

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
1.	<b>Rosuvastatin Tablets IP 10 mg</b>	Each film coated tablet contains: Rosuvastatin Calcium equivalent to Rosuvastatin Excipients Colour:Approved colour used in tablets	IP	10 mg q.s.
2.	<b>Rosuvastatin Tablets IP 20 mg</b>	Each film coated tablet contains: Rosuvastatin Calcium equivalent to Rosuvastatin Excipients Colour:Approved colour used in tablets	IP	20 mg q.s.
3.	<b>Rosuvastatin Tablets IP 40 mg</b>	Each film coated tablet contains: Rosuvastatin Calcium IP equivalent to Rosuvastatin Excipients Colour:Approved colour used in tablets	IP	40 mg q.s.
4.	<b>Rosuvastatin &amp; Fenofibrate Tablets</b>	Each film coated tablet contains: Rosuvastatin Calcium equivalent to Rosuvastatin Fenofibrate Excipients Colour:Approved colour used in tablets	IP IP	10 mg 160 mg q.s.
5.	<b>Telmisartan Tablets IP 20 mg</b>	Each film coated tablets contains: Telmisartan Excipients Colour:Approved colour used in tablets	IP	20 mg q.s.
6.	<b>Telmisartan Tablets IP 40 mg</b>	Each uncoated tablets contains: Telmisartan Excipients Colour: Approved colour used in tablets	IP	40 mg q.s.
7.	<b>Telmisartan &amp; Amlodipine Tablets IP</b>	Each uncoated tablet contains: Telmisartan Amlodipine Besylate equivalent to Amlodipine Excipients Colour:Approved colour used in tablets	IP IP	40 mg 5 mg q.s.
8.	<b>Telmisartan &amp; Amlodipine Tablets IP</b>	Each film coated tablet contains: Telmisartan Amlodipine Besylate equivalent to Amlodipine Excipients Colour:Approved colour used in tablets	IP IP	80 mg 5 mg q.s.
9.	<b>Metoprolol Succinate Prolonged Release Tablets IP 25 mg</b>	Each film coated prolonged release tablet contains: Metoprolol Succinate equivalent to Metoprolol Tartrate Excipients Colour: Approved colour used in tablets	IP	23.75 mg 25 mg q.s.
10.	<b>Metoprolol Succinate Prolonged Release Tablets IP 50 mg</b>	Each film coated prolonged release tablet contains: Metoprolol Succinate equivalent to Metoprolol Tartrate Excipients Colour: Approved colour used in tablets	IP	47.5 mg 50 mg q.s.
11.	<b>Azithromycin Tablets IP 250 mg</b>	Each film coated tablet contains: Azithromycin (as dihydrate) Eq. to Azithromycin Excipients Colour: Approved colour used in tablets	IP	250 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
12.	<b>Azithromycin Tablets IP 500 mg</b>	Each film coated tablet contains: Azithromycin (as dihydrate) Eq. to Azithromycin Excipients Colour: Approved colour used in tablets	IP	500 mg q.s.
13.	<b>Calcium Citrate Maleate with Vitamin D3 Tablets</b>	Each film coated tablet contains: Calcium Citrate Maleate Eq. to elemental Calcium Vitamin D3 Excipients Colour: Approved colour used in tablets	IP	250 mg 200 IU q.s.
14.	<b>Calcium with Vitamin D3 Tablets</b>	Each film coated tablet contains: 1.25g of Calcium Carbonate from an organic source (Oyster shell) Eq. to elemental calcium Vitamin D3 Excipients Colour: Approved colour used in tablets	IP IP	500 mg 200 IU q.s.
15.	<b>Clarithromycin Tablets IP 250mg</b>	Each film coated tablets contains: Clarithromycin Excipients Colour: Approved colour used in tablets	IP	250 mg q.s.
16.	<b>Clarithromycin Tablets IP 500mg</b>	Each film coated tablets contains: Clarithromycin Excipients Colour: Approved colour used in tablets	IP	500 mg q.s.
17.	<b>Diclofenac Potassium, Paracetamol &amp; Serratiopeptidase Tablets</b>	Each film coated tablets contains: Diclofenac Potassium Paracetamol Serratiopeptidase (As enteric coated granules) (Eq. to enzyme activity 20,000units) Excipients Colour: Approved colour used in tablets	IP IP IP	50 mg 325 mg 10 mg  q. s
18.	<b>Aceclofenac, Paracetamol &amp; Serratiopeptidase Tablets</b>	Each film coated tablets contains: Aceclofenac Paracetamol Serratiopeptidase (As enteric coated granules) (Eq. to enzyme activity 30,000units) Excipients Colour: Approved colour used in tablets	IP IP IP	100 mg 325 mg 15 mg  q. s
19.	<b>Montelukast Sodium &amp; Levocetirizine Hydrochloride Tablets IP</b>	Each uncoated tablets contains: Montelukast Sodium Eq. to Montelukast Levocetirizine Hydrochloride Excipients Colour: Approved colour used in tablets	IP IP	10 mg 5 mg q.s.
