GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As

FORM 20

[See Rule 61 (1)] **Change of Pharmacist w.e.f 10/12/2025**

[Licence to sell, stock or exhibit or offer for sale, or distribute] drugs by retail other than those specified in [Schedules C, C(1) and X]

1.

1 **CH.BHADRA REDDY, (DIRECTOR**)

2 **CH.MAHENDAR REDDY, (DIRECTOR**)

of **M/S.MALLAREDDY MEDICALS(A UNIT OF MALLAREDDY HEALTHCARE PRIVATE LIMITED)**is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs other than those specified in [Schedules C, C (1) and X] of the Drugs and Cosmetics Rules 1945, *and to operate a pharmacy on the premises situated at **D.NO.2-18/2,GROUND FLOOR,SY.NO.138,C/O MALLAREDDY HOSPITAL,SURARAM,JEEDIMETLA,HYDERABAD,SURARAM VILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI DIST,TELANGANA,INDIA.,SURARAM VILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE, INDIA,TELANGANA,INDIA** Subject to the conditions specified below and to provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder.

2.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

3.Name(s) of qualified person(s) in charge

1 **SURESH KUMAR NETHI, D PHARMACY**, **RegNo. TG076627, DT: 29/11/2025**

2 **VADTHYA JYOTHI, B.PHARMACY**, **RegNo. TS001992, DT: 08/02/2019**

4.Categories of drugs : Drugs Other than those specified in [Schedules C, C(1) and X]

Licence No: **TS/MDL/2024-112709**

Date:03/01/2026

Next License Retention Fee Due Dt:01/01/2029

Digitally Signed By
KANDADI PRABHAKAR
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:03-01-2026 13:15:14 PM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
4. No drug shall be sold unless such drug is purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As

FORM 21

[See Rule 61 (2)]

Change of Pharmacist w.e.f 10/12/2025

(Licence to sell, stock or exhibit or offer for sale,or distribute) by retail drugs specified in Schedules C and C (1)(excluding those specified in Schedule X)

1.

1 **CH.BHADRA REDDY, (DIRECTOR**)

2 **CH.MAHENDAR REDDY, (DIRECTOR**)

of **M/S.MALLAREDDY MEDICALS(A UNIT OF MALLAREDDY HEALTHCARE PRIVATE LIMITED)**is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs specified in Schedules C and C (1)[excluding those specified in Schedule X] of the Drugs and Cosmetics Rules 1945, *and to operate a pharmacy on the premises situated at **D.NO.2-18/2,GROUND FLOOR,SY.NO.138,C/O MALLAREDDY HOSPITAL,SURARAM,JEEDIMETLA,HYDERABAD,SURARAM VILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI DIST,TELANGANA,INDIA.,SURARAM VILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE, INDIA,TELANGANA,INDIA**Subject to the conditions specified below and to provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder.

2.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

3.Name(s) of qualified person(s) in charge

1 **SURESH KUMAR NETHI, D PHARMACY**, **RegNo. TG076627, DT: 29/11/2025**

2 **VADTHYA JYOTHI, B.PHARMACY**, **RegNo. TS001992, DT: 08/02/2019**

4.Categories of drugs : Drugs specified in Schedules C and C (1)[excluding those specified in Schedule X]

Licence No: **TS/MDL/2024-112709**

Date:03/01/2026

Next License Retention Fee Due Dt:01/01/2029

Digitally Signed By
KANDADI PRABHAKAR
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:03-01-2026 13:15:14 PM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
4. No drug shall be sold unless such drug is purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As
CH.MAHENDER REDDY (DIRECTOR)

FORM 20

Registered Pharmacist
R.SRIKANTH (D.PHARMACY)

[See Rule 61 (1)]

[Licence to sell, stock or exhibit or offer for sale, or distribute] drugs by retail other than those specified in [Schedules C, C(1) and X]

1.

1 **CH.BADRA REDDT, (DIRECTOR**)

2 **CH.MAHENDER REDDY, (DIRECTOR**)

of **M/s M/S MALLAREDDY MEDICALS (A UNIT OF MALLAREDDY HEALTHCARE PRIVATE LIMITED)** is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs other than those specified in [Schedules C, C (1) and X] of the Drugs and Cosmetics Rules 1945, *and to operate a pharmacy on the premises situated at **M/S.MALLAREDDY MEDICALS A UNIT OF MALLAREDDY HEALTHCARE PRIVATELIMITED,C-BLOCK SITUATED AT D.NO.2-18/2,GROUND FLOOR,SY NO.138,C/O MALLAREDDY HOSPITAL,SURARAM,JEEDIMETLA,HYDERABAD,SURARAMVILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI DIST,TELANGANA,IN, Suraram(V), Quthbullapur(M), MEDCHAL - MALKAJGIRI(Dist.),Telangana,India** Subject to the conditions specified below and to provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder.

2.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

3.Name(s) of qualified person(s) in charge

1 **R.SREEDHAR, D.PHARMACY**, **RegNo. TS048592, DT: 04/08/2023**

2 **R.SRIKANTH, D.PHARMACY**, **RegNo. TS013973, DT: 20/11/2020**

4.Categories of drugs : Drugs Other than those specified in [Schedules C, C(1) and X]

Licence No: **TS/MDL/2024-113024**

Date:09/01/2024

Next License Retention Fee Due Dt:08/01/2029

Digitally Signed By
VIJAYA GOPAL MAYURI
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:09-01-2024 08:17:16 AM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
4. No drug shall be sold unless such drug is purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As
CH.MAHENDER REDDY (DIRECTOR)

FORM 21

Registered Pharmacist
R.SRIKANTH (D.PHARMACY)

[See Rule 61 (2)]

(Licence to sell, stock or exhibit or offer for sale,or distribute) by retail drugs specified in Schedules C and C (1)(excluding those specified in Schedule X)

1.

