LOXIER PHARMA

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
1	Sertaconazole Nitrate Cream	Composition:				
		Sertaconazole Nitrate	BP	2%	w/w	
		Preservative: Benzyl Alcohol	IP	1%	w/w	
		Cream base		q.s		APPR
2	Mometasone Furoate & Fusidic Acid Cream	Composition:				
		Mometasone Furoate	IP	0.1%	w/w	
		Fusidic Acid	IP	2%	w/w	
		Cream base		q.s		APPR
3	Clobetasol Propionate Cream IP	Composition :				
		Clobetasol Propionate	IP	0.05%	w/w	
		Chlorocresol (as preservative)	IP	0.1%	w/w	
		Cream base		q.s		APPR
4	Hydroquinone, Tretinoin & Mometasone Furoate Cream	Composition:				
		Hydroquinone	IP	2%	w/w	
		Tretinoin	USP	0.025%	w/w	
		Mometasone Furoate	IP	0.1%	w/w	
		Cream base		q.s		APPR
5	Mometasone Furoate Cream IP	Composition:				
		Mometasone Furoate	IP	0.1%	w/w	
		Chlorocresol (as preservative)	IP	0.1%	w/w	
		Cream base		q.s		APPR
6	Fusidic Acid Cream IP	Composition:				
		Fusidic Acid	IP	2%	w/w	
		Cream base		q.s		APPR
7	Mometasone Furoate & Terbinafine HCl Cream	Composition:				
		Mometasone Furoate	IP	0.1%	w/w	
		Terbinafine HCl	BP	1%	w/w	
		Cream base		q.s		APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
8	Betamethasone & Miconazole Cream	Composition:				
		Betamethasone Dipropionate Eq. to Betamethasone	IP	0.05%	w/w	
		Miconazole Nitrate Eq. to Miconazole	IP	2%	w/w	
		Cream base		q.s		APPR
9	Linseed Oil, Diclofenac	Composition:				
		Linseed Oil	BP	3%	w/w	

	Diethylamine, Methyl Salicylate & Menthol Gel	Diclofenac Diethylamine Eq. to Diclofenac Sodium	BP	1%	w/w	
		Methyl Salicylate	IP	10%	w/w	
		Menthol	IP	5%	w/w	
		Benzyl Alcohol (As Preservative)	IP	1%	w/w	
		In a water washable base		q.s		APPR
10	Pantoprazole Tablets IP 40mg	Each enteric coated tablet contains :				
		Pantoprazole Sodium Eq. to Pantoprazole	IP	40	mg	
		Excipients		q.s		
		Approved colour used				
11	Aceclofenac & Paracetamol Tablets	Each film coated tablet contains :				
		Aceclofenac	IP	100	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
12	Ofloxacin & Ornidazole Tablets	Each film coated tablet contains :				
		Ofloxacin	IP	200	mg	
		Ornidazole	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR
13	Ofloxacin Tablets IP	Each film coated tablet contains :				
		Ofloxacin	IP	200	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
14	Azithromycin Tablets IP 250mg	Each film coated tablet contains :				
		Azithromycin	IP			
		Eq. to Anhydrous Azithromycin		250	mg	
		Excipients		q.s		
		Approved colour used				
15	Azithromycin Tablets IP 500mg	Each film coated tablet contains :				
		Azithromycin Eq. to Anhydrous Azithromycin	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR
16	Rabeprazole Tablets IP 20mg	Each enteric coated tablet contains :				
		Rabeprazole Sodium	IP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR
17	Levocetirizine & Montelukast Tablets	Each film coated tablet contains :				
		Levocetirizine Hydrochloride	IP	5	mg	
		Montelukast Sodium Eq. to Montelukast	IP	10	mg	
		Excipients		q.s		APPR

		Approved colour used				
18	Levofloxacin Tablets IP 250mg	Each film coated tablet contains :				
		Levofloxacin Hemihydrate Eq. to Levofloxacin	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APPR
19	Levofloxacin Tablets IP 500mg	Each film coated tablet contains :				
		Levofloxacin Hemihydrate Eq. to Levofloxacin	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
20	Cefixime Tablets IP 100mg	Each film coated tablet contains :				
		Cefixime Eq. to Anhydrous Cefixime	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR
21	Cefixime Tablets IP 200mg	Each film coated tablet contains :				
		Cefixime Eq. to Anhydrous Cefixime	IP	200	mg	
		Excipients		q.s		
		Approved colour used				APPR
22	Cefpodoxime Tablets IP 100mg	Each film coated tablet contains :				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR
23	Cefpodoxime Tablets IP 200mg	Each film coated tablet contains :				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	200	mg	
		Excipients		q.s		
		Approved colour used				
24	Cefuroxime Axetil Tablets IP 250mg	Each film coated tablet contains :				
		Cefuroxime Axetil Eq. to Cefuroxime	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APPR
25	Cefuroxime Axetil Tablets IP 500mg	Each film coated tablet contains :				
		Cefuroxime Axetil Eq. to Cefuroxime	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
26	Cefixime & Ofloxacin Tablets	Each film coated tablet contains :				
		Cefixime Eq. to Anhydrous Cefixime	IP	200	mg	
		Ofloxacin	IP	200	mg	
		Excipients		q.s		
		Approved colour used				APPR
27		Each film coated tablet contains :				

		Cefixime	IP			
	Cefixime & Potassium Clavulanate Tablets	Eq. to Anhydrous Cefixime		200	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APPR
28	Cefpodoxime & Potassium Clavulanate Tablets	Each film coated tablet contains :				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	200	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APPR
29	Amoxycillin & Potassium Clavulanate Tablets IP	Each film coated tablet contains :				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	500	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		q.s		
		Approved colour used				
30	Cefixime Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	50	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
31	Cefixime Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	100	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				
32	Cefpodoxime Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	50	mg	
		Excipients		q.s		APPR
		In a flavoured base				
		Approved colour used				
33	Cefpodoxime Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	100	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR
34	Amoxycillin Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	125	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR
35	Amoxycillin Oral	Each 5ml of the reconstituted suspension contains:				

	Suspension IP	Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
36	Cephalexin Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	125	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR
37	Cefaclor Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Cefaclor	IP	125	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR
38	Cefixime & Potassium Clavulanate Oral Suspension	Each 5ml of the reconstituted suspension contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	50	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	31.25	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR
39	Cefpodoxime & Potassium Clavulanate Oral Suspension	Each 5ml of the reconstituted suspension contains:				
		Cefpodoxime Proxetil Eq . to Cefpodoxime	IP	50	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	31.25	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANT ITY	UNIT	APP
40	Omeprazole Capsules IP	Each hard gelatin capsule contains:				
		Omeprazole (As enteric coated pellets)	IP	20	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
41	Omeprazole & Domperidone Capsules	Each hard gelatin capsule contains:				
		Omeprazole (As enteric coated pellets)	IP	20	mg	
		Domperidone (As enteric coated pellets)	IP	10	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
42	Pantoprazole Sodium	Each hard gelatin capsule contains:				

	(EC)	Pantoprazole Sodium	IP			
	& Domperidone (SR) Capsules	Eq. to Pantoprazole (As enteric Coated pellets)		40	mg	
		Domperidone (As sustained release pellets)	IP	30	mg	
		Excipients		q.s		
		Approved colour used in empty capsule shells.				APPR
43	Calcium Dobesilate Monohydrate Capsules	Each hard gelatin capsule contains:				
		Calcium Dobesilate Monohydrate	IP	500	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR
44	Rabeprazole Sodium (EC) & Domperidone (SR) Capsules	Each hard gelatin capsule contains:				
		Rabeprazole Sodium (As enteric Coated pellets)	IP	20	mg	
		Domperidone (As sustained release pellets)	IP	30	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
45	Pantoprazole Sodium (EC) & Itopride HCl (SR) Capsules	Each hard gelatin capsule contains:				
		Pantoprazole Sodium Eq. to Pantoprazole (As enteric Coated pellets)	IP	40	mg	
		Itopride HCl (As sustained release pellets)		150	mg	
		Excipients		q.s		
		Approved colour used in empty capsule shells.				APPR
46	Rabeprazole Sodium(EC) & Itopride(SR) Capsules	Each hard gelatin capsule contains:				
		Rabeprazole Sodium (As enteric Coated pellets)	IP	20	mg	
		Itopride HCl (As sustained release pellets)		150	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
47		Each hard gelatin capsule contains:				
	Esomeprazole Magnesium (EC) & Domperidone (SR) Capsules	Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (As enteric Coated pellets)	IP	40	mg	
		Domperidone (As sustained release pellets)	IP	30	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
48	Esomeprazole Magnesium (EC) & Itopride HCl (SR) Capsules	Each hard gelatin capsule contains:				
		Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (As enteric Coated pellets)	IP	40	mg	
		Itopride HCl (As sustained release pellets)		150	mg	

		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
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S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFICATION**	**QUANTITY**	**UNIT**	**APP**
49	Itraconazole Capsules	Each hard gelatin capsule contains:				
		Itraconazole Pellets Eq. to Itraconazole	BP	100	mg	
		Excipients		q.s		
		Approved colour used in empty capsule shell.				APPR
50	Amoxycillin Capsules IP 250mg	Each hard gelatin capsule contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR
51	Amoxycillin Capsules IP 500mg	Each hard gelatin capsule contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	500	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR
52	Cephalexin Capsules IP 250mg	Each hard gelatin capsule contains:				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	250	mg	
		Excipients		q.s		APPR
		Approve colour used in empty capsule shell.				
53	Cephalexin Capsules IP 500mg	Each hard gelatin capsule contains:				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	500	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell				APPR
54	Amoxycillin & Lactobacillus Capsules	Each hard gelatin capsule contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	mg	
		Lactobacillus		1.66	billion spores	
		Excipients		q.s		
		Approved colours used in capsule shells.				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFICATION**	**QUANTITY**	**UNIT**	**APP**
55	Amoxycillin & Lactobacillus Capsules	Each hard gelatin capsule contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	500	mg	
		Lactobacillus		1.66	billion spores	
		Excipients		q.s		
		Approved colours used in capsule shells.				APPR
56	Pregabalin & Methycobalamin Capsules	Each hard gelatin capsule contains:				
		Pregabalin	IP	75	mg	
		Methylcobalamin	USP	750	mcg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR
57	Methycobalamin, Alpha	Each hard gelatin capsule contains:				

	Lipoic Acid,Thiamine Mononitrate, Pyridoxine HCl & Folic Acid Capsules	Methylcobalamin	USP	1500	mcg	
		Alpha Lipoic Acid	USP	100	mg	
		Thiamine Mononitrate	IP	10	mg	
		Pyridoxine Hydrochloride	IP	3	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR
58		Each hard gelatin capsule contains:				
		Cefixime	IP			
	Cefixime & Erdosteine Capsules	Eq. to Anhydrous Cefixime		200	mg	
		Erdosteine		300	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell.				APPR
59	Ambroxol Hydrochloride, Levosalbutamol & Guaiphenesin Syrup	Each 5 ml contains :				
		Ambroxol Hydrchloride	IP	30	mg	
		Levosalbutamol Sulphate Eq. to Levosalbutamol	IP	1	mg	
		Guaiphenesin	IP	50	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used.				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
60	Ambroxol HCl, Terbutaline Sulphate, Guaiphenesin & Menthol Syrup	Each 5 ml contains :				
		Ambroxol HCl	IP	15	mg	
		Terbutaline Sulphate	IP	1.25	mg	
		Guaiphenesin	IP	50	mg	
		Menthol	IP	2.5	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used.				APPR
61	Ofloxacin Oral Suspension IP 50mg/5ml	Each 5 ml contains :				
		Ofloxacin	IP	50	mg	
		In a flavoured base		q.s		
		Approved colour used.				APPR
62	Ofloxacin Oral Suspension IP 100mg/5ml	Each 5 ml contains :				
		Ofloxacin	IP	100	mg	
		In a flavoured base		q.s		
		Approved colour used.				APPR
63	Levocetirizine & Montelukast Syrup	Each 5 ml contains :				
		Levocetirizine Dihydrochloride	IP	2.5	mg	
		Montelukast Sodium Eq. to Montelukast	IP	4	mg	
		In a flavoured syrupy base		q.s.		
		Approved colour used.				APPR
64		Each 5 ml contains :				
		Albendazole	IP	200	mg	
	Albendazole Oral Suspension IP	In a flavoured base		q.s		APPR
		Approved colour used				
65	Liquid Paraffin, Milk of Magnesia & Sodium Picosulphate Suspension	Each 5 ml contains :				
		Liquid Paraffin	IP	1.25	ml	
		Milk of Magnesia	IP	3.75	ml	
		Sodium Picosulphate	BP	3.33	mg	

