
GENERAL NOTICE ALGEMENE KENNISGEWING

NOTICE 559 OF 2007

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 No. 101 OF 1965)

1. The applicant shall ensure that the medicine is manufactured and controlled in terms of 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 No. 101 of 1965).
5. The registration of this medicine shall be subject to regular review 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 registration dossier is subject to review at intervals as determined by Council.
8. A post-registration inspection must be conducted in the first production batch of the locally manufactured product.
9. A post-registration inspection must be conducted on the first production batch manufactured by each local manufacturer.
10. A post-registration inspection must be conducted on the first production batch of the imported product.
11. Marketing of the product may only commence following a satisfactory post-registration inspection report.
12. One sample of every batch, together with four copies of the protocol for testing of the bulk lot and filling lot, and six copies of the certificate of release issued by a competent authority in the country in which the product was manufactured, must be submitted to the Council for lot release purposes.
13. The expiry date allocated shall be modified by adding a statement that the virus strains are currently recommended for South African usage in the specific year.
14. The strains of the master seed viruses must be approved by the Department of Health for each year.

MRF 15	
Registration number:	37/21.10/0374
Name of medicine:	OVIDREL 250 ug
Dosage form:	INJECTION
Active ingredients:	EACH VIAL CONTAINS: CHORIOGONADOTROPIN ALFA 250,0 ug
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	SERONO SOUTH AFRICA (PTY) LTD
Manufacturer:	INDUSTRIA FARMACEUTICA SERONO S.p.A., BARI, ITALY
Packer:	INDUSTRIA FARMACEUTICA SERONO S.p.A., BARI, ITALY
Laboratory: FPRC:	INDUSTRIA FARMACEUTICA SERONO S.p.A., BARI, ITALY
Shelf-life:	24 months
Date of registration:	13 APRIL 2007

MRF 15	
Registration number:	36/30.3/0057
Name of medicine:	ALBUMIN HUMAN 20 % OCTAPHARMA
Dosage form:	SOLUTION
Active ingredients:	EACH 1 000,0 ml SOLUTION CONTAINS: HUMAN ALBUMIN 200,0 g
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	JE ECKARD
Manufacturer:	OCTAPHARMA PHARMAZEUTIKA, VIENNA, AUSTRIA
Packer:	OCTAPHARMA PHARMAZEUTIKA, VIENNA, AUSTRIA
Laboratory: FPRC:	OCTAPHARMA PHARMAZEUTIKA, VIENNA, AUSTRIA
Shelf-life:	36 months
Date of registration:	13 APRIL 2007

MRF 15

11 MAY 2007

3074.7/0239

Name of medicine: NOVODOL SYRUP

Dosage form: SYRUP

Active ingredients: EACH 5.0 ml SYRUP CONTAINS:
PARACETAMOL 120.0 mg

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

Applicant: MEDICINE DEVELOPERS INTERNATIONAL c.c.

Manufacturer: PHARMANOVA (PVT) LTD, HARARE, ZIMBABWE

Packer: PHARMANOVA (PVT) LTD, HARARE, ZIMBABWE

Laboratory: FPRC: PHARMANOVA (PVT) LTD, HARARE, ZIMBABWE
FPRR: MEDICINE DEVELOPERS INTERNATIONAL c.c, MENLO PARK, PRETORIA

Shelf-life: 24 months

Date of registration 13 APRIL 2007

MRF 16

Registration number: A38/1.2/0618

Name of medicine: CAMOX-SERTRALINE 50 mg

Dosage form: TABLET

Active ingredients: EACH TABLET CONTAINS:
SERTRALINE HYDROCHLORIDE EQUIVALENT TO
SERTRALINE 50.0 mg

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

Applicant: CAMOX PHARMACEUTICALS (PTY) LTD

Manufacturer: INTAS PHARMACEUTICALS LTD, AHMEDABAD,
GUJARAT, INDIA

Packer: INTAS PHARMACEUTICALS LTD, AHMEDABAD,
GUJARAT, INDIA

Laboratory: FPRC: INTAS PHARMACEUTICALS LTD, AHMEDABAD,
GUJARAT, INDIA
SOUTH AFRICAN BUREAU OF STANDARDS,
GROENKLOOF, PRETORIA, RSA
INSTITUTE FOR PHARMACEUTICAL
SERVICES, SILVERTONDALE, RSA
M&L LABORATORIES, ORMONDE, JOHANNESBURG

