GENNEX LABORATORIES LIMITED
you can trust

Ref: GLL/BSE/2025-26/Jul-

Date: July 28, 2025

To
The General Manager,
Corporate Relations Department,
BSE Limited
Phiroze Jeejeebhoy Towers,
Dalal Street, **Mumbai – 400001**.
Maharashtra State, India.
Script Code: 531739

To
The Listing Manager,
The Ahmedabad Stock Exchange Limited
A-2, Kamdhenu Complex, Opp. Sahajanand College, 120 Feet Ring Road, Panjara Pol, Ambawadi, **Ahmedabad - 380015**.
Gujarat State, India.
Script Code:

To
The Calcutta Stock Exchange Limited,
#7, Lyons Range, Murgighata,
Dalhousie, **Kolkata - 700001**,
West Bengal State, India.
Scrip Code: 26178

Dear Sir/Madam,

Sub: Approval of Products from Drug Control Administration, Government of Andhra Pradesh for our Subsidiary Company, Deccan Remedies Limited;
Ref: BSE Security ID: GENNEX, Script Code: 531739

We are pleased to inform that our Subsidiary Company, M/s Deccan Remedies Limited, has received Approval from the Drug Control Administration, Government of Andhra Pradesh granting Licence for manufacture of Products under Form 25, Drug Licence No.31/VP/AP/2014/B/R/CC, valid upto 28.05.2028.

List of Products approved for manufacture and sale (Domestic and Export purpose) details as follows:

S.No.	Product Name	Pharmacopoeias
1	Pantoprazole Sodium	IP/USP
2	Pregabalin – IP	IP
3	Pantoprazole Sodium Seaquihydrate	Ph.Eur
4	Allopurinol – IP/USP/ Ph.Eur	IP/USP/Ph.Eur
5	Clopidogrel bisulphate	USP/IP
6	Escitalopram Oxalate	USP/IP
7	Fexofenadine Hydrochloride	USP/Ph.Eur/JP/IP
8	Irbesartan	USP/IP
9	Lamivudine	USP/Ph.Eur/IP
10	Olmesartan Medoxomile	USP/Ph.Eur/IHS
11	Rabeprazole Sodium	USP/JP/IP
12	Sertraline Hydrochloride	USP/Ph.Eur/IP
13	Telmisartan	USP/Ph.Eur/IP
14	Tenofovir Disoproxil Fumarate	USP/Ph.Eur/IP
15	Guaifenesin	IP/h.Eur/USP/BP
16	Methocarbamol	USP/IP
17	Phenazopyridine Hydrochloride	USP
18	Fluconazole	IP/BP/USP/EP
19	Minoxidil	EP/IP/USP

Dinesh Kumar Kejriwal
Digitally signed by Dinesh Kumar Kejriwal
Date: 2025.07.28 13:51:52 +05'30'

Gennex Laboratories Limited
Office: 'Akash Ganga' 3rd Floor, Plot NO.144, Srinagar colony, Hyderabad-500073, T.S. India | Phone: +91-40-67334400 (30 Lines), Fax: +91-40-67334433
Factory: Sy.No.133, IDA Bollaram, Jinnaram Mandal, Sangareddy Dist – 502325, Telangana, India | Tel: +91-08458 279406, Telefax: +91-08454 279516
Info@gennexlab.com, www.gennexlab.com ▌CIN :L L24230TG1990PLC011168

20	Ketoconazole	IP/BP/USP/EP
21	Flupentixol Hydrochloride	BP/IHS
22	Melitracen Hydrochloride	IHS
23	Domperidone Maleate	IP/EP
24	Etoricoxib	IP
25	Nitazoxanide	IHS
26	Voriconazole	IP/EP/BP/USP
27	Tapentadol Hydrochloride	Ph.Eur
28	Acyclovir	IP/USP
29	Triclabendazole	IHS
30	Ezetimibe	IP/USP
31	Carvedilol	IP/USP/Ph.Eur
32	Oxcarbazepine	Ph.Eur/USP
33	Atorvastatin Calcium	USP/IP
34	Mesna	BP/IP/Ph.Eur/USP
35	Dolutegravir Sodium	IP
36	Lovocarnitine	Ph.Eur/USP
37	Montelukast Sodium	USP/Ph.Eur/IP
38	Moxifloxacin Hydrochloride	IP/Ph.Eur/USP
39	Apixaban	IHS
40	Carisoprodol	IP/USP
41	Gliclazide	IP/Ph.Eur
42	Posaconazole	IHS

Drug Licence Approval received from the DCA- Government of Andhra Pradesh for the above Products is enclosed with this letter for your information and for taking the same on record.

