

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
01	Cyproheptadine HCl & Tricholine citrate Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Cyproheptadine HCl	IP	2	mg	
		Eq. to Cyproheptadine HCl Anhydrous				
		Tricholine Citrate		275	mg	
		In a flavoured sorbitol base (70%)_syrup base				
Approved Color used						
02	Ferrous Ascorbate & Folic Acid Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Ferrous Ascorbate		30	mg	
		Eq. to elemental Iron				
		Folic Acid	IP	550	mcg	
		In a Flavoured vehicle base		q.s.		
Approved Color used						
Appropriate overages added						
03	Levocetirizine Dihydrochloride Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Levocetirizine Dihydrochloride	IP	2.5	mg	
		In a flavoured syrup base		q.s.		
04	Paracetamol Oral Suspension IP	Each 5 ml contains:				Approved on 28.09.2017
		Paracetamol	IP	250	mg	
		In a Flavoured vehicle base		q.s.		
		Approved Color used				
05	Terbutaline, Ambroxol HCl, Guaiphenesin & Menthol Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Terbutaline sulphate	IP	1.25	mg	
		Ambroxol HCl	IP	15	mg	
		Guaiphenesin	IP	50	mg	
		Menthol	IP	2.5	mg	
		In a Sugar free base		q.s.		
Approved Color used						
06	Levocetirizine Dihydrochloride & Ambroxol HCl Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Levocetirizine Dihydrochloride	IP	5	mg	
		Ambroxol HCl	IP	30	mg	
		In a flavoured syrup base		q.s.		
07	Paracetamol Oral Suspension IP	Each 5 ml contains:				Approved on 28.09.2017
		Paracetamol	IP	125	mg	
		In a Flavoured vehicle base		q.s.		
		Approved Color used				
08	Fexofenadine Hydrochloride Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Fexofenadine Hydrochloride	IP	30	mg	
		In a Flavoured vehicle base		q.s.		
		Approved Color used				
09	Hydroxyzine Hydrochloride Oral solution IP	Each 5 ml contains:				Approved on 28.09.2017
		Hydroxyzine Hydrochloride	IP	10	mg	
		Excipients		q.s.		
		In a Flavoured syrupy base				
		Approved Color used				

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION			Reference
-------	---------------------------------------	-------------	--	--	-----------

10	Sucralfate & Oxetacaine Suspension	Each 10 ml contains:			Approved on 28.09.2017	
		Sucralfate	USP	1000		mg
		Oxetacaine	BP	20		mg
		Excipients		q.s		
		In a Flavoured vehicle base				
		Approved colours used				

11	Mefenamic Acid & Paracetamol Suspension	Each 5 ml contains:			Approved on 28.09.2017	
		Mefenamic Acid	IP	50		mg
		Paracetamol	IP	125		mg
		Excipients		q.s.		
		In a Flavoured vehicle base				
		Approved Color used				

12	Cyproheptadine HCl & Tricholine citrate Drops	Each ml contains:			Approved on 28.09.2017	
		Cyproheptadine HCl	IP			
		Eq. to Cyproheptadine HCl Anhydrous		1.5		mg
		Tricholine Citrate		55		mg
		In a flavoured sorbitol (70%) syrup base				q.s.
		Approved Colors used				

13	Ambroxol HCl, Levosalbutamol & Guaiphenesin Syrup	Each 5 ml contains:			Approved on 28.09.2017	
		Ambroxol HCl	IP	30		mg
		Levosalbutamol Sulphate	IP			
		Eq. to Levosalbutamol		1		mg
		Guaiphenesin	IP	50		mg
		In a flavoured sorbitol base				q.s.
		Approved Color used				

14	Paracetamol Phenylephrine Hydrochloride Chlorpheniramine Maleate Sodium Citrate & Menthol syrup	Each 5 ml contains:			Approved on 28.09.2017	
		Paracetamol	IP	125		mg
		Phenylephrine Hydrochloride	IP	5		mg
		Chlorpheniramine Maleate	IP	1		mg
		Sodium Citrate	IP	60		mg
		Menthol	IP	1		mg
		Excipients		q.s		
In a flavoured sorbitol base						

15	Dextromethorphan Hydrobromide, Phenylephrine Hydrochloride, Chlorpheniramine Maleate & Guaiphenesin Syrup (BANNED)	Each 5 ml contains:			Approved on 28.09.2017	
		Dextromethorphan Hydrobromide	IP	10		mg
		Phenylephrine Hydrochloride	IP	5		mg
		Chlorpheniramine Maleate	IP	2		mg
		Guaiphenesin	IP	50		mg
		Excipients		q.s.		
		In a flavoured syrup base				
		Approved Color used				

16	Dextromethorphan Hydrobromide,	Each 5 ml contains:			
		Dextromethorphan Hydrobromide	IP	10	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION			Reference		
	Phenylephrine Hydrochloride & Chlorpheniramine Maleate Syrup	Phenylephrine Hydrochloride	IP	5	mg	Approved on 28.09.2017	
		Chlorpheniramine Maleate	IP	2	mg		
		Excipients			q.s.		
		In a flavoured syrup base					
		Approved Color used					