Shelf-life: FPRR: CAMOX PHARMACEUTICALS, AMALGAM,
JOHANNESBURG, RSA

Date of registration: 24 months
13 APRIL 2007

MPE 15

Registration number:	Registration number:
Name of medicine:	Name of medicine:
Dosage form:	Dosage form:
Active ingredients:	Active ingredients:
Conditions of registration:	Conditions of registration:
Applicant:	Applicant:
Manufacturer:	Manufacturer:
Packer:	Packer:
Laboratory:	Laboratory:
FPRC:	FPRC:
Shelf-life:	Shelf-life:
Date of registration:	Date of registration:

A38/7.1.3/0066

CAMOX-SERTRALINE 100 mg

TABLET

EACH TABLET CONTAINS:
 SERTRALINE HYDROCHLORIDE EQUIVALENT TO
 SERTRALINE 100,0 mg

1, 2, 3, 4, 5, 6

CAMOX PHARMACEUTICALS (PTY) LTD

INTAS PHARMACEUTICALS LTD, AHMEDABAD,
 GUJARAT, INDIA

INTAS PHARMACEUTICALS LTD, AHMEDABAD
 GUJARAT, INDIA

FPRC:

INTAS PHARMACEUTICALS LTD, AHMEDABAD,
 GUJARAT, INDIA
 SOUTH AFRICAN BUREAU OF STANDARDS,
 GROENKLOOF, PRETORIA, RSA
 INSTITUTE FOR PHARMACEUTICAL SERVICES,
 SILVERTONDALE, RSA
 M&L LABORATORIES, ORMONDE, JOHANNESBURG

FPRC:

CAMOX PHARMACEUTICALS, AMALGAM,
 JOHANNESBURG, RSA

Shelf-life:

24 months

Date of registration:

13 APRIL 2007

Registration number:

A38/7.1.3/0066

ZESTOZIDE 10

TABLET

EACH TABLET CONTAINS:
 LISINOPRIL DIHYDRATE EQUIVALENT TO
 LISINOPRIL 10,0 mg
 HYDROCHLOROTHIAZIDE 12,5 mg

Conditions of registration:

1, 2, 3, 4, 5, 6

Applicant:

XIXIA PHARMACEUTICALS (PTY) LTD

Manufacturer:

LABORATORIOS LESVI SL, BARCELONA, SPAIN

Packer:

LABORATORIOS LESVI SL, BARCELONA, SPAIN
 MERCK FARMA y QUIMICA S.A., BARCELONA, SPAIN
 GENERICS UK LTD, STATION CLOSE,
 HERTFORDSHIRE, UK
 GERARD LABORATORIES, DUBLIN, IRELAND
 MERCK PHARMACEUTICAL MANUFACTURING,
 WADEVILLE, GERMISTON

Laboratory: FPRC:

LABORATORIOS LESVI SL, BARCELONA, SPAIN
 MERCK FARMA y QUIMICA S.A., BARCELONA, SPAIN
 GENERICS UK LTD, STATION CLOSE,
 HERTFORDSHIRE, UK
 GERARD LABORATORIES, DUBLIN, IRELAND
 MERCK PHARMACEUTICAL MANUFACTURING,
 WADEVILLE, GERMISTON
 RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY,
 NORTH WEST UNIVERSITY, POTCHEFSTROOM

FPRC:

XIXIA PHARMACEUTICALS, MODDERFORTEIN

Shelf-life:

36 months

Date of registration:

13 APRIL 2007

MRF 15

Registration	A39/7.1.3/0067
Name of me	ZESTOZIDE 20
Dosage form	TABLET
Active ingred	EACH TABLET CONTAINS: LISINOPRIL DIHYDRATE EQUIVALENT TO 20,0 mg HYDROCHLOROTHIAZIDE 12,5 mg
Co	1, 2, 3, 4, 5, 6
Ap	XIXIA PHARMACEUTICALS (PTY) LTD
Ma	LABORATORIOS LESVI SL, BARCELONA, SPAIN
Pa	LABORATORIOS LESVI SL, BARCELONA, SPAIN MERCK FARMA y QUIMICA S.A., BARCELONA, SPAIN GENERICS UK LTD, STATION CLOSE, HERTFORDSHIRE, UK GERARD LABORATORIES, DUBLIN, IRELAND MERCK PHARMACEUTICAL MANUFACTURING, WADEVILLE, GERMISTON LABORATORIOS LESVI SL, BARCELONA, SPAIN MERCK FARMA y QUIMICA S.A., BARCELONA, SPAIN GENERICS UK LTD, STATION CLOSE, HERTFORDSHIRE, UK GERARD LABORATORIES, DUBLIN, IRELAND MERCK PHARMACEUTICAL MANUFACTURING, WADEVILLE, GERMISTON RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH WEST UNIVERSITY, POTCHEFSTROOM
	XIXIA PHARMACEUTICALS, MODDERFORTEIN
	36 months
	13 April 2007

