



Govt. Approved Testing Lab

RESULTS OF ANALYSIS

Approval No. WH (023) / 14

CERTIFICATE OF ANALYSIS

Swiss Federal Act on Medicinal Products and Medical Devices (MPA)

Submitted By	: Proton Pharma Kavaci Mah. Kovak Sok. No8/24, Kavacik Ticaret Merkezi, Istanbul	Report No.	GOHSP- 220412144
		Received On	15.01.2025
Sample	: Stanoxin 10mg	Mfg.Lic.No:	04.2019-127/22
Manufactured By	: Proton Pharma	Exp Date:	09/2027
Supplied By	: Proton Pharma	Batch	
Batch No.	: 01823	Size:	10 tablets
Mfg. Date	: N/A		

Description:	A white, flat-faced, bevel-edged tablet.		
Average weight:	N/A		
Identification:	Positive for Stanozolol		
Uniformity of content:	99.9%-101.1% of the average contains (Limit : 90% to 110%)		
Disintegration:	6-9min (Limit: NMT 40 min)		
Assay:	Each tablet, on average, contains:		
	Result	Claim	Limit
Stanozolol	10.7 mg	10mg	9-11mg

Remark:Based on the evaluation conducted by the undersigned, it is determined that the sample in question meets the required quality and/or quantity standards as outlined in the relevant regulations and guidelines. The details supporting this conclusion are provided below.

Date: 20-01-2025

Person In-charge



• The results provided are specific to the samples tested and the applicable parameters. No endorsement of the product is either implied or suggested.
• The test certificate, either in part or full, cannot be used as evidence in legal proceedings.
• Drug and cosmetic samples will be destroyed six months after testing, while other non-perishable samples will be discarded within two weeks from the certificate issuance date.
• The liability of our analytical division is strictly limited to the invoiced amount. Any samples not collected by us, unless otherwise specified, are excluded from this liability. The reported results are only applicable to the specific sample(s) tested.