20.	<b>Aceclofenac &amp; Paracetamol Tablets IP</b>	Each film coated tablets contains: Aceclofenac Paracetamol Excipients Colour: Approved colour used in tablets	IP IP	100 mg 325 mg q. s
21.	<b>Itraconazole Capsules IP 100 mg</b>	Each hard gelatin capsules contains: Itraconazole (As pellets) Excipients Approved colour used in empty capsule shells	IP	100 mg  q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
22.	<b>Itraconazole Capsules IP 200 mg</b>	Each hard gelatin capsules contains: Itraconazole (As pellets) Excipients Approved colour used in empty capsule shells	IP	200 mg  q.s.
23.	<b>Omeprazole &amp; Domperidone Capsules IP</b>	Each hard gelatin capsules contains: Omeprazole (As enteric coated granules) Domperidone Excipients Approved colour used in empty capsule shells	IP  IP	20 mg  10 mg q.s.
24.	<b>Pregabalin &amp; Methylcobalamin Capsules IP</b>	Each hard gelatin capsules contains: Pregabalin Methylcobalamin Excipients Approved colour used in empty capsule shells	IP IP	75 mg 750 mcg q.s.
25.	<b>Lactic Acid Bacillus, B-Complex with Vitamins Capsules</b>	Each hard gelatin contains: Lactic Acid Bacillus Acidophilus  Folic Acid Niacinamide Calcium Pantothenate Vitamin B1 Vitamin B2 Vitamin B6 Vitamin B12 Vitamin C Excipients Approved colour used in empty capsule shells (Appropriate Overages added of vitamins to compensate )	  IP IP IP IP IP IP IP IP	2000 lacs Spores  300 mcg 26 mg 5 mg 2 mg 3 mg 1.5 mg 1 mcg 50 mg q. s
26.	<b>Rabeprazole Sodium (EC) &amp; Domperidone (SR) Capsules</b>	Each hard gelatin capsules contains: Rabeprazole Sodium (As enteric coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empty capsule shells	IP  IP	20 mg  30 mg  q.s.
27.	<b>Rabeprazole Sodium (EC) &amp; Levosulpiride (SR) Capsules</b>	Each hard gelatin capsules contains: Rabeprazole Sodium (As enteric coated pellets) Levosulpiride (As sustained release pellets) Excipients Approved colour used in empty capsule shells	IP  IP	20 mg  75 mg q.s.
28.	<b>Esomeprazole (EC) &amp; Domperidone (SR) Capsules</b>	Each hard gelatin capsules contains: Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (As enteric coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empty capsule shells	IP  IP	20 mg  30 mg  q.s.
29.	<b>Esomeprazole (EC) &amp; Itopride Hydrochloride (SR) Capsules</b>	Each hard gelatin capsules contains: Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (As enteric coated pellets) Itopride Hydrochloride (As sustained release pellets) Excipients Approved colour used in empty capsule shells	IP  IP	40 mg  150 mg  q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
<b>30.</b>	<b>Esomeprazole (EC) &amp; Domperidone (SR) Capsules</b>	Each hard gelatin capsules contains: Esomeprazole Magnesium Trihydrate Eq. to Esomeprazole (As enteric coated pellets) Domperidone (As sustained release pellets) Excipients Approved colour used in empty capsule shells	IP  IP	40 mg  30 mg  q.s.
<b>31.</b>	<b>Pantoprazole Gastro Resistant &amp; Domperidone Prolonged Release Capsules IP</b>	Each hard gelatin capsules contains: Pantoprazole Sodium Sesquihydrate Eq. to Pantoprazole (As gastro resistant pellets) Domperidone (As prolonged release form) Excipients Approved colour used in empty capsule shells	IP  IP	40 mg  30 mg  q.s.
<b>32.</b>	<b>Omeprazole Gastro Resistant Capsules IP 20mg</b>	Each hard gelatin capsules contains: Omeprazole (As gastro resistant granules) Excipients Approved colour used in empty capsule shells	IP	20 mg  q.s.
<b>33.</b>	<b>Dextromethorphan Hydrobromide, Phenylephrine Hydrochloride &amp; Chlorpheniramine Maleate Syrup</b>	Eah 5 ml Contains: Dextromethorphan Hydrobromide Phenylephrine Hydrochloride Chlorpheniramine Maleate In a flavoured syrupy base Colour: Approved colour used	IP IP IP	10 mg 5 mg 2 mg q.s.
