



FINAL APPROVAL LIST

CB Healthcare

Works : Vill. Messa Tibba, Tehsil. Nalagarh, Distt : Solan (H.P.) 174 101

S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
1	Fungal Diastase With Papain Syrup	Each 5ml Contains Fungal Diastase (1:800) (Derived From aspergillus Oryzae Digest not less than 10 gm of Cooked Starch) Papain Flavoured Syrupy Base Colour: Caramel	I.P. I.P.	12.50 mg 7.5 mg q.s.		01-09-2017
2	Levocetirizine Hydrochloride & Montelukast Sodium Syrup	Each 5 ml Contains Levocetirizine Hydrochloride Montelukast Sodium Eq.to Montelukast Flavoured Syrupy Base Approved Colour Used	I.P. I.P.	2.5 mg 4 mg q.s.		01-09-2017
3	Codeine Phosphate & Triprolidine Hcl. Syrup	Each 5ml Contains Codeine Phosphate Triprolidine Hydrochloride Flavoured Syrupy Base Approved Colour Used	I.P. I.P.	10 mg 1.25 mg q.s.		01-09-2017
5	Dextromethorphan Hbr. Phenylephrine Hcl. & Chlorpheniramine Maleate Syrup	Each 5ml Contains Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Chlorpheniramine Maleate Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	10 mg 5 mg 2 mg q.s.		01-09-2017
6	Cyproheptadine Hcl , Tricholine Citrate & Sorbitol Syrup	Each 5ml Contains Cyproheptadine Hydrochloride Eq.to anhydrous Cyproheptadine Hcl Tricholine Citrate (65 % w/v) Sorbitol 70 % (Non Crystallizing) Flavoured Syrupy Base Approved Colour Used	I.P.	2 mg 275 mg q.s. q.s.		01-09-2017
7	Cetirizine , Dextromethorphan Hbr. & Phenylephrine Hcl. Syrup	Each 5ml Contains Cetirizine Hydrochloride Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	5 mg 5 mg 2.5 mg q.s.		01-09-2017
8	Ofloxacin , Metronidazole & Simethicone Suspension	Each 5ml Contains Ofloxacin Metronidazole Benzoate Eq.to Metronidazole Simethicone Flavoured Palatable Base Colour: Sunset Yellow FCf	I.P. I.P. I.P.	50 mg 120 mg 10 mg q.s.		01-09-2017
9	Chlorpheniramine Maleate & Dextromethorphan Hbr. Syrup	Each 5 ml Contains Chlorpheniramine Maleate Dextromethorphan Hydrobromide Flavoured Syrupy Base Approved Colour Used	I.P. I.P.	4 mg 10 mg q.s.		01-09-2017
10	Calcium Carbonate & Vitamin D3 Suspension	Each 5ml Contains Calcium Carbonate Eq.to Elemental Calcium Vitamin D3 Flavoured Palatable Base Approved Colour Used	I.P. I.P.	625 mg 250 mg 125 IU q.s.		01-09-2017



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11	Azithromycin Tablets IP	Each Film Coated Tablet Contains Azithromycin Dihydrate Eq.to. Anhydrous Azithromycin Excipients Approved Colour Used	I.P.	250 mg q.s.		01-09-2017
12	Azithromycin Tablets IP	Each Film Coated Tablet Contains Azithromycin Dihydrate Eq.to. Anhydrous Azithromycin Excipients Approved Colour Used	I.P.	500 mg q.s.		01-09-2017
13	Aceclofenac & Diacerein Tablet	Each Uncoated Tablet Contains Aceclofenac Diacerein Excipients Approved Colour Used	I.P. I.P.	100 mg 50 mg q.s.		01-09-20217
14	Alprazolam Tablet IP	Each Uncoated Tablet Contains Alprazolam Excipients Approved Colour Used	I.P.	0.5 mg q.s.		01-09-2027
15	Ofloxacin & Ornidazole Tablets I.P.	Each Film Coated Tablet Contains Ofloxacin Ornidazole Excipients Approved Colour Used	I.P. I.P.	200 mg 500 mg q.s.		01-09-20217
16	Diclofenac potassium, Paracetamol & Serratiopeptidase Tablet	Each Film Coated tablet Contain Diclofenac Potassium Paracetamol Serratiopeptidase (Enteric Coated Granules) Excipients Approved Colour Used	I.P. I.P. I.P.	50 mg 325 mg 10 mg q.s.		01-09-2017
17	Diclofenac potassium, Paracetamol & Chlorzoxazone Tablet	Each Uncoated tablet Contain Diclofenac Potassium Paracetamol Chlorzoxazone Excipients Approved Colour Used	I.P. I.P. U.S.P.	50 mg 325 mg 250 mg q.s.		01-09-2017
18	Terbutaline Sulphate, Bromhexine Hcl with Chlorpheniramine Maleate Tablets	Each Uncoated Tablet Contains Terbutaline Sulphate Bromhexine Hydrochloride Chlorpheniramine Maleate Excipients Colour: Erythrosine	I.P. I.P. I.P.	2.5 mg 16 mg 4 mg q.s.		01-09-2017
19	Aceclofenac & Paracetamol Tablet	Each Uncoated tablet Contain Aceclofenac Paracetamol Excipients Approved Colour Used	I.P. I.P.	100 mg 325 mg q.s.		01-09-2017
21	Prolonged Release Tramadol Tablets IP	Each uncoated prolonged release tablet contains: Tramadol Hydrochloride (As prolonged release) Excipients	I.P.	100 mg		01-09-2017



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22	Aceclofenac, Paracetamol & Thiocholchicoside Tables	Each Film Coated tablet Contain Aceclofenac Paracetamol Thiocholchicoside Excipients Approved Colour Used	I.P. I.P. I.P.	100 mg 325 mg 4 mg q.s.		01-09-2017
23	Lycopene with Multivitamins and Multiminerals Capsule	Each Hard Gelatin Capsule Contain Vitamin A (as Acetate) Vitamin E Vitamin C (Coated) Zinc Sulphate Copper Sulphate Lycopene 10 % Manganese Sulphate Beta Carotene(10 % Dispersible) Selenium (As Selenium Dioxide) Approved Colour Used in Empty Capsule Shell	IP IP IP IP USP USP USP USP USP	5000 IU 15 mg 75 mg 7.5 mg 1 mg 2000 mcg 1.5 mg 10 mg 150 mcg		01-09-2017
24	Methylcobalamin, Alpha Lipoic Acid , Folic Acid, Thiamine Hcl, & Pyridoxine Hcl Capsule	Each Hard Gelatin Capsule Contain Methylcobalamin Alpha Lipoic Acid Folic Acid Thiamine Hydrochloride Pyridoxine Hydrochloride Excipients Approved Colour Used in Empty Capsule Shell	I.P. I.P. I.P. I.P. I.P.	1500 mcg 100 mg 1.5 mg 10 mg 3 mg q.s.		01-09-2017
25	Lycopene, Lutein Beta Carotene, Vitamin B-Complex , Iron Lysine Hcl , Minerals With Zinc Sulphate Capsules	Each Hard Gelatin Capsule Contains Lycopene 6% Lutein Beta Carotene (10 % Dispersion) Vitamin B6 Vitamin B12 Vitamin D3 Folic Acid Niacinamide Biotin Lysine Hcl Ferrous Fumarate Dibasic Calcium Phosphate Zinc Sulphate Monohydrate Copper Sulphate Magnesium Hydroxide Magnesium Sulphate Chromium Chloride Sodium selenate Approved colour used in Empty Capsule Shell	USP USP USP IP IP IP IP IP IP USP USP IP IP IP USP IP USP USP BP	5 mg 6 mg 10 mg 1 mg 5 mcg 200 IU 500 mcg 25 mg 10 mcg 50 mg 45 mg 75 mg 10 mg 500 mcg 10 mg 500 mcg 200 mcg 100 mcg		01-09-2017
26	Pregabalin & Methylcobalamin Capsules I.P.	Each Hard Gelatin Capsule Contain Pregabalin Methylcobalamin Excipients Approved Colour Used in Empty Capsule Shell	I.P. I.P.	75 mg 750 mcg q.s.		01-09-2017
27	Gabapentin & Methylcobalamin	Each Hard Gelatin Capsule Contain Gabapentin	U.S.P.	300 mg		01-09-2017



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	Capsule	Methylcobalamin Excipients Approved Colour Used in Empty Capsule Shell	I.P.	500 mcg q.s.		
28	Omeprazole Gastro Resistant Capsule IP	Each Hard Gelatin Capsule Contains Omeprazole (As Enteric coated Pallets) Approved Colour used in Empty capsule shell	I.P.	20 mg		01-09-2017
29	Dicyclomine Hcl., Acetaminophen & Tramadol Hcl. Capsules	Each Hard Gelatin Capsule Contains Dicyclomine Hydrochloride Acetaminophen Tramadol Hydrochloride Excipients Approved Colour Used in Empty Capsule Shell	I.P. I.P. I.P.	10 mg 325 mg 50 mg		01-09-2017
30	Omeprazole & Domperidone Capsule	Each Hard Gelatin Capsule Contains Omeprazole (As Enteric coated Pallets) Domperidone Approved Colour used in Empty capsule shell	I.P. I.P.	20 mg 10 mg		01-09-2017
31						
32	Enteric Coated Rabeprazole Sodium & Itopride Hcl Sustained Release Capsule I.P.	Each Hard Gelatin Capsule Contains Rabeprazole Sodium (As Enteric Coated Pallets) Itopride Hydrochloride (As Sustained Release Pallets) Excipients Approved Colour used un Empty Capsule Shell	I.P. I.P.	20 mg 150 mg q.s		01-09-2017
33	Enteric Coated Rabeprazole Sodium & Aceclofenac Sustained Release Capsule	Each Hard Gelatin Capsule Contains Rabeprazole Sodium (As Enteric Coated Pallets) Aceclofenac (As Sustained Release Pallets) Excipients Approved Colour used un Empty Capsule Shell	I.P. I.P.	20 mg 200 mg q.s		01-09-2017
34	Enteric Coated Rabeprazole Sodium & Levosulpiride Sustained Release Capsule	Each Hard Gelatin Capsule Contains Rabeprazole Sodium (As Enteric Coated Pallets) Levosulpiride (As Sustained Release Pallets) Excipients Approved Colour used un Empty Capsule Shell	I.P. I.P.	20 mg 75 mg q.s		01-09-2017
35	Enteric Coated Pantoprazole & Domperidone Sustained Release Capsule I.P.	Each Hard Gelatin Capsule Contains Pantoprazole Sodium Eq. To. Pantoprazole (As Enteric Coated Pallets) Domperidone (As Sustained Release Pallets) Excipients Approved Colour used un Empty Capsule Shell	I.P. I.P.	40 mg 30 mg q.s		01-09-2017



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37	Aceclofenac Tablets IP	Each Film Coated tablet Contains Aceclofenac Excipients Approved Colour Used	I.P.	100 mg q.s.		01-09-2017
38	Paracetamol, Phenylephrine Hcl & Chlorpheniramine Maleate Suspension	Each 5ml Contains Paracetamol Phenylephrine Hydrochloride Chlorpheniramine Maleate Flavoured Palatable Base Approved Colour Used	I.P. I.P. I.P.	125 mg 2.5 mg 1 mg q.s.		01-09-2017
39	Paracetamol, Phenylephrine Hcl & Chlorpheniramine Maleate Suspension	Each 5ml Contains Paracetamol Phenylephrine Hydrochloride Chlorpheniramine Maleate Flavoured Palatable Base Approved Colour Used	I.P. I.P. I.P.	250 mg 5 mg 2 mg q.s.		01-09-2017
40	Terbutaline sulphate, Bromhexine Hcl, Guaiphenesin & Menthol Syrup	Each 10 ml Contains Terbutaline sulphate Bromhexine Hydrochloride Guaiphenesin Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	2.5 mg 8 mg 100 mg 3 mg q.s.		01-09-2017
42	Piracetam Syrup	Each 5 ml contains Piracetam In a Flavoured Syrupy Base Colour :Erythrosine	I.P.	500 mg		01-09-2017

43	Paracetamol Paediatric Oral Suspension IP	Each 5ml Contains Paracetamol Flavoured Palatable Base Approved Colour Used	I.P.	125 mg q.s.		01-09-2017
44	Paracetamol Paediatric Oral Suspension IP	Each 5ml Contains Paracetamol Flavoured Palatable Base Approved Colour Used	I.P.	250 mg q.s.		01-09-2017
45	Aceclofenac & Paracetamol Suspension	Each 5 ml Contains Aceclofenac Paracetamol Flavoured Palatable Base Approved Colour Used	I.P. I.P.	50 mg 125 mg q.s.		01-09-2017
46	Cetirizine Hcl , Paracetamol & Phenylephrine Hcl Suspension	Each 5 ml Contains Cetirizine Hydrochloride Paracetamol Phenylephrine Hydrochloride Flavoured Palatable Base Approved Colour Used	I.P. I.P. I.P.	2.5 mg 125 mg 2.5 mg q.s.		01-09-2017
47	Ambroxol Hcl, Guaiphenesin, Chlorpheniramine Maleate , Phenylephrine Hcl & Menthol Syrup	Each 5 ml Contains Ambroxol Hydrochloride Guaiphenesin Chlorpheniramine Maleate Phenylephrine Hydrochloride Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P. I.P.	15 mg 50 mg 2 mg 5 mg 1 mg q.s.		01-09-2017



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48	Ofloxacin Suspension IP	Each 5ml Contains Ofloxacin Flavoured Palatable Base Colour: Sunset Yellow FCf	I.P.	50 mg q.s.		01-09-2017
49	Pantoprazole Tablet IP	Each Enteric Coated Tablet Contains Pantoprazole Sodium Sesquihydrate Equivalent to Pantoprazole Excipients Approved colour used	I.P.	40 mg q.s.		01-09-2017
50	Pantoprazole & Domperidone Tablet	Each Enteric Coated Tablet Contains Pantoprazole Sodium Sesquihydrate Equivalent to Pantoprazole Domperidone Excipients Approved colour used	I.P. I.P.	40 mg 10 mg q.s.		01-09-2017
51	Mefenamic Acid & Dicyclomine Hcl Tablets I.P.	Each Uncoated Tablet Contains Mefenamic Acid Dicyclomine Hydrochloride Excipients Approved Colour Used	I.P. I.P.	250 mg 10 mg q.s.		01-09-2017
52	Levofloxacin & Ornidazole Suspension	Each 5ml Contains Levofloxacin Hemihydrate Eq. to Levofloxacin Ornidazole Flavoured Palatable Base Colour: Sunset Yellow FCF	I.P. I.P.	125 mg 125 mg q.s.		01-09-2017
53	Fexofenadine with Montelukast Tablets	Each Film coated tablet contains Fexofenadine Hydrochloride Montelukast Sodium Eq.to Montelukast Excipients Approved Colour Used	I.P. I.P.	120 mg 10 mg q.s.		01-09-2017
54	Aceclofenac Sustained Release Tablet	Each Film Coated Sustained Release Tablet Contains Aceclofenac Excipients Approved Colour Used	I.P.	200 mg q.s.		01-09-2017
55	Cetirizine , Dextromethorphan Hbr & Phenylephrine Hcl Syrup	Each 5ml Contains Cetirizine Hydrochloride Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	5 mg 5 mg 2.5 mg q.s.		01-09-2017
56	Pantoprazole & Ondansetron Hcl. Tablet	Each Enteric Coated Tablet Contains Pantoprazole Sodium Eq. To Pantoprazole Ondansetron Hydrochloride Eq. to Ondansetron Excipients Approved Colour Used	I.P. I.P.	40 mg 4 mg q.s.		01-09-2017
57	Aceclofenac & Drotaverine Hcl. Tablets	Each Film coated tablet contains Aceclofenac Drotaverine Hydrochloride Eq. to Drotaverine Excipients Approved Colour Used	I.P. I.P.	100 mg 80 mg q.s.		01-09-2017
58	Ciprofloxacin Hcl Tablets IP	Each Film Coated Tablet Contains Ciprofloxacin Hydrochloride	I.P.			01-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	500 mg	Eq.to Ciprofloxacin Excipients Approved Colour Used		500 mg q.s.		
59	Ciprofloxacin Hcl Tablets IP 250 mg	Each Film Coated Tablet Contains Ciprofloxacin Hydrochloride Eq.to Ciprofloxacin Excipients Approved Colour Used	I.P.	250 mg q.s.		01-09-2017
60	Levofloxacin Tablets IP 500 mg	Each Film Coated Tablet Contains Levofloxacin Hemihydrate Eq.to Levofloxacin Excipients Approved Colour used	I.P.	500 mg q.s.		01-09-2017
61	Levofloxacin Tablets IP 250 mg	Each Film Coated Tablet Contains Levofloxacin Hemihydrate Eq.to Levofloxacin Excipients Approved Colour used	I.P.	250 mg q.s.		01-09-2017
62	Ofloxacin Tablets IP 200 mg	Each Film Coated Tablet Contains Ofloxacin Excipients Approved Colour Used	I.P.	200 mg q.s.		01-09-2017
63	Cetirizine Hcl. With Phenylephrine Hcl. Tablets	Each Uncoated Tablet Contains Cetirizine Hydrochloride Phenylephrine Hydrochloride Excipients Colour Ponceau 4R	I.P. I.P.	10 mg 10 mg q.s.		01-09-2017
64	Levocetirizine DI- Hcl Tablets IP	Each Film Coated Tablet Contains Levocetirizine Dihydrochloride Excipients Approved Colour Used	I.P.	5 mg q.s.		01-09-2017
65	Febuxostat Tablet	Each Film coated tablet contains Febuxostat Excipients Approved Colour Used		40 mg q.s.		01-09-2017
66	Cetirizine Hcl. Tablets IP 10 mg	Each Film Coated Tablet Contains Cetirizine Hydrochloride Excipients Approved colour Used	I.P.	10 mg q.s.		01-09-2017
67	Fexofenadine Hcl. Tablets IP 120 mg	Each Film Coated Tablet Contains Fexofenadine Hydrochloride Excipients Approved Colour Used	I.P.	120 mg q.s.		01-09-2017
68	Febuxostat Tablet	Each Film coated tablet contains Ebuxostat Excipients Approved Colour Used		80 mg q.s.		01-09-2017
69	Diclofenac Potassium & Serratiopeptidase Tablet	Each Enteric Coated Tablet Contains Diclofenac Potassium Serratiopeptidase Excipients Approved Colour Used	B.P. I.P.	50 mg 10 mg q.s.		01-09-2017
71	Nimesulide & Paracetamol Tablets	Each uncoated tablet contains Nimesulide Paracetamol Excipients Approved Colour Used	B.P. I.P.	325 mg 100 mg q.s.		01-09-2017



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73	Nimesulide Tablets	Each uncoated tablet contains Nimesulide Excipients Approved Colour Used	B.P.	100 mg q.s.		01-09-2017
74	Methylprednisolone Tablets IP	Each uncoated tablet contains Methylprednisolone Excipients	I.P.	4 mg q.s.		01-09-2017
75	Methylprednisolone Tablets IP	Each uncoated tablet contains Methylprednisolone Excipients	I.P.	16 mg q.s.		01-09-2017
76	Ferrous Ascorbate Folic Acid & Cyanocobalamin Suspension	Each 5ml of suspension contains Ferrous Ascorbate eq. to Elemental Iron Folic Acid Cyanocobalamin triturate eq. to Cyanocobalamin Flavoured Palatable Base Approved Colour Used	I.P. I.P.	30 mg 0.5 mg 2.5 mcg q.s.		01-09-2017
77	Deflazacort Tablets	Each uncoated tablet contains Deflazacort Excipients		6 mg q.s.		01-09-2017
78	Ondansetron Dispersible Tablets	Each uncoated Dispersible tablet contains Ondansetron Hydrochloride Eq.to Ondansetron Excipients	I.P.	4 mg q.s.		01-09-2017
79	Ambroxol Hcl, Terbutaline sulphate , Guaiphenesin & Menthol Syrup	Each 5ml Contains Ambroxol Hydrochloride Terbutaline sulphate Guaiphenesin Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	15 mg 1.25 mg 50 mg 2.5 mg q.s.		01-09-2017
80	Telmisartan Tablets I.P. 40 mg	Each uncoated tablet contains Telmisartan Excipients	I.P.	40 mg q.s.		01-09-2017
81	Amlodipine Besylate Tablet I.P. 5mg	Each uncoated tablet contains Amlodipine Besylate Eq.to Amlodipine Excipients	I.P.	5 mg q.s.		01-09-2017
82	Phenylephrine Hcl & Chlorpheniramine Maleate Syrup IP	Each 5 ml Contains Phenylephrine Hydrochloride Chlorpheniramine Maleate Flavoured Syrupy Base Approved Colour Used	I.P. I.P.	5 mg 2 mg q.s.		01-09-2017
83	Terbutaline sulphate, Bromhexine Hcl., Guaiphenesin & Menthol Syrup	Each 5ml Contains Terbutaline sulphate Bromhexine Hydrochloride Guaiphenesin Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	1.25 mg 2 mg 50 mg 0.5 mg q.s.		01-09-2017
84	Albendazole & Ivermectin Tablets	Each Uncoated Chewable Tablet Contains Albendazole Ivermectin Excipients	I.P. I.P.	400 mg 6 mg q.s.		01-09-2017



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		Approved Colour & Flavour Used				
85	Albendazole & Ivermectin Tablets	Each Uncoated Chewable Tablet Contains Albendazole Ivermectin Excipients Approved Colour & Flavour Used	I.P. I.P.	400 mg 12 mg		01-09-2017
86	Torseamide Tablets IP 10 mg	Each Uncoated Tablet Contains Torsemide Excipients	U.S.P.	10 mg		01-09-2017
87	Fluconazole Tablets I.P. 150 mg	Each uncoated tablet contains Fluconazole Excipients Approved Colour Used	I.P.	150 mg q.s.		01-09-2017
88	Fluconazole Tablets I.P. 200 mg	Each uncoated tablet contains Fluconazole Excipients Approved Colour Used	I.P.	200 mg q.s.		01-09-2017
89	Mefenamic Acid & Paracetamol Suspension	Each 5 ml Contains Mefenamic Acid Paracetamol Flavoured Palatable Base Approved Colour Used	I.P. I.P.	50 mg 125 mg q.s.		01-09-2017
90	Deflazacort Oral Suspension	Each 5 ml Contains Deflazacort Flavoured Palatable Base Approved Colour Used		6 mg q.s.		01-09-2017
91	Calcium Carbonate & Vitamin D3 Tablet I.P.	Each Film Coated Tablet Contains Calcium Carbonate Eq.to Elemental Calcium Vitamin D3 Excipients Approved Colour Used	I.P. I.P.	1250 mg 500 mg 250 I.U. q.s.		01-09-2017
92	Sucralfate Suspension	Each 5 ml Contains Sucralfate Flavoured Palatable Base Colour: Sunset Yellow FCF	U.S.P.	500 mg q.s.		01-09-2017
93	Haematinic Syrup of Iron, Folic Acid , Vitamin B12, Manganese & Copper	Each 15 ml Contains Ferric Ammonium Citrate Eq.to Elemental Iron Vitamin B12 Folic Acid Manganese Sulphate Cupric Sulphate Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. U.S.P. U.S.P.	160 mg 32.8 mg 7.5 mcg 0.5 mg 30 mcg 30 mcg q.s.		01-09-2017
94	Dextromethorphan Hbr., Chlorpheniramine Maleate & Menthol Syrup	Each 5ml Contains Dextromethorphan Hydrobromide Chlorpheniramine Maleate Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	10 mg 4 mg 2.5 mg q.s.		01-09-2017
95	Calcium Citrate Malate , Vitamin D3 with Magnesium Hydroxide Tablet	Each Film Coated Tablet Contains Calcium Citrate Malate Eq.to elemental Calcium Vitamin D3 Magnesium Hydroxide Eq.to elemental Magnesium Excipients	I.P. B.P.	250 mg 250 IU 50 mg q.s.		01-09-2017



