



**nexttec**<sup>TM</sup>  
DNA isolation systems

**Protocol**

**DNA Isolation  
from Blood (200 µl)  
by nexttec<sup>TM</sup> 1-Step**

- nexttec<sup>TM</sup> cleanColumns -

<b>Cat. No. 50N.010</b>	395010N
<b>Cat. No. 50N.050</b>	395020N
<b>Cat. No. 50N.250</b>	395025N

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. 50N.010	Art.No. 50N.050	Art.No. 50N.250
Buffer R	2 ml	10 ml	38 ml
Proteinase K	0.3 ml	1.5 ml	6 ml
Prep Solution	6 ml	20 ml	100 ml
Solution B	2 x 10 ml	2 x 50 ml	2 x 200 ml
DTT	0.2 ml	0.2 ml	0.5 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](mailto:service@nexttec.biz) for detailed information.

## Storage Conditions

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

**Buffer R** 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**

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™ cleanColumns**

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

- **Preheat a thermomixer to 56 °C**

## Protocol

### Lysis

L1	Transfer <b>200 µl of EDTA</b> or <b>heparinized blood</b> to a reaction tube. Add <b>600 µl cold Solution B</b> . Invert tube several times to mix the contents thoroughly ( <u>don't vortex!</u> ).
L2	<b>Incubate 5 min on ice.</b> Centrifuge the mixture ( <b>3 min, 6,000 x g</b> ), remove <b>700 µl</b> carefully and discard the supernatant.
L3	Add <b>500 µl cold Solution B</b> . Invert tube several times to mix the contents thoroughly ( <u>don't vortex!</u> ).
L4	<b>Incubate 5 min on ice.</b> Centrifuge the mixture ( <b>3 min, 6,000 x g</b> ), remove <b>550 µl</b> carefully and discard the supernatant.
L5	Add <b>125 µl Buffer R</b> , <b>20 µl Proteinase K</b> and <b>1.5 µl DTT*</b> to each sample. Resuspend pellet by pipetting up and down ( <b>3 times</b> ) or by vortexing. Incubate with shaking ( <b>56 °C, 1200 rpm, 30 min</b> ) in a thermomixer.

\*For Pre-Mixes see Technical Section.

### Purification of DNA

P	Transfer <b>100 µl</b> of the lysate to an <b>equilibrated nexttec™ cleanColumn</b> . Incubate for <b>3 min</b> at <b>room temperature</b> . Centrifuge at <b>700 x g</b> for <b>1 min</b> .
	<b>The eluate contains the purified DNA!!</b>

### Notes:

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

LR	<b>Lysis Buffer LR:</b>	1 sample	<50 samples*	>50 samples*
	<b>Buffer R</b>	125 µl	125 µl x (n+3)	125 µl x (n+5)
	<b>Proteinase K</b>	20 µl	20 µl x (n+3)	20 µl x (n+5)
	<b>DTT</b>	1.5 µl	1.5 µl x (n+3)	1.5 µl x (n+5)
Mix by vortexing. Add <b>146 µl</b> of <b>Buffer LR</b> to each sample (L5). The <b>Lysis Buffer LR</b> is stable for 1 working day if stored at <b>+2 °C to +8 °C</b> .				

\*n= samples [e.g. 22 samples: Buffer R: 125 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](mailto: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](http://www.nexttec.biz)

Email: [service@nexttec.biz](mailto: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](http://www.nexttec.biz).

### **Distributor**

**Biozym**  
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[www.biozym.com](http://www.biozym.com)

**Biozym Scientific GmbH**  
Tel.: 05152 / 9020, Fax: 05152 / 2070  
Mail: [support@biozym.com](mailto:support@biozym.com)

**Biozym Biotech Trading GmbH**  
Tel.: 01 / 334 0156 0, Fax: 01 / 334 0156 88  
Mail: [support@biozym.com](mailto:support@biozym.com)