<b>34.</b>	<b>Montelukast &amp; Levocetirizine Dihydrochloride Syrup</b>	Eah 5 ml Contains: Montelukast Sodium Montelukast Levocetirizine Dihydrochloride In a flavoured syrupy base Colour: Approved colour used	IP  IP	4 mg 2.5 mg q.s.
<b>35.</b>	<b>Levosaltbutamol Sulphate, Ambroxol Hydrochloride &amp; Guaiphenesin Syrup</b>	Eah 5 ml contains: Levosaltbutamol Sulphate Eq. to Levosaltbutamol Ambroxol Hydrochloride Guaiphenesin In a flavoured base Colour: Approved colour used	IP  IP  IP	1 mg 30 mg 50 mg q.s.
<b>36.</b>	<b>Terbutaline Sulphate, Ambroxol Hydrochloride, Guaiphenesin &amp; Menthol Syrup</b>	Each 5 ml contains: Terbutaline Sulphate Ambroxol Hydrochloride Guaiphenesin Menthol In a non-syrupy base Colour: Approved colour used	IP IP IP IP	1.25 mg 15 mg 50 mg 1.5 mg q.s.
<b>37.</b>	<b>Digestive Enzyme Syrup</b>	Each 5 ml contains Diastase (1:1200) (Fungal diastase derived from Aspergillus Oryzae digests not less than 60 gm of cooked starch). Pepsin (1:3000) (Digests not less than 30 gm of coagulated egg albumin) in a flavoured syrupy base Colour: Approved colour used.	IP  IP	50 mg  10 mg  q.s.
<b>38.</b>	<b>Mefenamic Acid &amp; Paracetamol Suspension</b>	Eah 5 ml (One teaspoonful) Contains: Mefenamic Acid Paracetamol In a flavoured syrupy base Colour: Approved colour Used	IP IP	100 mg 250 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
39.	<b>Megaldrate &amp; Simethicone Syrup+</b>	Eah 5 ml (One teaspoonful) Contains: Megaldrate Simethicone In a flavoured syrupy base Colour: Approved colour Used	IP IP	480mg 20mg q.s.
40.	<b>Megaldrate, Simethicone &amp; Oxetacaine Suspension</b>	Eah 5 ml Contains: Megaldrate Simethicone Oxetacaine Sorbitol base Colour: Approved colour Used	IP IP BP	540 mg 50 mg 10 mg q.s.
41.	<b>Cyproheptadine and Tricholine Citrate Syrup</b>	Each 5 ml contains: Cyproheptadine Hydrochloride Tricholine Citrate Solution(65 %) In a flavoured syrupy base Colour: Approved colour Used	IP	2 mg 275 mg q.s.
42.	<b>Disodium Hydrogen Citrate Syrup</b>	Each 5 ml contains: Disodium Hydrogen Citrate Flavoured Syrup base Colour: Approved colour Used	BP	1.38gm q.s.
43.	<b>Paracetamol, Phenylephrine Hydrochloride &amp; Chlorpheniramine Maleate Drops</b>	Eah ml contains: Paracetamol Phenylephrine Hydrochloride Chlorpheniramine Maleate In a flaoured non-syrupy base Colour: Approved colour Used	IP IP IP	125 mg 5 mg 1 mg q.s.
44.	<b>Ibuprofen &amp; Paracetamol Suspension</b>	Eah 5 ml contains: Ibuprofen Paracetamol Flavoured syrup base Colour: Approved colour used	IP IP	100 mg 162.5 mg q.s.
45.	<b>Cefixime Oral Suspension IP 50 mg</b>	Each 5 ml (After reconstitution) contains: Cefixime (as Trihydrate) Eq. to anhydrous Cefixime Excipients Colour: Approved Colour used	IP	50 mg q.s.
46.	<b>Cefixime Oral Suspension IP 100 mg</b>	Each 5 ml (After reconstitution) contains: Cefixime (as Trihydrate) Eq. to anhydrous Cefixime Excipients Colour: Approved Colour used	IP	100 mg q.s.
47.	<b>Cefpodoxime Oral suspension IP 50 mg</b>	Each 5 ml (After reconstitution ) contains: Cefpodoxime Proxetil Eq. to Cefpodoxime Excipients Colour: Approved Colour used	IP	50 mg q.s.
48.	<b>Cefpodoxime Oral suspension IP 100 mg</b>	Each 5 ml (After reconstitution) contains: Cefpodoxime Proxetil Eq. to Cefpodoxime Excipients Colour: Approved Colour used	IP	100 mg q.s.