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		Approved Colour Used				
96	Linezolid Tablets IP	Each Film Coated Tablet Contains Linezolid Excipients Approved Colour Used	I.P.	600 mg q.s.		01-09-2017
97	Paracetamol, Phenylephrine Hcl., Diphenhydramine Hcl. & Caffeine Tablets	Each Uncoated Tablet Contains Paracetamol Phenylephrine Hydrochloride Diphenhydramine Hydrochloride Caffeine (Anhydrous) Excipients Approved Colour Used	I.P. I.P. I.P. I.P.	325 mg 5 mg 25 mg 30 mg q.s.		01-09-2017
98	Iron, Folic Acid & Cyanocobalamin Syrup	Each 15 ml Contains Ferric Ammonium Citrate Eq.to Elemental Iron Cyanocobalamin Folic Acid Sorbitol Solution 70 % Flavoured Syrupy Base Colour: Caramel	I.P. I.P. I.P. I.P.	160 mg 32.8 mg 7.5 mcg 0.5 mg q.s. q.s.		01-09-2017
99	Levocetirizine Di Hcl, Phenylephrine Hcl. & Paracetamol Tablet	Each Uncoated Tablet Contains Levocetirizine Dihydrochloride Phenylephrine Hydrochloride Paracetamol Excipients Approved Colour Used	I.P. I.P. I.P.	5 mg 5 mg 325 mg q.s.		01-09-2017
100	Paracetamol Cetirizine Hcl Phenylephrine Hcl. & Tablet	Each Uncoated Tablet Contains Paracetamol Cetirizine Hydrochloride Phenylephrine Hydrochloride Excipients Approved Colour Used	I.P. I.P. I.P.	325 mg 5 mg 5 mg q.s.		01-09-2017
101	Sucralfate & Oxetacaine Suspension	Each 10 ml Contains Sucralfate Oxetacaine Flavoured Palatable Base Colour: Sunset Yellow FCF	U.S.P. B.P.	1 gm 10 mg q.s.		01-09-2017
102	Artemether & Lumafantrine Tablets	Each Uncoated Tablet Contains Artemether Lumafantrine Ph. Int. Excipients	I.P.	80 mg 480 mg q.s.		01-09-2017
103	Ferric Ammonium Citrate , Folic Acid , Cyanocobalamin & Sorbitol Syrup	Each 5 ml Contains Ferric Ammonium Citrate Eq.to Elemental Iron Cyanocobalamin Folic Acid Sorbitol Solution (70 %) (Non Crystallizing) Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	110 mg 22.5 mg 15 mcg 1.5 mg q.s.		01-09-2017
104	Artemether & Lumafantrine Tablets	Each Uncoated Dispersible Tablet Contains Artemether Lumafantrine Ph. Int.	I.P.	20 mg 120 mg		01-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients		q.s.		
105	Lornoxicam & Paracetamol Tablets	Each Film Coated Tablet Contains Lornoxicam Paracetamol Excipients Approved Colour Used	I.P.	8 mg 325 mg q.s.		01-09-2017
106	Levocetirizine & Montelukast Tablets I.P.	Each Film Coated Tablet Contains Levocetirizine Dihydrochloride Montelukast Sodium Eq. to Montelukast Excipients Approved Colour Used	I.P. I.P.	5 mg 10 mg q.s.		01-09-2017
107	Paracetamol & Dicyclomine Hcl. Tablet	Each Uncoated Tablet Contains Paracetamol Dicyclomine Hydrochloride Excipients Approved Colour Used	I.P. I.P.	325 mg 20 mg q.s.		01-09-2017
108	Aceclofenac, Paracetamol & Serratiopeptidase Tablet	Each Film Coated tablet Contain Aceclofenac Paracetamol Serratiopeptidase (Enteric Coated Granules) Approved Colour Used	I.P. I.P. I.P.	100 mg 325 mg 15 mg q.s.		01-09-2017
109	Tranexamic Acid With Mefenamic Acid Tablets	Each Film coated tablet contains Tranexamic Acid Mefenamic Acid Excipients Approved Colour Used	I.P. I.P.	500 mg 250 mg q.s.		01-09-2017
110	Ambroxol Hcl, Levosalbutamol & Guaiphenesin Syrup	Each 5 ml contains Ambroxol hydrochloride Levosulbutamol Sulphate Eq.to Levosalbutamol Guaiphenesin Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	30 mg 1 mg 50 mg q.s.		01-09-2017
111	Multivitamin With Protein Syrup	Each 15 ml contain ferric Ammonium Citrate Vitamin B1 Vitamin B2 Vitamin B6 Vitamin B 12 Protein Hydrolysate 20 % Niacinamide D-Panthenol L-Lysine Hcl Manganese Sulphate Zinc Sulphate Copper Sulphate Sorbitol Solution 70 % flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P. I.P. I.P. I.P. I.P. U.S.P. U.S.P. I.P. B.P.	165 mg 2 mg 2 mg 1 mg 7.5 mcg 1000 mg 15 mg 1.5 mg 30 mg 1 mg 2 mg 2 mg q.s.		01-09-2017
112		Each 5 ml Contains				01-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Vitamin B-Complex With L-Lysine Syrup	Thiamine Hydrochloride Riboflavin Sodium Phosphate Pyridoxine Hydrochloride Vitamin B12 D-Panthenol Niacinamide L-Lysine hydrochloride Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P. I.P. I.P. U.S.P.	0.5 mg 0.5 mg 0.25 mg 0.25 mcg 0.5 mg 7.50 mg 30 mg q.s.		
113	Calcium Carbonate Zinc , Vitamin D3 & Magnesium Suspension	Each 5ml Contains Calcium Carbonate Eq.ro Elemental Calcium Zinc Gluconate Eq.to Elemental Zinc Vitamin D3 Magnesium Hydroxide Eq.to Magnesium Flavoured Palatable Base Approved Colour Used	I.P. U.S.P. I.P. I.P.	625 mg 250 mg 14 mg 2 mg 200 I.U. 180 mg 75 mg q.s.		01-09-2017
114	Aceclofenac, Paracetamol & Thiocholchicoside Tablet	Each Film Coated tablet Contains Aceclofenac Paracetamol Thiocholchicoside Excipients Approved Colour Used	I.P. I.P. I.P.	100 mg 325 mg 8 mg q.s.		01-09-2017
115	Lycopene, Antioxidant, Multivitamins & Multiminerals Syrup	Each 10ml contains: Lycopene 6% Vitamin A (as palmitate) Vitamin E Acetate Ascorbic Acid Sodium Selenite eq. to Selenium Zinc Gluconate eq. to Zinc Manganese Gluconate eq. to Manganese Potassium Iodide eq. to Iodine Cupric Sulphate eq. to Copper Thiamine Hydrochloride Rivoflavin Pyridoxine Hcl In a flavoured Syrupy Base	U.S.P. I.P. I.P. I.P. U.S.P. U.S.P. U.S.P. I.P. U.S.P. I.P. I.P. I.P.	1000 mcg 2500 IU 10 IU 50 mg 35 mcg 3 mg 2 mg 100 mcg 500 mcg 2 mg 3 mg 1.5 mg q.s.		01-09-2017
116	Dextromethorphan Hbr. Phenylephrine Hcl & Chlorpheniramine Maleate Syrup	Each 5 ml Contains Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Chlorpheniramine Maleate Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	15 mg 5 mg 2 mg q.s.		01-09-2017
117	Trypsin, Bromelain & Rutoside Trihydrate & Aceclofenac Tablets	Each Enteric Coated Tablet Contains Trypsin Bromelain Rutoside Trihydrate Eq.to Rutoside Aceclofenac Excipients Approved Colour Used	B.P. B.P. I.P.	48 mg 90 mg 100 mg 100 mg q.s.		01-09-2017
119	Trypsin, Bromelain Rutoside Trihydrate & Diclofenac Sodium	Each Enteric Coated Tablet Contains Trypsin Bromelain	I.P.	48 mg		01-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Tablets	Rutoside Trihydrate Eq.to Rutoside Diclofenac Sodium Excipients Approved Colour Used	B.P. I.P.	90 mg 100 mg 50 mg q.s.		
120	Trypsin, Bromelain & Rutoside Trihydrate Tablets	Each Enteric Coated Tablet Contains Trypsin Bromelain Rutoside Trihydrate Eq.to Rutoside Excipients Approved Colour Used	B.P. B.P.	48 mg 90 mg 100 mg q.s.		01-09-2017
121	Trypsin-Chymotrypsin Tablets	Each Enteric Coated Tablet Contain 10,000 Armour units of enzymatic Activity supplied by a purified concentrate which has specific trypsin and Chymotrypsin activity in a ratio of approximately six to one Excipients Approved Colour Used		q.s.		01-09-2017
122	Trypsin-Chymotrypsin, with Diclofenac Potassium Tablets	Each Enteric Coated Tablet Contain 50000 Armour units of enzymatic Activity supplied by a purified concentrate which has specific trypsin and Chymotrypsin activity in a ratio of approximately six to one Diclofenac Potassium Excipients Approved Colour Used	B.P.	50 mg q.s.		01-09-2017
123	Trypsin-Chymotrypsin, Diclofenac Potassium & Paracetamol Tablets	Each Enteric Coated Tablet Contain 50000 Armour units of enzymatic Activity supplied by a purified concentrate which has specific trypsin and Chymotrypsin activity in a ratio of approximately six to one Diclofenac Potassium Paracetamol Excipients Approved Colour Used	B.P. I.P.	50 mg 325 mg q.s.		01-09-2017
124	Ferrous Ascorbate, Folic Acid & Zinc Sulphate Tablets	Each Film Coated Tablets Contains Ferrous Ascorbate eq. To Elemental Iron Folic acid Zinc Sulphate Monohydrate eq. To Elemental Zinc Excipients Approved colour Used	I.P U.S.P.	100 mg 1.5 mg 22.5 mg q.s		01-09-2017
125	Ferrous Ascorbate, Folic Acid Tablets	Each Film Coated Tablets Contains Ferrous Ascorbate eq. To Elemental Iron Folic acid Excipients Approved Colour Used	I.P.	100 mg 1.5 mg q.s.		01-09-2017
126	Piracetam Tablets	Each Film coated tablet contains Piracetam Excipients Approved Colour Used	I.P.	800 mg q.s.		01-09-2017
127	01-09-2017					



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
128	Gabapentin & Nortriptyline Tablets	Each Film coated tablet contains Gabapentin Nortriptyline Hydrochloride Eq.to Nortriptyline Excipients Approved Colour Used	U.S.P. I.P.	400 mg 10 mg q.s.		01-09-2017
129	Pregabalin & Nortriptyline Tablets	Each Film coated tablet contains Pregabalin Nortriptyline Hydrochloride Eq.to Nortriptyline Excipients Approved Colour Used	I.P. I.P.	75 mg 10 mg q.s.		01-09-2017
130	Levofloxacin & Ornidazole Tablets	Each Film coated tablet contains Levofloxacin Hemihydrate Eq.to Levofloxacin Ornidazole Excipients Colour: Red oxide of Iron	I.P. I.P.	250 mg 500 mg q.s.		01-09-2017
131	Citicoline Tablets I.P. 500 MG	Each Film coated tablet contains Citicoline Sodium Eq.to Citicoline Excipients Approved Colour Used	I.P.	500 mg q.s.		01-09-2017
132	Citicoline With Piracetam Tablets	Each Film coated tablet contains Citicoline Sodium Eq.to Citicoline Piracetam Excipients Approved Colour Used	I.P. I.P.	500 mg 800 mg q.s.		01-09-2017
133	Methylcobalamin & Gabapentin Tablets	Each Film Coated Tablet Contains Methylcobalamin Gabapentin Excipients Colour: Red Oxide Of Iron	U.S.P.	500 mcg 300 mg q.s.		01-09-2017
134	Aceclofenac & Thiocolchicoside Tablet	Each Uncoated Tablet Contains Aceclofenac Thiocolchicoside Excipients Approved Colour Used	I.P. I.P.	100 mg 4 mg q.s.		01-09-2017
135	Tricholine Citrate & Sorbitol Solution	Each 10 ml. Contain: Tricholine citrate (65 % w/v) Sorbitol solution (70 %) (Non crystallising) In a flavour syrupy base Colour sunset yellow FCF	I.P.	550 mg q.s.		01-09-2017
136	Ferrous Ascorbate & Folic Acid Suspension	Each 5 ml Contains Ferrous Ascorbate Eq.to elemental Iron Folic Acid Flavoured Syrupy Base Colour: Caramel	I.P.	30 mg 550 mcg q.s.		01-09-2017
137	Terbutaline sulphate, Bromhexine Hcl, Guaiphenesin & Menthol Syrup	Each 5ml Contains Terbutaline sulphate Bromhexine Hydrochloride Guaiphenesin Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	1.25 mg 4 mg 50 mg 2.5 mg q.s.		01-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
138	Ambroxol Hcl, Levocetirizine Hcl Guaiphenesin & Menthol Syrup	Each 5ml Contains Ambroxol hydrochloride Levocetirizine Hydrochloride Guaiphenesin Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	15 mg 2.5 mg 50 mg 1 mg		01-09-2017
139	Glucosamine Sulphate Potassium Chloride Diacerein & Methyl Sulphonyl Methane Tablets	Each Film Coated Tablet Contains Glucosamine Sulphate Potassium Chloride Eq. To Glucosamine Diacerein Methyl Sulphonyl Methane Excipients Approved Colour Used	U.S.P. I.P. U.S.P.	750 MG 446 mg 50 mg 250 mg q.s.		01-09-2017
140	Aceclofenac, Paracetamol & Serratiopeptidase Tablet	Each Film Coated tablet Contain Aceclofenac Paracetamol Serratiopeptidase (Enteric Coated Granules) Excipients Approved Colour Used	I.P. I.P. I.P.	100 mg 325 mg 10 mg q.s.		01-09-2017
141	Aceclofenac, Paracetamol & Chlorzoxazone Tablet	Each Uncoated tablet Contain Aceclofenac Paracetamol Chlorzoxazone Excipients	I.P. I.P. U.S.P	100 mg 325 mg 250 mg q.s.		01-09-2017
142	Aceclofenac & Paracetamol Tablet	Each Film coated tablet Contain Aceclofenac Paracetamol Excipients Approved Colour Used	I.P. I.P.	100 mg 325 mg q.s.		01-09-2017
143	Diclofenac Potassium & Paracetamol Tablet	Each Uncoated Tablet Contains Paracetamol Diclofenac Potassium Excipients	I.P. B.P.	325 mg 50 mg q.s.		01-09-2017
144	Multi Vitamin & Multi mineral Syrup	Each 5 ml contains Zinc Sulphate Nicotinamide L-Lysine Hcl D-Panthenol Thiamine Hcl Riboflavin Sodium Phosphate Pyridoxine Hcl Copper Sulphate Potassium iodide Sodium Selenate Pentahydrate Eq.to selenium Cyanocobalamin Vitamin A (as Palmitate) Cholecalciferol Vitamin E (as Di A -Tocopheryl Acetate) Flavoured Syrupy Base Approved Colour Used	IP IP USP IP IP IP IP IP BP IP IP IP IP IP	22.2 mg 7.5 mg 5.0 mg 1.25 mg 0.75 mg 0.75 mg 0.5 mg 100 mcg 50 mcg 10 mcg 0.5 mcg 1250 IU 100 IU 2.5 IU q.s.		13-09-2017
145	Fexofenadine Hcl. Tablets I.P. 180 mg	Each Film Coated Tablet Contains Fexofenadine Hydrochloride Excipients Approved Colour Used	I.P.	180 mg q.s.		13-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
146	Methylprednisolone Tablets	Each uncoated tablet contains Methylprednisolone Excipients	I.P.	8 mg q.s.		13-09-2017
147	Ondansetron Orally Disintegrating Tablet I.P.	Each Uncoated Orally Disintegrating Tablets Contain Ondansetron Hydrochloride Eq.to Ondansetron Excipients	I.P.	4 mg q.s.		13-09-2017
148	Flunarizine Tablet B.P. 10 mg	Each uncoated tablet contains Flunarizine (As Flunarizine Dihydrochloride) Excipients Approved Colour Used	B.P.	10 mg q.s.		13-09-2017
149	Cinnarizine & Domperidone Tablets	Each uncoated tablet contains Cinnarizine Domperidone Excipients Approved Colour Used	I.P. I.P.	20 mg 15 mg q.s.		13-09-2017
150	Betamethasone Sodium Phosphate Tablets I.P.	Each Uncoated Tablet Contains Betamethasone Sodium Phosphate Eq. to Betamethasone Excipients Approved Colour Used	I.P.	0.5 mg q.s.		13-09-2017
151	Betamethasone Sodium Phosphate Tablets I.P.	Each Uncoated Tablet Contains Betamethasone Sodium Phosphate Eq. to Betamethasone Excipients Approved Colour Used	I.P.	1 mg q.s.		13-09-2017
152	Calcium Carbonate , Calcitriol & Zinc Tablet	Each Film Coated Tablet Contains Calcium Carbonate Eq.to Elemental Calcium Calcitriol Zinc Sulphate monohydrate Eq.to elemental Zinc Excipients Approved Colour Used	I.P B.P. U.S.P.	1250 mg 500 mg 0.25 mcg 7.5 mg q.s.		13-09-2017
153	Salbutamol Sulphate , Ambroxol Hcl, Guaiphenesin & Menthol Syrup	Each 5ml Contains Salbutamol Sulphate Eq.to Salbutamol Ambroxol hydrochloride Guaiphenesin Menthol Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P. I.P.	1mg 15 mg 50 mg 1 mg q.s		13-09-2017
154	Sulphamethoxazole & Pyrimethamine Tablets	Each Uncoated Tablet Contains Sulphamethoxazole Pyrimethamine Excipients Approved Colour Used	I.P. I.P.	500 mg 25 mg q.s.		13-09-2017
155	Zinc Gluconate Oral Solution	Each 5 ml Contains Zinc Gluconate Eq.to Elemental Zinc Flavoured Syrupy Base Approved Colour Used	U.S.P.	20 mg q.s.		13-09-2017
157	Drotaverine Hcl. & Mefenamic	Each Uncoated Tablet Contains Drotaverine Hydrochloride	I.P.			13-09-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Acid Tablets	Eq.to Drotaverine Mefenamic Acid Excipients Approved Colour Used	I.P.	80 mg 250 mg q.s.		
158	Serratiopeptidase Tablets I.P.	Each Enteric Coated Tablet Contain Serratiopeptidase Excipients Approved Colour Used	I.P.	10 mg q.s.		13-09-2017
160	Methylcobalamin, L-Methylfolate & Pyridoxal 5- Phosphate Tablets	Each Film Coated Tablet Contains Methylcobalamin L-Methylfolate Pyridoxal 5-Phosphate Excipients Approved colour Used	I.P.	1500 mcg 1 mg 0.5 mg q.s.		13-09-20217
161	Itraconazole Capsules 100 mg	Each Hard Gelatin Capsule contains: Itraconazole (in the form of Pellets) Excipients Approved colour used in Empty Capsule Shell	B.P.	100 mg q.s.		13-09-20217
162	Dextromethorphan Hbr. Ambroxol Hcl., Chlorpheniramine Maleate, Phenylephrine Hcl. And Menthol Syrup	Each 5 ml Contains Dextromethorphan Hydrobromide Ambroxol Hydrochloride Chlorpheniramine Maleate Phenylephrine Hydrochloride Menthol Flavoured Syrupy Base Colour: Ponceau 4R Supra & Sunset Yellow Supra	I.P. I.P. I.P. I.P. I.P.	10 mg 15 mg 2 mg 5 mg 1.5 mg q.s.		13-09-20217
163	Methylcobalamin, Alpha Lipoic Acid, Benfotiamine Vitamin B6, Calcium Pantothenate, Nicotinamide & Folic Acid Capsule	Each Hard Gelatin Capsule Contains Methylcobalamin Alpha Lipoic Acid Benfotiamine Vitamin B6 Calcium Pantothenate Nicotinamide Folic Acid Excipients Approved Colour Used In Empty Capsule Shell	U.S.P. I.P. i.p. I.P. I.P.	750 mcg 100 mg 7.5 mg 1.5 mg 25 mg 50 mg 0.75 mg q.s.		13-09-20217
164	Paracetamol Suspension I.P.	Each 5 ml Contains Paracetamol Excipients Flavoured Palatable Base Approved Colour Used	I.P.	120 mg q.s.		13-09-20217
165	Disodium Hydrogen Citrate Syrup	Each 5ml Contains Di Sodium Hydrogen Citrate Flavoured Syrupy Base Approved Colour Used	B.P.	1.40 gm q.s		13-09-20217
166	Mefenamic Acid Suspension	Each 5ml Contains Mefenamic Acid Flavoured Palatable Base Approved Colour Used	I.P.	100 mg q.s.		13-09-20217
167	Ambroxol Hcl, Terbutaline Sulphate & Guaiphenesin Syrup	Each 10 ml Contains Ambroxol Hydrochloride Terbutaline sulphate Guaiphenesin Flavoured Syrupy Base Approved Colour Used	I.P. I.P. I.P.	30 mg 2.5 mg 100 mg q.s.		13-09-20217



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
168	Mefenamic Acid & Paracetamol Suspension	Each 5 ml contains Mefenamic Acid Paracetamol Flavoured Base Colour Sunset Yellow FCF	I.P. I.P.	100 mg 250 mg q.s.		13-09-20217
169	Sucralfate & Oxetacaine Suspension	Each 10 ml Contains Sucralfate Oxetacaine Flavoured Palatable Base Colour: Sunset Yellow FCF	U.S.P. B.P.	1 gm 20 mg q.s.		13-09-20217
170	Diclofenac Potassium, Paracetamol & Chlorzoxazone Tablets	Each Film Coated Tablet Contains Diclofenac Potassium Paracetamol Chlorzoxazone Excipients Approved Colour Used	B.P. I.P. U.S.P. q.s.	50 mg 325 mg 250 mg		22-12-2017
171	Tapentadol Hydrochloride Tablet 50 mg	Each Film Coated Tablet Contain Tapentadol Hydrochloride Eq.to Tapentadol Excipients Approved Colour Used	I.P.	50 mg q.s.		25-10-2018
172	Tapentadol Hydrochloride Tablet 100 mg	Each Film Coated Tablet Contain Tapentadol Hydrochloride Eq.to Tapentadol Excipients Approved Colour Used	I.P.	100 mg q.s.		25-10-2018
173	Amisulpride Tablets I.P. 100mg	Each Uncoated tablet contains Amisulpride Excipients Colour Approved Colour Used	I.P.	100mg q.s.		25-10-2018
174	Levocloperastine Fendizoate Oral suspension	Each 5ml Contains Levocloperastine Fendizoate Eq. to Levocloperastine Excipients Colour Approved colour Used		20 mg q.s.		25-10-2018
175	Fluoxetine Capsule I.P. 20mg	Each Hard Gelation capsule Contains Fluoxetine Hydrochloride Eq. to Fluoxetine Excipients Approved colour used in Empty capsule shell	I.P.	20mg q.s.		25-10-2018
176	Meloxicam Tablets B.P. 7.5mg	Each Uncoated Tablet Contains Meloxicam Excipients Colour Approved colour Used	B.P.	7.5mg q.s.		25-10-2018
177	Deflazacort Tablets 24 mg	Each Uncoated Tablet Contains Deflazacort Excipients Colour Approved colour Used		24mg q.s.		25-10-2018
178	Deflazacort Tablets 18 mg	Each Uncoated Tablet Contains Deflazacort Excipients Colour Approved colour Used		18mg q.s.		25-10-2018
179	Amitriptyline Hydrochloride Tablets I.P. 25mg	Each Film Coated Tablet Contains Amitriptyline Hydrochloride Excipients Colour: approved Colour Used	I.P.	25mg q.s.		25-10-2018
180	Hydroxyzine Oral Solution I.P.	Each 5 ml Contains Hydroxyzine Hydrochloride Flavored Syrupy Base Colour: Approved Colour Used	I.P.	10 mg q.s.		25-10-2018



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
181	Etodolac & Thiocolchicoside Tablets	Each Film Coated Tablet Contains Etodolac Thiocolchicoside Excipients Approved Colour Used	I.P. I.P.	400 mg 4 mg q.s.		25-10-2018
182	Deflazacort Tablets 30 mg	Each Uncoated Tablet Contains Deflazacort Excipients		30 mg q.s.		25-10-2018
183	Etoricoxib & Paracetamol Tablets	Each Film Coated Tablet Contains Paracetamol Etoricoxib Excipients Colour; Approved Colour Used	I.P. I.P.	325mg 60mg q.s.		25-10-2018
184	Ferrous Ascorbate, Folic Acid Methylcobalamin and Zinc tablets	Each Film Coated Tablet Contains Ferrous Ascorbate Eq.to Elemental Iron Folic Acid Zinc Sulphate Monohydrate Eq.to Elemental Zinc Methylcobalamin Excipients Colour Approved Colour Used	I.P. I.P. I.P.	100mg 1.1 mg 22.5 mg 1.5 mg q.s.		25-10-2018
185	Methylcobalamin, Calcium , Calcitriol ,Folic Acid , Vitamin B6 & Zinc Capsule	Each Hard Gelation capsule Contains Methylcobalamin Calcium Carbonate Eq.to Elemental Calcium Folic Acid Calcitriol Vitamin B6 Zinc Sulphate Monohydrate Eq.to Elemental Zinc Excipients Approved colour used In empty capsule shell	I.P. I.P. I.P. I.P. I.P.	500 mcg 250mg 1.5 mg 0.25 mcg 3 mg 7.5 mg q.s.		25-10-2018
186	Etoricoxib Tablets I.P. 60mg	Each Film Coated Tablet Contains Etoricoxib Excipients Colour: Approved Colour Used	I.P.	60mg q.s.		25-10-2018
187	Etoricoxib Tablets I.P. 120mg	Each Film Coated Tablet Contains Etoricoxib Excipients Colour: Approved Colour Used	I.P.	120mg q.s.		25-10-2018
188	Mirtazapine Tablets I.P. 15 mg	Each Film Coated Tablet Contains Mirtazapine Excipients Colour: approved Colour Used	I.P.	15mg q.s.		25-10-2018
189	Chlorpromazine Tablets I.P. 100mg	Each Film Coated Tablet Contains Chlorpromazine Hydrochloride Excipients Colour: approved Colour Used	I.P.	100mg q.s.		25-10-2018
190	Clozapine Tablets I.P. 100 mg	Each Uncoated tablet contains Clozapine Excipients Colour Approved Colour Used	I.P.	100mg q.s.		25-10-2018
191	Phenytoin Sodium Tablets I.P. 100mg	Each Film Coated Tablet Contains Phenytoin Sodium Excipients Colour Approved Colour Used	I.P.	100mg q.s.		25-10-2018
192	Sertraline Hydrochloride Tablets I.P. 100mg	Each Uncoated Tablet Contains Sertraline Hydrochloride E.q.to Sertraline	I.P.	100mg		25-10-2018



