

Genomic DNA from blood

User manual

NucleoSpin® Dx Blood

CE

IVD

In Vitro Diagnostic Medical Device

REF

740899.50, 740899.250



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1 Components

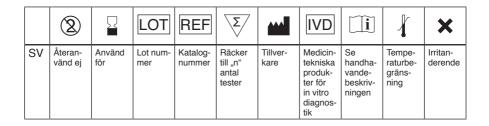
1.1 Kit contents

		NucleoSpin® Dx Blood		
REF		50 preps 740899.50	250 preps 740899.250	
Buffer B3	BUF B3	13 mL	65 mL	
Wash Buffer BW	BUF BW	30 mL	2 x 75 mL	
Wash Buffer B5 (Concentrate)*	BUF B5 conc.	7 mL	2 x 20 mL	
Elution Buffer BE**	BUF BE	13 mL	60 mL	
Proteinase Buffer PB	BUF PB	1.8 mL	8 mL	
Proteinase K (lyophilized)*	Proteinase K	30 mg	2 x 75 mg	
NucleoSpin® Dx Blood Columns (red rings - plus Collection Tubes)	Dx Blood Columns	50	250	
Collection Tubes (2 mL)	Collection Tubes	3 x 50	3 x 250	
Lysis Tubes (1.5 mL)	Lysis Tubes	50	5 x 50	
Elution Tubes (1.5 mL)	Elution Tubes	50	5 x 50	
User manual	ĺĺį	1	1	

^{*} For preparation of working solutions and storage conditions see section 3.

^{**} Composition of Elution Buffer BE: 5 mM Tris/HCl, pH 8.5

	(3)	Ω	LOT	REF	Σ	AAA	IVD	[]	1	•
	(8)		LOT	NEF			IVD		1	^
EN	Do not reuse	Use by	Batch code	Cata- logue number	Contains sufficient for <n> tests</n>	Manuf- acturer	In Vitro Diag- nostic Medical Device	Consult Instruc- tions for use	Tempe- rature limitation	Irritant
DE	Nicht zur Wieder- verwen- dung	Ver- wendbar bis	Char- genbe- zeich- nung	Bestell- nummer	Inhalt ausrei- chend für <n> Tests</n>	Herstel- ler	In vitro-Di- agnosti- kum	Ge- brauchs- anwei- sung beach- ten	Tempe- ratur- begren- zung	Reizend
ES	No reuti- lizar	Fecha de cadu- cidad	Código de lote	Número de catá- logo	Contenido suficiente para <n> ensayos</n>	Fabri- cante	Pro- ducto sanitario para diagnós- tico in vitro	Consul- te las instruc- ciones de uso	Límite de tempe- ratura	Irritante
IT	Non riutiliz- zare	Utilizza- re entro	Codice del lotto	Numero di cata- logo	Conte- nuto suf- ficiente per "n" saggi	Fabbri- cante	Dispositivo medicodiagnostico in vitro	Consultare le istruzioni per l'uso	Limiti di tempe- ratura	Irritant
FR	Ne pas réutiliser	Utiliser jusque	Code du lot	Réfé- rence du catalo- gue	Contenu suffisant pour "n" tests	Fabri- cant	Dis- positif médical de dia- gnostic in vitro	Consulter les instructions d'utilisation	Limites de tempé- rature	Irritante
NL	Niet opnieuw gebrui- ken	Houd- baar tot	Lot num- mer	Cata- logus nummer	Inhoud vol- doende voor "n" testen	Fabri- kant	Medisch hulpmid- del voor in vitro diagnos- tiek	Raad- pleeg de ge- bruiks- aanwij- zing	Tempe- ratuurli- miet	Irritie- rend
DA	Må ikke genbru- ges	Holdbar til	Lotnum- mer	Katalog- nummer	Inde- holder tilsttræk- keligt til "n" test	Produ- cent	Medi- cinsk udstyr til in vitro- diagno- stik	Se brugsan- visning	Tempe- raturbe- græns- ning	Lokalirri- terende
EL	Μην κάνετε επα-να- ληπτική χρήση	Ημεφο- μηνία λήξης	Αφιθμός Παφτί- δας	Αφιθμός καταλό- γου	Περιε- χόμενο επαρκές για «ν» εξετά- σεις	Κατα- σκευα- στής	In Vitrο Διαγνω- στικό Ιατφο- τεχνολο- γικό πφοϊόν	Συμβου- λευτείτε τις οδηγίες χρήσης	Περιο- οισμοί θερμο- κοασίας	Διαβοω- τιχό
PT	Não reutilizar	Prazo de vali- dade	Código do lote	Referên- cia de catálogo	Conteú- do su- ficiente para "n" ensaios	Fabri- cante	Dispo- sitivo médico para dia- gnóstico in vitro	Consulte as in- struções de utili- zação	Limites de tempe- ratura	Irritante



