

# Easy Protein Quantitative Kit (Bradford)

Cat. No. DQ101

Storage: Coomassie brilliant blue solution at 2-8°C in dark for two years; BSA Standard Solution at -20°C for two years Description

*Easy* Protein Quantitative Kit is a ready-to-use modified Bradford Coomassie-binding, colorimetric method for protein quantification. Under acidic condition, Coomassie Brilliant Blue G-250 binds to proteins providing an immediate shift in absorption maximum from 465 nm to 596 nm and a color change from brown to blue.

## Measurement Range

total protein concentration of 50-1000 µg/ml

### Kit Contents

Coomassie Brilliant Blue Solution	100 ml
BSA Standard Solution (0.22 mg/ml)	4×1 ml

#### **Procedures**

- 1. Prior to use, equilibrate Coomassie Brilliant Blue Solution to room temperature and gently invert to mix well.
- 2. Transfer 0, 10, 30, 50, 70, 90, 100  $\mu$ l of BSA Standard Solution (0.22 mg/ml) into seven of 1.5 ml microfuge tubes, and add H<sub>2</sub>O to a final volume of 100  $\mu$ l.
- 3. Transfer protein sample into a new 1.5 ml microfuge tube, and add H<sub>2</sub>O to a final volume of 100 μl.
- 4. Pipette 1.0 ml Coomassie Brilliant Blue Solution into each tube, mix thoroughly and incubate at room temperature for 5-10 minutes.
- 5. Measure the absorbance at 595 nm by spectrophotometer and record the value. Use the absorbance of sample without BSA as a blank control.
- 6. Plot the standard curve and calculate protein concentration in sample. Dilute the sample and re-measure it if the protein concentration falls out of the range of the standard curve.
- 7. The above procedures can be performed with microtiter-plate with 1/10 of the original volume.

## Notes

- Coomassie Brilliant Blue G-250 can strongly bind to quartz cuvette, we recommend to use glass or plastic cuvettes.
- Measurement should be performed from low to high concentration. Do not wash cuvette during this procedure.
- The maximum of color shift occurs during 5-10 minutes after adding Coomassie brilliant blue. The most reliable data should be obtained during this time period.
- Use duplicate or triplicate data for the standard cure.

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