DEPARTMENT OF HEALTH

MEDICINES AND RELATED SUBSTANCES ACT, 1965 (ACT No. 101 OF 1965)

SCHEDULES

The Minister of Health has, in terms of section 22A(2) of the Medicines and Related Substances Act, 1965 (Act No. 101 of 1965), on the recommendation of the South African Health Products Regulatory Authority (SAHPRA), made and updated the Schedules.


- Words in bold and in square brackets (e.g. [Gamma benzene hexachloride] in Schedule 1), indicate omission from a Schedule
- Words underlined with a solid line (e.g. Gamma benzene hexachloride), indicate insertions in a Schedule.

SCHEDULE

In these Schedules, "the Act" means the Medicines and Related Substances Act, 1965 (Act No.101 of 1965)

Note: Where an alternative schedule(s) is included in natural parentheses at any point of an inscription, this is provided to indicate one or more alternative scheduling designation/s. This is for information only and shall not be used in the interpretation of such inscription.

SCHEDULE 1

a. All substances referred to in this Schedule are excluded when specifically packed, labelled, sold and used for –
   (i) industrial purposes including the manufacture or compounding of consumer items or products which have no pharmacological action or medicinal purpose; and
   (ii) analytical laboratory purposes.

b. All preparations of substances or mixtures of such substances containing or purporting to contain any substance referred to in this Schedule and includes the following:
(i) The salts and esters of such substances, where the existence of such salts and esters is possible; and
(ii) all preparations and mixtures of such substances where such preparations and mixtures are not expressly excluded.

c. In terms of section 22A(4)(a)(v) of the Act, a practitioner, nurse or a person registered under the Health Professions Act, 1974 (Act No. 56 of 1974) other than a medical practitioner or dentist may prescribe and supply, only within his/her scope of practice and subject to the indication for use of such substances and medicines and to the conditions determined by the Authority, to patients under his/her care, the Schedule 1 substances and medicines provided for in the Annexures to this Schedule published in the Gazette in terms of the Act.

(i) Annexure 1A: Emergency Care Provider (Paramedic);
Annexure 1B: Emergency Care Provider (Emergency Care Practitioner);
Annexure 1C: Basic Ambulance Assistant
Annexure 1D: Ambulance Emergency Assistant
Annexure 1E: Emergency Care Technician
Annexure 1F: Emergency Care Assistant

(ii) Annexure 2: Dental Therapist;
(iii) Annexure 3: Optometrist.
(iv) Annexure 4: Podiatrist
(v) Annexure 5: Oral hygienists

Dequalinium

(a) when intended for oral topical use, as oral solutions or lozenges;
(b) except when intended for human vaginal use (S2)

Phenylephrine

a. when intended for oral dosage forms, nasal dosage forms, or ophthalmic dosage forms containing more than 0.2 percent (S1)

b. except ophthalmic preparations containing 0.2 percent or less. (S0)

c. except when intended for injection (S4)
ANNEXURE 5: ORAL HYGIENISTS

Oral hygienists registered with the Health Professions Council of South Africa (HPCSA) in terms of the Health Professions Act, 1974 (Act 56 of 1974)

<table>
<thead>
<tr>
<th>ORAL HYGIENISTS</th>
<th>LOCAL ANAESTHETIC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Substance</td>
<td>Lignocaine/Lidocaine hydrochloride</td>
</tr>
<tr>
<td>Indication</td>
<td>Dental surface anaesthesia (excluding injectables)</td>
</tr>
<tr>
<td>Route of Administration</td>
<td>Topical</td>
</tr>
</tbody>
</table>

| TOPICAL FLUORIDES | | |
|-------------------|------------------|
| Substance         | -                |
| Indication        | Applicable to dentistry |
| Route of administration | Topical |