17	Potassium Citrate, Magnesium Citrate & Vitamin B6 Oral solution	Each 5 ml contains:			Approved on 28.09.2017	
		Potassium Citrate	IP	1100		mg
		Magnesium Citrate	USP	375		mg
		Pyridoxine HCl (Vitamin B6)	IP	20		mg
		Excipients				q.s.
		(Each ml. contains approx.1m.Eq.Magnesium Ion, 2 m.Eq. Potassium Ion, 3 m.Eq. Citrate Ion & 4 mg Pyridoxine HCl)				
		In a flavoured vehicle base				q.s.
Approved Color used						

18	Cetirizine Hydrochloride, Dextromethorphan Hydrobromide & Ambroxol HCl syrup	Each 5 ml contains:			Approved on 28.09.2017	
		Cetirizine Hydrochloride	IP	5		mg
		Dextromethorphan Hydrobromide	IP	10		mg
		Ambroxol Hydrochloride	IP	15		mg
		Excipients				q.s.
		In a flavoured Sugar free base				
Approved Color used						

19	Calcium Citrate, Vitamin D3, L-Lysine HCl, Zinc & Magnesium Hydroxide Oral Suspension	Each 5 ml contains:			Approved on 28.09.2017	
		Calcium Citrate Eq. to Calcium	USP	100		mg
		Vitamin D3	IP	200		IU
		Lysine HCl	USP	50		mg
		Zinc (As Zinc Sulphate IP)		4		mg
		Magnesium Hydroxide	IP	50		mg
		In a flavoured vehicle base				q.s.
Approved Colors used						

20	Magaldrate & Simethicone Suspension USP	Each 5 ml contains:			Approved on 28.09.2017	
		Magaldrate (Anhydrous)	IP	400		mg
		Simethicone	IP	20		mg
		Excipients				q.s.
		In a flavoured vehicle base				
Approved Colors used						

21	Diastase & pepsin syrup	Each 5 ml contains:			Approved on 28.09.2017	
		Diastase (1:1200) (Diastase derived from gillius Oryzae. Digest not less than 60 g of cooked starch)	IP	50		mg
		Pepsin (1:3000)	IP	10		mg

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
		(Digest not less than 30 g of coagulated egg albumin)				
		In a flavoured Syrupy base				

22	Calcium Carbonate, Magnesium Hydroxide, Zinc Gluconate & Vitamin D3 Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Calcium Carbonate	IP	625	mg	
		Magnesium Hydroxide	IP	180	mg	
		Zinc Gluconate	USP	14	mg	
		Vitamin D3	IP	200	IU	
		Appropriate overages of vitamin added				
		In a flavoured vehicle base				
		Approved Colors used "For prophylactic use only"				

23	Dextromethorphan HBr, Phenylephrine HCl, Chlorpheniramine Maleate & Menthol Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Dextromethorphan Hydrobromide	IP	10	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		Menthol	IP	0.5	mg	
		Excipients				
		In a flavoured syrup base				
		Approved Colours used				

24	Chlorpheniramine Maleate, Phenylephrine Hydrochloride & Paracetamol Drops	Each ml contains:				Approved on 28.09.2017
		Chlorpheniramine Maleate	IP	1	mg	
		Phenylephrine Hydrochloride	IP	2.5	mg	
		Paracetamol	IP	125	mg	
		In a flavoured syrupy base				
Approved Colour used						

25	Chlorpheniramine Maleate & Phenylephrine Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Chlorpheniramine Maleate	IP	2	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		In a flavoured syrup base				
Approved Color used						

26	Alpha amylase, Papain, Dill Oil, Anise oil & Caraway Oil Drops	Each 5 ml contains:				Approved on 28.09.2017
		Alpha amylase (1:2000)	IP	20	mg	
		Digest not less than 40 mg of cooked starch)				
		Papain	IP	10	mg	
		Dill Oil	BP	2	mg	
		Anise oil	BP	2	mg	
		Caraway Oil	BP	2	mg	
		In a flavoured syrupy base				
Approved Colours used						

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
27	Calcium Citrate, Vitamin D3, L-Lysine HCl, Zinc & Magnesium Hydroxide Oral Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Calcium Citrate Eq. to Calcium	USP	100	mg	
		Vitamin D3	IP	200	IU	
		Lysine HCl	USP	50	mg	
		Zinc (As Zinc Sulphate IP)		4	mg	
		Magnesium Hydroxide	IP	50	mg	
		In a flavoured vehicle base		q.s.		
Approved Colors used						
28	Levocetirizine & Montelukast Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Levocetirizine Dihydrochloride	IP	2.5	mg	
		Montelukast sodium Eq. to Montelukast	IP	4	mg	
		In a flavoured syrup base		q.s.		
29	Terbutaline Sulphate, Guaiphenesin, Bromhexine HCl & Menthol Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.25	mg	
		Guaiphenesin	IP	50	mg	
		Bromhexine HCl	IP	4	mg	
		Menthol	IP	2.5	mg	
		Excipients		q.s		
		In a flavoured syrupy base				
Approved Color used						
30	Aceclofenac Paracetamol Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Aceclofenac	IP	50	mg	
		Paracetamol	IP	250	mg	
		Excipients		q.s		
In a flavoured syrup base						
31	Dextromethorphan HBr, Phenylephrine HCl, Chlorpheniramine Maleate Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Dextromethorphan Hydrobromide	IP	10	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		Excipients		q.s		
		In a flavoured syrup base				
Approved Colours used						
32	Simethicone, Dill Oil & Fennel Oil Drops	Each ml contains:				Approved on 28.09.2017
		Simethicone	IP	40	mg	
		Dill Oil	BP	0.005	ml	
		Fennel Oil	BP	0.0007	ml	
		Excipients		q.s		
In a flavoured syrup						
33	Ondansetron Oral solution IP	Each 5 ml contains:				Approved on 28.09.2017
		Ondansetron HCl Eq. to Ondansetron	IP	2	mg	
		In a flavoured vehicle base		.		