	Suspension	In a flavoured base		q.s		
		Approved colour used.				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
66	Ferrous Ascorbate & Folic Acid Suspension	Each 5 ml contains :				
		Ferrous Ascorbate Eq. to Elemental Iron		30	mg	
		Folic Acid	IP	550	mcg	
		In a flavoured base		q.s		
		Approved colour used.				APPR
67	Chloramphenicol Oral Suspension IP	Each 5 ml contains :				
		Chloramphenicol Palmitate Eq. to Chloramphenicol	APPROVED	125	mg	
		In a flavoured base	APPROVED	q.s		
		Approved colour used.				APPR
68	Cyproheptadine Hydrocholride Syrup IP	Each 5 ml contains :				
		Cyproheptadine Hydrocholride (Anhydrous)	IP	2	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used.				APPR
69	Paracetamol Tablets IP 500mg	Each uncoated tablet contains:				
		Paracetamol	IP	500	mg	
		Excipients		q.s		APPR
70	Paracetamol Tablets IP 650mg	Each uncoated tablet contains				
		Paracetamol	IP	650	mg	
		Excipients		q.s		APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
71	Drotaverine Hydrochloride Tablets IP 80mg	Each film coated tablet contains:				
		Drotaverine Hydrochloride	IP	80	mg	
		Excipients		q.s		
		Approved colour used				APPR
72	Lamotrigine Dispersible Tablets IP 100mg	Each uncoated dispersible tablet contains:				
		Lamotrigine	IP	100	mg	
		Excipients		q.s		APPR
73	Ursodeoxycholic Acid Tablets IP	Each film coated tablet contains:				
		Ursodeoxycholic Acid	IP	300	mg	
		Excipients		q.s		
		Approved colour used				APPR
74	Escitalopram Tablets IP 10mg	Each film coated tablet contains:				
		Escitalopram Oxalate Eq. to Escitalopram	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
75	Rosuvastatin Tablets IP 10mg	Each film coated tablet contains:				
		Rosuvastatin Calcium Eq. to Rosuvastatin	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
76	Rosuvastatin Tablets	Each film coated tablet contains:				

	IP 20mg	Rosuvastatin Calcium Eq. to Rosuvastatin	IP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
77	Loratadine Tablets USP	Each uncoated tablet contains:				
		Loratadine	USP	10	mg	
		Excipients		q.s		APPR
78	Folic Acid Tablets IP 5mg	Each uncoated tablet contains:				
		Folic Acid	IP	5	mg	
		Excipients		q.s		APPR
79	Betahistine Tablets IP 8mg	Each uncoated tablet contains:				
		Betahistine Hydrochloride	IP	8	mg	
		Excipients		q.s		APPR
80	Aceclofenac Tablets IP 100mg	Each film coated tablet contains:				
		Aceclofenac	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR
81	Amlodipine Tablets IP	Each uncoated tablet contains:				
		Amlodipine Besylate Eq. to Amlodipine	IP	2.5	mg	
		Excipients		q.s		
		Approved colour used				APPR
82	Amlodipine Tablets IP 5mg	Each uncoated tablet contains:				
		Amlodipine Besylate Eq. to Amlodipine	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
83	Roxithromycin Tablets IP 150mg	Each film coated tablet contains:				
		Roxithromycin	IP	150	mg	
		Excipients		q.s		
		Approved colour used				APPR
84	Albendazole Tablets IP 400mg	Each uncoated chewable tablet contains:				
		Albendazole	IP	400	mg	
		Excipients		q.s		APPR
85	Ondansetron Tablets IP 4mg	Each uncoated tablet contains:				
		Ondansetron HCl Eq. to Ondansetron	IP	4	mg	
		Excipients		q.s		
		Approved colour used				APPR
86	Ondansetron Orally Disintegrating Tablets IP 4mg	Each uncoated orally disintegrating tablet contains:				
		Ondansetron HCl Eq. to Ondansetron	IP	4	mg	
		Excipients		q.s		
		Approved colour used				APPR
87	Cinnarizine Tablets IP 25mg	Each uncoated tablet contains:				
		Cinnarizine	IP	25	mg	
		Excipients		q.s		APPR

88	Ciprofloxacin Tablets IP 250mg	Each film coated tablet contains:				
		Ciprofloxacin HCl Eq. to Ciprofloxacin	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
89	Ciprofloxacin Tablets IP 500mg	Each film coated tablet contains:				
		Ciprofloxacin HCl Eq. to Ciprofloxacin	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR
90	Olanzapine Tablets IP 5mg	Each film coated tablet contains:				
		Olanzapine	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
91	Olanzapine Tablets IP 10mg	Each film coated tablet contains:				
		Olanzapine	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
92	Linezolid Tablets IP 600mg	Each film coated tablet contains:				
		Linezolid	IP	600	mg	
		Excipients		q.s		
		Approved colour used				APPR
93	Cetirizine Tablets IP 10mg	Each film coated tablet contains:				
		Cetirizine Hydrochloride	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
94	Glimepiride Tablets IP 1mg	Each uncoated tablet contains:				
		Glimepiride	IP	1	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
95	Glimepiride Tablets IP 2mg	Each uncoated tablet contains:				
		Glimepiride	IP	2	mg	
		Excipients		q.s		APPR
96	Racecadotril Sachets IP 10mg	Each sachet contains:				
		Racecadotril	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
97	Escitalopram Tablets IP 5mg	Each film coated tablet contains:				
		Escitalopram Oxalate Eq. to Escitalopram	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
98	Cinnarizine Tablets IP	Each uncoated tablet contains:				
		Cinnarizine	IP	75	mg	
		Excipients		q.s		APPR
99	Metformin Hydrochloride SR Tablets IP 500mg	Each film coated sustained release tablet contains:				
		Metformin Hydrochloride	IP	500	mg	
		Excipients		q.s		

	Tablets IP 500mg	Approved colour used				APPR
100	Thiocolchicoside & Aceclofenac Tablets	Each film coated tablet contains :				
		Thiocolchicoside	IP	4	mg	
		Aceclofenac	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
101		Each film coated tablet contains :				
	Thiocolchicoside & Aceclofenac Tablets	Thiocolchicoside	IP	8	mg	
		Aceclofenac	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR
102	Diclofenac Potassium & Paracetamol Tablets	Each film coated tablet contains :				
		Diclofenac Potassium	BP	50	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
103	Beclomethasone Dipropionate Cream	Composition :				
		Beclomethasone Dipropionate	IP	0.025%	w/w	
		Cream base		q.s		APPR
104	Terbinafine HCl Cream	Composition :				
		Terbinafine HCl	BP	1%	w/w	
		Preservative: Benzyl Alcohol	IP	1%	w/w	
		Cream base		q.s		APPR
105	Ambroxol Hydrochloride SR & Levocetirizine Tablets	Each film coated tablet contains :				
		Levocetirizine Dihydrochloride	IP	5	mg	
		Ambroxol Hydrochloride (In Sustained release form)	IP	75	mg	
		Excipients		q.s		
		Approved colour used				APPR
106	Metoprolol Succinate Extended Release Tablets USP	Each extended release film coated tablet contains :				
		Metoprolol Succinate	USP	11.875	mg mg	
		Eq. to Metoprolol Tartrate		12.5		
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
107	Metoprolol Succinate Extended Release Tablets USP	Each extended release film coated tablet contains :				
		Metoprolol Succinate	USP	23.75	mg mg	
		Eq. to Metoprolol Tartrate		25		
		Excipients		q.s		APPR
		Approved colour used				
108	Metoprolol Succinate Extended Release Tablets USP	Each extended release film coated tablet contains :				
		Metoprolol Succinate	USP	47.50	mg mg	
		Eq. to Metoprolol Tartrate		50		
		Excipients		q.s		
		Approved colour used				APPR

109	Metoprolol Succinate (ER) & Telmisartan Tablets	Each film coated tablet contains :				
		Metoprolol Succinate Eq. to Metoprolol Tartrate (As Extended Release)	USP	50	mg	
		Telmisartan	IP	40	mg	
		Excipients		q.s		
		Approved colour used				APPR
110	Metoprolol Succinate (ER) & Ramipril Tablets	Each film coated tablet contains :				
		Metoprolol Succinate Eq. to Metoprolol Tartrate (As Extended Release)	USP	50	mg	
		Ramipril	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
111	Metformin Hydrochloride SR Tablets IP 1000mg	Each film coated sustained release tablet contains:				
		Metformin Hydrochloride	IP	1000	mg	
		Excipients		q.s		
		Approved colour used				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANT ITY**	**UNIT**	**APP**
112	Glucosamine Sulphate Potassium Chloride, Methyl Sulphonyl Methane & Diacerein Tablets	Each film coated tablet contains:				
		Glucosamine Sulphate Potassium Chloride (eq. to Glucosamine 444 mg)	USP	750	mg	
		Methyl Sulphonyl Methane	USP	250	mg	
		Diacerein	IP	50	mg	
		Excipients		q.s		
		Approved colour used				
113	Calcium Citrate Malate, Cholecalciferol & Folic Acid Tablets	Each film coated tablet contains:				
		Calcium Citrate Malate Eq. to Elemental Calcium		250	mg	
		Cholecalciferol	IP	100	IU	
		Folic Acid	IP	50	mcg	
		Excipients		q.s		
		Approved colour used				APPR
114	Citicoline Tablets IP	Each film coated tablet contains:				
		Citicoline Sodium Eq. to Citicoline	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR
115	Pregabalin SR & Methylcobalamin Tablets	Each film coated tablet contains:				
		Pregabalin (In sustained release form)	IP	75	mg	
		Methylcobalamin	USP	1500	mcg	
		Excipients		q.s		
		Approved colour used				APPR
116	Ivermectin Tablets	Each uncoated tablet contains:				
		Ivermectin	IP	3	mg	
		Excipients		q.s		APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
117	Ivermectin Tablets	Each uncoated tablet contains:				
		Ivermectin	IP	6	mg	
		Excipients		q.s		APPR
118	Ivermectin Tablets	Each uncoated tablet contains:				
		Ivermectin	IP	9	mg	
		Excipients		q.s		APPR
119	Ivermectin Tablets	Each uncoated tablet contains:				
		Ivermectin	IP	12	mg	
		Excipients		q.s		APPR
120	Chlorhexidine Mouthwash IP	Composition :				
		Chlorhexidine Gluconate Solution Eq. to Chlorhexidine Gluconate	IP	0.2%	w/v	
		In a pleasantly flavoured base				
		Approved colour used				APPR
121	Losartan Potassium Tablets IP 50mg	Each film coated tablet contains:				
		Losartan Potassium	IP	50	mg	
		Excipients		q.s		
		Approved colour used				APPR
122	Fluconazole Tablets IP 200mg	Each uncoated tablet contains:				
		Fluconazole	IP	200	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
123	Etoricoxib Tablets IP 120mg	Each film coated tablet contains:				
		Etoricoxib	IP	120	mg	
		Excipients		q.s		
		Approved colour used				APPR
124	Etoricoxib & Paracetamol Tablets	Each film coated tablet contains:				
		Etoricoxib	IP	60	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
125		Each film coated tablet contains:				
	Ibuprofen & Paracetamol Tablets	Ibuprofen	IP	400	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
126	Metoprolol Succinate (ER) & Amlodipine Tablets	Each film coated tablet contains :				
		Metoprolol Succinate Eq. to Metoprolol Tartrate (As Extended Release)	USP	50	mg	
		Amlodipine Besylate Eq. to Amlodipine	IP	5	mg	
		Excipients		q.s		
		Approved colour used				

