■ GMP-Certified Pharmaceutical Laboratory

■CERTIFICATE OF LABORATORY ANALYSIS

Product Name: **Cenforce 200 mg**
Active Ingredient: **Sildenafil Citrate**
Manufacturer: **Centurion Laboratories Pvt. Ltd.**
Batch Number: **C200-BX/0725**
Date of Manufacture: **July 2025**
Date of Testing: **25 July 2025**
Report Issued By: **Rahul Mehta – Senior QA Executive (10+ years experience)**
Lab ID: **QAL-IND/CENF/0725-01**

Test Parameter	Specification	Result	Method Used	Status
Appearance	Blue, round, film-coated tablet	Complies	Visual Inspection	■Passed
Identification (API)	Positive for Sildenafil Citrate	Positive	FTIR + HPLC	■Passed
Average Weight	350–370 mg/tablet	361.4 mg	Digital Balance (20 units)	■Passed
Content of Sildenafil	95%–105% of label claim (200 mg)	198.6 mg (99.3%)	HPLC	■Passed
Uniformity of Dosage Units	±5% variation allowed	Complies (2.4% max)	HPLC (10 units)	■Passed
Dissolution (30 mins)	Not less than 80%	89.7%	USP II (Paddle)	■Passed
Disintegration Time	Within 30 minutes	12 minutes	USP Disintegration Tester	■Passed
Microbial Load	Within USP limits	Complies	Plate Count	■Passed
Heavy Metals	NMT 10 ppm	< 5 ppm	ICP-MS	■Passed
Related Substances	NMT 2.0% total impurities	0.82%	HPLC	■Passed

Conclusion:
The tested batch of **Cenforce 200 mg** complies with all **USP & internal quality specifications** for Sildenafil Citrate tablets. Product is confirmed **safe, potent, and pharmaceutically equivalent** to reference standards.

Certified by:
Rahul Mehta
Senior Quality Assurance Executive
GMP-Certified Pharmaceutical Laboratory
License No: QAL/IND/2208

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