MRF 15

registration number:	A39/11.5/0261
Name of medicine:	LAXADOR SENNA 7,5 mg TABLETS
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: CALCIUM SENNOSIDES EQUIVALENT TO 7,5 mg SENNOSIDES
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	PHARMACARE LIMITED
Manufacturer:	PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE, WILSONIA, EAST LONDON
Packer:	PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE, WILSONIA, EAST LONDON
Laboratory: FPRC:	SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA, RSA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
FPRC/FPRR	PHARMACARE LIMITED, KORSTEN, PORT ELIZABETH, RSA ASPEN PHARMACARE, WILSONIA, EAST LONDON
Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007

MRF 15

Registration number:	A39/20.2.3/0325
Name of medicine	ANTIB-4
Dosage form	TABLET
Active ingredients:	EACH TABLET CONTAINS: RIFAMPICIN 150,0 mg ISONIAZID 75,0 mg PYRAZINAMIDE 400,0 mg ETHAMBUTOL HYDROCHLORIDE 275,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	MEDICINE DEVELOPERS INTERNATIONAL cc
Manufacturer:	RUSAN PHARMA LTD, GANDHIDHAM-KUTCH, INDIA
Packer:	RUSAN PHARMA LTD, GANDHIDHAM-KUTCH, INDIA
Laboratory:	FPRC: RUSAN PHARMA LTD, GANDHIDHAM-KUTCH, INDIA CONSULTING CHEMICAL LABORATORIES, STAR STREET, BOKSBURG, RSA
Shelf-life:	MDI cc, MENLOPARK, PRETORIA, RSA 24 months
Date of registration:	2 FEBRUARY 2007

MRF 15

Registration number:	A39/20.2.8/0345
Name of medicine:	IMMUNAC 200 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: ACYCLOVIR 200,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	CAMOX PHARMACEUTICALS (PTY) LTD
Manufacturer:	MEDREICH STERILAB LTD, VIRGONAR BANGALORE, INDIA
Packer:	MEDREICH STERILAB LTD, VIRGONAR BANGALORE, INDIA
Laboratory:	FPRC: MEDREICH STERILAB LTD, VIRGONAR BANGALORE, INDIA SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA M&L LABORATORIES, ORMONDE, JOHANNESBURG
Shelf-life:	FPRR: CAMOX PHARMACEUTICALS, AMALGAM, JOHANNESBURG, RSA 24 months (provisional)
Date of registration	13 APRIL 2007

MRF 15	MRF 15
Registration number:	A39/5.8/0450
Name of medicine:	EFFERCO-C
Dosage form:	TABLET
Active ingredients:	EACH EFFERVESCENT TABLET CONTAINS: PARACETAMOL 500,0 mg SODIUM ASCORBATE EQUIVALENT TO 250,0 mg VITAMIN C 250,0 mg CHLORPHENAMINE MALEATE 2,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	PHARMA DYNAMICS (PTY) LTD
Manufacturer:	E-PHARMA TRENTO S.p.A., TRENTO, ITALY
Packer:	E-PHARMA TRENTO S.p.A., TRENTO, ITALY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA PHARMACEUTICAL ENTERPRISES, N'DABENI, RSA
Laboratory: FPRC:	E-PHARMA TRENTO S.p.A., TRENTO, ITALY TECHNIKON LABORATORIES, ROBERTVILLE, FLORIDA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR:	PHARMA DYNAMICS, SILVERWOOD, WESTLAKE
Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007

Registration number:	
Name of medicine:	
Dosage form:	
Active ingredients:	
Conditions of registration:	
Applicant:	CAMOX PHARMACEUTICALS (PTY) LTD
Manufacturer:	MEDREICH STERILAB LTD, VIRGONAR BANGALORE, INDIA
Packer:	MEDREICH STERILAB LTD, VIRGONAR BANGALORE, INDIA
Laboratory: FPRC:	MEDREICH STERILAB LTD, VIRGONAR BANGALORE, INDIA SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA M&L LABORATORIES, ORMONDE, JOHANNESBURG
FPRR:	CAMOX PHARMACEUTICALS, AMALGAM, JOHANNESBURG, RSA
Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007

