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

ARRANGEMENT OF REGULATIONS

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- 2. Application of relevant Acts

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HEALTH SHOP PERMIT

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- 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

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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

- These Regulations may be cited as the Medicines and Allied Title Substances (Health Shops) Regulations, 2016.
 - 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

- (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.

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(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)

APPLICATION	N FOR A HEALTH	SHOPPERMIT	
Please complete in block letters	Shaded fields for official use	Application No.	
	only	Date and Time	
Information Required	Information P	rovided	1
	PART I	S.A.	
	CULARS OF APPL	ICANT	
 (a) Name of business entity 	/		
(b) Registration No.			
Type of business entity			
Business premises			
(a) Plot No:			
(b) Street:	4 7		
(c) Telephone No:			
(d) Fax No:	Y THE STATE OF THE		
(e) Mobile No:			
(f) Email address	1		
(g) Village			
(h) Chief			9/-
(i) Town			
(j) District			
(k) Province			
PROPOSED	LOCATION OF HI	EALTH SHOP	
4. Name of health shop:			
5. Physical Address			
6. Postal Address			
PARTICULAR	S OF HEALTH SHO	OP DISPENSER	
7. Name:	*		

	Registration No:	
9.	Date of Issue:	1
10.	Signature:	aupaou –
-	PARTICULARS OF RESPONSIBLE	PERSON
	(a) Name	
	(b) Registration No.	
	(c) Date of issue:	
	(d) Signature:	
8.	Attachments	
	(a) Valid Practicing Certificate for the	Responsible Person
Ì	(b) Sketch of the floor plan of the pre	emises
1		
best o	of my knowledge and belief. I understand or the application void and that if approva	in this application is correct and truthful to the that submission of false information shall lis granted, it shall be revoked and the permit
best of rende revok	of my knowledge and belief. I understand or the application void and that if approva	I that submission of false information shall I is granted, it shall be revoked and the permit of the Applicant
best of rende revok	of my knowledge and belief. I understand or the application void and that if approva ted. culars of the Person signing on behalf	I that submission of false information shall I is granted, it shall be revoked and the permit of the Applicant Designation
best of rende revok	of my knowledge and belief. I understand or the application void and that if approva ted. iculars of the Person signing on behalf	I that submission of false information shall I is granted, it shall be revoked and the permit of the Applicant Designation
best of rende revok	of my knowledge and belief. I understand or the application void and that if approva ted. iculars of the Person signing on behalf Name	I that submission of false information shall I is granted, it shall be revoked and the permi of the Applicant Designation
best or render revok	of my knowledge and belief. I understand or the application void and that if approvated. Iculars of the Person signing on behalf Name Signature	of the Applicant Designation Date
Parti	of my knowledge and belief. I understander the application void and that if approvated. I culars of the Person signing on behalf Name Signature OFFICIAL USE ONLY	that submission of false information shall l is granted, it shall be revoked and the permit of the Applicant Designation Date
FOR Payer	of my knowledge and belief. I understand on the application void and that if approvated. I culars of the Person signing on behalf Name Signature OFFICIAL USE ONLY of Submission: ication Number: ment Receipt Number:	that submission of false information shall l is granted, it shall be revoked and the permit of the Applicant Designation Date
render revok Parti FOR Date Appli	of my knowledge and belief. I understander the application void and that if approvated. Iculars of the Person signing on behalf Name Signature OFFICIAL USE ONLY of Submission: ication Number: ication Accepted (Proceed for Inspection)	that submission of false information shall l is granted, it shall be revoked and the permi of the Applicant Designation Date
render revok Parti FOR Date Appli	of my knowledge and belief. I understander the application void and that if approvated. Iculars of the Person signing on behalf Name Signature OFFICIAL USE ONLY of Submission: ication Number: ication Accepted (Proceed for Inspection)	that submission of false information shall l is granted, it shall be revoked and the permit of the Applicant Designation Date
render revok Parti FOR Date Appli	of my knowledge and belief. I understander the application void and that if approvated. Iculars of the Person signing on behalf Name Signature OFFICIAL USE ONLY of Submission: ication Number: ication Accepted (Proceed for Inspection)	that submission of false information shall l is granted, it shall be revoked and the permit of the Applicant Designation Date
FOR Date Appli	of my knowledge and belief. I understander the application void and that if approvated. Iculars of the Person signing on behalf Name Signature OFFICIAL USE ONLY of Submission: ication Number: ication Accepted (Proceed for Inspection)	that submission of false information shall l is granted, it shall be revoked and the permi of the Applicant Designation Date

