GOVERNMENT OF ZAMBIA

STATUTORY INSTRUMENT No. 12 of 2016

The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

The Medicines and Allied Substances (Health Shops) Regulations, 2016

f Arrangement of f Regulations

PART I

PRELIMINARY

- 1. Title
- 2. Application of relevant Acts

PART II

HEALTH SHOP PERMIT

- 3. Application for permit
- 4. Request for information
- 5. Rejection of application for permit
- 6. Issuance of permit
- 7. Application for renewal of permit
- 8. Transfer of permit
- 9. Amendment of permit
- 10. Application for duplicate permit
- 11. Suspension of permit
- 12. Revocation of permit
- 13. Location of health shop

- 14. Sale of medicine
- 15. Identity of health shop
- 16. Dispensing of medicines in health shop
- 17. Storage of medicine

PART III

GENERAL PROVISIONS

18. Register of health shop permits

IN EXERCISE of the powers contained in section *thirty* of the Medicines and Allied Substances Act, 2013, the following Regulations are made:

PART I

PRELIMINARY

- 1. These Regulations may be cited as the Medicines and Allied Substances (Health Shops) Regulations, 2016.
- Title
- 2. In these Regulations, unless the context otherwise requires —

Interpretation

- "authorised supplier" means a holder of a pharmaceutical licence issued under section *thirty-four* of the Act;
- "health shop dispenser" means a person responsible for managing the health shop and has undergone training approved by the Authority;
- "patient pack" means a quantity of medicines sufficient to treat a single patient for a specified condition;
- "permit" means a health shop permit issued under section *thirty* of the Act;
- "re-packing of medicines" means the act of removing a preparation from its original primary container and placing it into a patient pack, but does not include the act of cutting of a blister pack;
- " responsible person " means a pharmacist or pharmacy technologist; and
- " supervising pharmacist " means a pharmacist providing supervisory services to a health shop.

PART II

HEALTH SHOP PERMIT

3. (1) A person shall apply to the Authority for a permit in Form I set out in the First Schedule.

Application for permit

- (2) The Authority shall, within fourteen days of the receipt of an application, notify the applicant of the decision of the Authority in respect of the application.
- (3) A separate application shall be made and a separate permit issued in respect of each premises.
- (4) The Authority may inspect the premises in respect of which an application for a permit is made to determine if the applicant meets the requirements of the Act and the guidelines issued by the Authority.

Request for information

4. The Authority may request an applicant to submit information in relation to an application in Form II set out in the First Schedule.

Rejection of application for permit

- 5. (1) The Authority shall reject an application for a permit if -
 - (a) the applicant fails to comply with any condition precedent to the issue of the permit;
 - (b) the permit issued to the applicant was revoked by the Authority within a period of two years preceding the date of the application; or
 - (c) the applicant is convicted of an offence under the Act or any other relevant written law.
- (2) The Authority shall, where it rejects an application under subregulation (1), inform the applicant within seven days of the decision in Form III set out in the First Schedule.

Issuance of permit

- 6. (1) The Authority shall, where the applicant meets the requirements of the guidelines issued by the Authority and the Act, issue a permit in Form IV set out in the First Schedule.
 - (2) A permit shall be valid for two years from the date of issue.
- (3) A health shop shall be managed by a health shop dispenser under the supervision of a responsible person.

Application for renewal of permit

- 7. (1) An application for the renewal of a permit shall be made to the Authority in Form V set out in the First Schedule.
- (2) The Authority shall, within fourteen days of the receipt of an application for the renewal of a permit, grant the application for the renewal of the permit if the applicant meets the requirements of the Act and the guidelines issued by the Authority and has complied with the terms and conditions of the permit.
- (3) The Authority shall, where it renews a permit, issue a new permit to the applicant.
- (4) A permit that is not renewed by the Authority lapses on its date of expiry.

Transfer of permit

- 8. (1) A permit shall be used solely by the holder and is not transferable to any other person without the prior approval of the Authority.
- (2) An application for approval to transfer a permit shall be made to the Authority in Form VI set out in the First Schedule.

