

Product List

1	Sertaconazole	Composition:				
	Nitrate	Sertaconazole Nitrate	BP	2%	w/w	
	Cream	Preservative:				APPROVED
		Benzyl Alcohol	IP	1%	w/w	
		Cream base		q.s		
2	Mometasone	Composition:				
	Furoate &	Mometasone Furoate	IP	0.1%	w/w	
	Fusidic Acid	Fusidic Acid	IP	2%	w/w	APPROVED
	Cream	Cream base		q.s		
3	Clobetasol	Composition:		1		
	Propionate	Clobetasol Propionate	IP	0.05%	w/w	
	Cream IP	Chlorocresol	IP	0.1%	w/w	APPROVED
		(as preservative)				111110 (22
		Cream base		q.s		
4	Hydroquinone,	Composition:				
	Tretinoin &	Hydroquinone	IP	2%	w/w	
	Mometasone	Tretinoin	USP	0.025%	w/w	APPROVED
	Furoate Cream	Mometasone Furoate	IP	0.1%	w/w	AITROVED
		Cream base		q.s		
5	Mometasone	Composition:				
	Furoate Cream IP	Mometasone Furoate	IP	0.1%	w/w	
		Chlorocresol	IP	0.1%	w/w	
		(as preservative)				APPROVED
		Cream base		q.s		
6	Fusidic Acid	Composition:	•		-	
	Cream IP	Fusidic Acid	IP	2%	w/w	APPROVED
	Crount II	Cream base		q.s	**/ **	
7	M			1"		
/	Mometasone	Composition: Mometasone Furoate	IP	0.1%	w/w	
	Furoate & Terbinafine HCl	Terbinafine HCl	BP	1%		APPROVED
	Cream	Cream base	DP		w/w	
		L		q.s		
8	Betamethasone	Composition:		1 1		
	& Miconazole	Betamethasone				
	Cream	Dipropionate	IP	0.050	,	
		Eq. to Betamethasone		0.05%	w/w	APPROVED
		Miconazole Nitrate	IP	201	,	
		Eq. to Miconazole		2%	w/w	
		Cream base		q.s		

9	Linseed Oil,	Composition:					
	diclofenac	Linseed Oil	BP	3%	w/w		
	diethylamine,	Diclofenac Diethylamine	BP/IP				
	Methyl	Eq. to Diclofenac Sodium		1%	w/w		
	Salicylate &	Methyl Salicylate	IP	10%	w/w	APPROVED	
	Menthol	Menthol	IP	5%	w/w		
	Gel	Benzyl Alcohol	IP	1%	w/w		
	361	(As Preservative)					
		In a water washable base		q.s			
10	Pantoprazole	Each enteric coated tablet co					
	Tablets IP 40mg	Pantoprazole Sodium	IP				
		Eq. to Pantoprazole		40	mg	APPROVED	
		Excipients		q.s		AFFRUVED	
		Approved colour used					
11	Aceclofenac &	Each film coated tablet cont					
	Paracetamol	Aceclofenac	IP	100	mg	APPROVED	
	Tablets	Paracetamol	IP	325	mg	AITROVED	
		Excipients		q.s			
		Approved colour used					
12	Ofloxacin &	Ofloxacin & Each film coated tablet contains :					
	Ornidazole	Ofloxacin	IP	200	mg		
	Tablets ip	Ornidazole	IP	500	mg	APPROVED	
	1	Excipients		q.s	- U	ATTROVED	
		Approved colour used		1			
13	Ofloxacin	Each film coated tablet cont	tains :				
	Tablets IP	Ofloxacin	IP	200	mg		
		Excipients		q.s	8	APPROVED	
		Approved colour used		1.~		APPROVED	
		**					
14	Azithromycin	Each film coated tablet cont					
	Tablets IP 250mg	Azithromycin	IP			APPROVED	
		Eq. to Anhydrous		250			
		Azithromycin		250	mg		
		Excipients		q.s			
		Approved colour used					
15	Azithromycin	Each film coated tablet cont	tains :				
	Tablets IP 500mg	Azithromycin	IP				
		Eq. to Anhydrous				APPROVED	
		Azithromycin		500	mg	III I I I I I I I I I I I I I I I I I	
		Excipients		q.s			
		Approved colour used		-			
16	Rabeprazole	Each enteric coated tablet co	ontains :				
	Tablets IP 20mg	Rabeprazole Sodium	IP	20	mg		
	8	Excipients	**		****5	APPROVED	
		Approved colour used		q.s	<u> </u>		
		Approved colour used					

Montelukast Tablets IP	17	Levocetirizine &	Each film coated tablet conta	ains :			
Tablets IP							
Montelukast Sodium IP 10 mg Eq. to Montelukast IP 10 mg Excipients q.s Approved colour used				IP	5	mg	
Excipients				IP			APPROVED
Excipients			Eq. to Montelukast		10	mg	
Approved colour used			Excipients		q.s		
Tablets IP 250mg			Approved colour used				
Tablets IP 250mg	18	Levofloxacin	Each film coated tablet conta	ains :			
Eq. to Levofloxacin 250 mg APPROVED			Levofloxacin Hemihydrate	IP			
Excipients					250	mg	APPROVED
Approved colour used Each film coated tablet contains : Levofloxacin Eq. to Levo			Excipients		q.s		MINOVED
Tablets IP 500mg Levofloxacin Hemihydrate			Approved colour used				
Tablets IP 500mg Levofloxacin Hemihydrate Eq. to Levofloxacin	19	Levofloxacin	Each film coated tablet conta				
Eq. to Levofloxacin			Levofloxacin Hemihydrate	IP			
Excipients					500	mg	APPROVED
Approved colour used Each film coated tablet contains : Tablets IP 100mg Eq. to Anhydrous Cefixime IP 100 mg Excipients Q.s Approved colour used Cefixime					q.s		MIKOVED
Cefixime Tablets IP 100mg			-				
Tablets IP 100mg	20	C-6-:	11				
100mg	20						
Excipients q.s Approved colour used				ır	100	ma	APPROVED
Approved colour used Cefixime Each film coated tablet contains : Cefixime IP 200mg Eq. to Anhydrous Cefixime 200 mg Excipients Q.s Approved colour used		Tooms	1 2			mg	
Cefixime Tablets IP			*		4.5		
Tablets IP 200mg			**				
200mg Eq. to Anhydrous Cefixime 200 mg Excipients q.s Approved colour used	21				1		
Excipients q.s APPROVED				IP	200		
Approved colour used 22		200mg	<u> </u>			mg	APPROVED
Cefpodoxime Each film coated tablet contains : Cefpodoxime Proxetil IP Eq. to Cefpodoxime 100 mg Excipients Approved colour used Approved Approve			*		q.s		
Tablets IP 100mg			**				
100mg Eq. to Cefpodoxime 100 mg Excipients q.s Approved colour used	22	Cefpodoxime		ns:			
Excipients q.s APPROVED Approved colour used				IP	100		
Approved colour used		100mg				mg	APPROVED
					q.s		
23 Cefpodoxime Each film coated tablet contains:	22	C C 1 ·		ne :			
Cerpodoxime	23	-					
Fa to Cefnodovime 200 mg				ır	200	mg	
Excipients Q.S APPROVED		200Hig	<u> </u>			5	APPROVED
Approved colour used			*	l	1		
24 Cefuroxime Each film coated tablet contains :	24	Cefuroxime	**				
Axetil Tablets Cefuroxime Axetil IP							A DDD CYTES
IP 250mg Eq. to Cefuroxime 250 mg APPROVED					250	mg	APPKOVED
Excipients q.s					q.s		
Approved colour used			Approved colour used				

25	Cefuroxime	Each film coated tablet contai				
	Axetil Tablets	Cefuroxime Axetil	IP			
	IP 500mg	Eq. to Cefuroxime	_	500	mg	A PPP OVER
	n cooms	Excipients		q.s	8	APPROVED
		Approved colour used		4.5		
		Approved colour used				
26	Cefixime &	Each film coated tablet contains				
	Ofloxacin	Cefixime	IP			
	Tablets	Eq. to Anhydrous Cefixime		200	mg	A PPP OVER
		Ofloxacin	IP	200	mg	APPROVED
		Excipients		q.s		
		Approved colour used				
27	Cefixime &	Each film coated tablet contains				
	Potassium	Cefixime	IP			
	Clavulanate	Eq. to Anhydrous Cefixime		200	mg	
	Tablets	Potassium Clavulanate Diluted	IP			APPROVED
		Eq. to Clavulanic Acid		125	mg	
		Excipients		q.s		
		Approved colour used				
28	Cefpodoxime &	Each film coated tablet contains				
20	Potassium	Cefpodoxime Proxetil	· IP			
	Clavulanate	Eq. to Cefpodoxime	ır	200	mg	
	Tablets	Potassium Clavulanate Diluted	IP	200	mg	
		Eq. to Clavulanic Acid	11	125	mg	APPROVED
		Excipients		q.s	1115	
		Approved colour used		1	l	
20		1 ^^				
29	Amoxycillin &	Each film coated tablet contains		1	Г	
	Potassium Clavulanate	Amoxycillin Trihydrate	IP	500		
	Tablets IP	Eq. to Amoxycillin Potassium Clavulanate Diluted	ID	500	mg	APPROVED
	Tablets IF	Eq. to Clavulanic Acid	IP	125	ma	III NO VED
		Excipients		+	mg	
		Approved colour used		q.s		
		1 **				
30	Cefixime Oral	Each 5ml of the reconstituted sus	spension co	ontains:		
	Suspension IP	Cefixime	IP			
		Eq. to Anhydrous Cefixime		50	mg	APPROVED
		Excipients		q.s		APPROVED
		In a flavoured base				
		Approved colour used				
31	Cefixime Oral	Each 5ml of the reconstituted	s:			
	Suspension IP	Cefixime	IP			
	•	Eq. to Anhydrous Cefixime		100	mg	APPROVED
		Excipients		q.s	J	ALLKUYED
		In a flavoured base		1	1	
		Approved colour used				
		11pproved colour used				

32	Cefpodoxime	Each 5ml of the reconstituted	suspensio	n contains	S:	
	Oral Suspension	Cefpodoxime Proxetil	IP			
	IP	Eq. to Cefpodoxime		50	mg	A PRO CAMER
		Excipients		q.s		APPROVED
		In a flavoured base				
		Approved colour used				
33	Cefpodoxime	Each 5ml of the reconstituted	suspensio	n contains	3:	
	Oral Suspension	Cefpodoxime Proxetil	IP			
	ΙΡ̈́	Eq. to Cefpodoxime		100	mg	APPROVED
		Excipients		q.s		ATTROVED
		In a flavoured base	I			
		Approved colour used				
24	A •11•	11				
34	Amoxycillin	Each 5ml of the reconstituted		n contains	3:	
	Oral Suspension	Amoxycillin Trihydrate	IP	125	ma	
	IP	Eq. to Amoxycillin Excipients			mg	
		In a flavoured base		q.s		APPROVED
		Approved colour used				
35	A movyoillin	Each 5ml of the reconstituted				
33	Amoxycillin Oral Suspension		IP		· ·	
	IP	Amoxycillin Trihydrate	IP	250	ma	
	П	Eq. to Amoxycillin Excipients			mg	A PRO CATED
		In a flavoured base		q.s		APPROVED
		Approved colour used				
36	Cephalexin Oral	Each 5ml of the reconstituted	suspensio	n contains	3:	
	Suspension IP	Cephalexin	IP			
		Eq. to Anhydrous				
		Cephalexin		125	mg	APPROVED
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				
37	Cefaclor Oral	Each 5ml of the reconstituted	suspensio	n contains	3:	
	Suspension IP	Cefaclor	IP	125	mg	
	Suspension n	Excipients		q.s		A PRIN COMME
		In a flavoured base		<u>. ^ l</u>		APPROVED
		Approved colour used				
38	Cefixime &	Each 5ml of the reconstituted	suspensio	n contains	3:	
	Potassium	Cefixime	IP			
	Clavulanate	Eq. to Anhydrous Cefixime		50	mg	
	Oral Suspension	Potassium Clavulanate				
	1	Diluted	IP			APPROVED
		Eq. to Clavulanic Acid	_	31.25	mg	
		Excipients		q.s		
		In a flavoured base	<u>l</u>	1.~		
		Approved colour used				
		PP10 (Ca Colour about				

39	Cefpodoxime &	Each 5ml of the reconstituted	suspensio	:				
	Potassium	Cefpodoxime Proxetil	IP					
	Clavulanate	Eq . to Cefpodoxime		50	mg			
	Oral Suspension	Potassium Clavulanate	TD			A DDD OVED		
		Diluted Eq. to Clavulanic Acid	IP	31.25	ma	APPROVED		
		Excipients			mg	-		
		In a flavoured base		q.s		-		
		Approved colour used				-		
40		11	. •			1		
40	Omeprazole	Each hard gelatin capsule con		1 20 1		-		
	Capsules IP	Omeprazole	IΡ	20	mg			
		(As enteric coated pellets)				APPROVED		
		Excipients	-4rr	q.s		-		
			Approved colours used in empty capsule shell.					
41	Omeprazole &	Each hard gelatin capsule con	tains:					
	Domperidone	Omeprazole	IP	20	mg			
	Capsules	(As enteric coated pellets)						
		Domperidone	IP	10	mg			
		(As enteric coated pellets)				APPROVED ANOTHER APPROVIS HERE		
		Excipients		q.s		-		
		Approved colours used in emp	oty capsul	e shell.				
42	Pantoprazole	Each hard gelatin capsule con	tains:					
	~ ~~~	Dontonrozolo Codium	ID					
	Sodium (EC)	Pantoprazole Sodium	IP					
	& Domperidone	Eq. to Pantoprazole	IP	40	mg			
	& Domperidone (SR)	Eq. to Pantoprazole (As enteric Coated pellets)			mg			
	& Domperidone	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone	IP	40	mg mg	APPROVED		
	& Domperidone (SR)	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release				APPROVED		
	& Domperidone (SR)	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets)		30		APPROVED		
	& Domperidone (SR)	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients	IP	30 q.s		APPROVED		
	& Domperidone (SR) Capsules	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empression	IP	30 q.s		APPROVED		
43	& Domperidone (SR) Capsules	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empty	IP	30 q.s		APPROVED		
43	& Domperidone (SR) Capsules Calcium Dobesilate	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empter Each hard gelatin capsule control Calcium Dobesilate	IP ty capsule	30 q.s shells.		APPROVED		
43	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empirical entering control colour	IP	q.s shells.		APPROVED		
43	& Domperidone (SR) Capsules Calcium Dobesilate	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empter search hard gelatin capsule contact Calcium Dobesilate Monohydrate Excipients	IP ty capsule tains:	30 q.s shells.	mg			
43	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empirical entering control colour	IP ty capsule tains:	30 q.s shells.	mg			
43	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empter search hard gelatin capsule contact Calcium Dobesilate Monohydrate Excipients	IP tains: IP capsule s	30 q.s shells.	mg			
	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate Capsules	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empiral Each hard gelatin capsule con Calcium Dobesilate Monohydrate Excipients Approve colour used in empty	IP tains: IP capsule s	30 q.s shells.	mg			
	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate Capsules Rabeprazole Sodium (EC) & Domperidone	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in emptement of the pellets o	IP ty capsule tains: IP capsule :	q.s shells.	mg			
	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate Capsules Rabeprazole Sodium (EC) & Domperidone (SR)	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empter services and pellets Each hard gelatin capsule control Calcium Dobesilate Monohydrate Excipients Approve colour used in empty Each hard gelatin capsule control Rabeprazole Sodium (As enteric Coated pellets) Domperidone	IP ty capsule tains: IP capsule :	q.s shells.	mg			
	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate Capsules Rabeprazole Sodium (EC) & Domperidone	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empter services and services are services are services and services are services ar	IP tains: IP capsule: tains: IP	30 q.s shells. 500 q.s shell.	mg	APPROVED		
	& Domperidone (SR) Capsules Calcium Dobesilate Monohydrate Capsules Rabeprazole Sodium (EC) & Domperidone (SR)	Eq. to Pantoprazole (As enteric Coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empter services and pellets Each hard gelatin capsule control Calcium Dobesilate Monohydrate Excipients Approve colour used in empty Each hard gelatin capsule control Rabeprazole Sodium (As enteric Coated pellets) Domperidone	IP ty capsule tains: IP capsule : IIP IIP	30 q.s shells. 500 q.s shell. 20 30 q.s	mg	APPROVED		

45	Pantoprazole	Each hard gelatin capsule conta	ains:			
	Sodium (EC)	Pantoprazole Sodium	IP			
	& Itopride HCl	Eq. to Pantoprazole		40	mg	
	(SR)	(As enteric Coated pellets)				
	Capsules	Itopride HCl		150	mg	APPROVED
		(As sustained release pellets)				
		Excipients		q.s		
		Approved colour used in empty	y capsule	shells.		
46	Rabeprazole	Each hard gelatin capsule conta	ains:			
	Sodium(EC) &	Rabeprazole Sodium	IP	20	mg	
	Itopride(SR)	(As enteric Coated pellets)				
	Capsules	Itopride HCl		150	mg	
		(As sustained release pellets)				APPROVED
		Excipients		q.s		
		Approved colours used in emp	ty capsule	e shell.		
47	Esomeprazole	Each hard gelatin capsule conta	ains.			
1,	Magnesium	Esomeprazole Magnesium	illis.			
	(EC) &	Trihydrate	IP			
	Domperidone Domperidone	Eq. to Esomeprazole	11	40	mg	
	(SR)	(As enteric Coated pellets)		10	IIIg	
	Capsules	Domperidone	IP	30	mg	APPROVED
	1.1	(As sustained release pellets)			1118	
		Excipients		q.s		
		Approved colours used in emp	ty capsule			
48	Esomeprazole	Each hard gelatin capsule conta	ains.			
	Magnesium	Esomeprazole Magnesium				
	(EC) &	Trihydrate	IP			
	Itopride HCl	Eq. to Esomeprazole		40	mg	
	(SR)	(As enteric Coated pellets)			8	APPROVED
	Capsules	Itopride HCl		150	mg	
	•	(As sustained release pellets)				
		Excipients		q.s		
		Approved colours used in emp				
49	Itraconazole	Each hard gelatin capsule conta	ains:			
	Capsules	Itraconazole Pellets				
	capsares	Eq. to Itraconazole	BP	100	mg	A DDD OVED
		Excipients		q.s		APPROVED
		Approved colour used in empty	y capsule			
50	Amoxycillin	Each hard gelatin capsule conta				
	Capsules IP	Amoxycillin Trihydrate	IP	I		
	250mg	Eq. to Amoxycillin	"	250	mg	APPROVED
		Excipients Excipients		q.s	5	
		Approve colour used in empty	capsule s			
\Box			- apouto b		<u> </u>	

51	Amoxycillin	Each hard gelatin capsule con	tains:			
	Capsules IP	Amoxycillin Trihydrate	IP			
	500mg	Eq. to Amoxycillin		500	mg	APPROVED
		Excipients		q.s		MINOVE
		Approve colour used in empty		hell.		
52	Cephalexin	Each hard gelatin capsule con				
	Capsules IP	Cephalexin	IP			
	250mg	Eq. to Anhydrous		250	ma	APPROVED
		Cephalexin Excipients		250	mg	
		Approve colour used in empty	z cansule sl	q.s hell		
53	Cephalexin	Each hard gelatin capsule con				
33	Capsules IP	Cephalexin	IP			
	500mg	Eq. to Anhydrous	IL			
	zoomg	Cephalexin		500	mg	APPROVED
		Excipients		q.s	mg	
		Approve colour used in empty	z cansule sl			
			•	iicii		
54	Amoxycillin &	Each hard gelatin capsule con				
	Lactobacillus	Amoxycillin Trihydrate	IP	250		
	Capsules	Eq. to Amoxycillin		250	mg billion	
		Lactobacillus		1.66	spores	APPROVED
		Excipients		q.s		
		Approved colours used in cap	sule shells.			
55	Amoxycillin &	Each hard gelatin capsule con	tains:			
	Lactobacillus	Amoxycillin Trihydrate	IP			
	Capsules	Eq. to Amoxycillin		500	mg	
		Lactobacillus		1.66	billion spores	APPROVED
		Excipients		q.s		
		Approved colours used in cap	sule shells.			
56	Pregabalin &	Each hard gelatin capsule con	tains:			
	Methycobalamin	Pregabalin	IP	75	mg	
	Capsules IP	Methylcobalamin	IP	750	mcg	APPROVED
		Excipients		q.s		
		Approve colour used in empty	capsule sl	hell.		
57	Methycobalamin,	Each hard gelatin capsule con	tains:			
	Alpha Lipoic	Methylcobalamin	IP	1500	mcg	
	Acid,Thiamine	Alpha Lipoic Acid	USP	100	mg	
	Mononitrate,	Thiamine Mononitrate	IP	10	mg	
	Pyridoxine HCl & Folic Acid	Pyridoxine Hydrochloride	IP	3	mg	APPROVED
	Capsules	Folic Acid	IP	1.5	mg	
	Capsares	Excipients		q.s		

58	Cefixime &	Each hard gelatin capsule con	ntains:			
	Erdosteine	Cefixime	IP			
	Capsules	Eq. to Anhydrous Cefixime		200	mg	
		Erdosteine		300	mg	APPROVED
		Excipients		q.s		
		Approve colour used in empt	ty capsule			
59	Ambroxol	Each 5 ml contains:				
	Hydrochloride,	Ambroxol Hydrochloride	IP	30	mg	
	Levosalbutamol &	Levosalbutamol Sulphate	IP			
	Guaiphenesin	Eq. to Levosalbutamol		1	mg	
	Syrup	Guaiphenesin	IP	50	mg	APPROVED
		In a flavoured syrupy base		q.s		
		Approved colour used.	•			
60	Ambroxol HCl,	Each 5 ml contains:				
	Terbutaline	Ambroxol HCl	IP	15	mg	
	Sulphate,	Terbutaline Sulphate	IP	1.25	mg	
	Guaiphenesin &	Guaiphenesin	IP	50	mg	APPROVED
	Menthol Syrup	Menthol	IP	2.5	mg	
		In a flavoured syrupy base		q.s	6	
		Approved colour used.		1		
61	Od .	Each 5 ml contains :				
01	Ofloxacin	Ofloxacin	IP	50	ma	
	Oral Suspension	In a flavoured base	п		mg	APPROVED
	IP 50mg/5ml			q.s		
		Approved colour used.				
62	Ofloxacin	Each 5 ml contains :		100		
	Oral Suspension	Ofloxacin	IP	100	mg	APPROVED
	IP 100mg/5ml	In a flavoured base		q.s		
		Approved colour used.				
63	Levocetirizine	Each 5 ml contains:				
	& Montelukast	Levocetirizine				
	Syrup	Dihydrochloride	IP	2.5	mg	
		Montelukast Sodium	IP			
		Eq. to Montelukast		4	mg	APPROVED
		In a flavoured syrupy base Approved colour used.		q.s.		
61	A 11 1 1	Each 5 ml contains :				
64	Albendazole	Albendazole	IP	200	mg	
	Oral Suspension IP	In a flavoured base	11	q.s	mg	APPROVED
	IΡ	Approved colour used		7''		
65	Liquid Paraffin,	Each 5 ml contains :				
	Milk of	Liquid Paraffin	IP	1.25	ml	
	Magnesia &	Milk of Magnesia	IP	3.75	ml	ADDOCATED
	Sodium	Sodium Picosulphate	BP	3.33	mg	APPROVED
	Picosulphate	In a flavoured base	~-	q.s	۵	
	Suspension	Approved colour used.		7.0		
		Approved colour used.				

66	Ferrous	Each 5 ml contains:					
	Ascorbate &	Ferrous Ascorbate					
	Folic Acid	Eq. to Elemental Iron		30	mg	APPROVED	
	Suspension	Folic Acid	IP	550	mcg		
		In a flavoured base		q.s			
		Approved colour used.	I		-		
67	Chloramphenicol	Each 5 ml contains :					
	Oral Suspension	Chloramphenicol Palmitate	IP				
	IP	Eq. to Chloramphenicol		125	mg		
		In a flavoured base		q.s		APPROVED	
		Approved colour used.	•	•			
68	Cyproheptadine	Each 5 ml contains:					
	Hydrocholride	Cyproheptadine					
	Syrup IP	Hydrocholride (Anhydrous)	IP	2	mg		
		In a flavoured syrupy base		q.s		APPROVED	
		Approved colour used.					
69	Paracetamol	Each uncoated tablet contains	s:				
	Tablets IP 500mg	Paracetamol	IP	500	mg		
		Excipients		q.s		APPROVED	
		-		1			
70	Paracetamol	Each uncoated tablet contains		•		A PROCEED	
	Tablets IP 650mg	Paracetamol	IP	650	mg	APPROVED	
		Excipients		q.s			
71	Drotaverine	Each film coated tablet contain	ins:				
	Hydrochloride	Drotaverine					
	Tablets IP	Hydrochloride	IP	80	mg	APPROVED	
	80mg	Excipients		q.s			
		Approved colour used	1				
72	Lamotrigine	Each uncoated dispersible tab				APPROVED	
	Dispersible	Lamotrigine	IP	100	mg	111110 (22	
	Tablets IP 100mg	Excipients		q.s			
73	Ursodeoxycholic	Each film coated tablet contain		200			
	Acid Tablets IP	Ursodeoxycholic Acid	IΡ	300	mg	APPROVED	
		Excipients		q.s		MINOVED	
7.4	T 1. 1	Approved colour used					
74	Escitalopram	Each film coated tablet contain	IP	<u> </u>			
	Tablets IP	Escitalopram Oxalate Eq. to Escitalopram	IP	10	mg	APPROVED	
	10mg	Excipients Excipients		q.s	nig		
		Approved colour used		7.5			
75	Rosuvastatin	Each film coated tablet contain	ins:				
'	Tablets IP	Rosuvastatin Calcium	IP				
	10mg	Eq. to Rosuvastatin	11	10	mg	ADDOCATO	
	105	Excipients		q.s	8	APPROVED	
		Approved colour used		7.5			
l		F P10 : 00 001 0000					

Tablets IP 20mg	76	Rosuvastatin	Each film coated tablet cont	ains:			
20mg			Rosuvastatin Calcium	IP			APPROVED
Approved colour used		20mg	Eq. to Rosuvastatin		20	mg	APPROVED
Tablets USP		_	Excipients		q.s		
Tablets USP			Approved colour used	•			
Excipients	77	Loratadine	Each uncoated tablet contain				
Excipients		Tablets USP	Loratadine	USP	10	mg	APPROVED
Tablets IP 5mg			Excipients		q.s		APPROVED
Tablets IP 5mg							
Tablets IP 5mg	78	Folic Acid	Each uncoated tablet contain	ns:			
Excipients					5	mg	APPROVED
Page		Tuoious II Sing				1115	
Rablets IP 8mg	70	Databiatina	-	ic.	1		
Excipients	19				Q	ma	APPROVED
So		Tablets II ollig	·	111		nig	
Tablets IP 100mg			•	•	q.s		
Excipients Q.S APPROVED	80				100	***	
Approved colour used Each uncoated tablet contains: Amlodipine Besylate IP Eq. to Amlodipine 2.5 mg Excipients q.s Approved colour used Each uncoated tablet contains: Approved colour used Each uncoated tablet contains: Amlodipine Besylate IP Eq. to Amlodipine Besylate IP Eq. to Amlodipine Besylate IP Eq. to Amlodipine S mg Excipients q.s Approved colour used Each uncoated tablet contains: Approved colour used Each film coated tablet contains: Roxithromycin IP 150 mg Excipients q.s Approved colour used Each film coated tablet contains: Roxithromycin IP 150 mg Excipients q.s Approved colour used Each uncoated chewable tablet contains: Albendazole Tablets IP 400mg Excipients q.s Sondansetron Tablets IP 4mg Each uncoated tablet contains: Ondansetron IT IP Eq. to Ondansetron IP Eq. to Ondansetron IP Each uncoated tablet contains: Ondansetron IP Each uncoated tablet contains: Ondansetron IP Eq. to Ondansetron IP Excipients q.s Approved Each uncoated tablet contains: Ondansetron IP Eq. to Ondansetron IP Excipients q.s Approved Each uncoated tablet contains: Ondansetron IP Eq. to Ondansetron IP Excipients q.s Approved Each uncoated tablet contains: Ondansetron IP Eq. to Ondansetron IP Excipients q.s Approved		Tablets IP 100mg		IP		mg	APPROVED
Amlodipine Each uncoated tablet contains: Amlodipine Besylate IP Eq. to Amlodipine Excipients Q.s Approved colour used			-		q.s		
Amlodipine Besylate Eq. to Amlodipine Excipients Q.S Approved colour used	Q1	Amladinina		ne:			
Eq. to Amlodipine Excipients q.s Approved colour used	01						
Excipients q.s Approved colour used				117	2.5	ma	
Approved colour used Each uncoated tablet contains:						mg	APPROVED
Amlodipine Each uncoated tablet contains: Amlodipine Besylate IP Eq. to Amlodipine S mg Approved			-		4.5		
Tablets IP 5mg	00						
Eq. to Amlodipine 5 mg Excipients q.s Approved colour used	82			•	I		
Excipients Q.S APPROVED		Tablets IP 5mg		IP	5	ma	
Approved colour used 83 Roxithromycin Tablets IP 150mg Each film coated tablet contains: Roxithromycin IP 150 mg Excipients q.s Approved Approved colour used 84 Albendazole Tablets IP 400mg Excipients q.s 85 Ondansetron Tablets IP 4mg 86 Ondansetron Orally Disintegrating Tablets IP 4mg 86 Ondansetron Orally Disintegrating Tablets IP 4mg 86 Each uncoated doesnot on the contains on the contains of the cont						nig	APPROVED
Roxithromycin Each film coated tablet contains: Roxithromycin IP 150 mg Excipients q.s Approved colour used			-		q.s		
Tablets IP 150mg Roxithromycin IP 150 mg Excipients q.s Approved colour used 84 Albendazole Tablets IP 400mg Each uncoated chewable tablet contains: 85 Ondansetron Tablets IP 4mg Each uncoated tablet contains: 86 Ondansetron Orally Disintegrating Tablets IP 4mg 86 Condansetron Orally Disintegrating Tablets IP 4mg 86 Each uncoated orally disintegrating tablet contains: 87 Ondansetron Ondansetron Undansetron Quality Ondansetron Undansetron Un	-02	D 1.1					
Excipients q.s Approved colour used Albendazole Tablets IP 400mg Each uncoated chewable tablet contains: Albendazole IP 400 mg Excipients q.s	83				150	ı	
Approved colour used S4		Tablets IP 150mg		IP		mg	APPROVED
Second			-		q.s		
Tablets IP 400mg Albendazole Excipients Ondansetron Tablets IP 4mg Eq. to Ondansetron Excipients Ondansetron Excipients Ondansetron Excipients Approved colour used Ondansetron Orally Disintegrating Tablets IP 4mg Eq. to Ondansetron Ondansetron Orally Disintegrating Tablets IP 4mg Excipients Each uncoated orally disintegrating tablet contains: Ondansetron Ondansetron Ondansetron Ondansetron Eq. to Ondansetron Ondansetron Ondansetron Eq. to Ondansetron Eq. to Ondansetron	<u></u>						
Robinstron Each uncoated tablet contains: Ondansetron Eq. to Ondansetron Excipients Q.S	84			•			A DDD CYMP
Solution Each uncoated tablet contains: Ondansetron HCl Eq. to Ondansetron 4 mg Excipients Approved colour used		Tablets IP 400mg		IP	400	mg	APPROVED
Tablets IP 4mg Ondansetron HCl Eq. to Ondansetron Excipients Approved colour used Ondansetron Orally Disintegrating Tablets IP 4mg Eq. to Ondansetron Orally Disintegrating Tablets IP 4mg Excipients Ondansetron			Excipients		q.s		
Eq. to Ondansetron 4 mg APPROVED Excipients q.s Approved colour used 86 Ondansetron Orally Disintegrating Tablets IP 4mg Eq. to Ondansetron 4 mg APPROVED APPROVED APPROVED APPROVED	85	Ondansetron	Each uncoated tablet contain	ns:	-	Ī	
Excipients q.s Approved colour used 86 Ondansetron		Tablets IP 4mg	Ondansetron HCl	IP			
Excipients q.s Approved colour used 86 Ondansetron		_	Eq. to Ondansetron		4	mg	APPROVED
86 Ondansetron Corally Disintegrating Tablets IP 4mg Condansetron HCl IP Eq. to Ondansetron HCl Eq. to Ondansetron HCl Eq. to Ondansetron HCl Excipients IP 4mg Excipients IP 4mg Excipients IP 4mg APPROVED					q.s		
Orally Ondansetron HCl IP Disintegrating Eq. to Ondansetron 4 mg Tablets IP 4mg Excipients q.s			Approved colour used				
Orally Ondansetron HCl IP Disintegrating Eq. to Ondansetron 4 mg Tablets IP 4mg Excipients q.s	86	Ondansetron	Each uncoated orally disinte	grating tal	olet contai	ns:	
Disintegrating Eq. to Ondansetron 4 mg Tablets IP 4mg Excipients q.s							
Tablets IP 4mg Excipients q.s					4	mg	APPROVED
					q.s		III I III I III
Tr transfer and the last transfer and the last transfer and the last transfer and t			Approved colour used	•			

