GRV PHARMA™
A Unit of : GRV International Business Consultancy LLP

Product List

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S.No	Generic Name	Composition	Ph.Ref	Claim
1	**Dicyclomine Hydrochloride & Paracetamol Tablets**	**Each uncoated tablet Contains :**		
		Dicyclomine Hydrochloride	IP	20 mg.
		Paracetamol	IP	325 mg.
		Excipients		q.s.
2	**Cetirizine Tablets IP**	**Each uncoated tablet contains :**		
		Cetirizine Hydrochloride	IP	10 mg.
		Excipients		q.s.
3	**Diphenhydramine HCl. , Ammonium Chloride & Sodium Citrate Syrup**	**Each 5 ml. contains :**		
		Diphenhydramine Hydrochloride	IP	14.08mg.
		Ammonium Chloride	IP	138 mg.
		Sodium Citrate	IP	57.03mg.
		Menthol	IP	q.s.
		Flavoured Base		q.s.
		Approved Colour used		
4	**Gliclazide Tablets IP**	**Each uncoated tablet contains:**		
		Gliclazide	IP	80 mg.
		Excipients		q.s.
5	**Gliclazide & Metformin Hydrochloride Tablets**	**Each uncoated tablet contains:**		
		Gliclazide	IP	80 mg.
		Metformin Hydrochloride	IP	500 mg.
		Excipients		q.s.
6	**Bromhexine Hydrochloride, Terbutaline Sulphate , Guaiphenesine Syrup**	**Each 10 ml. contains:**		
		Bromhexine Hydrochloride	IP	8 mg.
		Terbutaline Sulphate	IP	2.50 mg.
		Guaiphenesin	IP	100mg.
		Menthol	IP	q.s.
		Flavoured Base		q.s.
		Approved Colour used		
7	**Albendazole Tablets IP 400 mg**	**Each uncoated chewable tablet contains :**		
		Albendazole	IP	400 mg.
		Excipients		q.s.
		Approved colour used		
8	**Albendazole Oral Suspension IP**	**Each 5 ml. contains:**		
		Albendazole	IP	200 mg.
		Flavoured Palatable base		q.s.
		Approved Colour used		
9	**Nimesulide Oral Dispersible Tablets 100 mg**	**Each uncoated flavoured dispersible tablet contains:**		
		Nimesulide	BP	100 mg.
		Approved Colour used		
		''Not for use below 12 years of age''		

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10	**Nimesulide Oral Dispersible Tablets 200 mg**	**Each uncoated flavoured dispersible tablet contains:**		
		Nimesulide	BP	200 mg.
		Approved Colour used		
		''Not for use below 12 years of age''		
11	**Nimesulide Mouth Dissolving Tablets 100 mg**	**Each uncoated mouth dissolving tablet contains:**		
		Nimesulide	BP	100 mg.
		Excipients		q.s.
		Approved Colour used		
		''Not for use below 12 years of age''		
12	**Nimesulide Mouth Dissolving Tablets 200 mg**	**Each uncoated mouth dissolving tablet contains:**		
		Nimesulide	BP	200 mg.
		Excipients		q.s.
		Approved Colour used		
		''Not for use below 12 years of age''		
13	**Nimesulide Tablets 100 mg**	**Each uncoated tablet contains:**		
		Nimesulide	BP	100 mg.
		Excipients		q.s.
		Approved Colour used		
		''Not for use below 12 years of age''		
14	**Nimesulide & Paracetamol Tablets**	**Each uncoated tablet contains :**		
		Nimesulide	BP	100 mg.
		Paracetamol	IP	325 mg
		Excipients	q.s.	
		''Not for use below 12 years of age''		
15	**Pantoprazole Gastro-resistant Tablets IP**	**Each enteric coated tablet contains:**		
		Pantoprazole Sodium	IP	
		Equivalent to Pantoprazole		40 mg.
		Excipients		q.s.
		Approved colour used		
16	**Pantoprazole Sodium & Domperidone Tablets**	**Each enteric coated tablet contains:**		
		Pantoprazole Sodium	IP	
		Equiv. to Pantoprazole		40 mg.
		Domperidone	IP	10 mg.
		Excipients		
		Approved colour used		
17	**Pantoprazole Sodium & Domperidone Tablets**	**Each enteric coated tablet contains:**		
		Pantoprazole Sodium	IP	
		Equiv. to Pantoprazole		20 mg.
		Domperidone	IP	10 mg.
		Excipients		q.s.
		Approved colour used		

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18	**Pantoprazole Sodium Enteric coated & Domperidone Sustained Release Capsules**	**Each hard gelatin capsule contains:** Pantoprazole Sodium Eq. to Pantoprazole (as enteric coated pellets) Domperidone (As sustained release pellets) Colours: Approved colours used in empty shells & Pellets	 IP IP 	 40 mg. 30 mg.
19	**Alprazolam Tablets IP 0.25 mg**	**Each uncoated tablet contains:** Alprazolam Excipients Approved colour used	 IP 	 0.25 mg. q.s.
20	**Alprazolam Tablets IP 0.50 mg**	**Each uncoated tablet contains:** Alprazolam Excipients Approved colour used	 IP 	 0.50 mg. q.s.
21	**Alprazolam & Propranolol Tablets**	**Each uncoated tablet contains:** Alprazolam Propranolol Hydrochloride Excipients Approved colour used	 IP IP 	 0.25 mg. 20 mg. q.s.
22	**Mefenamic Acid & Dicyclomine Hydrochloride Tablets IP**	**Each Uncoated Tablet contains :** Mefenamic Acid Dicyclomine Hydrochloride Excipients Approved colour used	 IP IP 	 250 mg. 20 mg. q.s.
23	**Omeprazole Capsules IP**	**Each hard gelatin capsule contains :** Omeprazole (As enteric coated pellets) Approved colour used in empty shell	 IP 	 20 mg.
24	**Omeprazole & Domperidone Capsules**	**Each hard gelatin capsule contains :** Omeprazole (As enteric coated pellets) Domperidone Approved colour used in empty shell	 IP IP 	 20 mg. 10 mg.
25	**Lactulose Solution USP**	**Each 15 ml contains:** Lactulose Concentrate Equivalent to Lactulose In palatable base	 USP 	 10 gm. q.s.

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26	**Cyproheptadine HCl. & Tricholine Citrate Syrup**	Each 5 ml. contains: Cyproheptadine Hydrochloride Tricholine Citrate Sorbitol(70% Solution) Flavoured Base Approved colour used	 IP IP 	 2 mg. 275 mg. q.s. q.s.
27	**Cyproheptadine HCl. Syrup**	**Each 5 ml. contains:** Cyproheptadine Hydrochloride Flavoured Base Approved colour used	 IP 	 2 mg. q.s.
28	**Cyproheptadine HCl & Tricholine Citrate Drops**	**Each ml. Contains :** Cyproheptadine Hydrochloride Tricholine Citrate (65%) Flavoured base Approved Colour used	 IP 	 1.50 mg. 55 mg. q.s.
29	**Magaldrate & Simethicone Oral Suspension**	**Each 5ml. contains :** Magaldrate Simethicone Flavoured base Flavoured base	 IP IP 	 480 mg. 20 mg. q.s.
30	**Magaldrate, Simethicone & Oxetacaine Suspension**	**Each 5ml. contains :** Magaldrate Simethicone Oxetacaine Flavoured base Colour : Approved colour used	 IP IP BP 	 540 mg. 50 mg. 10 mg. q.s.
31	**Fluconazole Capsule IP 150 mg**	**Each hard gelatin capsule Contains:** Fluconazole Approved colour used in empty shell of hard gelatin capsules	 IP 	 150 mg.
32	**Fluconazole Tablets IP 150 mg**	**Each uncoated tablet contains :** Fluconazole Excipients	 IP 	 150 mg. q.s.
33	**Ciprofloxacin HCl. & Tinidazole Tablets**	**Each film coated tablet contains:** Ciprofloxacin Hydrochloride Eq. to Ciprofloxacin Tinidazole Excipients Approved colour used	 IP IP 	 500 mg. 600 mg. q.s.

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34	**Ofloxacin & Ornidazole Suspension**	**Each 5ml. Contains:**		
		Ofloxacin	IP	50mg.
		Ornidazole	IP	125mg.
		Flavoured base	q.s.	
		Approved colour used		
35	**Ofloxacin & Ornidazole Tablets**	**Each film coated tablet contains:**		
		Ofloxacin	IP	200 mg.
		Ornidazole	IP	500 mg.
		Excipients		q.s.
		Approved colour used		
36	**Nitazoxanide & Ofloxacin Tablets**	**Each film coated tablet contains:**		
		Nitazoxanide		500 mg.
		Ofloxacin	IP	200 mg.
		Excipients		q.s.
		Approved colour used		
37	**Nitazoxanide & Ofloxacin Suspension**	**Each 5ml. contains:**		
		Nitazoxanide		100 mg.
		Ofloxacin	IP	50 mg.
		Flavoured base		q.s.
		Approved Colour used		
38	**Diclofenac Potassium , Paracetamol & Magnesium Trisilicate Tablets**	**Each uncoated tablet contains:**		
		Diclofenac Potassium	BP	50 mg.
		Paracetamol	IP	325 mg.
		Magnesium Trisilicate	IP	100 mg.
		Approved Colour used		
39	**Diclofenac Potassium , Paracetamol & Chlorzoxazone Tablets**	**Each uncoated tablet contains**		
		Diclofenac Potassium	BP	50 mg.
		Paracetamol	IP	325 mg.
		Chlorzoxazone	USP	500 mg.
		Approved colour used		
40	**Diclofenac Potassium & Paracetamol Tablets**	**Each uncoated tablet contains :**		
		Diclofenac Potassium	BP	50 mg.
		Paracetamol	IP	325 mg.
		Excipients		q.s.
41	**Diclofenac Potassium , Paracetamol & Chlorzoxazone Tablets**	**Each uncoated tablet contains**		
		Diclofenac Potassium	BP	50 mg.
		Paracetamol	IP	325 mg.
		Chlorzoxazone	USP	250 mg.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
42	**Paracetamol , Phenylephrine HCl., & Chlorpheniramine Maleate Tablets**	**Each Uncoated Tablet Contains :** Paracetamol Phenylephrine Hydrochloride Chlorpheniramine Maleate Excipients Approved colour used	 IP IP IP	 325mg. 10 mg. 2 mg. q.s.
43	**Paracetamol , Phenylephrine HCl., Chlorpheniramine Maleate & Sodium Citrate Suspension**	**Each 5 ml. Contains :** Paracetamol Phenylephrine Hydrochloride Sodium Citrate Chlorpheniramine Maleate Flavoured base Approved colour used	 IP IP IP IP	 250mg. 2.50mg. 60 mg. 1 mg. q.s.
44	**Paracetamol, Chlorpheniramine Maleate & Phenylephrine Hydrochloride Oral Solution**	**Each 5 ml. Contains :** Paracetamol Phenylephrine Hydrochloride Chlorpheniramine Maleate Flavoured base Approved colour used	 IP IP IP	 125mg. 5 mg. 1 mg. q.s.
45	**Paracetamol, Caffeine Anhydrous, Phenylephrine HCl. & Chlorpheniramine Maleate Tablets**	**Each uncoated Tablet Contains :** Paracetamol Caffeine Anhydrous Phenylephrine Hydrochloride Chlorpheniramine Maleate Excipients Approved colour used	 IP IP IP IP	 325mg. 25 mg. 5 mg. 2 mg. q.s.
46	**Paracetamol Tablets IP 500 mg**	**Each Uncoated Tablet Contains :** Paracetamol Excipients	 IP	 500mg. q.s.
47	**Paracetamol Tablets IP 650 mg**	**Each uncoated tablet contains :** Paracetamol Excipients	 IP	 650 mg. q.s.
48	**Paracetamol Paediatric Oral Suspension IP 125 mg**	**Each 5 ml. contains :** Paracetamol Flavoured Base Approved colour used	 IP	 125 mg. q.s.
49	**Paracetamol Paediatric Oral Suspension IP 250 mg**	**Each 5 ml. contains :** Paracetamol Flavoured Base Approved colour used	 IP	 250 mg. q.s.

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50	**Paracetamol Syrup IP**	**Each 5 ml. Contains :** Paracetamol Flavoured Syrupy Base Approved Colour used	IP	125 mg q.s.
51	**THEOPAR-120 Paracetamol Oral Suspension BP**	**Each 5 ml. Contains :** Paracetamol Flavoured Base Approved colour used	BP	120 mg. q.s.
52	**Paracetamol Oral Drops**	**Each ml contains:** Paracetamol In Flavoured Base Approved colour used	IP	100mg. q.s.
53	**Codeine Phosphate & Chlorpheniramine Maleate Syrup**	**Each 5ml. contains:** Codeine Phosphate Chlorpheniramine Maleate Flavoured Base Approved colour used	IP IP	10 mg. 4 mg. q.s.
54	**Codeine Phosphate & Chlorpheniramine Maleate Syrup**	**Each 5ml. contains:** Codeine Phosphate Chlorpheniramine Maleate Menthol Flavoured Sugar free base Approved colour used	IP IP IP	10 mg. 4 mg. q.s. q.s.
55	**Mefenamic Acid & Dicyclomine Hydrochloride Tablets IP**	**Each uncoated tablet contains :** Mefenamic Acid Dicyclomine Hydrochloride Excipients Approved colour used	IP IP	250mg. 10 mg. q.s.
56	**Levocetirizine Tablets IP 5 mg**	**Each film-coated tablet contains:** Levocetirizine Dihydrochloride Excipients Approved colour used	IP	5mg. q.s.
57	**Levocetirizine Syrup 2.5 mg**	**Each 5 ml. contains :** Levocetirizine Dihydrochloride Flavoured Base Approved colour used	IP	2.50mg. q.s.

