

## Protein Chemiluminescence Reagents (Cat. E208080)

Product Information:	
Contents:	Reagent-A (10mL 12x) and Reagent-B (10mL 12x), 1xTBST (100mL)
Catalog Number:	E208080
Size:	<b>120 ml</b> (1200 cm <sup>2</sup> )
Storage Conditions: Stored at 2-8°C	
Description:	Design for the detection of antibodies conjugated to Horseradish Peroxidase (HRP) in western blotting, suitable for both PVDF and NC membranes. Reagent A con- tains the Luminol and ECL enhancer. Reagent B contains the Hydrogen Peroxide (H <sub>2</sub> O <sub>2</sub> ) and buffer stabilizer. The working solution (A&B mixture) is stable up to 24 hours at ambient temperature.

### For research use only.

# **Procedure:**

- 1. Prepare 1x working solution: mix 100ul reagent-A, 100ul reagent-B and 1ml 1x TBST/PBST for a mini-gel membrane (8 cm x 10 cm). The final volume of detection reagent mixture is around 0.1 ml/cm<sup>2</sup>.
- 2. Place the membranes (protein side up) on a clean surface. Drain off the excess wash buffer.
- 3. Pipette the 1x working solution onto the membrane. The solution should cover the entire surface of the membrane.
- 4. Incubate for 5 minutes at room temperature without agitation.
- 5. Chemiluminescent detection:

Drain off excess working solution and place the blots (protein side up) on a clean surface. Directly expose the membrane on a chemiluminescent / fluorescent imager or wrap up the membrane for x-ray film development.

## Tech Tips:

- 1. Working solution mixture is stable up to 24 hours at ambient temperature.
- 2. Optimization range of primary antibody: 1/3,000-1/5,000.
- 3. Optimization range of secondary antibody (HRP-labeled): 1/30,000-1/50,000.
- 4. Optimization range of working solution mixture: 100-500ul of reagents (A/B) into 1 ml TBST/PBST to increase the sensitivity.

#### Precautions and Disclaimer:

This product and procedure described are intended for R&D use only. Purchase of this product does not convey a license to perform any patented process.

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