



**CHOLESTEROL REAGENT**  
**C103-01**

<b>Contents</b>	<b>Product No.</b>	<b>Package</b>
Cholesterol Reagent	C103-01	1 x 250 mL

**REAGENT PREPARATION**

The Catachem Cholesterol Reagent is packaged ready for use.  
No preparation is required.

**REAGENT STORAGE AND STABILITY**

Store the unopened reagent at 2-8°C.  
When stored as directed, the reagent is stable until the expiration date stated on the label.

***NOT FOR USE IN UNPROFESSIONAL SETTINGS***

FOR TECHNICAL ASSISTANCE:  
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# CHOLESTEROL REAGENT C103-01 MANUAL/AUTOMATED APPLICATION

## Intended Use

For the quantitative determination of Cholesterol in serum.

## Clinical Significance

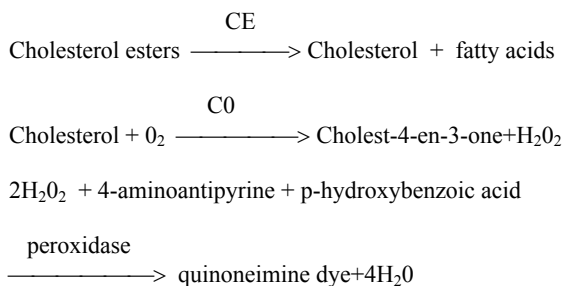
Determinations of Cholesterol are primarily used for diagnosing atherosclerosis and heart disease, liver disease and renal disease, as well as for monitoring the causes and treatment. (1)

## Method History

Enzymatic Cholesterol methods were introduced in the early nineteen seventies when Cholesterol Oxidase (EC1.1.3.6) and Cholesterol Esterase (EC 3.1.1.13) became available. Allain, et al, (1) and Roeschlau, et al (2) described the use of a single enzymatic reagent for the assay of Cholesterol in serum. Later, L.P.Leon, et al (3) automated these methods for use in continuous flow systems. Catachem's enzymatic Cholesterol method described below uses a combination of microbial Cholesterol Oxidase, pancreatic Cholesterol Esterase of animal source and a colorimetric end-point reaction based upon the work of Trinder. (4)

## Method Principle

Cholesterol esters present in the serum sample are hydrolyzed into free Cholesterol and fatty acids by pancreatic Cholesterol esterase (CE). The free Cholesterol formed in the presence of oxygen, is then oxidized to cholest-4-en-3-one by Cholesterol Oxidase (CO) with concomitant production of hydrogen peroxide. The hydrogen peroxide produced is quantitatively determined by coupling 4-aminoantipyrine with p-hydroxybenzoic acid (5) where a quinoneimine dye with maximum absorption at 550nm is produced. The intensity of the dye color thus produced is directly proportional to the concentration of the total Cholesterol in the serum sample. The reaction scheme illustrates the reactions that occur in this method.



## Reagent Content

Active ingredients in the reagents will be approximately as follows:

### Cholesterol Reagent

Each liter contains:

Peroxidase	≥1600 U
Cholesterol Oxidase (microbial)	≥150 U
Cholesterol Esterase (pancreatic)	≥125 U
4-aminoantipyrine	0.6 mM
p-hydroxybenzoic acid	18.7 mM
Buffer and non reactive ingredients	

## Precaution

Avoid contact of the reagent with skin and eyes. Should contact occur, wash affected area with plenty of cold water. **DO NOT PIPETTE REAGENTS BY MOUTH.**

## Preparation Of Working Reagent

Catachem Cholesterol reagent is packaged in ready-to-use form. No preparation is required.

## Storage And Stability

Store the Cholesterol reagent at 2-8°C. When stored as directed, the reagent is stable until the expiration date stated on the label. When stored as directed, the Working Cholesterol Reagent is stable for 12 months at 2-8°C and 12 weeks at room temperature. The Catachem Cholesterol Reagent has been tested to reflect shipping conditions and is stable for the lifespan of the product if frozen up to 5 times or upon reaching temperatures up to 40°C for up to one week.

## Specimen Collection And Preparation

Fresh non-hemolyzed serum is the specimen of choice. No preservatives are necessary. Cholesterol in human serum samples is stable for one week at 2-8 °C and for six months at -20 °C.