Registration number:	A39/14/0399	Registration number:	A39/20/2.0/0610
Name of medicine:	ZYLIN	Name of medicine:	VARI-LAMIVUDINE 150 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: SERTRALINE HYDROCHLORIDE EQUIVALENT TO SERTRALINE 50.0 mg	Active ingredients:	EACH TABLET CONTAINS: LAMIVUDINE 150.0 mg
Conditions of registration	1, 2, 3, 4, 5, 6	Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	ZYDUS HEALTHCARE SA (PTY) LTD	Applicant:	LEBASI PHARMACEUTICALS CC
Manufacturer	ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA	Manufacturer:	VARICHEM PHARMACEUTICALS (PVT) LTD, HARARE, ZIMBABWE
Packer:	ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA	Packer:	VARICHEM PHARMACEUTICALS (PVT) LTD, HARARE, ZIMBABWE
Laboratory: FPRC	ZYDUS CADILA HEALTHCARE LTD, SANAND, AHMEDABAD, INDIA INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH- WEST UNIVERSITY, POTCHEFSTROOM	Laboratory:	FPRC: VARICHEM PHARMACEUTICALS (PVT) LTD, HARARE, ZIMBABWE RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM INSTITUTE FOR PHARMACEUTICAL SERVICES, SILVERTONDALE, RSA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRC:	ZYDUS HEALTHCARE SA, VAN DER HOFF PARK, POTCHEFSTROOM	FPRR	LEBASI PHARMACEUTICALS, POTCHEFSTROOM
Shelf-life:	24 months (provisional)	Shelf-life:	24 months
Date of registration:	13 APRIL 2007	Date of registration:	13 APRIL 2007

MRF 15

Registration number: A407.1/0077
 Name of medicine: STAMLATE 5
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 AMLODIPINE MALEATE EQUIVALENT TO
 AMLODIPINE 5,0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6
 Applicant: DR. REDDY'S LABORATORIES (PTY) LTD
 Manufacturer: DR. REDDY'S LABORATORIES LTD, RANGA REDDY,
 ANDHRA PRADESH, INDIA
 Packer: DR. REDDY'S LABORATORIES LTD, RANGA REDDY,
 ANDHRA PRADESH, INDIA
 Laboratory: FPRC: DR. REDDY'S LABORATORIES LTD, RANGA REDDY,
 ANDHRA PRADESH, INDIA
 INSTITUTE FOR PHARMACEUTICAL SERVICES,
 SILVERTONDALE, RSA
 FPRR: DR. REDDY'S LABORATORIES, ROSEBANK,
 JOHANNESBURG
 Shelf-life: 24 months
 Date of registration: 13 APRIL 2007

MRF 15

Registration number: A407.1/0078
 Name of medicine: STAMLATE 10
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS:
 AMLODIPINE MALEATE EQUIVALENT TO
 AMLODIPINE 10,0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6
 Applicant: DR. REDDY'S LABORATORIES (PTY) LTD
 Manufacturer: DR. REDDY'S LABORATORIES LTD, RANGA REDDY,
 ANDHRA PRADESH, INDIA
 Packer: DR. REDDY'S LABORATORIES LTD, RANGA REDDY,
 ANDHRA PRADESH, INDIA
 Laboratory: FPRC: DR. REDDY'S LABORATORIES LTD, RANGA REDDY,
 ANDHRA PRADESH, INDIA
 INSTITUTE FOR PHARMACEUTICAL SERVICES,
 SILVERTONDALE, RSA
 FPRR: DR. REDDY'S LABORATORIES, ROSEBANK,
 JOHANNESBURG
 Shelf-life: 24 months
 Date of registration: 13 APRIL 2007