49.	<b>Cefixime and Ofloxacin oral suspension</b>	Each 5 ml of reconstitution contains: Cefixime (as Trihydrate) Eq. to anhydrous Cefixime Ofloxacin Excipients Colour: Approved Colour used	IP IP	50 mg 50 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
50.	<b>Cefixime with Lactic Acid Bacillus Oral Suspension</b>	Each 5 ml of reconstitution contains: Cefixime (as Trihydrate) Eq. to anhydrous Cefixime Lactic Acid Bacillus Excipients Colour: Approved Colour used	IP	50 mg 60 million spores q.s.
51.	<b>Cefixime with Lactic Acid Bacillus Oral Suspension</b>	Each 5 ml of reconstitution contains: Cefixime (as Trihydrate) Eq. to anhydrous Cefixime Lactic Acid Bacillus Excipients Colour: Approved Colour used	IP	100 mg 60 million spores q.s.
52.	<b>Permethrin Lotion</b>	Permethrin Perfume Lotion Base.	IH	5 % w/v q.s.
53.	<b>Povidone Iodine, Metronidazole &amp; Sucralfate Ointment</b>	Povidone Iodine (Available Iodine 0.5% w/w) Metronidazole Sucralfate Water soluble ointment base	IP IP IP	5.00% w/w 1.00% w/w 7.00% w/w q.s.
54.	<b>Betamethasone, Clindamycin &amp; Nicotinamide Cream</b>	Betamethasone Valerate Eq to Betamethasone Clindamycin Phosphate Eq. to Clindamycin Nicotinamide Cream Base	IP IP IP	0.1% w/w 1.0% w/w 4.0% w/w q.s.
55.	<b>Mupirocin Ointment IP</b>	Mupirocin in a ointment base	IP	2.0% w/w q.s.
56.	<b>Povidone Iodine Ointment USP 5.0% w/w</b>	Povidone Iodine (Available Iodine) Water soluble ointment base	IP	5.0% w/w 0.5% w/w q.s.
57.	<b>Povidone Iodine Ointment USP 10.0% w/w</b>	Povidone Iodine (Available Iodine) Water soluble ointment base	IP	10.0% w/w 1.0% w/w q.s.
58.	<b>Diclofenac Gel</b>	Diclofenac Diethylamine Eq. Diclofenac Sodium Gel base	BP	1.16% w/w 1.0% w/w q.s.
59.	<b>Hydroquinone, Tretinoin &amp; Mometasone Furoate Cream</b>	Hydroquinone Tretinoin Mometasone Furoate Cream Base	USP USP IP	2.0% w/w 0.025% w/w 0.1% w/w q.s.
60.	<b>Roxithromycin Tablets IP 150mg</b>	Each film coated tablets contains: Roxithromycin (Anhydrous) Excipients Colour: Approved colour used in tablets	IP	150 mg q.s.
61.	<b>Methylcobalamin Orally Disintegrating Tablets</b>	Each mouth dissolving tablet contains: Methylcobalamin Excipients	IP	1500 mcg q.s.
62.	<b>Desloratadine Tablets 5 mg</b>	Each film coated tablet contains: Desloratadine Excipients Approved colour used in tablets	IH	5 mg q.s.
63.	<b>Montelukast &amp; Bilastine Tablets</b>	Each film coated tablet contains: Montelukast Sodium equivalent to Montelukast Bilastine Excipients Colour: Approved colour used in tablets	IP	10 mg 20 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
<b>64.</b>	<b>Ursodeoxycholic Acid Tablets IP 150mg</b>	Each uncoated tablet contains: Ursodeoxycholic Acid Excipients Colour: Approved colour used in tablets	IP	150 mg q.s.
<b>65.</b>	<b>Ursodeoxycholic Acid Tablets IP 300mg</b>	Each uncoated tablet contains: Ursodeoxycholic Acid Excipients Approved colour used in tablets	IP	300 mg q.s.
<b>66.</b>	<b>Methylcobalamin, Alpha Lipoic Acid, Folic Acid, Pyridoxine &amp; Vitamin D3 Tablets</b>	Each film coated tablet contains: Methylcobalamin Alpha Lipoic Acid Folic Acid Pyridoxine HCl Vitamin D3 Excipients Approved colour used in tablets	IP USP IP IP IP	1500 mcg 100 mg 1.5 mg 3 mg 1000 IU q.s.
<b>67.</b>	<b>Prochlorperazine Maleate &amp; Acetaminophen Tablets</b>	Each uncoated tablet contains: Prochlorperazine Maleate Acetaminophen Excipients	IP IP	5 mg 650 mg q.s.
<b>68.</b>	<b>Rosuvastatin, Asprin &amp; Clopidogrel Capsules</b>	Each hard gelatin capsules contains: Rosuvastatin Calcium Eq. to Rosuvastatin (As pellets) Asprin (As gastro-resistant pellets) Clopidogrel Bisulphate eq. to Clopidogrel (As pellets) Excipients Approved colour used in empty capsule shells & Pellets	IP  IP  IP	10 mg  75 mg  75 mg  q.s.
