GOVERNMENT NOTICES • GOEWERMENTSKENNISGEWINGS

DEPARTMENT OF HEALTH

NO. R. 1375 18 DECEMBER 2020

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.

This Schedule amends the Schedules as inserted by Government Notice R.509 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 24727, 10 April 2003; substituted by Government Notice R.935 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 31387, 5 September 2008; and amended by Government Notice R.1230 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 32838, 31 December 2009; Government Notice R.227 (Medicines and Related Substances Act: Schedules)in Government Gazette 35149, 15 March 2012; Government Notice R.674 (Medicines and Related Substances Act, 1965; Schedules) in Government Gazette 36827, 13 September 2013, Government Notice R.690 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 36850, 20 September 2013, Government Notice R.104 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 37318. 11 February 2014; Government Notice R.352 (Medicines and Related Substances Act, 1965; Schedules) in, Government Gazette 37622, 8 May 2014; Government Notice R.234 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 38586, 20 March 2015; Government Notice R.254 (Medicines and Related Substances Act, 1965: Schedules) in Government Gezette 39815, 15 March 2016; Government Notice R.254 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 40041, 03 June 2016: Government Notice No.748 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 41009, 28 July 2017; Government Notice No.1261 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 41256, 17 November 2017; Government Notice No.1262 (Medicines and Related Substances Act, 1965; Schedules) in Government Gazette 42052, 23 November 2018 and Government Notice No.755 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 42477, 23 May 2019; Government Notice No.R219 (Medicines and Related Substances Act, 1985: Schedules) in Government Gazette 430151, 28 February 2020 and Government Notice No.R586 (Medicines and Related Substances Act, 1965: Schedules) in Government Gazette 43347, 22 May 2020 using the following convention:

- 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. <u>Gamma benzene hexachloride</u>), 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

- All substances referred to in this Schedule are excluded when specifically packed, labelled, sold and used for –
 - 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:
 - The salts and esters of such substances, where the existence of such salts and esters is possible; and
 - 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

Arsenic;

- a. in oral dosage forms in concentrations equivalent to 0,01 percent or less of arsenic trioxide; (S2)
- b. except when intended for injection. (S4)

Benzydamine,

- a. preparations and mixtures containing [0,15 percent or less of benzydamine, when] intended for
 use as a mouth rinse or for topical application in the mouth and throat; provided that the total dose
 swallowed does not exceed 36 milligrams of benzydamine per day. (S3)
- except preparations and mixtures containing 3 percent or less of benzydamine, when intended for application to the skin (S0) [(S3)]; or
- except preparations and mixtures containing 3 milligrams or less of benzydamine per throat lozenge: Provided that the total dose swallowed does not exceed 36 milligrams of benzydamine per day and the pack size does not exceed 16 lozenges. (S0)
- d. except when indicated for human vaginal use. (S2)

Flurbiprofen,

- a. when in the form of lozenges, indicated for the relief of pain associated with sore throats, subject to:
 - (i) a maximum of 8,75 milligrams per lozenge;
 - (ii) a maximum treatment period of 3 days; and
 - (iii) a maximum pack size of 15 lozenges. (S3)
- except when intended for application to the skin and indicated for the symptomatic relief of localised pain and inflammation, provided that in the case of application by transdermal patch;
 - (i) use is restricted to adults and children 12 years and older; and
 - (ii) the treatment period is limited to a maximum of 4 weeks (S0)

Schedule 1

- except when intended for the treatment of post-traumatic conditions, for a maximum period of 5 days; (S2)
- d. except when intended for ophthalmic use. (S4)

ANNEXURE 1A: EMERGENCY CARE PROVIDER (PARAMEDIC)

PARAMEDIC (National Diploma in Emergency Medical Care graduates only) registered with Health Professions Council of South Africa

PARAMEDIC (National D	iploma in Emergency Medical Care graduates only)
WATER	. Mater for injection
Substance	: Water for injection
Indication	: Diluent
Route of Administration	; Parenteral
WATER	
Substance	: Water for irrigation
Indication	: Wound and dressing irrigation
Route of Administration	: Solution

ANNEXURE 1B: EMERGENCY CARE PROVIDER (EMERGENCY CARE PRACTITIONER)

EMERGENCY CARE PRACTITIONER (Bachelor of Technology Degree in Emergency Medical Care) registered with Health Professions Council of South Africa

EMERGENCY CARE PR	ACTITIONER (B Tech: Emergency Medical Care)
WATER Substance Indication	: Water for injection : Diluent
Route of Administration	: Parenteral
WATER Substance	: Water for irrigation
Indication	: Wound and dressing irrigation
Route of Administration	: Solution

ANNEXURE 1C: BASIC AMBULANCE ASSISTANT

BASIC AMBULANCE ASSISTANT registered with Health Professions Council of South Africa

BASIC AMBULANCE ASSISTANT registered with Health Professions Council of South Africa		
WATER		
Substance	: Water for injection	
Indication	: Diluent	
Route of Administration	: Parenteral	
WATER		
Substance	: Water for irrigation	
Indication	: Wound and dressing irrigation	
Route of Administration	: Solution	

ANNEXURE 1D: AMBULANCE EMERGENCY ASSISTANT

AMBULANCE EMERGENCY ASSISTANT registered with Health Professions Council of South Africa

