



nexttecTM
DNA isolation systems

Protocol

DNA Isolation from Tissue & Cells by nexttecTM 1-Step

- nexttecTM cleanPlate96 -

| | |
|-------------------------|---------|
| Cat. No. 10N.901 | 391091N |
| Cat. No. 10N.902 | 391092N |
| Cat. No. 10N.904 | 391094N |
| Cat. No. 10N.924 | 391096N |

Version 3.0

For research only

Biozym
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Principle

nexttec™ 1-Step is the easiest handling and fastest DNA purification system containing a single buffer system and a one-step DNA purification after lysis.

Proteins, detergents and low molecular weight compounds are retained by the nexttec™ sorbent. DNA passes through the nexttec™ cleanColumn during a short, one-step purification procedure.

The obtained DNA is suitable for all common enzymatic reactions (restriction digests, real-time PCR, PCR, genotyping etc.).

Kit contents

The kit contains all necessary reagents for lysis and subsequent DNA purification.

| Component | Art.No. 10N.901 | Art.No. 10N.902 | Art.No. 10N.904 | Art.No. 10N.924 |
|--------------------------------------|--------------------|--------------------|--------------------|--------------------|
| Buffer G | 15 ml | 42 ml | 65 ml | 400 ml |
| Proteinase K | 1.5 ml | 3 ml | 4.5 ml | 72 ml |
| Prep Solution | 40 ml | 100 ml | 150 ml | 2 x 450 ml |
| DTT (optional for special protocols) | 0.5 ml | 1.0 ml | 1 ml | 5 ml |
| nexttec™ cleanPlates96 | 1 | 2 | 4 | 24 |
| nexttec™ deep-well plate | 3 | 6 | 12 | 72 |
| Sealing tapes | 3 | 6 | 12 | 72 |
| Alu sealing tapes | 2 | 4 | 8 | 48 |

nexttec™ service

To extend the application range to samples, which are difficult to lyse by the standard procedure, it is recommended to include optional components in the lysis buffer and to optimize the lysis conditions. Please get in contact with service@nexttec.biz for detailed information.

Storage Conditions

During shipment all kit components are stable at room temperature. After arrival, store **Prep Solution** and **Proteinase K** at **+2 °C to +8 °C**. After first opening freeze **DTT** at **-18 °C to -25 °C**.

Buffer G and **nexttec™ cleanPlates96** are stored at **room temperature (+20 °C to +25 °C)**. If properly stored, see expiration date for the stability of the kit.

Safety Information

Proteinase K Danger



H334

P304+P341, P342+P311

DTT Warning



H315, H319

P280, P305+P351+P338, P321, P362, P332+P313,

P337+P313

Hazard Statements

H315 Causes skin irritation

H319 Causes serious eye irritation

H334 May cause allergy or asthma symptoms or breathing difficulties if inhaled

Precautionary Statements

| | |
|-----------------|---|
| P280 | Wear protective gloves/protective clothing/eye protection/ face protection |
| P342+P311 | If experiencing respiratory symptoms: Call a POISON CENTER or doctor/physician. |
| P305+P351+ P338 | IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing |
| P304+P341 | IF INHALED: if breathing is difficult, remove victim to fresh air and keep at rest in a position comfortable for breathing |
| P321 | Specific treatment (see on this label) |
| P362 | Take off contaminated clothing and wash before reuse |
| P332+P313 | If skin irritation occurs: Get medical advice/attention |
| P337+P313 | If eye irritation persists: Get medical advice/attention |

When working with chemicals, always wear a suitable lab coat, disposable gloves and protective goggles. For more information please consult the appropriate material safety data sheets (MSDS).