1.2 Reagents, consumables, and equipment to be supplied by user

Reagents

 96–100% ethanol (to adjust DNA binding conditions and to prepare Wash Buffer B5)

Consumables

 Disposable pipet tips (aerosol barrier pipet tips are recommended to avoid cross-contamination)

Equipment

- Manual pipettors
- · Centrifuge for microcentrifuge tubes
- Vortex mixer
- Thermal heating block or water bath (for samples lysis at 70°C)
- Personal protection equipment (e.g., lab coat, gloves, goggles)

1.3 About this user manual

It is strongly recommended that first-time users of the **NucleoSpin® Dx Blood** kit read the detailed protocol sections of this user manual. Experienced users, however, may refer to the Protocol-at-a-glance instead. The Protocol-at-a-glance is designed to be used only as a supplemental tool for quick referencing while performing the purification procedure.

All technical literature is available on the internet at www.mn-net.com.

Please contact Technical Service regarding information about changes of the current user manual compared to previous revisions.

2 Product description

2.1 Intended use

The **NucleoSpin® Dx Blood** kit is a generic system for the isolation and purification of genomic DNA from human whole blood samples for subsequent *in vitro* diagnostic purposes. The kit can be used with fresh and frozen human whole blood treated with EDTA, citrate, or heparin, from common blood collection systems.

The kit is designed to be used with any downstream application employing enzymatic amplification and detection of DNA (e.g., PCR). Any diagnostic results generated using the DNA isolated with the **NucleoSpin® Dx Blood** kit in conjunction with an *in vitro* diagnostic assay should be interpreted with regard to additional clinical or laboratory findings.

To minimize irregularities in diagnostic results, suitable controls for downstream applications (e.g., extraction controls, positive/negative controls) should be used.

The **NucleoSpin® Dx Blood** kit is intended for use by professional users such as technicians and physicians experienced and trained in molecular biological techniques including experience with whole blood samples and DNA isolation.

The **NucleoSpin® Dx Blood** kit does not provide a diagnostic result. It is the sole responsibility of the user to use and validate the kit in conjunction with a downstream *in vitro* diagnostic assay.

2.2 Product use limitations

The **NucleoSpin® Dx Blood** kit is not for use with tissue or stool samples, cell-free body fluids such as plasma, serum, urine, or cerebrospinal fluid. The kit performance has not been evaluated with buffy coat, cultured or isolated cells, swabs, dried blood spots, and viral DNA. The kit is also not specified for the isolation and purification of bacterial, fungal, or parasite nucleic acids.

2.3 Quality control

In accordance with MACHEREY-NAGEL's Quality Management System, each lot of **NucleoSpin® Dx Blood** kit is tested against predetermined specifications to ensure consistent product quality.

2.4 Introduction and kit specifications

NucleoSpin® Dx Blood is based on well-established NucleoSpin® silica-membrane technology and provides an easy way to isolate genomic DNA from 200 μL of whole blood samples. Purified DNA is ready-to-use for downstream PCR amplification.

The NucleoSpin® Dx Blood procedure is based on a series of simple steps:

First, the blood samples are lysed in the presence of chaotropic salts and Proteinase K. The genomic DNA in the lysate is then bound to a **NucleoSpin® Dx Blood Column**. Subsequently, the membrane with bound nucleic acids is washed and finally highly pure genomic DNA is eluted.