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58	**Levocetirizine Dihydrochloride & Ambroxol Hydrochloride Syrup**	**Each 5 ml. contains :** Levocetirizine Dihydrochloride Ambroxol Hydrochloride Flavoured Base Approved colour used	 IP IP 	 2.50 mg. 30 mg. q.s.
59	**Levocetirizine Dihydrochloride & Ambroxol Hydrochloride Syrup**	**Each 5ml. contains:** Levocetirizine Dihydrochloride Ambroxol Hydrochloride In a Palatable sugar & Sodium free base Approved colour used	 IP IP 	 2.5 mg. 30 mg. q.s.
60	**Levocetirizine Dihydrochloride & Ambroxol Hydrochloride Tablets**	**Each filmcoated tablet contains :** Levocetirizine Dihydrochloride Ambroxol Hydrochloride Excipients Approved colour used	 IP IP 	 5 mg. 60 mg. q.s.
61	**Levocetirizine Dihydrochloride, Phenylephrine HCl, Ambroxol Hydrochloride & Paracetamol Tablets**	**Each film coated tablet contains:** Levocetirizine Dihydrochloride Phenylephrine Hydrochloride Ambroxol Hydrochloride Paracetamol Excipients Approved Colour used	 IP IP IP IP 	 5 mg. 5 mg. 60 mg. 325mg. q.s.
62	**Levocetirizine Dihydrochloride, Phenylephrine HCl, Ambroxol Hydrochloride & Paracetamol Tablets**	**Each film coated tablet contains:** Levocetirizine Dihydrochloride Phenylephrine Hydrochloride Ambroxol Hydrochloride Paracetamol Excipients Approved colour used	 IP IP IP IP 	 2.5 mg. 5 mg. 60 mg. 325mg. q.s.
63	**Levocetirizine Dihydrochloride Dispersible Tablets 5 mg**	**Each flavoured dispersible tablet contains:** Levocetirizine Dihydrochloride Excipients Approved colour used	 IP 	 5 mg. q.s.
64	**Montelukast Sodium & Levocetirizine Dihydrochloride Tablets**	**Each uncoated bilayered tablet contains :** Montelukast Sodium Equiv. to Montelukast Levocetirizine Dihydrochloride Excipients Approved colour used	 IP IP 	 10 mg. 5 mg. q.s.

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65	**Montelukast Sodium & Levocetirizine Dihydrochloride Tablets**	**Each film-coated tablet contains :**		
		Montelukast Sodium	IP	
		Equiv. to Montelukast		10 mg.
		Levocetirizine Dihydrochloride	IP	5 mg.
		Excipients		q.s.
		Approved colour used		
66	**Levocetirizine Dihydrochloride, Phenylephrine HCl. & Ambroxol Hydrochloride Tablets**	**Each uncoated tablet contains:**		
		Levocetirizine Dihydrochloride	IP	5 mg.
		Phenylephrine Hydrochloride	IP	10 mg.
		Ambroxol Hydrochloride	IP	60 mg.
		Excipients		q.s.
		Approved colour used		
67	**Terbutaline Sulphate , Etofyline & Ambroxol Hydrochloride Tablets**	**Each uncoated tablet contains :**		
		Terbutaline Sulphate	IP	2.50 mg.
		Etofylline	BP	100 mg.
		Ambroxol Hydrochloride	IP	30 mg.
		Approved colour used		
68	**Cinnarizine Tablets IP**	**Each uncoated tablet contains :**		
		Cinnarizine	IP	25 mg.
		Excipients		q.s.
		Approved colour used		
69	**Cinnarizine & Domperidone Tablets**	**Each uncoated tablet contains :**		
		Cinnarizine	IP	20 mg.
		Domperidone Maleate	IP	
		Equiv. to Domperidone		15 mg.
		Excipients		q.s.
		Approved colour used		
70	**Ondansetron Orally Disintegrating Tablets IP 4 mg.**	**Each uncoated mouth dissolving tablet contains:**		
		Ondansetron	IP	4 mg
		Excipients		q.s.
		Approved colour used		
71	**Ondansetron Oral Solution IP**	**Each 5ml. contains:**		
		Ondansetron Hydrochloride	IP	
		Equiv. to Ondansetron		2 mg.
		Flavoured Base		q.s.
		Approved colour used		
72	**Dried Aluminium Hydroxide,Magnesium Hydroxide & Simethicone Oral Suspension**	**Each 5 ml. Contains :**		
		Dried Aluminium Hydroxide Gel	IP	200 mg.
		Magnesium Hydroxide	IP	200 mg.
		Simethicone	IP	25 mg.
		Flavoured Base		q.s.
		Colour : Approved colour used		

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73	**Liquid Paraffin & Milk of Magnesia Laxative**	**Each 15 ml. Contains :**		
		Liquid Paraffin	IP	3.75ml.
		Milk of Magnesia	IP	11.25ml.
		Flavoured non syrupy base		q.s.
		Approved colour used		
74	**Alprazolam & Fluoxetine Hydrochloride Tablets**	**Each Uncoated Tablet Contains :**		
		Alprazolam	IP	0.25 mg.
		Fluoxetine Hydrochloride	IP	
		Equiv. to Fluoxetine		20 mg.
		Excipients		q.s.
75	**Clidinium Bromide, Chlordiazepoxide & Dicyclomine Hydrochloride Tablets**	**Each Film Coated Tablet Contains :**		
		Clidinium Bromide	USP	2.50 mg.
		Chlordiazepoxide	IP	5 mg.
		Dicyclomine Hydrochloride.	IP	10 mg.
		Excipients		q.s.
		Approved colour used		
76	**Aceclofenac & Paracetamol Tablets**	**Each film-coated tablet contains :**		
		Aceclofenac	IP	100 mg.
		Paracetamol	IP	325 mg.
		Excipients		q.s.
		Approved colour used		
77	**Aceclofenac , Paracetamol & Chlorzoxazone Tablets**	**Each film-coated tablet contains:**		
		Aceclofenac	IP	100mg.
		Paracetamol	IP	325mg.
		Chlorzoxazone	USP	250mg.
		Excipients		q.s.
		Approved colour used		
78	**Thiocolchicoside & Aceclofenac Tablets**	**Each film-coated tablet contains :**		
		Thiocolchicoside	IP	4 mg.
		Aceclofenac	IP	100 mg
		Excipients		q.s.
		Approved colour used		
79	**Thiocolchicoside & Aceclofenac Tablets**	**Each film-coated tablet contains :**		
		Thiocolchicoside	IP	8 mg.
		Aceclofenac	IP	100 mg
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
80	**Aceclofenac , Paracetamol & Chlorzoxazone Tablets**	**Each film coated tablet contains:** Aceclofenac Paracetamol Chlorzoxazone Excipients Approved colour used	 IP IP USP 	 100mg. 325mg. 500mg. q.s.
81	**Gabapentin Capsules IP 300 mg**	**Each Hard Gelatin Capsule Contains :** Gabapentin Approved Colour used in capsule shell.	 IP 	 300 mg.
82	**Gabapentin Capsules IP 400 mg**	**Each Hard Gelatin Capsule Contains :** Gabapentin Approved Colour used in capsule shell.	 IP 	 400 mg.
83	**Diazepam Tablets IP 5 mg**	**Each uncoated tablet contains** Diazepam Excipients	 IP 	 5 mg. q.s.
84	**Diazepam Tablets IP 10 mg**	**Each uncoated tablet contains** Diazepam Excipients	 IP 	 10 mg. q.s.
85	**Diazepam & Propranolol Hydrochloride Tablets**	**Each uncoated tablet contains:** Diazepam Propranolol Hydrochloride Excipients	 IP IP 	 2.5 mg. 20 mg. q.s.
86	**Disodium Hydrogen Citrate Syrup**	**Each 5 ml. Contains :** Disodium Hydrogen Citrate Flavoured Base Approved colour used	 BP 	 1.25 g. q.s.
87	**Disodium Hydrogen Citrate Syrup**	**Each 5 ml. Contains :** Disodium Hydrogen Citrate Flavoured Base Approved colour used	 BP 	 1.53 g. q.s.
88	**Diphenhydramine HCl. , Ammonium Chloride, Sodium Citrate & Citric Acid Syrup**	**Each 5 ml. Contains :** Diphenhydramine Hydrochloride Ammonium Chloride Sodium Citrate Citric Acid Menthol Flavoured Base Apprved Colour used	 IP IP IP IP IP 	 12.50mg. 130mg. 60mg. 40mg. q.s. q.s.

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89	**Tranexamic Acid & Mefenamic Acid Tablets**	**Each film coated tablet contains :**		
		Tranexamic Acid	IP	500 mg.
		Mefenamic Acid	IP	250 mg.
		Excipients		q.s.
		Approved Colour used		
90	**Etamsylate & Tranexamic Acid Tablets**	**Each film coated tablet contains :**		
		Etamsylate	BP	250mg.
		Tranexamic Acid	IP	250mg.
		Excipients		q.s.
		Colour : Titanium Dioxide IP		
91	**Etamsylate Tablets 250 mg.**	**Each uncoated tablet contains :**		
		Etamsylate	BP	250 mg.
		Excipients		q.s.
		Colour : Ponceau 4R		
92	**Paracetamol, Phenylephrine Hydrochloride, Caffeine Anhydrous & Cetirizine Hydrochloride Tablets**	**Each Uncoated Tablet Contains :**		
		Paracetamol	IP	325 mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Caffeine Anhydrous	IP	25 mg.
		Cetirizine Hydrochloride	IP	5 mg.
		Excipients		q.s.
93	**Levofloxacin & Ornidazole Suspension**	**Each 5 ml. Contains :**		
		Levofloxacin Hemihydrate	IP	
		Equiv. to Levofloxacin		125mg.
		Ornidazole	IP	125 mg.
		Flavoured Base		q.s.
		Approved Colour used		
94	**Levofloxacin & Ornidazole Tablets**	**Each film coated tablet contains :**		
		Levofloxacin Hemihydrate	IP	
		Equiv. to Levofloxacin		250 mg.
		Ornidazole	IP	500 mg.
		Excipients		q.s.
		Approved Colour used		
95	**Levofloxacin & Ambroxol Hydrochloride (Sustained Release) Bilayer Tablets**	**Each uncoated bilayered tablet contains :**		
		Levofloxacin Hemihydrate	IP	
		Equiv. to Levofloxacin		500 mg.
		Ambroxol Hydrochloride	IP	75 mg.
		(As sustained release form)		
		Excipients		q.s.
		Approved colour used		

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96	**Levofloxacin & Ambroxol Hydrochloride (Sustained Release) Bilayer Tablets**	**Each uncoated bilayered tablet contains :** Levofloxacin Hemihydrate Equiv. to Levofloxacin Ambroxol Hydrochloride (As sustained release form) Excipients Approved colour used	 IP IP 	 250 mg. 75 mg. q.s.
97	**Paracetamol & Tramadol Hydrochloride Tablets USP**	**Each uncoated tablet contains :** Paracetamol Tramadol Hydrochloride Excipients	 IP IP 	 325 mg. 37.50 mg. q.s.
98	**Tramodol hydrochloride sustained release Tablets IP 100 mg.**	**Each film coated tablets contains:** Tramadol Hydrochloride (In a controlled release system) Excipients Approved colour used	 IP 	 100 mg
99	**Paracetamol & Tramadol Hydrochloride Tablets USP**	**Each uncoated tablet contains :** Tramadol Hydrochloride Paracetamol Excipients	 IP IP 	 50 mg. 325 mg. q.s.
100	**Rabeprazole Gastro-resistant Tablets IP**	**Each enteric coated tablet contains :** Rabeprazole Sodium Excipients Approved colour used	 IP 	 20 mg. q.s.
101	**Rabeprazole Sodium & Domperidone Tablets**	**Each enteric coated tablet contains:** Rabeprazole Sodium Domperidone Excipients Approved colour used	 IP IP 	 20 mg. 10 mg. q.s.
102	**Rabeprazole Sodium Enteric coated & Domperidone Sustained Release Capsules**	**Each hard gelatin capsule contains :** Rabeprazole sodium (As enteric coated pellets) Domperidone (As sustained release pellets) Approved colour used in empty capsule shell & pellets	 IP IP 	 20 mg. 30 mg.
103	**Rabeprazole Sodium Enteric Coated & Domperidone Sustained Release Capsules**	**Each hard gelatin capsule contains:** Rabeprazole Sodium (as enteric coated pellets) Domperidone (As sustained release pellets) Domperidone (As immediate release pellets) Approved colours are used in empty capsules shell & pellets	 IP IP IP 	 20 mg. 20 mg. 10 mg.

