

# **HOMOCYSTEINE 2 REAGENT ENZYMATIC ASSAY**

***Dual Vial Liquid Stable***

Diazyme's Homocysteine 2 Reagent Enzymatic Assay features convenient ready to use reagent, calibrators and controls for the quantitative determination of total L-homocysteine in serum or plasma. Diazyme's proprietary Enzyme Cycling methodology is an excellent choice for cost conscious laboratories of all sizes due to a wide variety of instrument specific packaging options. The assay requires minimal patient sample and provides fast, accurate and precise results. A wide variety of reliable instrument parameters means the assay is readily available for installation on most automated clinical chemistry analyzers.

## ***DIAZYME HOMOCYSTEIN 2 REAGENT ASSAY ADVANTAGES***

- Award winning Homocysteine recognized by the American Association of Clinical Chemistry (AACC) for outstanding contribution to scientific research
- Innovative enzyme cycling based technology for accurate and reliable results
- Excellent correlation to HPLC and immunochemical methods
- No "carry over" issues with iron or lipase reagents
- Test renal patients with confidence since there is no interference from cystathionine which affects some other less specific methods
- Wide range of instrument parameters available for facilitating and simplifying implementation
- Liquid stable format requires no reagent preparation saving time and reducing sample handling

## ***REGULATORY STATUS***

510(k) Cleared

Health Canada Registered



## ***AVAILABLE INSTRUMENT SPECIFIC PACKAGING***

- Roche
- Beckman
- Hitachi
- AU Series



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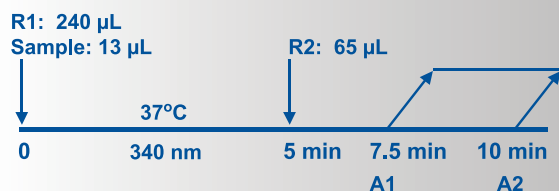
Dual Vial Liquid Stable



## ASSAY SPECIFICATIONS

<b>Method</b>	Diazyme Patented Enzyme Cycling
<b>Sample Type &amp; Volume</b>	<ul style="list-style-type: none"><li>• Serum</li><li>• Plasma</li><li>- EDTA</li><li>- Li-heparin</li></ul> Sample Volume 13 µL
<b>Method Correlation</b>	N = 40 y-intercept = 1.05 Slope = 0.94 R <sup>2</sup> = 0.99
<b>Linear Range</b>	Up to 50 µmol/L
<b>LOD</b>	0.4 µmol/L
<b>Calibration Levels</b>	5-Point Calibration
<b>Reagent On-Board Stability</b>	Opened: At least 60 days (Analyzer Dependent)

### Homocysteine 2 Reagent Assay Procedure\*



\*Analyzer Dependent

### Two Reagent System

Parameter questions for Enzymatic Homocysteine 2 Reagent Assay should be addressed to Diazyme technical support. Please call 858.455.4768 or email [support@diazyme.com](mailto:support@diazyme.com)

1. Vilaseca et al. *Clin. Chem.* 43: 690-692 (1997)
2. Faure-Delanef et al. *Am. J. Hum. Genet.* 60: 999-1001 (1997)

## ASSAY PRECISION

Precision studies were conducted according to the NCCLS EP-5 protocol. Four HCY serum samples containing 7.0, 12.0, 15.6, and 29.0 µM HCY were tested.

HCY Concentration	7 µM	12 µM	15.6 µM	29 µM
<b>Within-Run Imprecision CV% N = 20</b>	4.5	1.87	3.04	2.4
<b>Total Imprecision CV% N = 30</b>	5.87	4.88	5.51	2.57

## ASSAY INTERFERENCE

An interference study was performed by testing a serum sample spiked with varied concentrations of endogenous substances. The following substances normally present in the serum produced less than 10% deviation when tested at the stated concentrations:

Bilirubin:	40 mg/dL
Triglycerides:	1000 mg/dL
Hemoglobin:	500 mg/dL
Bilirubin Conjugate:	40 mg/dL
Ascorbic Acid:	10 mM
Cystathionine:	100 µM**

\*\*The concentrations tested are about 5-10 times higher than the normal range of serum levels.

## REFERENCE RANGE

In most of the U.S. clinical laboratories, 15 µmol/L is used as the cut-off value for normal level of Hcy for adults.<sup>1-2</sup> In Europe, 12 µmol/L is used as the cut-off value. However, each laboratory is recommended to establish a range of normal values for the population in their region.

## DIAZYME LABORATORIES

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