Samples

The kit can be used with 200 µL fresh or frozen human whole blood treated with EDTA, citrate, or heparin, from common blood collection systems. Cryoprecipitates formed during thawing of frozen samples may clog the **NucleoSpin® Dx Blood Column**. If such precipitates are visible avoid aspirating them when loading the lysate to the binding column.

Typically, 200 μ L human whole blood will yield 3–5 μ g genomic DNA, depending on the white blood cell count of the sample.

A selection of suitable blood collection devices is shown below:

Table 1: Selection of suitable blood collection systems		
Blood collecting system	Manufacturer	
S-Monovette® Li-Heparin	Sarstedt	
S-Monovette® EDTA	Sarstedt	
S-Monovette® Citrat	Sarstedt	
VACUETTE® EDTA	GREINER BIO-ONE	
BD VACUTAINER® K2E	BD	
K3 EDTA	DELTA LAB	
K2 EDTA	APTACA	

Table 2: Kit specifications at a glance		
Parameter	NucleoSpin® Dx Blood	
Sample material Fresh and frozen human whole blood treated EDTA, citrate, or heparin, from common be collection systems		
Sample volume	200 μL	
Typical DNA yield	3–5 µg depending on white blood cell count	
Typical DNA quality	Ratio A ₂₆₀ /A ₂₈₀ 1.7–1.9 Ratio A ₂₆₀ /A ₂₃₀ 1.8–2.3	
Elution volume	50–200 μL	
Typical DNA concentration	40–60 ng/μL	
Processing	Centrifugation	

2.5 Elution procedures

DNA is eluted from the <code>NucleoSpin®</code> Dx Blood Columns with 50 to 200 μ L Elution Buffer BE. Overall DNA yield increases with increasing elution volume, whereas the DNA concentration decreases (see Figure 1).

Typically, up to 10 μ L of the eluate can be used as template in a 50 μ L PCR mix without affecting PCR performance. It is recommended storing eluted DNA at -20 °C. Several freeze-thaw cycles will not interfere with most downstream applications.

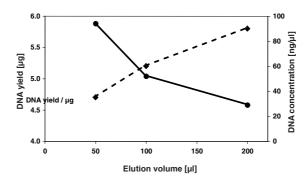


Figure 1: Impact of elution volume on overall DNA yield (dashed line) and concentration (solid line). Elution was performed with 50, 100, and 200 μL.

3 Storage conditions and preparation of working solutions

Attention:

- Check all components for damages after receiving the kit. If kit contents, like buffer bottles or blister packages are damaged, contact MACHEREY-NAGEL technical support and customer service, or your local distributor.
- · Do not use damaged kit components.
- Upon arrival the NucleoSpin® Dx Blood kit should be stored at room temperature (18–25°C). It is NOT required to open the kit on delivery and remove individual components for separate storage.
- NucleoSpin® Dx Blood Columns can be used until the expiration date specified on the kit box.

Before starting the NucleoSpin® Dx Blood protocol prepare the following:

- Wash Buffer B5: Add the indicated volume of ethanol (96–100 %, see table below or on the bottle) to Wash Buffer B5 Concentrate. Mark the label of the bottle to indicate that ethanol was added. Store Wash Buffer B5 at room temperature (18–25 °C) until the expiration date.
- Lyophilized Proteinase K can be stored at room temperature (18–25 °C) until
 the expiration date without decrease in performance. Before first use of the
 kit, add the indicated volume of Proteinase Buffer PB to dissolve lyophilized
 Proteinase K. Reconstituted Proteinase K should be stored at -20 °C for up to 6
 months, but only until the expiration date.
- During storage, especially at low temperatures, a white precipitate may form in Buffer B3 and Buffer BW. Such precipitates can be easily dissolved by incubating the bottle at 70 °C for 5 min before use.

	NucleoSpin [®] Dx Blood			
	50 preps 250 preps			
REF	740899.50	740899.250		
Wash Buffer B5 (Concentrate)	7 mL Add 28 mL ethanol	2 x 20 mL Add 80 mL ethanol to each bottle		
Proteinase K	30 mg Add 1.35 mL Proteinase Buffer	2 x 75 mg Add 3.35 mL Proteinase Buffer to each vial		

4 Safety instructions – risk and safety phrases

The following components of the **NucleoBond**® kits contain hazardous contents. Wear gloves and goggles and follow the safety instructions given in this section.