WATER		
Substance	: Water for injection	
Indication	: Diluent	
Route of Administration	; Parenteral	
WATER		
Substance	: Water for irrigation	
Indication	: Wound and dressing irrigation	
Route of Administration	: Solution	

ANNEXURE 1E: EMERGENCY CARE TECHNICIAN

EMERGENCY CARE TECHNICIAN registered with Health Professions Council of South Africa

WATER		
Substance	: Water for injection	
Indication	: Diluent	
Route of Administration	: Parenteral	
WATER		
Substance	: Water for irrigation	

Schedule 1

EMERGENCY CARE TE	CHN	ICIAN registered with Health Professions Council of South Africa
Indication	:	Wound and dressing irrigation
Route of Administration		Solution

ANNEXURE 1F: EMERGENCY CARE ASSISTANT

EMERGENCY CARE ASSISTANT registered with Health Professions Council of South Africa

EMERGENCY CARE AS	SISTANT registered with Health Profess	sions Council of South Africa
WATER		
Substance	: Water for injection	
Indication	: Diluent	
Route of Administration	: Parenteral	-
WATER		
Substance	: Water for irrigation	
Indication	: Wound and dressing irrigation	The second section is a second
Route of Administration	: Solution	

ANNEXURE 4: PODIATRIST

PODIATRIST [(B.Tech degree in Podiatry)] registered with the Health Professions Council of South Africa in terms of the Health Professions Act, 1974 (Act 56 of 1974)

- END SCHEDULE 1 -

SCHEDULE 2

- All substances referred to in this Schedule are excluded when specifically packed, labeled, sold and used for --
 - 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.
- 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:
 - The salts and esters of such substances, where the existence of such salts and esters is possible; and
 - 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 18:	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

Arsenic:

- <u>except [preparations]</u> in oral dosage forms containing the equivalent of 0,01 percent or less of arsenic trioxide; (S1)
- b. except when intended for injection. (S4)

Benzydamine,

a. when intended for use human vaginal use; (S3)

- except preparations and mixtures containing 3 percent or less of benzydamine, when intended for application to the skin; (S0)
- except preparations and mixtures intended for use as a mouth rinse or for topical application in the mouth and throat; provided that the total dose swallowed does not exceed 36 milligrams of benzydamine per day; (S1)
- d. except preparations and mixtures containing 3 milligrams or less of benzydamine per throat lozenge; Provided that the total dose swallowed does not exceed 36 milligrams of benzydamine per day and the pack size does not exceed 16 lozenges. (S0)*

Clonidine when intended for the [treatment] prevention of migraine. (S3)

Doxycycline,

a. when intended and labelled for the chemoprophylaxis of malaria in those aged 8 years and older
 [for periods not exceeding 4 months of continuous use]. (S4)

Flurbiprofen,

- a. when intended for the treatment of post-traumatic conditions, for a maximum period of 5 days; (S3)
- except when in the form of lozenges, indicated for the relief of pain associated with sore throats, subject to:
 - (i) a maximum of 8,75 milligrams per lozenge;
 - (ii) a maximum treatment period of 3 days; and
 - (iii)a maximum pack size of 15 lozenges. (S1)
- c. except when intended for application to the skin and indicated for the symptomatic relief of localised pain and inflammation, provided that in the case of application by transdermal patch:
 - (i) use is restricted to adults and children 12 years and older; and
 - (ii) the treatment period is limited to a maximum of 4 weeks [S1] (S0)
- d. except when intended for ophthalmic use. (S4)

olopatidine.

ANNEXURE 1A: EMERGENCY CARE PROVIDER (PARAMEDIC)

PARAMEDIC (National Diploma in Emergency Medical Care graduates only) registered with Health Professions Council of South Africa

PARAMEDIC(National Di	ploma in Emergency Medical Care graduates only)
SELECTIVE \$2 AGONIST	rs -
Substance	: Salbutamol
Indication	; Bronchodilator
Route of Administration	: Aerosol
NON-STEROIDAL ANTI-	NFLAMMATORY
Substance	: Ibuprofen
Indication	: Analgesic/ Anti-inflammatory
Route of Administration	: Oral
ANALGESIC	
Substance	: Paracetamol
Indication	: Analgesic/ Anti-pyrexia
Route of Administration	: Oral

ANNEXURE 1B: EMERGENCY CARE PROVIDER (EMERGENCY CARE PRACTITIONER)

EMERGENCY CARE PRACTITIONER

(Bachelor of Technology Degree in Emergency Medical Care) registered with Health Professions Council of South Africa

EMERGENCY CARE PR	
(B Tech: Emergency Med	cal Care)
SELECTIVE B2 AGONIST	S THE STATE OF THE
Substance	: Salbutamol
Indication	; Bronchodilator
Route of Administration	: Aerosol
ANTI-SPASMODIC	
Substance	: Hyoscine butylbromide
Indication	; Anti-spasmodic .
Route of Administration	: Oral
ANTI-PROPULSIVE	
Substance	: Loperamide
Indication	: Symptomatic management of diarrhoea in adults
Route of Administration	: Oral