87	Cinnarizine	Each uncoated tablet contain	ns:			
	Tablets IP 25mg	Cinnarizine	IP	25	mg	APPROVED
		Excipients		q.s		
88	Cinneflavasin	Each film coated tablet cont	oing:			
00	Ciprofloxacin Tablets IP 250mg	Ciprofloxacin HCl	IP			
	Tablets II 250llig	Eq. to Ciprofloxacin	11	250	mg	
		Excipients			mg	APPROVED
		Approved colour used		q.s		
89	Ciprofloxacin	Each film coated tablet cont	aine.			
0)	Tablets IP 500mg	Ciprofloxacin HCl	IP			
	Tablets II 300ling	Eq. to Ciprofloxacin	11	500	mg	A PPP OVER
		Excipients		q.s	5	APPROVED
		Approved colour used		4.5		
90	Olanzapine	Each film coated tablet cont				
	Tablets IP 5mg	Olanzapine	IP	5	mg	APPROVED
		Excipients		q.s		III NO VEE
		Approved colour used				
91	Olanzapine	Each film coated tablet cont	ains:			
71	Tablets IP 10mg	Olanzapine Olanzapine	IP	10	mg	
	Tuoicus II Tomig	Excipients		q.s	1115	APPROVED
		Approved colour used		4.5		
02	r ' 1' 1	Each film coated tablet cont				
92	Linezolid	Linezolid	IP IP	600	mg	
	Tablets IP 600mg	Excipients	-	q.s	1115	APPROVED DOUBLE
		Approved colour used		1		92 & 248
93		Each film coated tablet cont	tains:			
	Cetirizine Tablets	Cetirizine Hydrochloride	IP	10	mg	
	IP 10mg	Excipients		q.s		APPROVED
		Approved colour used				
94	Glimepiride	Each uncoated tablet contain	ns:			
,	Tablets IP 1mg	Glimepiride	IP	1	mg	
	Twelves in Time	Excipients		q.s		APPROVED
		Encipients		4.5		
95	Glimepiride	Each uncoated tablet contain	ns:	•		
	Tablets IP 2mg	Glimepiride	IP	2	mg	ADDROVED
		Excipients		q.s		APPROVED
		-		-		
96	Racecadotril	Each sachet contains:				
	Sachets IP 10mg	Racecadotril	IP	10	mg	APPROVED
		Excipients		q.s		
		Approved colour used	· I			
97	E14-1	Each film accept tablet	oing.			
9/	Escitalopram	Each film coated tablet cont	IP	1		
	Tablets IP 5mg	Escitalopram Oxalate	IP	5	ma	
		Eq. to Escitalopram Excipients	1		mg	ADDDOVED
		Approved colour used		q.s		APPROVED
		Approved colour used				

98	Cinnarizine	Each uncoated tablet contains	s:			
	Tablets IP	Cinnarizine	IP	75	mg	APPROVED
		Excipients		q.s		
99	Metformin	Each film coated sustained re	lease tabl	et contains	:	
	Hydrochloride	Metformin Hydrochloride	IP	500	mg	
	SR Tablets IP	Excipients		q.s	- 8	APPROVED
	500mg	Approved colour used	1	1		AITROVED
100	Thiocolchicoside	Each film coated tablet conta	ins:			
	& Aceclofenac	Thiocolchicoside	IP	4	mg	
	Tablets	Aceclofenac	IP	100	mg	APPROVED
		Excipients		q.s		
		Approved colour used				
101	Thiocolchicoside	Each film coated tablet conta	ins :			
	& Aceclofenac	Thiocolchicoside	IP	8	mg	ADDROVED
	Tablets	Aceclofenac	IP	100	mg	APPROVED
		Excipients		q.s		
		Approved colour used	1.			
102	Diclofenac					
102	Potassium &	Each film coated tablet conta Diclofenac Potassium	BP	50	mg	
	Paracetamol	Paracetamol	IP	325	mg	A POPO CANADA
	Tablets	Excipients		q.s	1115	APPROVED
		Approved colour used		4.5		
103	D 1 4	Composition :				
103	Beclomethasone	Beclomethasone	1	1		
	Dipropionate Cream	Dipropionate	IP	0.025%	w/w	APPROVED
	Clean	Cream base	11	q.s	VV / VV	
104	T 1' C' HOI			4.5		
104	Terbinafine HCl Cream IP/N	Composition : Terbinafine HCl	IP	1%	w/w	
	Cream IP/N	Preservative:		170	**/ **	APPROVED
		Benzyl Alcohol	IP	1%	w/w	
		Cream base		q.s		
105	Ambroxol	Each film coated tablet conta	ins :			
	Hydrochloride	Levocetirizine				
	SR	Dihydrochloride	IP	5	mg	APPROVED
	& Levocetirizine	Ambroxol Hydrochloride	IP	75	mg	
	Tablets	(In Sustained release form)				
		Excipients		q.s		
		Approved colour used				
106	Metoprolol	Each extended release film co	oated table	et contains	:	
	Succinate	Metoprolol Succinate	USP	11.875	mg	
	Extended Release	Eq. to Metoprolol Tartrate		12.5	mg	APPROVED
	Tablets USP	Excipients		q.s		
		Approved colour used				

107	Metoprolol	Each extended release film co	ated table	:		
	Succinate	Metoprolol Succinate	USP	23.75	mg	
	Extended Release	Eq. to Metoprolol Tartrate		25	mg	APPROVED
	Tablets USP	Excipients		q.s		- '
		Approved colour used				
108	Metoprolol	Each extended release film co	ated table	t contains	:	
	Succinate	Metoprolol Succinate	USP	47.50	mg	APPROVED
	Extended Release	Eq. to Metoprolol Tartrate		50	mg	APPROVED
	Tablets USP	Excipients		q.s		
		Approved colour used				
109	Metoprolol	Each film coated tablet contain	ins:			
107	Succinate (ER) &	Metoprolol Succinate	USP			
	Telmisartan	Eq. to Metoprolol Tartrate	CDI	50	mg	
	Tablets	(As Extended Release)		50	1115	APPROVED
		Telmisartan	IP	40	mg	11110,22
		Excipients		q.s		
		Approved colour used	l		ı	
110	Metoprolol	Each film coated tablet contain	ins :			
110	Succinate (ER) &	Metoprolol Succinate	USP			
	Ramipril	Eq. to Metoprolol Tartrate	0.01	50	mg	
	Tablets	(As Extended Release)			8	
		Ramipril	IP	5	mg	APPROVED
		Excipients		q.s		
		Approved colour used			1	
111	Metformin	Each film coated sustained re	lease table	et contains:	:	
	Hydrochloride	Metformin Hydrochloride	IP	1000	mg	APPROVED
	SR Tablets IP	Excipients		q.s		
	1000mg	Approved colour used		'		
112	Glucosamine	Each film coated tablet contain	ns:			
	Sulphate	Glucosamine Sulphate				
	Potassium	Potassium Chloride	USP	750	mg	
	Chloride, Methyl	(eq. to Glucosamine 444				
	Sulphonyl	mg)	****	2.50		APPROVED
	Methane &	Methyl Sulphonyl Methane	USP	250	mg	
	Diacerein Tablets	Diacerein	IP	50	mg	
		Excipients		q.s		
		Approved colour used				
113	Calcium Citrate	Each film coated tablet contain	ns:	1		
	Malate,	Calcium Citrate Malate		250		
	Cholecalciferol	Eq. to Elemental Calcium		250	mg	
	& Folic Acid	Cholecalciferol	IP	100	IU	APPROVED
	Tablets	Folic Acid	IP	50	mcg	
		Excipients		q.s		
		Approved colour used				

114	Citicoline	Each film coated tablet conta	ins:			
	Tablets IP	Citicoline Sodium	IP			
		Eq. to Citicoline		500	mg	APPROVED
		Excipients		q.s		ATROVED
		Approved colour used				
			•		1	
115	Pregabalin SR &	Each film coated tablet conta	ins: IP	75		
	Methylcobalamin	Pregabalin (In sustained release form)	IP	75	mg	
	Tablets	Methylcobalamin	USP/IP	1500	mcg	
		Excipients	CSI7II	q.s	meg	APPROVED
		Approved colour used	1	1	1	
116	Ivermectin	Each uncoated tablet contains	\.			
110	Tablets	Ivermectin	IP	3	mg	APPROVED
	1 401015	Excipients	- 11	q.s	mg	
				4.5		
117	Ivermectin	Each uncoated tablet contains				APPROVED
	Tablets	Ivermectin	IP	6	mg	
		Excipients		q.s		
118	Ivermectin	Each uncoated tablet contains				A PPD CAVED
	Tablets	Ivermectin	IP	9	mg	APPROVED
		Excipients		q.s		
119	Ivermectin					
	Tablets	Each uncoated tablet contains Ivermectin	IP	12	mg	APPROVED
		Excipients		q.s	- U	
120	Chlorhexidine	Composition:				
120	Mouthwash IP	Chlorhexidine Gluconate				
	Wodinwash n	Solution Solution	IP			A PPD CAVED
		Eq. to Chlorhexidine				APPROVED
		Gluconate		0.2%	w/v	
		In a pleasantly flavoured base)			
		Approved colour used				
121	Losartan	Each film coated tablet conta	ins:			
121	Potassium	Losartan Potassium	IP	50	mg	
	Tablets IP 50mg	Excipients		q.s	8	APPROVED
	8	Approved colour used	1	1	1	
122	Fluconazole	Each uncoated tablet contains	1.			
122	Tablets IP 200mg	Fluconazole	IP	200	mg	
	radicts if 200mg	Excipients	11	q.s	mg	A PPP CATED
		_		4.5		APPROVED
		Approved colour used				
123	Etoricoxib	Each film coated tablet conta	ins:			
	Tablets IP 120mg	Etoricoxib	IP	120	mg	
		Excipients		q.s		APPROVED
		Approved colour used	1			

124	Etoricoxib &	Each film coated tablet contain	ins:					
	Paracetamol	Etoricoxib	IP	60	mg			
	Tablets	Paracetamol	IP	325	mg	APPROVED		
		Excipients		q.s				
		Approved colour used	<u> </u>	<u> </u>				
125	Ibuprofen &	Each film coated tablet contain	ins:					
	Paracetamol	Ibuprofen	IP	400	mg			
	Tablets	Paracetamol	IP	325	mg	APPROVED		
		Excipients		q.s		MINOVED		
		Approved colour used		1 1				
126	Metoprolol	Each film coated tablet conta	ins :					
	Succinate (ER) &	Metoprolol Succinate	USP					
	Amlodipine	Eq. to Metoprolol Tartrate		50	mg			
	Tablets	(As Extended Release)						
		Amlodipine Besylate	IP	_		APPROVED		
		Eq. to Amlodipine		5	mg	AFFROVED		
		Excipients Approved colour used		q.s				
		Approved colour used						
127	Metformin	Each film coated tablet contain	ing					
127	Hydrochloride	Metformin Hydrochloride	IP	1000	ma			
	(SR) &	(In Sustained release form)	II	1000	mg			
	Glimepiride	Glimepiride	IP	1	mg	APPROVED		
	Tablets	Excipients		q.s				
		Approved colour used	l	1				
128	Metformin	Each film coated tablet contain	ins:					
	Hydrochloride	Metformin Hydrochloride	IP	1000	mg			
	(SR) &	(In Sustained release form)						
	Glimepiride Tablets	Glimepiride	IP	2	mg	APPROVED		
	Tablets	Excipients		q.s		AFFROVED		
		Approved colour used		•				
129	Ursodeoxycholic	Each film coated tablet contain	ins:					
	Acid Tablets IP	Ursodeoxycholic Acid	IP	150	mg			
		Excipients		q.s		APPROVED		
		Approved colour used		1		MINOVED		
130	I acoute:-	Each film coated tablet conta	ine:		<u> </u>			
130	Losartan Potassium	Losartan Potassium	IP	50	mg			
	1 Otassium &	Hydrochlorothiazide	IP	12.5	mg	APPROVIEW		
	Hydrochlorothiaz	Excipients		q.s	15	APPROVED		
	ide Tablets IP	Approved colour used		1 4·5				
		Approved colour used						

131	Metoprolol	Each film coated extended re	lease table	et contains	s :	
	Succinate	Metoprolol Succinate	IP	95	mg	
	Extended Release	Eq. to Metoprolol Tartrate		100	mg	APPROVED
	Tablets USP	Excipients		q.s		ATTROVED
		Approved colour used	1	1		
132	Tt'	Each film coated tablet conta	ing			
132	Levetiracetam Tablets USP	Levetiracetam	IP	250	- ma	
	Tablets USF		IF		mg	APPROVED
		Excipients		q.s		
100	· ·	Approved colour used				
133	Lornoxicam Tablets	Each film coated tablet contains:				
	Tablets	Lornoxicam	IP	4	mg	APPROVED
		Excipients		q.s		IN THO VED
		Approved colour used				
134	Metoprolol	Each film coated tablet contains:				
	Succinate (ER) &	Metoprolol Succinate	USP			
	Telmisartan	Eq. to Metoprolol Tartrate		25	mg	
	Tablets	(As Extended Release)		40		APPROVED
		Telmisartan	IP	40	mg	ATTROVED
		Excipients		q.s		
		Approved colour used				
135	Metoprolol	Each film coated tablet contains:				
	Succinate (ER) &	Metoprolol Succinate	USP			
	Amlodipine	Eq. to Metoprolol Tartrate		25	mg	
	Tablets	(As Extended Release)				APPROVED
		Amlodipine Besylate	IP	5	ma	ATTROVED
		Eq. to Amlodipine Excipients		q.s	mg	
		Approved colour used		4.5		
		**				
136	Amoxycillin &	Each film coated tablet contains:				
	Potassium Clavulanate	Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	ma	
	Tablets IP	Potassium Clavulanate Diluted	IP	230	mg	
	Tuolots II	Eq. to Clavulanic Acid	11	125	mg	APPROVED
		Excipients		q.s	8	
		Approved colour used	ı	1 1		
137	Amoxycillin &	Each uncoated dispersible tablet	containe.			
131	Potassium	Amoxycillin Trihydrate	IP			
	Clavulanate	Eq. to Amoxycillin		200	mg	
	Dispersible Tablets	Potassium Clavulanate Diluted	IP			APPROVED
	•	Eq. to Clavulanic Acid		28.5	mg	
		Excipients		q.s		
138	Amoxycillin &	Each film coated tablet conta	ins:		Ī	
	Potassium	Amoxycillin Trihydrate	IP			
	Clavulanate	Eq. to Amoxycillin	*	875	mg	
	Tablets IP	Potassium Clavulanate			- 0	
		Diluted	IP			APPROVED
		Eq. to Clavulanic Acid		125	mg	
		Excipients		q.s		
		Approved colour used		7.0		
		1-pproved colour about				

139	Amoxycillin	Each uncoated dispersible tal				
	Dispersible	Amoxycillin Trihydrate	IP			APPROVED
	Tablets IP 250mg	Eq. to Amoxycillin		250	mg	ATROVED
		Excipients		q.s		
140	Amoxycillin	Each uncoated dispersible tal	blet contain	ns:		
	Dispersible	Amoxycillin Trihydrate	IP			
	Tablets IP 125mg	Eq. to Amoxycillin		125	mg	APPROVED
		Excipients		q.s		
141	Cefixime &	Each film coated tablet conta	ins:			
	Lactic Acid	Cefixime	ΙP			
	Bacillus Tablets	Eq. to Anhydrous				
		Cefixime		100	mg	
		Lactic Acid Bacillus		2.5	billion	
					spores	APPROVED
		Excipients		q.s		
		Approved colour used				
142	Cefixime &	Each film coated tablet conta	ins:			
	Lactic Acid	Cefixime	IP			
	Bacillus Tablets	Eq. to Anhydrous				
		Cefixime		200	mg	
		Lactic Acid Bacillus		2.5	billion	APPROVED
					spores	AFFROVED
		Excipients		q.s		
		Approved colour used				
143	Cefpodoxime &	Each film coated tablet conta	ins:			
	Ofloxacin	Cefpodoxime Proxetil	IP			
	Tablets	Eq. to Cefpodoxime		200	mg	
		Ofloxacin	IP	200	mg	APPROVED
		Excipients		q.s		
		Approved colour used				
144	Fexofenadine	Each film coated tablet conta	ins:			
	Tablets IP 180mg	Fexofenadine	-	100		
		Hydrochloride	IP	180	mg	APPROVED
		Excipients		q.s		
		Approved colour used		•		
145	Artemether &	Each uncoated tablet contain	s:			
	Lumefantrine	Artemether	IP	80	mg	
	Tablets	Lumefantrine		480	mg	APPROVED
		Excipients		q.s		
146	Atorvastatin	Each film coated tablet conta	ins:			
	Tablets IP 10mg	Atorvastatin Calcium	IP			
	3	Eq. to Atorvastatin		10	mg	
		Excipients		q.s		APPROVED
		Approved colour used				

Tablets IP 20mg	147	Atorvastatin	Each film coated tablet cont	ch film coated tablet contains:					
Excipients		Tablets IP 20mg		IP					
Approved colour used 148		<u>_</u>	-		20	mg			
Atorvastatin Tablets IP 40mg Eq. to Atorvastatin Excipients Atorvastatin Eq. to			Excipients		q.s		APPROVED		
Tablets IP 40mg Atorvastatin Calcium Eq. to Atorvastatin Excipients Approved colour used 149		-	Approved colour used	-					
Eq. to Atorvastatin Excipients Approved colour used 149	148	Atorvastatin	Each film coated tablet cont	ains:					
Excipients		Tablets IP 40mg		IP					
Approved colour used 149			Eq. to Atorvastatin		40	mg			
Atorvastatin Tablets IP 80mg Atorvastatin Calcium IP 80 mg Eq. to Atorvastatin Each film coated tablet contains:			Excipients		q.s		APPROVED		
Tablets IP 80mg Atorvastatin Calcium Eq. to Atorvastatin Excipients Approved colour used Each uncoated tablet contains: Rupatadine Fumarate Tablets Fundatine Eq. to Rupatadine Eq. to Rupatadine Eq. to Rupatadine Eq. to Rupatadine Excipients Approved colour used Excipients Approved colour used Each film coated tablet contains: Levocetirizine Tablets IP 5mg Excipients IP Some Excipients Approved Excipients ApproveD APPROVED APPROVED			Approved colour used						
Eq. to Atorvastatin 80 mg Excipients q.s Approved colour used 150 Rupatadine Fumarate Fumarate Tablets Eq. to Rupatadine Eq. to Rupatadine Fundatine Tablets Approved colour used 151 Levocetirizine Tablets IP 5mg Each film coated tablet contains: Levocetirizine Hydrochloride IP 5 mg Excipients Q.s ApproveD	149	Atorvastatin		ontains:					
Excipients		Tablets IP 80mg		IP					
Excipients q.s Approved colour used			Eq. to Atorvastatin			mg	APPROVED		
Rupatadine Fumarate Rupatadine Fumarate Tablets Eq. to Rupatadine 10 mg Excipients Q.s					q.s				
Fumarate Tablets Rupatadine Fumarate Eq. to Rupatadine Excipients Approved colour used Levocetirizine Tablets IP 5mg Levocetirizine Hydrochloride Excipients IP 10 mg APPROVED APPROVED APPROVED APPROVED APPROVED			**						
Tablets	150	-		_					
Excipients q.s APPROVED Excipients q.s				IP					
Approved colour used Levocetirizine Tablets IP 5mg Each film coated tablet contains: Levocetirizine Hydrochloride IP 5 mg Excipients APPROVED		Tablets			10	mg	APPROVED		
151 Levocetirizine Tablets IP 5mg Each film coated tablet contains: Levocetirizine Hydrochloride IP 5 mg Excipients APPROVED			Excipients		q.s				
Tablets IP 5mg Levocetirizine Hydrochloride IP 5 mg Excipients q.s			Approved colour used		1	•			
Hydrochloride IP 5 mg Excipients q.s	151	Levocetirizine	Each film coated tablet co	ontains :					
Excipients q.s APPROVED		Tablets IP 5mg	Levocetirizine						
APPROVED			Hydrochloride	IP	5	mg			
Approved colour used			Excipients		q.s		ADDDOVED		
			Approved colour used	Approved colour used					
152 Desloratadine Each uncoated mouth dissolving tablet contains :	152	Desloratadine	Each uncoated mouth dis	solving tab		ns:			
Mouth Dissolving Desloratadine 5 mg APPROVED		Mouth Dissolving	Desloratadine		5	mg	APPROVED		
Tablets Excipients q.s		Tablets	Excipients		q.s		ALLKOVED		
Approved colour used			Approved colour used	1	1				
153 Methylprednisolone Each uncoated tablet contains:	153	Methylprednisolone	Each uncoated tablet con	tains:					
Tablets IP 4mg Methylprednisolone IP 4 mg APPROVED				IP	4	mg	APPROVED		
Excipients q.s		_	Excipients		q.s				
154 Methylprednisolone Each uncoated tablet contains:	15/	Methylprednisolone	Each uncoated tablet con						
Tablets IP 16mg Methylprednisolone IP 16 mg APPROVED	154			ID	16	ma	APPROVED		
Excipients q.s	134	Tablets IP 16mg	Methylprednisolone	ır	10	mg			

155	Fexofenadine	Each film coated tablet contain				
	Hydrochloride &	Fexofenadine				APPROVED
	Montelukast	Hydrochloride	IP			APPROVED
	Sodium	Eq. to Fexofenadine		120	mg	
	Tablets	Montelukast Sodium	IP			
		Eq. to Montelukast		10	mg	
		Excipients		q.s		
		Approved colour used				
156	Ferrous Ascorbate,	Each film coated tablet contain	ins::			
	Folic Acid & Zinc	Ferrous Ascorbate				
	Sulphate	Eq. to Elemental Iron		100	mg	
	Tablets	Folic Acid	IP	1500	mcg	
		Zinc Sulphate Monohydrate	IP	61.8	mg	APPROVED
		(Eq. to Elemental Zinc 22.5 mg)				
		Excipients		q.s		
		Approved colour used	<u> </u>			
157	Voglibose Mouth	<u>l</u>				
137	Dissolving	Each uncoated mouth dissolv Voglibose	IP	0.2	ma	
	Tablets		11		mg	APPROVED
	Tuolets	Excipients		q.s		
158	Voglibose Mouth					
	Dissolving	Voglibose	IP	0.3	mg	APPROVED
	Tablets	Excipients		q.s		
159	Calcium With	Each film coated tablet contain	ins:			
	Vitamin D	Calcium Carbonate	IP			
	Tablets USP/IP	Eq. to Elemental Calcium		500	mg	APPROVED
		Vitamin D3	IP	250	IU	AITKOVED
		Excipients		q.s		
		Approved colour used	<u>I</u>	L		
160	Terbinafine	Each uncoated tablet contains:				
	Tablets USP/IP	Terbinafine HCl	BP\IP			
		Eq. to Terbinafine		250	mg	APPROVED
		Excipients		q.s		
161	Etizolam	Each film coated tablet contains:				
	Tablets	Etizolam		0.25	mg	
		Excipients		q.s		APPROVED
		Approved colour used				
162	Etizolam	Each film coated tablet contains:				
	Tablets	Etizolam		0.5	mg	
		Excipients		q.s		APPROVED
		Approved colour used	ı	1	1	
		<u> </u>				

163	Metformin	Each film coated tablet contains	:			
	Hydrochloride (SR) &	Metformin Hydrochloride (In Sustained release form)	IP	500	mg	
	Glimepiride	Glimepiride	IP	1	mg	
	Tablets	Excipients		q.s		APPROVED
		Approved colour used				
164	Metformin	Each film coated tablet contains	:			
	Hydrochloride	Metformin Hydrochloride	IP	500	mg	
	(SR) & Glimepiride	(In Sustained release form)	TD.			APPROVED
	Tablets	Glimepiride	IP	2	mg	MIKOVED
	Tuoreus	Excipients		q.s		
		Approved colour used				
165	Amisulpride	Each uncoated tablet contains:				
	Tablets IP 50mg	Amisulpride	IP	50	mg	APPROVED
		Excipients		q.s		
166	Amisulpride	Each uncoated tablet contains:				
	Tablets IP 100mg	Amisulpride	IP	100	mg	APPROVED
		Excipients		q.s		
167	Amisulpride	Each uncoated tablet contains:				APPROVED
	Tablets IP 200mg	Amisulpride	IP	200	mg	
		Excipients		q.s		
168	Amisulpride	Each uncoated tablet contains:				
	Tablets IP 300mg	Amisulpride	IP	300	mg	APPROVED
		Excipients		q.s		
169	Amisulpride	Each uncoated tablet contains:				
	Tablets IP 400mg	Amisulpride	IP	400	mg	APPROVED
		Excipients		q.s		
170	Flupentixol	Each film coated tablet contains	:			
	Tablets	Flupentixol Dihydrochloride	BP			
		Eq. to Flupentixol		0.5	mg	APPROVED
		Excipients		q.s		APPROVED
		Approved colour used				
171	Calcium &	Each uncoated tablet contains:				
	Vitamin D Tablets	Calcium Citrate Malate				
		Eq. to Elemental Calcium	TD.	250	mg	APPROVED
		Vitamin D3	IP	200	IU	ALLKOVED
		Excipients		q.s		
		Approved colour used				

172	Pioglitazone	Each film coated tablet contains	s:			
	Tablets IP 15mg	Pioglitazone Hydrochloride	IP			
		Eq. to Pioglitazone		15	mg	
	(NOTE:- Package insert as per notification of	Excipients		q.s		
	ministry of H & FW No.:-GSR-520(E) Date:-31/07/2013)	Approved colour used				APPROVED
173	Pioglitazone	Each film coated tablet contains	s:			
	Tablets IP 30mg	Pioglitazone Hydrochloride	IP			
	(NOTE:- Package insert	Eq. to Pioglitazone		30	mg	A POPO CAMPA
	as per notification of	Excipients		q.s		APPROVED
	ministry of H & FW	Approved colour used	l	I	<u> </u>	
	No.:-GSR-520(E) Date:-31/07/2013)					
174	Hydroxyzine	Each film coated tablet contains	s:			
1,1	Tablets IP 10mg	Hydroxyzine Hydrochloride	IP	10	mg	
		Excipients		q.s		
		Approved colour used		•		APPROVED
		1 11				
175	Hydroxyzine	Each film coated tablet contains		1		
	Tablets IP 25mg	Hydroxyzine Hydrochloride	IP	25	mg	
		Excipients		q.s		APPROVED
		Approved colour used				MINOVED
176	Chloramphenicol	Each hard gelatin capsule conta	ins:			
	Capsules IP 250mg	Chloramphenicol	IP	250	mg	
		Excipients		q.s		APPROVED
		Approved colours used in empt	y capsule sl	hell.		
177	Chloramphenicol	Each hard gelatin capsule conta	ins:			
	Capsules IP 500mg	Chloramphenicol	IP	500	mg	
		Excipients		q.s		A POPO CATALO
		Approved colours used in empt	y capsule sl	hell.	1	APPROVED
170	Catistain E 11 /	Eash unagated told to out?			<u> </u>	
178	Cetirizine Tablets IP 10mg	Each uncoated tablet contains:	ID	10	400 -	APPROVED
	n ronig	Cetirizine Hydrochloride	IP	10	mg	
		Excipients	<u> </u>	q.s	<u> </u>	
179	Esomeprazole	Each enteric coated tablet contain	ns:			
	Tablets IP 20mg	Esomeprazole Magnesium				
		Trihydrate Eq. to Esomeprazole	IP	20	ma	ADDDOVED
		Eq. to Esomeprazole Excipients			mg	APPROVED
		*	<u> </u>	q.s		
		Approved colour used				
180	Esomeprazole	Each enteric coated tablet contai	ns:			
	Tablets IP 30mg	Esomeprazole Magnesium				
		Trihydrate	IP	20		ADDDOVED
		Eq. to Esomeprazole		30	mg	APPROVED
		Excipients		q.s		
		Approved colour used				

181	Esomeprazole	Each enteric coated tablet contain				
	Tablets IP 40mg	Esomeprazole Magnesium				
		Trihydrate Eq. to Esomeprazole	IP	40	ma	APPROVED
		Excipients		_	mg	ATTROVED
		*		q.s		
		Approved colour used				
182	Ramipril	Each uncoated tablet contains:				
	Tablets IP 1.25mg	Ramipril	IP	1.25	mg	APPROVED
		Excipients		q.s		
183	Ramipril	Each uncoated tablet contains:				
	Tablets IP 2.5mg	Ramipril	IP	2.5	mg	APPROVED
		Excipients		q.s		
184	Ramipril	Each uncoated tablet contains:				
	Tablets IP 5mg	Ramipril	IP	5	mg	APPROVED
		Excipients		q.s		
185	Ramipril	Each uncoated tablet contains:				
	Tablets IP 10mg	Ramipril	IP	10	mg	APPROVED
	_	Excipients		q.s		
186	Ascorbic Acid	Each uncoated tablet contains:				
	Tablets IP 500mg	Ascorbic Acid	IP	500	mg	APPROVED
		Excipients		q.s		
187	Sertraline	Each film coated tablet contains:				
	Tablets IP 25mg	Sertraline Hydrochloride	IP			
		Eq. to Sertraline		25	mg	APPROVED
		Excipients		q.s		MIKOVED
		Approved colour used				
188	Sertraline	Each film coated tablet contains:				
	Tablets IP 50mg	Sertraline Hydrochloride	IP			
		Eq. to Sertraline		50	mg	
		Excipients		q.s		APPROVED
		Approved colour used				
189	Sertraline	Each film coated tablet contains:				
	Tablets IP 100mg	Sertraline Hydrochloride	IP			
		Eq. to Sertraline		100	mg	APPROVED
		Excipients		q.s		AFFKUVED
		Approved colour used				

190	Pregabalin	Each uncoated tablet contains:						
	Sustained Release	Pregabalin	IP	75	mg			
	Tablets	(In sustained release form)				APPROVED		
		Excipients		q.s				
191	Pregabalin	Each uncoated tablet contains:						
	Sustained Release	Pregabalin	IP	150	mg			
	Tablets	(In sustained release form)				APPROVED		
		Excipients		q.s				
192	Etoricoxib	Each film coated tablet contains:						
	Tablets IP 60mg	Etoricoxib	IP	60	mg	APPROVED		
		Excipients		q.s		APPROVED		
		Approved colour used	L					
193	Etoricoxib	Each film coated tablet contains:						
175	Tablets IP 90mg	Etoricoxib	IP	90	mg			
	C	Excipients		q.s				
		Approved colour used		1		APPROVED		
		Approved colour used						
194	Amitriptyline							
	Tablets IP 10mg	Amitriptyline Hydrochloride	IP	10	mg			
		Excipients		q.s		APPROVED		
		Approved colour used				APPROVED		
195	Amitriptyline	Each film coated tablet contains:						
	Tablets IP 25mg	Amitriptyline Hydrochloride	IP	25	mg			
		Excipients		q.s		APPROVED		
		Approved colour used			ı	APPROVED		
196	A	Each film coated tablet contains:						
190	Amitriptyline Tablets IP 50mg	Amitriptyline Hydrochloride	IP	50	ma			
	radicts ir 30mg	Excipients	II		mg			
				q.s		APPROVED		
		Approved colour used						
197	Gabapentin	Each film coated tablet contains:						
	Tablets IP	Gabapentin	IP	300	mg			
		Excipients		q.s		APPROVED		
		Approved colour used	l		1			
198	Aceclofenac SR	Each film coated sustained relea	se tablet cor	ntains:				
	Tablets	Aceclofenac	IP	200	mg	ADDDOVED		
		Excipients		q.s		APPROVED		
		Approved colour used						
199	Duloxetine	Each film coated tablet contains:						
	Hydrochloride	Duloxetine Hydrochloride	USP/IP					
	Tablets	Eq. to Duloxetine		20	mg	APPROVED		
		Excipients		q.s				
		Approved colour used	1	1	1			

200	Duloxetine	Each film coated tablet contains	s:			
	Hydrochloride	Duloxetine Hydrochloride	Usp/IP			
	Tablets	Eq. to Duloxetine		30	mg	
		Excipients		q.s		APPROVED
		Approved colour used	•			
201	Duloxetine	Each film coated tablet contains	s:			
	Hydrochloride	Duloxetine Hydrochloride	USP			
	Tablets	Eq. to Duloxetine		40	mg	
		Excipients		q.s		APPROVED
		Approved colour used			1	
202	Duloxetine	Each film coated tablet contains				
202	Hydrochloride	Duloxetine Hydrochloride	USP			
	Tablets	Eq. to Duloxetine		60	mg	APPROVED
		Excipients		q.s		
		Approved colour used			1	
203	Levocetirizine &	Each uncoated dispersible table	et contains :			
	Montelukast	Levocetirizine Hydrochloride	IP	2.5	mg	
	Dispersible	Montelukast Sodium	IP			
	Tablets	Eq. to Montelukast		4	mg	APPROVED
		Excipients		q.s		
		Approved colour used			1	
204	Desvenlafaxine	Each uncoated extended release				
	Extended Release	Desvenlafaxine Succinate				APPROVED
	Tablets	Eq. to Desvenlafaxine		50	mg	
		Excipients		q.s		
205	Desvenlafaxine	Each uncoated extended release	tablet conta	ins:		
	Extended Release	Desvenlafaxine Succinate		100		APPROVED
	Tablets	Eq. to Desvenlafaxine		100	mg	AFFROVED
		Excipients		q.s		
206	Albendazole &	Each uncoated tablet contains:				
	Ivermectin Tablets	Albendazole	IP	400	mg	A PROP CATE
		Ivermectin	IP	6	mg	APPROVED
		Excipients		q.s		
207	Diclofenac					
	Potassium &	Diclofenac Potassium	BP	50	mg	
	Paracetamol	Paracetamol	IP	325	mg	
	Tablets	Excipients		q.s		APPROVED
		Approved colour used	'			

208	Methylcobalamin,	Each film coated tablet contains	:			
	Alpha Lipoic	Methylcobalamin	USP/IP	1500	mcg	
	Acid, Vitamin D3,	Alpha Lipoic Acid	USP	100	mg	
	Pyridoxine	Vitamin D3	IP	1000	IU	
	Hydrochloride &	Pyridoxine Hydrochloride	IP	3	mg	APPROVED
	Folic Acid Tablets	Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colour used				
209	Clarithromycin	Each 5ml of reconstituted suspe	nsion contai	ns:		
	Oral Suspension	Clarithromycin	IP	125	mg	
		Excipients		q.s		
		In a flavoured base	- I I			APPROVED
		Approved colour used				
210	m.1. 1	T 1			L	
210	Telmisartan Tablets IP 20mg	Each uncoated tablet contains:	l m	20		APPROVED
	Tablets IP 20mg	Telmisartan	IP	20	mg	ATTROVED
		Excipients		q.s		
211	Telmisartan	Each uncoated tablet contains:				
	Tablets IP 40mg	Telmisartan	IP	40	mg	APPROVED
		Excipients		q.s		
212	Telmisartan	Each uncoated tablet contains:	•			
	Tablets IP 80mg	Telmisartan	IP	80	mg	APPROVED
	C	Excipients		q.s	8	
212	T. 1	E-1				
213	Telmisartan & Amlodipine	Each uncoated tablet contains:	III)	40		
	Tablets IP	Telmisartan	IP	40	mg	
	1401013 11	Amlodipine Besylate Eq. to Amlodipine	IP	5	ma	APPROVED
		Excipients	1		mg	
		Approved colour used		q.s		
		**				
214	Clobetasol	Composition:	TD.	0.050/		
	Propionate Cream IP	Clobetasol Propionate	IP	0.05%	w/w	
	п	Chlorocresol (as Preservative)	IP	0.1%	w/w	APPROVED
		In a Cream base				
215	Prochlorperazine	Each uncoated tablet contains:	T	_		A DDD OVED
	Tablets IP 5mg	Prochlorperazine Maleate	IP	5	mg	APPROVED
		Excipients		q.s		
216	Prochlorperazine	Each uncoated tablet contains:				
	Tablets IP 25mg	Prochlorperazine Maleate	IP	25	mg	APPROVED
		Excipients		q.s		
217	Diclofenac Gel BP	Composition:				
	_ ioioionae Goi Bi	Diclofenac Diethylamine	BP	1.16%	w/w	APPROVED
		Eq. to Diclofenac Sodium		1%	w/w	
		Gel Base				
		I .				