<b>69.</b>	<b>Aspirin Gastro-resistant and Rosuvastatin Capsules IP</b>	Each hard gelatin capsules contains: Asprin (As enteric coated pellets ) Rosuvastatin Calcium Eq. to Rosuvastatin (As pellets) Excipients Approved colour used in empty capsule shells & Pellets	IP  IP	75 mg  20 mg q.s.
<b>70.</b>	<b>Aspirin Gastro-resistant and Rosuvastatin Capsules IP</b>	Each hard gelatin capsules contains: Asprin (As enteric coated pellets ) Rosuvastatin Calcium Eq. to Rosuvastatin (As pellets) Excipients Approved colour used in empty capsule shells & Pellets	IP  IP	150 mg  20 mg q.s.
<b>71.</b>	<b>B- Complex, L-Lysine Hydrochloride Syrup</b>	Each 15 ml contains: L-Lysine Hydrochloride Niacinamide Thiamine Hydrochloride Riboflavin Sodium Phosphate eq. to Riboflavin D-Panthenol Pyridoxine Hydrochloride Cyanocobalamin Flavoured syrupbase Colour: Approved colour used	USP IP IP IP  IP IP IP	50 mg 25 mg 2 mg  2.5 mg 2.5 mg 1 mg 1mcg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
72.	<b>Luliconazole Cream IP 1.0% w/w</b>	Luliconazole Benzyl Alcohol (As a preservative) Cream base	IP IP	1.0% w/w 1.0% w/w  q.s.
73.	<b>Diclofenac Diethylamine, Methyl Salicylate, Linseed Oil &amp; Menthol Gel</b>	Diclofenac Diethylamine equipment to Diclofenac Sodium Methyl Salicylate Linseed Oil Menthol Benzyl Alcohol (As a preservative) Gel base	IP  IP BP IP IP	1.16% w/w 1.0% w/w 10.0% w/w 3.0% w/w 5.0% w/w 1.0% w/w q.s.
74.	<b>Desonide Cream 0.05% w/w</b>	Desonide <b>Preservatives:</b> Potassium Sorbate IP and Sorbic Acid IP Combination equipment to Sorbic Acid In a cream base	USP	0.05% w/w   0.175% w/w q.s.
75.	<b>Gliclazide &amp; Metformin Hydrochloride Tablets</b>	Each uncoated tablet contains: Gliclazide Metformin Hydrochloride Excipients	IP IP	80 mg 500 mg q.s.
76.	<b>Vildagliptin Tablets 50 mg</b>	Each uncoated tablet contains: Vildagliptin Excipients		50 mg q.s.
77.	Nitrofurantoin Tablets BP 20 mg  <b>FURACILIN</b>	Each uncoated tablet contains: Nitrofurantoin Excipients	BP	20 mg q.s.
78.	<b>Calcium, Vitamin D3, Methylcobalamin, L- Methylfolate Calcium &amp; Pyridoxal-5-Phosphate Tablets</b>	Each film coated tablets contains: Calcium Carbonate Eq. to Elemental Calcium Cholecalciferol (Vitamin D3) (As Stabilized Form) Mecobalamin (Methylcobalamin) L-Methylfolate Calcium Pyridoxal-5-Phosphate Excipients Colour: Approved colour used in tablets	IP  IP  IP	1250 mg 500 mg 2000 IU  1500 mcg 1 mg 20 mg q.s.
79.	<b>Glimepiride &amp; Metformin Hydrochloride (ER) Tablets</b>	Each uncoated bilayered tablet contains: Glimepiride Metformin HCl (As extended release) Excipients Colour: Approved colour used in tablets	IP IP	2 mg 500 mg  q.s.
80.	<b>Voglibose &amp; Metformin Hydrochloride (SR) Tablets</b>	Each uncoated bilayered tablet contains: Voglibose Metformin Hydrochloride (As sustained release form) Excipients Colour: Approved colour used in tablets	IP IP	0.3 mg 500 mg  q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
<b>81.</b>	<b>Glimepiride &amp; Metformin Hydrochloride (SR) Tablets</b>	Each uncoated bilayered tablet contains: Glimepiride Metformin Hydrochloride (As sustained release form) Excipients Colour: Approved colour used in tablets	IP IP	1 mg 1000 mg  q.s.
<b>82.</b>	<b>Glimepiride &amp; Metformin Hydrochloride (SR) Tablets</b>	Each uncoated bilayered tablet contains: Glimepiride Metformin Hydrochloride (As sustained release form) Excipients Colour: Approved colour used in tablets	IP IP	2 mg 1000 mg  q.s.