EMERGENCY CARE PRA (B Tech: Emergency Medi		
NON-STEROIDAL ANTI-I	NFLAMMATORY.	
Substance	: Ibuprofen	
Indication	: Analgesic/ Anti-inflammatory	
Route of Administration	: Oral	
ANALGESIC		
Substance	: Paracetamol	
Indication	: Analgesic/ Anti-pyrexia	
Route of Administration	: Oral	

ANNEXURE 1C: BASIC AMBULANCE ASSISTANT

BASIC AMBULANCE ASSISTANT registered with Health Professions Council of South Africa

*ANTI-CHOLINERGIC		
Substance		Ipratropium bromide
Indication	1	Inhalant Bronchodilator (atropine derivative anti-cholinergic)
Route of Administration	;	Respirator Solution
*SELECTIVE \$2 AGONIS	TS	
Substance	1	Salbutamol
Indication	:	Bronchodilator
Route of Administration		Aerosol

ANNEXURE 1D: AMBULANCE EMERGENCY ASSISTANT

AMBULANCE EMERGENCY ASSISTANT registered with Health Professions Council of South Africa

AMBULANCE EMERGEN	ICY A	SSISTANT registered with Health Professions Council of South Africa
ANTI-CHOLINERGIC		
Substance	1	Ipratropium bromide
Indication	4	Inhalant Bronchodilator (atropine derivative anti-cholinergic)
Route of Administration		Respirator Solution
SELECTIVE B2 AGONIST	S	
Substance	:	Salbutamol
Indication		Bronchodilator
Route of Administration	-	Aerosol

ANNEXURE 1E: EMERGENCY CARE TECHNICIAN

EMERGENCY CARE TECHNICIAN registered with Health Professions Council of South Africa

EMERGENCY CARE TEC	HNIC	AN registered with Health Professions Council of South Africa
ANTI-CHOLINERGIC		
Substance		Ipratropium bromide
Indication	:	Inhalant Bronchodilator (atropine derivative anti-cholinergic)
Route of Administration	:	Respirator Solution
SELECTIVE \$2 AGONIST	S	
Substance	:	Salbutamol
Indication	:	Bronchodilator
Route of Administration		Aerosol

ANNEXURE 1F: EMERGENCY CARE ASSISTANT

EMERGENCY CARE ASSISTANT registered with Health Professions Council of South Africa

EMERGENCY CARE ASS	SISTA	NT registered with Health Professions Council of South Africa
ANTI-CHOLINERGIC	PERSON	
Substance		Ipratropium bromide
Indication	1	Inhalant Bronchodilator (atropine derivative anti-cholinergic)
Route of Administration		Respirator Solution ·
SELECTIVE 82 AGONIST	S	
Substance		Salbutamol
Indication	:	Bronchodilator
Route of Administration		Aerosol

ANNEXURE 4: PODIATRIST

PODIATRIST [(B.Tech degree in Podiatry)] registered with the Health Professions Council of South Africa in terms of the Health Professions Act, 1974 (Act 56 of 1974)

- END SCHEDULE 2-

SCHEDULE 3

- All substances referred to in this Schedule are excluded when specifically packed, labelled, sold and used for --
 - 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.
- 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:
 - The salts and esters of such substances, where the existence of such salts and esters is possible; and
 - 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);
	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

Benzydamine, except preparations and mixtures-

- a. containing 3 percent or less of benzydamine, when intended for application to the skin; (S0) [(S1)]
- [containing 0,15 percent or less of benzydamine, when] intended for use as a mouth rinse or for topical application in the mouth and throat; provided that the total dose swallowed does not exceed 36 milligrams of benzydamine per day; (S1)

- c. containing 3 milligrams or less of benzydamine per throat lozenge: Provided that the total dose swallowed does not exceed 36 milligrams of benzydamine per day and the pack size does not exceed 16 lozenges. (S0)"
- d. intended for human vaginal use. (S2)

Clonidine except when intended for the [treatment] prevention of migraine. (S2)

Dexketoprofen trometamol.

(-)-6 epigallocatechin gallate

Flurbiprofen, except

- a. when intended for the treatment of post-traumatic conditions, for a maximum period of 5 days; (S3)
- when in the form of lozenges, indicated for the relief of pain associated with sore throats, subject to:
 - (i). a maximum of 8,75 milligrams per lozenge;
 - (ii). a maximum treatment period of 3 days; and
 - (iii). a maximum pack size of 15 lozenges. (S1)
- b. except when intended for application to the skin and indicated for the symptomatic relief of localised pain and inflammation, provided that in the case of application by transdermal patch:
 - (i) use is restricted to adults and children 12 years and older; and
 - (ii) the treatment period is limited to a maximum of 4 weeks. [(S1)] (S0)
- c. when intended for the treatment of post-traumatic conditions, for a maximum period of 5 days;
 (S2)
- d. when intended for ophthalmic use; (S4)

Nicardipine.

Umeclidinium.

