CERTIFICATE OF ANALYSIS

BUPROWELL SR 150 TABLETS

kinorac

BATCH NUMBER
HT-24247A

A. R. NUMBER
FP240666

RELEASE DATE
23/11/2024

PART A: PRODUCT PARTICULARS

Product name	**BUPROWELL SR 150 tab**
Source document	**Certificate of Analysis of Finished Good**
Source reference	Drug & Cosmetic Act, 1940; Form 39 Rule 150-E(F), as stated in the source

PART B: TEST RESULTS

SR. NO.	TEST	RESULT	SPECIFICATION
5	Dissolution	Not provided	1 hr: NMT 40% 4 hr: NLT 40% 8 hr: NLT 60% 10 hr: NLT 70%
6	Uniformity of dosage unit	Min: 102.04% Max: 104.16% Avg: 102.91%	AV: NMT 15
7	Assay	Bupropion Hydrochloride: 102.6% (153.9 mg)	Each film-coated sustained-release tablet contains: Bupropion Hydrochloride USP 150 mg (90.0% to 110.0%) (135.0 mg to 165.0 mg)

CONCLUSION RECORDED IN SOURCE

The source report states that the sample complies with IP / BP / USP / In-house requirements and is of standard quality as defined in the Act and the rules made thereunder.