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients Colour Approved Colour Used		q.s.		
193	Tapentadol Hydrochloride Extended Release Tablet	Each Film Coated Extended Release Tablet Contain Tapentadol Hydrochloride Eq.to Tapentadol Excipients Approved Colour Used	I.P.	50 mg q.s.		25-10-2018
194	Norfloxacin & Tinidazole Tablets	Each Film Coated Tablet Contains Norfloxacin Tinidazole Excipients Colour: Tartrazine & Titanium Dioxide IP	I.P. I.P.	400mg 600mg q.s.		25-10-2018
195	Cetirizine HCL Syrup I.P.	Each 5ml Contains Cetirizine Hydrochloride Flavoured Syrupy Base Colour Approved Colour Used	I.P.	5mg q.s.		16-01-2019
196	Esomeprazole (E.C.) & Domperidone (S.R.) Capsules	Each hard Gelatin capsule contains: Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (as enteric coated pellets) Domperidone (As Sustained Release Pellets) Approved colour used in empty capsule shell & pellets	I.P. I.P.	40mg 30 mg		16-01-2019
197	Metformin HCL. (S.R.) & Voglibose Tablets	Each Uncoated Bilayered Tablet contains Metformin Hydrochloride (As Sustained Release) Voglibose Excipients Colour Approved Colour Used	I.P. I.P.	500mg 0.2 mg q.s.		16-01-2019
198	Metformin HCL. (S.R.) & Voglibose Tablets	Each Uncoated Bilayered Tablet Contains Metformin Hydrochloride (As Sustained Release) Voglibose Excipients Colour Approved Colour Used	I.P. I.P.	500mg 0.3 mg q.s.		16-01-2019
199	Pregabalin (SR) & Methylcobalamin Tablets	Each Film coated Tablet Contains Pregabalin (As Sustained Release) Methylcobalamin Excipients Colour : Approved Colour Used	I.P. I.P.	150mg 1500mcg q.s.		16-01-2019
200	Telmisartan & Hydrochlorothiazide Tablets I.P.	Each Uncoated Tablet Contains Telmisartan Hydrochlorothiazide Excipients Approved colour Used	I.P. I.P.	40 mg 12.5 mg q.s.		16-01-2019
201	Atenolol & Amlodipine Tablets	Each Uncoated Tablet Contains Atenolol Amlodipine Besylate Eq. to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	50mg 5 mg q.s.		16-01-2019
202	Dextromethorphan Hbr. , Ammonium Chloride and Bromhexine Hcl. Syrup	Each 5 ml Contains Dextromethorphan Hydrobromide Bromhexine Hydrochloride Ammonium Chloride Flavored Syrupy Base	I.P. I.P. I.P.	5 mg 4 mg 50 mg q.s.		16-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Colour: Approved Colour Used				
203	Flavoxate Hydrochloride Tablets I.P. 200mg	Each Film Coated tablets Contains Flavoxate Hydrochloride Excipients Colour: Titanium Dioxide IP	I.P.	200mg q.s.		16-01-2019
204	Olmesartan Tablets I.P. 20 mg	Each film coated tablets contains Olmesartan Medoxomil Excipients Colour: Titanium Dioxide IP	I.P.	20 mg q.s.		16-01-2019
205	Olmesartan Tablets I.P. 40 mg	Each film coated tablets contains Olmesartan Medoxomil Excipients Colour: Titanium Dioxide IP	I.P.	40 mg q.s.		16-01-2019
206	Nitrofurantoin Sustained Release Tablets I.P. 100 mg	Each Film Coated Sustained Release Tablet contains Nitrofurantoin Excipients Colour: Approved Colour Used	I.P.	100mg q.s.		16-01-2019
207	Rosuvastatin Tablets I.P. 10 mg	Each Film Coated Tablets Contains Rosuvastatin Calcium Eq. to Rosuvastatin Excipients Colour Approved Colour Used	I.P.	10mg q.s.		16-01-2019
208	Rosuvastatin Tablets I.P. 5 mg	Each Film Coated Tablets Contains Rosuvastatin Calcium Eq. to Rosuvastatin Excipients Colour Approved Colour Used	I.P.	5mg q.s.		16-01-2019
209	Pregabalin (SR) & Methylcobalamin Tablets	Each Film coated Tablet Contains Pregabalin (As Sustained Release) Methylcobalamin Excipients Colour :Approved Colour Used	I.P. I.P.	75mg 1500mcg q.s.		16-01-2019
210	Glimepiride & Metformin Hydrochloride (Sustained release) Tablets	Each Uncoated tablet contains: Glimepiride Metformin Hydrochloride (As Sustained release) Excipients Colour: Approved Colour Used	I.P. I.P.	1 mg 500mg q.s.		16-01-2019
211	Voglibose Mouth Dissolving Tablets 0.2 mg	Each Uncoated Mouth Dissolving tablet contains: Voglibose Excipients Colour: Approved Colour Used	I.P	0.2mg q.s.		16-01-2019
212	Voglibose Mouth Dissolving Tablets 0.3 mg	Each Uncoated Mouth Dissolving tablet contains: Voglibose Excipients Colour: Approved Colour Used	I.P	0.3mg q.s.		16-01-2019
213	Metformin Hydrochloride Sustained Release Tablets I.P. 500 mg	Each film coated sustained release tablet contains: Metformin Hydrochloride Excipients Colour: Red Oxide Of Iron	I.P.	500 mg q.s.		16-01-2019
214	Metformin Hydrochloride Sustained Release Tablets I.P. 1000 mg	Each film coated sustained release tablet contains: Metformin Hydrochloride Excipients Colour: Red Oxide Of Iron	I.P.	1000 mg q.s.		16-01-2019
215		Each Uncoated tablet contains:				16-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Glimepiride & Metformin Hydrochloride (Sustained release) Tablets	Glimepiride Metformin Hydrochloride (As Sustained release) Excipients Colour: Approved Colour Used	I.P. I.P.	2 mg 500mg q.s.		
216	Diphenhydramine HCL, Ammonium Chloride, Sodium Citrate & Menthol Syrup	Each 5ml Contains Diphenhydramine Hydrochloride Ammonium Chloride Sodium Citrate Menthol Flavoured Syrupy Base Colour : Approved Colour Used	I.P. I.P. I.P. I.P.	14.08mg 138 mg 57.03 mg 1.14 mg q.s.		16-01-2019
217	Atorvastatin Calcium & Fenofibrate Tablets	Each film coated tablet contains: Atorvastatin Calcium Eq. to Atorvastatin Fenofibrate Excipients Colour: Approved Colour Used	I.P. I.P.	10mg 160 mg q.s.		16-01-2019
218	Rosuvastatin & Fenofibrate Tablets I.P.	Each Film Coated Tablets Contains Rosuvastatin Calcium Eq. to Rosuvastatin Fenofibrate Excipients Approved Colour Used	I.P. I.P.	10mg 160 mg q.s.		16-01-2019
219	Olanzapine & Fluoxetine Tablets	Each Uncoated Tablet Contains Olanzapine Fluoxetine Hydrochloride Eq.to Fluoxetine Excipients Colour: Approved Colour Used	I.P. I.P.	5 mg 20mg q.s.		16-01-2019
220	Olanzapine Mouth Dissolving Tablets	Each Uncoated Mouth Dissolving Tablets Contain Olanzapine Excipients Colour: Approved Colour Used	I.P.	2.5 mg q.s.		16-01-2019
221	Sodium Valproate Oral Solution I.P.	Each 5 ml Contains Sodium Valproate Flavoured base Colour: Ponceau 4R	I.P.	200mg q.s.		16-01-2019
222	Telmisartan Tablets I.P. 80 mg	Each Uncoated Tablet Contains Telmisartan Excipients Approved colour Used	I.P.	80 mg q.s.		16-01-2019
223	Clopidogrel Tablets I.P. 75 mg	Each Film Coated Tablets Contains Clopidogrel Bisulphate Eq.to Clopidogrel Excipients Approved Colour Used	I.P.	75 mg q.s.		16-01-2019
224	Lorazepam Tablets I.P. 2mg	Each Uncoated tablet contains Lorazepam Excipients Colour Approved Colour Used	I.P.	2 mg q.s.		16-01-2019
225	Citicoline & Piracetam Syrup	Each 5ml Contains Citicoline Sodium Eq. to Citicoline Piracetam Flavoured Base Colour: Approved Colour Used	I.P. I.P.	500mg 400mg q.s.		16-01-2019
226	Telmisartan &	Each uncoated tablet contains: Telmisartan	I.P.	40mg		16-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Amlodipine Tablets I.P.	Amlodipine Besylate Eq. to Amlodipine Excipients: Colour: Approved Colour Used	I.P.	5 mg q.s.		
227	Paracetamol Tablets I.P. 500 mg	Each Uncoated Tablets Contains Paracetamol Excipients	I.P.	500mg q.s.		16-01-2019
228	Paracetamol Tablets I.P. 650 mg	Each Uncoated Tablets Contains Paracetamol Excipients	I.P.	650 mg q.s.		16-01-2019
229	Atorvastatin Tablets I.P. 10 mg	Each film coated tablet contains: Atorvastatin Calcium Eq.to Atorvastatin Excipients Colour : Approved Colour Used	I.P.	10 mg q.s.		16-01-2019
230	Atorvastatin Tablets I.P. 20 mg	Each film coated tablet contains: Atorvastatin Calcium Eq.to Atorvastatin Excipients Colour Titanium Dioxide I.P.	I.P.	20 mg q.s.		16-01-2019
231	Atorvastatin Tablets I.P. 40 mg	Each film coated tablet contains: Atorvastatin Calcium Eq.to Atorvastatin Excipients Colour : Approved Colour Used	I.P.	40 mg q.s.		16-01-2019
232	Olanzapine & Fluoxetine Tablets I.P.	Each Uncoated Tablet Contains Olanzapine Fluoxetine Hydrochloride Eq.to Fluoxetine Excipients Colour: Approved Colour Used	I.P. I.P.	10 mg 20mg q.s.		16-01-2019
233	Telmisartan & Metoprolol (E.R.) Tablets	Each film coated tablet contains: Telmisartan Metoprolol Succinate Eq. to Metoprolol Tartrate (as extended release) Excipients: Colour: Red Oxide Of Iron	I.P. I.P.	40 mg 50 mg q.s.		16-01-2019
236	Doxycycline & lactic Acid Bacillus capsule	Each Hard Gelatin capsule Contains Doxycycline Hyclate Eq.to Doxycycline (As Pellets) Lactic Acid bacillus (As enteric Coated Pellets) Excipients Approved Colour Used in Empty Capsule Shell	I.P.	100mg 5 Billion Spore q.s.		16-01-2019
237	Nebivolol Tablet I.P. 5 mg	Each Uncoated Tablet Contains Nebivolol Hydrochloride Eq.to Nebivolol Excipients Colour : Approved Colour Used	I.P.	5 mg q.s.		16-01-2019
238	Terbinafine Tablets I.P. 250 mg	Each Uncoated Tablets Contains Terbinafine Hydrochloride Eq.to Terbinafine Excipients Colour Approved Colour Used	I.P.	250mg q.s		16-01-2019
239	Torsemide Tablet I.P.	Each Uncoated Tablet Contains Torsemide Excipients Colour Approved Colour Used	I.P.	5 mg q.s.		16-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
240	Amlodipine Tablets I.P. 10 mg	Each Uncoated Tablets Contains Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P.	10 mg q.s.		16-01-2019
241	S-Amlodipine Tablets I.P. 2.5 mg	Each Uncoated Tablets Contains S-Amlodipine Besylate Eq.to S-Amlodipine Excipients Colour Approved Colour Used	I.P.	2.5mg q.s.		16-01-2019
242	S-Amlodipine Tablets I.P. 5 mg	Each Uncoated Tablets Contains S-Amlodipine Besylate Eq.to S-Amlodipine Excipients Colour Approved Colour Used	I.P.	5mg q.s.		16-01-2019
243	Nebivolol & Amlodipine Tablets	Each Uncoated Tablet Contains Nebivolol Hydrochloride Eq.to Nebivolol Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	5 mg 5 mg q.s.		16-01-2019
244	Nebivolol & S- Amlodipine Tablets	Each Uncoated Tablet Contains Nebivolol Hydrochloride Eq.to Nebivolol S- Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	5 mg 2.5 mg q.s.		16-01-2019
245	Atorvastatin & Aspirin (E.C.) Capsules I.P.	Each Hard Gelatin Capsule Contains Atorvastatin Calcium Eq. to Atorvastatin Aspirin (as enteric coated Pellets) Excipients Approved Colour Used in capsule shell	I.P. I.P.	10mg 75 mg q.s.		16-01-2019
246	Esomeprazole (E.C.) & Levosulpiride (S.R.)Capsule	Each Hard Gelatin capsule Contains Esomeprazole Magnesium Trihydrate Eq.to Esomeprazole (As Enteric Coated Pellets) Levosulpiride (As Sustained Release Pellets) Excipients Approved Colour Used in Capsule Shell	I.P.	40 mg 75 mg q.s		16-01-2019
247	Rosuvastatin & Aspirin (E.C.) Capsule I.P.	Each Hard Gelatin Capsule Contains Rosuvastatin Calcium Eq.to Rosuvastatin (As Pellets) Aspirin (As Enteric Coated pellets) Excipients Approved Colour in Empty Capsule shell	I.P. I.P.	10 mg 75 mg q.s.		16-01-2019
249	Sertraline Hydrochloride Tablet I.P. 25 mg	Each Uncoated Tablets Contains Sertraline Hydrochloride Eq.to Sertraline Excipients Approved Colour Used	I.P.	25 mg q.s.		15-08-2018
250	Sertraline Hydrochloride Tablet I.P. 50 mg	Each Uncoated Tablets Contains Sertraline Hydrochloride Eq.to Sertraline Excipients Approved Colour Used	I.P.	50 mg q.s.		15-08-2018



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
253	Rabeprazole 20 mg Capsules	Each Hard Gelatin Capsule Contains Rabeprazole Sodium (As Enteric Coated Pellets) Excipients Approved Colour used in empty capsule shell	I.P.	20 mg q.s.		15-08-2018
254	Desloratadine Tablets U.S.P. 10 mg	Each Film Coated Tablet Contain Desloratadine Excipients Approved Colour Used	U.S.P.	10 mg q.s.		15-08-2018
257	Lactulose Oral Solution U.S.P.	Each 15 Contains Lactulose Concentrate Eq.to Lactulose In a flavoured Base	U.S.P.	10 gm q.s.		15-08-2018
258	Diclofenac Sodium & Serratiopeptidase Tablets	Each Enteric Coated Tablet Contains Diclofenac Sodium Serratiopeptidase Excipients Approved Colour Used	IP IP	50mg 10mg q.s.		15-08-2018
259	Aceclofenac & Drotaverine Hcl. Tablets	Each Film Coated Tablet Contains Aceclofenac Drotaverine Hydrochloride Excipients Approved Colour Used	I.P. I.P.	100mg 80mg q.s.		15-08-2018
260	Diclofenac Sodium & Chlorzoxazone Tablets	Each Film Coated Tablet Contains Diclofenac Sodium Chlorzoxazone Excipients Approved Colour Used	I.P. U.S.P.	50 mg 500mg		15-08-2018
262	Escitalopram Tablets I.P. 10mg	Each Film Coated Tablet Contains Escitalopram Oxalate Eq.to Escitalopram Excipients Approved Colour Used	I.P.	10 mg q.s.		15-08-2018
263	Escitalopram Tablets I.P. 20 mg	Each Film Coated Tablet Contains Escitalopram Oxalate Eq.to Escitalopram Excipients Approved Colour Used	I.P.	20 mg q.s.		15-08-2018
264	Olanzapine Tablets I.P. 2.5mg	Each Uncoated Tablet Contains Olanzapine Excipients Approved Colour Used	I.P.	2.5 mg q.s.		15-08-2018
265	Olanzapine Tablets I.P.5 mg	Each Uncoated Tablet Contains Olanzapine Excipients Approved Colour Used	I.P.	5 mg q.s.		15-08-2018
266	Olanzapine Tablets I.P. 10mg	Each Uncoated Tablet Contains Olanzapine Excipients Approved Colour Used	I.P.	10 mg q.s.		15-08-2018
268	Labetalol Hydrochloride Tablets IP 100 mg	Each Film Coated Tablet Contains Labetalol Hydrochloride Excipients Approved Colour Used	I.P.	100 mg q.s.		15-08-2018
269	Cyproheptadine Hcl. Syrup I.P.	Each 5 ml Contains Cyproheptadine Hydrochloride Eq.to Anhydrous Cyproheptadine In a Flavoured Syrupy Base Approved Colour Used	I.P.	2 mg q.s.		15-08-2018
270	Phenylephrine Hcl., Chlorpheniramine maleate	Each 5ml Contains Phenylephrine Hydrochloride	IP	5 mg		15-08-2018



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	Paracetamol, Sodium Citrate & Menthol syrup	Chlorpheniramine maleate Paracetamol Sodium Citrate Menthol Flavoured Syrupy Base Approved Colour Used	IP IP IP IP	0.5 mg 125 mg 60 mg 1 mg q.s		
271	Sodium Picosulphate , Milk of Magnesia & Liquid Paraffin Suspension	Each 5 ml Contains Liquid Paraffin Milk of Magnesia Sodium Picosulphate Colour: Erythrosine Supra	I.P. I.P. I.P.	1.25 ml 3.75 ml 3.33 mg		15-08-2018
272	Ursodeoxycholic Acid Tablets I.P. 300 mg	Each Film Coated Tablet Contains Ursodeoxycholic Acid Excipients Approved colour Used	I.P.	300 mg q.s.		16-08-2018
273	Glimepiride & Metformin Hcl (Sustained Release) Tablets I.P.	Each Uncoated Bilayered Tablet Contains Glimepiride Metformin Hydrochloride (As Sustained Release) Excipients Colour Approve Colour Use	I.P. I.P.	1mg 500 mg q.s.		11-09-2020
274	Glimepiride & Metformin Hcl (Sustained Release) Tablets I.P.	Each Uncoated Bilayered Tablet Contains Glimepiride Metformin Hydrochloride (As Sustained Release) Excipients Colour Approve Colour Use	I.P. I.P.	2mg 500 mg q.s.		11-09-2020
275	Etoricoxib Tablets I.P. 90 mg	Each Film Coated tablets Etoricoxib Excipients Colour Approved Colour Used	I.P.	90 mg q.s.		11-09-2020
276	Losartan Potassium & Hydrochlorothiazide Tablets I.P.	Each Film Coated Tablets Contains Losartan Potassium Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P. I.P.	50 mg 12.5 mg q.s.		11-09-2020
277	Montelukast & Levocetirizine Dispersible Tablets	Each Uncoated Dispersible Tablet Contains:- Montelukast Sodium Eq. to Montelukast Levocetirizine Hydrochloride Excipients Approved colour used.	I.P. I.P.	5mg 2.5 mg q.s.		11-09-2020
278	Paracetamol , Phenylephrine Hydrochloride & Chlorpheniramine Maleate Tablets	Each Uncoated Tablets Contains Paracetamol Phenylephrine hydrochloride Chlorpheniramine Maleate Excipients Colour approved colour Used	I.P. I.P. I.P.	500mg 10 mg 2 mg q.s.		11-09-2020
279	Paracetamol & Mefenamic Acid Tablets	Each Uncoated Tablets Contains Paracetamol Mefenamic Acid Excipients Colour Approved Colour Used	I.P. I.P.	325 mg 500mg		11-09-2020
280	Metformin Hcl (S.R.) & Glimepiride Tablets I.P.	Each Uncoated bilayer Tablet Contains Metformin Hydrochloride As Sustained Release) Glimepiride	I.P. I.P.	1000 mg 2 mg q.s.		11-09-2020



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients Colour Approved Colour Used				
281	Metformin Hcl (S.R.) & Glimepiride Tablets I.P.	Each Uncoated bilayer Tablet Contains Metformin Hydrochloride As Sustained Release) Glimepiride Excipients Colour Approved Colour Used	I.P. I.P.	1000 mg 1 mg q.s.		11-09-2020
282	Vildagliptin Tablets I.P. 50 mg	Each Uncoated Tablets Contains Vildagliptin Excipients Colour approved Colour Used	I.P.	50 mg q.s.		11-09-2020
283	Lornoxicam & Paracetamol Tablets	Each film coated tablet contains:- Lornoxicam Paracetamol Excipients Approved colour used.	I.P. I.P.	4 mg 325 mg q.s.		11-09-2020
284	Metformin Hydrochloride Tablets I.P. 500 mg	Each Uncoated Tablets Contains Metformin Hydrochloride Excipients Colour Approved Colour Used	I.P.	500 mg q.s.		11-09-2020
285	Metformin Hydrochloride Tablets I.P. 1000 mg	Each Uncoated Tablets Contains Metformin Hydrochloride Excipients Colour Approved Colour Used	I.P.	1000 mg q.s.		11-09-2020
286	Vildagliptin & Metformin Hcl Tablets I.P.	Each Film Coated Tablets Contains Vildagliptin Metformin Hydrochloride Excipients Colour approved Colour Used	I.P. I.P.	50 mg 500mg q.s.		11-09-2020
287	Vildagliptin & Metformin Hcl Tablets I.P.	Each Film Coated Tablets Contains Vildagliptin Metformin Hydrochloride Excipients Colour approved Colour Used	I.P. I.P.	50 mg 850mg q.s.		11-09-2020
288	Vildagliptin & Metformin Hcl. Tablets I.P.	Each Film Coated Tablets Contains Vildagliptin Metformin Hydrochloride Excipients Colour approved Colour Used	I.P. I.P.	50 mg 1000mg q.s.		11-09-2020
289	Levocetirizine Hydrochloride Syrup	Each 5ml Contains Levocetirizine Hydrochloride Flavored Syrupy Base Colour approved Colour used	I.P.	2.5mg q.s.		11-09-2020
290	Telmisartan & Amlodipine Tablets I.P.	Each Film Coated tablet contains Telmisartan Amlodipine Besylate Eq. to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	40 mg 5 mg q.s.		11-09-2020
291	Etodolac & Thiocolchicoside Tablets	Each film coated tablet contains:- Etodolac Thiocolchicoside Excipients Approved colour used.	I.P. I.P.	300 mg 8 mg q.s.		11-09-2020
292	Ursodeoxycholic Acid Tablets 150mg I.P.	Each Uncoated Tablet Contains Ursodeoxycholic Acid Excipients Approved Colour Used	I.P.	150mg		11-09-2020
293		Each Uncoated tablet Contains				11-09-2020



