Cholinesterase FS*

Order Information

Cat. No.	Kit s	ıze			
1 1401 99 10 021	R1	5 x 20 mL	+	R2	1 x 25 mL
1 1401 99 10 930	R1	4 x 20 mL	+	R2	2 x 10 mL
1 1401 99 90 314	R1	10 x 20 mL	+	R2	2 x 30 mL

Intended Use

Diagnostic reagent for quantitative in vitro determination of cholinesterase (CHE) in serum or plasma on photometric systems.

Summary

Cholinesterases (CHE) are a group of enzymes preferably splitting choline and thiocholine esters. The denomination Serum Cholinesterase and Pseudocholinesterase are also commonly used. The CHE measured in serum and plasma is synthesized in the liver and is determined in diagnosis of liver diseases, nephrotic syndrome and intestinal diseases with loss of protein (exudative enteropathy). Strongly decreased values can indicate intoxication by pesticides. Measurement of CHE is also a part of pre-operative diagnostics as CHE is needed for the inactivation of muscle relaxants often used in surgeries. [1]

Method

Kinetic photometric test, optimized method according to the recommendation of the German Society of Clinical Chemistry (DGKC).

Cholinesterase hydrolyses butyrylthiocholine under release of butyric acid and thiocholine. Thiocholine reduces yellow potassium hexacyanoferrate (III) to colorless potassium hexacyanoferrate (II). The decrease of absorbance is measured at 405 nm.

Cholinesterase				
Butyrylthiocholine + H ₂ O	Thiocholine + Butyrate			
2 Thiocholine + 2 [Fe(CN) ₆] ³⁻ + H ₂ O	—► Choline + 2 [Fe(CN) ₆] ⁴⁻ + H ₂ O			

Reagents

Components and Concentrations

R1:	Pyrophosphate	pH 7.6	95 mmol/L
	Potassium hexacyanoferrate (III)		2.5 mmol/L
R2:	Butyrylthiocholine		75 mmol/L

Storage and Stability

The reagents are stable up to the end of the indicated month of expiry, if stored at $2-8^{\circ}\text{C}$ and contamination is avoided. Do not freeze the reagents and protect them from light.

Warnings and Precautions

- Reagent 1: Danger. Contains Tetrasodium pyrophosphate-10-hydrate. H318 Causes serious eye damage. P280 Wear protective gloves/protective clothing/eye protection. P305+P351+P338 If in eyes: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. P310 Immediately call a poison center or doctor/physician.
- In very rare cases, samples of patients with gammopathy might give falsified results [2].
- Please refer to the safety data sheets and take the necessary precautions for the use of laboratory reagents. For diagnostic purposes, the results should always be assessed with the patient's medical history, clinical examinations and other findings.
- For professional use only.

Waste Management

Refer to local legal requirements.

Reagent Preparation

The reagent is ready to use.

Materials Required

General laboratory equipment

Specimen

Serum or heparin plasma

Stability [1,3]:

1 week at $15-25^{\circ}$ C 2 week at $2-8^{\circ}$ C 6 months at -20° C

Only freeze once. Discard contaminated specimens.

Assav Procedure

Applications for automated systems are available on request.

Wavelength Hg 405 nm Optical path 1 cm Temperature 37°C

Measurement Against reagent blank

	Blank	Sample or calibrator			
Sample or calibrator	-	20 μL			
Dist. Water	20 µL	<u>-</u> Î			
Reagent 1	1000 μL	1000 μL			
Mix, incubate for approx. 3 min., then add:					
Reagent 2	250 µL	250 μL			
Mix, read absorbance ((A) after 2 m	in and start stop watch.			
Read absorbance (A) again after 1, 2 and 3 minutes.					

Calculation

With factor

 Δ A/min x 68500 = CHE activity [U/L]

With calibrator

CHE [U/L] = $\frac{\Delta A/min. Sample}{\Delta A/min. Calibrator} x Conc. Calibrator [U/L]$

Conversion Factor

Cholinesterase [kU/L] x 16.67 = Cholinesterase [µkat/L]

Calibrators and Controls

DiaSys TruCal U is recommended for calibration. TruCal U calibrator values have been made traceable to the molar extinction coefficient. Use DiaSys TruLab N and P for internal quality control. Each laboratory should establish corrective action in case of deviations in control recovery.

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	Cat. No.	Ki	t size)
TruCal U	5 9100 99 83 063	20	Х	3 mL
	5 9100 99 83 064	6	Х	3 mL
TruLab N	5 9000 99 83 062	20	Х	5 mL
	5 9000 99 83 061	6	Х	5 mL
TruLab P	5 9050 99 83 062	20	Х	5 mL
	5 9050 99 83 061	6	Х	5 mL

Performance Characteristics

Limit of detection**

Data evaluated on BioMajesty® JCA-BM6010/C

Exemplary data mentioned below may slightly differ in case of deviating measurement conditions.

Measuring range up to 19 kU/L. When values exceed this range, samples should be diluted 1 + 5 with NaCl solution (9 g/L) and the result multiplied by 6.

0.04 kU/L

Interfering substance	Interferences ≤ 10% up to	
Ascorbic acid	30 mg/dL	
Bilirubin (conjugated)	54 mg/dL	
Bilirubin (unconjugated)	42 mg/dL	
Hemoglobin	500 mg/dL	
Lipemia (triglycerides)	1000 mg/dL	
For further information on interfering substances refer to Young DS. [4]		

Precision			
Within run (n=20)	Sample 1	Sample 2	Sample 3
Mean [kU/L]	4.34	5.75	6.90
CV [%]	1.13	1.08	0.972
Between day (n=20)	Sample 1	Sample 2	Sample 3
Mean [kU/L]	4.22	4.88	6.91
CV [%]	0.887	1.46	1.69

Method comparison (n=100)		
Test x	Competitor Cholinesterase	
Test y	DiaSys Cholinesterase FS	
Slope	1.000	
Intercept	–0.240 kU/L	
Coefficient of correlation	0.9996	

lowest measurable activity which can be distinguished from zero; mean + 3 SD (n = 20) of an analyte free specimen.

Reference Range

As follows [3]:

Women 3.93 - 10.8 kU/L 65.5 - 180 µkat/L 4.62 - 11.5 kU/L 77.0 - 192 µkat/L Men

Each laboratory should check if the reference ranges are transferable to its own patient population and determine own reference ranges if necessary.

Literature

- 1. Hallbach J, Klinische Chemie für den Einstieg. 1st ed Stuttgart: Thieme;2001. p. 143-4.
- Bakker AJ, Mücke M. Gammopathy interference in clinical chemistry assays: mechanisms, detection and prevention. ClinChemLabMed 2007;45(9):1240-1243.
- Recommendations of the German Society for Clinical Chemistry. Standardization of methods for the estimation of enzyme activities in biological fluids: Standard method for the determination of Cholinesterase activity. J Clin Chem Clin Biochem 1992;30:163-70.
- Young DS. Effects of Drugs on Clinical Laboratory Tests. 5th ed. Volume 1 and 2. Washington, DC: The American Association for Clinical Chemistry Press 2000.







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December 2019/1

^{*} Fluid Stable