						APPR
127	Metformin Hydrochloride (SR) & Glimepiride Tablets	Each film coated tablet contains:				
		Metformin Hydrochloride (In Sustained release form)	IP	1000	mg	
		Glimepiride	IP	1	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
128	Metformin Hydrochloride (SR) & Glimepiride Tablets	Each film coated tablet contains:				
		Metformin Hydrochloride (In Sustained release form)	IP	1000	mg	
		Glimepiride	IP	2	mg	
		Excipients		q.s		
		Approved colour used				APPR
129	Ursodeoxycholic Acid Tablets IP	Each film coated tablet contains:				
		Ursodeoxycholic Acid	IP	150	mg	
		Excipients		q.s		
		Approved colour used				APPR
130	Losartan Potassium	Each film coated tablet contains:				
		Losartan Potassium	IP	50	mg	
	& Hydrochlorothiazide Tablets IP	Hydrochlorothiazide	IP	12.5	mg	
		Excipients		q.s		
		Approved colour used				APPR
131	Metoprolol Succinate Extended Release Tablets USP	Each film coated extended release tablet contains :				
		Metoprolol Succinate Eq. to Metoprolol Tartrate	IP	95 100	mg mg	
		Excipients		q.s		
		Approved colour used				APPR
132	Levetiracetam Tablets USP	Each film coated tablet contains :				
		Levetiracetam	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
133	Lornoxicam Tablets	Each film coated tablet contains:				
		Lornoxicam	IP	4	mg	
		Excipients		q.s		
		Approved colour used				APPR
134	Metoprolol Succinate (ER) & Telmisartan Tablets	Each film coated tablet contains:				
		Metoprolol Succinate Eq. to Metoprolol Tartrate (As Extended Release)	USP	25	mg	
		Telmisartan	IP	40	mg	
		Excipients		q.s		
		Approved colour used				APPR
135	Metoprolol Succinate	Each film coated tablet contains:				

	(ER) & Amlodipine Tablets	Metoprolol Succinate Eq. to Metoprolol Tartrate (As Extended Release)	USP	25	mg	
		Amlodipine Besylate Eq. to Amlodipine	IP	5	mg	
		Excipients		q.s		APPR
		Approved colour used				
136	Amoxycillin & Potassium Clavulanate Tablets IP	Each film coated tablet contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APPR
137	Amoxycillin & Potassium Clavulanate Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	200	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	28.5	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
138	Amoxycillin & Potassium Clavulanate Tablets IP	Each film coated tablet contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	875	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APPR
139	Amoxycillin Dispersible Tablets IP 250mg	Each uncoated dispersible tablet contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	mg	
		Excipients		q.s		APPR
140	Amoxycillin Dispersible Tablets IP 125mg	Each uncoated dispersible tablet contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	125	mg	
		Excipients		q.s		APPR
141		Each film coated tablet contains:				
		Cefixime	IP			
	Cefixime & Lactic Acid Bacillus Tablets	Eq. to Anhydrous Cefixime		100	mg	
		Lactic Acid Bacillus		2.5	billion spores	
		Excipients		q.s		
		Approved colour used				APPR
142	Cefixime & Lactic Acid Bacillus Tablets	Each film coated tablet contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	200	mg	
		Lactic Acid Bacillus		2.5	billion spores	
		Excipients		q.s		

		Approved colour used				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANT ITY**	**UNIT**	**APP**
143	Cefpodoxime & Ofloxacin Tablets	Each film coated tablet contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	200	mg	
		Ofloxacin	IP	200	mg	
		Excipients		q.s		
		Approved colour used				APPR
144	Fexofenadine Tablets IP 180mg	Each film coated tablet contains:				
		Fexofenadine Hydrochloride	IP	180	mg	
		Excipients		q.s		
		Approved colour used				APPR
145	Artemether & Lumefantrine Tablets	Each uncoated tablet contains:				
		Artemether	IP	80	mg	
		Lumefantrine		480	mg	
		Excipients		q.s		APPR
146	Atorvastatin Tablets IP 10mg	Each film coated tablet contains:				
		Atorvastatin Calcium Eq. to Atorvastatin	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
147	Atorvastatin Tablets IP 20mg	Each film coated tablet contains:				
		Atorvastatin Calcium Eq. to Atorvastatin	IP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR
148	Atorvastatin Tablets IP 40mg	Each film coated tablet contains:				
		Atorvastatin Calcium Eq. to Atorvastatin	IP	40	mg	
		Excipients		q.s		
		Approved colour used				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANTI TY**	**UNIT**	**APP**
149	Atorvastatin Tablets IP 80mg	Each film coated tablet contains:				
		Atorvastatin Calcium Eq. to Atorvastatin	IP	80	mg	
		Excipients		q.s		
		Approved colour used				APPR
150	Rupatadine Fumarate Tablets	Each uncoated tablet contains :				
		Rupatadine Fumarate Eq. to Rupatadine	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
151	Levocetirizine Tablets IP 5mg	Each film coated tablet contains :				
		Levocetirizine Hydrochloride	IP	5	mg	
		Excipients		q.s		

		Approved colour used				APPR
152	Desloratadine Mouth Dissolving Tablets	Each uncoated mouth dissolving tablet contains :				
		Desloratadine		5	mg	
		Excipients		q.s		
		Approved colour used				APPR
153	Methylprednisolone Tablets IP 4mg	Each uncoated tablet contains:				
		Methylprednisolone	IP	4	mg	APPR
		Excipients		q.s		
154	Methylprednisolone Tablets IP 16mg	Each uncoated tablet contains:				
		Methylprednisolone	IP	16	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
155	Fexofenadine Hydrochloride & Montelukast Sodium Tablets	Each film coated tablet contains:				
		Fexofenadine Hydrochloride Eq. to Fexofenadine	IP	120	mg	
		Montelukast Sodium Eq. to Montelukast	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
156	Ferrous Ascorbate, Folic Acid & Zinc Sulphate Tablete	Each film coated tablet contains::				
		Ferrous Ascorbate Eq. to Elemental Iron		100	mg	
		Folic Acid	IP	1500	mcg	
		Zinc Sulphate Monohydrate (Eq. to Elemental Zinc 22.5 mg)	IP	61.8	mg	
		Excipients		q.s		
		Approved colour used				APPR
157	Voglibose Mouth Dissolving Tablets	Each uncoated mouth dissolving tablet contains:				
		Voglibose	IP	0.2	mg	
		Excipients		q.s		APPR
158	Voglibose Mouth Dissolving Tablets	Each uncoated mouth dissolving tablet contains :				
		Voglibose	IP	0.3	mg	
		Excipients		q.s		APPR
159	Calcium With Vitamin D	Each film coated tablet contains:				
		Calcium Carbonate	IP			
	Tablets USP	Eq. to Elemental Calcium		500	mg	
		Vitamin D3	IP	250	IU	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
160	S	Each uncoated tablet contains:				
		Terbinafine HCl Eq. to Terbinafine	BP	250	mg	
		Excipients		q.s		APPR
161	Etizolam Tablets	Each film coated tablet contains:				

		Etizolam		0.25	mg	
		Excipients		q.s		
		Approved colour used				APPR
162	Etizolam Tablets	Each film coated tablet contains:				
		Etizolam		0.5	mg	
		Excipients		q.s		
		Approved colour used				APPR
163	Metformin Hydrochloride (SR) & Glimepiride Tablets	Each film coated tablet contains:				
		Metformin Hydrochloride (In Sustained release form)	IP	500	mg	
		Glimepiride	IP	1	mg	
		Excipients		q.s		
		Approved colour used				APPR
164	Metformin Hydrochloride (SR) & Glimepiride Tablets	Each film coated tablet contains:				
		Metformin Hydrochloride (In Sustained release form)	IP	500	mg	
		Glimepiride	IP	2	mg	
		Excipients		q.s		
		Approved colour used				APPR
165	Amisulpride Tablets IP 50mg	Each uncoated tablet contains:				
		Amisulpride	IP	50	mg	APPR
		Excipients		q.s		

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
166	Amisulpride Tablets IP 100mg	Each uncoated tablet contains:				
		Amisulpride	IP	100	mg	
		Excipients		q.s		APPR
167	Amisulpride Tablets IP 200mg	Each uncoated tablet contains:				APPR
		Amisulpride	IP	200	mg	
		Excipients		q.s		
168	Amisulpride Tablets IP 300mg	Each uncoated tablet contains:				
		Amisulpride	IP	300	mg	
		Excipients		q.s		APPR
169	Amisulpride Tablets IP 400mg	Each uncoated tablet contains:				
		Amisulpride	IP	400	mg	
		Excipients		q.s		APPR
170	Flupentixol Tablets	Each film coated tablet contains:				
		Flupentixol Dihydrochloride Eq. to Flupentixol	BP	0.5	mg	
		Excipients		q.s		
		Approved colour used				APPR
171	Calcium & Vitamin D Tablets	Each uncoated tablet contains:				
		Calcium Citrate Malate Eq. to Elemental Calcium		250	mg	
		Vitamin D3	IP	200	IU	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
172	Pioglitazone Tablets IP 15mg (NOTE:- Package insert as per notification of ministry of H & FW No.:-GSR-520(E) Date:- 31/07/2013)	Each film coated tablet contains:				
		Pioglitazone Hydrochloride Eq. to Pioglitazone	IP	15	mg	
		Excipients		q.s		
		Approved colour used				APPR
173	Pioglitazone Tablets IP 30mg (NOTE:- Package insert as per notification of ministry of H & FW No.:-GSR-520(E) Date:- 31/07/2013)	Each film coated tablet contains:				
		Pioglitazone Hydrochloride Eq. to Pioglitazone	IP	30	mg	
		Excipients		q.s		
		Approved colour used				APPR
174	Hydroxyzine Tablets IP 10mg	Each film coated tablet contains:				
		Hydroxyzine Hydrochloride	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
175	Hydroxyzine Tablets IP 25mg	Each film coated tablet contains:				
		Hydroxyzine Hydrochloride	IP	25	mg	
		Excipients		q.s		
		Approved colour used				APPR
176	Chloramphenicol Capsules IP 250mg	Each hard gelatin capsule contains:				
		Chloramphenicol	IP	250	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
177	Chloramphenicol Capsules IP 500mg	Each hard gelatin capsule contains:				
		Chloramphenicol	IP	500	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
178	Cetirizine Tablets IP 10mg	Each uncoated tablet contains:				
		Cetirizine Hydrochloride	IP	10	mg	
		Excipients		q.s		APPR
179	Esomeprazole Tablets IP 20mg	Each enteric coated tablet contains:				
		Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole	IP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR
180	Esomeprazole Tablets IP 30mg	Each enteric coated tablet contains:				
		Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole	IP	30	mg	
		Excipients		q.s		
		Approved colour used				APPR
181	Esomeprazole Tablets	Each enteric coated tablet contains:				

	IP 40mg	Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole	IP	40	mg	
		Excipients		q.s		
		Approved colour used				APPR
182	Ramipril Tablets IP 1.25mg	Each uncoated tablet contains:				
		Ramipril	IP	1.25	mg	
		Excipients		q.s		APPR
183	Ramipril Tablets IP 2.5mg	Each uncoated tablet contains:				
		Ramipril	IP	2.5	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
184	Ramipril Tablets IP 5mg	Each uncoated tablet contains:				
		Ramipril	IP	5	mg	
		Excipients		q.s		APPR
185	Ramipril Tablets IP 10mg	Each uncoated tablet contains:				
		Ramipril	IP	10	mg	
		Excipients		q.s		APPR
186	Ascorbic Acid Tablets IP 500mg	Each uncoated tablet contains:				
		Ascorbic Acid	IP	500	mg	
		Excipients		q.s		APPR
187	Sertraline Tablets IP 25mg	Each film coated tablet contains:				
		Sertraline Hydrochloride Eq. to Sertraline	IP	25	mg	
		Excipients		q.s		
		Approved colour used				APPR
188	Sertraline Tablets IP 50mg	Each film coated tablet contains:				
		Sertraline Hydrochloride Eq. to Sertraline	IP	50	mg	
		Excipients		q.s		
		Approved colour used				APPR
189	Sertraline Tablets IP 100mg	Each film coated tablet contains:				
		Sertraline Hydrochloride Eq. to Sertraline	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
190	Pregabalin Sustained Release Tablets	Each uncoated tablet contains:				
		Pregabalin (In sustained release form)	IP	75	mg	
		Excipients		q.s		APPR
191	Pregabalin Sustained Release Tablets	Each uncoated tablet contains:				
		Pregabalin (In sustained release form)	IP	150	mg	
		Excipients		q.s		APPR
192	Etoricoxib Tablets IP 60mg	Each film coated tablet contains:				
		Etoricoxib	IP	60	mg	

