



Special Protocol DNA Isolation Drosophila

by nexttec™ 1<sup>-Step</sup>

- nexttec™ cleanColumns -

Cat. No. 10N.010 391010N Cat. No. 10N.050 391020N Cat. No. 10N.250 391025N

Version 3.0

For research only





# <u>Principle</u>

nexttec<sup>TM</sup> 1<sup>-Step</sup> is the easiest handling and fastest DNA purification system containing a <u>sin-</u><u>gle</u> buffer system and a <u>one-step</u> DNA purification after lysis.

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

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

# Kit contents

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

Component	Art. No. 10N.010	Art. No. 10N.050	Art. No. 10N.250
Buffer G	1x 2.1 ml	1x 10.5 ml	1x 42 ml
Proteinase K	1x 0,15 ml	1x 0,75 ml	1x 3 ml
Prep Solution	1x 6 ml	1x 20 ml	1x 100 ml
DTT (1,4- Dithio-DL-threitol)	1x 0.5 ml	1x 0.5 ml	1x 0.5 ml
nexttec <sup>™</sup> cleanColumns	10	50	250
Waste collection tubes	10	50	250
DNA collection tubes	10	50	250

## <u>nexttec<sup>™</sup> service</u>

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 time. Please get in contact with <u>service@nexttec.biz</u> for detailed information.

# **Storage Conditions**

During shipment all kit components are stable at room temperature. After arrival, **Proteinase K solution**, **Buffer G** and **Prep Solution** must be stored **at +2°C to +8°C**.

Store **DTT solution** after first opening **at -18°C to -25°C.** nexttec<sup>™</sup> 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**

Proteinase K	Danger
(!)	

H334 P304+P341, P342+P311



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 pro-
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. Re- move 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).

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# Before starting

# • Equilibrate nexttec<sup>™</sup> cleanColumns

E1	Add <b>350 µl Prep Solution</b> to a <b>nexttec<sup>™</sup> 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<sup>™</sup> cleanColumn</b> into a new DNA collection tube. Use equilibrated nexttec <sup>™</sup> 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**

<u>Lysis</u>			
L1	Transfer a frozen or anesthetised fly ( <i>D. melanogaster</i> ) to a reaction tube.		
L2	Add <b>140 µl Buffer G</b> , <b>10 µl Proteinase K and 1,5 µl DTT</b> * to each sample. Incubate the sample with shaking ( <b>56°C</b> , <b>1200 rpm</b> , <b>30 min to overnight)</b> in a thermomixer.		
*For Pre-Mixes see Technical Section.			
Purification of DNA			
	Transfer <b>100 µI</b> of the lysate to an <b>equilibrated nexttec™ cleanColumn</b> .		
Р	Incubate for <b>3 min</b> at <b>room temperature</b> .		
	Centrifuge at <b>700x g</b> for <b>1 min.</b>		
	The eluate contains the purified DNA!!		

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# **Technical Section**

## • Preparation of Lysis Pre-Mix

	Lysis Buffer LG:	1 sample	<50 samples*	>50 samples*
	Buffer G	140 µl	140 µl x (n+3)	140 µl x (n+5)
LG	Proteinase K	10 µl	10 µl x (n+3)	10 µl x (n+5)
	DTT	1.5 µl	1.5 µl x (n+3)	1.5 µl x (n+5)
	Mix by vortexing. Add <b>150 µl of Buffer LG</b> to each sample (L2). The <b>Lysis Buffer LG</b> is stable for <b>1</b> 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<sup>™</sup> 1<sup>-Step</sup> 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 concentration with standards, e.g. in ethidium bromide stained agarose gels.

Please notice:

The use of absorption measurement at 260nm (A<sub>260</sub>) in a spectrophotometer (e.g. NanoDrop<sup>®</sup>) for determination of DNA concentration <u>is system related</u> 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<sup>TM</sup> 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

nexttec<sup>TM</sup> 1<sup>-Step</sup> 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<sup>™</sup> 1<sup>-Step</sup> DNA Isolation Kit for a specific application as the performance characteristic of this kit has not been verified to a specific organism.

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## **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<sup>™</sup> 1<sup>-Step</sup> products.

### **Contact Information**

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### **Ordering Information**

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

### **Distributor**



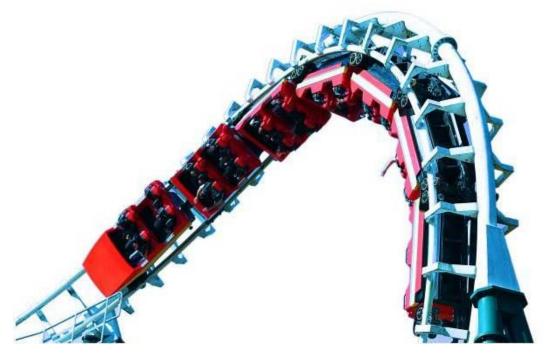
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