**CERTIFICATE OF ANALYSIS**

**CERTIFICATE No:**  CAN/2015-16/XYZ

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| **PRODUCT** | NEVIRAPINE TABLETS USP 200 mg | | REF. NO: NDQD201508156 |
| **DATE RECEIVED:** 06.08.2015 | **LABEL CLAIM:** | Each uncoated tablet contains: Nevirapine U.S.P. 200 mg. | |
| **BATCH NO.:** NE2014053-B | **PRESENTATION:** | White colored, caplet shaped, biconvex faced tablets, single scored on both faces and embossed "C" and "35" on either sides of the score on one face, packed in a white plastic multi-dose container carrying 60 tablets in a unit box. | |
| **MGF. DATE:** Jul. 2014 | **MANUFACTURER:** | AUROBINDO Pharma LImited. | |
| **EXP. DATE:** Jun. 2016 | **ADDRESS:** | Unit III, Survey No. 313, Bachupally Village, Quthubullapur Mandal, Ranga Reddy District (A.P.), INDIA. | |
| **CLIENT REF NO:** | **CLIENT:** | National AIDS and STI Control Programme P.O. BOX 19361 Nairobi | |
|  | **TEST(S) REQUESTED:** | Uniformity of Weight, Identification, Dissolution, Assay | |

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| **TEST** | **METHOD** | **COMPEDIA** | **SPECIFICATION** | **DETERMINED** | **REMARKS** |
| **Uniformity of Weight** | Weight | B.P. 2012 Vol. V App XII C | NMT 2 tablets deviate by more than 5% from mean weight | None Deviates | **COMPLIES** |
| **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 5.4 +/- 10% min. present in both the assay sample and standard preparations | **COMPLIES** |
| **Dissolution** | HPLC | Adopted In-House Method | No tablet less than 80% [n = 6] | 100.6% (RSD = 1.9%; n = 6) | **COMPLIES** |
| **Assay** | HPLC | Adopted In-House Method | 90.0 - 110.0% | 98.2% (RSD = 0.8%; n = 9) | **COMPLIES:** |

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

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| ANALYST: | DR. E. MUTUA |  | DATE:01-10-2015 |
| REVIEWER: | DR. G. WANG'ANG'A |  | DATE:19-10-2015 |
| DIRECTOR: | HEZEKIAH CHEPKWONY |  | DATE:11-05-2015 |