4.1 Risk and safety phrases

Component	Hazard contents	Hazard symbol	Risk phrases	Safety phrases
Inhalt	Gefahrstoff	Gefahrstoff- symbol	R-Sätze	S-Sätze
B3	Guanidine hydrochloride Guanidinhydrochlorid	X Xn*	22-36	26-39
BW	Guanidine hydrochloride Guanidinhydrochlorid	X Xn*	10-22-36	16-26-39
Proteinase K	Proteinase K, lyophilized Proteinase K, lyophilisiert	X Xn	36/37/38-42	22-26- 36/37

Risk phrases

R 10	Flammable.
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Entzündlich.

R 22 Harmful if swallowed.

Gesundheitsschädlich beim Verschlucken.

R 36 Irritating to eyes. *Reizt die Augen.*

R 36/37/38 Irritating to eyes, respiratory system, and skin.

Reizt die Augen, Atmungsorgane und die Haut.

R 42 May cause sensitization by inhalation

Sensibilisierung durch Einatmen möglich.

Safety phrases

S 16 Keep, away from source of ignition – No smoking!

Von Zündquellen fernhalten.

S 22 Do not breathe dust.

Staub nicht einatmen.

S 26 In case of contact with eyes, rinse immediately with plenty of water and seek

medical advice.

Bei Berührung mit den Augen gründlich mit Wasser abspülen und Arzt konsultieren.

^{*} Hazard labeling not necessary if quantity per bottle below 125 g or mL (certificate of exemption according to 67/548/EEC Art. 25, 1999/45/EC Art. 12 and German GefStoffV § 20 (3) and TRGS 200 7.1). For further information see Material Safety Data Sheet.

S 36/37 Wear suitable protective clothing and gloves.

Bei der Arbeit geeignete Schutzhandschuhe und Schutzkleidung tragen.

S 39 Wear eye / face protection.

Schutzbrille/Gesichtsschutz tragen.

4.2 GHS classification

Only harmful features need not be labeled with H and P phrases until 125 mL or 125 g. Mindergefährliche Eigenschaften müssen bis 125 mL oder 125 g nicht mit H- und P-Sätzen gekennzeichnet werden.

Component	Hazard contents	GHS symb	ool	Hazard phrases	Precaution phrases
Inhalt	Gefahrstoff	GHS Symbo	ol	H-Sätze	P-Sätze
B3	Guanidine hydrochloride 36–50 % Guanidinhydrochlorid 36–50 %	<₩	Warning Achtung	302-319	280-301+312- 305+351+338- 330-337+313
BW	Guanidine hydrochloride 36–50 % + isopropanol 20–50 % Guanidinhydrochlorid 36–50 % + Isopropanol 20–50 %	(A) (I)	Warning Achtung	226-302- 319	210-233-280- 301+312- 305+351+338- 330-337+313- 403+235
Proteinase K	Proteinase K, lyophilized Proteinase K, lyophilisiert	/ 4 \ / 4 \	Danger <i>Gefahr</i>	334	261-304+341- 342+311

Hazard phrases

H 226	Flammable liquid and vapour. Flüssigkeit und Dampf entzündbar.
H 302	Harmful if swallowed. Gesundheitsschädlich bei Verschlucken.
H 319	Causes serious eye irritations. Verursacht schwere Augenreizung.
H 334	May cause allergy or asthma symptoms or breathing difficulties if inhaled. Kann bei Einatmen Allergie, asthmaartige Symptome oder Atembeschwerden verursachen.

Precaution phrases

Keep away from heat/sparks/open flames/hot surfaces. – No smoking. Von Hitze/Funken/offener Flamme/heißen Oberflächen fernhalten. – Nicht rauchen.
Keep container tightly closed. Behälter dicht verschlossen halten.
Avoid breathing dust. Einatmen von Staub vermeiden.
Wear protective gloves/eye protection. Schutzhandschuhe/Augenschutz tragen.

