



LP Calibration Verification / Linearity Test Kit

INTENDED USE

VALIDATE LP Calibration Verification / Linearity Test Kit (U.S. Patent #7,244,618 and Canadian Patent #2,579,001) solutions are intended for *in vitro* diagnostic use in the quantitative determination of linearity, calibration verification and verification of reportable range in automated, semi-automated and manual instrument systems for the following analytes: human high density lipoprotein (HDL) and human low density lipoprotein (LDL). Test kits also contain APOA and APOB as structural elements of the HDL and LDL micelles.

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

Each **VALIDATE LP** Calibration Verification / Linearity Test Kit contains human HDL and human LDL in a human serum protein base. Multiple levels are provided to establish the relationship between theoretical and actual performance of each of the included analytes. The **VALIDATE LP** 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:

Human HDL, human LDL, APOA and APOB in a human serum protein base.

Nonreactive Ingredients:

Sodium Azide, preservatives and stabilizers.

Precautions and Warnings:

For In Vitro Diagnostic Use

Disposal of all waste material should be in accordance with local guidelines.

WARNING: Potentially Biohazardous

Human source material is considered potentially biohazardous. Material of human origin used in the manufacture of this test kit was tested at the donor level using FDA or CE approved methods and found to be non-reactive for HBV, HCV and HIV. Because no test method can offer complete assurance that infectious agents are absent, these specimens should be handled and treated as potentially infectious.

WARNING: Sodium Azide

Sodium Azide may react with lead and copper plumbing to form potentially explosive metal azides. On disposal, flush with a large volume of water to prevent azide build-up.

STORAGE AND STABILITY

VALIDATE LP Calibration Verification / Linearity Test Kits are 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 bottle and storage container when handled according to instructions.

PREPARATION

Prior to use, remove the **VALIDATE LP** 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.

The **VALIDATE LP** Calibration Verification / Linearity Test Kit 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 each consecutive level. 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:

Mean Recovered Values

Level 1	51
Level 2	164
Level 3	275
Level 4	388
Level 5	501

Using Level 2 and Level 3 recovered values to calculate the Delta, the above data produces the following:

$$\text{Level 3} - \text{Level 2} = \text{Delta, or } (275 - 164 = 111)$$

$$\begin{aligned} \text{Level 1 Theoretical} &= \text{Level 2 Recovered} - \text{Delta, or } (164 - 111 = 53) \\ \text{Level 4 Theoretical} &= \text{Level 3 Recovered} + \text{Delta, or } (275 + 111 = 386) \\ \text{Level 5 Theoretical} &= \text{Level 4 Theoretical} + \text{Delta, or } (386 + 111 = 497) \end{aligned}$$

Using the delta between Level 2 and Level 3, the theoretical value for each level would be:

Level	Theoretical (x-axis)	Recovered (y-axis)
1	53	51
2	164	164
3	275	275
4	386	388
5	497	501

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:

$$\begin{aligned} \text{Level 2 Theoretical} &= 0.75 * (\text{Level 1}) + 0.25 * (\text{Level 5}) \\ \text{Level 3 Theoretical} &= 0.5 * (\text{Level 1}) + 0.5 * (\text{Level 5}) \\ \text{Level 4 Theoretical} &= 0.25 * (\text{Level 1}) + 0.75 * (\text{Level 5}) \end{aligned}$$

Using the recovered values for Level 1 (51) and Level 5 (501), the following applies:

$$\begin{aligned} \text{Level 2 Theoretical} &= 0.75 * (51) + 0.25 * (501) = 163.5 \\ \text{Level 3 Theoretical} &= 0.5 * (51) + 0.5 * (501) = 276 \\ \text{Level 4 Theoretical} &= 0.25 * (51) + 0.75 * (501) = 388.5 \end{aligned}$$

Level	Theoretical (x-axis)	Recovered (y-axis)
1	51	51
2	163.5	164
3	276	275
4	388.5	388
5	501	501

After the 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 Maine Standards Company. 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 LP Calibration Verification / Linearity Test Kit solutions are not intended for use as routine quality control materials or as calibration materials.

EXPECTED VALUES

VALIDATE LP Calibration Verification / Linearity Test Kits are manufactured such that a linear relationship exists among Levels 1 through 5. The reagent manufacturer's recommended diluent can be used to make dilutions of Level 1 to obtain a result lower than Level 1, if needed.

TRACEABILITY

VALIDATE Calibration Verification / Linearity Test Kit solutions are tested during manufacturing with standards traceable to National Institute for Standards and Technology (NIST) Standard Reference Material (SRM), where available. For analytes where NIST materials are not available, primary analytical standards are used.

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 Values by Level 501ro						
Analyte	Units	1	2	3	4	5
HDL	mg/dL	3	32	62	91	120
LDL	mg/dL	3	140	277	413	550

ORDERING INFORMATION

ORDER NO.: 501ro

VALIDATE LP

Calibration Verification / Linearity Test Kit 5 x 3 mL

For technical assistance or to place an order, please call:
800-377-9684 or
207-892-1300
Fax 207-892-2266

Please allow 5 to 7 days for delivery.



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Repeating text 'VALIDATE' used as a background watermark.

CE Symbols – This product fulfills the requirements of the European Directive 98/79/EC for in vitro medical devices. The following symbols may be used where applicable in labeling for Maine Standards Company products:

- LOT Lot Number
Expiration Date
Manufacturer
Storage Temperature
IVD In Vitro Diagnostic Medical Device
REF Catalog Number
Insert
Biological Risk



For a list of countries in which VALIDATE® is registered see: www.mainestandards.com/ce

A worksheet to assist with manually calculating theoretical values can be found at www.mainestandards.com/Products or by calling Customer Support at 1-800-377-9684