Schedule 3

ANNEXURE 1A: EMERGENCY CARE PROVIDER (PARAMEDIC)

PARAMEDIC (National Diploma in Emergency Medical Care graduates only) registered with Health Professions Council of South Africa

ANALGESIC	
Substance	; Paracetamol
Indication	: Analgesic/ Anti-pyrexia
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Ringers Lactate
Indication	: Plasma expanders
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Polygeline
Indication	: Plasma expanders
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Balanced Salt Solution
Indication	: Plasma expanders
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Sodium Bicarbonate 8,5 %
Indication	: Metabolic acidosis
Route of Administration	: Parenteral

ANNEXURE 1B: EMERGENCY CARE PROVIDER (EMERGENCY CARE PRACTITIONER)

EMERGENCY CARE PRACTITIONER

(Bachelor of Technology Degree in Emergency Medical Care) registered with Health Professions Council of South Africa

EMERGENCY CARE PR	ACTITIONER(B Tech: Emergency Medical Care)
ANALGESIC	
Substance	; Paracetamol
Indication	: Analgesic/ Anti-pyrexia
Route of Administration	: Parenteral
*ANTI-SPASMODIC	
Substance	: Hyoscine butylbromide
Indication	: Anti-spasmodic
Route of Administration	: Parenteral
**ARTERIAL SMOOTH M	IUSCLE AGENT
Substance	: Hydralazine
Indication	: Hypertension in pregnancy
Route of Administration	: Oral
BETA BLOCKER	
Substance	; Labetalol
Indication	: Hypertension in pregnancy
Route of Administration	; Parenteral
*CALCIUM CHANNEL BL	OCKER
Substance	: Nifedipine
Indication	: Hypertension in pregnancy
Route of Administration	: Parenteral
**CLASS III ANTI-ARRHY	THMIC
Substance	: Sotalol
Indication	: Anti-arrhythmic
Route of Administration	: Oral/ Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Ringers Lactate
Indication	; Plasma expanders
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Polygeline
Indication	: Plasma expanders
Route of Administration	: Parenteral

Schedule 3

EMERGENC! CARE PR	4CIIII	ONER(B Tech: Emergency Medical Care)	
PLASMA SUBSTITUTES	AND C	COLLOID SOLUTIONS	
Substance	:	Balanced Salt Solution	
Indication	:	Plasma expanders	
Route of Administration		Parenteral	
PLASMA SUBSTITUTES	AND C	COLLOID SOLUTIONS	
Substance	1	Sodium Bicarbonate 8,5 %	
Indication	:	Metabolic acidosis	
Route of Administration	:	Parenteral	
**VASODILATOR			
Substance	:	Isosorbide dinitrate	
Indication		Acute pulmonary syndrome/ Acute pulmonary oedema	
Route of Administration		Parenteral	

ANNEXURE 1C: BASIC AMBULANCE ASSISTANT

BASIC AMBULANCE ASSISTANT registered with Health Professions Council of South Africa

*SELECTIVE B2 AGONIST	TS				
Substance		Salbutamol			
Indication:		Bronchodilator			
Route of Administration		Inhalant			

ANNEXURE 1D: AMBULANCE EMERGENCY ASSISTANT

AMBULANCE EMERGENCY ASSISTANT registered with Health Professions Council of South Africa

		SSISTANT registered with Heal	
PLASMA SUBSTITUTES	AND C	COLLOID SOLUTIONS	
Substance	- 0	Dextran	
Indication		Plasma expanders	
Route of Administration	:	Parenteral	

PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Hydroxyethyl Starch
Indication	: Plasma expanders
Route of Administration:	Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Sodium chloride
Indication	: Plasma expanders
Route of Administration:	Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Ringers Lactate
Indication	: Plasma expanders
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Polygeline
Indication	: Plasma expanders
Route of Administration	: Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Balanced Salt Solution
Indication	: Piasma expanders
Route of Administration	; Parenteral
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS
Substance	: Sodium Bicarbonate 8,5 %
Indication	: Metabolic acidosis
Route of Administration	: Parenteral
*CARBOHYDRATES	
Substance	: Dextrose
Indication	: Nutrition / Acute Symptomatic Hypoglycaemic Treatment in adults and paediatrics
Route of Administration	: Parenteral
*CO-ENZYME	
Substance	: Thiamine (Vitamin B1)
Indication	Nutritional supplement/ Vitamin B (Emergency treatment of Wernicke's encephalopathy and Beriberi)
Route of Administration	; Parenteral
OTHER MINERAL SUPP	EMENTS
Substance	: Magnesium sulphate
Indication	: Mineral supplement; prevention and control of seizures and hypertension in toxaemia of pregnancy
Route of Administration	; Parenteral

SELECTIVE \$2 AGONIST	S		
Substance		Salbutamol	
Indication:	1	Bronchodilator	
Route of Administration	:	Inhalant	

ANNEXURE 1E: EMERGENCY CARE TECHNICIAN

EMERGENCY CARE TECHNICIAN registered with Health Professions Council of South Africa

PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS	
Substance	: Dextran	
Indication	: Plasma expanders	
Route of Administration	: Parenteral	
PLASMA SUBSTITUTES	AND COLLOID SOLUTIONS	
Substance	: Hydroxyethyl Starch	
Indication	: Plasma expanders	
Route of Administration:	Parenteral	
PLASMA SUBSTITUTES	ND COLLOID SOLUTIONS	
Substance	: Sodium chloride	
Indication	: Plasma expanders	
Route of Administration :	<u>Parenteral</u>	
PLASMA SUBSTITUTES	ND COLLOID SOLUTIONS	
Substance,	: Ringers Lactate	
Indication	: Plasma expanders	
Route of Administration	: Parenteral	
PLASMA SUBSTITUTES	ND COLLOID SOLUTIONS	
Substance	: Polygeline	
Indication	: Plasma expanders	
Route of Administration	: Parenteral	

