

Ammonia Colorimetric Assay Kit

Product Information

Common Name

Ammonia

Cat.No. Kit-0085

Product Overview

Ammonia Assay Kit provides a rapid, simple and sensitive colorimetric method for the quantitation of ammonia/ammonium concentration in biological samples such as serum, plasma and urine. The assay is based on a chromogenic reaction to produce a blue product upon the reaction of ammonia with our sensor. The intensity of color produced is proportional to the concentration of ammonia in the sample, which can be measured at absorbance 660-670 nm. This kit provides a simple assay to detect as little as 4 μ M ammonia in a 150 μ L assay volume.

Description

Ammonia (NH_3)/ammonium salt is an important source of nitrogen. In human, ammonia is produced through amino acid deamination and converted to urea through the urea cycle, which is then eliminated in urine. Ammonia levels in the blood rise when the liver is not able to convert ammonia to urea. Hyperammonemia, the elevated levels of ammonia in the blood, has been found in liver dysfunction (cirrhosis), while hypoammonemia has been associated with defects in the urea cycle enzymes (e.g. ornithine transcarbamylase). An ammonia test is usually important in clinical diagnostics to check how well the liver is working or the success of treatment for severe liver disease.

Usage

For research use only (RUO)

Storage

Store at -20°C and avoid light.

Size

200 tests

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Kit Components

Component A: Assay Buffer I 1 bottle (10 mL)

Component B: Assay Buffer II 1 bottle (10 mL)

Component C: Ammonium Chloride Standard (1.0 M) 1 vial (0.2mL)

Detection method Colorimetric

Assay Protocol

Note: Thaw all the kit components to room temperature before starting the experiment.

1. Prepare serial dilutions of Ammonium Chloride (0 to 1mM) solutions:

1.1 Add 1 μ L of 1.0 M Ammonium Chloride Standard (Component C) to 999 μ L DPBS to generate 1.0 mM standard ammonium chloride solution.

1.2 Take 300 μ L of 1.0 mM standard to perform 1:3 serial dilutions to get 300, 100, 30, 10, 3, 1, and 0 μ M standard ammonium chloride solutions.

1.3 Add Ammonium Chloride Standards and ammonia containing test samples into a 96-well clear bottom microplate as described in Tables 1 and 2.

Table 1. Layout of standards and test samples in a clear bottom 96-well microplate:

BL BL TS TS

AS 1 AS 1

AS 2 AS 2

AS 3 AS 3

AS 4 AS 4

AS 5 AS 5

AS 6 AS 6

AS 7 AS 7

Note: AS= Ammonium Chloride Standards, BL=Blank Control, TS=Test Samples.

Table 2. Reagent composition for each well:

Ammonium Chloride Standard Blank Control Test Sample

Serial dilutions*: 50 μ L DPBS: 50 μ L 50 μ L

*Note: Add the serially diluted ammonium chloride standards from 1 to 1000 μ M into wells from AS1 to AS7 in duplicate.

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2. Run Ammonium Chloride Assay:

2.1 Add 50 μ L of Assay Buffer I (Component A) to each well of the Ammonium Chloride Standard, blank control, and test samples (see Step 1.3) so that the total assay volume is 100 μ L/well.

Note: For a 384-well plate, add 25 μ L sample, 25 μ L of Assay Buffer I per well.

2.2 Incubate the reaction for 5 minutes at room temperature.

2.3 Add 50 μ L of Assay Buffer II (Component B) to each well so that the total assay volume is 150 μ L/well.

Note: For a 384-well plate, add 25 μ L Assay Buffer II (Component B) to each well.

2.4 Incubate at room temperature for 30-60 minutes, and monitor the absorbance increase at 660-670 nm using an absorbance microplate reader.

Note: The color turns to yellow after Assay Buffer II (Component B) is added, and the wells with Ammonium Chloride Standard or samples will show bluish green color after incubation. The intensity of the color will reach the maximum in 30-60 minutes.

Analysis

The absorbance in blank wells (with DPBS only) is used as a control, and is subtracted from the values for those wells with Ammonium Chloride Standards.

Note: The absorbance background is subtracted from the absorbance intensity value of the wells for each data point.
