

## User manuals

**NucleoSnap® Finisher Midi** 740434 / .10 / .50

**NucleoSnap® Finisher Maxi** 740435 / .10 / .50



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 of this kit.

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

Important 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



[qr.mn-net.com/qr/\(241\)740434](https://qr.mn-net.com/qr/(241)740434)

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

### QR-Code user manual



[qr.mn-net.com/qr/\(IFU\)740434](https://qr.mn-net.com/qr/(IFU)740434)

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.01)  
Plasmid DNA concentration and desalination

|                             |                                                                                     | Midi<br>NucleoSnap®<br>Finisher Midi                                | Maxi<br>NucleoSnap®<br>Finisher Maxi                                |
|-----------------------------|-------------------------------------------------------------------------------------|---------------------------------------------------------------------|---------------------------------------------------------------------|
| 1 Adjust binding conditions |    | 2.5 mL Buffer FB<br>Mix                                             | 7.5 mL Buffer FB<br>Mix                                             |
| 2 Filtrate precipitate      |    | Connect NucleoSnap®<br>Column to vacuum<br>Load sample<br>-0.3 bar* | Connect NucleoSnap®<br>Column to vacuum<br>Load sample<br>-0.3 bar* |
| 3 Wash precipitate          |    | 2 mL Buffer A4<br>-0.3 bar*                                         | 4 mL Buffer A4<br>-0.3 bar*                                         |
| 4 Separate column           |                                                                                     | Remove upper part and<br>discard                                    | Remove upper part and<br>discard                                    |
| 5 Dry silica membrane       |   | 11,000 x g, 1 min                                                   | 11,000 x g, 1 min                                                   |
| 6 Elute DNA                 |  | 200 µL H <sub>2</sub> O-EF<br>11,000 x g, 1 min                     | 500 µL H <sub>2</sub> O-EF<br>11,000 x g, 1 min                     |

\* Reduction of atmospheric pressure. Use of a vacuum regulator (e.g., NucleoVac Vacuum Regulator) to prevent a reduced recovery due to higher vacuum forces is highly recommended, see ordering information.

## 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](http://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.

## Contact MN

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