

## DATASHEET

### BCA Protein Assay Kit

## Product overview

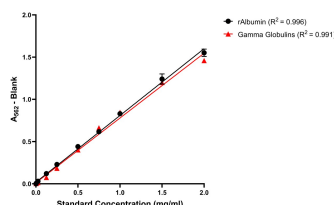
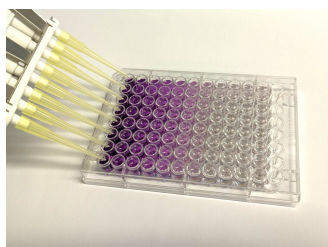
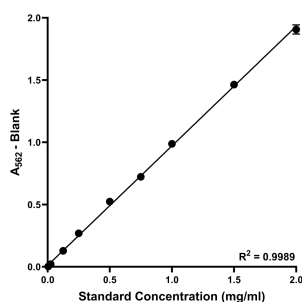
<b>Name</b>	BCA Protein Assay Kit
<b>Cat No</b>	HB7109
<b>Biological description</b>	Simple, rapid, detergent tolerant kit for determining the concentration of proteins in solution. This kit is optimized to measure protein concentrations from 0.02 to 2mg/mL.

Key features of the BCA Protein Assay Kit:

- **Detergent compatible** - compatible with detergent concentrations up to 5%
- **Quick** - 30 minute incubation time means total assay time of around 45 minutes.
- **Wide assay range** - can measure protein concentrations from 0.02 to 2 mg/ml
- **Stability** - kit is stable at room temperature

<b>Biological action</b>	Kit
<b>Description</b>	Simple, rapid, detergent tolerant kit for measuring the concentration of proteins in solution (0.02 to 2mg/mL).

## Images



## Biological Data

### Application notes



#### BCA Assay Protocol PDF

Before using the kit for the first time add 5ml of buffer to the protein standard to create a 2mg/ml solution. Use this to create a dilution series of 0, 0.001, 0.005, 0.025, 0.125, 0.25, 0.5, 0.75, 1, 1.5 and 2mg/ml samples (please see the [pdf protocol](#) for precisely how to do this). For best accuracy use the same buffer as your proteins of interest. The assay can either be carried out in microplate or tube format.

#### Microplate Protocol

1. Add 25µl of each standard to a microplate well alongside the unknown samples. Ideally this

- should be carried out in triplicate.
2. Prepare the working reagent by mixing 50 parts reagent A with 1 part reagent B. For easy calculation add 200µl reagent A and 4µl reagent B per well.
  3. Add 200µl of working reagent to each well and incubate for 30 minutes at 37°C
  4. Let the plate cool to room temperature for 1-2 minutes.
  5. Measure absorbance at 562nm using a microplate reader. This should be ideally carried out within 40 minutes of the start of the assay to maximise accuracy.
  6. Subtract the absorbance of the 0mg/ml samples from all measurements then construct a standard curve using the sample data.
  7. Use the standard curve to calculate the protein concentration of the unknown samples.

#### Tube Protocol

1. Add 100µl of each standard to labelled test-tubes alongside the unknown samples. Ideally this should be carried out in triplicate.
2. Prepare the working reagent by mixing 50 parts reagent A with 1 part reagent B. For easy calculation add 2ml reagent A and 40µl reagent B per tube.
3. Add 2ml of working reagent to each tube and mix thoroughly.
4. Incubate for 30 minutes at 37°C. Alternatively the incubation can be carried out for 60 minutes at room temperature.
5. Cool all tubes to room temperature for 1-2 minutes.
6. Measure absorbance at 562nm using a spectrophotometer ensuring that all measurements are made within 10 minutes. Prior to taking the measurements ensure the spectrophotometer is blanked using a cuvette filled with dH<sub>2</sub>O.
7. Subtract the absorbance of the 0mg/ml samples from all measurements then construct a standard curve using the sample data.
8. Use the standard curve to calculate the protein concentration of the unknown samples.

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## Solubility & Handling

### Storage instructions Important

Room temperature

This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

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## Chemical Data

### Kit contents

- Reagent A
  - Reagent B
  - BCA Protein Standard
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