1 **CH.BADRA REDDT, (DIRECTOR**)

2 **CH.MAHENDER REDDY, (DIRECTOR**)

of **M/s M/S MALLAREDDY MEDICALS (A UNIT OF MALLAREDDY HEALTHCARE PRIVATE LIMITED)** is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs specified in Schedules C and C (1)[excluding those specified in Schedule X] of the Drugs and Cosmetics Rules 1945, *and to operate a pharmacy on the premises situated at **M/S.MALLAREDDY MEDICALS A UNIT OF MALLAREDDY HEALTHCARE PRIVATELIMITED,C-BLOCK SITUATED AT D.NO.2-18/2,GROUND FLOOR,SY NO.138,C/O MALLAREDDY HOSPITAL,SURARAM,JEEDIMETLA,HYDERABAD,SURARAMVILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI DIST,TELANGANA,IN, Suraram(V), Quthbullapur(M), MEDCHAL - MALKAJGIRI(Dist.),Telangana,India**
Subject to the conditions specified below and to provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder.

2.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

3.Name(s) of qualified person(s) in charge

1 **R.SREEDHAR, D.PHARMACY**, **RegNo. TS048592, DT: 04/08/2023**

2 **R.SRIKANTH, D.PHARMACY**, **RegNo. TS013973, DT: 20/11/2020**

4.Categories of drugs : Drugs specified in Schedules C and C (1)[excluding those specified in Schedule X]

Licence No: **TS/MDL/2024-113024**

Date:09/01/2024

Next License Retention Fee Due Dt:08/01/2029

Digitally Signed By
VIJAYA GOPAL MAYURI
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:09-01-2024 08:17:16 AM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
4. No drug shall be sold unless such drug is purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As

FORM 20

[See Rule 61 (1)]

Change of Pharmacist w.e.f 05/05/2025

[Licence to sell, stock or exhibit or offer for sale, or distribute] drugs by retail other than those specified in [Schedules C, C(1) and X]

1.

1 **CH.BHADRA REDDY, (DIRECTOR**)

2 **CH.MAHENDER REDDY, (DIRECTOR**)

of **M/S MALLAREDDY MEDICALS(A UNIT OF MALLAREDDY HEALTHCARE PRIVATE LIMITED)**is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs other than those specified in [Schedules C, C (1) and X] of the Drugs and Cosmetics Rules 1945, *and to operate a pharmacy on the premises situated at **D.NO.2-18/2,GROUNDFLOOR,SY.NO.138,C/O MALLAREDDY HOSPITAL,SURARAM,JEEDIMETLA,HYDERABAD.,SURARAM VILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE, INDIA,TELANGANA,INDIA** Subject to the conditions specified below and to provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder.

2.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

3.Name(s) of qualified person(s) in charge

1 **PALLE SAILOO, B.PHARMACY**, **RegNo. TS028633, DT: 28/06/2022**

2 **MD.SADDAM HUSSAIN, B.PHARMACY**, **RegNo. TS035647, DT: 21/11/2022**

4.Categories of drugs : Drugs Other than those specified in [Schedules C, C(1) and X]

Licence No: **98/RR1/AP/2011**

Date:03/06/2025

Next License Retention Fee Due Dt:25/06/2030

Digitally Signed By
KANDADI PRABHAKAR
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:03-06-2025 14:05:38 PM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
4. No drug shall be sold unless such drug is purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As

FORM 21

[See Rule 61 (2)]

Change of Pharmacist w.e.f 05/05/2025

(Licence to sell, stock or exhibit or offer for sale,or distribute) by retail drugs specified in Schedules C and C (1)(excluding those specified in Schedule X)

1.

1 **CH.BHADRA REDDY, (DIRECTOR**)

2 **CH.MAHENDER REDDY, (DIRECTOR**)

of **M/S MALLAREDDY MEDICALS(A UNIT OF MALLAREDDY HEALTHCARE PRIVATE LIMITED)**is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs specified in Schedules C and C (1)[excluding those specified in Schedule X] of the Drugs and Cosmetics Rules 1945, *and to operate a pharmacy on the premises situated at **D.NO.2-18/2,GROUNDFLOOR,SY.NO.138,C/O MALLAREDDY HOSPITAL,SURARAM,JEEDIMETLA,HYDERABAD.,SURARAM VILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE, INDIA,TELANGANA,INDIA**Subject to the conditions specified below and to provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder.

2.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

3.Name(s) of qualified person(s) in charge

1 **PALLE SAILOO, B.PHARMACY**, **RegNo. TS028633, DT: 28/06/2022**

2 **MD.SADDAM HUSSAIN, B.PHARMACY**, **RegNo. TS035647, DT: 21/11/2022**

4.Categories of drugs : Drugs specified in Schedules C and C (1)[excluding those specified in Schedule X]

Licence No: **98/RR1/AP/2011**

Date:03/06/2025

Next License Retention Fee Due Dt:25/06/2030

Digitally Signed By
KANDADI PRABHAKAR
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:03-06-2025 14:05:38 PM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
4. No drug shall be sold unless such drug is purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

GOVERNMENT OF TELANGANA
DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As
CH.BHADRA REDDY (DIRECTOR)

FORM 20F

Registered Pharmacist
CHERLIPELLI MOUNIKA
(B.PHARMACY)

[See Rule 61 (3)]

Licence to sell, stock or exhibit for sale or distribute by drugs by retail drugs specified in [Schedules X].