Thanking you,

Yours faithfully
For Gennex Laboratories Limited

Dinesh Kumar Kejriwal
Digitally signed by Dinesh Kumar Kejriwal
Date: 2025.07.28 13:52:12 +05'30'

Dinesh Kumar Kejriwal
Company Secretary & Compliance Officer

Encl: a/a.

ORIGINAL

GOVERNMENT OF ANDHRA PRADSH
DRUGSCONTROLADMINISTRATION

Change of Constitution
w.e.f. 29-05-2023

FORM-25

[SeeRule70]

License to manufacture for sale(or for distribution) of drugs other than those specified In Schedules C,C(1) and X)

Number of license and date of issue **31/VP/AP/2014/B/R/CC Dt.14/11/2014**

1. M/s. **DECCAN REMEDIES LIMITED**
is hereby licensed to manufacture the following categories of drugs being drugs other than those specified in [Schedules, C, C(1) and X] to the Drugs and Cosmetics Rules,1945, on the premises situated at **Plot No:58A, Road No. :01, J.N Pharma City, Parawada, Anakapalli - 531021, ANDHRA PRADESH, INDIA** under the direction and supervision of the following competent technical staff.

(a) Competent Technical Staff (Names):Over Leaf

(b) Names of drugs (Each Item to be separately specified): LIST ENCLOSED

2. The license authorities the sale by way of wholesale dealing and storage for sale by the licensee of the drugs manufactured under the license, subject to the conditions applicable to license for sale.

3. The license shall be in force from **29-05-2023 TO 28-05-2028**

4. The license is subject to the conditions stated below and to such other conditions as may be specified in the Rules for the time being in force under the Drugs and Cosmetics Act, 1940.

Signature
Designation
(Licensing Authority)

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2023.06.01 14:26:17 +05'30'

CONDITIONS OF LICENCE

1. This license and any certificate of renewal in force shall be kept on the approved premises and shall be produced at the request of an Inspector appointed under the Drugs and Cosmetics Act,1940.

2. Any change in the [competent technical staff] named in the license shall be forthwith reported to the Licensing Authority.

3. If the licensee wants to manufacture for sale additional items of drugs not included above, he should apply to the Licensing Authority for the necessary endorsement as provided in Rule 69 (5). This license will be deemed to extend to the categories soendorsed.

4. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the license. Where any change in the constitution of the firm takes place, the current license shall be deemed to be valid for A maximum period of three months from the date on which the change takes place unless, in the meantime, afresh license has been taken from the Licensing Authority in the name of the firm with the changed constitution.

(a) COMPETENT TECHNICAL STAFF(NAMES)

1.Mr. T. Murali Krishna	B.Sc	MANUFACTURING CHEMIST
2.Mr. K.V.M.Phani Kumar	B.Sc	MANUFACTURING CHEMIST
3.Mrs. Punem Shilpa	M.Pahrmacy	ANALYTICAL CHEMIST
4.Mr. Srigisetty Prasad	M.Sc	ANALYTICAL CHEMIST

CONSTITUTION OF THE FIRM (LIMITED)

1) Mr. M. Khaleequr Rahaman	–	Managing Director
2) Mr. Arihant Baid	–	Director
3) Mr. Gopalakrishnan T M	–	Director
4) Mr. Laxmipat Baid	–	Director
5) Mr. Harrsha Addala	–	Director

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2023.06.01 14:26:32 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL ADMINISTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From
M.B.R. Prasad, M.Pharm, M.Phil, A.I.C.
Director & Licencing Authority
O/o. the Director General,
Drugs and Copyrights,
Drugs Control Administration,
SMC Campus Gunadala,
Vijayawada, NTR dist- 520008,
Andhra Pradesh, India.

To
M/s. Deccan Remedies Limited,
Plot No.58 A,Road No.01,
J.N Pharma City,
Parawada,
Anakapalli District – 531021,
Andhra Pradesh, India.

Sirs,

Sub:- Drugs and Cosmetics Act, 1940 and Rules made there under – Change of Constitution (W.e.f. 29.05.2023) and Firm Name Change in License in Form-25 – Regarding.

Ref:- Your Application, dated: Nil

With reference to your application cited, I forward herewith the Drug Manufacturing License in Form-25 bearing **No.31/VP/AP/2014/B/R/CC** dt: 14.11.2014 for manufacture of the following products and the License is valid from 29-05-2023 to 28-05-2028.