Registration number:	A40/26/0352	Registration number:	A40/26/0353
Name of medicine:	DOXORUBICIN "EBEWE" 10 mg	Name of medicine:	DOXORUBICIN "EBEWE" 50 mg
Dosage form:	SOLUTION	Dosage form:	SOLUTION
Active ingredients:	EACH VIAL CONTAINS: DOXORUBICIN HYDROCHLORIDE 2,0 mg/ml	Active ingredients:	EACH VIAL CONTAINS: DOXORUBICIN HYDROCHLORIDE 2,0 mg/ml
Conditions of registration:	1, 2, 3, 4, 5, 6	Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	PHARMAFRICA (PTY) LTD	Applicant:	PHARMAFRICA (PTY) LTD
Manufacturer:	EBEWE PHARMA Ges.m.b.H., UNTERACH, AUSTRIA	Manufacturer:	EBEWE PHARMA Ges.m.b.H., UNTERACH, AUSTRIA
Packer:	EBEWE PHARMA Ges.m.b.H., UNTERACH, AUSTRIA	Packer:	EBEWE PHARMA Ges.m.b.H., UNTERACH, AUSTRIA
Laboratory:	FPRC: EBEWE PHARMA Ges.m.b.H., UNTERACH, AUSTRIA MIKROBIOLOGISCHES PRUFLABOR, INNSBRUCK, AUSTRIA LABOR L+SAG, BAD BOCKLET, GERMANY SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG	Laboratory:	FPRC: EBEWE PHARMA Ges.m.b.H., UNTERACH, AUSTRIA MIKROBIOLOGISCHES PRUFLABOR, INNSBRUCK, AUSTRIA LABOR L+SAG, BAD BOCKLET, GERMANY SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA CONSULTING CHEMICAL LABORATORIES, ATLASVILLE, BOKSBURG
FPRR:	PHARMAFRICA, NEW CENTRE, JOHANNESBURG	FPRR:	PHARMAFRICA, NEW CENTRE, JOHANNESBURG
Shelf-life:	24 months (provisional)	Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007	Date of registration:	13 APRIL 2007

MRF 15

Registration number: A407.5/0399
 Name of medicine: MEDPRO SIMVASTATIN 10
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS: SIMVASTATIN 10.0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6
 Applicant: CIPLA MEDPRO (PTY) LTD
 Manufacturer: CIPLA LTD, KURKUMBH, PUNE, INDIA
 Packer: CIPLA LTD, KURKUMBH, PUNE, INDIA
 Laboratory: CIPLA LTD, KURKUMBH, PUNE, INDIA
 FPRC: CIPLA MEDPRO, ROSENPARK, BELLVILLE
 FPRR:
 Shelf-life: 24 months
 Date of registration: 13 APRIL 2007

MRF 15

Registration number: A407.5/0400
 Name of medicine: MEDPRO SIMVASTATIN 20
 Dosage form: TABLET
 Active ingredients: EACH TABLET CONTAINS SIMVASTATIN 20.0 mg
 Conditions of registration: 1, 2, 3, 4, 5, 6
 Applicant: CIPLA MEDPRO (PTY) LTD
 Manufacturer: CIPLA LTD, KURKUMBH, PUNE, INDIA
 Packer: CIPLA LTD, KURKUMBH, PUNE, INDIA
 Laboratory: CIPLA LTD, KURKUMBH, PUNE, INDIA
 FPRC: CIPLA MEDPRO, ROSENPARK, BELLVILLE
 FPRR:
 Shelf-life: 24 months
 Date of registration: 13 APRIL 2007

Registration number:	A40/15.4/0511	Registration number:	A40/20.2.3/0535
Name of medicine:	TRAVOPROST/TIMOLOL ALCON	Name of medicine:	RIFINAH 300 FC
Dosage form:	DROPS	Dosage form:	TABLET
Active ingredients:	EACH 1,0 ml SOLUTION CONTAINS: TRAVOPROST 40,0 ug TIMOLOL MALEATE EQUIVALENT TO TIMOLOL 5,0 mg	Active ingredients:	EACH TABLET CONTAINS: RIFAMPICIN 300,0 mg ISONIAZID 150,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6	Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	ALCON LABORATORIES (S.A.) (PTY) LTD	Applicant:	AVENTIS PHARMA (PTY) LTD
Manufacturer:	S.A. ALCON-COUVREUR N.V., PUURS, BELGIUM	Manufacturer:	AVENTIS PHARMA, WALTLOO, PRETORIA
Packer:	S.A. ALCON-COUVREUR N.V., PUURS, BELGIUM	Packer:	AVENTIS PHARMA, WALTLOO, PRETORIA
Laboratory:	S.A. ALCON-COUVREUR N.V., PUURS, BELGIUM RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM	Laboratory:	FPRC: SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY, NORTH-WEST UNIVERSITY, POTCHEFSTROOM
	FPRR:		FPRC/FPRR: AVENTIS PHARMA, WALTLOO, PRETORIA
Shelf-life:	36 months	Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007	Date of registration:	13 APRIL 2007

