

# PRODUCT LIST OF TRIREME LIFE SCIENCES PVT. LTD.

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NO: HFW- H (Drugs) 286/06,Health and Family Welfare Department,Himachal Pradesh

Sr. No.	Generic Name	Composition	PH. Ref.	Strength	STATUS OF PRODUCTS
1	<b>OLANZAPINE TABLETS 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	5 mg	
		Excipients		q.s.	
2	<b>OLANZAPINE TABLETS 10 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	10 mg	
		Excipients		q.s.	
3	<b>ESCITALOPRAM &amp; CLONAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	IP		
		eq. to Escitalopram		10mg	
		Clonazepam	IP	0.5mg	
		Excipients		q.s.	
		Approved colour used			
4	<b>PROPRANOLOL HYDROCHLORIDE EXTENDED RELEASE CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Propranolol Hydrochloride	IP	40 mg	
		(as extended release pellets)			
		Excipients		q.s.	
		approved colour used in empty capsule shells			
5	<b>TRIHEXYPHENIDYL HCl TABLETS IP 2MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
6	<b>PROPRANOLOL HCl TABLETS IP 40MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Propranolol Hcl	IP	40mg	
		Excipients		q.s.	
7	<b>CLOZAPINE TABLETS IP 100 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clozapine	IP	100 mg	
		Excipients		q.s.	
8	<b>AMISULPRIDE TABLETS 200MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Amisulpride	IP	200mg	
		Excipients		q.s.	
9	<b>LOSARTAN POTASSIUM TABLETS IP 50 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Losartan Potassium	IP	50 mg	
		Excipients		q.s.	
		Approved colour used			
10	<b>CLOZAPINE TABLETS IP 50 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clozapine	IP	50 mg	
		Excipients		q.s.	
11	<b>PROPRANOLOL HCl TABLETS 20MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Propranolol Hcl	IP	20mg	
		Excipients		q.s.	
12	<b>RAMIPRIL TABLETS IP 2.5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Ramipril	IP	2.5mg	
		Excipients		q.s.	
13	<b>RAMIPRIL TABLETS IP 5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Ramipril	IP	5mg	
		Excipients		q.s.	

14	<b>BETAHISTINE HYDROCHLORIDE TABLETS IP 8MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Betahistine Hydrochloride	IP	8 mg	
		Excipients		q.s.	
15	<b>CLOPIDOGREL TABLETS IP 75MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clopidogrel Bisulphate	IP		
		eq. to Clopidogrel		75 mg	
		Excipients		q.s.	
		Approved colour used			
16	<b>ESCITALOPRAM TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	IP		
		eq to Escitalopram		10mg	
		Excipients		qs	
		Approved colour used			
17	<b>MIRTAZAPINE MOUTH DISSOLVING TABLETS 15 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Mirtazapine	IP	15 mg	
		Excipients		q.s.	
18	<b>MIRTAZAPINE MOUTH DISSOLVING TABLETS 30 MG</b>	Each uncoated mouth dissolving tablets contains:			<b>APPROVED</b>
		Mirtazapine	IP	30mg	
		Excipients		q.s.	
19	<b>CLOBAZAM MOUTH DISSOLVING TABLETS 10MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Clobazam	IP	10 mg	
		Excipients		q.s.	
20	<b>CLOBAZAM MOUTH DISSOLVING TABLETS 20MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Clobazam	IP	20 mg	
		Excipients		q.s.	
21	<b>OLANZAPINE MOUTH DISSOLVING TABLETS 5 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	5 mg	
		Excipients		q.s.	
22	<b>OLANZAPINE MOUTH DISSOLVING TABLETS 10MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	10 mg	
		Excipients		q.s.	
23	<b>AMITRIPTYLINE HYDROCHLORIDE TABLETS 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Amitriptyline Hydrochloride	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
24	<b>AMITRIPTYLINE HYDROCHLORIDE TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Amitriptyline Hydrochloride	IP	10 mg	
		Excipients			
		Approved colour used		q.s.	
25	<b>AMITRIPTYLINE &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Chlordiazepoxide	IP	5mg	
		Amitriptyline Hydrochloride	IP		
		eq. to Amitriptyline		12.5mg	
		Excipients		q.s.	
		Approved colour used			

26	<b>AMITRIPTYLINE &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Chlordiazepoxide	IP	10mg	
		Amitriptyline Hydrochloride	IP		
		eq. to Amitriptyline		25mg	
		Excipients		q.s.	
		Approved colour used			
27	<b>DIAZEPAM TABLETS IP 2MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Diazepam	IP	2mg	
		Excipients		qs	
28	<b>SODIUM VAPROATE &amp; VALPROIC ACID TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sodium Valproate	IP	200 mg	
		Valproic Acid	IP	87 mg	
		(Both together corresponds to			
		Sodium Valproate IP 300 mg)			
		Excipients		q.s.	
29	<b>CLONAZEPAM TABLETS IP 0.5 MG</b>	Approved colour used			<b>APPROVED</b>
		Each uncoated tablet contains:			
		Clonazepam	IP	0.5mg	
		Excipients		q.s.	
30	<b>CLONAZEPAM TABLETS IP 1 MG</b>	Approved colour used			<b>APPROVED</b>
		Each uncoated tablet contains:			
		Clonazepam	IP	1mg	
		Excipients		q.s.	
31	<b>SERTRALINE HCL TABLETS IP 50MG</b>	Approved colour used			<b>APPROVED</b>
		Each film coated tablet contains:			
		Sertraline HCl	IP	50mg	
		Excipients		q.s.	
32	<b>QUETIAPINE TABLETS IP 100MG</b>	Approved colour used			<b>APPROVED</b>
		Each film coated tablet contains:			
		Quetiapine Fumatate	IP		
		eqv. to Quetiapine		100 mg	
		Excipients		q.s.	
33	<b>CLOPIDOGREL &amp;ASPIRIN TABLETS</b>	Approved colour used			<b>APPROVED</b>
		Each enteric coated tablet contains:			
		Clopidogrel Bisulphate	IP		
		eq. to Clopidogrel		75 mg	
		Aspirin (as enteric coated granules)	IP	75 mg	
		Excipients		q.s.	
34	<b>GLIMEPIRIDE &amp; METFORMIN HYDROCHLORIDE (SR) TABLETS</b>	Approved colour used			<b>APPROVED</b>
		Each uncoated bilayered tablet			
		contains:			
		Glimepiride	IP	1 mg	
		Metformin Hydrochloride	IP	500 mg	
		(as sustained release form)			
35	<b>RISPERIDONE &amp; TRIHENYPHENIDYL TABLETS</b>	Excipients		q.s.	<b>APPROVED</b>
		Each uncoated tablet contains:			
		Risperidone	BP	3 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
35	<b>RISPERIDONE &amp; TRIHENYPHENIDYL TABLETS</b>	Approved colour used			<b>APPROVED</b>

36	<b>CIPROFLOXACIN &amp;TINIDAZOLE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ciprofloxacin Hydrochloride	IP		
		eq. to Ciprofloxacin		500 mg	
		Tinidazole	IP	600mg	
		Excipients		q.s.	
		Approved colour used			
37	<b>AMOXAPINE TABLETS 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Amoxapine	USP	50 mg	
		Excipients		q.s.	
38	<b>IMIPRAMINE HCL TABLETS 75 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Imipramine Hcl	IP	75mg	
		Excipients		qs	
		Approved colour used			
39	<b>LEVETIRACETAM TABLETS 250 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	250mg	
		Excipients		q.s.	
		Approved colour used			
40	<b>LEVETIRACETAM TABLETS 500 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	500mg	
		Excipients		q.s.	
		Approved colour used			
41	<b>FLUNARIZINE TABLETS 10MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Flunarizine Dihydrochloride	B.P		
		eq. to Flunarizine		10 mg	
		Excipients		q.s.	
42	<b>PIRACETAM TABLETS 800 MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Piracetam	IP	800mg	
		Excipients		q.s.	
		Approved colour used			
43	<b>NITRAZEPAM TABLETS IP 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Nitrazepam	IP	5mg	
		Excipients		qs	
44	<b>NITRAZEPAM TABLETS IP 10 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Nitrazepam	IP	10mg	
		Excipients		qs	
45	<b>PAROXETINE CONTROLLED RELEASE TABLETS IP 12.5 MG</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Paroxetine Hcl	IP		
		eq. to Paroxetine		12.5mg	
		Excipients		q.s.	
		Approved colour used			
46	<b>PAROXETINE CONTROLLED RELEASE TABLETS IP 25 MG</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Paroxetine Hcl	IP		
		Eq. to Paroxetine		25mg	
		Excipients		q.s.	
		Approved colour used			
47	<b>CARBAMAZEPINE CONTROLLED RELEASE TABLETS IP 200 MG</b>	Each uncoated controlled-release tablet contains:			<b>APPROVED</b>
		Carbamazepine	IP	200 mg	
		Excipients		q.s.	
		Approved colour used			

48	<b>NORTRIPTYLINE TABLETS IP 25 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Nortriptyline Hcl	IP		
		eq. to Nortriptyline		25mg	
		Excipients		qs	
		Approved colour used			
49	<b>OLANZAPINE MOUTH DISSOLVING TABLETS 15 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	15 mg	
		Excipients		q.s.	
50	<b>OLANZAPINE MOUTH DISSOLVING TABLETS 7.5 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	7.5 mg	
		Excipients		q.s.	
51	<b>ESCITALOPRAM TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	IP		
		eq. to Escitalopram		20mg	
		Excipients		q.s.	
		Approved colour used			
52	<b>RISPERIDONE TABLETS 1MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	BP	1 mg	
		Excipients		q.s.	
53	<b>RISPERIDONE TABLETS 2MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	BP	2 mg	
		Excipients		q.s.	
54	<b>RISPERIDONE TABLETS 3MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	BP	3mg	
		Excipients		q.s.	
55	<b>RISPERIDONE TABLETS 4MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	BP	4mg	
		Excipients		q.s.	
56	<b>DIVALPROEX EXTENDED RELEASE TABLETS IP 500MG</b>	Each film coated extended release tablet contains:			<b>APPROVED</b>
		Divalproex Sodium	IP		
		eq. to Valproic Acid		500 mg	
		Excipients		q.s.	
		Approved colour used			
57	<b>AZITHROMYCIN TABLETS IP 500MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Azithromycin Dihydrate	IP		
		eq. to Azithromycin (Anhydrous)		500 mg	
		Excipients		q.s.	
		Approved colour used			
58	<b>TRIFLUOPERAZINE HCl &amp; TRIHEXYPHENIDYL HCl TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Trifluoperazine Hydrochloride	IP		
		eqv. to Trifluoperazine		5 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Approved colour used		q.s.	
59	<b>OLANZAPINE MOUTH DISSOLVING TABLETS 20 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	20 mg	
		Excipients		q.s.	
60	<b>PIRACETAM TABLETS 400 MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Piracetam	IP	400mg	
		Excipients		q.s.	
		Approved colour used			

61	<b>PIRACETAM TABLETS 1200 MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Piracetam	IP	1200mg	
		Excipients		q.s.	
		Approved colour used			
62	<b>HALOPERIDOL DISPERSIBLE TABLETS 5 MG</b>	Each uncoated dispersible tablet contains:			<b>APPROVED</b>
		Haloperidol	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
63	<b>DIVALPROEX EXTENDED RELEASE TABLETS IP 250MG</b>	Each film coated extended release tablet contains:			<b>APPROVED</b>
		Divalproex Sodium	IP		
		eq. to Valproic Acid		250 mg	
		Excipients		q.s.	
		Approved colour used			
64	<b>DOTHIEPIN HCL TABLETS 25 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Dothiepin Hcl	IP	25mg	
		Excipients		q.s.	
		Approved colour used			
65	<b>DOTHIEPIN HCL TABLETS 50 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Dothiepin Hcl	IP	50mg	
		Excipients		q.s.	
		Approved colour used			
66	<b>SODIUM VAPROATE &amp; VALPROIC ACID TABLETS</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Sodium Valproate	IP	133.5 mg	
		Valproic Acid	IP	58 mg	
		(Both together corresponds to Sodium Valproate IP 200 mg)			
		Excipients		q.s.	
		Approved colour used			
67	<b>CARBAMAZEPINE CONTROLLED RELEASE TABLETS 200 MG</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Carbamazepine	IP	200 mg	
		Excipients		q.s.	
		Approved colour used			
68	<b>CARBAMAZEPINE TABLETS 200 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Carbamazepine	IP	200 mg	
		Excipients		q.s.	
69	<b>CARBAMAZEPINE CONTROLLED RELEASE TABLETS 300 MG</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Carbamazepine	IP	300 mg	
		Excipients		q.s.	
		Approved colour used			
70	<b>CARBAMAZEPINE CONTROLLED RELEASE TABLETS 400 MG</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Carbamazepine	IP	400 mg	
		Excipients		q.s.	
		Approved colour used			
71	<b>RISPERIDONE &amp; TRIHXYPHENIDYL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	BP	4 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
		Approved colour used			

72	<b>RISPERIDONE &amp; TRIHEXYPHENIDYL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	BP	2 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
73	<b>OLANZAPINE MOUTH DISSOLVING TABLETS 2.5 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	2.5 mg	
		Excipients		q.s.	
74	<b>AZITHROMYCIN TABLETS IP 250MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Azithromycin Dihydrate	IP		
		eq. to Azithromycin (Anhydrous)		250 mg	
		Excipients		q.s.	
		Approved colour used			
75	<b>LITHIUM CARBONATE SUSTAINED RELEASE TABLETS IP 400MG</b>	Each uncoated sustained release tablet contains:			<b>APPROVED</b>
		Lithium Carbonate	IP	400mg	
		Excipients		q.s.	
76	<b>RABEPRAZOLE GASTRO RESISTANT TABLETS IP 20MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Rabeprazole Sodium	IP	20 mg	
		Excipients		q.s.	
		Approved colour used			
77	<b>AMITRIPTYLINE HYDROCHLORIDE TABLETS IP 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Amitriptyline Hydrochloride	IP	25 mg	
		Excipients		q.s.	
		Approved colour used			
78	<b>CLOBAZAM TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clobazam	IP	10 mg	
		Excipients		q.s.	
		Approved colour used			
79	<b>VENLAFAXINE EXTENDED RELEASE TABLETS 150 MG</b>	Each uncoated extended release tablet contains:			<b>APPROVED</b>
		venlafaxine Hcl	BP		
		eq to venlafaxine		150mg	
		Excipients		q.s.	
80	<b>BETAHISTINE HYDROCHLORIDE TABLETS IP 16MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Betahistine Hydrochloride	IP	16 mg	
		Excipients		q.s.	
		Approved colour used			
81	<b>ETORICOXIB TABLETS IP 90 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Etoricoxib	IP	90 mg	
		Excipients		q.s.	
		Approved colour used			
82	<b>PARACETAMOL TABLETS IP 650MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	IP	650 mg	
		Excipients		q.s.	
83	<b>LEVOFLOXACIN TABLETS IP 500MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levofloxacin Hemihydrate	IP		
		eq. to Levofloxacin		500 mg	
		Excipients		q.s.	
84	<b>QUETIAPINE TABLETS 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Quetiapine Fumate	IP		
		eqv. to Quetiapine		25 mg	
		Excipients		q.s.	
		Approved colour used			



85	<b>QUETIAPINE TABLETS 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Quetiapine Fumatate	IP		
		eqv. to Quetiapine		50 mg	
		Excipients		q.s.	
		Approved colour used			
86	<b>QUETIAPINE TABLETS 200 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Quetiapine Fumatate			
		eqv. to Quetiapine	IP	200 mg	
		Excipients		q.s.	
		Approved colour used			
87	<b>SERTRALINE HCL TABLETS IP 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sertraline Hcl	IP	100mg	
		Excipients		q.s.	
		Approved colour used			
88	<b>DULOXETINE GASTRO RESISTANT TABLETS IP 20MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Duloxetine Hcl	IP		
		eq. to Duloxetine		20mg	
		Excipients		q.s.	
		Approved colour used			
89	<b>DULOXETINE GASTRO RESISTANT TABLETS IP 30MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Duloxetine Hcl	IP		
		eq. to Duloxetine		30mg	
		Excipients		q.s.	
		Approved colour used			
90	<b>DULOXETINE GASTRO RESISTANT TABLETS IP 60MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Duloxetine Hcl	IP		
		eq. to Duloxetine		60mg	
		Excipients		q.s.	
		Approved colour used			
91	<b>GLIMEPIRIDE &amp; METFORMIN HYDROCHLORIDE (SR) TABLETS</b>	Each uncoated bilayered tablet contains			<b>APPROVED</b>
		Glimepiride	IP	2 mg	
		Metformin Hydrochloride (as sustained release form)	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			
92	<b>SERTRALINE HCL &amp; ALPRAZOLAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sertraline Hydrochloride	IP	50mg	
		Alprazolam	IP	0.5mg	
		Excipients		q.s.	
		Approved colour used			
93	<b>ALPRAZOLAM TABLETS IP 0.5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	0.5 mg	
		Excipients		q.s.	
		Approved colour used			
94	<b>LEVOFLOXACIN TABLETS IP 250MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levofloxacin Hemihydrate	IP		
		eq. to Levofloxacin		250 mg	
		Excipients		q.s.	
		Approved colour used			
95	<b>LEVOFLOXACIN TABLETS IP 750MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levofloxacin Hemihydrate	IP		
		eq. to Levofloxacin		750 mg	
		Excipients		q.s.	
		Approved colour used			



96	ZIPRASIDONE CAPSULES 20 MG	Each hard gelatin capsule contains:			APPROVED
		Ziprasidone HCl Monohydrate	IP		
		eq. to Ziprasidone		20mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shell.			
97	ZIPRASIDONE CAPSULES 40 MG	Each hard gelatin capsule contains:			APPROVED
		Ziprasidone Hcl Monohydrate	IP		
		eq. to Ziprosidone		40mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shell.			
98	GLIMEPIRIDE TABLETS IP 1MG	Each uncoated tablet contains			APPROVED
		Glimepiride	IP	1mg	
		Excipients		q.s.	
99	GLIMEPIRIDE TABLETS IP 2MG	Each uncoated tablet contains			APPROVED
		Glimepiride	IP	2mg	
		Excipients		q.s.	
100	PANTOPRAZOLE & DOMPERIDONE TABLETS	Each enteric coated tablet contains:			APPROVED
		Pantoprazole Sodium Sesquihydrate	IP		
		eq. to Pantoprazole		40 mg	
		Domperidone	IP	10mg	
		Excipients		q.s.	
		Approved colour used			
101	PANTOPRAZOLE GASTRO RESISTANT TABLETS IP 40MG	Each enteric coated tablet contains:			APPROVED
		Pantoprazole Sodium Sesquihydrate	IP		
		eq. to Pantoprazole		40 mg	
		Excipients		q.s.	
		Approved colour used			
102	THEOPHYLLINE SUSTAINED-RELEASE TABLETS 200 MG	Each uncoated sustained-release tablet contains:			APPROVED
		Theophylline (anhydrous)	IP	200 mg	
		Excipients		q.s.	
103	QUETIAPINE SUSTAINED RELEASE TABLETS 100MG	Each film coated sustained release tablet contains:			APPROVED
		Quetiapine Fumatate	IP		
		eq. to Quetiapine		100 mg	
		Excipients		q.s.	
		Approved colour used			
104	AMITRIPTYLINE HYDROCHLORIDE TABLETS 75MG	Each film coated tablet contains:			APPROVED
		Amitriptyline Hydrochloride	IP	75 mg	
		Excipients		q.s.	
		Approved colour used			
105	DICLOFENAC SODIUM , PARACETAMOL & CHLORZOXAZONE TABLETS	Each uncoated tablet contains:			APPROVED
		Diclofenac Sodium	IP	50 mg	
		Paracetamol	IP	325 mg	
		Chlorzoxazone	USP	500mg	
		Excipients		q.s.	
		Approved colour used			
106	CHLORPROMAZINE HYDROCHLORIDE TABLETS IP 100MG	Each uncoated tablet contains:			APPROVED
		Chlorpromazine Hydrochloride	IP	100mg	
		Excipients		q.s.	
		Approved colour used			

107	<b>LUTEIN , LYCOPENE &amp; ASTAXANTHIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Natural carotenoids rich in			
		Lutein/Zeaxanthin		3.2mg	
		(containing not less than 3 mg of			
		Lutein/Zeaxanthin)			
		Lycopene	USP	2mg	
		Astaxanthin		2mg	
		Excipients		q.s.	
		Approved colour used			
108	<b>GLUCOSAMINE HCL &amp; CHONDROITIN SULPHATE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Glucosamine Hydrochloride	IP	750 mg	
		Chondroitin Sulphate		400 mg	
		Excipients		q.s.	
		Approved colour used			
109	<b>TOPIRAMATE TABLETS IP 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Topiramate	IP	50 mg	
		Excipients		q.s.	
		Approved colour used			
110	<b>RISPERIDONE &amp; TRIHEXYPHENIDYL MOUTH DISSOLVING TABLETS</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Risperidone	BP	3 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
		Approved colour used			
111	<b>ALPRAZOLAM TABLETS IP 0.25 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	0.25 mg	
		Excipients		q.s.	
		Approved colour used			
112	<b>CLOMIPRAMINE HYDROCHLORIDE EXTENDED RELEASE TABLETS 75MG</b>	Each film coated extended release tablet contains:			<b>APPROVED</b>
		Chlomipramine Hydrochloride	IP	75mg	
		Excipients		q.s.	
		Approved colour used			
113	<b>ATORVASTATIN TABLETS 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Atorvastatin Calcium	IP		
		eq. to Atorvastatin		5mg	
		Excipients		q.s.	
		Approved colour used			
114	<b>AMISULPRIDE TABLETS IP 100MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Amisulpride	IP	100mg	
		Excipients		q.s.	
115	<b>TOLPERISONE HCL &amp; DICLOFENAC SODIUM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tolperisone Hcl		150mg	
		Diclofenac Sodium	IP	50mg	
		Excipients		qs	
		Approved colour used			
116	<b>NAPROXEN TABLETS IP 250MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Naproxen	IP	250mg	
		Excipients		q.s.	
		Approved colour used			
117	<b>RIZATRIPTAN TABLETS IP 10MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Rizatriptan Benzoate	IP		
		equivalent to Rizatriptan		10mg	
		Excipients		q.s.	
		Approved colour used			