NAME OF THE PRODUCTS

1. Pantoprazole Sodium	–	IP/USP
2. Pregabalin	–	IP
3. Pantoprazole Sodium Seaquihydrate	–	Ph.Eur
4. Allopurinol	–	IP/USP/ Ph.Eur
5. Clopidogrel bisulphate	–	USP/IP
6. Escitalopram Oxalate	–	USP/IP
7. Fexofenadine Hydrochloride	–	USP/Ph.Eur/JP/IP
8. Irbesartan	–	USP/IP
9. Lamivudine	–	USP/Ph.Eur/IP
10. Olmesartan Medoxomile	–	USP/Ph.Eur/IHS
11. Rabeprazole Sodium	–	USP/JP/IP
12. Sertraline Hydrochloride	–	USP/Ph.Eur/IP
13. Telmisartan	–	USP/Ph.Eur/IP
14. Tenofovir Disoproxil Fumarate	–	USP/Ph.Eur/IP
15. Guaifenesin	–	IP/Ph.Eur/USP/BP
16. Methocarbamol	–	IP/ USP
17. Phenazopyridine Hydrochloride	–	USP

The above additional products are granted subject to the following conditions.

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2023.06.01 14:26:47 +05'30'

DIRECTOR & LICENSING AUTHORITY
DRUGS CONTROL ADMINISTRATION

Change of Constitution (w.e.f:19.07.2020) by M/s. Deccan Remedies Limited, Plot No.58 A,Road No.01, J.N Pharma City , Parawada, Anakapalli District – 531021, AP in Form- 25 bearing No. **31/VP/AP/2014/B/R/CC** dt: 14.11.2014 valid upto : 28-05-2028.

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1. You are furthere informed that on failure to manufacture any of the drugs approved herwith without a reasonable cause during the licensing period the matter will be reviewed and such drugs are liable for deletion.

2. The provisions of Drugs Price Conrol Order, 2013 shall e complied with.

3. You should ensure that the drugs approved herewith shall not make any false/ misleading/ objectionable claims and should not be an imitation or resemble any other drugs in respect of design, colour combination etc., and shall comply with all the provisions relating to the laeling of Drugs.

4. Specific permission for each export order need not be obtained. However, the details of export by the exporter shall be furnished immediately after completion of each export in the format mentioned below.

Sl. No:	Date of export	Name of the Drugs exported	Batch No. of the drug	Quantity of the drug	Name of the Importing country.
1.	2.	3.	4.	5.	6.

5. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

6. The specifications/standards asked for by the importing country/firm shall be complied with while exporting the drugs.

7. The federal regulations of the importing country shall be fulfilled while exporting/ supplying the drugs.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th Mal Morar, Gwalior – 6 In case of Narcotics/ Physhotropic Drugs, you are also directed to approach the Narcotic Commissioner of India, 19th The Mal Morar, Gwalior-6 so far as the provisions of the NDPS Act and the Rules are concerned in the matter.

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued here with are liable for suspension/cancellation for which you may take this as a notice under rule 85(2) of the Drugs and Cosmetics Rules. You are therefore requested to plan your production accordingly.

Yours faithfully,

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2023.06.01 14:27:02 +05'30'

DIRECTOR & LICENSING AUTHORITY
DRUGS CONTROL ADMINISTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINSTRATION

From:

M.B.R. Prasad, M. Pharm, M. Phil, AIC,
Director & Licensing Authority
O/o. Director General,
Drugs and Copyrights,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada – 520 008.

To:

M/s. DECCAN REMEDIES LIMITED,
Plot No.58A, Road No:01,
Jawaharlal Nehru Pharma City,
Parawada Mandal, Anakapalli District,
Andhra Pradesh, India – 531 021.

Sir,

Sub: Drugs and Cosmetics Act 1940 and Rules made there under- Approval of additional Product in Form – 25 – Regarding.

Ref: Your application date. 26.07.2024.

@@@

With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug Licenses in **Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated. 14.11.2014, Valid up to 28.05.2028**. for Domestic and Export Purpose.

NAMES OF THE PRODUCTS

1. **Fluconazole - IP/BP/USP/EP For Domestic & Export**
2. **Minoxidil - EP/IP/USP For Domestic & Export**
3. **Ketoconazole - IP/BP/USP/EP For Domestic & Export**

The above additional product is granted subject to the following conditions.

1. You are informed that on failure to manufacture any of the drugs approved herewith without cause during the licensing period the matter will be reviewed, and such are liable for deletion.
2. The provisions of the drug price control Order, 2013 shall be complied with.
3. You should ensure that the drugs approved here with shall not make, any false/misleading / objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating to the labeling of drugs.

Digitally signed by
BABU RAJENDRA
PRASAD MEESALA
Date: 2024.10.26
18:15:24 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

Approval of (03) Additional Products approved to M/s. Deccan Remedies Limited, Plot No. 58A, JN Pharma City, Parawada Mandal, Anakapalli District, Andhra Pradesh. Under drug license in Form – 25 bearing No. **31/VP/AP/2014/B/R/CC**, dated: **14.11.2014**, valid up to **28.05.2028** for Domestic and Export purpose.