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104	**Ofloxacin & Metronidazole Suspension**	**Each 5 ml. contains :**		
		Ofloxacin	IP	50 mg.
		Metronidazole Benzoate	IP	
		Equiv. to Metronidazole		120 mg.
		In flavoured base		q.s.
		Approved colour used		
105	**Ofloxacin, Metronidazole & Simethicone Suspension**	Each 5 ml. Contains :		
		Ofloxacin	IP	50 mg.
		Metronidazole Benzoate	IP	
		Equiv. to Metronidazole		120 mg.
		Simethicone	IP	10 mg.
		In flavoured base		q.s.
		Approved colour used		
106	**Dried Aluminium Hydroxide,Magnesium Hydroxide, Simethicone & Sodium Carboxymethyl Cellulose Oral Suspension**	**Each 10 ml. contains :**		
		Dried Aluminium Hydroxide Gel	IP	830 mg.
		Magnesium Hydroxide	IP	185 mg.
		Simethicone	IP	50 mg.
		Sodium Carboxymethyl Cellulose	IP	100 mg.
		Sorbitol Solution 70 %	IP	1.25 g.
		(Non Crystallizing)		
		Flavoured non syrupy base		q.s.
		Approved Colour		
107	**Dried Aluminium Hydroxide,Magnesium Hydroxide & Simethicone Oral Suspension**	**Each 5 ml. contains :**		
		Dried Aluminium Hydroxide Gel	IP	200 mg.
		Magnesium Hydroxide	IP	200 mg.
		Simethicone	IP	25 mg.
		Sorbitol Solution 70 %	IP	1.25 g.
		(Non Crystallizing)		
		Flavoured non syrupy base		q.s.
		Approved Colour		
108	**Ofloxacin & Tinidazole Tablets**	**Each film coated tablet contains :**		
		Ofloxacin	IP	200 mg.
		Tinidazole	IP	600 mg.
		Excipients		q.s.
		Approved colour used		
109	**Amitriptyline Hydrochloride & Chlordiazepoxide Tablets**	**Each film coated tablet contains:**		
		Amitriptyline Hydrochloride	IP	
		Eq. to Amitriptyline		12.50mg.
		Chlordiazepoxide	IP	5 mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
110	**Amitriptyline Hydrochloride & Chlordiazepoxide Tablets**	**Each film coated tablet contains:**		
		Amitriptyline Hydrochloride	IP	
		Eq. to Amitriptyline		25 mg.
		Chlordiazepoxide	IP	10 mg.
		Excipients		q.s.
		Approved colour used		
111	**Sildenafil Tablets IP 50 mg**	**Each film-coated tablet contains :**		
		Sildenafil Citrate	IP	
		Equivalent to Sildenafil		50 mg
		Excipients		q.s.
		Approved colour used		
112	**Sildenafil Tablets IP 100 mg**	**Each film-coated tablet contains :**		
		Sildenafil Citrate	IP	
		Equivalent to Sildenafil		100 mg
		Excipients		q.s.
		Approved colour used		
113	**Clotrimazole, Tinidazole,Povidone Iodine Vaginal Tablets**	**Each uncoated vaginal tablet contains:**		
		Clotrimazole	IP	200 mg.
		Tinidazole	IP	600 mg.
		Povidone Iodine	IP	200 mg.
		Excipients		q.s.
114	**Pantoprazole Sodium & Domperidone Capsules**	**Each hard gelatin capsule contains :**		
		Pantoprazole Sodium	IP	
		Equiv. to Pantoprazole		20 mg.
		(As enteric coated pellets)		
		Domperidone	IP	10 mg.
		Excipients		
		Approved colours used in empty shells of hard gelatin capsules		
115	**Pantoprazole Gastro-resistant Tablets IP 20 mg**	**Each enteric coated tablet contains:**		
		Pantoprazole Sodium	IP	
		Equivalent to Pantoprazole		20 mg.
		Excipients		q.s.
		Approved colour used		
116	**Disulfiram Tablets IP 250 mg**	**Each uncoated tablet contains :**		
		Disulfiram	IP	250 mg.
		Excipients		q.s.
117	**Disulfiram Tablets IP 500 mg**	**Each uncoated tablet contains :**		
		Disulfiram	IP	500 mg.
		Excipients		q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
118	**Telmisartan Tablets IP**	**Each uncoated tablet contains :**		
		Telmisartan	IP	40 mg.
		Excipients		q.s.
119	**Telmisartan & Hydrochlorothiazide Tablets IP**	**Each uncoated tablet contains :**		
		Telmisartan	IP	40 mg.
		Hydrochlorothiazide	IP	12.5mg.
		Excipients		q.s.
		Approved colour used		
120	**Telmisartan & Amlodipine Besilate Tablets IP**	**Each uncoated tablet contains :**		
		Telmisartan	IP	40 mg.
		Amlodipine Besilate	IP	
		Equivalent to Amlodipine		5 mg.
		Excipients		q.s.
		Approved colour used		
121	**Amlodipine Besilate & Enalopril Maleate Tablets**	**Each uncoated tablet contains:**		
		Amlodipine Besilate	IP	
		Equiv. to Amlodipine		5 mg.
		Enalapril Maleate	IP	5 mg.
		Excipients		q.s.
122	**Enalapril Maleate Tablets IP 2.5 mg**	Each uncoated tablet contains:		
		Enalapril Maleate	IP	2.5 mg.
		Excipients		q.s.
123	**Enalapril Maleate Tablets IP 5 mg**	Each uncoated tablet contains:		
		Enalapril Maleate	IP	5 mg.
		Excipients		q.s.
124	**Amlodipine Besilate Tablets IP 5 mg**	Each uncoated tablet contains:		
		Amlodipine Besilate	IP	
		Eq. to Amlodipine		5 mg.
		Excipients		q.s.
125	**Amlodipine Besilate Tablets IP 2.5 mg**	Each uncoated tablet contains:		
		Amlodipine Besilate	IP	
		Eq. to Amlodipine		2.5 mg.
		Excipients		q.s.
126	**Amlodipine & Atenolol Tablets**	**Each uncoated tablet contains:**		
		Amlodipine Besilate	IP	
		Equiv. to Amlodipine		5 mg.
		Atenolol	IP	50 mg.
		Excipients		q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
127	**Losartan Potassium & Amlodipine Tablets IP**	**Each film coated tablet contains:**		
		Losartan Potassium	IP	50 mg.
		Amlodipine Besilate	IP	
		Equiv. to Amlodipine		5 mg.
		Excipients		q.s.
		Approved colour used		
128	**Losartan Potassium Tablets IP 25 mg.**	**Each film-coated tablet contains :**		
		Losartan Potassium	IP	25 mg.
		Excipients		q.s.
		Approved colour used		
129	**Losartan Potassium Tablets IP 50 mg.**	**Each film-coated tablet contains :**		
		Losartan Potassium	IP	50 mg.
		Excipients		q.s.
		Approved colour used		
130	**Losartan Potassium & Hydrochlorothiazide Tablets IP**	**Each film-coated tablet contains :**		
		Losartan Potassium	IP	50 mg.
		Hydrochlorothiazide	IP	12.5mg.
		Excipients		q.s.
		Approved colour used		
131	**Ergotamine, Caffeine, Paracetamol & Prochlorperazine Maleate Tablets**	**Each uncoated tablet contains:**		
		Ergotamine Tartrate	IP	1 mg.
		Caffeine(Monohydrate)	IP	100 mg.
		Paracetamol	IP	250 mg.
		Prochlorperazine Maleate	IP	2.50 mg.
		Excipients		q.s.
		Approved colour used		
132	**Isoxsuprine Hydrochloride sustained release tablet 40 mg**	**Each uncoated sustained release tablet contains :**		
		Isoxsuprine Hydrochloride	IP	40 mg.
		Excipients		q.s.
133	**Chlorpheniramine Maleate, Ammonium Chloride & Sodium Citrate Syrup**	**Each 5 ml. contains**		
		Chlorpheniramine Maleate	IP	2.5 mg
		Ammonium Chloride	IP	125 mg
		Sodium Citrate	IP	55 mg
		Flavoured Base		q.s.
		Approved colour used		
134	**Betamethasone Sodium Phosphate Tablets IP 0.5mg**	**Each uncoated tablet contains:**		
		Betamethasone Sodium Phosphate	IP	
		Equiv. to Betamethasone		0.5 mg.
		Excipients		q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
135	**Betamethasone Sodium Phosphate Tablets IP 1mg**	**Each uncoated tablet contains:**		
		Betamethasone Sodium Phosphate	IP	
		Equiv. to Betamethasone		1 mg.
		Excipients		q.s.
136	**Betamethasone Sodium Phosphate Oral Drops**	**Each ml. (Approx. 20 Drops) Contains:**		
		Betamethasone Sodium Phosphate	IP	
		Equiv. to Betamethasone		0.50 mg.
		Flavoured Base		q.s.
		Approved colour used		
137	**Betahistine Tablets IP 8 mg.**	**Each uncoated tablet contains :**		
		Betahistine Dihydrochloride	IP	8 mg.
		Excipients		q.s.
		Approved colour used		
138	**Betahistine Tablets IP 16 mg.**	**Each uncoated tablet contains :**		
		Betahistine Dihydrochloride	IP	16 mg.
		Excipients		q.s.
139	**Trihexyphenidyl Hydrochloride Tablets IP 2 mg**	**Each Uncoated Tablet Contains:**		
		Trihexyphenidyl Hydrochloride	IP	2 mg.
		Excipients		q.s.
140	**Trifluoperazine Hydrochloride & Trihexyphenidyl Hydrochloride Tablets**	**Each Uncoated tablet contains:**		
		Trifluoperazine Hydrochloride	IP	
		Equiv. to Trifluoperazine		5 mg.
		Trihexyphenidyl Hydrochloride	IP	2 mg.
		Excipients	q.s.	
		Approved colour used		
141	**Aceclofenac Tablets IP 100 mg**	Each film coated tablet contains:		
		Aceclofenac	IP	100mg.
		Excipients		q.s.
		Colour:Titanium Dioxide IP		
142	**Aceclofenac Sustained Release Tablets 200 mg**	**Each film coated sustained release tablet contains:**		
		Aceclofenac	IP	200 mg.
		Excipients		q.s.
		Colour : Titanium Dioxide IP		
143	**Aceclofenac & Paracetamol Suspension**	**Each 5 ml. contains :**		
		Aceclofenac	IP	50 mg.
		Paracetamol	IP	125 mg.
		Flavoured Syrupy Base		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
144	**Mefenamic Acid & Paracetamol Suspension**	**Each 5ml. contains:**		
		Mefenamic Acid	IP	50 mg.
		Paracetamol	IP	125 mg.
		Flavoured base		q.s.
		Approved Colour used		
145	**Clonazepam Tablets IP 0.25 mg**	**Each Film Coated Tablet Contains :**		
		Clonazepam	IP	0.25mg.
		Excipients		q.s.
		Approved Colour used		
146	**Clonazepam Tablets IP 0.50 mg**	**Each Film Coated Tablet Contains :**		
		Clonazepam	IP	0.50mg.
		Excipients		q.s.
		Approved Colour used		
147	**Clonazepam Mouth Dissolving Tablets**	**Each uncoated mouth dissolving tablet contains :**		
		Clonazepam	IP	0.25mg.
		Excipients		q.s.
148	**Escitalopram Oxalate & Clonazepam Tablets IP**	**Each film coated tablet contains**		
		Escitalopram Oxalate	IP	
		eq. to Escitalopram		5 mg
		Clonazepam	IP	0.5 mg
		Excipients		q.s.
		Approved colour used		
149	**Escitalopram Oxalate & Clonazepam Tablets IP**	**Each film coated tablet contains**		
		Escitalopram Oxalate	IP	
		eq. to Escitalopram		10 mg
		Clonazepam	IP	0.5 mg
		Excipients		q.s.
		Approved colour used		
150	**Promethazine Hydrochloride & Paracetamol Syrup**	**Each 5ml. contains:**		
		Promethazine Hydrochloride	IP	5 mg.
		Paracetamol	IP	125mg.
		Flavoured base		q.s.
		Approved Colour used		
151	**Promethazine HCl, Pholcodine & Phenylephrine HCl Syrup**	**Each 5 ml. contains :**		
		Promethazine Hydrochloride	IP	1.5 mg.
		Pholcodine	IP	1.5 mg.
		Phenylephrine Hydrochloride	IP	2.5 mg.
		Flavoured base		q.s.
		Approved Colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
152	**Promethazine HCl. & Pholcodine Syrup**	**Each 5 ml. contains :**		
		Promethazine Hydrochloride	IP	1.5 mg.
		Pholcodine	IP	1.5 mg.
		Flavoured base		q.s.
		Approved colour used		
153	**Metoprolol Tartrate Tablets IP**	**Each uncoated tablet contains:**		
		Metoprolol Tartrate	IP	25 mg.
		Excipients		q.s.
154	**Metoprolol Succinate Extended Release Tablets IP**	**Each film-coated extended release tablet contains:**		
		Metoprolol Succinate	IP	23.75mg.
		Equivalent to Metoprolol Tartrate		25 mg.
		Excipients		q.s.
		Approved colour used		
155	**Metoprolol Succinate Extended Release Tablets IP**	**Each film-coated extended release tablet contains:**		
		Metoprolol Succinate	IP	47.5 mg.
		Equivalent to Metoprolol Tartrate		50 mg.
		Excipients		q.s.
		Approved colour used		
156	**Potassium Citrate Monohydrate & Magnesium Citarte Syrup**	**Each 5 ml. contains:**		
		Potassium Citrate Monohydrate	IP	1100mg.
		Magnesium Citrate	USP	375mg.
		Flavoured Sorbitol Base		q.s.
		Approved colour used		
157	**Potassium Citrate & Citric Acid Oral Solution USP**	**Each 5 ml. Contains :**		
		Potassium Citrate	IP	1100mg.
		Citric Acid	IP	334mg.
		Flavoured Sugar free Base		q.s.
		Approved colour used		
158	**Diclofenac Sodium Sustained Release Tablets**	**Each Sustained release film coated tablet contains:**		
		Diclofenac Sodium	IP	100mg.
		Approved colour used		
159	**Diclofenac Sodium, Paracetamol & Magnesium Trisilicate Tablets**	**Each uncoated tablet contains:**		
		Diclofenac Sodium	IP	50 mg.
		Paracetamol	IP	325mg.
		Magnesium Trisilicate	IP	100mg.
		Excipients		q.s.
		Approved colour used		
160	**Diclofenac Sodium, Paracetamol & Chlorzoxazone Tablets**	**Each film-coated tablet contains:**		
		Diclofenac Sodium	IP	50 mg.
		Paracetamol	IP	325mg.
		Chlorzoxazone	USP	250mg.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
161	**Ibuprofen & Paracetamol Suspension**	**Each 5ml. contains:** Ibuprofen Paracetamol Flavoured base Approved colour used	 IP IP	 100mg. 162.5mg. q.s.
162	**Deflazacort Oral Suspension 6 mg.**	**Each 5 ml contains :** Deflazacort Flavoured Syrupy base Approved colour used		 6 mg q.s.
163	**Deflazacort Tablets 6 mg**	**Each uncoated tablets contains :** Deflazacort Exicipents		 6 mg q.s.
164	**Deflazacort Tablets 30 mg**	**Each uncoated tablets contains :** Deflazacort Exicipents		 30 mg q.s.
165	**Atorvastatin Tablets IP 10 mg**	**Each Film coated tablet contains:** Atorvastatin Calcium Equivalent to Atorvastatin Excipients Approved colour used	 IP	 10 mg q.s.
166	**Atorvastatin Tablets IP 20 mg**	**Each film-coated tablet contains :** Atorvastatin calcium Equiv. to Atorvastatin Excipients Approved colour used	 IP	 20 mg. q.s.
167	**Simvastatin Tablets IP 10 mg**	**Each film coated tablet contains :** Simvastatin Excipients Approved colour used	 IP	 10 mg. q.s.
168	**Simvastatin Tablets IP 20 mg**	**Each film coated tablet contains :** Simvastatin Excipients Approved colour used	 IP	 20 mg. q.s.
169	**Torsemide Tablets IP10 mg**	**Each uncoated tablet contains:** Torsemide Equiv. to Torsemide (Anhydrous)	 IP	 10 mg.
170	**Torsemide Tablets IP 20 mg**	**Each uncoated tablet contains:** Torsemide Equiv. to Torsemide (Anhydrous) Excipients Approved colour used	 IP	 20 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
171	**Torsemide Tablets 40 mg**	**Each uncoated tablet contains:**		
		Torsemide	USP	
		Equiv. to Torsemide (Anhydrous)		40 mg.
		Excipients		q.s.
		Approved colour used		
172	**Sucralfate & Oxetacaine Suspension**	**Each 10 ml. contains :**		
		Sucralfate	USP	1 gm.
		Oxetacaine	BP	20 mg.
		Flavoured sorbitol base		q.s.
		Approved colour used		
173	**Sucralfate Oral Suspension**	**Each 10 ml. contains :**		
		Sucralfate	USP	1 gm.
		Flavoured base		
		Approved colour used		
174	**Sodium Picosulphate Syrup**	**Each 5 ml contains**		
		Sodium Picosulphate	BP	5 mg
		In a palatable Sorbitol Base		q.s.
		Approved colour used		
175	**Alprazolam & Sertraline Hydrochloride Tablets**	**Each uncoated tablets contains:**		
		Alprazolam	IP	0.5 mg
		Sertraline Hydrochloride	IP	
		Equiv. to Sertraline		50 mg
176	**Ambroxol Hydrochloride , Terbutaline Sulphate & Guaiphenesin Syrup**	**Each 5 ml. contains:**		
		Ambroxol Hydrochloride	IP	15 mg.
		Terbutaline Sulphate	IP	1.25mg.
		Guaiphenesin	IP	50 mg.
		Menthol	IP	q.s.
		Flavoured base		q.s.
		Approved colour used		
177	**Alprazolam Tablets IP 1 mg**	**Each uncoated tablet contains:**		
		Alprazolam	IP	1 mg.
		Excipients		q.s.
		Approved colour used		
179	**Drotaverine Hydrochloride & Mefenamic Acid Tabs.**	**Each film coated tablet contains:**		
		Drotaverine Hydrochloride		80 mg.
		Mefenamic Acid	IP	250mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
180	**Paracetamol & Caffeine Anhydrous Tablets**	**Each uncoated tablet contains:** Paracetamol Caffeine Anhydrous Excipients	 IP IP	 325mg. 50 mg.
181	**Dothiepin Hydrochloride Tablets 25 mg**	**Each film coated tablet contains:** Dothiepin Hydrochloride Excipients Approved colour used	 IP	 25 mg. q.s.
182	**Dothiepin Hydrochloride Tablets 75 mg**	**Each film coated tablet contains:** Dothiepin Hydrochloride Excipients Approved colour used	 IP	 75 mg. q.s.
183	**Metformin HCl ER & Glimepiride Tablets**	**Each uncoated bilayered tablet contains:** Metformin Hydrochloride (In Extended release form) Glimepiride Colour : Approved colour used Excipients	 IP IP	 500 mg. 1 mg. q.s.
184	**Metformin HCl ER & Glimepiride Tablets**	**Each uncoated bilayered tablet contains:** Metformin Hydrochloride (In Extended release form) Glimepiride Colour : Approved colour used Excipients	 IP IP	 500 mg. 2 mg. q.s.
185	**Metformin HCl SR & Glimepiride Tablets**	**Each uncoated bilayered tablet contains:** Metformin Hydrochloride (As sustained release) Glimepiride Colour : Red Oxide of Iron Excipients	 IP IP	 500 mg. 1 mg. q.s.
186	**Metformin HCl SR & Glimepiride Tablets**	**Each uncoated bilayered tablet contains:** Metformin Hydrochloride (As sustained release) Glimepiride Colour : Yellow Oxide of Iron Excipients	 IP IP	 500 mg. 2 mg. q.s.
187	**Diclofenac Potassium , Paracetamol & Chlorzoxazone Tablets**	**Each filmcoated tablet contains** Diclofenac Potassium Paracetamol Chlorzoxazone Approved colour used	 BP IP USP	 50 mg. 325 mg. 250 mg.

