

M-Fast **SDS-PAGE Gel Solutions (WF):**

Use for science research only. Not for Therapeutic or Diagnostic Use.

(Catalog #: WF-150 (15%); WF-100(10%); WF-075(7.5%))

Size: Gel solution-A (250mL) Gel solution-B (250mL)

Storage: **2-8°C.**

- Solution-A & B contain Acrylamide-Bis, SDS, Tris-HCl, TEMED and Stabilizers, used to cast your own polyacrylamide gels. (Note*: **Not includes the fresh 10%APS**)

*** Protocol for preparing a mini-SDS-PAGE gel (8x10cm):**

| A mini SDS-PAGE gel (8x10) | Resolver | Stacker |
|----------------------------|----------|---------|
| Solution- A | 4mL | 0.5mL |
| Solution- B | 4mL | 1.5mL |
| Total Volume | 8mL | 2mL |
| 10% APS | 50uL | 20uL |

1. Set the casting frames (clamp two glass plates in the casting frames) on the casting stands.
Note*: Pre-warm the gel solution-A and -B at room temperature for 10-20 minutes prior to use.
2. Prepare 10% APS (fresh) before casting gels. Dissolve 0.2 g of APS in 2 mL of deionized water.
3. Prepare the resolving gel solution by mixing **4mL gel solution-A and 4mL solution-B.**
4. Add **50uL 10% fresh APS** to the combined resolving gel solution and mix well.
5. Pipet the gel solution into the gap between the glass plates. Fill the cassette to 0.5–1 cm below the bottom of the teeth on the comb. ****Immediately prepare and pour the stacking solution. It is not necessary to overlay water/butanol/isopropanol on the top of the resolving gel.**
6. Prepare the stacking gel solution by mixing **500uL solution-A and 1.5mL solution-B.**
7. Add **20uL 10% fresh APS** to the combined stacking gel solution and mix well.
8. Pipet the stacking gel solution into the cassette and filling to the top of the short plates. Insert the comb without trapping air under the teeth.
9. Wait for 20-30 minutes and the gel is ready to use for protein electrophoresis.

| Related products: | Size: | Cat. No. | Price | Recipe: |
|-------------------------------------|-------|----------|-------|---|
| 10x Tris-Glycine SDS running buffer | 500mL | P-3500 | \$19 | 25mM Tris, PH8.3, 192mM glycine, 0.1%SDS |
| 2x Tris-Glycine SDS Sample buffer | 50mL | PS-50 | \$19 | 100mM Tris-HCl, PH6.8, 40%glycerol, 2%SDS, 0.01%Bromophenol blue (add 2-mercaptoethanol before use) |
| 10x Sharper Protein running buffer | 500mL | B5020 | \$55 | Used for small Proteins (<100kDa)/peptide electrophoreses |
| APS (power) | 50g | APS-50 | \$19 | 10%APS= 0.2 g of APS in 2 mLof deionized water |

Additional informations:

| Acrylamide (%) | M.W. Range | Thickness of the gel | Volumes of the gel solution |
|----------------|------------------|----------------------|-----------------------------|
| 7.5% | 50 kDa - 500 kDa | 0.75mm | 6.0 mL |
| 10% | 20 kDa - 300 kDa | 1.0 mm | 10 mL |
| 15% | 3 kDa - 100 kDa | 1.5 mm | 12 mL |

Precautions and Disclaimer: This product and procedure described are intended for R&D use only. Purchase of this product does not convey a license to perform any patented process.