Substance		Balanced Salt Solution
Indication		Plasma expanders
Route of Administration		Parenteral
PLASMA SUBSTITUTES A	ND C	COLLOID SOLUTIONS
Substance	:	Sodium Bicarbonate 8,5 %
Indication	:	Metabolic acidosis
Route of Administration	:	Parenteral
CARBOHYDRATES		
Substance	1	Dextrose
Indication	:	Nutrition / Acute Symptomatic Hypoglycaemic Treatment in adults and
		paediatrics
Route of Administration	<u> </u>	Parenteral
*CO-ENZYME		
Substance	1	Thiamine (Vitamin 81)
Indication	:	Nutritional supplement/ Vitamin B (Emergency treatment of Wernicke's encephalopathy and Beriberi)
Route of Administration	:	Parenteral Parenteral
OTHER MINERAL SUPPLE	MEN	<u>TS</u>
Substance		Magnesium sulphate
Indication	1	Mineral supplement; prevention and control of seizures and
		hypertension in toxaemia of pregnancy, Ventricular anti-arrhythmic
Route of Administration	1	Parenteral
ORGANIC NITRATES		
Substance	:	Glyceryl trinitrate
Indication	1	Vasodilator
Route of Administration	;	Oral
SELECTIVE 82 AGONISTS		
Substance	:	Salbutamol
Outstatice		
Indication:	:	Bronchodilator

ANNEXURE 1F: EMERGENCY CARE ASSISTANT

EMERGENCY CARE ASSISTANT registered with Health Professions Council of South Africa

PLASMA SUBSTITUTES	AND	COLLOID SOLUTIONS	
Substance		Dextran	
Indication	1	Plasma expanders	
Route of Administration	;	Parenteral	
PLASMA SUBSTITUTES	AND	COLLOID SOLUTIONS	
Substance		Hydroxyethyl Starch	
Indication	1	Plasma expanders	
Route of Administration:		Parenteral	
PLASMA SUBSTITUTES	AND	COLLOID SOLUTIONS	
Substance		Sodium chloride	
Indication	1	Plasma expanders	
Route of Administration:		Parenteral	
PLASMA SUBSTITUTES	AND	COLLOID SOLUTIONS	
Substance	<u>:</u>	Ringers Lactate	
Indication	j	Plasma expanders	
Route of Administration		Parenteral Parenteral	
PLASMA SUBSTITUTES	AND	COLLOID SOLUTIONS	
Substance		Polygetine	
Indication	<u> </u>	Plasma expanders	
Route of Administration	<u>i</u>	Parenteral	
PLASMA SUBSTITUTES	AND (COLLOID SOLUTIONS	
Substance	:	Balanced Salt Solution	
Indication	:	Plasma expanders	
Route of Administration		Parenteral	医自己性不足的
PLASMA SUBSTITUTES	AND (COLLOID SOLUTIONS	
Substance		Sodium Bicarbonate 8,5 %	
Indication	1	Metabolic acidosis	
Route of Administration		Parenteral	

CARBOHYDRATES		
Substance	1	Dextrose
Indication		Nutrition / Acute Symptomatic Hypoglycaemic Treatment in adults and paediatrics
Route of Administration	:	Parenteral
*CO-ENZYME	THE	
Substance	:	Thiamine (Vitamin B1)
Indication	- :	Nutritional supplement/ Vitamin B (Emergency treatment of Wernicke's encephalopathy and Beriberi).
Route of Administration		Parenteral
OTHER MINERAL SUPPL	EMEN	ITS
Substance	;	Magnesium sulphate
Indication	1	Mineral supplement; prevention and control of seizures and hypertension in toxaemia of pregnancy
Route of Administration		Parenteral
SELECTIVE B2 AGONIST	S	
Substance	:	Salbutamol
Indication:		Bronchodilator
Route of Administration	:	Inhalant

ANNEXURE 4: PODIATRIST

PODIATRIST [(B.Tech degree in Podiatry)] registered with the Health Professions Council of South Africa in terms of the Health Professions Act, 1974 (Act 56 of 1974)

- END SCHEDULE 3 -

Schedule 4

SCHEDULE 4

- All substances referred to in this Schedule are excluded when specifically packed, labelled, sold and used for –
 - 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.
- 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:
 - (ii) The salts and esters of such substances, where the existence of such salts and esters is possible; and
 - (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);
	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:	Podlatrist

Abemaciclib.

Acalabrutinib.

Arsenic:

- a. when intended for injection;
- b. except in oral dosage form. (S1, S2)

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Schedule 4	
Schedula	
COLICADIO 4	

Atezolizumab.

Baloxavir.

Baricitinib.

Benralizumab.

Besifloxacin.

Brolucizumab.

Carglumic.

Crisaborole.

Crisanlizumab.

Dimethyl fumarate.