- (3) The Authority shall, within thirty days of receipt of an application for the transfer of a permit, approve the transfer if the applicant meets the requirements of the Act, and issue the transferee with a permit.
- (4) The Authority shall reject an application for the transfer of a permit if the applicant fails to comply with the conditions for the grant of the permit, the provisions of the Act and the guidelines issued by the Authority.
- (5) The Authority shall, where it rejects an application to transfer a permit under subregulation (4)—
 - (a) inform the applicant in Form III set out in the First Schedule; and
 - (b) suspend or revoke the permit.
 - 9. (1) The Authority may amend a permit where—

Amendment of permit

- (a) some other person succeeds to the interest in the business belonging to the holder of the permit; or
- (b) the name of the business changes.
- (2) An application for the amendment of a permit shall be made in Form VII set out in the First Schedule.
- (3) The Authority shall communicate its decision to the permit holder within fourteen days of receipt of the application for the permit.
- (4) The Authority shall, where it approves the amendment of a permit, issue the applicant with a new permit.
- 10. A person shall, where that person's permit is lost, damaged or defaced, apply to the Authority for a duplicate permit in Form VIII set out in the First Schedule.

Application for duplicate permit

11. (1) The Authority shall suspend a permit if—

Suspension of permit

- (a) the holder operates the health shop under insanitary conditions:
- (b) the holder obtains or sells medicine from unauthorised suppliers or stocks and sells

unauthorised products;

- (c) the health shop in respect of which it was issued contravenes the prescribed standards;
- (d) the health shop is not managed or controlled by a responsible person determined by the Authority;

- (e) the responsible person fails to maintain the required records on medicines;
- (f) the health shop stocks and sells medicines that are not on the prescribed list; or
- (g) the holder contravenes the terms and conditions of the permit, the provisions of the Act or any other relevant written law.
- (2) The Authority shall, before suspending a permit, give notice to the holder of the intention to suspend the permit and request the holder to show cause, within a specified period, why the permit should not be suspended.
- (3) A notice of intention to suspend a permit shall be in Form IX set out in the First Schedule.
- (4) The Authority shall suspend a permit if the holder of the permit fails to take remedial measures within the period specified in the notice issued under sub-regulation (2).
- (5) A notice of the suspension of a permit shall be in Form X set out in the First Schedule.
- (6) The product affected by the suspension of the permit shall be quarantined at the cost of the permit holder during the period of the suspension of the permit.

Revocation of permit

- 12. (1) The Authority shall revoke a permit if the holder—
 - (a) contravenes the provisions of the Act or any other relevant written law or breaches the terms or conditions of the permit;
 - (b) fails to take corrective measures following the suspension of the permit within the specified period;
 - (c) changes the health shop premises without authorisation; or
 - (d) obtained the permit by fraud or deliberate or negligent submission of false information or statements.
- (2) The Authority shall, before revoking a permit, give notice to the holder of the permit of the intention to revoke the permit and request the holder to show cause, within a specified period, why the permit should not be revoked.
- (3) A notice of the intention to revoke a permit shall be in Form IX set out in the First Schedule.
- (4) The Authority shall revoke a permit if the holder fails to take remedial measures during the period specified by the Authority.

- (5) A notice of the revocation of a permit shall be in Form X set out in the First Schedule.
- (6) The Authority shall, where it determines that the holder stocks medicines in the health shop under insanitary conditions, direct the holder to dispose of the medicine.
- (7) A holder of a permit shall, where the permit is revoked, quarantine the products on the premises and dispose of the products as directed by the Authority at the holder's cost.
- 13. (1) The Authority shall, in considering an application for a permit, prioritise the submissions filed by applicants in the following areas:

Location of health shop

- (a) rural areas and districts where access by the members of the public to medicines is limited; and
- (b) peripheral areas of big towns or cities, where access to medicines is limited.
- (2) The Authority shall, when considering the grant of a permit, take into account the availability of dispensing facilities in the area with respect to which the permit relates.
- 14. (1) A holder of a permit shall only sell the medicines prescribed in the Second Schedule.

Sale of medicines

- (2) A health shop shall not stock for sale prescription only medicines and pharmacy sale medicines which are not specified in the prescribed list for health shops.
- 15. A health shop shall affix a standard logo for purposes of identifying the health shop as specified in the guidelines issued by the Authority.