1.

1 CH.BHADRA REDDY, (DIRECTOR)

2 CH.MAHENDER REDDY, (DIRECTOR)

of **M/s MALLAREDDY MEDICALS OT PHARMACY** is hereby [licensed to sell, stock or exhibit or offer for sale, or distribute] by retail drugs specified in Schedules X to the Drugs and Cosmetics Rules 1945, on the premises situated at **M/S MALLAREDY MEDICALS-OTPHARMACY,FIRSTFLOOR,SIITUATED AT SYNO-138,D.NO-2-18/2,C/O MALLAREDDY HOSPITAL,SURARAM,SURARAMVILLAGE,QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI DIST,TELANGANA INDIA-,, Suraram(V), Quthbullapur(M), MEDCHAL - MALKAJGIRI(Dist.),Telangana State,India**

2. Names of drugs:[SCHEDULES X Drugs].

3.The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the conditions of licence and the provisions of the Drugs and Cosmetics Act, 1940 (23 of 1940) and the Drugs and Cosmetics Rules,1945 shall be assessed not less than once in three years or as needed as per risk based approach.

4.Name(s) of qualified person-in-charge.

1 CHERLIPELLI MOUNIKA, B.PHARMACY, RegNo. TS052501, DT: 02/09/2023

5.The license is subject to the conditions stated below and the provisions of the Drugs and Cosmetics Act,1940 and the Rules,made thereunder

Licence No: **TG/MDL/2025-140052**

Date:14/10/2025

License Retention Fee Due Dt:13/10/2030

Digitally Signed By
KANDADI PRABHAKAR
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:14-10-2025 13:38:02 PM

This Document is Digitally Signed. Signature is not required

Conditions Of Licence

---

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall report to the Licensing Authority any change in the qualified staff in charge within one month of such change.
3. No drug shall be stocked or sold unless such drug has been purchased under cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.
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 licence. Where any change in the constitution of the firm takes place, the current licence 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, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

Digitally Signed By
KANDADI PRABHAKAR
Licensing Authority
Assistant Director
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:14-10-2025 13:38:02 PM

DCA
DRUGS CONTROL ADMINISTRATION
TELANGANA

DRUGS CONTROL ADMINISTRATION
Government of Telangana

LICENCE RETENTION FEE RECEIPT

Dated:22/04/2022

This is to certify that the Licence of the firm **M/s MALLAREDDY INSTITUTE OF DENTAL SCIENCES (A UNIT OF CHANDRAMMA EDUCATIONAL SOCIETY)** situated at **SY NO.46.47.48, GROUND FLOOR, SURARAM X ROADS, JEEDIMETLA, QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI (DIST.), SURARAM (VILLAGE), QUTHBULLAPUR (MANDAL), MEDCHAL-MALKAJGIRI (DIST.),TELANGANA, INDIA** bearing Licence No. **TG/15/03/2017-23604, 23605** in the statutory **Form 20 &Form 21** granted/ renewed on **08/04/2022** whose validity would get expired by **07/04/2027,** is hereby permitted to be retained with the extended validity upto **07/04/2027**.with the exiting Constitution and Qualified person as on date.

Retention fees/ with penalty for an amount of Rs. **3060.00** vide challan no:**6100884794** and Bank Transaction No.:**115570833** has been paid on **22/04/2022** by the licensee in accordance with provisions of Rule 63(1) of the Drugs and Cosmetics Rules, 1945.

This is Electronically Generated Receipt

GRANT OF LICENCE

GOVERNMENT OF TELANGANA

DRUGS CONTROL ADMINISTRATION

Partners/Directors/Gpa/As
DR.CH.BHADRAREDDY,PRESIDENT,
,Society

FORM 20

[See Rule 61 (1)]

Registered Pharmacist
B.CHANDANA and REGNO=60522/A1

LICENCE TO SELL,STOCK OR EXHIBIT [OR OFFER] FOR SALE OR DISTRIBUTE DRUGS BY RETAIL OTHER THAN THOSE SPECIFIED IN [SCHEDULES C,C(1) AND X]

1. DR.CH.BHADRAREDDY,PRESIDENT, , ,CH.KALPANA.VICE PRESIDENT, , CH.MAHENDER REDDY,SECRETARY, , P.VASANTHA,JOINT-SECRETARY , P.ANJI REDDY, TREASURER, , CH.SHALINI, ORG-SECRETARY, , CH.PREETHIREDDY,JOINT-SECRETARY Society of **M/S.MALLAREDDY INSTITUTE OF DENTAL SCIENCES PHARMACY (A UNIT OF CHANDRAMMA EDUCATIONAL SOCIETY)**

is here by licenced to sell,stock or exhibit [or offer] for sale or distribute by retail drugs other thanthose specified in [Schedules C and C(1) and X] of the Drugs and Cosmetics Rules 1945,* and to operate a pharmacy on the premises situated at **SY NO.46.47.48,GROUND FLOOR,SURAM X ROAD,, JEEDIMETLA,QUTHBULLAPUR MANDAL, , MEDCHAL DIST,Rangareddy_cir1 (Dist)** subject to the conditions specified below and to the provisions of the Drugs and Cosmetics Act, 1940 and the rules thereunder.

