# CERTIFICATE OF ANALYSIS

## BUPROWELL SR 150 TABLETS

HIGH STREET PHARMA

**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<br>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%<br>4 hr: NLT 40%<br>8 hr: NLT 60%<br>10 hr: NLT 70% |
| 6 | Uniformity of dosage unit | Min: 102.04%<br>Max: 104.16%<br>Avg: 102.91% | AV: NMT 15 |
| 7 | Assay | Bupropion Hydrochloride:<br>102.6% (153.9 mg) | Each film-coated sustained-release tablet contains:<br>Bupropion Hydrochloride USP 150 mg<br>(90.0% to 110.0%)<br>(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.