

MACHEREY-NAGEL

Plasmid purification for transfection

scientifix

PO Box 662 South Yarra
VIC 3141 Australia

www.scientifix.com.au

Free call: 1800 007 900
T: +61(0)3 8540 5900
F: +61(0)3 9543 7827
E: info@scientifix.com.au

Bioanalysis



NucleoSnap Plasmid Midi

NEW generation of plasmid midi preparation

- New column design for vacuum processing of large sample volumes
- Transfection-grade plasmid DNA for sensitive downstream applications
- Isolate up to 700 μ g plasmid DNA in only 35 minutes



NEW
column design
for VACUUM
processing

MACHEREY-NAGEL

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Since 1911

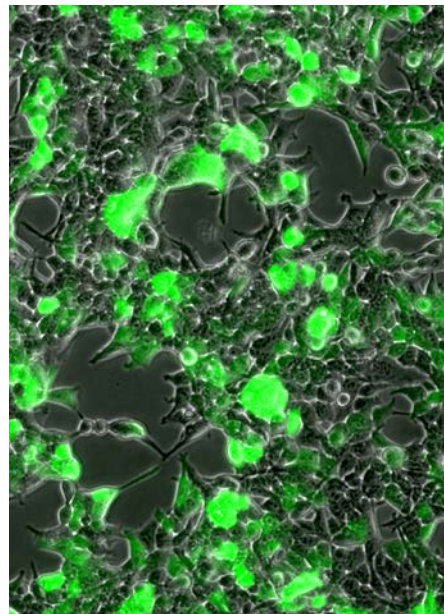
NucleoSnap Plasmid Midi

The fastest way to isolate transfection-grade plasmid DNA

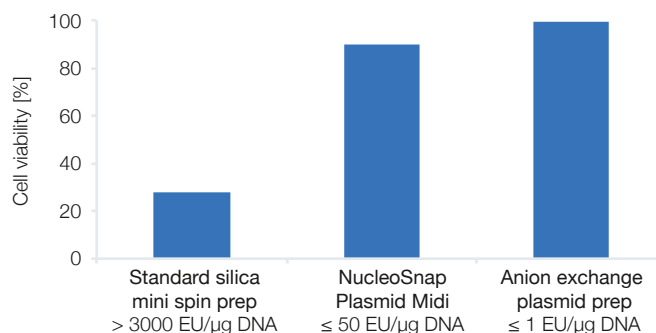
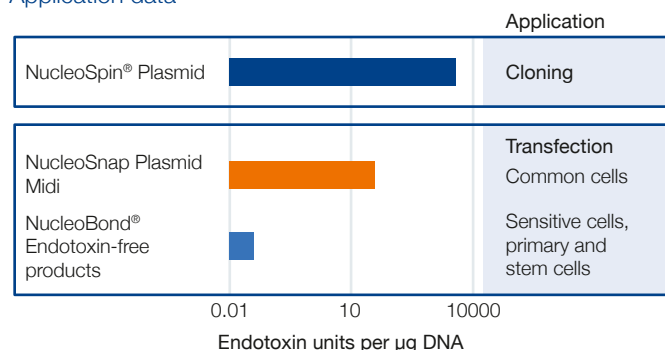
Endotoxins are co-purified during plasmid preparations from bacterial lysates. Since they interfere with eukaryotic cell survival, endotoxin reduction is essential prior to cell transfection. MACHEREY-NAGEL has developed a new column format and a novel buffer chemistry to enable vacuum-processed isolation of transfection-grade plasmid DNA in a midi format.

Product at a glance

| | |
|------------------------------------|--|
| Technology | Silica-membrane technology |
| Format | Snap-off column, vacuum processing |
| Lysate clarification | Large filter spin columns (2 min centrifugation) |
| Sample material | Typically 50 mL <i>E. coli</i> culture (OD ₆₀₀ = 5) |
| Vector size | ≤ 25 kbp |
| Typical yield | 400–700 µg (50 mL culture, OD ₆₀₀ = 4, high-copy plasmid) |
| A ₂₆₀ /A ₂₈₀ | 1.8–1.9 |
| Preparation time | 35 min/6 preps |
| Endotoxin level | < 50 EU/µg DNA |
| Binding capacity | 1.5 mg |



Application data



Endotoxin levels appropriate for individual applications

A quantitative chromogenic LAL-test was used to assess endotoxin content. As indicated, the content of endotoxin is strongly depended on the technology of plasmid purification. Low endotoxin levels were detected after purification with NucleoSnap Plasmid Midi resulting in a plasmid solution directly appropriate for transfection of common cells.

High cell viabilities of eukaryotic cells

Eukaryotic Huh-7 cells were transfected with Lipofectamin 2000 and 2.5 µg plasmid DNA (pCMV-GFP, kindly provided by PlasmidFactory GmbH & Co. KG, Bielefeld, Germany). The plasmid DNA was prepared with a standard silica mini spin prep (such as NucleoSpin® Plasmid), NucleoSnap Plasmid Midi and an anion exchange DNA isolation kit (such as NucleoBond® Xtra Midi EF).

Ordering information

| Product | Specifications | Preps / Pack of | REF |
|---|--|---------------------------|------------------------|
| NucleoSnap Plasmid Midi | Kit for the isolation of up to 700 µg transfection-grade plasmid DNA | 10 / 50 | 740494.10 / .50 |
| NucleoVac 24 Vacuum Manifold | Vacuum manifold for processing NucleoSnap or NucleoSpin® columns | 1 | 740299 |
| NucleoVac Mini Adapter | Disposable adapters for processing NucleoSpin® or NucleoSnap columns on the NucleoVac 24 Vacuum Manifold | 100 | 740297.100 |
| NucleoVac Stop-cock | Stop-cocks for processing samples with different flow rates on the NucleoVac 24 Vacuum Manifold | 24 | 740298.24 |
| NucleoSpin® Plasmid Transfection-grade | Mini spin kit for the isolation of transfection-grade plasmid DNA | 10 / 50 / 250 | 740490.10 / .50 / .250 |
| NucleoSpin® 96 Plasmid Transfection-grade | For isolation of transfection-grade plasmid DNA in 96-well format | 1 x 96 / 4 x 96 / 24 x 96 | 740491.1 / .4 / .24 |

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MACHEREY-NAGEL



MACHEREY-NAGEL GmbH & Co. KG · Neumann-Neander-Str. 6–8 · 52355 Düren · Germany

DE / International:

Tel.: +49 24 21 969-0

Fax: +49 24 21 969-199

E-mail: info@mn-net.com

CH:

Tel.: +41 62 388 55 00

Fax: +41 62 388 55 05

E-mail: sales-ch@mn-net.com

FR:

Tel.: +33 388 68 22 68

Fax: +33 388 51 76 88

E-mail: sales-fr@mn-net.com

US:

Tel.: +1 484 821 0984

Fax: +1 484 821 1272

E-mail: sales-us@mn-net.com