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOEIAL TITLE	COMPOSITION				Reference
		Approved Color used				
34	Paracetamol Phenylephrine Chlorpheniramine Maleate Sodium Citrate & Menthol syrup	Each 5 ml contains:				Approved on 28.09.2017
		Paracetamol	IP	125	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	0.5	mg	
		Sodium Citrate (Anhydrous)	IP	60	mg	
		Menthol	IP	1	mg	
		Approved Color used				
35	Azithromycin Oral Suspension IP	Each 5 ml contains:				Approved on 28.09.2017
		Azithromycin Dihydrate	IP			
		Eq. to Azithromycin Anhydrous		200	mg	
		In a flavoured vehicle base		q.s.		
		Approved Color used				
36	Sodium Picosulphate Oral solution BP	Each 5 ml contains:				Approved on 28.09.2017
		Sodium Picosulphate	BP	5	mg	
		In a palatable sorbitol base		.		
		Approved Color used				
37	Levetiracetam Oral solution	Each 5 ml contains:				Approved on 28.09.2017
		Levetiracetam	BP	100	mg	
		Methylparaben (As Preservative)	IP	1.35	mg	
		Propylparaben (As Preservative)	IP	0.15	mg	
		In a flavoured vehicle base		q.s.		
		Approved Color used				
38	Sodium Feredetate,Folic Acid & Vitamin B12 Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Sodium Feredetate	BP	231	mg	
		Folic Acid	IP	1.5	mg	
		Vitamin B12	IP	15	mcg	
		In a flavoured syrupy base		q.s.		
		Approved Color used				
Appropriate overage of vitamin added						
"For therapeutic use only"						
39	Magaldrate & Simethicone Oral Suspension USP	Each 5 ml contains:				Approved on 28.09.2017
		Magaldrate (Anhydrous)	IP	480	mg	
		Simethicone	IP	20	mg	
		Excipients		q.s		
		In a flavoured vehicle base				
40	Domperidone Suspension IP	Each ml contains:				Approved on 28.09.2017
		Domperidone	IP	1	mg	
		Excipients		q.s.		
		In a flavoured vehicle base				
41	Cholecalciferol Solution USP	Each ml contains:				Approved on 28.09.2017
		Cholecalciferol (As Stabilized)	IP	400	IU	
		In a flavoured vehicle base		q.s.		

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION				Reference
42	Oxetacaine, Aluminium Hydroxide & Magnesium Hydroxide Gel	Each 5 ml contains:				Approved on 28.09.2017
		Oxetacaine	BP	10	mg	
		Aluminium Hydroxide	IP	291	mg	
		Magnesium Hydroxide	IP	98	mg	
		Excipients		q.s		
		In a flavoured vehicle base				
43	Chlorpheniramine Maleate, Dextromethorphan Hydrobromide & Phenylephrine Hydrochloride Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Chlorpheniramine Maleate	IP	2	mg	
		Dextromethorphan Hydrobromide	IP	15	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Excipients		q.s.		
		In a flavoured vehicle base				
44	Sodium Citrate, Chlorpheniramine Maleate, Phenylephrine Hydrochloride & Paracetamol syrup	Each 5 ml contains:				Approved on 28.09.2017
		Sodium Citrate	IP	120	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Paracetamol	IP	250	mg	
		Excipients q.s. In a flavoured vehicle base				
45	Mefenamic & Paracetamol Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Mefenamic	IP	100	mg	
		Paracetamol	IP	250	mg	
		Excipients		q.s		
		In a flavored syrup base				
46	Carbonyl Iron, Folic Acid, Zinc & Vitamin B12 Syrup	Each 5 ml contains:				Approved on 28.09.2017
		Carbonyl Iron Eq. to elemental iron		50	mg	
		Folic Acid	IP	500	mcg	
		Zinc Sulphate Monohydrate	IP	11	mg	
		Vitamin B12	IP	6	mcg	
		Excipients q.s.				
47	Deflazacort Suspension	Each 5 ml contains:				Approved on 28.09.2017
		Deflazacort		6	mg	
		In a flavoured vehicle base			q.s.	
48	Cholecalciferol Solution USP	Each ml contains:				Approved on 28.09.2017
		Cholecalciferol (As Stabilized)	IP	800	IU	
		excipients		q.s.		
		In a flavoured vehicle base				
		Approved Color used				
49	Hydroxyzine Hydrochloride Oral solution IP	Each ml contains:				Approved on 28.09.2017
		Hydroxyzine Hydrochloride	IP	6	mg	
		Excipients		q.s.		
		In a Flavoured syrupy base				
		Approved Color used				

