

Acetaminophen L3K®

FOR THE QUANTITATIVE MEASUREMENT OF ACETAMINOPHEN METHOD: ENZYMATIC (ACYL AMIDOHYDROLASE); ENDPOINT

Acetaminophen (paracetamol) measurements are used in the diagnosis and treatment of acetaminophen overdose toxicity.

SEKISUI'S Acetaminophen L3K[®] assay uses a spectrophotometric method that is rapid, reliable, convenient and specific for acetaminophen. It is intended for use with serum and plasma.

Features	
 Two part stable liquid A single level liquid acetaminophen calibrator is included in the kit 	 Reduced NAC interference Suitable for use with serum and lithium heparin plasma
Benefits	
 Easy to use, no additional reagent preparation required Convenient and cost effective 	Confidence in resultsFlexible sample types to meet different laboratory needs
Performance Characteristics	
Precision • Within-Run: ≤1.5% • Total Precision: ≤2.9% Accuracy SERUM(a) • Slope: 1.064 • Intercept: 1.1 µg/mL (7.0 µmol/L) • Correlation Coefficient: 0.9998 PLASMA(b) • Slope: 0.999 • Intercept: -0.3 µg/mL (-2.2 µmol/L) • Correlation Coefficient: 0.9999	 No Significant Interferences Up to Levels Indicated Hemoglobin: 200 mg/dL (31 μmol/L) Conjugated Bilirubin: 2 mg/dL (23.7 μmol/L) Unconjugated Bilirubin: 2 mg/dL (34.2 μmol/L) Ascorbic Acid: 3000 μg/dL (170 μmol/L) N-Acetylcysteine: 1500 mg/L (9.2 mmol/L) Intralipid: 200 mg/dL (600 mg/dL (6.8 mmol/L) Simulated Triglycerides) Reference Range⁽¹⁾ Therapeutic Concentration: 10-30 μg/mL (66-199 μmol/L) Toxic Concentration: >200 μg/mL (1324 μmol/L)
Linearity • 0.6-377.5 μg/mL (4-2500 μmol/L)	
 (a) The performance of this method (y) was compared with the performance with a similar acetaminophen method (x) on a Roche/Hitachi® 717 analyzer. (b) The performance of this method with plasma (y) was compared with the performance of this method with serum (x) on a Roche/Hitachi® 717 analyzer. 	

Ordering Information		
	Configuration	Catalog Number
ACETAMINOPHEN L3K®	R1 3 x 10 mL R2 6 x 10 mL 1 x 5 mL (Calibrator)	506-30

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