

A. Reagents:

– **Coomassie Brilliant Blue (CBBR-250) stain:**

0.1% CBBR-250, 10% acetic acid, 50% methanol.

1. Stir for 4 hours to overnight.
 2. Filter through filter paper.
- 100 ml/a minigel is required.

– **Destain:**

12% methanol, 7% acetic acid
200-400 ml/a minigel is required.

– **Reservation solution:**

5% acetic acid

B. Method:

1. After electrophoresis, place the gel in CBBR-250 stain in amount to cover the gel (100 ml for a typical minigel).
2. Agitate the gel for 5 to 20 min at room temperature.
3. Discard the used CBBR-250 stain.
4. Rinse the gel with a small amount of destain, and discard the destain solution.
5. Add ~ 100 ml of destain and agitate it until a suitable background is achieved.
6. Store the gel in 5% acetic acid.

Proteomics & Metabolomics Unit
proteomics.lab@57357.org

Children's Cancer Hospital 57357
Cairo, Egypt

<https://www.57357.org/proteomics-unit/>