Enzymatic creatinine assays have less interference than older Jaffe creatinine assays. A study from leading clinical journal shows that interference with Jaffe creatinine assays may lead to inaccuracies in estimated glomerular filtration rates that are clinically important, especially in children and neonates.¹

Diazyme’s Creatinine Liquid Reagent assay is intended for the in vitro quantitative determination of creatinine in serum and urine. The assay is cost effective and provides outstanding reagent stability combined with the added convenience of instrument specific packing for several major instrument families.

**DIAZYME CREATININE LIQUID REAGENT ASSAY ADVANTAGES**

- Accurate measurement of creatinine with enzymatic method
- Reduced interferences and no cuvette staining as seen in Jaffe method
- Assay is traceable to NIST material (IDMS)
- Measuring range:  
  - Serum: 0.14-13.56 mg/dL  
  - Urine: 0.14-141.25 mg/dL
- Liquid stable reagent, calibrator and controls requires no reagent preparation
- Wide range of instrument parameters available for simplifying implementation

**REGULATORY STATUS**

510(k) Cleared; EU: CE (IVD)
**ASSAY SPECIFICATIONS**

<table>
<thead>
<tr>
<th>Method</th>
<th>Enzymatic Assay</th>
</tr>
</thead>
</table>
| Sample Type & Volume | • Serum  
• Urine  
Sample Volume 8 μL |
| Method Correlation | Serum:  
N = 55  
y-intercept = 0.0643  
Slope = 0.9467  
R² = 0.9981  

Urine:  
N = 51  
y-intercept = -0.0518  
Slope = 1.0002  
R² = 0.9968 |
| Linearity | Serum:  
0.14 - 13.56 mg/dL  
(12 - 1200 μmol/L)  

Urine:  
0.14 - 141.25 mg/dL  
(12 - 12500 μmol/L) |
| LOD | 12 μmol/L (0.14 mg/dL) |
| Calibration Levels | 1-Point Calibration |
| Traceability | Standard traceable NIST’s SRM 914a |
| Reagent On-Board Stability | Opened:  
4 weeks when stored at 2-8°C |

**Creatinine Assay Procedure**

1. **R1**: 270 μL  
Sample: 8 μL
2. **R2**: 90 μL
3. 37°C
4. 5 min  
550 nm  
10 min
5. A1  
A2

*Analyzer Dependent

Parameter questions for Enzymatic Creatinine Assay should be addressed to Diazyme technical support. Please call 858.455.4768 or email support@diazyme.com

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**ASSAY PRECISION**

The assay was evaluated according to Clinical Laboratory Standards Institute EPS-A guidelines. Four serum specimens were tested on a Hitachi 917 twice daily, in duplicates over 20 days.

<table>
<thead>
<tr>
<th>Serum Testing</th>
<th>Within-Run Precision (80 Data Points)</th>
</tr>
</thead>
</table>
| Mean mg/dL (μM) | 0.74 (65.4)  
1.38 (122.3)  
4.04 (357.5)  
10.28 (908.7) |
| SD mg/dL (μM) | 0.015 (1.3)  
0.015 (1.37)  
0.029 (2.54)  
0.015 (1.3) |
| CV% | 2.1%  
1.1%  
0.7%  
0.1% |

**Total Precision (80 Data Points)**

<table>
<thead>
<tr>
<th>Serum Testing</th>
<th>Mean mg/dL (μM)</th>
<th>SD mg/dL (μM)</th>
<th>CV%</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.75 mg/dL (66.3 μM)</td>
<td>0.74 (65.4)</td>
<td>0.015 (1.3)</td>
<td>2.1%</td>
</tr>
<tr>
<td>1.41 mg/dL (125 μM)</td>
<td>1.38 (122.3)</td>
<td>0.015 (1.37)</td>
<td>1.1%</td>
</tr>
<tr>
<td>4.11 mg/dL (346 μM)</td>
<td>4.04 (357.5)</td>
<td>0.029 (2.54)</td>
<td>0.7%</td>
</tr>
<tr>
<td>10.28 mg/dL (908.7 μM)</td>
<td>10.28 (908.7)</td>
<td>0.015 (1.3)</td>
<td>0.1%</td>
</tr>
</tbody>
</table>

**Urine Testing**

<table>
<thead>
<tr>
<th>Level 1</th>
<th>Mean mg/dL (μM)</th>
<th>SD mg/dL (μM)</th>
<th>CV%</th>
</tr>
</thead>
<tbody>
<tr>
<td>29.09 (2572)</td>
<td>0.1 (8.84)</td>
<td>0.36%</td>
<td></td>
</tr>
<tr>
<td>87.1 (7711)</td>
<td>0.27 (23.60)</td>
<td>0.31%</td>
<td></td>
</tr>
<tr>
<td>196.7 (17407)</td>
<td>0.90 (79.71)</td>
<td>0.46%</td>
<td></td>
</tr>
</tbody>
</table>

**Urine Testing**

<table>
<thead>
<tr>
<th>Total Precision (20 Data Points)</th>
<th>Level 1</th>
<th>Level 2</th>
<th>Level 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean mg/dL (μM)</td>
<td>29.86 (2640)</td>
<td>49.86 (4592)</td>
<td>195 (17265)</td>
</tr>
<tr>
<td>SD mg/dL (μM)</td>
<td>0.79 (69.8)</td>
<td>0.67 (59.2)</td>
<td>1.19 (105.2)</td>
</tr>
<tr>
<td>CV%</td>
<td>2.64%</td>
<td>0.76%</td>
<td>0.60%</td>
</tr>
</tbody>
</table>

**ASSAY INTERFERENCE**

Interference for the Diazyme Creatinine Assay was evaluated on the Hitachi 917. The following substances normally present in serum produced less than 10% deviation at the listed concentrations:

- Triglyceride: up to 1000 mg/dL
- Ascorbic Acid: up to 10 mM
- Bilirubin (Conjugate): up to 30 mg/dL
- Bilirubin: up to 40 mg/dL
- Hemoglobin: up to 500 mg/dL

The following substances normally present in urine produced less than 10% deviation at the listed concentrations:

- Triglyceride: up to 1000 mg/dL
- Ascorbic Acid: up to 10 mM
- Bilirubin (Conjugate): up to 40 mg/dL
- Bilirubin: up to 40 mg/dL
- Hemoglobin: up to 1000 mg/dL

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