

## Cholesterol/Cholesteryl Ester Quantitation Kit II

(Catalog #K623-100; 100 assays; Store at -20°C)

### I. Introduction:

Cholesterol is an essential molecule in all animal life. It has been involved in both normal development and diseases. The Cholesterol/Cholesteryl Ester Quantitation Kit provides a simple method for sensitive quantification of free cholesterol, cholesteryl esters, or both using a colorimetric method. Most of the cholesterol in blood is in the form of cholesteryl ester. These esters can be hydrolyzed to free cholesterol under the appropriate conditions. In the assay, free cholesterol is oxidized by cholesterol dehydrogenase to generate NADH which reacts with a sensitive probe resulting in strong absorbance at 450 nm. The assay can detect free or total cholesterol depending upon whether esterase is utilized to hydrolyze cholesteryl esters present. Cholesteryl ester can be determined by subtracting the value of free cholesterol from the total cholesterol (cholesterol plus cholesteryl esters). The probe in this kit is more stable, sensitive and specific. The assay can tolerate interferences from various samples significantly.

### II. Kit Contents:

Component	Volume	Cap Code	Part No.
Cholesterol Assay Buffer	25 ml	WM	K623-100-1
Substrate Mix (lyophilized)	1 vial	Red	K623-100-2
Enzyme Mix (lyophilized)	1 vial	Green	K623-100-3
Esterase (lyophilized)	1 vial	Blue	K623-100-4
Cholesterol Standard (2 µg/µl)	100 µl	Yellow	K623-100-5

### III. Storage and Handling:

Store kit at -20°C, protect from light. Allow Assay Buffer to warm to room temperature before use. Keep enzymes and cholesterol standard on ice while using. Read the entire protocol before perform the assay.

### IV. Reagent Preparation:

**Substrate Mix:** Reconstitute with 220 µl of Assay Buffer and mix thoroughly. The solution is stable for 2 months at 4°C.

**Enzyme Mix:** Dissolve in 220 µl Cholesterol Assay Buffer before use. Aliquot and store at -20°C. Use within two months.

**Esterase:** Dissolve in 220 µl Cholesterol Assay Buffer before use. Aliquot and store at -20°C. Use within two months.

### V. Cholesterol Assay Protocol:

- Standard Curve Preparations:** Dilute the Cholesterol Standard to 0.25 µg/µl by adding 20 µl of the Cholesterol Standard to 140 µl of Cholesterol Assay Buffer, mix well. Add 0, 4, 8, 12, 16, 20 µl into a series of wells. Adjust volume to 50 µl/well with Cholesterol Assay Buffer to generate 0, 1, 2, 3, 4, 5 µg/well of the Cholesterol Standard.
- Sample Preparation:** Serum should be diluted 10X with Cholesterol Assay Buffer, then use 2-20 µl for each testing. For cells or tissue samples, extract 10<sup>8</sup> cells or 10 mg tissue with 200 µl CHCl<sub>3</sub>:IPA:NP-40 (7:11:0.1) in a microhomogenizer. Centrifuge the extract for 5 min at 15,000xg. Transfer the liquid phase to a new tube and air dry at 50°C then place samples under vacuum for 30 min to remove any remaining solvent. Dissolve dried lipids with 200 µl of Cholesterol Assay Buffer by sonicating or vortexing until homogeneous (OK if the solution becomes cloudy). The extraction procedure can be scaled up if larger amounts of sample are desired. Use 1- 50 µl of extract per assay. Adjust volume to 50 µl/well with Cholesterol Assay Buffer. For unknown samples, we suggest testing different amounts of sample to ensure that readings are within the limits of the standard curve.
- Ester Hydrolysis:** Add 2 µl Esterase to each standard and samples for which the total cholesterol value is desired. (See Notes a and b in step 4). Incubate 30 min at 37°C.

- Reaction Mix Preparation:** Mix enough reagent for the number of assays performed: For each well, prepare a total 50 µl Reaction Mix containing:

44 µl Cholesterol Assay Buffer  
2 µl Substrate Mix  
2 µl Cholesterol Enzyme Mix

Mix. Add 50 µl of the Reaction Mix to each well containing standard or samples.

### Notes:

a) Cholesterol Esterase hydrolyzes cholesteryl ester to cholesterol. If you want to detect free cholesterol only, omit the Cholesterol Esterase in the reaction. In the presence of Cholesterol Esterase, the assay detects both free cholesterol and cholesteryl esters. If you want to determine Cholesteryl Ester only, subtract the value of free cholesterol from the value of total cholesterol (Cholesterol and Cholesteryl Ester).

b) **The Cholesterol Standard contains a mixture of free cholesterol and cholesterol esters in a similar ration of serum. Cholesterol Esterase must be added to the standard reaction to convert all cholesterol in the standard.**

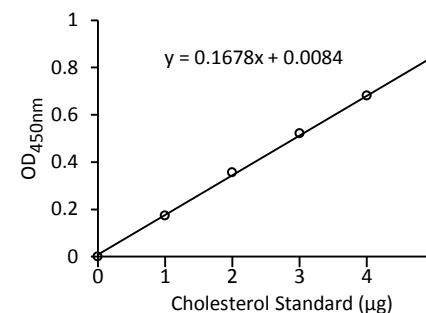
- Incubate the reaction for 30 min at 37°C, protect from light.
- Measure absorbance at 450 nm in a microplate reader.
- Calculations:** Subtract the 0 standard background reading from all readings. Plot the standard curve. Apply sample readings to the standard curve. Cholesterol concentration in samples can then be calculated:

$$C = A/V \text{ (}\mu\text{g}/\mu\text{l)}$$

Where: **A** = amount of cholesterol determined from Standard Curve (in µg).

