

## Product Information

### Molecular Weight Marker (M.W. 14,000–66,000)

Catalog Number **SDS7**  
Storage Temperature 2–8 °C

#### Product Description

Polyacrylamide gel electrophoresis (PAGE) in the presence of the anionic detergent, sodium dodecyl sulfate (SDS), has proven to be a useful tool for the separation of protein subunits and the determination of their molecular masses. The proteins supplied in this mixture provide a molecular mass range common to most proteins and their subunits.

This marker is a lyophilized mixture of seven proteins (see Table 1). The mixture has been formulated to yield well-defined bands, which after SDS-PAGE and staining with Brilliant Blue R (Catalog Number B8647) are approximately equal in color intensity.

**Table 1.**  
Protein Mixture in SDS7

Protein	Approximate Molecular Mass (Da)
Albumin, bovine	66,000
Albumin, egg	45,000
Glyceraldehyde-3-phosphate Dehydrogenase, rabbit muscle	36,000
Carbonic Anhydrase, bovine	29,000
Trypsinogen, bovine pancreas	24,000
Trypsin Inhibitor, soybean	20,000
α-Lactalbumin, bovine milk	14,200

#### Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

#### Preparation Instructions

##### Instructions for SDS Laemmli PAGE System only

1× Sample Buffer (62.5 mM Tris-HCl, pH 6.8, containing 2% SDS, 5% 2-mercaptoethanol, 10% glycerol, and 0.002% bromophenol blue) – Prepared by mixing Laemmli Sample Buffer, 2× concentrate (Catalog Number S3401) with an equal volume of water.

Molecular Weight Marker – Add 1.5 ml of 1× Sample Buffer to the vial. Mix by inversion and then vortex for 5 seconds to complete solubilization. Aliquot and freeze at –20 °C or below.

#### Storage/Stability

Store the lyophilized product at 2–8 °C. Store the reconstituted Molecular Weight Marker in aliquots at –20 °C or below.

#### Procedure

1. Incubate a thawed aliquot of the reconstituted Molecular Weight Marker in a boiling water bath for 60 seconds immediately before application of the marker to the gel.

2. Recommended sample volumes:

Standard size gel (16 × 14 cm) – 10 µl/well

Mini-gel (10 × 10 cm) – 5 µl/well

#### Reference

Laemmli, U.K., *Nature*, **227**, 680 (1970).

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