Genomic DNA

User manuals

NucleoSpin® 8 Tissue **740740 / .5**

NucleoSpin® 8 Tissue Core Kit **740453.4**

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 product 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)740740

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)740740

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 is a short protocol and does not replace the full manual!

Protocol at a glance

NucleoSpin® 8 Tissue – centrifuge processing

Please check the user manual if your centrifuge and setup meets the requirements for centrifuge processing of 8-well strips.

1 Prepare samples	2 × 0.5 cm mouse tail or up to 20 mg tissue, 10^6 cultured cells, or bacteria
2 Lyse samples	180 µL T1 25 µL Proteinase K Mix 56 °C, ≥ 6 h
3 Adjust DNA binding conditions	200 µL BQ1 200 µL ethanol (96 – 100 %) Mix
4 Transfer lysates to NucleoSpin® Tissue Binding Strips	
5 Bind DNA to silica membrane of the NucleoSpin® Tissue Binding Strips	5,600 x *g*, 10 min
6 Wash silica membrane	500 µL BW 5,600 x *g*, 2 min 700 µL B5 5,600 x *g*, 4 min
7 Dry silica membrane	70 °C, 10 min
8 Elute DNA	100 µL BE (70 °C) 5,600 x *g*, 2 min Optional: Repeat elution step once.

Protocol at a glance
NucleoSpin® 8 Tissue – vacuum processing

1 Prepare samples	2 × 0.5 cm mouse tail or up to 20 mg tissue, 10^6 cultured cells, or bacteria
2 Lyse samples	180 µL T1 25 µL Proteinase K Mix 56 °C, ≥ 6 h
3 Adjust DNA binding conditions	200 µL BQ1 200 µL ethanol (96 – 100 %) Mix Prepare the NucleoVac 96 Vacuum Manifold
4 Transfer lysates to NucleoSpin® Tissue Binding Strips	
5 Bind DNA to silica membrane of the NucleoSpin® Tissue Binding Strips	-0.2 bar*, 5 min
6 Wash silica membrane	600 µL BW 900 µL B5 900 µL B5 -0.2 bar*, 5 min each step Remove MN Wash Plate
7 Dry silica membrane	-0.6 bar*, 10 min
8 Elute DNA	100 µL BE (70 °C) -0.4 bar*, 2 min Optional: Repeat elution step once

** Reduction of atmospheric pressure*

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.

Contact MN

Germany and international

MACHEREY-NAGEL GmbH & Co. KG
Valencienner Str. 11 · 52355 Düren · Germany
Tel.: +49 24 21 969-0
Toll-free: 0800 26 16 000 (Germany only)
E-mail: info@mn-net.com

Technical support Bioanalysis

Tel.: +49 24 21 969-333
E-mail: support@mn-net.com

USA

MACHEREY-NAGEL Inc.
924 Marcon Blvd. · Suite 102 · Allentown PA, 18109 · USA
Toll-free: 888 321 6224 (MACH)
E-mail: sales-us@mn-net.com

France

MACHEREY-NAGEL SAS
1, rue Gutenberg – BP135 · 67720 Hoerdt Cedex · France
Tel.: +33 388 68 22 68
E-mail: sales-fr@mn-net.com

MACHEREY-NAGEL SAS (Société par Actions Simplifiée) au capital de 186600 €
Siret 379 859 531 00020 · RCS Strasbourg B379859531
N° intracommunautaire FR04 379 859 531

Switzerland

MACHEREY-NAGEL AG
Hirsackerstr. 7 · 4702 Oensingen · Switzerland
Tel.: +41 62 388 55 00
E-mail: sales-ch@mn-net.com

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