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Form II (Regulation 4)



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

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

REQUEST FOR INFORMATION

To:	
Address:	
Application No:	
You are requested to furnish the following information or documents in respec	t 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 be treated as invalid and shall be rejected.	application will
Dated this day of	20
Director-General	
	OFFICIAL

STAMP

Form III (Regulations 5(2) and 8(5))



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

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

Director-General

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8(5))

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Form IV (Regulation 6(1))



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

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

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

HEALTH SHOP PERMIT

legistration No.:
ermit No.: HS/
his is to certify that (Name of Health Shop)
f (Physical Address)
ealth shop
Iame of Responsible person:
he conditions of the health shop permit are overleaf.
Valid until 20
Director-General

OFFICIAL STAMP

Conditions for Health Shop Permit

- 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.
- 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.

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pity.

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)

10.0	APPLICATIO	N FOR RENEWAL	OFPERMIT	
Please complete in block letters		Shaded fields for official use	Application No.	
		only	Date and Time	
_	rmation Required	Information Pr	rovided	
1.	Permit No.			I V
2.	Registration No.			
3.	Name of permit holder			-
	(a) Plot No:			
	(b) Street:			
-	(c) Telephone No:			
	(d) Fax No:			
	(e) Mobile No:			11
	(f) Email address			1
	(g) Village			+ 1
	(h) Chief			100
	(i) Town	-8-		
	(j) District			
	(k) Province			
5.	Appendix			V
	Annual Report (a) Monthly records of qua (b) Monthly records of nar (c) Monthly records of pre (d) Monthly records of me	mes and receipts from	om authorized suppliers Is/Ps dispensed	
	me of Applicant (individual or au	the first of the second of the second of the		
FOR	OFFICIAL USE ONLY			
Recei Amou	ved by:			
Serial	No. of application:		OFFICIAL STAMP	

Form VI (Regulation 8(2)) (To be completed in triplicate)



THE ZAMBIAMEDICINES REGULATORY AUTHORITY

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

Please complete in block letters Information Required		Shaded fields for official use	Application No.		
		only	Date and Time		
			Information Provided	1	
	PART	ICULARS OF APPLI	ICANT		
1.	(a) Name of business entity	/			
	(b) Registration No.			\neg	
2.	Type of entity				
3.	Business premises				
	(a) Plot No:				
	(b) Street:				
	(c) Telephone No:				
	(d) Fax No:				
	(e) Mobile No:				
	(f) Email address				
	(g) Village				
	(h) Chief				
	(i) Town				
	(j) District				
	(k) Province				
	PARTIC	CULARS OF TRANS	FEREE		
1.	(a) Name of business ent	itv			
	(b) Registration No.				
2.	Type of entity			\neg	
3.	Business premises				
	(a) Plot No:				
	(b) Street:				
	(c) Postal address				
	(d) Telephone No:				
	(e) Fax No:				
	(f) Mobile No:				
	(g) Email address				
	(h) Village				
	(i) Town				

	(j) District		
1	(k) Province		ī
4.	Appendix		_
	Contract of sale or acquisition of business b and the proposed permit holder	etween the current permit holder	
	DECLARATION ANI	SIGNATURE	Ī
best rend revo	clare that all the information I have stated in the of my knowledge and belief. I understand that the application void and that if approval is sked. ticulars of the Person signing on behalf of	at submission of false information shall granted, it shall be revoked and the permi	
	Name	Designation	•••
	Signature	Date	•••
FOF	R OFFICIAL USE ONLY		
App Payr	e of Submission: lication Number: ment Receipt Number: lication Accepted (Proceed for Inspection):		
App	lication Rejected (Notify Applicant)		
*******		OFFICIAL STAMP	

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 AN	MENDMENT OF A H	EALTH SHOP PERMIT		
Please complete in block letters		Shaded fields for official use	Application No.		
		only	Date and Time		
Infor	mation Required	Information Pro	vided	1	
	PART	ICULARS OF APPLI	CANT		
1.	(a) Name of business entity			T	
	(b) Registration No.				
2.	Type of entity				
3.	Business premises				
7	(a) Plot No:	1			
	(b) Street:				
	(c) Postal address				
	(d) Telephone No:		100 1100 1100		
	(e) Fax No:				
	(f) Mobile No:				
	(g) Email address				
	(h) Village				
	(i) Town				
	(j) District				
	(k) Province			. 10	
4.	PARTICULARS OF AMENI	DMENT DESC	CRIPTION OF AMENDMEN	NT(S)	
	1.				
	2.				
	3.				