Doxycycline, except

- a. when intended and labelled for the chemoprophylaxis of malaria in those aged 8 years and older
 [for periods not exceeding 4 months of continuous use]. (S2)
- in preparations thereof for the treatment of animals and registered in terms of the provisions of the Fertilizers, Farm Feeds, Agricultural Remedies and Stock Remedies Act, 1947 (Act 36 of 1947).

Doravirine.

Durvalumab.

Erenumab.

Flurbiprofen,

- a. when intended for ophthalmic use; (S4)
- except when in the form of lozenges, indicated for the relief of pain associated with sore throats, subject to:
 - (i) a maximum of 8,75 milligrams per lozenge;
 - (ii) a maximum treatment period of 3 days; and
 - (iii) a maximum pack size of 15 lozenges. (S1)
- c. except when intended for application to the skin, provided that in the case of application by transdermal patch:
 - (i) use is restricted to adults and children 12 years and older; and
 - (ii) the treatment period is limited to a maximum of 4 weeks. [(S1)] (S0)
- except when intended for the treatment of post-traumatic conditions, for a maximum period of 5 days; (S2)

Schedule 4

Follitropin delta.

Frovatriptan.

Galcanezumab.

Human normal immunoglobulin.

Lenvatinib.

Lokivetmab.

Ranolazine.

Saroglitazar magnesium.

Semaglutide.

Simoctogog alfa.

Siponimod.

ANNEXURE 1A: EMERGENCY CARE PROVIDER (PARAMEDIC)

PARAMEDIC (National Diploma in Emergency Medical Care graduates only) registered with the Health Professions Council of South Africa

*ANTI-FIBRINOLYTIC		
Substance	: Tranexamic acid	
Indication	: Major haemorrhage in trauma	
Route of Administration	; Parenteral	
**OXYTOCIN		
Substance	: Oxytocin	
Indication	: Post-partum haemorrhage	
Route of Administration	: Parenteral	
CORTICOSTEROID		
Substance	: Prednisolone	
Indication	: Glucocorticoid/ Steroidal anti-inflammatory	
Route of Administration	: Parenteral	
LOCAL ANAESTHETIC		
Substance	: Lignocaine hydrochloride	
Indication	: Local anaesthesia	
Route of Administration	: Parenteral	

ANNEXURE 18: EMERGENCY CARE PROVIDER (EMERGENCY CARE PRACTITIONER)

EMERGENCY CARE PRACTITIONER

(Bachelor of Technology Degree in Emergency Medical Care) registered with the Health Professions Council of South Africa

EMERGENCY CARE PRA	CTITIONER (B Tech: Emergency Medical Care)
**CORTICOSTEROID	
Substance	: Betamethasone
Indication	: Pre-term birth
Route of Administration	: Parenteral
**CORTICOSTEROID	
Substance	: Dexamethasone
Indication	: Pre-term birth
Route of Administration	: Parenteral
**DIRECT THROMBIN INH	IBITOR
Substance	: Bivalirudin
Indication	: Adjunct in percutaneous coronary angioplasty
Route of Administration	: Parenteral
**DOPAMINERGIC	
Substance	: Dopamine
Indication	: Haemodynamic support
Route of Administration	: Parenteral
**ADRENERGIC	
Substance	; Dobutamine
Indication	: Haemodynamic support
Route of Administration	: Parenteral
*ANTICHOLINESTERASE	
Substance	: Neostigmine
Indication	: Reversal of neuromuscular blockade
Route of Administration	: Parenteral
*CHOLINESTERASE INHIE	BITOR
Substance	: Sugammadex
Indication	: Reversal of neuromuscular blockade
Route of Administration	: Parenteral
*SEROTONIN ANTAGONIS	
Substance	: Ondansetron
Indication	: Post-operative nausea and vomiting
Route of Administration	: Parenteral

Schedule 4

*ANTI-FIBRINOLYTIC	
Substance	: Tranexamic acid .
Indication	: Major haemorrhage in trauma
Route of Administration	: Parenteral
OXYTOCIN	
Substance	: Oxytocin
Indication	: Post-partum haemorrhage
Route of Administration	: Parenteral
CORTICOSTEROID	
Substance	: Prednisolone
Indication	: Glucocorticoid/ Steroidal anti-inflammatory
Route of Administration	: Parenteral
LOCAL ANAESTHETIC	
Substance	: Lignocaine hydrochloride
Indication	: Local anaesthesia
Route of Administration	: Parenteral

ANNEXURE 1C: BASIC AMBULANCE ASSISTANT

BASIC AMBULANCE ASSISTANT registered with Health Professions Council of South Africa

BASIC AMBULANCE AS	SISTA	NT registered with Hea
*SELECTIVE B2 AGONIS	TS	
Substance		Fenoterol
Indication	:	Bronchodilator
Route of Administration	- :	Parenteral

ANNEXURE 1D: AMBULANCE EMERGENCY ASSISTANT

AMBULANCE EMERGENCY ASSISTANT registered with Health Professions Council of South Africa