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMA COEIAL TITLE	COMPOSITION			Reference
50	Piracetam Syrup	<b>Each 5 ml contains:</b>			Approved on 28.09.2017
		Piracetam	IP	500 mg	
		In a Flavoured syrupy base		q.s.	
		Approved Color used			
51	Simethicone , Magnesium Hydroxide, Dried Aluminium Hydroxide gel, Carboxy methylcellulose sodium oral suspension USP	<b>Each 10 ml contains:</b>			Approved on 28.09.2017
		Simethicone Emulsion Eq. to Simethicone	USP	50 mg	
		Magnesium Hydroxide (As Paste)	IP	185 mg	
		Dried Aluminium Hydroxide gel (As paste)	IP	830 mg	
		Carboxy methyl cellulose sodium	IP	100 mg	
		In a flavoured sorbitol base		q.s.	
Approved Color used					
52	Sucralfate Suspension	<b>Each 10 ml contains:</b>			Approved on 28.09.2017
		Sucralfate	USP	1000 mg	
		In a Flavoured vehicle base		q.s.	
		Approved Color used			
53	Azithromycin Oral Suspension IP	<b>Each 5 ml contains:</b>			Approved on 28.09.2017
		Azithromycin Dihydrate Eq. to Azithromycin Anhydrous	IP	100 mg	
		In a flavoured vehicle base		q.s.	
		Approved Color used			
54	Ambroxol HCl, Guaiphenesin & Terbutaline Sulphate Syrup	<b>Each 10 ml contains:</b>			Approved on 28.09.2017
		Ambroxol HCl	IP	30 mg	
		Guaiphenesin	IP	100 mg	
		Terbutaline	IP	2.5 mg	
		excipients		q.s.	
Approved Color used					
55	Paracetamol, Phenylephrine Hydrochloride & Chlorpheniramine Maleate syrup	<b>Each 5 ml contains:</b>			Approved on 28.09.2017
		Paracetamol	IP	250 mg	
		Phenylephrine Hydrochloride	IP	5 mg	
		Chlorpheniramine Maleate	IP	2 mg	
		In a flavoured syrupy base		q.s.	
Approved Color used					
56	Mefenamic Acid Suspension	<b>Each 5 ml contains:</b>			Approved on 28.09.2017
		Mefenamic Acid	IP	100 mg	
		In a Flavoured vehicle base		q.s.	
		Approved Color used			
57	Terbutaline, Ambroxol HCl, Guaiphenesin & Menthol Syrup	<b>Each 5 ml contains:</b>			Approved on 28.09.2017
		Terbutaline	IP	1.25 mg	
		Ambroxol HCl	IP	30 mg	
		Guaiphenesin	IP	50 mg	
		Menthol	IP	2.5 mg	
In a Flavoured vehicle base					



**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
		Approved Color used				
58	Terbutaline Sulphate, Ambroxol HCl, Guaiphenesin & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.25	mg	
		Ambroxol HCl	IP	30	mg	
		Guaiphenesin	IP	50	mg	
		Menthol	IP	0.5	mg	
		In a Flavoured vehicle base		q.s.		
		Approved Color used				
59	Dextromethorphan Hydrobromide, Cetirizine, Phenylephrine Hydrochloride & Guaiphenesin Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Dextromethorphan Hydrobromide	IP	10	mg	
		Cetirizine HCl	IP	5	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Guaiphenesin	IP	50	mg	
		In a flavoured sugar free base		q.s.		
		Approved Color used				
60	Dextromethorphan Hydrobromide, Chlorpheniramine Maleate Phenylephrine Hydrochloride & Guaiphenesin Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Dextromethorphan Hydrobromide	IP	10	mg	
		Chlorpheniramine Maleate	IP	4	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Guaiphenesin	IP	100	mg	
		In a flavoured sorbitol base		q.s.		
		Approved Color used				
61	Dried Aluminium Hydroxide, Magnesium Hydroxide, Simethicone & Oxetacaine Suspension	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Dried Aluminium Hydroxide	IP	600	mg	
		Magnesium Hydroxide	IP	300	mg	
		Simethicone	IP	25	mg	
		Oxetacaine	BP	10	mg	
		In a flavoured sorbitol base		q.s.		
		Approved Color used				
62	Alumina, Magnesia & Simethicone Oral Suspension USP	<b>Each 10 ml contains:</b>				Approved on 28.09.2017
		Dried Aluminium Hydroxide	IP	250	mg	
		Magnesium Hydroxide	IP	250	mg	
		Simethicone	IP	25	mg	
		In a flavoured sorbitol base		q.s.		
		Approved Color used				
63	Metronidazole Benzoate Oral Suspension IP	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Metronidazole Benzoate	IP			
		Eq. to Metronidazole		200	mg	
		In an aqueous vehicle base		q.s.		
		Approved Color used				
64	Furazolidone Oral Suspension IP	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Furazolidone	IP	25	mg	
		In an aqueous vehicle base		q.s.		
		Approved Color used				