218	Clindamycin &	Composition:							
	Adapalene	Clindamycin Phosphate	IP						
	Gel	Eq. to Clindamycin		10	mg	APPROVED			
		Adapalene	BP	1	mg				
		Gel Base							
219	Cilnidipine	Each film coated tablet contains:							
	Tablets	Cilnidipine		10	mg				
		Excipients			ing in	APPROVED			
		-		q.s	1				
		Approved colour used	Approved colour used						
220	Gabapentin &	Each film coated tablet contains							
	Methylcobalamin	Gabapentin	IP	300	mg				
	Tablets	Methylcobalamin	IP	500	mcg	APPROVED			
		Excipients		q.s		APPROVED			
		Approved colour used							
221	Diclofenac	Each enteric coated tablet contain	ns:						
	Potassium &	Diclofenac Potassium	BP	50	mg				
	Serratiopeptidase	Serratiopeptidase (EC)	IP	10	mg				
	Tablets	(As 20,000 units of			8				
		Serratiopeptidase)				APPROVED			
		Excipients		q.s					
		Approved colour used							
222	Cefixime & Lactic	Each uncoated dispersible tablet	contains:						
	Acid Bacillus	Cefixime							
	Dispersible Tablets	Eq. to Anhydrous Cefixime		100	mg				
		Lactic Acid Bacillus		2.5	billion				
					spores				
		Excipients		q.s		APPROVED			
		Approved colour used							
223	Cefixime & Lactic	Each uncoated dispersible tablet	contains:						
	Acid Bacillus	Cefixime	IP						
	Dispersible Tablets	Eq. to Anhydrous Cefixime		200	mg				
		Lactic Acid Bacillus		2.5	billion	APPROVED			
					spores				
		Excipients		q.s					
224	Cefixime &	Each film coated tablet contain	ns:						
	Ofloxacin	Cefixime	IP						
	Tablets	Eq. to Anhydrous Cefixime		100	mg				
		Ofloxacin	IP	100	mg				
		Excipients		q.s	8	APPROVED			
		Approved colour used	1	1 4.5	1				
227	0.11.								
225	Cephalexin	Each uncoated dispersible tab							
	Dispersible	Cephalexin	IP			APPROVED			
	Tablets	Eq. to Anhydrous		250	400 -				
		Cephalexin		250	mg				
		Excipients		q.s					

226	Rabeprazole	Each hard gelatin capsule con	tains:			
220	Sodium (EC) &	Rabeprazole Sodium	IP	20	mg	
	Levosulpiride	(As enteric Coated pellets)	11	20	mg	
	(SR)	Levosulpiride		75	mg	
	Capsules	(As sustained release		, 5	5	APPROVED
	Capsules	pellets)				ATTROVED
		Excipients		q.s		
		Approved colours used in em	pty capsule			
227	Nimesulide &	Each uncoated tablet contains:				
	Paracetamol	Nimesulide	BP	100	mg	
	Tablets	Paracetamol	IP	325	mg	
		Excipients		q.s	5	
	(Not for Children	Excipients		4.5		
	below the age of					APPROVED
	12 years)					
228		Each film coated tablet contai	ne :	ı l		
220	Febuxostat	1	115 .	40	ma	
	Tablets	Febuxostat		40	mg	
		Excipients		q.s		APPROVED
		Approved colour used				
229	Febuxostat	Each film coated tablet contains:				
	Tablets	Febuxostat		80	mg	
		Excipients		q.s		APPROVED
		Approved colour used		_	•	
230	Febuxostat	Each film coated tablet contains:				
	Tablets	Febuxostat		120	mg	
		Excipients		q.s		APPROVED
		Approved colour used			1	
231	Datahistina	Each uncoated tablet contains :				
231	Betahistine Tablets IP 16mg	Betahistine Hydrochloride	IP	16	ma	APPROVED
	rablets if Tollig	·	II		mg	
		Excipients		q.s	<u> </u>	
232	Metaxalone &	Each uncoated tablet contains :				
	Diclofenac	Metaxalone		400	mg	APPROVED
	Potassium	Diclofenac Potassium	BP	50	mg	AFFROVED
	Tablets	Excipients		q.s		
233	Methylcobalamin,	Each hard gelatin capsule contain	ns:			
	Alpha Lipoic Acid,	Methylcobalamin	USP	1500	mcg	
	Pyridoxine	Alpha Lipoic Acid	USP	100	mg	
	Hydrochloride, Folic Acid,	Pyridoxine Hydrochloride	IP	3	mg	
	Benfotiamine,	Folic Acid	IP	1.5	mg	
	Biotin &	Benfotiamine	****	50	mg	APPROVED
	Chromium	Biotin	USP	5	mg	
	Capsules	Chromium Picolinate	USP	200		
	-	Eq. to Chromium		200	mcg	
		Excipients Approved colours used in ampty	aanaula aka	q.s		
		Approved colours used in empty	capsule she	žII.		

234	Pancreatin	Each hard gelatin capsules con						
	capsules	Pancreatin			mg			
		Corresponding to enzymatic		300		APPROVED		
		activity of						
		Lipase 25000 unit						
		Amylase 18000 unit						
		Protease 1000 unit						
		Excipients		q.s				
		Approved colours used						
234	Pancreatin	Each hard gelatin capsules con	ntains:					
	capsules	Pancreatin		I	mg			
	T	Corresponding to enzymatic		150	****5	APPROVED		
		activity of						
		Lipase 10000 unit						
		Amylase 8000 unit						
		Protease 600 unit						
		Excipients		q.s				
		Approved colours used	I					
236	Fluconazole Tablets	Each uncoated tablet contains:						
230	IP 150mg	Fluconazole	IP	150	ma			
	n 150mg		ır		mg	APPROVED		
		Excipients		q.s				
		Approved colours used						
237	Ketoconazole	Each uncoated tablet contains	:					
	Tablets IP 200mg	Ketoconazole	IP	200	mg	APPROVED		
		Excipients		q.s				
238	Lornoxicam	Each film coated tablet contain	18 .					
230	Tablets	Lornoxicam	IP	8	mg			
		Excipients		q.s	8	APPROVED		
		Approved colour used		1				
239	Methylprednisolone	Each film coated tablet contain	ns:	l.				
	Tablets IP	Methylprednisolone	IP	8	mg			
		Excipients		q.s		APPROVED		
		Approved colour used		1				
240	Telmisartan &	Each uncoated tablet contains:						
210	Hydrochlorothiazide	Telmisartan	IP	40	mg			
	Tablets IP	Hydrochlorothiazide	IP	12.5		APPROVED		
		Excipients	11	q.s	mg			
241	D to I	•	1	4.5	1			
241	Escitalopram Tablets IP	Each film coated tablet contain		1				
	20mg	Escitalopram Oxalate Eq. to Escitalopram	IP	20	mg			
	Zomg	Excipients			mg	APPROVED		
		Approved colour used		q.s	<u> </u>			
0:5		^^						
242	Secnidazole Tablets	Each film coated tablet contain		1 1				
	IP	Secnidazole Excipients	IP	1	gm	APPROVED		
		Approved colour used	1	q.s	<u> </u>	III NO (LD		
243	Voglibose &	Each uncoated tablet contains:						
243	Metformin HCl		IP	0.2	ma			
	Tablets	Voglibose			mg	APPROVED		
	1 401000	Metformin Hydrochloride	IP	500	mg			
		Excipients						

244	Voglibose &	Each uncoated tablet contains		A PRO OVER		
	Metformin HCl	Voglibose	IP	0.3	mg	APPROVED
	Tablets	Metformin Hydrochloride	IP	500	mg	
		Excipients		q.s		
245	Amlodipine	Each uncoated tablet contains:		_		
	Besylate &	Amlodipine Besylate	IP	_		
	Atenolol Tablets	Eq.to Amlodipine	ID	5	mg	APPROVED DOUBLE
	Tablets	Atenolol	IP	50	mg	245 & 288
		Excipients Approved colour used		q.s		
246	Losartan Potassium	Each film coated tablet contains	:			
	& Amlodipine	Losartan Potassium	IP	50	mg	
	Besylate	Amlodipine Besylate	IP			
	Tablets IP	Eq. to Amlodipine		5	mg	
		Excipients		q.s		APPROVED
		Approved colour used				
247	Ferrous Ascorbate	Each film coated tablet contains	:	1		
	& Folic Acid Tablets	Ferrous Ascorbate Eq. to Elemental Iron		100	mg	
		Folic Acid	IP	1.5	mg	APPROVED
		Excipients		q.s		
		Approved colour used				
248	Linezolid	Each film coated tablet contains				
	Tablets IP	Linezolid	IP	600	mg	
		Excipients		q.s		APPROVED
		Approved colours used				
249	Deflazacort	Each uncoated tablet contains:				
	Tablets	Deflazacort		1	mg	APPROVED
		Excipients		q.s		
250	Deflazacort	Each uncoated tablet contains :				
	Tablets	Deflazacort		6	mg	APPROVED
		Excipients		q.s		
251	Deflazacort	Each uncoated tablet contains:				!
	Tablets	Deflazacort		30	mg	APPROVED
		Excipients		q.s		
252	Levetiracetam	Each film coated tablet contains	:			
	Tablets USP	Levetiracetam	IP	500	mg	APPROVED
		Excipients		q.s		
253	Deflazacort	Each uncoated tablet contains:	1	1	<u> </u>	
233	Tablets	Deflazacort		18	mg	APPROVED
		Excipients		q.s	IIIg	
				1		
254	Deflazacort	Each 5ml contains:	_	1		
	Oral Suspension	Deflazacort		6	mg	ADDDOVED
		In a flavoured base		q.s		APPROVED
		Approved colour used				

255	Aceclofenac,	Each film coated tablet contain	ıs:			
	Paracetamol &	Aceclofenac	IP	100	mg	
	Serratiopeptidase	Paracetamol	IP	325	mg	
	Tablets	Serratiopeptidase (EC)	IP	10	mg	
		(as 20,000 units of			8	APPROVED
		Serratiopeptidase)				
		Excipients		q.s		
		Approved colour used	L L			
256	Aceclofenac,	Each film coated tablet contains				
200	Paracetamol &	Aceclofenac	IP	100	mg	
	Serratiopeptidase	Paracetamol	IP	325		
	Tablets	Serratiopeptidase (EC)	IP	15	mg	
		(as 30,000 units of	11	13	mg	APPROVED
		Serratiopeptidase)				
		Excipients		q.s		
		Approved colour used		4.5		
255	~ ~.	**				
257	Calcium Citrate Malate,	Each film coated tablet contains Calcium Citrate Malate	:	ı		
	Calcitriol,& zinc	Eq. to Elemental Calcium		250	mg	
	Tablets	Calcitriol	IP	0.25	mcg	
	Tablets	Zinc Sulphate Monohydrate	IP	0.25	mes	APPROVED
		Eq. to Elemental Zinc		7.5	mg	MIROVED
		Excipients		q.s		
		Approved colour used				
258	Doxylamine	Each enteric coated tablet ontain	ns:			
	Succinate,	Doxylamine Succinate	USP	10	mg	
	Pyridoxine HCl &	Pyridoxine HCl	IP	10	mg	
	Folic Acid Tablets	Folic Acid	IP	2.5	mg	
		Excipients		q.s		APPROVED
		Approved colour used	<u> </u>	7.~		
259	C-1-i Citt-	Each uncoated tablet contains:				
239	Calcium Citrate, Magnesium, Zinc	Calcium Citrate	USP	1000	mg	
	& Vitamin D3	Magnesium Hydroxide	IP	1000	mg	
	Tablets	Eq. to Elemental Magnesium	ır	100	mg	
	1401045	Zinc Sulphate Monohydrate	IP	100	mg	APPROVED
		Eq. to Elemental Zinc	11	4	mg	
		Vitamin D3	IP	200	IU	
		Excipients	<u> </u>	q.s	10	
0.00	E 4 1 :	-	l	1 7		
260	Ferrous Ascorbate & Folic Acid	Each film coated tablet contains	s: 			
	Tablets.	Ferrous Ascorbate Eq. to Elemental Iron		100	ma	
	radicus.	Folic Acid	IP	1.5	mg	APPROVED
		Excipients	II		mg	
		Approved colour used		q.s		
261	Calcium Citrate,	Each film coated tablet contains				
	Calcitriol & zinc	Calcium Citrate	USP	1000	mg	
	Tablets	Calcitriol	IP	0.25	mcg	ADDDOVED
		Zinc Sulphate	IP	7.5	mg	APPROVED
		Excipients		q.s		
		Approved colour used				

262	Levofloxacin	Each 5 ml contains:				
	suspension	Levofloxacin Hemihydrate	IP			
	_	Eq. to Levofloxacin		125	mg	APPROVED
		Excipients		q.s		
		Approved colour used			•	
263		Each enteric coated tablet contain	ns:			
	Pantoprazole	Pantoprazole Sodium	IP			
	Sodium &	Eq. to Pantoprazole		40	mg	
	Domperidone	Domperidone	IP	10	mg	
	Tablets	Excipients		q.s		APPROVED
		Approved colour used				
264	Aceclofenac,	Each film coated tablet contains:				
	Paracetamol &	Aceclofenac	IP	100	mg	
	Chlorzoxazone	Paracetamol	IP	325	mg	
	Tablets	Chlorzoxazone	USP	250	mg	APPROVED
		Excipients		q.s		
		Approved colour used		_		
265	Clobetasol	Composition:				
	Propionate,	Clobetasol Propionate	IP	0.05%	w/w	
	Neomycin &	Neomycin Sulphate	IP	0.5%	w/w	
	Miconazole Nitrate	Miconazole Nitrate	IP	2.0%	w/w	APPROVED
	Cream	Chlorocresol	IP	0.1%	w/w	
		(as Preservatives)				
		Cream Base		q.s		
266	Loratadine	Each uncoated dispersible tablet	contains:			
	Dispersible Tablets	Loratadine	USP	10	mg	APPROVED
		Excipients		q.s		
267	Povidon-Iodine	Composition:				
20,	Ointment USP	Povidon-Iodine	IP	5%	w/w	
		(Available Iodine 0.5% w/w)		2,0	,	APPROVED
		Water Soluble Ointment base.		l l		
268	Clindamycin &	Composition:				
	Niacinamide Gel.	Clindamycin		1%	w/w	
		(as Clindamycin Phosphate IP)		170	**/ **	
		Niacinamide	IP	4%	w/w	APPROVED
		Gel base.				
269	Diclofenac					
209	Potassium,	Each film coated tablet contains:				
		Diclofenac Potassium	BP	50	mg	
	Paracetamol &	Paracetamol	IP	325	ma	
	Chlorzoxazone	Chlorzoxazone			mg	
	Tablets		USP	250	mg	APPROVED
		Excipients		q.s		
		Approved colour used				

270	Cyproheptadine HCl, Tricholine	Each 5 ml contains:						
	Citrate & Sorbitol	Cyproheptadine HCl						
	Syrup	(Anhydrous) Tricholine Citrate	IP	2	mg			
		Sorbitol Solution (70%)	IP	0.275	gm			
		Non Crystallising	IF IF	2	gm	APPROVED		
		In a flavoured Syrupy base		q.s				
		Approved colour used	ı					
271	Drotaverine HCl &	Each film coated tablet contains:						
2/1	Mefenamic Acid	Drotaverine HCl	IP	80	mg			
	Tablets	Mefenamic Acid	IP	250	mg	APPROVED		
		Excipients	-	q.s	5			
		Approved colour used		1				
272	A1 - £ 0	Each 5 ml contains:						
212	Aceclofenac & Paracetamol Oral	Aceclofenac	IP	50	mg			
	Suspension	Paracetamol	IP	125		APPROVED		
	_	In a flavoured base	II	-	mg			
		Approved colour used		q.s				
273	Magaldrate &	Each 5 ml contains:	T	1 400				
	Simethicone Oral Suspension USP	Magaldrate (Anhydrous)	IP	480	mg	APPROVED		
	Buspension est	Simethicone	IP	50	mg			
		In a flavoured base		q.s				
		Approved colour used						
274	Dicyclomine HCl	Each uncoated tablet contains:						
	& Paracetamol Tablets	Dicyclomine HCl	IP	20	mg	APPROVED		
	Tablets	Paracetamol	IP	325	mg	MIROVED		
		Excipients		q.s				
275	Terbutaline Sulphate,	Each 5ml contains:						
	_	Terbutaline Sulphate	IP	1.25	mg			
	Bromohexine HCl	Bromohexine Hydrochloride	IP	2	mg	ADDOVED		
	& Guaiphenesin	Guaiphenesin	IP	50	mg	APPROVED		
	Syrup	In a flavoured Syrupy base		q.s				
		Approved colour used		1				
27.6	T 1				<u> </u>			
276	Lopeamide hydrochloride	Each hard gelatin capsule contai Lopeamide hydrochloride	ns: IP	2	mg			
	capsules IP	Excipients Excipients	IP IP	q.s.	mg			
		Approved colour used		7.~.		APPROVED		
277	Povidon-Iodine &	Composition:						
	Metronidazole	Povidon-Iodine	IP	5%	w/w			
	Ointment	(Available Iodine 0.5% w/w)				APPROVED		
		Metronidazole	IP	1%	w/w			
		Water soluble ointment base.						

278	Clobetasol	Composition:				
	Propionate & Zinc	Clobetasol Propionate	IP	0.05%	w/w	
	Sulphate Cream	Zinc Sulphate	IP	2.5%	w/w	APPROVED
		Cream Base	1		•	
279	Piracetam	Each film coated tablet Contain				
	Tablets	Piracetam	IP	800	mg	APPROVED
		Excipients		q.s		AFFROVED
		Approved colour used			•	
280	Piracetam	Each 5ml contains:				
	Syrup	Piracetam	IP	500	mg	
		In a flavoured Syrupy base		q.s		APPROVED
		Approved colour used		1	1	
281	Tricholine Citrate	Each 10 ml Contains:			<u> </u>	
201	Syrup	Tricholine Citrate		0.55	gm	
	, ,	Sorbitol (70%)	IP	q.s	8	
		Non Crystallising		1		APPROVED
		In a flavoured Palatable base				
		Approved colour used				
282	Diclofenac					
	Potassium, Paracetamol &	Diclofenac Potassium	BP	50	mg	
		Paracetamol	IP	325	mg	
	Serratiopeptidase Tablets	Serratiopeptidase (EC)	IP	10	mg	APPROVED
	Tablets	(as 20,000 units of				MIKOVED
		Serratiopeptidase) Excipients		0.0		
		Approved colour used		q.s		
283	M-f A -: 1 0	Each uncoated tablet contains:				
283	Mefenamic Acid & Tranexamic Acid	Mefenamic Meson uncoated tablet contains:	IP	250	ma	
	Tablets	Tranexamic Acid	BP	500	mg	APPROVED
		Excipients	DI	q.s	mg	
284	Torsemide	Each uncoated tablet contains:		4.0		
204	Tablets IP	Torsemide	USP/IP	10	mg	
		Excipients	OSITI		III5	APPROVED
		Approved colour used		q.s	1	
		1 **				
285	Torsemide	Each uncoated tablet contains:	T105 75	•		
	Tablets IP	Torsemide	USP/IP	20	mg	APPROVED
		Excipients		q.s		1110 1110
		Approved colour used				
286	Sodium	Each 5 ml contains:				
	Picosulphate	Sodium Picosulphate	BP	5	mg	
	Syrup	In a Palatable Sorbitol base		q.s		APPROVED
		Approved colour used				

287	Potassium Nitrate	Composition:						
	& Fluoride	Potassium Nitrate	BP	5%	w/w			
	Medicated foaming Dental Gel	Sodium Monofluorophosphate Approved colour used	USP	0.7%	w/w	APPROVED		
288	Amlodipine &	Each uncoated tablet contains:						
	Atenolol Tablets	Amlodipine Besylate	IP					
		Eq. to Amlodipine		5	mg			
		Atenolol	IP	50	mg	APPROVED		
		Excipients		q.s		APPROVED		
		Approved colour used						
289		Each Sachet of 1 g. Contains:						
	Cholecalciferol	Cholecalciferol	IP	60,000	IU	APPROVED		
	Sachet	Excipients		q.s				
290	Cefixime	Each uncoated dispersible table	et contains:					
	Dispersible	Cefixime	IP			APPROVED		
	Tablets IP/NOt	Eq. to Anhydrous Cefixime		100	mg			
		Excipients		q.s				
291	Cefixime	Each uncoated dispersible table	Each uncoated dispersible tablet contains:					
271	Dispersible	Cefixime	IP					
	Tablets IP/NOT	Eq. to Anhydrous Cefixime		200	mg	APPROVED		
		Excipients		q.s				
292	Cefpodoxime	Each uncoated dispersible table	t contains:					
	Dispersible	Cefpodoxime Proxetil	IP					
	Tablets	Eq. to Cefpodoxime		100	mg	APPROVED		
		Excipients		q.s				
293	Cefpodoxime	Each uncoated dispersible table	t contains:					
	Dispersible	Cefpodoxime Proxetil	IP			APPROVED		
	Tablets	Eq. to Cefpodoxime		50	mg			
		Excipients		q.s				
294	Acetaminophen	Each sachet contains:						
	, phenylphrine HCL,	Acetaminophen	USP	650	mg			
	HCL, guaifenesin	phenylphrine HCL	IP	25	mg	APPROVED		
	sachets	guaifenesin	IP	10	mcg			
		Excipients Approved colour used		q.s				
		Approved colour used						
295	Carbonyl Iron,	Each hard gelatin capsule conta	ins:	T				
	Folic Acid & Zinc Capsules	Carbonyl Iron		60	ma			
	Capsules	Eq. to. Elemental Iron Folic Acid	IP	60	mg			
				1.5	mg			
		Zinc Sulphate Monohydrate (Eq. to elemental Zinc 22.5 mg)	IP	61.8	mg	APPROVED		
		Approved Colour used in empty	y Capsule s	hells.				

Dispersible Tablets	296	Cephalexin	Each uncoated dispersible table				
Excipients		Dispersible Tablets		IP			
Approved colour used					125	mg	
Cephalexin Dispersible Tablets Cephalexin Eq. to Anhydrous Cephalexin IP 250 mg APPROVED			Excipients		q.s		APPROVED
Dispersible Tablets			Approved colour used				
Eq. to Anhydrous Cephalexin	297	Cephalexin	Each uncoated dispersible table	t contains:			
Captal C		Dispersible Tablets					
Cephalexin Tablets			Eq. to Anhydrous Cephalexin		250	mg	APPROVED
Cephalexin			Excipients		q.s		
Eq. to Anhydrous Cephalexin	298			:			
Excipients		IP		IP			
Approved colour used					500	mg	APPROVED
Calcium & Vitamin D3			_		q.s		
Vitamin D3 Calcium Carbonate from an organic source (Oyster shell) Eq. to Elemental Calcium 250 mg Vitamin D3 IP 125 IU In a flavoured base Approved colour used			Approved colour used				
Suspension From an organic source (Oyster shell) Eq. to Elemental Calcium 250 mg Vitamin D3 IP 125 IU In a flavoured base Approved colour used Each uncoated tablet contains: Each uncoated tablet contains: Calcitriol Tablets Calcitriol IP 0.25 mcg Approved	299						
Calcium Citrate & USP 1000 mg Calcium Citrate & USP 11.25 ml In a flavoured base q.s. Approved colour used Solic Acid Syrup FOR THERAPEUTIC USE Each 15ml contains : Ferric Ammonium Citrate & IP 160 mg Calcium Citrate & IP 1							
Eq. to Elemental Calcium		Suspension					
Solution Calcium Citrate & Calcium Citrate Each uncoated tablet contains:					250		ADDDOVED
In a flavoured base Q.S. Approved colour used				TD.			AFFROVED
Approved colour used South Calcitriol Tablets Each uncoated tablet contains: Calcitriol Tablets Calcitriol IP 0.25 mcg Excipients Q.s IP 0.25 mcg Excipients Q.s IP 0.25 mcg Excipients Q.s IP II.25 ml In a flavoured base Q.s. Approved colour used IP II.25 ml In a flavoured base Q.s. Approved colour used IP II.25 ml IIP III.25 III IIP III.25 III IIII IIIII IIII IIIII IIII IIII IIII IIII IIII IIII IIII IIII IIIII IIII IIII IIII IIII IIII IIII IIII IIII IIIII IIII IIII IIII IIII IIII IIII IIII IIII IIIII IIII IIII IIII IIII IIII IIII IIII IIII IIIII IIII IIII IIIII IIIIII				IP		IU	
Calcitriol Tablets					q.s		
Calcitriol Tablets							
Calcitriol IP 0.25 mcg APPROVED	300						
Excipients Q.S APPROVED		Calcitriol Tablets	Calcium Citrate	USP	1000	mg	
Solution Excipients Q.S			Calcitriol	IP	0.25	mcg	APPROVED
Paraffin & Milk of Magnesia Laxative Oral Suspension Milk of Magnesia IP 11.25 ml In a flavoured base q.s. Approved colour used			Excipients		q.s		
Paraffin & Milk of Magnesia Laxative Oral Suspension Milk of Magnesia IP 11.25 ml In a flavoured base q.s. Approved colour used							
Magnesia Laxative Oral Suspension Milk of Magnesia IP 11.25 ml In a flavoured base Approved colour used Iron, Cyanocobalamin & Folic Acid Syrup Folic Acid Syrup Folic Acid IP 160 mg (Eq.to Elemental Iron 32mg) Cyanocobalamin IP 7.5 mcg Folic Acid IP 0.5 mg In a flavoured syrupy base Approved colour used Reclomethasone Dipropionate, Clotrimazole & Neomycin Cream Cream Milk of Magnesia IP 11.25 ml IP 160 mg APPROVED APPROVED APPROVED APPROVED APPROVED APPROVED APPROVED APPROVED	301		Each 15ml contains:				
Oral Suspension In a flavoured base Approved colour used In a flavoured base Approved colour used In a flavoured base Approved colour used			Light Liquid Paraffin	IP	3.75	ml	
Approved colour used Solition Cyanocobalamin & Folic Acid Syrup Each 15ml contains: Ferric Ammonium Citrate (Eq. to Elemental Iron 32mg) Cyanocobalamin IP 7.5 mcg Folic Acid IP 0.5 mg In a flavoured syrupy base Approved colour used Approved colour used Approved colour used Composition: Beclomethasone Dipropionate, Clotrimazole & Neomycin Cream Cream Chlorocresol IP 0.5% w/w Approved Approved IP 1% w/w w/w				IP	11.25	ml	A DDD OX/ED
Solic Acid Syrup FOR THERAPEUTIC USE Each 15ml contains : Ferric Ammonium Citrate (Eq.to Elemental Iron 32mg) Cyanocobalamin IP 7.5 mcg Folic Acid IP 0.5 mg In a flavoured syrupy base Approved colour used Approved colour used Composition : Beclomethasone Dipropionate, Clotrimazole & Neomycin Cream Cream Neomycin Chlorocresol IP 0.5% w/w Approved Approved IP Neomycin Chlorocresol IP 0.1% w/w Approved IP Neomycin Chlorocresol IP 0.1% W/w Approved IP IP IP IP IP IP IP I		Oral Suspension	In a flavoured base		q.s.		APPROVED
Cyanocobalamin & Folic Acid Syrup Folic Acid Syrup			Approved colour used				
Folic Acid Syrup Ferric Ammonium Citrate (Eq.to Elemental Iron 32mg) Cyanocobalamin IP 7.5 mcg Folic Acid IP 0.5 mg In a flavoured syrupy base Approved colour used Composition: Beclomethasone Dipropionate, Clotrimazole & Neomycin Cream Cream Remain IP 0.05 mg IP 0.05 mg IP 0.025% w/w Clotrimazole IP 1% w/w Neomycin Sulphate Eq. to Neomycin Chlorocresol IP 0.5% w/w APPROVED	302	Iron,	FOR THERAPEUTIC USE				
Cyanocobalamin IP 7.5 mcg Folic Acid IP 0.5 mg In a flavoured syrupy base Approved colour used			Each 15ml contains:				
Cyanocobalamin IP 7.5 mcg Folic Acid IP 0.5 mg In a flavoured syrupy base q.s. Approved colour used Composition: Beclomethasone Dipropionate, Clotrimazole & Neomycin Cream Cream Cream Cuthorial Iron 32Hg) IP 7.5 mcg IP 0.5 mg In a flavoured syrupy base q.s. Approved colour used Composition: Beclomethasone Dipropionate IP 0.025% w/w Clotrimazole IP 1% w/w Neomycin Sulphate IP Eq. to Neomycin 0.5% w/w Chlorocresol IP 0.1% w/w		Folic Acid Syrup		IP	160	mg	APPROVED
Folic Acid IP 0.5 mg In a flavoured syrupy base q.s. Approved colour used Composition: Beclomethasone Dipropionate, Clotrimazole & Neomycin Cream Cream Reproved Colour used Composition: Beclomethasone Dipropionate IP 0.025% w/w Clotrimazole IP 1% w/w Neomycin Sulphate IP Eq. to Neomycin 0.5% w/w Chlorocresol IP 0.1% w/w				ID	7.5	maa	
In a flavoured syrupy base q.s. Approved colour used 303							
Approved colour used Solution Approved colour used Approved colour used				11		mg	
Beclomethasone Composition : Beclomethasone Dipropionate IP 0.025% w/w				1	4.5.	l	
Dipropionate, Clotrimazole & Neomycin Cream Cream Dipropionate, Clotrimazole & IP 0.025% w/w Neomycin Sulphate IP Eq. to Neomycin Chlorocresol IP 0.5% w/w APPROVED APPROVED	202	Daglamathasana	**				
Clotrimazole & Clotrimazole IP 1% w/w	303			TD	0.025%	337/337	
Neomycin Neomycin Sulphate IP 1% W/W			1 1				
Cream Eq. to Neomycin 0.5% w/w Chlorocresol IP 0.1% w/w					1 70	W/W	
Chlorocresol IP 0.1% w/w		_		IF	0.5%	w/w	APPROVED
				ΙP			
			(as preservative)		2.1/0		
In a cream base							

304	Clobetasol	Composition:						
	Propionate &	Clobetasol Propionate	IP	0.05%	w/w			
	Salicylic Acid	Salicylic Acid	IP	6.0%	w/w	APPROVED		
	Ointment	In an Ointment base		q.s				
305	Vitamin E & Aloe	Composition:						
	Vera Cream	Aloe extract	IP	10%	w/w			
		Vitamin E Acetate	IP	0.5%	w/w	APPROVED		
		Moisturising Cream base		q.s				
306	Omega 3 Fatty	Each hard gelatin capsule conta	Each hard gelatin capsule contains:					
	Acid,	Alpha Lipoic Acid	USP	50	mg			
	Methylcobalamin,	Methylcobalamin	USP	750	mcg			
	Alpha Lipoic Acid, Chromium	Omega 3 Fatty Acid		500	mg			
	Polynicotinate &	Chromium Polynicotinate		200	mcg	APPROVED		
	Selenium Dioxide	Selenium Dioxide				ATROVED		
	Capsules	Mohohydrate	USP	75	mcg			
	1	Excipients		q.s				
		Approved colours used in empt	y capsule s	hell.				
307	Mefenamic Acid &	Each uncoated tablet contains:						
	Dicyclomine HCl	Mefenamic Acid	IP	250	mg			
	Tablets IP	Dicyclomine HCl	IP	10	mg	APPROVED		
		Excipients		q.s				
308	Ciprofloxacin HCl	Each film coated tablet contains						
	& Tinidazole	Ciprofloxacin HCl	IP					
	Tablets	Eq.to Ciprofloxacin		500	mg			
		Tinidazole	IP	600	mg	APPROVED		
		Excipients		q.s				
		Approved colour used						
309	Dried Aluminium	Each 5ml contains:						
	Hydroxide,	Dried Aluminium Hydroxide						
	Magnesium	Gel	IP	250	mg			
	Hydroxide,	Magnesium Hydroxide	IP	250	mg			
	Activated Dimethicone &	Activated Dimethicone	IP	50	mg	APPROVED		
	Sorbitol	Sorbitol Solution 70%	IP	0.65	gm			
	Suspension	(Non-crystallising)						
		In a flavoured base		q.s				
		Approved colour used						
310	Sodium Valproate	Each film coated controlled rele	ease tablet	contains :				
	& Valproic Acid	Sodium Valproate	IP	133	mg			
	Controlled Release	Valproic Acid	IP	58	mg	A PROPOSITION		
	Tablets	(Both together correspond to				APPROVED		
		Sodium Valproate 200 mg)						
		Excipients		q.s				
		Approved colour used						

311	Sodium Valproate	Each film coated controlled rele				
	& Valproic Acid	Sodium Valproate	IP	200	mg	
	Controlled Release	Valproic Acid	IP	87	mg	
	Tablets	(Both together correspond to				
		Sodium Valproate 300 mg)				APPROVED
		Excipients		q.s		MINO (ED
		Approved colour used				
312	Sodium Valproate	Each film coated controlled rele	ease tablet	contains:		
	& Valproic Acid	Sodium Valproate	IP	333	mg	
	Controlled Release	Valproic Acid	IP	145	mg	
	Tablets	(Both together correspond to				APPROVED
		Sodium Valproate 500 mg)				ATTROVED
		Excipients		q.s		
		Approved colour used				
313	Gliclazide &	Each uncoated tablet contains:				
	Metformin HCl	Gliclazide	IP	80	mg	
	Tablets	Metformin HCl	IP	500	mg	APPROVED
		Excipients		q.s		
214	0 10 .			I		
314	Sucralfate Suspension	Each 10 ml contains:	ID	1 1		
	Suspension	Sucralfate Lea flavoured base	IP	1	gm	APPROVED
		In a flavoured base		q.s		
		Approved colour used				
315	Disodium	Each 5ml contains:				
	Hydrogen Citrate	Disodium Hydrogen Citrate	BP	1.25	gm	
	Syrup	In a flavoured syrupy base		q.s.		APPROVED
		Approved colour used	•			
316	Fusidic Acid &	Composition:				
310	Beclomethasone	Fusidic Acid	IP	2%	w/w	
	Dipropionate	Beclomethasone	- 11	270	VV / VV	APPROVED
	Cream	Dipropionate	IP	0.025%	w/w	
		Cream base		q.s		
317	Chlorhexidine	Composition :	1			
311	Gluconate, Sodium	Chlorhexidine Gluconate				
	Fluoride & Zinc	Solution Solution	IP			
	Chloride	Eq.to Chlorhexidine				
	Mouthwash	Gluconate		0.20%	w/v	APPROVED
		Sodium Fluoride	IP	0.05%	w/v	
		Zinc Chloride	IP	0.09%	w/v	
		In a pleasantly flavoured base				
		Approved colour used				
318	L-Ornithine-L-	Each Sachet (5gm) contains:				
	Aspartate	L-Ornithine -L-Aspartate		3	gm	
1	Sachets	Excipients		q.s		APPROVED
		Lacipicitis		4.5		
	2	Approved colour used		4.5		

319	Terbutaline	Each 5ml contains:				
	Sulphate,	Terbutaline Sulphate	IP	1.25	mg	
	Bromhexine	Bromhexine Hydrochloride	IP	4	mg	
	Hydrochloride,	Guaiphenesin	IP	50	mg	
	Guaiphenesin &	Menthol	IP	2.5	mg	APPROVED
	Menthol Syrup	In a flavoured syrupy base		q.s		
		Approved colour used		4.5		
320	Paracetamol,	Each film coated tablet contains		1		
	Diclofenac	Paracetamol	IP	325	mg	
	Potassium & Chlorzoxazone	Diclofenac Potassium	BP	50	mg	
	Tablet	Chlorzoxazone	USP	500	mg	APPROVED
	Tablet	Excipients		q.s		MIROVED
		Approved colour used				
321	Rabeprazole	Each enteric coated tablet conta	ins:			
	Sodium &	Rabeprazole Sodium	IP	20	mg	
	Domperidone	Domperidone	IP	10	mg	APPROVED
	Tablets	Excipients	1	q.s	1115	
		Approved colour used		4.5		
200	GI: 1 : 0	1 **				
322	Clindamycin & Adapalene	Composition:	TD	1		
	Gel	Clindamycin Phosphate	IP	10/	w/w	
	Gei	Eq. to Clindamycin Adapalene	BP	1% 0.1%	W/W	APPROVED
		Gel base	DF		W/W	
		Preservatives :		q.s		
		Methyl Paraben	IP	0.1%	w/w	
		Phenoxyethanol	IP	0.25%	w/w w/w	
323	Beclomethasone	Composition:		I		
323	Dipropionate,	Beclomethasone				
	Miconazole Nitrate	Dipropionate	IP	0.025%	w/w	
	&	Miconazole Nitrate	IP	2.0%	w/w	
	Neomycin Sulphate	Neomycin Sulphate	IP	0.5%	w/w	APPROVED
	Cream	In a cream base	11		VV / VV	
		Preservatives :		q.s.		
		Methyl Paraben	IP	0.2%	w/w	
		Propyl Paraben	IP IP	0.2%	W/W W/W	
224	****			**-/*	,	
324	Ubidecarenone,	Each hard gelatin capsule contain		1.5-		
	L-Arginine,	Ubidecarenone (Coenzyme Q	USP	100	mg	
	Alpha Tocopheryl Acetate &	10)	ID	100		
	Selenium	L-Arginine	IP	100	mg	
	Capsules	Alpha Tocopheryl Acetate	IP	25	IU	APPROVED
	Сирошео	Selenium (As Sodium		100	mcg	
		selenate)		0.0		
		Excipients		q.s		
		Approved colours used in empty	y capsule	snen.		