118	<b>RAMIPRIL&amp; HYDROCHLOROTHIAZIDE TABLETS IP</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Ramipril	IP	10mg	
		Hydrochlorothiazide	IP	12.5mg	
		Excipients		q.s.	
		Approved colour used			
119	<b>CINNARIZINE TABLETS IP 25MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Cinnarizine	IP	25mg	
		Excipients		q.s.	
120	<b>PERINDOPRIL ERBUMINE &amp; AMLODIPINE BESYLATE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Perindopril Erbumine	BP	4mg	
		Amlodipine Besylate	IP		
		Eq. to Amlodipine		5mg	
		Excipients		q.s.	
		Approved colour used			
121	<b>FEXOFENADINE HCL AND MONTELUKAST TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fexofenadine Hydrochloride	IP	120 mg	
		Montelukast Sodium	IP		
		eqv. to Montelukast		10 mg	
		Excipients		q.s.	
		Approved colour used			
122	<b>ARIPIRAZOLE TABLETS IP 10 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aripiprazole	IP	10 mg	
		Excipients		q.s.	
		Colour: Approved colour used			
123	<b>ARIPIRAZOLE TABLETS IP 15 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aripiprazole	IP	15 mg	
		Excipients		q.s.	
		Colour: Approved colour used			
124	<b>SILDENAFIL TABLETS IP 100MG</b>	Each Film coated Tablet contains:			<b>APPROVED</b>
		Sildenafil Citrate	IP		
		eq. to Sildenafil		100 mg	
		Excipients		qs	
		Approved colour used			
125	<b>ESCITALOPRAM &amp;CLONAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	IP		
		eq to Escitalopram		10mg	
		Clonazepam	IP	0.25mg	
		Excipients		qs	
		colour:Approved colour used			
126	<b>PENFLURIDOL TABLETS</b>	Each film coated tablets contains:			<b>APPROVED</b>
		Penfluridol		20mg	
		Excipients		qs	
		Approved colour used			
127	<b>SUMATRIPTAN &amp; NAPROXEN TABLETS</b>	Each film coated tablets contains:			<b>APPROVED</b>
		Sumatriptan Succinate	IP		
		eq. to Sumatriptan		50mg	
		Naproxen	IP	275mg	
		Excipients		qs	
		Approved colour used			
128	<b>RISPERIDONE &amp; TRIHXYPHENIDYL HCL TABLETS</b>	Each film coated tablets contains:			<b>APPROVED</b>
		Risperidone	BP	2 mg	
		Trihexyphenidyl Hcl	IP	2 mg	
		Excipients		qs	
		Approved colour used			

129	<b>LAMOTRIGINE DISPERSIBLE TABLETS IP 25MG</b>	Each uncoated dispersible tablet			<b>APPROVED</b>
		contains:			
		Lamotrigine	IP	25 mg	
		Excipients		q.s.	
130	<b>LAMOTRIGINE DISPERSIBLE TABLETS IP 50MG</b>	Each uncoated dispersible tablet			<b>APPROVED</b>
		contains:			
		Lamotrigine	IP	50 mg	
		Excipients		q.s.	
131	<b>LAMOTRIGINE DISPERSIBLE TABLETS IP 100MG</b>	Each uncoated dispersible tablet			<b>APPROVED</b>
		contains:			
		Lamotrigine	IP	100 mg	
		Excipients		q.s.	
132	<b>LAMOTRIGINE TABLETS 25MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lamotrigine	IP	25 mg	
		Excipients		q.s.	
133	<b>LAMOTRIGINE TABLETS 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lamotrigine	IP	50 mg	
		Excipients		q.s.	
134	<b>LAMOTRIGINE TABLETS 100MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lamotrigine	IP	100 mg	
		Excipients		q.s.	
135	<b>TELMISARTAN &amp; HYDROCHLOROTHIAZIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	80 mg	
		Hydrochlorothiazide	IP	12.5 mg	
		Excipients		q.s.	
		Approved colour used			
136	<b>RIFAXIMIN TABLETS 550MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rifaximin	BP	550mg	
		Excipients		q.s.	
		Approved colour used			
137	<b>PROPRANOLOL &amp; ALPRAZOLAM TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Propranolol Hcl	IP	20 mg	
		Alprazolam	IP	0.25 mg	
		Excipients		q.s.	
138	<b>DOXYLAMINE SUCCINATE, PYRIDOXINE HCL &amp; FOLIC ACID TABLETS</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Doxylamine Succinate	BP	20mg	
		Pyridoxine Hydrochloride	IP	20mg	
		Folic Acid	IP	5mg	
		Excipients		q.s.	
		Approved colour used			
139	<b>MODAFINIL TABLETS 200MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Modafinil	BP	200mg	
		Excipients		q.s.	
140	<b>LORAZEPAM MOUTH DISSOLVING TABLETS 1MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Lorazepam	IP	1mg	
		Excipients		q.s.	
141	<b>LORAZEPAM MOUTH DISSOLVING TABLETS 2MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Lorazepam	IP	2mg	
		Excipients		q.s.	

142	<b>BLONANSERIN TABLETS 2MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Blonanserin		2mg	
		Excipients		q.s.	
143	<b>BLONANSERIN TABLETS 4MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Blonanserin		4mg	
		Excipients		q.s.	
144	<b>BLONANSERIN TABLETS 8MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Blonanserin		8mg	
		Excipients		q.s.	
145	<b>GLUCOSAMINE HCL , CHOLECALCEFEROL , TOCOPHEROL ACETATE, ASCORBIC ACID , BORON, SELENIUM , ZINC &amp; MANGANESE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Glucosamine HCl	USP	750mg	
		Cholecalciferol	IP	200IU	
		Tocopherol Acetate	IP	12.5mg	
		Ascorbic Acid	IP	50mg	
		Sodium Borate	BP		
		eq to Boron		0.5mg	
		Sodium Selenite	USP		
		eq to Selenium		70mcg	
		Zinc Sulphate Monohydrate	IP		
		eq to Zinc		3mg	
		Manganese Sulphate	BP		
		eq to Manganese		3mg	
		Excipients		q.s.	
		Approved colour used			
146	<b>EPERISONE HYDROCHLORIDE (SR) &amp; DICLOFENAC SODIUM (SR) CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Eperisone Hydrochloride		150 mg	
		(As Sustained Release Pellets)			
		Diclofenac Sodium	IP	100 mg	
		(As Sustained Release Pellets)			
		Excipients		q.s.	
		Approved colour used in empty Capsule shell			
147	<b>DICLOFENAC POTASSIUM, PARACETAMOL&amp; SERRATIOPEPTIDASE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Diclofenac Potassium	BP	50 mg	
		Paracetamol	IP	325 mg	
		Serratiopeptidase	IP	10mg	
		Excipients		q.s.	
		Approved colour used			
148	<b>CITICOLINE &amp; PIRACETAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Citicoline Sodium	IP		
		eq. to Citicoline		500mg	
		Piracetam	IP	800mg	
		Excipients		q.s.	
		Approved colour used			
149	<b>ESCITALOPRAM &amp;CLONAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	IP		
		eq to Escitalopram		5mg	
		Clonazepam	IP	0.25mg	
		Excipients		qs	
		Approved colour used			
150	<b>PREGABALIN CAPSULES IP 300 MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Pregabalin	IP	300 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			

151	<b>MONTELUKAST SODIUM TABLETS IP 10MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Montelukast Sodium	IP		
		eq to Montelukast		10mg	
		Excipients		q.s.	
		Approved colour used			
152	<b>CINNARIZINE &amp; DIMENHYDRINATE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Cinnarizine	IP	20mg	
		Dimenhydrinate	USP	40mg	
		Excipients		q.s.	
153	<b>CINNARIZINE &amp;DOMPERIDONE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Cinnarizine	IP	20mg	
		Domperidone Maleate	IP		
		eq. to Domperidone		15mg	
		Excipients		q.s.	
154	<b>DOMPERIDONE MALEATE &amp; NAPROXEN SODIUM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Domperidone Maleate	IP		
		eq. to Domperidone		10mg	
		Naproxen Sodium	IP	500mg	
		Excipients		q.s.	
		Approved colour used			
155	<b>LORAZEPAM TABLETS 0.5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lorazepam	IP	0.5mg	
		Excipients		q.s.	
156	<b>LORAZEPAM TABLETS 3MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lorazepam	IP	3mg	
		Excipients		q.s.	
157	<b>PREGABALIN AND NORTRIPTYLINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pregabalin	IP	75 mg	
		Nortriptyline Hydrochloride	IP		
		eq. to Nortriptyline		10 mg	
		Excipients		q.s.	
		Approved colour used			
158	<b>PROPRANOLOL HCL &amp; FLUNARIZINE CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Propranolol HCl	I.P	40 mg	
		(As Sustained Release Pellets)			
		Flunarizine Dihydrochloride	B.P		
		eq. to Flunarizine		5 mg	
		Excipients		q.s.	
		Approved colour use in empty			
		Capsule shell			
159	<b>OLANZAPINE TABLETS IP 2.5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	2.5 mg	
		Excipients		q.s.	
		Approved colour used			
160	<b>OLANZAPINE TABLETS IP 5.0MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	5.0 mg	
		Excipients		q.s.	
		Approved colour used			
161	<b>OLANZAPINE TABLETS 15MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	15 mg	
		Excipients		q.s.	
		Approved colour used			

162	HYDROCHLOROTHIAZIDE, AMLODIPINE & TELMISARTAN TABLETS	Each film coated tablet contains:			APPROVED
		Hydrochlorothiazide	IP	12.5 mg	
		Amlodipine Besilate	IP		
		eq. to Amlodipine		5 mg	
		Telmisartan	IP	40 mg	
		Excipients		q.s.	
		Approved colour used			
163	CHLORPROMAZINE HCl & TRIHEXYPHENIDYL HCl TABLETS	Each film coated tablet contains:			APPROVED
		Chlorpromazine Hydrochloride	IP	200 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
164	ACETAZOLAMIDE TABLETS IP 250MG	Each uncoated tablet contains:			APPROVED
		Acetazolamide	IP	250 mg	
		Excipients		q.s.	
165	LITHIUM CARBONATE PROLONGED RELEASE TABLETS IP 300MG	Each uncoated prolonged release tablet contains:			APPROVED
		Lithium Carbonate	IP	300mg	
		Excipients		q.s.	
166	LITHIUM CARBONATE EXTENDED RELEASE TABLETS 600MG	Each uncoated extended release tablet contains:			APPROVED
		Lithium Carbonate	IP	600 mg	
		Excipients		q.s.	
167	ENALAPRIL MALEATE & HYDROCHLOROTHIAZIDE TABLETS	Each uncoated tablet contains:			APPROVED
		Enalapril Maleate	IP	10 mg	
		Hydrochlorothiazide	IP	25 mg	
		Excipients		q.s.	
		Approved colour used			
168	METOPROLOL SUCCINATE (E.R) & TELMISARTAN TABLETS	Each film coated tablet contains:			APPROVED
		Metoprolol Succinate	IP	47.5mg	
		eq. to Metoprolol Tartrate		50 mg	
		(Exetended Release)			
		Telmisartan	IP	40 mg	
		Excipients		q.s.	
		Approved colour used			
169	FLUOXETINE HYDROCHLORIDE TABLETS IP 40MG	Each film coated tablet contains:			APPROVED
		Fluoxetine Hydrochloride	IP		
		eqv. to Fluoxetine		40mg	
		Excipients		q.s.	
		Approved colour used			
170	GABAPENTIN TABLETS IP 100MG	Each film coated tablet contains:			APPROVED
		Gabapentin	IP	100 mg	
		Excipients		q.s.	
		Approved colour used			
171	DOMPERIDONE MALEATE & NAPROXEN SODIUM TABLETS	Each film coated tablet contains:			APPROVED
		Domperidone Maleate	IP		
		eq. to Domperidone		10 mg	
		Naproxen	IP	250mg	
		Excipients		q.s.	
172	RISPERIDONE TABLETS 2MG	Each film coated tablet contains:			APPROVED
		Risperidone	BP	2 mg	
		Excipients		q.s.	
		Approved colour used			



173	<b>SODIUM PICOSULPHATE TABLETS 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sodium Picosulphate	BP	5mg	
		Excipients		q.s.	
		Approved colour used			
174	<b>SODIUM PICOSULPHATE TABLETS 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sodium Picosulphate	BP	10mg	
		Excipients		q.s.	
		Approved colour used			
175	<b>GLIMEPIRIDE, METFORMIN HYDROCHLORIDE &amp; VOGLIBOSE TABLETS</b>	Each uncoated bilayered tablet contains:			<b>APPROVED</b>
		Glimepiride	IP	1mg	
		Metformin Hydrochloride (as sustained release form)	IP	500 mg	
		Voglibose	IP	0.2 mg	
		Excipients		q.s.	
		Approved colour used			
176	<b>GLIMEPIRIDE, METFORMIN HYDROCHLORIDE &amp; VOGLIBOSE TABLETS</b>	Each uncoated bilayered tablet contains:			<b>APPROVED</b>
		Glimepiride	IP	2mg	
		Metformin Hydrochloride (as sustained release form)	IP	500 mg	
		Voglibose	IP	0.2 mg	
		Excipients		q.s.	
		Approved colour used			
177	<b>CHLORPROMAZINE HYDROCHLORIDE TABLETS IP 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Chlorpromazine Hydrochloride	IP	50mg	
		Excipients		q.s.	
		Approved colour used			
178	<b>AMISULPRIDE TABLETS IP 300MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Amisulpride	IP	300mg	
		Excipients		q.s.	
179	<b>AMISULPRIDE TABLETS IP 400MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Amisulpride	IP	400mg	
		Excipients		q.s.	
180	<b>EPERISONE &amp; PARACETAMOL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Eperisone Hydrochloride		50 mg	
		Paracetamol	IP	325 mg	
		Excipients		q.s.	
181	<b>ESCITALOPRAM &amp; CLONAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	IP		
		eq. to Escitalopram		5 mg	
		Clonazepam	IP	0.5mg	
		Excipients		qs	
		Approved colour used			
182	<b>LITHIUM CARBONATE TABLETS IP 300MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lithium Carbonate	IP	300mg	
		Excipients		q.s.	
183	<b>CLOMIPRAMINE HYDROCHLORIDE TABLETS 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		tablet contains:			
		Chlomipramine Hydrochloride	IP	50mg	
		Excipients		q.s.	
		Approved colour used			

184	<b>SERTRALINE HCL TABLETS IP 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sertraline Hydrochloride	IP	25mg	
		Excipients		q.s.	
		Approved colour used			
185	<b>DULOXETINE GASTRO RESISTANT TABLETS IP 40MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Duloxetine Hydrochloride	IP		
		eq to Duloxetine		40mg	
		Excipients		q.s.	
		Approved colour used			
186	<b>ZOLPIDEM TARTRATE TABLETS IP 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Zolpidem Tartrate	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
187	<b>ZOLPIDEM TARTRATE TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Zolpidem Tartrate	IP	10 mg	
		Excipients		q.s.	
		Approved colour used			
188	<b>PAROXETINE CONTROLLED RELEASE TABLETS 37.5 MG</b>	Each film coated controlled release tablet contains:			<b>APPROVED</b>
		Paroxetine Hydrochloride	IP		
		eq. to Paroxetine		37.5mg	
		Excipients		q.s.	
		Approved colour used			
189	<b>LEVETIRACETAM TABLETS 750 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	750mg	
		Excipients		q.s.	
		Approved colour used			
190	<b>TRANEXMIC ACID , ETAMSYLATE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Tranexmic Acid	IP	250 mg	
		Etamsylate	BP	250 mg	
		Excipients		q.s.	
191	<b>CALCIUM CARBONATE &amp; ALFACALCIDOL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium carbonate	IP		
		eq. to elemental calcium		500mg	
		Alfacalcidol	IP	0.25mcg	
		Approved colour used			
192	<b>AMISULPRIDE TABLETS 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Amisulpride	IP	50mg	
		Excipients		q.s.	
193	<b>TRIFLUOPERAZINE HCl &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Trifluoperazine Hydrochloride	IP		
		eq. to Trifluoperazine		10 mg	
		Chlordiazepoxide	IP	1 mg	
		Approved colour used		q.s.	
194	<b>CAMYLOFIN HCL &amp; NIMESULIDE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Camylofin Dihydrochloride		50mg	
		Nimesulide	BP	100 mg	
		Excipients		qs	
		Approved Colour Used			
195	<b>LOSARTAN POTASSIUM TABLETS 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Losartan Potassium	IP	25 mg	
		Excipients		q.s.	
		Approved colour used			

196	<b>TELMISARTAN &amp; HYDROCHLOROTHIAZIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	40 mg	
		Hydrochlorothiazide	IP	12.5mg	
		Excipients		q.s.	
		Approved colour used			
197	<b>CIPROFLOXACIN TABLETS IP 250 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ciprofloxacin Hydrochloride	IP		
		eq. to Ciprofloxacin		250 mg	
		Excipients		q.s.	
		Approved colour used			
198	<b>PANTOPRAZOLE (EC) &amp; DOMPERIDONE (SR) CAPSULES</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Pantoprazole Sodium Sesquihydrate	IP		
		eq. to Pantoprazole		40 mg	
		(as enteric coated pellets)			
		Domperidone	IP	30mg	
		(as sustained release pellets)			
		Excipients		q.s.	
		Approved colour used in empty capsule shells			
199	<b>MIRTAZAPINE MOUTH DISSOLVING TABLETS 7.5 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Mirtazapine	IP	7.5 mg	
		Excipients		q.s.	
		Approved colour used			
200	<b>FLUPENTIXOL HYDROCHLORIDE TABLETS 0.5 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Flupentixol Hydrochloride	BP		
		eq. to Flupentixol		0.5 mg	
		Excipients		q.s.	
		Approved colour used			
201	<b>LORAZEPAM TABLETS IP 1MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lorazepam	IP	1mg	
		Excipients		q.s.	
		Approved colour used			
202	<b>MIRTAZAPINE MOUTH DISSOLVING TABLETS 45 MG</b>	Each uncoated mouth dissolving tablets contains:			<b>APPROVED</b>
		Mirtazapine	IP	45mg	
		Excipients		q.s.	
203	<b>LORAZEPAM TABLETS IP 2MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Lorazepam	IP	2mg	
		Excipients		q.s.	
204	<b>CHLORPHENIRAMINE MALEATE TABLETS 4 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Chlorpheniramine Maleate	IP	4 mg	
		Excipients		q.s.	
		Approved colour used			
205	<b>DICLOFENAC SODIUM &amp; PARACETAMOL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Diclofenac Sodium	IP	50 mg	
		Paracetamol	IP	325 mg	
		Excipients		q.s.	
		Approved colour used			
206	<b>NIMESULIDE TABLETS 100MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Nimesulide	BP	100 mg	
		Excipients		q.s.	