		Excipients		q.s		
		Approved colour used				APPR
193	Etoricoxib Tablets IP 90mg	Each film coated tablet contains:				
		Etoricoxib	IP	90	mg	
		Excipients		q.s		
		Approved colour used				APPR
194	Amitriptyline Tablets IP 10mg	Each film coated tablet contains:				
		Amitriptyline Hydrochloride	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
195	Amitriptyline Tablets IP 25mg	Each film coated tablet contains:				
		Amitriptyline Hydrochloride	IP	25	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
196	Amitriptyline Tablets IP 50mg	Each film coated tablet contains:				
		Amitriptyline Hydrochloride	IP	50	mg	
		Excipients		q.s		
		Approved colour used				APPR
197	Gabapentin Tablets IP	Each film coated tablet contains:				
		Gabapentin	IP	300	mg	
		Excipients		q.s		
		Approved colour used				APPR
198	Aceclofenac SR Tablets	Each film coated sustained release tablet contains:				
		Aceclofenac	IP	200	mg	
		Excipients		q.s		
		Approved colour used				APPR
199	Duloxetine Hydrochloride Tablets	Each film coated tablet contains:				
		Duloxetine Hydrochloride Eq. to Duloxetine	USP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR
200	Duloxetine Hydrochloride Tablets	Each film coated tablet contains:				
		Duloxetine Hydrochloride Eq. to Duloxetine	USP	30	mg	
		Excipients		q.s		
		Approved colour used				APPR
201	Duloxetine Hydrochloride Tablets	Each film coated tablet contains:				
		Duloxetine Hydrochloride Eq. to Duloxetine	USP	40	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
202	Duloxetine	Each film coated tablet contains:				

	Hydrochloride Tablets	Duloxetine Hydrochloride Eq. to Duloxetine	USP	60	mg	
		Excipients		q.s		APPR
		Approved colour used				
203	Levocetirizine & Montelukast Dispersible Tablets	Each uncoated dispersible tablet contains :				
		Levocetirizine Hydrochloride	IP	2.5	mg	
		Montelukast Sodium Eq. to Montelukast	IP	4	mg	
		Excipients		q.s		
		Approved colour used				APPR
204	Desvenlafaxine Extended Release Tablets	Each uncoated extended release tablet contains:				
		Desvenlafaxine Succinate Eq. to Desvenlafaxine		50	mg	
		Excipients		q.s		APPR
205	Desvenlafaxine Extended Release Tablets	Each uncoated extended release tablet contains:				
		Desvenlafaxine Succinate Eq. to Desvenlafaxine		100	mg	
		Excipients		q.s		APPR
206	Albendazole & Ivermectin Tablets	Each uncoated tablet contains :				
		Albendazole	IP	400	mg	
		Ivermectin	IP	6	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
207	Diclofenac Potassium & Paracetamol Tablets	Each uncoated tablet contains :				
		Diclofenac Potassium	BP	50	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
208		Each film coated tablet contains:				
	Methylcobalamin, Alpha Lipoic Acid,Vitamin D3, Pyridoxine Hydrochloride & Folic Acid Tablets	Methylcobalamin	USP	1500	mcg	
		Alpha Lipoic Acid	USP	100	mg	
		Vitamin D3	IP	1000	IU	
		Pyridoxine Hydrochloride	IP	3	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colour used				APPR
209	Clarithromycin Oral Suspension	Each 5ml of reconstituted suspension contains:				
		Clarithromycin	IP	125	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR
210	Telmisartan Tablets IP 20mg	Each uncoated tablet contains:				
		Telmisartan	IP	20	mg	
		Excipients		q.s		APPR
211	Telmisartan Tablets IP 40mg	Each uncoated tablet contains:				
		Telmisartan	IP	40	mg	
		Excipients		q.s		APPR
212	Telmisartan Tablets IP 80mg	Each uncoated tablet contains:				
		Telmisartan	IP	80	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
213	Telmisartan & Amlodipine Tablets	Each uncoated tablet contains:				
		Telmisartan	IP	40	mg	
		Amlodipine Besylate Eq. to Amlodipine	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
214	Clobetasol Propionate Cream IP	Composition:				
		Clobetasol Propionate	IP	0.05%	w/w	
		Chlorocresol (as Preservative)	IP	0.1%	w/w	APPR
		In a Cream base				
215	Prochlorperazine Tablets IP 5mg	Each uncoated tablet contains:				
		Prochlorperazine Maleate	IP	5	mg	
		Excipients		q.s		APPR
216	Prochlorperazine Tablets IP 25mg	Each uncoated tablet contains:				
		Prochlorperazine Maleate	IP	25	mg	
		Excipients		q.s		APPR
217	Diclofenac Gel BP	Composition:				
		Diclofenac Diethylamine Eq. to Diclofenac Sodium	BP	1.16% 1%	w/w w/w	
		Gel Base				APPR
218	Clindamycin & Adapalene Gel	Composition:				
		Clindamycin Phosphate Eq. to Clindamycin	IP	10	mg	
		Adapalene	BP	1	mg	
		Gel Base				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
219	Cilnidipine Tablets	Each film coated tablet contains:				
		Cilnidipine		10	mg	
		Excipients		q.s		
		Approved colour used				APPR
220	Gabapentin & Methylcobalamin Tablets	Each film coated tablet contains :				
		Gabapentin	IP	300	mg	
		Methylcobalamin	USP	500	mcg	
		Excipients		q.s		
		Approved colour used				APPR
221	Diclofenac Potassium &	Each enteric coated tablet contains:				
		Diclofenac Potassium	BP	50	mg	
		Serratiopeptidase (EC)	IP	10	mg	
	Serratiopeptidase Tablets	(As 20,000 units of Serratiopeptidase)				APPR
		Excipients		q.s		
		Approved colour used				
222	Cefixime & Lactic Acid Bacillus Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	100	mg	
		Lactic Acid Bacillus		2.5	billion spores	
		Excipients		q.s		
		Approved colour used				APPR
223	Cefixime & Lactic Acid	Each uncoated dispersible tablet contains:				

	Bacillus Dispersible Tablets	Cefixime Eq. to Anhydrous Cefixime	IP	200	mg	
		Lactic Acid Bacillus		2.5	billion spores	
		Excipients		q.s		APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANT ITY**	**UNIT**	**APP**
224	Cefixime & Ofloxacin Tablets	Each film coated tablet contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	100	mg	
		Ofloxacin	IP	100	mg	
		Excipients		q.s		
		Approved colour used				APPR
225	Cephalexin Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	250	mg	
		Excipients		q.s		APPR
226	Rabeprazole Sodium (EC) & Levosulpiride	Each hard gelatin capsule contains:				
		Rabeprazole Sodium (As enteric Coated pellets)	IP	20	mg	
	(SR) Capsules	Levosulpiride (As sustained release pellets)		75	mg	APPR
		Excipients		q.s		
		Approved colours used in empty capsule shell.				
227	Nimesulide & Paracetamol Tablets (Not for Children below the age of 12 years)	Each uncoated tablet contains:				
		Nimesulide	BP	100	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		APPR
228	Febuxostat Tablets	Each film coated tablet contains :				
		Febuxostat		40	mg	
		Excipients		q.s		
		Approved colour used				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANTI TY**	**UNIT**	**APP**
229	Febuxostat Tablets	Each film coated tablet contains:				
		Febuxostat		80	mg	
		Excipients		q.s		
		Approved colour used				APPR
230	Febuxostat Tablets	Each film coated tablet contains:				
		Febuxostat		120	mg	
		Excipients		q.s		
		Approved colour used				APPR
231	Betahistine Tablets IP 16mg	Each uncoated tablet contains :				
		Betahistine Hydrochloride	IP	16	mg	
		Excipients		q.s		APPR
232	Metaxalone & Diclofenac Potassium	Each uncoated tablet contains :				
		Metaxalone		400	mg	

	Tablets	Diclofenac Potassium	BP	50	mg	
		Excipients		q.s		APPR
233		Each hard gelatin capsule contains:				
	Methylcobalamin, Alpha Lipoic Acid, Pyridoxine Hydrochloride, Folic Acid, Benfotiamine, Biotin & Chromium Capsules	Methylcobalamin	USP	1500	mcg	
		Alpha Lipoic Acid	USP	100	mg	
		Pyridoxine Hydrochloride	IP	3	mg	
		Folic Acid	IP	1.5	mg	
		Benfotiamine		50	mg	
		Biotin	USP	5	mg	
		Chromium Picolinate Eq. to Chromium	USP	200	mcg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
234	Fluconazole Tablets IP 50mg	Each uncoated tablet contains:				
		Fluconazole	IP	50	mg	
		Excipients		q.s		
		Approved colours used				APPR
235	Fluconazole Tablets IP 100mg	Each uncoated tablet contains:				
		Fluconazole	IP	100	mg	
		Excipients		q.s		
		Approved colours used				APPR
236	Fluconazole Tablets IP 150mg	Each uncoated tablet contains:				
		Fluconazole	IP	150	mg	
		Excipients		q.s		
		Approved colours used				APPR
237	Ketoconazole Tablets IP 200mg	Each uncoated tablet contains :				
		Ketoconazole	IP	200	mg	
		Excipients		q.s		APPR
238	Lornoxicam Tablets	Each film coated tablet contains :				
		Lornoxicam	IP	8	mg	
		Excipients		q.s		
		Approved colour used				APPR
239	Methylprednisolone Tablets IP	Each film coated tablet contains:				
		Methylprednisolone	IP	8	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
240	Telmisartan & Hydrochlorothiazide Tablets	Each uncoated tablet contains:				
		Telmisartan	IP	40	mg	
		Hydrochlorothiazide	IP	12.5	mg	
		Excipients		q.s		APPR
241	Escitalopram Tablets IP 20mg	Each film coated tablet contains:				
		Escitalopram Oxalate Eq. to Escitalopram	IP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR
242	Secnidazole Tablets IP	Each film coated tablet contains :				
		Secnidazole	IP	1	gm	
		Excipients		q.s		
		Approved colour used				APPR
243	Voglibose & Metformin	Each uncoated tablet contains:				

	HCl Tablets	Voglibose	IP	0.2	mg	
		Metformin Hydrochloride	IP	500	mg	
		Excipients				APPR
244	Voglibose & Metformin HCl Tablets	Each uncoated tablet contains :				
		Voglibose	IP	0.3	mg	
		Metformin Hydrochloride	IP	500	mg	
		Excipients		q.s		APPR
245		Each uncoated tablet contains:				
	Amlodipine Besylate & Atenolol Tablets	Amlodipine Besylate Eq.to Amlodipine	IP	5	mg	
		Atenolol	IP	50	mg	
		Excipients		q.s		A
		Approved colour used				

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
246	Losartan Potassium & Amlodipine Besylate Tablets IP	Each film coated tablet contains:				
		Losartan Potassium	IP	50	mg	
		Amlodipine Besylate Eq. to Amlodipine	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
247	Ferrous Ascorbate & Folic Acid Tablets	Each film coated tablet contains:				
		Ferrous Ascorbate Eq. to Elemental Iron		100	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colour used				APPR
248	Linezolid Tablets IP	Each film coated tablet contains :				
		Linezolid	IP	600	mg	
		Excipients		q.s		
		Approved colours used				APPR
249	Deflazacort Tablets	Each uncoated tablet contains:				
		Deflazacort		1	mg	
		Excipients		q.s		APPR
250	Deflazacort Tablets	Each uncoated tablet contains :				
		Deflazacort		6	mg	
		Excipients		q.s		APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
251	Deflazacort Tablets	Each uncoated tablet contains:				
		Deflazacort		30	mg	
		Excipients		q.s		APPR
252	Levetiracetam Tablets USP	Each film coated tablet contains:				
		Levetiracetam	IP	500	mg	
		Excipients		q.s		APPR
253	Deflazacort Tablets	Each uncoated tablet contains:				
		Deflazacort		18	mg	
		Excipients		q.s		APPR
254	Deflazacort Oral Suspension	Each 5ml contains :				
		Deflazacort		6	mg	
		In a flavoured base		q.s		

