



**Protocol**  
**Plasmid DNA Isolation**  
**from Bacteria (E.coli)**  
**by nexttec™ 1<sup>-Step</sup>**

**- nexttec™ cleanColumns -**

**Cat. No. 30N.010**

**Cat. No. 30N.050**

**Cat. No. 30N.250**

**Version 1.0**

**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 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.

<b>Component</b>	<b>Art. No. 30N.010</b>	<b>Art. No. 30N.050</b>	<b>Art. No. 30N.250</b>
<b>Buffer P</b>	12 ml	12 ml	55 ml
<b>Protease</b>	0.5 ml	0.5 ml	2.0 ml
<b>Prep Solution</b>	6 ml	20 ml	100 ml
<b>RNase A</b>	1.0 ml	1.0 ml	5.0 ml
<b>Lysozyme</b>	12.5 mg (0.5 ml)	12.5 mg (0.5 ml)	37.5 mg (1.5 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 of samples, which are difficult to lyse by the standard procedure, it is recommended to include optional components in the lysis buffer and optimize the lysis time. 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, **Buffer P, Protease, Prep Solution, RNase A and Lysozyme must be stored at +2°C to +8°C.**

nexttec™ cleanColumns can be stored at **room temperature (+20°C to +25°C)**. If properly stored, see expiration date for the stability of the kit.

## **Safety Information**

**Protease** Xn R 36/37/38 R42/43; S 23-24-26 36/37

### Risk Phrases

**R 42/43:** May cause sensitisation by inhalation and skin contact.  
**R 36/37/38:** Irritating to eyes, respiratory system and skin.

### Safety Phrases

**S 23:** Do not breathe gas/fumes/vapour/spray.  
**S 24:** Avoid contact with skin.  
**S 26:** In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.  
**S 36/37:** Wear suitable protective clothing and gloves.

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>350x 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 block thermostat to 90°C**
- **Dissolve Lysozyme**  
Add **0.5 ml** (30.010 / 30.050) or **1.5 ml** (30.250) of **purified water** to the **lyophilized lysozyme** (final concentration: 25 mg/ml).  
Mix thoroughly by vortexing and store at **-18°C to -25°C**.

### Protocol

## Lysis

L1	Transfer <b>1 ml</b> overnight <b>bacteria culture</b> into a reaction tube. Centrifuge ( <b>6,000x g, 1 min</b> ), remove and discard supernatant.
L2	Add <b>130 µl Buffer P, 5 µl Protease</b> and <b>3 µl Lysozyme*</b> to the cell pellet. Dissolve cell pellet by vortexing or pipetting up and down. Incubate for <b>5 min</b> at <b>room temperature</b> .
L3	Transfer the tube to a preheated block thermostat. Incubate ( <b>90°C, 1 min</b> ), then cool down the solution to <b>room temperature</b> .
L4	Add <b>15 µl RNase A</b> to the lysate and mix gently ( <u>do not vortex!</u> ) Incubate for <b>3 min</b> at <b>room temperature</b> .

\*For Pre-Mixes see Technical Section.

## Purification of DNA

P	Transfer the complete <b>lysate</b> onto an <b>equilibrated nexttec™ cleanColumn</b> (use 1 ml or 200 µl wide bore pipette tips). Incubate for <b>3 min</b> at <b>room temperature</b> . Centrifuge at <b>700x 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 buffers (Pre-Mixes)**

LP	<b>Lysis Buffer LP:</b>	1 sample	<50 samples*	>50 samples*
	<b>Buffer P</b>	130 µl	130 µl x (n+3)	130 µl x (n+5)
	<b>Protease</b>	5 µl	5 µl x (n+3)	5 µl x (n+5)
	<b>Lysozyme</b>	3 µl	3 µl x (n+3)	3 µl x (n+5)
Mix by vortexing. Add <b>138 µl of Buffer LP1</b> to each sample (L2). The <b>Lysis Buffer LP</b> is stable for 1 working day, if stored <b>at +2°C to +8°C</b> .				

\*n= samples [e.g. 22 samples: Buffer G: 140 µl x (22+3)]

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

We recommend determining the DNA concentration:

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

Please notice:

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

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 of 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.



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