Ref: GLL/BSE/2026-27/Sep- Date: September 08, 2026

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 Additional 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 following products as additional item 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	Caffeine	IP,BP,PhEUR,JP,USP
2	Duloxetine Hydrochloride	IP,BP,PhEUR,USP
3	Mebeverine Hydrochloride	IP,BP,PhEUR
4	Phenylephrine Hydrochloride	IP,BP,PhEUR,JP,USP
5	Trimebutine Maleate	IP,BP,PhEUR,JP
6	Oxyclozanide	IP,BP,InHouse

In addition to above, M/s Deccan Remedies Limited, has also received Approval from the Drug Control Administration, Government of Andhra Pradesh granting Test Licence for manufacture of following drugs for purpose of examination, test or analysis under Form 29, valid for three years from the date of issue.

S.No.	Product Name	Pharmacopoeias
1	Ranolazine	IH.
2	Cinnarizine	IH/Ph. Eur.
3	Sacubitril	IH.
4	Triclabendazole	USP/BP.

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

ARIHANT BAID
Managing Director
DIN#01171845

ARIHANT BAID Digitally signed by ARIHANT BAID Date: 2026.09.08 17:00:07 +05'30'

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

Grant of License for Additional Products

Form 25

From

M.Pandu Ranga Prasad
Director & Licensing Authority(FAC),
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan remedies Ltd, Plot No-58/A,Road No-1,JNPharmacity,Parawada,Anakapalli,AP.India.

Sir,

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

Ref: Application dated : 11/08/2026

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
Caffeine	Export & Domestic Only	IP,BP,PhEUR,JP,USP	NA
Duloxetine Hydrochloride	Export & Domestic Only	IP,BP,PhEUR,USP	NA
Mebeverine Hydrochloride	Export & Domestic Only	IP,BP,PhEUR	NA

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

Signature Not Verified
Digitally Signed.
Name: PANDURANGA PRASAD MADDALA
Date: 02-Sep-2026 11:52:10

Single Desk Approval Ref ID : SDDHW238260254

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

Approval of (Three) Additional Product to Deccan remedies Ltd, Plot No-58/A,Road No-1,JNPharmacity,Parawada,Anakapalli,AP.India. -in Form 25 bearing No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

Signature Not Verified
Digitally Signed.
Name: PANDURANGA PRASAD MADDALA
Date: 02-Sep-2016 11:52:10

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.

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,

Signature Not Verified
Digitally Signed.
Name: PANDURANGA PRASAD MADDALA
Date: 02-Sep-2016 11:52:10

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

Grant of License for Additional Products

Form 25

From

M.Pandu Ranga Prasad
Director & Licensing Authority(FAC),
O/o. The Director General,
Drugs Control Administration,
Siddhartha Medical College Campus,
Gunadala, Vijayawada - 520 008,
Andhra Pradesh, India.

To

Deccan remedies Ltd, Plot No-58/A,Road No-1,JNPharmacity,Parawada,Anakapalli,AP.India.

Sir,

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

Ref: Application dated : 12/08/2026

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
Phenylephrine Hydrochloride	Export & Domestic Only	IP,BP,PhEUR,JP,USP	NA
Trimebutine Maleate	Export & Domestic Only	IP,BP,PhEUR,JP	NA
Oxyclozanide	Export & Domestic Only	IP,BP,InHouse	NA

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

Signature Not Verified
Digitally Signed.
Name: PANDURANGA PRASAD MADDALA
Date: 01-Sep-2026 17:49:03

Single Desk Approval Ref ID : SDDHW238260243

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

Approval of (Three) Additional Product to Deccan remedies Ltd, Plot No-58/A,Road No-1,JNPharmacity,Parawada,Anakapalli,AP.India. -in Form 25 bearing No.31/VP/AP/2014/B/R/CC dated 14/11/2014 valid up to 28/05/2028

Signature Not Verified
Digitally Signed.
Name: PANDURANGA PRASAD MADDALA
Date: 01-Sep-2016 17:49:03

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.

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,

Signature Not Verified
Digitally Signed.
Name: PANDURANGA PRASAD MADDALA
Date: 01-Sep-2016 17:49:03

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

From:

M. Pandu Ranga Prasad,
Director (FAC) & Licensing Authority
O/o. The Director General,
Drugs Control Administration,
Hanumanpet, Vijayawada - 520003,
Vijayawada – 520003,
Andhra Pradesh, INDIA.

To:

M/s. DECCAN REMEDIES LIMITED,
Plot No: 58/A, Road No.01,
J.N. Pharma City,
Parawada Mandal,
Anakapalli District – 531021,
Andhra Pradesh, INDIA

Sir,

Sub:-Drugs and Cosmetics Act, 1940 and Rules made there under - **Grant of Test License in Form-29**– Regarding.

Ref:- Your Application Date: 10.07.2026.

*****

With reference to your letter cited above, I am forwarding herewith the Test License in Form – 29 for the purpose of Examination, test for Chemical and Instrumental analysis.

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

This Product(s) manufactured against this Test License is not for any commercial use and shall be used for Bio/clinical studies subject to the grant of BE/CT permission from the DCG(I) office.

The label of the container shall bear the words "Not of medical use" and "For test or analysis only" in bold letters.

You are required to ensure the quantity of the drug(s) manufactured on basis of the above NOC is shall be supplied to the formulation manufacturer for the specified purpose and no part of it is diverted for domestic sale in India though a declaration.

You shall ensure that in the event of Non- Materialization of formulation development the same shall be intimated to the Licensing Authority and the manufacturer shall ensure physical destruction of such stocks in the presence of State Licensing Authority.

Yours faithfully,

Digitally signed by
PANDURANGA PRASAD
MADDALA
Date: 2026.09.08 11:47:41
+05'30'

DIRECTOR (FAC) AND LICENSING AUTHORITY
DRUGS CONTROL ADMINISTRATION

Grant of Test License

FORM-29
(See Rule 89)

LICENSE TO MANUFACTURE DRUGS FOR PURPOSE OF EXAMINATION, TEST OR ANALYSIS

1 Sri. A.V.R. Harrsha, Director of M/s. DECCAN REMEDIES LIMITED, is hereby licensed to manufacture the drugs specified below for purpose of Examination, Test or Analysis at Plot No: 58/A, Road No.01, J.N.Pharma City, Parawada Mandal, Anakapalli District – 531021 , Andhra Pradesh, India.

2 This license is subject to the conditions prescribed in part VIII of the Drugs and Cosmetics Rules 1945.

3 This License shall be in force for **Three Years** from the date of issue.

NAME(S) OF THE PRODUCT(S)

S.No	Name of the Drug	Specifications
1	Ranolazine	IH.
2	Cinnarizine	IH/Ph. Eur.
3	Sacubitril	IH.
4	Triclabendazole	USP/BP.

Digitally signed by PANDURANGA PRASAD MADDALA
Date: 2026.09.08 11:47:52 +05'30'

DIRECTOR (FAC) AND LICENSING AUTHORITY
DRUGS CONTROL ADMINISTRATION