207	<b>NIMESULIDE&amp; PARACETAMOL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Nimesulide	BP	100mg	
		Paracetamol	IP	325mg	
		Excipients		q.s.	
		Approved colour used			
208	<b>CETIRIZINE HYDROCHLORIDE TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Cetirizine Hydrochloride	IP	10 mg	
		Excipients		q.s.	
		Approved colour used			
209	<b>PARACETAMOL TABLETS IP 500MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	IP	500mg	
		Excipients		q.s.	
210	<b>ONDANSETRON MOUTH DISSOLVING TABLETS IP 4MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Ondansetron Hydrochloride	IP		
		eq. to Ondansetron		4 mg	
		Excipients		q.s.	
		Approved colour used			
211	<b>NIACINAMIDE,VIT A, CHOLECALCIFEROL, VIT.E, VIT B1, RIBOFLAVIN, PYRIDOXINE HCL , LECITHIN &amp;VIT D3 GRANULES</b>	Each gram contains			<b>APPROVED</b>
		Vitamin A	IP	400 IU	
		Cholecalciferol	IP	50 IU	
		Vitamin E Acetate	IP	5 mg	
		Lecithin	IP	800 mg	
		Vitamin B1	IP	2 mg	
		Riboflavin	IP	2 mg	
		Pyridoxine HCl	IP	1 mg	
		Niacinamide	IP	5 mg	
		Excipients		q.s.	
212	<b>IBUPROFEN TABLETS IP 200MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ibuprofen	IP	200mg	
		Excipients		q.s.	
		Approved colour used			
213	<b>IBUPROFEN TABLETS IP 400MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ibuprofen	IP	400mg	
		Excipients		q.s.	
		Approved colour used			
214	<b>LITHIUM CARBONATE SUSTAINED RELEASE TABLETS IP 450MG</b>	Each uncoated sustained release tablet contains:			<b>APPROVED</b>
		Lithium Carbonate	IP	450mg	
		Excipients		q.s.	
215	<b>CLOMIFENE TABLETS IP 50 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clomifene Citrate	IP	50 mg	
		Excipients		q.s.	
216	<b>LEVOCETIRIZINE DIHYDROCHLORIDE TABLETS IP 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levocetirizine Dihydrochloride	IP	5mg	
		Excipients		q.s.	
		Approved colour used			
217	<b>ATORVASTATIN TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Atorvastatin Calcium	IP		
		eq. to Atorvastatin		10mg	
		Excipients		q.s.	
		Approved colour used			

218	ATORVASTATIN TABLETS IP 20MG	Each film coated contains:			APPROVED
		Atorvastatin Calcium	IP		
		eq. to Atorvastatin		20mg	
		Excipients		q.s.	
		Approved colour used			
219	SERRATIOPEPTIDASE TABLETS IP 10MG	Each enteric coated tablet contains:			APPROVED
		Serratiopeptidase	IP	10 mg	
		(As enteric coated granules 20000 Units)			
		Excipients		q.s.	
		Approved colour used			
220	LOSARTAN POTASSIUM & HYDROCHLOROTHIAZIDE TABLETS	Each film coated tablet contains:			APPROVED
		Losartan Potassium	IP	50 mg	
		Hydrochlorothiazide	IP	12.5mg	
		Excipients		q.s.	
		Approved colour used			
221	PRAMIPEXOLE TABLETS 0.5MG	Each uncoated tablet contains:			APPROVED
		Pramipexole Dihydrochloride	USP	0.5mg	
		Excipients		q.s.	
222	PRAMIPEXOLE TABLETS 0.25MG	Each uncoated tablet contains:			APPROVED
		Pramipexole Dihydrochloride	USP	0.25mg	
		Excipients		q.s.	
223	PIROXICAM TABLETS 20MG	Each uncoated tablets contains:			APPROVED
		Piroxicam	IP	20mg	
		Excipients		q.s.	
224	ACECLOFENAC & PARACETAMOL TABLETS	Each film coated tablet contains:			APPROVED
		Aceclofenac	IP	100 mg	
		Paracetamol	IP	325 mg	
		Excipients		q.s.	
		Approved colour used			
225	CLOZAPINE TABLETS IP 25 MG	Each uncoated tablet contains:			APPROVED
		Clozapine	IP	25mg	
		Excipients		q.s.	
226	OXCARBAZEPINE TABLETS IP 300MG	Each film coated tablet contains:			APPROVED
		Oxcarbazepine	IP	300 mg	
		Excipients		q.s.	
		Approved colour used			
227	FLUOXETINE TABLETS IP 20MG	Each film coated tablet contains:			APPROVED
		Fluoxetine Hydrochloride	IP		
		eq. to fluoxetine		20mg	
		Excipients		q.s.	
		Approved colour used			
228	PYRAZINAMIDE TABLETS IP 500 MG	Each uncoated tablet contains:			APPROVED
		Pyrazinamide	IP	500mg	
		Excipients		qs	
229	PYRAZINAMIDE TABLETS IP 750 MG	Each uncoated tablet contains:			APPROVED
		Pyrazinamide	IP	750mg	
		Excipients		qs	
230	BACLOFEN TABLETS IP 10MG	Each uncoated tablet contains:			APPROVED
		Baclofen	IP	10 mg	
		Excipients		q.s.	
231	CHLORDIAZEPOXIDE TABLETS IP 25MG	Each film coated tablet contains:			APPROVED
		Chlordiazepoxide	IP	25 mg	
		Excipients		q.s.	
		Approved colour used			

232	<b>CHLORDIAZEPOXIDE TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Chlordiazepoxide	IP	10 mg	
		Excipients		q.s.	
		Approved colour used			
233	<b>CLOMIFENE CITRATE TABLETS 100 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clomifene citrate	IP	100mg	
		Excipients		q.s.	
234	<b>CLOMIFENE CITRATE TABLETS IP 25 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clomifene citrate	IP	25mg	
		Excipients		q.s.	
235	<b>QUETIAPINE SUSTAIN RELEASE TABLETS 300MG</b>	Each film coated sustain release tablet contains:			<b>APPROVED</b>
		Quetiapine fumarate	IP		
		eq. to Quetiapine		300mg	
		Excipients		q.s.	
		Approved colour used			
236	<b>QUETIAPINE SUSTAIN RELEASE TABLETS 400MG</b>	Each film coated sustain release tablet contains:			<b>APPROVED</b>
		Quetiapine fumarate	IP		
		eq. to Quetiapine		400mg	
		Excipients		q.s.	
		Approved colour used			
237	<b>OLMESARTAN MEDOXOMIL TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olmесartan Medoxomil	IP	20mg	
		Excipients		q.s.	
		Colour: Titanium Dioxide			
238	<b>METFORMIN HYDROCHLORIDE SUSTAIN RELEASE TABLETS 1000 MG</b>	Each uncoated sustain release tablet contains:			<b>APPROVED</b>
		Metformine Hydrochloride	IP	1000 mg	
		Excipients		q.s.	
239	<b>DEFLAZACORT TABLET 6 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Deflazacort		6mg	
		Excipients		qs	
240	<b>DIAZEPAM TABLETS IP 5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Diazepam	IP	5mg	
		Excipients		qs	
241	<b>IMIPRAMINE HCL TABLETS IP 25 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Imipramine Hcl	IP	25mg	
		Excipients		qs	
		Approved colour used			
242	<b>VOGLIBOSE TABLETS IP 0.2 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Voglibose		0.2 mg	
		Excipients		q.s.	
243	<b>VOGLIBOSE TABLETS IP 0.3 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Voglibose		0.3 mg	
		Excipients		q.s.	
244	<b>OFLOXACIN TABLETS IP 200 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ofloxacin	IP	200mg	
		Excipients		qs	
		Approved colour used			
245	<b>OFLOXACIN TABLETS IP 400 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ofloxacin	IP	400mg	
		Excipients		qs	
		Approved colour used			

246	<b>DICLOFENAC POTASSIUM &amp;SERRATIOPEPTIDASE TABLETS</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Diclofenac Potassium	BP	50mg	
		Serratiopeptidase (20,000 unit )	IP	10mg	
		Excipients		q.s.	
		Approved colour used			
247	<b>TOPIRAMATE TABLETS IP 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Topiramate	IP	25 mg	
		Excipients		q.s.	
		Approved colour used			
248	<b>DONEPEZIL HCL TABLETS IP 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Donepezil Hydrochloride	IP	5mg	
		Excipients		q.s.	
		Approved colour used			
249	<b>DONEPEZIL HCL &amp; MEMANTINE HCL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Donepezil Hydrochloride	IP	5mg	
		Memantine Hydrochloride	USP	5mg	
		Excipients		q.s.	
		Approved colour used			
250	<b>QUETIAPINE SUSTAINED RELEASE TABLETS 200MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Quetiapine Fumatate	IP		
		eq. to Quetiapine		200 mg	
		Excipients		q.s.	
		Approved colour used			
251	<b>BROMOCRIPTINE TABLETS IP 2.5MG</b>	Each uncoated tablets contains:			<b>APPROVED</b>
		Bromocriptine	IP	2.5mg	
		Excipients		q.s.	
252	<b>BROMOCRIPTINE TABLETS 1.25MG</b>	Each uncoated tablets contains:			<b>APPROVED</b>
		Bromocriptine	IP	1.25mg	
		Excipients		q.s.	
253	<b>GLIMEPIRIDE TABLETS 4MG</b>	Each uncoated tablet contains			<b>APPROVED</b>
		Glimepiride	IP	4mg	
		Excipients		q.s.	
254	<b>OXCARBAZEPINE TABLETS IP 600MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Oxcarbazepine	IP	600 mg	
		Excipients		q.s.	
255	<b>OLANZAPINE &amp;FLUOXETINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	5 mg	
		Fluoxetine Hcl	IP		
		eq. to fluoxetine		20mg	
		Excipients		q.s.	
		Approved colour used			
256	<b>HYDROCHLOROTHIAZIDE &amp; OLMESARTAN MEDOXOMIL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Hydrochlorothiazide	IP	12.5 mg	
		Olmesartan Medoxomil	IP	40mg	
		Excipients		q.s.	
		Approved colour used			
257	<b>LEVOSULPIRIDE TABLETS 25 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levosulpiride		25 mg	
		Excipients		q.s.	
		Approved colour used			



258	TRANEXAMIC ACID TABLETS IP 500MG	Each uncoated tablet contains:			APPROVED
		Tranexamic Acid	IP	500mg	
		Excipients		q.s.	
259	ROXITHROMYCIN TABLETS IP 150MG	Each film coated tablet contains:			APPROVED
		Roxithromycin	IP	150mg	
		Excipients		q.s.	
260	NORFLOXACIN TABLETS IP 400 MG	Approved colour used			APPROVED
		Each film coated tablet contains:			
		Norfloxacin	IP	400mg	
261	SODIUM VALPROATE TABLETS IP 200MG	Excipients			APPROVED
		Approved colour used			
		Each film coated tablet contains:			
262	MEMANTINE HCL TABLETS 10 MG	Sodium Valproate	IP	200 mg	APPROVED
		Excipients		q.s.	
		Approved colour used			
263	MEMANTINE HCL TABLETS 10 MG	Each film coated tablet contains:			APPROVED
		Memantine Hcl	USP	10mg	
		Excipients		q.s.	
264	DOXYLAMINE SUCCINATE & PYRIDOXINE HCL TABLETS	Approved colour used			APPROVED
		Each enteric coated tablet contains:			
		Doxylamine Succinate	BP	20mg	
265	ARIPIRAZOLE TABLETS IP 10 MG	Pyridoxine Hydrochloride	IP	20mg	APPROVED
		Excipients		q.s.	
		Approved colour used			
266	ARIPIRAZOLE TABLETS IP 15 MG	Each uncoated tablet contains:			APPROVED
		Aripiprazole	IP	10mg	
		Excipients		q.s.	
267	DICLOFENAC SODIUM & TIZANIDINE TABLETS	Each uncoated tablet contains:			APPROVED
		Aripiprazole	IP	15mg	
		Excipients		q.s.	
268	DEFLAZACORT TABLETS 30MG	Diclofenac Sodium	IP	50mg	APPROVED
		Tizanidine Hydrochloride	IP		
		eq. to Tizanidine		2mg	
269	CHOLECALCIFEROL SACHET 1GM	Excipients		q.s.	APPROVED
		Each uncoated tablet contains:			
		Deflazacort		30 mg	
270	OXCARBAZEPINE TABLETS IP 150MG	Excipients		qs	APPROVED
		Each 1gm Sachet contains:			
		Cholecalciferol	IP	60000 I.U.	
271	PROMETHAZINE HYDROCHLORIDE TABLETS IP 25MG	Excipients		qs	APPROVED
		Each uncoated tablet contains:			
		Oxcarbazepine	IP	150 mg	
271	PROMETHAZINE HYDROCHLORIDE TABLETS IP 25MG	Excipients		q.s.	APPROVED
		Each film coated tablet contains:			
		Promethazine Hydrochloride	IP	25mg	
271	PROMETHAZINE HYDROCHLORIDE TABLETS IP 25MG	Excipients		q.s.	APPROVED
		Approved colour used			
		Each film coated tablet contains:			

272	<b>LACTULOSE &amp; ISPAGHULA HUSK GRANULES</b>	Each 15 gm sachet contains:			<b>APPROVED</b>
		Lactulose concentrate	IP		
		eq. to Lactulose		10mg	
		Ispaghula Husk	IP	3.5gm	
		Excipients		q.s.	
		Approved colour used			
273	<b>PIOGLITAZONE TABLETS IP 15MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Pioglitazone hydrochloride	IP		
		eq to pioglitazone		15mg	
		Excipients		q.s.	
274	<b>HALOPERIDOL TABLETS 0.25MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Haloperidol	IP	0.25mg	
		Excipients		q.s.	
275	<b>VOGLIBOSE &amp; METFORMIN HYDROCHLORIDE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Voglibose	IP	0.2 mg	
		Metformin Hydrochloride	IP	500 mg	
		Excipients		q.s.	
276	<b>GLICLAZIDE &amp; METFORMIN HYDROCHLORIDE HCL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Gliclazide	IP	80 mg	
		Metformin Hydrochloride	IP	500 mg	
		Excipients		q.s.	
277	<b>CLONAZEPAM TABLETS 0.25 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clonazepam	IP	0.25mg	
		Excipients		qs	
278	<b>ETIZOLAM TABLETS 0.25 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Etizolam		0.25mg	
		Excipients		q.s.	
279	<b>ETIZOLAM TABLETS 0.5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Etizolam		0.5mg	
		Excipients		q.s.	
280	<b>DISULFIRAM TABLETS 250 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Disulfiram	IP	250mg	
		Excipients		q.s.	
281	<b>NORTRIPTYLINE TABLETS IP 10 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Nortriptyline Hydrochloride	IP		
		eq. to Nortriptyline		10mg	
		Excipients		qs	
		Approved colour used			
282	<b>PROCYCLIDINE TABLETS IP 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Procyclidine Hydrochloride	IP	5mg	
		Excipients		qs	
283	<b>NORTRIPTYLINE TABLETS 50 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Nortriptyline Hydrochloride	IP		
		eq. to Nortriptyline		50mg	
		Excipients		qs	
		Approved colour used			
284	<b>PYRIDOXINE HCL, FOLIC ACID , THIAMINE MONONITRATE , ALPHA LIPOIC ACID &amp; MECOBALAMINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pyridoxine Hydrochloride	IP	3mg	
		Folic Acid	IP	1.5mg	
		Alpha Lipoic Acid	USP	100mg	
		Mecobalamine	USP	1500mcg	
		Thiamine Mono Nitrate	IP	10mg	
		Excipients		q.s.	
		Approved colour used			

285	<b>DIAZEPAM TABLETS IP 10MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Diazepam	IP	10 mg	
		Excipients		qs	
286	<b>FLUVOXAMINE TABLETS IP 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fluvoxamine maleate	IP	100 mg	
		Excipients		q.s.	
287	<b>FLUVOXAMINE TABLETS IP 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fluvoxamine maleate	IP	50 mg	
		Excipients		q.s.	
288	<b>MECOBALAMIN, ALPHA LIPOIC ACID, PYRIDOXINE HCl &amp; FOLIC ACID TABLETS</b>	Each film coated tablets contains:			<b>APPROVED</b>
		Mecobalamin	USP	1500 mcg	
		Alpha Lipoic Acid	USP	100 mg	
		Pyridoxine Hydrochloride	IP	3 mg	
		Folic Acid	IP	1.5 mg	
		Excipients		q.s.	
		Approved colour used			
289	<b>DESVENLAFAXINE EXTENDED RELEASE TABLETS 50MG</b>	Each uncoated extended release tablet contains:			<b>APPROVED</b>
		Desvenlafaxine Succinate			
		eq. to Desvenlafaxine		50mg	
		Excipients		q.s.	
		Approved colour used			
290	<b>DESVENLAFAXINE EXTENDED RELEASE TABLETS 100MG</b>	Each uncoated extended release tablet contains:			<b>APPROVED</b>
		Desvenlafaxine Succinate			
		eq to Desvenlafaxine		100mg	
		Excipients		q.s.	
		Approved colour used			
291	<b>ARIPIRAZOLE TABLETS IP 20MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Aripiprazole	IP	20 mg	
		Excipients		q.s.	
292	<b>CLOMIPRAMINE HYDROCHLORIDE TABLETS 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Chlomipramine Hydrochloride	IP	25mg	
		Excipients		q.s.	
293	<b>ETAMSYLATE TABLETS 500 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Etamsylate	BP	500mg	
		Excipients		q.s.	
294	<b>DICLOFENAC POTASSIUM &amp; SERRATIOPEPTIDASE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Diclofenac Potassium	BP	50mg	
		Serratiopeptidase(as enteric coated) (20,000 unit )	IP	10mg	
		Excipients		q.s.	
		Approved colour used			
295	<b>ROSUVASTATIN TABLETS IP 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium	IP		
		eqv. to Rosuvastatin		5 mg	
		Excipients		q.s.	
		Approved colour used			

296	<b>ATORVASTATIN TABLETS IP 40MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Atorvastatin Calcium	IP		
		eq. to Atorvastatin		40mg	
		Excipients		q.s.	
		Approved colour used			
297	<b>SILDENAFIL TABLETS IP 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sildenafil Citrate	IP		
		eq. to sildenafil		25mg	
		Excipients		q.s.	
		Approved colour used			
298	<b>SILDENAFIL TABLETS IP 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sildenafil Citrate	IP		
		eq. to sildenafil		50mg	
		Excipients		q.s.	
		Approved colour used			
299	<b>TADALAFIL TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tadalafil	IP	20mg	
		Excipients		q.s.	
		Approved colour used			
300	<b>ROSUVASTATIN TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium	IP		
		eq. to Rosuvastatin		10 mg	
		Excipients		q.s.	
		Colour: Titanium Dioxide			
301	<b>METFORMIN HYDROCHLORIDE SUSTAIN RELEASE TABLETS IP 500MG</b>	Each uncoated sustain release tablet contains:			<b>APPROVED</b>
		Metformin Hydrochloride	IP	500 mg	
		Excipients		q.s.	
302	<b>METFORMIN HYDROCHLORIDE SUSTAIN RELEASE TABLETS IP 850MG</b>	Each uncoated sustain release tablet contains:			<b>APPROVED</b>
		Metformin Hydrochloride	IP	850 mg	
		Excipients		q.s.	
303	<b>GLIMEPIRIDE &amp; METFORMIN HYDROCHLORIDE (SR) TABLETS</b>	Each uncoated bilayered tablet contains:			<b>APPROVED</b>
		Glimepiride	IP	1 mg	
		Metformin Hydrochloride (as sustained release form)	IP	1000 mg	
		Excipients		q.s.	
		Approved colour used			
304	<b>GLIMEPIRIDE &amp; METFORMIN HYDROCHLORIDE (SR) TABLETS</b>	Each uncoated bilayered tablet contains:			<b>APPROVED</b>
		Glimepiride	IP	2 mg	
		Metformin Hydrochloride (as sustained release form)	IP	1000 mg	
		Excipients		q.s.	
		Approved colour used			
305	<b>SULFASALAZINE DELAYED RELEASE TABLETS 500MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Sulfasalazine	USP	500mg	
		Excipients		q.s.	
		Approved colour used			
306	<b>LEVOCETIRIZINE DIHYDROCHLORIDE &amp; MONTELUKAST TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levocetirizine Dihydrochloride	IP	5mg	
		Montelukast Sodium	IP		
		eqv. to Montelukast		10 mg	
		Excipients		q.s.	

307	<b>PARACETAMOL &amp; IBUPROFEN TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	IP	325 mg	
		Ibuprofen	IP	400mg	
		Excipients		q.s.	
		Approved colour used			
308	<b>OLANZAPINE &amp; FLUOXETINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	10mg	
		Fluoxetine Hcl	IP		
		eq. to fluoxetine		20mg	
		Excipients		q.s.	
		Approved colour used			
309	<b>DICLOFENAC POTASSIUM &amp; METAXALONE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Diclofenac Potassium	BP	50mg	
		Metaxalone		400mg	
		Excipients		q.s.	
		Approved colour used			
310	<b>TOLPERISONE HYDROCHLORIDE SR TABLETS 450 MG</b>	Each film coated Tablets contains:			<b>APPROVED</b>
		Tolperisone Hydrochloride		450mg	
		Excipients		q.s.	
		Approved colour used			
311	<b>ETORICOXIB &amp; PARACETAMOL TABLES</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Etoricoxib	IP	60mg	
		Paracetamol	IP	325mg	
		Excipients		q.s.	
		Approved colour used			
312	<b>GLUCOSAMINE HCL &amp; DIACERIN TABLETS</b>	Each film coated Tablet contains:			<b>APPROVED</b>
		Glucosamine HCl	BP		
		eq. to glucosamine		446mg	
		Diacerin	IP	50mg	
		Excipients		q.s.	
		Approved colour used			
313	<b>GLUCOSAMINE HCL TABLETS</b>	Each film coated Tablet contains:			<b>APPROVED</b>
		Glucosamine HCl	BP	750mg	
		Excipients		q.s.	
		Approved colour used			
314	<b>ZINC SULPHATE TABLETS 10MG</b>	Each film coated Tablet contains:			<b>APPROVED</b>
		Zinc Sulphate	IP	10mg	
		Excipients		qs	
		Approved colour used			
315	<b>METOPROLOL SUCCINATE SUSTAINED RELEASE TABLETS IP 50 MG</b>	Each uncoated sustained release tablet contains:			<b>APPROVED</b>
		Metoprolol Succinate	IP		
		eq. to Metoprolol Tartrate		50 mg	
		Excipients		q.s.	
316	<b>CAMYLOFIN HCL &amp; PARACETAMOL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Camylofin Dihydrochloride		25mg	
		Paracetamol	IP	300 mg	
		Excipients		qs	
		Approved Colour Used			
317	<b>TIANEPTINE SODIUM TABLETS 12.5 MG</b>	Each uncoated sustained release tablet contains:			<b>APPROVED</b>
		Tianeptine Sodium	BP	12.5mg	
		Excipients		q.s.	

