# **AMMONIA SYSTEM PACK**

# (KINETIC METHOD)

B Auto 200, Unicorn 230, Unicorn 120 & Bonavera Chem 200, Beaconic chem 200, Beaconic B200, Beaconic analyzer 120, Bonavera chem 100 (Fully Auto Biochemistry Analyzer)

| Code  | Product Name        | Pack Size     |
|-------|---------------------|---------------|
| BA204 | Ammonia System Pack | 1x16 + 1x4 ml |

#### INTENDED USE

The reagent kit is intended for the  ${\it In \ Vitro}$  quantitative determination of Ammonia.

#### SUMMARY

Ammonia (NH<sub>3</sub>) is a reagent kit used for the quantitative determination of ammonia in plasma, based on enzymatic method using glutamate dehydrogenase (GLDH) enzyme.

#### PRINCIPLE

Ammonia reacts with  $\alpha\text{-ketoglutarate}$  to form glutamate in presence of glutamate dehydrogenase. NADH is oxidized to NAD+ in this reaction, which is measured as decrease in absorbance at 340 nm. The rate of decrease in absorbance at 340 nm is directly proportional to plasma ammonia concentration.

 $NH_3 + \alpha$ -ketoglutarate +  $NAD_{GLDH}$  Glutamate +  $NAD_{GLDH}$ 

### REAGENT COMPOSITION

Reagent 1: Ammonia Reagent 1
Reagent 2: Ammonia Reagent 2

Reagent 3 : Ammonia Standard (500  $\mu g/dl$ )

# REAGENT PREPARATION

Ready to use

# STORAGE & STABILITY

The reagent kit should be stored at  $+2 - +8^{\circ}C$  and is stable till the expiry date indicated on the label.

On board stability: Min. 30 days if refrigerated (+8-+14°C) and not contaminated.

# SPECIMEN COLLECTION AND HANDLING

EDTA plasma or Heparinized plasma.

Blood is collected from a stasis-free vein and stored in an ice bath.

The plasma is then separated within 30 min. Ammonia assay should be carried out immediately. The plasma may be stored for 2 hour at  $+2^{\circ}\text{C}-+8^{\circ}\text{C}$ .

### LINEARITY

This procedure is linear up to 1500  $\mu$ g/dl. If value exceeds this limit dilute the sample with normal saline (NaCl 0.9%) and repeat the assay Multiply result by dilution factor.

### CALIBRATION

Calibration with the Ammonia Standard provide in the kit is recommended.



### QUALITY CONTROL

For accuracy, it is advised to run known controls with each assay.

#### LIMITATION

- 1. Anticoagulants having ammonium ions should not be used because of extreme sensitivity of the color reaction to ammonia.
- 2. Reaction is linear up to 1500  $\mu$ g/dl. For higher values, dilute the sample with normal saline and perform the assay. Multiply the final result by dilution factor to get the real value.
- 3. The working reagent is considered unsatisfactory and should not be used if the absorbance is less than 0.700 at  $340\,\mathrm{nm}$  against distilled water.
- $4.\,Do\,not\,use\,strongly\,hemolysed\,samples.$

### **UNIT CONVERSION**

Ammonia (µg/dl)x 0.588= Ammonia (µmol/L)

### **EXPECTED VALUES**

Plasma: 17-90 µg/dl

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. Data obtained in your laboratory may differ from these values.

 $\begin{array}{ll} \mbox{Limit of quantification} & : 10 \, \mu \mbox{g/dl} \\ \mbox{Linearity} & : 1500 \, \mu \mbox{g/dl} \\ \mbox{Measuring range} & : 10 \cdot 1500 \, \mu \mbox{g/dl} \\ \end{array}$ 

Precision

| Intra-assay precision<br>Within run (n=20) | Mean<br>(U/L) | SD<br>(U/L) | CV<br>(%) |
|--|---------------|-------------|-----------|
| Sample 1                                   | 105           | 1.96        | 1.87      |
| Sample 2                                   | 152           | 1.46        | 0.96      |
| Inter-assay precision<br>Run to run (n=20) | Mean<br>(U/L) | SD<br>(U/L) | CV<br>(%) |
| Sample 1                                   | 97            | 1.23        | 1.27      |

# COMPARISON

A comparison between Ammonia System Pack (y) and commercially available tests (x) using 40 samples gave following results:

y = 1.0174x - 1.3868

r=0.999

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### **INTERFERENCES**

Following substances do not interfere:

Ascorbic acid UP TO 40 mg/dl ,Bilirubin up to 20 mg/dl,triglyceride up to 700 mg/dl & hemolysed samples should not be used as erythrocytes contain level of ammonia 3 times that of plasma .

### WARNING AND PRECAUTIONS

For *in vitro* diagnostic use. To be handled by entitled and professionally educated person. MSDS will be provided on request.

### WASTE MANAGEMENT

Please refer to local legal requirements.

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| Ammonia        |  |
|----------------|--|
| Ammonia        |  |
| 340 nm         |  |
| 630 nm         |  |
| Fixed Time     |  |
| 20             |  |
| 30             |  |
| 1              |  |
| μg/dL          |  |
| 10             |  |
| 1500 μΙ        |  |
| 20 μΙ          |  |
| 160 μl         |  |
| 40 μΙ          |  |
| -              |  |
| 1500 μg/dl     |  |
| -              |  |
| 2 Point linear |  |
| 2              |  |
| Water          |  |
| 0.00           |  |
| 500 μg/dl      |  |
|                |  |

# NOTES

Clinical diagnosis should not be made on findings of a single test result, but should integrate both clinical and laboratory data.

# **BIBLIOGRAPHY**

- 1. Dewan, J.G., Biochem J., 1938; 32:1378.
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# Symbols Used On Labels

REF

Catalogue Number



Manufacturer



See Instruction for Use



Lot Number



Content



Storage Temperature



**Expiry Date** 



In Vitro Diagnostics

BEA/24/AMM/SB/IFU Ver-02 05/10/2024



