GOVERNMENT GAZETTE

OF THE

REPUBLIC OF NAMIBIA

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Government Notices

MINISTRY OF HEALTH AND SOCIAL SERVICES

No. 196 2008

MEDICINES AND RELATED SUBSTANCES CONTROL ACT, 1965:
REGISTRATION OF CERTAIN MEDICINES

In terms of section 17(a) of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965), I give notice that the medicines set out in the Schedule are registered subject to the following conditions:

(a) the manufacture and control of the medicine must be in accordance with the existing good manufacturing practices as required by the World Health Organization;

(b) in order to assess compliance with paragraph (a), inspections and investigations may be carried out, at such times as the council may consider necessary, by inspectors authorized in terms of section 26 of the Act to do so;

(c) the information contained in the package insert must, by the manufacturer of the medicine concerned, be updated so as to conform with the package insert approved by the Medical Control Council so as to ensure that such insert provides accurate information to the user of the medication concerned;

(d) the registration of medicine is subject to regular review regarding its quality, safety and efficacy, and the Council may, as it considers necessary, vary the conditions of registration of the medicine;

(e) the first two production batches must be validated in accordance with the detailed process validation protocol which was submitted at the time of the application for registration;

(f) a validation report must be submitted to the council within one month from the date of completion of the validation referred to in paragraph (e); and

(g) the Council may review the registration dossier at such intervals as the Council may determine.

J. ≠GAESEB
REGISTRAR OF MEDICINES

Windhoek, 29 July 2008

SCHEDULE

S/N	APPLICANT	REGISTERED NAME	APPROVED NAME OF ACTIVE(S)	DOSSAGE FORM	STRENGTH/DOSE UNIT	REGIST. NO.	REG. DATE
1	Pfizer Laboratories (Pty) Ltd	Caduet 5 mg / 40 mg	Amlodipine Besylate, Atorvastatin Calcium	Tablet	Each tablet contains Amlodipine Besylate equiv. to Amlodipine 5,0 mg , Atorvastatin Calcium equiv. to Atorvastatin 40,0 mg	08/7.0/0091	18/6/08
2	Pfizer Laboratories (Pty) Ltd	Caduet 5 mg / 80 mg	Amlodipine Besylate, Atorvastatin Calcium	Tablet	Each tablet contains Amlodipine Besylate equiv. to Amlodipine 5,0 mg , Atorvastatin Calcium equiv. to Atorvastatin 80,0 mg	08/7.0/0092	18/6/08
3	Pfizer Laboratories (Pty) Ltd	Caduet 10 mg / 40 mg	Amlodipine Besylate, Atorvastatin Calcium	Tablet	Each tablet contains Amlodipine Besylate equiv. to Amlodipine 10,0 mg , Atorvastatin Calcium equiv. to Atorvastatin 40,0 mg	08/7.0/0093	18/6/08
4	Pfizer Laboratories (Pty) Ltd	Caduet 10 mg / 80 mg	Amlodipine Besylate, Atorvastatin Calcium	Tablet	Each tablet contains Amlodipine Besylate equiv. to Amlodipine 10,0 mg , Atorvastatin Calcium equiv. to Atorvastatin 80,0 mg	08/7.0/0094	18/6/08
5	GlaxoSmithKline S.A. (Pty) Ltd	Augmentin ES 600	Amoxycillin Trihydrate / Potassium Clavulanate	Powder for suspension	Each 5ml suspension contains; Amoxycillin Trihydrate equiv. to Amoxycillin 600,0 mg, Potassium Clavulanate equiv. to Clavulanic Acid 42,9 mg	08/20.1.2/0095	18/6/08
6	Dr Reddys Laboratories (Pty) Ltd	Lamitor-100	Lamotrigine	Tablet	Each tablet contains Lamotrigine 100,0 mg	08/2.5/0096	18/6/08
7	Dr Reddys Laboratories (Pty) Ltd	Lamitor-50	Lamotrigine	Tablet	Each tablet contains Lamotrigine 50,0 mg	08/2.5/0097	18/6/08
8	Dr Reddys Laboratories (Pty) Ltd	Lamitor-25	Lamotrigine	Tablet	Each tablet contains Lamotrigine 25,0 mg	08/2.5/0098	18/6/08
9	Dr Reddys Laboratories (Pty) Ltd	Cifloc 250 mg	Ciprofloxacin Hydrochloride	Tablet	Each tablet contains Ciprofloxacin Hydrochloride equivalent to Ciprofloxacin 250,0 mg	08/20.1.1/0099	18/6/08
10	Dr Reddys Laboratories (Pty) Ltd	Cifloc 500 mg	Ciprofloxacin Hydrochloride	Tablet	Each tablet contains Ciprofloxacin Hydrochloride equivalent to Ciprofloxacin 500,0 mg	08/20.1.1/0100	18/6/08
11	Pharma Dynamics (Pty) Ltd	Serrapress 20	Paroxetine Hydrochloride	Tablet	Each film coated tablet contains Paroxetine Hydrochloride corres. to Paroxetine 20,0 mg	08/1.2/0101	18/6/08