318	<b>ACECLOFENAC, PARACETAMOL &amp;CHLORZOXAZONE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aceclofenac	IP	100 mg	
		Chlorzoxazone	USP	500mg	
		Paracetamol	IP	325 mg	
		Excipients		q.s.	
		Approved colour used			
319	<b>GLICLAZIDE (M.R.) &amp; METFORMIN HYDROCHLORIDE (E.R.) TABLETS</b>	Each uncoated bilayered tablet contains			<b>APPROVED</b>
		Gliclazide (as M.R.)	IP	60 mg	
		Metformin Hydrochloride (as E.R.)	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			
320	<b>PIOGLITAZONE &amp; METFORMIN ER TABLETS</b>	Each uncoated bilayered tablet contains:			<b>APPROVED</b>
		Pioglitazone hydrochloride	IP		
		eq. to Pioglitazone		15mg	
		Metformin HCl (as Sustained Release form)	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			
321	<b>PREGABALIN (S.R.) &amp; METHYLCOBALAMIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pregabalin (S.R.)	IP	150 mg	
		Methylcobalamin	USP	1500mcg	
		Excipients		q.s.	
		Approved colour used			
322	<b>ETIZOLAM TABLETS 1MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Etizolam		1mg	
		Excipients		q.s.	
323	<b>ACAMPROSATE CALCIUM TABLETS 333MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Acamprosate Calcium	IP	333mg	
		Excipients		qs	
		Approved colour used			
324	<b>IMIPRAMINE HCL&amp;DIAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Imipramine Hcl	IP	50 mg	
		Diazepam	IP	5 mg	
		Excipients		qs	
		Approved colour used			
325	<b>DICYCLOMINE&amp; PARACETAMOL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Dicyclomine Hcl	IP	20mg	
		Paracetamol	IP	325mg	
		Excipients		qs	
		Approved colour used			
326	<b>CALCIUM CITRATE &amp;CALCITRIOL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium Citrate	USP		
		eq. to elemental calcium		252mg	
		Calcitriol	IP	0.25mcg	
		Excipients		q.s.	
		Approved colour used			
327	<b>GLIMEPIRIDE &amp; METFORMIN HYDROCHLORIDE (SR) TABLETS</b>	Each uncoated bilayered tablet contains			<b>APPROVED</b>
		Glimepiride	IP	3mg	
		Metformin Hydrochloride (as sustained release form)	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			

328	<b>ALPRAZOLAM &amp;MELATONIN TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	0.5mg	
		Melatonin	BP	3mg	
		Excipients		q.s.	
		Approved colour used			
329	<b>ALPRAZOLAM &amp; MELATONIN TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	0.25mg	
		Melatonin	BP	3mg	
		Excipients		q.s.	
		Approved colour used			
330	<b>METHYLCOBALAMIN SUBLINGUAL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Methylcobalamin	USP	1500mcg	
		Excipients		q.s.	
331	<b>DICLOFENAC POTASSIUM, PARACETAMOL &amp; SERRATIOPEPTIDASE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Diclofenac Potassium	BP	50 mg	
		Paracetamol	IP	325 mg	
		Serratiopeptidase	IP	15mg	
		Excipients		q.s.	
332	<b>BIOTIN,CALCIUM PANTOTHENATE, ACETYLCYSTEINE&amp; MINERALS TABLETS</b>	Approved colour used			<b>APPROVED</b>
		Each film coated tablet contains:			
		Biotin	USP	10mg	
		Acetylcysteine	USP	50mg	
		Calcium Pantothenate	IP	100mg	
		Sodium Selenite	USP		
		eq. to elemental selenium		65mcg	
		Cupric Oxide			
		eq. to elemental copper		3mg	
		Zinc Oxide	IP		
		eq. to elemental zinc		22.5mg	
		Excipients		q.s.	
333	<b>DOMPERIDONE &amp; CINNARIZINE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Domperidone	IP	15mg	
		Cinnarizine	IP	20mg	
		Excipients		q.s.	
334	<b>RISPERIDONE MOUTH DISSOLVING TABLET 2 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Risperidone	BP	2 mg	
		Excipients		q.s.	
335	<b>RESPERIDONE MOUTH DISSOLVING TABLET 4 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Risperidone	BP	4 mg	
		Excipients		q.s.	
336	<b>CLONAZEPAM MOUTH DISSOLVING TABLETS 0.25 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Clonazepam	IP	0.25mg	
		Excipients		q.s.	
337	<b>CLONAZEPAM MOUTH DISSOLVING TABLETS 0.5 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Clonazepam	IP	0.5mg	
		Excipients		q.s.	
338	<b>CLONAZEPAM MOUTH DISSOLVING TABLETS 1 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Clonazepam	IP	1mg	
		Excipients		q.s.	



339	<b>CLONAZEPAM MOUTH DISSOLVING TABLETS 2 MG</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Clonazepam	IP	2mg	
		Excipients		q.s.	
340	<b>RUTIN, ASCORBIC ACID,VITAMIN K3, VITAMIN D-3 &amp; ADRENOCHROME MONOSEMICARBAZONE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rutin		100mg	
		Ascorbic Acid	IP	15mg	
		Menadione Sodium Bisulphate(vit k3)		20mg	
		Cholecalciferol(vit d3)	IP	300IU	
		Adrenochrome Monosemicarbazone		1.0mg	
		Excipients		q.s.	
		Approved colour used			
341	<b>PROPRANOLOL&amp; DIAZEPAM TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Propranolol Hcl	IP	20mg	
		Diazepam	IP	2.5mg	
		Excipients		q.s.	
342	<b>BETAMETHASONE TABLETS IP 0.5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Betamethasone	IP	0.5mg	
		Excipients		q.s.	
		Approved colour used			
343	<b>FEXOFENADINE HYDROCHLORIDE TABLETS IP 120MG</b>	Each film coated contains:			<b>APPROVED</b>
		Fexofenadine Hydrochloride	IP	120mg	
		Excipients		qs	
		Approved colour used			
344	<b>FEXOFENADINE HYDROCHLORIDE TABLETS IP 180MG</b>	Each film coated contains:			<b>APPROVED</b>
		Fexofenadine Hydrochloride	IP	180mg	
		Excipients		qs	
		Approved colour used			
345	<b>FLUPIRTINE MALEATE CAPSULES 100MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Flupirtine Maleate		100mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shells			
346	<b>HYDROCHLOROTHIAZIDE TABLETS IP 12.5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Hydrochlorothiazide	IP	12.5mg	
		Excipients		qs	
		Approved colour used			
347	<b>HYDROCHLOROTHIAZIDE &amp; AMLODIPINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Hydrochlorothiazide	IP	12.5 mg	
		Amlodipine Besilate	IP		
		eq. to Amlodipine		5.0mg	
		Excipients		q.s.	
		Approved colour used			
348	<b>ASPIRIN TABLETS IP 75MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aspirin	IP	75mg	
		Excipients		q.s.	
		Approved colour used			
349	<b>FINASTERIDE TABLETS 1MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Finasteride	IP	1 mg	
		Excipients		q.s.	
		Approved colour used			

350	<b>TERBINAFINE HYDROCHLORIDE TABLETS 250 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Terbinafine Hydrochloride	BP		
		eq. to Terbinafine		250mg	
		Excipients		qs	
		Approved colour used			
351	<b>PROMETHAZINE HYDROCHLORIDE TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Promethazine Hydrochloride	IP	10mg	
		Excipients		q.s.	
		Approved colour used			
352	<b>VOGLIBOSE &amp; METFORMIN HYDROCHLORIDE (SR) TABLETS</b>	Each uncoated bilayered tablet contains:			<b>APPROVED</b>
		Voglibose	IP	0.3mg	
		Metformin Hydrochloride (as sustained release form)	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			
353	<b>ISOXSUPRINE TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Isoxsuprine Hydrochloride	IP	20mg	
		Excipients		q.s.	
		Approved colour used			
354	<b>GABAPENTIN &amp; METHYLCOBALAMIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Gabapentin	IP	100 mg	
		Methylcobalamin	USP	500mcg	
		Excipients		q.s.	
		Approved colour used			
355	<b>TELMISARTAN, AMLODIPINE &amp; HYDROCHLOROTHIAZIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	40 mg	
		Amlodipine Besilate	IP		
		eq. to Amlodipine		5mg	
		Hydrochlorothiazide	IP	12.5mg	
		Excipients		q.s.	
		Approved colour used			
356	<b>PAROXETINE (C.R.) &amp; CLONAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Paroxetine Hydrochloride	IP		
		eq. to Paroxetine (C.R.)		12.5mg	
		Clonazepam	IP	0.5mg	
		Excipients		q.s.	
		Approved colour used			
357	<b>GLICLAZIDE (M.R.) &amp; METFORMIN HYDROCHLORIDE (E.R.) TABLETS</b>	Each uncoated bilayered tablet contains			<b>APPROVED</b>
		Gliclazide (M.R.)	IP	80 mg	
		Metformin Hydrochloride (as extended-release form)	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			
358	<b>MEBEVERINE HYDROCHLORIDE &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Mebeverine Hydrochloride	IP	135 mg	
		Chlordiazepoxide	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
359	<b>CLIDINIUM BROMIDE &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clidinium bromide	USP	2.5 mg	
		Chlordiazepoxide	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			

360	<b>CHOLECALCIFEROL TABLETS IP 1MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Cholecalciferol	IP	1 mg	
		Excipients		q.s.	
361	<b>SUMATRIPTAN &amp; NAPROXEN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Naproxen	IP	500mg	
		Sumatriptan Succinate	IP		
		eq to Sumatriptan		85mg	
		Approved colour used			
362	<b>SUMATRIPTAN TABLETS 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sumatriptan Succinate			
		eq. to Sumatriptan	IP	50mg	
		Approved colour used			
363	<b>SUMATRIPTAN TABLETS 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Sumatriptan Succinate			
		eq. to Sumatriptan	IP	100mg	
		Approved colour used			
364	<b>PIOGLITAZONE&amp; METFORMIN HCL (ER) TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Pioglitazone hydrochloride	IP		
		eq. to pioglitazone		15mg	
		Metformin HCl (ER)	IP	1000 mg	
		Excipients		q.s.	
365	<b>AMANTADINE HYDROCHLORIDE CAPSULES IP 100MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Amantadine hydrochloride	IP	100 mg	
		Excipients		qs	
		Approved colour used in empty capsule shell			
366	<b>METOPROLOL SUCCINATE EXTENDED RELEASE TABLETS IP 25MG</b>	Each film coated tablet extended release tablet contains:			<b>APPROVED</b>
		Metoprolol Succinate	IP	23.75mg	
		eqv. to Metoprolol Tartrate		25 mg	
		Excipients		q.s.	
		Approved colour used			
367	<b>METOPROLOL SUCCINATE (E.R.)&amp; AMLODIPINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Metoprolol Succinate	IP	47.5mg	
		eqv. to Metoprolol Tartrate		50 mg	
		(as extended Release)			
		Amlodipine Besilate	IP		
		eqv. to Amlodipine		5 mg	
		Excipients		q.s.	
		Approved colour used			
368	<b>DOXOFYLLINE TABLETS IP 400MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Doxofylline	IP	400mg	
		Excipients		q.s.	
369	<b>THEOPHYLLINE C.R. &amp;MONTELUKAST TABLETS</b>	Each film coated bilayered tablet contains:			<b>APPROVED</b>
		Theophylline (anhydrous)	IP	400mg	
		(as controlled release form)			
		Montelukast sodium	IP		
		eq. to Montelukast		10mg	
		Excipients		q.s.	
		Approved colour used			

370	<b>DOXOFYLLINE &amp; MONTELUKAST TABLETS</b>	Each film coated bilayered tablet			<b>APPROVED</b>
		contains:			
		Doxofylline	IP	400mg	
		(as extended release)			
		Montelukast sodium	IP		
		eq. to Montelukast		10mg	
		Excipients		q.s.	
		Approved colour used			
371	<b>GLICLAZIDE TABLETS 40MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Gliclazide	IP	40mg	
		Excipients		q.s.	
		Approved colour used			
372	<b>GLICLAZIDE TABLETS IP 80MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Gliclazide	IP	80mg	
		Excipients		q.s.	
		Approved colour used			
373	<b>PREDNISOLONE TABLETS IP 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		prednisolone	IP	5mg	
		Excipients		q.s.	
374	<b>PREDNISOLONE TABLETS IP 10 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		prednisolone	IP	10mg	
		Excipients		q.s.	
375	<b>PREDNISOLONE TABLETS IP 20 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		prednisolone	IP	20mg	
		Excipients		q.s.	
376	<b>NIMESULIDE &amp; TIZANIDINE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Nimesulide	IP	100mg	
		Tizanidine Hcl	IP		
		eq to Tizanidine		2 mg	
		Excipients		q.s.	
377	<b>TRIFLUOPERAZINE HCl &amp; TRIHEXYPHENIDYL HCl TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Trifluoperazine Hydrochloride	IP		
		eqv. to Trifluoperazine		2.5mg	
		Trihexyphenidyl Hydrochloride	IP	1 mg	
		Excipients		q.s.	
378	<b>ARIPIRAZOLE TABLETS IP 30MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aripiprazole	IP	30 mg	
		Excipients		q.s.	
		Approved colour used			
379	<b>TELMISARTAN TABLETS IP 20MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	20 mg	
		Excipients		q.s.	
380	<b>TELMISARTAN TABLETS IP 40MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	40 mg	
		Excipients		q.s.	
381	<b>TELMISARTON TABLETS IP 80MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Telmisarton	IP	80 mg	
		Excipients		q.s.	
382	<b>LINEZOLID TABLETS IP 600MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Linezolid	IP	600 mg	
		Excipients		q.s.	
383	<b>HYDROCHLOROTHIAZIDE TABLETS 12.5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Hydrochlorothiazide	IP	12.5 mg	
		Excipients		q.s.	

384	<b>NALTREXONE HCL TABLETS IP 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Naltrexone Hcl	IP	50 mg	
		Excipients		q.s.	
		Approved colour used			
385	<b>NAPROXEN SUSTAINED RELEASE TABLETS IP 375MG</b>	Each film coated sustained release tablet contains			<b>APPROVED</b>
		Naproxen	IP	375 mg	
		Excipients		q.s.	
		Approved colour used			
386	<b>LACTITOL MONOHYDRATE &amp; ISPAGHULA HUSK GRANULES</b>	Each 15 gm granules contain:			<b>APPROVED</b>
		Lactitol Monohydrate	USP	10 gm	
		Ispaghula Husk	IP	3.5gm	
		Excipients		q.s.	
387	<b>RIFAXIMIN TABLETS 400MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rifaximin	BP	400 mg	
		Excipients		q.s.	
		Approved colour used			
388	<b>THEOPHYLLINE PROLONGED RELEASE TABLETS 200MG</b>	Each uncoated prolonged-release tablet contains:			<b>APPROVED</b>
		Theophylline (anhydrous)	IP	200 mg	
		Excipients		q.s.	
389	<b>TAPENTADOL HYDROCHLORIDE TABLETS 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tapentadol Hydrochloride	IP	50 mg	
		eqv. to Tapentadol			
		Excipients		q.s.	
390	<b>TAPENTADOL HYDROCHLORIDE TABLETS 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tapentadol Hydrochloride	IP	100 mg	
		eqv. to Tapentadol			
		Excipients		q.s.	
391	<b>RIFAXIMIN TABLETS 200MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rifaximin	BP	200 mg	
		Excipients		q.s.	
		Approved colour used			
392	<b>AZITHROMYCIN DISPERSIBLE TABLETS 100MG</b>	Each uncoated dispersible tablet			<b>APPROVED</b>
		Contains:			
		Azithromycin Dihydrate	IP		
		eqv. Azithromycin Anhydrous		100 mg	
393	<b>AMISULPRIDE SUSTAINED RELEASE TABLETS 300MG</b>	Each uncoated sustained-release tablet contains:			<b>APPROVED</b>
		Amisulpride	IP	300 mg	
		Excipients		q.s.	
394	<b>METOPROLOL SUCCINATE (E.R) &amp; AMLODIPINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Metoprolol Succinate	IP		
		eqv. to Metoprolol Tartrate (Exetended Release)		25 mg	
		Amlodipine Besilate	IP		
		eqv. to Amlodipine		2.5 mg	
		Excipients		q.s.	
		Approved colour used			

395	<b>CALCIUM, MAGNESIUM, VITAMIN D3 &amp; MINERALS TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium Carbonate	IP	1250 mg	
		Magnesium Hydroxide	IP	25 mg	
		Manganese Sulphate	BP	0.1 mg	
		Pyridoxine Hydrochloride	IP	1.5 mg	
		Folic Acid	IP	0.5 mg	
		Cholecalciferol	IP	200 IU	
		Zinc Sulphate Monohydrate	IP	40 mg	
		Copper Sulphate	BP	0.1 mg	
		Excipients		q.s.	
		Approved colour used			
396	<b>METOPROLOL SUCCINATE (E.R) &amp; TELMISARTAN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Metoprolol Succinate	IP		
		eqv. to Metoprolol Tartrate (Exetended Release)		25 mg	
		Telmisartan	IP	40 mg	
		Excipients		q.s.	
		Approved colour used			
397	<b>FLUCONAZOLE TABLTs I.P.150MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fluconazole	IP	150 mg	
		Excipients		q.s.	
		Approved colour used			
398	<b>VINPOCETINE TABLETS 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		vinpocetine	USP	5 mg	
		Excipients		q.s.	
		Approved colour used			
399	<b>PARACETAMOL &amp; TRAMADOL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	IP	325 MG	
		Tramadol Hcl	IP	37.5 MG	
		Excipients		q.s.	
400	<b>BUPROPION HYDROCHLORIDE EXTENDED RELEASE TABLETS 150MG</b>	Each film coated extended release tablet Contains:			<b>APPROVED</b>
		Bupropion Hydrochloride	USP	150 mg	
		Excipients		q.s.	
		Approved colour used			
401	<b>BUPROPION HYDROCHLORIDE EXTENDED RELEASE TABLETS 300MG</b>	Each film coated extended release tablet Contains:			<b>APPROVED</b>
		Bupropion Hydrochloride	USP	300 mg	
		Excipients		q.s.	
		Approved colour used			
402	<b>ACECLOFENAC AND THIOLCHICOSIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aceclofenac	IP	100 mg	
		Thiocolchicoside	IP	4 mg	
		Excipients		q.s.	
		Approved colour used			
403	<b>FEXOFENADINE AND MONTELUKAST CHEWABLE TABLETS</b>	Each uncoated chewable tablet contains:			<b>APPROVED</b>
		Fexofenadine Hydrochloride	IP		
		eq. to Fexofenadine		120 mg	
		Montelukast Sodium	IP		
		eq. to Montelukast		10 mg	
		Excipients		q.s.	

404	<b>CALCIUM CITRATE MALEATE, VITAMIN D3 &amp; FOLIC ACID TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium Citrate Maleate			
		eq. to elemental Calcium		250 mg	
		Vitamin D3	IP	100 IU	
		(as Colecalciferol Concentrate)			
		Folic Acid	IP	50 mcg	
		Excipients		q.s.	
		Approved colour used			
405	<b>ROSUVASTATIN &amp; FENOFIBRATE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium	IP		
		eq. to Rosuvastatin		10 mg	
		Fenofibrate	IP	160 mg	
		Excipients		q.s.	
		Approved colour used			
406	<b>ARMODAFINIL TABLETS 50 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Armodafinil		50 mg	
		Excipients		q.s.	
407	<b>ARMODAFINIL TABLETS 150MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Armodafinil		150 mg	
		Excipients		q.s.	
408	<b>ARMODAFINIL TABLETS 250MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Armodafinil		250 mg	
		Excipients		q.s.	
409	<b>TAPENTADOL TABLETS 75MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tapentadol Hydrochloride	IP		
		eq. to Tapentadol		75 mg	
		Excipients		q.s.	
		Approved colour used			
410	<b>PAROXETINE TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Paroxetine Hydrochloride	IP		
		eq. to Paroxetine		20 mg	
		Excipients		q.s.	
		Approved colour used			
411	<b>TIAPRIDE TABLETS 25MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tiapride Hydrochloride			
		eqv. to Tiapride		25 mg	
		Excipients		q.s.	
		Approved colour used			
412	<b>TIAPRIDE TABLETS 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tiapride Hydrochloride			
		eqv. to Tiapride		50 mg	
		Excipients		q.s.	
		Approved colour used			
413	<b>TIAPRIDE TABLETS 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tiapride Hydrochloride			
		eqv. to Tiapride		100 mg	
		Excipients		q.s.	
		Approved colour used			
414	<b>TELMISARTAN &amp; AMLODIPINE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	40 mg	
		Amlodipine Besilate	IP		
		eqv. to Amlodipine		10 mg	
		Excipients		q.s.	