- END SCHEDULE 1 -
SCHEDULE 2

a. All substances referred to in this Schedule are excluded when specifically packed, labeled, sold and used for –
   (i) industrial purposes including the manufacture or compounding of consumer items or products which have no pharmacological action or medicinal purpose; and
   (ii) analytical laboratory purposes.

b. All preparations of substances or mixtures of such substances containing or purporting to contain any substance referred to in this Schedule and includes the following:
   (i) The salts and esters of such substances, where the existence of such salts and esters is possible; and
   (ii) all preparations and mixtures of such substances where such preparations and mixtures are not expressly excluded.

c. In terms of section 22A(5)(f) of the Act, a practitioner, nurse or a person registered under the Health Professions Act, 1974 (Act 56 of 1974) other than a medical practitioner or dentist may prescribe and supply, only within their scope of practice and subject to the indication for use of such substances and medicines and to the conditions determined by the Authority, to patients under his/her care, the Schedule 2 substances and medicines provided for in the Annexures to this Schedule published in the Gazette in terms of the Act.

   (i) Annexure 1A: Emergency Care Provider (Paramedic);
   Annexure 1B: Emergency Care Provider (Emergency Care Practitioner);
   Annexure 1C: Basic Ambulance Assistant
   Annexure 1D: Ambulance Emergency Assistant
   Annexure 1E: Emergency Care Technician
   Annexure 1F: Emergency Care Assistant
   (ii) Annexure 2: Dental Therapist;
   (iii) Annexure 3: Optometrist.
   (iv) Annexure 4: Podiatrist

Dequalinium

(a) when intended for human vaginal use;
(b) except when intended for oral topical use, as oral solutions or lozenges (31)
Estriol.
   a. When intended for human vaginal use
   b. except when intended for oral contraception; (S3)
   c. except when intended for hormone replacement therapy; (S4)
   d. except when intended for veterinary use (S4)

Rizatriptan, when in oral solid dosage forms providing 5 mg or less and presented as packs of
no more than 2 oral solid dosage forms, indicated for the acute relief of migraine attacks, with or
without aura, in patients previously diagnosed by a medical practitioner and initiated on
treatment with rizatriptan (S4)

ANNEXURE 4: PODIATRIST

PODIATRIST registered with the Health Professions Council of South Africa in terms of the
Health Professions Act, 1974 (Act 56 of 1974)

<table>
<thead>
<tr>
<th>PODIATRIST</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anti-inflammatories</td>
</tr>
<tr>
<td>Substance     : Diclofenac sodium and Ibuprofen</td>
</tr>
<tr>
<td>Indication    : Pain management</td>
</tr>
<tr>
<td>Route of Administration  : Oral</td>
</tr>
</tbody>
</table>

END SCHEDULE 2

SCHEDULE 3

a. All substances referred to in this Schedule are excluded when specifically packed,
   labelled, sold and used for —
   (i) industrial purposes including the manufacture or compounding of consumer items
      or products which have no pharmacological action or medicinal purpose; and
   (ii) analytical laboratory purposes.

b. All preparations of substances or mixtures of such substances containing or purporting
to contain any substance referred to in this Schedule and includes the following:
   (i) The salts and esters of such substances, where the existence of such salts and
       esters is possible; and
   (ii) all preparations and mixtures of such substances where such preparations and
       mixtures are not expressly excluded.
c. In terms of section 22A(5)(f) of the Act, a practitioner, nurse or a person registered under the Health Professions Act, 1974 (Act 56 of 1974) other than a medical practitioner or dentist may prescribe and supply, only within his/her scope of practice and subject to the indication for use of such substances and medicines and to the conditions determined by the Authority, to patients under his/her care, the Schedule 3 substances and medicines provided for in the Annexures to this Schedule published in the Gazette in terms of the Act.

