

Genomic DNA from Blood

User Manual

NucleoSpin® Blood

NucleoSpin® Blood L

NucleoSpin® Blood XL

NucleoSpin® Blood QuickPure

January 2011 / Rev. 11

Genomic DNA Purification from Blood

Protocol-at-a-glance (Rev. 11)







































	Mini	Midi	Maxi	Mini
	NucleoSpin® Blood	NucleoSpin® Blood L	NucleoSpin® Blood XL	NucleoSpin® Blood QuickPure
1 Lyse blood samples	 200 µL blood 25 µL Pro.K 200 µL B3 Mix  70 °C 10–15 min	 2 mL blood 150 µL Pro.K 2 mL BQ1 Mix  56 °C 10–15 min	 10 mL blood 500 µL Pro.K 10 mL BQ1 Mix  56 °C 10–15 min	 200 µL blood 25 µL Pro.K 200 µL BQ1 Mix  70 °C 10–15 min
2 Adjust DNA binding conditions	210 µL ethanol	2 mL ethanol	10 mL ethanol	200 µL ethanol
3 Bind DNA	 Load all  11,000 x g 1 min	 Load 3 mL  4,500 x g 3 min	 Load 15 mL  4,000 x g 3 min	 Load all  11,000 x g 1 min
	—	Load 3 mL of residue  4,500 x g 5 min	Load 15 mL of residue  4,000 x g 3 min	—
4 Wash silica membrane	 500 µL BW 600 µL B5	 2 mL BQ2 2 mL BQ2	 7.5 mL BQ2 7.5 mL BQ2	 350 µL BQ2
1st wash	 11,000 x g 1 min	 4,500 x g 2 min	 4,000 x g 2 min	 11,000 x g 3 min
2nd wash	 11,000 x g 1 min	 4,500 x g 10 min	 4,000 x g 10 min	—
5 Dry silica membrane	 11,000 x g 1 min	Drying is performed during centrifugation of the last washing step	Drying is performed during centrifugation of the last washing step	Drying is performed during centrifugation of the last washing step
6 Elute highly pure DNA	 100 µL BE (70 °C) RT 1 min  11,000 x g 1 min	 200 µL BE (70 °C) RT 2 min  4,500 x g 2 min	 500 µL - 2000 µL BE (70 °C) RT 2 min  4,000 x g 2 min	 50 µL BE (70 °C) RT 1 min  11,000 x g 1 min

Table of contents

1	Components	4
1.1	Kit contents	4
1.2	Reagents, consumables, and equipment to be supplied by user	8
1.3	About this User Manual	8
2	Product description	9
2.1	The basic principle	9
2.2	Kit specifications	9
2.3	Storage of blood samples	10
2.4	Elution procedures	10
3	Storage conditions and preparation of working solutions	12
4	Safety instructions – risk and safety phrases	14
5	Protocols for DNA purification from whole blood	15
5.1	Genomic DNA purification with NucleoSpin® Blood	15
5.2	Genomic DNA purification with NucleoSpin® Blood L	17
5.3	Genomic DNA purification with NucleoSpin® Blood XL	20
5.4	Genomic DNA purification with NucleoSpin® Blood QuickPure	23
6	Appendix	25
6.1	Troubleshooting	25
6.2	Ordering information	28
6.3	Reference	28
6.4	Product use restriction/warranty	29

1 Components

1.1 Kit contents

NucleoSpin® Blood			
REF	10 preps 740951.10	50 preps 740951.50	250 preps 740951.250
Buffer B1*	3.2 mL	10 mL	50 mL
Buffer B2*	0.8 mL	2.5 mL	12.5 mL
Wash Buffer BW	6 mL	30 mL	2 x 75 mL
Wash Buffer B5 (Concentrate)*	4 mL	7 mL	2 x 20 mL
Elution Buffer BE**	4 mL	13 mL	60 mL
Proteinase K (lyophilized)*	6 mg	30 mg	2 x 75 mg
Proteinase Buffer PB	0.8 mL	1.8 mL	8 mL
NucleoSpin® Blood Columns (red rings - plus Collection Tubes)	10	50	250
Collection Tubes (2 mL)	20	100	500
Labels for Lysis Buffer B3	1	1	1
User Manual	1	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

