Body Fluids 2 Calibration Verification / Linearity Test Kit

INTENDED USE
VALIDATE Body Fluids 2 Calibration Verification / Linearity Test Kit solutions are intended for *in vitro* diagnostic use in the determination of linearity, calibration verification and verification of reportable range in automated instrument systems for the following analytes: Calcium (CA), Chloride (CL), Potassium (K), Sodium (NA), Total Bilirubin (TBIL) and Uric Acid (UA).

VALIDATE Test Kits are non-automated and intended for laboratory professional use only.

Each test kit consists of one bottle each of Levels 1 through 5. Each bottle contains 3.0 milliliters. There exists a linear relationship among Levels 1 through 5.

SUMMARY
For each VALIDATE Calibration Verification / Linearity Test Kit, multiple levels are provided to establish the relationship between theoretical and actual performance of each of the included analytes. The VALIDATE Calibration Verification / Linearity Test Kit will assist in the documentation of linearity, calibration verification and verification of linear range required by many inspection agencies. The solutions will also provide assistance when troubleshooting instrument systems, reagent problems and calibration anomalies.

REAGENTS
Reactive Ingredients:
Purified chemicals and constituents of human and/or animal source in human ascites fluid.

Nonreactive Ingredients:
Preservatives and stabilizers.

Precautions and Warnings:
For In Vitro Diagnostic Use
Disposal of all waste material should be in accordance with local guidelines.
Full SDS is available at www.mainestandards.com\doc-search.
Consult this SDS for detailed safety information including handling spills.

WARNING: Potentially Biohazardous
Human source material is considered potentially biohazardous. Use the Centers for Disease Control (CDC) recommended universal precautions for handling VALIDATE products.

STORAGE AND STABILITY
The **VALIDATE Body Fluids 2** Calibration Verification / Linearity Test Kit is stored at -10° to -25°C. ***Do NOT store in a frost-free freezer.*** Test kits are stable until the expiration date printed on the storage container when handled according to instructions. Do not use the Kit past its expiration date. ***A maximum of four (4) freeze-thaw cycles is recommended.***

PREPARATION
Prior to use, remove the **VALIDATE Body Fluids 2** Calibration Verification / Linearity Test Kit from storage and allow to come to room temperature (18° to 25°C). Invert gently several times before dispensing.

To maximize stability, it is recommended that exposure to room temperature be minimized. Tightly cap opened bottles and return to -10° to -25°C immediately after dispensing.

Discard any solutions that appear to have gross bacterial contamination.

VALIDATE Calibration Verification / Linearity Test Kits should be treated in the same manner as patient samples. If dilutions or pre-treatment are required as part of the testing procedure, follow the manufacturer's instructions.

ASSAY
Analyze each level in replicates. If following the CLSI EP6 guidelines for linearity, use a random analytical sequence to assay each level.

CALCULATION OF RESULTS
VALIDATE Calibration Verification / Linearity material is prepared in a manner such that an equal distance (delta) exists between Levels 1 through 5. This dilution scheme is consistent with the CLSI EP6 recommendation for preparing linearity sets.

Two examples for calculating the theoretical values of Levels 1 through 5 are provided below.

Example 1:
Choose two consecutive levels and calculate the delta between the recovered values. The following example demonstrates the use of the delta between Levels 2 and 3 to calculate the theoretical value for Levels 1, 4, and 5.

Level 3 – Level 2 = Delta

Level 1 Theoretical = Level 2 Recovered – Delta
Level 4 Theoretical = Level 3 Recovered + Delta
Level 5 Theoretical = Level 4 Theoretical + Delta

NOTE: The user can select the calculated delta between any two consecutive levels to calculate the theoretical values. Typically, the user should choose an area of recovery known to be linear for the method being studied.