2.The licences shall be in force From **10/04/2017 TO 09/04/2022**

3.Name(s) of qualified person(s) in charge

B.CHANDANA , RegNo. 60522/A1 , DT:02/01/2010(D.O.J: 10/04/2017)

4.Categories of drugs : Drugs Other than those specified in [Schedules C, C(1) and X]

Licence No:**TG/15/03/2017-23604**

Date: 12/04/2017

Licensing Authority

Assistant Director & Licensing Authority
Drugs Control Administration
Medchal District,Telangana State

Assistant Director & Licensing Authority Drugs Control Administration District

* Delete if not applicable

Conditions Of Licence

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall comply with the provisions of the Drugs and Cosmetics Act,1940 and the Rules thereunder for the time being in force.
3. The licensee shall report to the licencing authority any change in the qualified staff incharge within one month of such change
4. No drug shall be sold unless such drug is purchased under cash or Credit memo from a duly licensed dealer or a duly licensed manufacturer.
5. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence 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,a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution

Firm_no	Circle_code = Circle_Name	Area_code = Area_Name	
273639	15=Rangareddy_cir1	03=Jeedimetla (sales)	NIC

GRANT OF LICENCE

GOVERNMENT OF TELANGANA

DRUGS CONTROL ADMINISTRATION

FORM 21

[See Rule 61 (2)]

Partners/Directors/Gpa/As
DR.CH.BHADRAREDDY,PRESIDENT, ,Society

Registered Pharmacist
B.CHANDANA and REGNO=60522/A1

LICENCE TO SELL,STOCK OR EXHIBIT[OR OFFER]FOR SALEOR DISTRIBUTE DRUGS BY RETAIL DRUGS SPECIFIED IN SCHEDULES C AND C(1)[EXCLUDING THOSE SPECIFIED IN SCH.X].

1.DR.CH.BHADRAREDDY,PRESIDENT, , CH.KALPANA.VICE PRESIDENT, , CH.MAHENDER REDDY,SECRETARY, , P.VASANTHA,JOINT-SECRETARY , P.ANJI REDDY, TREASURER, , CH.SHALINI, ORG-SECRETARY, , CH.PREETHIREDDY,JOINT-SECRETARY Society **of**

M/S.MALLAREDDY INSTITUTE OF DENTAL SCIENCES PHARMACY (A UNIT OF CHANDRAMMA EDUCATIONAL SOCIETY)

Is here by licenced to sell, stock or exhibit [or offer] for sale or distribute by retail the following categories of Drugs specified in Schedules C and C(1)[Excluding those specified in sch.X] to the Drugs and Cosmetics Rules 1945 and to operate a pharmacy on the premises situated at **SY NO.46.47.48,GROUND FLOOR,SURAM X ROAD,,JEEDIMETLA,QUTHBULLAPUR MANDAL,,**
MEDCHAL DIST, Rangareddy_cir1 (Dist) subject to the conditions specified below and to the provisions of the Drugs and Cosmetics Act, 1940 and the rules thereunder.

2. The licences shall be in force From **10/04/2017** TO **09/04/2022**

3. Name(s) of qualified person(s) in charge:
B.CHANDANA , RegNo. 60522/A1 , DT:02/01/2010(D.O.J: 10/04/2017)

4. Categories of drugs : Drugs Specified In Schedules C And C(1)[Excluding Those Specified In Sch.X].

Licence No:**TG/15/03/2017-23605**

Date: 12/04/2017

Licensing Authority
Assistant Director & Licensing Authority
Drugs Control Administration
Medchal District,Telangana State

Conditions Of Licence

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.
2. The licensee shall report to the licencing authority any change in the qualified staff incharge within one month of such change.
3. [**] Omitted by GSR 17(E),dt. 7-1-1986 (w.e.f. 7-1-1986).
4. If the licensee wants to sell stock or exhibit for sale or distribute, during the currency of the licence,additiol categories of Drugs listed in [Schedules C and C(1)[Excluding those specified in Sch: X] but not included in this licence, you should apply to the licensing authority for the necessary permission. This licence will be deemed to extend to the categories of drugs in respect of which such permission is given.This permission shall be endorsed on the licence by the licensing authority.
5. No drug shall be sold unless such drug is purchased under cash or Credit memo from a duly licensed dealer or a duly licensed manufacturer.
6. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from date on which the change takes place unless ,in the meantime, a fresh licence has been taken from the Licensing Authority in the name of thefirm with the changed constitution

Firm_No	Circle_code = Circle_Name	Area_code = Area_Name	
273639	15=Rangareddy_cir1	03=Jeedimetla (sales)	N I C

DRUGS CONTROL ADMINISTRATION
Government of Telangana

L.Dis.No. NDPS-2/TS/MDL/2023-100974 Dated:02/01/2026

To,
M/s MALLAREDDY MEDICALS (O.T PHARMACY) ,
Address **SY.NO.138, D.NO.2-18/2, C/O.MALLAREDDY HOSPITAL, SURARAM MAIN ROAD, SURARAM, QUTHBULLAPUR, MEDCHAL, SURARAM VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE,INDIA**

Sir,

Sub:T.S.N.D.P.S.Rules,1986 – Renewal of NDPS License in Form NDPS-2 – Regarding.