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Nimesulide ,Paracetamol , Phenylephrine Hcl. & Caffeine Tablets	Nimesulide Paracetamol Phenylephrine Hydrochloride Caffeine Excipients Colour Approved Colour Used	B.P. I.P. I.P. I.P.	100mg 325mg 10mg 25mg q.s.		
294	Carbonyl Iron , Folic Acid & Zinc Capsule	Each Hard Gelatine Capsule contains Elemental Iron (In the form of Carbonyl iron) Folic Acid Zinc Sulphate Monohydrate (Eq.to 22.5 mg Elemental Zinc) Excipients Approved Colour used in Empty Capsule shell	I.P. I.P.	50 mg 0.5 mg 61.8 mg q.s.		11-09-2020
295	Nimesulide, Phenylephrine Hcl.,Caffeine & Cetirizine Hcl. Tablets	Each Uncoated tablet Contain Nimesulide Phenylephrine Hydrochloride Caffeine (Anhydrous) Cetirizine Hydrochloride Excipients Colour Approved Colour Used	B.P. I.P. I.P. I.P.	100 mg 10 mg 30 mg 5 mg q.s.		11-09-2020
296	Tamsulosin Hydrochloride Prolonged- Release Capsules IP	Each Hard gelatin Capsule Contains Tamsulosin Hydrochloride (As Prolonged Release pellets) Excipients Approved Colour Used in empty capsule shell	I.P.	0.4 mg q.s.		11-09-2020
297	Amlodipine, Telmisartan & Hydrochlorothiazide Tablets.	Each Uncoated tablet contains:- Amlodipine Besylate Eq. to Amlodipine Telmisartan Hydrochlorothiazide Excipients Colour: Approved Colour used	I.P. I.P. I.P.	5 mg 40 mg 12.5 mg q.s.		11-09-2020
298	Calcitriol , Calcium & Zinc Capsules	Each Hard Gelatin Capsule Contains:- Calcitriol Calcium Carbonate Eq. to elemental Calcium Zinc Sulphate Monohydrate Eq. to Zinc Excipients Approved colour used in capsule shell.	I.P. I.P. I.P.	0.25 200 mg 7.5 mg q.s.		11-09-2020
299	Thiocolchicoside & Etoricoxib Tablets	Each film coated tablet contains:- Thiocolchicoside Etoricoxib Excipients Approved colour used.	I.P. I.P.	4 mg 60 mg q.s.		11-09-2020
300	Etodolac & Thiocolchicoside Tablets	Each film coated tablet contains:- Etodolac Thiocolchicoside Excipients Approved colour used.	I.P. I.P.	300 mg 4 mg q.s.		11-09-2020
301	Loratadine Tablets USP 10mg	Each Uncoated tablet Contains Loratadine Excipients Approved Colour used	U.S.P.	10mg		11-09-2020
302	Telmisartan & Hydrochlorothiazide Tablets IP	Each Un coated tablet Contains Telmisartan Hydrochlorothiazide Excipients	I.P. I.P.	80 mg 12.5 mg		11-09-2020



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Approved Colour Used				
303	Lornoxicam Tablets 4 mg	Each Film Coated tablet Contains Lornoxicam Excipients Approved Colour Used		4 mg q.s.		11-09-2020
304	Lornoxicam Tablets 8 mg	Each Film Coated tablet Contains Lornoxicam Excipients Approved Colour Used		8 mg q.s.		11-09-2020
305	Levocetirizine Hydrochloride Tablets I.P. 10 mg	Each Film Coated tablet Contains Levocetirizine Hydrochloride Excipients Approved Colour Used	I.P.	10 mg q.s.		11-09-2020
306	Lansoprazole Orally Disintegrating Tablets 15 mg	Each Uncoated Tablet Contains: Lansoprazole (as enteric coated pellets) Excipients Colour: Sunset Yellow FCF	I.P.	15mg q.s		19-01-2019
307	Paracetamol Sustained Release Tablets 1000mg	Each Uncoated Bilayer Tablets Contains Paracetamol (as immediate release) Paracetamol (as sustained release) Excipients Colour: Approved Colour Used	I.P. I.P.	300 mg 700 mg q.s.		19-01-2019
308	Sildenafil Citrate Tablets I.P. 50 mg	Each Film Coated Tablet Contains Sildenafil Citrate Eq.to Sildenafil Excipients Colour: Approved Colour used	I.P.	50 mg q.s		19-01-2019
309	Sildenafil Citrate Tablets I.P. 100 mg	Each Film Coated Tablet Contains Sildenafil Citrate Eq.to Sildenafil Excipients Colour: Approved Colour used	I.P.	100 mg q.s		19-01-2019
310	Silodosin Capsules 4 mg	Each Hard Gelatine Capsule Contain Silodosin Excipients Approved Colour used in empty capsule shell		4mg q.s.		19-01-2019
311	Silodosin Capsules 8 mg	Each Hard Gelatine Capsule Contain Silodosin Excipients Approved Colour used in empty capsule shell		5mg q.s.		19-01-2019
312	Levocetirizine Hcl. & Montelukast Sodium Dispersible tablets	Each Uncoated Dispersible Tablet Contains Levocetirizine Hydrochloride Montelukast Sodium Eq.to Montelukast Excipients Colour Approved Colour Used	I.P. I.P.	2.5 mg 4 mg q.s.		19-01-2019
313	Flupirtine Maleate Capsules 100mg	Each Hard Gelatine Capsule Contain Flupirtine Excipients Approved Colour Used in Empty Capsule shell		100 mg q.s.		19-01-2019
314	Metformin HCL. (S.R.) , Glimepiride & Pioglitazone	Each Uncoated Bilayer Tablet Contains Metformin Hydrochloride (As Sustained Release)	I.P.	500 mg		19-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Tablets	Glimepiride Pioglitazone Hydrochloride Eq.to Pioglitazone Excipients Colour Approved Colour Used	I.P. I.P.	2 mg 15 mg q.s.		
315	Metformin HCL. (S.R.) , Glimepiride & Pioglitazone Tablets	Each Uncoated Bilayer Tablet Contains Metformin Hydrochloride (As Sustained Release) Glimepiride Pioglitazone Hydrochloride Eq.to Pioglitazone Excipients Colour Approved Colour Used	I.P. I.P. I.P.	500 mg 1 mg 15 mg q.s.		19-01-2019
316	Enteric Coated Rabeprazole Sodium & Domperidone Sustained Release Capsule	Each Hard Gelatin Capsule Contains Rabeprazole Sodium (As Enteric Coated Pallets) Domperidone (As Sustained Release Pallets) Excipients Approved Colour used un Empty Capsule Shell	I.P. I.P.	20 mg 30 mg q.s		01-09-2017
317	Montelukast Sodium & Desloratadine Tablets	Each Film Coated tablet Contains Montelukast Sodium Eq. to Montelukast Desloratadine Excipients Colour Approved Colour Used	I.P.	10mg 5 mg q.s.		19-01-2019
318	Cetirizine Hcl., Phenylephrine Hcl. & Paracetamol Suspension	Each 5ml Contains Cetirizine Hydrochloride Phenylephrine Hydrochloride Paracetamol Flavoured base Colour Approved Colour Used	I.P. I.P. I.P.	2.5 mg 2.5 mg 250 mg q.s.		19-01-2019
319	Losartan Tablets I.P. 50 mg	Each Film coated Tablet Contains Losartan Potassium Excipients Colour Approved Colour Used	I.P.	50 mg q.s.		19-01-2019
320	Losartan Tablets I.P. 25 mg	Each Film coated Tablet Contains Losartan Potassium Excipients Colour Approved Colour Used	I.P.	25 mg q.s.		19-01-2019
321	Acarbose Tablet IP 50 mg	Each Uncoated tablet Contains Acarbose Excipients Colour Approved Colour Used	I.P.	50 mg q.s.		19-01-2019
322	Acarbose Tablet IP 100 mg	Each Uncoated tablet Contains Acarbose Excipients Colour Approved Colour Used	I.P.	100 mg q.s.		19-01-2019
323	Losartan & Amlodipine Tablets I.P.	Each Film Coated Tablet Contains Losartan Potassium Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	50 mg 5mg q.s.		19-01-2019
324	Amisulpride Tablet IP 200 mg	Each Uncoated tablet Contains Amisulpride Excipients Colour Approved Colour Used	I.P.	200 mg q.s.		19-01-2019
325	Gabapentin tablets IP 300 mg	Each Film Coated Tablet Contains Gabapentin Excipients	I.P.	300 mg q.s.		19-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Colour Approved Colour Used				
326	Metoprolol Prolonged Release Tablets IP 25mg	Each Film coated Prolonged release Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate Excipients Colour Approved colour Used	I.P.	23.75 mg 25mg q.s.		19-01-2019
327	Metoprolol Prolonged Release Tablets IP 50mg	Each Film coated Prolonged release Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate Excipients Colour Approved colour Used	I.P.	47.5 mg 50 mg q.s.		19-01-2019
328	Levetiracetam Prolonged Release Tablets IP 500mg	Each Film Coated Prolonged Release Tablet Contains Levetiracetam (As Prolonged Release) Excipients Colour Approved Colour Used	I.P.	500mg q.s.		19-01-2019
329	Levetiracetam Tablets IP 250mg	Each Film Coated Tablet Contains Levetiracetam Excipients Colour Approved Colour Used	I.P.	250mg q.s.		19-01-2019
330	Telmisartan Tablet IP 20mg	Each Uncoated Tablet Contains Telmisartan Excipients Colour Approved Colour Used	I.P.	20 mg q.s.		19-01-2019
331	Rosuvastatin Tablets IP 20 mg	Each Film Coated Tablet Contains Rosuvastatin Calcium Eq.to Rosuvastatin Excipients Colour Approved Colour Used	I.P.	20 mg q.s.		19-01-2019
332	Olmesartan Medoxomil & Hydrochlorothiazide Tablet I.P.	Each Film Coated Tablet Contains Olmesartan Medoxomil Hydrochlorothiazide ` ` Excipients Colour: Approved Colour Used	I.P. I.P.	20 mg 12.5 mg q.s.		19-01-2019
333	Clopidogrel & Aspirin Tablets	Each Film Coated Tablet Contains Clopidogrel Bisulphate Eq.to Clopidogrel Aspirin (As Enteric Coated Granules) Excipients Colour approved Colour Used	I.P. I.P.	75 mg 150 mg q.s.		19-01-2019
334	Gabapentin & Methylcobalamin Tablets	Each Film Coated Tablet Contains Gabapentin Methylcobalamin Excipients Colour: Approved Colour Used	I.P. I.P.	100mg 500 mcg q.s.		19-01-2019
335	Glimepiride Tablets IP 1 mg	Each uncoated tablet contains: Glimepiride Excipients Colour : Approved Colour Used	I.P.	1 mg q.s.		19-01-2019
336	Glimepiride Tablets IP 2 mg	Each uncoated tablet contains: Glimepiride Excipients Colour : Approved Colour Used	I.P.	2 mg q.s.		19-01-2019
337	Paracetamol Sustained Release Tablets	Each uncoated Sustained release tablet contains: Paracetamol	I.P.	650 mg		19-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients Colour : Approved Colour Used		q.s.		
338	Drotaverine Hydrochloride Tablets I.P. 40mg	Each uncoated tablet contains: Drotaverine Hydrochloride Excipients Colour : Approved Colour Used	I.P.	40 mg q.s.		19-01-2019
339	Drotaverine Hydrochloride Tablets I.P. 80 mg	Each uncoated tablet contains: Drotaverine Hydrochloride Excipients Colour : Approved Colour Used	I.P.	80 mg q.s.		19-01-2019
340	Hydroxyzine Hydrochloride Tablets IP 10 mg	Each Film Coated Tablet contains: Hydroxyzine Hydrochloride Excipients Colour : Approved Colour Used	I.P.	10 mg q.s.		19-01-2019
341	Glimepiride, Metformin Hydrochloride (S.R.) & Voglibose Tablets	Each Uncoated Bilayered Tablet Contains: Glimepiride Metformin Hydrochloride (In Sustained release Form) Voglibose Excipients: Colour: Approved Colour	I.P. I.P. I.P.	1 mg 500 mg 0.2 mg q.s		19-01-2019
342	Glimepiride, Metformin Hydrochloride (S.R.) & Voglibose Tablets	Each Uncoated Bilayered Tablet Contains: Glimepiride Metformin Hydrochloride (In Sustained release Form) Voglibose Excipients: Colour: Approved Colour	I.P. I.P. I.P.	2 mg 500 mg 0.2 mg q.s		19-01-2019
343	Venlafaxine Hydrochloride Tablets	Each uncoated tablet contains: Venlafaxine Hydrochloride Eq. to Venlafaxine Excipients: Colour: Approved Colour	U.S.P.	75 mg q.s.		19-01-2019
344	Calcium citrate with Vitamin D3, Magnesium & Zinc Tablets	Each Uncoated Tablet Contains: Calcium Citrate Vitamin D3 Magnesium Hydroxide Eq. to elemental Magnesium Zinc Sulphate monohydrate Eq. to elemental Zinc Excipients Colour: Approved Colour	U.S.P I.P. I.P. I.P.	1000 mg 200 IU 100 mg 4 mg q.s.		19-01-2019
345	Paroxetine Prolonged Release Tablets I.P. 12.5 mg	Each Extended release tablet contains: Paroxetine Hydrochloride Hemihydrate Eq. to Paroxetine Excipients: Colour: Approved Colour Used	I.P.	12.5 mg q.s		19-01-2019
346	Paroxetine Prolonged Release Tablets I.P. 25 mg	Each Extended release tablet contains: Paroxetine Hydrochloride Hemihydrate Eq. to Paroxetine Excipients: Colour: Approved Colour Used	I.P.	25 mg q.s		19-01-2019
347	Nortriptyline Hcl., Pregabalin & Methylcobalamin Tablets	Each Film Coated Tablet Contains Nortriptyline Hydrochloride Eq.to Nortriptyline Pregabalin Methylcobalamin Excipients Colour: Approved Colour Used	I.P. I.P. I.P.	10mg 75 mg 1500mcg q.s.		19-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
348	Paracetamol Tablets IP 500 mg	Each Uncoated tablet contains Paracetamol Excipients	I.P.	500 mg q.s.		19-01-2019
349	Desvenlafaxine Extended Release Tablets 50 mg	Each Film coated Extended Release Tablet Contains Desvenlafaxine Succinate Eq. to Desvenlafaxine Excipients Colours : Ferric oxide usp –NF yellow ferric oxide USP NF red & titanium dioxide IP		50 mg q.s		19-01-2019
350	Desvenlafaxine Extended Release Tablets 100 mg	Each Film coated Extended Release Tablet Contains Desvenlafaxine Succinate Eq. to Desvenlafaxine Excipients Colours : Ferric oxide usp –NF yellow ferric oxide USP NF red & titanium dioxide IP		100 mg q.s.		19-01-2019
351	Dried Aluminium Hydroxide Gel, Magnesium Hydroxide & Simethicone Suspension	Each 5 ml contains Dried Aluminium Hydroxide Gel Magnesium Hydroxide Simethicone Excipients Colour: Erythrosine Supra	I.P. I.P. I.P.	250 mg 250 mg 50 mg q.s.		19-01-2019
352	Glimepiride Tablets I.P. 3 mg	Each uncoated tablet contains Glimepiride Excipients Colour : Approved Colour Used	I.P.	3 mg q.s.		19-01-2019
353	Glimepiride Tablets I.P. 4 mg	Each uncoated tablet contains Glimepiride Excipients Colour : Approved Colour Used	I.P.	4 mg q.s.		19-01-2019
354	Valsartan Tablets I.P. 80 mg	Each Film coated tablet contains Valsartan Excipients Colour : Approved Colour Used	I.P.	80 mg q.s.		19-01-2019
355	Megaldrate , Oxetacaine & Simethicone Oral Suspension	Each 5ml Contains Megaldrate (Anhydrous) Oxetacaine Simethicone In a palatable non syrupy base Colour: Erythromycin Supra	I.P. B.P. I.P.	400mg 10 mg 20 mg		19-01-2019
356	Teneligliptin Tablets I.P. 20 mg	Each Uncoated tablet Contains Teneligliptin Hydrobromide Hydrate Eq.to Teneligliptin Excipients Colour Approved Colour Used	I.P.	20 mg q.s.		11-09-2020
357	Levofloxacin & Ambroxol Hydrochloride (Sustained Release)Tablets	Each Uncoated bilayered Tablet contains Levofloxacin Hemihydrate Eq. to. Levofloxacin Ambroxol Hydrochloride (in Sustained Release Form) Excipients Approved Colour Used	I.P. I.P.	500mg 75 mg q.s.		11-09-2020
358	Glibenclamide & Metformin Hcl. Tablets IP	Each Uncoated Tablet Contains Glibenclamide Metformin Hydrochloride Excipients	I.P. I.P.	5mg 500mg q.s		11-09-2020



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Approved Colour Used				
359	Glibenclamide & Metformin Hcl. Tablets IP	Each Uncoated Tablet Contains Glibenclamide Metformin Hydrochloride Excipients Approved Colour Used	I.P. I.P.	2.5mg 400mg q.s		11-09-2020
360	Atorvastatin & Ezetimibe Tablets	Each Film Coated tablet Contains Atorvastatin Calcium Eq.to Atorvastatin Ezetimibe Excipients Approved Colour Used	I.P. I.P.	10 mg 10 mg s.s.		11-09-2020
361	Zinc Sulphate Tablets BP 50mg	Each film coated tablet contains: Zinc Sulphate Monohydrate Eq. to Elemental Zinc Excipients Colour: Titanium Dioxide IP	I.P.	137.232mg 50 mg q.s.		11-09-2020
362	Zinc Acetate Tablets	Each film coated tablet contains: Zinc Acetate USP(as dehydrate) Eq. to Elemental Zinc Excipients Colour: Titanium Dioxide IP	U.S.P.	50 mg q.s.		11-09-2020
363	Vitamin C, Zinc Citrate & Selenium Chewable Tablets	Each uncoated chewable tablet contains: Ascorbic Acid Sodium Ascorbate Eq. to Ascorbic Acid Zinc Citrate USP Eq. to Elemental Zinc Selenium Dioxide Monohydrate USP Eq. to Elemental Selenium Excipients Colour: Sunset Yellow FCF	I.P. I.P. U.S.P. U.S.P.	100 mg 450 mg 400 mg 2.2 mg 60 mcg q.s.		11-09-2020
364	Vitamin C, Sodium Ascorbate & Zinc Chewable Tablets	Each Uncoated Chewable Tablet Contains: Ascorbic Acid Sodium Ascorbate Eq. to Ascorbic Acid Zinc Citrate Eq. to Elemental Zinc Excipients Colour: Sunset Yellow FCF	I.P. I.P. U.S.P.	100 mg 450 mg 400mg 5 mg q.s.		11-09-2020
365	Vitamin C Chewable Tablets	Each uncoated Chewable Tablet Contains: Ascorbic Acid Sodium Ascorbate Eq. to Ascorbic Acid Excipients Colour: Sunset Yellow FCF	I.P. I.P.	100 mg 450 mg 400 mg q.s.		11-09-2020
366	Dexchlorpheniramine Maleate Tablets I.P. 2 mg	Each Uncoated tablets Contains Dexchlorpheniramine Maleate Excipients Colour Approved Colour Used	I.P.	2mg q.s.		18-03-2021
367	Hydroxyzine Hydrochloride Tablets I.P.	Each film Coated Tablets Contains: Hydroxyzine Hydrochloride Excipients Colour: Approved colour Used	I.P.	25 mg q.s.		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
368	Moxifloxacin Tablets 400mg	Each Film Coated tablets Contains Moxifloxacin Excipients Clour Approved Colour Used		400mg q.s.		18-03-2021
369	Ketorolac Tromethamine Tablets I.P.	Each Film Coated Tablets Contains: Ketorolac Tromethamine Excipients Colour: Approved Colour used	I.P.	10 mg q.s.		18-03-2021
370	Azithromycin Suspension I.P.	Each 5 ml Contains : Azithromycin Dihydrate Eq.to. Anhydrous Azithromycin Flavoured syrup base Colour: Approved colour used	I.P	200 mg q.s.		18-03-2021
371	Terbinafine Tablets I.P. 500 mg	Each Uncoated Tablet Contains: Terbinafine Hydrochloride Eq. to Terbinafine Excipients Colour: Approved colour Used	I.P.	500 mg q.s.		18-03-2021
372	Piroxicam Tablets I.P. 20 mg	Each Uncoated Tablet Contains : Piroxicam Excipients Colour: Approved colour used	I.P.	20 mg q.s.		18-03-2021
373	Esomeprazole Gastro-Resistant Tablets I.P. 20 mg	Each Gastro-Resistant Tablets Contains : Esomeprazole Magnesium Eq. to Esomeprazole Excipients Colour: Approved colour used	I.P.	20 mg q.s.		18-03-2021
374	Ofloxacin Tablets I.P. 400 mg	Each Film Coated Tablet Contains: Ofloxacin Excipients Colour: Approved colour Used	I.P.	400 mg q.s.		18-03-2021
375	Doxofylline (S.R.) & Montelukast Tablets	Each Uncoated Bilayer Tablets Contains Doxofylline (as Sustained Release) Montelukast Sodium Eq.to Montelukast Excipients Colour Approved Colour Used	I.P. I.P.	400mg 10mg q.s.		18-03-2021
376	Pregabalin Capsules I.P. 300 mg	Each Hard Gelatin Capsule Contains: Pregabalin Excipients Colour: Approved colour used in empty Capsule shell	I.P.	300 mg q.s.		18-03-2021
377	Losartan Potassium & Amlodipine Tablets	Each film coated Tablet Contains:- Losartan Potassium Amlodipine Besylate Eq. to Amlodipine Excipients Approved colour used.	I.P. I.P.	25 mg 5 mg q.s.		18-03-2021
378	Losartan Potassium & Hydrochlorothiazide Tablets	Each Film Coated Tablets Contains Losartan Potassium Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P. I.P.	25 mg 12.5mg q.s.		18-03-2021
379	Citicoline Sodium & Piracetam Tablets	Each Film coated Tablet Contains : Citicoline Sodium Eq. to Citicoline Piracetam Excipients	I.P. I.P.	500 mg 400 mg q.s.		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
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		Colour: Approved colour used				
380	Telmisartan & Chlorthalidone Tablets	Each Uncoated tablet Contains Telmisartan Chlorthalidone Excipients Colour Approved Colour Used	I.P. I.P.	40 mg 12.5 mg q.s.		18-03-2021
381	Olmesartan & Hydrochlorothiazide Tablets IP	Each Film Coated tablets Contains Olmesartan Medoxomil Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P. I.P.	40mg 12.5mg q.s.		18-03-2021
382	Trypsin , Bromelain Rutoside Tablets	Each Enteric Coated Tablet Contains Trypsin Bromelain Rutoside Trihydrate Eq.to Rutoside Excipients Colour Approved Colour Used	B.P. B.P.	96 mg 180 mg 200mg q.s.		18-03-2021
383	Dapagliflozin Tablets 5 mg	Each film coated Tablet Contains:- Dapagliflozin Propanediol Monohydrate Equivalent to Dapagliflozin Excipients Approved colour used.		5mg q.s.		18-03-2021