A40/1.2/0564

Registration number:

A40/1.2/0564

Name of medicine:

CILORAM

Dosage form:

TABLET

Active ingredients:

EACH TABLET CONTAINS:
CITALOPRAM HYDROBROMIDE EQUIVALENT  20,0 mg
CITALOPRAM

Conditions of registration:

1, 2, 3, 4, 5, 6

Applicant:

AUROBINDO PHARMA (PTY) LTD

Manufacturer:

AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT,
ANDHRA PRADESH, INDIA

Packer:

AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT,
ANDHRA PRADESH, INDIA

Laboratory:

AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT,
ANDHRA PRADESH, INDIA

FPRC:

AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG,
RSA

Shelf-life:

24 months (provisional)

Date of registration:

13 APRIL 2007

A40/5.7.1/0588

Registration number:

A40/5.7.1/0588

Name of medicine:

FEXO-FAST 120

Dosage form:

TABLET

Active ingredients:

EACH TABLET CONTAINS:
FEXOFENADINE HYDROCHLORIDE 120,0 mg

Conditions of registration:

1, 2, 3, 4, 5, 6

Applicant:

SANDOZ (PTY) LTD

Manufacturer:

SANDOZ PRIVATE LTD, NAVI MUMBAI, INDIA

Packer:

SANDOZ PRIVATE LTD, NAVI MUMBAI, INDIA

Laboratory:

SANDOZ PRIVATE LTD, NAVI MUMBAI, INDIA
ANALYTICON, TERENURE, KEMPTON PARK
SOUTH AFRICAN BUREAU OF STANDARDS,
GROENKLOOF, PRETORIA

FPRC/FPRR:

SANDOZ, SPARTAN, KEMPTON PARK

Shelf-life:

24 months (provisional)

Date of registration:

13 APRIL 2007

MRF 15	
Registration number:	A40/5.7.1/0589
Name of medicine:	FEXO-FAST 180
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: FEXOFENADINE HYDROCHLORIDE 180,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	SANDOZ (PTY) LTD
Manufacturer:	SANDOZ PRIVATE LTD, NAVI MUMBAI, INDIA
Packer:	SANDOZ PRIVATE LTD, NAVI MUMBAI, INDIA
Laboratory:	FPRC: SANDOZ PRIVATE LTD, NAVI MUMBAI, INDIA ANALYTICON, TERENURE, KEMPTON PARK SOUTH AFRICAN BUREAU OF STANDARDS, GROENKLOOF, PRETORIA
FPRC/FPRR	SANDOZ, SPARTAN, KEMPTON PARK
Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007
Name of medicine:	AURO-METFORMIN 500 mg
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: METFORMIN HYDROCHLORIDE 500,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	F AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Shelf life:	F AUROBINDO PHARMA, ROSEBANK, JOHANNESBURG 24 months (provisional)
Date of registration:	13 APRIL 2007

MRF 15

Registration number:	A40/21.2/0639	Registration number:	A40/21.2/0640
Name of medicine:	AURO-METFORMIN 850 mg	Name of medicine:	AURO-METFORMIN 1 000 mg
Dosage form:	TABLET	Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: METFORMIN HYDROCHLORIDE 850.0 mg	Active ingredients:	EACH TABLET CONTAINS: METFORMIN HYDROCHLORIDE 1000.0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6	Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	AUROBINDO PHARMA (PTY) LTD	Applicant:	AUROBINDO PHARMA (PTY) LTD
Manufacturer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Manufacturer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Packer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Packer:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
Laboratory:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA	Laboratory:	AUROBINDO PHARMA LTD, RANGA REDDY DISTRICT, ANDHRA PRADESH, INDIA
		FPRC:	FPRC:
		FPRR:	FPRR:
Shelf-life:	24 months (provisional)	Shelf-life:	24 months (provisional)
Date of registration:	13 APRIL 2007	Date of registration:	13 APRIL 2007

MRF 15

REGISTRASIEINOMMER:

A49120/2.010081

Name of medicine:

VIREAD

Dosage form:

TABLET

Active ingredients:

EACH TABLET CONTAINS:
TENOFIVIR DISOPROXIL FUMARATE 300.0 mg

Conditions of registration:

