

CERTIFICATE OF ANALYSIS

STP Prime CCS Comp Cal Cartridge Lot Number 16354046 Ref Number 52427 Exp Date 2017-10-26

Date of Manufacture 2016-10-26

| Cal A L/N 163 | 00042 | Cal B L/N 16300034 | | Flush L/N 16316042 | | |
|-----------------|----------------|---------------------|-----------|--------------------|-------|-----------|
| S | Spec Pass/Fail | Spec | Pass/Fail | | Spec | Pass/Fail |
| pH 7. | .353 N/A | pH 6.837-6.8 | 343 Pass | pCO2 mmHg | 51.6 | N/A |
| pCO2 mmHg 2 | 24.6 N/A | Na mmol/L 68.0-69 | .0 Pass | pO2 mmHg | 101.8 | N/A |
| Na mmol/L 132. | 1-133.9 Pass | K mmol/L 9.92-10. | 08 Pass | | | |
| K mmol/L 3.9 | 7-4.03 Pass | CI mmol/L 45.6-46 | .4 Pass | | | |
| CI mmol/L 105.2 | 2-106.8 Pass | iCa mmol/L 2.17-2.2 | 23 Pass | | | |
| iCa mmol/L 1.08 | 8-1.12 Pass | Glu mg/dL 197-20 | 3 Pass | | | |
| Glu mg/dL 79 | 9-81 Pass | Lac mmol/L 9.5-10. | 5 Pass | | | |
| Lac mmol/L 1.8 | 8-2.2 Pass | | | | | |

Reference L/N 16300021

Spec Pass/Fail KCI 1.8-2.2 Pass

This certifies that this product was manufactured and tested at Nova Biomedical Corporation, Waltham MA 02454 U. S. A. in accordance with 21 CRF-Part 820, Quality System Regulation (QSR), Current Good Manufacturing Practice (cGMP) and EN ISO 13485:2012, Medical Devices, Quality Management Systems for in Vitro Diagnostic devices, and conforms to the indicated test specifications. All listed analytes are traceable to NIST SRM Materials.

Joyce ansara Approval

QC Supervisor

Title

1/10/2017 Date