Identity of health shop

16. Dispensing of medicines in a health shop shall be in accordance with the guidelines for dispensing of medicines in a health shop issued by the Authority.

Dispensing of medicines in health shop

17. The storage of medicines in a health shop shall be in the patient pack size and under conditions stipulated by the manufacturer.

Storage of medicines

PART III

GENERAL PROVISIONS

Register of health shop permits

- 18. (1) The Authority shall keep and maintain a register of health shop permits in Form XI set out in the First Schedule.
- (2) The register referred to in subregulation (1) shall be kept at the offices of the Authority and shall be open to inspection by the public at such times and upon payment of an inspection fee as prescribed in the Medicines and Allied Substances (Fees) Regulations, 2015.

FIRST SCHEDULE

(Regulations 3, 4, 5, 6, 7, 8, 9, 10, 11 and 12)

Form I (Regulation 3 (1)) (To be completed in triplicate)



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

	01 1 1 0 11		
Please complete in block letters	Shaded fields for official use	Application No.	
	only	Date and Time	-
Information Required	Information P	rovided	1
PARTIC	PART I CULARS OF APPL	ICANT	
1. (a) Name of business entity		acia (1	
(b) Registration No.	1		
2. Type of business entity			
3. Business premises			
(a) Plot No:	+		
(b) Street:			
(c) Telephone No:	1		
(d) Fax No:	15-		
(e) Mobile No:			
(f) Email address			
(g) Village			
(h) Chief			
(1) Iown			
(j) District			
(k) Province			
	OCATION OF H	EALTH SHOP	
4. Name of health shop:			
Physical Address			
6. Postal Address	· E		
	OFHEALTH SHO	OP DISPENSER	
7. Name:			

8. 9.		
- 10	Registration No: Date of Issue:	
10.	Signature:	
	PARTICULARS OF RESPONSIBLE I	PERSON
	(a) Name	
	(b) Registration No.	
	(c) Date of issue:	
	(d) Signature:	
8.	Attachments	
	(a) Valid Practicing Certificate for the	e Responsible Person
	(b) Sketch of the floor plan of the pre	emises
ı		
		I that submission of false information shall I is granted, it shall be revoked and the permit
revol		l is granted, it shall be revoked and the permit of the Applicant
revol	iculars of the Person signing on behalf Name	l is granted, it shall be revoked and the permit of the Applicant Designation
revol	ced.	l is granted, it shall be revoked and the permit of the Applicant Designation
Parti	sed. iculars of the Person signing on behalf Name	l is granted, it shall be revoked and the permit of the Applicant Designation
FOR Date Appl Payn	Name Signature OFFICIALUSE ONLY of Submission: ication Number: nent Receipt Number:	l is granted, it shall be revoked and the permit of the Applicant Designation Date
FOR Date Appl Payn Appl	Name Signature OFFICIAL USE ONLY of Submission: ication Number: nent Receipt Number: ication Accepted (Proceed for Inspection)	l is granted, it shall be revoked and the permit of the Applicant Designation Date
FOR Date Appl Payn Appl	Name Signature OFFICIAL USE ONLY of Submission: ication Number: nent Receipt Number: ication Accepted (Proceed for Inspection)	l is granted, it shall be revoked and the permit of the Applicant Designation Date

STAMP



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013

The Medicines and Allied Substances (Health Shops) Regulations, 2016

REQUEST FOR INFORMATION

<i>To:</i>
Address:
Application No:
You are requested to furnish the following information or documents in respect of your
application for
(a)
(b)
(e)
(d)
within days of this notice.
If you fail to furnish the requested information within the stipulated period, your application will be treated as invalid and shall be rejected.
Dated this
Director-General
OFFICIAL



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

(1) Here insert the full names and address of the applicant

(2) Here insert the reference No. of the application

The Medicines and Allied Substances (Health Shops) Regulations, 2016

NOTICE OF REJECTION	
IN THE MATTER OF (2)notified that your application for a health the Authority on the following grounds:	
(a)(b)	
(c)	
(d)	
Dated thisday of	, 20
	OFFICIAL STAMP