Example 2:
Theoretical values can be determined using the recovered values for Levels 1 and 5. Using this method, the following formulas apply:

Level 2 Theoretical = 0.75 * (Level 1) + 0.25 * (Level 5)
Level 3 Theoretical = 0.5 * (Level 1) + 0.5 * (Level 5)
Level 4 Theoretical = 0.25 * (Level 1) + 0.75 * (Level 5)

After theoretical values are calculated, for each analyte plot the expected (Theoretical) value on the x-axis versus the Recovered value on the y-axis using standard linear graph paper. If the system is linear, the plot should approximate a straight line. The point at which the line is no longer straight can be used to determine the limit of linearity or the reportable range.

Data reduction is available from LGC Clinical Diagnostics. Commercially available linear regression software may also be used. The software should provide data point display and x-y graphical presentation. Linear regression should be interpreted using standard statistical analysis and the results should be compared with the instrument manufacturer's claims for linearity or with individual laboratory performance requirements. The degree of acceptable nonlinearity is an individual judgment based on methodology, clinical significance and medical decision levels of the test analyte.

LIMITATIONS
VALIDATE Calibration Verification / Linearity Test Kit solutions are not intended for use as routine quality control materials or as calibration materials.

EXPECTED VALUES
VALIDATE Calibration Verification / Linearity Test Kits are manufactured such that an equal distance (delta) exists between levels as recommended by CLSI EP6 for assessing linearity. As the distance between levels is equal, any two levels can be held to be 'true' when assayed and the theoretical values for each of the other three levels can be calculated allowing this test kit to be used on multiple automated instrument systems.

The reagent manufacturer's recommended diluent can be used to make dilutions of the low level to obtain a result lower than that level, if needed.

TYPICAL VALUES

Actual results obtained may vary depending on instrumentation, methodology and assay temperature. Results may also be dependent on the accuracy of the instrument / reagent system calibration. The degree of acceptable nonlinearity is an individual judgment based on methodology, clinical significance and medical decision levels of the test analyte.

Typical recovered values are calculated based on CLSI EP6 equal distance (delta) manufacturing protocols where two recovered values can be held as "True", and other levels calculated based on equal distance. **VALIDATE** typically uses Level 1 and Level 3 to calculate additional target values.

Typical Recovered Values on Roche COBAS®

206ro Lot: 10830817

Analyte	Units	Level 1	Level 2	Level 3	Level 4	Level 5
CA	mg/dL	1.6	6.0	10.4	14.7	19.1
CL	mmol/L	54	72	91	109	128
K	mmol/L	3.0	4.5	6.0	7.5	9.0
NA	mmol/L	61	88	115	142	169
TBIL	mg/dL	0.3	7.6	14.9	22.2	29.5
UA	mg/dL	0.46	6.5	12.6	18.6	24.7

206ro Lot: 10830817

Analyte	SI Units	Level 1	Level 2	Level 3	Level 4	Level 5
CA	mmol/L	0.400	1.500	2.600	3.675	4.775
CL	mmol/L	54	72	91	109	128
K	mmol/L	3.0	4.5	6.0	7.5	9.0
NA	mmol/L	61	88	115	142	169
TBIL	µmol/L	5.1	130.0	254.8	379.6	504.5
UA	µmol/L	27.4	386.6	749.4	1106.3	1469.2

The following symbols may be used where applicable in labeling for LGC Clinical Diagnostics products:

LOT **Lot Number**

Expiration Date

Manufacturer

Temperature Limit

IVD ***In Vitro* Diagnostic Medical Device**

REF **Catalog Number**

Instructions for Use

Biological Risk

Do Not Reuse

Caution

CE EU REP Wellkang Ltd, Enterprise Hub, NW Business Complex, 1 Beraghmore Rd, Derry, BT48 8SE, Northern Ireland

Most VALIDATE products are registered in the EU. For availability within EU nations, check with your distributor or contact cdx-sales@lgcgroup.com

ORDERING INFORMATION

ORDER NO.: 206ro

VALIDATE Body Fluids 2
Calibration Verification / Linearity Test Kit:
5 x 3 mL

Any serious incident occurring in relation to the use of this device shall be reported to the Manufacturer as well as the Competent Authority of the member state in which the user/patient is established.

CONTACT INFORMATION:

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Issue Date	Modification
Feb 26, 2026	Removed porcine statement due to design change