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
65	Levofloxacin Oral Suspension IP	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Levofloxacin Hemihydrate	IP	125	mg	
		Eq. to Levofloxacin				
		In an aqueous vehicle base				
		excipients		q.s.		
		Approved Color used				
66	Paracetamol, Phenylephrine Hydrochloride, Chlorpheniramine Maleate & Sodium Citrate syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Paracetamol	IP	250	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		Sodium Citrate	IP	60	mg	
		excipients	q.s.			
		In a flavoured suspensionl base				
		Approved Color used				
67	Cyproheptadine HCl & Tricholine citrate Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Cyproheptadine HCl	IP	2	mg	
		Eq. to Cyproheptadine HCl Anhydrous				
		Tricholine Citrate		275	mg	
		In a flavoured sorbitol base (70%)_syrup base				
		Approved Color used				
68	Ofloxacin Oral Suspension IP	<b>Each 10 ml contains:</b>				Approved on 28.09.2017
		Ofloxacin	IP	100	mg	
		excipients		q.s.		
		In a Flavoured vehicle base		q.s.		
				Approved Color used		
69	Loratadine Oral suspension USP	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Loratadine	USP	5	mg	
		excipients		q.s.		
		In a Flavoured syrupy base		q.s.		
				Approved Color used		
70	Dried Aluminium Hydroxide, Magnesium Hydroxide & Simethicone Oral Suspension	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Dried Aluminium Hydroxide (Added as Aluminium Hydroxide paste )	IP	250	mg	
		Magnesium Hydroxide (Added as Magnesium Hydroxide paste )	IP	250	mg	
		Simethicone	IP	50	mg	
		Sorbitol Solution 70%(Non-Crystallising)	IP	1.25	mg	
		Approved Color used				
71	Albendazole & Ivermectin Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Albendazole	IP	200	mg	
		Ivermectin	IP	3	mg	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
		In a Flavoured syrupy base		q.s.		
		Approved Color used				
72	Ferrous Bisglycinate, Folic Acid, Cyanocobalamin & Zinc Sulphate Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Ferrous Bisglycinate		50	mg	
		Eq. to elemental iron				
		Folic Acid	IP	0.5	mg	
		Cyanocobalamin	IP	2.5	mcg	
		Zinc Sulphate	IP	7.0	mg	
		In a flavoured Syrupy base		q.s.		
		Approved Color used				
73	Chlorpheniramine Maleate, Phenylephrine Hydrochloride & Paracetamol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Chlorpheniramine Maleate	IP	1	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Paracetamol	IP	125	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
74	Terbutaline Sulphate, Bromhexine HCl & Guaiphenesin Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.25	mg	
		Bromhexine HCl	IP	2	mg	
		Guaiphenesin	IP	50	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
75	Calcium Carbonate & Vitamin D3 Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Calcium Carbonate	IP	250	mg	
		Vitamin D3	IP	200	IU	
		In a flavoured syrupy base		q.s.		
		Appropriate overages of vitamins added				
		Approved Color used				
		"For prophylactic use only"				
76	Chlorpheniramine Maleate, Dextromethorphan & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Chlorpheniramine Maleate	IP	4	mg	
		Dextromethorphan Hydrobromide	IP	10	mg	
		Menthol	IP	0.1	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
77	Dextromethorphan HBr, Phenylephrine HCl, Chlorpheniramine Maleate Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Dextromethorphan Hydrobromide	IP	10	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		Excipients		q.s.		
		In a flavoured syrup base				
		Approved Colours used				
78	Lycopene with Multivitamin & Multimineral Syrup	<b>Each 10 ml contains:</b>				Approved on
		Lycopene (6%)	USP	2000	mcg	
		Lysine Hydrochloride	USP	10	mg	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION				Reference
		Thiamine Hydrochloride	IP	2	mg	28.09.2017
		Riboflavin Sodium Phospahte	IP	2	mg	
		Pyridoxine HCl	IP	1	mg	
		Vitamin A Palmitate	IP	2500	IU	
		Alpha Tocopheryl Acetate	IP	5	IU	
		Ascorbic Acid	IP	40	mg	
		Zinc Sulphate Monohydrate	IP	2	mg	
		Potassium Iodide	IP	30	mcg	
		D-Panthenol	IP	3	mg	
		Cholecalciferol	IP	150	IU	
		Selenium (As Sodium Selenite BP)		20	mcg	
		Folic Acid	IP	150	mcg	
		Chromium Chloride Hexahydrate	USP	16	mcg	
		Manganese Chloride	USP	16	mcg	
		Niacinamide	IP	10	mg	
		Ferrous Gluconate	IP	40	mg	
		In a flavoured syrup base		q.s.		
		Appropriate overages of vitamins added				
		Approved Color used				
		"For prophylactic use only"				
79	Lycopene with Multivitamin & Multimineral Syrup	<b>Each 15 ml contains:</b>				Approved on 28.09.2017
		Lycopene (10%)	USP	1000	mcg	
		Folic Acid	IP	1.5	mg	
		Zinc Sulphate Monohydrate	IP			
		Eq. to elemental Zinc		3	mg	
		Vitamin B1	IP	2	mg	
		Vitamin B2	IP	3	mg	
		Vitamin B6	IP	1.5	mg	
		Manganese Chloride	USP	10	mcg	
		Sorbitol Solution (70%) Non-Crystallising	IP	1	gm	
		In a flavoured syrup base		q.s.		
		Appropriate overages of vitamins added				
		Approved Color used				
		"For prophylactic use only"				
80	Cetirizine Di-hydrochloride, Phenylephrine hydrochloride & Paracetamol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Cetirizine Di-hydrochloride	IP	2.5	mg	
		Phenylephrine hydrochloride	IP	2.5	mg	
		Paracetamol	IP	125	mg	
		In a flavoured Syrupy base		q.s.		
		Approved Color used				
81	Terbutaline Sulphate, Bromhexine HCl, Guaiphenesin & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.25	mg	
		Bromhexine HCl	IP	4	mg	
		Guaiphenesin	IP	50	mg	
		Menthol	IP	1	mg	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
		In a flavoured syrup base		q.s.		
		Approved Color used				
82	Iron, Vitamin B12, Folic Acid & Lysine Hydrochloride syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Carbonyl Iron Eq. to elemental iron		100	mg	
		Vitamin B12	IP	5	mcg	
		Folic Acid	IP	500	mcg	
		Lysine Hydrochloride	USP	100	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
		Childern above 1 year of age Dose: One teaspoon once a day				
83	Iron, Vitamin B12, Folic Acid & Lysine Hydrochloride Drops	<b>Each ml contains:</b>				Approved on 28.09.2017
		Carbonyl Iron Eq. to elemental iron		20	mg	
		Vitamin B12	IP	1	mcg	
		Folic Acid	IP	100	mcg	
		Lysine Hydrochloride	USP	20	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
		For infants below 1 year of age Dose: One teaspoon once a day				
84	Terbutaline Sulphate, Guaiphenesin, Ambroxol HCl, Cetirizine Dihydrochloride & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.25	mg	
		Guaiphenesin	IP	50	mg	
		Ambroxol HCl	IP	15	mg	
		Cetirizine Dihydrochloride	IP	2.5	mg	
		Menthol	IP	1.5	mg	
		excipients		q.s.		
		In a flavoured syrup base Approved Color used				
85	Chlorpheniramine Maleate, Phenylephrine HCl, Zinc Gluconate, Dextromethorphan & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Dextromethorphan Hydrobromide	IP	10	mg	
		Chlorpheniramine Maleate	IP	2	mg	
		Phenylephrine HCl	IP	5	mg	
		Zinc Gluconate Eq. to elemental Zinc	USP	7.5	mg	
		Menthol	IP	1	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
86	Terbutaline Sulphate, Guaiphenesin, Bromhexine HCl & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.2	mg	
		Guaiphenesin	IP	50	mg	
		Bromhexine HCl	IP	4	mg	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION				Reference
		Menthol	IP	1.5	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
87	Terbutaline Sulphate, Ammonium Chloride, Bromhexine HCl & Menthol Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Terbutaline Sulphate	IP	1.25	mg	
		Ammonium Chloride	IP	50	mg	
		Bromhexine HCl	IP	4	mg	
		Menthol	IP	2	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
88	Chlorpheniramine Maleate, Dextromethorphan HBr, Phenylephrine Hydrochloride & Menthol Syrup  (BANNED)	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Chlorpheniramine Maleate	IP	2	mg	
		Dextromethorphan Hydrobromide	IP	10	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Menthol	IP	0.5	mg	
		In a flavoured syrup base		q.s.		
		Approved Color used				
89	Piracetam Syrup	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Piracetam	IP	500	mg	
		In a Flavoured syrupy base		q.s.		
		Approved Color used				
90	Cetirizine HCl, Phenylephrine Hydrochloride, Dextromethorphan Hydrobromide & Menthol Syrup (BANNED)	<b>Each 5 ml contains:</b>				Approved on 28.09.2017
		Cetirizine HCl	IP	5	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Dextromethorphan Hydrobromide	IP	10	mg	
		Menthol	IP	1.5	mg	
		In a flavoured sugar free base		q.s.		
		Approved Color used				
91	Vitamin A,Vitamin E,Vitamin C, Vitamin D3, Vitamin B1, Vitamin B2, Vitamin B6, Vitamin B12, Selenium, Calcium D-Pantothenate, D-Panthenol, Zinc , Chromium ,Manganese Molybdenum & Iodine Syrup	<b>Each 15 ml contains:</b>				Approved on 28.09.2017
		Vitamin A (as palmitate)	IP	2500	IU	
		Vitamin E (as acetate)	IP	7.5	IU	
		Vitamin C	IP	40	mg	
		Vitamin D3	IP	200	IU	
		Vitamin B1	IP	2.25	mg	
		Vitamin B2	IP	2.5	mg	
		Vitamin B6	IP	1.0	mg	
		Vitamin B12	IP	2.5	mcg	
		Selenium (Selenium Dioxide USP)		10	mcg	
		Calcium D-Pantothenate	IP	2.5	Mg	
		D-Panthenol	IP	1.25	Mg	
		Zinc (As Zinc Gluconate USP)		1	mg	
		Chromium (As Chromium Chloride Hexahydrate USP)		0.8	mcg	
		Manganese (As Manganese ChlorideUSP)		0.8	mcg	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION			Reference	
		Molybdenum (As Sodium Molybdate IP)		0.8	mcg	
		Iodine (As Potassium Iodide IP)		50	mcg	
		excipients		q.s.		
		In a flavoured syrup base				
		Appropriate overages of vitamins added				
		Approved Color used				
		“For therapeutic use only”				
		Dose: one table spoonful twice daily or as directed by the physician				