P 301+312	IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell. Bei Verschlucken: Bei Unwohlsein Giftinformationszentrum oder Arzt anrufen.
P 304+341	IF INHALED: If breathing is difficult, remove to fresh air and keep at rest in a position comfortable for breathing. Bei Einatmen: Bei Atembeschwerden an die frische Luft bringen und in einer Position ruhigstellen, die das Atmen erleichtert.
P 305+351+338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. Bei Kontakt mit den Augen: Einige Minuten lang behutsam mit Wasser spülen. Vorhandene Kontaktlinsen nach Möglichkeit entfernen. Weiter spülen.
P 330	Rinse mouth. Mund ausspülen.
P 337+313	lf eye irritation persists: Get medical advice / attention. Bei anhaltender Augenreizung: Ärztlichen Rat einholen / ärztliche Hilfe hinzuziehen.
P 342+311	If experiencing respiratory symptoms: Call a POISON CENTER or doctor / physician. Bei Symptomen der Atemwege: Giftinformationszentrum oder Arzt anrufen.
P 403+235	Store in a well ventilated place. Keep cool. An einem gut belüfteten Ort lagern. Kühl halten.

For further information please see Material Safety Data Sheets (www.mn-net.com). Weiterführende Informationen finden Sie in den Sicherheitsdatenblättern (www.mn-net.com).

When working with the **NucleoSpin® Dx Blood** kit wear suitable protective clothing (e.g., lab coat, disposable gloves, and protective goggles). For more information consult the appropriate Material Safety Data Sheets (MSDS available online at *http://www.mn-net.com/msds*).

Caution: Guanidine hydrochloride in Lysis Buffer B3 and Wash Buffer BW can form highly reactive compounds when combined with bleach! Thus, do not add bleach or acidic solutions directly to the sample preparation waste.

The waste generated with the **NucleoSpin® Dx Blood** kit has not been tested for residual infectious material. A contamination of the liquid waste with residual infectious material is highly unlikely due to strong denaturing lysis buffer and Proteinase K treatment but it cannot be excluded completely. Therefore, liquid waste must be considered infectious and should be handled and discarded according to local safety regulations.

5 Genomic DNA purification with NucleoSpin® Dx Blood

The procedure below provides instructions for processing a single blood sample. However, several samples can be processed at the same time; the number depends on the capacity of the microcentrifuge used.

Before starting the preparation:

- · Check if Buffer B5 and Proteinase K were prepared according to section 3.
- Check that 96–100% ethanol (denatured or non-denatured) is available to adjust DNA binding conditions.
- Set an incubator (e.g., heating block) or water bath to 70 °C.
- Equilibrate the blood samples to room temperature (18–25 °C). Make sure that the samples are mixed well.
- If a precipitate has formed in Lysis Buffer B3 or Buffer BW, incubate the buffer at 70 °C until the precipitate is dissolved.
- Generally, do not mix reagents and columns from different kits and lots.
- Equilibrate Elution Buffer BE to room temperature (18–25 °C).
- Do not add Proteinase K solution directly to Lysis Buffer B3. Proteinase K has
 to be mixed with the blood sample before addition of Buffer B3.
- All centrifugation steps should be carried out at room temperature (18–25 °C).

5.1 Protocol-at-a-glance

Supplemental protocol-overview:

Carefully read the detailed protocol (section 5.2) before starting the procedure.

Lyse blood	1	25 μL Proteinase K	
samples	2	200 μL blood	
	3	200 μL B3, mix	
	4	RT, 5 min	٧
	5	7 °C, 10 min, mix	
	6	2,000 x g, 1 s	
Adjust DNA binding conditions	7	210 μL ethanol, mix	9
			Ū
	8	2,000 x g, 1 s	
Bind DNA	9	Load lysate	
	10	11,000 x g, 1 min	
	11	Transfer the NucleoSpin® Dx Blood Column to a new Collection Tube	
Wash silica membrane	12	500 μL BW	
	13	11,000 x g, 1 min	3
	14	Transfer the NucleoSpin® Dx Blood Column to a new Collection Tube	
	15	600 μL B5	
	16	11,000 x g, 1 min	
Dry silica membrane	17	Transfer the NucleoSpin® Dx Blood Column to a new Collection Tube	Ò
	18	11,000 x g, 1 min	
Elute DNA	19	Transfer the NucleoSpin® Dx Blood Column to an Elution Tube	
	20	50 – 200 μL BE	
	21	11,000 x g, 1 min	

