10x Tris-Glycine PAGE Running Buffer

Christopher Bartley

Abstract

Western blot running buffer.

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Protocol

Step 1.
Fill 1L pyrex bottle with 700mL dH20

Step 2.
Add 30.2g Tris base

Step 3.
Add 144.2g glycine

Step 4.
pH solution to 8.80 after disolution of tris and glycine

Step 5.
Add 10g SDS (1% final)

Step 6.
Fill to 1L with dH20