ProtoGel® 40%

- 37.5:1 Acrylamide:Bisacrylamide
- Ultra-Pure, Stabilized Solution
- Consistently Crystal Clear Gels
- Zero Fluorescence

national diagnostics

METHOD FOR SDS-PAGE

1. Gel Formulation—Laemmli SDS-PAGE

Use the chart below to determine the volumes of reagents required for desired gel composition. If the percentage gel you are running is not included in the table, use the formula below to calculate the volumes of ProtoGel, ProtoGel Resolving Buffer and other reagents needed.

Volumes of Solution Components for Common Gel Percentages (mL)

Using Premixed 4X Resolving Buffer or 1.5M Tris-HCl

Gel %		-(OR-	
6	ProtoGel 40%: 4X Resolving Buffer: Deionized H ₂ O:	15.0 25.0 58.9	ProtoGel 40%: 1.5 M Tris-HCl, pH 8.8: 10% SDS: Deionized H ₂ O:	15.0 25.0 1.0 57.9
8	ProtoGel 40%: 4X Resolving Buffer: Deionized H ₂ O:	20.0 25.0 53.9	ProtoGel 40%: 1.5 M Tris-HCl, pH 8.8: 10% SDS: Deionized H ₂ O:	20.0 25.0 1.0 52.9
10	ProtoGel 40%: 4X Resolving Buffer: Deionized H ₂ O:	25.0 25.0 48.9	ProtoGel 40%: 1.5 M Tris-HCl, pH 8.8: 10% SDS: Deionized H ₂ O:	25.0 25.0 1.0 47.9
12	ProtoGel 40%: 4X Resolving Buffer: Deionized H ₂ O:	30.0 25.0 43.9	ProtoGel 40%: 1.5 M Tris-HCl, pH 8.8: 10% SDS: Deionized H ₂ O:	30.0 25.0 1.0 42.9
15	ProtoGel 40%: 4X Resolving Buffer: Deionized H ₂ O:	37.5 25.0 36.4	ProtoGel 40%: 1.5 M Tris-HCl, pH 8.8: 10% SDS: Deionized H ₂ O:	37.5 25.0 1.0 35.4

Note: The amount of ProtoGel Resolving Buffer is always the same, regardless of percentage of monomer in the gel (25mL of ProtoGel Resolving Buffer per 100mL gel casting solution).

The volume of ProtoGel required for gel casting solutions for any volume and acrylamide concentration may be calculated from the following formula:

$$V_{p} = (XV_{t}) / 40$$

where V_p = volume of 40% ProtoGel, X = percent monomer desired in gel and V_t = total volume of gel casting solution.

EXAMPLE: To make 100 mL of a 10% percent monomer gel, the amount of ProtoGel to add is calculated as follows:

$$V_p = (10 * 100) / 40 = 25.0$$

2. Add Initiators and Cast Gel

For optimal results, degas gel solution for 10 minutes under vacuum aspiration prior to initiation with ammonium persulfate and TEMED. Add 1.0 mL of 10% (w/v) ammonium persulfate for every 100 mL of gel casting solution. Swirl gently to mix. Add 0.1 mL of TEMED for every 100 mL of gel casting solution. Pour the solution into the gel casting cassette. The gel should begin to set in 10-20 minutes. To provide a sharp interface, overlay the gel with water saturated n-butanol during polymerization. Flush butanol away with water just before casting the stacking gel (below).

3. Pour Stacking Gel

Use ProtoGel Stacking Buffer to make 10 mL of a 4% stacking gel as follows:

ProtoGel: 1.00 mL
ProtoGel Stacking Buffer: 2.50 mL
Deionized Water: 6.44 mL

Afterwards, add 0.05 mL 10% ammonium persulfate and 0.01 mL TEMED. Gel will begin to set in 20 minutes.

Note: A solution of 0.5M Tris-HCl, 0.4% SDS at pH 6.8 may be substitued for ProtoGel Stacking Buffer.

4. Select Running Buffer

Laemmli SDS-PAGE - 1X Tris-Glycine-SDS (EC-870) is the most sutiable tank buffer for most SDS-PAGE applications.

Small Protein SDS-PAGE (<20kD) - National Diagnostics' unique Tris-Tricine-SDS running buffer (EC-869) helps resolve smaller proteins without requiring the full Schagger-Von Jagow protocol.

Order Numbers for this Protocol

ProtoGel 40%	450 mL
Order No. EC-891	1 liter
4X Resolving Buffer	450 mL
Order No. EC-892	1 liter
4X Stacking Buffer	200 mL
Order No. EC-893	
10X Tris-Glycine-SDS Bufffer	1 liter
Order No. EC-870	4 liter
10X Tris-Tricine-SDS Buffer	1 liter
Order No. EC-869	
2X Protein Loading Buffer Blue	10 X 1 mL
Order No. EC-886	
5X Protein Loading Buffer	10 X 1mL
Order No. EC-887	
Ammonium Persulfate	25 g
Order No. EC-504	100 g
TEMED	25 mL
Order No. EC-503	

For Additional Information & Order Placement:

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