5.2 Procedure

- 1. Pipette **25 µL Proteinase K** solution into a Lysis Tube (1.5 mL, provided).
- 2. Add 200 µL blood sample into the Lysis Tube. Mix.
- 3. Add 200 µL Buffer B3 to the Lysis Tube, close the lid, and mix by pulse-vortexing vigorously for 10 s.

Do not premix Buffer B3 and Proteinase K!

- 4. Incubate at **room temperature** (18–25 °C) for **5 min** (±1 min).
- Incubate the Lysis Tube at 70 °C (±2 °C) for 10 min (±1 min). After incubation mix by pulse-vortexing vigorously for 5 s.
- 6. **Briefly centrifuge** Lysis Tube (approx. 1 s at 2,000 x *g*) to remove drops from the lid (short spin only).
- 7. Add **210 \muL ethanol** (96–100 %) to the sample. Close the lid and mix by pulse-vortexing for 5 s.

Make sure that the ethanol and the lysate is mixed well.

- 8. **Briefly centrifuge** Lysis Tube (approx. 1 s at 2,000 x *g*) to remove drops from the lid (short spin only).
- Carefully load the entire lysate to the NucleoSpin® Dx Blood Column placed in a Collection Tube and close the lid.
- 10. Centrifuge 1 min at 11,000 x g.

If the lysate is not completely drawn through the membrane, repeat the centrifugation at higher g-force (15,000–20,800 x g for 1 min). If the lysate still does not pass the membrane completely, discard the sample and repeat the isolation with new sample material.

- Place the NucleoSpin® Dx Blood Column into a new Collection Tube (2 mL, provided) and discard the Collection Tube with flow-through from the previous step.
- Open the NucleoSpin[®] Dx Blood Column and add 500 μL Buffer BW to the column. Close the lid.

Note: Make sure that residual lysate is washed away with Buffer BW.

- 13. Centrifuge 1 min at 11,000 x g.
- Place the NucleoSpin® Dx Blood Column into a new Collection Tube (2 mL, provided) and discard the Collection Tube with flow-through from the previous step.

 Open the NucleoSpin® Dx Blood Column and add 600 μL Buffer B5 to the column. Close the lid.

Note: Make sure that residual wash buffer from the previous step is washed away with Buffer B5.

- 16. Centrifuge 1 min at 11,000 x g.
- Place the NucleoSpin® Dx Blood Column into a new Collection Tube (2 mL, provided) and discard the Collection Tube with flow-through from the previous step.
- 18. **Centrifuge 1 min** at **11,000 x** *g*.

 Residual ethanol is removed during this step.
- 19. Place the **NucleoSpin® Dx Blood Column** in a clean Elution Tube (1.5 mL, provided) and discard the Collection Tube from the previous step.
- Open the NucleoSpin® Dx Blood Column and add 50–200 μL Buffer BE directly onto the center of the membrane.
- 21. Centrifuge 1 min at 11,000 x g to elute the DNA from the column.

6 Appendix

6.1 Troubleshooting

Problem Possible cause and suggestions

Low concentration of white blood cells in sample

 The DNA yield depends on the number of white blood cells per sample. Blood samples with low white blood cell count yield low DNA amounts.

Incomplete sample lysis

- Inhomogeneous blood sample or blood clots within the sample:
 Make sure that blood samples are collected following the
 instructions of the manufacturer of the blood collection tube.
 Make sure that only blood which can be easily transferred by
 pipetting is used as sample material. If necessary, homogenize
 the blood sample before use.
- Sample not thoroughly mixed with Proteinase K and lysis buffer.
 The mixture has to be vortexed vigorously immediately after addition of Lysis Buffer B3.