92	Vitamin A,Thiamine Hydrochloride,Riboflavin, Pyridoxine HCl,Vitamin B12, Vitamin C,Vitamin D3, Vitamin E acetate, Niacinamide, D-Panthenol, Biotin,Zinc ,Iodine , Chromium & Molybdenum Syrup	<b>Each 15 ml contains:</b>				Approved on 28.09.2017
		Vitamin A Palmitate	IP	2500	IU	
		Thiamine Hydrochloride	IP	1.5	mg	
		Riboflavin Sodium Phosphate Eq. to Riboflavin	IP	1.7	mg	
		Pyridoxine HCl	IP	1.5	mg	
		Vitamin B12	IP	1	mcg	
		Vitamin C	IP	50	mg	
		Vitamin D3	IP	200	IU	
		Vitamin E acetate	IP	10	IU	
		Niacinamide		20	mg	
		D-Panthenol	IP	5	mg	
		Biotin	USP	30	mcg	
		Zinc (As Zinc Gluconate USP)		3	mg	
		Iodine (As Potassium Iodide IP)		150	mcg	
		Manganese (As Manganese Chloride USP)		2.5	mg	
		Chromium (As Chromium Chloride Hexahydrate USP)		25	mcg	
		Molybdenum (As Sodium Molybdate IP)		25	mcg	
		Sorbitol Solution (70%) Non-Crystallising	IP	7.5	gm	
excipients		q.s.				
In a flavoured syrup base						
Approved Color used						