		Approved colour used				APPR
255	Aceclofenac, Paracetamol & Serratiopeptidase Tablets	Each film coated tablet contains:				
		Aceclofenac	IP	100	mg	
		Paracetamol	IP	325	mg	
		Serratiopeptidase (EC) (as 20,000 units of Serratiopeptidase)	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
256	Aceclofenac, Paracetamol & Serratiopeptidase Tablets	Each film coated tablet contains:				
		Aceclofenac	IP	100	mg	
		Paracetamol	IP	325	mg	
		Serratiopeptidase (EC) (as 30,000 units of Serratiopeptidase)	IP	15	mg	
		Excipients		q.s		APP
		Approved colour used				
257	Calcium Citrate Malate, Calcitriol,& zinc Tablets	Each film coated tablet contains:				
		Calcium Citrate Malate Eq. to Elemental Calcium		250	mg	
		Calcitriol	IP	0.25	mcg	
		Zinc Sulphate Monohydrate Eq. to Elemental Zinc	IP	7.5	mg	
		Excipients		q.s		
		Approved colour used				APP
258	Doxylamine Succinate, Pyridoxine HCl & Folic Acid Tablets	Each enteric coated tablet ontains:				
		Doxylamine Succinate	USP	10	mg	
		Pyridoxine HCl	IP	10	mg	
		Folic Acid	IP	2.5	mg	
		Excipients		q.s		
		Approved colour used				
259	Calcium Citrate, Magnesium, Zinc & Vitamin D3 Tablets	Each uncoated tablet contains:				
		Calcium Citrate	USP	1000	mg	
		Magnesium Hydroxide Eq. to Elemental Magnesium	IP	100	mg	
		Zinc Sulphate Monohydrate Eq. to Elemental Zinc	IP	4	mg	
		Vitamin D3	IP	200	IU	
		Excipients		q.s		APP

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
260	Ferrous Ascorbate & Folic Acid Tablets.	Each film coated tablet contains:				
		Ferrous Ascorbate Eq. to Elemental Iron		100	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colour used				APP
261	Calcium Citrate, Calcitriol & zinc Tablets	Each film coated tablet contains:				
		Calcium Citrate	USP	1000	mg	
		Calcitriol	IP	0.25	mcg	
		Zinc Sulphate	IP	7.5	mg	
		Excipients		q.s		
		Approved colour used				APP
262	Levofloxacin	Each 5 ml contains:				

	suspension	Levofloxacin Hemihydrate Eq. to Levofloxacin	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APP
263	Pantoprazole Sodium & Domperidone Tablets	Each enteric coated tablet ontains:				
		Pantoprazole Sodium Eq. to Pantoprazole	IP	40	mg	
		Domperidone	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APP
264	Aceclofenac, Paracetamol & Chlorzoxazone Tablets	Each film coated tablet contains:				
		Aceclofenac	IP	100	mg	
		Paracetamol	IP	325	mg	
		Chlorzoxazone	USP	250	mg	
		Excipients		q.s		
		Approved colour used				APP
265	Clobetasol Propionate, Neomycin & Miconazole Nitrate Cream	Composition:				
		Clobetasol Propionate	IP	0.05%	w/w	
		Neomycin Sulphate	IP	0.5%	w/w	
		Miconazole Nitrate	IP	2.0%	w/w	
		Chlorocresol (as Preservatives)	IP	0.1%	w/w	
		Cream Base		q.s		APP

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
266	Loratadine Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Loratadine	USP	10	mg	
		Excipients		q.s		APP
267	Povidon-Iodine Ointment USP	Composition:				
		Povidon-Iodine (Available Iodine 0.5% w/w)	IP	5%	w/w	
		Water Soluble Ointment base.				APP
268	Clindamycin & Niacinamide Gel.	Composition:				
		Clindamycin (as Clindamycin Phosphate IP)		1%	w/w	
		Niacinamide	IP	4%	w/w	
		Gel base.				APP
269	Diclofenac Potassium, Paracetamol & Chlorzoxazone Tablets	Each film coated tablet contains:				
		Diclofenac Potassium	BP	50	mg	
		Paracetamol	IP	325	mg	
		Chlorzoxazone	USP	250	mg	
		Excipients		q.s		
		Approved colour used				APP
270	Cyproheptadine HCl, Tricholine Citrate & Sorbitol Syrup	Each 5 ml contains:				
		Cyproheptadine HCl (Anhydrous)	IP	2	mg	
		Tricholine Citrate		0.275	gm	
		Sorbitol Solution (70%) Non Crystallising	IP	2	gm	
		In a flavoured Syrupy base		q.s		
		Approved colour used				APP

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
271	Drotaverine HCl &	Each film coated tablet contains:				

	Mefenamic Acid Tablets	Drotaverine HCl	IP	80	mg	
		Mefenamic Acid	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APP
272	Aceclofenac & Paracetamol Oral Suspension	Each 5 ml contains:				
		Aceclofenac	IP	50	mg	
		Paracetamol	IP	125	mg	
		In a flavoured base		q.s		
		Approved colour used				APP
273	Magaldrate & Simethicone Oral Suspension USP	Each 5 ml contains:				
		Magaldrate (Anhydrous)	IP	480	mg	
		Simethicone	IP	50	mg	
		In a flavoured base		q.s		
		Approved colour used				APP
274	Dicyclomine HCl & Paracetamol Tablets	Each uncoated tablet contains:				
		Dicyclomine HCl	IP	20	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		APP
275	Terbutaline Sulphate, Bromohexine HCl & Guaiphenesin Syrup	Each 5ml contains :				
		Terbutaline Sulphate	IP	1.25	mg	
		Bromohexine Hydrochloride	IP	2	mg	
		Guaiphenesin	IP	50	mg	
		In a flavoured Syrupy base		q.s		
		Approved colour used				APP

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
276	Tannic Acid Glycerin, Potassium Iodide, Iodine, Menthol, Thymol & Glycerin, Gumpent	Composition:				
		Tannic Acid Glycerin		27%	w/v	
		Potassium Iodide	IP	0.05%	w/v	
		Iodine	IP	0.03%	w/v	
		Menthol	IP	0.05%	w/v	
		Thymol	IP	0.033%	w/v	
		Glycerin	IP	72%	w/v	APP
277	Povidon-Iodine & Metronidazole Ointment	Composition:				
		Povidon-Iodine (Available Iodine 0.5% w/w)	IP	5%	w/w	
		Metronidazole	IP	1%	w/w	
		Water soluble ointment base.				APP
278	Clobetasol Propionate & Zinc Sulphate Cream	Composition:				
		Clobetasol Propionate	IP	0.05%	w/w	
		Zinc Sulphate	IP	2.5%	w/w	
		Cream Base				APP
279	Piracetam Tablets	Each film coated tablet Contains:				
		Piracetam	IP	800	mg	
		Excipients		q.s		
		Approved colour used				APP
280	Piracetam Syrup	Each 5ml contains :				
		Piracetam	IP	500	mg	
		In a flavoured Syrupy base		q.s		
		Approved colour used				APP
281	Tricholine Citrate	Each 10 ml Contains:				

	Syrup	Tricholine Citrate		0.55	gm	
		Sorbitol (70%) Non Crystallising	IP	q.s		
		In a flavoured Palatable base				
		Approved colour used				APP

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
282	Diclofenac Potassium, Paracetamol & Serratiopeptidase Tablets	Each film coated tablet contains:				
		Diclofenac Potassium	IP	50	mg	
		Paracetamol	IP	325	mg	
		Serratiopeptidase (EC) (as 20,000 units of Serratiopeptidase)	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APP
283	Mefenamic Acid & Tranexamic Acid Tablets	Each uncoated tablet contains:				
		Mefenamic	IP	250	mg	
		Tranexamic Acid	BP	500	mg	
		Excipients		q.s		APP
284	Torsemide Tablets	Each uncoated tablet contains:				
		Torsemide	USP	10	mg	
		Excipients		q.s		
		Approved colour used				APP
285	Torsemide Tablets	Each uncoated tablet contains:				
		Torsemide	USP	20	mg	
		Excipients		q.s		
		Approved colour used				APP
286	Sodium Picosulphate Syrup	Each 5 ml contains:				
		Sodium Picosulphate	BP	5	mg	
		In a Palatable Sorbitol base		q.s		
		Approved colour used				APP
287	Potassium Nitrate & Fluoride Medicated foaming Dental Gel	Composition:				
		Potassium Nitrate	BP	5%	w/w	
		Sodium Monofluorophosphate	USP	0.7%	w/w	
		Approved colour used				APP

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
288	Amlodipine & Atenolol Tablets	Each uncoated tablet contains:				
		Amlodipine Besylate Eq. to Amlodipine	IP	5	mg	
		Atenolol	IP	50	mg	
		Excipients		q.s		
		Approved colour used				APP
289	Cholecalciferol Sachet	Each Sachet of 1 g. Contains:				
		Cholecalciferol	IP	60,000	IU	
		Excipients		q.s		APP
290	Cefixime Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	100	mg	
		Excipients		q.s		APP
291	Cefixime Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	200	mg	

		Excipients		q.s		APP
292	Cefpodoxime Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	100	mg	
		Excipients		q.s		APP
293	Cefpodoxime Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	50	mg	
		Excipients		q.s		APP

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
294	Sodium Feredetate, Folic Acid & Vitamin B12 Tablets	Each film coated tablet contains:				
		Sodium Feredetate Eq. to Elemental Iron	BP	231 33	mg mg	
		Folic Acid	IP	1.5	mg	
		Vitamin B12	IP	15	mcg	
		Excipients		q.s		APP
		Approved colour used				
295	Carbonyl Iron , Folic Acid & Zinc Capsules	Each hard gelatin capsule contains:				
		Carbonyl Iron Eq. to. Elemental Iron		60	mg	
		Folic Acid	IP	1.5	mg	
		Zinc Sulphate Monohydrate (Eq. to elemental Zinc 22.5 mg)	IP	61.8	mg	
		Approved Colour used in empty Capsule shells.				APP
296	Cephalexin Dispersible Tablets	Each uncoated dispersible tablet contains				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APP
297	Cephalexin Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	250	mg	
		Excipients		q.s		APP
298	Cephalexin Tablets IP	Each film coated tablet contains:				
		Cephalexin Eq. to Anhydrous Cephalexin	IP	500	mg	
		Excipients		q.s		
		Approved colour used				APP

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	API
299	Calcium & Vitamin D3 Suspension	Each 5 ml contains:				
		Calcium Carbonate from an organic source (Oyster shell) Eq. to Elemental Calcium		250	mg	
		Vitamin D3	IP	125	IU	
		In a flavoured base		q.s		
		Approved colour used				APP
300		Each uncoated tablet contains:				
	Calcium Citrate & Calcitriol Tablets	Calcium Citrate	USP	1000	mg	
		Calcitriol	IP	0.25	mcg	

		Excipients		q.s		APP
301	Light Liquid Paraffin & Milk of Magnesia Laxative Oral Suspension	Each 15ml contains :				
		Light Liquid Paraffin	IP	3.75	ml	
		Milk of Magnesia	IP	11.25	ml	
		In a flavoured base		q.s.		
		Approved colour used				APP
302	Iron, Cyanocobalamin & Folic Acid Syrup	**FOR THERAPEUTIC USE**				
		Each 15ml contains :				
		Ferric Ammonium Citrate (Eq.to Elemental Iron 32mg)	IP	160	mg	
		Cyanocobalamin	IP	7.5	mcg	
		Folic Acid	IP	0.5	mg	
		In a flavoured syrupy base		q.s.		
		Approved colour used				APP
303	Beclomethasone Dipropionate, Clotrimazole & Neomycin Cream	Composition :				
		Beclomethasone Dipropionate	IP	0.025%	w/w	
		Clotrimazole	IP	1%	w/w	
		Neomycin Sulphate Eq. to Neomycin	IP	0.5%	w/w	
		Chlorocresol (as preservative)	IP	0.1%	w/w	
		In a cream base				APP