*ADRENERGIC		
Substance	1	Adrenaline / Epinephrine
Indication	:	Sympathomimetic catecholamine
Route of Administration	1	Parenteral
*CORTICOSTEROIDS		
Substance		Methylprednisolone
Indication	;	Glucocorticoid / Steroidal Anti-Inflammatory
Route of Administration		Oral
*CORTICOSTEROIDS		
Substance	:	Hydrocortisone
Indication		Glucocorticoid / Steroidal Anti-Inflammatory
Route of Administration		Parenteral
*HYPERGLYCAEMIC AG	ENT	
Substance		Glucagon
Indication		Hyperglycaemic agent
Route of Administration		Parenteral
*OPIOID ANTAGONIST	NV.	
Substance	1	Naloxone hydrochloride
Indication	1	Opioid Antagonist / Narcotic Antagonist
Route of Administration		Parenteral
OPIOID ANTAGONIST		
Substance		Nitrous oxide
Indication	;	Analgesic Gas
Route of Administration	:	Inhalant (50:50 combination with Medical Oxygen)
SELECTIVE B2 AGONIST	S	
Substance		Fenoterol
Indication		Bronchodilator
Route of Administration		Parenteral
LOCAL ANAESTHETIC		
Substance	: 1	ignocaine hydrochloride
Indication	: L	ocal anaesthesia
Route of Administration		Parenteral

ANNEXURE 1E: EMERGENCY CARE TECHNICIAN

EMERGENCY CARE TECHNICIAN registered with Health Professions Council of South Africa

ADRENERGIC		
Substance		Adrenatine / Epinephrine
Indication	1	Sympathomimetic catecholamine
Route of Administration	:	Parenteral
CORTICOSTEROIDS		
Substance	1	Methylprednisolone
Indication	1	Glucocorticoid / Steroidal Anti-Inflammatory
Route of Administration	i	Oral
CORTICOSTEROIDS		
Substance		Hydrocortisone
Indication	1	Glucocorticoid / Steroidal Anti-Inflammatory
Route of Administration		Parenteral
HYPERGLYCAEMIC AGE	NT	
Substance	:	Glucagon
Indication	1	Hyperglycaemic agent
Route of Administration		Parenteral
ANTI-ARRHYTHMICS		
Substance		Amiodarone
Indication		Class III Anti-arrhythmic / Atrial & Ventricular
Route of Administration	:	Parenteral
*ANTI-EMETIC		
Substance		Metoclopramide monohydrochloride
Indication		Propulsive Anti-emetic / Dopamine Antagonist
Route of Administration	1	Parenteral
SELECTIVE B2 AGONIST	S	
Substance	1	Salbutamol
Indication:	-1	Bronchodilator
Route of Administration		Parenteral
SELECTIVE \$2 AGONIST	S	
Substance	1	Fenoteral
Indication	1	Brenchodilator
Route of Administration :		Parenteral

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ANTI-CHOLINERGIC	
Substance	: Atropine
Indication	: Competitive Anti-Cholinergic, Bradycardia, Anti-arrhythmic
Route of Administration	: Parenteral
OPIOID ANTAGONIST	
Substance	: Naloxone hydrochloride
Indication	: Opioid Antagonist / Narcotic Antagonist
Route of Administration	; Parenteral
OPIOID ANTAGONIST	
Substance	: Nitrous oxide
Indication	: Analgesic Gas
Route of Administration	: Inhalant (50:50 combination with Medical Oxygen)
**OXYTOCIN	
Substance	: Oxytocin
Indication	: Post-partum haemorrhage
Route of Administration	: Parenteral
CORTICOSTEROID	
Substance	: Prednisolone
Indication	: Glucocorticoid/ Steroidal anti-inflammatory
Route of Administration	: Parenteral
LOCAL ANAESTHETIC	
Substance	; Lignocaine hydrochloride
Indication	: Local anaesthesia
Route of Administration	: Parenteral

ANNEXURE 1F: EMERGENCY CARE ASSISTANT

EMERGENCY CARE ASSISTANT registered with Health Professions Council of South Africa

EMERGENCY CARE AS	SISTAN	IT registered with Health Professions Council of South Africa
*ADRENERGIC	+	
Substance		Adrenaline / Epinephrine
Indication		Sympathomimetic catecholamine
Route of Administration	;	Parenteral

CORTICOSTEROIDS		and the second s
Substance		Methylprednisolone
Indication	:	Glucocorticoid / Steroidal Anti-Inflammatory
Route of Administration	<u>:</u>	Oral
CORTICOSTEROIDS		
Substance		Hydrocortisone
Indication	1	Glucocorticoid / Steroidal Anti-Inflammatory
Route of Administration	:	Parenteral
HYPERGLYCAEMIC AGEN	T	
Substance	1	Glucagon
Indication		Hyperglycaemic agent
Route of Administration	:	Parenteral
ANTI-CHOLINERGIC		
Substance	3	Atropine
Indication		Competitive Anti-Cholinergic, Bradycardia, Anti-arrhythmic
Route of Administration		Parenteral
OPIOID ANTAGONIST		
Substance	:	Naloxone hydrochloride
Indication		Opioid Antagonist / Narcotic Antagonist
Route of Administration		Parenteral
OPIOID ANTAGONIST		
Substance		Nitrous oxide
Indication		Analgesic Gas
Route of Administration		Inhalant (50:50 combination with Medical Oxygen)
SELECTIVE B2 AGONISTS		
Substance		Fenoterol
Indication	1	Bronchodilator
Route of Administration		Parenteral

ANNEXURE 4: PODIATRIST

PODIATRIST [(B.Tech degree in Podiatry)] registered with the Health Professions Council of South Africa in terms of the Health Professions Act, 1974 (Act 56 of 1974)

- 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:
 - The salts and esters of such substances, where the existence of such salts and esters is possible; and
 - 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);
 - Annexure 1B: Emergency Care Provider (Emergency Care Practitioner),
 - 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 **

Armodafanil.