S.No	Generic Name	Composition	Ph.Ref	Claim
188	**Voglibose Dispersible Tablets IP 0.2 mg**	**Each uncoated dispersible tablet contains :** Voglibose Excipients	 IP 	 0.2 mg q.s.
189	**Voglibose Dispersible Tablets IP 0.3 mg**	**Each uncoated dispersible tablet contains :** Voglibose Excipients	 IP 	 0.3 mg q.s.
190	**Voglibose & Metformin Hydrochloride Tablets**	**Each uncoated tablet contains :** Voglibose Metformin Hydrochloride Excipients	 IP 	 0.2 mg 500 mg q.s.
191	**Metformin HCl SR Tablets IP 500 mg.**	**Each uncoated sustained release tablet contains:** Metformin Hydrochloride Excipients	 IP 	 500 mg. q.s.
192	**Metformin HCl SR Tablets IP 1000 mg**	**Each uncoated sustained release tablet contains:** Metformin Hydrochloride Excipients	 IP 	 1000mg. q.s.
193	**Gliclazide & Metformin HCl SR Tablets**	**Each uncoated Bilayer tablet contains :** Metformin Hydrochloride (As sustained release) Gliclazide Approved colour used Excipients	 IP BP 	 500 mg. 80 mg. q.s.
194	**Cholecalciferol Granules**	**Each sachet of 1 Gm contains :** Cholecalciferol Excipients Overages of Vitamins added	 IP 	 60,000 IU q.s.
195	**Racecadotril Granules**	**Each sachet contains:** Racecadotril Excipients	 BP 	 10 mg. q.s.
196	**Racecadotril Dispersible Tablets**	**Each uncoated dispersible tablet contains :** Racecadotril Excipients Approved colour used	 BP 	 10 mg. q.s.
197	**Oral rehydration Salt IP**	**Each pack(21gm.) contains:** Sodium Chloride Potassium Chloride Sodium Citrate Dextrose(Anhydrous) Excipients The oral rehydration Therapy	 IP IP IP IP 	 2.6 g. 1.5 g. 2.6 g. 13.5 g. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
198	**Oral rehydration Salt IP**	**Each pack(21.8 gm.) contains:**		
		Sodium Chloride	IP	2.6 g.
		Potassium Chloride	IP	1.5 g.
		Sodium Citrate	IP	2.9 g.
		Dextrose(Anhydrous)	IP	13.5 g.
		Excipients		q.s.
		The oral rehydration Therapy		
199	**Fluoxetine Hydrochloride Capsules IP 20 mg**	Each hard gelatin capsule contains:		
		Fluoxetine Hydrochloride	IP	
		Equiv. to Fluoxetine		20 mg.
		Excipients		q.s.
		Approved colour used in empty capsule shell		
200	**Hydroxyzine Hydrochloride Tablets USP**	Each film coated tablet contains:		
		Hydroxyzine Hydrochloride	USP	25 mg.
		Excipients		q.s.
		Approved colour used		
201	**Dicyclomine Hydrochloride & Acetaminophen Capsules**	**Each hard gelatin capsule contains:**		
		Dicyclomine Hydrochloride	IP	20 mg.
		Acetaminophen	IP	325 mg.
		Approved colours used in empty capsule shell		
202	**Aciclovir Tablets IP 800 mg**	**Each Uncoated Tablet Contains:**		
		Aciclovir	IP	800 mg.
		Excipients		q.s.
203	**Risperidone Tablets BP 2 mg**	**Each film coated tablet contains :**		
		Risperidone	BP	2 mg.
		Excipients		q.s.
		Approved colours used		
204	**Domperidone Maleate Suspension**	**Each ml. contains :**		
		Domperidone Maleate	IP	1 mg.
		Flavoured base		q.s.
		Approved colour used		
205	**Dried Aluminium Hydroxide,Magnesium Hydroxide & Activated Dimethicone Oral Suspension**	**Each 10 ml. Contains :**		
		Dried Aluminium Hydroxide Gel	IP	200 mg.
		Magnesium Hydroxide	IP	200 mg.
		Activated Dimethicone	IP	25 mg.
		Flavoured Base		q.s.
		Colour : Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
206	**Bromhexine Hydrochloride, Terbutaline Sulphate , Guaiphenesine Syrup**	**Each 10 ml. contains:**		
		Bromhexine Hydrochloride	IP	4 mg.
		Terbutaline Sulphate	IP	2.50 mg.
		Guaiphenesin	IP	100mg.
		Menthol	IP	q.s.
		Flavoured Base		q.s.
		Approved Colour used		
207	**Paracetamol , Phenylephrine HCl., Chlorpheniramine Maleate & Sodium Citrate Syrup**	**Each 5 ml. Contains :**		
		Paracetamol	IP	125mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Sodium Citrate	IP	60 mg.
		Chlorpheniramine Maleate	IP	1 mg.
		Flavoured base		q.s.
		Approved colour used		
208	**Nimesulide , Paracetamol & Chlorzoxazone Tablets**	**Each uncoated tablet contains :**		
		Nimesulide	BP	100 mg.
		Paracetamol	IP	325 mg.
		Chlorzoxazone	USP	375 mg.
		Excipients		q.s.
		Approved colour used		
209	**Cetirizine HCl., Dextromethorphan HBr, Phenylephrine HCl & Zinc Gluconate Syrup**	**Each 5 ml contains :**		
		Cetirizine Hydrochloride	IP	2.5 mg.
		Dextromethorphan Hydrobromide	IP	5 mg.
		Phenylephrine Hydrochloride	IP	2.5 mg.
		Zinc Gluconate	USP	7.5 mg.
		Menthol	IP	q.s.
		Flavoured base		q.s.
		Approved colour used		
210	**Cetirizine HCl., Dextromethorphan HBr, Phenylephrine HCl, Zinc Gluconate & Paracetamol Syrup**	**Each 5 ml contains:**		
		Cetirizine Hydrochloride	IP	2.5 mg.
		Dextromethorphan Hydrobromide	IP	7.5 mg.
		Phenylephrine Hydrochloride	IP	5.0 mg.
		Zinc Gluconate	USP	
		Eq. to Elemental Zinc		7.5 mg.
		Paracetamol	IP	125mg.
		Menthol	IP	q.s.
		Flavoured base		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
211	**Prazosin Hydrochloride Tablets IP 5mg.**	**Each film coated sustained release tablet contains :**		
		Prazosin Hydrochloride	IP	
		Equivalent to Prazosin		5 mg.
		Excipients		q.s.
		Approved colour used		
212	**Prazosin Hydrochloride Tablets IP 2.5mg.**	**Each film coated sustained release tablet contains :**		
		Prazosin Hydrochloride	IP	
		Equivalent to Prazosin		2.5 mg.
		Excipients		q.s.
		Approved colour used		
213	**Phenylephrine HCl, Paracetamol & Cetirizine Hydrochloride Syrup**	**Each 5 ml. contains:**		
		Phenylephrine Hydrochloride	IP	5 mg.
		Paracetamol	IP	125 mg.
		Cetirizine Hydrochloride	IP	5 mg.
		Flavoured base		q.s.
		Approved colour used		
214	**Carisoprodol Tablets IP 350 mg**	**Each uncoated tablet contains:**		
		Carisoprodol	IP	350 mg.
		Excipients		q.s.
		Approved colour used		
215	**Paracetamol Bromhexine Hydrochloride, Chlorpheniramine Maleate, Phenylephrine HCl & Guaiphenesin Tablets**	**Each film coated tablet Contains:**		
		Paracetamol	IP	325mg
		Bromhexine Hydrochloride	IP	8 mg.
		Chlorpheniramine Maleate	IP	2 mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Guaiphenesin	IP	100mg.
		Excipients		q.s.
		Approved colour used		
216	**Oxetacaine, Aluminium Hydroxide, Magnesium Hydroxide Suspension**	**Each 5 ml. contains:**		
		Oxetacaine	BP	10 mg.
		Aluminium Hydroxide		0.291 g.
		(Added as Dried Aluminium		
		Hydroxide gel)	IP	380 mg.
		Magnesium Hydroxide	IP	98 mg.
		In Palatable Sugar free base		q.s.
		Approved colour used		
217	**Phenobarbitone Tablets IP 30 mg**	**Each uncoated tablet contains:**		
		Phenobarbitone	IP	30 mg.
		Excipients		q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
218	**Phenobarbitone Tablets IP 60 mg**	**Each uncoated tablet contains:**		
		Phenobarbitone	IP	60 mg.
		Excipients		q.s.
219	**Cetirizine Hydrochloride Phenylephrine HCl, Zinc Gluconate & Paracetamol Syrup**	**Each 5 ml. contains :**		
		Cetirizine Hydrochloride	IP	2.5 mg.
		Phenylephrine Hydrochloride	IP	2.5 mg.
		Zinc Gluconate	USP	7.5 mg.
		Paracetamol	IP	125 mg.
		Flavoured mentholated base		q.s.
		Approved colour used		
220	**Cetirizine Hydrochloride Phenylephrine HCl, Zinc Gluconate & Paracetamol Syrup**	**Each 5 ml. contains:**		
		Cetirizine Hydrochloride	IP	2.5 mg.
		Phenylephrine Hydrochloride	IP	5 mg
		Zinc Gluconate	USP	26.14mg.
		Eq. to Elemental Zinc		3.75 mg.
		Paracetamol	IP	250 mg.
		Flavoured mentholated base		q.s.
		Approved colour used		
221	**Atenolol Tablets IP 50 mg**	**Each uncoated tablet contains:**		
		Atenolol	IP	50 mg.
		Excipients		q.s.
222	**Levocetirizine Dihydrochloride Phenylephrine HCl, Paracetamol & Caffeine Tablets**	**Each uncoated tablet contains :**		
		Levocetirizine Dihydrochloride	IP	2.5 mg.
		Phenylephrine hydrochloride	IP	10 mg.
		Paracetamol	IP	325 mg.
		Caffeine (Anhydrous)	IP	30 mg.
		Excipients		q.s.
		Approved colour used		
223	**Olanzapine Tablets IP 2.50 mg**	**Each film coated tablet contains :**		
		Olanzapine	IP	2.50 mg.
		Excipients		q.s.
		Approved colour used		
224	**Olanzapine Tablets IP 5.0 mg**	**Each film coated tablet contains :**		
		Olanzapine	IP	5.0 mg.
		Excipients		q.s.
		Approved colour used		
225	**Olanzapine Tablets IP 7.50 mg**	**Each film coated tablet contains :**		
		Olanzapine	IP	7.50 mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
226	**Cetirizine Syrup IP**	**Each 5 ml contains :**		
		Cetirizine Hydrochloride	IP	5 mg.
		Excipients		q.s.
		Approved colour used		
227	**Carvedilol Tablets IP**	**Each uncoated tablet contains :**		
		Carvedilol	IP	3.125 mg.
		Excipients		q.s.
228	**Haloperidol Tablets IP 5 mg.**	**Each uncoated Tablets contains:**		
		Haloperidol	IP	5 mg.
		Excipients		q.s.
		Approved colour used		
229	**Haloperidol Tablets IP 10 mg.**	**Each uncoated Tablets contains:**		
		Haloperidol	IP	10 mg.
		Excipients		q.s.
		Approved colour used		
230	**Bromhexine Hcl, Dextromethorphan Hbr & Ammonium Chloride Syrup**	**Each 10 ml. contains :**		
		Bromhexine Hydrochloride	IP	8 mg.
		Dextromethorphan Hydrobromide	IP	10 mg.
		Ammonium Chloride	IP	100 mg.
		Menthol	IP	q.s.
		Flavoured Base		q.s.
		Approved colour used		
231	**Bromhexine Hcl, Dextromethorphan Hbr & Ammonium Chloride Syrup**	**Each 5 ml. contains :**		
		Bromhexine Hydrochloride	IP	4 mg.
		Dextromethorphan Hydrobromide	IP	5 mg.
		Ammonium Chloride	IP	50 mg.
		Menthol	IP	q.s.
		Flavoured Base		q.s.
		Approved colour used		
232	**Bromhexine Hcl, Dextromethorphan Hbr, Guaiphenesin & Chlorpheniaramine Maleate Tablets**	**Each uncoated tablet contains:**		
		Bromhexine Hydrochloride	IP	8 mg.
		Dextromethorphan Hydrobromide	IP	10 mg.
		Guaiphenesin	IP	100 mg.
		Chlorpheniramine Maleate	IP	2 mg.
		Excipients		q.s.
		Approved colour used		
233	**Levosulpiride Tablets 25 mg**	**Each uncoated tablet contains :**		
		Levosulpiride		25 mg.
		Excipients		q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
234	**Citicoline Tablets 500 mg.**	**Each filmcoated tablet contains:**		
		Citicoline Sodium		
		Equivalent to Citicoline		500 mg.
		Excipients		q.s.
		Approved colour used		
235	**Nimodipine Tablets BP 30 mg**	**Each filmcoated tablet contains:**		
		Nimodipine	BP	30 mg.
		Excipients		q.s.
		Approved colour used		
236	**Dextromethorphan Hbr., Phenylephrine Hcl. & Chlorpheniramine Maleate Syrup.**	**Each 5ml Contains:**		
		Dextromethorphan Hydrobromide	IP	10 mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Chlorpheniramine Maleate	IP	2 mg.
		Flavoured Base		q.s.
		Approved colour used		
237	**Dextromethorphan Hydrobromide & Chlorpheniramine Maleate Syrup**	**Each 5 ml. Contains :**		
		Dextromethorphan Hydrobromide	IP	10 mg.
		Chlorpheniramine Maleate	IP	2 mg.
		Flavoured Base		q.s.
		Approved colour used		
238	**Acebrophylline Capsules 100 mg.**	**Each hard gelatin capsule contains :**		
		Acebrophylline		100 mg.
		Excipients		q.s.
		Approved colour used in empty capsule shells		
239	**Montelukast Sodium & Bambuterol Hydrochloride Tablets**	**Each film coated tablet contains :**		
		Montelukast Sodium	IP	
		Equiv. to Montelukast		10 mg.
		Bambuterol Hydrochloride	IP	10 mg.
		Excipients		q.s.
		Apprved colour used		
240	**Ivermectin & Albendazole Tablets.**	**Each uncoated tablet contains :**		
		Ivermectin	IP	6 mg.
		Albendazole	IP	400 mg.
		Excipients		q.s.
241	**Phenylephrine HCl., Chlorpheniramine Maleate Paracetamol & Sodium Citrate Syrup**	**Each 5 ml contains :**		
		Phenylephrine Hydrochloride	IP	5 mg.
		Chlorpheniramine Maleate	IP	0.5 mg.
		Paracetamol	IP	125 mg.
		Sodium Citrate	IP	60 mg.
		Mentholated surupy base		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
242	**Ambroxol HCl., Guaiphenesin & Levosalbutamol Syrup**	**Each 5 ml. contains:**		
		Ambroxol Hydrochloride	IP	30 mg
		Guaiphenesin	IP	50 mg.
		Levosalbutamol Sulphate	IP	
		Equiv. to Levosalbutamol		1 mg.
		Flavoured base		q.s.
		Approved colour used		
243	**Sulfamethoxazole and Trimethoprim Oral Suspension USP**	**Each teaspoonful (5 mL) contains:**		
		Sulfamethoxazole	USP	200mg.
		Trimethoprim	USP	40 mg.
		Flavoured Syrupy Base		
		Approved Colour used		
244	**Linezolid Tablets IP 600 mg**	**Each film coated tablet contains**		
		Linezolid	IP	600 mg
		Excipients		q.s.
		Approved colour used		
245	**Esomeprazole Magnesium (Enteric Coated) & Domperidone (Sustained Release) Capsules**	**Each hard gelatin capsule contains :**		
		Esomeprazole Magnesium	IP	
		eq. to Esomeprazole		40 mg
		(as enteric coated pellets)		
		Domperidone maleate	IP	
		Eq. to Domperidone		30 mg
		(As sustained release pellets)		
		Colours: Approved colours used in empty shells & Pellets		
246	**Esomeprazole Magnesium Tablets IP**	**Each enteric coated tablet contains :**		
		Esomeprazole Magnesium (As Trihydrate)	IP	
		eq. to Esomeprazole		40 mg
		Excipients		q.s.
		Approved colour used		
247	**Doxofylline & Ambroxol Hydrochloride Tablets**	**Each uncoated tablet contains :**		
		Doxofylline	IP	400 mg
		Ambroxol Hydrochloride	IP	30 mg
		Excipients		q.s.
248	**Febuxostat Tablets 40 mg**	**Each film coated tablet contains**		
		Febuxostat		40 mg
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
249	**Febuxostat Tablets 80 mg**	**Each film coated tablet contains** Febuxostat Excipients Approved colour used		 80 mg q.s.
250	**Lornoxicam & Paracetamol Tablets**	**Each film coated tablet contains** Lornoxicam Paracetamol Excipients Approved colour used	 IP 	 8 mg 325 mg q.s.
251	**Citicoline & Piracetam Tablets**	**Each film coated tablet contains:** Citicoline Sodium Equivalent to Citicoline Piracetam Excipients Approved colour used	 IP 	 500 mg 400 mg q.s.
252	**Doxofylline Tablets IP 400 mg.**	**Each uncoated tablet contains :** Doxofylline Excipients	 IP 	 400 mg q.s.
253	**Ketoconazole Tablets IP 200 mg**	**Each uncoated tablet contains:** Ketoconazole Excipients	 IP 	 200 mg q.s.
254	**Diethylcarbamazine Tablets IP 50 mg**	**Each uncoated tablet contains:** Diethylcarbamazine Citrate Excipients Approved colour used	 IP 	 50 mg q.s.
255	**Diethylcarbamazine Tablets IP 100 mg**	**Each uncoated tablet contains:** Diethylcarbamazine Citrate Excipients Approved colour used	 IP 	 100 mg q.s.
256	**Ispaghula husk & Mebeverine hydrochloride Sachet**	**Each sachet contains** Ispaghula husk Mebeverine hydrochloride	 BP IP	 3.5 gm 135 mg
257	**Cetirizine Tablets IP**	**Each filmcoated tablet contains :** Cetirizine Hydrochloride Excipients	 IP 	 10 mg. q.s.
258	**Diclofenac Potassium , Paracetamol & Chlorzoxazone Tablets**	**Each filmcoated tablet contains** Diclofenac Potassium Paracetamol Chlorzoxazone Approved colour used	 BP IP USP 	 50 mg. 325 mg. 500 mg.