93	Lycopene (6%),Vitamin A , Vitamin E ,Vitamin C Selenium,Zinc ,Manganese, Iodine, Copper, Vitamin B1, Vitamin B2 & Vitamin B6 Syrup	<b>Each 10 ml contains:</b>				Approved on 28.09.2017
		Lycopene (6%)	USP	1000	mcg	
		Vitamin A (as acetate)	IP	2500	IU	
		Vitamin E (as acetate)	IP	10	IU	
		Vitamin C	IP	50	mg	
		Selenium (Sodium Selenite USP)		35	mcg	
		Zinc (As Zinc Gluconate USP)		3	mg	
		Manganese (As Manganese Gluconate USP)		2	mg	
		Iodine (As Potassium Iodide IP)		100	mcg	
		Copper (As Cupric Sulphate USP)		500	mcg	
		Vitamin B1	IP	2	mg	
		Vitamin B2	IP	3	mg	
		Vitamin B6	IP	1.5	mg	
		In a flavoured syrup base				

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPEIAL TITLE	COMPOSITION				Reference
		Appropriate overages of vitamins added				
		Approved Color used				
		"For prophylactic use only"				
94	Terbutaline, Guaiphenesin ,Ambroxol HCl & Menthol Syrup	<b>Each 5 ml contains:</b> Terbutaline IP 1.25 mg Guaiphenesin IP 50 mg Ambroxol HCl IP 15 mg Menthol IP 1 mg In a flavoured syrup base q.s. Approved Color used				Approved on 28.09.2017
95	Protein Hydrosylate, Ferric Ammonium Citrate, Vitamin B1, Vitamin B6,Vitamin B12,Folic Acid,vit.B2Niacinamide, D-Panthenol, Calcium Gluconate & Sorbitol solution Syrup	<b>Each 15 ml contains:</b> Protein Hydrosylate IP 1 gm Ferric Ammonium Citrate IP 200 mg Vitamin B1 IP 2.5 mg Vitamin B2 IP 2.5 mg Vitamin B6 IP 1.5 mg Vitamin B12 IP 2.5 mg Folic Acid IP 0.5 Mg Niacinamide IP 25 mg D-Panthenol IP 5 mg Calcium Gluconate IP 0.5 mg Sorbitol Solution (70%) Non-Crystallising IP 0.5 gm excipients q.s. In a flavoured syrup base				Approved on 28.09.2017
96	Terbutaline Sulphate, Guaiphenesin,Bromhexine HCl & Menthol Syrup	<b>Each 5 ml contains:</b> Terbutaline Sulphate IP 1.25 mg Guaiphenesin IP 50 mg Bromhexine HCl IP 8 mg Menthol IP 1 mg excipients q.s. In a flavoured syrup base Approved Color used				Approved on 28.09.2017
97	Ofloxacin Suspension IP 50 mg	<b>Each 5 ml contains:</b> Ofloxacin IP 50 mg In a flavoured vehicle base q.s. Approved color used				28.09.2017
98	Ofloxacin, Metronidazole & Simethicone Oral Suspension	<b>Each 5 ml contains:</b> Ofloxacin IP 50 mg Metronidazole Benzoate Eq. to Metronidazole 120 mg Simethicone (Added as emulsion form) IP 10 mg In a flavoured palatable base Approved color used				Approved on 25.05.2017