1.1 Kit contents *continued*

NucleoSpin® Blood L	
REF	20 preps 740954.20
Lysis Buffer BQ1	45 mL
Wash Buffer BQ2 (Concentrate)*	20 mL
Elution Buffer BE**	10 mL
Proteinase K (lyophilized)*	60 mg
Proteinase Buffer PB	3.6 mL
NucleoSpin® Blood L Columns (plus Collection Tubes)	20
Collection Tubes (15 mL)	20
User Manual	1

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

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

1.1 Kit contents *continued*

NucleoSpin® Blood XL		
REF	10 preps 740950.10	50 preps 740950.50
Lysis Buffer BQ1	125 mL	3 x 200 mL
Wash Buffer BQ2 (Concentrate)*	50 mL	4 x 50 mL
Elution Buffer BE**	25 mL	75 mL
Proteinase K (lyophilized)*	126 mg	5 x 126 mg
Proteinase Buffer PB	8 mL	35 mL
NucleoSpin® Blood XL Columns (plus Collection Tubes)	10	50
Collection Tubes (50 mL)	10	50
User Manual	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

1.1 Kit contents *continued*

NucleoSpin® Blood QuickPure			
REF	10 preps 740569.10	50 preps 740569.50	250 preps 740569.250
Lysis Buffer BQ1	4 mL	12.5 mL	62.5 mL
Wash Buffer BQ2 (Concentrate)*	3 mL	7 mL	2 x 20 mL
Elution Buffer BE**	4 mL	13 mL	60 mL
Proteinase K (lyophilized)*	6 mg	30 mg	2 x 75 mg
Proteinase Buffer PB	0.8 mL	1.8 mL	8 mL
NucleoSpin® Blood QuickPure Columns (dark red rings - plus Collection Tubes)	10	50	250
Collection Tubes (2 mL)	10	50	250
User Manual	1	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

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

Reagents

- 96–100 % ethanol
- Phosphate-buffered saline (PBS) may be required for some samples

Consumables

- 1.5 mL microcentrifuge tubes (NucleoSpin® Blood/ QuickPure), 15 mL (NucleoSpin® Blood L), or 50 mL centrifuge tubes (NucleoSpin® Blood XL), for sample lysis and DNA elution
- Disposable pipette tips

Equipment

- Manual pipettors
- Centrifuge for microcentrifuge tubes (NucleoSpin® Blood/QuickPure), centrifuge for 15 mL (NucleoSpin® Blood L) or 50 mL (NucleoSpin® Blood XL) centrifuge tubes, with a swing-bucket rotor
- Vortex mixer
- Thermal heating block (NucleoSpin® Blood/QuickPure) or water bath (NucleoSpin® Blood L/XL)
- Personal protection equipment (lab coat, gloves, goggles)

1.3 About this User Manual

It is strongly recommended reading the detailed protocol sections of this User Manual if the **NucleoSpin® Blood** kit is used for the first time. 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.

2 Product description

2.1 The basic principle

With the **NucleoSpin® Blood** method, genomic DNA is prepared from whole blood, cultured cells, serum, plasma, or other body fluids. Lysis is achieved by incubation of whole blood in a solution containing large amounts of chaotropic ions in the presence of Proteinase K. Appropriate conditions for binding of DNA to the silica membrane of the corresponding **NucleoSpin® Blood Columns** are achieved by addition of ethanol to the lysate. The binding process is reversible and specific to nucleic acids. Washing steps efficiently remove contaminations. With the **NucleoSpin® Blood QuickPure** kit, contaminations are removed by a single wash step. Pure genomic DNA is finally eluted under low ionic strength conditions in a slightly alkaline elution buffer.

