

Assay kits

BCA Protein Assay Kit

Bicinchoninic Acid Protein Assay Kit (BCA)

Detergent compatible assay reagent to measure protein concentration

Protocol

IMPORTANT NOTES – Before you begin

- ✓ BCA assay kit offers a fast colorimetric detection and quantification method of total protein content even in the presence of detergents.
- ✓ This kit is based on the reduction of Cu²⁺ to Cu¹⁺ by protein in alkaline solution; monovalent copper ions produced are detected in a concentration-dependent manner. Bicinchoninic acid (BCA) chelates with the reduced copper Cu¹⁺ and form a water-soluble purple reaction complex that exhibits a strong absorbance at 562 nm.
- ✓ Absorbance is linear over a wide range of protein concentrations between 25-2000 µg/mL.
- ✓ Protein concentrations are estimated with reference to a commonly used protein standard; the kit also includes Bovine Serum Albumin (BSA) at 1.5 mg/mL as a protein standard for a convenient preparation of protein concentration standard curves.

NOTE: During long term storage or upon shipping in cold weather, Reagent A or B may precipitate; we recommend to gently warming and stirring the solution to dissolve precipitates.

For additional information and protocols
(optimization, scaling, co-transfection...) tips, troubleshooting or other applications

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Any questions?

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BCA Protein Assay Kit | Specifications

Package content	BCA2500 BCA Reagent A (500mL), Copper Solution B (15mL) BSA 5x1mL (1500µg/mL) Number of assays (96-well plate): 2500 Number of assays (1mL cuvettes): 500
Shipping conditions	The kit is shipped at Room Temperature.
Storage conditions	Store at 4°C
Shelf life	1 year from the date of purchase when properly stored and handled
Important notice	For research use only. Not for use in diagnostic procedures.

Application and Protocols

1. General Considerations

- We recommend using a 50:1 ratio between Reagent A and B. Different ratios were demonstrated to impair protein detection.
- Detection of small protein amount can be performed by increasing time of incubation at 37°C.

2. Solution preparation

Standard Solution.

Prepare a range of concentrations from 1500 µg/mL to 50 µg/mL by serial dilutions according to the protocol below.

Prepare 7 tubes containing H₂O or sample buffer, refer to the illustration and table below:

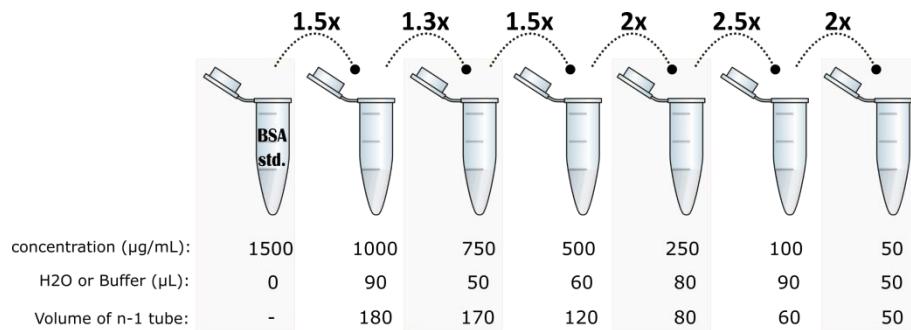


Figure 1: Recommendations for performing serial dilutions using standard BSA.

Vial	Volume of BSA (µL)	Volume of Diluent (µL)	Final BSA concentration (µg/mL)
1	280 of 1500 µg/mL stock	0	1500
2	180 of vial 1	90	1000
3	170 of vial 2	50	750
4	120 of vial 3	60	500
5	80 of vial 4	80	250
6	60 of vial 5	90	100
7	50 of vial 6	50	50
8 (blank)	0		0

Table 1: Volumes to consider for preparing BSA standards

Working Solution (WR).

Prepare a working solution by mixing 50 parts of BCA reagent A with 1 part of copper solution B.

Refer to the table 1 below to prepare solution for 1 mL cuvette or 96-well plate (200 μ L). .