S.No	Generic Name	Composition	Ph.Ref	Claim
259	**Diclofenac Potassium & Paracetamol Tablets**	**Each filmcoated tablet contains**		
		Diclofenac Potassium	BP	50 mg.
		Paracetamol	IP	325 mg.
		Excipients		q.s.
260	**Acebrophylline Syrup**	**Each 5 ml contains**		
		Acebrophylline		50 mg
		Flavoured base		q.s.
		Approved colour used		
261	**Dried Aluminium Hydroxide Gel, Magnesium Hydroxide & Activated Dimethicone Suspension**	**Each 5 ml. contains :**		
		Dried Aluminium Hydroxide Gel	IP	250 mg.
		Magnesium Hydroxide	IP	250 mg.
		Activated Dimethicone	IP	50 mg.
		Sorbitol Solution 70 %	IP	1.25 g.
		(Non Crystallizing)		
		Flavoured base		q.s.
		Colour : Approved colour used		
262	**Dicyclomine Hydrochloride & Paracetamol Tablets**	**Each uncoated tablet contains :**		
		Dicyclomine Hydrochloride	IP	20 mg.
		Paracetamol	IP	500 mg.
		Excipients		q.s.
263	**Dicyclomine Hydrochloride & Acetaminophen Capsules**	**Each hard gelatin capsule contains:**		
		Dicyclomine Hydrochloride	IP	20 mg.
		Acetaminophen	IP	500 mg.
		Approved colours used in empty capsule shell		
264	**Paracetamol,Phenylephrine Hydrochloride, Caffeine Anhydrous & Diphenhydramine Hydrochloride Tablets**	**Each uncoated tablet contains :**		
		Paracetamol	IP	500 mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Caffeine (Anhydrous)	IP	30 mg.
		Diphenhydramine Hydrochloride	IP	25 mg.
		Excipients		q.s.
265	**Levodropropizine & Chlorpheniramine Maleate Syrup**	**Each 5ml contains :**		
		Levodropropizine	IP	30 mg.
		Chlorpheniramine Maleate	IP	2 mg.
		Flavoured Base		q.s.
		Approved colour used		
266	**Ambroxol Hydrochloride & Guaiphenesin Tablets**	**Each film coated tablet contain :**		
		Ambroxol Hydrochloride	IP	30 mg.
		Guaiphenesin	IP	200 mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
267	**Levocetirizine Dihydrochloride & Ambroxol Hydrochloride SR Tablets**	**Each uncoated bilayered tablet contains:** Levocetirizine Dihydrochloride Ambroxol Hydrochloride (As sustained release form) Excipients Approved colour used	 IP IP 	 5 mg. 75 mg. q.s.
268	**Mefenamic Acid & Paracetamol Suspension**	**Each 5ml. contains:** Mefenamic Acid Paracetamol Flavoured base Approved Colour used	 IP IP 	 100 mg. 250 mg. q.s.
269	**Mefenamic Acid Suspension**	**Each 5ml. contains:** Mefenamic Acid Flavoured base Approved Colour used	 IP 	 100 mg. q.s.
270	**Codeine Phosphate & Triprolidine Hydrochloride Syrup**	**Each 5ml contains** Codeine Phosphate Triprolidine Hydrochloride Flavoured base Approved colour used	 IP IP 	 10 mg. 1.25 mg q.s.
271	**Thiocolchicoside, Aceclofenac & Paracetamol Tablets**	**Each film-coated tablet contains :** Thiocolchicoside Aceclofenac Paracetamol Excipients Approved colour used	 IP IP IP 	 4 mg. 100 mg 325 mg. q.s.
272	**Etoricoxib Tablets IP 60 mg.**	**Each film coated tablet contains :** Etoricoxib Excipients Approved colour used	 IP 	 60 mg. q.s.
273	**Etoricoxib Tablets IP 90 mg.**	**Each film coated tablet contains :** Etoricoxib Excipients Approved colour used	 IP 	 90 mg. q.s.
274	**Etoricoxib Tablets IP 120 mg.**	**Each film coated tablet contains :** Etoricoxib Excipients Approved colour used	 IP 	 120 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
275	**Etoricoxib & Thiocolchicoside Tablets**	**Each film coated tablet contains :**		
		Etoricoxib	IP	60 mg.
		Thiocolchicoside	IP	4 mg.
		Excipients		q.s.
		Approved colour used		
277	**Rifaximin Tablets 200 mg.**	**Each film coated tablet contains :**		
		Rifaximin	BP	200 mg.
		Excipients		q.s.
		Approved colour used		
278	**Rifaximin Tablets 400 mg.**	**Each film coated tablet contains :**		
		Rifaximin	BP	400 mg.
		Excipients		q.s.
		Approved colour used		
279	**Rifaximin Tablets 550 mg.**	**Each film coated tablet contains :**		
		Rifaximin	BP	550 mg.
		Excipients		q.s.
		Approved colour used		
280	**Tolperisone Hydrochloride Tablets 150 mg.**	**Each film coated tablet contains :**		
		Tolperisone Hydrochloride		150 mg.
		Excipients		q.s.
		Approved colour used		
281	**Etoricoxib & Paracetamol Tablets**	**Each film coated tablet contains:**		
		Etoricoxib	IP	60 mg.
		Paracetamol	IP	325 mg.
		Excipients		q.s.
		Approved colour used		
282	**Telmisartan Tablets IP 40mg.**	**Each film coated tablet contains:**		
		Telmisartan	IP	40 mg.
		Excipients		q.s.
		Approved colour used		
283	**Telmisartan & Hydrochlorothiazide Tablets USP**	**Each film coated tablet contains:**		
		Telmisartan	IP	40 mg.
		Hydrochlorothiazide	IP	12.5mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
284	**Telmisartan &**	**Each film coated tablet contains:**		
	Amlodipine Besilate	Telmisartan	IP	40 mg.
	Tablets	Amlodipine Besilate	IP	
		Equivalent to Amlodipine		5 mg.
		Excipients		q.s.
		Approved colour used		
285	**Montelukast Sodium**	**Each uncoated dispersible tablet contains :**		
	& Levocetirizine	Montelukast Sodium	IP	
	Dihydrochloride	Equiv. to Montelukast		4 mg.
	Dispersible Tablets	Levocetirizine Dihydrochloride	IP	2.5 mg.
		Excipients		q.s.
		Approved colour used		
286	**Paracetamol,**	**Each uncoated Tablet Contains :**		
	Caffeine Anhydrous,	Paracetamol	IP	325mg.
	Phenylephrine HCl. &	Caffeine Anhydrous	IP	25 mg.
	Chlorpheniramine Maleate	Phenylephrine Hydrochloride	IP	5 mg.
	Tablets	Chlorpheniramine Maleate	IP	2 mg.
		Excipients		q.s.
287	**Pregabalin with**	**Each hard gelatin capsule contains :**		
	Methylcobalamin Capsules	Pregabalin	IP	75 mg
		Methylcobalamin		750 mcg
		Excipients		q.s.
		Approved colour used in empty capsule shells		
		Appropriate overages added to compensate loss on storage		
288	**Metformin HCl ER &**	**Each uncoated bilayered tablet contains:**		
	Glimepiride Tablets	Metformin Hydrochloride	IP	1000 mg.
		(In Extended release form)		
		Glimepiride	IP	1 mg.
		Colour : Approved colour used		
		Excipients		q.s.
289	**Metformin HCl ER &**	**Each uncoated bilayered tablet contains:**		
	Glimepiride Tablets	Metformin Hydrochloride	IP	1000 mg.
		(In Extended release form)		
		Glimepiride	IP	2 mg.
		Colour : Approved colour used		
		Excipients		q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
290	**Glimepiride ,Voglibose & Metformin ER Tablets**	**Each uncoated bilayered tablet contains:**		
		Glimepiride	IP	1 mg.
		Voglibose	IP	0.2 mg.
		Metformin Hydrochloride	IP	500 mg.
		(In Sustained release form)		
		Colour : Approved colour used		
		Excipients		q.s.
291	**Glimepiride ,Voglibose & Metformin ER Tablets**	**Each uncoated bilayered tablet contains:**		
		Glimepiride	IP	2 mg.
		Voglibose	IP	0.2 mg.
		Metformin Hydrochloride	IP	500 mg.
		(In Sustained release form)		
		Colour : Approved colour used		
		Excipients		q.s.
292	**Montelukast Sodium & Fexofenadine Hydrochloride Tablets**	**Each film coated tablet contains :**		
		Montelukast Sodium	IP	
		Equiv. to Montelukast		10 mg.
		Fexofenadine Hydrochloride	IP	120 mg.
		Excipients		q.s.
		Apprved colour used		
293	**Potassium Citrate Monohydrate, Magnesium Citarte & Vitamin B6 Oral Solution**	**Each 5 ml. contains:**		
		Potassium Citrate Monohydrate	IP	1100 mg.
		Magnesium Citrate	USP	375 mg.
		Vitamin B6	IP	20 mg.
		Flavoured Sorbitol Base		q.s.
		Approved colour used		
294	**Flavoxate Hydrochloride & Ofloxacin Tablets**	**Each film coated tablet contains :**		
		Flavoxate Hydrochloride	BP	200 mg.
		Ofloxacin	IP	200 mg.
		Excipients		q.s.
		Apprved colour used		
295	**Paracetamol, Tramadol Hydrochloride & Domperidone Tablets**	**Each uncoated tablet contains :**		
		Paracetamol	IP	325 mg.
		Tramadol Hydrochloride	IP	37.50 mg.
		Domperidone	IP	10 mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
296	**Acebrophylline , Terbutaline Sulphate & Guaiphenesin Syrup**	**Each 5 ml contains :**		
		Acebrophylline		50 mg.
		Terbutaline Sulphate	IP	1.25 mg.
		Guaiphenesin	IP	50 mg.
		Mentholated Syrupy base		q.s.
		Approved colour used		
297	**Montelukast Sodium, Levocetirizine Dihydrochloride & Ambroxol Hydrochloride SR Tablets**	**Each film-coated tablet contains :**		
		Montelukast Sodium	IP	
		Equiv. to Montelukast		10 mg.
		Levocetirizine Dihydrochloride	IP	5 mg.
		Ambroxol Hydrochloride	IP	75 mg.
		(In Sustained release form)		
		Excipients		q.s.
		Approved colour used		
298	**Ofloxacin, Ornidazole & Lactic Acid Bacillus Tablets**	**Each film coated tablet contains:**		
		Ofloxacin	IP	200 mg.
		Ornidazole	IP	500 mg.
		Lactic Acid Bacillus Viable Spores		60 Million
		Excipients		q.s.
		Approved colour used		
299	**Gabapentin & Nortriptyline Hydrochloride Tablets**	**Each film coated tablet contains:**		
		Gabapentin	IP	400 mg.
		Nortriptyline Hydrochloride	IP	
		Equiv. to Nortriptyline		10 mg.
		Excipients		q.s.
		Approved colour used		
300	**Ursodeoxycholic Acid Sustained Release Tablets**	**Each film coated tablet contains:**		
		Ursodeoxycholic Acid	IP	450 mg.
		(In Sustained release form)		
		Excipients		q.s.
		Approved colour used		
301	**Montelukast Sodium, Levocetirizine Dihydrochloride & Acebrophylline SR Tablets**	**Each film-coated tablet contains :**		
		Montelukast Sodium	IP	
		Equiv. to Montelukast		10 mg.
		Levocetirizine Dihydrochloride	IP	5 mg.
		Acebrophylline		200 mg.
		(In Sustained release form)		
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
302	**Paracetamol & Domperidone Maleate Suspension**	**Each 5 ml contains**		
		Paracetamol	IP	125 mg.
		Domperidone Maleate	IP	5 mg.
		Flavoured Base		q.s.
		Approved colour used		
303	**Paracetamol & Dicyclomine Hydrochloride Suspension**	**Each 5 ml contains**		
		Paracetamol	IP	125 mg.
		Dicyclomine Hydrochloride	IP	5 mg.
		Flavoured Base		q.s.
		Approved colour used		
304	**Ambroxol Hydrochloride , Guaiphenesin & Levosalbutamol Drops**	**Each ml. contains:**		
		Ambroxol Hydrochloride	IP	7. 5 mg
		Guaiphenesin	IP	12.5 mg
		Levosalbutamol Sulphate	IP	
		Equiv. to Levosalbutamol		0.25 mg
		Flavoured base		q.s.
		Approved colour used		
305	**Potassium Citrate , Citric Acid & Sodium Citrate Oral Solution**	**Each 5 ml. Contains :**		
		Potassium Citrate	IP	550 mg.
		Citric Acid	IP	334 mg.
		Sodium Citrate	IP	500 mg.
		Flavoured Sugar free Base		q.s.
		Approved colour used		
309	**Paracetamol, Phenylephrine Hydrochloride & Chlorpheniramine Maleate Drops**	**Each ml. Contains :**		
		Paracetamol	IP	125 mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Chlorpheniramine Maleate	IP	1 mg.
		Flavoured base		q.s.
		Approved colour used		
310	**Paracetamol, Phenylephrine Hydrochloride & Chlorpheniramine Maleate Drops**	**Each ml. Contains :**		
		Paracetamol	IP	125 mg.
		Phenylephrine Hydrochloride	IP	2.5 mg.
		Chlorpheniramine Maleate	IP	1 mg.
		Flavoured base		q.s.
		Approved colour used		
311	**Dicyclomine Hydrochloride & Activated Dimethicone Drops**	**Each ml. Contains :**		
		Dicyclomine Hydrochloride	IP	10 mg.
		Activated Dimethicone	IP	40 mg.
		Flavoured base		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
312	**Domperidone Suspension IP**	**Each ml. contains :** Domperidone Flavoured base Approved colour used	 IP 	 1 mg. q.s.
313	**Hydroxyzine Hydrochloride Oral Solution USP**	**Each ml. contains :** Hydroxyzine Hydrochloride Flavoured base Approved colour used	 IP 	 6 mg. q.s.
314	**Ambroxol Hydrochloride Drops**	**Each ml. contains :** Ambroxol Hydrochloride Flavoured base Approved colour used	 IP 	 7.5 mg. q.s.
315	**Tamsulosin Hydrochloride (Modified Release) & Finasteride Tablets**	**Each film coated tablet contains** Tamsulosin Hydrochloride (in a modified release form) Finasteride Excipients Colour : Approved colours used	 IP IP 	 0.4 mg 5mg q.s.