2.2 Kit specifications

- **NucleoSpin® Blood** kits are designed for the rapid isolation of highly pure genomic DNA from whole blood, serum, plasma, or other body fluids. It is also possible to purify viral DNA (e.g., HBV) from blood samples. As viral DNA co-purifies with cellular DNA, we recommend using cell-free samples (serum or plasma) to prepare pure viral DNA.
- The **NucleoSpin® Blood QuickPure** kit is designed for ultra-fast small-scale purification of highly pure genomic DNA from whole blood, serum, plasma, or other body fluids. The number of washing and drying steps is reduced from 3 to 1! Therefore, the hands-on time is less than 10 min.
- DNA can be purified successfully from blood samples treated with EDTA, citrate, or heparin. If leukocyte rich materials like buffy coat are used, apply smaller volumes and dilute the samples with sterile PBS (dissolve 8 g NaCl, 0.2 g KCl, 1.44 g Na₂HPO₄, and 0.24 g KH₂PO₄ in 800 mL H₂O. Adjust pH to 7.4 with HCl. Add H₂O to 1 liter).
- The kits allow purification of highly pure genomic DNA with an A₂₆₀/A₂₈₀ ratio between 1.60 and 1.90 and a typical concentration of 40–60 ng per µL for the **NucleoSpin® Blood** kit, 80–120 ng per µL for the **NucleoSpin® Blood QuickPure** kit and 200–300 ng per µL for the **NucleoSpin® Blood L/XL** kits.
- The obtained DNA is ready-to-use for subsequent reactions like PCR, Southern blotting, or any kind of enzymatic reactions.

Table 1: Kit specifications at a glance

Parameter	Blood	Blood L	Blood XL	Blood QuickPure
Sample material	Up to 200 µL / 5 x 10 ⁶ cells	Up to 2 mL / 2 x 10 ⁷ cells	Up to 10 mL / 1 x 10 ⁸ cells	Up to 200 µL / 5 x 10 ⁶ cells
Typical yield	4–6 µg	40–60 µg	200–300 µg	4–6 µg
Elution volume	100 µL	200 µL	500–2000 µL	50 µL
Binding capacity	60 µg	250 µg	700 µg	50 µg
Preparation time	30 min /prep	1 h /prep	1 h /prep	< 10 min /prep
Format	Mini spin column	Midi spin column	Maxi spin column	Mini spin column

2.3 Storage of blood samples

For the isolation of genomic DNA from blood treated with anticoagulants (heparin, citrate, or EDTA) using a **NucleoSpin® Blood** kit the blood samples can be stored at room temperature, +4 °C, or frozen.

Blood samples stored at room temperature or +4 °C for up to several days or weeks, respectively, will still allow DNA isolation. However, DNA yield and quality will slowly decrease due to prolonged storage of blood samples under these conditions.

Blood stored frozen for years is well suited for DNA isolation.

Highest yields and quality of DNA are obtained from fresh blood.

2.4 Elution procedures

It is possible to adapt elution method and volume of elution buffer to the subsequent application of interest. In addition to the standard method (recovery rate about 70–90%) there are several modifications possible. Use elution buffer preheated to 70 °C for one of the following procedures:

- **High yield:** Perform two elution steps with the volume indicated in the individual protocol. About 90–100 % of bound nucleic acid can be eluted.
- **High concentration:** Perform one elution step with 60 % of the volume indicated in the individual protocol. Concentration of DNA will be higher than with standard elution (**NucleoSpin® Blood**: ca. 130 %; **NucleoSpin® Blood QuickPure**: ca. 150 %; **NucleoSpin® Blood L**: ca. 140 %; **NucleoSpin® Blood XL**: ca. 115 %). Maximal yield of bound nucleic acid is about 80 %.

- **High yield and high concentration:** Apply half the volume of elution buffer as indicated in the individual protocol, incubate for 3 min and centrifuge. Apply a second aliquot of elution buffer, incubate and centrifuge again. Thus, about 85–100 % of bound nucleic acid is eluted in the standard elution volume at a high concentration.
- **Convenient elution:** For convenience, elution buffer of ambient temperature may be used. This will result in a lower yield (approximately 20 %) compared to elution with preheated elution buffer.

Elution may also be performed with Tris-EDTA-buffer (TE) of pH equal or higher than 8. This will increase DNA stability especially during long term and/or multi use storage at 4 °C or ambient temperature by inhibition of omnipresent DNases. However, EDTA interferes, depending on the final concentration, with certain downstream applications.