Recombinant human epidermal growth factor (rhEGF).

Schedule 5

ANNEXURE 1A: EMERGENCY CARE PROVIDER (PARAMEDIC)

PARAMEDIC (National Diploma in Emergency Medical Care graduates only) registered with the Health Professions Council of South Africa

PARAMEDIC (National D	iploma i	Emergency Medical Care gradua	ates)	
*INDUCTION AGENTS				
Substance		Ketamine		
Indication	:	Analgesia		
Route of Administration		Parenteral		

ANNEXURE 1E: EMERGENCY CARE TECHNICIAN

EMERGENCY CARE TECHNICIAN registered with Health Professions Council of South Africa

BENZODIAZEPINE DERI	VATIV	
Substance	:	Diazepam
Indication	:	Anti-convulsant/Sedative/Hypnotic
Route of Administration		<u>Parenteral</u>
*BENZODIAZEPINE DER	IVATIV	<u>re</u>
Substance		Midazolam
Indication		Antic-convulsant/Sedative/Hypnotic
Route of Administration		Parenteral
*BENZODIAZEPINE DER	IVATIV	<u>(E</u>
Substance	1	Lorazepam
Indication		Anti-convulsant/Sedative/Hypnotic
Route of Administration		<u>Parenteral</u>
BENZODIAZEPINE ANTA	GONIS	
Substance		Flumazenil
Indication	:	Benzodiazepine Antagonist
Route of Administration	- ;	Parenteral
NON-SELECTIVE ANTIHI	STAM	NE
Substance	4:15	Promethazine
Indication	1	Antihistamine
Route of Administration	Yalle.	Parenteral

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Schedule 5

BENZODIAZEPINE DERI	VATIVE		
Substance		Diazepam	
Indication		Anti-convulsant/Sedative/Hypnotic	
Route of Administration		Parenteral	
*BENZODIAZEPINE DER	IVATIVE		
Substance		Midazolam	
Indication		Antic-convulsant/Sedative/Hypnotic	
Route of Administration	:	Parenteral	
*BENZODIAZEPINE DER	IVATIVE		
Substance		Lorazepam	
Indication	1	Anti-convulsant/Sedative/Hypnotic	
Route of Administration	<u>:</u>	Parenteral	
BENZODIAZEPINE ANTA	GONIST		
Substance	•	Flumazenil	A Comment
Indication		Benzodiazepine Antagonist	
Route of Administration		Parenteral	

- END SCHEDULE 5 -

Schedule 6

SCHEDULE 6

- 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 (unless expressly excluded or unless listed in another Schedule):
 - the isomers of such substances, where the existence of such isomers is possible within the chemical designation;
 - 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.
- 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 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 6 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 1E: Emergency Care Technician

Dexamfetamine (Dexamphetamine) in medicines registered in terms of the Act and intended for the treatment of Attention-Deficit Hyperactivity Disorder. (S7)

ANNEXURE 1A: EMERGENCY CARE PROVIDER (PARAMEDIC)

PARAMEDIC (National Diploma in Emergency Medical Care graduates only) registered with the Health Professions Council of South Africa

PARAMEDIC (National D	iploma	in Emergency Medical Care graduates)
**ANALGESIC		
Substance		Fentanyl
Indication	:	Opioid/ Narcotic
Route of Administration	1	Parenteral Parenteral

ANNEXURE 1B: EMERGENCY CARE PROVIDER (EMERGENCY CARE PRACTITIONER)

EMERGENCY CARE PRACTITIONER

(Bachelor of Technology Degree in Emergency Medical Care) registered with the Health Professions Council of South Africa

EMERGENCY CARE PR	ACTIT	ONER		
(Bachelor of Technology I	Degree	in Emergency Medical Care) registered with the	Health Pro	fessions Council
of South Africa	The state of the			
ANALGESIC				
Substance		Fentanyl		
Indication '		Opioid/ Narcotic/ Induction of Anaesthesia		
Route of Administration		Parenteral		

ANNEXURE 1E: EMERGENCY CARE TECHNICIAN

EMERGENCY CARE TECHNICIAN registered with Health Professions Council of South Africa

*ANALGESICS				
Substance	:	Morphine sulphate		
Indication		Opioid/Narcotic		
Route of Administration		Parenteral		

Schedule 7

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):

- the isomers of such substances, where the existence of such isomers is possible within the chemical designation;
- 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.

Dexamfetamine (Dexamphetamine) except in medicines registered in terms of the Act and intended for the treatment of Attention-Deficit Hyperactivity Disorder (S6) [and its salts; preparations thereof. (S8)]

- END SCHEDULE 7 -

SCHEDULE 8

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):

- 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 such isomers of esters and 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.

[Dexamfetamine (Dexamphetamine) and it salts; preparations thereof. (S7)]

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

DR ZWELINI LAWRENCE MKHIZE, MP

MINISTER OF HEALTH

DATE: 23/11/20 20