Before starting

- **Equilibrate nexttec™ cleanPlates96**

| | |
|----|---|
| E1 | Add 350 µl Prep Solution onto each well of a nexttec™ cleanPlate96 . Incubate for at least 5 min at room temperature and centrifuge at 350 x g for 1 min or apply vacuum for 30 to 60 sec to remove excess buffer. |
| E2 | Discard the Waste deep-well plate. Place the nexttec™ cleanPlate96 onto a new deep-well plate. Use equilibrated nexttec™ cleanPlates96 or store closed at +2 °C to +8 °C and use within one week . |

- **Preheat an incubator to 56 °C**

Protocol

Lysis

| | |
|----|--|
| L1 | Transfer tissue (5-20 mg fresh weight) or cells (1-2 x 10⁶) to a reaction tube. Centrifuge cell suspensions (2,000 x g, 10 min), remove and discard the supernatant. |
| L2 | Add 140 µl Buffer G, 10 µl Proteinase K and <i>optional 1.5 µl DTT*</i> to each sample. Close deep-well plate using an alu sealing tape. Incubate the sample with shaking (56 °C, 1200 rpm, 30 min to overnight). |

*For Pre-Mixes see Technical Section.

Purification of DNA

| | |
|---|--|
| P | Transfer 100 µl of the lysate to an equilibrated nexttec™ cleanPlate96 . Incubate for 3 min at room temperature . Centrifuge at 700 x g for 1 min or apply vacuum for 1 min . |
| The eluate contains the purified DNA!! | |

Notes:

Technical Section**Preparation of Lysis Pre-Mixes**

| | | | | | |
|--|-------------------------|----------|---------|----------|----------|
| LG | Lysis Buffer LG: | 1 sample | 1 plate | 2 plates | 4 plates |
| | Buffer G | 140 µl | 15.4 ml | 30.8 ml | 61.6 ml |
| | Proteinase K | 10 µl | 1.1 ml | 2.2 ml | 4.4 ml |
| | DTT | 1.5 µl | 165 µl | 330 µl | 660 µl |
| Mix by vortexing. Add 150 µl of Buffer LG to each sample (L2). The Lysis Buffer LG is stable for 1 working day if stored at +2 °C to +8 °C . | | | | | |

- **Determination of DNA concentration in nexttec™ 1-Step DNA preparations**

We recommend to determine the DNA concentration:

- Using the fluorescent dye Picogreen® or similar.
- Comparing the fluorescence intensity of DNA bands of unknown concentrations with standards, e.g. in ethidium bromide stained agarose gels.

Please notice:

The use of absorption measurement at 260 nm (A_{260}) in a spectrophotometer (e.g. NanoDrop®) for determination of DNA concentration is system related not recommended.

For details and possible workarounds for your specific application please contact:

service@nexttec.biz.

- **Centrifugation**

For centrifugation at 350x g or 700x g use the settings for relative centrifugal force (RCF) of your centrifuge. Alternatively measure the distance of the nexttec™ cleanPlate96 to the centre of your rotor and calculate the necessary rotations per minute (e.g. rpm = $299.07 \times \sqrt{350 / r}$; r=radius in cm).

- **Vacuum Application**

Assemble the vacuum manifold as described in the manual. The flow rate should be between 15 L/min and 150 L/min.

A regulation of vacuum is not necessary.

Product Use Restriction

nexttec™ 1-Step DNA Isolation Kit components were developed, designed and sold **for research purposes only**. They are suitable for in vitro uses only. No claim or representation is intended for use to identify any specific organism or for clinical use.

It is the responsibility of the user to verify the use of the nexttec™ 1-Step DNA Isolation Kit for a specific application as the performance characteristic of this kit has not been verified to a specific organism.

Troubleshooting, FAQ and Special Applications

Product claims are subject to change. Therefore, please, visit our website or contact our technical service team for troubleshooting guide, up-to-date protocols and latest applications on nexttec™ 1-Step products.

Contact Information

Web: www.nexttec.biz

Email: service@nexttec.biz

Tel: +49 (0) 8250 / 92 790 32

Fax: +49 (0) 8250 / 92 790 99

Ordering Information

For ordering information please visit our website www.nexttec.biz.

Distributor

Biozym
SCIENCE IS OUR BUSINESS
www.biozym.com

Biozym Scientific GmbH
Tel.: 05152 / 9020, Fax: 05152 / 2070
Mail: support@biozym.com

Biozym Biotech Trading GmbH
Tel.: 01 / 334 0156 0, Fax: 01 / 334 0156 88
Mail: support@biozym.com



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