

## **PRODUCT INFORMATION** BDE641, BRADFORD REAGENT

Direction for use Protein Quantitation by Bradford Method.

## BDE641 Bradford Reagent Biotechnology Grade

**Storage:** Cold (4<sup>O</sup>C)

## Instructions:

- 1. Into 4 separate microcentrifuge tubes, aliquot 5, 10, 15 and 20ul of 0.5mg/ml BSA solution. Bring the volume of each to 100ul with 0.15N NaCl.
- 2. Into 1 tube, aliquot 100ul 0.15N NaCl. This will serve as a blank.
- 3. Add to each tube, 1ml Bradford Reagent and vortex. Allow to stand at room temperature for 2 minutes.
- 4. Determine A<sub>595</sub> nm using a 1ml microcuvette. Generate a standard curve by plotting absorbance at 595nm versus protein concentration.
- 5. For the unknown sample, repeat step 1-4 using the unknown in place of the BSA. Plot the A595nm and use the standard curve as a reference to determine the concentration of the unknown sample.

If after the initial assay, the unknown protein concentration is too high, dilute the protein or assay a smaller aliquot of the unknown.

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