415	<b>TELMISARTAN &amp; AMLODIPINE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	IP	40 mg	
		Amlodipine Besilate	IP		
		eqv. to Amlodipine		5 mg	
		Excipients		q.s.	
416	<b>MODAFINIL TABLETS 50 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Modafinil	BP	50 mg	
		Excipients		q.s.	
		Approved colour used			
417	<b>MODAFINIL TABLETS 100 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Modafinil	BP	100 mg	
		Excipients		q.s.	
		Approved colour used			
418	<b>GABAPENTIN TABLETS IP 300 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Gabapentin	IP	300 mg	
		Excipients		q.s.	
		Approved colour used			
419	<b>FERROUS ASCORBATE, CYANOCOBALAMIN, FOLIC ACID &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ferrous Ascorbate			
		eqv. to Elemental Iron		100 mg	
		Cyanocobalamin	IP	15 mcg	
		Folic Acid	IP	1.5 mg	
		Zinc Sulphate Monohydrate	IP		
		eqv. To Elemental Zinc		22.5 mg	
		Excipients		q.s.	
		Approved colour used			
420	<b>ILAPRAZOLE TABLETS</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Ilaprazole	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
421	<b>ILAPRAZOLE TABLETS</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Ilaprazole	IP	10 mg	
		Excipients		q.s.	
		Approved colour used			
422	<b>RAMIPRIL &amp; HYDROCHLOROTHIAZIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ramipril	IP	5 mg	
		Hydrochlorothiazide	IP	12.5 mg	
		Excipients		q.s.	
423	<b>THIAMINE TABLETS IP 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Thiamine Hydrochloride	IP	50 mg	
		Excipients		q.s.	
424	<b>CHOLECALCIFEROL TABLETS IP 0.25 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Cholecalciferol	IP	0.25 mg	
		Excipients		q.s.	
		Approved colour used			
425	<b>MONTELUKAST &amp; DESLORATADINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Montelukast Sodium	IP		
		eq. to Montelukast		10 mg	
		Desloratadine		5 mg	
		Excipients		q.s.	
		Approved colour used			

426	<b>RACECADOTRIL CAPSULES IP 100MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Racecadotril	IP	100 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
427	<b>VOGLIBOSE MOUTH DISSOLVING TABLETS</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Voglibose	IP	0.2 mg	
		Excipients		q.s.	
428	<b>VOGLIBOSE MOUTH DISSOLVING TABLETS</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Voglibose	IP	0.3 mg	
		Excipients		q.s.	
429	<b>LEVOCETIRIZINE DIHYDROCHLORIDE &amp; MONTELUKAST DISPERSIBLE TABLETS</b>	Each uncoated dispersible tablet contains:			<b>APPROVED</b>
		Levocetirizine Dihydrochloride	IP	2.5 mg	
		Montelukast Sodium	IP		
		eqv. to Montelukast		5 mg	
		Excipients		q.s.	
430	<b>MINOCYCLINE HYDROCHLORIDE TABLETS 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Minocycline Hydrochloride	USP		
		eqv. to Minocycline		100 mg	
		Excipients		q.s.	
		Approved colour used			
431	<b>MINOCYCLINE HYDROCHLORIDE TABLETS 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Minocycline Hydrochloride	USP		
		eqv. to Minocycline		50 mg	
		Excipients		q.s.	
		Approved colour used			
432	<b>KETOROLAC TROMETHAMINE DISPERSIBLE TABLETS</b>	Each uncoated dispersible tablet contains:			<b>APPROVED</b>
		Ketorolac Tromethamine	I.P.	10 mg	
		Excipients			
		Approved colour used			
433	<b>ACEBROPHYLLINE CAPSULES 100MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Acebrophylline		100 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
434	<b>RABEPRAZOLE SODIUM (EC) &amp; LEVOSULPIRIDE (SR) CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Rabeprazole Sodium (As enteric coated pellets)	IP	20 mg	
		Levosulpiride (As sustained release pellets)		75 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
435	<b>ESOMEPRAZOLE (EC) &amp; DOMPERIDONE (SR) CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Esomeprazole Magnesium (As enteric coated pellets)	IP	20 mg	
		Domperidone (As sustained release pellets)	IP	30 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			

436	<b>PROPRANOLOL SUSTAINED RELEASE TABLETS 80 MG</b>	Each uncoated sustained-release			<b>APPROVED</b>
		tablet contains:			
		Propranolol Hydrochloride	IP	80 mg	
		Excipients		q.s.	
437	<b>OXCARBAZEPINE SUSTAINED RELEASE TABLETS 300MG</b>	Each film coated sustained-release			<b>APPROVED</b>
		tablet contains:			
		Oxcarbazepine	IP	300 mg	
		Excipients		q.s.	
		Approved colour used			
438	<b>OXCARBAZEPINE SUSTAINED RELEASE TABLETS 600MG</b>	Each film coated sustained-release			<b>APPROVED</b>
		tablet contains:			
		Oxcarbazepine	IP	600 mg	
		Excipients		q.s.	
		Approved colour used			
439	<b>AMISULPRIDE SUSTAINED RELEASE TABLETS 200MG</b>	Each uncoated sustained			<b>APPROVED</b>
		release tablet contains:			
		Amisulpride	IP	200 mg	
		Excipients			
440	<b>AMISULPRIDE SUSTAINED RELEASE TABLETS 400MG</b>	Each uncoated sustained			<b>APPROVED</b>
		release tablet contains:			
		Amisulpride	IP	400 mg	
		Excipients			
441	<b>GLICLAZIDE(MR)&amp; METFORMIN HCL (E.R.) TABLETS</b>	Each uncoated bilayered tablet			<b>APPROVED</b>
		contains:			
		Gliclazide(MR)	IP	40 mg	
		Metformin hcl	IP	500 mg	
		(extended release form)		q.s.	
		Excipients			
442	<b>MEBEVERINE HYDROCHLORIDE &amp; ISPAGHULA HUSK GRANULES</b>	Each sachet contains:			<b>APPROVED</b>
		Mebeverine Hydrochloride	IP	135 mg	
		Ispaghula husk	IP	3.5 gm	
		Excipients		q.s.	
443	<b>MEBEVERINE HYDROCHLORIDE &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Mebeverine Hydrochloride	IP	135 mg	
		Chlordiazepoxide	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
444	<b>MECOBALAMIN EXTENDED RELEASE TABLETS 1500 MCG</b>	Each uncoated extended-release tablet			<b>APPROVED</b>
		contains:			
		Mecobalamin	USP	1500 mcg	
		Excipients		q.s.	
445	<b>BACLOFEN EXTENDED RELEASE CAPSULES 30MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Baclofen	IP	30 mg	
		(As extended release form)			
		Excipients		q.s.	
		Approved colour used in empty			
		capsule Shell			
446	<b>TERBUTALINE SULPHATE &amp; DOXOFYLLIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Terbutaline sulphate	IP	5 mg	
		Doxofylline	IP	400 mg	
		Excipients		q.s.	
		Approved colour used			

447	<b>PYRIDOXINE HCl, CYANOCOBALAMIN, NICOTINAMIDE, FOLIC ACID &amp; CHROMIUM PICOLINATE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pyridoxine HCl	IP	3 mg	
		Nicotinamide	IP	100 mg	
		Cyanocobalamin	IP	15 mcg	
		Folic Acid	IP	500 mcg	
		Chromium Picolinate	USP	250 mcg	
		Excipients		q.s.	
		Approved colour used			
448	<b>MULTIVITAMIN, MULTIMINERAL &amp; ANTIOXIDANT TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Methylcobalamin	USP	500 mcg	
		Vitamin A Concentrate	IP	5000 IU	
		Vitamin D3	IP	400 IU	
		Vitamin E Acetate	IP	25 mg	
		Clacium Pantothenate	IP	50 mg	
		L-Glutamic Acid	IP	50 mg	
		Vitamin B1	IP	10 mg	
		Vitamin B2	IP	10 mg	
		Vitamin B6	IP	3 mg	
		Niacinamide	IP	50 mg	
		Vitamin C	IP	75 mg	
		Biotin	USP	260 mcg	
		Folic Acid	IP	1 mg	
		Zinc Sulphate	IP	60 mg	
		Chromium Picolinate	USP	150 mcg	
		Sodium Selenite	USP		
		eq. to selenium		100 mcg	
		Manganese Sulphate	BP	200 mcg	
		Magnesium Hydroxide	IP	10 mg	
		Copper Sulphate	BP	200 mcg	
		Excipients		q.s.	
		Approved colour used			
449	<b>LEVOCETIRIZINE DIHYDROCHLORIDE &amp; MONTELUKAST TABLETS</b>	Each uncoated dispersible tablet contains:			<b>APPROVED</b>
		Levocetirizine Dihydrochloride	IP	2.5 mg	
		Montelukast Sodium	IP		
		eqv. to Montelukast		4 mg	
		Excipients		q.s.	
450	<b>FEBUXOSTAT TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Febuxostat		80 mg	
		Excipients		q.s.	
		Approved colour used			
451	<b>RABEPRAZOLE SODIUM &amp; ACECLOFENAC (SR) CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Rabeprazole Sodium	IP	20 mg	
		(as enteric coated pellets)			
		Aceclofenac	IP	200 mg	
		(As sustained release pellets)			
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
452	<b>CALCIUM CITRATE MALEATE &amp; VITAMIN D3 TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium Citrate Maleate			
		eqv. to elemental Calcium		250 mg	
		Vitamin D3	IP	200 IU	
		Excipients		q.s.	
		Approved colour used			

453	<b>ORLISTAT CAPSULES 120MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Orlistat		120 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
454	<b>MULTIVITAMIN, MULTIMINERAL &amp; ANTIOXIDANT TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		β-Carotene	USP	20.66 mg	
		Vitamin E Acetate	IP	25 IU	
		Vitamin B6	IP	1.5 mg	
		Vitamin B1	IP	10 mg	
		Methylcobalamin	USP	500 mcg	
		Folic Acid	IP	1.5 mg	
		Alpha Lipoic Acid	USP	50 mg	
		Calcium Pantothenate	IP	10 mg	
		Selenium Dioxide		163.6 mcg	
		Excipients		q.s.	
		Approved colour used			
455	<b>BISACODYL GASTRO RESISTANT TABLETS IP 5MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Bisacodyl	IP	5 mg	
		Excipients		q.s.	
		Approved colour used			
456	<b>CHLORPROMAZINE HCL, TRIFLUOPERAZINE HCL &amp; TRIHEXYPHENIDYL HCL</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Chlorpromazine HCl	IP	50mg	
		Trifluoperazine HCl	IP		
		eq. to Trifluoperazine		5mg	
		Trihexyphenidyl HCl	IP	2mg	
		Excipients		q.s.	
		Approved colour used			
457	<b>LORNOXICAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Lornoxicam		4 mg	
		Excipients		q.s.	
		Approved colour used			
458	<b>LORNOXICAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Lornoxicam		8 mg	
		Excipients		q.s.	
		Approved colour used			
459	<b>ALPRAZOLAM PROLONGED- RELEASE TABLETS IP 0.5 MG</b>	Each uncoated prolonged-release tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	0.5 mg	
		Excipients		q.s.	
		Approved colour used			
460	<b>ALPRAZOLAM PROLONGED- RELEASE TABLETS IP 1 MG</b>	Each uncoated prolonged-release tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	1 mg	
		Excipients		q.s.	
		Approved colour used			
461	<b>ALPRAZOLAM PROLONGED- RELEASE TABLETS IP 1.5 MG</b>	Each uncoated prolonged-release tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	1.5 mg	
		Excipients		q.s.	
		Approved colour used			

462	<b>ESOMEPRAZOLE (EC) &amp; DOMPERIDONE (SR) CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Esomeprazole Magnesium	IP		
		eqv. to Esomeprazole		40 mg	
		(As enteric coated pellets)			
		Domperidone	IP	30 mg	
		(As sustained release pellets)			
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
463	<b>ROSUVASTATIN TABLETS 40MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium	IP		
		eq. to Rosuvastatin		40 mg	
		Excipients		q.s.	
		Approved colour used			
464	<b>GLICLAZIDE (M.R.) &amp; METFORMIN HYDROCHLORIDE (E.R.) TABLETS</b>	Each uncoated bilayered tablet contains			<b>APPROVED</b>
		Gliclazide (M.R.)	IP	30 mg	
		Metformin Hydrochloride	IP	500 mg	
		(Extended-release form)			
		Excipients		q.s.	
		Approved colour used			
465	<b>PERINDOPRIL ARGININE TABLETS 2.5 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Perindopril Arginine eq. to Perindopril		2.5 mg	
		Excipients		q.s.	
		Approved colour used			
466	<b>PERINDOPRIL ARGININE TABLETS 5 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Perindopril Arginine eq. to Perindopril		5 mg	
		Excipients		q.s.	
		Approved colour used			
467	<b>TRAMADOL HYDROCHLORIDE TABLETS 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Tramadol Hydrochloride	IP	50 mg	
		Excipients		q.s.	
468	<b>TRAMADOL HYDROCHLORIDE TABLETS 100MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Tramadol Hydrochloride	IP	100 mg	
		Excipients		q.s.	
469	<b>ORLISTAT CAPSULES 60MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Orlistat		60 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
470	<b>CHOLECALCIFEROL TABLETS 60000 IU</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Cholecalciferol (as stabilized)	IP	60000 IU	
		Excipients		q.s.	
		Approved colour used			
471	<b>METHYLCOBALAMIN, ALPHA LIPOIC ACID, PYRIDOXINE HCl, VITAMIN D3 AND FOLIC ACID TABLETS</b>	Each film coated tablets contains:			<b>APPROVED</b>
		Methylcobalamin	USP	1500 mcg	
		Alpha Lipoic Acid	USP	100 mg	
		Pyridoxine Hydrochloride	IP	3 mg	
		Vitamin D3	IP	1000 IU	
		Folic Acid	IP	1.5 mg	
		Excipients		q.s.	
		Approved colour used			

472	<b>MIRTAZAPINE TABLETS IP 30 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Mirtazapine	IP	30 mg	
		(As Hemihydrate)			
		Excipients		q.s.	
		Approved colour used			
473	<b>MIRTAZAPINE TABLETS IP 45 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Mirtazapine	IP	45 mg	
		(As Hemihydrate)			
		Excipients		q.s.	
		Approved colour used			
474	<b>CLOBAZAM TABLETS IP 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clobazam	IP	5 mg	
		Excipients		q.s.	
475	<b>CLOBAZAM TABLETS IP 10 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clobazam	IP	10 mg	
		Excipients		q.s.	
476	<b>CLOBAZAM TABLETS IP 20 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clobazam	IP	20 mg	
		Excipients		q.s.	
477	<b>TOLPERISONE HYDROCHLORIDE TABLETS 150 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tolperisone Hydrochloride		150 mg	
		Excipients		q.s.	
		Approved colour used			
478	<b>CITICOLINE TABLETS IP 250MG</b>	Each film coated tablet contains			<b>APPROVED</b>
		Citicoline Sodium	IP		
		Eq to Citicoline		250mg	
		Excipients		q.s.	
		Approved colour used			
479	<b>CITICOLINE TABLETS IP 500MG</b>	Each film coated tablet contains			<b>APPROVED</b>
		Citicoline Sodium	IP		
		eq. to Citicoline		500mg	
		Excipients		q.s.	
		Approved colour used			
480	<b>MODAFINIL TABLETS 100MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Modafinil	BP	100mg	
		Excipients		q.s.	
481	<b>OLANZAPINE TABLETS IP 10 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olanzapine	IP	10 mg	
		Excipients		q.s.	
		Approved colour used			
482	<b>VOGLIBOSE &amp; METFORMIN HYDROCHLORIDE TABLETS</b>	Each uncoated bilayered tablet			<b>APPROVED</b>
		contains:			
		Voglibose	IP	0.2 mg	
		Metformin Hydrochloride	IP	500 mg	
		(as sustained release form)			
		Excipients		q.s.	
		Approved colour used			
483	<b>METHYLPREDNISOLONE TABLETS IP 4 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Methylprednisolone	IP	4 mg	
		Excipients		q.s.	
		Approved colour used			



484	<b>METHYLPREDNISOLONE TABLETS 8MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Methylprednisolone	IP	8 mg	
		Excipients		q.s.	
		Approved colour used			
485	<b>METHYLPREDNISOLONE TABLETS IP 16 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Methylprednisolone	IP	16 mg	
		Excipients		q.s.	
		Approved colour used			
486	<b>PREGABALIN CAPSULES IP 50 MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Pregabalin	IP	50 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
487	<b>IMIPRAMINE HCL TABLETS IP 50MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Imipramine Hcl	IP	50mg	
		Excipients		qs	
		Approved colour used			
488	<b>EPERISONE HCL SR CAPSULES 150MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Eperisone Hcl		150mg	
		(as modified dosage form)			
		Excipients		qs	
		approved colour used in empty capsules shell			
489	<b>TRIHEXYPHENIDYL HCl TABLETS IP 5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Trihexyphenidyl Hydrochloride	IP	5 mg	
		Excipients		q.s.	
490	<b>FOLIC ACID TABLETS IP 5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Folic Acid	IP	5mg	
		Excipients		q.s.	
491	<b>CALCIUM CITRATE , CALCITRIOL &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium citrate	USP	1000mg	
		Calcitriol	IP	0.25 mcg	
		Zinc Sulphate Monohydrate	IP	17.6mg	
		Excipients		q.s.	
		Approved colour used			
492	<b>VENLAFAXINE EXTENDED RELEASE TABLETS 37.5 MG</b>	Each uncoated extended release tablet contains:			<b>APPROVED</b>
		venlafaxine Hcl	BP		
		eq to venlafaxine		37.5mg	
		Excipients		q.s.	
		Approved colour used			
493	<b>VENLAFAXINE EXTENDED RELEASE TABLETS 75 MG</b>	Each uncoated extended release tablet contains:			<b>APPROVED</b>
		venlafaxine Hcl	BP		
		eq to venlafaxine		75mg	
		Excipients		q.s.	
		Approved colour used			
494	<b>RISPERIDONE &amp; TRIHEXYPHENIDYL MOUTH DISSOLVING TABLETS</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Risperidone	BP	4 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	

495	<b>RISPERIDONE &amp; TRIHEXYPHENIDYL MOUTH DISSOLVING TABLETS</b>	Each uncoated mouth dissolving tablet contains:			<b>APPROVED</b>
		Risperidone	BP	2 mg	
		Trihexyphenidyl Hydrochloride	IP	2 mg	
		Excipients		q.s.	
496	<b>PROPRANOLOL HCL TABLETS 10MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Propranolol Hcl	IP	10mg	
		Excipients		q.s.	
497	<b>GABAPENTIN CAPSULES IP 400MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Gabapentin	IP	400 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shell			
498	<b>GABAPENTIN CAPSULES IP 300MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Gabapentin	IP	300 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shell			
499	<b>DICLOFENAC SODIUM TABLETS 50MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Diclofenac Sodium	IP	50 mg	
		Excipients		q.s.	
500	<b>ALBENDAZOLE TABLETS IP 400MG</b>	Each uncoated chewable tablet contains:			<b>APPROVED</b>
		Albendazole	IP	400 mg	
		Excipients		q.s.	
501	<b>PHENYTOIN TABLETS IP 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Phenytoin Sodium	IP	100 mg	
		Excipients		q.s.	
		Approved colour used			
502	<b>CALCIUM CARBONATE, VIT. D3 &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		calcium carbonate	I.P.		
		eq. to elemental calcium		500mg	
		Vitamin D3	I.P.	250IU	
		Zinc Sulphate Monohydrate	I.P.	61.8mg	
		Excipients		q.s.	
503	<b>PIRACETAM TABLETS 800 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Piracetam	I.P.	800mg	
		Excipients		q.s.	
		Approved colour used			
504	<b>ACECLOFENAC, PARACETAMOL &amp; CHLORZOXAZONE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aceclofenac	I.P.	100 mg	
		Chlorzoxazone	U.S.P.	250mg	
		Paracetamol	I.P.	325 mg	
		Excipients		q.s.	
505	<b>PREGABALIN (S.R.) AND NORTRIPTYLINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pregabalin	I.P.	75 mg	
		(as sustain release form)			
		Nortriptyline Hydrochloride	I.P.		
		eq. to Nortriptyline		10 mg	
		Excipients		q.s.	
505		Approved colour used			

