E-Gel[™] 1 Kb Plus DNA Ladder

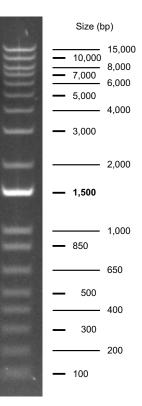
PRODUCT INFORMATION SHEET

Pub. No. MAN0000772

K	Contents	Catalog No. 10488090	Amount 100 applications	() Kit contents
	Storage	 Product is shipped at ambient temperature. Store at room temperature or at 4°C for up to 6 months, or at -20°C for long term storage. 		

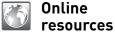
Product description

- The Invitrogen[™] E-Gel[™] 1 Kb Plus DNA Ladder is designed for sizing and quantification of double stranded DNA on 1.2% E-Gel[™] agarose gels.
- The E-Gel[™] 1 Kb Plus DNA Ladder consists of 18 individual chromatography-purified DNA fragments ranging in size from 100 bp to 15,000 bp.
- A reference band at 1,500 bp is included for easy orientation.
- The ladder is supplied with 1X E-Gel[™] Sample Loading Buffer for sample DNA.



Rev. A.0

 Visit our product pages for additional information and protocols.



- Go online to view related DNA ladders and markers.
- For support, visit thermofisher.com/support.

Required materials

- E-Gel[™] EX or E-Gel[™] Agarose Gel with SYBR[™] Safe (See **Choosing the right** DNA ladder for your E-Gel[™] agarose gel)
- TE Buffer (Cat. No. AM9858)
- Ultrapure[™] DNase/RNase-Free Distillated Water (Cat. No. 10977023)



Important guidelines

- Do not heat the E-Gel[™] 1 Kb Plus DNA Ladder before loading.
- Load the same volume of DNA sample and DNA ladder.
- For quantification, adjust the concentration of the sample to equalize it approximately with the amount of DNA in the nearest band of the ladder.
- Dilute sample DNA in TE buffer to avoid degradation of DNA sample.
- ⑦ Choosing the right DNA ladder for your E-Gel™ agarose gel
- Troubleshooting
- U Limited product warranty and disclaimer details





invitrogen

Prepare DNA ladders and samples for electrophoresis

This protocol provides a brief description of how to use the DNA ladder with E-Gel[™] agarose gels. For detailed instructions on using specific types of E-Gel[™] agarose gels, go to thermofisher.com or contact Technical Support.

Step			Action		
1		Prepare DNA ladder	 a. Thaw, mix and briefly centrifuge DNA ladder before use. b. Prepare DNA ladder. For E-Gel[™] EX Agarose Gels, mix 2 μL of DNA ladder with 18 μL of water. For E-Gel[™] Agarose Gels, mix and use the ladder without dilution. For E-Gel[™] 48 Agarose Gels, mix 2 μL of DNA ladder with 13 μL of water. 		
2		Prepare samples	a. Dilute your sample 2- to 10-fold with TE Buffer (Cat. No. AM9858), 1X E-Gel [™] Sample Loading Buffer (Cat No. 10482055), or water. b. Mix gently.		
3		Load samples and DNA ladders	 a. Load DNA ladders and DNA samples into the appropriate wells of the E-Gel[™] agarose gel. Add 20 μL for E-Gel[™] and E-Gel[™] EX Agarose Gels. Add 15 μL for E-Gel[™] 48 Agarose Gels. b. Add water to any empty wells, so that all wells contain an equal volume of liquid. 		
		Perform electrophoresis	a. Choose the appropriate E-Gel [™] run protocol for your gel type based on the electrophoresis device being used.		
4			Gel type Program Recommended run time		
			E-Gel [™] Power Snap Electrophoresis Device (Cat. No. G8100)		
			E-Gel [™] EX Agarose Gel (1%) E-Gel EX 4 1-2% 15 min (20 min max)		
			E-Gel [™] Agarose Gel (0.8%, 1.2%, 2%) E-Gel 0.8-2% 26 min (40 min max)		
			E-Gel [™] E-Base [™] Device		
			E-Gel [™] 48 Agarose Gel (4%) EG 20 min		
			b. Run the program to start electrophoresis.		
		Visualize agarose gel	Visualize DNA ladder and samples.		
5			 Use the E-Gel[™] Power Snap Camera (Cat. No. G8200), E-Gel[™] Imager (Cat. No. 466612), or other blue light imager to detect DNA bands stained with SYBR[™] stains. 		
			 UV transilluminator to detect DNA bands stained with ethidium bromide. 		

For support, visit thermofisher.com/support.

Thermo Fisher

SCIENTIFIC