

ELISABETH PHARMACON Ltd. Rokycanova 4437/5

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EliDNATM PS Green Plus Instructions for Use

Package:

Ref. No. Package ED02 1 ml ED02s 50 µl

Storage:

The dye should be stored at 4 °C and protected from light. When stored under these conditions, the dye will retain full activity until the expiration date indicated on the label.

Product description

EliDNATM PS Green Plus is a new generation of a highly sensitive green fluorescent nucleic acid dye intended for gel staining. The dye is non-toxic and non-mutagenic. Therefore, making it a safer alternative to ethidium bromide. EliDNATM PS Green Plus can be used with a standard UV transilluminator (300 nm) or with instruments using visible light (~500 nm). Gels stained with EliDNATM PS Green Plus are compatible with various downstream applications such as gel extraction, cloning and many others.

Protocol

Prestaining:

Prepare the desired volume of agarose gel solution according to your protocol. For a 100 ml agarose solution add 4-6 μ l of **EliDNATM PS Green Plus** and mix it properly. Allow the gel solution with the dye to cool down to ~60 °C before casting it into the gel tray. After the gel has solidified, load the samples and perform electrophoresis. Image the gel with a UV transilluminator or with a visible light with ~500 nm using yellow or green filter.

Poststaining:

Prepare and run the agarose gel according to your protocol. For 100 ml of buffer or distilled water use 20-30 μ l of **EliDNATM PS Green Plus**. Store the staining solution at room temperature in the dark. Use a suitable container and place the gel into the solution. Recommended staining time is 10-20 min. Depending on the thickness of the gel and its concentration, staining time and the volume of the dye may be adjusted.



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Troubleshooting

- If you experience low fluorescence signal of DNA bands, try to add more dye. Signal intensity depends on the number of bands and their concentration.
- We do not recommend repeated melting of gel containing the dye. This may lead to reduced signal intensity.
- Smeared DNA bands may be caused by excessive DNA concentration. Reduce the amount of loaded DNA or perform poststaining.
- If there are any discrepancies in bands migration, try to reduce the amount of loaded DNA or use less dye.

Related products

REF	Name of product	UV light	Blue LED	Green/blue LED	In gel	Post-st.	Loading	dsDNA	ssDNA	RNA
ED01	EliDNA [™] PS Green	✓	✓	✓	✓	✓	Х	1	√ ¹	√ ¹
ED02	EliDNA [™] PS Green Plus	1	✓	✓	✓	1	Х	✓	1	√
ED03	EliDNA [™] PS Green Ultra²	1	✓	✓	✓	✓	Х	✓	✓	√
ED04	EliDNA [™] PS Red	✓	Х	Х	✓	✓	Х	✓	✓	√
ED05	EliDNA [™] LD Green	✓	✓	✓	Х	Х	✓	✓	✓	√
ED06	FliDNA™ LD Red		Х	Х	Х	Х	1	1	1	

¹ Since ssDNA and RNA are single stranded, you may experience a lower signal intensity.

Manufacturer:

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Catalog number



Batch code



Use by (last day of month)



Upper limit of temperature



Manufacturer



Contains sufficient "N" tests

²The dye with the highest sensitivity.