

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

A Geno Technology, Inc. (USA) brand name

# Tris-Acetate SDS Buffer Kit (for Tris-Acetate Gels)

### For SDS Polyacrylamide Gel Electrophoresis

## (Cat. #786-589)



#### **INTRODUCTION**

Tris-Acetate SDS Buffer Kit is a complete buffer kit to resolve high molecular weight proteins (36-400 kDa) under denaturing conditions on Tris/Acetate gels

#### ITEM(S) SUPPLIED

Description	Cat. #786-589
Tris-Acetate/SDS Running Buffer [20X]	2 x 250 ml
DTT [ 1M](15.4mg)	1 vial
Protein Antioxidant [200X]	15 ml
LDS Sample Loading Buffer [4X]	10 ml

#### **STORAGE CONDITIONS**

The kit is shipped at ambient temperature. Store at 4°C.

#### **PREPARATION BEFORE USE**

- 1. Add 100  $\mu l$  deionized water to DTT vial. Mix well to dissolve and store it at 20°C after use.
- 2. Add 50 ml of Tris-Acetate/SDS Running Buffer to 950 ml of deionized water to obtain 1X Tris-Acetate/SDS Running Buffer.

#### PROTOCOL

#### Loading Sample Preparation

- 1. Bring the LDS Sample Loading Buffer [4X] to room temperature before use.
- 2. Prepare reduced or non-reduced protein samples as below:

Reagent	Reduced Protein Sample	Non-reduced Protein Sample
Protein Sample	<i>x</i> μl	<i>x</i> μl
LDS Sample Loading Buffer [4X]	2.5 μl	2.5 μl
DTT [ 1M](15.4mg)	0.5 μl	-

Make up the final volume of sample to 10  $\mu l$  with deionized water.

- 3. Vortex the tube to mix the contents.
- Heat both the reduced and non-reduced samples at 70°C for 10 minutes and let it cool.
- 5. Centrifuge the sample tubes and load the samples on gel to run SDS-PAGE.

#### SDS-PAGE with Tris-Acetate Gels and Tris-Acetate/SDS Running Buffer

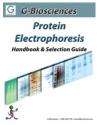
 For reducing SDS-PAGE add 1 ml Protein Antioxidant [200X] to 200 ml of 1X Tris-Acetate/SDS Running Buffer and fill the upper chamber of electrophoresis unit.

**Note:** Reducing and non-reducing samples are preferably run in different gels. For non-reducing sample run, Protein Antioxidant is not added to the running buffer in the cathodic chamber of electrophoresis unit. If reducing and non-reducing samples are run on same gel for some reason, then do not use Protein Antioxidant.

 Load the samples and perform electrophoresis at constant voltage of 150 V (Approx. run time for gel is 1 hr).

#### **RELATED PRODUCTS**

Download our Protein Electrophoresis Handbook.



http://info2.gbiosciences.com/complete-protein-electrophoresis-handbook

For other related products, visit our website at <u>www.GBiosciences.com</u> or contact us.

Last saved: 1/30/2017 CMH



## www.GBiosciences.com