Serum High-Density Lipoprotein Cholesterol (HDL-C):

The principle role of HDL in the lipid metabolism is the uptake and transport of cholesterol from peripheral tissues to the liver through a process known as reveres cholesterol transport.

The principle:

This reagent is only used for treatment of specimens before the determination of HDL-C with a reagent for total cholesterol. Low density lipoprotein (LDL), very low density lipoprotein (VLDL) and chylomicrons from specimens are precipitated by phosphotungstic acid (PTA) and magnesium chloride. HDL-C obtained in the supernatant after centrifugation is then measured with a total cholesterol reagent

Procedure:

Do not treat standard (vial R₂) enclosed in kit:

Solution	Macro-method	Micro-method	
Specimen	1 ml	0.5 ml	
Precipitant	100 μl	50 μl	

Mix vigorously, let stand for 10 min. at room temperature and centrifuge for 15 min. at 3500-4000 RPM. with BIOLAB total cholesterol CHOD-PAP or equivalent: Let stand reagents and supernatants at room temperature. Calibrate with standard enclosed in the kit or pre- treated series calibrator.

Solution	Blank	Standard	Sample
Reagent	ı ml	ı ml	ıml
DW	25 µl		
Standard		25µl	
Serum			25µl

Mix well the test tubes and let stand for 5 min. at 37°C or 10 min. at room temperature. Record absorbance at 500 nm. (480-520 nm) against the reagent blank.

1.1= standard remaining undiluted, 1.1 factor takes into account dilution of the specimen during the precipitation step.

Serum Low-Density Lipoprotein Cholesterol (LDL-C):

Serum LDL concentration can be calculated by the following equation

LDL = TC - (HDL + TG/5)

Serum Very Low-Density Lipoprotein (VLDL):

Serum VLDL concentration was calculated by dividing serum TG/5.

VLDL=TG/5