CLINICAL CHEMISTRY REAGENTS

Urea L3K®

FOR THE QUANTITATIVE MEASUREMENT OF UREA

METHOD: ENZYMATIC (UREASE); RATE

Urea measurements are used as an aid to monitor and diagnose renal disease.

The Sekisui Urea L3K[®] assay employs the urease method with a stabilized NADH cofactor analog for improved performance. It is intended for the measurement of urea in serum.

Features:		
 One part stable liquid No significant interference displayed from samples with elevated levels of hemolysis, icterus or lipemia 	• Applicable to multiple chemistry platforms	
Benefits:		
 Easy to use, no additional reagent preparation required 	 Flexible testing, well suited for use with fully automated procedures 	

• Reduces the need for sample dilutions

Performance Characteristics

Precision

- Within-Run: ≤2.9%
- Total Precision: ≤4.0%

Accuracy^(a)

- Slope: 1.00
- Intercept: -0.1 mg/dL (-0.04 mmol/L)
- Correlation Coefficient: 0.9951

Linearity

• 4 - 150 mg/dL (1.4 - 53.6 mmol/L)

No Significant Interferences Up to Levels Indicated

- Hemoglobin: 1000 mg/dL (155 µmol/L)
- Bilirubin: 40 mg/dL (684 µmol/L)
- Intralipid: 1000 mg/dL (3000 mg/dL (33.9 mmol/L) Simulated Triglycerides)

Reference Range⁽¹⁾

- Urea: 11 37 mg/dL (3.9 13.2 mmol/L)
- Urea Nitrogen: 5 17 mg/dL (1.8 6.1 mmol/L)

(a) The performance of this method (y) was compared with the performance of a similar method (x) on a Roche/Hitachi® 717 analyzer.

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Ordering information

	Configuration	Catalog Number
UREA L3K®	R1 6 x 30mL	283-30
DC-Cal Calibrator	5 x 3mL	SE-035
DC-Trol Level 1	10 x 5mL	SM-052
DC-Trol Level 2	10 x 5mL	SM-056

(1) Kaplan, L.A. and Pesce, A.J., Clinical Chemistry - Theory, Analysis, and Correlation, Third Edition. Mosby Year-Book Inc., St. Louis, p. 500 (1996).



Experience + Technology + Portfolio + Support = CHEMABILITY

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