Number of assays 96-well plate (200 μL)	Cuvette (1mL)	Volumes		
		BCA Reagent A	Copper solution B	Total Working solution (WR)
1	-	196 μ L	4 μ L	0.2 mL
5	1	980 μ L	20 μ L	1.0 mL
25	5	4.9 mL	0.1 mL	5.0 mL
50	10	9.8 mL	0.2 mL	10.0 mL
100	20	19.6 mL	0.4 mL	20.0 mL

Table 2: Volumes to consider for preparing Working Solution

3. Microplate procedure – General protocol for 96-well plate

1. Add **25 μ L** of each standard point to a 96-well plate
2. Add **25 μ L** of sample to 96-well plate.

NOTE: We recommend performing at least duplicate; for concentrated samples, dilute 5x or 10x your sample in sample buffer or in H₂O.

3. Prepare a blank with **25 μ L** of sample buffer or H₂O.
4. Add **200 μ L** of Working Solution to each sample, standard and blank wells
5. Incubate **1 H at 37°C.**
6. Read absorbance at **562 nm**

NOTE: Wavelengths from 540-595 nm can also be used.

7. Subtract background fluorescence of the blank from all other values.
8. Calculate the amount of protein present in samples.

4. General protocol for 1mL Test Tubes

1. Add **100 μ L** of each standard point into 1.5 mL tube
2. Add **100 μ L** of sample into 1.5 mL tube
3. Prepare a blank with **100 μ L** of sample buffer or H₂O.
4. Add **1 mL** of Working Solution to each sample, standard and blank wells
5. Incubate **1 H at 37°C.**
6. Read absorbance at **562 nm**

NOTE: Wavelengths from 540-595 nm can also be used.

7. Subtract background fluorescence of the blank from all other values.
8. Calculate the amount of protein present in samples.

5. Results

Standard curve

Create a standard curve by plotting Absorbance over protein standard amount ($\mu\text{g/mL}$).

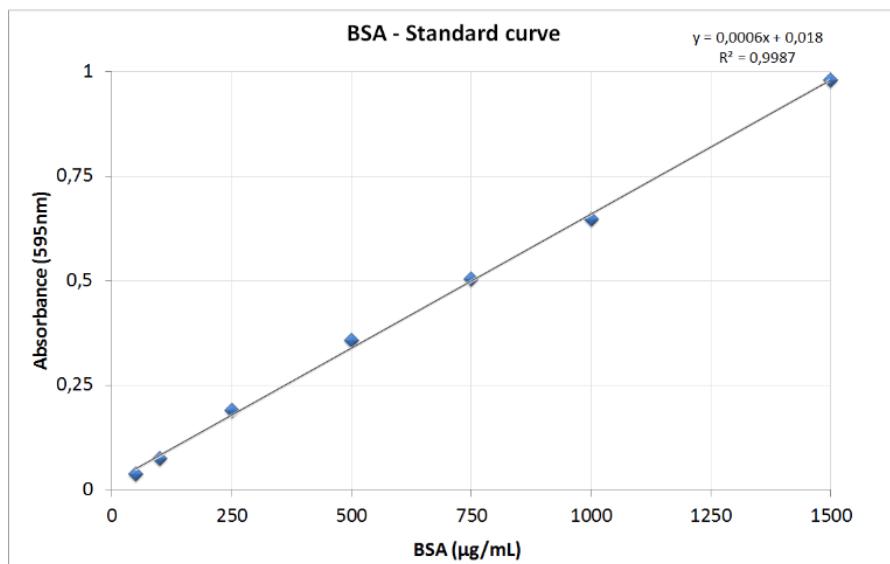


Figure 2: Standard curve realized using serial dilutions of standard BSA

Use the standard curve to determine the sample protein concentration.

6. Interfering compounds and compatible substances

The BCA protein assay kit is detergent-tolerant as demonstrated by the experiment below.