316	**Telmisartan & Cilnidipine Tablets**	**Each film coated tablet contains:** Telmisartan Cilnidipine Excipients Colour : Approved colours used	 IP IP 	 40 mg 10 mg q.s.
317	**Telmisartan & Cilnidipine Tablets**	**Each film coated tablet contains:** Telmisartan Cilnidipine Excipients Colour : Approved colours used	 IP IP 	 80 mg 10 mg q.s.
318	**Montelukast Sodium & Doxofylline SR Tablets**	**Each uncoated bilayered tablet contains :** Montelukast Sodium Equiv. to Montelukast Doxofylline (As sustained release) Excipients Approved colour used	 IP IP 	 10 mg. 400 mg q.s.
319	**Enteric Coated Rabeprazole Sodium & Levosulpiride Sustained Release Capsules**	**Each hard gelatin capsule contains :** Rabeprazole Sodium (As enteric coated pellets) Levosulpiride (As sustained release pellets Excipients Approved colour used in empty capsule shells & Pellets	 IP 	 20 mg 75 mg q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
320	**Torsemide Tablets 5 mg**	**Each uncoated tablet contains:** Torsemide Equiv. to Torsemide (Anhydrous) Approved colour used	USP	 5 mg.
321	**Deflazacort Tablets 18 mg**	**Each uncoated tablets contains :** Deflazacort Exicipents		 18 mg q.s.
322	**Nitroglycerin Controlled Release Tablets**	**Each uncoated tablet contains :** Diluted Nitroglycerin Equivalent to Nitroglycerin (In a controlled release system) Excipients	IP	 2.6 mg. q.s.
323	**Ergotamine Tablets IP 1 mg.**	**Each uncoated tablet contains:** Ergotamine Tartrate Excipients Approved colour used	IP	 1 mg. q.s.
324	**Voglibose Tablets IP 0.2 mg.**	**Each uncoated tablet contains:** Voglibose Excipients	IP	 0.2 mg. q.s.
325	**Voglibose Tablets IP 0.3 mg.**	**Each uncoated tablet contains:** Voglibose Excipients Approved colour used		 0.3 mg. q.s.
326	**Montelukast Sodium Tablets IP 5 mg.**	**Each film coated tablet contains :** Montelukast Sodium Equivalent to Montelukast Excipients Apprved colour used	IP	 5 mg. q.s.
327	**Montelukast Sodium Tablets IP 10 mg.**	**Each film coated tablet contains :** Montelukast Sodium Equivalent to Montelukast Excipients Apprved colour used	IP	 10 mg. q.s.
328	**Levocetirizine Tablets IP 10 mg**	**Each film-coated tablet contains:** Levocetirizine Dihydrochloride Excipients Approved colour used	IP	 10 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
329	**Mefenamic Acid & Paracetamol Tablets**	**Each uncoated tablet contains :**		
		Mefenamic Acid	IP	500 mg.
		Paracetamol	IP	325 mg.
		Excipients		q.s.
		Approved colour used		
330	**Ondansetron Orally Disintegrating Tablets IP 8 mg.**	**Each uncoated mouth dissolving tablet contains:**		
		Ondansetron	IP	8 mg
		Excipients		q.s.
		Approved colour used		
331	**Rosuvastatin Tablets IP 5 mg.**	**Each film coated tablet contains :**		
		Rosuvastatin Calcium	IP	
		Equivalent to Rosuvastatin		5 mg.
		Excipients		q.s.
		Approved colour used		
332	**Rosuvastatin Tablets IP 10 mg.**	**Each film coated tablet contains :**		
		Rosuvastatin Calcium	IP	
		Equivalent to Rosuvastatin		10 mg.
		Excipients		q.s.
		Approved colour used		
333	**Rosuvastatin Tablets IP 20 mg.**	**Each film coated tablet contains :**		
		Rosuvastatin Calcium	IP	
		Equivalent to Rosuvastatin		20 mg.
		Excipients		q.s.
		Approved colour used		
334	**Olmesartan Medoxomil Tablets IP 20 mg**	**Each film coated tablet contains :**		
		Olmesartan Medoxomil	IP	20 mg.
		Excipients		q.s.
		Approved colour used		
335	**Olmesartan Medoxomil Tablets IP 40 mg**	**Each film coated tablet contains :**		
		Olmesartan Medoxomil	IP	40 mg.
		Excipients		q.s.
		Approved colour used		
336	**Pregabalin Capsules IP 75 mg**	**Each hard gelatin capsule contains :**		
		Pregabalin	IP	75 mg.
		Excipients		q.s.
		Approved colour used in empty capsule shells		
337	**Paracetamol, Chlorpheniramine Maleate & Phenylephrine Hydrochloride Oral Solution**	**Each 5 ml. Contains :**		
		Paracetamol	IP	250 mg.
		Phenylephrine Hydrochloride	IP	5 mg.
		Chlorpheniramine Maleate	IP	2 mg.
		Flavoured base		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
338	**Sucralfate & Oxetacaine Oral Suspension**	**Each 10 ml. contains :** Sucralfate Oxetacaine Flavoured sorbitol base Approved colour used	 USP BP 	 1 gm. 10 mg. q.s.
339	**Fexofenadine Hydrochloride Tablets IP 120 mg**	**Each film coated tablet contains :** Fexofenadine Hydrochloride Excipients Approved colour used	 IP 	 120 mg. q.s.
340	**Fexofenadine Hydrochloride Tablets IP 180 mg**	**Each film coated tablet contains :** Fexofenadine Hydrochloride Excipients Approved colour used	 IP 	 180 mg. q.s.
341	**Hydroxychloroquine Tablets IP 200 mg**	**Each film coated tablet contains :** Hydroxychloroquine Sulphate Excipients Approved colour used	 IP 	 200 mg. q.s.
342	**Bromhexine Hydrochloride Syrup 4 mg**	**Each 5ml contains :** Bromhexine Hydrochloride Flavoured base Approved colour used	 IP 	 4 mg. q.s.
343	**Torsemide Tablets IP 100 mg**	**Each uncoated tablet contains:** Torsemide Excipients	 IP 	 100mg. q.s.
344	**Doxofylline & Terbutaline Sulphate Tablets**	**Each uncoated tablet contains :** Doxofylline Terbutaline Sulphate Excipients Approved colour used	 IP IP 	 400 mg 5 mg q.s.
345	**Magaldrate & Simethicone Oral Suspension IP** **XOM-MPS**	**Each 5ml. contains :** Magaldrate Simethicone Flavoured base Approved colour used	 IP IP 	 480 mg. 20 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
346	**Aluminium Hydroxide , Magnesium Hydroxide & Simethicone Oral Suspension IP**	**Each 5 ml. contains :**		
		Dried Aluminium Hydroxide Gel	IP	250 mg.
		Magnesium Hydroxide	IP	250 mg.
		Simethicone	IP	50 mg.
		Sorbitol Solution 70 % (Non Crystallizing)	IP	q.s.
		Flavoured base		q.s.
		Colour : Approved colour used		
347	**Aluminium Hydroxide , Magnesium Hydroxide & Simethicone Oral Suspension IP**	**Each 5 ml. contains :**		
		Dried Aluminium Hydroxide Gel	IP	200 mg.
		Magnesium Hydroxide	IP	200 mg.
		Simethicone	IP	25 mg.
		Sorbitol Solution 70 % (Non Crystallizing)	IP	q.s.
		Flavoured base		q.s.
		Colour : Approved colour used		
348	**Drotaverine Hydrochloride & Aceclofenac Tablets**	**Each film coated tablet contains:**		
		Drotaverine Hydrochloride	IP	80 mg.
		Aceclofenac	IP	100mg.
		Excipients		q.s.
		Approved colour used		
349	**Propranolol Hydrochloride Sustained-release & Flunarizine Dihydrochloride Capsules**	**Each hard gelatin capsule contains :**		
		Propranolol Hydrochloride (As Sustained-release pellets)	IP	40 mg
		Flunarizine Dihydrochloride	BP	
		Eq. to Flunarizine		5 mg.
		Excipients		q.s.
		Approved colour used in empty capsule shells & Pellets		
350	**Propranolol Hydrochloride Sustained-release & Flunarizine Dihydrochloride Capsules**	**Each hard gelatin capsule contains :**		
		Propranolol Hydrochloride (As Sustained-release pellets)	IP	40 mg
		Flunarizine Dihydrochloride	BP	
		Eq. to Flunarizine		10 mg.
		Excipients		q.s.
		Approved colour used in empty capsule shells & Pellets		
351	**Diclofenac Potassium & Chlorzoxazone Tablets**	**Each film coated tablet contains**		
		Diclofenac Potassium	BP	50 mg.
		Chlorzoxazone	USP	500 mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
352	**Dicyclomine Hydrochloride, Aluminium Hydroxide , Magnesium Hydroxide & Simethicone Oral Suspension**	**Each 5 ml. contains :** Dicyclomine Hydrochloride Dried Aluminium Hydroxide Gel Magnesium Hydroxide Simethicone Flavoured base Colour : Approved colour used	 IP IP IP IP	 2.5 mg. 200 mg. 100 mg. 20 mg. q.s.
353	**Etoricoxib & Thiocolchicoside Tablets**	**Each film coated tablet contains :** Etoricoxib Thiocolchicoside Excipients Approved colour used	 IP IP	 60 mg. 8 mg. q.s.
354	**Acebrophylline Sustained release Tablets 200 mg.**	**Each film coated sustained release tablet contains :** Acebrophylline Excipients Approved colour used		 200 mg. q.s.
355	**Theophylline Prolonged-release Tablets IP 200 mg.**	**Each uncoated tablet contains :** Theophylline (Anhydrous) (In Prolonged-release form) Excipients	 IP	 200 mg. q.s.
356	**Theophylline Prolonged-release Tablets IP 400 mg.**	**Each uncoated tablet contains :** Theophylline (Anhydrous) (In Prolonged-release form) Excipients Approved colour used	 IP	 400 mg. q.s.
357	**Enteric Coated Pantoprazole Sodium & Levosulpiride Sustained release Capsules**	**Each hard gelatin capsule contains :** Pantoprazole Sodium Equivalent to Pantoprazole (As enteric coated pellets) Levosulpiride (As sustained release pellets Excipients Approved colour used in empty capsule shells & Pellets	 IP	 40 mg. 75 mg. q.s.
358	**Finasteride Tablets IP 1mg.**	**Each film coated tablet contains** Finasteride Excipients Colour : Approved colours used	 IP	 1 mg. q.s.
359	**Diacerein Capsules IP 50 mg.**	**Each hard gelatin capsule contains :** Diacerein Excipients Approved colours used in capsule shell.	 IP	 50 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
360	**Ibuprofen & Paracetamol Tablets IP**	**Each uncoated tablet contains :** Ibuprofen Paracetamol Excipients	 IP IP	 400 mg. 325 mg. q.s.
361	**Levocloperastine Fendizoate Oral Suspension**	**Each 5ml of suspension contains:** Levocloperastine Fendizoate Equivalent to Levocloperastine Hydrochloride Flavoured base Approved colour & flavour used		 35.4 mg. 20 mg. q.s.
362	**Tapentadol Hydrochloride Tablets 50 mg.**	**Each film coated tablet contains :** Tapentadol Hydrochloride Eq. to Tapentadol Excipients Approved colour used	 IP	 50 mg. q.s.
363	**Tapentadol Hydrochloride Tablets 75 mg.**	**Each film coated tablet contains :** Tapentadol Hydrochloride Eq. to Tapentadol Excipients Approved colour used	 IP	 75 mg. q.s.
364	**Tapentadol Hydrochloride Tablets 100 mg.**	**Each film coated tablet contains :** Tapentadol Hydrochloride Eq. to Tapentadol Excipients Approved colour used	 IP	 100 mg. q.s.
365	**Tapentadol Extended-release Tablets 50 mg.**	**Each film coated extended-release tablet contains :** Tapentadol Hydrochloride Eq. to Tapentadol Excipients Approved colour used	 IP	 50 mg. q.s.
366	**Clopidogrel Tablets IP 75 mg.**	**Each film coated tablet contains :** Clopidogrel Bisulphate Eq. to Clopidogrel Excipients Approved colour used	 IP	 75 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
367	**Clopidogrel Tablets IP 150 mg.**	**Each film coated tablet contains :**		
		Clopidogrel Bisulphate	IP	
		Eq. to Clopidogrel		150 mg.
		Excipients		q.s.
		Approved colour used		
368	**Clopidogrel 75 mg. & Aspirin 75 mg. Tablets**	**Each film coated bilayered tablet contains :**		
		Clopidogrel Bisulphate	IP	
		Eq. to Clopidogrel		75 mg.
		Aspirin	IP	75 mg.
		Excipients		q.s.
		Approved colour used		
369	**Clopidogrel 150 mg. & Aspirin 75mg. Tablets**	**Each film coated bilayered tablet contains :**		
		Clopidogrel Bisulphate	IP	
		Eq. to Clopidogrel		150mg.
		Aspirin	IP	75 mg.
		Excipients		q.s.
		Approved colour used		
370	**Clopidogrel 75 mg. & Aspirin 75 mg. Capsules**	**Each hard gelatin capsule contains :**		
		Clopidogrel Bisulphate	IP	
		Eq. to Clopidogrel		75 mg.
		Aspirin	IP	75 mg.
		(As enteric coated form)		
		Excipients		q.s.
		Approved colour used in empty capsule shells		
371	**Clopidogrel 75 mg. & Aspirin 150 mg. Capsules**	**Each hard gelatin capsule contains :**		
		Clopidogrel Bisulphate	IP	
		Eq. to Clopidogrel		75 mg.
		Aspirin	IP	150 mg.
		(As enteric coated form)		
		Excipients		q.s.
		Approved colour used in empty capsule shells		
372	**Linagliptin Tablets 5 mg.**	**Each film coated tablet contains :**		
		Linagliptin		5 mg.
		Excipients		q.s.
		Approved colour used		

