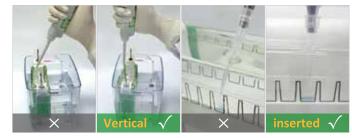
### 4. Sample loading.

You can use 10  $\mu$ l pipette tip to load samples into the wells easily. Make sure that the loading tip is inserted into the sample well vertically for optimum result and to avoid inserting the tip into the groove on the back plate.



### 5. Running conditions.

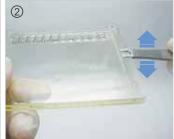
Voltage	Started Current	Finished Current	Run Time per Gel
140 V	75-100 mA	30-50 mA	45-55 min

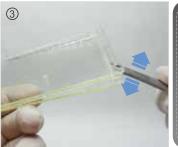
The running time may be varied depending on your power supply and the gel percentage.

#### 6. Open the cassette with an opener or a screw driver to remove the gel.

GenScript's eStain 2.0 Protein Staining System (Cat No. L02016) can help you to get better results.

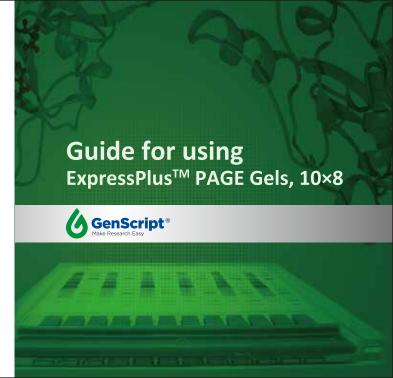






For more support please call local office number or in the U.S., call 1-877-436-7274 (1-732-885-9188), also you can visit www.genscript.com

Open the cassette with GenScript's opener or a screw driver.



# 1. Choose the appropriate gels for your protein electrophoresis analysis using the charts given below.

ercentage	Separation Range	8-16%	4-20%	4-12%	8%	10%
-16%	160-10 KDa	250kDa			250kDa	250kDa
4-20%	250-10 KDa	150kDa	250kDa	250kDa	LANCE DO	150kDa
4-12%	250-20 KDa	100kDa	150kDa		1300SM	100kDa
8%	180-20 KDa	80kDa	160kDa	150kDa	100kDa	80kDa
10%	160-20 KDa	60kDa				- COLON
12%	120-6.5 KDa	50kDa	80kDa	100kDa	DULLIA	Indicate and in contract of
		MANAGEMENT,	MOLDO.	80kDa	60kDa	50kDa
wells	volume/well	luddis@au	50kDa			Ath Da
10	80 µl	304Da	141.0	60kDa	SOLDa.	
12	60 µI	- Carrier	-AUKUR-	-		30kDa
15	40 µl	25500	30kDa	Indication.	40L0a	
		-	-	L-ACADA		25ADa
. Prepare gel tank and		20kOu	25kDa		30kDa	
uffer. Using MOPS or MES		45kDa	20kDa	30kDn	-	20kDa
	AGE running	C-MADa	ASADa	orana.	0.000	MADA
uffer (Cat No. M00138).		-	dian.	- CONTRACT	mankin.	Name of the local division in the local divi
, 546 1				20k0a	20k0a	infékba-

### ★ Compatible gel tanks:

Bio-Rad Mini-PROTEAN (II/ 3/ Tetra System) Hoefer Mighty Small (SE 250/ SE 260)

Sample preparation	Reagent	Volume	
	Protein sample	xμl	
	5 × loading buffer	2 μΙ	
	Deionized H <sub>2</sub> O	to 8 µI	
	Total Volume	10 µl	

### SDS-PAGE:

Heat the sample at 100°C for 10 min, spin down sample before loading.

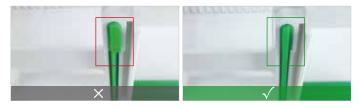
## Native PAGE:

Don't heat the sample.

3. Peel the tape at the bottom of the gel plate, remove the comb gently, then insert the gel into the gel running apparatus and add running buffer.



Turn around the gasket to fit the gel plate when using Bio-Rad's gel tanks



Adapters provided in the gel package are needed since the thickness of Express Plus PAGE Gel cassette is smaller than that of Invitrogen NuPAGE gel cassette.

