



INCES[®] G-Biosciences + 1-800-628-7730 + 1-314-991-6034 + <u>technical@GBiosciences.com</u>

A Geno Technology, Inc. (USA) brand name

Coomassie Brilliant Blue G-250 Stain

For Staining of Polyacrylamide Gels

(Cat. # 786-497)



INTRODUCTION

Coomassie Brilliant Blue G-250 Stain is based on a colloidal Coomassie stain. Coomassie Brilliant Blue G-250 Stain stains proteins, with high band visibility. The staining of gels with Coomassie Brilliant Blue G-250 Stain allows the examination of protein bands even during the staining process. After the staining process, the band intensity may be further enhanced by de-staining the stained gel in our Coomassie Brilliant Blue G-250 Stain has a sensitivity of staining 8-10ng protein/band, i.e. 8-10ng of BSA is visible in 4-20% SDS acrylamide gels. The supplied volume is suitable for 40-50 Mini gels (8 x 10cm) & 20-25 large gels (12 x 15cm).

ITEM(S) SUPPLIED (Cat. # 786-497)

Description	Size
Coomassie Brilliant Blue G-250 Stain	1L

STORAGE CONDITION

It is shipped at ambient temperature. Store it at room temperature, upon arrival and is stable for 1 year when stored and used properly.

PROTOCOL FOR STAINING THE POLYACRYLAMIDE GELS:

- Wash gel 2-3 times, 5 minutes each in deionized water to remove SDS present in the gel. Each wash should be in large volumes of water. For gels without SDS, a single wash in deionized water is sufficient.
 NOTE: Isoelectric focusing gels require prefixing in 20% trichloroacetic acid for 30 minutes, followed by extensive washing to remove TCA.
- Remove all free water from the gel. Add an adequate amount of Coomassie Brilliant Blue G-250 stain to cover the gel. Gently shake the gel in stain for 1 hour. Protein bands will be visible within 3-5 minutes and reach a maximum intensity within 1 hour. Longer incubation may be performed.
- Rinse the stained gel in a large volume of deionized water, 2-3 times for 5 minutes each. Rinsing in deionized water enhances the intensity of the protein bands. Rinse gel in Coomassie Brilliant Blue Destaining Solution (Catalog # 786-499) or 30% Methanol, until desired resolution is attained. Store the stained gel in deionized water.

NOTE: If background staining is noticed, it is indicative of residual SDS in the gel. Rinsing the gel extensively in deionized water will remove the background staining.

RELATED PRODUCTS

Download our Protein Electrophoresis Handbook



http://info.gbiosciences.com/complete-protein-electrophoresis-handbook/ For other related products, visit our website at <u>www.GBiosciences.com</u> or contact us.

Last saved: 8/16/2012 CMH



www.GBiosciences.com