### LYPHOZYME

# **UREA (BERTHELOT METHOD)**

Code	Product Name	Pack Size
LP007A	Lyphozyme Urea	100 ml
LP007B	Lyphozyme Urea	200 ml
LP007C	Lyphozyme Urea	500 ml
LP007E	Lyphozyme Urea	1000 ml

#### Intended Use

Diagnostic reagent for quantitative in vitro determination of Urea in human serum, plasma and urine.

#### **Clinical Significance**

Urea is the end product of the protein metabolism. It is synthesized in the liver from the ammonia produced by the catabolism of amino acids. It is transported by the blood to the kidneys from where it is excreted. Increased levels are found in renal diseases, urinary obstructions, shock, congestive heart failure and burns. Decreased levels are found in liver failure and pregnancy.

#### Principle

Urease hydrolyzes urea to ammonia and  $\text{CO}_2$ . The ammonia further reacts with a phenolic chromogen and hypochlorite to form a green coloured complex. Intensity of the colour formed is directly proportional to the amount of urea present in the sample.

#### Reaction:

### Reagent Composition

Reagent 1: Urea Enzyme Reagent

Ready to use

Reagent Preparation

Reagent is liquid, ready to use.

### Materials Required But Not Provided

- Clean & Dry container.
- Laboratory Glass Pipettes or Micropipettes & Tips
- $\ \, {\sf Colorimeter}\, {\sf or}\, {\sf Bio-Chemistry}\, {\sf Analyzer}.$

### Samples

Serum, plasma, Urine. Dilute urine specimen 1 + 49 with distilled water before the assay (results  $\times$  50). Urea is reported to be stable in serum for 5 days at +2-+8°C.

# $Preparation \, of \, Reagent \, \& \, Stability$

Bring all the reagent to room temperature.



#### **Working Reagent**

- Dissolve the enzyme reagent (Reagent 1) in Deionised water as per volume indicated on the vial.
- Reagent 2 is ready to use. Allow the reagents to stand for 5 minutes at R.T for equilibriation.
- Store the kit at +2°-+8 °C away from light.
- Working reagent after reconstitution is stable for 75 days when stored at +2 +8°C.
- Reagent 2 Once open is stable for six months when stored At+2-+8  $^{\circ}\text{C}.$

#### Stability And Storage

The unopened reagents are stable till the expiry date stated on the bottle an it label when stored at +2-+8°C.

#### Specimen Collection And Handling

Use unheamolytic serum or plasma (EDTA, heparin) and urine. It is recommended to follow NCCLS procedures (or similar standardized conditions).

### Stability In Serum:

 $5 \, \text{days}$  :  $at + 2 - +8 \,^{\circ}\text{C}$  Discard contaminated specimens.

#### Calibration

Calibration with the Urea standard provided in the kit is recommended.

### **Quality Control**

It's recommended to run normal and abnormal control sera to validate reagent performance.

### **Expected Values**

 Serum / Plasma
 : 10 - 50 mg/dl

 Urine
 : 15 - 30 mg/24 hrs

It is recommended that each laboratory verify this range or derives reference interval for the population it serves.

# Performance Data

Data contained within this section is representative of performance on Beacon system. Data obtained in your laboratory may differ from these values.

### Precision

Intra-assay precision Within run (n=20)	Mean (mg/dl)	SD (mg/dl)	CV (%)
Sample 1	45.09	2.24	2.27
Sample 2	124.79	2.75	4.98

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Inter-assay precision	Mean	SD	CV
Run to run (n=20)	(mg/dl)	(mg/dl)	(%)
Sample 1	42.2	1.86	4.41

#### Comparison

A comparison between Beacon Urea (y) and a commercially available test (x) using 20 samples gave following results :

y = 0.99 x - 0.4829 mg/dl

r = 0.999

#### Interferences

- Lipemia (intralipid 20 g/L) does not interfere.
- Bilirubin (40 mg/dL) does not interfere Hemoglobin (>2 g/L) may affect the results.
- Other drugs and substances may interfere.

### Warning And Precautions

For in vitro diagnostic use. To be handled by entitled and professionally educated person.

### Waste Management

 $Please\ refer\ to\ local\ legal\ requirements.$ 

#### Procedure:

(S) and Test (T):

Addition Sequence	Reagent Blank	Standard	Sample	
Working Reagent	1000 μΙ	1000 μΙ	1000 μΙ	
Standard	-	10 µl	1	
Sample	-	-	10 μΙ	
Mix and incubate 5 min. At +37°C. Then add				
Chromogen Reagent	1000 μΙ	1000 μΙ	1000 μΙ	

Mix and incubate for 5 minutes at  $+37^{\circ}$ C. Measure the absorbance of standard (Abs. S) and test (Abs. T) against reagent blank at 630 nm within 60 minutes.

## Calculation

Urea (mg/dl) = 
$$\frac{\text{Abs. of T}}{\text{Abs. of S}} \times 40$$

### **Assay Parameters For Photometers**

Mode	End point
Wavelength 1 (nm)	630
Sample Volume (µl)	10
Reagent Volume (μΙ)	1000
Incubation time (min.)	5 + 5
Incubation temp. (°C)	37
Normal Low (mg/dl)	10
Normal High (mg/dl)	50
Linearity Low (mg/dl)	1
Linearity High (mg/dl)	300
Standard Concentration	40 mg/dl
Blank with	Reagent
Unit	mg/dl

### References

- 1. Leonard, P. L., Persaud, J., Motwani, R.: Clin. Chim. Acta 35, 409, 1971.
- 2.Tietz Textbook of Clinical Chemistry and Molecular diagnostics. Burtis, C.A., Ashwood, E.R., Bruns, D.E.; 5th edition, WB Saunders Company, 2012.

### Symbols Used On Labels

REF

Catalogue Number

Manufacturer

Lot Number

 $\mathbf{i}$ 

See Instruction for Use

Storage Temperature

CONT

**Expiry Date** 

Content

IVD

In Vitro Diagnostics