4. Specific permission for each export order need not be obtained. However, the details of export by the Exporter shall be furnished immediately after completion of each export in the format mentioned Below.

Sl. No:	Date of export	Name of the Drugs Exported	Batch No. of the Drug	Quantity of the Drug	Name of the Importing Country.
1	2	3	4	5	6

5. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

6. The specifications/standards asked for by the importing country/firm shall be complied with while exporting the drugs.

7. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

8. The Products are approved subject to conducting necessary stability studies.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th Mal Morar, Gwalior – 6 so far as the provisions of the NDPS Act and Rules are concerned in the matter.

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued here with are liable for suspension/cancellation for which you may take this as a notice under rule 85(2) of the drugs and cosmetics rules. You are therefore requested to plan your production accordingly.

Yours faithfully,

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2024.10.26 18:15:39 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINSTRATION

From:

M.B.R. Prasad, M. Pharm, M. Phil, AIC,
Director & Licensing Authority
O/o. Director General,
Drugs and Copyrights,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada – 520 008.

To:

M/s. DECCAN REMEDIES LIMITED,
Plot No.58A, Road No:01,
Jawaharlal Nehru Pharma City,
Parawada Mandal, Anakapalli District,
Andhra Pradesh, India – 531 021.

Sir,

Sub: Drugs and Cosmetics Act 1940 and Rules made there under- Approval of additional Product in Form – 25 – Regarding.

Ref: Your application date. 26.07.2024.

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With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug Licenses in **Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated. 14.11.2014, Valid up to 28.05.2028**. for Domestic and Export Purpose.

NAMES OF THE PRODUCTS

1. **Flupentixol Hydrochloride - BP/IHS For Domestic & Export**
2. **Melitracen Hydrochloride - IHS For Domestic & Export**
3. **Domperidone Maleate - IP/EP For Domestic & Export**

The above additional product is granted subject to the following conditions.

1. You are informed that on failure to manufacture any of the drugs approved herewith without cause during the licensing period the matter will be reviewed, and such are liable for deletion.
2. The provisions of the drug price control Order, 2013 shall be complied with.
3. You should ensure that the drugs approved here with shall not make, any false/misleading / objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating to the labeling of drugs.

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2024.10.26 18:18:06 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

Approval of (03) Additional Products approved to M/s. Deccan Remedies Limited, Plot No. 58A, JN Pharma City, Parawada Mandal, Anakapalli District, Andhra Pradesh. Under drug license in Form – 25 bearing No. **31/VP/AP/2014/B/R/CC**, dated: **14.11.2014**, valid up to **28.05.2028** for Domestic and Export purpose.

4. Specific permission for each export order need not be obtained. However, the details of export by the Exporter shall be furnished immediately after completion of each export in the format mentioned Below.

Sl. No:	Date of export	Name of the Drugs Exported	Batch No. of the Drug	Quantity of the Drug	Name of the Importing Country.
1	2	3	4	5	6

5. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

6. The specifications/standards asked for by the importing country/firm shall be complied with while exporting the drugs.

7. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

8. The Products are approved subject to conducting necessary stability studies.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th Mal Morar, Gwalior – 6 so far as the provisions of the NDPS Act and Rules are concerned in the matter.

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued here with are liable for suspension/cancellation for which you may take this as a notice under rule 85(2) of the drugs and cosmetics rules. You are therefore requested to plan your production accordingly.

Yours faithfully,

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2024.10.26 18:18:14 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINSTRATION

From:

M.B.R. Prasad, M. Pharm, M. Phil, AIC,
Director & Licensing Authority
O/o. Director General,
Drugs and Copyrights,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada – 520 008.

To:

M/s. DECCAN REMEDIES LIMITED,
Plot No.58A, Road No:01,
Jawaharlal Nehru Pharma City,
Parawada Mandal, Anakapalli District,
Andhra Pradesh, India – 531 021.

Sir,

Sub: Drugs and Cosmetics Act 1940 and Rules made there under- Approval of additional Products in Form – 25 – Regarding.

Ref: Your application date. 13.03.2025.

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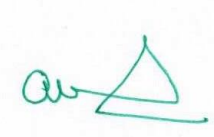
With reference to your application cited, you are hereby permitted to manufacture the following product as additional items under your Drug Licenses in **Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated. 14.11.2014, Valid up to 28.05.2028**. for Domestic and Export Purpose.

