

CH 50 | Liposome Immunoassay

Screening for Complement Activity in human serum

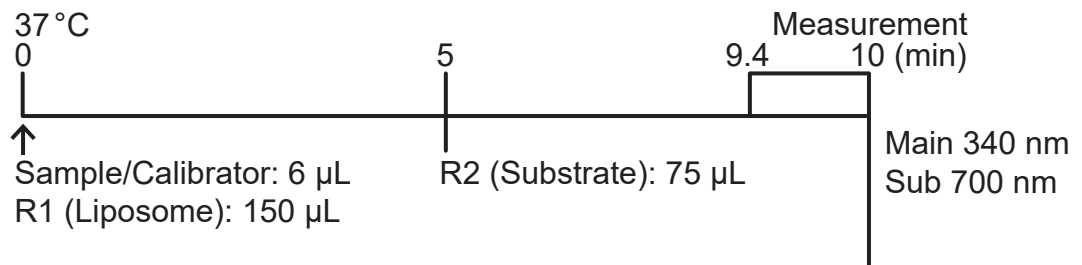
- Liposome immunoassay, stable and homogeneous
- Applicable to automated analyzers
- Precise, accurate
- Extended calibration stability
- Good correlation with Mayer’s hemolytic method

■ Principle

Complement in the sample is activated by the antigen-antibody complexes on the liposomes. The activated complement breaks the membrane of the liposomes. The enzyme glucose-6-phosphate dehydrogenase (G6PDH) contained in the liposome reacts with NAD and glucose-6-phosphate (G6P) in the reagent. During this enzyme reaction, the NAD is reduced to NADH. As a result of this reduction, absorbance at 340 nm increases. This is proportional to the CH50 activity.

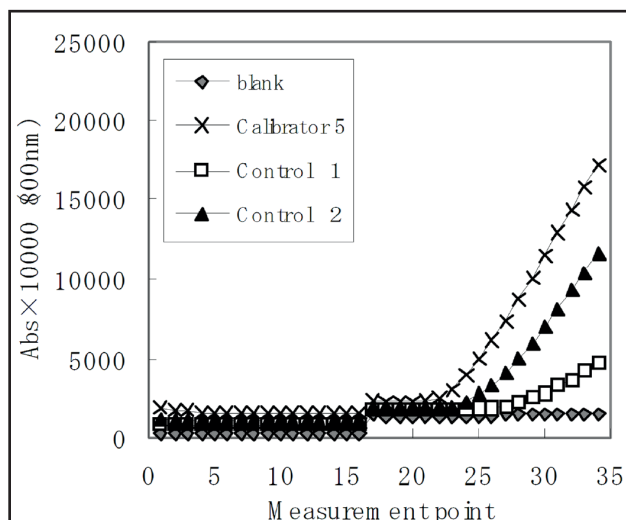
■ Procedure

Standard Procedure (Hitachi 917s)



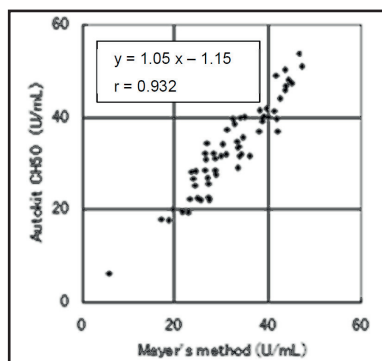
■ Reaction

Reaction time course

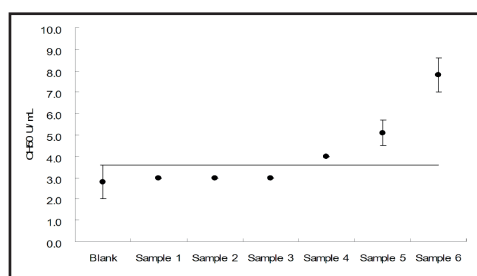


Range The measurable range is 10 – 60 U/mL

Correlation



Sensitivity 4U/mL



Interference Ascorbic acid concentrations up to 50 mg/dL, hemoglobin concentrations up to 500 mg/dL and bilirubin concentrations up to 40 mg/dL do not have a significant effect on the Autokit CH50 assay.

CE Applications	Aeroset	AU640	Hitachi 902
	Architect c8000	AU2700	Hitachi 904
	Architect c16000	Cobas6000	Hitachi 911
	AU400	Cobas8000	Hitachi 912
	AU600	Dimension	Konelab 30/60i

Ordering

Code No.	Product	Content
995-40801	Autokit CH50	R1: 2 x 20 mL R2: 1 x for 20 mL R2a: 1 x 20 mL
997-43801	CH50 Calibrator	CAL: 5 Conc. x for 0.5 mL
991-43701	Complement Control	CONTROL H: 10 x for 0.5 mL CONTROL L: 10 x for 0.5 mL