(i) Annexure 1A: Emergency Care Provider (Paramedic);
(ii) Annexure 1B: Emergency Care Provider (Emergency Care Practitioner);
(iii) Annexure 2: Dental Therapist;
(iv) Annexure 3: Optometrist.
(v) Annexure 4: Podiatrist

Estradiol,

a. when intended for oral contraception
b. except when intended for human vaginal use (S2);
c. except when intended for hormone replacement therapy, (S4)
d. except when intended for veterinary use (S4)

Folinic acid (leucovorin)

Levalbuterol

- END SCHEDULE 3 -

SCHEDULE 4

a. All substances referred to in this Schedule are excluded when specifically packed, labelled, sold and used for –
   (i) industrial purposes including the manufacture or compounding of consumer items or products which have no pharmacological action or medicinal purpose; and
   (ii) analytical laboratory purposes.

b. All preparations of substances or mixtures of such substances containing or purporting to contain any substance referred to in this Schedule and includes the following:
   (i) The salts and esters of such substances, where the existence of such salts and esters is possible; and

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(iii) all preparations and mixtures of such substances where such preparations and mixtures are not expressly excluded.

c. In terms of section 22A(5)(f) of the Act, a practitioner, nurse or a person registered under the Health Professions Act, 1974 (Act 56 of 1974) other than a medical practitioner or dentist may prescribe and supply, only within his/her scope of practice and subject to the indication for use of such substances and medicines and to the conditions determined by the Authority, to patients under his/her care, the Schedule 4 substances and medicines provided for in the Annexures to this Schedule published in the Gazette in terms of the Act.

(i) Annexure 1A: Emergency Care Provider (Paramedic);
(ii) Annexure 1B: Emergency Care Provider (Emergency Care Practitioner);
(iii) Annexure 1C: Basic Ambulance Assistant
(iv) Annexure 1D: Ambulance Emergency Assistant
(v) Annexure 1E: Emergency Care Technician
(vi) Annexure 1F: Emergency Care Assistant
(vii) Annexure 2: Dental Therapist;
(viii) Annexure 3: Optometrist.
(ix) Annexure 4: Podiatrist

Alectinib
Alpelisib
Apalutamide
Asciminib
Bedinvetmab
Bictegravir
Cabotegravir
Cabozantinib
Casirivimab
Dacomitinib
Dapivirine
Darolutamide
[Dequalinium]

Entrectinib

Estriol

a. when intended for hormone replacement therapy
b. when intended for veterinary use
c. except when intended for oral contraception: (S3)
  d. except when intended for human vaginal use (S2);

Faricimab

Fremenezumab

Glucagon

Guselkumab

Icatibant

Idebenone

Imdevimab

Inclisiran

Itopride

Leremovir

Linagliptin

Molnupiravir

Neratinib

Noradrenaline (noradrenaline)

Olaparib

Phenylephrine

a. when intended for injection
b. except ophthalmic preparations containing 0.2 percent or less. (S0)
c. except for oral dosage forms, nasal dosage forms, or ophthalmic dosage forms containing more than 0.2 percent (S1)
Polatuzumab
Pralsetinib
Pretomanid
Recombinant human epidermal growth factor (rhEGF)
Remdesivir
Revenfacin
Risdiplam
Rizatriptan, except when in oral solid dosage forms providing 5 mg or less and presented as packs of no more than 2 oral solid dosage forms, indicated for the acute relief of migraine attacks, with or without aura, in patients previously diagnosed by a medical practitioner and initiated on treatment with rizatriptan (S2)
Safinamide
Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) vaccine
Satralizumab
Selexipag
Tafamidis
Tivozanib
Tozinameran
Turoctocog Alpha
Upadacitinib
Zofenopril

ANNEXURE 3: OPTOMETRISTS

OPTOMETRIST (Bachelors degree in Optometry – B OPTOM) registered with the Health Professions Council of South Africa in terms of the Health Professions Act, 1974 (Act 56 of 1974) and recognised by the Health Professions Council of South Africa as an authorised prescriber.