384	Dapagliflozin Tablets 10 mg	Each film coated Tablet Contains:- Dapagliflozin Propanediol Monohydrate Equivalent to Dapagliflozin Excipients Approved colour used.		10 mg q.s.		18-03-2021
385	Tolperisone Hydrochloride Tablets 150mg	Each Film Coated Tablet Contains Tolperisone Hydrochloride Excipients Colour Approved Colour Used		150 mg q.s.		18-03-2021
386	Tolperisone Hydrochloride & Paracetamol Tablets	Each Film Coated Tablet Contains Tolperisone Hydrochloride Paracetamol Excipients Colour Approved Colour Used	I.P.	150 mg 325 mg q.s.		18-03-2021
387	Naproxen Tablets I.P. 250 mg	Each Uncoated Tablets Contains: Naproxen Excipients Colour: Approved Colour Used	I.P.	250 mg q.s.		18-03-2021
388	Doxofylline Tablets I.P. 400mg	Each Uncoated tablets Contains Doxofylline Excipients Colour Approved Colour Used	I.P.	400mg q.s.		18-03-2021
389	Doxofylline & Ambroxol Hcl. Tablets	Each Uncoated tablets Contains Doxofylline Ambroxol Hydrochloride Excipients Colour Approved Colour Used	I.P. I.P.	400mg 30mg q.s.		18-03-2021
390	Dexrabeprazole & Domperidone S.R. Capsule	Each Hard Gelatin Capsule Contains: Dexrabeprazole (As enteric coated Pellets) Domperidone (As sustained Release Pellets) Excipients Colour: Approved colour used in	I.P.	10 mg 30 mg q.s.		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		empty Capsule shell				
391	Atorvastatin & Enteric Coated Aspirin Capsules I.P.	Each Hard Gelatin Capsule Contains: Atorvastatin Calcium Eq. to Atorvastatin (as Pellets) Aspirin (as Enteric Coated) Pellets) Excipients Colour: Approved colour used in Capsule	I.P. I.P.	10 mg 150 mg q.s.		18-03-2021
392	Methylcobalamin & Calcium & Calcitriol, Folic Acid, Pyridoxine & Zinc Capsules	Each Hard Gelatin Capsule Contains: Methylcobalamin Calcium Calcitriol Folic Acid Pyridoxine Hcl. Zinc Excipients Colour: Approved colour used in Capsule Shells	J.P. I.P. I.P. I.P. I.P. I.P.	500 mcg 250 mg 0.25 mcg 1.5 mg 3 mg 7.5 mg q.s.		18-03-2021
393	Pyridoxine Hcl., Nicotinamide, Cyanocobalamin & Lysine Syrup	Each 5ml Contains Pyridoxine Hydrochloride Nicotinamide Cyanocobalamin Lysine Hydrochloride Flavoured Syrup Base Colour approved Colour Used	I.P. I.P. I.P. U.S.P.	0.75mg 15 mg 2 mcg 37.5 mg q.s.		18-03-2021
394	Ketoconazole Tablets I.P.	Each uncoated Tablet Contains : Ketoconazole Excipients Colour: Approved colour used	I.P.	200 mg q.s.		18-03-2021
395	Megaldrate & Simethicone Oral Suspension I.P.	Each 5 ml Contains: Megaldrate Simethicone In a flavored Syrup Base Colour: Approved colour Used	I.P. I.P.	400 mg 60 mg q.s.		18-03-2021
396	Dried Aluminium Hydroxide, Magnesium Hydroxide, Simethicone And Oxetacaine Syrup.	Each 5 ml contains: Dried Aluminum Hydroxide Gel e.q to Aluminum Hydroxide Magnesium Hydroxide Simethicone Oxetacaine Excipients In a flavoured Syrup Base Colour: Approved colour Used	I.P. I.P. I.P. B.P.	600 mg 300 mg 25 mg 10 mg q.s.		18-03-2021
397	Methylcobalamin, & Alpha Lipoic Acid & Folic Acid, Biotin, Pyridoxine Capsules	Each Hard Gelatin Capsule Contains: Methylcobalamin Alpha Lipoic Acid Folic Acid Biotin Pyridoxine Hydrochloride Excipients Colour: Approved colour used in empty Capsule Shells	I.P. U.S.P. I.P. B.P. I.P.	1500 mcg 200 mcg 1.5 mg 30 mcg 3 mg q.s.		18-03-2021
398	Methylcobalamin, & Pyridoxine, Folic Acid Capsules	Each Hard Gelatin Capsule Contains: Methylcobalamin Pyridoxine Hydrochloride Folic Acid Excipients Colour: Approved colour used in Capsule Shells	J.P. I.P. I.P.	1500 mcg 1.5 mg 5 mg q.s.		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
399	Levocetirizine Hcl. & Ambroxol Hcl. Syrup	Each 5 ml Contains: Levocetirizine Dihydrochloride Ambroxol Hydrochloride In a flavored Syrup Base Colour: Approved colour Used	I.P. I.P.	2.5 mg 30 mg q.s.		18-03-2021
400	Levocetirizine Di-Hcl. & Ambroxol Hcl. Syrup	Each 5 ml Contains: Levocetirizine Dihydrochloride Ambroxol Hydrochloride In a flavored Syrup Base Colour: Approved colour Used	I.P. I.P.	5 mg 30 mg q.s.		18-03-2021
401	Cetirizine Hcl. & Ambroxol Hcl. Syrup	Each 5 ml Contains: Cetirizine Hydrochloride Ambroxol Hydrochloride In a flavored Syrup Base Colour: Approved colour Used	I.P. I.P.	5 mg 30 mg q.s.		18-03-2021
402	Cetirizine Hcl. & Ambroxol Hcl. Syrup	Each 5 ml Contains: Cetirizine Hydrochloride Ambroxol Hydrochloride In a flavored Syrup Base Colour: Approved colour Used	I.P. I.P.	10 mg 30 mg q.s.		18-03-2021
403	Levofloxacin Tablets I.P. 750 mg	Each Film Coated Sustained release Levofloxacin Hemihydrate Eq. to Levofloxacin Excipients Colour Approved Colour Used	I.P.	750mg q.s.		18-03-2021
404	Enteric Coated Pantoprazole & Levosulpiride (S.R.) Capsule	Each Hard Gelatin Capsule Contains Pantoprazole Sodium Eq. To Pantoprazole (as Enteric Coated Pellets) Levosulpiride (As Sustained Release Pellets) Excipients Approved Colour Used in Empty Capsule Shells & Pellets	I.P.	40 mg 75 mg q.s.		18-03-2021
405	Betahistine Di Hydrochloride Sustained Release Tablets 24 mg	Each Film Coated Sustained Release Tablet Contains: Betahistine Di Hydrochloride Excipients Colour: Approved colour Used	I.P.	24 mg q.s.		18-03-2021
406	Betahistine Tablets I.P.16 mg	Each Uncoated Tablet Contains: Betahistine Di Hydrochloride Excipients Colour: Approved colour Used	I.P.	16 mg q.s.		18-03-2021
407	Betahistine Tablets I.P 8 mg	Each Uncoated Tablet Contains: Betahistine Di Hydrochloride Excipients Colour: Approved colour Used	I.P.	8 mg q.s.		18-03-2021
408	Pregabalin Capsules I.P. 75 mg	Each Hard Gelatin Capsule Contains: Pregabalin Excipients Colour: Approved colour used in empty Capsule shell	I.P.	75 mg q.s.		18-03-2021
409	Clindamycin Capsules I.P. 300 mg	Each Hard Gelatin Capsule Contains: Clindamycin Hydrochloride Eq. to Clindamycin Excipients Colour: Approved colour used in empty Capsule shell	I.P.	300 mg q.s.		18-03-2021
410	Tolperisone Hydrochloride	Each Film Coated Sustained Release Tablet Contains				18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Sustained Release Tablets 450mg	Tolperisone Hydrochloride Excipients Colour Approved Colour Used		450 mg q.s.		
411	Amitriptyline Hcl. Tablets IP 10mg	Each Film Coated Tablet Contains Amitriptyline Hydrochloride Excipients Colour Approved Colour Used	I.P.	10mg q.s.		18-03-2021
412	Ivabradine Tablets 5 mg	Each Film Coated tablet Contains Ivabradine Hydrochloride Eq.to Ivabradine Excipients Colour approved Colour Used		5mg q.s.		18-03-2021
413	Hydroxychloroquine Sulphate Tablets 400mg	Each Film Coated Tablet Contains Hydroxychloroquine Sulphate Excipients Colour Approved Colour Used	I.P.	400 mg q.s.		18-03-2021
414	Hydroxychloroquine sulphate Tablets I.P. 200mg	Each Film Coated Tablet Contains Hydroxychloroquine Sulphate Excipients Colour Approved Colour Used	I.P.	200 mg q.s.		18-03-2021
415	Metoprolol (E.R.) 50 mg & Amlodipine 5 mg Tablets I.P.	Each Uncoated Bilayer Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended release) Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	47.5mg 50mg 5 mg q.s.		18-03-2021
416	Metoprolol (E.R.) 25 mg & Amlodipine 2.5 mg Tablets I.P.	Each Uncoated Bilayer tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended release) Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	23.75mg 25 mg 2.5 mg q.s.		18-03-2021
417	Teneligliptin & Metformin Hcl (S.R.) Tablets I.P.	Each Uncoated Bilayer Tablet Contains Metformin Hydrochloride (As Sustained Release) Teneligliptin Hydrobromide Hydrate Eq.to Teneligliptin Excipients Colour Approved Colour Used	I.P. I.P.	500 mg 20 mg q.s.		18-03-2021
418	Teneligliptin & Metformin Hcl. (S.R.) Tablets I.P.	Each Uncoated Bilayer Tablets Contains Metformin Hydrochloride (As Sustained Release) Teneligliptin Hydrobromide Hydrate Eq.to Teneligliptin Excipients Colour Approved Colour Used	I.P. I.P.	1000 mg 20 mg q.s.		18-03-2021
419	Ambroxol Hcl. (S.R.) & Levocetirizine Hcl. Tablets	Each Uncoated Tablet contains Ambroxol Hydrochloride (As Sustained Release) Levocetirizine Hydrochloride Excipients Colour Approved Colour Used	I.P. I.P.	75mg 5mg q.s.		18-03-2021
420		Each Uncoated Bilayer Tablets				18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Metformin Hcl. (S.R.) & Glimepiride Tablets I.P.	Contains Metformin Hydrochloride (as Sustained release) Glimepiride Excipients Colour Approved Colour Used	I.P. I.P.	500mg 3 mg q.s.		
421	Rifaximin Tablets 200mg	Each Film Coated tablets Rifaximin Ph. Eur. Excipients Colour Approved Colour Used		200mg q.s.		18-03-2021
422	Rifaximin Tablets 400mg	Each Film Coated tablets Contains Rifaximin Ph. Eur. Excipients Colour Approved Colour Used		400mg q.s.		18-03-2021
423	Rifaximin Tablets 550mg	Each Film Coated tablets Contains Rifaximin Ph. Eur. Excipients Colour Approved Colour Used		550mg q.s.		18-03-2021
424	Teneligliptin & Metformin Hcl. (S.R.) Tablets I.P.	Each Uncoated Tablet Contains Metformin Hydrochloride (As Sustained Release) Teneligliptin Hydrobromide Hydrate Eq.to Teneligliptin Excipients Colour Approved Colour Used	I.P. I.P.	500mg 20mg q.s.		18-03-2021
425	Aceclofenac & Thiocholchicoside Tablets	Each Film Coated tablets Contains Aceclofenac Thiocholchicoside Excipients Colour approved Colour Used	I.P. I.P.	100 mg 8 mg q.s.		18-03-2021
426	Amlodipine, Telmisartan & Hydrochlorothiazide Tablets.	Each Uncoated tablet contains:- Amlodipine Besylate Eq. to Amlodipine Telmisartan Hydrochlorothiazide Excipients Colour: Approved Colour used	I.P. I.P. I.P.	2.5mg 20 mg 12.5 mg q.s.		18-03-2021
427	Amlodipine, Telmisartan & Hydrochlorothiazide Tablets.	Each Uncoated tablet contains:- Amlodipine Besylate Eq. to Amlodipine Telmisartan Hydrochlorothiazide Excipients Colour: Approved Colour used	I.P. I.P. I.P.	10 mg 80 mg 25 mg q.s.		18-03-2021
428	Etoricoxib & Thiocholchicoside Tablets	Each Film Coated tablets Contains Etoricoxib Thiocholchicoside Excipients Colour Approved Colour Used	I.P. I.P.	60 mg 8 mg q.s.		18-03-2021
429	Lornoxicam & Thiocholchicoside Tablets	Each Film Coated tablets Contains Lornoxicam Thiocholchicoside Excipients Colour Approved Colour Used	I.P.	4 mg 4 mg q.s.		18-03-2021
430	Lornoxicam & Thiocholchicoside Tablets	Each Film Coated tablets Contains Lornoxicam Thiocholchicoside Excipients Colour Approved Colour Used	I.P.	8 mg 8 mg q.s.		18-03-2021
431	Etodolac Sustained Release Tablets I.P. 600 mg	Each Film Coated Sustained release tablets Contains Etodolac	I.P.	600 mg		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients Colour Approved Colour Used		q.s.		
432	Etodolac Sustained Release Tablets I.P. 200 mg	Each Film Coated Sustained release tablets Contains Etodolac Excipients Colour Approved Colour Used	I.P.	200 mg q.s.		18-03-2021
433	Rabeprazole & Domperidone Tablets	Each Gastro-Resistant Tablets Contains : Rabeprazole Sodium Domperidone Excipients Colour: Approved colour used	I.P. I.P.	20 mg 10 mg q.s.		18-03-2021
434	Etodolac Extended Release Tablets 400 mg	Each Film Coated Extended Release Tablets Contains Etodolac Excipients Colour Approved Colour Used	I.P.	400 mg q.s.		18-03-2021
435	Faropenem Tablets 200mg	Each Film Coated tablets Contains Faropenem Sodium Eq.to Faropenem Excipients Colour Approved Colour Used	J.P.	200mg q.s.		18-03-2021
436	Faropenem Extended Release Tablets 300mg	Each Film Coated Extended Release Tablets Contains Faropenem Sodium Eq.to Faropenem Excipients Colour Approved Colour Used	J.P.	300mg q.s.		18-03-2021
437	Fluconazole Tablet I.P. 400 mg	Each Uncoated Tablet Contains: Fluconazole Excipients Colour: Approved Colour Used	I.P.	400 mg q.s.		18-03-2021
438	Methylcobalamin & Alpha Lipoic Acid & Pyridoxine Hcl. & Thiamine Hcl., Calcium Picolinate Tablets	Each Film coated Tablet Contains : Methylcobalamin Alpha Lipoic Acid Pyridoxine Hydrochloride Thiamine Hydrochloride Calcium Picolinate Excipients Colour: Approved colour used	I.P. U.S.P. I.P. I.P.	500 mcg 100 mg 1.5 mg 2 mg 200 mg q.s.		18-03-2021
439	Trypsin- Chymotrypsin, Aceclofenac & Paracetamol Tablets	Each Enteric Coated Tablet Contain 50000 Armour units of enzymatic Activity supplied by a purified concentrate which has specific trypsin and Chymotrypsin activity in a ratio of approximately six to one Aceclofenac Paracetamol Excipients Colour Approved Colour Used	I.P. I.P.	100 mg 325 mg q.s.		18-03-2021
440	Acebrophylline Capsules 100 mg	Each Hard Gelatin Capsules Contains : Acebrophylline Excipients Colour: Approved colour used in empty Capsule shell		100 mg q.s.		18-03-2021
441	Olmesartan Medoxomil , Amlodipine & Hydrochlorothiazide Tablets	Each Film Coated Tablet Contains Olmesartan Medoxomil Amlodipine Besylate Eq.to Amlodipine	I.P. I.P.	20 mg 5 mg		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P.	12.5 mg q.s.		
442	Olmesartan Medoxomil , Amlodipine & Hydrochlorothiazide Tablets	Each Film Coated Tablet Contains Olmesartan Medoxomil Amlodipine Besylate Eq.to Amlodipine Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P. I.P. I.P.	40 mg 5 mg 12.5 mg q.s.		18-03-2021
443	Nebivolol & (S-) Amlodipine Tablets	Each Film Coated Tablet Contains Nebivolol Hydrochloride Eq.to Nebivolol (S-) Amlodipine Besylate Eq.to (S-) Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	5mg 2.5 mg		18-03-2021
444	(S-) Amlodipine 2.5 mg & Atenolol 25 mg Tablets	Each Uncoated Tablet Contains (S-) Amlodipine Besylate Eq.to (S-) Amlodipine Atenolol Excipients Colour Approved Colour Used	I.P. I.P.	2.5 mg 25 mg q.s.		18-03-2021
445	(S-) Amlodipine 2.5 mg & Atenolol 50 mg Tablets	Each Uncoated Tablet Contains (S-) Amlodipine Besylate Eq.to (S-) Amlodipine Atenolol Excipients Colour Approved Colour Used	I.P. I.P.	2.5 mg 50 mg q.s.		18-03-2021
446	(S-) Amlodipine 5 mg & Atenolol 50 mg Tablets	Each Uncoated Tablet Contains (S-) Amlodipine Besylate Eq.to (S-) Amlodipine Atenolol Excipients Colour Approved Colour Used	I.P. I.P.	5 mg 50 mg q.s.		18-03-2021
447	Ketorolac Tromethamine Dispersible Tablets 10mg	Each Uncoated Dispersible Tablet Contains Ketorolac Tromethamine Excipients Colour: Approved Colour used	I.P.	10 mg q.s.		18-03-2021
448	Prazosin Tablet I.P. 0.5 mg	Each Uncoated tablet Contains Prazosin Hydrochloride E.q.to Prazosin Excipients Colour Approved Colour Used	I.P.	0.5 mg q.s.		18-03-2021
449	Prazosin Tablet I.P. 1 mg	Each Uncoated tablet Contains Prazosin Hydrochloride E.q.to Prazosin Excipients Colour Approved Colour Used	I.P.	1 mg q.s.		18-03-2021
450	Prazosin Tablet I.P. 2 mg	Each Uncoated tablet Contains Prazosin Hydrochloride E.q.to Prazosin Excipients Colour Approved Colour Used	I.P.	2 mg q.s.		18-03-2021
451	Prazosin Tablet I.P. 5 mg	Each Uncoated Tablet Contains Prazosin Hydrochloride E.q.to Prazosin Excipients Colour Approved Colour Used	I.P.	5 mg q.s.		18-03-2021
452	Voriconazole Tablets I.P.	Each Film Coated Tablet Contains Voriconazole	I.P.	50 mg		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	50 mg	Excipients Colour Approved Colour Used		q.s.		
453	Voriconazole Tablets I.P. 200 mg	Each Film Coated Tablet Contains Voriconazole Excipients Colour Approved Colour Used	I.P.	200 mg q.s.		18-03-2021
454	Ranolazine Extended Release Tablets 375 mg	Each Film Coated Extended Release Tablet Contains Ranolazine Excipients Colour Approved Colour Used		375 mg q.s.		18-03-2021
455	Ranolazine Extended Release Tablets 750 mg	Each Film Coated Extended Release Tablet Contains Ranolazine Excipients Colour Approved Colour Used		750 mg q.s.		18-03-2021
456	Ranolazine Extended Release Tablets 1000 mg	Each Film Coated Extended Release Tablet Contains Ranolazine Excipients Colour Approved Colour Used		1000 mg q.s.		18-03-2021
457	Ranolazine Extended Release Tablets 500 mg	Each Film Coated Extended Release Tablet Contains Ranolazine Excipients Colour Approved Colour Used		500 mg q.s.		18-03-2021
458	Sevelamer Tablets 400 mg	Each Film Coated Tablet Contains Sevelamer Carbonate Excipients Colour : Approved Colour Used		400 mg q.s.		18-03-2021
459	Sevelamer Tablets 800 mg	Each Film Coated Tablet Contains Sevelamer Carbonate Excipients Colour : Approved Colour Used		800 mg q.s.		18-03-2021
460	Etizolam Tablet 0.25 mg	Each Uncoated Tablet Contains Etizolam Excipients Colour Approved Colour Used		0.25 mg q.s.		18-03-2021
461	Etizolam Tablet 0.5 mg	Each Uncoated Tablet Contains Etizolam Excipients Colour Approved Colour Used		0.5 mg q.s.		18-03-2021
462	Fexofenadine Hcl. Suspension	Each 5 ml Contains : Fexofenadine Hydrochloride Flavored base Colour: Approved colour used	I.P.	30 mg q.s.		18-03-2021
463	Citicoline & Methylcobalamin Tablets	Each Film Coated Tablet Contains Citicoline Sodium Eq.to Citicoline Methylcobalamin Excipients Colour Approved Colour Used	I.P. I.P.	500 mg 750 mcg q.s.		18-03-2021
464	Metformin Hcl (S.R.) & Glimepiride Tablets	Each Uncoated Bilayer Tablets Contains Metformin Hydrochloride (as Sustained release) Glimepiride Excipients Colour Approved Colour Used	I.P. I.P.	500mg 4 mg q.s.		18-03-2021
465	Saxagliptin Tablets 2.5 mg	Each Film Coated Tablet Contains Saxagliptin Hydrochloride Eq.to Saxagliptin		2.5 mg		18-03-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients Colour Approved Colour Used		q.s.		
466	Saxagliptin Tablets 5 mg	Each Film Coated Tablet Contains Saxagliptin Hydrochloride Eq.to Saxagliptin Excipients Colour Approved Colour Used		5 mg q.s.		18-03-2021
467	Saxagliptin & Metformin Hcl. (S.R.) Tablets	Each Film Coated Tablet Contains Saxagliptin Hydrochloride Eq.to Saxagliptin Metformin Hydrochloride (As Sustained Release) Excipients Colour Approved Colour Used	I.P.	5 mg 500mg q.s.		18-03-2021
468	Saxagliptin & Metformin Hcl. (S.R.) Tablets	Each Film Coated Tablet Contains Saxagliptin Hydrochloride Eq.to Saxagliptin Metformin Hydrochloride (As Sustained Release) Excipients Colour Approved Colour Used	I.P.	2.5 mg 1000mg q.s.		18-03-2021
469	Flavoxate Hydrochloride & Ofloxacin Tablets	Each Film Coated Tablet Contains Flavoxate Hydrochloride Ofloxacin Excipients Colour Approved Colour Used	I.P. I.P.	200 mg 200 mg q.s.		18-03-2021
470	Vildagliptin Tablets I.P. 100 mg	Each Uncoated tablet Contains Vildagliptin Excipients Colour Approved Colour Used	I.P.	100 mg q.s.		18-03-2021
471	Aceclofenac & Thiocholchicoside Tablets	Each Film Coated tablets Contains Aceclofenac Thiocholchicoside Excipients Colour Approved Colour Used	I.P. I.P.	100 mg 4 mg q.s.		18-03-2021
472	Zinc Acetate Oral Solution	Each 5 ml Contains : Zinc Acetate (as Dihydrate) Eq. to Elemental Zinc Flavoured Syrup Base Colour: Approved colour used	U.S.P.	20 mg q.s.		05-05-2021
473	Flupirtine Maleate & Paracetamol Tablets	Each Film Coated Tablet Contains Flupirtine Maleate Paracetamol Excipients Colour Approved Colour Used	I.P.	100 mg 325mg q.s.		05-05-2021
474	Acebrophylline, Guaiphenesin & Terbutaline Sulphate Syrup	Each 5ml Contains Acebrophylline Guaiphenesin Terbutaline Sulphate Flavoured Syrup Base Colour Approved Colour Used	I.P. I.P.	50 mg 50 mg 1.25 mg q.s.		05-05-2021
475	Levodropropizine & Chlorpheniramine Maleate Syrup	Each 5ml Contains Levodropropizine Chlorpheniramine Maleate In a Flavoured Base Colour Approved Colour Used	I.P. I.P.	30 mg 2 mg q.s.		05-05-2021
476	Ferrous Bisglycinate, Zinc Bisglycinate , Folic Acid & Methylcobalamin Tablets	Each Film Coated Tablet Contains Ferrous Bisglycinate Eq.to Elemental Iron Zinc Bisglycinate Eq.to Elemental Zinc Folic Acid	I.P.	60 mg 15 mg 1 mg		05-05-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Methylcobalamin Excipients Colour Approved Colour Used	I.P.	500 mcg q.s.		
477	Metformin Hcl. Tablets I.P. 850 mg	Each Uncoated Tablet contains Metformin Hydrochloride Excipients Approved colour used.	I.P.	850 mg q.s.		05-05-2021
478	Metformin Hcl. S.R. Tablets I.P. 850 mg	Each Uncoated Sustained Release Tablet Contains Metformin Hydrochloride Excipients Approved colour used.	I.P.	850 mg q.s.		05-05-2021
479	Metaxalone & Diclofenac Potassium Tablets	Each Uncoated Tablet Contains Metaxalone Diclofenac Potassium Excipients Colour Approved Colour Used	U.S.P. B.P.	400 mg 50 mg q.s.		05-05-2021
480	Ivermectin Tablets USP 12 mg	Each Uncoated Tablet Contains Ivermectin Excipients Colour Approved Colour Used	U.S.P.	12 mg q.s.		05-05-2021
481	Flupirtine Maleate Sustained Release Tablets 400 mg	Each Film Coated Sustained Release Tablet contains Flupirtine Maleate Excipients Colour approved Colour Used		400 mg q.s.		05-05-2021
482	Flupirtine Maleate Capsules 100mg	Each Hard Gelatin capsule Contains Flupirtine Maleate Excipients Approved Colour Used in Empty Capsule Shells		100 mg q.s.		05-05-2021
483	Tapentadol Hcl. Tablets 50 mg	Each Film Coated Tablet Contains Tapentadol Hydrochloride Excipients Colour Approved Colour Used	I.P.	50 mg q.s.		05-05-2021
484	Tapentadol Hcl. Tablets 75 mg	Each Film Coated Tablet Contains Tapentadol Hydrochloride Excipients Colour Approved Colour Used	I.P.	75 mg q.s.		05-05-2021
485	Tapentadol Hcl. Tablets 100 mg	Each Film Coated Tablet Contains Tapentadol Hydrochloride Excipients Colour Approved Colour Used	I.P.	100 mg q.s.		05-05-2021
486	Trypsin Chymotrypsin Tablets	Each Enteric Coated Tablets Contains 200000 Armour Unit of Enzymatic Activity Supplied by a Purified Concentrate which has Specific Trypsin & Chymotrypsin activity in a Ratio of Approximately Six to One Excipients Colour Approved Colour Used		q.s.		05-05-2021
487	Flupentixol Tablets B.P. 0.5mg	Each Uncoated Tablet Contains Flupentixol Dihydrochloride Eq.to Flupentixol Excipients Colour approved Colour Used	B.P.	0.5 mg q.s.		05-05-2021
488	Ticagrelor Tablets 90 mg	Each Film Coated tablet Contains Ticagrelor Excipients Colour Approved Colour Used	I.P.	90 mg q.s.		05-05-2021
489	Fluconazole Tablets I.P. 300mg	Each Uncoated Tablet Contains Fluconazole	I.P.	300 mg		05-05-2021