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
304	Clobetasol Propionate & Salicylic Acid Ointment	Composition:				
		Clobetasol Propionate	IP	0.05%	w/w	
		Salicylic Acid	IP	6.0%	w/w	
		In an Ointment base		q.s		APPR
305	Vitamin E & Aloe Vera Cream	Composition:				
		Aloe extract	IP	10%	w/w	
		Vitamin E Acetate	IP	0.5%	w/w	
		Moisturising Cream base		q.s		APPR
306	Omega 3 Fatty Acid,	Each hard gelatin capsule contains:				
		Alpha Lipoic Acid	USP	50	mg	
		Methylcobalamin	USP	750	mcg	
	Methylcobalamin, Alpha Lipoic Acid, Chromium Polynicotinate & Selenium Dioxide Capsules	Omega 3 Fatty Acid		500	mg	
		Chromium Polynicotinate		200	mcg	
		Selenium Dioxide Mohohydrate	USP	75	mcg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
307	Mefenamic Acid & Dicyclomine HCl Tablets	Each uncoated tablet contains:				
		Mefenamic Acid	IP	250	mg	
		Dicyclomine HCl	IP	10	mg	
		Excipients		q.s		APPR
308	Ciprofloxacin HCl & Tinidazole Tablets	Each film coated tablet contains:				
		Ciprofloxacin HCl Eq.to Ciprofloxacin	IP	500	mg	
		Tinidazole	IP	600	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP

309	Dried Aluminium Hydroxide, Magnesium Hydroxide, Activated Dimethicone & Sorbitol Suspension	Each 5ml contains :				
		Dried Aluminium Hydroxide Gel	IP	250	mg	
		Magnesium Hydroxide	IP	250	mg	
		Activated Dimethicone	IP	50	mg	
		Sorbitol Solution 70% (Non-crystallising)	IP	0.65	gm	
		In a flavoured base		q.s		
		Approved colour used				APPR
310	Sodium Valproate & Valproic Acid Controlled Release Tablets	Each film coated controlled release tablet contains :				
		Sodium Valproate	IP	133	mg	
		Valproic Acid (Both together correspond to Sodium Valproate 200 mg)	IP	58	mg	APPR
		Excipients		q.s		
		Approved colour used				
311	Sodium Valproate & Valproic Acid Controlled Release Tablets	Each film coated controlled release tablet contains :				
		Sodium Valproate	IP	200	mg	
		Valproic Acid (Both together correspond to Sodium Valproate 300 mg)	IP	87	mg	
		Excipients		q.s		
		Approved colour used				APPR
312	Sodium Valproate & Valproic Acid Controlled Release Tablets	Each film coated controlled release tablet contains :				
		Sodium Valproate	IP	333	mg	
		Valproic Acid (Both together correspond to Sodium Valproate 500 mg)	IP	145	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
313	Gliclazide & Metformin HCl Tablets	Each uncoated tablet contains:				
		Gliclazide	IP	80	mg	
		Metformin HCl	IP	500	mg	
		Excipients		q.s		APPR
314	Sucralfate Suspension	Each 10 ml contains:				
		Sucralfate	USP	1	gm	
		In a flavoured base		q.s		
		Approved colour used				APPR
315	Disodium Hydrogen Citrate Syrup	Each 5ml contains :				
		Disodium Hydrogen Citrate	BP	1.25	gm	
		In a flavoured syrupy base		q.s.		
		Approved colour used				APPR
316	Fusidic Acid & Beclomethasone Dipropionate Cream	Composition :				
		Fusidic Acid	IP	2%	w/w	
		Beclomethasone Dipropionate	IP	0.025%	w/w	APPR
		Cream base		q.s		
317	Chlorhexidine Gluconate, Sodium Fluoride & Zinc Chloride Mouthwash	Composition :				
		Chlorhexidine Gluconate Solution Eq.to Chlorhexidine Gluconate	IP	0.20%	w/v	
		Sodium Fluoride	IP	0.05%	w/v	

		Zinc Chloride	IP	0.09%	w/v	
		In a pleasantly flavoured base				
		Approved colour used				APPR
318	L-Ornithine-L-Aspartate Sachets	Each Sachet (5gm) contains:				
		L-Ornithine -L-Aspartate		3	gm	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
319	Terbutaline Sulphate, Bromhexine Hydrochloride, Guaiphenesin & Menthol Syrup	Each 5ml contains :				
		Terbutaline Sulphate	IP	1.25	mg	
		Bromhexine Hydrochloride	IP	4	mg	
		Guaiphenesin	IP	50	mg	
		Menthol	IP	2.5	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used				APPR
320	Paracetamol, Diclofenac Potassium & Chlorzoxazone Tablet	Each film coated tablet contains :				
		Paracetamol	IP	325	mg	
		Diclofenac Potassium	BP	50	mg	
		Chlorzoxazone	USP	500	mg	
		Excipients		q.s		
		Approved colour used				APPR
321	Rabeprazole Sodium & Domperidone Tablets	Each enteric coated tablet contains:				
		Rabeprazole Sodium	IP	20	mg	
		Domperidone	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
322	Clindamycin &	Composition :				
	Adapalene Gel	Clindamycin Phosphate Eq. to Clindamycin	IP	1%	w/w	
		Adapalene	BP	0.1%	w/w	
		Gel base		q.s		
		Preservatives :				
		Methyl Paraben Phenoxyethanol	IP IP	0.1% 0.25%	w/w w/w	APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
323	Beclomethasone Dipropionate, Miconazole Nitrate & Neomycin Sulphate Cream	Composition:				
		Beclomethasone Dipropionate	IP	0.025%	w/w	
		Miconazole Nitrate	IP	2.0%	w/w	
		Neomycin Sulphate	IP	0.5%	w/w	
		In a cream base		q.s.		
		Preservatives :				
		Methyl Paraben Propyl Paraben	IP IP	0.2% 0.1%	w/w w/w	APPR
324	Ubidecarenone, L-Arginine, Alpha Tocopheryl Acetate & Selenium Capsules	Each hard gelatin capsule contains:				
		Ubidecarenone (Coenzyme Q 10)	USP	100	mg	
		L-Arginine	IP	100	mg	
		Alpha Tocopheryl Acetate	IP	25	IU	
		Selenium (As Sodium selenate)		100	mcg	
		Excipients		q.s		

		Approved colours used in empty capsule shell.				APPR
325	Sucralfate & Oxetacaine Suspension	Each 10 ml contains :				
		Sucralfate	USP	1000	mg	
		Oxetacaine	BP	20	mg	
		In a flavoured base		q.s		
		Approved colour used				APPR
326	Diclofenac Potassium, Paracetamol & Serratiopeptidase Tablets	Each film coated tablet contains:				
		Diclofenac Potassium	BP	50	mg	
		Paracetamol	IP	325	mg	
		Serratiopeptidase (EC) (As 30,000 units of Serratiopeptidase)	IP	15	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
327	Lyophilized Saccharomyces Boulardii, Lactic Acid Bacillus & Racecadotril Sachets	Each sachet contains:				
		Lyophilized Saccharomyces Boulardii (Corresponding to 125 mg of Yeast)		2.5	billion	
		Lactic Acid Bacillus		100	million spores	
		Racecadotril	IP	10	mg	
		Excipients		q.s		APPR
328	Lyophilized Saccharomyces Boulardii, Lactic Acid Bacillus & Zinc Sachets	Each sachet contains:				
		Lyophilized Saccharomyces Boulardii (Corresponding to 125 mg of Yeast)		2.5	billion	
		Lactic Acid Bacillus		150	million spores	
		Elemental Zinc (As Zinc Lactate)		7.5	mg	
		Excipients		q.s		APPR
329	Cefpodoxime Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	200	mg	
		Excipients		q.s		APPR
330	Cefixime Dispersible Tablets	Each uncoated dispersible tablet contains:				
		Cefixime Eq. to Anhydrous Cefixime	IP	50	mg	
		Excipients		qs		
		Approved colour used				APPR
331	Cefpodoxime Dispersible Tablets.	Each uncoated dispersible tablet contains:				
		Cefpodoxime Proxetil Eq. to Cefpodoxime	IP	50	mg	
		Excipients		qs		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
332	Ondansetron Oral	Each 5ml contains :				

	Solution IP .	Ondansetron Hydrochloride Eq. to Ondansetron	IP	2	mg	
		In a flavoured syrupy base		q.s.		
		Approved colour used				APPR
333	Magaldrate & Simethicone Oral Suspension USP	Each 5ml contains :				
		Magaldrate (Anhydrous)	IP	400	mg	
		Simethicone	IP	20	mg	
		In a flavoured sugar free base				
		Approved colour used				APPR
334	Lactulose Solution USP	Each 15ml contains :				
		Lactulose (As Lactulose Concentrate USP)		10	gm	
		Palatable base		q.s		APPR
335	Azithromycin Oral Suspension IP	Each 5ml contains :				
		Azithromycin Dihydrate Eq. to Anhydrous Azithromycin	IP	200	mg	
		In a flavoured base		q.s		
		Approved colour used				APPR
336	Trioxsalen Tablets USP 25 mg	Each film coated tablet contains :				
		Trioxsalen	USP	25	mg	
		Excipients		q.s		
		Approved colour used				APPR
337	Racecadotril Sachets IP 30mg .	Each sachet contains:				
		Racecadotril	IP	30	mg	
		Excipients		q.s		
		Approved colour used				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFICATION**	**QUANTITY**	**UNIT**	**APP**
338	Roxithromycin Tablets IP 300mg	Each film coated tablet contains:				
		Roxithromycin	IP	300	mg	
		Excipients		q.s		
		Approved colour used				APPR
339	Amlodipine Tablets IP 10mg	Each uncoated tablet contains:				
		Amlodipine Besylate Eq. to Amlodipine	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
340	Lamotrigine Dispersible Tablets IP 25mg	Each uncoated dispersible tablet contains:				
		Lamotrigine	IP	25	mg	
		Excipients		q.s		APPR
341	Lamotrigine Dispersible Tablets IP 50mg	Each uncoated dispersible tablet contains:				
		Lamotrigine	IP	50	mg	
		Excipients		q.s		APPR
342	Fexofenadine Tablets IP 120mg.	Each film coated tablet contains :				
		Fexofenadine Hydrochloride	IP	120	mg	
		Excipients		q.s		
		Approved colour used				APPR
343	Montelukast Tablets IP 10mg	Each film coated tablet contains:				
		Montelukast Sodium Eq. to Montelukast	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
344	Montelukast Tablets IP 5mg	Each film coated tablet contains:				
		Montelukast Sodium Eq. to Montelukast	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
345	Finasteride Tablets IP	Each film coated tablet contains:				
		Finasteride	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
346	Ketoconazole 2% Cream	Composition :				
		Ketoconazole	IP	2%	w/w	
		In a cream base				APPR
347	Clarithromycin Tablets IP 250mg	Each uncoated tablet contains:				
		Clarithromycin	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APPR
348	Clarithromycin Tablets IP 500mg	Each uncoated tablet contains:				
		Clarithromycin	IP	500	mg	
		Excipients		q.s		APPR
349	Ranitidine Tablets IP 150mg .	Each film coated tablet contains :				
		Ranitidine Hydrochloride Eq. to Ranitidine	IP	150	mg	
		Excipients		q.s		
		Approved colour used				APPR

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S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
350	Lornoxicam & Paracetamol Tablets	Each film coated tablet contains:				
		Lornoxicam	IP	4	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
351	Lornoxicam & Paracetamol Tablets	Each film coated tablet contains:				
		Lornoxicam	IP	8	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				APPR
352	Etodolac & Paracetamol Tablets	Each film coated tablet contains:				
		Etodolac	IP	400	mg	
		Paracetamol	IP	325	mg	
		Excipients		q.s		
		Approved colour used				REJE
353	Mupirocin Ointment IP	Composition :				
		Mupirocin	IP	2%	w/w	
		Ointment base		q.s		APPR
354	Tacrolimus Ointment	Composition :				
		Tacrolimus	IP	0.1%	w/w	
		Ointment base		q.s		APPR
355	Montelukast & Desloratadine Tablet	Each filmcoated tablet contains :				
		Montelukast Sodium Eq. to Montelukast	IP	10	mg	
		Desloratadine		5	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
356	Rabeprazole Sodium (EC) & Aceclofenac (SR) Capsules	Each hard gelatin capsule contains:				
		Rabeprazole Sodium (As enteric Coated pellets)	IP	20	mg	
		Aceclofenac (As sustained release pellets)	IP	200	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
357	Paracetamol, Phenylephrine HCl, Caffeine & Diphenhydramine HCl Tablets	Each film coated tablet contains:				
		Paracetamol	IP	325	mg	
		Phenylephrine HCl	IP	5	mg	
		Caffeine (Anhydrous)	IP	30	mg	
		Diphenhydramine HCl	IP	25	mg	
		Excipients		q.s		
		Approved colour used				APPR
358	Cefadroxil Capsules IP 500mg	Each hard gelatin capsule contains:				
		Cefadroxil Eq .to Anhydrous Cefadroxil	IP	500	mg	
		Excipients		q.s		
		Approve colour used in empty capsule shell				APPR
359	Cholecalciferol Chewable Tablets	Each uncoated chewable tablet contains:				
		Cholecalciferol	IP	60,000	IU	
		Excipients		q.s		
		Approved colour used				APPR
360	Cefuroxime Axetil & Potassium Clavulanate Tablets	Each film coated tablet contains :				
		Cefuroxime Axetil Eq. to Cefuroxime	IP	500	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
361	Terbutaline Sulphate, Bromhexine Hydrochloride & Guaiphenesin Syrup	Each 5ml contains :				
		Terbutaline Sulphate	IP	1.25	mg	
		Bromhexine Hydrochloride	IP	2	mg	
		Guaiphenesin	IP	25	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used				APPR
362	Clindamycin Phosphate Gel USP	Composition:				
		Clindamycin Phosphate Eq. to Clindamycin	IP	1%	w/w	
		Gel base		q.s		
		Preservatives :				
		Methyl Paraben Propyl Paraben	IP IP	0.2% 0.02%	w/w w/w	APPR
363	Ketorolac Tromethamine Tablets IP 10mg	Each film coated tablet contains:				
		Ketorolac Tromethamine	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR

364	Dextromethorphan Hydrobromide, Chlorpheniramine Maleate, Guaiphenesin & Ammonium Chloride Syrup.	Each 5ml contains :				
		Dextromethorphan Hydrobromide	IP	5	mg	
		Chlorpheniramine Maleate	IP	2.5	mg	
		Guaiphenesin	IP	50	mg	
		Ammonium Chloride	IP	60	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used				APPR
365	Amoxycillin & Potassium Clavulanate Oral Suspension IP	Each 5ml of the reconstituted suspension contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	200	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	28.5	mg	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
366	Amoxycillin & Potassium	Each 5ml of the reconstituted suspension contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	400	mg	
	Clavulanate Oral Suspension IP	Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	57	mg	APPR
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				
367	Ciprofloxacin Hydrochloride Powder 10% w/w. (for Veterinary use Only)	Composition:				
		Ciprofloxacin Hydrochloride eq. to Anhydrous Ciprofloxacin	IP	10%	w/w	
		Excipients		q.s		APPR
368	Levofloxacin Water Soluble Powder 10% w/w (for Veterinary use Only)	Each gram Contains:				
		Levofloxacin Hemihydrate eq. to Levofloxacin	IP	100	mg	
		Excipients		q.s		APPR
369	Amoxycillin, Potassium Clavulanate & Lactic Acid Bacillus Tablets	Each film coated tablet contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	500	mg	
		Potassium Clavulanate Diluted Eq. to Clavulanic Acid	IP	125	mg	
		Lactic Acid Bacillus		60	million spores	
		Excipients		q.s		
		Approved colour used				APPR
370	Paracetamol Paediatric Oral Suspension IP	Each 5ml contains :				
		Paracetamol	IP	125	mg	
		In a flavoured base		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP

371	Paracetamol Paediatric Oral Suspension IP	Each 5ml contains :				
		Paracetamol	IP	250	mg	
		In a flavoured base		q.s		
		Approved colour used				APPR
372		Each film coated tablet contains:				
	Etoricoxib & Thiocolchicoside Tablets	Etoricoxib	IP	60	mg	
		Thiocolchicoside	IP	4	mg	APPR
		Excipients		q.s		
		Approved colour used				
373	Etoricoxib & Thiocolchicoside Tablets	Each film coated tablet contains:				
		Etoricoxib	IP	60	mg	
		Thiocolchicoside	IP	8	mg	
		Excipients		q.s		
		Approved colour used				APPR
374	Cetirizine Hydrochloride, Paracetamol & Phenylephrine Hydrochloride Tablets.	Each uncoated tablet contains:				
		Cetirizine Hydrochloride	IP	5	mg	
		Paracetamol	IP	325	mg	
		Phenylephrine Hydrochloride	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
375	Dextromethorphan Hydrobromide, Phenylephrine Hydrochloride & Chlorpheniramine Maleate Syrup	Each 5ml contains :				
		Dextromethorphan Hydrobromide	IP	10	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used				APPR
376	Nimesulide Dispersible Tablets (Not for Children below the age of 12 years)	Each uncoated dispersible tablet contains:				APPR
		Nimesulide	BP	100	mg	
		Excipients		q.s		

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
377	Paracetamol, Phenylephrine Hydrochloride & Chlorpheniramine Maleate Suspension	Each 5ml contains :				
		Paracetamol	IP	250	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		In a flavoured base				
		Approved colour used				APPR
378		Each 5ml contains :				
	Paracetamol, Phenylephrine Hydrochloride & Chlorpheniramine Maleate Suspension	Paracetamol	IP	125	mg	
		Phenylephrine Hydrochloride	IP	2.5	mg	
		Chlorpheniramine Maleate	IP	1	mg	
		In a flavoured base				
		Approved colour used				APPR
379	Amoxycillin, Dicloxacillin & Lactobacillus Capsules	Each hard gelatin capsule contains:				
		Amoxycillin Trihydrate Eq. to Amoxycillin	IP	250	mg	
		Dicloxacillin Sodium Eq. to Dicloxacillin	IP	250	mg	
		Lactobacillus		2.5	million spores	
		Excipients		q.s		

		Approved colours used in empty capsule shell.				APPR
380	Methylcobalamin, Alpha Lipoic Acid, Pyridoxine Hydrochloride & Folic Acid Capsules	Each hard gelatin capsule contains:				
		Methylcobalamin	USP	1500	mcg	
		Alpha Lipoic Acid	USP	100	mg	
		Pyridoxine Hydrochloride	IP	3	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shell.				APPR
381	Finasteride Tablets IP	Each film coated tablet contains :				
		Finasteride	IP	1	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
382	Ofloxacin & Ornidazole Suspension	Each 5ml contains :				
		Ofloxacin	IP	50	mg	
		Ornidazole	IP	125	mg	
		In a flavoured base		q.s		APPR
		Approved colour used				Bann
383	Paracetamol, Levocetirizine Hydrochloride, Phenylephrine &	Each film coated tablet contains:				
		Paracetamol	IP	325	mg	
		Levocetirizine Hydrochloride	IP	2.5	mg	
		Phenylephrine Hydrochloride	IP	10	mg	
	Caffeine Tablets	Caffeine (Anhydrous)	IP	30	mg	
		Excipients		q.s		
		Approved colour used				APPR
384	Lycopene With Multi Vitamin & Minerals Syrup	FOR PROPHYLACTIC USE				
		Each 10 ml contains:				
		Lycopene (10%)	USP	1000	mcg	
		Vitamin A concentrate (Oily form) (as Palmitate)	IP	2500	IU	
		Vitamin E (as Aacetate)	IP	10	IU	
		Ascorbic Acid (Vitamin C)	IP	50	mg	
		Thiamine HCl (Vitamin B1)	IP	2	mg	
		Riboflavin (Vitamin B2)	IP	3	mg	
		PyridoxineHCl (Vitamin B6)	IP	1.5	mg	
		Selenium (as Sodium Selenate)		35	mcg	
		Zinc (as Zinc Gluconate USP)		3	mg	
		Manganese (as Manganese Gluconate USP)		2	mg	
		Iodine (as Potassium Iodide IP)		100	mcg	
		Copper(as CupricSulphateUSP)		500	mcg	
		In a flavoured syrupy base		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
385	Ofloxacin, Ornidazole, Terbinafine Hydrochloride & Clobetasol Propionate Cream	Composition:				
		Ofloxacin	IP	0.75%	w/w	
		Ornidazole	IP	2%	w/w	
		Terbinafine Hydrochloride	BP	1%	w/w	
		Clobetasol Propionate	IP	0.05%	w/w	

		Methylparaben (As Preservative)	IP	0.2%	w/w	
		Propylparaben (As Preservative)	IP	0.02%	w/w	APPR
		In a non-greasy base				Bann
386	Azithromycin Oral Suspension IP	Each 5ml contains :				
		Azithromycin Dihydrate	IP			
		Eq.to Anhydrous Azithromycin		100	mg	APPR
		In a flavoured base		q.s		
		Approved colour used				
387	Paracetamol & Mefenamic Acid Suspension	Each 5ml contains :				
		Paracetamol	IP	125	mg	
		Mefenamic Acid	IP	50	mg	
		In a flavoured base		q.s		
		Approved colour used				APPR
388	Pantoprazole Sodium & Domperidone Tablets	Each enteric coated tablet contains:				
		Pantoprazole Sodium Eq. to Pantoprazole	IP	20	mg	
		Domperidone	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
389	S-Amlodipine Tablets IP 2.5 mg	Each uncoated tablet contains:				APPR
		S-Amlodipine Besylate Eq. to S-Amlodipine	IP	2.5	mg	
		Excipients		q.s		

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
390	S-Amlodipine Tablets IP 5mg	Each uncoated tablet contains:				
		S-Amlodipine Besylate Eq. to S-Amlodipine	IP	5	mg	
		Excipients		q.s		APPR
391	Paracetamol, Chlorpheniramine Maleate, Phenylepherine, Sodium Citrate & Menthol Syrup	Each 5ml contains :				
		Paracetamol	IP	125	mg	
		Chlorpheniramine Maleate	IP	0.5	mg	
		Phenylepherine HCl	IP	5	mg	
		Sodium Citrate	IP	60	mg	
		Menthol	IP	1	mg	
		In a flavoured syrupy base		q.s		
		Approved colour used				APPR
392	Levofloxacin Tablets IP 750mg	Each film coated tablet contains				
		Levofloxacin Hemihydrate	IP			
		Eq. to Levofloxacin		750	mg	APPR
		Excipients		q.s		
		Approved colour used				
393	Carbonyl Iron, Folic Acid, Vitamin B12, Vitamin C & Zinc Capsules	FOR THERAPEUTIC USE				
		Each hard gelatin Capsule Contains: (In pellets form)				
		Carbonyl Iron Eq. to. Elemental Iron		100	mg	
		Zinc Sulphate Monohydrate	IP	61.8	mg	
		Vitamin B12	IP	15	mcg	
		Folic Acid	IP	1.5	mg	
		Vitamin C	IP	75	mg	
		Excipients		q.s		

		Approved Colours used in empty Capsule shells.				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANT ITY**	**UNIT**	**APP**
394	Carbonyl Iron, Folic Acid, Zinc & Vitamin C Capsules	Each hard gelatin Capsule Contains:				
		(In the form of pellets)				
		Carbonyl Iron Eq. to Elemental Iron		100	mg	
		Folic Acid	IP	1.5	mg	
		Zinc Sulphate Monohydrate (Eq. to Elemental Zinc 22.5mg)	IP	61.8	mg	
		Vitamin B12	IP	15	mcg	
		Excipients		q.s		
		Approved Colours used in empty Capsule shells.				APPR
395	Levofloxacin & Bromhexine HCl Solution (for Veterinary use Only)	Each ml Contains				
		Levofloxacin Hemihydrate eq.to Levofloxacin	IP	100	mg	
		Bromhexine HCl	IP	7.5	mg	
		In a suiatable vehicle base		q.s		APPR
396	Rosuvastatin	Each film coated tablet contains:				
	Tablets IP 5mg	Rosuvastatin Calcium Eq. to Rosuvastatin	IP	5	mg	
		Excipients		q.s		APPR
		Approved colour used				
397	Benfotiamine, Methylcobalamin, Alpha Lipoic Acid, Vitamin B6, Inositol & Folic Acid Capsules	Each hard gelatin capsule contains:				
		Benfotiamine		100	mg	
		Methylcobalamin	USP	1000	mcg	
		Alpha Lipoic Acid	USP	200	mg	
		Vitamin B6	IP	3	mg	
		Inositol	USP	100	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colour used in empty capsule shell.				APPR
PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945						
S. No.	**GENERIC NAME & DOSAGE FORM**	**COMPOSITION**	**SPECIFI CATION**	**QUANT ITY**	**UNIT**	**APP**
398	Telmisartan & Hydrochlorothiazi de Tablets	Each uncoated tablet contains:				
		Telmisartan	IP	80	mg	
		Hydrochlorothiazide	IP	12.5	mg	
		Excipients		q.s		APPR
399	Luliconazole Cream	Composition :				
		Luliconazole		1%	w/w	
		Preservative: Benzyl Alcohol	IP	1%	w/w	
		Cream base		q.s		APPR
400	Ebastine Tablets IP 10mg	Each film coated tablet contains:				
		Ebastine	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
401	Ebastine Tablets IP 20mg	Each film coated tablet contains:				
		Ebastine	IP	20	mg	
		Excipients		q.s		
		Approved colour used				APPR
402	Nadifloxacin,	Composition :				
		Nadifloxacin	IP	1%	w/w	

