEMISTRY DEPARTMENT GROENKLOOF, PRETORIA, RSA
FPRR:	ALKEM LABORATORIES (PTY) LTD, POTCHEFSTROOM, RSA	FPRC/FPRR:	GLAXOSMITHKLINE SOUTH AFRICA (PTY) LTD, EPPING, CAPE TOWN, RSA	FPRR:	MACLEODS PHARMACEUTICALS SA (PTY) LTD, WOODMEAD, JOHANNESBURG, RSA
Shelf-life:	24 months	Shelf-life:	36 months stored at 2 – 8 °C	Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012	Date of registration:	27 JULY 2012

MRF 15

Registration number:	45/20.2.8/0766
Name of medicine:	CIPLA LAMIVUDINE 30 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LAMIVUDINE 30,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	CIPLA MEDPRO (PTY) LTD
Manufacturer:	CIPLA LTD, UNIT IV, VERNA, GOA, INDIA
Packer:	CIPLA LTD, UNIT IV, VERNA, GOA, INDIA
Laboratory: FPRC:	CIPLA LTD, UNIT IV, VERNA, GOA, INDIA
FPRR:	CIPLA MEDPRO (PTY) LTD, ROSENPARK, BELLVILLE, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF15

Registration number:	45/20.2.8/0767
Name of medicine:	MEDPRO LAMIVUDINE 30 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LAMIVUDINE 30,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	MEDPRO PHARMACEUTICA (PTY) LTD
Manufacturer:	CIPLA LTD, UNIT IV, VERNA, GOA, INDIA
Packer:	CIPLA LTD, UNIT IV, VERNA, GOA, INDIA
Laboratory: FPRC:	CIPLA LTD, UNIT IV, VERNA, GOA, INDIA
FPRR:	MEDPRO PHARMACEUTICA (PTY) LTD, ROSENPARK, BELLVILLE, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15

Registration number:	45/20.2.8/0815
Name of medicine:	NEVIRAPINE MEDPRO 50 mg TABLETS FOR ORAL SUSPENSION
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: NEVIRAPINE 50,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7, 8
Applicant:	CIPLA MEDPRO (PTY) LTD
Manufacturer:	CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA
Packer:	CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA
Laboratory: FPRC:	CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA
FPRR:	CIPLA MEDPRO (PTY) LTD, ROSENPARK, BELLVILLE, RSA
Shelf-life:	24 months (Provisional)
Date of registration:	27 JULY 2012

MRF 15

Registration number: 45/20.2.8/0816

Name of medicine: NEVIRAPINE MEDPRO 100 mg TABLETS FOR ORAL SUSPENSION

Dosage form: TABLET

Active ingredients: EACH TABLET CONTAINS: NEVIRAPINE 100,0 mg

Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8

Applicant: CIPLA MEDPRO (PTY) LTD

Manufacturer: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

Packer: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

Laboratory, FPRC: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

FPRR: CIPLA MEDPRO (PTY) LTD, ROSEN PARK, BELLVILLE, RSA

Shelf-life: 24 months (Provisional)

Date of registration: 27 JULY 2012

MRF15

Registration number: 45/20.2.8/0817

Name of medicine: NEVIMUNE 50 mg TABLETS FOR ORAL SUSPENSION

Dosage form: TABLET

Active ingredients: EACH TABLET CONTAINS: NEVIRAPINE 50,0 mg

Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8

Applicant: CIPLA MEDPRO (PTY) LTD

Manufacturer: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

Packer: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

Laboratory, FPRC: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

FPRR: CIPLA MEDPRO (PTY) LTD, ROSEN PARK, BELLVILLE, RSA

Shelf-life: 24 months (Provisional)

Date of registration: 27 JULY 2012

MRF 15

Registration number: 45/20.2.8/0818

Name of medicine: NEVIMUNE 100 mg TABLETS FOR ORAL SUSPENSION

Dosage form: TABLET

Active ingredients: EACH TABLET CONTAINS: NEVIRAPINE 100,0 mg

Conditions of registration: 1, 2, 3, 4, 5, 6, 7, 8

Applicant: CIPLA MEDPRO (PTY) LTD

Manufacturer: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

Packer: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

Laboratory, FPRC: CIPLA LTD, UNIT II, PATALGANGA DISTRICT RAIGAD, MAHARASHTRA, INDIA

FPRR: CIPLA MEDPRO (PTY) LTD, ROSEN PARK, BELLVILLE, RSA

Shelf-life: 24 months (Provisional)

Date of registration: 27 JULY 2012

MRF 15

Registration number:	46/20.2.8/0040
Name of medicine:	TENOFOVIR WINTHROP
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: TENOFOVIR DISOPROXIL FUMARATE 300,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD
Manufacturer:	HETERO DRUGS LIMITED, UNIT III, JEEDIMETLA, HYDERABAD, INDIA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Packer:	HETERO DRUGS LIMITED, UNIT III, JEEDIMETLA, HYDERABAD, INDIA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory:	FPRC: HETERO DRUGS LIMITED, UNIT III, JEEDIMETLA, HYDERABAD, INDIA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
	FPRR: WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
	WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA
Shelf-life:	36 months
Date of registration:	27 JULY 2012

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Registration number:	46/20.2.8/0039
Name of medicine:	EFAVIRENZ WINTHROP
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: EFAVIRENZ 600,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6, 7
Applicant:	WINTHROP PHARMACEUTICALS (PTY) LTD
Manufacturer:	HETERO DRUGS LIMITED, UNIT III, JEEDIMETLA, HYDERABAD, INDIA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Packer:	HETERO DRUGS LIMITED, UNIT III, JEEDIMETLA, HYDERABAD, INDIA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
Laboratory:	FPRC: HETERO DRUGS LIMITED, UNIT III, JEEDIMETLA, HYDERABAD, INDIA WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
	FPRR: WINTHROP PHARMACEUTICALS (PTY) LTD, WALTLOO, PRETORIA, RSA
	WINTHROP PHARMACEUTICALS (PTY) LTD, MIDRAND, RSA
Shelf-life:	36 months
Date of registration:	27 JULY 2012