QR-Code Safety Data Sheet

Please find a digital version of the safety data sheets by following the link below:



www.mn-net.com/sds



We strongly recommend to carefully read the detailed protocol section of the product's user manual. If you have any questions about the protocol or product,

please contact our Technical Support.

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User manuals

NucleoSpin® Gel and PCR Clean-up

740609/.10/ .50/.250



Dear valued customer,

Thank you for choosing MACHEREY-NAGEL for your application. We have attached a short protocol for your review.

To obtain the best results we recommend to follow the detailed protocol available online, especially when you are a first time user

All important links to the above mentioned product are listed in this leaflet.

information regarding product components, specifications, safety instructions, and processing protocols, can all be found on the product website and accessed easily via the QR code.

QR-Code product website



DNA Clean-Up

qr.mn-net.com/qr/(241)740609

Use the following QR code or the link below for direct access to the user manual.

QR-Code user manual



gr.mn-net.com/gr/(IFU)740609

It is strongly recommended to read the detailed protocol section of the user manual if using the kit for the first time. However, experienced users may refer to the protocol at a glance. The protocol at a glance is designed to be used only as a supplemental tool for quick reference while performing the purification procedure.

We are constantly improving our products and we reserve the right to make changes or additions to protocols. Please check for updated revisions for previously downloaded manuals.

This leaflet does not replace the full manual!





Protocol at a glance (Rev. 08) PCR clean-up, gel extraction

	PCR clean-up	Gel extraction	DNA clean-up (with SDS)	Single stranded DNA clean-up
1 PCR clean-up, DNA clean-up, or single stranded DNA clean-up: Adjust binding condition Gel extraction: Excise DNA fragment/ solubilize gel slice	200 µL NTI/ 100 µL PCR	200 µL NTI/ 100 mg gel 50 °C 5-10 min	500 μL NTB / 100 μL sample	200 μL NTC/ 100 μL sample
2 Bind DNA			11,000 x g 30 s	
3 Wash silica membrane			700 µL NT3 11,000 x g 30 s Recommended: 2 nd wash 700 µL NT3 11,000 x g 30 s	
4 Dry silica membrane			11,000 x <i>g</i> 1 min	
5 Elute DNA			15-30 μL NE RT 1 min 11,000 x <i>g</i> 1 min	