**CERTIFICATE OF ANALYSIS**

**CERTIFICATE No:** CAN/2015-16/713

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| **PRODUCT** | MYLOVASC (TM)- 2.5 TABLETS | | REF. NO: NDQD201512635 |
| **DATE RECEIVED:** 15.12.2015 | **LABEL CLAIM:** | Each uncoated tablet contains: S-Amlodipine besilate equivalent to S-Amlodipine 2.5 mg | |
| **BATCH NO.:** 14SAMT 001 | **PRESENTATION:** | Off white coloured, circular shaped, biconvex faced tablets, plain on both sides, packed in a blister strip of 10 tablets and 3 such strips in a unit box | |
| **MGF. DATE:** Dec. 2014 | **MANUFACTURER:** | BAFNA Pharmaceuticals Ltd. | |
| **EXP. DATE:**  Nov. 2017 | **ADDRESS:** | 147, Madhavaram Redhills High Rd, Grantlyon Village,Chennai-600052, INDIA. | |
| **CLIENT REF NO:** | **CLIENT:** | PHARMA SPECIALITIES Ltd P.O. Box 49146- 00100 GPO, Nairobi | |
|  | **TEST(S) REQUESTED:** | Uniformity of Content, Identification, Dissolution, Assay | |

**RESULTS**

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| **TEST** | **METHOD** | **COMPEDIA** | **SPECIFICATION** | **DETERMINED** | **REMARKS** |
| **Uniformity of Content** | Weight | USP 38 NF 33 Page 2209 | L2: Acceptance Value of the 30 dosage units is < 15 (n=30) | AV=11 (n=30) | **COMPLIES** |
| **Identification** | HPLC | USP 38 NF 33 Page 2209 | RT of the component peak in the assay sample preparation corresponds to that in the assay standard preparation | Super-imposable peak at RT 10.1 -/+ 10% min. present in both the assay sample and standard preparations | **COMPLIES** |
| **Dissolution** | UV | USP 38 NF 33 Page 2209 | No tablet less than 80.0% (n=6) | 101.8% (RSD=5.9%, n=6) | **COMPLIES** |
| **Assay** | HPLC | USP 38 NF 33 Page 2209 | 90.0 - 110.0% | 90.9% (RSD=0.9%, n=6) | **COMPLIES** |

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

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| ANALYST: | DR. E. MUTUA |  | DATE:04-03-2016 |
| ANALYST: | DR. E. TANUI |  | DATE:14-03-2016 |
| DIRECTOR: | HEZEKIAH CHEPKWONY |  | DATE:06-13-2016 |