

Protein Sample Loading Buffer

(Denaturing, Reducing, 5×)

S750308

Store at 2-8°C short term (3 months). Store at -20°C long term (12 months).

Introduction:

This product is designed for protein sample preparation in SDS-PAGE electrophoresis. Its novel formulation ensures a neutral pH after boiling, preventing protein degradation. It features fast thawing from -20°C and eliminates unpleasant odors by replacing traditional reducing agents (e.g., β -mercaptoethanol or DTT) with an advanced odorless alternative.

Key Features:

1. Prevents Degradation: Neutral pH after boiling avoids protein degradation, ideal for phosphorylated proteins.
2. Rapid Settlement: Samples sink quickly to the bottom of wells without floating.
3. Fast Thawing: Rapid melting from -20°C without viscosity.
4. Odorless: Contains a novel reducing agent—no β -mercaptoethanol or DTT, no unpleasant smell.

Protocol:

1. Add 1 volume of loading buffer to 4 volumes of protein sample (1:4 ratio) and mix thoroughly.
2. Heat the mixture in a boiling water bath for 5–10 minutes.
3. Centrifuge at high speed for 5 minutes. Use the supernatant for electrophoresis.
 - 3.1 Note: If no insoluble debris is observed after adding loading buffer, brief centrifugation is sufficient to collect the mixture.
 - 3.2 If condensation forms on the tube lid after heating, gently flick the tube to mix condensation with the sample before centrifugation to ensure uniform loading concentration.

Usage method:

1. Gels loaded with this buffer may exhibit yellowing or streaks in silver staining due to the reducing agent.
2. Not suitable for non-denaturing gel electrophoresis as it contains denaturing agents.
3. Wear a lab coat and disposable gloves for safety.
4. For research use only.