12	Pharma Dynamics (Pty) Ltd	Serrapress 30	Paroxetine Hydrochloride	Tablet	Each film coated tablet contains Paroxetine Hydrochloride corres. to Paroxetine 30,0 mg	08/1.2/0102	18/6/08
13	Pharma Dynamics (Pty) Ltd	Carvetrend 25	Carvedilol	Tablet	Each tablet contains Carvedilol 25,0 mg	08/7.1.3/0103	18/6/08
14	Pharma Dynamics (Pty) Ltd	Carvetrend 12,5	Carvedilol	Tablet	Each tablet contains Carvedilol 12,5 mg	08/7.1.3/0104	18/6/08
15	Pharma Dynamics (Pty) Ltd	Carvetrend 6,25	Carvedilol	Tablet	Each tablet contains Carvedilol 6,25 mg	08/7.1.3/0105	18/6/08
16	Specpharm (Pty) Limited	Spec-Oxytocin 10	Synthetic Oxytocin	Injection	Each ampoule contains Synthetic Oxytocin 10 IU	08/19/0106	18/6/08
17	Specpharm (Pty) Limited	Spec-Oxytocin 5 IU	Synthetic Oxytocin	Injection	Each 1ml ampoule contains Synthetic Oxytocin 5 IU	08/19/0107	18/6/08
18	Sandoz (Pty) Ltd	Clozapine-Hexal 25 mg	Clozapine	Tablet	Each tablet contains Clozapine 25,0 mg	08/2.6.5/0108	18/6/08
19	Sandoz (Pty) Ltd	Clozapine-Hexal 50 mg	Clozapine	Tablet	Each tablet contains Clozapine 50,0 mg	08/2.6.5/0109	18/6/08
20	Sandoz (Pty) Ltd	Clozapine-Hexal 100 mg	Clozapine	Tablet	Each tablet contains Clozapine 100,0 mg	08/2.6.5/0110	18/6/08
21	Sandoz (Pty) Ltd	Clozex 25	Clozapine	Tablet	Each tablet contains Clozapine 25,0 mg	08/2.6.5/0111	18/6/08
22	Sandoz (Pty) Ltd	Clozex 50	Clozapine	Tablet	Each tablet contains Clozapine 50,0 mg	08/2.6.5/0112	18/6/08
23	Sandoz (Pty) Ltd	Clozex 100	Clozapine	Tablet	Each tablet contains Clozapine 100,0 mg	08/2.6.5/0113	18/6/08
24	Sandoz (Pty) Ltd	Parax 20	Paroxetine Hydrochloride	Tablet	Each tablet contains Paroxetene Hydrochloride Anhydrate equiv. to Paroxetene Anhydrate 40,0 mg	08/1.2/0114	18/6/08