NAMES OF THE PRODUCTS

1. **Etoricoxib - IP - For Domestic & Export**
2. **Nitazoxanide - IH - For Domestic & Export**
3. **Voriconazole - IP/EP/BP/USP For Domestic & Export**

The above additional product is granted subject to the following conditions.

1. You are informed that on failure to manufacture any of the drugs approved herewith without cause during the licensing period the matter will be reviewed, and such are liable for deletion.
2. The provisions of the drug price control Order, 2013 shall be complied with.
3. You should ensure that the drugs approved here with shall not make, any false/misleading / objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating to the labeling of drugs.



Digitally signed by
BABU RAJENDRA
PRASAD MEESALA
Date: 2025.05.21
17:42:43 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

Approval of (03) Additional Products approved to M/s. Deccan Remedies Limited, Plot No. 58A, JN Pharma City, Parawada Mandal, Anakapalli District, Andhra Pradesh. Under drug license in Form – 25 bearing No. **31/VP/AP/2014/B/R/CC**, dated: **14.11.2014**, valid up to **28.05.2028** for Domestic and Export purpose.

4. Specific permission for each export order need not be obtained. However, the details of export by the Exporter shall be furnished immediately after completion of each export in the format mentioned Below.

Sl. No:	Date of export	Name of the Drugs Exported	Batch No. of the Drug	Quantity of the Drug	Name of the Importing Country.
1	2	3	4	5	6

5. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

6. The specifications/standards asked for by the importing country/firm shall be complied with while exporting the drugs.

7. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

8. The Products are approved subject to conducting necessary stability studies.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th Mal Morar, Gwalior – 6 so far as the provisions of the NDPS Act and Rules are concerned in the matter.

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued here with are liable for suspension/cancellation for which you may take this as a notice under rule 85(2) of the drugs and cosmetics rules. You are therefore requested to plan your production accordingly.

Yours faithfully,

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2025.05.21 17:42:53 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINSTRATION

From:

M.B.R. Prasad, M. Pharm, M. Phil, AIC,
Director & Licensing Authority
O/o. Director General,
Drugs and Copyrights,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada – 520 008.

To:

M/s. DECCAN REMEDIES LIMITED,
Plot No.58A, Road No:01,
Jawaharlal Nehru Pharma City,
Parawada Mandal, Anakapalli District,
Andhra Pradesh, India – 531 021.

Sir,

Sub: Drugs and Cosmetics Act 1940 and Rules made there under- Approval of additional Product in Form – 25 – Regarding.

Ref: Your application date. 17.03.2025.

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With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug Licenses in **Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated. 14.11.2014, Valid up to 28.05.2028**. **For Domestic and Export Purpose**.

NAME OF THE PRODUCT

Tapentadol Hydrochloride -Ph.Eur

The above additional product is granted subject to the following conditions.

1. You are informed that on failure to manufacture any of the drugs approved herewith without cause during the licensing period the matter will be reviewed, and such are liable for deletion.
2. The provisions of the drug price control Order, 2013 shall be complied with.
3. You should ensure that the drugs approved here with shall not make, any false/misleading / objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating to the labeling of drugs.

Digitally signed by
BABU RAJENDRA
PRASAD MEESALA
Date: 2025.05.21
17:34:12 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

Approval of (01) Additional Products approved to M/s. Deccan Remedies Limited, Plot No. 58A, JN Pharma City, Parawada Mandal, Anakapalli District, Andhra Pradesh. Under drug license in Form – 25 bearing No. **31/VP/AP/2014/B/R/CC**, dated: **14.11.2014**, valid up to **28.05.2028** for Domestic and Export purpose.

4. Specific permission for each export order need not be obtained. However, the details of export by the Exporter shall be furnished immediately after completion of each export in the format mentioned Below.

Sl. No:	Date of export	Name of the Drugs Exported	Batch No. of the Drug	Quantity of the Drug	Name of the Importing Country.
1	2	3	4	5	6

5. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

6. The specifications/standards asked for by the importing country/firm shall be complied with while exporting the drugs.

7. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

8. The Products are approved subject to conducting necessary stability studies.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th Mal Morar, Gwalior – 6 so far as the provisions of the NDPS Act and Rules are concerned in the matter.

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued here with are liable for suspension/cancellation for which you may take this as a notice under rule 85(2) of the drugs and cosmetics rules. You are therefore requested to plan your production accordingly.

Yours faithfully,

Digitally signed by BABU RAJENDRA PRASAD MEESALA
Date: 2025.05.21 17:34:21 +05'30'

DIRECTOR AND LICENSING AUTHORITY
DRUGS CONTROL AMINSTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From

M B R Prasad, M. Pharm, M. Phil, A.I.C.
Director & Licensing Authority,
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan Remedies Limited, Plot No. 58/A,
Road No. 01, JN Pharma city, Parawada,
Anakapalli-531021, Andhra Pradesh.