**V** = volume of sample added into the reaction well (in µl).

Cholesterol molecular weight: 386.65. 1µg/µl = 100 mg/dL.



**Figure: Cholesterol Standard Curve:** Cholesterol/Cholesteryl Ester was quantified according to the kit instructions. Background from the 0 standard reading (without cholesterol) has been subtracted from all readings.

### RELATED PRODUCTS:

- Cholesterol Assay Kit
- Triglyceride Assay Kit
- Fatty Acid Assay Kit
- HDL/LDL Assay Kit
- CETP/PLTP Assay Kits
- Glycerol Assay Kit

**FOR RESEARCH USE ONLY! Not to be used on humans.**

## GENERAL TROUBLESHOOTING GUIDE:

Problems	Cause	Solution
Assay not working	<ul style="list-style-type: none"> <li>• Use of ice-cold assay buffer</li> <li>• Omission of a step in the protocol</li> <li>• Plate read at incorrect wavelength</li> <li>• Use of a different 96-well plate</li> </ul>	<ul style="list-style-type: none"> <li>• Assay buffer must be at room temperature</li> <li>• Refer and follow the data sheet precisely</li> <li>• Check the wavelength in the data sheet and the filter settings of the instrument</li> <li>• Fluorescence: Black plates (clear bottoms) ; Luminescence: White plates ; Colorimeters: Clear plates</li> </ul>
Samples with erratic readings	<ul style="list-style-type: none"> <li>• Use of an incompatible sample type</li> <li>• Samples prepared in a different buffer</li> <li>• Samples were not deproteinized (if indicated in datasheet)</li> <li>• Cell/ tissue samples were not completely homogenized</li> <li>• Samples used after multiple free-thaw cycles</li> <li>• Presence of interfering substance in the sample</li> <li>• Use of old or inappropriately stored samples</li> </ul>	<ul style="list-style-type: none"> <li>• Refer data sheet for details about incompatible samples</li> <li>• Use the assay buffer provided in the kit or refer data sheet for instructions</li> <li>• Use the 10 kDa spin cut-off filter or PCA precipitation as indicated</li> <li>• Use Dounce homogenizer (increase the number of strokes); observe for lysis under microscope</li> <li>• Aliquot and freeze samples if needed to use multiple times</li> <li>• Troubleshoot if needed, deproteinize samples</li> <li>• Use fresh samples or store at correct temperatures till use</li> </ul>
Lower/ Higher readings in Samples and Standards	<ul style="list-style-type: none"> <li>• Improperly thawed components</li> <li>• Use of expired kit or improperly stored reagents</li> <li>• Allowing the reagents to sit for extended times on ice</li> <li>• Incorrect incubation times or temperatures</li> <li>• Incorrect volumes used</li> </ul>	<ul style="list-style-type: none"> <li>• Thaw all components completely and mix gently before use</li> <li>• Always check the expiry date and store the components appropriately</li> <li>• Always thaw and prepare fresh reaction mix before use</li> <li>• Refer datasheet &amp; verify correct incubation times and temperatures</li> <li>• Use calibrated pipettes and aliquot correctly</li> </ul>
Readings do not follow a linear pattern for Standard curve	<ul style="list-style-type: none"> <li>• Use of partially thawed components</li> <li>• Pipetting errors in the standard</li> <li>• Pipetting errors in the reaction mix</li> <li>• Air bubbles formed in well</li> <li>• Standard stock is at an incorrect concentration</li> <li>• Calculation errors</li> <li>• Substituting reagents from older kits/ lots</li> </ul>	<ul style="list-style-type: none"> <li>• Thaw and resuspend all components before preparing the reaction mix</li> <li>• Avoid pipetting small volumes</li> <li>• Prepare a master reaction mix whenever possible</li> <li>• Pipette gently against the wall of the tubes</li> <li>• Always refer the dilutions in the data sheet</li> <li>• Recheck calculations after referring the data sheet</li> <li>• Use fresh components from the same kit</li> </ul>
Unanticipated results	<ul style="list-style-type: none"> <li>• Measured at incorrect wavelength</li> <li>• Samples contain interfering substances</li> <li>• Use of incompatible sample type</li> <li>• Sample readings above/below the linear range</li> </ul>	<ul style="list-style-type: none"> <li>• Check the equipment and the filter setting</li> <li>• Troubleshoot if it interferes with the kit</li> <li>• Refer data sheet to check if sample is compatible with the kit or optimization is needed</li> <li>• Concentrate/ Dilute sample so as to be in the linear range</li> </ul>
<p><b>Note:</b> The most probable list of causes is under each problem section. Causes/ Solutions may overlap with other problems.</p>		