506	<b>VOGLIBOSE &amp; METFORMIN HYDROCHLORIDE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Voglibose	I.P.	0.3 mg	
		Metformin Hydrochloride	I.P.	500 mg	
		Excipients		q.s.	
		Approved colour used			
507	<b>BACLOFEN EXTENDED RELEASE TABLETS 30MG</b>	Each film coated extended release tablet contains:			<b>APPROVED</b>
		Baclofen	I.P.	30 mg	
		Excipients		q.s.	
		Approved colour used			
508	<b>PIRACETAM TABLETS 400 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Piracetam	I.P.	400mg	
		Excipients		q.s.	
		Approved colour used			
509	<b>PIRACETAM TABLETS 1200 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Piracetam	I.P.	1200mg	
		Excipients		q.s.	
		Approved colour used			
510	<b>HALOPERIDOL DISPERSIBLE TABLETS 0.25 MG</b>	Each uncoated dispersible tablet contains:			<b>APPROVED</b>
		Haloperidol	I.P.	0.25 mg	
		Excipients		q.s.	
511	<b>HALOPERIDOL DISPERSIBLE TABLETS 10MG</b>	Each uncoated dispersible tablet contains:			<b>APPROVED</b>
		Haloperidol	I.P.	10 mg	
		Excipients		q.s.	
512	<b>FLUOXETINE TABLETS 60MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fluoxetine Hydrochloride	I.P.		
		eq. to fluoxetine		60mg	
		Excipients		q.s.	
		Approved colour used			
513	<b>CALCIUM CARBONATE, VIT. D3 &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		calcium carbonate	I.P.		
		eq. to elemental calcium		500mg	
		Vitamin D3	I.P.	250IU	
		Zinc Sulphate Monohydrate	I.P.		
		eq to elemental Zinc		4mg	
		Excipients		q.s.	
		Approved colour used			
514	<b>CLOBAZAM TABLETS IP 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clobazam	I.P.	5 mg	
		Excipients		q.s.	
		Approved colour used			
515	<b>CLOBAZAM TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clobazam	I.P.	20 mg	
		Excipients		q.s.	
		Approved colour used			
517	<b>GABAPENTIN TABLETS IP 400 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Gabapentin	I.P.	400 mg	
		Excipients		q.s.	
		Approved colour used			

518	<b>FLUVOXAMINE SR TABLETS 100MG</b>	Each film coated sustained release			<b>APPROVED</b>
		tablet contains:			
		Fluvoxamine maleate	IP	100 mg	
		Excipients		q.s.	
		Approved colour used			
519	<b>CLIDINIUM BROMIDE , CHLORDIAZEPOXIDE&amp; DICYCLOMINE HCI TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clidinium Bromide	USP	2.5mg	
		Chlordiazepoxide	I.P.	5mg	
		Dicyclomine HCl	I.P.	10mg	
		Excipients		q.s.	
		Approved colour used			
520	<b>RISPERIDONE MOUTH DISSOLVING TABLETS 1 MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Risperidone	B.P.	1 mg	
		Excipients		q.s.	
521	<b>ROSUVASTATIN TABLETS IP 20MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium	I.P.		
		eq. to Rosuvastatin		20 mg	
		Excipients		q.s.	
		Approved colour used			
522	<b>ATORVASTATIN &amp; FENOFIBRATE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Atorvastatin Calcium	I.P.		
		eq. to Atorvastatin		10 mg	
		Fenofibrate	I.P.	160 mg	
		Excipients		q.s.	
		Approved colour used			
523	<b>OMEPRAZOLE (EC) &amp; DOMPERIDONE (SR) CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Omeprazole	I.P.	20 mg	
		(As enteric coated pellets)			
		Domperidone	I.P.	30 mg	
		(As sustained release pellets)			
		Excipients		q.s.	
		Approved colour used in empty			
		capsule Shells			
524	<b>TADALAFIL TABLETS IP 10MG</b>	Each Film coated Tablet contains:			<b>APPROVED</b>
		Tadalafil	I.P.	10mg	
		Excipients		qs	
		Approved colour used			
525	<b>PIROXICAM TABLETS 10MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Piroxicam	I.P.	10mg	
		Excipients		q.s.	
526	<b>TELMISARTAN &amp; CHLORTHALIDONE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	I.P.	40mg	
		Chlorthalidone	I.P.	12.5mg	
		Excipients		q.s.	
		Approved colour used			
527	<b>TELMISARTAN &amp; CHLORTHALIDONE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	I.P.	80mg	
		Chlorthalidone	I.P.	12.5mg	
		Excipients		q.s.	
		Approved colour used			

528	<b>OLMESARTAN MEDOXOMIL &amp; CHLORTHALIDONE TABLETS</b>	Each Film coated Tablet contains:			<b>APPROVED</b>
		Olmesartan Medoxomil	I.P.	20mg	
		Chlorthalidone	I.P.	12.5mg	
		Excipients		q.s.	
		Approved colour used			
529	<b>OLMESARTAN MEDOXOMIL &amp; CHLORTHALIDONE TABLETS</b>	Each Film coated Tablet contains:			<b>APPROVED</b>
		Olmesartan Medoxomil	I.P.	40mg	
		Chlorthalidone	I.P.	12.5mg	
		Excipients		q.s.	
		Approved colour used			
530	<b>PHENYLEPHRINE HCl , CAFFEINE , DIPHENHYDRAMINE HYDROCHLORIDE &amp;PARACETAMOL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Phenylephrine Hcl	I.P.	5 mg	
		Caffeine (anhydrous)	I.P.	30mg	
		Diphenhydramine Hydrochloride	I.P.	25mg	
		Paracetamol	I.P.	500 mg	
		Excipients		q.s.	
		Approved colour used			
531	<b>ACETAMINOPHEN, DICLOFENAC POTASSIUM&amp; CHLORZOXAZONE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Acetaminophen	I.P.	325mg	
		Diclofenac Potassium	B.P.	50mg	
		Chlorzoxazone	U.S.P.	250mg	
		Excipients		q.s.	
		Approved colour used			
532	<b>PARACETAMOL &amp; DICYCLOMINE HCl TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	I.P.	500mg	
		Dicyclomine Hydrochloride	I.P.	20mg	
		Excipients		q.s.	
		Approved colour used			
533	<b>GLICLAZIDE SUSTAINED RELEASE TABLETS 30MG</b>	Each uncoated sustained release tablet contains:			<b>APPROVED</b>
		Gliclazide	I.P.	30mg	
		Excipients		q.s.	
		Approved colour used			
534	<b>GLICLAZIDE SUSTAINED RELEASE TABLETS 60MG</b>	Each uncoated sustained release tablet contains:			<b>APPROVED</b>
		Gliclazide	I.P.	60mg	
		Excipients		q.s.	
		Approved colour used			
535	<b>TERBINAFINE TABLETS 500 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Terbinafine Hydrochloride	B.P.		
		eq. to Terbinafine		500mg	
		Excipients		qs	
536	<b>CLONAZEPAM TABLETS IP 2 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Clonazepam	IP	2 mg	
		Excipients		q.s.	
537	<b>DIVALPROEX EXTENDED RELEASE TABLETS IP 750MG</b>	Each film coated extended release tablet contains:			<b>APPROVED</b>
		Divalproex Sodium	I.P.		
		eq. to Valproic Acid		750 mg	
		Excipients		q.s.	
		Approved colour used			

538	<b>DONEPEZIL TABLETS IP 10MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Donepezil Hydrochloride	I.P.	10mg	
		Excipients		q.s.	
		Approved colour used			
539	<b>MIRTAZAPINE TABLETS IP 15 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Mirtazapine	I.P.	15 mg	
		Excipients		q.s.	
		Approved colour used			
540	<b>MIRTAZAPINE TABLETS IP 30 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Mirtazapine	I.P.	30 mg	
		Excipients		q.s.	
		Approved colour used			
541	<b>PREGABALIN SUSTAINED RELEASE TABLETS 75 MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Pregabalin	I.P.	75 mg	
		Excipients		q.s.	
		Approved colour used			
542	<b>PHENYTOIN SODIUM EXTENDED RELEASE TABLETS 300MG</b>	Each film coated extended release tablet contains:			<b>APPROVED</b>
		Phenytoin Sodium	I.P.	300 mg	
		Excipients		q.s.	
		Approved colour used			
543	<b>ALPRAZOLAM TABLETS IP 1 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Alprazolam	IP	1mg	
		Excipients		q.s.	
		Approved colour used			
544	<b>TOPIRAMATE TABLETS IP 100MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Topiramate	IP	100 mg	
		Excipients		q.s.	
		Approved colour used			
545	<b>LEVETIRACETAM SUSTAINED RELEASE TABLETS 500 MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	500 mg	
		Excipients		q.s.	
		Approved colour used			
546	<b>LEVETIRACETAM SUSTAINED RELEASE TABLETS 750 MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	750 mg	
		Excipients		q.s.	
		Approved colour used			
547	<b>LEVETIRACETAM SUSTAINED RELEASE TABLETS 1000 MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	1000 mg	
		Excipients		q.s.	
		Approved colour used			
548	<b>LEVETIRACETAM TABLETS 1000 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levetiracetam	IP	1000 mg	
		Excipients		q.s.	
		Approved colour used			
549	<b>FLUVOXAMINE SUSTAINED RELEASE TABLETS 50MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Fluvoxamine maleate	I.P.	50 mg	
		Excipients		q.s.	
		Approved colour used			

550	<b>CLOBAZAM MOUTH DISSOLVING TABLETS 5MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Clobazam	I.P.	5 mg	
		Excipients		q.s.	
551	<b>ARIPIRAZOLE MOUTH DISSOLVING TABLETS 15 MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Aripiprazole	IP	15mg	
		Excipients		q.s.	
552	<b>RISPERIDONE MOUTH DISSOLVING TABLETS 3MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Risperidone	B.P.	3 mg	
		Excipients		q.s.	
		Approved colour used			
553	<b>RISPERIDONE MOUTH DISSOLVING TABLETS 0.5MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Risperidone	B.P.	0.5 mg	
		Excipients		q.s.	
		Approved colour used			
554	<b>OXCARBAZEPINE DISPERSIBLE TABLETS 150MG</b>	Each uncoated dispersible tablet			<b>APPROVED</b>
		contains:			
		Oxcarbazepine	IP	150 mg	
		Excipients		q.s.	
		Approved colour used			
555	<b>OXCARBAZEPINE DISPERSIBLE TABLETS 300MG</b>	Each uncoated dispersible tablet			<b>APPROVED</b>
		contains:			
		Oxcarbazepine	IP	300 mg	
		Excipients		q.s.	
		Approved colour used			
556	<b>ETIZOLAM MOUTH DISSOLVING TABLETS 0.25 MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Etizolam		0.25 mg	<b>APPROVED</b>
		Excipients		q.s.	
557	<b>ETIZOLAM MOUTH DISSOLVING TABLETS 0.5 MG</b>	Each uncoated mouth dissolving			<b>APPROVED</b>
		tablet contains:			
		Etizolam		0.5 mg	<b>APPROVED</b>
		Excipients		q.s.	
558	<b>TORSEMIDE TABLETS IP 10 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Torsemide	I.P.	10 mg	
		Excipients		q.s.	
559	<b>TORSEMIDE TABLETS IP 20 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Torsemide	I.P.	20 mg	
		Excipients		q.s.	
560	<b>PREGABALIN (S.R.) &amp; METHYLCOBALAMIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pregabalin (S.R.)	I.P.	75 mg	
		Methylcobalamin	U.S.P.	750mcg	
		Excipients		q.s.	
		Approved colour used			
561	<b>FLUPENTIXOL TABLETS 1MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Flupentixol Hydrochloride	B.P.		
		eq. to Flupentixol		1 mg	
		Excipients		q.s.	
		Approved colour used			



562	ESCITALOPRAM & ETIZOLAM TABLETS	Each film coated tablet contains:			APPROVED
		Escitalopram Oxalate	I.P.		
		eq. to Escitalopram		5 mg	
		Etizolam		0.5 mg	
		Excipients		q.s.	
		Approved colour used			
563	ESCITALOPRAM & ETIZOLAM TABLETS	Each film coated tablet contains:			APPROVED
		Escitalopram Oxalate	I.P.		
		eq. to Escitalopram		10 mg	
		Etizolam		0.5 mg	
		Excipients		q.s.	
		Approved colour used			
564	ETIZOLAM & PROPRANOLOL HCl TABLETS	Each film coated tablet contains:			APPROVED
		propranolol Hydrochloride	I.P.	20 mg	
		Etizolam		0.5 mg	
		Excipients		q.s.	
		Approved colour used			
565	IMIPRAMINE HYDROCHLORIDE & CHLORDIAZEPOXIDE TABLETS	Each film coated tablet contains:			APPROVED
		Imipramine Hydrochloride	I.P.	25mg	
		Chlordiazepoxide	I.P.	10mg	
		Excipients		q.s.	
		Approved colour used			
566	PROCHLORPERAZINE MALEATE TABLETS IP 5 MG	Each uncoated tablet contains:			APPROVED
		Prochlorperazine Maleate	I.P.	5 mg	
		Excipients		q.s.	
567	PROCHLORPERAZINE MALEATE MOUTH DISSOLVING TABLETS 5 MG	Each uncoated mouth dissolving tablet contains:			APPROVED
		Prochlorperazine Maleate	I.P.	5 mg	
		Excipients		q.s.	
568	FERROUS ASCORBATE, FOLIC ACID & ZINC TABLETS	Each film coated tablet contains:			APPROVED
		Ferrous Ascorbate			
		eq. to elemental Iron		100 mg	
		Folic Acid	I.P.	1.5 mg	
		Zinc Sulphate Monohydrate	I.P.		
		eq. to elemental Zinc		22.5 mg	
		Excipients		q.s.	
		Approved colour used			
569	TRIFLUOPERAZINE TABLETS IP 5 MG	Each film coated tablet contains:			APPROVED
		Trifluoperazine Hydrochloride	I.P.		
		eq. to Trifluoperazine		5 mg	
		Excipients		q.s.	
		Approved colour used			
570	DESLORATADINE TABLETS 5 MG	Each film coated tablet contains:			APPROVED
		Desloratadine		5 mg	
		Excipients		q.s.	
		Approved colour used			
571	OLANZAPINE TABLETS IP 2.5 MG	Each uncoated tablet contains:			APPROVED
		Olanzapine	I.P.	2.5 mg	
		Excipients		q.s.	
572	OLANZAPINE TABLETS IP 7.5 MG	Each uncoated tablet contains:			APPROVED
		Olanzapine	I.P.	7.5 mg	
		Excipients		q.s.	

573	<b>OLANZAPINE TABLETS IP 15 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Olanzapine	I.P.	15 mg	
		Excipients		q.s.	
574	<b>ARIPIRAZOLE TABLETS IP 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Aripiprazole	I.P.	5mg	
		Excipients		q.s.	
575	<b>OFLOXACIN &amp; ORNIDAZOLE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ofloxacin	I.P.	200mg	
		Ornidazole	I.P.	500mg	
		Excipients		q.s.	
		Approved colour used			
576	<b>ESOMEPRAZOLE GASTRO RESISTANT TABLETS IP 40MG</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Esomeprazole Magnesium Trihydrate	IP		
		eq. to Esomeprazole		40 mg	
		Excipients		q.s.	
		Approved colour used			
577	<b>QUETIAPINE SUSTAINED RELEASE TABLETS 50 MG</b>	Each film coated sustained release tablet contains:			<b>APPROVED</b>
		Quetiapine Fumatate	I.P.		
		eq. to Quetiapine		50 mg	
		Excipients		q.s.	
		Approved colour used			
578	<b>PARACETAMOL, CAFFEINE, CPM&amp;PHENYLEPHRINE HCL TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	I.P.	325 mg	
		Caffeine (Anhydrous)	I.P.	30mg	
		Chlorpheniramine Maleate	I.P.	4mg	
		Phenylephrine Hydrochloride	I.P.	5 mg	
		Excipients		q.s.	
579	<b>PARACETAMOL, CAFFEINE, IBUPROFEN TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	I.P.	325 mg	
		Caffeine (Anhydrous)	I.P.	25mg	
		Ibuprofen	I.P.	400mg	
		Excipients		q.s.	
580	<b>NIMESULIDE, PARACETAMOL, CAFFEINE, CETIRIZINE HCL &amp;PHENYLEPHRINE HCL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Nimesulide	B.P.	100mg	
		Caffeine (Anhydrous)	I.P.	30mg	
		paracetamol	I.P.	325mg	
		Cetirizine Hydrochloride	I.P.	5 mg	
		phenylephrine Hydrochloride	I.P.	10 mg	
		Excipients		q.s.	
581	<b>PARACETAMOL, CAFFEINE, LEVOCETIRIZINE HCL &amp;PHENYLEPHRINE HCL TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Caffeine (Anhydrous)	I.P.	30mg	
		paracetamol	I.P.	325mg	
		Levocetirizine Hydrochloride	I.P.	2.5mg	
		Phenylephrine Hydrochloride	I.P.	10 mg	
		Excipients		q.s.	
		Approved colour used			

582	<b>GLIMEPIRIDE, PIOGLITAZONE &amp; METFORMIN HYDROCHLORIDE (ER) TABLETS</b>	Each uncoated bilayered tablet			<b>APPROVED</b>
		contains:			
		Glimepiride	I.P.	1 mg	
		Pioglitazone Hcl	I.P.		
		eq. to Pioglitazone		15mg	
		Metformin Hydrochloride (as extended release )	I.P.	500 mg	
		Excipients		q.s.	
		Approved colour used			
583	<b>GLIMEPIRIDE, PIOGLITAZONE &amp; METFORMIN HYDROCHLORIDE (ER) TABLETS</b>	Each uncoated bilayered tablet			<b>APPROVED</b>
		contains:			
		Glimepiride	I.P.	2mg	
		Pioglitazone Hcl	I.P.		
		eq. to Pioglitazone		15mg	
		Metformin Hydrochloride (as extended release )	I.P.	500 mg	
		Excipients		q.s.	
		Approved colour used			
584	<b>NIMESULIDE &amp; TIZANIDINE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Nimesulide	B.P.	100mg	
		Tizanidine Hcl	I.P.		
		eq. to Tizanidine		2 mg	
		Excipients		q.s.	
585	<b>CLOPIDOGREL &amp;ASPIRIN TABLETS</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Clopidogrel Bisulphate	I.P.		
		eq. to Clopidogrel		75 mg	
		Aspirin	I.P.	150 mg	
		Excipients		q.s.	
		Approved colour used			
586	<b>GABAPENTIN &amp; NORTRIPTYLINE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Gabapentin	I.P.	400 mg	
		Nortriptyline Hydrochloride	I.P.		
		eq. to Nortriptyline		10mg	
		Excipients		q.s.	
		Approved colour used			
587	<b>RISPERIDONE TABLETS 0.5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Risperidone	B.P.	0.5 mg	
		Excipients		q.s.	
588	<b>PAROXETINE (C.R.) &amp; CLONAZEPAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Paroxetine Hydrochloride	I.P.		
		eq. to Paroxetine (C.R.)		25mg	
		Clonazepam	I.P.	0.5mg	
		Excipients		q.s.	
		Approved colour used			
589	<b>PARACETAMOL, PHENYLEPHRINE HCl, CETIRIZINE HCl &amp; DEXTROMETHORPHAN HBr TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	I.P.	325 mg	
		Phenylephrine Hydrochloride	I.P.	5 mg	
		Cetirizine Hydrochloride	I.P.	10mg	
		Dextromethorphan Hydrobromide	I.P.	10mg	
		Excipients		q.s.	
		Approved colour used			

590	<b>OLMESARTAN MEDOXOMIL , CHLORTHALIDONE &amp; AMLODIPINE TABLETS</b>	Each Film coated Tablet contains:			<b>APPROVED</b>
		Olmесartan Medoxomil	I.P.	20mg	
		Chlorthalidone	I.P.	12.5mg	
		Amlodipine Besilate	I.P.		
		eq. to Amlodipine		5mg	
		Excipients		q.s.	
		Approved colour used			
591	<b>TELMISARTAN , CHLORTHALIDONE &amp; AMLODIPINE TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Telmisartan	I.P.	40mg	
		Chlorthalidone	I.P.	12.5mg	
		Amlodipine Besilate	I.P.		
		eq. to Amlodipine		5mg	
		Excipients		q.s.	
		Approved colour used			
592	<b>FLUPENTIXOL TABLETS BP 3MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Flupentixol Hydrochloride	B.P.		
		eq. to Flupentixol		3 mg	
		Excipients		q.s.	
		Approved colour used			
593	<b>FLUOXETINE HYDROCHLORIDE &amp; ALPRAZOLAM TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fluoxetine Hydrochloride	I.P.		
		eq. to Fluoxetine		20 mg	
		Alprazolam	I.P.	0.25 mg	
		Excipients		q.s.	
		Approved colour used			
594	<b>ETIZOLAM &amp;PROPRANOLOL HCl TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		propranolol Hydrochloride	I.P.	20 mg	
		Etizolam		0.5 mg	
		Excipients		q.s.	
		Approved colour used			
595	<b>IMIPRAMINE HYDROCHLORIDE &amp; CHLORDIAZEPOXIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Imipramine Hydrochloride	I.P.	25mg	
		Chlordiazepoxide	I.P.	10mg	
		Excipients		q.s.	
		Approved colour used			
596	<b>PARACETAMOL, CHLORPHENIRAMINE MALEATE, CAFFEINE &amp; PHENYLEPHRINE HCl TABLETS</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	I.P.	500mg	
		Caffeine (anhydrous)	I.P.	30mg	
		Chlorpheniramine Maleate	I.P.	2mg	
		Phenylephrine Hydrochloride	I.P.	5mg	
		Excipients		q.s.	
		Approved colour used			
597	<b>NORTRIPTYLINE TABLETS 75 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Nortriptyline Hydrochloride	I.P.		
		eq. to Nortriptyline		75mg	
		Excipients		qs	
		Approved colour used			
598	<b>DOTHIEPIN HCL TABLETS 75 MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Dothiepin Hcl	I.P.	75mg	
		Excipients		q.s.	
		Approved colour used			