1, 2, 3, 4, 5, 6

Applicant:

PHARMACARE LIMITED

Manufacturer:

ALTANA PHARMA ORANIENBURG GmbH,
ORANIENBURG, GERMANY
PATHEON INC, MISSISSAUGA, ONTARIO, CANADA
PHARMACARE LTD, KORSTEN, PORT ELIZABETH

Packer:

ALTANA PHARMA ORANIENBURG GmbH,
ORANIENBURG, GERMANY
PATHEON INC, MISSISSAUGA, ONTARIO, CANADA
PHARMACARE LTD, KORSTEN, PORT ELIZABETH
CARDINAL HEALTH GERMANY GmbH, SCHORNDORF,
GERMANY
GILEAD SCIENCES INC, SAN DIMAS, CALIFORNIA, USA
GILEAD SCIENCES LTD, DUBLIN, IRELAND

Laboratory:

FPRC:

ALTANA PHARMA ORANIENBURG GmbH,
ORANIENBURG, GERMANY
PATHEON INC, MISSISSAUGA, ONTARIO, CANADA
PATHEON INC, BURLINGTON, ONTARIO, CANADA
RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY,
NORTH-WEST UNIVERSITY, POTCHEFSTROOM
GILEAD SCIENCES INC, SAN DIMAS, CALIFORNIA, USA
GILEAD SCIENCES LTD, DUBLIN, IRELAND

FPRC/FPRR:

PHARMACARE LTD, KORSTEN, PORT ELIZABETH

Shelf-life:

36 months (packed in HDPE containers)
24 months (packed in Alu/Alu blisters)
24 weeks (bulk product transported in HDPE containers)

Date of registration:

13 APRIL 2007

41/20.2.8/0171

TRUVADA

TABLET

EACH TABLET CONTAINS:

EMTRICITABINE 200,0 mg
TENOFIVIR DISOPROXIL FUMARATE 300,0 mg

Conditions of registration:

1, 2, 3, 4, 5, 6

Applicant:

PHARMACARE LIMITED

Manufacturer:

ALTANA PHARMA ORANIENBURG GmbH,
ORANIENBURG, GERMANY
PATHEON INC, MISSISSAUGA, ONTARIO, CANADA
PHARMACARE LTD, KORSTEN, PORT ELIZABETH

Packer:

ALTANA PHARMA ORANIENBURG GmbH,
ORANIENBURG, GERMANY
PATHEON INC, MISSISSAUGA, ONTARIO, CANADA
PHARMACARE LTD, KORSTEN, PORT ELIZABETH
CARDINAL HEALTH GERMANY GmbH, SCHORNDORF,
GERMANY
GILEAD SCIENCES INC, SAN DIMAS, CALIFORNIA, USA
GILEAD SCIENCES LTD, DUBLIN, IRELAND

Laboratory:

FPRC:

ALTANA PHARMA ORANIENBURG GmbH,
ORANIENBURG, GERMANY
PATHEON INC, MISSISSAUGA, ONTARIO, CANADA
PATHEON INC, BURLINGTON, ONTARIO, CANADA
RESEARCH INSTITUTE FOR INDUSTRIAL PHARMACY,
NORTH-WEST UNIVERSITY, POTCHEFSTROOM
GILEAD SCIENCES INC, SAN DIMAS, CALIFORNIA, USA
GILEAD SCIENCES LTD, DUBLIN, IRELAND

FPRC/FPRR:

PHARMACARE LTD, KORSTEN, PORT ELIZABETH

Shelf-life:

24 months(provisional)

Date of registration:

13 APRIL 2007

ANNEX 4C

Registration number:	41/20.2.8/0229
Name of medicine:	LAVOS TABLETS
Dosage form:	TABLET
Active ingredients:	EACH TABLET CONTAINS: LAMIVUDINE 150,0 mg
Conditions of registration:	1, 2, 3, 4, 5, 6
Applicant:	ADCOCK INGRAM LIMITED
Manufacturer:	ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON
Packer:	ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON ADCOCK INGRAM LTD, AEROTON, JOHANNESBURG
Laboratory:	FPRC,FPRR: ADCOCK INGRAM HEALTHCARE, WADEVILLE, GERMISTON ADCOCK INGRAM LTD, AEROTON, JOHANNESBURG
Shelf-life:	24 months (provisional)
Date of registration:	13 April 2007