325	Sucralfate &	Each 10 ml contains:				
	Oxetacaine Suspension	Sucralfate	USP/I P	1000	mg	
	_	Oxetacaine	BP	20	mg	APPROVED
		In a flavoured base		q.s		
		Approved colour used				
326	Diclofenac	Each film coated tablet contains:	:			
	Potassium,	Diclofenac Potassium	BP	50	mg	
	Paracetamol &	Paracetamol	IP	325	mg	APPROVED
	Serratiopeptidase Tablets	Serratiopeptidase (EC)	IP	15	mg	ATTROVED
	Tuolets	(As 30,000 units of Serratiopeptidase)				
		Excipients		q.s		
		Approved colour used		4.5	1	
327	Lyophilized				1	
321	Saccharomyces	Each sachet contains:		1	1	
	Boulardii, Lactic	Lyophilized Saccharomyces Boulardii		2.5	billion	
	Acid Bacillus &	(Corresponding to 125 mg of		2.3	Omon	
	Racecadotril	Yeast)				APPROVED
	Sachets	Lactic Acid Bacillus		100	million	
				1.0	spores	
		Racecadotril	IP	10	mg	
		Excipients		q.s		
328	Lyophilized	Each sachet contains:				
	Saccharomyces Boulardii, Lactic	Lyophilized Saccharomyces		2.5	1 -11-	
	Acid Bacillus &	Boulardii (Corresponding to 125 mg of		2.5	billion	
	Zinc Sachets	Yeast)				
		Lactic Acid Bacillus		150	million	APPROVED
					spores	
		Elemental Zinc (As Zinc Lactate)		7.5	mg	
		Excipients		a s		
		*	<u> </u>	q.s		
329	Cefpodoxime	Each uncoated dispersible tablet		<u> </u>		
	Dispersible Tablets	Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	200	mg	APPROVED
		Excipients		q.s	mg	
200	G	1		_ 1	1 1	
330	Cefixime	Each uncoated dispersible tablet		1		
	Dispersible Tablets	Cefixime Eq. to Anhydrous Cefixime	IP	50	mg	
		Excipients Excipients		qs	mg	APPROVED
		Approved colour used	L	1"		
331	Cefpodoxime	Each uncoated dispersible tablet	contains		1	
331	Dispersible	Cefpodoxime Proxetil	IP		1	
	Tablets.	Eq. to Cefpodoxime	111	50	mg	APPROVED
		Excipients		qs		
		Approved colour used	1			
		•				

332	On don sotron Orol	Each 5ml contains :				
332	Ondansetron Oral Solution IP		1			
	Solution IP	Ondansetron Hydrochloride Eq. to Ondansetron	IP	2	ma	
	•	In a flavoured syrupy base	IF	_	mg	APPROVED
		Approved colour used		q.s.		
333	Magaldrate &	Each 5ml contains:				
	Simethicone Oral	Magaldrate (Anhydrous)	IP	400	mg	
	Suspension USP	Simethicone	IP	20	mg	A PRO CATED
		In a flavoured sugar free base				APPROVED
		Approved colour used				
334	Lactulose	Each 15ml contains:				
334	Solution USP	Lactulose		10		
	Solution OSF	(As Lactulose Concentrate		10	gm	APPROVED
		USP)				
		Palatable base		q.s		
				4.5		
335	Azithromycin Oral	Each 5ml contains:	.	_		
	Suspension IP	Azithromycin Dihydrate	IP			
		Eq. to Anhydrous		200		APPROVED
		Azithromycin		200	mg	APPROVED
		In a flavoured base		q.s		
		Approved colour used				
336	Trioxsalen	Each film coated tablet contain	ıs:			_
	Tablets USP	Trioxsalen	USP	25	mg	
	25 mg	Excipients		q.s	8	APPROVED
		Approved colour used		4.5		
337	Ketoprofen	Each hard gelatin capsule conta				
	Capsules	Ketoprofen		150	mg	APPROVED
		Excipients		q.s		ATTROVED
		Approved colour used in hard	gelatin cap	sule shells.		
338	Roxithromycin	Each film coated tablet contain	s:			
	Tablets IP 300mg	Roxithromycin	IP	300	mg	
	C	Excipients		q.s		APPROVED
		Approved colour used		4.5		
339	Amlodipine	Each uncoated tablet contains:	1			
	Tablets IP 10mg	Amlodipine Besylate	IP	10		
		Eq. to Amlodipine		10	mg	APPROVED
		Excipients		q.s		ATTROVED
		Approved colour used				
340	Iron (III)-	Each 1ml contains:				
- '*	hydroxide	Iron (III)- hydroxide	IP		mg	APPROVED
	polymaltose	polymaltose eq. to elemental			0	
	complex syrup	iron		10		
		In flavoured syrupy base		q.s		
		Approved colour used		+ -		
	_	**	<u> </u>			
341	Iron (III)-	Each chewable tablet contains:				A DDD CYTED
	hydroxide	Iron (III)- hydroxide	IP		mg	APPROVED
	polymaltose	polymaltose eq. to elemental		10		
	complex syrup	iron		10		
		LHyciniente	1	q.s	i l	
		Excipients Approved colour used		4.5		

342	Fexofenadine	Each film coated tablet contains	:			
5.2	Tablets IP 120mg.	Fexofenadine Hydrochloride	IP	120	mg	
	Č	Excipients		q.s		APPROVED
		Approved colour used		4.5		
343	Montelukast	Each film coated tablet contains	•			
343	Tablets IP 10mg	Montelukast Sodium	IP			
	140101011111111111111111111111111111111	Eq. to Montelukast	11	10	mg	
		Excipients		q.s	8	APPROVED
		Approved colour used		1		
344	Montelukast	Each film coated tablet contains				
311	Tablets IP 5mg	Montelukast Sodium	IP			
		Eq. to Montelukast	1	5	mg	
		Excipients		q.s		APPROVED
		Approved colour used	<u> </u>	1 1		
345	Finasteride	Each film coated tablet contains	:			
	Tablets IP	Finasteride	IP	5	mg	
		Excipients		q.s	8	APPROVED
		Approved colour used		1 1	1	
346	Ketoconazole 2%					
340	Cream	Composition : Ketoconazole	IP	2%	/	APPROVED
	Cicam	In a cream base	IP	2%	w/w	111110 (22
		in a cream base				
347	Clarithromycin	Each uncoated tablet contains:				
	Tablets IP 250mg	Clarithromycin	IP	250	mg	
		Excipients		q.s		APPROVED
		Approved colour used				
348	Clarithromycin	Each uncoated tablet contains:				
	Tablets IP 500mg	Clarithromycin	IP	500	mg	APPROVED
		Excipients		q.s		
341	Iron (III)-	Each chewable tablet contains:		•		
341	hydroxide	Iron (III)- hydroxide	IP		mg	APPROVED
	polymaltose	polymaltose eq. to elemental	11		mg	
	complex syrup	iron		10		
		Excipients		q.s		
		Approved colour used		-		
		^^				
250		T 1 (1)			ı	
350	Lornoxicam &	Each film coated tablet contains				
	Paracetamol Tablets	Lornoxicam	IP IP	4	mg	APPROVED
	1 autots	Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				
351	Lornoxicam &	Each film coated tablet contains		-		
	Paracetamol	Lornoxicam	IP	8	mg	A DDD OVED
	Tablets	Paracetamol	IP	325	mg	APPROVED
		Excipients		q.s		
		Approved colour used				

352	Itraconazole	Each hard gelatin capsule conta	ins:			
	capsules	Itraconazole	BP	200	mg	REJECTED
		Excipients		q.s		
		Approved colour used	•			
353	Mupirocin	Composition:				
333	Ointment IP	Mupirocin .	IP	2%	w/w	APPROVED
		Ointment base		q.s	.,,,,	
354	Tacrolimus	Commonition	I			
334	Ointment	Composition : Tacrolimus	IP	0.1%	w/w	APPROVED
	Omunent	Ointment base	11	+	W/W	
				q.s		
355	Montelukast &	Each filmcoated tablet contains		T T		
	Desloratadine Tablet	Montelukast Sodium Eq. to Montelukast	IP	10	ma	APPROVED
	rabiet	Desloratadine		5	mg mg	AFFROVED
		Excipients		q.s	mg	
		Approved colour used	I	T"		
356	Rabeprazole	Each hard gelatin capsule conta	ins:			
	Sodium (EC) & Aceclofenac	Rabeprazole Sodium (As enteric Coated pellets)	IP	20	mg	
	(SR) Capsules	Aceclofenac (As sustained release pellets)	IP	200	mg	APPROVED
		Excipients		q.s		
		Approved colours used in empt	y capsule s	shell.		
357	Paracetamol,	Each film coated tablet contains	s:			
	Phenylephrine	Paracetamol	IP	325	mg	
	HCl, Caffeine &	Phenylephrine HCl	IP	5	mg	
	Diphenhydramine	Caffeine (Anhydrous)	IP	30	mg	APPROVED
	HCl	Diphenhydramine HCl	IP	25	mg	
	Tablets	Excipients		q.s		
		Approved colour used	•	•		
358	Cefadroxil	Each hard gelatin capsule conta	ins:			
	Capsules IP 500mg	Cefadroxil Eq .to Anhydrous Cefadroxil	IP	500	mg	APPROVED
		Excipients		q.s	8	
		Approve colour used in empty	capsule she			
250	Cl 1 1 'C 1	11				
359	Cholecalciferol Chewable Tablets	Each uncoated chewable tablet Cholecalciferol	IP	60,000	пт	
	Chewavie Taviels	Excipients	IP	+ ' +	IU	APPROVED
		*		q.s		
		Approved colour used				
360	Cefuroxime Axetil	Each film coated tablet contains				
	& Potassium	Cefuroxime Axetil	IP			
	Clavulanate	Eq. to Cefuroxime		500	mg	
	Tablets	Potassium Clavulanate	ID.			APPROVED
		Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		+	mg	
		Approved colour used	l	q.s		
		ripproved colour used				

361	Terbutaline	Each 5ml contains:				
	Sulphate, Bromhexine	Terbutaline Sulphate	IP	1.25	mg	
	Hydrochloride &	Bromhexine Hydrochloride	IP	2	mg	
	Guaiphenesin	Guaiphenesin	IP	25	mg	APPROVED
	Syrup	In a flavoured syrupy base		q.s		
		Approved colour used				
362	Clindamycin	Composition:				
	Phosphate Gel USP	Clindamycin Phosphate Eq. to Clindamycin	IP	1%	***/***	
	Ger USP	Gel base		q.s	w/w	APPROVED
		Preservatives :		1		
		Methyl Paraben	IP	0.2%	w/w	
		Propyl Paraben	IP	0.02%	w/w	
363	Ketorolac	Each film coated tablet contains				
	Tromethamine Tablets IP 10mg	Ketorolac Tromethamine	IP	10	mg	APPROVED
	rablets if Tollig	Excipients		q.s		MI NO VED
		Approved colour used				
364	Dextromethorphan	Each 5ml contains:	1		L	
	Hydrobromide, Chlorpheniramine	Dextromethorphan Hydrobromide	IP	5	mg	
	Maleate,	Chlorpheniramine Maleate	IP	2.5	mg	
	Guaiphenesin &	Guaiphenesin	IP	50	mg	
	Ammonium	Ammonium Chloride	IP	60	mg	BANNED Notification No-930
	Chloride	In a flavoured syrupy base		q.s		Notification 110-250
	Syrup.	Approved colour used	I.	1		
365	Amoxycillin &	Each 5ml of the reconstituted so				
	Potassium Clavulanate Oral	Amoxycillin Trihydrate Eq. to Amoxycillin	IP	200	mg	APPROVED
	Suspension IP	Potassium Clavulanate				
		Diluted Eq. to Clavulanic Acid	IP	28.5	ma	
		Excipients Excipients		q.s	mg	
		In a flavoured base		1		
		Approved colour used				
366	Amoxycillin &	Each 5ml of the reconstituted so	uspension	contains:		
	Potassium	Amoxycillin Trihydrate	IP	400		
	Clavulanate Oral Suspension IP	Eq. to Amoxycillin Potassium Clavulanate		400	mg	
	Suspension II	Diluted	IP			APPROVED
		Eq. to Clavulanic Acid		57	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				
367	Ciprofloxacin	Composition:				
	Hydrochloride	Ciprofloxacin Hydrochloride	IP			
	Powder 10% w/w. (for Veterinary use	eq. to Anhydrous Ciprofloxacin		10%	w/w	APPROVED
	Only)	Excipients		q.s	**/ **	
	•	r · ···	l	T.~		

368	Levofloxacin	Each gram Contains:				
	Water Soluble	Levofloxacin Hemihydrate	IP			
	Powder 10% w/w	eq. to Levofloxacin		100	mg	APPROVED
	(for Veterinary use	Excipients		q.s		
	Only)	1		1		
369	Amoxycillin,	Each film coated tablet contains	:			
	Potassium					
	Clavulanate &	Amoxycillin Trihydrate	IP			
	Lactic Acid	Eq. to Amoxycillin		500	mg	
	Bacillus	Potassium Clavulanate	ID			APPROVED
	Tablets	Diluted Eq. to Clavulanic Acid	IP	125	ma	APPROVED
		Lactic Acid Bacillus		60	mg million	
		Lactic Acid Bacinus		00	spores	
		Excipients		q.s		
		Approved colour used				
270	D 1					
370	Paracetamol Paediatric Oral	Each 5ml contains : Paracetamol	IP	125	ma	
	Suspension IP	In a flavoured base	II	q.s	mg	APPROVED
	Suspension ii	Approved colour used		4.5		
		Tippio vod esisur used				
371	Paracetamol	Each 5ml contains:				
	Paediatric Oral	Paracetamol	IP	250	mg	
	Suspension IP	In a flavoured base		q.s		APPROVED
		Approved colour used		•	•	
372	Etoricoxib &	Each film coated tablet contains				
372	Thiocolchicoside	Etoricoxib	IP	60	mg	
	Tablets	Thiocolchicoside	IP	4	mg	APPROVED
		Excipients		q.s	8	
		Approved colour used		1		
373	Etoricoxib &	Each film coated tablet contains		1		
373	Thiocolchicoside	Etoricoxib	IP	60	mg	
	Tablets	Thiocolchicoside	IP	8	mg	APPROVED
		Excipients		q.s	mg	
		Approved colour used		4.5		
274	Q .: : :	**				
374	Cetirizine Hydrochloride,	Each uncoated tablet contains:				
	•	Cetirizine Hydrochloride	IP	5	mg	
	Paracetamol &	Paracetamol	IP	325	mg	
	Phenylephrine	Phenylephrine Hydrochloride	IP	10	mg	APPROVED
	Hydrochloride Tablets.	Excipients		q.s		
	i autets.	Approved colour used	l		' 	
375	Dextromethorphan	Each 5ml contains:				
	Hydrobromide,					
	Phenylephrine	Dextromethorphan				
	Hydrochloride &	Hydrobromide	IP	10	mg	APPROVED
	Chlorpheniramine	Phenylephrine Hydrochloride	IP	5	mg	
	Maleate	Chlorpheniramine Maleate	IP	2	mg	
	Syrup	In a flavoured syrupy base		q.s		
	2 . I .	Approved colour used				
			-			

376	Ibuprofen &	Each film coated tablet contains	3:				
	diphenyhydramine	Ibuprofen	IP	200	mg		
	citrate Tablets	diphenyhydramine citrate		38	mg	APPROVED	
		Excipients		q.s			
		Approved colour used					
		**					
377	Paracetamol,	Each 5ml contains:					
	Phenylephrine	Paracetamol	IP	250	mg		
	Hydrochloride &	Phenylephrine Hydrochloride	IP	5	mg		
	Chlorpheniramine Maleate	Chlorpheniramine Maleate	IP	2	mg	APPROVED	
	Suspension	In a flavoured base		AIIROVED			
	Бизрензіон	Approved colour used					
378	Paracetamol,	Each 5ml contains:					
	Phenylephrine	Paracetamol	IP	125	mg		
	Hydrochloride &	Phenylephrine Hydrochloride	IP	2.5	mg		
	Chlorpheniramine	Chlorpheniramine Maleate	IP	1	mg	APPROVED	
	Maleate Suspension	In a flavoured base	L				
		Approved colour used					
270	۸ '11'						
379	Amoxycillin,	Each hard gelatin capsule conta					
	Dicloxacillin & Lactobacillus	Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	ma		
	Capsules	Dicloxacillin Sodium	IP	230	mg		
	Сирзинез	Eq. to Dicloxacillin	II	250	mg	APPROVED	
		Lactobacillus		2.5	million		
				2.0	spores		
		Excipients		q.s			
		Approved colours used in empt					
380	Methylcobalamin,	Each hard gelatin capsule conta	ine:		Ī		
300	Alpha Lipoic Acid,	Methylcobalamin	USP/I	1500	mcg		
	Pyridoxine	Wedryleobalanini	P	1300	meg		
	Hydrochloride &	Alpha Lipoic Acid	USP	100	mg	APPROVED	
	Folic Acid	Pyridoxine Hydrochloride	IP	3	mg		
	Capsules	Folic Acid	IP	1.5	mg		
		Excipients		q.s	3		
		Approved colours used in empt	<u> </u>				
201	D' ('1 m 11 :						
381	Finasteride Tablets	Each film coated tablet contains					
	IP	Finasteride	IP	1	mg		
		Excipients		q.s		APPROVED	
		Approved colour used					
382	Ofloxacin &	Each 5ml contains:					
	Ornidazole	Ofloxacin	IP	50	mg		
	Suspension	Ornidazole	IP	125	mg	APPROVED	
	•	In a flavoured base		q.s	6	Banned	
		Approved colour used		4.5			
		Approved colour used					

383	Paracetamol,	Each film coated tablet contains:				
	Levocetirizine	Paracetamol	IP	325	mg	
	Hydrochloride,	Levocetirizine Hydrochloride	IP	2.5	mg	BANNED
	Phenylephrine &	Phenylephrine Hydrochloride	IP	10	mg	
	Caffeine Tablets	Caffeine (Anhydrous)	IP	30		
			II		mg	
		Excipients		q.s		
		Approved colour used				
384	Lycopene	FOR PROPHYLACTIC USE				
	With Multi	Each 10 ml contains:		1	_	
	Vitamin &	Lycopene (10%)	USP	1000	mcg	
	Minerals	Vitamin A concentrate	IP	2500	11 1	
	Syrup	(Oily form) (as Palmitate)	IP	2500	IU IU	
		Vitamin E (as Aacetate) Ascorbic Acid (Vitamin C)	IP	10 50	+	
		Thiamine HCl (Vitamin B1)	IP	2	mg mg	APPROVED
		Riboflavin (Vitamin B2)	IP	3	mg	ATTROVED
		PyridoxineHCl (Vitamin B6)	IP	1.5	mg	
		Selenium (as Sodium Selenate)	-	35	mcg	
		Zinc (as Zinc Gluconate USP)		3	mg	
		Manganese		2	mg	
		(as Manganese Gluconate USP)			8	
		Iodine (as Potassium Iodide IP)		100	mcg	
		Copper(as CupricSulphateUSP)		500	mcg	
		In a flavoured syrupy base		q.s		
		Approved colour used				
385	Ofloxacin,	Composition:				
	Ornidazole,	Ofloxacin	IP	0.75%	w/w	
	Terbinafine	Ornidazole	IP	2%	w/w	
	Hydrochloride & Clobetasol	Terbinafine Hydrochloride	BP	1%	w/w	
	Propionate Cream	Clobetasol Propionate	IP	0.05%	w/w	APPROVED
	1 Topionate Cream	Methylparaben	IP	0.2%	w/w	Banned
		(As Preservative)				
		Propylparaben	IP	0.02%	w/w	
		(As Preservative)				
		In a non-greasy base				
386	Azithromycin Oral	Each 5ml contains:				
	Suspension IP	Azithromycin Dihydrate	IP			
	1	Eq.to Anhydrous				
		Azithromycin		100	mg	APPROVED
		In a flavoured base		q.s	-	
		Approved colour used		<u>. ^ </u>		
387	Paracetamol &	Each 5ml contains :			I	
301	Mefenamic Acid	Paracetamol	IP	125	mg	
	Suspension	Mefenamic Acid	IP	50	mg	
	¥ 1	In a flavoured base	11		mg	APPROVED
				q.s		
		Approved colour used				

388	Pantoprazole	Each enteric coated tablet conta				
	Sodium &	Pantoprazole Sodium	IP			
	Domperidone	Eq. to Pantoprazole		20	mg	APPROVED
	Tablets	Domperidone	IP	10	mg	
		Excipients		q.s		
		Approved colour used				
389	S-Amlodipine	Each uncoated tablet contains:				APPROVED
	Tablets IP	S-Amlodipine Besylate	IP			
	2.5 mg	Eq. to S-Amlodipine		2.5	mg	
		Excipients		q.s		
390	S-Amlodipine	Each uncoated tablet contains:		•		
390	Tablets IP 5mg	S-Amlodipine Besylate	IP			
	Tuolous II omg	Eq. to S-Amlodipine	п	5	mg	APPROVED
		Excipients		q.s	5	
391	Paracetamol,	-		7	<u> </u>	
391	Chlorpheniramine	Each 5ml contains:	ID	105		
	Maleate,	Paracetamol	IP	125	mg	
	Phenylepherine,	Chlorpheniramine Maleate	IP	0.5	mg	
	Sodium Citrate &	Phenylepherine HCl	IP	5	mg	APPROVED
	Menthol Syrup	Sodium Citrate	IP	60	mg	
		Menthol	IP	1	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used				
392	Levofloxacin	Each film coated tablet contain	S			
	Tablets IP 750mg	Levofloxacin Hemihydrate	IP			
		Eq. to Levofloxacin		750	mg	APPROVED
		Excipients		q.s		
		Approved colour used				
393	Carbonyl Iron,	FOR THERAPEUTIC USE				
	Folic Acid,	Each hard gelatin Capsule Con-	tains:			
	Vitamin B12,	(In pellets form)				
	Vitamin C & Zinc	Carbonyl Iron				
	Capsules	Eq. to. Elemental Iron		100	mg	A PRO CATED
		Zinc Sulphate Monohydrate	IP	61.8	mg	APPROVED
		Vitamin B12	IP	15	mcg	
		Folic Acid	IP	1.5	mg	
		Vitamin C	IP	75	mg	
		Excipients		q.s		
		Approved Colours used in emp	ty Capsule	shells.		
394	Carbonyl Iron,	Each hard gelatin Capsule Con	tains:			
	Folic Acid, Zinc &		•			
	Vitamin B-12	(In the form of pellets)				
	Capsules	Carbonyl Iron				
	Capsules	Eq. to Elemental Iron		100	mg	
		Folic Acid	IP	1.5	mg	
		Zinc Sulphate Monohydrate	IP	61.8	mg	APPROVED
		(Eq. to Elemental Zinc				
		22.5mg)				
		Vitamin B12	IP	15	mcg	
		Excipients		q.s		
		Approved Colours used in emp	ty Capsule	shells.		
		·			1	

395	Levofloxacin &	Each ml Contains				
373	Bromhexine HCl	Levofloxacin Hemihydrate	IP		1	
	Solution	eq.to Levofloxacin	"	100	mg	
	(for Veterinary use	Bromhexine HCl	IP	7.5		APPROVED
	Only)	In a suiatable vehicle base	11		mg	
	- 3/	ili a sufatable vellicle base		q.s		
396	Rosuvastatin	Each film coated tablet contain	ns:			
	Tablets IP	Rosuvastatin Calcium	IP			
	5mg	Eq. to Rosuvastatin		5	mg	APPROVED
		Excipients		q.s		
		Approved colour used				
397	Benfotiamine,	Each hard gelatin capsule cont	ains:			
	Methylcobalamin,	Benfotiamine		100	mg	
	Alpha Lipoic Acid,	Methylcobalamin	USP	1000	mcg	
	Vitamin B6,	Alpha Lipoic Acid	USP	200	mg	APPROVED
	Inositol	Vitamin B6	IP	3	mg	
	& Folic Acid	Inositol	USP	100	mg	
	Capsules	Folic Acid	IP	1.5	mg	
		Excipients	+	q.s	5	
		Approved colour used in empt	y cansule sh			
			y capsure sir	icii.		
<mark>398</mark>	Telmisartan &	Each uncoated tablet contains:	_	1		
	Hydrochlorothiazi	Telmisartan	IP	80	mg	
	de Tablets	Hydrochlorothiazide	IP	12.5	mg	APPROVED
		Excipients		q.s		
		1				
399	Luliconazole	Composition:	1	T		
	Cream	Luliconazole		1%	w/w	
		Preservative:				APPROVED
		Benzyl Alcohol	IP	1%	W/W	APPROVED
		Cream base		q.s		
400	Ebastine Tablets	Each film coated tablet contain	ns:			
	IP 10mg	Ebastine	IP	10	mg	APPROVED
		Excipients		q.s		
		Approved colour used		4.5		
404					1	
401	Ebastine Tablets	Each film coated tablet contain		20		
	IP 20mg	Ebastine	IP	20	mg	APPROVED
		Excipients		q.s		ALLKOVED
		Approved colour used				
402	Nadifloxacin,	Composition:				
	Miconazole	Nadifloxacin	IP	1%	w/w	
	Nitrate &	Miconazole Nitrate	IP	2%	w/w	
	Mometasone	Mometasone Furoate	IP	0.1%	w/w	APPROVED
1	Furoate Cream	Cream base		q.s	//	
	T di odio Citalii					
402		Each proported tablet and				
403	Olmesartan	Each uncoated tablet contains:		40		
403	Olmesartan Medoxomil &	Olmesartan Medoxomil	IP	40	mg	APPROVED
403	Olmesartan Medoxomil & Hydrochlorothiazi	Olmesartan Medoxomil Hydrochlorothiazide	IP IP	12.5	mg mg	APPROVED
403	Olmesartan Medoxomil &	Olmesartan Medoxomil				APPROVED

404	Serratiopeptidase	Each enteric film coated tablet	contains:					
	Tablets IP	Serratiopeptidase	IP	5	mg			
	5mg	(10,000 Serratiopeptidase						
		units)				APPROVED		
		Excipients		q.s				
		Approved colour used						
405	Serratiopeptidase	Each enteric film coated tablet	contains:					
	Tablets IP	Serratiopeptidase	IP	10	mg			
	10mg	(20,000 Serratiopeptidase				APPROVED		
		units)						
		Excipients		q.s				
		Approved colour used						
406	Lactic Acid	Each uncoated mouth dissolvin	g tablet co	ntains:				
	Bacillus mouth	Lactic Acid Bacillus		60	million			
	dissolving Tablets				spores	APPROVED		
		Excipients		q.s				
407	Lactic Acid	Each 5ml of the reconstituted s	uspension	contains:				
107	Bacillus Oral	Lactic Acid Bacillus	dispension	60	million			
	Suspension	Eactic Field Bachius		00	spores			
	•	Excipients		q.s	1	APPROVED		
		In a flavoured base	II.					
		Approved colour used	pproved colour used					
408	Ofloxacin	Each film coated tablet contains	a•			1		
New	Tablets IP 400mg	Ofloxacin	s. IP	400	ma	APPROVED		
1	ruotets ir 100mg	Excipients	11		mg			
		Approved colour used		q.s	1			
		11pproved corour asca						
409	Flunarizine	Each uncoated tablet contains:						
New	Dihydrochloride	Flunarizine Dihydrochloride	BP			APPROVED		
2	Tablets	Eq. to Flunarizine		5	mg			
		Excipients		q.s				
410	Flunarizine	Each uncoated tablet contains:						
New	Dihydrochloride	Flunarizine Dihydrochloride	BP			APPROVED		
3	Tablets	Eq. to Flunarizine		10	mg			
		Excipients		q.s				
411	Etodolac	Each film ageted tablet as at in			1	1		
New	Tablets IP 400mg	Each film coated tablet contains Etodolac	S: IP	400	ma	APPROVED		
4	1 autois 11° 400111g	Excipients Excipients	IP	400	mg	MINOTED		
		Approved colour used	<u> </u>	q.s	1			
		11pproved colour used						
412	Isotretinoin	Each hard gelatin capsule conta						
New	Capsules	Isotretinoin	IP	20	mg	APPROVED		
5		Excipients	1	q.s				
412	1	Approved colours used in empt	* *	snells.				
413 Now	Atorvastatin &	Each film coated tablet contains	,	1		APPROVED		
New 6	Fenofibrate Tablets	Atorvastatin Calcium	IP	10	ma	AFFRUVED		
0	1 autets	Eq. to Atorvastatin Fenofibrate	IP	160	mg mg			
		Excipients	IP	_	mg			
		Approved colour used	I	q.s				
		ripproved colour used						