5. EXIST	ING PROPOSED AMENDMENT	T REASONS FOR AMENDMENT
-		
	ppendix	
Relevant	documents relating to proposed amendr	ment as required by the Authroity
DECLARA	ATION AND SIGNATURE	
I declare th knowledge	at all the information I have stated is con and belief.	rrect and truthful to the best of my
Particulars	of the Person signing on behalf of the	e Applicant
	Name	Designation
	Signature	Date
FOR OFFIC	CIAL USE ONLY	
Date of Subn	nission:	
Application	Number:	
Payment Rec	ceipt Number:	
Application .	Accepted (Proceed for Inspection):	
	Rejected (Notify Applicant)	
		25.00.00
		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	1
Name of business entity		ity		
2.	Permit No.			
3.	Registration No.			
4.				
5.	Affidavit of loss of perm	nit		
1	Name		Designation	***********
+				
Signature FOR OFFICIAL USE ONLY			Date	
Date	of Submission:			
Appli	cation Number:			******
Paymo	ent Receipt Number:			
		Inspection):		
	cation Accepted (Proceed for			
	cation Accepted (Proceed for cation Rejected (Notify Appli			
	이 마음을 많아서 있다. 그리고 있는데 가지 않는데 되었다.			

Form IX (Regulation 11(3) and 12(3))



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

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 holder of permit (2) Here insert the Permit No.	IN THE MATTER OF (2) you are notified that the Authority intends to *suspend/revoke your permit on the following grounds: (a)
	(b)
(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 this
(4) Signature of Director- General	(4)

*Delete as appropriate

OFFICIAL STAMP

(1) Here insert the full names and address of holder of permit

(2) Here insert the Permit No.

Form X (Regulation 11 (5) and 12 (5))



THE ZAMBIA MEDICINES REGULATORY AUTHORITY

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 HEALTH SHOP PERMIT

To (1)	

IN THE MATTER OF (2)	ou are notified that following grounds:
(a)	
(b)	
(c)	
(d)	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
Dated thisday of	20
(4) Director-General	OFFICIAL STAMP

*Delete as appropriate

Form XI (Regulation 19)



ZAMBIA MEDICINES REGULATORY AUTHORITY

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.	12				
7.					

SECOND SCHEDULE

(Regulation 14)

	Item	Strength	Pack Size
	Medicines for Asthma		
	Salbutamol tablets	2mg	Patient Pack
	Salbutamol Inhaler	100mcg/dose	Patient Pack
	Antibiotics		
	Amoxycillin tablets/capsules	250mg	Patient Pack
	Amoxycillin oral suspension	125mg/5ml	Patient Pack
5	Co-trimoxazole tablets	400/80 mg	Patient Pack
5	Co-trimoxazole suspension	200/40mg/5ml	Patient Pack
7	Doxycycline capsules/tablets	100mg	Patient Pack
3	Metronidazole tablets	200mg	Patient Pack
)	Tetracycline Hyclate Ointment	1%	Patient Pack
0	Silver sulfadiazine cream	10g	Patient Pack
	Antihelmentics		
1	Albendazole tablets	400mg	Patient Pack
	Anti-inflammatory/Analgesics		
2	Ibuprofen tablets	200mg	Patient Pack
3	Hydrocortisone ointment/cream	1%	Patient Pack
4	Paracetamol tablets	100mg, 500mg	Patient Pack
5	Acetylsalicylic acid (Aspirin) tablets	300mg	Patient Pack
-	Anti-fungal Agents		
6	Nystatin oral suspension	50mg/5ml, 100,000 UI/ml	Patient Pack
7	Clotrimazole cream	1%, 10%	Patient Pack
8	Clotrimazole vaginal tablets	100mg, 500mg	Patient Pack

	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		T
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 [мн/101/16/1] 1

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