



Protocol

DNA Isolation from Plant by nexttec[™] 1^{-Step}

- nexttec™ cleanColumns -

Cat. No. 47N.010 394010N

Cat. No. 47N.050 394020N

Cat. No. 47N.250 394025N

Version 3.0

For research only





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 nexttecTM sorbent. DNA passes through the nexttecTM cleanColumn during a short, <u>one-step</u> 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. 47N.010 | Art.No. 47N.050 | Art.No. 47N.250 |
|-------------------------------|--------------------|--------------------|--------------------|
| Buffer F | 4.2 ml | 21 ml | 84 ml |
| Protease | 0.3 ml | 1.5 ml | 6 ml |
| Prep Solution | 6 ml | 20 ml | 100 ml |
| Solution H | 3 ml | 15 ml | 75 ml |
| SDS | 0.5 ml | 0.75 ml | 3 ml |
| DTT (1,4- Dithio-DL-threitol) | 0.1 ml | 0.25 ml | 1.0 ml |
| RNase A | 0.2 ml | 1 ml | 4 ml |
| nexttec™ cleanColumns | 10 | 50 | 250 |
| Waste collection tubes | 10 | 50 | 250 |
| DNA collection tubes | 10 | 50 | 250 |

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**, **Solution H**, **Protease** and **RNase A at +2 °C to +8 °C**. After first opening freeze **DTT** at -18 °C to -25 °C.

SDS, Buffer F and nexttec[™] cleanColumns are stored at room temperature (+20 °C to +25 °C). If properly stored, see expiration date for the stability of the kit.

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Safety Information

Protease Danger H315, H318, H334, H335

P261, P280, P342+P311, P305+P351+P338

DTT Warning H315, H319

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

P337+P313

SDS Danger H315, H318

P305+P351+P338

Hazard Statements

H315 Causes skin irritation

H318 Causes serious eye damage

H319 Causes serious eye irritation

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

H335 May cause respiratory irritation

Precautionary Statements

P261 Avoid breathing dust/fume/gas/mist/vapors/spray

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

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).

Technical Support





Before starting

• Equilibrate nexttec[™] cleanColumns

| E1 | Add 350 µl Prep Solution to a nexttec[™] cleanColumn . Incubate for at least 5 min at room temperature. Centrifuge at 350 x g for 1 min to remove excess buffer. |
|----|---|
| E2 | Discard the Waste collection tube. Place the nexttec [™] cleanColumn into a new DNA collection tube. Use equilibrated nexttec [™] cleanColumns or store closed at +2 °C to +8 °C and use within one week. |

Preheat a thermomixer to 60 °C



Protocol

Lysis

A: Without homogenising (using leave pieces)

| L1 | Transfer 3 -10 pieces of leaf tissue (5x5 mm) to a reaction tube. |
|----|---|
| L2 | Add 280 Buffer F and 20 µl Protease* . Incubate with shaking (60 °C, 1200 rpm, 30 min to overnight) in a thermomixer |

B: With homogenising

| L1 | Transfer 50 mg plant leaf tissue to a reaction tube. |
|----|---|
| L2 | Add 185 µl Solution H, 3 µl DTT and 12 µl RNase* and homogenise tissue in a bead mill or by grinding with a pestle. |
| L3 | Add 70 µl Buffer F , 20 µl Protease and 10 µl SDS*. Incubate with shaking (60 °C, 1200 rpm, 30 min) in a thermomixer. |
| L4 | Centrifuge the lysate (10,000 x g, 1 min). Use clear supernatant for DNA purification. |

C: Plant seeds

| Add 280 µl Buffer F, 20 µl Protease and 3 µl DTT*. Incubate with shaking (60 °C, 1200 rpm, 30 min) in a thermomixer. Centrifuge the lysate at 10,000 x g for 1 min | L1 | Transfer up to 20 mg seed meal, grist or crude squashed seeds to a reaction tube. |
|--|----|---|
| Centrifuge the lysate at 10,000 x g for 1 min | L2 | · |
| Use the clear supernatant for DNA purification. | L3 | Centrifuge the lysate at 10,000 x g for 1 min Use the clear supernatant for DNA purification. |

^{*}For Pre-Mixes see Technical Section.