S.No	Generic Name	Composition	Ph.Ref	Claim
373	**Terbinafine Tablets IP 250 mg.**	**Each uncoated tablet contains :** Terbinafine Hydrochloride Eq. to Terbinafine Excipients	 IP 	 250mg. q.s.
374	**Terbinafine Tablets IP 500 mg.**	**Each uncoated tablet contains :** Terbinafine Hydrochloride Eq. to Terbinafine Excipients	 IP 	 500mg. q.s.
375	**Esomeprazole Magnesium Tablets IP 20 mg.**	**Each enteric coated tablet contains :** Esomeprazole Magnesium (Trihydrate) eq. to Esomeprazole Excipients Approved colour used	 IP 	 20 mg q.s.
376	**Omeprazole Capsules IP 40 mg.**	**Each hard gelatin capsule contains :** Omeprazole (As enteric coated pellets) Approved colour used in empty shell	 IP 	 40 mg.
377	**Esomeprazole Gastro-resistant Capsules IP 20 mg**	**Each hard gelatin capsule contains :** Esomeprazole Magnesium (Trihydrate) eq. to Esomeprazole (as Gastro-resitant pellets) Colours: Approved colours used in empty shells & Pellets	 IP 	 20 mg
378	**Esomeprazole Gastro-resistant Capsules IP 40 mg**	**Each hard gelatin capsule contains :** Esomeprazole Magnesium (Trihydrate) eq. to Esomeprazole (as Gastro-resitant pellets) Colours: Approved colours used in empty shells & Pellets	 IP 	 40 mg
379	**Piroxicam Dispersible Tablets 20 mg.**	**Each dispersible uncoated tablet contains :** Piroxicam Excipients	 IP 	 20 mg. q.s.
380	**Itraconazole Capsules 100 mg.**	**Each hard gelatin capsule contains :** Itraconazole Pellets eq. to Itraconazole Excipients Colours: Approved colours used in empty shells & Pellets	 BP 	 100 mg q.s.
381	**Itraconazole Capsules 200 mg.**	**Each hard gelatin capsule contains :** Itraconazole Pellets eq. to Itraconazole Excipients Colours: Approved colours used in empty shells & Pellets	 BP 	 200 mg q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
382	**Omeprazole Capsules IP 20mg.** **XOM-20**	**Each hard gelatin capsule contains :** Omeprazole (As enteric coated pellets) Approved colour used in empty shell	IP	20 mg.
383	**Esomeprazole Gastro-resistant Capsules IP 40 mg** **HYPOSIL 40**	**Each hard gelatin capsule contains :** Esomeprazole Magnesium (Trihydrate) eq. to Esomeprazole (as Gastro-resitant pellets) Colours: Approved colours used in empty shells & Pellets	IP	40 mg
384	**Piroxicam Dispersible Tablets 20 mg.** **XPRAM 20 DT**	**Each dispersible uncoated tablet contains :** Piroxicam Excipients	IP	20 mg. q.s.
385	**Telmisartan Tablets IP 20mg.**	**Each film coated tablet contains:** Telmisartan Excipients Approved colour used	IP	20 mg. q.s.
386	**Telmisartan Tablets IP 80mg.**	**Each film coated tablet contains:** Telmisartan Excipients Approved colour used	IP	80 mg. q.s.
387	**Mefenamic Acid Tablets 500 mg.** **MEFNISPAS**	**Each film coated tablet contains:** Mefenamic Acid Excipients Approved colour used	USP	500 mg. q.s.
388	**Atorvastatin Tablets 40 mg** **ATORNIX-40**	**Each film-coated tablet contains :** Atorvastatin calcium Equiv. to Atorvastatin Excipients Approved colour used	USP	40 mg. q.s.
389	**Metformin Hydrochloride Extended-release Tablets USP 750 mg.** **METFORNIX ER 750**	**Each film-coated Extended-release tablet contains:** Metformin Hydrochloride Excipients Approved colour used	USP	750 mg. q.s.
390	**Metformin Hydrochloride Tablets USP 500 mg.** **METFORNIX 500**	**Each film-coated tablet contains:** Metformin Hydrochloride Excipients Approved colour used	USP	500 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
391	**Metformin Hydrochloride Tablets USP 850 mg. METFORNIX 850**	**Each film-coated tablet contains:** Metformin Hydrochloride Excipients Approved colour used	USP	850 mg. q.s.
392	**Tadalafil Tablets USP 20 mg. TADANIX-20**	**Each film-coated tablet contains:** Tadalafil Excipients Approved colour used	USP	20 mg. q.s.
393	**Lansoprazole Delayed-release Capsules USP 30 mg. UNILENZ-30**	**Each hard gelatin capsule contains :** Lansoprazole (As enetric coated granules) Approved colour used in shells & granules	USP	30 mg.
394	**Diclofenac Sodium Delayed-release Tablets USP 50 mg. DICLONIX-50**	**Each enteric coated tablet contains:** Diclofenac Sodium Excipients Approved colour used	USP	50 mg. q.s.
395	**Metronidazole Tablets USP 500 mg. TONIZOL-500**	**Each film-coated tablet contains:** Metronidazole Excipients Approved colour used	USP	500 mg. q.s.
396	**Pregabalin Capsules 50 mg PREGANIX-50**	**Each hard gelatin capsule contains :** Pregabalin Excipients Approved colour used in empty capsule shells		50 mg. q.s.
397	**Meloxicam Tablets USP 15 mg. MELOXINEX-15**	**Each uncoated tablet contains:** Meloxicam Excipients	USP	15 mg. q.s.
398	**Clarithromycin Tablets USP 250 mg. KLATH-250**	**Each film coated tablet contains:** Clarithromycin Excipients Approved colour used	USP	250 mg. q.s.
399	**Clarithromycin Tablets USP 500 mg. KLATH-500**	**Each film coated tablet contains:** Clarithromycin Excipients Approved colour used	USP	500 mg. q.s.
400	**Amlodipine Besilate Tablets USP 10 mg AMLONIX-10**	**Each uncoated tablet contains:** Amlodipine Besilate Eq. to Amlodipine Excipients	USP	10 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
401	**Cilostazol Tablets USP 50 mg.** **PLATAZOL-50**	**Each uncoated tablet contains:** Cilostazol Excipients	 USP	 50 mg. q.s.
402	**Cilostazol Tablets USP 100 mg.** **PLATAZOL-100**	**Each uncoated tablet contains:** Cilostazol Excipients	 USP	 100 mg. q.s.
403	**Glimepiride Tablets USP 1 mg.** **GLI-1**	**Each uncoated tablet contains:** Glimepiride Excipients Approved colour used	 USP	 1 mg. q.s.
404	**Glimepiride Tablets USP 2 mg.** **GLI-2**	**Each uncoated tablet contains:** Glimepiride Excipients Approved colour used	 USP	 2 mg. q.s.
405	**Bromhexine Hydrochloride &** **Phenylephrine Hydrochloride** **Tablets** **BROHEXINEX-P**	**Each uncoated tablet contains:** Bromhexine Hydrochloride Phenylephrine Hydrochloride Excipients	 USP USP	 8 mg. 5 mg. q.s.
406	**Paracetamol and** **Domperidone Tablets** **U-MOL D**	**Each uncoated tablet contains:** Acetaminophen (Paracetamol) Domperidone Excipients	 USP USP	 325 mg. 10 mg. q.s.
407	**Allopurinol Tablets USP 100 mg.** **UPRINOL-100**	**Each uncoated tablet contains:** Allopurinol Excipients	 USP	 100 mg. q.s.
408	**Allopurinol Tablets USP 300 mg.** **UPRINOL-300**	**Each uncoated tablet contains:** Allopurinol Excipients	 USP	 300 mg. q.s.
409	**Metoclopramide Hydrochloride** **Tablets USP 10 mg.** **UNINORM**	**Each uncoated tablet contains:** Metoclopramide Hydrochloride equivalent to anhydrous Metoclopramide Hydrochloride Excipient	 USP	 10 mg. q.s.
410	**Paracetamol and** **Metoclopramide Hydrochloride** **Tablets** **UNINORM-P**	**Each uncoated tablet contains:** Acetaminophen (Paracetamol) Metoclopramide Hydrochloride equivalent to anhydrous Metoclopramide Hydrochloride Excipient	 USP USP	 325 mg. 10 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
411	**Omeprazole Capsules BP** **XOM-20**	**Each hard gelatin capsule contains :** Omeprazole (As enteric coated pellets) Approved colour used in empty shell	BP	20 mg.
412	**Piroxicam Dispersible Tablets 20 mg.** **XPRAM 20 DT**	**Each dispersible uncoated tablet contains :** Piroxicam Excipients	USP	20 mg. q.s.
413	**Ciprofloxacin Hydrochloride Tablets BP 250 mg** **LIDIK-250**	**Each film coated tablet contains:** Ciprofloxacin Hydrochloride Eq. to Ciprofloxacin Excipients Approved colour used	BP	 250 mg. q.s.
414	**Ciprofloxacin Hydrochloride Tablets BP 500 mg** **LIDIK-500**	**Each film coated tablet contains:** Ciprofloxacin Hydrochloride Eq. to Ciprofloxacin Excipients Approved colour used	BP	 500 mg. q.s.
415	**Levofloxacin Tablets USP 500 mg** **LE-FOX**	**Each film coated tablet contains :** Levofloxacin Hemihydrate Equiv. to Levofloxacin Excipients Approved colour used	USP	 500 mg. q.s.
416	**Lansoprazole Delayed-release Capsules BP 30 mg.** **UNILENZ-30**	**Each hard gelatin capsule contains :** Lansoprazole (As enetric coated granules) Approved colour used in shells & granules	BP	30 mg.
417	**Cetirizine Tablets BP** **SETOR**	**Each filmcoated tablet contains :** Cetirizine Hydrochloride Excipients Approved colour used	BP	10 mg. q.s.
418	**Atorvastatin Tablets USP 10 mg** **ATORNIX-10**	**Each film-coated tablet contains :** Atorvastatin calcium Equiv. to Atorvastatin Excipients Approved colour used	USP	 10 mg. q.s.
419	**Atorvastatin Tablets USP 20 mg** **ATORNIX-20**	**Each film-coated tablet contains :** Atorvastatin calcium Equiv. to Atorvastatin Excipients Approved colour used	USP	 20 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
420	**Atorvastatin Tablets USP 40 mg ATORNIX-40**	**Each film-coated tablet contains :** Atorvastatin calcium Equiv. to Atorvastatin Excipients Approved colour used	USP	 40 mg. q.s.
421	**Pregabalin Capsules 50 mg PREGANIX-50**	**Each hard gelatin capsule contains :** Pregabalin Excipients Approved colour used in empty capsule shells	BP	50 mg. q.s.
422	**Pregabalin Capsules 75 mg PREGANIX**	**Each hard gelatin capsule contains :** Pregabalin Excipients Approved colour used in empty capsule shells	BP	75 mg. q.s.
423	**Meloxicam Tablets BP 15 mg. MELOXINEX-15**	**Each uncoated tablet contains:** Meloxicam Excipients	BP	15 mg. q.s.
424	**Diclofenac Tablets BP 50 mg. DICLONIX-50**	**Each enteric coated tablet contains:** Diclofenac Sodium Excipients Approved colour used	BP	50 mg. q.s.
425	**Itraconazole Capsules BP 100 mg. IT-Nix 100**	**Each hard gelatin capsule contains :** Itraconazole Pellets eq. to Itraconazole Excipients Colours: Approved colours used in empty shells & Pellets	BP	 100 mg q.s.
426	**Itraconazole Capsules BP 200 mg. IT-Nix 200**	**Each hard gelatin capsule contains :** Itraconazole Pellets eq. to Itraconazole Excipients Colours: Approved colours used in empty shells & Pellets	BP	 200 mg q.s.
427	**Esomeprazole Delayed-release Capsules USP 40 mg HYPOSIL 40**	**Each hard gelatin capsule contains :** Esomeprazole Magnesium eq. to Esomeprazole (as Gastro-resitant pellets) Colours: Approved colours used in empty shells & Pellets	USP	 40 mg
428	**Etoricoxib Tablets 60 mg. ETORNIX-60**	**Each film coated tablet contains :** Etoricoxib Excipients Approved colour used		 60 mg. q.s.