599	<b>SODIUM VAPROATE &amp; VALPROIC ACID TABLETS</b>	Each film coated controlled release			<b>APPROVED</b>
		tablet contains:			
		Sodium Valproate	I.P.	333 mg	
		Valproic Acid	I.P.	145 mg	
		(Both together corresponds to			
		Sodium Valproate IP 500 mg)			
		Excipients		q.s.	
		Approved colour used			
600	<b>HALOPERIDOL TABLETS IP 1.5MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Haloperidol	I.P.	1.5mg	
		Excipients		q.s.	
601	<b>FLUNARIZINE TABLETS 5 MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Flunarizine Dihydrochloride	B.P.		
		eq. to Flunarizine		5 mg	
		Excipients		q.s.	
602	<b>DEFLAZACORT TABLETS 30MG</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Deflazacort		30 mg	
		Excipients		qs	
603	<b>GLICLAZIDE (M.R.) &amp; METFORMIN HYDROCHLORIDE (E.R.) TABLETS</b>	Each uncoated bilayered tablet contains			<b>APPROVED</b>
		Gliclazide (as M.R.)	I.P.	60 mg	
		Metformin Hydrochloride	I.P.	500 mg	
		(as E.R.)			
		Excipients		q.s.	
		Approved colour used			
604	<b>GLIBENCLAMIDE &amp; METFORMIN HYDROCHLORIDE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Glibenclamide	I.P.	5mg	
		Metformin Hydrochloride	I.P.	500 mg	
		Excipients		q.s.	
		Approved colour used			
605	<b>ESCITALOPRAM TABLETS IP 5MG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Escitalopram Oxalate	I.P.		
		eq. to Escitalopram		5mg	
		Excipients		qs	
		Approved colour used			
606	<b>PIOGLITAZONE &amp; METFORMIN HCL(SR) TABLETS</b>	Each uncoated bilayered tablet			<b>APPROVED</b>
		contains:			
		Pioglitazone hydrochloride	I.P.		
		eq. to Pioglitazone		15mg	
		Metformin HCl	I.P.	500 mg	
		(as Sustained Release form)			
		Excipients		q.s.	
		Approved colour used			
607	<b>GABAPENTIN &amp; METHYLCOBALAMIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Gabapentin	I.P.	300 mg	
		Methylcobalamin	U.S.P.	500mcg	
		Excipients		q.s.	
		Approved colour used			
608	<b>FERROUS ASCORBATE, FOLIC ACID &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ferrous Ascorbate		100 mg	
		Folic Acid	I.P.	1.5 mg	
		Zinc Sulphate Monohydrate	I.P.		
		eq. to elemental Zinc		22.5 mg	
		Excipients		q.s.	
		Approved colour used			

609	<b>CALCIUM CARBONATE, VIT. D3 &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		calcium carbonate	I.P.		
		eq. to elemental calcium		500mg	
		Vitamin D3	I.P.	250IU	
		Zinc Sulphate Monohydrate	I.P.	61.8mg	
		Excipients		q.s.	
		Approved colour used			
610	<b>CALCIUM CARBONATE &amp; VIT. D3 TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium carbonate	I.P.		
		eq. to elemental calcium		500mg	
		Vitamin D3	I.P.	250IU	
		Excipients		q.s.	
		Approved colour used			
611	<b>DOXYLAMINE SUCCINATE, PYRIDOXINE HCL &amp; FOLIC ACID TABLETS</b>	Each enteric coated tablet contains:			<b>APPROVED</b>
		Doxylamine Succinate	B.P.	10mg	
		Pyridoxine Hydrochloride	I.P.	10mg	
		Folic Acid	I.P.	2.5mg	
		Excipients		q.s.	
		Approved colour used			
612	<b>METHYLCOBALAMIN, ALPHA LIPOIC ACID, PYRIDOXINE HCI AND FOLIC ACID TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Methylcobalamin	U.S.P.	1500 mcg	
		Alpha Lipoic Acid	U.S.P.	100 mg	
		Pyridoxine Hydrochloride	I.P.	3 mg	
		Folic Acid	I.P.	1.5 mg	
		Excipients		q.s.	
		Approved colour used			
613	<b>METHYLCOBALAMIN TABLETS 500 MCG</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Methylcobalamin	U.S.P.	500mcg	
		Excipients		q.s.	
		Approved colour used			
617	<b>ACECLOFENAC, PARACETAMOL &amp; SERRATIOPEPTIDASE TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Aceclofenac	I.P.	100 mg	
		Serratiopeptidase	I.P.	15mg	
		(30,000 ENZYME UNIT)			
		Paracetamol	I.P.	325 mg	
		Excipients		q.s.	
		Approved colour used			
618	<b>CALCIUM CITRATE, MAGNESIUM ,VIT D3 &amp; ZINC TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Calcium Citrate	U.S.P.	1000 mg	
		Magnesium Hydroxide	I.P.		
		eq. to Magnesium		100 mg	
		Vit. D3	I.P.	250 IU	
		Zinc Sulphate Monohydrate	I.P.		
		eq. to elemental Zinc		4mg	
		Excipients		q.s.	
		Approved colour used			
619	<b>PREGABALIN (S.R.) &amp; METHYLCOBALAMIN TABLETS</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Pregabalin (S.R.)	I.P.	75 mg	
		Methylcobalamin	U.S.P.	1500mcg	
		Excipients		q.s.	
		Approved colour used			

620	LOXAPINE CAPSULES 10MG	Each hard gelatin capsule contains:			APPROVED
		Loxapine Succinate	U.S.P.		
		eq. to Loxapine		10 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shell.			
621	LOXAPINE CAPSULES 25MG	Each hard gelatin capsule contains:			APPROVED
		Loxapine Succinate	U.S.P.		
		eq. to Loxapine		25 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shell.			
622	RABEPRAZOLE SODIUM (EC) & DOMPERIDONE (SR) CAPSULES	Each hard gelatin capsule contains:			APPROVED
		Rabeprazole Sodium	I.P.	20 mg	
		(As enteric coated pellets)			
		Domperidone	I.P.	30mg	
		(As sustained release pellets)			
		Excipients		q.s.	
		Approved colour used in empty capsule Shells			
623	FLUOXETINE CAPSULES IP 20 MG	Each hard gelatin capsule contains:			APPROVED
		Fluoxetine Hydrochloride	I.P.		
		eq. to Fluoxetine		20 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			
624	FLUOXETINE CAPSULES 60 MG	Each hard gelatin capsule contains:			APPROVED
		Fluoxetine Hydrochloride	I.P.		
		eq. to Fluoxetine		60 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			
625	RABEPRAZOLE SODIUM (EC) & DOMPERIDONE CAPSULES	Each hard gelatin capsule contains:			APPROVED
		Rabeprazole Sodium	I.P.	20 mg	
		(As enteric coated pellets)			
		Domperidone	I.P.	10 mg	
		(As pellets)			
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
626	FLUOXETINE CAPSULES IP 40 MG	Each hard gelatin capsule contains:			APPROVED
		Fluoxetine Hydrochloride	I.P.		
		eq. to Fluoxetine		40 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells			
627	PREGABALIN CAPSULES IP 75 MG	Each hard gelatin capsule contains:			APPROVED
		Pregabalin	I.P.	75 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells			



628	<b>PROPRANOLOL HCL(S.R) &amp; FLUNARIZINE CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Propranolol HCl	I.P.	40 mg	
		(As Sustened Release Pellets)			
		Flunarizine Dihydrochloride	B.P.		
		eq. to Flunarizine		10 mg	
		Approved colour used in empty capsule shells			
629	<b>OMEPRAZOLE GASTRO RESISTANT CAPSULES IP 20MG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Omeprazole	I.P.	20mg	
		(as enteric coated pellets)			
		Excipients		q.s.	
		Approved colour used in empty capsule shells			
630	<b>CARBONYL IRON , FOLIC ACID, CYANOCOBALAMIN &amp; ZINC CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Carbonyl Iron eq. to Elemental Iron		100 mg	
		Folic Acid	I.P.	1.5 mg	
		Cyanocobalamin	I.P.	15 mcg	
		Zinc Sulphate Monohydrate	I.P.	61.8 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			
631	<b>FLAVONOL GLYCOSIDES CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Ginkgo Biloba (50:1)	B.P.	60 mg	
		(Containing 14.4mg of Flavonol Glycosides)			
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			
632	<b>ALPHA LIPOIC ACID , BENFOTIAMINE, CHROMIUM PICOTINATE, METHYLCOBALAMIN &amp; INOSITOL CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Methylcobalamin	U.S.P.	1500 mcg	
		Alpha Lipoic Acid	U.S.P.	100 mg	
		Benfotiamine		150 mg	
		Chromium Polynicotinate	U.S.P.	200 mcg	
		Inositol	B.P.	150 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shells			
633	<b>GABAPENTIN &amp; METHYLCOBALAMIN CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Gabapentin	I.P.	300 mg	
		Methylcobalamin	U.S.P.	750mcg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells			
634	<b>GABAPENTIN &amp; METHYLCOBALAMIN CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Gabapentin	I.P.	400 mg	
		Methylcobalamin	U.S.P.	750mcg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells			
635	<b>ALFACALCIDOL CAPSULES 0.25 MCG</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Alfacalcidol	I.P.	0.25mcg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			

638	<b>PREGABALIN &amp; METHYLCOBALAMIN CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Pregabalin	I.P.	75 mg	
		Methylcobalamin	U.S.P.	750mcg	
		Excipients		q.s.	
		Approved colour used in empty capsule Shell			
639	<b>B-COMPLEX FORTE WITH VITAMIN C CAPSULES</b>	Each hard gelatin capsules contains:			<b>APPROVED</b>
		Thiamine mononitrate	I.P.	10mg	
		Riboflavin	I.P.	10mg	
		Pyridoxine Hcl	I.P.	3mg	
		Vitamin B12	I.P.	15mcg	
		Niacinamide	I.P.	100mg	
		Calcium pantothenate	I.P.	50mg	
		Folic acid	I.P.	1.5mg	
		Biotin	U.S.P.	100mcg	
		Ascorbic acid(coated)	I.P.	150mg	
		approved colour used in empty capsule shell.			
640	<b>CARBONYL IRON , FOLIC ACID &amp; ZINC CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Carbonyl Iron eq. to Elemental Iron		50 mg	
		Folic Acid	I.P.	0.5 mg	
		Zinc Sulphate Monohydrate	I.P.	61.8 mg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			
641	<b>PREGABALIN &amp; METHYLCOBALAMIN CAPSULES</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Pregabalin	I.P.	150 mg	
		Methylcobalamin	U.S.P.	750mcg	
		Excipients		q.s.	
		Approved colour used in empty capsule shells.			
642	<b>Ambroxol &amp; Cetirizine Hcl yrup</b>	Each 5ml contains :-			<b>APPROVED</b>
		Ambroxol	IP	30mg	
		Cetirizine Hcl	IP	10mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colour used			
643	<b>Ondansetron Oral Solution</b>	Each 5 ml contains			<b>APPROVED</b>
		Ondansetron Hydrochloride eq. to Ondansetron	IP	2mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colour used			
644	<b>Ambroxol, levosalbutamol &amp; Guaifenesin Syrup</b>	Each 5ml contains :-			<b>APPROVED</b>
		Ambroxol Hcl	IP	30mg	
		Levosaltbutamol Sulphate	IP	1mg	
		Guaifenesin	IP	15mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colour used			
645	<b>Albendazole Oral Suspension IP</b>	Each 5ml contains :-			<b>APPROVED</b>
		Albendazole	IP	200 mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colour used			

646	<b>Ambroxol HCL, Guaiphenesin, Terbutaline &amp; Menthol Syrup</b>	Each 5 ml contains :-			<b>APPROVED</b>
		Ambroxol HCL	IP	30mg	
		Guaiphenesin	IP	100mg	
		Terbutaline Sulphate	IP	2.5mg	
		Menthol	IP	2.5mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colour used			
647	<b>Mecobalamin, Pyridoxine, Nicotinamide,&amp; Benfotiamine Tablets</b>	Each film coated tablet contains :			<b>APPROVED</b>
		Mecobalamin	U.S.P.	750mcg	
		Pyridoxine	I.P.	1.5mg	
		Nicotinamide	I.P.	45mg	
		Benfotiamine		150mg	
		Excipients		q.s	
		Approved colour used in coating.			
648	<b>Rabeprazole Sodium(EC) &amp; Itopride(SR) Capsules</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Rabeprazole Sodium (As enteric Coated pellets )	I.P.	20mg	
		Itopride HCl (As sustained release pellets )		150mg	
		Excipients		q.s	
		Approved colours used in empty			
649	<b>Itraconazole Capsules</b>	Each hard gelatin capsules contains:			<b>APPROVED</b>
		Itraconazole Pellets Eq. to Itraconazole	B.P.	100mg	
		Excipients		q.s	
		Approved colour used in empty			
650	<b>Esomeprazole Magnesium (EC) &amp; Itopride HCl (SR) Capsules</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (As enteric Coated pellets )	I.P.	40mg	
		Itopride HCl (As sustained release pellets )		150mg	
		Excipients		q.s	
		Approved colours used in empty			
651	<b>Chloramphenicol Capsules IP</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Chloramphenicol	I.P.	500 mg	
		Excipients		q.s	
		Approved colours used in empty			
652	<b>Iron Amino Acid ,Folic Acid, Vit-B12 &amp; Zinc Picolinate SYRUP</b>	Each 5ml contains :-			<b>APPROVED</b>
		Iron Amino Acid Chelate AS Ferrous Bisglycinate eq. to elemental iron		15mg	
		Folic Acid	IP	0.25mg	
		Vit-B12	IP	3.75mg	
		Zinc Picolinate eq. to Elemental Zinc	BP	5.5mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colours used			

653	<b>Elemental Iron , Vit-B12 &amp; Vit-B6 SYRUP</b>	Each 5ml contains :-			<b>APPROVED</b>
		Elemental Iron in a Complex of iron polymatose aq. Solution		50mg	
		Vit-B12	IP	5mcg	
		Vit-B6	IP	2mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colours used			
654	<b>Dicyclomine, Dried Aluminium hydroxide,Light magnesium oxide &amp; Simethicone Suspension</b>	Each 5ml contains :-			<b>APPROVED</b>
		Dicyclomine	IP	2.5mg	
		Dried Aluminium hydroxide	USP	200mg	
		Light magnesium oxide	IP	100mg	
		Simethicone	USP	20mg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colours used			
655	<b>Ferrous Ascorbate &amp; Folic acid Suspension</b>	Each 5ml contains :-			<b>APPROVED</b>
		Ferrous Ascorbate Eq. to elemental Iron		30mg	
		Folic acid	IP	550mcg	
		Excipients		q.s.	
		A Flavoured Syrupy base			
		Approved colours used			
657	<b>Pioglitazone Tablets I.P.</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Pioglitazone Hydrochloride eq. to Pioglitazone	I.P.	30mg	
		Excipients		q.s	
		Colour: approved colour used.			
658	<b>Paracetamol, Phenylephrine HCL, Caffeine, Diphenhydramine HCL Tablets</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Paracetamol	I.P.	500mg	
		Phenylephrine HCL	I.P.	5mg	
		Caffeine (anhydrous)	I.P.	30mg	
		Diphenhydramine HCL	I.P.	25mg	
		Excipients		q.s	
		Colour : approved colour used.			
659	<b>Benfotiamine, Pyridoxine HCL, Methylcobalamin, Folic Acid, &amp; Pregabalin Capsules</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Benfotiamine		7.5mg	
		Pyridoxine Hydrochloride	I.P.	1.5mg	
		Methylcobalamin	U.S.P.	750mcg	
		Folic Acid	I.P.	0.75mg	
		Pregabalin	I.P.	150mg	
		Excipients		q.s	
		Approved colour used in empty			
660	<b>Levocetirizine &amp; Ambroxol HCL Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Levocetirizine Dihydrochloride	I.P.	5mg	
		Ambroxol HCL (In sustained released form)	I.P.	75mg	
		Excipients		q.s	
		Colour: Approved colour used in			
661	<b>Ivermectin &amp; Albendazole Tablets</b>	Each uncoated chewable tablet			<b>APPROVED</b>
		Ivermectin	B.P.	6mg	
		Albendazole	I.P.	400mg	
		Excipients		q.s	
		Colour : Approved colour used.			

662	<b>Ciprofloxacin Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ciprofloxacin Hydrochloride Eq. to Ciprofloxacin	I.P.	500mg	
		Excipients		q.s	
		Approved colour used in coating			
663	<b>Carvedilol Tablets I.P.</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Carvedilol	I.P.	6.25mg	
		Excipients		q.s	
		Approved colour used.			
664	<b>Carvedilol I.P. Tablets</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Carvedilol	I.P.	12.5mg	
		Excipients		q.s	
		Approved colour used.			
665	<b>Amlodipine Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Amlodipine Besylate eq. to Amlodipine	I.P.	5mg	
		Excipients		q.s	
		Approved colour used in coating			
666	<b>Amlodipine Besylate &amp; Atenolol Tablets</b>	Each Film coated tablet contains:			<b>APPROVED</b>
		Amlodipine Besylate eq. to Amlodipine	I.P.	5mg	
		Atenolol	I.P.	50mg	
		Excipients		q.s	
		Approved colour used in coating.			
667	<b>Ondansetron Orally Disintegrating Tablets I.P.</b>	Each uncoated orally disintegrating			<b>APPROVED</b>
		Ondansetron Hydrochloride Eq. to Ondansetron	I.P.	8mg	
		Excipients		q.s.	
		Colour: approved colour used.			
668	<b>Rosuvastatin Calcium Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium Eq. to Rosuvastatin	I.P.	5mg	
		Excipients		q.s.	
		Colour: approved colour used in			
669	<b>Rosuvastatin Calcium Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium Eq. to Rosuvastatin	I.P.	10mg	
		Excipients		q.s.	
		Colour: approved colour used in coating.			
670	<b>Thiocolchicoside &amp; Aceclofenac Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Thiocolchicoside	I.P.	8mg	
		Aceclofenac	I.P.	100mg	
		Excipients		q.s	
		Approved colour used in coating.			
671	<b>S(-) Amlodipine Besylate Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		S(-) Amlodipine Besylate eq. to S-Amlodipine	I.P.	5mg	
		Excipients		q.s	
		Approved colour used in coating			

672	<b>Esomeprazole Magnesium (SR) &amp; Levosulpiride (ER) Capsules</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Esomeprazole (As sustained release coated pellets)	I.P.	40mg	
		Levosulpiride(As extended release pellets )		75mg	
		Excipients		q.s	
		Approved colours used in empty			
673	<b>Aceclofenac Sustained Release Tablets</b>	Each uncoated sustained release			<b>APPROVED</b>
		Aceclofenac	I.P.	200mg	
		Excipients		q.s	
		Approved colour used.			
674	<b>Deflazacort Tablets</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Deflazacort		18mg	
		Excipients		q.s	
		Approved colour used.			
675	<b>Flavoxate B.P. Tablets</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Flavoxate	B.P.	200mg	
		Excipients		q.s	
		Approved colour used.			
676	<b>Nitrofurantoin S R Tablets</b>	Each uncoated sustained release			<b>APPROVED</b>
		Nitrofurantoin	I.P.	100mg	
		Excipients		q.s	
		Approved colour used.			
678	<b>Etodolac I.P. Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Etodolac	I.P.	400mg	
		Excipients		q.s	
		Approved colour used in coating.			
679	<b>Orlistat Capsules</b>	Each hard gelatin capsules contains :			<b>APPROVED</b>
		Orlistat		120mg	
		Excipients		q.s	
		Approved colour used in empty			
680	<b>Etoricoxib Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Etoricoxib	I.P.	120mg	
		Excipients		q.s	
		Approved colour used in coating.			
681	<b>Glibenclamide &amp; Metformin Hydrochloride (SR) Tablets</b>	Each uncoated bilayered tablet			<b>APPROVED</b>
		Glibenclamide	I.P.	5mg	
		Metformin Hydrochloride (as Sustained release form)	I.P.	850mg	
		Excipients		q.s	
		Approved colour used.			
682	<b>Mefenamic Acid &amp; Tranexamic Acid Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Mefenamic Acid	I.P.	250mg	
		Tranexamic Acid	I.P.	500mg	
		Excipients		q.s.	
		Approved colour used in coating.			
683	<b>Paracetamol SR Tablets B.P.</b>	Each uncoated sustained release			<b>APPROVED</b>
		Paracetamol	I.P.	1000mg	
		Excipients		q.s	
		Approved colour used.			
684	<b>Ursodeoxycholic acid (SR) Tablets</b>	Each uncoated sustained release			<b>APPROVED</b>
		Ursodeoxycholic acid	I.P.	300mg	
		Excipients		q.s	
		Approved colour used.			