Sir,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder–Approval of Additional Products in Form-25 - Regarding.

Ref: Application dated : 27/06/2025

With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug License in Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated 14/11/2014 valid up to 28/05/2028

NAME OF THE DRUG

Name of the product	Market	Specifications	Composition
Acyclovir	Export & Domestic Only	IP	NA
Acyclovir	Export & Domestic Only	USP	NA
Triclabendazole	Export & Domestic Only	InHouse	NA
Ezetimibe	Export & Domestic Only	IP	NA
Ezetimibe	Export & Domestic Only	USP	NA

The above additional products is permitted subject to the following conditions.

1. Specific permission for each export order need not be obtained. However, the details of exports by the exporter shall be furnished immediately after completion of each export in the format mentioned below.

Sl. No.	Date of export	Names of the drugs Exported	Batch No. of the drug.	Quantity of the drug.	Name of the importing Country.
1.	2.	3.	4.	5.	6.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEEGALA
Date: 24-Jul-2025 19:46:27

Single Desk Approval Ref ID : SDDHW238250062

Approval of (Five) Additional Product to Deccan Remedies Limited, Plot No. 58/A, Road No. 01, JN Pharma city, Parawada, Anakapalli-531021, Andhra Pradesh. -in Form-25 baring No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

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2. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

3. The specification/standards asked for by the Importing Country/Firm shall be complied with while exporting the drugs.

4. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

5. You are further informed that on failure to manufacture any of the drugs approved here with without a reasonable cause during the licensing period the matter will be reviewed and such drugs are liable for deletion.

6. The provisions of Drugs Price Control order, 2013 shall be complied with.

7. The Product are approved subject to conducting necessary Stability Studies

8. You should ensure that the drugs approved here with shall not make any false/Misleading objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating the labelling of Drugs.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th The Mal Morar, Gwalior-6 so far as the provisions of the NDPS Act and the Rules are concerned in the matter.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEESALA
Date: 24-Jul-2025 19:46:27

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued herewith are liable for suspension/cancellation for which you may take this as a notice under Rule 85 (2) of the Drugs and Cosmetics Rules. You are therefore requested to plan your production accordingly

Yours faithfully,

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

Signature Not Verified

Digitally Signed.
Name: BABU RAJENDRA
PRASAD MEESALA
Date: 24-Jul-2025 19:46:27

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From

M B R Prasad, M. Pharm, M. Phil, A.I.C.
Director & Licensing Authority,
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan Remedies Limited, Plot No. 58/A,
Road No. 01, JN Pharma city, Parawada,
Anakapalli-531021, Andhra Pradesh.

Sir,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder–Approval of Additional Products in Form-25 - Regarding.

Ref: Application dated : 27/06/2025

With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug License in Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated 14/11/2014 valid up to 28/05/2028

NAME OF THE DRUG

Name of the product	Market	Specifications	Composition
Carvedilol	Export & Domestic Only	IP	NA
Oxcarbazepine	Export & Domestic Only	PhEUR	NA
Atorvastatin Calcium	Export & Domestic Only	USP	NA
Oxcarbazepine	Export & Domestic Only	USP	NA
Atorvastatin Calcium	Export & Domestic Only	IP	NA
Carvedilol	Export & Domestic Only	USP	NA
Carvedilol	Export & Domestic Only	PhEUR	NA

The above additional products is permitted subject to the following conditions:

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD VINEESALA
Date: 24-Jul-2025 19:09:56

1. Specific permission for each export order need not be obtained. However, the details of exports by the exporter shall be furnished immediately after completion of each export in the format mentioned below.

Sl. No.	Date of export	Names of the drugs Exported	Batch No. of the drug.	Quantity of the drug.	Name of the importing Country.
1.	2.	3.	4.	5.	6.

Approval of (Seven) Additional Product to Deccan Remedies Limited, Plot No. 58/A, Road No. 01, JN Pharma city, Parawada, Anakapalli-531021, Andhra Pradesh. -in Form-25 baring No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

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2. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

3. The specification/standards asked for by the Importing Country/Firm shall be complied with while exporting the drugs.

4. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

5. You are further informed that on failure to manufacture any of the drugs approved here with without a reasonable cause during the licensing period the matter will be reviewed and such drugs are liable for deletion.