<b>83.</b>	<b>Deflazacort Tablets 6 mg</b>	Each uncoated tablet contains: Deflazacort Excipients Colour: Approved colour used in tablets	IH	6 mg q.s.
<b>84.</b>	<b>Cilnidipine Tablets IP 5 mg</b>	Each film coated tablet contains: Cilnidipine Excipients Approved colour used in tablets	IP	5 mg q.s.
<b>85.</b>	<b>Cilnidipine Tablets IP 10 mg</b>	Each film coated tablet contains: Cilnidipine Excipients Approved colour used in tablets	IP	10 mg q.s.
<b>86.</b>	<b>Telmisartan &amp; Metoprolol Succinate (ER) Tablets</b>	Each uncoated bilayered tablet contains: Telmisartan Metoprolol Succinate Eq. to Metoprolol Tartartrate (As extended release) Excipients Colour: Approved colour used in tablets	IP IP	40 mg 23.75 mg 25 mg q.s.
<b>87.</b>	<b>Telmisartan &amp; Metoprolol Succinate (ER) Tablets</b>	Each uncoated bilayered tablet contains: Telmisartan Metoprolol Succinate Eq. to Metoprolol Tartartrate (As extended release) Excipients Colour: Approved colour used in tablets	IP IP	40 mg 47.5 mg 50 mg q.s.
<b>88.</b>	<b>Sitagliptin Phosphate &amp; Metformin HCl Tablets</b>	Each film Coated tablets contains : Sitagliptin Phosphate monohydrate Eq. to Sitagliptin Metformin Hydrochloride Excipients Colour: Approved Colour Used in Tablets	IP IP	50 mg 500 mg q.s.
<b>89.</b>	<b>Amlodipine Tablets IP 5 mg</b>	Each uncoated tablets contains: Amlodipine Besylate Eq. to Amlodipine Excipients Approved colour used in tablets	IP	5 mg q.s.
<b>90.</b>	<b>Aceclofenac (SR) Tablets 200 mg</b>	Each film coated tablet contains: Aceclofenac (As sustained release form) Excipients Approved colour used in tablets	IP	200 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
91.	<b>Dapagliflozin &amp; Metformin Hydrochloride (ER) Tablets</b>	Each film coated tablet contains: Dapagliflozin Propanediol Monohydrate equivalent to Dapagliflozin Metformin Hydrochloride (As Extended Release ) Excipients Approved colour used in tablets	IP	10 mg 500 mg  q.s.
92.	<b>Acebrophylline Sustained Release Tablets 200 mg</b>	Each film coated sustained release tablet contains: Acebrophylline Excipients Colour: Approved colour used in tablets		200 mg q. s.
93.	<b>Febuxotat Tablet 40 mg</b>	Each film coated tablet contains: Febuxot Excipients Colour: Approved colour used in tablets	IP	40 mg q.s.
94.	<b>Pioglitazone Hydrochloride Tablets IP 15 mg</b>	Each uncoated tablet contains: Pioglitazone Hydrochloride Eq. to Pioglitazone Excipients	IP	15 mg q.s.
95.	<b>Pioglitazone Hydrochloride Tablets IP 30 mg</b>	Each uncoated tablet contains: Pioglitazone Hydrochloride Eq. to Pioglitazone Excipients	IP	30 mg q.s.
96.	<b>Voglibose Mouth Dissolving Tablets 0.2 mg</b>	Each uncoated mouth dissolving tablet contains: Voglibose Excipients	IP	0.2 mg q.s.
97.	<b>Voglibose Mouth Dissolving Tablets 0.3 mg</b>	Each uncoated mouth dissolving tablet contains: Voglibose Excipients	IP	0.3 mg
98.	<b>Hydroxychloroquine Tablets IP 200 mg</b>	Each film coated tablet contains: Hydroxychloroquine Sulphate Excipients Colour: Approved colour used in tablets	IP	200 mg q.s.
99.	<b>Hydroxychloroquine Tablets IP 300 mg</b>	Each film coated tablet contains: Hydroxychloroquine Sulphate Excipients Colour: Approved colour used in tablets	IP	300 mg q.s.
100.	<b>Hydroxychloroquine Tablets IP 400 mg</b>	Each film coated tablet contains: Hydroxychloroquine Sulphate Excipients Colour: Approved colour used in tablets	IP	400 mg q.s.
101.	<b>Teneligliptin Tablets 20 mg</b>	Each film coated tablet contains: Teneligliptin Hydrobromide Hydrate eq. to Teneligliptin Excipients Colour: Approved colour used in tablets		20 mg q.s.
