

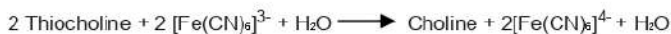
Reagent Kit for quantitative estimation of Cholinesterase (ChE) in Serum or Plasma.

DIAGNOSTIC SIGNIFICANCE:

Cholinesterases (ChE) are a group of enzymes preferably splitting choline and thiocholine esters. The names 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.

PRINCIPLE:

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.



SPECIMEN COLLECTION:

Serum or heparinised or EDTA plasma is suitable. Serum or plasma samples remain stable for 14 days at 2-8°C.

KIT PRESENTATION:

PACK SIZE	1 X 20 ml	1 X 40 ml	2 X 50 ml
R1- Cholinesterase (Buffer)	1 X 16 ml	1 X 32 ml	2 X 40 ml
R2- Cholinesterase (Substrate)	1 X 04 ml	1 X 08 ml	2 X 10 ml

WORKING REAGENT PREPARATION

Mixing 4 volumes of R1-Cholinesterase (Buffer) with 1 volume of R2- Cholinesterase (Substrate). i.e. 800 µl R1 + 200 µl R2.

REAGENT STORAGE AND STABILITY

All reagents are stable at 2-8°C until the expiry date stated on the label. Do not freeze the reagents and protect from light.

NORMAL VALUES:

- Female : 3930 – 10800 IU/L
- Male : 4620 – 11500 IU/L

Each laboratory should establish its own reference range.

SENSITIVITY / LIMIT OF DETECTION:

The lower limit of detection is 55 U/L

ASSAY PARAMETERS:

Reaction	: Kinetic	Sample Volume	: 20 µl
Wavelength	: 405 nm	R1 + R2 Volume	: 800 µl + 200 µl
Flow Cell Temp.	: 37°C	Factor	: 55000
Initial Delay	: 120 Sec	Reaction Slop	: Decreasing
Interval Time	: 60 Sec	Zero Setting	: Dist. Water
Read Time	: 180 Sec	Linearity	: 20,000
No. of Reading	: 03	Unit	: IU/L

PROCEDURE:

Addition Sequence	Test
R1- Cholinesterase (Buffer Reagent)	800 µl
R2- Cholinesterase (Substrate Reagent)	200 µl
Sample (Test)	20 µl

Mix & aspirate immediately and read first absorbance of test exactly at 120 seconds and then, **second, third and fourth** at an interval of 60 seconds at 405 nm. Determine the mean change in absorbance per minute. (ΔA/min) and calculate the test results.

CALCULATION:

Cholinesterase Activity (IU/L) = ΔA/min X 55000

LINEARITY:

This method is linear up to 20,000 IU/L. For values above 20,000 IU/L, dilute the sample suitably with 0.9 % saline, and repeat the assay. Apply correction due to dilution to arrive at a final result.

REFERENCES:

1. Recommendation 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.
2. Thomas L, Clinical laboratory diagnostics, 1st ed frankfurt: TH-Books Verlagsgesellschaft; 1998. P.65-71.
3. Hallbach J, Klinische Chemie fur den Einstieg. 1st ed Stuttgart: Thieme;2001. p.143-4.

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Expiry Date



In-Vitro Diagnostics Use



Storage



Mfg. Date



Batch Number



Catalogue Number



See Package Insert