A1C CONTROL LEVELS 1 & 2

Product Lot Number Expiration Date
NOD® A1c Control, Level 1 & 2 3220M002 November 2012

INTENDED USE
NOD® A1c Control is intended for use as quality control material to monitor the performance and precision of Hemoglobin A1c (HbA1c %) Immunoassay and HPLC test methods using protocols established in individual laboratories.

SUMMARY AND PRINCIPLE
Laboratories should run both quality control materials before any patient samples to ensure proper performance of instruments and reagents. NOD® A1c Control is provided at two levels – normal (L1) and abnormal (L2).

REAGENT
NOD® A1c Control is prepared from human whole blood to which stabilizers are added. The product is provided in liquid form for user convenience. No further dilution of the control solution is needed.

STORAGE AND STABILITY
Kits may be received thawed but should be stored frozen (-15°C to -25°C) immediately upon receipt. Unopened NOD® A1c Control is stable until the expiration date printed on the container when stored frozen.

Refrigerated OPEN OR CLOSED vials are stable for 180 days for use with immunoassay methods.

Refrigerated CLOSED OR CLOSED vials are stable for 30 days when used with HPLC methods.

(Walgreens, Little Clinic & MinuteClinic Sites Please Note: Commercial Mini refrigerator freezers do not freeze control materials to the temperature specified. Your controls should be treated as refrigerated immediately upon receipt. The control will expire 180 days from the date received when stored properly and tightly capped.)

Upon receiving the kit, mark the specific date refrigerated storage began on the vials; product is stable for 180 days when stored at 2-8°C in tightly closed containers. Aliquots made immediately from freshly open vials may be frozen one time and stored until expiration date printed on the container. Thawed aliquots cannot be refrozen. (Please Note) Dried red control residue in top of cap means vials are not being recapped properly.

PROCEDURE
NOD® A1c Control should be treated in the same manner as patient samples in accordance with instructions for testing determination method being used. No further dilution of the control solution is needed. Frozen control should be thawed at 2-8°C and mixed by gentle inversion several times prior to use. Refrigerated control should be mixed by gentle inversion prior to use. Do not shake vigorously. Do not warm up refrigerated control material before using and always clean any excess control material from dropper tip before tightly recapping a vial. NOD control vials should be used, wiped clean, recapped and back in the refrigerator in approximately 5 minutes or less to preserve refrigerated stability.

LIMITATIONS
Different values from those obtained with reagents available at the time of assay may be obtained as a result of changes in manufacturer’s reagents or lot-to-lot reagent variability. NOD® A1c Control should not be used past its expiration date or after improper handling. Microbial contamination will affect performance of this product.

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(REV 4-27-2011)
ANALYTE VALUES
In accordance with good laboratory practices, each laboratory should establish its own analyte means and acceptable performance ranges.

SPECIFIC PERFORMANCE CHARACTERISTICS
NOD® A1c Control is manufactured in accordance with industry guidelines and standards. To perform as intended, the control requires proper storage and handling as described in this package insert. Over time and repeated use the dropper tip can collect dried control debris if not wiped clean immediately after each use. If this debris is allowed to collect it can block your test assays collector channel and cause a low reading for the control material. If you are recording out of range low control values and a dropper vial cap shows evidence of dried red colored control material on the inside of the cap – Order a fresh control kit.

WARNING
Biological source material, treat as potentially infectious. All human plasma units used in manufacturing this product have been tested according to by FDA accepted methods and found non-reactive or negative for Hepatitis B Surface Antigen (HbsAg), HCV antibodies, and HIV-1/2 antibodies. This product may contain other human or animal source materials for which there are no approved tests and should be considered as potentially infectious for Hepatitis B (HBV), Hepatitis C (HCV), HIV-1, HIV-2, HTLV-I, HTLV-II, as well as any other infectious agents, and handled with the same precautions used in handling patient specimens.

Assigned Values and Ranges Lot #3220M002 (Representative Values)
(Containing Vial Lots L1 #3222M001 & L2 #3224M002)

<table>
<thead>
<tr>
<th>For In Vitro Diagnostic Use</th>
<th>LEVEL 1 – 3222M001</th>
<th>LEVEL 2 – 3224M002</th>
</tr>
</thead>
<tbody>
<tr>
<td>INSTRUMENT METHOD</td>
<td>UNITS</td>
<td>MEAN</td>
</tr>
<tr>
<td>BAYER A1cNOW Plus+</td>
<td>%</td>
<td>5.1</td>
</tr>
<tr>
<td>Siemens DCA 2000 “Vantage”</td>
<td>%</td>
<td>5.1</td>
</tr>
<tr>
<td>Tosoh G7</td>
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<tr>
<td>Tosoh G8</td>
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<tr>
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<td>Trinity Ultra2</td>
<td>%</td>
<td>5.3</td>
</tr>
<tr>
<td>Cobas Integra 400</td>
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</tr>
</tbody>
</table>

+Data not available at time of printing. Contact NOD for updated package insert values.

*Add your analyzers values by establishing your own internal value assignment if not already listed

Complementary NOD Liquid HbA1c Linearity is also available:
DIABETES A1c LINEARITY 4 LEVEL Part No: HbL-G04041-100