25	Sandoz (Pty) Ltd	Parax 40	Paroxetine Hydrochloride	Tablet	Each tablet contains Paroxetene Hydrochloride Anhydrate equiv. to Paroxetene Anhydrate 40,0 mg	08/1.2/0115	18/6/08
26	Sandoz (Pty) Ltd	CitaloHexal 20	Citalopram Hydrobromide	Tablet	Each film coated tablet contains Citalopram Hydrobromide equiv. to Citalopram 20,0 mg	08/1.2/0116	18/6/08
27	Ranbaxy (S.A.) (Pty) Ltd	Simayla Citalopram 40	Citalopram Hydrobromide	Tablet	Each film coated tablet contains Citalopram Hydrobromide equiv. to Citalopram 40,0 mg	08/1.2/0117	18/6/08
28	Allergan Pharmaceuticals (Pty) Ltd	Combigan	Brimodine Tartrate , Timolol Maleate	Opthalmic Solution	Each 1 ml solution contains: Brimodine Tartrate 2,0 mg, Timolol Maleate equiv. to Timolol 5,0 mg	08/15.4/0118	18/6/08
29	Allergan Pharmaceuticals (Pty) Ltd	Alphagan Purite Eye drops	Brimodine Tartrate	Opthalmic Solution	Each 1 ml solution contains: Brimodine Tartrate 1,5 mg,	08/15.4/0119	18/6/08
30	Novartis South Africa (Pty) Ltd	Enablex 7.5 mg	Darifenacin Hydrobromide	Tablet	Each tablet contains Darifenacin Hydrobromide equiv. to Darifenacin 7,5 mg	08/5.4/0120	18/6/08
31	Novartis South Africa (Pty) Ltd	Enablex 15 mg	Darifenacin Hydrobromide	Tablet	Each tablet contains Darifenacin Hydrobromide equiv. to Darifenacin 15,0 mg	08/5.4/0121	18/6/08
32	Novartis South Africa (Pty) Ltd	Exjade 125 mg	Deferasirox	Tablet	Each dispersible tablet contains Deferasirox 125,0 mg	08/27/0122	18/6/08
33	Novartis South Africa (Pty) Ltd	Exjade 250 mg	Deferasirox	Tablet	Each dispersible tablet contains Deferasirox 250,0 mg	08/27/0123	18/6/08
34	Novartis South Africa (Pty) Ltd	Exjade 500 mg	Deferasirox	Tablet	Each dispersible tablet contains Deferasirox 500,0 mg	08/27/0124	18/6/08
35	Novartis South Africa (Pty) Ltd	Cleevec 100	Imatinib Mesilate	Tablet	Each tablet contains Imatinib Mesilate equiv. to Imatinib 100,0 mg	08/34/0125	18/6/08
36	Novartis South Africa (Pty) Ltd	Cleevec 400	Imatinib Mesilate	Tablet	Each tablet contains Imatinib Mesilate equiv. to Imatinib 400,0 mg	08/34/0126	18/6/08

37	Novartis South Africa (Pty) Ltd	Sandostatin Lar 10 mg	Octreotide Acetate	Injection	Each vial contains Octreotide Acetate equiv. to Octreotide 10,0 mg	08/34/0127	18/6/08
38	Novartis South Africa (Pty) Ltd	Sandostatin Lar 30 mg	Octreotide Acetate	Injection	Each vial contains Octreotide Acetate equiv. to Octreotide 30,0 mg	08/34/0128	18/6/08
39	Novartis South Africa (Pty) Ltd	Aclasta	Zoledronic Acid Monohydrate	Infusion	Each 100 ml solution contains Zoledronic Acid Monohydrate equiv. to Zoledronic Acid 5,0 mg	08/34/0129	18/6/08
40	Novartis South Africa (Pty) Ltd	Zometa Concentrate for Solution for Infusion	Zoledronic Acid Monohydrate	Concentrate	Each 5 ml vial contains Zoledronic Acid Monohydrate equiv. to Zoledronic Acid 4,0 mg	08/34/0130	18/6/08
41	Novartis South Africa (Pty) Ltd	Tegretol 200	Carbamazepine	Tablet	Each tablet contains Carbamazepine 200,0 mg	08/2.5/0131	18/6/08
42	Sandoz (Pty) Ltd	Sandoz Topiramate 200	Topiramate	Tablet	Each film-coated tablet contains Topiramate 200,0 mg	08/2.5/0132	18/6/08
43	Sandoz (Pty) Ltd	Sandoz Topiramate 100	Topiramate	Tablet	Each film-coated tablet contains Topiramate 100,0 mg	08/2.5/0133	18/6/08
44	Sandoz (Pty) Ltd	Sandoz Topiramate 50	Topiramate	Tablet	Each film-coated tablet contains Topiramate 50,0 mg	08/2.5/0134	18/6/08
45	Sandoz (Pty) Ltd	Sandoz Topiramate 25	Topiramate	Tablet	Each film-coated tablet contains Topiramate 25,0 mg	08/2.5/0135	18/6/08
46	Sandoz (Pty) Ltd	Fexofast 120	Fexofenadine Hydrochloride	Tablet	Each film-coated tablet contains Fexofenadine Hydrochloride 120,0 mg	08/5.7.1/0136	18/6/08
47	Sandoz (Pty) Ltd	Fexofast 180	Fexofenadine Hydrochloride	Tablet	Each film-coated tablet contains Fexofenadine Hydrochloride 180,0 mg	08/5.7.1/0137	18/6/08
48	iNova Pharmaceuticals (Pty) Limited	Duro-Tuss Linctus	Bromhexine Hydrochloride, Salbutamol Sulphate	Linctus	Each 5 ml liquid contains Bromhexine Hydrochloride 4,0 mg, Salbutamol Sulphate equiv. to Salbutamol 2,0 mg	08/10.1/0138	18/6/08