Ref: Your application dated:

-x-x-x-

With reference to your application cited, I am herewith forwarding the License in N.D.P.S-2 bearing No.NDPS-2 **TS/MDL/2023-100974,dated:24/02/2023,** duly renewed for the period from: **01/01/2026** is valid upto **31/12/2026** for the quantities of the Narcotic Drugs and Psychotropic Substances mentioned in the License enclosed.

Digitally Signed By
G SREENIVAS
Deputy Director and Licensing Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:02-01-2026 14:03:57 PM

This Document is Digitally Signed. Signature is not required

FORM N.D.P.S-II
[Rule 93(1)]
LICENCE FOR THE MANUFACTURE POSSESSION AND OTHERWISE THAN ON PRESCRITION OF MANUFACTURED DRUGS BY DEALERS

Licence No.NDPS-2/:TS/MDL/2023-100974 Dated:02/01/2026

Licence is hereby granted to of M/s MALLAREDDY MEDICALS (O.T PHARMACY) of following the profession of Pharmacy at SY.NO.138, D.NO.2-18/2, C/O.MALLAREDDY HOSPITAL, SURARAM MAIN ROAD, SURARAM, QUTHBULLAPUR, MEDCHAL, SURARAM VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE,INDIA(hereinafter called the Licensee) authorizing him under and subject to the provisions of the Narcotics Drugs and Psychotropic Act, 1985 and the rules made thereunder to possess and sell or dispense, or prescription only, manufactured drugs at his shop/Dispensary situated at SY.NO.138, D.NO.2-18/2, C/O.MALLAREDDY HOSPITAL, SURARAM MAIN ROAD, SURARAM, QUTHBULLAPUR, MEDCHAL, SURARAM VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE,INDIA during the period commencing on 01/01/2026 and ending on 31/12/2026 on payment of a fee of Rs.50/- (in words Rupees Fifty only) and subject to the conditions hereinafter mentioned viz:-

1. The licensee shall purchase all manufactured drugs to be sold or dispensed under this license from a dealer in manufactured drugs licensed under the Telangana State NDPS Rules,1986 or under the corresponding rules for the time being in force in any part of India, or in accordance with condition. He shall not receive or have in his possession any manufactured, drugs which are not specified in this condition or which have been obtained otherwise than as permitted under this condition nor shall be possess then in quantities exceeding those specified below:-

(a) Coca derivative containing in the aggregate more than of Cocaine, -N.A.
(b) Opium derivatives containing in the aggregate more than – of Morphine, Diacetylmorphine or both – N.A.
(c) Medical hemp exceeding – in the case of extract and in the case of tinctures –N.A.

S.No	Drug Name	Strength	Pack Size	Quantity
1	Fentanly Citrate Inj	50 mcg/ml	10 ml	2000 amps
2	Fentanly Citrate Inj	50 mcg/ml	2 ml	10000 amps
3	Fentanyl Transdermal Patches	25 mcg/hr		500 patches
4	Fentanyl Transdermal Patches	50 mcg/hr		500 patches
5	Morphine Sulphate Inj	10 mg/ml	1 ml	2000 amps
6	Morphine sulphate Tabs	10 mg		5000 tablets

Digitally Signed By
G SREENIVAS
Deputy Director and Licensing Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:02-01-2026 14:03:57 PM

In the case of preparations and admixtures of coca derivatives and opium derivatives, the limit shall be fixed with reference to the Cocaine and Morphine contents, and not with reference to the quantity or bulk of the preparation, and the bottles, phials, packages other containers of these preparations or labels affixed to them shall plainly exhibit the actual quantity of the manufactured drugs present in each container or sufficient particulars to admit of the ready calculation of such quantity.

2.(a) The Licensee unless he is a Registered Medical Practitioner shall not keep, store, sell or dispense manufacture drugs in any place except in his dispensary prescribed above.

(b)If the Licensee, is a Registered Medical Practitioner, he may carry with him, from place to place manufactured drugs in quantities not exceeding these specified in condition 1 above.

3. The Licensee shall be responsible for the act and omissions of every person, appointed to officiate for him in carrying on the business of the said dispensary, and of all his servants as if the said acts and omissions were his own.

4.(1) The licensee shall not sell or dispense manufactured drugs except on a bonafied prescription given by himself, if he is a Registered Medical Practitioner or by any other Registered Medical Practitioner nor in longer quantity nor to any other person than may be specified in the prescription, provided the prescription is not given for the use of the prescribed himself.

(2) A prescription for the supply of manufactured drugs must comply with the following conditions:-

(a) The prescription shall be in writing and shall be dated and signed by a Registered Medical Practitioner with his full name and qualification and address and shall also specify the name and address of the person to whom it is given and the total quantity of the drug to be supplied thereof. If the drug to be supplied be coca derivatives the quantity should not contain more then 389 milligrams of cocaine, provided that the licensing authority may be special order authorize the supply of larger quantity considering the circumstances of the particular case.

(b) The prescription shall not be given for the use of the prescriber himself.

(c) A prescription given by a Registered Dentist shall be only for the purpose of dental treatment of and shall be marked "For Local Dental Treatment Only" and

(d) A prescription given by an Veterinary Surgeon shall be only for the purpose of treatment animals and shall be marked "For Animal Treatment only".

(3) When coca derivatives are to be sold or dispensed, the licensee shall see that the prescription is marked with the words "Not to be repeated" and shall not supply coca derivatives more than once on the same prescription except in pursuance of fresh directions only endorsed on the prescription by the approved practitioner by when it was originally issued and signed with his name in full and dated. Except under a special order made by the commissioner under rule of the Narcotic Drugs and Psychotropic Substances Rules the quantity so sold or dispensed at one time or to one the same person in the aggregate on any one day shall not contain more than 389 milligrams of Cocaine.

