

Nebivolol 5mg

Pharmaceutical Compound • Research Use Only

Report ID	PHX-B1012-26	Test Date	7 April 2026
Product	Nebivolol 5mg	Laboratory	Puralytix (Independent Third-Party)
Category	Pharmaceutical Compound	Test Method	HPLC-UV / LC-MS; ICP-MS (heavy metals)
Sample Form	Oral solid (tablet / capsule)	Label Claim (total)	5mg
Quantified (total)	5.06mg	HPLC Purity	99.5 %

COMPOUND ANALYSIS

Active compound	Label claim	Quantified	Result
Nebivolol	5mg	5.06mg	Pass

HEAVY METAL SCREEN (ICP-MS)

Lead (Pb)	Mercury (Hg)	Arsenic (As)	Cadmium (Cd)
ND (≤ 0.5 ppm)	ND (≤ 0.1 ppm)	ND (≤ 0.5 ppm)	ND (≤ 0.25 ppm)

ND = not detected at or above the stated method reporting limit (ppm).

RESULT

PASS — HPLC Purity 99.5% · Heavy metals: not detected

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