



DRUGS CONTROL ADMINISTRATION
Government of Telangana



LICENCE RETENTION CERTIFICATE

Dated:17/03/2025

This is to certify that the Licence of the firm **M/s HORIZON BIOLABS PRIVATE LIMITED** situated at **SY.NO.374,PLOT. NO.61/1, FIRST AND SECOND FLOORS,CIE,GANDHINAGAR EXPANSION, BALANAGAR,QUTHBULLAPUR MANDAL,MEDCHAL-MALKAJGIRI DISTRICT,HYDERABAD, TELANGANA-500037, TELANGANA,INDIA,BALANAGAR(VILLAGE), BALANAGAR(MANDAL), MEDCHAL - MALKAJGIRI(DIST.),TELANGANA STATE,INDIA** bearing Licence No. **3/ML/TS/2020/AL/G** in the statutory **Form 37** granted/ renewed on **27/04/2025** whose validity would get expired by **26/04/2030** , is hereby permitted to be retained with the extended validity upto **26/04/2030** .with the existing Constitution ,Technical Staff and Total Products as on date.

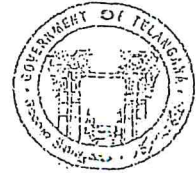
Retention fees/ with penalty for an amount of Rs. **7500.00** vide challan no:**6404544306** and Bank Transaction No.:**2039149470** has been paid on **17/03/2025** by the licensee in accordance with provisions of Rule 63(1) of the Drugs and Cosmetics Rules, 1945.



This is Electronically Generated Receipt Signature is not required



DRUGS CONTROL ADMINISTRATION
Government of Telangana



L.Dis.No.35441/ODLS/A3/2020.

Dated 28-04-2020.

To

M/s. Horizon BioLabs Private Limited,
Sy.No. 374, Plot.No. 61/1, First and Second Floors,
CIE., Gandhi Nagar Expansion, Balanagar,
Quthbullapur Mandal, Medchal-Malkajgiri District – 500 037,
Telangana, INDIA.

Sirs,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder – Grant of Approval in Form – 37 – Regarding.

Ref: Your application dated 12.02.2020.

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With reference to your application cited, I am herewith forwarding the Certificate of approval in Form – 37 bearing No. 3/ML/TS/2020/AL/G, Dated 27.04.2020 valid upto 26.04.2025 for the following Drugs and Cosmetics for carrying out the Tests.

(a) DRUGS OTHER THAN THOSE SPECIFIED IN SCHEDULE C AND C(1) AND ALSO EXCLUDING HOMEOPATHIC DRUGS.

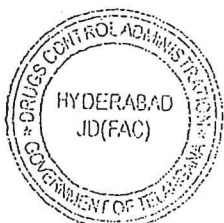
1. Drugs requiring the use of Ultraviolet / Infra Red or Chromatography.
2. Disinfectants

(b). DRUGS SPECIFIED IN SCHEDULE C AND C(1)

1. Sera, Vaccines, Antigens, Toxins, Antitoxins, Toxoids, Bacteriophages and Similar Immunological Products
2. Antibiotics
3. Vitamins
4. Parenteral Preparations
5. Drugs Requiring Microbiological Tests
6. Drugs requiring the use of Ultraviolet / Infra Red / Spectrophotometer or Chromatography.

(c) Cosmetics.

Yours faithfully,



B. Venkateshwarlu
28/04/2020
Dr. B. VENKTESHWARLU
JOINT DIRECTOR(FAC)

LICENSING & CONTROLLING AUTHORITY