**CERTIFICATE OF ANALYSIS**

**CERTIFICATE No:** CAN/2015-16/440

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| **PRODUCT** | CETAPAR ORAL SUSPENSION | | REF. NO: NDQD201512644 |
| **DATE RECEIVED:** 28.12.2015 | **LABEL CLAIM:** | Each 5 mL contains: Paracetamol BP 125 mg. | |
| **BATCH NO.:** LE5129 | **PRESENTATION:** | Pink coloured suspension for oral administration contained in a 60 mL amber coloured plastic bottle with a metallic cap, packed along with a 10 mL graduated plastic measuring cup in a unit box. | |
| **MGF. DATE:** Aug. 2015 | **MANUFACTURER:** | ZEST Pharma | |
| **EXP. DATE:**  Jul. 2017 | **ADDRESS:** | Plot No. 275, Sector 'F', Sanwer Road, Indore-452015, INDIA | |
| **CLIENT REF NO:** | **CLIENT:** | LIBERIA MEDICINES AND HEALTH PRODUCTS REGULATORY AUTHORITY Vp Road,Old Road, Sinkor, Monrovia, LIBERIA | |
|  | **TEST(S) REQUESTED:** | Identification, Assay, Acidity/Alkalinity | |

**RESULTS**

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| **TEST** | **METHOD** | **COMPEDIA** | **SPECIFICATION** | **DETERMINED** | **REMARKS** |
| **Identification** | HPLC | USP 38 NF 33 Page 2008 | RT of the major peak in the sample preparation corresponds to that in the standard preparation | Super-imposable peak at RT 4.7 -/+ 10% min. present in both the assay sample and standard preparations | **COMPLIES** |
| **Assay** | HPLC | USP 38 NF 33 Page 2008 | 90.0 - 110.0% | 92.4% (RSD=0.3%; n=9) | **COMPLIES** |
| **Acidity/Alkalinity** | pH | USP 38 NF 33 Page 2008 | 4.0 - 6.9 | 6.0 | **COMPLIES** |

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

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| ANALYST: | MR. F. NAULA |  | DATE:01-12-2016 |
| ANALYST: | MR. D. MOENGA |  | DATE:29-02-2016 |
| REVIEWER: | DR. G. WANG'ANG'A |  | DATE:02-03-2016 |
| ANALYST: | NICHOLAS MWAURA |  | DATE:03-07-2016 |
| ANALYST: | NICHOLAS MWAURA |  | DATE:03-07-2016 |
| DIRECTOR: | HEZEKIAH CHEPKWONY |  | DATE:03-09-2016 |