Purification of DNA from lysis A, B and C

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



Technical Section

Preparation of Lysis Pre-Mixes

• A: without homogenising (using leave pieces)

| | Lysis Buffer LF: | 1 sample | <50 samples* | >50 samples* |
|----|--|----------|---------------|---------------|
| | 280 μl x (n+5) | | | |
| LF | Protease | 20 μΙ | 20 μl x (n+3) | 20 μl x (n+5) |
| | Mix by vortexing. Add 300 μl of Buffer LF to each sample (L2). The Lysis Buffer LF is | | | |
| | stable for 1 working day if stored at +2 °C to +8 °C. | | | |

B: with homogenising

| | Lysis Buffer LF1: | 1 sample | <50 samples* | >50 samples* | |
|-----|--|----------|----------------|----------------|--|
| LF1 | Solution H | 185 µl | 185 µl x (n+3) | 185 μl x (n+5) | |
| | DTT | 3 µl | 3 µl x (n+3) | 3 μl x (n+5) | |
| | RNase | 12 µl | 12 µl x (n+3) | 12 μl x (n+5) | |
| | Mix by vortexing. Add 200 μl of Buffer LF1 to each sample (L2). The Lysis Buffer LF1 is | | | | |
| | stable for 1 working day if stored at +2 °C to +8 °C. | | | | |
| | Lysis Buffer LF2: | 1 sample | <50 samples* | >50 samples* | |
| LF2 | Buffer F | 70 µl | 70 μl x (n+3) | 70 μl x (n+5) | |
| | Protease | 20 μΙ | 20 μl x (n+3) | 20 μl x (n+5) | |
| | SDS | 10 µl | 10 μl x (n+3) | 10 μl x (n+5) | |
| | Mix by vortexing. Add 100 µl of Buffer LF2 to each sample (L3). The Lysis Buffer LF2 is stable for 1 working day if stored at +2 °C to +8 °C . | | | | |

• C: for plant seeds

| | Lysis Buffer LF: | 1 sample | <50 samples* | >50 samples* |
|----|--|----------|----------------|----------------|
| | Buffer F | 280 µl | 280 µl x (n+3) | 280 μl x (n+5) |
| LF | Protease | 20 µl | 20 μl x (n+3) | 20 μl x (n+5) |
| | DTT | 3 µl | 3 µl x (n+3) | 3 μl x (n+5) |
| | Mix by vortexing. Add 300 μl of Buffer LF to each sample (L2). The Lysis Buffer LF is | | | |
| | stable for 1 working day if stored at +2 °C to +8 °C. | | | |

^{*}n= samples [e.g. 22 samples: Buffer F: 280 µl x (22+3)]

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Plant Version 3.0

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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 (A₂₆₀) 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 nexttecTM cleanColumn to the centre of your rotor and calculate the necessary rotations per minute (e.g. rpm = 299.07 x

 $\sqrt{350}/r$; r=radius in cm).

Product Use Restriction

nexttecTM 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

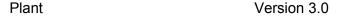
specific organism.

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 nexttecTM 1^{-Step} products.

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Contact Information

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Ordering Information

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

Distributor



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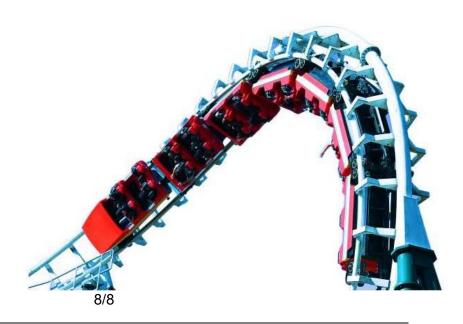


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