(4) Where Opium derivatives or medical hemp are to be sold or dispensed:

(a) If the prescription does not bear a subscription by a Registered Medical Practitioner stating that it is to be repeated and at what interval of time it is to be repeated and how may times it is to repeated, the Licensee shall sell the drugs once only on such prescription, and shall retain the prescription provided that the he shall first worn the person presenting the prescription that, unless it bears such a superscription as aforesaid, it will be retained.

(b) If the prescription bears a superscription as aforesaid, and it appears that Opium derivatives or medical hemp have already been sold on the prescription six times, or such number of times as the prescription is required to be repeated, or that the interval specified in the superscription has been elapsed since the prescription was lost dispensed, he shall not sell the drugs on such prescription unless it is further superscribed in that behalf by a Registered Medical Practitioner.

(5) The licensee shall mark on every prescription dispensed by him his name, the address of the premises at which and the date on which it was dispensed. In the case of every preparation made upon a prescription which contains manufactured drugs, the bottle or other receiptable or the wrapper or other covering in which such preparation is enclosed shall bear clearly marked upon it the amount and percentage of cocaine or morphine or diacetylmorphine or medicinal hamp contained in such preparation; provided that if the preparation in the form of uniformly divided dosal units, e.g.,pills, powders, tablets, capsules etc., it shall be sufficient if the bottle or other receiptable of the wrapper of other covering in which such preparation is enclosed bears clearly marked upon it the amount and percentage of cocaine of morphine contained in each such dosal unit.

(6) Where the prescription has to be returned to the person who presents it, the Licensee shall, on the first sale thereon, take and keep a copy of it, and on the occasion of each subsequent sale, note thereon the date of the sale and also sign and seal it.

(7) The Licensee may import, export or transport manufactured drugs by rail or inland post subject to the following conditions:-

(a) The parcel of manufactured drug when sent by a post shall be sent by registered parcel.

(aa) The parcel, whether sent by rail or by post, shall be insured.

(b) The parcel shall be covered by an authorization issued by competent authority at the place to which the parcel is addressed;

(c) The parcel shall be accompanied by a declaration showing the names of the consigner and the consignee, the contents, of the parcel in detail, the number and date of the authorization covering the import, export or transport, as the case may be, and the number of the licence if any, held by the consigner and the consignee.

The Licensee shall file and preserve for one year all prescriptions upon which manufactured drugs have been sold or dispensed by him, and shall produce such prescription alongwith this Licence and any manufactured drug that may be in his possession for inspection on demand by the Licensing Authority duly authorized by him.

The Licensee shall maintain a register in such form as may be approved by the Licensing Authority, wherein he shall from time to time record, in respect of the manufactured drugs dispensed by him, the full names and address of the Registered Medical Practitioners prescribing the drugs and of the persons for whom they are prescribed. The Licensee shall similarly record in the said register a true account of the kind and quantity of the manufactured drugs dispensed and the balance held by him in stock. The Licensee shall, before the seventh day of each calendar month, furnish to the Commissioner and Licensing Authority or such other officer as he may appoint in this behalf a copy of the entries made by him in the register during the proceeding calendar month

(8)(1)This licence may be cancelled or suspended by the Licensing Authority at any time;

(a) for non-payment of duty or fee payable by the Licensee.

(b) for default or violation by himself or by any servant or person acting on his behalf of any of the conditions specified in the licence or of the provisions of the Telangana State N.D.P.S.Rules, 1986;

(c) if the Licence, be convicted of any offence under the Narcotic Drugs and Psychotropic Substances Act, 1985 or under the law for the time being in force relating to excise revenue or of a breach of the peace or of any other criminal offence during the currency of the licence;

(d) if the Licensee infringes any of the conditions imposed on him by the Narcotic Drugs and Psychotropic Substances Act, 1985 or by the rules in force thereunder.

(e) After giving the licensee, 15 days notice, or if the licensee, desires to surrender his license, within 15 days from the receipt of such notice from him;

(2) When such licence is cancelled, suspended of surrendered, the licensee shall forthwith made over to the Licensing Authority or such other Officer as he may appoint, the Licence together with all the manufactured drugs in his possession.

(9) The Licensee shall be bound to purchase in such quantity not exceeding that which he is likely to sell in two months and at such rates as the Licensing Authority may direct, any manufactured drugs, that may be delivered to the Licensing Authority by any other Licensee who licence has expired or has been cancelled or suspended.

(10) All preparations containing not more than 0.1% of cocaine or 0.2% of Morphine and any preparation which the Central Government may by notification in the Gazette of India, made in pursuance of finding under article 8 of the Geneva Convention declare not be a manufactured drug, may be imported, exported, transported, possessed and sold without restriction.

Granted this on Date:02/01/2026

Digitally Signed By
G SREENIVAS
Deputy Director and Licensing Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:02-01-2026 14:03:57 PM

This Document is Digitally Signed. Signature is not required

DRUGS CONTROL ADMINISTRATION
Government of Telangana

L.Dis.No. NDPS-2/TS/MDL/2023-100974 Dated:02/01/2026

To,
M/s MALLAREDDY MEDICALS (O.T PHARMACY) ,
Address **SY.NO.138, D.NO.2-18/2, C/O.MALLAREDDY HOSPITAL, SURARAM MAIN ROAD, SURARAM, QUTHBULLAPUR, MEDCHAL, SURARAM VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE,INDIA**

Sir,

Sub:T.S.N.D.P.S.Rules,1986 – Renewal of NDPS License in Form NDPS-2 – Regarding.