<table>
<thead>
<tr>
<th>OPTOMETRISTS</th>
<th>ANTIBIOTICS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Substance</td>
<td>Fusidic acid</td>
</tr>
<tr>
<td>Indication</td>
<td>For Blepharitis and stye</td>
</tr>
<tr>
<td>Route of Administration</td>
<td>Topical drops or ointment</td>
</tr>
</tbody>
</table>
OPTOMETRISTS

ANTIBIOTICS
Substance : Neomycin
Indication : For Blepharitis only
Route of Administration : Topical drops or ointment

ANTIBIOTICS
Substance : Bacitracin
Indication : For Blepharitis only
Route of Administration : Ointment

ANTIBIOTICS
Substance : Polymyxin B
Indication : For Blepharitis only
Route of Administration : Ointment

PROSTAGLANDIN ANALOGUES (PGAs)
Substance : Latanoprost, Travoprost, Bimatoprost
Indication : Glaucoma
Route of Administration : Drops

— END SCHEDULE 4 —

SCHEDULE 5 AND SPECIFIED SCHEDULE 5

a. All preparations or mixtures of such substances containing or purporting to contain
substances that is chemically related and incorporates a structural fragment into its
structure that is similar to the structure of a listed substance and/or exhibits
pharmacodynamic properties similar to the listed substance referred to in this Schedule
include the following:

(i) The salts and esters of such substances, where the existence of such salts
and esters is possible; and

(ii) all preparations and mixtures of such substances where such preparations
and mixtures are not expressly excluded.

(iii) all homologues of listed substances (being any chemically related
substances that incorporate a structural fragment into their structures that is
similar to the structure of a listed substance and/or exhibit
pharmacodynamic properties similar to the listed substance in the
schedules), unless listed separately in the Schedules.
b. In terms of Section 22A(5)(f) of the Act, a practitioner, nurse or a person registered under the Health Professions Act, 1974, other than a medical practitioner or dentist, may prescribe and apply, only within his/her scope of practice and subject to the indication for use of such substances and medicines and to the conditions determined by the Medicines Control Council, to patients under his/her care, the Schedule 5 and Specified Schedule 5 substances and medicines provided for in the Annexures to this Schedule published in the Gazette in terms of the Act.

(i) Annexure 1A: Emergency Care Provider (Paramedic);
(ii) Annexure 1B: Emergency Care Provider (Emergency Care Practitioner);
(iii) Annexure 1E: Emergency Care Technician

c. Specified Schedule 5 substances listed in this schedule are subject to additional control in terms of section 22A of the Act as required under the provisions of the 1971 Convention on Psychotropic Substances and are denoted by **

Esketamine
Lemborexant
[Recombinant human epidermal growth factor (rhEGF)]

– END SCHEDULE 5 –

SCHEDULE 7

All preparations or mixture of such substances containing or purporting to contain substances referred to in this Schedule include the following (unless expressly excluded or unless listed in another Schedule):

(i) the isomers of such substances, where the existence of such isomers is possible within the chemical designation;

(ii) the esters and ethers of such substances and of the isomers referred to in (i), as well as the isomers of such esters and ethers, where the existence of isomers of such esters, or ethers is possible;

(iii) the salts of such substances and of the isomers referred to in (i), as well as the salts of the esters, ethers and isomers referred to in (ii), where the existence of such salts is possible;
(iv) the isomers of any of the salts referred to in (iii), where the existence of such isomers is possible;

(v) all preparations and mixtures of any of the above.

(vi) all homologues of listed substances (being any chemically related substances that incorporate a structural fragment into their structures that is similar to the structure of a listed substance and/or exhibit pharmacodynamic properties similar to the listed substance in the schedules), unless listed separately in the Schedules.

Borphine
Eutylone
Metonitazene
Norfentanyl

– END SCHEDULE 7 –

These Schedules as amended come into operation on the date of publication in the Government Gazette.

DR M. PHAAHLA, MP
MINISTER OF HEALTH
DATE: 06/10/2022