For optimal performance of isolated DNA in subsequent downstream applications we recommend elution with the supplied elution buffer and storage, especially long term, at -20 °C. Several freeze-thaw cycles will not interfere with most downstream applications.

Performance of long-range PCR (e.g., > 10 kbp) or detection sensitivity of trace amount of DNA species might be reduced after multiple freeze-thaw cycles or prolonged storage of eluted DNA at +4 °C or room temperature due to DNA shearing or adsorption to surfaces.

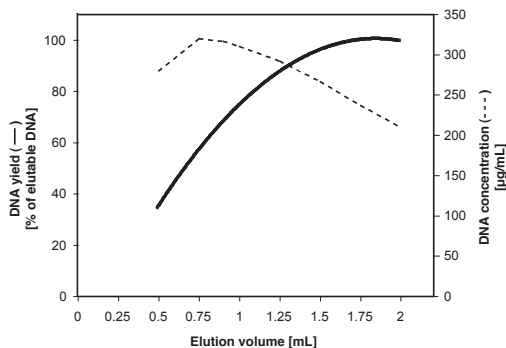


Figure 1: Dependence of DNA yield (solid line) and concentration (dashed line) on elution volume.

Genomic DNA was purified from 10 mL whole blood and eluted using different elution volumes as indicated. Highest DNA yield is obtained with 1.5-2.0 mL elution volume. Highest DNA concentration is obtained with approximately 0.75 mL elution volume. Furthermore, yield and concentration may vary as they depend on the kind of sample (blood, serum, plasma), type of blood sample (human or animal), and quality of the samples (fresh, old, frozen, clotted, etc.).

3 Storage conditions and preparation of working solutions

Attention:

Buffers B1, BQ1, and BW contain guanidine hydrochloride. Wear gloves and goggles!

All kit components can be stored at room temperature (18–25 °C) and are stable up to one year.

- During storage, especially at low temperatures, a white precipitate may form in Buffer T1, B1, BQ1, or B3. Such precipitates can be easily dissolved by incubating the bottle at 70 °C before use.

Before starting any **NucleoSpin® Blood** protocol prepare the following:

- **Lysis Buffer B3** (NucleoSpin® Blood): Transfer the total content of **Buffer B1** to **Buffer B2** and mix well. Place the labels for Lysis Buffer B3 on the bottle. The resulting Lysis Buffer B3 is stable for up to one year at room temperature.
- **Wash Buffer B5** (NucleoSpin® Blood): Add the indicated volume of ethanol (96–100 %) 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) for up to one year.

Wash Buffer BQ2 (NucleoSpin® Blood L / XL / QuickPure): Add the indicated volume of ethanol (96–100 %) to **Wash Buffer BQ2 Concentrate**. Mark the label of the bottle to indicate that ethanol was added. Store Wash Buffer BQ2 at room temperature (18–25 °C) for up to one year.

- **Proteinase K**: Add the indicated volume of Proteinase Buffer PB to dissolve lyophilized **Proteinase K**. Proteinase K solution is stable at -20 °C for up to 6 months.

NucleoSpin® Blood			
REF	10 preps 740951.10	50 preps 740951.50	250 preps 740951.250
Wash Buffer B5 (Concentrate)	4 mL Add 16 mL ethanol	7 mL Add 28 mL ethanol	2 x 20 mL Add 80 mL ethanol to each bottle
Proteinase K	6 mg Add 260 µL Proteinase Buffer	30 mg Add 1.35 mL Proteinase Buffer	2 x 75 mg Add 3.35 mL Proteinase Buffer to each vial

NucleoSpin® Blood L

	20 preps 740954.20
REF	
Wash Buffer BQ2 (Concentrate)	20 mL Add 80 mL ethanol
Proteinase K	60 mg Add 3.15 mL Proteinase Buffer