49	Novartis South Africa (Pty) Ltd	Trileptal 150	Oxcarbazepine	Tablet	Each tablet contains Oxcarbazepine 150,0 mg	08/2.5/0139	18/6/08
50	Adcock Ingram Ltd	Adco-Nevirapine	Nevirapine	Tablet	Each tablet contains Nevirapine 200,0 mg	08/20.2.8/0140	18/6/08
51	Adcock Ingram Ltd	Adco-Lamivudine	Lamivudine	Tablet	Each tablet contains Lamivudine 150,0 mg	08/20.2.8/0141	18/6/08
52	Adcock Ingram Ltd	Adco-Efavirenz	Efavirenz	Tablet	Each tablet contains Efavirenz 600,0 mg	08/20.2.8/0142	18/6/08
53	Adcock Ingram Ltd	Adco-Zidovudine	Zidovudine	Tablet	Each tablet contains Zidovudine 300,0 mg	08/20.2.8/0143	18/6/08
54	Pfizer Laboratories (Pty) Ltd	Benylin Children's	Diphenhydramine Hydrochloride, Ammonium Chloride	Syrup	Each 5ml syrup contains Diphenhydramine Hydrochloride 12,5 mg, Ammonium Chloride 125,0 mg	08/10.1/0144	18/6/08
55	Pfizer Laboratories (Pty) Ltd	Dalacin VC	Clindamycin Phosphate	Vag. Cream	Each 1 g cream contains Clindamycin Phosphate equiv. to Clindamycin 20,0 mg	08/20.1.1/0145	18/6/08
56	Pfizer Laboratories (Pty) Ltd	Xalacom	Latanoprost, Timolol Maleate	Eye Drops	Each 1ml solution contains Latanoprost 50,0 μg, Timolol Maleate equiv. to Timolol 5,0 mg	08/15.4/0146	18/6/08
57	Pfizer Laboratories (Pty) Ltd	Sutent 25 mg Capsules	Sunitinib Maleate	Capsules	Each capsule contains Sunitinib Maleate 25,0 mg	08/26/0147	18/6/08
58	Pfizer Laboratories (Pty) Ltd	Sutent 12,5 mg Capsules	Sunitinib Maleate	Capsules	Each capsule contains Sunitinib Maleate 12,5 mg	08/26/0148	18/6/08
59	Pfizer Laboratories (Pty) Ltd	Sutent 50,0 mg Capsules	Sunitinib Maleate	Capsules	Each capsule contains Sunitinib Maleate 50,0 mg	08/26/0149	18/6/08
60	Pfizer Laboratories (Pty) Ltd	Lyrica 25 mg	Pregabalin	Capsules	Each capsule contains Pregabalin 25,0 mg	08/2.5/0150	18/6/08

61	Pfizer Laboratories (Pty) Ltd	Lyrica 50 mg	Pregabalin	Capsules	Each capsule contains Pregabalin 50,0 mg	08/2.5/0151	18/6/08
62	Pfizer Laboratories (Pty) Ltd	Lyrica 75 mg	Pregabalin	Capsules	Each capsule contains Pregabalin 75,0 mg	08/2.5/0152	18/6/08
63	Pfizer Laboratories (Pty) Ltd	Lyrica 100 mg	Pregabalin	Capsules	Each capsule contains Pregabalin 100,0 mg	08/2.5/0153	18/6/08
64	Pfizer Laboratories (Pty) Ltd	Lyrica 150 mg	Pregabalin	Capsules	Each capsule contains Pregabalin 150,0 mg	08/2.5/0154	18/6/08
65	Pharma Dynamics (Pty) Ltd	Pharma Dynamics Amlodipine Besylate 5 mg	Amlodipine Besylate	Tablet	Each tablet contains Amlodipine Besylate equiv. to Amlodipine 5,0 mg	08/7.1/0155	18/6/08
66	Pharma Dynamics (Pty) Ltd	Pharma Dynamics Amlodipine Besylate 10 mg	Amlodipine Besylate	Tablet	Each tablet contains Amlodipine besylate equiv. to Amlodipine 10,0 mg	08/7.1/0156	18/6/08
67	Abbott Laboratories (Pty) Ltd	Aluvia 100/25	Lopinavir, Ritonavir	Tablet	Each film-coated tablet contains Lopinavir 100,0 mg , Ritonavir 25,0 mg	08/20.2.8/0157	18/6/08
68	Dr Reddys Laboratories (Pty) Ltd	DRL-Omeprazole 40	Omeprazole	Capsules	Each capsule contains Omeprazole 40,0 mg	08/11.4.3/0158	18/6/08
69	AstraZeneca Pharmaceuticals (Pty) Ltd	Pulmicort Nebulising Suspension 0,25 mg/ml	Budesonide	Suspension	Each 1 ml suspension contains Budesonide 0,25 mg	08/21.5.1/0159	18/6/08
70	AstraZeneca Pharmaceuticals (Pty) Ltd	Pulmicort Nebulising Suspension 0,5 mg/ml	Budesonide	Suspension	Each 1 ml suspension contains Budesonide 0,5 mg	08/21.5.1/0160	18/6/08
71	Pharmaplan (Pty) Ltd	Dazit Tablets	Desloratadine	Tablet	Each tablet contains Desloratadine 5,0 mg	08/5.7.1/0161	18/6/08
72	Servier Laboratories SA (Pty) Ltd	Coralan 7,5 mg Tablets	Ivabradine Hydrochloride	Tablet	Each tablet contains Ivabradine Hydrochloride equiv. to Ivabradine 7,5 mg	08/7.6/0162	18/6/08