New Capsules IP 100mg Excipients IP 100 mg Recipients	414	Ascorbic Acid	Each uncoated tablet contains:				
APPROVED APPROVED APPROVED	New	Tablets IP 100mg	Ascorbic Acid	IP	100	mg	APPROVED
New Capsules IP 75mg	7		Excipients		q.s		
New Capsules IP 75mg	415	Pregabalin	Each hard gelatin capsule conta	ins:			
Excipients					75	mg	APPROVED
Approved colours used in empty capsule shells.	8					8	
New Section Pergabalin IP 150 mg Excipients Approved colours used in empty capsule hells.				y capsule sl			
New Section Pergabalin IP 150 mg Excipients Approved colours used in empty capsule hells.	416	Dragabalin	Each hard colotin consula conto	ina			
Separation Provided Provide		•			150	ma	APPROVED
Approved colours used in empty capsule shells.		Capsules II 130IIIg		II		mg	
17				v capsule sl			
New Release Tablets IP 250mg Excipients Q.S.							
10		-			ontains:		A DDD OVED
Tablets IP 250mg				IP	250		APPROVED
Approved colour used Approved Composition:	10					mg	
Mometasone Furoate Lotion Furoate		Tablets IP 250mg	_		q.s		
New 11 Paracetamol, Phenylephrine Each film coated tablet contains: Phenylephrine HCl, Diphenhydramin e HCl & Caffeine Tablets Caffeine (Anhydrous) IP 30 mg Excipients Approved colour used P 100 mg Excipients Approved value Aceclofenac IP 100 mg Excipients Approved colour used P 4 mg Approved value Each film coated tablet contains: Approved value Approved value P 4 mg Approved value Each film coated tablet contains: Approved value Approved value P 4 mg Approved value Each film coated tablet contains: Approved value Each film coated tablet contains: Approved value Approved value P 4 mg Approved value Each film coated tablet contains: Approved value Each film value Each fi	110	Momotocomo	**				
Montestand Fundace Tablets Composition: Clarithromycin Dispersible Tablets Calamine & Light Lotion & Calamine & Light Lotion & Calamine & Light Lotion & Calamine & Light			-	ID	0.10/	/	APPROVED
Approved Approved		Turoate Lotion		IP		W/V	11110,22
New Phenylephrine HCl, Diphenhydramin e HCl & Caffeine Tablets Caffeine (Anhydrous) IP 25 mg Caffeine (Anhydrous) IP 30 mg Excipients Approved colour used Aceclofenac Tablets Aceclofenac Tablets Tablets Tablets Each film coated tablet contains: Approved colour used Aceclofenac Tablets Tablets Each film coated tablet contains: Approved colour used Approv		D . 1			q.s		
Piletylephine Parameteristic Param	-	· · · · · · · · · · · · · · · · · · ·			500	***	APPROVED
Diphenhydramine HCl IP 25 mg Caffeine Tablets Approved colour used 420 New 14 Tablets Accolofenac Drotaverine HCl Excipients Approved colour used 421 Thiocolchicoside Excipients Approved colour used 422 Clarithromycin Dispersible Tablets Approved colour used 422 Clarithromycin Dispersible Tablets 423 Calamine & Light Liquid Paraffin Lotion 424 Ketoconazole Lotion Accomposition: Camposition: Caffeine (Anhydrous) IP 30 mg Excipients Approved colour used APPROVED							IN THO VED
Caffeine (Anhydrous) IP 30 mg	12					_	
Caffeine Tablets							
Approved colour used Approved colour used Approved colour used Approved colour used Aceclofenac Tablets Aceclofenac IP 100 mg Excipients Approved colour used						6	
Aceclofenac Tablets			Approved colour used				
Aceclofenac Tablets	420	Drotaverine HCl &	Each film coated tablet contains	3:			
Accolofenac IP 100 mg Excipients q.s Approved colour used 421 Thiocolchicoside Tablets Tablets Thiocolchicoside Excipients q.s Approved colour used 422 Clarithromycin Dispersible Tablets Clarithromycin IP 125 mg Excipients q.s APPROVED 423 Calamine & Light Liquid Paraffin Lotion Calamine IP 8.0% w/v Light Liquid Paraffin IP 10.0% w/v In a lotion base q.s APPROVED 424 Ketoconazole Composition: Ketoconazole IP 2.0% w/v APPROVED					80	mg	APPROVED
Excipients q.s Approved colour used	14	Tablets	Aceclofenac	IP	100	mg	
Approved colour used Approved colour used					g.s		
Thiocolchicoside New Tablets Each film coated tablet contains: Thiocolchicoside IP 4 mg Excipients Approved colour used			-		1		
New 15			**				
Excipients Q.S Approved colour used					1		APPROVED
Excipients q.s Approved colour used		Tablets	Thiocolchicoside	IP	4	mg	APPROVED
APPROVED Clarithromycin Each uncoated dispersible tablet contains: Clarithromycin IP 125 mg Excipients Clarithromycin IP 125 mg Excipients Clarithromycin Excipients Excipients E	13		Excipients		q.s		
Dispersible Tablets Clarithromycin IP 125 mg Excipients Calamine & Light Liquid Paraffin Lotion Calamine IP 8.0% W/V Light Liquid Paraffin IP 10.0% Myv In a lotion base Composition: 424 Ketoconazole Lotion Ketoconazole IP 2.0% W/V APPROVED APPROVED			Approved colour used				
Dispersible Tablets Clarithromycin IP 125 mg Excipients 423 Calamine & Light Liquid Paraffin Lotion Calamine IP 8.0% W/V Light Liquid Paraffin IP 10.0% Myv In a lotion base 424 Ketoconazole Lotion Ketoconazole IP 2.0% Myv APPROVED APPROVED	422	Clarithromycin	Fach uncoated dispersible table	t contains:			APPROVED
Excipients q.s Calamine & Light Liquid Paraffin Lotion Calamine IP 8.0% w/v Light Liquid Paraffin IP 10.0% w/v In a lotion base q.s	722		1		125	mg	
Calamine & Light Composition: Calamine IP 8.0% w/v Light Liquid Paraffin IP 10.0% w/v In a lotion base q.s		•	•				
Liquid Paraffin Lotion Calamine IP 8.0% w/v Light Liquid Paraffin IP 10.0% w/v In a lotion base 424 Ketoconazole Lotion Ketoconazole IP 2.0% w/v APPROVED APPROVED			Excipients		q.s		
Lotion Light Liquid Paraffin IP 10.0% w/v In a lotion base q.s 424 Ketoconazole Lotion Ketoconazole IP 2.0% w/v APPROVED	423		Composition:				
APPROVED APPROVED							APPROVED
424 Ketoconazole Composition: Lotion Ketoconazole IP 2.0% w/v APPROVED		Lotion		IP		w/v	
Lotion Ketoconazole IP 2.0% w/v APPROVED			In a lotion base		q.s		
Lotion Ketoconazole IP 2.0% w/v APPROVED	424	Ketoconazole	Composition:				
In a lotion base q.s		Lotion	_	IP	2.0%	w/v	APPROVED
			In a lotion base		q.s		

425	Ketoconazole	Composition:						
	Solution	Ketoconazole	IP		2.0%)	w/v	APPROVED
		Approved colour used						
426	Povidone Iodine	Composition:						
120	Solution IP	Povidone Iodine	IP		1.0%		w/v	APPROVED
	Solution II	(0.1% w/v available Iodine)	11		1.070	,	W/V	
		Aqueous Base			q.s			-
125				-				A PRO OVER
427	Povidone Iodine	Composition:						APPROVED
	Solution IP	Povidone Iodine (0.2% w/v available Iodine)	IP		2.0%)	w/v	
		Aqueous Base			a c			-
		Aqueous Base			q.s			
428	Gliclazide Tablets II	Each uncoated tablet contains	s:					
	80 mg	Gliclazide	IP		80		mg	Approved
		Excipients			q.s			
429	Artemether &	Each uncoated tablet contains					1	
429	Lumefantrine Table		s. IP		40		ma	Approved
	Eumeranume Table		IP				mg	Approved
		Lumefantrine			240	,	mg	
		Excipients			q.s			
430	Artemether &	Each uncoated dispersible tal	olet co	ntains	:			
	Lumefantrine	Artemether	IP		40)	mg	Approved
	Dispersible Tablet	Lumefantrine			240		mg	
		Excipients			q.s			
431	Methylprednisolone	Each uncoated tablet contains	3.					
431	Tablets IP	Methylprednisolone	IP		8		ma	Approved
	Tuolets II	Excipients	п		q.s		mg	Прргочец
		Excipients			4.5			
432	Pantoprazole	Each hard gelatin capsule con	tains					
	Sodium(EC) &							
	Levosulpiride(SR)	Pantoprazole Sodium		IP				Approved
	Capsule	Eq. to Pantoprazole			40		mg	
		(As enteric Coated pellets)			75		ma	
		Levosulpiride (As sustained release pellets)			13		mg	
		Excipients			q.s			
		Approved colours used in emp	oty cap	sule s				
	D 0.11:	**						
122	Doxofylline Tablets	Each uncoated tablet contains:		40	n I		ma	Approved
433	IP 400 mg	Doxofylline Excipients	IP	40	IU]	mg	
<u> </u>	<u> </u>	Excipients	<u> </u>	q.s				
434	Streptococcus Faecali		ntains					Approved
	Clostridium	Streptococcus Faecalis			30		nillion	
	butyricum, Bacillus				2		nillion	
	mesentericus &	Bacillus Mesentericus			1	_	million	
	Lactic Acidbacillus	Editio Field Editing			50	r	million	
	(lactobacillus Sporogenes)	(lactobacillus Sporogenes)				+		
	C apsule	Excipients Approved colours used in	amet-	0052	q.s			
	C apsuic	Approved colours used in	empty	capsi	uie sne	ens.		

435	Olmesartan	Each film coated tablet conta	ine.					
433	Medoxomil Tablet IP	Olmesartan Medoxomil	IP		20		na	Approved
	20 mg		II			1	ng	прргочец
	20 mg	Excipients Approved colour used.			q.s			
		Approved colour used.						
436	Olmesartan	Each film coated tablet conta	ins:					
	Medoxomil Tablet IP	Olmesartan Medoxomil	IP		40	r	ng	Approved
	40 mg	Excipients			q.s			
		Approved colour used.			7.~			
437	Diacerein &	Each film coated tablet conta						
	Aceclofenac Tablet	Aceclofenac	IF		100	m	g	Approved
		Diacerein	IF	•	50	m	g	
		Excipients			q.s			
		Approved colour used.						
438	Sitagliptin Phosphate	Each film coated tablet conta	ains:					
	& Metformin	Sitagliptin Phosphate						
	Hydrochloride Tablets	Monohydrate	I	P				Approved
		Eq. to Sitagliptin			50	1	mg	
		Metformin Hydrochloride	I	P	500	1	mg	
		Excipients			q.s			
		Approved colour used.						
439	Diacerein Capsule IP	Each hard gelatin capsule con	itains					
	50 mg	Diacerein		IP	50		mg	Approved
	_	Excipients			q.s			
	-	Approved colours used in em	pty ca	psule		I		
440	Cetrimide Cream	Composition:						Approved
	0.5 % w/w	Cetrimide		ΙP	0.5	%	w/w	1
		Cream Base			q			
441					1 1			
441	Doxycycline Hydrochlorido	Each hard gelatin capsule con	itains	- TD				Annuarad
	Hydrochloride Capsule IP 100 mg	Doxycycline Hydrochloride		IP	100	. .		Approved
	Capsule IF 100 IIIg	Eq. to Doxycycline Excipients			100	1	ng	
				1.	q.s			
<u> </u>		Approved colours used in em	ipty ca	psuie	snens.			<u> </u>
442	Paroxetine	Each film coated sustained re	elease	tablet	t contai	ns:		
	Sustained Release	Paroxetine Hydrochloride					1	
	Tablets IP 25 mg	Hemihydrate		II	2		1	Approved
		Eq. to Paroxetine				25	mg	<u> </u>
		Excipients				q.s		
		Approved colour used.						
443	Paroxetine	Each film coated sustained re	elease	tablet	t contai	ns:		
	Sustained Release	Paroxetine Hydrochloride						Approved
	Tablets IP 12.5 mg	Hemihydrate		I	P			
		Eq. to Paroxetine				12.5	m	g
		Excipients				q.s		
		Approved colour used.						

444	Paroxetine Sustained	Each film coated sustained re				
	Release Tablets IP	Paroxetine Hydrochloride				
	37.5 mg	Hemihydrate	1	P		Approved
		Eq. to Paroxetine		37.5	5 mg	
		Excipients		q.	S	
		Approved colour used.	•		•	
445	Tranexamic Acid	Each film coated tablet conta	ine.			
113	Tablet IP 250 mg	Tranexamic Acid	IP	250	mg	Approved
		Excipients		q.s	1115	11
		Approved colour used.		4.5		
446	Tranexamic Acid	Each film coated tablet conta				
110	Tablet IP 500 mg	Tranexamic Acid	IP	500	mg	Approved
	Tuester in the ing	Excipients Excipients			nig	rippi oved
		Approved colour used.		q.s		
447	Atorvastatin	Each film coated tablet conta	_		_	
	Tablets IP	Atorvastatin Calcium	IP	_		Approved
		Eq. to Atorvastatin		5	mg	
		Excipients		q.s		
		Approved colour used.				
448	Pregabalin,	Each hard gelatin capsule con-	tains			
	Methylcobalamin,	Pregabalin	IP	75	Mg	
	Alpha Lipoic Acid	Methylcobalamin	IP	750	Mcg	
	,Pyridoxine HCL,	Alpha Lipoic Acid	USP	100	Mg	Approved
	Folic Acid Capsule	Pyridoxine Hydrochloride	IP	3	Mg	
		Folic Acid	IP	1.5	Mg	
		Excipients		q.s		
		Approved colours used in emp	pty capsu	e shells.		
449	Pregabalin,	Each hard gelatin capsule con-				
	Methylcobalamin,	Pregabalin	IP	150	mg	
	Alpha Lipoic Acid	Methylcobalamin	USP	750	mcg	_
	,Pyridoxine HCL,	Alpha Lipoic Acid	USP		mg	Approved
	Folic Acid Capsule	Pyridoxine Hydrochloride	IP	3	mg	
		Folic Acid	IP	1.5	mg	
		Excipients	****	q.s		
		Approved colours used in emp		e shells.		
450	Sodium Bicarbonate	Each film coated tablet conta		700		
	Tablet USP 500 mg	Sodium Bicarbonate	IP	500	mg	Rejected
		Excipients Approved colour used.		q.s		
451	Veries 1 TD 11	**	•			
451	Voriconazole Tablet IP 200 mg	Each film coated tablet conta		200		APPROVED
	ir 200 ilig	Voriconazole	IP	200	mg	AFFRUVED
		Excipients		q.s		
		Approved colour used.				
452	Voriconazole Tablet	Each film coated tablet conta	ins:			
	IP 50 mg	Voriconazole	IP	50	mg	APPROVED
		Excipients		q.s		
		Approved colour used.				

453	Hydroxyzine Oral	Each 5ml contains:				
433	Solution IP		IP	10	ma	APPROVED
	Solution ii	Hydroxyzine Hydrochloride In a flavoured syrupy base	IP		mg	MIROVED
				q.s		
		Approved colour used.				
454	Moxifloxacin	Each film coated tablet contains:	:			
	Hydrochloride Tablet	Moxifloxacin Hydrochloride	IP			
	400 mg	Eq. to Moxifloxacin		400	mg	APPROVED
	-	Excipients		q.s		
		Approved colour used.	1			
455						
455	Flavoxate	Each film coated tablet contains:		200		APPROVED
	Hydrochloride Tablets I.P	Flavoxate Hydrochloride Excipients	IP	200	mg	APPROVED
	Tablets 1.P	Approved colour used.		q.s		
		Approved colour used.				
456	Eplerenone	Each film coated tablet contains:				
	Tablets 25 mg	Eplerenone	IP	25	mg	APPROVED
		Excipients		q.s		
		Approved colour used.				
457	Eplerenone	Each film coated tablet contains:	:			
	Tablets 50 mg	Eplerenone	IP	50	mg	APPROVED
		Excipients		q.s		
		Approved colour used.				
450	(C) A 1 1' ' 0	Each Uncoated tablet contains:				
458	(S)- Amlodipine & Atenolol		ID			APPROVED
	Atenoioi Tablets	(S)-Amlodipine Besylate Eq. to (S)-Amlodipine	IP	2.5	mg	APPROVED
	Tablets	Atenolol	IP	25	mg	
		Excipients		q.s	mg .	
		Approved colour used.	1	1		
459	Cyanocobalamin,Chro	Each film coated tablet contains:				
	mium Picolinate, Folic	Cyanocobalamin	IP	15	mcg	
	Acid, Nicotinamide,	Chromium Picolinate	USP	250	mcg	
	Pyridoxine	Folic Acid	IP	1500	mcg	APPROVED
	HCL, Selenious Acid,	Nicotinamide	IP	100	mg	
	Zinc Sulphate	Pyridoxine Hydrochloride	IP	3	mg	
	Monohydrate Tablet	Selenious Acid	USP			
		Eq. to Elemental Selenium		100	mcg	
		Zinc Sulphate Monohydrate	IP	61.8	mg	
		(Eq. to Elemental Zinc 22.5mg)				
		Excipients		q.s		
		Approved colour used.				
450	Sildenafil Citrate	Each film coated tablet contains:				
	Tablets IP 100 mg	Sildenafil Citrate	IP			APPROVED
		Eq. to Sildenafil		100	mg	
		Excipients		q.s		
		Approved colour used.		*		
45 1	Sildenafil Citrate	Each film coated tablet contains:				
731	Tablets IP 50 mg	Sildenafil Citrate	IP			APPROVED
	radicts if 50 mg	Eq. to Sildenafil	II'	50	mg	MIROVED
		Excipients		q.s	1115	
		Approved colour used.	1	4.5		
		ripproved corour used.				

452	Levocetirizine &	Each 5ml contains:				
	Ambroxol	Levocetirizine Dihydrochloride	IP	2.5	mg	APPROVED
	Hydrochloride Syrup	Ambroxol Hydrochloride	IP	30	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used.				
453	Metronidazole	Each 5 ml contains:				
	Benzoate Oral	Metronidazole Benzoate	IP			APPROVED
	Suspension IP	Eq. to Metronidazole		200	mg	
		In a flavoured base		q.s		
		Approved colour used.				
454	Methylcobalamin,	Each film coated tablet contains				
	Vitamin A Acetate,	Methylcobalamin	IP	500	mcg	
	Vitamin D3, Vitamin E	Vitamin A Acetate	IP	5000	IU	
	Acetate, Vitamin B1,	Vitamin D3	IP	400	IU	
	Riboflavin, Pyridoxine HCL, Niacinamide,	Vitamin E Acetate	IP	25	IU	
	Calcium Pantothenate	Vitamin B1	IP	10	mg	
	,Vitamin C, Folic Acid,	Riboflavin	IP	10	mg	
	Biotin, L-Glutamic	Pyridoxine Hydrochloride	IP	3	mg	
	Acid,	Niacinamide	IP	50	mg	
	Dl-Methionine,	Calcium Pantothenate	IP	12.5	mg	
	Magnesium Oxide,	Vitamin C	IP	100	mg	APPROVED
	Manganese, Copper, Zinc Sulphate	Folic Acid	IP	1500	mcg	
	Monohydrate,	Biotin	USP	60	mcg	
	Chromium,	L-Glutamic Acid	BP	100	mg	
	,Selenious Acid Tablet	Dl-Methionine	BP	25	mg	
		Magnesium Oxide	IP	30	mg	
		Manganese Gluconate Eq. to Elemental Manganese	USP	2.5	mg	
		Cupric Sulphate	USP			
		Eq. to Elemental Copper		2.5	mg	
		Zinc Sulphate Monohydrate	IP			
		Eq. to Elemental Zinc		22.5	mg	
		Chromium Picolinate	USP	100		
		Eq. to Chromium	LICD	100	mcg	
		Selenious Acid Eq. to Elemental Selenium	USP	60	meg	
		Excipients		q.s	mcg	
		Approved colour used.		4.5	<u> </u>	
		ripproved colour used.				

455	Methylcobalamin,	Each film coated tablet contain	s:			
	Alpha Lipoic Acid , Inositol Vitamin A	Methylcobalamin	IP	1500	mcg	
	Acetate, Vitamin B1,	Alpha Lipoic Acid	USP	100	mg	
	, Pyridoxine HCL,	Inositol	USP	100	mg	
	Vitamin E Acetate,	Vitamin A Acetate	IP	5000	IU	
	Calcium Pantothenate,	Vitamin B1	IP	10	mg	APPROVED
	Folic Acid, Chromium,	Pyridoxine Hydrochloride	IP	1.5	mg	
	,Selenious Acid Tablet	Vitamin E Acetate	IP	25	IU	
		Calcium Pantothenate	IP	10	mg	
		Folic Acid	IP	1500	mcg	
		Chromium Picolinate	USP			
		Eq. to Chromium		200	mcg	
		Selenious Acid	USP	162.6		
		Eq. to Elemental Selenium		163.6	mcg	
		Excipients Approved colour used.		q.s		
		**				
456	Nifedipine & Lidocaine	Composition:				
	Cream	Nifedipine	IP	0.3 %	w/w	APPROVED
		Lidocaine	BP	1.5 %	w/w	
		Cream Base				
457	Metformin	Each uncoated tablet contains:				
	Hydrochloride	Metformin Hydrochloride	IP	500	mg	Approved
	Tablets IP 500mg	Excipients		q.s		
458	Acarbose	Each uncoated tablet contains:				
	Tablet IP 50 mg	Acarbose	IP	50	mg	Approved
		Excipients		q.s		
459	Aciclovir	Each uncoated tablet contains:			· · · · · · · · · · · · · · · · · · ·	Annound
459	Tablet IP 200 mg	Aciclovir	IP	200	ma	Approved
	Tablet II 200 mg	Excipients	ır		mg	
		*		q.s		
460	Aciclovir	Each uncoated tablet contains:	_			
	Tablet IP 400 mg	Aciclovir	IP	400	mg	Approved
		Excipients		q.s		
461	Aciclovir	Each uncoated tablet contains:				
	Tablet IP 800 mg	Aciclovir	IP	800	mg	Approved
		Excipients		q.s		
462	Allopurinol	Each uncoated tablet contains:	•	•		
402	Tablet IP 100 mg	Allopurinol	IP	100	mg	Approved
	ruotet ir 100 mg	Excipients	11	q.s	mg	прриотец
463	Quetiapine Fumarate	Each film coated tablet contain	Annewed			
403	Tablets IP 50 mg Quetiapine Fumarate IP Quetiapine Fumarate IP					Approved
	Tableto II 50 llig	Eq. to Quetiapine	II.	50	mg	
		Excipients Excipients		q.s	8	
		Approved colour used.	1			
		•				

464	Quetiapine Fumarate	Each film coated tablet contains	s:			
	Tablets IP 100 mg	Quetiapine Fumarate	IP			Approved
		Eq. to Quetiapine		100	mg	
		Excipients		q.s		
		Approved colour used.				
465	Olanzapine	Each film coated tablet contains	s:			
	Tablets IP 2.5 mg	Olanzapine	IP	2.5	mg	Approved
		Excipients		q.s		
		Approved colour used.				
466	Oxcarbazepine	Each film coated tablet contains	s:			
	Tablets IP 450 mg	Oxcarbazepine	IP	450	mg	Approved
		Excipients		q.s		
		Approved colour used.	•			
167	0	Each film agated tablet contains				
467	Oxcarbazepine Tablets IP 300 mg	Each film coated tablet contains	s: IP	200		Approved
	Tablets II 500 mg	Oxcarbazepine	IP	300	mg	Approved
		Excipients Approved colour used.		q.s		-
		Approved colour used.				
468	r					
	Hydrochloride	Propranolol Hydrochloride	IP	40	mg	Approved
	Tablets IP 40 mg	Excipients		q.s		
469	C111	Each film coated tablet contains			1	Ammunud
409	Clopidogrel Tablet IP 75 mg	Clopidogrel bisulphate	s. IP	1	1	Approved
	rablet if 73 mg	Eq. to Clopidogrel	II	75	mg	
		Excipients		q.s		
		Approved colour used.		1 1		1
470	Tapentadol	Each film coated tablet contains				
	Hydrochloride	Tapentadole Hydrochloride	IP			Approved
	Tablet 50 mg	Eq. to Tapentadole		50	mg	
		Excipients		q.s		
		Approved colour used.				
471	Tapentadol	Each film coated tablet contains		1	1	
	Hydrochloride	Tapentadole Hydrochloride	IP	100	ma	Approved
	Tablet 100 mg	Eq. to Tapentadole Excipients			mg	-
		Approved colour used.	1	q.s		1
472	Clindamycin	**	inc:			
7/2	Capsule IP 150 mg	Each hard gelatin capsule conta Clindamycin Hydrochloride	IP			Approved
	Supposite if 150 mg	Eq. to Clindamycin	l IF	150	mg	Approved
		Excipients	-	q.s	1115	
		Approved colour used in empty	capsule		1	
450			poure			
473	Clotrimazole	Composition:		4.6		
	Cream IP	Clotrimazole	IP	1.0 %	w/w	Approved
		Preservative:		1.0.07	,	
		Benzyl Alcohol	IP	1.0 %	w/w	
		Cream Base				

474	Permethrin	Composition:				
•	Cream	Per methrin	[5.0 %	w/w	Approved
:	: :	Pre servative:	 			
:	:	Formaldehyde Solution	BP	0.1 %	w/w	
	: :	Cream Base		.b.=====		
	~	·				
475	Prebiotic & Probiotic	Each 1gm sachet contains:				
:	Sachet	Probiotic(CMP 5)	: :			
	<u> </u>	Total probiotic not less than		1.25 billion	Ì	
-		(Lactobacillus acidophilus,		CFU		_
		Lactobacillus rhamnosus,				Approved
Ē	<u> </u>	Bifidobacterium Longum,	: :			
:	:	Saccharomyces boulardii)	: :			
:		Fructo Oligosaccharides	<u> </u>	100	mg	
: : ;	- - -	Excipients	<u> </u>	q.s	<u> </u>	
476	Omeprazole &	Each hard gelatin capsule conta	ins:			
:	Domperidone	Omeprazole	IP	20	mg	"
-	Capsules IP	(As enteric coated pellets)				Approved
-		Domperidone	IP	10	mg	
		(As immediate release pellets)				
	:	Excipients	1	q.s		
		Approved colours used in emp	tv cansu	.J	·	
	: 	I ripproved colours used in emp				L
						F
477	: :	Composition:				
	Ethyl alcohol & glycerine	Ethyl alcohol 95%	ΙP	70%	v/v	
	(hand sanitizer)	Glycerine	ΙΡ	2%	V/V	Approved
	-	Perfume Gel Base	! 	q.s	 	
	-	Terrume der Base	::	q.s		
470	-	[C				
478	-	Composition:	: TS T		-1	
	Isopropyl alcohol With	Isopropyl alcohol	IΡ	70 %	v/v	
:	Chlorhexidine Gluconate	Chlorhexidine Gluconate			· - 	
:	Hand Rub		IΡ	2.5%	İ	Approved
	(hand sanitizer)	Solution Equi. To Chlorhexidine Gluconate		0.5%		прргочец
:	-	Cinomexianic Giaconate	<u> </u>	0.070	v/v	
-		Distilles Water		q.s		
	•				•	
479		Composition:				
:	: 	Ethyl alcohol 95%	IP	70	V/V	
-	Ethyl alcohol With Chlorhexidine Gluconate			%		
-	Hand Rub	Chlorhexidine Gluconate	IP:	2.5%		↓ • •
:	(hand sanitizer)	Solution Equi. To	-			Approved
	(Hand Samuzer)	Chlorhexidine Gluconate		0.5%	v/v	
:	i		ļ			I
:	: :	Distilles Water		q.s	1	
480	- - -	Composition:				
	Ethyl alcohol, Isopropyl	Ethyl alcohol 95%	IP	55	V/V	
	alcohol With Tocopheryl			<u>%</u>		
:	Acetate Hand Rub	Isopropyl alcohol	IΡ	10	v/v	
	(hand sanitizer)			%		Approved
		Tocopheryl Acetate	ΙΡ	0.05	v/v	
:		Distilles Water		q.s	7	1
-	-	•	i 1	1	i	i

481			Composition:				
	Isopropyl alcohol F Rub		Isopropyl alcohol	IP	70 %	V/V	Approved
	(hand sanitizer)		Aqueous Base		q.s		••
482	Cefixime,	Eac	h film coated tablet cor	ntains:			
	Azithromy		xime	IP			
	cin &	Eq.	to Anhydrous				
	Lactic Acid	Cef	xime		200	mg	
	Bacillus Tablets	Azi	hromycin	IP			
			o anhydrous		250	mg	APPROVED
		Lac	tic Acid Bacillus		2.5	billion spores	
			ipients		q.s		
		App	proved colour used				
483	Ofloxacin,	Con	position:				
	Ornidazole,	Oflo	xacin	IP	0.75%	w/w	
itraconazole & Clobetasol	Orni	dazole	IP	2%	w/w		
	Propionate Cream	Itrac	onazole	USP	1%	w/w	
	1	Clob	etasol Propionate	IP	0.05%	w/w	APPROVED
		Metl	nylparaben	IP	0.2%	w/w	
		(As	Preservative)				
			ylparaben	IP	0.02%	w/w	
	(A		Preservative)				
		In a	non-greasy base				
484	Citicoline &	Eac	h film coated tablet cor	ntains:			
	piracetam		coline Sodium	IP			
	Tablets		to Citicoline		500	mg	APPROVED
			cetam	IP	800	mg	
			ipients		q.s		
		App	roved colour used				
485	Rifaximin	Each	film coated tablet contai	ns:			
	Tablets		ximin	II	200	mg	Approved
			pients		q.s		
		Appı	oved colour used.				
486	Rifaximin		film coated tablet contai				
	Tablets		kimin	II		mg	Approved
			pients		q.s		
		App	oved colour used.				

487	Trypsin,	Each film coated tablet contain	ns:			
	Bromelain, Rutoside	Trypsin	IP	48	mg	
		Bromelain		90	mg	
	Trihydeate &	Rutoside Trihydeate	IP	100	mg	APPROVED
	Serratiopeptid ase Tablets	Serratiopeptidase (EC) (as 30,000 units of	IP	15	mg	ATTROVED
		Serratiopeptidase)				
		Excipients		q.s		
		Approved colour used				
488	Nortriptyline	Each film coated tablet cor	ntains:			
	& pregabalin	Nortriptyline				
	tablets	hydrochloride				APPROVED
		Eq. to	IP			APPROVED
		Nortriptyline		10	mg	
		Pregabalin	IP	75	mg	
		Excipients	- 11	+	mg	
				q.s		
		Approved colour used				
489	Nortriptyline	Each film coated tablet cor	ntains:			
	& Gabapentin	Nortriptyline				
	tablets	hydrochloride	-			APPROVED
		Eq. to	IP			ATTROVED
		Nortriptyline		10	mg	
		Gabapentin	IP	400	mg	
		Excipients		q.s		
		Approved colour used		1.5	L	
l l					<u> </u>	
490	Clobetasol	Composition:				
	Propionate,	Clobetasol Propionate	IP	0.05%	w/w	
	fusidic acid Cream	fusidic acid	IP	2%	w/w	APPROVED
		In a Cream base				
491	Trypsin,	Each enteric coated tablet of	contains:			
	Bromelain &	Trypsin	IP	48	mg	
	Rutoside	Bromelain		90	mg	
	Trihydeate	Rutoside Trihydeate	IP	100	mg	
	Tablets					APPROVED
		Excipients		q.s		
		Approved colour used				
492	Trypsin,	Each enteric coated tablet of	contains:			
	Bromelain,	Trypsin	IP	48	ma	
	Rutoside		ır		mg	
	Trihydeate &	Bromelain		90	mg	
	diclofenac	Rutoside Trihydeate	IP	100	mg	APPROVED
	potassium	diclofenac potassium	IP	50	mg	
	Tablets	Excipients		q.s		

Approved colour used		
ripproved colour used		Approved colour used

493.	Diclofenac Sodium Prolonged -	Each film coated prolonged - release	tablet cont	ains:		
	release Tablet IP	Diclofenac Sodium	IP	100	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
494.	Dicyclomine Hydrochloride	Each uncoated tablet contains:				Approved on
	Tablets IP	Dicyclomine Hydrochloride	IP	20	mg	17.05.2021
		Excipients		q.s		17.03.2021
405	To: 1 : 11 : 1	T. 1.5.1				1
495.	Dicyclomine Hydrochloride	Each 5 ml contains:	1 m 1	10	1	
	Oral solution IP	Dicyclomine Hydrochloride	IP	10	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
106	Diltiogom Hydrochloride	Each film coated tablet contains:				
496.	Diltiazem Hydrochloride Tablets IP	Diltiazem Hydrochloride	IP	30	-m~	A 1
	Tablets IP	-	IP		mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
497	Diltiazem Hydrochloride	Each film coated tablet contains:				
777	Tablets IP	Diltiazem Hydrochloride	IP	60	mg	Approved on
	Tablets II	Excipients	11		mg	17.05.2021
		Approved colour used		q.s		17.03.2021
		Approved colour used				
498.	Disulfiram Tablets IP	Each uncoated tablet contains:				
		Disulfiram	IP	250	mg	- Approved on
		Excipients		q.s		17.05.2021
		1		1	_	<u> </u>
499.	Disulfiram Tablets IP	Each uncoated tablet contains:				A 1
		Disulfiram	IP	500	mg	Approved on
		Excipients		q.s		17.05.2021
500.	Divalproex Sodium Gastro-	Each Gastro-resistant tablet contains:				
	resistant Tablets IP	Divalproex Sodium	IP			Annroyadan
		eq. to Valproic Acid		125	mg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used				
501.	Divalproex Sodium Gastro-	Each Gastro-resistant tablet contains:			_	
	resistant Tablets IP	Divalproex Sodium	IP			Approved on
		eq. to Valproic Acid		250	mg	17.05.2021
		Excipients		q.s		17.05.2021
l		Approved colour used				

502.	Divalproex Sodium Gastro-	Each Gastro-resistant tablet conta	ins:			
	resistant Tablets IP	Divalproex Sodium	IP			1
		eq. to Valproic Acid		500	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	l l	-		-
700	I					1
503.	Divalproex Sodium Prolonged-	Each film coated prolonged-relea	1	uns:	1	
	release Tablets IP	Divalproex Sodium	IP			Approved on
		eq. to Valproic Acid		125	mg	17.05.2021
		Excipients		q.s		
		Approved colour used				
504.	Divalproex Sodium Prolonged-	Each film coated prolonged-relea				
	release Tablets IP	Divalproex Sodium	IP			1
		eq. to Valproic Acid		500	mg	Approved or 17.05.2021
		Excipients		q.s		
ļ		Approved colour used		1		1
505.	Divalproex Sodium Prolonged-	Each film coated prolonged-relea	se tablet conta			
	release Tablets IP	Divalproex Sodium	IP			Approved or
		eq. to Valproic Acid		750	mg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used				
506.	Divalproex Sodium Prolonged-	Each film coated prolonged-relea	se tablet conta	ins:		1
	release Tablets IP	Divalproex Sodium	IP			Approved on
	Teleuse Tueleus II	eq. to Valproic Acid		1000	mg	
		Excipients		q.s	11.5	17.05.2021
		Approved colour used		4.5		
507.	Domperidone Maleate Tablets	Each uncoated tablet contains:			•	
	IP	Domperidone Maleate	IP			Approved on
		eq. to Domperidone		10	mg	17.05.2021
		Excipients		q.s		
508.	Domperidone Suspension IP	Each ml contains:				
500.	Domperidone Suspension if	Domperidone	IP	1	ma	
		-	IP		mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
509.	Donepezil Hydrochloride	Each film coated tablet contains:				
	Tablets IP	Donepezil Hydrochloride	IP	5	mg	Approved on
		Excipients		q.s	1	17.05.2021
		Approved colour used	I I	•	1	1