	Miconazole Nitrate & Mometasone Furoate Cream	Miconazole Nitrate	IP	2%	w/w	
		Mometasone Furoate	IP	0.1%	w/w	
		Cream base		q.s		APPR
403	Olmesartan Medoxomil & Hydrochlorothiazi de Tablets	Each uncoated tablet contains:				
		Olmesartan Medoxomil	IP	40	mg	
		Hydrochlorothiazide	IP	12.5	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
404	Serratiopeptidase Tablets IP 5mg	Each enteric film coated tablet contains:				
		Serratiopeptidase (10,000 Serratiopeptidase units)	IP	5	mg	
		Excipients		q.s		
		Approved colour used				APPR
405	Serratiopeptidase Tablets IP 10mg	Each enteric film coated tablet contains:				
		Serratiopeptidase (20,000 Serratiopeptidase units)	IP	10	mg	
		Excipients		q.s		
		Approved colour used				APPR
406	Lactic Acid Bacillus mouth dissolving Tablets	Each uncoated mouth dissolving tablet contains:				
		Lactic Acid Bacillus		60	million spores	
		Excipients		q.s		APPR
407	Lactic Acid Bacillus Oral Suspension	Each 5ml of the reconstituted suspension contains:				
		Lactic Acid Bacillus		60	million spores	
		Excipients		q.s		
		In a flavoured base				
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
1	Ofloxacin Tablets IP 400mg	Each film coated tablet contains:				
		Ofloxacin	IP	400	mg	
		Excipients		q.s		
		Approved colour used				APPR
2	Flunarizine Dihydrochloride Tablets	Each uncoated tablet contains:				
		Flunarizine Dihydrochloride Eq. to Flunarizine	BP	5	mg	
		Excipients		q.s		APPR
3	Flunarizine Dihydrochloride Tablets	Each uncoated tablet contains:				
		Flunarizine Dihydrochloride Eq. to Flunarizine	BP	10	mg	
		Excipients		q.s		APPR
4	Etodolac Tablets IP 400mg	Each film coated tablet contains:				
		Etodolac	IP	400	mg	
		Excipients		q.s		
		Approved colour used				APPR
5	Isotretinoin Capsules	Each hard gelatin capsule contains:				
		Isotretinoin	IP	20	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				APPR
6	Atorvastatin &	Each film coated tablet contains:				

	Fenofibrate Tablets	Atorvastatin Calcium Eq. to Atorvastatin	IP	10	mg	APPR
		Fenofibrate	IP	160	mg	
		Excipients		q.s		
		Approved colour used				
7	Ascorbic Acid Tablets IP 100mg	Each uncoated tablet contains:				
		Ascorbic Acid	IP	100	mg	
		Excipients		q.s		APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
8	Pregabalin Capsules IP 75mg	Each hard gelatin capsule contains:				
		Pregabalin	IP	75	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				APPR
9	Pregabalin Capsules IP 150mg	Each hard gelatin capsule contains:				
		Pregabalin	IP	150	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				APPR
10	Divalproex Sodium Extended Release Tablets IP 250mg	Each film coated extended release tablet contains:				
		Divalproex Sodium Eq. to Valproic Acid	IP	250	mg	
		Excipients		q.s		
		Approved colour used				APPR
11	Mometasone Furoate Lotion	Composition:				
		Mometasone Furoate	IP	0.1%	w/v	
		Lotion Base		q.s		APPR
12	Paracetamol, Phenylephrine HCl, Diphenhydramin e HCl & Caffeine Tablets	Each film coated tablet contains:				
		Paracetamol	IP	500	mg	
		Phenylephrine HCl	IP	5	mg	
		Diphenhydramine HCl	IP	25	mg	
		Caffeine (Anhydrous)	IP	30	mg	
		Excipients		q.s		
		Approved colour used				APPR
13		Each film coated tablet contains:				
		Drotaverine HCl	IP	80	mg	APPR
	Drotaverine HCl & Aceclofenac Tablets	Aceclofenac	IP	100	mg	
		Excipients		q.s		
		Approved colour used				
14	Thiocolchicoside Tablets	Each film coated tablet contains:				
		Thiocolchicoside	IP	4	mg	
		Excipients		q.s		
		Approved colour used				APPR

PACK SIZE AS PER SCHEDULE-P-1 OF DRUGS & COSMETICS ACT 1940 AND RULES 1945

S. No.	GENERIC NAME & DOSAGE FORM	COMPOSITION	SPECIFICATION	QUANTITY	UNIT	APP
15	Clarithromycin Dispersible Tablets	Each uncoated dispersible tablet contains:				APPR
		Clarithromycin	IP	125	mg	
		Excipients		q.s		
16	Calamine & Light Liquid Paraffin Lotion	Composition:				
		Calamine	IP	8.0%	w/v	
		Light Liquid Paraffin	IP	10.0%	w/v	
		In a lotion base		q.s		APPR
17	Ketoconazole Lotion	Composition:				
		Ketoconazole	IP	2.0%	w/v	
		In a lotion base		q.s		APPR

18	Ketoconazole Solution	Composition:				
		Ketoconazole	IP	2.0%	w/v	
		Approved colour used				APPR
19	Povidone Iodine Solution IP	Composition:				
		Povidone Iodine (0.1% w/v available Iodine)	IP	1.0%	w/v	
		Aqueous Base		q.s		APPR
20	Povidone Iodine Solution IP	Composition:				APPR
		Povidone Iodine (0.2% w/v available Iodine)	IP	2.0%	w/v	
		Aqueous Base		q.s		

S. No.	GENERIC NAME /PHARMACOPAEIAL	COMPOSITION				App
1	Gliclazide Tablets IP 80 mg	Each uncoated tablet contains:				
		Gliclazide	IP	80	mg	
		Excipients		q.s		App
2	Artemether & Lumefantrine Tablet	Each uncoated tablet contains:				
		Artemether	IP	40	mg	
		Lumefantrine		240	mg	
		Excipients		q.s		App
3	Artemether & Lumefantrine Dispersible Tablet	Each uncoated dispersible tablet contains:				
		Artemether	IP	40	mg	
		Lumefantrine		240	mg	
		Excipients		q.s		App
4	Methylprednisolone Tablets IP	Each uncoated tablet contains:				
		Methylprednisolone	IP	8	mg	
		Excipients		q.s		App
5	Pantoprazole Sodium(EC) & Levosulpride(SR) Capsule	Each hard gelatin capsule contains				
		Pantoprazole Sodium Eq. to Pantoprazole (As enteric Coated pellets)	IP	40	mg	
		Levosulpiride (As sustained release pellets)		75	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				App
6	Doxyfylline Tablets IP 400 mg	Each uncoated tablet contains:				App
		Doxyfylline	IP	400	mg	
		Excipients		q.s		
7	Streptococcus Faecalis, Clostridium butyricum, Bacillus mesentericus & Lactic Acid bacillus (lactobacillus Sporogenes) C apsule	Each hard gelatin capsule contains				
		Streptococcus Faecalis		30	million	
		Clostridium Butyricum		2	million	
		Bacillus Mesentericus		1	million	
		Lactic Acid Bacillus (lactobacillus Sporogenes)		50	million	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				App

S. No.	GENERIC NAME /PHARMACOPAEIAL	COMPOSITION				REF
1	Olmesartan Medoxomil Tablet IP 20 mg	Each film coated tablet contains:				
		Olmesartan Medoxomil	IP	20	mg	
		Excipients		q.s		
		Approved colour used.				App
2	Olmesartan Medoxomil Tablet IP 40 mg	Each film coated tablet contains:				
		Olmesartan Medoxomil	IP	40	mg	
		Excipients		q.s		
		Approved colour used.				App
3	Diacerein & Aceclofenac	Each film coated tablet contains:				

	Tablet	Aceclofenac	IP	100	mg	
		Diacerein	IP	50	mg	
		Excipients		q.s		
		Approved colour used.				Apr
4	Sitagliptin Phosphate & Metformin Hydrochloride Tablets	Each film coated tablet contains:				
		Sitagliptin Phosphate Monohydrate Eq. to Sitagliptin	IP	50	mg	
		Metformin Hydrochloride	IP	500	mg	
		Excipients		q.s		
		Approved colour used.				Apr
5	Diacerein Capsule IP 50 mg	Each hard gelatin capsule contains				
		Diacerein	IP	50	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				Apr
6	Cetrimide Cream 0.5 % w/w	Composition:				Apr
		Cetrimide	IP	0.5 %	w/w	
		Cream Base		q.s		
7	Doxycycline Hydrochloride Capsule IP 100 mg	Each hard gelatin capsule contains				
		Doxycycline Hydrochloride Eq. to Doxycycline	IP	100	mg	
		Excipients		q.s		Apr
		Approved colours used in empty capsule shells.				
8	Paroxetine Sustained Release Tablets IP 25 mg	Each film coated sustained release tablet contains:				
		Paroxetine Hydrochloride Hemihydrate Eq. to Paroxetine	IP	25	mg	
		Excipients		q.s		
		Approved colour used.				
9	Paroxetine Sustained Release Tablets IP 12.5 mg	Each film coated sustained release tablet contains:				
		Paroxetine Hydrochloride Hemihydrate Eq. to Paroxetine	IP	12.5	mg	
		Excipients		q.s		
		Approved colour used.				
10	Paroxetine Sustained Release Tablets IP 37.5 mg	Each film coated sustained release tablet contains:				
		Paroxetine Hydrochloride Hemihydrate Eq. to Paroxetine	IP	37.5	mg	
		Excipients		q.s		
		Approved colour used.				
1	Tranexamic Acid Tablet IP 250 mg	Each film coated tablet contains:				
		Tranexamic Acid	IP	250	mg	
		Excipients		q.s		
		Approved colour used.				Apr
2	Tranexamic Acid Tablet IP 500 mg	Each film coated tablet contains:				
		Tranexamic Acid	IP	500	mg	
		Excipients		q.s		
		Approved colour used.				Apr
3	Atorvastatin Tablets IP	Each film coated tablet contains:				
		Atorvastatin Calcium Eq. to Atorvastatin	IP	5	mg	
		Excipients		q.s		
		Approved colour used.				Apr
4	Pregabalin, Methylcobalamin,	Each hard gelatin capsule contains				
		Pregabalin	IP	75	Mg	
	Alpha Lipoic Acid ,Pyridoxine HCL, Folic	Methylcobalamin	USP	750	Mcg	
		Alpha Lipoic Acid	USP	100	Mg	

	,Pyridoxine HCL, Folic Acid Capsule	Pyridoxine Hydrochloride	IP	3	Mg	
		Folic Acid	IP	1.5	Mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				**Appr**
5	Pregabalin, Methylcobalamin, Alpha Lipoic Acid ,Pyridoxine HCL, Folic Acid Capsule	Each hard gelatin capsule contains				
		Pregabalin	IP	150	mg	
		Methylcobalamin	USP	750	mcg	
		Alpha Lipoic Acid	USP	100	mg	
		Pyridoxine Hydrochloride	IP	3	mg	
		Folic Acid	IP	1.5	mg	
		Excipients		q.s		
		Approved colours used in empty capsule shells.				**Appr**
6	Sodium Bicarbonate Tablet USP 500 mg	Each film coated tablet contains:				
		Sodium Bicarbonate	IP	500	mg	
		Excipients		q.s		
		Approved colour used.				

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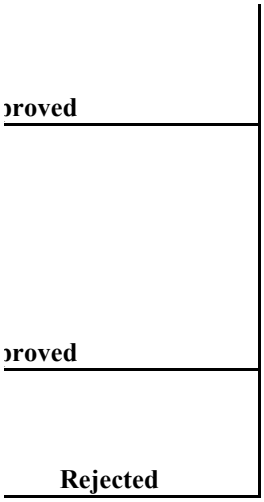
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