73	Servier Laboratories SA (Pty) Ltd	Coralan 5 mg Tablets	Ivabradine Hydrochloride	Tablet	Each tablet contains Ivabradine Hydrochloride equiv. to Ivabradine 5,0 mg	08/7.6/0163	18/6/08
74	Biogaran South Africa (Pty) Ltd	Caranor 7,5 mg Tablets	Ivabradine Hydrochloride	Tablet	Each tablet contains Ivabradine Hydrochloride equiv. to Ivabradine 7,5 mg	08/7.6/0164	18/6/08
75	Biogaran South Africa (Pty) Ltd	Caranor 5 mg Tablets	Ivabradine Hydrochloride	Tablet	Each tablet contains Ivabradine Hydrochloride equiv. to Ivabradine 5,0 mg	08/7.6/0165	18/6/08
76	Sandoz (Pty) Ltd	Sandoz Cefpodoxime 100	Cefpodoxime Proxetil	Tablet	Each tablet contains Cefpodoxime Proxetil equiv. to Cefpodoxime 100,0 mg	08/20.1.1/0166	18/6/08
77	Sandoz (Pty) Ltd	Sandoz Cefpodoxime 40mg/ 5ml	Cefpodoxime Proxetil	Suspension	Each 5ml suspension contains Cefpodoxime Proxetil equiv. to Cefpodoxime 40,0 mg	08/20.1.1/0167	18/6/08
78	AstraZeneca Pharmaceuticals (Pty) Ltd	Zoladex 10,8	Goserelin Acetate	Injection	Each Zoladex Depot contains Goserelin Acetate equiv. to Goserelin Base 10,8 mg	08/21.10/0168	18/6/08
79	Adcock Ingram Ltd	Rilace 1.25	Ramipril	Capsules	Each capsule contains Ramipril 1,25 mg	08/7.1.3/0169	18/6/08
80	Adcock Ingram Ltd	Rilace 2.5	Ramipril	Capsules	Each capsule contains Ramipril 2.5 mg	08/7.1.3/0170	18/6/08
81	Adcock Ingram Ltd	Rilace 5	Ramipril	Capsules	Each capsule contains Ramipril 5 mg	08/7.1.3/0171	18/6/08
82	Adcock Ingram Ltd	Rilace 10	Ramipril	Capsules	Each capsule contains Ramipril 10 mg	08/7.1.3/0172	18/6/08
83	Adcock Ingram Ltd	Tramgesic Flashtabs	Tramadol Hydrochloride	Tablet	Each tablet contains Tramadol Hydrochloride 50,0 mg	08/2.9/0173	18/6/08
84	Adcock Ingram Ltd	Adco-Fem 35	Cyproterone Acetate, Ethynyloestradol	Tablet	Each tablet contains Cyproterone Acetate 2,0 mg, Ethynyloestradiol 0,035 mg	08/21.8.2/0174	18/6/08

MINISTRY OF HEALTH AND SOCIAL SERVICES

No. 197 2008

REGULATIONS RELATING TO SCOPE OF PRACTICE OF ORAL HYGIENE: MEDICAL AND DENTAL ACT, 2004

Under section 59 of the Medical and Dental Act, 2004 (Act No. 10 of 2004), and on the recommendation of the Medical and Dental Council of Namibia, I have made the regulations set out in the Schedule.