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Excipients Colour: Approved Colour Used		q.s.		
490	Dicyclomine Hcl. & Simethicone Suspension	Each ml Contains Dicyclomine Hydrochloride Simethicone Flavoured Syrup Base Colour: Approved Colour Used	I.P. U.S.P.	10 mg 40 mg q.s.		05-05-2021
491	Torsemide Tablets I.P. 20 mg	Each Uncoated Tablet contains Torsemide Excipients Colour Approved Colour Used	I.P.	20 mg q.s.		05-05-2021
492	Torsemide Tablets I.P. 100 mg	Each Uncoated Tablet contains Torsemide Excipients Colour Approved Colour Used	I.P.	100 mg q.s.		05-05-2021
493	Ilaprazole Tablets 5 mg	Each Enteric Coated Tablet Contains Ilaprazole Excipients Colour Approved Colour Used	I.P.	5 mg q.s.		05-05-2021
494	Ilaprazole Tablets 10 mg	Each Enteric Coated Tablet Contains Ilaprazole Excipients Colour Approved Colour Used	I.P.	10 mg q.s.		05-05-2021
495	Doxycycline Hyclate & Lactic Acid Bacillus Capsules	Each Hard Gelatin Capsule Contains Doxycycline Hyclate Eq.to. Doxycycline Lactic Acid Bacillus Excipients Approved Colour Used in Empty Capsule Shell	I.P.	100 mg 5 Billion Spore q.s.		05-05-2021
496	Betahistine Sustained Release Tablets 48 mg	Each Film Coated Tablet Contains Betahistine Hydrochloride Excipients Colour Approved Colour Used	I.P.	48 mg q.s.		05-05-2021
497	Tadalafil Tablets I.P. 20 mg	Each Film Coated Tablet Contains Tadalafil Excipients Colour Approved Colour Used	I.P.	20 mg q.s.		05-05-2021
498	Acebrophylline (S.R.) , Fexofenadine Hcl. & Montelukast Tablets	Each Film Coated Tablet Contains Acebrophylline (As Sustained Release form) Fexofenadine Hydrochloride Montelukast Sodium Eq.to Montelukast Excipients Colour Approved Colour Used	I.P. I.P.	100 mg 120 mg 10 mg q.s.		05-05-2021
499	Bisacodyl Gastro Resistant Tablets I.P. 5mg	Each Enteric Coated tablet Contains Bisacodyl Excipients Colour Approved Colour Used	I.P.	5 mg q.s.		05-05-2021
500	Paracetamol & Caffeine Tablets I.P.	Each Uncoated Tablet Contains Paracetamol Caffeine (Anhydrous) Excipients Colour Approved Colour Used	I.P. I.P.	650 mg 50 mg q.s.		03-02-2022
501	Paracetamol, Dextromethorphan Hbr., Phenylephrine Hcl., Levocetirizine Hcl. & Caffeine Tablets	Each Uncoated Tablet Contains Paracetamol Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Levocetirizine Hydrochloride. Caffeine (Anhydrous) Excipients	I.P. I.P. I.P. I.P.	325 mg 5 mg 5 mg 5 mg 30 mg q.s.		03-02-2022



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CB Healthcare

Works : Vill. Messa Tibba, Tehsil. Nalagarh, Distt : Solan (H.P.) 174 101

S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Colour Approved colour Used				
502	Aceclofenac, Paracetamol, Cetirizine Hcl., Phenylephrine Hcl. & Caffeine Tablets	Each Uncoated Tablet Contains Aceclofenac Paracetamol Cetirizine Hydrochloride Phenylephrine Hydrochloride Caffeine (Anhydrous) Excipients Colour Approved colour Used	I.P. I.P. I.P. I.P. I.P.	100 mg 325 mg 10 mg 5 mg 25 mg q.s.		03-02-2022
503	Levocetirizine Hcl. & Montelukast Suspension	Each 5ml Contains Levocetirizine Hydrochloride Montelukast Sodium Eq.to Montelukast Flavoured Base Approved Colour Use	I.P. I.P.	2.5 mg 4 mg q.s.		03-02-2022
504	Aceclofenac & Paracetamol Tablets	Each Uncoated Tablet Contains Aceclofenac Paracetamol Excipients Approved Colour Used	I.P. I.P.	100 mg 325 mg q.s.		03-02-2022
505	Dextromethorphan Hbr., Cetirizine Hcl., Phenylephrine Hcl., Ambroxol Hcl. & Guaiphenesin Tablets	Each Uncoated Tablet Contains Dextromethorphan Hydrobromide Cetirizine Hydrochloride Phenylephrine Hydrochloride Ambroxol Hydrochloride Guaiphenesin Excipients Approved Colour Used	I.P. I.P. I.P. I.P. I.P.	15 mg 5 mg 10 mg 30 mg 100 mg q.s.		03-02-2022
506	Ambroxol Hcl. Levosalbutamol & Guaiphenesin Syrup	Each 5ml Contains Ambroxol Hydrochloride Levosaltbutamol Sulphate Eq.to Levosalbutamol Guaiphenesin Flavoured Syrup Base Approved Colour Used	I.P. I.P. I.P.	15 mg 0.5 mg 50 mg q.s.		03-02-2022
507	Dextromethorphan Hbr., Phenylephrine Hcl. & Chlorpheniramine Maleate Tablets	Each Uncoated tablet Contains Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Chlorpheniramine Maleate Excipients Colour Approved colour Used	I.P. I.P. I.P.	10 mg 5 mg 2 mg q.s		03-02-2022
508	Paracetamol, Phenylephrine Hcl., Ambroxol Hcl., Guaiphenesin & Cetirizine Hcl. Tablets	Each Uncoated Tablet Contains Paracetamol Phenylephrine Hydrochloride Ambroxol Hydrochloride Guaiphenesin Cetirizine Hydrochloride Excipients Colour Approved Colour Used	I.P. I.P. I.P. I.P. I.P.	325 mg 5 mg 15 mg 50 mg 5 mg q.s.		03-02-2022
509	Dapagliflozin & Metformin Hcl.(Sustained Release) Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Sustained Release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P.	500 mg 5 mg q.s		03-02-2022
510	Dapagliflozin & Metformin Hcl. Sustained Release Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Sustained Release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin	I.P.	500 mg 10 mg q.s		03-02-2022



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
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		Excipients Colour Approved colour Used				
511	Levocetirizine Hcl. , Montelukast & Ambroxol Hcl. (S.R.) Tablets	Each Uncoated Bilayer Tablet Contains Levocetirizine Hydrochloride Montelukast Sodium Eq.to Montelukast Ambroxol Hydrochloride (as Sustained Release) Excipients Colour Approved Colour Used	I.P. I.P. I.P.	5 mg 10mg 75mg q.s.		01-04-2022
512	Ergotamine Tartrate ,Caffeine Paracetamol & Prochlorperazine Maleate Tablets	Each Uncoated Tablet Contains Ergotamine Tartrate Caffeine (Monohydrate) Paracetamol Prochlorperazine Maleate Excipients Colour Approved Colour Used	I.P. I.P. I.P. I.P.	1 mg 100 mg 250 mg 2.5 mg q.s.		01-04-2022
513	Phenylephrine Hcl. & Chlorpheniramine Maleate Sustained Release Capsules	Each Hard Gelatine Sustained Release Capsule Contains Phenylephrine Hydrochloride Chlorpheniramine Maleate Excipients Colour: Approved Colour Used in Empty Hard gelatine Capsule shells & pellets	I.P. I.P.	20 mg 8 mg q.s.		01-04-2022

514	Cabergoline Tablets I.P. 0.5mg	Each Uncoated Tablet Contains Cabergoline Excipients Colour Approved Colour Used	I.P.	0.5 mg q.s.		01-04-2022
515	Topiramate Tablets I.P. 25 mg	Each Film Coated Tablet Contains Topiramate Excipients Colour Approved Colour Used	I.P.	25mg q.s.		01-04-2022
516	Topiramate Tablets I.P.50 mg	Each Film Coated Tablet Contains Topiramate Excipients Colour Approved Colour Used	I.P.	50mg q.s.		01-04-2022
517	Topiramate Tablets I.P. 100mg	Each Film Coated Tablet Contains Topiramate Excipients Colour Approved Colour Used	I.P.	100mg q.s.		01-04-2022
518	Telmisartan 40 mg & Hydrochlorothiazide 12.5mg Tablets I.P.	Each Uncoated Bilayer Tablet Contains Telmisartan Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P. I.P.	40 mg 12.5m g q.s.		12-05-2022
519	Telmisartan 80 mg & Hydrochlorothiazide 12.5mg Tablets I.P.	Each Uncoated Bilayer Tablet Contains Telmisartan Hydrochlorothiazide Excipients Colour Approved Colour Used	I.P. I.P.	80 mg 12.5m g q.s.		12-05-2022
520	Albendazole Tablet I.P.	Each Uncoated Chewable Tablet Contains Albendazole Excipients Colour Approved Colour Used	I.P.	400mg q.s.		12-05-2022



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
521	Telmisartan 40 mg & Amlodipine 5mg Tablets I.P.	Each Uncoated Bilayer Tablet Contains Telmisartan Amlodipine Besylate Eq.to Amlodipine Excipients Colour Approved Colour Used	I.P. I.P.	40 mg 5mg q.s.		12-05-2022
522	Alpha Galactosidase, Lipase, Alpha Amylase , Protease & Lactase Capsules	Each Hard Gelatin capsule Contains Alpha Galactosidase Lipase Enzyme Alpha Amylase Enzyme (1:800) Protease Enzyme Lactase Enzyme Excipients Colour: Approved Colour Used in empty capsule shell	I.P. U.S.P.	150 mg 3 mg 50 mg 25 mg 7 mg q.s.		12-05-2022
523	Rosuvastatin Calcium , Clopidogrel & Aspirin Capsules	Each Hard Gelatin Capsule Contains Rosuvastatin Calcium Eq.to Rosuvastatin (as pellets) Clopidogrel Bisulphate Eq.to Clopidogrel (as pellets) Aspirin (As Enteric Coated pellets) Excipients Colour: Approved Colour Used in empty Capsule shells & pellets	I.P. I.P. I.P.	10 mg 75 mg 75mg q.s.		12-05-2022
524	Potassium Citrate, Magnesium Citrate & Vitamin B6 Syrup	Each 5ml Contains Potassium Citrate monohydrate Magnesium Citrate Vitamin B6 Flavoured Syrup Base Colour Approved Colour Used	I.P. U.S.P. I.P.	1100mg 375 mg 20 mg q.s.		22-06-2022
525	Methylcobalamin, Alpha Lipoic Acid ,Benfotiamine, Inositol, Folic Acid, Chromium Picolinate & Selenium Dioxide Capsules	Each hard Gelatin Capsule Contains Methylcobalamin Alpha Lipoic Acid Benfotiamine Inositol Folic Acid Chromium Picolinate Selenium Dioxide Excipients Approved Colour Used in Empty Hard Gelatin Capsule Shells	I.P. U.S.P. B.P I.P. U.S.P. U.S.P.	1500 mcg 100 mg 50 mg 100 mg 0.5 mg 65 mcg 18 mcg q.s.		11-07-2022
526	Nimesulide, Paracetamol & Serratiopeptidase Tablets	Each Film Coated Tablet Contains Nimesulide Paracetamol Serratiopeptidase Excipients Approved Colour Used	B.P. I.P. I.P. q.s.	50 mg 325 mg 10 mg		22-12-2017
527	Ursodeoxycholic Acid Sustained Release Tablets 450 mg	Each Film Coated Sustained Release Tablet Contains Ursodeoxycholic Acid Excipients Colour approved Colour Used	I.P.	450 mg q.s.		05-05-2021
528	Paracetamol Bilayer Tablets 1000 mg	Each Uncoated Bilayer Tablets Contains Paracetamol (as immediate release)	I.P I.P	325 mg 675 mg q.s		As per last approval 19.01.2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
		Paracetamol (as immediate release) Excipients Colour: Approved Colour Used				
529	Dextrabe[prazole Sodium Enteric Coated Tablets 5 mg	Each Enteric coated Tablet Contains Dextrabe[prazole Sodium Excipients Approved Colour Used		5 mg q.s		DCGI Approval No./Date 84/13-06-2007
530	Dextrabe[prazole Sodium Enteric Coated Tablets 10 mg	Each Enteric coated Tablet Contains Dextrabe[prazole Sodium Excipients Approved Colour Used		5 mg q.s		DCGI Approval No./Date 84/13-06-2007
531	Desloratadine , Ambroxol Hcl. Guaiphenesin & Menthol Syrup	Each 5ml Contains Desloratadine Ambroxol Hydrochloride Guaiphenesin Menthol Flavoured Syrup Base Colour Approved Colour Used	U.S.P I.P I.P I.P	2.5 mg 30 mg 50 mg 1 mg q.s		DCGI Approval No./Date 282/02-03-2005
532	Nitrazepam Tablet IP	Each Uncoated tablet Contains: Nitrazepam Excipient Colour : Approved Colour Used	I.P.	10 mg q.s		I.P. 2022 Page No. 3071
533	Trihexyphenidyl Hydrochloride Tablets I.P.	Each Uncoated Tablet Contains Trihexyphenidyl Hydrochloride Excipients Colour: Approved Colour Used	I.P.	2 mg q.s.		I.P. 2022 Page No. 1598
534	Trihexyphenidyl Hydrochloride Tablets I.P.	Each Uncoated Tablet Contains Trihexyphenidyl Hydrochloride Excipients Colour: Approved Colour Used	I.P.	5 mg q.s.		I.P. 2022 Page No. 1598
535	Acebrophylline (S.R.) , Fexofenadine Hcl. & Montelukast Tablets	Each Film Coated Tablet Contains Acebrophylline (As Sustained Release form) Fexofenadine Hydrochloride Montelukast Sodium Eq.to Montelukast Excipients Colour Approved Colour Used	I.P. I.P.	200 mg 120 mg 10 mg q.s.		Lediz 3 D Tablets Mfd. By Medoz Pharmaceuticals Pvt Ltd. Manpura Baddi Solan H.P.
536	Levetiracetam Tablets IP 750mg	Each Film Coated Tablet Contains Levetiracetam Excipients Colour Approved Colour Used	I.P.	750mg q.s.		I.P. 2022 Page No. 2733
537	Ferrous Bisglycinate , Folic Acid ,Vitamin B12 & Zinc Syrup	Each 5ml Contains Iron Amino Acid Chelate Ferrous Bisglycinate Eq.to Elemental iron Folic Acid Vitamin B12 Zinc Picolinate Eq.to Elemental Zinc Flavoured Syrup Base Colour: Approved Colour Used	I.P. I.P.	75 mg 15 mg 0.25 mg 3.75 mcg 26 mg 5.5 mg q.s.		DCGI Approval No./Date 1144 / 22-12-2011
538	Aceclofenac Sustained Release 200mg & Thiocolchicoside Sustained Release 8 mg	Each Hard Gelatin capsule contains Aceclofenac (As Sustained release Pellets) Thiocolchicoside	I.P. I.P.	200 mg 8 mg		DCGI Approval No. / Date 1147 / 04-01-2012



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
	Capsules	(As Sustained release Pellets) Excipients Approved Colour Used in empty capsule shells		q.s.		
539	Tadalafil & Dapoxetine Tablets	Each Film Coated tablet Contains Tadalafil Dapoxetine Hydrochloride Eq.to Dapoxetine Excipients Colour Approved Colour Used	I.P. I.P.	20 mg 30 mg q.s.		DCGI Approval No./Date 1321 / 30-07-2019
540	Sodium Alginate , Sodium bicarbonate & Calcium Carbonate Suspension	Each 5 ml contains Sodium Alginate Sodium bicarbonate Calcium Carbonate Flavoured Base Colour: Approved Colour Used	I.P. I.P. I.P.	250 mg 133.5 mg 80 mg q.s.		DCGI Approval No./Date 08 / 09-04-2020
541	Empagliflozin & Linagliptin Tablets	Each Film Coated tablet Contains Empagliflozin Linagliptin Excipients Colour Approved Colour Used		25 mg 5 mg q.s.		DCGI Approval No. / Date 1273/ 16-10-2017
542	Empagliflozin & Linagliptin Tablets	Each Film Coated tablet Contains Empagliflozin Linagliptin Excipients Colour Approved Colour Used		10 mg 5 mg q.s.		DCGI Approval No. / Date 1273/ 16-10-2017
543	Ferrous Ascorbate Folic Acid & Methylcobalamin Suspension	Each 5ml contains Ferrous Ascorbate Eq.to Elemental Iron Folic Acid Methylcobalamin Flavoured Base Colour: Approved Colour Used	I.P. I.P. I.P.	10 mg 0.5 mg 500 mcg q.s.		AORFER-XT Suspension Mfd. By. Athens Life Sciences Kala Amb Sirmor H.P.
544	Naproxen Tablets I.P. 500 mg	Each Uncoated Tablets Contains: Naproxen Excipients Colour: Approved Colour Used	I.P.	500 mg q.s.		IP 2022 Page No. 3033
545	Acetaminophen & Dicyclomine Hcl. Capsules	Each Hard Gelatin Capsule Contains Acetaminophen Dicyclomine Hydrochloride Excipients Colour Approved Colour Used in Empty Capsule shell	I.P. I.P.	325 mg 10 mg q.s.		Petmol- Spas Mfd. By. Euphoria India pharmaceuticals Baaddi Solan H.P
546	Aceclofenac & Rabeprazole Capsule	Each Hard Gelatin Capsule Contains Aceclofenac (As Sustained release Pellets Rabeprazole sodium (As Sustained release Pellets) Excipients Colour Approved Colour Used in Empty Capsule shell	I.P. I.P.	200 mg 10 mg		DCGI Approval No. / Date 963/ 01-07-2009
1.	Dapagliflozin & Metformin Hcl. Extended Release Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Extended Release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P.	1000 mg 5 mg q.s	Self	DCGI Approval No./Date 1272 / 16-10-2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
2.	Dapagliflozin & Metformin Hcl. Extended Release Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Extended Release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P.	1000 mg 10 mg q.s	Self	DCGI Approval No./Date 1272 / 16-10-2017
3.	Metformin Sustained Release Tablets I.P. 500 mg	Each Uncoated Sustained Release Tablet Contains Metformin Hydrochloride Excipients Colour Approved Colour Used	I.P.	500 mg q.s.	Self	I.P. 2018 Page No. 2546
4.	Metformin Sustained Release Tablets I.P. 1000 mg	Each Uncoated Sustained Release Tablet Contains Metformin Hydrochloride Excipients Colour Approved Colour Used	I.P.	1000 mg q.s.	Self	I.P. 2018 Page No. 2546
1.	Dapagliflozin & Metformin Hcl.(Immediate Release) Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Immediate Release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P.	1000 mg 5 mg q.s	Self	DCGI Approval No./Date 1297 / 03-10-2018
2.	Itraconazole Capsules BP 100 mg	Each Hard Gelatin Capsule Contains Itraconazole (As Pellets) Excipients Approved Colour Used in empty Hard Gelatine Capsule shells	B.P.	100 mg q.s.	Self	BP 2022 Vol. III Page No. 856
3.	Itraconazole Capsules BP 200 mg	Each Hard Gelatin Capsule Contains Itraconazole (As Pellets) Excipients	B.P.	200 mg q.s.	Self	BP 2022 Vol. III Page No. 856
1.1	Naproxen & Domperidone Tablets	Each Film Coated tablet Contains Naproxen Domperidone Maleate Eq.to Domperidone Excipients Colour Approved Colour Used	I.P. I.P.	500 mg 10 mg q.s.	Self	Naproford-D 500 Tab Mfd. By. Logos Pharma Nalagarh H.P.
1.2	Atorvastatin Calcium 10mg & Fenofibrate 145 mg Tablets	Each Film Coated Tablet Contains Atorvastatin Calcium Eq.to Atorvastatin Fenofibrate Excipients Colour Approved Colour Used	I.P. I.P.	10 mg 145 mg q.s.	Self	DCGI Approval No./Date 157 / 30-10-2007
1.3	Tofisopam Tablet 50mg	Each Uncoated Tablet Contains Tofisopam Excipients Colour Approved Colour Used	JP	50 mg q.s	Self	DCGI Approval No./Date 08 / 13-04-2010
1.4	Tofisopam Tablet 100 mg	Each Uncoated Tablet Contains Tofisopam Excipients Colour Approved Colour Used	JP	100 mg q.s	Self	DCGI Approval No./Date 08 / 13-04-2010



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
1.5	Disodium Hydrogen citrate Syrup	Each 5ml Contains Disodium Hydrogen Citrate Flavoured Base Colour Approved Colour Used	B.P.	1.25 gm q.s.	Self	Alkamute Syrup Mfd. By Athens Life Sciences Kala Amb H.P.
1.6	Tofacitinib Tablets 5 mg	Each Film Coated Tablets Contains Tofacitinib Citrate Eq.to Tofacitinib Excipients Colour Approved Colour Used		8.08 mg 5 mg q.s.		DCGI Approval No./Date 04 / 01-04-2016
1.7	Esomeprazole 40 mg Tablets (with sodium bicarbonate as buffer)	Each Film Coated tablet Contains Esomeprazole Magnesium Trihydrate Eq.to esomeprazole Excipients Colour Approved Colour Used	I.P.	40 mg q.s	Self	DCGI Approval No./Date 135 / 14-05-2008
1.8	Naproxen & Sumatriptan Tablets	Each Film Coated tablet Contains Naproxen Sumatriptan Excipients Colour Approved Colour Used	I.P. I.P.	500 mg 85 mg q.s.	Self	DCGI Approval No./Date 129 / 03-08-2009
2.1	Azithromycin Oral Suspension IP	Each 5 ml Contains : Azithromycin Dihydrate Eq.to. Anhydrous Azithromycin Flavoured Syrup base Colour: Approved colour used	IP	100 mg q.s.	Self	I.P. 2018 Page No. 1311
2.2	Sodium Bicarbonate Tablets USP 500 mg	Each Film Coated Tablet Contains Sodium Bicarbonate Excipients Colour Approved Colour Used	IP	500 mg q.s.	Self	USP 43 Page No. 4062
2.3	Simvastatin Tablets IP 10 mg	Each Film Coated Tablet contains Simvastatin Excipients Colour Approved Colour Used	IP	10 mg q.s.	Self	IP 2018 Page No. 3195
2.4	Simvastatin Tablets IP 20 mg	Each Film Coated Tablet contains Simvastatin Excipients Colour Approved Colour Used	IP	20 mg q.s.	Self	IP 2018 Page No. 3195
2.5	Levetiracetam Tablets IP 1000 mg	Each Film Coated Tablet Contains Levetiracetam Excipients Colour Approved Colour Used	IP	1000 mg q.s.	Self	IP 2018 Page No. 2411
2.6	Levetiracetam Oral Solution	Each ml Contains Levetiracetam Flavoured Base Colour Approved Colour Used	IP	100 mg q.s	Self	DCGI No. / Date 05-05-2011
3.1	Dapagliflozin & Metformin Hcl.(Extended release) Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Extended release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P.	1000 mg 5 mg q.s	Self	DCGI Approval No./Date 1272 / 16.10.2017



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
3.2	Dapagliflozin & Metformin Hcl.(Extended release) Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Extended release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P.	1000 mg 10 mg q.s	Self	DCGI Approval No./Date 1272 / 16.10.2017
3.3	Metformin Sustained Release Tablets I.P. 500 mg	Each Uncoated Sustained Release Tablet Contains Metformin Hydrochloride Excipients Colour Approved Colour Used	I.P.	500 mg q.s.	Self	I.P. 218 Page No. 2546
3.4	Metformin Sustained Release Tablets I.P. 1000 mg	Each Uncoated Sustained Release Tablet Contains Metformin Hydrochloride Excipients Colour Approved Colour Used	I.P.	1000 mg q.s.	Self	I.P. 218 Page No. 2546
4.1	Dapagliflozin & Metformin Hcl.(Immediate release) Tablets	Each Film Coated Tablet Contains Metformin Hydrochloride (As Immediate release) Dapagliflozin Propanediol Monohydrate Eq.to Dapagliflozin Excipients Colour Approved colour Used	I.P	1000 mg 5 mg q.s	Self	DCGI Approval No./Date 1297 / 03-10-2018
4.2	Itraconazole Capsules BP 100 mg	Each Hard Gelatin Capsule contains: Itraconazole (As Pellets) Excipients Approved colour used in Empty Hard Gelatine Capsule Shells	B.P	100 mg q.s.	Self	BP 2022 Vol. III Page No. 856
4.3	Itraconazole Capsules BP 200 mg	Each Hard Gelatin Capsule contains: Itraconazole (in the form of Pellets) Excipients Approved colour used in Empty Hard Gelatin Capsule Shells	B.P.	200 mg q.s.	Self	BP 2022 Vol. III Page No. 856
4.4	Acetylcysteine & Ambroxol Hydrochloride Capsules	Each Hard Gelatin Capsule Contains Acetylcysteine Ambroxol Hydrochloride Excipients Approved Colour Used in empty Hard	B.P. I.P	300 mg 60 mg q.s.	Self	RT-O Capsule Mfd. By Swiss Garnier Biotech Pvt Ltd. Una H.P.
5.1	Paracetamol Bilayer Tablets 1000mg	Each Uncoated Bilayer Tablets Contains Paracetamol (as immediate release) Paracetamol (as sustained release)	I.P. I.P.	325 mg 675 mg q.s.	Self	As per Last Approval 19-01-2019



