



nexttecTM
DNA isolation systems

Protocol

DNA Isolation

from Plant

by nexttecTM 1-Step

- nexttecTM cleanColumns -

Cat. No. 47N.010 394010N

Cat. No. 47N.050 394020N

Cat. No. 47N.250 394025N

Version 3.0

For research only

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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. 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.

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).

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

L1	Transfer up to 20 mg seed meal, grist or crude squashed seeds to a reaction tube.
L2	Add 280 µl Buffer F , 20 µl Protease and 3 µl DTT* . Incubate with shaking (60 °C, 1200 rpm, 30 min) in a thermomixer.
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

P	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)**

LF	<u>Lysis Buffer LF:</u>	1 sample	<50 samples*	>50 samples*
	Buffer F	280 µl	280 µl x (n+3)	280 µl x (n+5)
	Protease	20 µl	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**

LF1	<u>Lysis Buffer LF1:</u>	1 sample	<50 samples*	>50 samples*
	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 .				
LF2	<u>Lysis Buffer LF2:</u>	1 sample	<50 samples*	>50 samples*
	Buffer F	70 µl	70 µl x (n+3)	70 µl x (n+5)
	Protease	20 µl	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**

LF	<u>Lysis Buffer LF:</u>	1 sample	<50 samples*	>50 samples*
	Buffer F	280 µl	280 µl x (n+3)	280 µl x (n+5)
	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)]

- **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™ cleanColumn 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).

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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