510.	Donepezil Hydrochloride	Each film coated tablet contains:					
	Tablets IP	Donepezil Hydrochloride	IP	10	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used			_ L	1	
511.	Dothiepin Hydrochloride Each hard gelatin capsule contains:						
	Capsules IP	Dothiepin Hydrochloride	IP	25	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used			•		
512.	Dothiepin Hydrochloride	Each film coated contains:			1		
	Tablets IP	Dothiepin Hydrochloride	IP	25	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used					
512	D-41::- H111:-1-	Each film coated contains:					
513.	Dothiepin Hydrochloride Tablets IP	Dothiepin Hydrochloride	ID	50	1	Approved on 17.05.2021	
	Tablets IP	1 2	IP	50	mg		
		Excipients		q.s		17.03.2021	
		Approved colour used					
514.	Dothiepin Hydrochloride	Each film coated contains:					
51	Tablets IP	Dothiepin Hydrochloride	IP	75	mg	Approved on	
		Excipients		q.s	1115	17.05.2021	
		Approved colour used		4.5		- 17.03.2021	
						ı	
515.	Dothiepin Hydrochloride	Oothiepin Hydrochloride Each film coated contains:					
	Tablets IP	Dothiepin Hydrochloride	IP	100	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used			•		
516.	1 3	Each film coated contains:			1		
	Tablets IP	Dothiepin Hydrochloride	IP	150	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used					
517	Danish Hadaa dhadda	Each hand calatin consula contain					
517.	Doxepin Hydrochloride Capsules IP	Each hard gelatin capsule contain Doxepin Hydrochloride	IP		1	_	
	Capsules IP			25	ma	Approved on	
		eq. to Doxepin			mg	17.05.2021	
		Excipients		q.s		_	
		Approved colour used				1	
518.	Doxepin Hydrochloride	Each hard gelatin capsule contain	s:				
210.	Capsules IP	Doxepin Hydrochloride	IP			1	
	Caponico II	eq. to Doxepin	"	50	mg	Approved on	
		Excipients		q.s	5	17.05.2021	
		Approved colour used			1	1	

519.	Doxepin Hydrochloride	Each hard gelatin capsule contains:					
	Capsules IP	Doxepin Hydrochloride	IP			A mmmay and a mm	
		eq. to Doxepin		75	mg	Approved or 17.05.2021	
		Excipients		q.s		17.03.2021	
		Approved colour used			•		
520.	Doxycycline Hydrochloride	Each hard gelatin capsule contains:					
320.	Capsules IP	Doxycycline Hydrochloride	IP			-	
	Capsules IF	eq. to Doxycycline	IL	50	ma	Approved or	
		Excipients			mg	17.05.2021	
		Approved colour used		q.s		-	
						1	
521.	Doxycycline Hydrochloride	Each hard gelatin capsule contains:					
	Capsules IP	Doxycycline Hydrochloride	IP			Approved or	
		eq. to Doxycycline		100	mg	17.05.2021	
		Excipients		q.s			
		Approved colour used					
523.	Doxycycline Hydrochloride	Each hard gelatin capsule contains:					
323.	Capsules IP	Doxycycline Hydrochloride	IP			Approved on 17.05.2021	
	Cupsules II	eq. to Doxycycline	1 1	200	mg		
		Excipients		q.s	mg		
		Approved colour used		4.5		<u> </u>	
	<u> </u>	rippio ved colour used					
524.	Doxycycline Monohydrate	Each uncoated dispersible tablet co	ntains:				
	Tablets IP	Doxycycline Monohydrate	IP			Approved or	
		eq. to Doxycycline		100	mg	17.05.2021	
		Excipients		q.s			
525	D.,	Each film coated tablet contains:				<u> </u>	
525.	Drotaverine Hydrochloride		I ID	40	<u> </u>		
	Tablets IP	Drotaverine Hydrochloride	IP	40	mg	Approved or	
		Excipients		q.s		17.05.2021	
		Approved colour used					
526.	Econazole Nitrate Cream IP	IP Composition:				T	
		Econazole Nitrate	IP	1.0%	w/w	Approved or 17.05.2021	
		In cream base		q.s		17.05.2021	
527.	Enalapril Maleate Tablets IP	Each uncoated tablet contains:					
J41.	Emalapin Maicate Tablets II	Enalapril	IP	2.5	mg	Approved or	
		Excipients	11	q.s	mg	17.05.2021	
			ı	-	1	1	
528.	Enalapril Maleate Tablets IP	Each uncoated tablet contains:	 ,		1	Approved or	
		Enalapril	IP	5	mg	17.05.2021	
	1	Excipients		q.s	1	17.05.2021	

529.	Enalapril Maleate Tablets IP	Each uncoated tablet contains:				Ammoveden	
		Enalapril	IP	10	mg	- Approved on 17.05.2021	
		Excipients		q.s		17.03.2021	
530.	Enalapril Maleate Tablets IP	Each uncoated tablet contains:					
	•	Enalapril	IP	20	mg	Approved on	
		Excipients		q.s		- 17.05.2021	
531.	Enalapril Maleate and	nd Each uncoated tablet contains:					
331.	Hydrochlorothiazide Tablets IP	Enalapril	IP	2.5	mg	Approved on	
	Try droemorounazide Tublets II	Hydrochlorothiazide	IP	12.5	mg	17.05.2021	
		Excipients	11	q.s	Ilig	- 17.03.2021	
		Excipients		4.5			
532.	Enalapril Maleate and	Each uncoated tablet contains:					
	Hydrochlorothiazide Tablets IP	Enalapril	IP	5	mg	Approved on	
		Hydrochlorothiazide	IP	12.5	mg	17.05.2021	
		Excipients		q.s		1	
533.	Enalapril Maleate and	Each uncoated tablet contains:			_		
	Hydrochlorothiazide Tablets IP	Enalapril	IP	10	mg	Approved on	
		Hydrochlorothiazide	IP	25	mg	17.05.2021	
		Excipients		q.s			
534.	Ergotamine Tartrate Tablets IP	Each uncoated tablet contains:					
0011	Eigotamine Tarate Tablets in	Ergotamine Tartrate	IP	1	mg	Approved on	
		Excipients		q.s	8	17.05.2021	
					1		
535.	Erythromycin Gastro-resistant	Each Gastro-resistant tablet contains:					
	Tablets IP	Erythromycin	IP	250	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used					
						1	
536.	Erythromycin Stearate Tablets	Each film coated tablet contains:			1	_	
	IP	Erythromycin Stearate	IP	107		Approved on	
		eq. to Erythromycin		125	mg	17.05.2021	
		Excipients		q.s		- 17.03.2021	
		Approved colour used					
537.	Erythromycin Stearate Tablets	Each film coated tablet contains:					
331.	IP	Erythromycin Stearate	IP			1	
		eq. to Erythromycin		250	mg	Approved on	
		Excipients		q.s	1115	17.05.2021	
			1	4.0	1	1	

538.	Escitalopram Oxalate Tablets	Each film coated tablet contains:				
	IP	Escitalopram Oxalate	IP] , , ,
		eq. to Escitalopram		5	mg	Approved on 17.05.2021
		Excipients		q.s		17.05.2021
		Approved colour used	I			_
	I					
539.	Escitalopram Oxalate Tablets	Each film coated tablet contains:				
	IP	Escitalopram Oxalate	IP			Approved on
		eq. to Escitalopram		10	mg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used				
540.	Escitalopram Oxalate Tablets	Each film coated tablet contains:			_	
	IP	Escitalopram Oxalate	IP			Approved on
		eq. to Escitalopram		20	mg	17.05.2021
		Excipients		q.s		
		Approved colour used				
~ 4.1	[n					
541.	Eslicarbazepine Acetate Tablets	Each film coated tablet contains:	m	200	1	┧
	IP	Eslicarbazepine Acetate	IP	200	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
542.	Eslicarbazepine Acetate Tablets	Tablets Each film coated tablet contains:				
3 12.	IP	Eslicarbazepine Acetate	IP	400	mg	Approved on
		Excipients		q.s	1115	17.05.2021
		Approved colour used		4.5		17.00.2021
	<u> </u>	Tipproved colour used				
543.	Eslicarbazepine Acetate Tablets	Each film coated tablet contains:				
	IP	Eslicarbazepine Acetate	IP	600	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	ll	<u>_</u>		
						1
544.	Eslicarbazepine Acetate Tablets	Each film coated tablet contains:				
	IP	Eslicarbazepine Acetate	IP	800	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	•			
545.	Etodolac Capsules IP	Each hard gelatin capsule contains:			_	
		Etodolac	IP	200	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				

546.	Esomeprazole Magnesium	Each Gastro-resistant capsules con	ntains:			
	Trihydrate Gastro-resistant	Esomeprazole Magnesium				1
	Capsules IP	Trihydrate	IP			Approved on
		eq. to Esomeprazole		40	mg	17.05.2021
		Excipients		q.s		1
		Approved colour used	I			1
547.	1 0	Each Gastro-resistant capsules con	ntains:		1	
	Trihydrate Gastro-resistant	Esomeprazole Magnesium				
	Capsules IP	Trihydrate	IP			Approved on
		eq. to Esomeprazole		20	mg	17.05.2021
		Excipients		q.s		
		Approved colour used				
548.	Etodolac Capsules IP	Each hard gelatin capsule contains				<u> </u>
5 10.	Etodolae Capsules II	Etodolac	IP	300	mg	Approved on
		Excipients	11		IIIg	17.05.2021
		Approved colour used		q.s		17.03.2021
		Approved coroni used				
549.	Etodolac Sustained-release	Each film coated sustained release	tablet contai	ns:		
	Tablets IP	Etodolac	IP	200	mg	Approved on 17.05.2021
		Excipients		q.s		
		Approved colour used	l			1
						1
550.	Etodolac Sustained-release	Each film coated sustained release				
550.	Etodolac Sustained-release Tablets IP	Etodolac	tablet contai	ns: 600	mg	Approved on
550.		Etodolac Excipients			mg	Approved on 17.05.2021
550.		Etodolac		600	mg	
550. 551.	Tablets IP	Etodolac Excipients		600	mg	
		Etodolac Excipients Approved colour used		600 q.s		17.05.2021
	Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac	IP	600 q.s	mg	17.05.2021 Approved on
	Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients	IP	600 q.s		17.05.2021
	Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac	IP	600 q.s		17.05.2021 Approved on
	Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients	IP	600 q.s		17.05.2021 Approved on
551.	Tablets IP Etodolac Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used	IP	600 q.s		17.05.2021 Approved on
551.	Tablets IP Etodolac Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Each film coated tablet contains:	IP IP	600 q.s 200 q.s	mg	17.05.2021 Approved on 17.05.2021
551.	Tablets IP Etodolac Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac	IP IP	200 q.s 300	mg	Approved on 17.05.2021 Approved on Approved on
551. 552.	Tablets IP Etodolac Tablets IP Etodolac Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Excipients Approved colour used	IP IP	200 q.s 300	mg	Approved on 17.05.2021 Approved on Approved on
551.	Tablets IP Etodolac Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used	IP IP	200 q.s 300 q.s	mg	Approved on 17.05.2021 Approved on 17.05.2021 Approved on 17.05.2021
551. 552.	Tablets IP Etodolac Tablets IP Etodolac Tablets IP	Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Each film coated tablet contains: Etodolac Excipients Approved colour used Excipients Approved colour used	IP IP	200 q.s 300	mg	Approved on 17.05.2021 Approved on Approved on

554.	Cilnidipine Tablets IP	Each film coated tablet contains:				
	_	Cilnidipine	IP	5	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
555.	Cilnidipine Tablets IP	Each film coated tablet contains:			_	
		Cilnidipine	IP	10	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
556.	CilnidipineTablets IP	Each film coated tablet contains:		20	1	A
		Cilnidipine	IP	20	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
557.	Cimetidine Tablets IP	Each uncoated tablet contains:				1
337.	Cimetidine Tablets II	Cimetidine	IP	200	ma	Approved on
		Excipients	11	q.s	mg	17.05.2021
		Excipients		4.5		
558.	Cimetidine Tablets IP	Each uncoated tablet contains:				
		Cimetidine	IP	400	mg	- Approved on
		Excipients		q.s		17.05.2021
	l	1		1	I	
559.	Cimetidine Tablets IP	Each uncoated tablet contains:				- Approved on
		Cimetidine	IP	800	mg	
		Excipients		q.s		17.05.2021
560.	Etodolac Tablets IP	Each film coated tablet contains:				
		Etodolac	IP	300	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	<u> </u>			1
561.	Cholecalciferol Tablet IP	Each film coated tablet contains:				
		Cholecalciferol	IP	0.25	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used		4.5		- 17,000,2021
		Approved colour used				
562.	Citalopram Hydrobromide	Each film coated tablet contains:				
302.	Tablets IP	Citalopram Hydrobromide	IP			1
	Tablets II	eq. to Citalopram	11	10	ma	Approved on
					mg	17.05.2021
		Excipients		q.s		_
		Approved colour used				
5.02	C'. 1 II 1 1 11	Early Classical College				1
563.	Citalopram Hydrobromide	Each film coated tablet contains:				4
	Tablets IP	Citalopram Hydrobromide	IP			Approved on
		eq. to Citalopram		40	mg	17.05.2021
		Excipients		q.s		17.03.2021
l l		Approved colour used			'	

564.	Citicoline Sodium Prolonged -	Each film coated prolonged release t	ablet conta	ains:		
	release Tablets IP	Citicoline Sodium	IP			1
		e.q to Citicoline		1000	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				-
565.	Citicoline Sodium Tablets IP	Each film coated tablet contains:				
		Citicoline Sodium	IP			Annroyadan
		e.q to Citicoline		100	mg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used				
						_
566.	Citicoline Sodium Tablets IP	Each film coated tablet contains:			_	
		Citicoline Sodium	IP			Approved on
		e.q to Citicoline		1205	mg	17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used				
						1
567.	Clobetasol Propionate Cream	Composition:	•	T	1	Approved on
	IP	Clobetasol Propionate	IP	0.05%	w/w	17.05.2021
		In a cream base		q.s		
7.60						1
568.	Clobetasol Propionate	Composition:		0.050/	1 ,	Approved on
	Ointment IP	Clobetasol Propionate	IP	0.05%	w/w	17.05.2021
		In a ointment base		q.s		
569.	Clomifene Citrate Tablets IP	Each uncoated tablet contains:				1
307.	Cionniche Citrate Tablets II	Clomifene Citrate	IP	25	ma	Approved on
		Excipients	11		mg	17.05.2021
		Excipients		q.s		
570.	Clomifene Citrate Tablets IP	Each uncoated tablet contains:				
		Clomifene Citrate	IP	50	mg	Approved on
		Excipients		q.s	8	17.05.2021
		r		1.~	1	
571.	Clonidine Hydrochloride and	Each uncoated tablet contains:				
	Chlorthalidone Tablets IP	Clonidine Hydrochloride	IP	0.1	Mg	Approved on
		Chlorthalidone	IP	15	Mg	17.05.2021
		Excipients		q.s		1
	<u> </u>	· •			1	1
572.	Clopidogrel Bisulphate IP	Each film coated tablet contains:				
	_	Clopidogrel Bisulphate	IP			Λ mmmo 1 -
		Eq. to Clopidogrel		75	mg	Approved on
		Excipients		q.s		17.05.2021
				1		

	Clotrimazole Cream IP	Composition:				Approved on
		Clotrimazole	IP	1.0%	w/w	17.05.2021
		In a cream base		q.s		17.03.2021
						_
574.	Clotrimazole Cream IP	Composition:			1	Approved or
		Clotrimazole	IP	2.0%	w/w	17.05.2021
		In a cream base		q.s		1710012021
575		Each uncoated tablet contains:				
575.	Cyproheptadine Hydrochloride Tablets IP	Cyproheptadine Hydrochloride	IP	1		A
	Tablets IP		IP	4	mg	Approved on 17.05.2021
		(Anhydrous)				17.03.2021
		Excipients		q.s		
576.	Dexamethasone Tablets IP	Each uncoated tablet contains:				
J.	1 401040 11	Dexamethasone	IP	0.5	mg	Approved or
		Excipients		q.s	III.S	17.05.2021
	L	Excipients		4.5		
577.	Dexchlorpheniramine Maleate	Each 5 ml contains:				
	Oral Solution IP	Dexchlorpheniramine Maleate	IP	2	mg	Approved or
		In a flavoured syrupy base		q.s		17.05.2021
		Approved colour used				-
578.	Dextromethorphan	Each 5 ml contains:				Approved or
	Hydrobromide Syrup IP	Dextromethorphan Hydrobromide	IP	13.5	mg	17.05.2021
		Excipients		0.0		17.05.2021
		_		q.s		
		Approved colour used		ų.s		-
		Approved colour used		q.s		1
579.	Diacerein Capsules IP	Approved colour used Each hard gelatin capsule contains:				Approved or
579.	Diacerein Capsules IP	Approved colour used Each hard gelatin capsule contains: Diacerein	IP	50	mg	Approved or 17.05.2021
579.	Diacerein Capsules IP	Approved colour used Each hard gelatin capsule contains: Diacerein Excipients	IP		mg	
579.	Diacerein Capsules IP	Approved colour used Each hard gelatin capsule contains: Diacerein	IP	50	mg	
		Approved colour used Each hard gelatin capsule contains: Diacerein Excipients Approved colour used	IP	50	mg	
579. 580.	Diacerein Capsules IP Diazoxide Tablets IP	Approved colour used Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains:		50 q.s		17.05.2021 Approved or
		Approved colour used Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains: Diazoxide	IP IP	50 q.s	mg	Approved on 17.05.2021 Approved on 17.05.2021
		Approved colour used Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains:		50 q.s		17.05.2021 Approved or
		Approved colour used Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains: Diazoxide		50 q.s		17.05.2021 Approved or 17.05.2021
580.	Diazoxide Tablets IP	Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains: Diazoxide Excipients		50 q.s		Approved or 17.05.2021 Approved or Approved or
580.	Diazoxide Tablets IP	Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains: Diazoxide Excipients Each uncoated tablet contains:	IP	50 q.s 50 q.s	mg	17.05.2021 Approved or 17.05.2021
580.	Diazoxide Tablets IP Atenolol Tablets IP	Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains: Diazoxide Excipients Each uncoated tablet contains: Atenolol Excipients	IP	50 q.s 50 q.s	mg	Approved or 17.05.2021 Approved or Approved or
580.	Diazoxide Tablets IP	Approved colour used Each hard gelatin capsule contains: Diacerein Excipients Approved colour used Each uncoated tablet contains: Diazoxide Excipients Each uncoated tablet contains: Atenolol	IP	50 q.s 50 q.s	mg	Approved or 17.05.2021 Approved or Approved or

583.	Atenolol and Chlorthalidone	Each film coated tablet contains:					
	Tablets IP	Atenolol	IP	25	mg	Approved on	
		Chlorthalidone	IP	6.25	mg	17.05.2021	
		Excipients		q.s		17.03.2021	
		Approved colour used					
584.	Atomolol and Chlombolidana	Each film coated tablet contains:					
384.	Atenolol and Chlorthalidone Tablets IP	Atenolol	I ID	<i>5</i> 0		<u> </u> 	
	Tablets IP		IP	50	mg	Approved or	
		Chlorthalidone	IP	12.5	mg	17.05.2021	
		Excipients Approved colour used		q.s		_	
	<u> </u>	Approved colour used					
585.	Atenolol and Chlorthalidone	hlorthalidone Each film coated tablet contains:					
	Tablets IP	Atenolol	IP	100	mg	Approved or	
		Chlorthalidone	IP	25	mg	17.05.2021	
		Excipients		q.s		17.05.2021	
		Approved colour used	1	_	•		
586.	Atorvastatin Calcium and Fenofibrate Tablets IP	Each film coated tablet contains:					
		Atorvastatin Calcium	IP				
		eq. to Atorvastatin		10	mg	Approved or	
		Fenofibrate	IP	72.5	mg	17.05.2021	
		Excipients		q.s			
		Approved colour used					
587.	Atorvastatin Calcium and Fenofibrate Tablets IP	rvastatin Calcium and Each film coated tablet contains:					
		Atorvastatin Calcium	IP				
		eq. to Atorvastatin		20	mg	Approved or	
		Fenofibrate	IP	145	mg	17.05.2021	
		Excipients		q.s			
		Approved colour used	<u> </u>	•			
588.	A -: 41	Each hard colotin concells contained					
300.	Azithromycin Capsules IP	Each hard gelatin capsule contains:	IP	500	ma	_	
		Azithromycin (Anhydrous) (as Azithromycin Dihydrate)	IF	300	mg	Approved or	
		Excipients		q.s		17.05.2021	
		Approved color used		-			
589.	Betahistine Hydrochloride	Each uncoated tablet contains:			1	Approved or	
	Tablets IP	Betahistine Hydrochloride	IP	50	mg	17.05.2021	
		Excipients		q.s			
590.	Betamethasone Tablets IP	methasone Tablets IP					
570.	Detailled addets if	Betamethasone	IP	0.5	mg	Approved or	
		Excipients	11	q.s	1118	17.05.2021	
	ı	1 *		1	1	1	
591.	Betamethasone Tablets IP	Each uncoated tablet contains:				Annroyed or	
	1	Betamethasone	IP	1.0	ma	- Approved on 17.05.2021	
		Detaineulasone	- 11	1.0	mg	17.05.2021	

592.	Betamethasone Dipropionate	Composition				
	Cream IP	Betamethasone Dipropionate	IP			Approved on
		eq. to Betamethasone		0.05%	w/w	17.05.2021
		In a cream base		q.s		
502	D-441	Composition				1
593.	Betamethasone Dipropionate	Composition	TD	1	1	Approved on
	Lotion IP	Betamethasone Dipropionate	IP	0.050/	/	Approved on
		eq. to Betamethasone		0.05%	w/w	17.05.2021
		In a lotion base		q.s		
594.	Betamethasone Dipropionate	Composition				
57	Ointment IP	Betamethasone Dipropionate	IP			Approved or
		eq. to Betamethasone		0.05%	w/w	17.05.2021
		In a ointment base		q.s	1	17.03.2021
				1		
595.	Bromhexine Hydrochloride	Each uncoated tablet contains:				Approved on
	Tablets IP	Bromhexine Hydrochloride	IP	4	mg	
		Excipients		q.s		17.05.2021
596.	Bromhexine Hydrochloride Tablets IP	Each uncoated tablet contains:				Approved on
		Bromhexine Hydrochloride	IP	8	mg	17.05.2021
		Excipients		q.s		17.03.2021
		1				ı
597.	Captopril and Hydrochlorothiazide Tablets IP	Each uncoated tablet contains:	-		1	
		Captopril	IP	25	mg	Approved on 17.05.2021
		Hydrochlorothiazide	IP	15	mg	
		Excipients				
500	C	Each uncoated tablet contains:				<u> </u>
598.	Captopril and		IP	50		Approved on
	Hydrochlorothiazide Tablets IP	Captopril Hydrochlorothiazide	IP IP	25	mg	17.05.2021
		Excipients	11	23	mg	
		Excipients				
599.	Captopril and	Each uncoated tablet contains:				
5,,,	Hydrochlorothiazide Tablets IP	Captopril	IP	25	mg	Approved on
	Try droemorounaziae Tablets II	Hydrochlorothiazide	IP	25	mg	17.05.2021
		Excipients				1710012021
		T				<u> </u>
600.	Carbamazepine Prolonged –	Each uncoated prolonged release table	t contain	ıs:		Approved on
	release Tablets IP	Carbamazepine	IP	100	mg	Approved on
		Excipients				17.05.2021
					•	
601.	Carbamazepine Prolonged –	Each uncoated prolonged release table	t contain	ıs:		Approved on
	release Tablets IP	Carbamazepine	IP	200	mg	17.05.2021
		Excipients				17.03.2021
	T	I m				T
602.	Carbamazepine Prolonged –	Each uncoated prolonged release table			T	Approved on
	release Tablets IP	Carbamazepine	IP	400	mg	17.05.2021
		Excipients				17.03.2021
605	I a	In a contract of				T
603.	Captopril Tablets IP	Each uncoated tablet contains:		10.7		Approved on
		Captopril	IP	12.5	mg	17.05.2021
		Excipients				

604.	Captopril Tablets IP	Each uncoated tablet contains:				Annroyadan
	1 1	Captopril	IP	25	mg	Approved on
		Excipients				17.05.2021
		1		I.		
605.	Captopril Tablets IP	Each uncoated tablet contains:				Approved on
		Captopril	IP	50	mg	17.05.2021
		Excipients				17.03.2021
						_
606.	Carbamazepine Tablets IP	Each uncoated tablet contains:			_	Approved on
		Carbamazepine	IP	400	mg	17.05.2021
		Excipients				17,00,2021
		T				1
607.	Carbidopa and Levodopa	Each uncoated Orally Disintegrating t		tains:		
	Orally Disintegrating Tablets IP	Carbidopa	IP			Approved on
		eq. to Carbidopa Anhydrous		25	mg	17.05.2021
		Levodopa	IP	100	mg	
		Excipients				
(00		E-1	-1-1 /	4-:		1
608.	Carbidopa and Levodopa	Each uncoated Orally Disintegrating t		tains:	1	4
	Orally Disintegrating Tablets IP	Carbidopa	IP	25		Approved on
		eq. to Carbidopa Anhydrous	TD	25	mg	17.05.2021
		Levodopa	IP	250	mg	
		Excipients				
609.	Carbidopa and Levodopa	Each uncoated Orally Disintegrating t	ahlet con	taine:		1
00).	Orally Disintegrating Tablets IP	Carbidopa	IP	tanis.		
	Orany Disintegrating Tablets IP	eq. to Carbidopa Anhydrous	1 11	10	mg	Approved on
		Levodopa	IP	100	mg	17.05.2021
		Excipients		100	1115	1
						<u> </u>
610.	Carbimazole Tablets IP	Each uncoated tablet contains:				
		Carbimazole	IP	5	mg	Approved on
		Excipients				17.05.2021
					1	
611.	Carbimazole Tablets IP	Each uncoated tablet contains:				
611.		Carbimazole	IP	20	mg	Approved on
		i Carbimazoie	I IP		1112	17.05.2021
			IP	20		17.03.2021
		Excipients	IP	20	8	17.03.2021
612	Cetirizine Tablets IP	Excipients	IP IP	20		17.03.2021
612.	Cetirizine Tablets IP	Excipients Each uncoated tablet contains:				Approved on
612.	Cetirizine Tablets IP	Excipients Each uncoated tablet contains: Cetirizine	IP IP	20	mg	
612.	Cetirizine Tablets IP	Excipients Each uncoated tablet contains:				Approved on
		Excipients Each uncoated tablet contains: Cetirizine Excipients				Approved on
612.	Cetirizine Tablets IP Chlorhexidine Mouthwash IP	Excipients Each uncoated tablet contains: Cetirizine Excipients Composition:	IP			Approved on
		Excipients Each uncoated tablet contains: Cetirizine Excipients Composition: Chlorhexidine Gluconate Solution		20	mg	Approved on 17.05.2021
		Excipients Each uncoated tablet contains: Cetirizine Excipients Composition: Chlorhexidine Gluconate Solution diluted to Chlorhexidine Gluconate	IP			Approved on 17.05.2021 Approved on
		Excipients Each uncoated tablet contains: Cetirizine Excipients Composition: Chlorhexidine Gluconate Solution	IP	20	mg	Approved on 17.05.2021
		Excipients Each uncoated tablet contains: Cetirizine Excipients Composition: Chlorhexidine Gluconate Solution diluted to Chlorhexidine Gluconate	IP	20	mg	Approved on 17.05.2021 Approved on

614.	Chlorhexidine Mouthwash IP	Composition:						
		Chlorhexidine Gluconate Solution	IP					
		diluted to Chlorhexidine Gluconate		0.5%	w/v	Approved on		
		In a pleasantly flavoured aqueous				17.05.2021		
		base						
		Approved colour used				_		
		Tippio ved colodi discu						
615.	Chlorhexidine Mouthwash IP	hexidine Mouthwash IP Composition:						
		Chlorhexidine Gluconate Solution	IP					
		diluted to Chlorhexidine Gluconate		4%	w/v	Approved on		
		In a pleasantly flavoured aqueous				17.05.2021		
		base						
		Approved colour used			•			
		La				T		
616.	Chlorhexidine Mouthwash IP	Composition:	T 100			_		
		Chlorhexidine Gluconate Solution	IP	200/	,			
		diluted to Chlorhexidine Gluconate		20%	w/v	Approved on		
		In a pleasantly flavoured aqueous				17.05.2021		
		base						
		Approved colour used						
617.	Chloroquine Phosphate	Each 5 ml contains:						
017.	Suspension IP	Chloroquine Phosphate	IP	80	mg	_		
	Suspension II	eq. to Chloroquine base	1 1	50	mg	Approved on		
		In a flavoured syrupy base		q.s	ıng	17.05.2021		
		Approved colour used		T**				
		II						
618.	Chloroquine Phosphate Tablets	Each film coated tablet contains:						
	IP	Chloroquine Phosphate	IP	250	mg	Ammoveden		
		eq. to Chloroquine		155	mg	Approved on 17.05.2021		
		Excipients		q.s		17.03.2021		
		Approved colour used			•			
619.	Chloroquine Sulphate Tablets	Each film coated tablet contains:	T 1	• 0 -	1	_		
	IP	Chloroquine Sulphate	IP	200	mg	Approved on		
		eq. to Chloroquine		147	mg	17.05.2021		
		Excipients		q.s		_		
		Approved colour used						
620.	Chloroquine Sulphate Syrup IP	Each 5 ml contains:				T		
020.	Chroroquine Surphate Syrup IF	Chloroquine Sulphate	IP	80	ma	-		
		eq. to Chloroquine base	l IL	67	mg mg	Approved on		
		In a syrupy base			mg	17.05.2021		
		Approved colour used	1	q.s		_		
		Approved colour used						

621.	Chloroquine Sulphate Syrup IP	Each 5 ml contains:					
		Chloroquine Sulphate	IP	80	mg	1	
		eq. to Chloroquine base		50	mg	Approved on	
		In a syrupy base		q.s	8	17.05.2021	
		Approved colour used		4.5		-	
		Approved colour used					
622.	Chlorothiazide Tablets IP	Each film coated tablet contains:					
		Chlorothiazide	IP	250	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used	<u> </u>		_1	1	
623.	Chlorothiazide Tablets IP	Each film coated tablet contains:					
		Chlorothiazide	IP	500	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour used	Approved colour used				
624.	Chlorothiazide Oral Suspension	Each ml contains:					
	IP	Chlorothiazide	IP	50	mg	Approved on	
		In a suspension base		q.s		17.05.2021	
		Approved colour used	•		•	1	
625.	Chlorothalidone Tablets IP	Each uncoated tablet contains:				Approved on	
		Chlorothalidone	IP	50	mg	17.05.2021	
		Excipients		q.s		17.03.2021	
626.	Abacavir Sulphate Tablets IP	Each film coated tablet contains:]	
		AbacavirSulphate	IP			Approved on	
		eq. to Abacavir		300	mg	17.05.2021	
		Excipients		q.s.		17.03.2021	
		Approved color used in film coating					
		I =				, 	
627.	Abacavir Sulphate Oral	Each ml contains:	T		T		
	Solution IP	AbacavirSulphate	IP			Approved on	
		eq. to Abacavir		20	mg	17.05.2021	
		Excipients		q.s			
		Approved colour used					
		F 1.C1 (1.11)				1 1	
(20	Abacavir Sulphate and	Each film coated tablet contains:]	
628.	Lamivudine Tablets IP	AbacavirSulphate	IP				
		eq. to Abacavir		600	mg	Approved on	
		Lamivudine	IP	300	mg	17.05.2021	
		Excipients		q.s.]	
		Approved color used in film coating]	
		•				•	

629.	Abacavir Sulphate,Lamivudine	Each film coated tablet contains:				
	and Zidovudine Tablets IP	AbacavirSulphate	IP			
		eq. to Abacavir		300	mg	
		Lamivudine	IP	150	mg	Approved on 17.05.2021
		Zidovudine	IP	300	mg	17.05.2021
		Excipients		q.s.		•
		Approved color used in film coating		1		
					1	
630.	Acarbose Tablets IP	Each uncoated tablet contains:		Annroyadan		
		Acarbose	IP	25	mg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
631.	Acarbose Tablets IP	Each uncoated tablet contains:				Approved on
		Acarbose	IP	50	mg	17.05.2021
		Excipients		q.s		17.00.2021
		I m				T 1
632.	Acarbose Tablets IP	Each uncoated tablet contains:		100		Approved on
		Acarbose	IP	100	mg	17.05.2021
		Excipients		q.s.		
633.	Acetazolamide Tablets IP	Each uncoated tablet contains:				
055.	Acetazolallide Tablets IF	Acetazolamide	IP	250	ma	Approved on
			IF		mg	17.05.2021
		Excipients		q.s.		
634.	Aciclovir Cream IP	Composition:				
051.	ACICIOVII CICAIII IP	Aciclovir	IP	5%	w/w	Approved on
		In a cream base	- 11	q.s.	VV/ VV	17.05.2021
		III a cream base		4.5.		
635.	Aciclovir Dispersible Tablets	Each uncoated dispersible tablet conta	ins:			
	IP	Aciclovir	IP	200	mg	Approved on
		Excipients		q.s.		17.05.2021
		· ·	1	1		l l
636.	Aciclovir Dispersible Tablets	Each uncoated dispersible tablet conta	ins:			A mm 1
	IP	Aciclovir	IP	400	mg	Approved on 17.05.2021
		Excipients		q.s.		17.03.2021
				•		
637.	Aciclovir Dispersible Tablets	Each uncoated dispersible tablet conta				Approved on
	IP	Aciclovir	IP	800	mg	17.05.2021
		Excipients		q.s.		17.03.2021
638.	Aciclovir Dispersible Tablets					Approved on
	IP	Aciclovir	IP	800	mg	17.05.2021
		Excipients		q.s.		-