685	<b>Ketorolac tromethamine Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ketorolac tromethamine	I.P.	10mg	
		Excipients		q.s	
		Approved colour used in coating.			
686	<b>Rosuvastatin &amp; Finofibrate Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Rosuvastatin Calcium eq. to. Rosuvastatin	I.P.	5mg	
		Finofibrate	I.P.	160mcg	
		Excipients		q.s.	
		Approved colour used in coating.			
687	<b>Hydroxychloroquine Sulphate Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Hydroxychloroquine Sulphate	I.P.	200mg	
		Excipients		q.s	
		Approved colour used in coating			
688	<b>Tizanidine HCL Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Tizanidine Hydrochloride eq. to Tizanidine	I.P.	2mg	
		Excipients		q.s	
		Approved colour used in coating			
689	<b>Atorvastatin calcium Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Atorvastatin calcium eq. to Atorvastatin	I.P.	40mg	
		Excipients		q.s	
		Approved colour used in coating			
690	<b>Atorvastatin calcium Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Atorvastatin calcium eq. to Atorvastatin	I.P.	80mg	
		Excipients		q.s	
		Approved colour used in coating			
691	<b>Carvedilol I.P. Tablets</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Carvedilol	I.P.	3.125mg	
		Excipients		q.s	
		Approved colour used.			
692	<b>Azithromycin Oral Suspension IP</b>	Each 1 ml contains :			<b>APPROVED</b>
		Azithromycin dihydrate eq. to anhydrous azithromycin	IP	40mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
693	<b>Azithromycin Oral Suspension IP</b>	Each 5 ml contains :			<b>APPROVED</b>
		Azithromycin dihydrate eq. to anhydrous azithromycin	IP	200mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
694	<b>Ofloxacin Oral Suspension IP</b>	Each 5 ml contains :			<b>APPROVED</b>
		Ofloxacin	IP	50mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
695	<b>Tranexamic Acid Tablets I.P.</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Tranexamic Acid	I.P.	250mg	
		Excipients		q.s	
		Approved colour used.			



696	<b>Terbutaline Sulphate ,Ambroxol, Guaiphenesin &amp; Menthol Syrup</b>	Each 5 ml contains :			<b>APPROVED</b>
		Terbutaline Sulphate	IP	1.25mg	
		Ambroxol HCL	IP	15mg	
		Guaiphenesin	IP	50mg	
		Menthol	IP	2mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
697	<b>Complex of iron, Vit-B12 &amp; Vit-B6 Syrup</b>	Each 5 ml contains :			<b>APPROVED</b>
		Elemental iron in a complex of iron polymatose eq. Solution		50mg	
		Vit-B12	IP	5mg	
		Vit-B6	IP	20mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
698	<b>Magaldrate &amp; Simethicone Oral Suspension</b>	Each 5 ml contains :			<b>APPROVED</b>
		Magaldrate (Anhydrous)	IP	400mg	
		Simethicone	USP	20mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
699	<b>Mefenamic Acid                      Oral Suspension</b>	Each 5 ml contains :			<b>APPROVED</b>
		Mefenamic Acid	IP	100mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
700	<b>Levocetirizine &amp; Ambroxol HCL Syrup</b>	Each 5 ml contains :			<b>APPROVED</b>
		Levocetirizine Dihydrochloride	IP	2.5mg	
		Ambroxol Hydrochloride	IP	30mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
701	<b>Albendazole Oral Suspension IP</b>	Each 5 ml contains :			<b>APPROVED</b>
		Albendazole	IP	200mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
702	<b>Itraconazole Capsules</b>	Each hard gelatin capsule contains:			<b>APPROVED</b>
		Itraconazole pellets                      eq. to	BP	200mg	
		Itraconazole (As enteric coated pellets)			
		Excipients		q.s	
		Approved colours used in empty			
703	<b>Levosaltbutamol Sulphate, Ambroxol HCL &amp; Guaiphenesin Syrup</b>	Each 5 ml contains:			<b>APPROVED</b>
		Levosaltbutamol Sulphate                      eq.	IP	0.5mg	
		to Levosaltbutamol			
		Ambroxol Hydrochloride	IP	15mg	
		Guaiphenesin	IP	50mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colours used.			

704	<b>Hydroxyzine Hydrochloride Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Hydroxyzine Hydrochloride	I.P.	25mg	
		Excipients		q.s	
		Approved colour used in coating			
705	<b>Ferrous Ascorbate &amp; Folic Acid Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ferrous Ascorbate eq. to elemental Iron	I.P.	100mg	
		Folic Acid	I.P.	1.5mg	
		Excipients		q.s	
		Approved colour used in coating.			
706	<b>Chlorpheniramine &amp; Dextromethorphan HBR Syrup</b>	Each 5 ml contains :			<b>APPROVED</b>
		Chlorpheniramine Maleate	I.P.	2.5mg	
		Dextromethorphan Hydrobromide	I.P.	10mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
707	<b>Deflazacort Oral Suspension</b>	Each 5 ml contains :			<b>APPROVED</b>
		Deflazacort		6mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
708	<b>Doxofylline Syrup</b>	Each 1 ml contains :			<b>APPROVED</b>
		Doxofylline		20mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
709	<b>Chlorhexidine Mouthwash I.P.</b>	Mouth Wash			<b>APPROVED</b>
		Chlorhexidine Gluconate Solution eq. to Chlorhexidine Gluconate	I.P.	20mg	
		Excipients		q.s	
		In a flavoured Syrupy base			
		Approved colour used			
710	<b>Olmisartan Medoxomil Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olmisartan Medoxomil	I.P.	20mg	
		Excipients		q.s	
		Approved colour used in coating.			
711	<b>Olmisartan Medoxomil Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Olmisartan Medoxomil	I.P.	40mg	
		Excipients		q.s	
		Approved colour used in coating.			
712	<b>Ambroxol HCl &amp; Doxofylline Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Ambroxol Hydrochloride	I.P.	30mg	
		Doxofylline	I.P.	400mg	
		Excipients		q.s	
		Approved colour used in coating.			
713	<b>Levosambutamol Sulphate, Ambroxol HCL &amp; Guaiphenesin Syrup</b>	Each 5 ml contains:			<b>APPROVED</b>
		Levosambutamol Sulphate eq. to Levosambutamol	IP	1mg	
		Ambroxol Hydrochloride	IP	30mg	
		Guaiphenesin	IP	50mg	
		Excipients		q.s	
		In a flavoured Syrupy base			

		Approved colours used.			
714	<b>Ondansetron Hydrochloride Oral Suspension I.P.</b>	Each 5 ml contains : Ondansetron Hydrochloride eq. to Ondansetron Excipients In a flavoured Syrupy base Approved colour used	I.P.	2mg q.s	<b>APPROVED</b>
715	<b>Amlodipine Besylate Tablets I.P.</b>	Each uncoated tablet contains: Amlodipine Besylate eq. to Amlodipine Excipients Approved colour used.	I.P.	5mg q.s	<b>APPROVED</b>
716	<b>Hydrochlorothiazide Tablets I.P.</b>	Each uncoated tablet contains: Hydrochlorothiazide Excipients Approved colour used.	I.P.	50mg q.s	<b>APPROVED</b>
717	<b>Citicoline Sodium &amp; Piracetam Tablets</b>	Each film coated tablet contains: Citicoline Sodium Eq. to Citicoline Piracetam Excipients Approved colour used in coating.	I.P. I.P.	500mg 400mg q.s	<b>APPROVED</b>
718	<b>Itraconazole Capsules B.P.</b>	Each hard gelatin capsule contains: Itraconazole Pellets Eq. to Itraconazole Excipients Approved colour used in empty	B.P.	100mg q.s	<b>APPROVED</b>
719	<b>Telmisartan &amp; Ramipril Tablets</b>	Each film coated tablet contains: Telmisartan Ramipril Excipients Approved colour used in Coating.	I.P. I.P.	40mg 5mg q.s	<b>APPROVED</b>
720	<b>Nevivolol &amp; Hydrochlorothiazide Tablets</b>	Each film coated tablet contains: Nevivolol Hydrochloride eq. to Nevivolol Hydrochlorothiazide Excipients Approved colour used in Coating.	I I.P. I.P.	5mg 12.5mg q.s	<b>APPROVED</b>
721	<b>Artemether &amp; Lumefantrine Tablets</b>	Each uncoated tablet contains: Artemether Lumefantrine Excipients Approved colour used.	I.P.	80mg 480mg q.s	<b>APPROVED</b>
722	<b>Thiocolchicoside &amp; Etoricoxib Tablets</b>	Each film coated tablet contains: Thiocolchicoside Etoricoxib Excipients Approved colour used in coating.	I.P. I.P.	4mg 60mg q.s	<b>APPROVED</b>
723	<b>Roxithromycin Tablets</b>	Each film coated tablet contains: Roxithromycin Excipients	I.P.	300mg q.s	<b>APPROVED</b>

		Approved colour used in coating.			
724	<b>Thiocolchicoside &amp; Etoricoxib Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Thiocolchicoside	I.P.	8mg	
		Etoricoxib	I.P.	60mg	
		Excipients		q.s	
		Approved colour used in coating.			
725	<b>Thiocolchicoside &amp; Etodolac Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Etodolac	I.P.	400mg	
		Thiocolchicoside	I.P.	4mg	
		Excipients		q.s	
		Approved colour used in coating.			
726	<b>Ondansetron Orally Disintegrating Tablets I.P.</b>	Each uncoated orally disintegrating			<b>APPROVED</b>
		Ondansetron Hydrochloride Eq. to Ondansetron	I.P.	16mg	
		Excipients		q.s.	
		Colour: approved colour used.			
727	<b>Fluconazole Tablets I.P.</b>	Each uncoated tablet contains:			<b>APPROVED</b>
		Fluconazole	I.P.	200mg	
		Excipients		q.s	
		Approved colour used.			
728	<b>Drotaverine HCl &amp; Mefenamic Acid Tablets</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Drotaverine Hydrochloride	I.P.	80mg	
		Mefenamic Acid	I.P.	250mg	
		Excipients		q.s	
		Approved colour used in coating.			
729	<b>Clarithromycin Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clarithromycin	I.P.	250mg	
		Excipients		q.s	
		Approved colour used in coating.			
730	<b>Clarithromycin Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Clarithromycin	I.P.	500mg	
		Excipients		q.s	
		Approved colour used in coating.			
731	<b>Letrozole Tablets U.S.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Letrozole	U.S.P.	2.5mg	
		Excipients		q.s	
		Approved colour used.			
732	<b>Metronidazole Benzoate Oral Suspension</b>	Each 5 ml contains			<b>APPROVED</b>
		Metronidazole Benzoate eq. to Metronidazole	I.P.	200mg	
		Excipients	q.s.		
		In A Flavoured Syrupy base			
		Approved colour used			
733	<b>Fexofinadine Hydrochloride Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fexofinadine Hydrochloride Eq. to Fexofinadine	I.P.	120mg	
		Excipients		q.s	
		Colour: Approved colour used in			
734	<b>Fexofinadine Hydrochloride Tablets I.P.</b>	Each film coated tablet contains:			<b>APPROVED</b>
		Fexofinadine Hydrochloride Eq. to Fexofinadine	I.P.	180mg	
		Excipients		q.s	

		Colour: Approved colour used in				
735	Tinidazole Tablets I.P.	Each film coated tablet contains:			APPROVED	
		Tinidazole	I.P.	500mg		
		Excipients		q.s		
		Colour: Approved colour used in coating				
736	Granisetron Hydrochloride Tablets	Each film coated tablet contains:				APPROVED
		Granisetron Hydrochloride eq. to Granisetron	I.P.	1mg		
		Excipients		q.s		
		Colour: Approved colour used in				
737	Lactulose Oral Solution U.S.P.	Each 5 ml contains				APPROVED
		Lactulose concentrate eq. to	I.P.	3.33 mg		
		Lactulose				
		Excipients		q.s.		
		In A Flavoured Syrupy base				
Approved colour used						
738	Lactulose Oral Solution U.S.P.	Each 15 ml contains				APPROVED
		Lactulose concentrate eq. to	I.P.	10 mg		
		Lactulose				
		Excipients		q.s.		
		In A Flavoured Syrupy base				
Approved colour used						
739	Cefixime Tablet IP	Each uncoated tablet contains:				APPROVED
		Cefixime Trihydrate	I.P.	200 mg		
		Equ. to Cefixime Anhydrous				
740	Cefixime Tablet IP	Each dispersible tablet contains:				APPROVED
		Cefixime Trihydrate	I.P.	200 mg		
		Equ. to Cefixime Anhydrous				
741	Cefixime Tablet	Each uncoated tablet contains:				APPROVED
		Cefixime Trihydrate	I.P.	100 mg		
		Equ. to Cefixime Anhydrous				
741	Cefixime Tablet	Each dispersible tablet contains:				APPROVED
		Cefixime Trihydrate	I.P.	100 mg		
		Equ. to Cefixime Anhydrous				
743	Cefixime and Potassium Clavulanate Tablet	Each film coated tablet contains:				APPROVED
		Cefixime Trihydrate	I.P.	200 mg		
		Equ. to Cefixime Anhydrous				
		Potassium Clavulanate Diluted	I.P.	125 mg		
		Equ. to Clavulanic Acid				
744	Cefixime and Dicloxacillin ER Tablets	Each film coated tablet contains:				APPROVED
		Cefixime Trihydrate	I.P.	200 mg		
		Equ. to Cefixime Anhydrous				
		Dicloxacillin Sodium	I.P.	500 mg		
		equ.to Dicloxacillin				
(as Extended release granules)						
745	Cefuroxime Axetil Tablet	Each film coated tablet contains:				APPROVED
		Cefuroxime Axetil	I.P.	250 mg		
		Equ to Cefuroxime Anhydrous				
746	Cefuroxime Axetil Tablet	Each film coated tablet contains:				APPROVED
		Cefuroxime Axetil	I.P.	500 mg		
		Equ to Cefuroxime Anhydrous				
747	Cefpodoxime Tablet	Each uncoated tablet contains				APPROVED
		Cefpodoxime Proxetil	I.P.	100 mg		

		Equ to Cefpodoxime Anhydrous	I.P.	100 mg	
748	Cefpodoxime Tablet	Each filmcoated tablet contains Cefpodoxime Proxetil Equ to Cefpodoxime Anhydrous	I.P.	200 mg	APPROVED
749	Cefpodoxime and Potassium Clavulanate Tablet	Each filmcoated tablet contains Cefpodoxime Proxetil Equ to Cefpodoxime Potassium Clavulanate Diluted Equ. to Clavulanic Acid	I.P. I.P.	200 mg 125 mg	APPROVED
750	Cefpodoxime and Ofloxacin Tablet	Each filmcoated tablet contains Cefpodoxime Proxetil Equ to Cefpodoxime Ofloxacin	I.P. I.P.	200 mg 100 mg	APPROVED
751	Cefpodoxime and Ofloxacin Tablet	Each filmcoated tablet contains Cefpodoxime Proxetil Equ to Cefpodoxime Ofloxacin	I.P. I.P.	200 mg 100 mg	APPROVED
752	Amoxycillin Trihydrate and Potassium Clavulanate Tablets	Each filmcoated tablet contains Amoxycillin Trihydrate Equ. to Amoxycillin Potassium Clavulanate Diluted Equ. to Clavulanic Acid	I.P. I.P.	875 mg 125mg	APPROVED
753		Each filmcoated tablet contains			
754	Cefuroxime and Potassium Clavulanate Tablets	Cefuroxime Axetil Equ to Cefuroxime Anhydrous Potassium Clavulanate Diluted Equ. to Clavulanic Acid	I.P. I.P.	500 mg 125 mg	APPROVED
755	Cefuroxime Axetil (ER) Tablet	Each filmcoated tablet contains Cefuroxime Axetil Equ to Cefuroxime Anhydrous (As extended release granules)	I.P.	500 mg	APPROVED
756	Cefixime and Potassium Clavulanate Tablets	Each filmcoated tablet contains Cefixime Trihydrate Equ. to Cefixime Anhydrous Potassium Clavulanate Diluted Equ. to Clavulanic Acid	I.P. I.P.	50 mg 31.5 mg	APPROVED
757	Cefixime and Cloxacillin Tablet	Each filmcoated tablet contains Cefixime Trihydrate Equ. to Cefixime Anhydrous Cloxacillin Sodium Equ. To Cloxacillin	I.P. I.P.	200 mg 500 mg	APPROVED
758	Cefixime, Cloxacillin amg Lactobacillus Tablet	Each filmcoated tablet contains Cefixime Trihydrate Equ. to Cefixime Anhydrous Cloxacillin Sodium Equ.to Cloxacillin Lactobacillus 90M.Spores	I.P. I.P.	200 mg 500 mg	APPROVED
759	Amoxycillin Trihydrate and Potassium Clavulanate Tablets	Each filmcoated tablet contains Amoxycillin Trihydrate Equ. to Amoxycillin Potassium Clavulanate Diluted	I.P.	500 mg	APPROVED

		Equ. to Clavulanic Acid	B.P.	125 mg	
760	Amoxycillin Trihydrate and Potassium Clavulanate Tablets	Each dispersible tablet contains :			APPROVED
		Amoxycillin Trihydrate	I.P.	400 mg	
		Equ. to Amoxycillin			
		Potassium Clavulanate Diluted	B.P.	57 mg	
		Equ. to Clavulanic Acid			
761	Amoxycillin Trihydrate and Potassium Clavulanate Kid Tablets	Each filmcoated tablet contains			APPROVED
		Amoxycillin Trihydrate	I.P.	200 mg	
		Equ. to Amoxycillin			
		Potassium Clavulanate Diluted	I.P.	28.5mg	
		Equ. to Clavulanic Acid			
762	Cefixime, and Lactobacillus Tablet	Each coated tablet contains			APPROVED
		Cefixime Trihydrate	I.P.	200 mg	
		Equ. to Cefixime Anhydrous			
		Lactobacillus 45M.Spores			
763	Cefixime Tablet	Each film coated tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	200 mg	
		Equ. to Cefixime Anhydrous			
764	Cefixime Tablet	Each filmcoated tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	100 mg	
		Equ. to Cefixime Anhydrous			
765	Cefixime and Moxifloxacin (SR) Tablet	Each filmcoated tablet contains:			APPROVED
		Cefixime Tryhydrate	I.P.	400 mg	
		Equ. to Cefixime Anhydrous			
		(As sustained-release granules)			
		Moxifloxacin HCl	I.P.	400 mg	
		equ. to Moxifloxacin			
(As sustained-release granules)					
766	Cefixime, Cloxacillin and Lactobacillus Tablet	Each filmcoated tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	100 mg	
		Equ. to Cefixime Anhydrous			
		Cloxacillin Sodium	I.P.	500 mg	
		Equ.to Cloxacillin			
Lactobacillus 45M.Spores					
767	Cefpodoxime Tablet	Each dispersible tablet contains:			APPROVED
		Cefpodoxime Proxetil	I.P.	200 mg	
		Equ to Cefpodoxime Anhydrous			
768	Cefpodoxime Dicloxacillin ER Tablets	Each film coated tablet contains:			APPROVED
		Cefpodoxime proxetil	I.P.	200 mg	
		Equ. to Cefpodoxime			
		Dicloxacillin Sodium	I.P.	500 mg	
		equ.to Dicloxacillin			
		(as Extended release granules)			
769	Cefpodoxime Dicloxacillin ER Tablets	Each filmcoated tablet contains:			APPROVED
		Cefpodoxime proxetil	I.P.	100 mg	
		Equ. to Cefpodoxime			
		Dicloxacillin Sodium	I.P.	500 mg	
		equ.to Dicloxacillin			
		(as Extended release granules)			
770	Cefpodoxime and Potassium Clavulanate Tablet	Each filmcoated tablet contains:			APPROVED
		Cefpodoxime Proxetil	I.P.	100 mg	
		Equ to Cefpodoxime			
		Potassium Clavulanate Diluted	I.P.	62.5 mg	

		Equ. to Clavulanic Acid		02.5 mg	
771	Cefixime and Lactobacillus dispersible Tablet	Each dispersible tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	100 mg	
		Equ. to Cefixime Anhydrous			
		Lactobacillus 2.5 B.Spores			
772	Cefixime, and Lactobacillus dispersible Tablet	Each dispersible tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	200 mg	
		Equ. to Cefixime Anhydrous			
		Lactobacillus 2.5 B.Spores			
773	Cefuroxime and Potassium Clavulanate Tablets	Each filmcoated tablet contains:			APPROVED
		Cefuroxime Axetil	I.P.	250 mg	
		Equ to Cefuroxime Anhydrous			
		Potassium Clavulanate Diluted	I.P.	125 mg	
		Equ. to Clavulanic Acid			
774	Amoxycillin Trihydrate, Potassium Clavulanate Tablets	Each filmcoated tablet contains:			APPROVED
		Amoxycillin Trihydrate	I.P.	250 mg	
		Equ. to Amoxycillin			
		Potassium Clavulanate Diluted	I.P.	125 mg	
		Equ. to Clavulanic Acid			
775	Amoxycillin Trihydrate, Potassium Clavulanate with Lactobacillus Tablets	Each filmcoated tablet contains:			APPROVED
		Amoxycillin Trihydrate	I.P.	500 mg	
		Equ. to Amoxycillin			
		Potassium Clavulanate Diluted	I.P.	125 mg	
		Equ. to Clavulanic Acid			
		Lactobacillus 2.5 B.Spores			
776	Cefixime trihydrate and Ofloxacin Tablet	Each filmcoated tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	200 mg	
		Equ. to Cefixime Anhydrous			
		Ofloxacin	I.P.	200 mg	
778	Cefixime Tablet	Each filmcoated tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	400 mg	
		Equ. to Cefixime Anhydrous			
779	Cefixime trihydrate and Azithromycin trihydrate Tablet	Each filmcoated tablet contains:			APPROVED
		Cefixime Trihydrate	I.P.	200 mg	
		Equ. to Cefixime Anhydrous			
		Azithromycin trihydrate	I.P.	250 mg	
		eq to any Azithromycin			