NucleoSpin® Blood XL

	10 preps 740950.10	50 preps 740950.50
REF		
Wash Buffer BQ2 (Concentrate)	50 mL Add 200 mL ethanol	4 x 50 mL Add 200 mL ethanol to each bottle
Proteinase K	126 mg Add 5.75 mL Proteinase Buffer	5 x 126 mg Add 5.75 mL Proteinase Buffer to each vial

NucleoSpin® Blood QuickPure

	10 preps 740569.10	50 preps 740569.50	250 preps 740569.250
REF			
Wash Buffer BQ2 (Concentrate)	3 mL Add 12 mL ethanol	7 mL Add 28 mL ethanol	2 x 20 mL Add 80 mL ethanol to each bottle
Proteinase K	6 mg Add 260 µL Proteinase Buffer	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 **NucleoSpin® Blood** kits contain hazardous contents.

Wear gloves and goggles and follow the safety instructions given in this section.

Component	Hazard contents	Hazard symbol		Risk phrases	Safety phrases
B1	Guanidine hydrochloride	✘ Xn*	Harmful if swallowed - Irritating to eyes and skin	R 22-36/38	
BQ1	Guanidine hydrochloride	✘ Xn*	Harmful if swallowed - Irritating to eyes and skin	R 22-36/38	
BW	Guanidine hydrochloride + isopropanol <25%	✘ Xn*	Flammable - Harmful if swallowed - Irritating to eyes and skin.	R 10-22-36/38	S 16-25
Proteinase K	Proteinase K, lyophilized	✘ Xn Xi**	Irritating to eyes, respiratory system and skin, may cause sensitization by inhalation	R 36/37/38-42	S 22-24-26-36/37

Risk phrases

R 10	Flammable
R 22	Harmful if swallowed
R 36/37/38	Irritating to eyes, respiratory system and skin
R 36/38	Irritating to eyes and skin
R 42	May cause sensitization by inhalation

Safety phrases

S 16	Keep away from sources of ignition - No smoking!
S 22	Do not breathe dust
S 24	Avoid contact with the skin
S 25	Avoid contact with the eyes
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

* 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.

**Hazard labeling not necessary if quantity per bottle below 25 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.

5 Protocols for DNA purification from whole blood

5.1 Genomic DNA purification with NucleoSpin® Blood

Before starting the preparation:

- Check if Buffer B3, Buffer B5, and Proteinase K were prepared according to section 3.
- Set an incubator or water bath to 70 °C.
- Preheat Elution Buffer BE to 70 °C.

1 Lyse blood sample

Pipette **25 µL Proteinase K** and up to **200 µL blood**, body fluid sample, or buffy coat from 1 mL blood (equilibrated to room temperature) into 1.5 mL microcentrifuge tubes (not provided).

For sample volumes less than 200 µL, add PBS to adjust the volume to 200 µL. If purifying DNA viruses, we recommend starting with 200 µL serum or plasma. If cultured cells are used, resuspend up to 5×10^6 cells in a final volume of 200 µL PBS.

Add **200 µL Buffer B3** to the samples and vortex the mixture vigorously (10–20 s).

Note: Vigorous mixing is important to obtain high yield and purity of DNA.

Incubate samples at **70 °C** for **10–15 min**.

The lysate should become brownish during incubation with Buffer B3. Increase incubation time with Proteinase K (up to 30 min) and vortex once or twice vigorously during incubation if processing older or clotted blood samples.



200 µL blood
+ 25 µL Proteinase K
+ 200 µL B3
Mix
70 °C
10–15 min

2 Adjust DNA binding conditions

Add **210 µL ethanol (96–100%)** to each sample and vortex again.



+ 210 µL ethanol
Mix

3 Bind DNA

For each preparation, take one **NucleoSpin® Blood Column** placed in a Collection Tube and load the sample. Centrifuge **1 min** at **11,000 x g**. If the samples are not drawn through the matrix completely, repeat the centrifugation at higher *g*-force (< 15,000 x *g*). Discard Collection Tube with flow-through.



Load lysate



**11,000 x g
1 min**

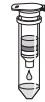
4 Wash silica membrane

1st wash

Place the NucleoSpin® Blood Column into a new Collection Tube (2 mL) and add **500 µL Buffer BW**. Centrifuge **1 min** at **11,000 x g**. Discard Collection Tube with flow-through.

