Certificate of Compliance

This is to certify that the
general requirements for the competence and consistent operation of reference material producers
of

LAUREL PHARMA LABS

at

PLOT NO. 35, IDA BALANAGAR, HYDERABAD, TELANGANA-500037, INDIA.

has been independently assessed and is
compliant with the requirements of:

ISO 17034:2016

While demonstrating technical competence in the field of
Reference Material Producer
Refer to the accompanying Scope of accreditation for information regarding the type of Materials to which this accreditation applies

AS PER APPENDIX-I

Certificate Number: UQ - 2026033017

Validity of this certificate can be verified at www.ukcertifications.org.uk/verify

Date of Certification	30th March 2026
1st Surveillance Audit Due	29th March 2027
2nd Surveillance Audit Due	29th March 2028
Certificate Expiry	29th March 2029

Authorised Signatory

Certificate of Compliance Appendix-I

Appendix to Certificate no.:- UQ-2026033017

Manufacturer :- LAUREL PHARMA LABS

Address :- PLOT NO. 35, IDA BALANAGAR, HYDERABAD, TELANGANA-500037, INDIA.

Category of Reference Material	Class or Type of Reference Material Produced (Include Range Where Applicable)	Methods or Techniques used in the RMP Laboratory (if Applicable)
Certified Reference Material (CRM) and Reference Materials (RM) :	• Organic Reference Materials • Active Pharmaceutical Ingredients and its process intermediates, impurities, metabolites, • labelled compounds. • Speciality chemicals & Fine chemicals	Pharmacopoeial or In-house methods • Chromatography HPLC, GC • Mass Spectrometry LC/MS. GC/MS • Nuclear Magnetic Resonance Spectrometry H1NMR, C13NMR • IR Spectroscopy • Thermo Gravimetric Analysis • Elemental Analysis • Loss on Drying, Moisture Content, PH, Chemical Assay, Sulphated ash.

Authorised Signatory