Director-General



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

The Medicines and Allied Substances (Health Shops) Regulations, 2016

HEALTH SHOPPERMIT

egistration No.: HS/	
ermit No.: HS/	
nis is to certify that (Name of Health Shop)	
(Physical Address)	
is registered to operat	
ame of Responsible person:	
ne conditions of the health shop permit are overleaf.	
Valid until	
Director-General	

OFFICIAL STAMP

Conditions for Health Shop Permit

- 1. Any change in the ownership, name and location of the health shop shall be approved by the Authority
- 2. The health shop shall only sell medicines that are on the prescribed list.
- 3. The premises and the manner in which the business is to be conducted must comply with the requirements of the Medicines and Allied Substances Act, No. 3 of 2013, and any other relevant written law.
- 4. The health shop permit is not transferable without the written approval of the Authority.
- 5. The health shop permit shall, upon grant, be displayed conspicuously at the front shop in a place visible to the public.

Form V (Regulation 7) (To be completed in triplicate)



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

APPLICATIO	N FOR RENEWAL	OFPERMIT	
Please complete in block letters	Shaded fields for official use only	Application No. Date and Time	
Information Required	Information Pr		_
1. Permit No.	Ingormanion I	0,1000	TV.
2. Registration No.	1		
3. Name of permit holder	1		
(a) Plot No:			
(b) Street:	17		
(c) Telephone No:			
(d) Fax No:			
(e) Mobile No:			
(f) Email address			
(g) Village			
(h) Chief	1		
(1) Town			90
(j) District			
(k) Province			0
5. Appendix	1		
Annual Report (a) Monthly records of qua (b) Monthly records of nar (c) Monthly records of pre (d) Monthly records of med	nes and receipts fro escriptions for PoM dicines stock-on-ha	om authorized suppliers As/Ps dispensed and	
Name of Applicant (individual or au	thorised representa	ative)	
Date:		gnature:	
FOR OFFICIAL USE ONLY			1
Received by:			
Serial No. of application:			
		STAMP	



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

	APPLICATION FOR	TRANSFER OF HE	ALTH SHOPPERMIT	
Please	complete in block letters	Shaded fields for official use only	Application No. Date and Time	
Inform	ation Required		Information Provided	V
	PART	ICULARS OF APPL	ICANT	
1.	(a) Name of business entity	/		
	(b) Registration No.			
2.	Type of entity			
3.	Business premises			
1	(a) Plot No:			
	(b) Street:			
	(c) Telephone No:			
	(d) Fax No:			
-	(e) Mobile No:			
	(f) Email address			
	(g) Village			- 1 -
1	(h) Chief			
	(i) Town			
	(j) District			
1	(k) Province			
	PARTIC	CULARS OF TRANS	SFEREE	
1.	(a) Name of business ent	ity		
	(b) Registration No.			_
2.	Type of entity			
3.	Business premises			
	(a) Plot No:			
	(b) Street:			
	(c) Postal address			
	(a) Telephone No:			
-	(e) Faxivo.			
	(f) Mobile No:			
	(g) Email address			
	(h) village			
	(1) IOWN			

	(j) District		
	(k) Province		
4.	Appendix		
	Contract of sale or acquisition of business between	een the current permit holder	
	and the proposed permit holder		
	DECLARATION AND SIG	CNATURE	
I doe	lare that all the information I have stated in this a		to the
		• •	
	of my knowledge and belief. I understand that sul		
	er the application void and that if approval is gran	ted, it shall be revoked and the p	ermit
revo	ked.		
Part	iculars of the Person signing on behalf of the	Annlicant	
1 41 0		жррпсанс	
•••••	Name	Designation	
		Designation Date	
FOR	Name		
	Name Signature COFFICIALUSE ONLY	Date	
Date	Name Signature OFFICIAL USE ONLY of Submission:	Date	
Date Appl	Name Signature OFFICIAL USE ONLY of Submission:	Date	
Date Appl Payr	Name Signature OFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number:	Date	
Date Appl Payr Appl	Name Signature COFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number: ication Accepted (Proceed for Inspection):	Date	
Date Appl Payr Appl	Name Signature OFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number:	Date	
Date Appl Payr Appl	Name Signature COFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number: ication Accepted (Proceed for Inspection):	Date	
Date Appl Payr Appl	Name Signature COFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number: ication Accepted (Proceed for Inspection):	Date	
Date Appl Payr Appl	Name Signature COFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number: ication Accepted (Proceed for Inspection):	Date	