Ref: Your application dated:

-x-x-x-

With reference to your application cited, I am herewith forwarding the License in N.D.P.S-2 bearing No.NDPS-2 **TS/MDL/2023-100974,dated:24/02/2023,** duly renewed for the period from: **01/01/2026** is valid upto **31/12/2026** for the quantities of the Narcotic Drugs and Psychotropic Substances mentioned in the License enclosed.

Digitally Signed By
G SREENIVAS
Deputy Director and Licensing Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:02-01-2026 14:03:57 PM

This Document is Digitally Signed. Signature is not required

FORM N.D.P.S-II
[Rule 93(1)]
LICENCE FOR THE MANUFACTURE POSSESSION AND OTHERWISE THAN ON PRESCRITION OF MANUFACTURED DRUGS BY DEALERS

Licence No.NDPS-2/:TS/MDL/2023-100974 Dated:02/01/2026

Licence is hereby granted to of M/s MALLAREDDY MEDICALS (O.T PHARMACY) of following the profession of Pharmacy at SY.NO.138, D.NO.2-18/2, C/O.MALLAREDDY HOSPITAL, SURARAM MAIN ROAD, SURARAM, QUTHBULLAPUR, MEDCHAL, SURARAM VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE,INDIA(hereinafter called the Licensee) authorizing him under and subject to the provisions of the Narcotics Drugs and Psychotropic Act, 1985 and the rules made thereunder to possess and sell or dispense, or prescription only, manufactured drugs at his shop/Dispensary situated at SY.NO.138, D.NO.2-18/2, C/O.MALLAREDDY HOSPITAL, SURARAM MAIN ROAD, SURARAM, QUTHBULLAPUR, MEDCHAL, SURARAM VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500055,TELANGANA STATE,INDIA during the period commencing on 01/01/2026 and ending on 31/12/2026 on payment of a fee of Rs.50/- (in words Rupees Fifty only) and subject to the conditions hereinafter mentioned viz:-

1. The licensee shall purchase all manufactured drugs to be sold or dispensed under this license from a dealer in manufactured drugs licensed under the Telangana State NDPS Rules,1986 or under the corresponding rules for the time being in force in any part of India, or in accordance with condition. He shall not receive or have in his possession any manufactured, drugs which are not specified in this condition or which have been obtained otherwise than as permitted under this condition nor shall be possess then in quantities exceeding those specified below:-

(a) Coca derivative containing in the aggregate more than of Cocaine, -N.A.
(b) Opium derivatives containing in the aggregate more than – of Morphine, Diacetylmorphine or both – N.A.
(c) Medical hemp exceeding – in the case of extract and in the case of tinctures –N.A.

S.No	Drug Name	Strength	Pack Size	Quantity
1	Fentanly Citrate Inj	50 mcg/ml	10 ml	2000 amps
2	Fentanly Citrate Inj	50 mcg/ml	2 ml	10000 amps
3	Fentanyl Transdermal Patches	25 mcg/hr		500 patches
4	Fentanyl Transdermal Patches	50 mcg/hr		500 patches
5	Morphine Sulphate Inj	10 mg/ml	1 ml	2000 amps
6	Morphine sulphate Tabs	10 mg		5000 tablets

Digitally Signed By
G SREENIVAS
Deputy Director and Licensing Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:02-01-2026 14:03:57 PM

In the case of preparations and admixtures of coca derivatives and opium derivatives, the limit shall be fixed with reference to the Cocaine and Morphine contents, and not with reference to the quantity or bulk of the preparation, and the bottles, phials, packages other containers of these preparations or labels affixed to them shall plainly exhibit the actual quantity of the manufactured drugs present in each container or sufficient particulars to admit of the ready calculation of such quantity.

2.(a) The Licensee unless he is a Registered Medical Practitioner shall not keep, store, sell or dispense manufacture drugs in any place except in his dispensary prescribed above.

(b)If the Licensee, is a Registered Medical Practitioner, he may carry with him, from place to place manufactured drugs in quantities not exceeding these specified in condition 1 above.

3. The Licensee shall be responsible for the act and omissions of every person, appointed to officiate for him in carrying on the business of the said dispensary, and of all his servants as if the said acts and omissions were his own.

4.(1) The licensee shall not sell or dispense manufactured drugs except on a bonafied prescription given by himself, if he is a Registered Medical Practitioner or by any other Registered Medical Practitioner nor in longer quantity nor to any other person than may be specified in the prescription, provided the prescription is not given for the use of the prescribed himself.

(2) A prescription for the supply of manufactured drugs must comply with the following conditions:-

(a) The prescription shall be in writing and shall be dated and signed by a Registered Medical Practitioner with his full name and qualification and address and shall also specify the name and address of the person to whom it is given and the total quantity of the drug to be supplied thereof. If the drug to be supplied be coca derivatives the quantity should not contain more then 389 milligrams of cocaine, provided that the licensing authority may be special order authorize the supply of larger quantity considering the circumstances of the particular case.

(b) The prescription shall not be given for the use of the prescriber himself.

(c) A prescription given by a Registered Dentist shall be only for the purpose of dental treatment of and shall be marked "For Local Dental Treatment Only" and

(d) A prescription given by an Veterinary Surgeon shall be only for the purpose of treatment animals and shall be marked "For Animal Treatment only".

