Quick Start Guide

mPAGE® Lux Bis-Tris Gel Casting System

mPAGE® Lux Curing Station mPAGE® Lux Bis-Tris Reagent Kit

FOR RESEARCH USE ONLY

Not for use in diagnostic procedures. Not for human or animal consumption.



Gel Solution Preparation

Important: Protect all solutions from light to prevent polymerization. Before casting, bring all solutions to room temperature.

Resolving Gel

Prepare Resolving Gel Solution by mixing Resolving Solution and Diluent.

- Use the Solution Volume Worksheet on page 3
 to calculate required mixing volumes. Resolving
 Gel Solution may be prepared in a large batch to
 make several gels sequentially.
- 2. Using a clean pipette, add required amount of Resolving Solution to provided black mixing tube or other opaque container.
- 3. Using a clean pipette, add required amount of Diluent to the same mixing tube.
- 4. Mix by turning container end over end. **Do not vortex**.

Stacking Gel

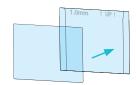
The Stacking Solution is ready to use directly from the bottle. See the Solution Volume Worksheet on page 3.

Important: Do not dilute Stacking Solution.



Gel Casting

- 1. Clean glass plates with mild detergent and rinse with DI water. Wipe with 70% Ethanol before use.
- 2. Assemble mPAGE® Gel Caster using mPAGE® Spacer Plate and mPAGE® Short Plate to form glass cassette. Ensure that both glass plates are aligned at the bottom of the caster frame before closing caster clamps.



Note: If using mPAGE® Lux Mask Short Plate, refer to full user guide for assembly instructions (available on the mPAGE® Lux product page at SigmaAldrich.com).

3. Power on mPAGE® Lux Curing Station by pressing the power button **③**. The Curing Station will initiate a self test. After self test is complete, the ready screen will appear.



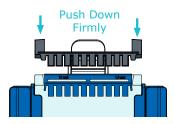
- Open door and place gel caster into Curing Station, aligning the Gel Caster behind the bumpers.
- 5. Using a clean 5 mL pipette, add prepared Resolving Gel Solution to the indicated fill line on the mPAGE® Gel Caster frame.
- Place **behind** bumpers
- 6. Using a clean 5 mL pipette, slowly add Stacking Solution up to the top of the short plate.
- 7. Slowly insert the mPAGE® comb at an angle to prevent air bubble formation under teeth.
- 8. Wipe solution spill-over from front of short plate.

9. With the hooks in the back, firmly push the mPAGE® Clip-on Mask down, over glass cassette until it completely covers the comb teeth. The mPAGE® Clip-on Mask prevents the gel from curing around the comb.

Note: Pushing front of plates during installation may cause gel to seep out.

Important: Mixing different well formats may result in improper well formation. Use:

- 10-well combs with 10-well mPAGE® Clip-on Mask
- 15-well combs with 15-well mPAGE® Clip-on Mask





10. Close Curing Station door and select gel thickness.



11. Press Start to begin gel curing.

Note: If casting multiple gels, the second gel caster can be assembled while the first is curing.

12. After curing is complete, open door and remove Gel Caster. Remove mPAGE® Clip-on Mask. Remove cassette from Gel Caster by lowering the tension clip to release the caster frame. Then open the sides of the frame and slide cassette out from top.

Option: After curing, The mPAGE® Clip-on Mask can be removed from the Gel Caster while it is in the mPAGE® Lux Curing Station.

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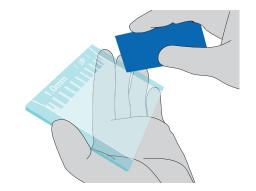
Use gel immediately or wrap the cassette in wet paper towel and store laying flat in a zip top bag or other airtight container at 2-8 °C for up to 2 weeks. Do not leave cassettes unwrapped as the gel will dry out.

How to Remove Gel After Electrophoresis

Remove the gel from glass cassette using a gel scraper. Cut along gel edges (as shown) to avoid gel tearing.

mPAGE® Lux Bis-Tris gels should ONLY be used with MOPS-SDS or MES-SDS running buffer. Bis-Tris gels are NOT compatible with Tris-Glycine running buffer. See the User Guide for recipes.

See complete mPAGE® Lux Casting System User Guide for process details, suggested electrophoresis and transfer conditions, and troubleshooting.



Solution Volume Worksheet

Using a dry erase marker, calculate the solution volumes in the table below.

For Resolving Solution

Gel %		Resolving Solution		Diluent	Total Volume
8%	2 mL x number of gels x gel thickness =	mL	3 mL + x number of gels x gel thickness =	mL :	mL mL
10%	2.5 mL x number of gels x gel thickness =	mL	2.5 mL * x number of gels x gel thickness =	mL :	= mL
12%	3 mL x number of gels x gel thickness =	mL	2 mL + x number of gels x gel thickness =	mL :	m L
13.5%	3.3 mL x number of gels x gel thickness =	mL	1.7 mL * x number of gels x gel thickness =	mL :	= mL

For Stacking Solution

Gel %	Stacking Solution			Volume Volume
5%	1.5 mL	x number of gels x gel thickness	=	mL

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