Form VII

(Regulation 9)

(To be completed in triplicate)



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

	APPLICATION FOR A	MENDMENT OF A H	EALTH SHOPPERMIT	
Please complete in block letters		Shaded fields for official use	Application No.	
		only	Date and Time	
Infor	mation Required	Information Pro	vided	V
	PART	ICULARS OF APPLI	CANT	
1.	(a) Name of business entity			
	(b) Registration No.			
2.	Type of entity			
3.	Business premises			
	(a) Plot No:			
	(b) Street:			
l t	(c) Postal address			
	(d) Telephone No:			
	(e) Fax No:			
	(f) Mobile No:			
Ιt	(g) Email address			
1	(h) Village			
1	(i) Town			7
1	(j) District			- 1
	(k) Province			
4.	PARTICULARS OF AMEN	DMENT DESC	CRIPTION OF AMENDM	IENT(S)
l 1	1.			
	2.			
l t	3.			-

5.	EXISTING	PROPOSED AMENDMENT	REASONS FOR AMENDMENT
	6. Appendix		
F		ents relating to proposed amendmen	t as required by the Authroity
		AND SIGNATURE	
	eclare that all the wledge and belie	e information I have stated is correctef.	t and truthful to the best of my
Part	iculars of the F	Person signing on behalf of the Ap	oplicant
		Name	Designation
	Siz	gnature	Date
FOF	OFFICIALUS	SE ONLY	
Date	of Submission:		
App!	lication Number		
Payr	nent Receipt Nu	mber:	
App.	lication Accepted	d (Proceed for Inspection):	
App!	lication Rejected	l (Notify Applicant)	
			OFFICIAL STAMP

Form VIII (Regulation 10) (To be completed in triplicate)



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

	APPLICAT	ION FOR DUPLICAT	TE PERMIT	
Please	complete in block letters	Shaded fields for official use	Application No.	
		only	Date and Time	
Information Required		Information Pro	vided	l V
1.	Name of business enti	ity		
2.	Permit No.			
3.	Registration No.			
4.	Address			
5.	Affidavit of loss of perm	nit		
	Name		Designation	
	Signature OFFICIAL USE ONLY		Date	
Date o	of Submission:			
Applie	cation Number:			
Payme	ent Receipt Number:			
Applie	cation Accepted (Proceed for	Inspection):		
Applie	cation Rejected (Notify Appli	cant)		
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			SIAN	ЛP



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

The Medicines and Allied Substances (Health Shops) Regulations, 2016

NOTICE OF INTENTION TO SUSPEND/REVOKE HEALTH SHOP PERMIT

(1) Here insert the full names and address of	<i>To</i> (1)
holder of permit	
(2) Here insert the Permit No.	IN THE MATTER OF (2)
	(a)
	(b)
	(c)
	(d)
(3) Here insert the number of days stipulated	Accordingly, you are requested to show cause why your permit should not be suspended/revoked and to take action to remedy the breaches set out in paragraphs
	Dated thisday of
(4) Signature of Director-	(4)
of Director- General	Director-General

*Delete as appropriate

OFFICIAL STAMP



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

The Medicines and Allied Substances (Health Shops) Regulations, 2016

NOTICE OF SUSPENSION OR REVOCATION OF

	HEALITISHOT LERVITI		
(1) Here insert the full	<i>To</i> (1)		
names and address of holder of			
permit			
(2) Here insert the	IN THE MATTER OF (2)		
Permit No.	the Authority intends to *suspend/revoke your permit on the following grounds:		
	(a)		
	(b)		
	(c)		
	(d)		
	Dated thisday of	20	
	_		
	(4)	OFFICIAL STAMP	
	Director-General		

^{*}Delete as appropriate



The Medicines and Allied Substances Act, 2013 (Act No. 3 of 2013)

	REGISTER OF HEALTH SHOP PERMITS				
No.	Name and Address of business	Permit Number	Registration number	Date of issue	Expiry Date
1.					
2.					
3.)	
4.					
5.)
6.					
7.					