(3) When coca derivatives are to be sold or dispensed, the licensee shall see that the prescription is marked with the words "Not to be repeated" and shall not supply coca derivatives more than once on the same prescription except in pursuance of fresh directions only endorsed on the prescription by the approved practitioner by when it was originally issued and signed with his name in full and dated. Except under a special order made by the commissioner under rule of the Narcotic Drugs and Psychotropic Substances Rules the quantity so sold or dispensed at one time or to one the same person in the aggregate on any one day shall not contain more than 389 milligrams of Cocaine.

(4) Where Opium derivatives or medical hemp are to be sold or dispensed:

(a) If the prescription does not bear a subscription by a Registered Medical Practitioner stating that it is to be repeated and at what interval of time it is to be repeated and how may times it is to repeated, the Licensee shall sell the drugs once only on such prescription, and shall retain the prescription provided that the he shall first worn the person presenting the prescription that, unless it bears such a superscription as aforesaid, it will be retained.

(b) If the prescription bears a superscription as aforesaid, and it appears that Opium derivatives or medical hemp have already been sold on the prescription six times, or such number of times as the prescription is required to be repeated, or that the interval specified in the superscription has been elapsed since the prescription was lost dispensed, he shall not sell the drugs on such prescription unless it is further superscribed in that behalf by a Registered Medical Practitioner.

(5) The licensee shall mark on every prescription dispensed by him his name, the address of the premises at which and the date on which it was dispensed. In the case of every preparation made upon a prescription which contains manufactured drugs, the bottle or other receiptable or the wrapper or other covering in which such preparation is enclosed shall bear clearly marked upon it the amount and percentage of cocaine or morphine or diacetylmorphine or medicinal hamp contained in such preparation; provided that if the preparation in the form of uniformly divided dosal units, e.g.,pills, powders, tablets, capsules etc., it shall be sufficient if the bottle or other receiptable of the wrapper of other covering in which such preparation is enclosed bears clearly marked upon it the amount and percentage of cocaine of morphine contained in each such dosal unit.

(6) Where the prescription has to be returned to the person who presents it, the Licensee shall, on the first sale thereon, take and keep a copy of it, and on the occasion of each subsequent sale, note thereon the date of the sale and also sign and seal it.

(7) The Licensee may import, export or transport manufactured drugs by rail or inland post subject to the following conditions:-

(a) The parcel of manufactured drug when sent by a post shall be sent by registered parcel.

(aa) The parcel, whether sent by rail or by post, shall be insured.

(b) The parcel shall be covered by an authorization issued by competent authority at the place to which the parcel is addressed;

(c) The parcel shall be accompanied by a declaration showing the names of the consigner and the consignee, the contents, of the parcel in detail, the number and date of the authorization covering the import, export or transport, as the case may be, and the number of the licence if any, held by the consigner and the consignee.

The Licensee shall file and preserve for one year all prescriptions upon which manufactured drugs have been sold or dispensed by him, and shall produce such prescription alongwith this Licence and any manufactured drug that may be in his possession for inspection on demand by the Licensing Authority duly authorized by him.

The Licensee shall maintain a register in such form as may be approved by the Licensing Authority, wherein he shall from time to time record, in respect of the manufactured drugs dispensed by him, the full names and address of the Registered Medical Practitioners prescribing the drugs and of the persons for whom they are prescribed. The Licensee shall similarly record in the said register a true account of the kind and quantity of the manufactured drugs dispensed and the balance held by him in stock. The Licensee shall, before the seventh day of each calendar month, furnish to the Commissioner and Licensing Authority or such other officer as he may appoint in this behalf a copy of the entries made by him in the register during the proceeding calendar month

(8)(1)This licence may be cancelled or suspended by the Licensing Authority at any time;

(a) for non-payment of duty or fee payable by the Licensee.

(b) for default or violation by himself or by any servant or person acting on his behalf of any of the conditions specified in the licence or of the provisions of the Telangana State N.D.P.S.Rules, 1986;

(c) if the Licence, be convicted of any offence under the Narcotic Drugs and Psychotropic Substances Act, 1985 or under the law for the time being in force relating to excise revenue or of a breach of the peace or of any other criminal offence during the currency of the licence;

(d) if the Licensee infringes any of the conditions imposed on him by the Narcotic Drugs and Psychotropic Substances Act, 1985 or by the rules in force thereunder.

(e) After giving the licensee, 15 days notice, or if the licensee, desires to surrender his license, within 15 days from the receipt of such notice from him;

(2) When such licence is cancelled, suspended of surrendered, the licensee shall forthwith made over to the Licensing Authority or such other Officer as he may appoint, the Licence together with all the manufactured drugs in his possession.

(9) The Licensee shall be bound to purchase in such quantity not exceeding that which he is likely to sell in two months and at such rates as the Licensing Authority may direct, any manufactured drugs, that may be delivered to the Licensing Authority by any other Licensee who licence has expired or has been cancelled or suspended.

(10) All preparations containing not more than 0.1% of cocaine or 0.2% of Morphine and any preparation which the Central Government may by notification in the Gazette of India, made in pursuance of finding under article 8 of the Geneva Convention declare not be a manufactured drug, may be imported, exported, transported, possessed and sold without restriction.

Granted this on Date:02/01/2026

Digitally Signed By
G SREENIVAS
Deputy Director and Licensing Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:02-01-2026 14:03:57 PM

This Document is Digitally Signed. Signature is not required