6. The provisions of Drugs Price Control order, 2013 shall be complied with.

7. The Product are approved subject to conducting necessary Stability Studies

8. You should ensure that the drugs approved here with shall not make any false/Misleading objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating the labelling of Drugs.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th The Mal Morar, Gwalior-6 Signature provisions of the NDPS Act and the Rules are concerned in the matter.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEESALA
Date: 24-Jul-2025 19:09:56

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued herewith are liable for suspension/cancellation for which you may take this as a notice under Rule 85 (2) of the Drugs and Cosmetics Rules. You are therefore requested to plan your production accordingly

Yours faithfully,

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

Signature Not Verified

Digitally Signed.
Name: BABU RAJENDRA
PRASAD MEESALA
Date: 24-Jul-2025 19:09:56

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From

M B R Prasad, M. Pharm, M. Phil, A.I.C.
Director & Licensing Authority,
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan Remedies Limited, Plot No. 58/A,
Road No. 01, JN Pharma city, Parawada,
Anakapalli-531021, Andhra Pradesh.

Sir,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder–Approval of Additional Products in Form-25 - Regarding.

Ref: Application dated : 27/06/2025

With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug License in Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated 14/11/2014 valid up to 28/05/2028

NAME OF THE DRUG

Name of the product	Market	Specifications	Composition
Mesna	Export & Domestic Only	IP	NA
Mesna	Export & Domestic Only	BP	NA
Levocarnitine	Export & Domestic Only	USP	NA
Dolutegravir Sodium	Export & Domestic Only	IP	NA
Mesna	Export & Domestic Only	PhEUR	NA
Mesna	Export & Domestic Only	USP	NA
Levocarnitine	Export & Domestic Only	PhEUR	NA

The above additional products is permitted subject to the following conditions:

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD VINEESHALA
Date: 24-Jul-2025 19:42:14

1. Specific permission for each export order need not be obtained. However, the details of exports by the exporter shall be furnished immediately after completion of each export in the format mentioned below.

Sl. No.	Date of export	Names of the drugs Exported	Batch No. of the drug.	Quantity of the drug.	Name of the importing Country.
1.	2.	3.	4.	5.	6.

Approval of (Seven) Additional Product to Deccan Remedies Limited, Plot No. 58/A, Road No. 01, JN Pharma city, Parawada, Anakapalli-531021, Andhra Pradesh. -in Form-25 baring No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

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2. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

3. The specification/standards asked for by the Importing Country/Firm shall be complied with while exporting the drugs.

4. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

5. You are further informed that on failure to manufacture any of the drugs approved here with without a reasonable cause during the licensing period the matter will be reviewed and such drugs are liable for deletion.

6. The provisions of Drugs Price Control order, 2013 shall be complied with.

7. The Product are approved subject to conducting necessary Stability Studies

8. You should ensure that the drugs approved here with shall not make any false/Misleading objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating the labelling of Drugs.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th The Mal Morar, Gwalior-6 Signature provisions of the NDPS Act and the Rules are concerned in the matter.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEESALA
Date: 24-Jul-2025 19:42:14

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued herewith are liable for suspension/cancellation for which you may take this as a notice under Rule 85 (2) of the Drugs and Cosmetics Rules. You are therefore requested to plan your production accordingly

Yours faithfully,

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

Signature Not Verified

Digitally Signed.
Name: BABU RAJENDRA
PRASAD MEESALA
Date: 24-Jul-2025 19:42:14

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From

M B R Prasad, M. Pharm, M. Phil, A.I.C.
Director & Licensing Authority,
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan Remedies Limited, Plot No. 58/A,
Road No. 01, JN Pharma city, Parawada,
Anakapalli-531021, Andhra Pradesh.

Sir,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder–Approval of Additional Products in Form-25 - Regarding.

Ref: Application dated : 27/06/2025

With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug License in Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated 14/11/2014 valid up to 28/05/2028

NAME OF THE DRUG

Name of the product	Market	Specifications	Composition
Montelukast Sodium	Export & Domestic Only	USP	NA
Moxifloxacin Hydrochloride	Export & Domestic Only	PhEUR	NA
Moxifloxacin Hydrochloride	Export & Domestic Only	USP	NA
Apixaban	Export & Domestic Only	InHouse	NA
Montelukast Sodium	Export & Domestic Only	IP	NA
Moxifloxacin Hydrochloride	Export & Domestic Only	IP	NA
Montelukast Sodium	Export & Domestic Only	PhEUR	NA

The above additional products is permitted subject to the following conditions.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEESALA
Date: 24-Jul-2025 19:18:25

Single Desk Approval Ref ID : SDDHW238250060

1. Specific permission for each export order need not be obtained. However, the details of exports by the exporter shall be furnished immediately after completion of each export in the format mentioned below.

Sl. No.	Date of export	Names of the drugs Exported	Batch No. of the drug.	Quantity of the drug.	Name of the importing Country.
1.	2.	3.	4.	5.	6.