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
5.2	Dexrabeprazole Sodium Enteric Coated Tablets 5 mg	Each Enteric Coated Tablet Contains Dexrabeprazole Sodium Excipients Colour: Approved Colour Used		5 mg q.s.	Self	DCGI Approval No. / Date 84 / 13-06-2007
5.3	Dexrabeprazole Sodium Enteric Coated Tablets 10 mg	Each Enteric Coated Tablet Contains Dexrabeprazole Sodium Excipients Colour: Approved Colour Used		10 mg q.s.	Self	DCGI Approval No. / Date 84 / 13-06-2007
5.4	Desloratadine , Ambroxol Hcl. Guaiphenesin & Menthol Syrup	Each 5ml Contains Desloratadine Ambroxol Hydrochloride Guaiphenesin Menthol Flavored Syrup Base Colour: Approved Colour Used	U.S.P. I.P. I.P. I.P.	2.5 mg 30 mg 50 mg 1 mg q.s.	Self	DCGI Approval No. / Date 282 / 02-03-2005
6.1	Mecobalamin Tablets	Each Uncoated Sublingual tablets Contains Mecobalamin Excipients Colour: Approved Colour Used	I.P.	1500mcg q.s.	Self	Nurokind Tablet Mfd. By: Mediforce Healthcare Pvt Ltd. Ponta Sahib Sirmor H.P.
6.2	Calcium Carbonate , Vitamin D3 , Methylcobalamin, L-Methylfolate Calcium, Pyridoxal -5 Phosphate Tablets	Each Film Coated Tablet Contains Calcium Carbonate Eq.to Elemental Calcium Vitamin D3 Methylcobalamin L-Methylfolate Calcium Pyridoxal 5-Phosphate Excipients Colour : Approved Colour Used	I.P. I.P. I.P.	1250 mg 500mg 2000 IU 1500mcg 1 mg 20 mg q.s.	Self	Eldecal-XT Tablets Mfd. By Dallas Drugs Pvt. Ltd. Baddi H.P.
6.3	Oxcarbazepaine Tablets I.P. 150mg	Each Uncoated Tablet Contains Oxcarbazepaine Excipients Colour : Approved Colour Used	I.P.	150 mg q.s.	Self	I.P. 2018 Page No. 2815
6.4	Oxcarbazepaine Tablets I.P. 300mg	Each Uncoated Tablet Contains Oxcarbazepaine Excipients Colour : Approved Colour Used	I.P.	300 mg q.s.	Self	I.P. 2018 Page No. 2815
6.5	Oxcarbazepaine Tablets I.P. 450mg	Each Uncoated Tablet Contains Oxcarbazepaine Excipients Colour : Approved colour Used	I.P.	450 mg q.s.	Self	I.P. 2018 Page No. 2815
6.6	Oxcarbazepaine Tablets I.P. 600mg	Each Uncoated Tablet Contains Oxcarbazepaine Excipients Colour : Approved Colour Used	I.P.	600 mg q.s.	Self	I.P. 2018 Page No. 2815
6.7	Sacubitril & Valsartan Tablets (200mg)	Each Film Coated Tablet Contains 97 mg Sacubitril and 103 mg Valsartan as Sodium Salt Complex Excipients Colour : Approved Colour Used		97+103 mg q.s.	Self	DCGI Approval No. / Date 11- 14/07/2016



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
6.8	Sacubitril & Valsartan Tablets (50 mg)	Each Film Coated Tablet Contains 24 mg Sacubitril and 26mg Valsartan as Sodium salt complex Excipients Colour : Approved Colour Used		24+ 26 mg q.s.	Self	DCGI Approval No. / Date 11- 14/07/2016
6.9	Sacubitril & Valsartan Tablets (100mg)	Each Film Coated Tablet Contains 49 mg Sacubitril and 51 mg Valsartan as Sodium salt complex Excipients Colour : Approved Colour Used		49+ 51 mg q.s.	Self	DCGI Approval No. / Date 11- 14/07/2016
6.10	Spironolactone Tablets IP 25 mg	Each Uncoated tablet Contains Spironolactone Excipients Colour : Approved Colour Used	I.P.	25 mg q.s.	Self	I.P. 2018 Page No. 3253
6.11	Spironolactone Tablets IP 50 mg	Each Uncoated tablet Contains Spironolactone Excipients Colour : Approved Colour Used	I.P.	50 mg q.s.	Self	I.P. 2018 Page No. 3253
6.12	Spironolactone Tablets IP 100 mg	Each Uncoated tablet Contains Spironolactone Excipients Colour : Approved Colour Used	I.P.	100 mg q.s.	Self	I.P. 2018 Page No. 3253
7.1	Trihexyphenidyl Hydrochloride Tablets I.P.	Each Uncoated Tablet Contains Trihexyphenidyl Hydrochloride Excipients Colour: Approved Colour Used	I.P.	2 mg q.s.	Self	I.P. 2018 Page No. 1344
7.2	Trihexyphenidyl Hydrochloride Tablets I.P.	Each Uncoated Tablet Contains Trihexyphenidyl Hydrochloride Excipients Colour: Approved Colour Used	I.P.	5 mg q.s.	Self	I.P. 2018 Page No. 1344
7.3	Acebrophylline (S.R.) , Fexofenadine Hcl. & Montelukast Tablets	Each Film Coated Tablet Contains Acebrophylline (As Sustained Release form) Fexofenadine Hydrochloride Montelukast Sodium Eq.to Montelukast	I.P. I.P.	200 mg 120 mg 10 mg	Self	Lediz 3 D Tablets Mfd. By Medoz Pharmaceuticals Pvt Ltd. Manpura Baddi Solan H.P.
7.4	Levetiracetam Tablets IP 750mg	Each Film Coated Tablet Contains Levetiracetam Excipients Colour Approved Colour Used	I.P.	750mg q.s.	Self	I.P. 2018 Page No. 2411
7.5	Ferrous Bisglycinate , Folic Acid ,Vitamin B12 & Zinc Syrup	Each 5ml Contains Iron Amino Acid Chelate Ferrous Bisglycinate Eq.to Elemental iron Folic Acid Vitamin B12 Zinc Picolinate Eq.to Elemental Zinc Flavoured Syrup Base Colour: Approved Colour Used	I.P. I.P.	75 mg 15 mg 0.25 mg 3.75 mcg 26 mg 5.5 mg q.s.	Self	DCGI Approval No./Date 1144 / 22-12-2011



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
7.6	Aceclofenac Sustained Release 200mg & Thiocolchicoside Sustained Release 8 mg Capsules	Each Hard Gelatin capsule contains Aceclofenac (As Sustained release Pellets) Thiocolchicoside (As Sustained release Pellets) Excipients Approved Colour Used in empty capsule shells	I.P. I.P.	200 mg 8 mg q.s.	Self	DCGI Approval No. / Date 1147 / 04-01-2012
7.7	Empagliflozin & Linagliptin Tablets	Each Film Coated tablet Contains Empagliflozin Linagliptin Excipients Colour Approved Colour Used		25 mg 5 mg q.s.	Self	DCGI Approval No. / Date 1273/ 16-10-2017
7.8	Empagliflozin & Linagliptin Tablets	Each Film Coated tablet Contains Empagliflozin Linagliptin Excipients Colour Approved Colour Used		10 mg 5 mg q.s.	Self	DCGI Approval No. / Date 1273/ 16-10-2017
7.9	Ferrous Ascorbate Folic Acid & Methylcobalamin Suspension	Each 5ml contains Ferrous Ascorbate Eq.to Elemental Iron Folic Acid Methylcobalamin Flavoured Base Colour: Approved Colour Used	I.P. I.P. I.P.	10 mg 0.5 mg 500 mcg q.s.	Self	AORFER-XT Suspension Mfd. By. Athens Life Sciences Kala Amb Sirmor H.P.
7.10	Naproxen Tablets I.P. 500 mg	Each Uncoated Tablets Contains: Naproxen Excipients Colour: Approved Colour Used	I.P.	500 mg q.s.	Self	IP 2022 Page No. 3033
7.11	Acetaminophen & Dicyclomine Hcl. Capsules	Each Hard Gelatin Capsule Contains Acetaminophen Dicyclomine Hydrochloride Excipients Colour Approved Colour Used in Empty Capsule shell	I.P. I.P.	325 mg 10 mg q.s.	Self	Petmol- Spas Mfd. By. Euphoria India pharmaceuticals Baddi Solan H.P
7.12	Aceclofenac & Rabeprazole Capsule	Each Hard Gelatin Capsule Contains Aceclofenac (As Sustained release Pellets Rabeprazole sodium (As Sustained release Pellets) Excipients Colour Approved Colour Used in Empty Capsule shell	I.P. I.P.	200 mg 10 mg	Self	DCGI Approval No. / Date 963/ 01-07-2009
8.1	Citicoline Sodium Syrup	Each 5 ml Contains : Citicoline sodium IP E.q to Citicoline In a Flavoured Syrupy Base Excipients Colour: Approved colour used.	I.P.	500 mg q.s.	Self	
8.2	Ranitidine Oral Solution IP	Each 10 ml Contains : Ranitidine Hydrochloride IP Eq.to Ranitidine In a Flavoured Syrupy Base Excipients Colour: Approved colour used.	IP	150 mg q.s	Self	



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
8.3	Telmisartan & Cilnidipine Tablets	Each Film Coated Tablet Contains : Telmisartan I.P Cilnidipine I.P Excipients Colour: Approved Colour Used		40 mg 10 mg q.s.	Self	
8.4	Teneligliptin & Metformin hydrochloride Extended Release Tablets	Each Uncoated Bilayered Tablets contains : Teneligliptin Hydrobromide Hydrate eq. to Teneligliptin Metformin Hydrochloride (As Extended Release) Excipients Colour: Approved Colour Used		20 mg 500 mg	Self	
9.1	Sodium Alginate , Sodium bicarbonate & Calcium Carbonate Suspension	Each 5 ml contains Sodium Alginate Sodium bicarbonate Calcium Carbonate Flavoured Base Colour: Approved Colour Used	I.P. I.P. I.P.	250 mg 133.5 mg 80 mg q.s.	Self	DCGI Approval No./Date 08 / 09-04-2020
9.2	Silodosin Capsules 8 mg	Each Hard Gelatine Capsule Contain Silodosin Excipients Approved Colour used in empty capsule shell		5mg q.s.	Self	DCGI Approval No./Date 69 / 23-06-2011
9.3	Dapoxetine Hydrochloride Tablets I.P	Each Film Coated tablet Contains Dapoxetine Hydrochloride IP equivalent to Dapoxetine Excipients Colour: Approved Colour Used	I.P	30 mg	Self	I.P 2018 Page no 1754
9.4	Linagliptin & Metformin hydrochloride Tablet	Each Film Coated Tablet contains Linagliptin Metformin Hydrochloride Excipients Colour: Approved Colour Used	IP	2.5 mg 500 mg q.s	Self	DCGI Approval No. Date 1159 / 26.07.2013
9.5	Linagliptin Tablet	Each Film Coated Tablet contains Linagliptin Excipients Colour: Approved Colour Used		5 mg q.s	Self	DCGI Approval No. Date 06 / 23.01.2012
9.6	Linagliptin & Metformin hydrochloride Tablet	Each Film Coated Tablet contains Linagliptin Metformin Hydrochloride Excipients Colour: Approved Colour Used	IP	2.5 mg 1000 mg q.s	Self	DCGI Approval No. Date 1159 / 26.07.2013
10.1	Cetirizine Hydrochloride , Phenylephrine Hydrochloride & Paracetamol Suspension	Each 5 ml contains Cetirizine Hydrochloride Phenylephrine Hydrochloride Paracetamol In a Flavoured Syrupy Base Colour: Approved Colour Used	I.P I.P I.P	2 mg 5 mg 250 mg q.s	Self	NA



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
10.2	Cetirizine Hydrochloride , Phenylephrine Hydrochloride & Paracetamol Suspension	Each 5 ml contains Cetirizine Hydrochloride Phenylephrine Hydrochloride Paracetamol In a Flavoured Syrupy Base Colour: Approved Colour Used	I.P I.P I.P	2 mg 2.5 mg 125 mg q.s	Self	NA
10.3	Roxithromycin Tablet IP	Each Film Coated tablet Contains Roxithromycin IP Excipients Colour: Approved Colour Used	I.P	150 mg	Self	NA
11.1	Sildenafil Citrate & Dapoxetine Hydrochloride Tablets I.P	Each Film Coated Tablet Contains Sildenafil Citrate Dapoxetine Excipients Colour: Approved Colour Used	IP IP	100 mg 60 mg q.s	Self	DCGI Approval No. Date 02 / 10.02.2020
11.2	Sildenafil Citrate & Dapoxetine Hydrochloride Tablets I.P	Each Film Coated Tablet Contains Sildenafil Citrate Dapoxetine Excipients Colour: Approved Colour Used	IP IP	50 mg 60 mg q.s	Self	DCGI Approval No. Date 02 / 10.02.2020
11.3	Bilastin & Montelukast Tablets	Each Film Coated Bilayer Tablet Contains Bilastin Montelukast Sodium e.q to Montelukast IP Excipients Colour: Approved Colour Used	I.P	20 mg 10 mg q.s	Self	DCGI Approval No. Date 06 / 09.03.2020
12.1	Rosuvastatin Calcium , Clopidogrel & Aspirin Capsules	Each Hard Gelatin Capsule Contains Rosuvastatin Calcium Eq.to Rosuvastatin (as pellets) Clopidogrel Bisulphate Eq.to Clopidogrel (as pellets) Aspirin (As Enteric Coated pellets) Excipients Colour: Approved Colour Used in empty Capsule shells & pellets	I.P. I.P. I.P.	20 mg 75 mg 75mg q.s.	Self	VAIROSE GOLD 20 CAPSULE B.NO-PC-40100 MFG DT. Feb.2024 EXP.DT.Jan.2026 Mfd By. Pharmaroots Healthcare Khasra No.411, Village Tipra, P.O Barotiwala, Tehsil-Baddi Distt. Solan (H.P) 174101
12.2	Levocloperastine Fendizoate Oral suspension	Each 5ml Contains Levocloperastine Fendizoate Eq. to Levocloperastine Excipients Colour Approved colour Used		35.4 mg 20 mg q.s.	Self	DCGI Approval No/Date 177 / 29.09.2010
12.3	Rosuvastatin & Aspirin Capsule	Each Hard Gelatin Capsule Contains Rosuvastatin Calcium IP Eq.to Rosuvastatin (As Granules) Aspirin IP (As Gastro-resistance tablet) Excipients Approved Colour Used	I.P. I.P.	10 mg 75 mg q.s.	Self	DCGI Approval No/Date 1101 / 30.03.2011



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
12.4	Lactulose Solution USP	Each 15 ml Contain: Lactulose concentrate USP eq.to Lactulose 66 % Excipients Colour: Approved Colour Used	USP	10 mg q.s	Self	EASYLAX L Oral solution B.No-EAL23022 MFG DT. Oct.2023 EXP.DT.jan.2023 Mfd. By: Tirupati Medicare LTD Nahan Road Ponta Sahib, Distt Sirmour 173025 (H.P)
13.1	Atorvastatin Calcium , Clopidogrel & Aspirin Capsules	Each Hard Gelatin Capsule Contains Atorvastatin Calcium Eq.to Atorvastatin (as pellets) Clopidogrel Bisulphate Eq.to Clopidogrel (as pellets) Aspirin (As Enteric Coated pellets) Excipients Colour: Approved Colour Used in empty Capsule shells & pellets	I.P. I.P. I.P.	20 mg 75 mg 75mg q.s.	Self	
13.2	Atorvastatin Calcium , Clopidogrel & Aspirin Capsules	Each Hard Gelatin Capsule Contains Atorvastatin Calcium Eq.to Atorvastatin (as pellets) Clopidogrel Bisulphate Eq.to Clopidogrel (as pellets) Aspirin (As Enteric Coated pellets) Excipients Colour: Approved Colour Used in empty Capsule shells & pellets	I.P. I.P. I.P.	10 mg 75 mg 75mg q.s.	Self	
13.3	Teneligliptin & Metformin hydrochloride Extended Release Tablets	Each Uncoated Bilayered Tablets Contains: Teneligliptin Hydrbromide Hydrate e.q to Teneligliptin Metformin Hydrochloride (As Extended release) Excipients Colour: Approved Colour Used		20 mg 1000 mg	Self	
13.4	Dapagliflozin & Metformin HCL.(Extended release) Tablets	Each Film Coated Tablet Contains: Metformin Hydrochloride (As Extended release) Dapagliflozin Propanediol Monohydrate E.q to Dapagliflozin Excipients Colour: Approved Colour Used	I.P	500 mg 5 mg q.s	Self	



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
13.5	Dapagliflozin & Metformin HCL.(Extended release) Tablets	Each Film Coated Tablet Contains: Metformin Hydrochloride (As Extended release) Dapagliflozin Propanediol Monohydrate E.q to Dapagliflozin Excipients Colour: Approved Colour Used	I.P	500 mg 10 mg q.s	Self	
13.6	Sodium Picosulfate Oral Solution BP	Each 5 ml Contains : Sodium Picosulfate BP In a flavoured Syrupy base Approved Colour Used	BP	5 mg q.s	Self	
13.7	Aluminium Hydroxide , Magnesium Hydroxide & Oxetacaine Gel	Each 5 ml (one teaspoonful) Contains Oxetacaine Aluminium Hydroxide (Added as Aluminium Hydroxide paste e.q.v to 0.380 g of dried Aluminium Hydroxide gel) Magnesium Hydroxide (Added as magnesium Hydroxide paste) Approved Colour Used	IP IP IP	10 mg 0.291 mg 98 mg	Self	
14.1	Polyethylene Glycol, Sodium Chloride, Sodium Hydrogen Carbonate & Potassium Chloride concentrate for oral solution	Each 25ml of Solution Contains: Macrogol 3350 (Polyethylene Glycol) Sodium Chloride Sodium Hydrogen Carbonate Potassium Chloride In a Flavoured Syrupy Base Colour: Approved Colour Used	USP IP IP IP	13.125 gm 0.3507 gm 0.1785 gm 0.0466 gm q.s	Self	MEGPEG Syrup B.NO- WDL-060 Mfg. Dt-05/2024 Exp Dt-04/2026 Mfd By: Wellmed pharma Plot no 9 A, Industrial Area Phase (3), Tahiwal Distt
14.2	Rosuvastatin Calcium & Clopidogrel Capsules	Each hard Gelatin Capsule Contain Rosuvastatin (as pellets) Clopidogrel (As pellets) Excipients Colour: Approved Colour Used in empty Capsule Shell & Pellets	IP IP	20 mg 75 mg	Self	VIROSE CV 20/75 Capsule B.NO-PC-40346 MFG.Dt-07/2024 Exp. Dt-06/2026 Nfd By: Pharmaroots Healthcare Khasra No-411 Vill. Tipra, P.O-Barotiwal
14.3	Sitagliptin Phosphate Tablet IP & Metformin Hydrochloride Tablet Ip Combipack	Each Combipack Strip Contains: (A) Sitagliptin Phosphate film coated tablets Each Film Coated Tablet Contains: Sitagliptin phosphate monohydrate e.q to 100 mg of Sitagliptin Excipients Colour: Approved Colour Used (B) Metformin Hydrochloride sustained release tablet IP. Each uncoated tablet contains 1000 mg of metformin hydrochloride IP Excipients Colour: Approved Colour Used	IP IP	100 mg q.s 1000 mg q.s	Self	DCGI Approval No./Date 1183 / 04.06.2014



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
14.4	Drotaverine Hcl. & Mefenamic Acid Tablets	Each Uncoated Tablet Contains Drotaverine Hydrochloride Eq.to Drotaverine Mefenamic Acid Excipients Approved Colour Used	I.P. I.P.	80 mg 250 mg q.s.	Self	ASTODROM-M b.No 21A2010 Mfg Dt-09/2021 Exp.Dt-08/2023 Mfg.By ASTAM Health Care PVT. Ltd Vill. Theda The Nalagarh Distt Solan-174101 (H.P)
14.5	Teneligliptin Tablets I.P. 20 mg	Each Uncoated tablet Contains Teneligliptin Hydrobromide Hydrate Eq.to Teneligliptin Excipients Colour Approved Colour Used	I.P.	20 mg q.s.	Self	I.P 2022 Volume III Page No 3737
14.6	Gliclazide MR (30mg) & Metformin (500mg) SR Tablet	Each Uncoated Bilayered Tablet Contains: Gliclazide (As Modified Release form Metformin Hydrochloride (As sustained Release form Excipients Colour: Approved Colour Used.	IP IP	30 mg 500 mg	Self	DCGI Approval No./Date 484 / 27.04.2005
14.7	Gliclazide MR (60mg) & Metformin (500mg) SR Tablet	Each Uncoated Bilayered Tablet Contains: Gliclazide (As Modified Release form Metformin Hydrochloride (As sustained Release form Excipients Colour: Approved Colour Used.	IP IP	60 mg 500 mg	Self	DCGI Approval No./Date 484 / 27.04.2005
14.8	Aciclofenac, Paracetamol & Trypsin-Chymotrypsin Tablets	Each Film Coated tablet Contains: Aciclofenac Paracetamol Trypsin-Chymotrypsin 150,000AU of enzyme activity (as entric coated granules) Excipients	IP IP	100 mg 325 mg	Self	CHYMOCET b.NO vt21158 MFG DT-11/2022 EXP DT-04/2024 MFD.BY Saphnix Life Science(AWHO, GMP Certified Company) Village-Barotiwalla, Ponta Sahib, Distt. Sirmour H.P-173025
14.9	Telmisartan 40 mg & Chlorthalidone 12.5 mg& Metoprolol succinate 50 mg ER Tablets	Each Film Coated Bilayer Tablet Contains: Metoprolol Succinate E.q to Metoprolol Tartrate (As Extended Release) Chlorthalidone Telmisartan Excipients Colour: Approved Colour Used	IP IP IP	47.50 mg 12.5 mg 40 mg q.s	Self	DCGI Approval No./Date 7 / 17.03.2020



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
15.1	Methylcobalamine, Alpha Lipoic Acid, Inositol, Folic Acid, Chromium Polynicotinate, Selenium Dioxide & Benfotiamine Capsules	Each Hard Gelatin Capsule: Methylcobalamin Alpha Lipoic Acid Inositol Folic Acid Chromium Polynicotinate Selenium Dioxide Benfotiamine Excipients Colour: Approved Colour Used in Empty Capsule Shell.	IP USP IP	1500 mcg 100 mg 100 mg 1.5 mg 200 mcg 55 mcg 150 mg q.s	Self	
15.2	Pantoprazole Gastro Resistant Tablet IP	Each Gastro-resistant Tablet Contains: Pantoprazole Sodium E.q to Pantoprazole Excipients Colour: Approved Colour Used	IP	40 mg	Self	
15.3	Esomeprazole Gastro-Resistant Tablets I.P 40 mg	Each Gastro-Resistant Tablet Contains: Esomeprazole Magnesium Trihydrate E.q to Esomeprazole Excipients Colour: Approved Colour Used	IP	40 mg q.s	Self	
15.4	Torsemide & Spironolactone Tablet	Each Uncoated Tablet Contains: Torsemide Spironolactone Excipients Colour: Approved Colour Used	IP IP	20 mg 50 mg q.s	Self	
15.5	Nimesulide & Paracetamol Tablets	Each uncoated tablet contains: Nimesulide Paracetamol Excipients Colour: Approved Colour Used	BP IP	100 mg 325 mg	Self	



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
15.6	Lycopene with Multivitamin, Multimineral and Antioxidant Capsules	Each Hard Gelatin Contains: Lycopene 10 % Methylcobalamin Grape Seed Extract 60 % Green Tea Extract 60 % Vitamin A Thiamine Mononitrate Riboflavin Pyridoxine Hydrochloride Niacinamide Folic Acid Ascorbic Acid Calcium Pantothenate Vitamin E Vitamin D3 Copper Sulphate E.q to copper Manganese Sulphate E.q to Manganese Magnesium Sulphate E.q to Magnesium Zinc Sulphate E.q to Zinc Selenium Dioxide E.q to Selenium Potassium Iodide E.q to Iodine Potassium Sulphate E.q to Potassium Excipients Colour: Approved Colour Used in Empty Capsule Shell	USP IP IP IP IP IP IP IP IP IP IP IP BP IP IP USP IP BP	4000 MCG 500 MCG 10MG 10 MG 2500IU 10MG 10 MG 3 MG 20 MG 5 MG 50 MG 5 MG 10 MG 200 IU 0.5 MG 0.5 MG 3 MG 10 MG 70 MCG 0.1 MG 2 mg	Self	
15.7	Diclofenac Potassium, Paracetamol & Serratiopeptidase Tablets	Each Film Coated Tablet Contains: Diclofenac Potassium Paracetamol Serratiopeptidase Excipients Colour: Approved Colour Used	BP IP IP	50 mg 325 mg 15 mg q.s	Self	
16.1	Nimesulide Phenylephrine Hcl. & Levocetirizine Hcl Tablets	Each Uncoated tablet Contains Nimesulide Phenylephrine Hydrochloride Levocetirizine Hydrochloride Excipients Colour Approved colour Used	B.P. I.P. I.P.	100 mg 5 mg 5 mg q.s.	Self	Fyzonim Cold Tablet Mfd. Unispeed Pharmaceuticals Pvt Ltd. Baddi H.P.
16.2	Ambroxol Hcl. Guaiphenesin & Levosalbutamol Drops	Each ml contains Levosulbutamol Sulphate Eq.to Levosalbutamol Ambroxol Hydrochloride Guaiphenesin Flavoured Syrup base Colour Approved colour Used	I.P. I.P. I.P.	0.25 mg 7.5 mg 12.5 mg q.s.	Self	AMROX-LS Drops MFD.By. Leeford Healthcare Ltd. Baddi H.P.
16.3	Paracetamol, Chlorpheniramine Maleate , Phenylephrine Hcl. & Guaiphenesin Tablets	Each Uncoated Tablet Contains Paracetamol Chlorpheniramine Maleate Phenylephrine Hydrochloride Guaiphenesin Excipients	I.P. I.P. I.P. I.P.	325mg 2 mg 5 mg 50 mg q.s.	Self	NT-COLD Tablet Mfd.By: Venus Biosciences Jharmajhri Baddi H.P.



