**CERTIFICATE OF ANALYSIS**

**CERTIFICATE No:** 0

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| **PRODUCT** | ORNILORÂ® TABLETS | | REF. NO: NDQD201509289 |
| **DATE RECEIVED:** 22.09.2015 | **LABEL CLAIM:** | Each film coated tablet contains: Ornidazole 500 mg, Excipients q.s. | |
| **BATCH NO.:** UOR1501 | **PRESENTATION:** | White coloured, caplet shaped, biconvex faced tablets, single scored on one face and plain on the other, packed in aluminium foil blister strips of 10 tablets each and 3 such strips in a unit box. | |
| **MGF. DATE:** Feb.2015 | **MANUFACTURER:** | CORAL Laboratories Ltd. | |
| **EXP. DATE:**  Jan.2018 | **ADDRESS:** | Plot No. 27-28, Pharmacity, Selaqui,Dehradun, Uttarakhand-248 197, INDIA. | |
| **CLIENT REF NO:** | **CLIENT:** | LORDS Healthcare Limited P. O. Box 49397 - 00100, Nairobi, KENYA | |
| - | **TEST(S) REQUESTED:** | Uniformity of Weight, Identification, Dissolution, Assay | |

**RESULTS**

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| **TEST** | **METHOD** | **COMPEDIA** | **SPECIFICATION** | **DETERMINED** | **REMARKS** |
| **Uniformity of Weight** | Weight | BP 2016 Vol. V App XII C | NMT 2 tablets deviate by more than 5% from mean tablet weight | 3 Deviate [-5.1%; -5.1% & -5.2%] | **DOES NOT COMPLY** |
| **Identification** | HPLC | Adopted In-House Method | RT of the major peak in the assay sample preparation corresponds to that in the assay standard preparation | Super-imposable peak at RT 4.9 -/+ 10% min. present in both the assay sample and standard preparations | **COMPLIES** |
| **Dissolution** | HPLC | Adopted In-House Method | No less than 80% | 102.4% (RSD=5.2%; n=6) | **COMPLIES** |
| **Assay** | HPLC | Adopted In-House Method | 90.0 -110.0% | 100.1% (RSD=1.9%; n=9) | **COMPLIES** |

**CONCLUSION:** The product complies with the specifications for the tests performed.

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| ANALYST: | DR . L. KARANJA |  | DATE:29-02-2016 |
| ANALYST: | DR . N. MWAURA |  | DATE:29-02-2016 |
| ANALYST: | NICHOLAS MWAURA |  | DATE:03-01-2016 |
| DIRECTOR: | HEZEKIAH CHEPKWONY |  | DATE:03-01-2016 |