SECOND SCHEDULE

(Regulation 14)

ItemStrengthMedicines for Asthma2mg1Salbutamol tablets2mg2Salbutamol Inhaler100mcg/doseAntibiotics3Amoxycillin tablets/capsules250mg4Amoxycillin oral suspension125mg/5ml5Co-trimoxazole tablets400/80 mg6Co-trimoxazole suspension200/40mg/5m7Doxycycline capsules/tablets100mg8Metronidazole tablets200mg9Tetracycline Hyclate Ointment1%10Silver sulfadiazine cream10gAntihelmentics11Albendazole tablets400mgAnti-inflammatory/Analgesics200mg13Hydrocortisone ointment/cream1%14Paracetamol tablets200mg15Acetylsalicylic acid (Aspirin) tablets300mg15Acetylsalicylic acid (Aspirin) tablets300mg16Nystatin oral suspension50mg/5ml, 1017Clotrimazole cream1%, 10%18Clotrimazole vaginal tablets100mg, 500m		Pack Size
Antibiotics 250mg 250mg 4 Amoxycillin tablets/capsules 250mg 125mg/5ml 5 Co-trimoxazole tablets 400/80 mg 6 Co-trimoxazole suspension 200/40mg/5m 7 Doxycycline capsules/tablets 100mg 8 Metronidazole tablets 200mg 9 Tetracycline Hyclate Ointment 1% 10 Silver sulfadiazine cream 10g Anti-inflammatory/Analgesics 12 Ibuprofen tablets 200mg 13 Hydrocortisone ointment/cream 1% Acetylsalicylic acid (Aspirin) tablets 300mg 15 Acetylsalicylic acid (Aspirin) tablets 300mg 17 Clotrimazole cream 1%, 10% 10 10 10 10 10 10 10		T went stage
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Amoxycillin tablets/capsules Amoxycillin oral suspension 125mg/5ml 200/80 mg Co-trimoxazole tablets Doxycycline capsules/tablets Metronidazole tablets Tetracycline Hyclate Ointment Silver sulfadiazine cream Antihelmentics Anti-inflammatory/Analgesics I buprofen tablets Domg Hydrocortisone ointment/cream Hydrocortisone ointment/cream Anti-fungal Agents Nystatin oral suspension Clotrimazole cream 250mg 125mg/5ml 400/80 mg 200mg 100mg 400mg 400mg 400mg 400mg 400mg 400mg Anti-inflammatory/Analgesics 100mg, 500mg 50mg/5ml, 100 17 Clotrimazole cream 1%, 10%		1
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	00,000 UI/ml	Patient Pack
18 Clotrimazole vaginal tablets 100mg 500m		Patient Pack
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	Anti-malarials		
19	Artemether-Lumefantrine tablets	20/120mg	Patient Pack
	Laxatives		
20	Bisacodyl tablets	5mg	Patient Pack
		•	•
	Anti-histamines		
21	Cetirizine hydrochloride tablets	10mg	Patient Pack
22	Cetirizine hydrochloride oral solution	5mg/5ml	Patient Pack
23	Chlorpheniramine Maleate tablets	4mg	Patient Pack
24	Chlorpheniramine Maleate syrup	2mg/5ml	Patient Pack
	Oral Contraceptives		
25	Ethinylestardiol + Northisterone tablets	0.03mg/0.3mg	Patient Pack
26	Ethinylestardiol + Levonogestrel tablets	0.03mg/0.15mg	Patient Pack
	Minerals/Vitamins		
27	Vitamin B Complex tablets		Patient Pack
28	Zinc Sulfate tablets	20mg	Patient Pack
	Fluids and Electrolytes		
29	Normal Saline IV		0.90% 1 Liters
30	Ringers Lactate IV		1Litres

PART B

All general sale medicines.

Note: Patient pack means a quantity of medicines sufficient to treat a single patient for a specified condition.

Dr. J. Kasonde, Minister of Heath

Lusaka 27th January, 2016 [MH/101/16/1]