

**Sample Name** Ondansetron Orally Disintegrating Tablets-IP. **Mfg Lic No** [REDACTED]  
**Submitted By** Quality Control Advisor. (MMDSL) Shillong. **Report No** [REDACTED]  
**Address** DHS LaitumkhrahNHM, EastKhasi Halls Meghalaya. **Receipt Date** 29-Apr-24  
**Mfg By** N.S. **Report Date** 15-May-24  
**Supplied By** N.S. **Ref No** N.S.  
**Batch No** MMDSL/QC-0207 **Mfg. Date** 02/2024 **Exp Date** 01/2026 **Batch Size** N.S. **Sample Qty** 60 TABS.

### TEST RESULTS

Date / Period of Performance of test 29/04/2024 to 15/05/2024.

Reference to protocol :- I.P-2022.

Description :- White round biconvex uncoated Orally Disintegrating tablets.

Identification (by HPLC) :- Complies.

Average weight :- 198.03mg.

Uniformity of content (by HPLC) :- Within limit.

Dissolution (by U.V) :- Complies (89.6% to 94.4%) Limit:- NLT 80%+ 5%

Related Substances (by HPLC) :- Complies.

Disintegration time :- Within 30 seconds (Limit:-NMT-30 seconds)

Assay (by HPLC) :- Each uncoated Orally Disintegrating tablet contains:-

| Contents of                                    | Obtd./Av.wt. | Claim  | LIMIT  |        | Method |
|--|--------------|--------|--------|--------|--------|
|  |              |        | Lower  | Upper  |        |
| Ondansetron Hydrochloride<br>Eq.to Ondansetron | :- 7.91mg    | 8.0mg. | 7.6mg. | 8.4mg. | IP.    |

NOTE:- SAMPLE CONSUMED IN TESTING.

Report: In Opinion of the undersigned, The sample as defined in the Act and the rules made there per IP.