R. N. KAMWI
MINISTER OF HEALTH AND SOCIAL SERVICES Windhoek, 29 July 2008

SCHEDULE

Definitions

1. In these regulations, unless the context otherwise indicates, a word or expression defined in the Act has that meaning, and -

"the Act" means the Medical and Dental Act, 2004 (Act No. 10 of 2004).

Scope of practice of oral hygiene

2. an oral hygienist may perform the following acts -

(a) the provisional examination and charting of conditions of the mouth, with particulars reference to the face, soft tissues, teeth, occlusion and periodontium;

(b) the scaling, roof planning and polishing of teeth, including the trimming and polishing of restorations;

(c) the performing of dental radiography;

(d) the topical application of agents appropriate to the practice of oral hygiene, including caries preventive agents, tooth-desensitizing agents, surface anaesthetics and plague-controlling agents;

(e) the application and removal of periodontal packs;

(f) the taking of impressions and the casting of study and primary work models:

(g) the placement of temporary fillings as an emergency measure prior to the referral to a dentist or dental therapist;

(h) the temporary cementing of inlays, crowns and bridges;

(i) the placement of glass ionomers cement on sensitive dentine abrasion lesions;

(j) the placement of soft lining in dentures as tissue conditioners;

(k) the taking of cytological smears, including for the purpose of testing for candida infection;

(l) the discussion of treatment options with, and the explaining of treatment procedures to patients;

(m) the discussion of treatment fees;

(n) the performing of the following functions in orthodontics -

(i) cephalometric tracing;

(ii) the relief of trauma caused by intra- and extra-oral appliances, including the cutting of the distal arch wires and the removal of the complete arch wire;

(iii) the placement of pre-activated orthodontic appliances and the removal of orthodontic attachments, bands, appliances and arch wires;

(iv) the fitting of orthodontic retainers; and

(o) the administering of applicable local analgesia appropriate to the scope of the profession of oral hygiene.

Performing of professional acts by student in oral hygiene

3. A student in oral hygiene may perform, as part of his or her education, tuition and training, and on the instructions, and under the direct supervision, of a dentist or an oral hygienist, any of the acts prescribed by regulation 2.

MINISTRY OF HEALTH AND SOCIAL SERVICES

No. 198 2008

REGULATIONS RELATING TO OWNERSHIP OF PHARMACY BY PRIVATE HOSPITAL: PHARMACY ACT, 2004

Under section 66 of the Pharmacy Act, 2004 (Act No. 9 of 2004), read with section 37 of that Act, on the recommendation of the Pharmacy Council of Namibia, I have made the regulations set out in the Schedule.

R. N. KAMWI
MINISTER OF HEALTH AND SOCIAL SERVICES Windhoek, 30 July 2008

SCHEDULE

Definitions

1. In these regulations, unless the context otherwise indicates, a word or expression defined in the Act has that meaning, and -

"hospital" means a state hospital established under section 2 of the Hospitals and Health Facilities Act, or a private hospital registered under section 23 of that Act;

"Hospitals and Health Facilities Act" means the Hospitals and Health Facilities Act, 1994 (Act No. 36 of 1994);

"Medicines and Related Substances Control Act" means the Medicines and Related Substances Control Act, 2003 (Act No. 13 of 2003); and

"the Act" means the Pharmacy Act, 2004 (Act No. 9 of 2004).

Person other than pharmacist, private company or close corporation who may own pharmacy

2. Subject to the Act, a hospital may own a community pharmacy.

Conditions subject whereto private hospital may own and conduct pharmacy

3. (1) A community pharmacy referred to in subregulation (2) must -

(a) be licensed as a private health facility under section 31 of the Hospitals and Health Facilities Act; and

(b) situate on the premises where the hospital is situated.

(2) A hospital referred to in regulation 2 –

(a) must conduct its community pharmacy in accordance with the Act, the Medicines and Related Substances Control Act and the Hospitals and Health Facilities Act; and

(b) may not -

(i) have any financial or other interest in any community pharmacy other than the community pharmacy of which it is the owner; or

(ii) conduct business as a wholesale pharmacist referred to in section 35(26) of the Act or as a manufacturer as defined in section 1 of the Medicines and Related Substances Control Act, or have any financial or other interest in such wholesale pharmacy or manufacturer.