639.	Aciclovir Oral Suspension IP	Each 5 ml contains:					
		Aciclovir	IP	200	mg	Approved on	
		In a flavoured syrupy base		q.s.		17.05.2021	
		Approved colour used				1	
						1	
640.	Aciclovir Oral Suspension IP	Each 5 ml contains:					
		Aciclovir	IP	400	mg	Approved on	
		In a flavoured syrupy base		q.s.		17.05.2021	
		Approved colour used]	
641.	Allopurinol Tablets IP	Each uncoated tablet contains:				Approved on	
		Allopurinol	IP	100	mg	17.05.2021	
		Excipients		q.s.		17.03.2021	
642.	Allopurinol Tablets IP	Each uncoated tablet contains:				Approved on	
		Allopurinol	IP	300	mg	17.05.2021	
		Excipients		q.s.		17,00,2021	
						1	
643.	Amiloride Hydrochloride	Each uncoated tablet contains:	T T		1		
	Tablets IP	Amiloride Hydrochloride	IP			Approved on	
		eq. to Anhydrous Amiloride Hydrochloride		5	mg	17.05.2021	
		Excipients Excipients			mg		
		Excipients		q.s.			
644.	Amiloride Hydrochloride	Each uncoated tablet contains:					
	Tablets IP	Amiloride Hydrochloride	IP			-	
		eq. to Anhydrous				Approved on	
		Amiloride Hydrochloride		2.5	mg	17.05.2021	
		Hydrochlorothiazide	IP	25	mg]	
		Excipients		q.s.			
					•		
645.	Amiloride Hydrochloride &	Each uncoated tablet contains:					
	Hydrochlorothiazide Tablets IP	Amiloride Hydrochloride	IP				
		eq. to Anhydrous		_		Approved on	
		Amiloride Hydrochloride		5	mg	17.05.2021	
		Hydrochlorothiazide	IP	50	mg	_	
		Excipients		q.s.			
646.	Amlodipine Besylate and	Each hard gelatin capsule contains:				1	
040.	Benazepril Hydrochloride	Amlodipine Besylate	IP			-	
	Capsules IP	eq. to Amlodipine	IP	2.5		Approved on	
	Capsules II		TD.	2.5	mg	17.05.2021	
		Benazepril Hydrochloride	IP	10	mg	-	
		Excipients		q.s.		1	
		Approved color used in empty					

647.	Amlodipine Besylate and	Each hard gelatin capsule contains:					
	Benazepril Hydrochloride	Amlodipine Besylate	IP			1	
	Capsules IP	eq. to Amlodipine		5	mg	Approved on	
		Benazepril Hydrochloride	IP	20	mg	17.05.2021	
		Excipients		q.s.		1	
		Approved color used in empty		1	7		
						<u> </u>	
648.	S-Amlodipine Besylate Tablets	Tablets Each uncoated tablet contains:					
	IP	S-Amlodipine Besylate	IP			Approved on	
		eq. to S-Amlodipine		1.25	mg	17.05.2021	
		Excipients		q.s.		_	
		Zacipiento		4.5.		<u> </u>	
649.	Ascorbic Acid Tablets IP	Each uncoated chewable tablet contai	ns:				
		Ascorbic Acid	IP	50	mg	Approved on	
		Excipients		q.s.	1	17.05.2021	
650.	Ascorbic Acid Tablets IP	Each uncoated chewable tablet contain	ns:				
	Ascorbic Acid Tablets II	Ascorbic Acid	IP	100	mg	Approved on	
		Excipients		q.s.	1	17.05.2021	
		· ·	1				
651.	Ascorbic Acid Tablets IP	corbic Acid Tablets IP Each uncoated chewable tablet contains:					
		Ascorbic Acid	IP	500	mg	Approved on 17.05.2021	
		Excipients		q.s.		17.05.2021	
652.	Aspirin Gastro- resistant	Each enteric coated (Gastro - resistant	Approved on				
	Tablets IP	Aspirin	IP	75	mg	17.05.2021	
		Excipients		q.s.			
		Approved color used in coating					
	<u> </u>	<u> </u>					
653.	Aspirin Gastro- resistant	Each enteric coated (Gastro - resistant	t) tablet co	ntains:			
	Tablets IP	Aspirin	IP	150	mg	Approved on	
	Tablets IP					17.05.2021	
				q.s.			
		Excipients Approved color used in coating		q.s.		1	
		Excipients		q.s.			
654	Pizatrintan Tablata ID	Excipients Approved color used in coating		q.s.			
654.	Rizatriptan Tablets IP	Excipients Approved color used in coating Each uncoated tablet contains:	ID	q.s.			
654.	Rizatriptan Tablets IP	Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate	IP		ma		
654.	Rizatriptan Tablets IP	Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan	IP	5	mg		
654.	Rizatriptan Tablets IP	Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan Excipients	IP		mg	Approved on	
654.	Rizatriptan Tablets IP	Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan	IP	5	mg	Approved on	
		Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan Excipients Approved colour to be used	IP	5	mg	Approved on	
654.	Rizatriptan Tablets IP Rizatriptan Tablets IP	Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan Excipients Approved colour to be used Each uncoated tablet contains:		5	mg	Approved on 17.05.2021	
		Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan Excipients Approved colour to be used Each uncoated tablet contains: Rizatriptan Benzoate	IP IP	5		Approved on 17.05.2021 Approved on	
		Excipients Approved color used in coating Each uncoated tablet contains: Rizatriptan Benzoate eq. to Rizatriptan Excipients Approved colour to be used Each uncoated tablet contains:		5 q.s	mg	Approved on	

656.	Ebastine Tablets IP	Each film coated tablet contains:					
		Ebastine	IP	10	mg	Approved on	
		Excipients		q.s	1	17.05.2021	
		Approved colour to be used in film co	oating			1	
657.	Ebastine Tablets IP	Each film coated tablet contains:					
		Ebastine	IP	20	mg	Approved on	
		Excipients		q.s		17.05.2021	
		Approved colour to be used in film co	oating				
658.	Nortriptyline Tablets IP	Each film coated tablet contains:	_	1			
		Nortriptyline Hydrochloride	IP			Approved on	
		eq. to Nortriptyline		10	mg	17.05.2021	
		Excipients		q.s		17.03.2021	
		Approved colour to be used in film co	oating]	
- W O						1	
659.	Lamotrigine Sustained release	Each uncoated sustained release table			T	1	
	Tablets IP	Lamotrigine	IP	200	mg	Approved on	
		(As Sustained Release)				17.05.2021	
		Excipients		q.s			
((0)	D 11' C 1 D						
660.	Pregabalin Capsules IP	Each hard gelatin capsule contains:	I ID	50	1	Approved	
		Pregabalin	IP	50	mg	on	
		Excipients	1	q.s		17.05.2021	
		Approved colour to be used in empty	capsule s	shells.			
661.	Pregabalin Capsules IP	Each hard gelatin capsule contains:					
	garanasa - arpanasa - a	Pregabalin	IP	75	mg	Approved	
		Excipients		q.s	1115	on	
		Approved colour to be used in empty	cansule s			17.05.2021	
		ripproved colour to be used in empty	eupsure s				
662.	Pregabalin Capsules IP	Each hard gelatin capsule contains:					
		Pregabalin	IP	150	mg	Approved	
		Excipients		q.s		on	
		Approved colour to be used in empty	capsule s			17.05.2021	
	<u> </u>	1 17				1	
663.	Pregabalin Capsules IP	Each hard gelatin capsule contains:				Δ 1	
		Pregabalin	IP	300	mg	Approved	
		Excipients		q.s		on 17.05.2021	
		Approved colour to be used in empty	capsule s			17.05.2021	
	I						
664.	Gabapentin Capsules IP	Each hard gelatin capsule contains:				Approved	
		Gabapentin	IP	100	mg	Approved	
		Excipients		q.s		on 17.05.2021	
		Approved colour to be used in empty	capsule s	hells.	•	17.03.2021	
		<u> </u>					

665.	Gabapentin Capsules IP	Each hard gelatin capsule contains	:			Approved	
		Gabapentin	IP	300	mg	on	
		Excipients		q.s		17.05.2021	
		Approved colour to be used in emp	pty capsule s	hells.		17.03.2021	
666.	Gabapentin Capsules IP	Each hard gelatin capsule contains					
000.	Gabapentin Capsules II	Gabapentin	IP	400	ma	Approved	
		Excipients	IL		mg	on	
		Approved colour to be used in emp	pty capsule s	q.s hells.		17.05.2021	
		11	1 7 1				
667.	Doxepin Capsules IP	Each hard gelatin capsule contains	:				
		Doxepin Hydrochloride	IP			Approved	
		eq. to Doxepin		25	mg	on	
		Excipients		q.s		17.05.2021	
		Approved colour to be used in emp	pty capsule si	_	1		
						L	
668.	Doxepin Capsules IP	Each hard gelatin capsule contains	:				
		Doxepin Hydrochloride	IP			Approved	
		eq. to Doxepin		50	mg	on	
		Excipients		q.s		17.05.2021	
		Approved colour to be used in emp	pty capsule s	hells.	ľ		
669.	Dovonin Conquies ID	Each hard colotin consula contains				1	
009.	Doxepin Capsules IP	tepin Capsules IP Each hard gelatin capsule contains: Doxepin Hydrochloride IP					
		eq. to Doxepin	11	75	ma	Approved	
		Excipients			mg	on 17.05.2021	
		Approved colour to be used in emp	nty cansule si	q.s hells		17.03.2021	
		ripproved colour to be used in emp	pty cupsuic si				
670.	Oxazepam Tablets IP	Each uncoated tablet contains:				Approved	
	-	Oxazepam	IP	10	mg	on	
		Excipients		q.s		17.05.2021	
671.	Oxazepam Tablets IP	Each uncoated tablet contains:				A 1	
0/1.	Ovarcham ranicis it		IP	15	ma	Approved	
		Oxazepam Excipients	ır		mg	on 17.05.2021	
		Excipients		q.s		17.03.2021	
672.	Oxazepam Tablets IP	Each uncoated tablet contains:				Approved	
		Oxazepam	IP	30	mg	on	
		Excipients		q.s		17.05.2021	
(72	Linia and Tall (ID	F-1					
673	Lisinopril Tablets IP	Each uncoated tablet contains:				Approved	
		Lisinopril	IP	_		on	
		eq. to Anhydrous Lisinopril		5	mg	17.05.2021	
		Excipients		q.s			

674.	Lisinopril Tablets IP	Each uncoated tablet contains:				Approved	
		Lisinopril	IP			on	
		eq. to Anhydrous Lisinopril		10	mg	17.05.2021	
		Excipients		q.s		17.03.2021	
675	V-1	Each film acoust dealer contains				1	
675.	Valsartan &	Each film coated tablet contains:					
	Hydrochlorothiazide Tablets IP	Valsartan	IP	80	mg	Approved	
		Hydrochlorothiazide	IP	12.5	mg	on 17.05.2021	
		Excipients	4:	q.s		17.05.2021	
		Approved colour to be used in film	n coating				
676.	Valsartan &	Each film coated tablet contains:					
	Hydrochlorothiazide Tablets IP	Valsartan	IP	160	mg	Approved	
		Hydrochlorothiazide	IP	12.5	mg	on	
İ		Excipients		q.s		17.05.2021	
		Approved colour to be used in film	n coating		1		
(77	D (' C (' 1 1	Each film coated sustained release	4-1-1-44-:			1	
677.	Paroxetine Sustained- release			ns:	T		
	Tablets IP	Paroxetine Hydrochloride	IP	27.5	***	Approved	
		eq. to Paroxetine		37.5	mg	on 17.05.2021	
		Excipients Approved colour to be used in film	acetina	q.s		17.03.2021	
		Approved colour to be used in film	1 Coating				
678.	Glipizide Tablets IP	Each uncoated tablet contains:				Approved	
		Glipizide	IP	5	mg	on	
		Excipients		q.s		17.05.2021	
679.	Terazosin Tablets IP	Each uncoated tablet contains:	pated tablet contains:				
077.	Terazosiii Tablets II	Terazosin Hydrochloride	IP			Approved	
		eq. to Terazosin	"	1	mg	on	
		Excipients		q.s	1115	17.05.2021	
		1				1	
680.	Terazosin Tablets IP	Each uncoated tablet contains:	1 1		T	Approved	
		Terazosin Hydrochloride	IP			on	
		eq. to Terazosin		2	mg	17.05.2021	
				0.0			
		Excipients		q.s		L	
681.	Terazosin Tablets IP	Excipients Each uncoated tablet contains:		q.s			
681.	Terazosin Tablets IP	Each uncoated tablet contains:	IP	q.s			
681.	Terazosin Tablets IP	Each uncoated tablet contains: Terazosin Hydrochloride	IP	4	mg	on	
681.	Terazosin Tablets IP	Each uncoated tablet contains:	IP	_	mg	on	
		Each uncoated tablet contains: Terazosin Hydrochloride eq. to Terazosin Excipients	IP	4	mg	on	
681.	Terazosin Tablets IP Terazosin Tablets IP	Each uncoated tablet contains: Terazosin Hydrochloride eq. to Terazosin Excipients Each uncoated tablet contains:		4	mg	on 17.05.2021	
		Each uncoated tablet contains: Terazosin Hydrochloride eq. to Terazosin Excipients	IP IP	4	mg	- Approved on 17.05.2021 - Approved on	

683.	Acarbose Tablets IP	Each uncoated tablet contains:		Approved			
		Acarbose	IP	25	mg	on	
		Excipients		q.s		17.05.2021	
684.	Acarbose Tablets IP	Each uncoated tablet contains:			•	Approved	
		Acarbose	IP	50	mg	on	
		Excipients		q.s		17.05.2021	
605		IB 1					
685.	Acarbose Tablets IP	Each uncoated tablet contains:	I ID	100	1	Approved	
		Acarbose	IP	100	mg	on 17.05.2021	
		Excipients		q.s		17.05.2021	
686.	Pregabalin(SR)&	Each Film Coated tablets contains:					
000.	Methylcobalamin Tablets	Pregabalin (SR)	IP	100	mg	Approved	
	Trioniy leoculumini Tuestets	Methylcobalamin	IP	1500	mcg	on	
		Excipients		q.s	lineg	17.05.2021	
		Colour:Approved colour used		4.5			
		Colour approved colour asea					
687.	Acebrophylline Syrup	Each ml contains:					
		Acebrophylline		10	mg	Approved	
		In flavoured syrupy base		q.s	-	on	
		Colour :approved colour used			_L	17.05.2021	
688.	Citicoline syrup	Each 5 ml contains:					
		Citicoline Sodium	IP			Approved	
		eq to Citicoline		500	mg	on	
		In flavoured syrupy base		q.s		17.05.2021	
		Colour :approved colour used					
100							
689.	Atorvastatin Calcium &	Each film coated tablets contains :					
	EzetimibeTablets	Atorvastatin Calcium	IP	20		Approved	
		eq to Atorvastatin	TD.	20	mg	on	
		Ezetimibe	IP	10	mg	17.05.2021	
		Excipients		q.s			
		Colour :approved colour used					
690.	Dried Aluminium Hydroxide	Each 5 ml contains :					
090.	Gel, Magnesium Hydroxide	Dried Aluminium Hydroxide Gel	IP	600	ma	-	
	Oxetacain & Simethicone	Magnesium Hydroxide Magnesium Hydroxide	IP	300	mg mg	Approved	
	suspension	Simethicone	IP	25	mg	on	
		Oxetacain	BP	10	mg	17.05.2021	
		In flavoured syrupy base	DI	q.s	IIIg	17.03.2021	
		Colour :approved colour used		4.5			
		Colour .approved colour used					

691.	Propranolol Hcl SR Capsules	Each hard gelatin capsule contains:				Approved
		Propranolol Hydrochloride (SR)	IP	80	mg	on
		Excipients		q.s		17.05.2021
		Colour:Approved colour used in em	pty capsule	shell	•	17.03.2021
						1
692.	Clindamycin Hcl Capsules IP	Each hard gelatin capsule contains:				
		Clindamycin Hcl	IP			Approved
		Eq to Clindamycin		80	mg	on
		Excipients		q.s		on 17.05.2021 Approved on 17.05.2021
		Colour:Approved colour used in em	pty capsule	shell		
693.	Aceclofenac &	Each film coated tablets contains :				
	Tizanidine Tablets	Aceclofenac	IP	200	mg	Annroved
		Tizanidine Hcl	IP			
		Eq to Tizanidine		2	mg	on
		Excipients		q.s		17.03.2021
		Colour :approved colour used				
		T.				
694.	Dexrabeprazole (EC) &	Each hard gelatin capsule contains:			_	
	Domperidone (SR) Capsules	Dexrabeprazole		10	mg	
		(as enteric coated pellets)				Approved on 17.05.2021
		Domperidone	IP	30	mg	
		(sustained release pellets)				17.05.2021
		Excipients		q.s		
		Colour:Approved colour used in em	pty capsule	shell		
605			tains:			
695.	Eperisone Hcl Tablets	Each film coated tablet contains:				
695.	Eperisone Hcl Tablets	Each film coated tablet contains: Eperisone Hcl	IP	50	mg	
695.	Eperisone Hcl Tablets		IP		mg	on
695.	Eperisone Hcl Tablets	Eperisone Hcl		50 q.s	mg	- Approved - on - 17.05.2021
		Eperisone Hcl Excipients Approved colour to be used in film	coating		mg	on
695. 696.	Dexrabeprazole Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain	coating	q.s		on 17.05.2021
		Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium	coating	q.s 5	mg	on
	Dexrabeprazole Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients	coating	q.s		on 17.05.2021 Approved
	Dexrabeprazole Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium	coating	q.s 5		on 17.05.2021 Approved on
	Dexrabeprazole Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients	coating	q.s 5		on 17.05.2021 Approved on
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains:	coating	q.s 5 q.s	mg	on 17.05.2021 Approved on 17.05.2021
696.	Dexrabeprazole Sodium Tablets	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam	coating as:	q.s 5 q.s		on 17.05.2021 - Approved on 17.05.2021
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium	coating	q.s 5 q.s	mg	on 17.05.2021 Approved on 17.05.2021 Approved on
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium eq to Citicoline	coating as:	q.s 5 q.s 400 500	mg	on 17.05.2021 Approved on 17.05.2021 Approved on
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium eq to Citicoline In flavoured syrupy base	coating as:	q.s 5 q.s	mg	on 17.05.2021 Approved on 17.05.2021 Approved on
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium eq to Citicoline	coating as:	q.s 5 q.s 400 500	mg	on 17.05.2021 Approved on 17.05.2021 Approved on
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium Syrup	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium eq to Citicoline In flavoured syrupy base	coating as:	q.s 5 q.s 400 500	mg	on 17.05.2021 - Approved on 17.05.2021 - Approved on 17.05.2021
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium eq to Citicoline In flavoured syrupy base Colour :approved colour used Each 5 ml contains:	coating is: IP IP IP	q.s 5 q.s 400 500 q.s	mg mg mg	on 17.05.2021 - Approved on 17.05.2021 - Approved on 17.05.2021
696.	Dexrabeprazole Sodium Tablets Piracetam & Citicoline Sodium Syrup	Eperisone Hcl Excipients Approved colour to be used in film Each Enteric Coated Tablets contain Dexrabeprazole Sodium Excipients Colour:Approved colour used Each 5 ml contains: Piracetam Citicoline Sodium eq to Citicoline In flavoured syrupy base Colour :approved colour used	coating as:	q.s 5 q.s 400 500	mg	on 17.05.2021 Approved on 17.05.2021

699.	Betahistine Hcl (SR) Tablets	Each Sustained Release uncoarted ta	ablets contai	ns:		
		Betahistine Hcl	IP	48	mg	Approved
		(As Sustained Release)				on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				
700	Diacerin & Aceclofenac tablets	Each film coated tablets contains:				
		Diacerin	IP	50	mg	Approved
		Aceclofenac	IP	100	mg	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				
=0.1	I					1
701.	Bromhexine Hydrochloride	Each 5 ml contains :			T	Approved
	Syrup	Bromhexine Hydrochloride	IP	4	mg	on
		In flavoured syrupy base		q.s		17.05.2021
		Colour :approved colour used				
702.	Sodium picosulphate Tablets	Each uncoated tablets contains :				
	Sourain preosurphate Tublets	Sodium picosulphate	BP	10	mg	Approved
		Excipients	DI	q.s	1115	on
		Colour :approved colour used		4.5		17.05.2021
	<u> </u>	Colour improved colour used				
703.	Acebrophylline SR Tablets	ets Each film coated sustained release tablet contains:				
		Acebrophylline	BP	200	mg	Approved
		(as sustained Relase)				on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				
704.	Acebrophylline ER Tablets	Each film coated Extended release ta	ahlet contain	16.		
701.	Ticorophymic Ex Tuolous	Acebrophylline	BP	400	mg	Approved
		(as extended Relase)		100	1115	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				
705.	Tranexamic Acid &	Each film coated tablets contains :				
	Ethamsylate Tablets	Tranexamic Acid	IP	250	mg	Approved
		Ethamsylate	IP	250	mg	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				
706.	Tranexamic Acid SR Tablets	Each film coated Sustained release	Cableta Cont	oine:		1
700.	Tranexamic Acid SK Tablets			750	ma	Approved
		Tranexamic Acid	IP		mg	on
		Excipients Colour conproved colour used		q.s		17.05.2021
		Colour :approved colour used				

707.	Lacosamide Tablets	Each flim coated Tablets Contains:				Ammovod
		Lacosamide		100	mg	Approved
		Excipients		q.s		on 17.05.2021
		Colour :approved colour used				17.05.2021
						l
708.	Lacosamide Tablets	Each flim coated Tablets Contains:			Approved	
		Lacosamide		150	mg	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				17.03.2021
709.	Lacosamide Tablets	Each flim coated Tablets Contains:				Approved
		Lacosamide		200	mg	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				17.03.2021
710.	Levocetirizine	Each 5 ml contains:				
	Dihydrochloride Syrup	Levocetirizine				Approved
	(for the use in children 2 years	Dihydrochloride	IP	2.5	mg	on
	and above	In flavoured syrupy base		q.s		17.05.2021
		Colour :approved colour used]
711.	Propranolol Hcl SR Tablets	Each film coated Sustained release T	ablets Con	ıtains:		Approved
		Propranolol Hyrochloride	IP	80	mg	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				17.03.2021
712.	Propranolol Hcl SR Tablets	Each film coated Sustained release T	ablets Con	itains:	_	Approved
		Propranolol Hyrochloride	IP	160	mg	on
		Excipients		q.s		17.05.2021
		Colour :approved colour used				17.03.2021
713.	Ursodeoxycholic Acid SR	Each hard gelatin capsule contains:		•		
	Capsules	Ursodeoxycholic Acid	IP	450	mg	Approved
		(sustained release Fome)				on
		Excipients		q.s		17.05.2021
		Colour:Approved colour used in em	pty capsule	e shell		
						1
714.	Diacerin Capsules	Each hard gelatin capsule contains:	r	1		Approved
		Diacerin	IP	50	mg	on
		Excipients		q.s		17.05.2021
		Colour:Approved colour used in em	pty capsule	e shell		
715	TT 1 1 1 A 11	Early 5 and a marks				1
715.	Ursodeoxycholic Acid	Each 5 ml contains :	m	107	1	Approved
	Suspension	Ursodeoxycholic Acid	IP	125	mg	on
		In flavoured suspension base		q.s		17.05.2021
Colour :approved colour used						1

716.	Lacosamide Tablets	Each flim coated Tablets Contains	:	1		Approved	
		Lacosamide		50	mg	on	
		Excipients		q.s		17.05.202	
		Colour :approved colour used					
717.	Loxapine Oral Concentration	Each ml Contains:					
, 1 , .	Solution	Loxapine Succinate	USP			Approved	
	201411011	Eq to Loxapine		25	mg	on	
		In a flavoured base		q.s	1115	17.05.202	
		Colour :approved colour used		1"			
718.	Eperisone Hydrochloride	Each Hard Gelatin capsule contain	is:	T		Approved	
	Sustained Release Capsule	Eperisone Hydrochloride		75	mg	on	
		In a flavoured base		q.s		17.05.202	
		Colour :approved colour used					
719.	Eperisone Hydrochloride	Each Hard Gelatin capsule contain	ıs:				
	Sustained Release Capsule	Eperisone Hydrochloride		150	mg	Approved	
		In a flavoured base		q.s		on	
		Colour :approved colour used	I	1		17.05.202	
720	Mathylachalamin Fach Hard Calatin capsula contains:						
720.	Methylcobalamin	Each Hard Gelatin capsule contain		1500	1		
	Benfotiamine,	Methylcobalamin	IP	1500	mcg	Approved on	
	Alpha Lipoic Acid, Vitamin	Alpha Lipoic acid	IP	200	mg		
	B6,Biotin & Folic Acid						
		Benfotiamine		50	mg		
	Capsules	W. D.	TIOD	2			
		Vitamin B6	USP	3	mg	17.05.202	
		Biotin	USP	5	mg	17.05.202	
		Folic Acid	IP	1.5	mg		
		Excipients Colour removed colour read		q.s			
		Colour :approved colour used					
721.	Betahistine Dihydrochloride	Each Sustained Release tablets con	ntains :				
	SR Tablets	Betahistine Dihydrochloride	IP	24	mg	Approved	
		Excipients		q.s		on 17.05.202	
		Colour :approved colour used	•			17.03.202	
722.	Betahistine Dihydrochloride	Each Sustained Release tablets con	ntains ·				
144.	SR Tablets	Betahistine Dihydrochloride	IP	32	mg	Approved	
	211 1401010	Excipients	11	q.s	****5	on	
		Colour :approved colour used		1. ~		17.05.202	
702		ID 16 1					
723.	Levocloperastine Fendizoate	Each 5 ml contains:		25.4	1		
	Suspension	Levocloperastine Fendizoate		35.4	mg	Approved	
		Eq to Levocloperastine Hcl		20	mg	on	
		In a suspension base		q.s		17.05.2021	

	Colour :approved colour used	

724.	Activated Dimethicone,	Each 5 ml contains:						
	Magnesium Hydroxide, Dried	Activated Dimethicone	IP	50	mg			
	Aluminium Hydroxide,Sorbitol	Magnesium Hydroxide	IP	250	mg	1		
	suspension	Dried Aluminium Hydroxide	IP	250	mg	Approved on		
		Sorbitol(70%)	IP	1.25	mg	17.05.2021		
		Excipients		q.s				
		Approved colour used	<u> </u>	•	1			
725.	Diacerin & Glucosamine	Each Film Coated Tablets Contains	s:					
	Tablets	Diacerin	IP	50	mg	1		
		Glucosamine Sulphate		500	mg	Approved on 17.05.2021		
		Excipients		q.s		17.03.2021		
		Approved colour used			•			
726.	Norfloxacin & Tinidazole	Each Film Coated Tablets Contains	:					
	Tablets	Norfloxacin	IP	400	mg	Annroyadan		
		Tinidazole	IP	600	mg	Approved on 17.05.2021		
		Excipients		q.s		17.03.2021		
		Approved colour used						
727.	Fluconazole Tablets IP	Each Uncoated Tablets Contains:						
		Fluconazole	IP	400	mg	Approved on 17.05.2021		
		Excipients		q.s				
		Approved colour used						
728.	Fluoxetine Capsules IP	Each Hard gelatin capsules contain			•			
		Fluoxetine Hydrochloride	IP			Approved on		
		Eq to fluoxetine		60	mg	17.05.2021		
		Excipients		q.s				
		Approved colour used						
729.	Rosuvastatin & Fenofibrate	Each Film Coated Tablets Contains						
	Tablets	Rosuvastatin Calcium	IP					
		Eq toRosuvastatin		10	mg	Approved on		
		Fenofibrate	BP	160	mg	17.05.2021		
		Excipients		q.s				
		Approved colour used						
720						 _		
730.	Amlodipine & Telmisartan	Each Film Coated Tablets Contains						
	Tablets	Amlodipine Besilate	IP	10				
		Eq. to Amlodipine		10	mg	Approved on		
		Telmisartan Hydrochloride	IP	40	mg	17.05.2021		
		Excipients		q.s				
		Approved colour used						

731.	Lornoxicam &	Each Film Coated Tablets Contains	:			
	Thiocholchicoside Tablets	Lornoxicam	IP	8	mg	Annroyadan
		Thiocholchicoside	IP	4	mg	- Approved on - 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used	1			
						T
732.	Amlodipine & Telmisartan	Each Film Coated Tablets Contains				
	Tablets	Amlodipine Besilate	IP			
		Eq. to Amlodipine		5	mg	
		Telmisartan Hydrochloride	IP	80	mg	17.05.2021
		Excipients		q.s		
		Approved colour used				
700	G G					T
733.	Sparfloxacin Tablets	Each film coated tablets contains:	T T	• • • •	1	
		Sparfloxacin		200	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
734.	Tolperisone tablets	Each film coated tablets contains :				
		Tolperisone Hcl		150	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used			II.	_
	<u> </u>					1
735.	Nimodipine Tablets	Each film coated tablets contains:				
		Nimodipine	BP	30	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	,		•	
						I
736.	Omeprazole Magnesium	Each Enteric Coated tablets contain			1	
	Tablets	Omeprazole Magnesium	BP			Approved on
		Eq. to omeprazole		10	mg	17.05.2021
		Excipients		q.s		17,00,2021
		Approved colour used				
737.	Amlodipine, Olmesartan	Each Film Coated Tablets Contains	·•			
131.	Medoxomil &	Amlodipine Besilate	IP			+
	Hydrochlorothiazide	Eq. to Amlodipine	11	10	ma	
	Tablet	Olmesartan Medoxomil	IP	40	mg	Approved on
	Taulet		IP		mg	17.05.2021
		Hydrochlorothiazide	IP	25	mg	
		Excipients Approved colour year		q.s		_
		Approved colour used				

738.	Alfuzocin Hcl. (ER)	Each Film Coated Tablets Contains	s:			
	& Dutasteride tablet	Alfuzocin Hcl (ER)	IP	10	mg	1
		Dutasteride	BP	0.5	mg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used	'		1	
739.	I 1 ' ' 1 77 11 4					<u> </u>
139.	Levosulpiride Tablets	Each uncoated tablets contains:	- DD	25	<u> </u>	1
		Levosulpiride	BP	25	mg	Approved on 17.05.2021
		Excipients		q.s		17.05.2021
		Approved colour used				
740.	Levosulpiride Tablets	Each uncoated tablets contains:				
		Levosulpiride	BP	75	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used			I	
						1
741.	Levosulpiride Tablets	Each uncoated tablets contains:				
		Levosulpiride	BP	150	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
	T					
742.	Levosulpiride Tablets	Each uncoated tablets contains:			1	
		Levosulpiride	BP	200	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
743.	Allopurinol SR Capsules	Each hard gelatin sustained release	capsules cor	ntains		
		Allopurinol	IP	250	mg	1
		(as sustain release)				Approved on
						17.05.2021
		Excipients		q.s		17.05.2021
		Excipients Approved colour used in empty ca	psule shell	q.s		_
		Approved colour used in empty car	psule shell	q.s		
744.	Primaquine Tablets IP	Approved colour used in empty car Each film coated tablets contains:	-	q.s		
744.	Primaquine Tablets IP	Approved colour used in empty car Each film coated tablets contains: Primaquine Phosphate	psule shell	q.s		Approved on
744.	Primaquine Tablets IP	Approved colour used in empty cap Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine	-	q.s 45	mg	Approved on 17.05.2021
744.	Primaquine Tablets IP	Approved colour used in empty car Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine Excipients	-	-	mg	Approved on 17.05.2021
744.	Primaquine Tablets IP	Approved colour used in empty cap Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine	-	45	mg	
		Approved colour used in empty car Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine Excipients Approved colour used	IP	45 q.s	mg	
744.	Primaquine Tablets IP Salbutamol SR tablets	Approved colour used in empty car Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine Excipients Approved colour used Each film coated sustained release	IP tablets contain	45 q.s	mg	
		Approved colour used in empty car Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine Excipients Approved colour used Each film coated sustained release Salbutamol Sulphate	IP	45 q.s ins:		17.05.2021 Approved on
		Approved colour used in empty car Each film coated tablets contains: Primaquine Phosphate Eq. to Primaquine Excipients Approved colour used Each film coated sustained release	IP tablets contain	45 q.s	mg	17.05.2021

746.	Salbutamol tablets	Each film coated tablets contains	S:					
		Salbutamol Sulphate	BP			A		
		Eq.to Salbutamol		4	mg	Approved on 17.05.2021		
		Excipients		q.s		17.03.2021		
		Approved colour used	1					
747.	Acamprosate Calcium (EC)							
	Tablets	Acamprosate Calcium	IP	333	mg	Approved on		
		Eq. to Acamprosate	IP	330	mg			
		Excipients		q.s		17.03.2021		
		Approved colour used				Approved on 17.05.2021 Approved on 17.05.2021 Approved on 17.05.2021 Approved on 17.05.2021		
748.	RifaximinTablets	Each film coated tablets contains			T			
		Rifaximin	BP	550	mg			
		Excipients		q.s		17.05.2021		
		Approved colour used						
740	D'C ' ' T 11 '	T 1 6'1						
749.	RifaximinTablets	Each film coated tablets contains		200				
		Rifaximin	BP	200	mg			
		Excipients		q.s		17.05.2021		
		Approved colour used						
750.	RifaximinTablets	Each film coated tablets contains	·					
, 5 0.		Rifaximin	BP	400	mg	Approved on		
		Excipients		q.s	8			
		Approved colour used		1		-		
751.	RifaximinTablets	Each uncoated dispersible Table	ts Contains :					
		Rifaximin	BP	200	mg	Approved on		
		Excipients		q.s		17.05.2021		
		Approved colour used	1					
752.	Ketoprofen &	Each Film Coated Tablets Conta						
	Thiocholchicside Tablets	Ketoprofen	IP	50	mg	Approved on		
		Thiocholchicoside	IP	4	mg	17.05.2021		
		Excipients		q.s		17.03.2021		
		Approved colour used						
753.	Vildagliptin & Metformin Hcl	Each uncoated bilayred (SR) tab	lets contains:			_[
	SR Tablets	Vildagliptin	IP	50	Mg	Approved on		
		Metfortin Hcl(as SR)	IP	850	Mg	17.05.2021		
		Excipients		q.s		17.05.2021		
		Approved colour used						