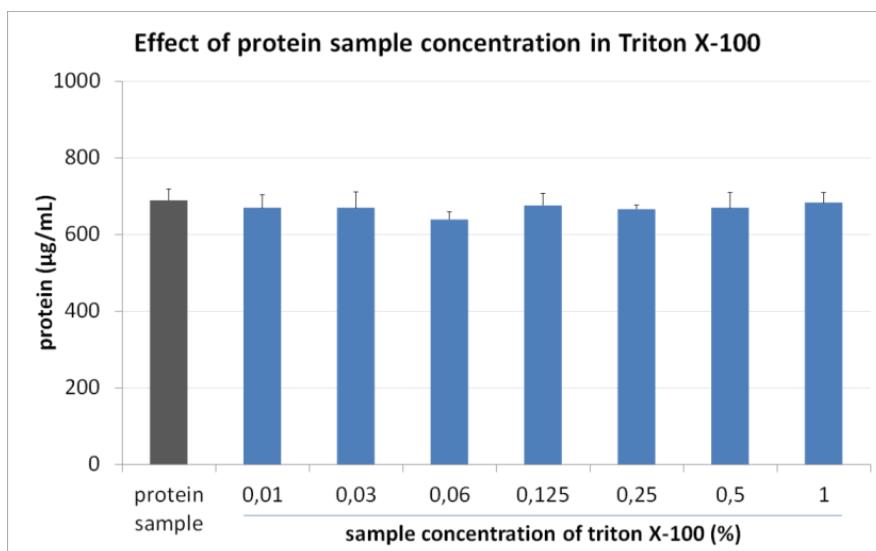


Figure 3: Triton X-100 concentration effect on BCA quantification

Results demonstrated that, as opposed to other quantification methods (Bradford), up to 1% triton X-100 in sample does not interfere with protein quantification using BCA assay kit

The following table list compounds that are compatible with BCA – Protein Assay Kit and their maximal non-interfering concentrations:

Compound	Maximal non interfering concentration
2-Mercaptoethanol	0.01 %
Acteone	10 %
CaCl	10 mM
CHAPS	5 %
DMF	10 %
DMSO	10 %
EDTA	50 mM
Ethanol	10 %
Glycerol	10 %
HCl	100 mM
Imidazole, pH 7.0	50 mM
NaCl	10 mM
MOPS	100 mM
NP40	5 %
SDS	5 %
Sucrose	40 %
TBP	10 mM
TCEP	2 mM
TFA	0.005 %
Thiourea	500 mM
Tris	0.5 M
Triton™ X-100	5 %
TWEEN®	5 %
Urea	3 M

Table 3: Compatible compounds and their maximal non-interfering concentrations.

Substances such as Asorbic acid, EGTA, Iron, Hydrogen Peroxide, Tyrosine, Uric Acid, Phenol Red, Creatinin are known to interfere with BCA – Protein Assay Kit, even when used at small concentrations.

These lists are not exhaustive. For an optimal reading, we recommend assaying the protein of interest in ultrapure water alone; dialysis or protein precipitation may also be used to remove interfering substance.

Additional products for Protein Dosage

- **Bradford Protein Assay** for determining the concentration of proteins
- **FluoProdige Assay Kit** for protein and peptide quantification

Purchaser Notification

Limited License

The purchase of the BCA assay kit grants the purchaser a non-transferable, non-exclusive license to use the kit and/or its separate and included components (as listed in this protocol). This reagent is intended for in-house research only by the buyer. Such use is limited to the transfection of nucleic acids as described in the product manual. In addition, research only use means that this kit and all of its contents are excluded, without limitation, from resale, repackaging, or use for the making or selling of any commercial product or service without the written approval of OZ Biosciences. Separate licenses are available from OZ Biosciences for the express purpose of non-research use or applications of the BCA assay kit. To inquire about such licenses, or to obtain authorization to transfer or use the enclosed material, contact us at OZ Biosciences. Buyers may end this License at any time by returning all BCA assay kit reagents and documentation to OZ Biosciences, or by destroying all BCA assay kit components. Purchasers are advised to contact OZ Biosciences with the notification that a BCA assay kit is being returned in order to be reimbursed and/or to definitely terminate a license for internal research use only granted through the purchase of the kit(s). This document covers entirely the terms of the BCA assay kit research only license, and does not grant any other express or implied license. The laws of the French Government shall govern the interpretation and enforcement of the terms of this License.

Product Use Limitations

BCA assay kit and all of its components are developed, designed, intended, and sold for research use only. They are not to be used for human diagnostic or included/used in any drug intended for human use. All care and attention should be exercised in the use of the kit components by following proper research laboratory practices.

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