Approval of (Seven) Additional Product to Deccan Remedies Limited, Plot No. 58/A, Road No. 01, JN Pharma city, Parawada, Anakapalli-531021, Andhra Pradesh. -in Form-25 baring No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

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2. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

3. The specification/standards asked for by the Importing Country/Firm shall be complied with while exporting the drugs.

4. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

5. You are further informed that on failure to manufacture any of the drugs approved here with without a reasonable cause during the licensing period the matter will be reviewed and such drugs are liable for deletion.

6. The provisions of Drugs Price Control order, 2013 shall be complied with.

7. The Product are approved subject to conducting necessary Stability Studies

8. You should ensure that the drugs approved here with shall not make any false/Misleading objectionable claims and should not be an imitation of trade mark of any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating the labelling of Drugs.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th The Mal Morar, Gwalior-6 so far as the provisions of the NDPS Act and the Rules are concerned in the matter.

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued herewith are liable for suspension/cancellation for which you may take this as a notice under Rule 85 (2) of the Drugs and Cosmetics Rules. You are therefore requested to plan your production accordingly

Yours faithfully,

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

Signature Not Verified

Digitally Signed.
Name: BABU RAJENDRA PRASAD MEESALA
Date: 24-Jul-2025 19:18:25

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From

M B R Prasad, M. Pharm, M. Phil, A.I.C.
Director & Licensing Authority,
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan Remedies Limited, Plot No. 58/A,
Road No. 01, JN Pharma city, Parawada,
Anakapalli-531021, Andhra Pradesh.

Sir,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder–Approval of Additional Products in Form-25 - Regarding.

Ref: Application dated : 27/06/2025

With reference to your application cited, you are hereby permitted to manufacture the following product as additional item under your Drug License in Form-25 bearing No.31/VP/AP/2014/B/R/CC, dated 14/11/2014 valid up to 28/05/2028

NAME OF THE DRUG

Name of the product	Market	Specifications	Composition
Posaconazole	Export & Domestic Only	InHouse	NA
Carisoprodol	Export & Domestic Only	USP	NA
Gliclazide	Export & Domestic Only	IP	NA
Gliclazide	Export & Domestic Only	PhEUR	NA
Carisoprodol	Export & Domestic Only	IP	NA

The above additional products is permitted subject to the following conditions.

1. Specific permission for each export order need not be obtained. However, the details of exports by the exporter shall be furnished immediately after completion of each export in the format mentioned below.

Sl. No.	Date of export	Names of the drugs Exported	Batch No. of the drug.	Quantity of the drug.	Name of the importing Country.
1.	2.	3.	4.	5.	6.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEEGALA
Date: 24-Jul-2025 19:50:05

Single Desk Approval Ref ID : SDDHW238250063

Approval of (Five) Additional Product to Deccan Remedies Limited, Plot No. 58/A, Road No. 01, JN Pharma city, Parawada, Anakapalli-531021, Andhra Pradesh. -in Form-25 baring No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

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2. Detailed particulars of rejects/returned goods if any shall be furnished to this office at once for the purpose of issuing necessary orders in such cases. Till such time, the goods shall not be altered/disposed of in any other manner.

3. The specification/standards asked for by the Importing Country/Firm shall be complied with while exporting the drugs.

4. The federal regulations of the importing country shall be fulfilled while exporting/supplying the drugs.

5. You are further informed that on failure to manufacture any of the drugs approved here with without a reasonable cause during the licensing period the matter will be reviewed and such drugs are liable for deletion.

6. The provisions of Drugs Price Control order, 2013 shall be complied with.

7. The Product are approved subject to conducting necessary Stability Studies

8. You should ensure that the drugs approved here with shall not make any false/Misleading objectionable claims and should not be an imitation or resemble any other drug in respect of design, colour combination etc., and shall comply with all the provisions relating the labelling of Drugs.

In case of Narcotics/Psychotropic Drugs, you are also directed to approach The Narcotic Commissioner of India, 19th The Mal Morar, Gwalior-6 so far as the provisions of the NDPS Act and the Rules are concerned in the matter.

Signature Not Verified
Digitally Signed.
Name: BABU RAJENDRA PRASAD MEESALA
Date: 24-Jul-2026 19:50:05

Further, you are informed that non-compliance of any of the conditions mentioned above the matter will be reviewed and the licenses/permissions issued herewith are liable for suspension/cancellation for which you may take this as a notice under Rule 85 (2) of the Drugs and Cosmetics Rules. You are therefore requested to plan your production accordingly

Yours faithfully,

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

Signature Not Verified

Digitally Signed.
Name: BABU RAJENDRA
PRASAD MEESALA
Date: 24-Jul-2025 19:50:05