754.	Vildagliptin & Metformin Hcl	Each uncoated bilayred (SR) tablets co	ntains:			
	SR Tablets	Vildagliptin	IP	50	Mg	Approved on
		Metfortin Hcl(as SR)	IP	500	Mg	17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used				
755.	SilodosinTablets	Each film coated tablets Contains :		T	r	
		Silodosin		4	Mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used				
756.	Sevelamer Carbonate Tablets	Each film coated tablets Contains :				
		Sevelamer Carbonate		400	Mg	Approved on
		Excipients		q.s	8	17.05.2021
		Approved colour used		1	l .	17.05.2021
<u> </u>	L					
757.	Sevelamer Carbonate Tablets	Each film coated tablets Contains:				
		Sevelamer Carbonate		800	Mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	•		•]
758.	Alfuzocin Prolonged released	Each prolonged release film coated tab				Approved on
	Tablets BP	Alfuzocin	IP	10	Mg	17.05.2021
		Excipients		q.s		
		Approved coloured used				
759.	Teneligliptin hydrobromide	Each film coated tablet contains:				
137.	Hydrate Tablets	Teneligliptin hydrobromide Hydrate				Approved on
	Trydrate Tublets	eq to Teneligliptin		20	Mg	17.05.2021
		eq to renengipuii		20	1415	17.00.2021
		Excipients		as		
		Excipients Approved coloured used		q.s		
		Excipients Approved coloured used		q.s		
760.	Paracetamol (SR) Tablets		contain			
760.	Paracetamol (SR) Tablets	Approved coloured used	contain		mg	Approved on
760.	Paracetamol (SR) Tablets	Approved coloured used Each uncoated sustained release tablet Paracetamol (as sustained release)		s:	mg	Approved on 17.05.2021
760.	Paracetamol (SR) Tablets	Approved coloured used Each uncoated sustained release tablet Paracetamol		s:	mg	
	` '	Approved coloured used Each uncoated sustained release tablet Paracetamol (as sustained release) Excipients		s: 650	mg	
760. 761.	Paracetamol (SR) Tablets Tapentadol HCl Tablets	Approved coloured used Each uncoated sustained release tablet Paracetamol (as sustained release) Excipients Each film coated tablet contains:	IP	s: 650	mg	17.05.2021
	` '	Approved coloured used Each uncoated sustained release tablet Paracetamol (as sustained release) Excipients Each film coated tablet contains: Tapentadol Hydrochloride		s: 650		
	` '	Approved coloured used Each uncoated sustained release tablet Paracetamol (as sustained release) Excipients Each film coated tablet contains:	IP	s: 650 q.s	mg	17.05.2021 Approved on

762.	Tapentadol HCl Tablets	Each film coated tablet contains:				
		Tapentadol Hydrochloride	IP			Approved on
		eq. to Tapentadol	-	150	mg	17.05.2021
		Excipients		q.s		
		Approved coloured used	I		1	
763.	Sitagliptin Phosphate	Each film coated tablet contains:				
	& Metformin HCl Tablets	Sitagliptin Phosphate	IP			Approved on
		eq. to Sitagliptin		50	mg	17.05.2021
		Metformin Hydrochloride	IP	500	mg	
		Excipients		q.s		
		Approved coloured used				
764.	Sitagliptin Phosphate	Each film coated tablet contains:				
	& Metformin HCl Tablets	Sitagliptin Phosphate	IP			A 1
		eq. to Sitagliptin		50	mg	Approved on 17.05.2021
		Metformin Hydrochloride	IP	1000	mg	17.05.2021
		Excipients		q.s		
		Approved coloured used				
765.	Methylcobalamin, Alpha	Each hard gelatin capsule contains :	: 	Т	Т	
	Lipoic Acid, Benfotiamine, Vitamin B6, Calcium	Methylcobalamin	IP	750	mcg	
	Pantothenate, Nicotinamide	Alpha Lipoic Acid	USP	100	mg	
	& Folic Acid Capsules	Benfotiamine		7.5	mg	Approved on
	•	Vitamin B6	IP	1.5	mg	17.05.2021
		Calcium Pantothenate	IP	25	mg	17.03.2021
		Nicotinamide	IP	50	mg	
		Folic Acid	IP	0.75	mg	
		Excipients		q.s		1
		Approved coloured used	-		'	
766.	Tolperisone HCl & Etodolac	Each film coated tablet contains:				
	Tablets	Tolperisone Hydrochloride		150	mg	Approved on
		Etodolac	IP	400	mg	17.05.2021
		Excipients		q.s		
		Approved coloured used				
767.	Torsemide & Amiloride HCl	Each uncoated tablet contains:				
, 5,.	Tablets	Torsemide	IP	10	ma	
		Amiloride Hydrochloride	IP	10	mg	Approved on
		eq. to. Anhydrous Amiloride	11			17.05.2021
		Hydrochloride		5	mg	
		Excipients		q.s]

	Tablets	G:				
		Cinnarizine	IP	20	mg	Approved on
		Dimenhydrinate	BP	40	mg	17.05.2021
		Excipients		q.s		
		Approved coloured used				
7.00	C' '' 0 D' 1 1' '	F 1 C1				
769.	Cinnarizine & Dimenhydrinate	Each film coated tablet contains:		20	1	. 1
	Tablets	Cinnarizine	IP	20	mg	Approved on
		Dimenhydrinate	BP	40	mg	17.05.2021
		Excipients		q.s		
		Approved coloured used				
770.	Glimepiride,	Each uncoated bilayered tablet co	nntains:			
,,0.	Atorvastatin & Metformin HCl	Glimepiride	IP I	1	mg	
	(SR) Tablets	Atorvastatin Calcium	IP		III'S	Approved on
	(210) 140100	eq. to. Atorvastatin	11	10	ma	Approved on
		Metformin Hydrochloride	IP	500	mg mg	17.05.2021
		(as sustained release)	1	300	mg	
		Excipients Excipients		q.s		17.05.2021
		Excipions		4.5		
771.	Glimepiride,	Each uncoated bilayered tablet co	ntains:			
	Atorvastatin & Metformin HCl	Glimepiride	IP	2	mg	
	(SR) Tablets	Atorvastatin Calcium	IP			A 1
		eq. to. Atorvastatin		10	mg	Approved on 17.05.2021
		Metformin Hydrochloride	IP	500	mg	17.05.2021
		(as sustained release)				
		Excipients		q.s		
	,				•	
772.	Pregabalin &	Each hard gelatin capsule contain	ns:			
	Methylcobalamin capsules IP	Pregabalin	IP	75	mg	Approved on
		Methylcobalamin	IP	500	mcg	Approved on 17.05.2021
		Excipients		q.s		17.03.2021
		Approved colour used in empty	capsule shell			
773.	Pregabalin capsules IP	Each hard gelatin capsule contains	•			
113.	1 regavanni capsules n	Pregabalin	IP	150	mg	A 1
		Excipients	11		mg	Approved on 17.05.2021
		Approved colour used in empty ca	uncula chall	q.s		17.03.2021
		Approved colour used in empty ca	ipsuie sileii			
774.	Pregabalin capsules IP	Each hard gelatin capsule contains	:			
		Pregabalin	IP	300	mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used in empty ca	nsule shell	1		

775.	Ambroxol & Cetirizine Hcl	Each 5ml contains :-				
	Syrup	Ambroxol	IP	30	mg	1
		Cetirizine Hcl	IP	10	mg	Annroyadan
		Excipients		q.s		Approved on 17.05.2021
		Flavoured Syrupy base		•		17.05.2021
		Approved colour used				1
		Tipproved colour used				
776.	Ambroxol, Guaiphenesin,	Each 5ml contains :-				
770.	Terbutaline Sulphate &	Ambroxol HCL	IP	30	ma	-
		Guaiphenesin	IP	100	mg mg	1
	Menthol Syrup	Terbutaline Sulphate	IP	2.5	mg	1
		Menthol	IP	2.5	mg	Approved on
		Excipients		q.s		17.05.2021
		Flavoured Syrupy base	<u> </u>			
		Approved colour used				-
777.	Mecobalamin, Pyridoxine,	Each film coated tablet contains :				
	Nicotinamide,& Benfotiamine	Mecobalamin	IP	750	mcg	1
	Tablets	Pyridoxine	IP	1.5	mg	Approved or
		Nicotinamide	IP	45	mg	17.05.2021
		Benfotiamine		150	mg	17.03.2021
		Excipients		q.s		_
		Approved coloured used				
778.	Gabapentin &	Each film coated tablet contains :				
	Methylcobalamin Tablets	Gabapentin	IP	600	mg	1
		(As Sustained Release)				Approved on
		Methylcobalamin	IP	1500	mcg	17.05.2021
		Excipients				17.03.2021
		Approved colour used		q.s		1
		Approved colour used				
779.	Glibenclamide & Metformin	Each uncoated bilayred tablets c	contains:			
	Hydrochloride (SR) Tablets	Glibenclamide	IP	5	Mg	A
		Metfortin Hcl(as SR)	IP	500	Mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used		1		
700	G 0.D	E-1 Cl 1 11 1				T
780.	Cinnarizine & Domperidone	Each film coated tablet contains			1	
	Tablets	Cinnarizine	IP	20	mg	Approved on
		Domperidone	IP	15	mg	17.05.2021
		Excipients		q.s		
		Approved coloured used				
781.	Nitrofurantoin (S R) Tablets	Each uncoated sustained realese	tablet contains:			_
701.	Theorem (D K) Tables			100	ma	Approved
		Nitrofurantoin	IP		mg	on
		Excipients		q.s		17.05.2021
		Approved colour used.				1

782.	Glibenclamide & Metformin	Each uncoated bilayred tablets con-				
	Hydrochloride (SR) Tablets	Glibenclamide	IP	5	Mg	A managed an
		Metfortin Hcl(as SR)	IP	850	Mg	Approved on
		Excipients		q.s		17.05.2021
		Approved colour used	L	_	_1	
						l
783.	Doxofylline Syrup	Each 1 ml contains:			_	
		Doxofylline		20	mg	Approved
		Excipients		q.s		on
		In a flavoured Syrupy base				17.05.2021
		Approved colour used.				
784.	Piracetam & Citicoline Sodium	Each film coated tablet contains:				
704.	Tablets	Piracetam	IP	400	mg	-
	Tablets	Citicoline Sodium	IP	400	mg	Approved
		eq to Citicoline	11	500	ma	on
		Excipients			mg	17.05.2021
		Colour :approved colour used		q.s		-
		Colour .approved colour used				
785.	Beclomethasone Dipropinate	Composition:				
	& Neomycin Sulphate cream	Beclomethasone Dipropinate	IP	0.025%	W/W	Approved on
	, ,	Neomycin Sulphate	IP	0.5%	W/W	31.05.2021
		In a gel base		q.s		
			I			
786.	Halobetasol Propionate &	Composition:				
	Salicylic Acid ointment	Halobetasol Propionate		0.5	mg	Approved on
		Salicylic Acid	IP	30	mg	31.05.2021
		In a Cream base		q.s		
			•	•		
787.	Halobetasol Propionate &	Composition:				
	Salicylic Acid ointment	Halobetasol Propionate		0.05%	W/W	Approved on
		Salicylic Acid	IP	6.00%	W/W	31.05.2021
		In a Cream base		q.s		
788.	Minoxidil Topical lotion	Composition:				
700.	Willioxidii Topicai Iodoli	Minoxidil Topical	IP	20	Mg	Approved on
		In a lotion base	- 1		IVIg	31.05.2021
		In a rotton oase		q.s		<u> </u>
789.	Minoxidil Topical solution	Each ml contains:				Approved on
		Minoxidil Topical	IP	5.00%	W/W	Approved on
		In a base		q.s		31.05.2021
7 00	B 11 A 11 O1					
790.	Povidone Iodine Ointment	Composition:		1007	******	Approved on
		Povidone Iodine	IP	10%	W/W	31.05.2021
		In a Ointment base		q.s		

791.	Sucralfate, Metronidazole &	Composition:				
	Povidone Iodine Ointment	Sucralfate	IP	70	mg	, ,
		Metronidazole	IP	10	mg	Approved on
		Povidone Iodine	IP	50	mg	31.05.2021
		In a Ointment base		q.s		
			l	1	I .	
792.	Tacrolimus Ointment	Composition:				A managed on
		Tacrolimus	IP	0.03%	w/w	Approved on
		In a Ointment base		q.s		31.05.2021
			•	•	•	
793.	Thiocolchicosode, Methyl	Composition:		_		
	Salicylate, Menthol & Absolute	Thiocolchicosode	IP	2.5	mg	
	Alcohol Gel	Methyl Salicylate,	IP	100	mg	Approved on
		Menthol	IP	50	mg	31.05.2021
		Absolute Alcohol		50	mg]
		In a Gel base		q.s		
						r
794.		Each hard gelatin capsule contains:		.		
	Clindamycin Capsules IP	Clindamycin hydrochloride	IP			A
		Eq. to Clindamycin		300	Mg	Approved on
		Excipients		q.s		31.05.2021
		Approved colour used in empty caps	ule shell		•	
						<u> </u>
795.	Benzoyl Peroxide gel	Composition:				
		hydrous benzoyl peroxide	IP			Approved on
		eq.to anhydrous benzoyl peroxide		10.0%	W/W	31.05.2021
		In a gel base		q.s		
						T
796.	Benzoyl Peroxide Gel	Composition:		T		
		hydrous benzoyl peroxide	IP			Approved on
		eq.to anhydrous benzoyl peroxide		2.50%	W/W	31.05.2021
		In a gel base		q.s		
705	D 1D 11 C	la				
797.	Benzoyl Peroxide Gel	Composition:	T -	T	_	_
		hydrous benzoyl peroxide	IP			Approved on
		eq.to anhydrous benzoyl peroxide		5.00%	W/W	31.05.2021
		In a gel base		q.s		
700	Disclass 0 1' 0 1' 1	Each film coated Sustained Release	tablet see	toing		<u> </u>
798.	Divalproex Sodium Sustained Release Tablets IP	Divalproex Sodium	IP IP	tailis.	T	
	ixelease Lablets II	Eq. to Valproic Acid	"	500	mg	Approved on
		Excipients		q.s	1115	31.05.2021
		Approved colour used		1 7.~		
799.	Nortriptyline Tablets IP	Each film coated tablet contains:				
		Nortriptyline Hydrochloride	IP	4.0		Approved on
		Eq. to Nortriptyline		10	mg	31.05.2021
		Excipients		q.s		

800.	Nortriptyline Tablets IP	Each film coated tablet contains:				
		Nortriptyline Hydrochloride	IP			Approved on
		Eq. to Nortriptyline		25	mg	31.05.2021
		Excipients		q.s		
801.	Etodolac & Thiocolchicoside	Each film coated tablet contains:	, ,			
	Tablets	Etodolac	IP	400	mg	Approved on
		Thiocolchicoside	IP	8	mg	31.05.2021
		Excipients		q.s		
902	I 114 1 C114-	Each 5ml contains:				Τ
802.	Levosalbutamol Sulphate, Ambroxol Hcl & Guaiphenesin	Levosalbutamol Sulphate	IP		1	-
	Syrup	Eq. to Levosalbutamol	IF	1	mg	
	Sylup	Ambroxol Hcl	IP	30	mg	
		Guaiphenesin	IP	50	mg	Approved on
		Excipients	11	q.s	mg	31.05.2021
		In a flavoured syrupy base		4.5		1
		Approved colours used				•
	<u> </u>	1				
803.	Diacerin & Aceclofenac	Each film coated tablet contains:				
	Tablets	Diacerin	IP	50	mg	
		Aceclofenac	IP	100	mg	Approved on
		Excipients		q.s		31.05.2021
		Approved colour used		1		
		FF				
804.	Moxifloxacin Hydrochloride	Each film coated tablet contains:				
	Tablets	Moxifloxacin Hydrochloride	IP			Approved on
		Eq. to Moxifloxacin		400	mg	31.05.2021
		Excipients		q.s		
			•		•	
805.	Ketoprofen & Thiocolchicoside	Each film coated tablet contains:				
	Tablets	Ketoprofen	IP	100	mg	Approved on
		Thiocolchicoside	IP	8	mg	31.05.2021
		Excipients		q.s		
906		Each film ageted tablet container				
806.	Cyanocobalamin, Chromium Picolinate, Folic Acid,	Each film coated tablet contains:	IP	15	mag	
	Nicotinamlide, Pyridoxine,	Cyanocobalamin Chromium Picolinate	USP	250	mcg	
	Selenius Acid & Zinc Sulphate	Folic Acid	IP	1500	mcg	
	Tablets	Nicotinamlide	IP	100	mcg	
		Pyridoxine Hcl	IP	3	mg	Approved on
		Selenius Acid	USP	100	mg	31.05.2021
		Zinc Sulphate Monohydrate	IP	61.8	mcg	
		Eq. to Zinc	117	22.5	mg mg	
		Excipients		q.s	5	
		Approved colour used		-1'-	_1	
	1	11				l .

807.	Methylcobalamin, Calcium,	Each hard gelatin capsule contains:				
	Calcitriol, Pyridoxine Hcl &	Methylcobalamin	IP	1500	mcg	1
	Folic Acid Capsules	Calcium	IP	200	mg	1
		Calcitriol	IP	0.25	mcg	1 1
		Pyridoxine Hcl	IP	3	mg	- Approved on
		Folic Acid	IP	1.5	mg	31.05.2021
		Excipients		q.s		1
		Approved colour used in empty		1'~		1
		capsule shell				
		1				
808.	Liquid Paraffin, Milk of	Each 5 ml contains:				
	Magnesia & Sodium	Light Liquid Paraffin	IP	1.25	ml	
	Picosulphate Suspension	Milk of Magnesia	IP	3.75	ml	Approved on
	1	Sodium Picosulphate	BP	3.33	mg	31.05.2021
		In a flavoured suspension base		q.s	1115	31.03.2021
		Approved colour used	1	4.5		-
		Approved colour used				
809.	Amlodipine, Olmesartan	Each film coated tablet contains:				
009.	Medoxomil &	Amlodipine Besilate	IP			
	Hydrochlorothiazide Tablets	Eq. to Amlodipine	11	10	mg	
	Trydroemorodmazide Tablets	Olmesartan Medoxomil	IP	40	mg	Approved on
		Hydrochlorothiazide	IP	25		31.05.2021
		Excipients	11	q.s	mg	
		Approved colour used		4.5		-
		ripproved colour used				
810.	Allopurinol SR Capsules 250	Each hard gelatin sustained release c	ansule con	tains:		
010.	mg	Allopurinol (as SR)	IP	250	mg	-
	5	Excipients		q.s		Approved on
		Approved colour used in empty		1'~		31.05.2021
		capsule shell				
		capsare sheri				
811.	Etodolac & Thiocolchicoside	Each film coated tablet contains:				
011.	Tablets	Etodolac	IP	300	mg	Approved on
		Thiocolchicoside	IP	8	mg	31.05.2021
		Excipients		q.s		31.03.2021
<u> </u>		1	I L	1	I	<u>ı</u>
812.	Ketoprofen & Thiocolchicoside	Each film coated tablet contains:				
	Tablets	Ketoprofen	IP	50	mg	Approved on
		Thiocolchicoside	IP	4	mg	31.05.2021
		Excipients		q.s]
813.	Amantadine Hydrochloride	Each Hard gelatin capsule contains :				
	Capsules IP	Amantadine Hydrochloride	IP	100	mg	Approved on
		Excipients		q.s		31.05.2021
		Approved colour used in empty caps	sule shell			
814.	Rivaroxaban Tablets	Each Film Coated Tablet Contains:				
		Rivaroxaban		2.5	mg	Approved on
		Excipients		q.s		31.05.2021
		Approved colour used]
		Tipproved colour used				<u> </u>

815.	Rivaroxaban Tablets	Each Film Coated Tablet Contains:				
		Rivaroxaban		10	mg	Approved on
		Excipients		q.s		31.05.2021
		Approved colour used				
		1 11				•
816.	Rivaroxaban Tablets	Each Film Coated Tablet Contains:				
		Rivaroxaban		15	mg	Approved on
		Excipients		q.s	31.05.2021	
		Approved colour used				
	L	11				l
817.	Rivaroxaban Tablets	Each Film Coated Tablet Contains:				
		Rivaroxaban		20	mg	Approved or
		Excipients		q.s		31.05.2021
		Approved colour used	_	1		1
818.	Dapagliflozin Tablets	Each Film Coated Tablet Contains:				
		Dapagliflozin propanediol				1
		monohydrate				Approved or
		Eq. to Dapagliflozin		5	mg	31.05.2021
		Excipients		q.s		
		Approved colour used				
819.	Dapagliflozin Tablets	Each Film Coated Tablet Contains:				
		Dapagliflozin propanediol				
		monohydrate				Approved or
		Eq. to Dapagliflozin		10	mg	31.05.2021
		Excipients		q.s		
		Approved colour used				
820.	D	Each 5ml contains:				
820.	Posaconazole oral suspension	Posaconazole		200	Mα	_
		Sodium benzoate (as Preservative)	IP	10	Mg Mg	Approved or
		Excipients	11	q.s.	IVIg	31.05.2021
		Approved colours used.		4.5.		-
	<u> </u>	ripproved colours used.				
821.	Rosuvastatin & Aspirin	Each hard gelatin capsule contains:				
	Capsules	Rosuvastatin Calcium	IP			
	1	Eq. to Rosuvastatin		10	mg	
		(As Pellets)				Approved or
		Aspirin	IP	75	mg	31.05.2021
		(As enteric coated Pellets)			-	
		Excipients		q.s		
		Approved colour used in empty capsu	ıle shell			
822.	Rosuvastatin & Aspirin	Each hard gelatin capsule contains:				
	Capsules	Rosuvastatin Calcium	IP			
		Eq. to Rosuvastatin		5	mg	
		(As Pellets)				Approved or
		Aspirin	IP	75	mg	31.05.2021
		(As enteric coated Pellets)				-
		Excipients Approved colour used in empty caps:		q.s		

823.	Rosuvastatin & Aspirin	Each hard gelatin capsule contains:				
	Capsules	Rosuvastatin Calcium	IP			
		Eq. to Rosuvastatin		10	mg	
		(As Pellets)			8	Approved on
		Aspirin	IP	150	mg	31.05.2021
		(As enteric coated Pellets)		150	1115	31.03.2021
		Excipients		q.s		
		Approved colour used in empty capsul	a chall	4.5		
	<u></u>	Approved colour used in empty capsur	c silcii			
824.	Metformin Hcl (Extended	Each uncoated bilayered tablet contains	s:			
	release) & Teneligliptin	Teneligliptin Hydrobromide Hydrate	IΡ			
	Tablets	Eq to Teneligliptin		20	mg	Approved on
		Metformin Hydrochloride	IP	1000	_	31.05.2021
		·	11	1000	mg	31.03.2021
		(as extended release)				
		Excipients		q.s		
825.	1	Each film coated tablet contains:				
023.	Betahistine Tablets IP 16mg	Betahistine Hydrochloride	IP	16	mg	Approved
	Betainstine Tablets II Tollig	Excipients	11		mg	on
		Approved colour used.		q.s		14.07.2021
		Approved colour used.				
826.		Each film coated tablet contains:				4
	Betahistine Tablets IP 8mg	Betahistine Hydrochloride	IP	8	mg	Approved
		Excipients		q.s		on
			LL	-		14.07.2021
		Approved colour used.				
027		1 **				
827.	D. I. C. W. II. C. D.	Each film coated tablet contains:	ID	24		<u> </u>
827.	Betahistine Tablets IP	Each film coated tablet contains : Betahistine Hydrochloride	IP	24	mg	Approved
827.	Betahistine Tablets IP	Each film coated tablet contains: Betahistine Hydrochloride Excipients	IP	24 q.s	mg	Approved
827.	Betahistine Tablets IP	Each film coated tablet contains : Betahistine Hydrochloride	IP		mg	Approved
	Betahistine Tablets IP	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used.	IP		mg	Approved
827. 828.		Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains:		q.s		Approved on 14.07.2021
	Ofloxacin & Metronidazole	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin	IP		mg	Approved
		Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate		q.s	mg	Approved on 14.07.2021 Approved on
	Ofloxacin & Metronidazole	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole	IP	100 200		Approved on 14.07.2021 Approved
	Ofloxacin & Metronidazole	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base	IP	q.s	mg	Approved on 14.07.2021 Approved on
	Ofloxacin & Metronidazole	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole	IP	100 200	mg	Approved on 14.07.2021 Approved on
	Ofloxacin & Metronidazole	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains:	IP IP	100 200 q.s	mg	Approved on 14.07.2021 Approved on
828.	Ofloxacin & Metronidazole	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used.	IP	100 200	mg	Approved on 14.07.2021 Approved on
828.	Ofloxacin & Metronidazole Suspension	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains:	IP IP	100 200 q.s	mg	Approved on 14.07.2021 Approved on 14.07.2021
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol	IP IP	100 200 q.s	mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid	IP IP	100 200 q.s 250 100	mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used.	IP IP	100 200 q.s 250 100	mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension Ambroxol Hydrochloride,	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used. Each 5 ml contains:	IP IP IP	100 200 q.s 250 100 q.s	mg mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension Ambroxol Hydrochloride, Levocetirizine Hydrochloride,	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used. Each 5 ml contains: Ambroxol Hydrochloride	IP IP IP IP	100 200 q.s 250 100 q.s	mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on 14.07.2021
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension Ambroxol Hydrochloride,	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used. Each 5 ml contains: Ambroxol Hydrochloride Levocetirizine Hydrochloride	IP IP IP IP IP	200 q.s 250 100 q.s 15 2.5	mg mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension Ambroxol Hydrochloride, Levocetirizine Hydrochloride,	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used. Each 5 ml contains: Ambroxol Hydrochloride Levocetirizine Hydrochloride Guaiphenesin	IP IP IP IP IP	100 200 q.s 250 100 q.s	mg mg mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on 14.07.2021 Approved on 14.07.2021
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension Ambroxol Hydrochloride, Levocetirizine Hydrochloride,	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used. Each 5 ml contains: Ambroxol Hydrochloride Levocetirizine Hydrochloride	IP IP IP IP IP	200 q.s 250 100 q.s 15 2.5	mg mg mg mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on 14.07.2021 Approved Approved
828.	Ofloxacin & Metronidazole Suspension Paracetamol & Mefenamic Acid Suspension Ambroxol Hydrochloride, Levocetirizine Hydrochloride,	Each film coated tablet contains: Betahistine Hydrochloride Excipients Approved colour used. Each 5ml contains: Ofloxacin Metronidazole benzoate Eq. Metronidazole Flavoured suspension base Approved colour used. Each 5ml contains: Paracetamol Mefenamic Acid Flavoured suspension base Approved colour used. Each 5 ml contains: Ambroxol Hydrochloride Levocetirizine Hydrochloride Guaiphenesin	IP IP IP IP IP	250 100 q.s 250 100 q.s 15 2.5 50	mg mg mg mg mg mg mg	Approved on 14.07.2021 Approved on 14.07.2021 Approved on 14.07.2021 Approved on 14.07.2021

831	Paraetamol & Ibuprofen	Each 5 ml contains:				
051	suspension	Paraetamol	IP	125	mg	FDCs
	1	Ibuprofen	IP	100	mg	permitted
		Excipients		q.s		16.07.2015
		Approved colour used		1		
		1 **				
832.	Metformin Hcl (Extended	Each film tablet contains:				
	release) & Teneligliptin	Teneligliptin Hydrobromide Hydrate	IP			
	Tablets	Eq to Teneligliptin		20	mg	FDCs
		Metformin Hydrochloride	IP	500	mg	permitted
		Excipients		q.s		16.07.2015
		Likelpients		4.5		
833.	Diclofenac Diethylamine	Composition:				
033.	,Linseed Oil, Menthol, Methyl	Diclofenac Diethylamine	IP	1.16%	W/W	1
	Salicylate gel	Eq. to Diclofenac Sodium		1%	w/w	
	, g.	Linseed Oil	BP	3.00%	w/w	FDCs
		Methyl Salicylate	IP	10.00%	w/w	permitted
		Benzyl Alcohol(as preservative)	IP	1.00%	w/w	17.07.2015
		Menthol	IP	0.50%	w/w	1
		Gel base		q.s		
				1	•	•
834.	Methylcobalamin,	Each 5 ml contains:				
	Cholecalciferol, Zinc	Methylcobalamin	IP	500	mcg	
	Gluconate, Lysine HCL,	Cholecalciferol(Vitamin D3)	IP	200	IU	
	Nicotinamide, Biotin, Folic	Zinc Gluconate	USP	5	mg	EDC.
	Acid, Syrup	Lysine HCL	BP	150	mg	FDCs permitted
		Nicotinamide	IP	15	mg	17.07.2015
		Biotin	USP	20	mcg	17.07.2013
		Folic Acid	IP	140	mcg	
		In A Flavoured Syrupy Base		q.s		
		Approved colour used.				
835.	Ferrous Ascorbate, Folic Acid,	Each 5 ml contains:				
055.	Methylcobalamin Suspension	Methylcobalamin	IP	500	mcg	_
	Wiedry reobulariting Suspension	Ferrous ascorbate	11	30	Mg	FDCs
		Eq. to elemental iron		10	mg	permitted
		Folic Acid	IP	0.5	mg	17.07.2015
		In a flavoured syrupy base	- 11	q.s	****8	17.07.2013
		Approved colour used.		4.5		
	<u> </u>	rapproved colour used.				
836.	L-Lysin Monohydrochloride,	Each 5 ml contains:				
	Benfothiamine, Pyridoxine	L-Lysine Monohydrochloride	USP	375	Mg	
	Hydrochloride,	Benfothiamine		1.0	Mg	
	Methylcobalamin, D-	Pyridoxine Hydrochloride	IP	0.75	Mg	
	Panthenol, Zinc Sulphate	Methylcobalamin	IP	15	Mcg	FDCs
	Monohydrate, Biotin Syrup	D-Panthenol	IP	3	Mg	permitted
		Zinc Sulphate Monohydrate	IP	5	Mg	03.02.2016
		(Eq.To Elemental Zinc)			3	
		Biotin	USP	50	Mcg	
		In A Flavoured Syrupy Base		q.s		
		Approved Colour Used.				

837.	Ferrous Bis-Glycinate, Folic	Each 5 Ml Contains:					
037.	Acid, Zinc Bis-Glycinate,	Methylcobalamin	IP	250	mcg		
	Methylcobalamine Syrup	Ferrous Bis-Glycinate					
		Eq. To Elemental Iron		15	mg	FDCs	
		Folic Acid	IP	0.37	mg	permitted	
		Zinc Bis-Glycinate				17.07.2015	
		Eq.To Elemental Zinc		3.75	mg		
		In A Flavoured Syrupy Base		q.s			
		Approved Colour Used.					
838.	Tadalafil Tablets IP 10mg	Each film coated tablet contain:					
		Tadalafil	IP	10	Mg	IP-2018	
		Excipients		q.s		P.NO.3302	
		approved colour used.		1			
		approvide colour accu.					
839.	Tadalafil Tablets IP 20mg	Each film coated tablet contain:				IP-2018	
		Tadalafil	IP	20	Mg	P.NO.3302	
		Excipients		q.s		P.NO.3302	
		approved colour used.	I		I		
						1	
840.	Tadalafil Orally Disintegrating	ng Each Orally Disintegrating tablet contain:					
	Tablets	Tadalafil	IP	10	Mg	DCGI- 30.07.2014	
		Excipients		q.s		30.07.2014	
		approved colour used.					
841.	T- 1-1-61 O-11- Disinttim-	Each Orally Disints quating tablet or				1	
041.	Tadalafil Orally Disintegrating Tablets	Each Orally Disintegrating tablet co	1	20	M-	DCGI-	
	Tablets		IP	20	Mg	30.07.2014	
		Excipients		q.s		-	
		approved colour used.					
842.	Permethrin Soap 1% W/W	Composition:				Lanark	
0 12.	Termeum Soap 176 W/W	Permethrin	BP	1%	w/w	Laboratories	
		Excipients		q.s	***	Paonta Sahib	
		Base		q.s		(Market	
		Buse		4.5		Sample)	
		1		_1	l	1 /	
843.	Ketoconazole &Cetrimide	Composition:					
	Soap	Ketoconazole	IP	2%	w/w	FDCs-	
		Cetrimide	IP	0.5%	w/w	17.07.2015	
		Excipients		q.s		17.07.2013	
		Base		q.s]	
844.	Ketoconazole Soap	Composition:	1	1		Lanark	
		Ketoconazole	IP	2%	w/w	laboratories	
		Excipients		q.s		paonta sahib	
		Base		q.s		(market	
						sample)	

845.	Monosulfiram Soap IP	Composition:				
		Monosulfiram	IP	5.00%	w/w	Alkem
		Excipients		q.s		Laboratories
		Base		q.s		
846.	Ivermectin Soap	Composition:				Lanark
		Ivermectin	IP	1%	w/w	laboratories
		Excipients		q.s		paonta sahib
		Base		q.s		(market
						sample)
847.	Allegate Witnesser E	Commonition			1	
847.	Allantoin, Vitamin-E	Composition:	DD	0.200/	/	
	Acetate, Tea tree oil &	Allantoin	BP	0.20%	w/w	
	Titanium Dioxide Soap	Vitamin-E Acetate		0. 25%	w/w	FDCs-
	(As cleansing agent for	Tea tree oil	BP	0. 25%	w/w	15.03.2016
	acne.)	Titanium Dioxide	IP	0.50%	w/w	
		Excipients		q.s		
		Base		q.s		
		T				
848.	Allantoin, Vitamin-E	Composition:			1	
	Acetate & Tea tree oil Soap	Allantoin	BP	0.25%	w/w	
	(As cleansing agent for	Vitamin-E Acetate		0. 25%	w/w	FDCs-
	acne.)	Tea tree oil	BP	0. 50%	w/w	15.03.2016
		Excipients		q.s		
		Base		q.s		
849.	Nimesulide & Paracetamol	Each uncoated tablet contains:				
	Tablets	Nimesulide	BP	100	Mg	FDCs
	(Not for Children below the	Paracetamol	IP	350	mg	16.07.2015
	age of 12 years)	Excipients		q.s		



approved colour used.