No or poor DNA yield

 Proteinase K digestion not optimal. Never add Proteinase K directly to Lysis Buffer B3.

Reagents not applied properly

 Prepare buffers and Proteinase K solution according to instructions (section 3). Add ethanol to lysate before loading lysate on the column.

Unappropriate centrifugation

Do not extend centrifugation time and speed in step 6 and 8.
 Only use a short spin to remove droplets from the lid.

Suboptimal elution of DNA from the column

 Elution efficiency depends on elution buffer volume. For highest elution efficiency use 200 µL elution buffer; for highest DNA concentration use 50 µL elution buffer.

Problem

Possible cause and suggestions

Inhomogeneous blood sample

Clogged DNA binding column

Cryoprecipitate formed during thawing of frozen samples may clog the NucleoSpin® Dx Blood Column. If such precipitates are visible avoid aspirating them when loading the lysate to the binding column. Precipitates may also form in fresh blood samples. Make sure that the samples are mixed well. If the column clogs during the DNA binding step repeat the centrifugation at higher g-force (15,000–20,800 x g for 1 min).

Reagents not applied properly

 Prepare buffers and Proteinase K solution according to instructions (section 3). Add ethanol to lysate and mix before loading them on columns.

Incomplete sample lysis

Poor DNA quality

- Sample not thoroughly mixed with Proteinase K solution and lysis buffer. The mixture has to be vortexed vigorously immediately after addition of lysis buffer.
- Proteinase K digestion not optimal. Do not add Proteinase K directly to Lysis Buffer B3.

Old or clotted blood samples processed

 Make sure that only blood is used as sample material which can be easily transferred by pipetting. If necessary homogenize the blood sample before use.

Carry-over of ethanol

Suboptimal performance of genomic DNA in enzymatic reactions

- Be sure to remove all of ethanolic Buffer B5 before eluting the DNA. If the filling level of Wash Buffer B5 flow-through after the second wash reaches the column outlet for any reason, discard flow-through, place the column back into the Collection Tube, and centrifuge again.
- DNA eluates may contain traces of ethanol. However, no decrease in PCR performance was observed using DNA eluate of up to 20% of the PCR final volume as template (e.g., using 4 µL from 100 µL eluate as template in a 20 µL PCR). The maximum percentage of template volume in a PCR may vary depending on the robustness of the PCR system and has to be determined by the user.

Problem	Possible cause and suggestions
Suboptimal performance of genomic DNA in enzymatic reactions (continued)	Contamination of DNA with inhibitory substances If preparing DNA from old or clotted blood samples, make sure that only blood is used as sample material which can be easily transferred by pipetting. If necessary, homogenize the blood sample before use.

6.2 Ordering information

Product	REF	Pack of
NucleoSpin® Dx Blood	740899.50/.250	50/250

Visit www.mn-net.com for more detailed product information.

6.3 Product use restriction/warranty

The **NucleoSpin® Dx Blood** Kit is a generic system for the isolation and purification of genomic DNA from human whole blood samples for subsequent *in vitro* diagnostic purposes.

The kit is designed to be used with downstream applications employing enzymatic amplification and detection of DNA (e.g., PCR). Any and all diagnostic results generated using the DNA isolated with the NucleoSpin® Dx Blood kit in conjunction with a diagnostic assay should be interpreted with regard to additional clinical or laboratory findings. The NucleoSpin® Dx Blood Kit does not provide a diagnostic result. It is the sole responsibility of the user to use and validate the kit in conjunction with a downstream in vitro diagnostic assay. ONLY MACHEREY-NAGEL products specially labeled as IVD are suitable for *In vitro*-diagnostic use.

NucleoSpin® Dx Blood kit is intended for use by professional users such as technicians and physicians experienced in and trained in molecular biological techniques including experience with whole blood samples and DNA isolation. For safety instructions please refer to the respective chapter in the user manual. NucleoSpin® Dx Blood kit shall exclusively be used in an adequate test environment, i.e. a suitable laboratory setting.

The respective user is liable for any and all damages resulting from application of the NucleoSpin® Dx Blood kit for use deviating from the intended use as specified in the user manual.

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