102.	<b>Teneligliptin &amp; Metformin Hydrochloride Tablets (20 mg + 500 mg)</b>	Each film coated tablet contains: Teneligliptin Hydrobromide Hydrate eq. to Teneligliptin Metformin Hydrochloride (As Extended Release ) Excipients Colour: Approved colour used in tablets	IP	20 mg 500 mg  q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
<b>103.</b>	<b>Teneligliptin &amp; Metformin Hydrochloride Tablets (20 mg + 1000 mg)</b>	Each film coated tablet contains: Teneligliptin Hydrobromide Hydrate eq. to Teneligliptin Metformin Hydrochloride (As Extended Release ) Excipients Colour: Approved colour used in tablets	IP	20 mg 1000 mg  q.s.
<b>104.</b>	<b>Ticagrelor Tablets IP 90 mg</b>	Each film coated tablet contains: Ticagrelor Excipients Colour: Approved colour used in tablets	IP	90 mg q.s.
<b>105.</b>	<b>Glimepiride, Pioglitazone Hydrochloride &amp; Metformin Hydrochloride (ER) Tablets</b>	Each uncoated bilayered tablet contains: Glimepiride Pioglitazone Hydrochloride eq.to Pioglitazone Metformin Hydrochloride (As extended release) Excipients Colour: Approved colour used in tablets	IP IP  IP	1 mg 15 mg  500 mg  q.s.
<b>106.</b>	<b>Glimepiride, Pioglitazone Hydrochloride &amp; Metformin Hydrochloride (ER) Tablets</b>	Each uncoated bilayered tablet contains: Glimepiride Pioglitazone Hydrochloride eq.to Pioglitazone Metformin Hydrochloride (As extended release) Excipients Colour: Approved colour used in tablets	IP IP  IP	2 mg 15 mg  500 mg  q.s.
<b>107.</b>	<b>Multi Vitamin Mineral Syrup</b>	Each 5 ml contains: Zinc Sulphate Nicotinamide L-Lysine Hydrochloride D-Panthenol Thiamine Hydrochloride Riboflavin Phosphate Sodium Pyridoxine Hydrochloride Copper sulphate Potassium Iodide Sodium Selenite Pentahydrate Eq. to Selenium Cyanocobalamine Vitamin A (as palmitate) Cholecalciferol Vitamin E (As di $\alpha$ -Tocopheryl acetate ) Flavoured Syrupy Base Colour: Approved colour used	IP IP USP IP IP IP IP IP BP IP  IP IP IP IP	22.5 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
<b>108.</b>	<b>Minoxidil &amp; Finasteride Topical Solution</b>	Minoxidil Finasteride base	IP IP	5.00% w/v 0.1% w/v q.s.
<b>109.</b>	<b>Azithromycin &amp; Lactic Acid Bacillus Tablets</b>	Each film coated tablet contains: Azithromycin Dihydrate eq. to Azithromycin (Anhydrous) Lactic Acid Bacillus Excipients Colour: Approved colour used in tablets	IP	250 mg 60 M.spores q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
110.	<b>Azithromycin &amp; Lactic Acid Bacillus Tablets</b>	Each film coated tablet contains: Azithromycin Dihydrate eq. to Azithromycin (Anhydrous) Lactic Acid Bacillus Excipients Colour: Approved colour used in tablets	IP	500 mg 60 M.spores q.s.
111.	<b>Terbutaline Sulphate, Ambroxol Hydrochloride, Guaiphenesin &amp; Syrup</b>	Eah 5 ml contains: Terbutaline Ambroxol Hydrochloride Guaiphenesin Menthol Flavoured syrupy base Colour: Approved colour used in Tablets	IP IP IP IP	1.25 mg 15 mg 50 mg 2.5 mg q.s.
112.	<b>Metoprolol Succinate Prolonged Release &amp; Amlodipine Tablets IP</b>	Each uncoated bilayered tablet contains: Metoprolol Succinate eq. to Metoprolol Tartartrate (As extended release form) Amlodipine Besylate eq. to Amlodipine Excipients Colour: Approved colour used in tablets	IP  IP	47.5 mg 50 mg  5 mg q.s.
113.	<b>Metoprolol Succinate (ER) 50 mg &amp; Amlodipine Besylate 5 mg Tablets</b>	Each uncoated bilayered tablet contains: Metoprolol Succinate eq. to Metoprolol Tartartrate (As extended release form) Amlodipine Besylate eq. to Amlodipine Excipients Colour: Approved colour used in tablets	IP  IP	47.5 mg 50 mg  5 mg q.s.
114.	<b>Metoprolol Succinate Prolonged Release &amp; Amlodipine Tablets IP</b>	Each uncoated bilayered tablet contains: Metoprolol Succinate eq. to Metoprolol Tartartrate (As extended release form) Amlodipine Besylate eq. to Amlodipine Excipients Colour: Approved colour used in tablets	IP  IP	23.75 mg 25 mg  2.5 mg q.s.
