**CERTIFICATE OF ANALYSIS**

**CERTIFICATE No:** CAN/2015-16/581

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| **PRODUCT** | UNIVIR TABLETS | | REF. NO: NDQB201604854 |
| **DATE RECEIVED:** 07.04.2016 | **LABEL CLAIM:** | Each tablet contains: Acyclovir USP 400 mg. | |
| **BATCH NO.:** 620190 | **PRESENTATION:** | Off-white coloured, caplet shaped, biconvex faced tablet, embossed 'UCL' on one face and single scored on the other, packed in a blister strip of 10 tablets and 3 such strips in a unit box. | |
| **MGF. DATE:** Feb. 2016 | **MANUFACTURER:** | UNIVERSAL Corporation Ltd. | |
| **EXP. DATE:**  Jan. 2019 | **ADDRESS:** | Club Road, Plot No. 13777, P.O. Box 1748-00902, Kikuyu, KENYA. | |
| **CLIENT REF NO:** | **CLIENT:** | Kenya Medical Supplies Authority, P.O. Box 47715 - 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 | None Deviates | **COMPLIES** |
| **Identification** | HPLC | USP 38 NF 33 Page 2058 | RT of the Major Peak in the sample preparation corresponds to that in the standard preparation | Super-imposable peak at RT 6.6 -/+ 10% min. present in both the assay sample and standard preparations | **COMPLIES** |
| **Dissolution** | UV | USP 38 NF 33 Page 2058 | No Tablet Less than 90% [n=6] | 93.4% (RSD=1.5%;n=6) | **COMPLIES** |
| **Assay** | HPLC | USP 38 NF 33 Page 2058 | 90.0 - 110.0% | 100.2% (RSD=0.5%;n=9) | **COMPLIES** |

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

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| ANALYST: | DR. E. MUTUA |  | DATE:26-04-2016 |
| ANALYST: | DR. G. WANG'ANG'A |  | DATE:26-04-2016 |
| ANALYST: | EMMANUEL TANUI |  | DATE:04-27-2016 |
| DIRECTOR: | HEZEKIAH CHEPKWONY |  | DATE:05-09-2016 |