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
16.4	Ondansetron Oral Solution I.P.	Each 5ml Contains Ondansetron Hydrochloride Eq.to Ondansetron Flavoured Syrup Base Colour: Approved Colour Used	I.P.	2 mg q.s.	Self	I.P. 2018 Page No. 2790
16.5	Vitamin B-Complex with Vitamin C & Lysine Syrup	Each 5ml Contains Thiamine Mono nitrate Riboflavin Sodium Phosphate Nicotinic Acid Niacinamide D-Panthenol	I.P. I.P. I.P. I.P. I.P.	10 mg 5 mg 5 mg 45 mg 5 mg	Self	Manufactured By Bennet Pharma Manpura Baddi H.P.
16.6	Lactic Acid Bacillus with Vitamin B-Complex Vitamin C , Vitamin E & Zinc Capsules	Each Hard Gelatine Capsule Contain Vitamin C (as ascorbic acid Coated) Niacinamide Calcium Pantothenate Nicotinic Acid Vitamin E Ascorbate (As Powder)	I.P. I.P. I.P. I.P. I.P.	150 mg 75 mg 50 mg 25 mg 25 mg	Self	Manufactured By Bennet Pharma Manpura Baddi H.P.
16.7	Promethazine Syrup I.P.	Each 5ml Contains Promethazine Hydrochloride Flavoured Syrup Base Colour: Approved Colour Used	I.P.	5 mg q.s	Self	I.P. 2018 Page No. 3016
16.8	Sitagliptin Tablets I.P. 25 mg	Each Film Coated Tablet Contains Sitagliptin Phosphate Monohydrate Eq.to Sitagliptin Excipients Colour Approved colour Used	I.P.	25 mg q.s.	Self	I.P. 2018 Page No. 3190
16.9	Sitagliptin Tablets I.P. 50 mg	Each Film Coated Tablet Contains Sitagliptin Phosphate Monohydrate Eq.to Sitagliptin Excipients Colour Approved colour Used	I.P.	50 mg q.s.	Self	I.P. 2018 Page No. 3190
16.10	Sitagliptin Tablets I.P. 100 mg	Each Film Coated Tablet Contains Sitagliptin Phosphate Monohydrate Eq.to Sitagliptin Excipients Colour Approved colour Used	I.P.	100 mg q.s.	Self	I.P. 2018 Page No. 3190
16.11	Itraconazole Capsules USP 100mg	Each Hard Gelatin Capsule contains Itraconazole (As Pellets) Excipients Colour : Approved colour Used in Fast Capsule Shell & Pellets	USP	100 mg q.s.	Self	USP 43 Page No. 2468
16.12	Itraconazole Capsules USP 200mg	Each Hard Gelatin Capsule contains Itraconazole (As Pellets) Excipients Colour : Approved colour Used in Fast Capsule Shell & Pellets	USP	200 mg q.s.	Self	USP 43 Page No. 2468



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
16.13	Montelukast Tablets I.P. 10 mg	Each Film Coated tablet Contains Montelukast Sodium Eq.to Montelukast Excipients Colour: Approved Colour Used	I.P.	10 mg q.s.	Self	I.P. 2018 Page No. 2631
16.14	Ursodeoxycholic Acid Sustained Release Tablets 450 mg	Each Film Coated Sustained Release Tablet Contains Ursodeoxycholic Acid Excipients Colour approved Colour Used	I.P.	450 mg q.s.	Self	Last Approval Date 05-05-2021
16.15.	Sitagliptin & Metformin Hcl. Tablets	Each film Coated Tablet Contains Sitagliptin Phosphate Monohydrate Eq.to Sitagliptin Metformin Hydrochloride Excipients Colour Approved colour Used	I.P. I.P.	64.25 mg 50 mg 500mg q.s	Self	DCGI Approval No./Date 148 / 02-08-2010
16.16	Sitagliptin & Metformin Hcl. Tablets	Each film Coated Tablet Contains Sitagliptin Phosphate Monohydrate Eq.to Sitagliptin Metformin Hydrochloride Excipients Colour Approved colour Used	I.P. I.P.	64.25 mg 50 mg 850mg q.s	Self	DCGI Approval No./Date 148 / 02-08-2010
16.17	Sitagliptin & Metformin Hcl. Tablets	Each film Coated Tablet Contains Sitagliptin Phosphate Monohydrate Eq.to Sitagliptin Metformin Hydrochloride Excipients Colour Approved colour Used	I.P. I.P.	64.25 mg 50 mg 1000mg q.s	Self	DCGI Approval No./Date 148 / 02-08-2010
16.18	Metoprolol Prolonged Release Tablets I.P. 100 mg	Each Film Coated Prolonged Release Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate Excipients Colour Approved colour Used	I.P.	95 mg 100mg q.s.	Self	I.P. 2018 Page No. 2585
16.19	Metoprolol (E.R.) 25 mg & Amlodipine 5 mg Tablets I.P.	Each Uncoated Bilayer tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended release) Amlodipine Besylate Eq.to Amlodipine	I.P. I.P.	23.75mg 25 mg 5 mg	Self	I.P. 2018 Page No. 2585
16.20	Metoprolol 50 mg (Extended Release) & Chlorthalidone 12.5 mg Tablets	Each Film Coated Bilayered Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended Release) Chlorthalidone Excipients	I.P. I.P.	47.5 mg 50 mg 12.5 mg	Self	DCGI Approval No./Date 71 / 25-04-2009
16.21	Metoprolol 25 mg (Extended Release) & Chlorthalidone 6.25 mg Tablets	Each Film Coated Bilayered Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended Release) Chlorthalidone Excipients	I.P. I.P.	23.75mg 25 mg 6.25 mg	Self	DCGI Approval No./Date 71 / 25-04-2009



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
16.22	Metoprolol 25 mg (Extended Release) & Telmisartan 40 mg Tablets	Each Uncoated bilayered Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended Release) Telmisartan	I.P. I.P.	23.75mg 25 mg 40 mg	Self	DCGI Approval No./Date 140 / 16-07-2010
16.23	Metoprolol 50 mg (Extended Release) & Telmisartan 40 mg Tablets	Each Uncoated bilayered Tablet Contains Metoprolol Succinate Eq.to Metoprolol Tartrate (As Extended Release) Telmisartan	I.P. I.P.	47.5mg 50 mg 40 mg	Self	DCGI Approval No./Date 140 / 16-07-2010
16.24	Propranolol 40mg (S.R. Pellets) & Flunarizine 10mg capsules	Each Hard Gelatin Capsule Contains Propranolol Hydrochloride (As Sustained Release Pellets) Flunarizine Dihydrochloride (As Pellets)	I.P. B.P.	40 mg 10 mg q.s.	Self	DCGI Approval No./ Date 48 / 16-03-20210
16.25	Propranolol 40mg (S.R. Pellets) & Flunarizine 5mg capsules	Each Hard Gelatin Capsule Contains Propranolol Hydrochloride (As Sustained Release Pellets) Flunarizine Dihydrochloride (As Pellets)	I.P. B.P.	40 mg 5 mg q.s.	Self	DCGI Approval No./Date 48 / 16-03-20210
16.26	Paracetamol, Phenylephrine Hcl., Cetirizine Hcl., Dextromethorphan Hbr. Sodium Citrate & Menthol Suspension	Each 5 ML Contains Paracetamol Phenylephrine Hydrochloride Cetirizine Hydrochloride Dextromethorphan Hydrobromide Sodium Citrate Menthol Flavoured Base Colour: Approved Colour Used	I.P. I.P. I.P. I.P. I.P.	250 mg 5 mg 2.5 mg 5 mg 25 mg 1 mg q.s.	Self	Azify Suspension Mfd. By Unispeed Pharmaceuticals Baddi H.P.
16.27	Azithromycin Suspension I.P.	Each 5 ml Contains : Azithromycin Dihydrate Eq.to. Anhydrous Azithromycin Flavoured syrup base Colour: Approved colour used	I.P	200 mg q.s.	Self	I.P. 2018 Page No. 1311
17.1	Camylofin Di Hcl. & Paracetamol Tablets	Each Film Coated Tablet Contains Camylofin Dihydrochloride Paracetamol Excipients Colour Approved Colour Used.	I.P.	25 mg 300 mg q.s.	Self	Camypan Tablets Mfd. Elfin Drugs Pvt Ltd Nalagarh H.P.
17.2	Isoxsuprine Hydrochloride 40mg Sustained Release Tablets	Each Film Coated Sustained Release tablets Contains Isoxsuprine Hydrochloride Excipients Colour Approved Colour Used.	I.P.	40 mg q.s.	Self	DCGI No./ Date 292 / April 1989
17.3	Levetiracetam Tablets I.P. 500mg	Each Film Coated Tablet Contains Levetiracetam Excipients Colour: Approved Colour Used	I.P.	500mg q.s.	Self	I.P. 2018 Page No. 2411



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
17.4	Ibuprofen & Paracetamol Suspension	Each 5ml Contains Ibuprofen Paracetamol Flavoured Base Colour: Approved Colour Used.	I.P. I.P.	100 mg 162.5 mg q.s.	Self	IBULUS Suspension Mfd. By Magma Allianz Laboratories Ltd. Ponta Sahib Distt. Sirmor H.P.
17.5	Bisoprolol Fumarate Tablets 2.5 mg	Each Film Coated Tablet Contains Bisoprolol Fumarate Excipients Colour: Approved Colour Used.	I.P.	2.5 mg q.s.	Self	DCGI Approval Date 19-01-2001
17.6	Bisoprolol Fumarate Tablets 5 mg	Each Film Coated Tablet Contains Bisoprolol Fumarate Excipients Colour: Approved Colour Used.	I.P.	5 mg q.s.	Self	DCGI No. / Date 29 / 27-02-1992
18.1	Lycopene, Anti - Oxidant, Vitamins & Essential Minerals Syrup	Each 10 ml Contains : Lycopene (10%) Niacinamide Folic Acid Selenium (As Sodium Selenate) Zinc (As Zinc Gluconate USP) Copper (As Cupric Sulphate USP) Iodine (As Potassium Iodine I.P.) Vitamin B12 Vitamin B6 Pleasantly Flavoured Sorbitol base Approved colour used	USP I.P. I.P. USP USP I.P. I.P. I.P.	2000 mcg 25 mg 100 mcg 35 mcg 3 mg 500 mcg 100 mcg 1 mcg 1.5 mcg q.s.	Self	DCGI Approval Date 25-11-2024
18.2	Vitamin D3 Oral Solution	Each 5 ml Contains : Cholecalciferol IP (In Nano Droplet from) In a flavoured sugar free syrup base Colour: Tartrazine Supra Appropriate overages of vitamin added to compensate for loss on storage	I.P q.s.	60,000IU	Self	DCGI Approval Date 25-11-2024



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
19.1	Protein & Mineral Syrup	Each 5 ml Contains : Iodised peptone Eq. to 33 mcg of Iodine Magnesium Chloride I.P. Eq. to 0.8 mg of Magnesium Manganese Sulphate BP Eq. 0.33 mg of Manganese Sodium Metavanadate Zinc Sulphate I.P. Eq. to 2.5 mg of Zinc Pyridoxine Hydrochloride I.P. Cyanocobalamin I.P. Nicotinamide I.P. Palatable base	I.P. B.P. I.P. I.P. I.P. I.P.	0.322 mg 6.67 mg 1.33 mg 0.22 mg 10.71 mg 0.25 mg 0.167 mcg 3.33 mg q.s	Self	DCGI Approval Date 25-11-2024
19.2.	Tolperisone HCL & Diclofenac Sodium tablets	Each film coated tablets contains: Tolperisone HCL Diclofenac Sodium Excipients	I.P	150 mg 50 mg q.s.	Self	DCGI Approval Date 25-11-2024
19.3	Rosuvastatin and Aspirin Capsule	Each Hard Gelatin Capsule Contains: Rosuvastatin Calcium IP Eq. to Rosuvastatin Aspirin IP (As enteric coated) Excipients Colour: Approved Colour used in hard	I.P. I.P.	20 mg 75 mg q.s.	Self	DCGI Approval Date 25-11-2024
20.1	Ergotamine Tartrate and Caffeine tablets USP	Each uncoated tablet Contains : Ergotamine Tartrate Caffeine (Anhydrous) Excipients Approved colour used	I.P. I.P.	1.0 mg 100 mg q.s.	Self	APPROVAL COPY ON DATED 14/11/2024
20.2	Cholecalciferol Chewable 60,000 IU	Each uncoated chewable tablet Contains : Cholecalciferol IP (as stabilized) Excipients Colour: Lake Sunset yellow	I.P q.s.	60,000IU	Self	D3SMEAR Batch No.AT2406008B MFG: Jun.- 2024 EXP: Nov.-2025 Manufactured by: ADDII BIOTECH PVT. LTD. Vill.Kaundi, P.O. Thana, Teh. Baddi, Distt. Solan- 173 205 (H.P)



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
20.3	Ferrous Asparto Glycinate, L-Methyl Folate & Methylcobalamin Tablets	Each film coated tablet Contains : Ferrous Asparto Glycinate Eqv. to Elemental Iron L-Methyl Folate Methylcobalamin Approved colour used	I.P	100 mg 300 mg 500 mcg q.s.	Self	APPROVAL COPY ON DATED 26/06/2024
20.4	Methylcobalamin, Alpha Lipoic Acid, Inositol, Folic Acid, Pyridoxine Hydrochloride, Vitamin D3, Chromium Polynicotinate, Chromium Carbonate & Benfotiamine tablets	Each film coated tablets Contains: Methylcobalamin Alpha Lipoic Acid Inositol Folic Acid Pyridoxine Hydrochloride Vitamin D3 Chromium Polynicotinate Chromium Carbonate Benfotiamine Excipients Colour: Red Oxide of Iron & Titanium Dioxide I.P.	I.P. USP USP I.P. I.P. I.P. I.P.	1500 mcg 100 mg 100 mg 1.5 mg 3 mg 1000 IU 200 mcg 500 mg 150 mg q.s.	Self	NVLAX – CD3 Batch No. VBT24228 MFG, AUG 2024 EXP. JUL 2026 Vats Biotech Pvt. Ltd. (A GMP & GLP Certified Co.) Khasra No. 93/45/1, Dharanji urf Dnoptan, Shamti, Distt. Solan H.P. 173212.
21.1	Doxycycline & Lactic Acid Bacillus Tablets	Each film coated tablet Contains : Doxycycline HCL Lactic Acid Bacillus Excipients Approved colour used.	I.P.	100 mg 5 BS q.s.	Self	M/s Khushru Medicare Product Permission on dated 24.09.2024 Approved Copy Attached.
21.2	Azithromycin Dispersible Tablets 100 mg	Each uncoated Dispersible tablets Contains : Azithromycin Dihydrate Eq. to Anhydrous Azithromycin Excipients Colour: Approved colour used.	I.P.	100 mg q.s	Self	M/s Khushru Medicare Product Product Permission dated on 17.08.2021 Approved Copy Attached.
21.3	Acebrophylline SR Tablets 200 mg	Each uncoated sustained released tablets Contains : Acebrophylline Excipients Approved colour used		200 mg q.s	Self	M/s Khushru Medicare Product dated on 29.09.2021 Approved Copy Attached.
21.4	Etodolac Prolonged Released / Extended release / Sustained Release tablets IP	Each Film coated PR / ER / SR tablets Contains : Etodolac Excipients Approved colour used	I.P.	600 mg q.s.	Self	M/s Khushru Medicare Product Permission on dated 22.11.2024 Approved Copy Attached.



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
21.5	Azithromycin with Lactic Acid Bacillus Tablets	Each Film coated tablets Contains : Azithromycin Anhydrous (As Azithromycin Dihydrate) Lactic Acid Bacillus Excipients Approved colour used	I.P.	250 mg 60 million Spores q.s.	Self	M/s Bonsai Pharma Product Permission on dated 21.04.2023 Approved Copy Attached.
21.6	Azithromycin with Lactic Acid Bacillus Tablets	Each Film coated tablets Contains : Azithromycin Anhydrous (As Azithromycin Dihydrate) Lactic Acid Bacillus Excipients Approved colour used	I.P.	500 mg 60 million Spores q.s.	Self	M/s Bonsai Pharma Product Permission on dated 21.04.2023 Approved Copy Attached.
22.1	Protein & Mineral Syrup	Each 5 ml Contains : Iodised peptone Eq. to 33 mcg of Iodine Magnesium Chloride I.P. Eq. to 0.8 mg of Magnesium Manganese Sulphate BP Eq. 0.33 mg of Manganese Sodium Metavanadate Zinc Sulphate I.P. Eq. to 2.5 mg of Zinc Pyridoxine Hydrochloride I.P. Cyanocobalamin I.P. Nicotinamide I.P. Palatable base	I.P. B.P. I.P. I.P. I.P. I.P.	0.322 mg 6.67 mg 1.33 mg 0.22 mg 10.71 mg 0.25 mg 0.167 mcg 3.33 mg q.s	Self	Protein & Minerals Syrup. B.No. S244003C MFG. DT.: APR 2024. Exp. DT.: SEP 2025 Mfg. By: Unispeed Pharmaceuticals Pvt. Ltd. 445/2, Behind Wrigley's, Vill. Katha, Baddi, Solan (H.P.) 173205.
23.1	Trifluoperazine, Trihexyphenidyl Hydrochloride & Chlorpromazine Hydrochloride Tablets	Each uncoated tablets Contains : Trifluoperazine Hydrochloride Eq. to Trifluoperazine Trihexyphenidyl Hydrochloride Chlorpromazine Hydrochloride Excipients Colour: Approved colour used.	I.P. I.P. I.P.	5 mg 2 mg 50 mg q.s	Self	PHEX-TC Tablets B.NO. IGT231288 MFG DT-OCT-2023 Exp. DT-SEP-2025 Mfg. By: LIVE WELL HEALTHCARE (A Cap tab Group Inc.) Plot No. 80 – A Phase 1 Jharmari Baddi, Distt. Solan (H.P.)
23.2	Paracetamol & Thiocolchicoside tablets	Each uncoated tablets Contains : Paracetamol Thiocolchicoside Excipients Approved colour used.	I.P. I.P.	500 mg 8 mg q.s	Self	THIOMAX 8 Tablets B.NO. GT412012A MFG DT-DEC-2024 Exp. DT-NOV-2026 Mfg. By: Nikvin Healthcare (I) Pvt. Ltd. Baddi – Barotiwala Road, Behind Govt. Timber Depot. Vill. Katha Baddi. Distt. Solan (H.P.)



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
23.3	L-Carnitine L- tartrate Methylcobalamin & Folic Acid Tablets	Each film coated tablet contains: L-Carnitine L- Tartrate Methylcobalamin Folic Acid Excipients Colour; Approved colour used	USP IP IP	500 mg 1500 mcg 1.5 mg q.s.	Self	M/s Khushru Medicare Product Permission on dated 30.12.2024 Approved Copy Attached.
24.1	Mecobalamin, Alpha Lipoic Acid, Folic Acid, Pyridoxine Hydrochloride & Vitamin D3 Tablets	Each film coated tablets Contains: Alpha Lipoic Acid Folic Acid Mecobalamin Pyridoxine Hydrochloride Vitamin D3 Excipients Approved Colour used	U.S.P. I.P. I.P. I.P. I.P.	100 mg 1.5 mg 1500 mcg 3 mg 1000 IU q.s.	Self	MIG-D3 Batch No. AT2403142B MFG, MAR 2024 EXP. AUG 2025 ADDI BIOTECH PVT. LTD. Vill. Kaundi P.O. Thana, Teh. baddi, Distt. Solan H.P. 173205.
24.2	Calcium Aspartate, Calcium Orotate, Vitamin D3, Zinc Sulphate, Magnesium Hydroxide, Hydroxocobalamin & Folic Acid Tablets	Each film coated tablets Contains: Calcium Aspartate Calcium Orotate Vitamin D3 (as stabilized) Zinc Sulphate monohydrate Eq. to elemental zinc Magnesium Hydroxide Hydroxocobalamin Folic Acid Excipients Colour: Titanium Dioxide	I.P I.P I.P I.P. I.P. I.P.	500 mg 500 mg 400 IU 7.5 mg 50 mg 100 mcg 1.5 mg q.s.	Self	Osedrate Batch No. 141TOS016 MFG, OCT 2024 EXP. MAR 2026 Swiss Garniers Biotech Pvt. Ltd. 21, Industrial Area, Mehtpur Dist. Una H.P. 174315.
24.3	Trioxsalen Tablets USP	Each film coated tablets Contains: Trioxsalen Excipients Colour: Titanium Dioxide	U.S.P. I.P.	25 mg q.s	Self	Approval copy Attached Mfg By: BONSAI PHARMA SITUATED AT VILL. KISHANPURA, BADDI DIST. SOLAN (HP)
24.4	Bilastine Tablets 20 mg	Each Uncoated Tablet Contains Bilastine Excipients Colour Approved colour Used		20 mg q.s	Self	Bilawise-20 Batch No. PGT25-0717A MFG, MAR 2025 EXP. FEB 2027 Plena Remedies Plot No.-17, Industrial Area (Himuda) Bhatoli kalan, Distt. Solan (HP) 173205.



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S.N.	Generic Name	Composition	Ph. Ref.	Claim	Mfg. For	Approval by Baddi FDA
24.5	Bilastine Oral Solution 2.5 mg/mL	Each mL suspension contains: Bilastine Flavoured syrup Base		2.5 mg q.s.	Self	Bilastic Batch No. PL25078A MFG, FEB 2025 EXP. JAN 2027 Plena Remedies Plot No.-17, Industrial Area (Himuda) Bhatoli kalan, Distt. Solan (HP) 173205
24.6	Doxycycline & Lactic Acid Bacillus Capsules	Each hard gelatin capsules contains: Doxycycline Hydrochloride Eq to Doxycycline Lactic Acid Bacillus Excipient	I.P.	100 mg 5 million spores q.s.	Self	ALTODOX-LB Batch No. PGC24-332D MFG, SEP 2024 EXP. AUG 2026 Plena Remedies Plot No.-17, Industrial Area (Himuda) Bhatoli kalan, Distt. Solan (HP) 173205
24.7	Doxycycline & Lactic Acid Bacillus Tablets	Each film coated tablet contains: Doxycycline Hydrochloride Eq to Doxycycline Lactic Acid Bacillus Excipient	I.P.	100 mg 90 million spores	Self	DOXBACK-LB Batch No. PGT24-236B MFG, AUG 2024 EXP. JUL 2026 Plena Remedies Plot No.-17, Industrial Area (Himuda) Bhatoli kalan, Distt. Solan (HP) 173205
24.8	Telmisartan, Cilnidipine & Chlorthalidone tablets	Each film coated tablet contains: Telmisartan Cilnidipine Chlorthalidone Excipient	I.P. I.P. I.P.	40 mg 10 mg 6.25 mg	Self	Approval copy Attached Mfg By: KHUSHRU MEDICARE SITUATED AT VILL. MESSA TIBBA ,P.O. MANJHOLI NALAGARH DIST. SOLAN (HP)