115.	<b>Drotaverine Hydrochloride &amp; Aceclofenac Tablets</b>	Each film coated tablet contains: Drotaverine Hydrochloride Aceclofenac Excipients Approved colour used in tablets	IP IP	80 mg 100 mg q.s.
116.	<b>Ondansetron Oral Suspension IP</b>	Eah 5 ml contains: Ondansetron Hydrochloride Eq. to Ondansetron In a flavoured syrupy base Colour: Approved colour used	IP	2 mg q.s.
117.	<b>Metoprolol Succinate Prolonged Release 25 mg &amp; Amlodipine 2.5 mg Tablets IP</b>	Each uncoated bilayered tablet contains: Metoprolol Succinate eq. to Metoprolol Tartartrate (As extended release form) Amlodipine Besylate eq. to Amlodipine Excipients Colour: Approved colour used in tablets	IP  IP	23.75 mg 25 mg  2.5 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
118.	<b>Drotaverine Hydrochloride &amp; Aceclofenac Tablets</b>	Each film coated tablet contains: Drotaverine Hydrochloride Aceclofenac Excipients Approved colour used in tablets	IP IP	80 mg 100 mg q.s.
119.	<b>Ondansetron Orally Disintegration Tablets IP 4 mg</b>	Each uncoated mouth dissolving tablets contains: Ondansetron Hydrochloride Eq. to Ondansetron Excipients	IP	4 mg q.s.
120.	<b>Methylprednisolone Tablets IP 4mg</b>	Each film coated tablet contains: Methylprednisolone Excipients Colour: Approved colour used in tablets	IP	4 mg q.s.
121.	<b>Methylprednisolone Tablets IP 8mg</b>	Each film coated tablet contains: Methylprednisolone Excipients Colour: Approved colour used in tablets	IP	8 mg q.s.
122.	<b>Methylprednisolone Tablets IP 16mg</b>	Each film coated tablet contains: Methylprednisolone Excipients Colour: Approved colour used in tablets	IP	16 mg q.s.
123.	<b>Calcium Citrate, Calcitriol, Zinc &amp; Manganese Tablets</b>	Each film coated tablet contains: Calcium Citrate Calcitriol Zinc Sulphate Monohydrate Eq. to elemental Zinc Manganese (As Manganese Sulphate) Excipients Colour: Approved colour used in tablets	USP BP IP  IP	1000 mg 0.25 mcg  7.5 mg 40 mg q.s.
124.	<b>Nifedipine Sustained Release Tablets IP 30 mg</b>	Each sustained release film coated tablet contains: Nifedipine Excipients Colour: Approved Colour Used in Tablets	IP	30 mg q.s.
125.	<b>Acebrophylline &amp; N-Acetylcysteine Tablets</b>	Each film coated tablet contains: Acebrophylline N-Acetylcysteine Excipients Colour: Approved Colour Used in Tablets	BP	100 mg 600 mg q.s.
126.	<b>Ofloxacin &amp; Ornidazole Tablets IP</b>	Each film coated tablets contains: Ofloxacin Ornidazole Excipients Colour: Approved colour used in tablets	IP IP	200 mg 500 mg q.s.
127.	<b>Ondansetron Oral Suspension IP</b>	Eah 5 ml contains: Ondansetron Hydrochloride Eq. to Ondansetron In a flavoured syrupy base Colour: Approved colour used	IP	2 mg q.s.
128.	<b>Ibuprofen &amp; Paracetamol Suspension</b>	Eah 5 ml contains: Ibuprofen Paracetamol Flavoured syrup base Colour: Approved colour used	IP IP	100 mg 162.5 mg q.s.

**APPROVED PRODUCTS LIST M/S APM MEDICHEM PRIVATE LIMITED**

<b>Sr. No.</b>	<b>Generic Name</b>	<b>Composition</b>	<b>Ph. Ref.</b>	<b>Strength</b>
<b>129.</b>	<b>Ofloxacin &amp; Ornidazole Tablets IP</b>	Each film coated tablets contains: Ofloxacin Ornidazole Excipients Colour: Approved colour used in tablets	IP IP	200 mg 500 mg q.s.
<b>130.</b>	<b>Naproxen and Domperidone Tablets</b>	Each film coated tablet contains: Naproxen Domperidone Excipients Colours: Approved colour used in tablets	IP IP	250 mg 10 mg q.s.
<b>131.</b>	<b>Naproxen and Domperidone Tablets</b>	Each film coated tablet contains: Naproxen Domperidone Excipients Colours: Approved colour used in tablets	IP IP	500 mg 10 mg q.s.
<b>132.</b>	<b>Piracetam Tablets 800 mg</b>	Each uncoated tablet contains: Piracetam Excipients	IP	800 mg q.s.