## Interfering Substances

A number of substances have been reported to affect the accuracy of cholesterol methods using the Cholesterol Oxidase/peroxidase procedures.(6) A summary of the influence of drugs on clinical laboratory procedures may be found by consulting D.S. Young, et al. (7)

## Expected Values

Based on the recommendations of the NIH Conference to review the scientific significance of Cholesterol values on human subjects (8) the following risk cutoff levels for the American population should be observed:

Ideal normal serum Cholesterol	≤200 mg/dL
Borderline serum Cholesterol	200-239 mg/dL
High serum Cholesterol	≥240 mg/dL

These values serve as suggested reference points only. For veterinary samples, ranges will be dependent on the species under test. It is recommended that each laboratory establish the normal ranges for the species under study and for the geographic area in which the laboratory is located.

## Directions For Use

Catachem's Cholesterol method requires one single reagent. The reagent is ready for use and requires no preparation. The Working Reagent is stable twelve months aboard the analyzer at 2-8°C and capped while not in use.

## Procedure

Important: Read the entire procedural instructions before proceeding with assay.



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## Materials Required But Not Provided

Spectrophotometer  
 Match cuvettes 1 cm light path  
 Timer to time incubation time  
 Pipette 1.0 ml for reagent  
 Pipette 0.010 ml for sample

## Materials Provided

Cholesterol Reagent

## Analytical Parameters

Wavelength 550nm  
 Temperature 37°C  
 Pathlength 1 cm  
 Reaction Mode endpoint  
 Reaction Time 5 min  
 Reagent Volume 1.0 ml  
 Sample Volume 0.01 ml (10uL)  
 Total Volume 1.01 ml  
 Sample-to-reagent Ratio 1:100

## Assay Procedure

1. Pipette 1.0 ml of Cholesterol reagent into each of three cuvettes marked "calibrator", "sample", and "blank".
2. Pipette 0.01 ml (10 uL) of calibrator or sample into their respective cuvettes. Mix all cuvettes well.
3. Incubate all cuvettes for 5 minutes at 37°C.
4. Set spectrophotometer wavelength at 550nm and zero the instrument with the cuvette marked "blank".
5. Read the "calibrator" and "sample" absorbencies.
6. Calculate the cholesterol concentration (mg/dL) in the sample(s), as shown in results and calculations.

## Results And Calculations

$$\text{Cholesterol (mg/dL)} = \frac{\text{sample abs}}{\text{calibrator abs}} \times \text{calibrator (mg/dL)}$$

Example:

Sample absorbance = 0.300

Calibrator absorbance = 0.250

Calibrator (mg/dL) = 200

$$\text{Cholesterol (mg/dL)} = \frac{0.300}{0.250} \times 200 = 240 \text{ mg/dL}$$

## Quality Control

To monitor the performance of the Reagent and the procedure used we recommend the regular use of a normal and abnormal control serum.

## Method Performance Characteristics

**Sensitivity:** 0.0010-0.0020 absorbance units per mg/dL.

**Linear Range:** 0-600 mg/dL.

**Precision:** Within-run and day-to-day precision is summarized below.

## Cholesterol Precision Study

Cholest.	Within-Run		Total Precision	
	Mean	SD	SD	CV
mg/dL	mg/dL	%	Mg/dL	%
56	0.71	1.2	0.71	1.3
261	1.95	0.75	6.3	2.4
456	2.97	0.65	3.0	0.70

## Correlation

A comparison of this method using an automated analyzer and a reference method based on the CO and CE reaction resulted in the following regression statistics:

$$\begin{aligned}
 N &= 155 \\
 Y &= 1.003x + 4.3 \\
 r &= 0.998 \\
 S_{y.x} &= 5.7
 \end{aligned}$$

## References

1. Allain CC, Poon LS, Chan CSG, Richmond W, Fu PC. Clin Chem 20, 470 (1974).
2. Roeschlau P, Bernt E. Gruber W. Clin Chem Clin Biochem 12, 226 (1974).
3. Leon LP, Chu DK, Stasiw RO, Snyder LR. Advances in Automated Analysis. Technicon's International congress, 152-156 (1976).
4. Trinder P. Ann Clin Biochem. 6, 24 (1969).
5. Bardelli F, Giannini G, Meattini F, Prencipe L, Tarli P. Clin Chem 24, No. 12 (1978).
6. Pesce MA, Bodourian SH. Clin Chem 23, 757-760 (1977).
7. Young DS, Pestaner LC, Gibberman V. Clin Chem 21, No. 5 (1975).
8. Report of the National Cholesterol Education Program Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults. Arch of Int Med. Vol 148, No. 1 pp 36-69. Jan. 1988.