S.No	Generic Name	Composition	Ph.Ref	Claim
429	**Etoricoxib**	**Each film coated tablet contains :**		
	Tablets 90 mg.	Etoricoxib		90 mg.
	ETORNIX-90	Excipients		q.s.
		Approved colour used		
430	**Etoricoxib**	**Each film coated tablet contains :**		
	Tablets 120 mg.	Etoricoxib		120 mg.
	ETORNIX-120	Excipients		q.s.
		Approved colour used		
431	**Montelukast**	**Each film coated tablet contains :**		
	Tablets BP 10 mg.	Montelukast Sodium	BP	
		Equivalent to Montelukast		10 mg.
	NEUMONT-10	Excipients		q.s.
		Apprved colour used		
432	**Pantoprazole**	**Each enteric coated tablet contains:**		
	Gastro-resistant Tablets BP	Pantoprazole Sodium Sesquihydrate	BP	
	40 mg.	Equivalent to Pantoprazole		40 mg.
	PANTS-40	Excipients		q.s.
		Approved colour used		
433	**Ofloxacin & Ornidazole**	**Each film coated tablet contains:**		
	Tablets	Ofloxacin	BP	200 mg.
		Ornidazole		500 mg.
	FOX-OZ	Excipients		q.s.
		Approved colour used		
434	**Omeprazole &**	**Each hard gelatin capsule contains :**		
	Domperidone Capsules	Omeprazole	BP	20 mg.
		(As enteric coated pellets)		
	XOM-D	Domperidone	BP	10 mg.
		Approved colour used in empty shell		
435	**Montelukast Sodium**	**Each film-coated tablet contains :**		
	& Levocetirizine	Montelukast Sodium	BP	
	Dihydrochloride Tablets	Equiv. to Montelukast		10 mg.
		Levocetirizine Dihydrochloride	USP	5 mg.
	NEUMONT-LC	Excipients		q.s.
		Approved colour used		
436	**Pregabalin &**	**Each hard gelatin capsule contains :**		
	Mecobalamin Capsules	Pregabalin	BP	75 mg
		Mecobalamin	JP	750 mcg
		Excipients		q.s.
	PREGANIX-M	Approved colour used in empty capsule shells.		
		Appropriate overages added to compensate loss on storage.		
437	**Levofloxacin**	**Each film coated tablet contains :**		
	Tablets USP 750 mg	Levofloxacin Hemihydrate	USP	
		Equiv. to Levofloxacin		750 mg
	LE-FOX 750	Excipients		q.s.
		Approved colour used		