**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMA COEIAL TITLE	COMPOSITION				Reference
99	Vitamin D Oral Solution 60,000 IU	<b>Each 5 ml contains:</b>				Approved on 25.05.2017
		Cholecalciferol (In nano droplet form)	IP	60000	IU	
		In a flavoured sugar free base		q.s		
		Approved color used				
100	Divalproex Sodium Oral solution	<b>Each 5 ml contains:</b>				Approved on 25.05.2017
		Divalproex Sodium Eq. to Valproic acid	IP	250	mg	
		Flavoured syrupy base		q.s		
		Approved colour used				
101	Cholecalciferol (Vitamin D3) Solution USP	<b>Each 5 ml contains:</b>				Approved on 07.09.2017
		Cholecalciferol (As stabilized ) Vitamin D3	IP	1000	IU	
		Excipients		q.s		
		In a flavoured vehicle base				
102	Cyproheptadine Hcl Syrup IP	<b>Each 5 ml contains:</b>				Approved on 28.10.2017
		Cyproheptadine Hydrochloride Eq. to cyproheptadine Hcl anhydrous	IP	2	mg	
		In a flavoured base		q.s		
103	Divalproex Sodium Oral solution	<b>Each 5 ml contains:</b>				Approved on 21.04.18
		Divalproex Sodium Eq. to Valproic Acid	IP	500	mg	
		In a flavoured sugar free base		q.s		
104	Paracetamol , Phenylephrine Hydrochloride & Chlorpheniramine Maleate Syrup	<b>Each 5 ml contains:</b>				Approved on 10.12.18
		Paracetamol	IP	125	mg	
		Phenylephrine Hydrochloride	IP	5	mg	
		Chlorpheniramine Maleate	IP	2	mg	
105	Potassium Citrate & Magnesium Citrate Oral Solution	<b>Each 5 ml contains:</b>				Approved on 10.12.18
		Potassium Citrate	IP	1100	mg	
		Magnesium Citrate	USP	375	mg	
		In a flavoured sorbitol (70%) non crystallizing IP base				
106	Paracetamol, Phenylephrine Hcl & Chlorpheniramine Maleate Syrup	<b>Each 5 ml contains:</b>				Approved on 10.12.18
		Phenylephrine HCl	IP	2.5	mg	
		Chlorpheniramine Maleate	IP	1	mg	
		Paracetamol	IP	125	mg	
		Excipients		q.s		
In a flavoured vehicle syrupy base						
	Iron, Folic Acid & Vitamin	<b>Each 5 ml contains:</b>				
		Ferric Ammonium Citrate	IP	110	mg	

**LIST OF ADDITIONAL PRODUCTS PERMITTED TO :**

<b>GNOSIS PHARMACEUTICALS PVT.LTD.</b> Vill.-Moginand, Nahan road, Kala-Amb, Distt. Sirmour (H.P.) 173030			Issue Date	
			Total Additional Products	
License No.	25/25A: MNB/07/541	28/28A: MB/07/542	Valid up to	27.04.2022

S.NO.	GENERIC NAME/ PHARMACOPOEIAL TITLE	COMPOSITION				Reference
107	B12 Syrup	Eq. to elemental Iron 22.5 mg				Approved on 10.12.18
		Foile Acid	IP	1.5	mg	
		Vitamin B12	IP	15	mcg	
		Sorbitaol Solutuin (70%) (Non Crystallising )	IP	10%	w/v	
		Flavoured syrup base		q.s.		
108	Chlorhexidine Mouthwash IP	Each 5 ml reconstituted suspension contains:				11.02.2019
		Chlorhexidine Gluconate solution	IP			
		Eq. to Chlorhexidine Gluconate		0.2%	w/v	
		Excipients		q.s.		
109	Ambroxol HCl Syrup 30mg	Each 5 ml contains:				11.02.2019
		Ambroxol HCl	IP	30	mg	
		Excipients		q.s.		
		In a flavoured syrup base				
110	Paracetamol Paediatric Oral Suspension IP	Each ml contains:				11.02.2019
		Paracetamol	IP	100	mg	
		In a Flavoured vehicle base				
		Approved Color used				
111	Salbutamol Sulphate Syrup IP 2 mg	Each 5 ml contains:				11.02.2019
		Salbutamol Sulphate	IP			
		Eq. to Salbutamol		2	mg	
		In a flavoured syrupy base				
112	Aluminium, Magnesium & Simethicone Oral Suspension IP	<b>Each 10 ml contains:</b>				Approved on 11.02.2019
		Dried Aluminium Hydroxide	IP	250	mg	
		Magnesium Hydroxide	IP	250	mg	
		Simethicone	IP	25	mg	
		In a flavoured sorbitol base				
Approved Color used						
113	Lactulose Solution USP	Each 5 ml contains:				11.02.2019
		Lactulose	IP	3.335	g	
		In a flavoured sorbitol base				
		Approved colour used				