+ 500 µL BW

**11,000 x g
1 min**



2nd wash

Place the NucleoSpin® Blood Column into a new Collection Tube (2 mL) and add **600 µL Buffer B5**. Centrifuge **1 min** at **11,000 x g**. Discard flow-through and reuse Collection Tube.

+ 600 µL B5

**11,000 x g
1 min**

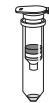


5 Dry silica membrane

Place the NucleoSpin® Blood Column back into the Collection Tube and centrifuge **1 min** at **11,000 x g**.

**11,000 x g
1 min**

Residual ethanol is removed during this step.



6 Elute highly pure DNA

Place the NucleoSpin® Blood Column in a 1.5 mL microcentrifuge tube (not provided) and add **100 µL preheated Buffer BE (70 °C)**. Dispense buffer directly onto the silica membrane. Incubate at **room temperature** for **1 min**. Centrifuge **1 min** at **11,000 x g**.

**+ 100 µL BE
(70 °C)**

**RT
1 min**



For alternative elution procedures see section 2.4.



**11,000 x g
1 min**

5.2 Genomic DNA purification with NucleoSpin® Blood L

Before starting the preparation:

- Check if Buffer BQ2 and Proteinase K were prepared according to section 3.
- Set an incubator or water bath to 56 °C.
- Preheat Elution Buffer BE to 70 °C.
- For centrifugation, a centrifuge with a **swing-out rotor** and appropriate buckets capable of reaching 4,000–4,500 x g is required.

1 Lyse blood sample

Pipette up to **2 mL blood** (or body fluid) sample (equilibrated to room temperature) and **150 µL Proteinase K** into a 15 mL tube (not provided).

If processing buffy coat, do not use more than 1 mL and add PBS to adjust the volume to 2 mL.

If cultured cells are used, resuspend up to 2×10^7 cells in a final volume of 2 mL PBS.

Add **2 mL Buffer BQ1** (if processing less than 2 mL blood, add one volume of Buffer BQ1) to the samples and vortex the mixture vigorously for 10 s.

Note: Vigorous mixing is important to obtain high yield and purity of DNA.

Incubate samples at **56 °C for 15 min**.

Let the samples cool down to room temperature before proceeding with addition of ethanol.

The lysate should become brownish during incubation with Buffer BQ1. Increase incubation time with Proteinase K (up to 20 min) and vortex once or twice during incubation if processing older or clotted blood samples.



2 mL blood
+ 150 µL Proteinase K
+ 2 mL BQ1
Mix
56 °C
15 min

2 Adjust DNA binding conditions

Add **2 mL ethanol (96–100 %)** (if processing less than 2 mL blood, add 1 volume of ethanol) to each sample and mix by inverting the tube 10 times.

**+ 2 mL
ethanol**

Mix

Note: High local ethanol concentration must be avoided by immediate mixing after addition.

Be sure that the lysate has cooled down to room temperature before loading it onto the column. Loading of hot lysate may lead to diminished yields.

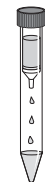
3 Bind DNA

For each preparation, take one **NucleoSpin® Blood L Column** placed in a Collection Tube and **load 3 mL of lysate**. Do not moisten the rims of the columns. Close the tubes with screw caps and centrifuge **3 min at 4,500 x g**.

Load 3 mL

4,500 x g
3 min

Usually the lysate will start to flow-through the columns even before centrifugation. This will not adversely affect DNA yield or purity. Keep NucleoSpin® Blood L Column in an upright position as liquid may pass through the ventilation slots on the rim of the column even if the caps are closed.



Load residue

4,500 x g
5 min

Load **all of the remaining lysate** in a second step to the respective NucleoSpin® Blood L Column, avoiding moistening the rim. Centrifuge **5 min at 4,500 x g**. Discard the flow-through and place the column back into the Collection Tube.



Remove the Collection Tube with the column carefully from the rotor to avoid that the flow-through comes in contact with the column outlet. Be sure to wipe off any spilled lysate from the Collection Tube before placing the column back.

4 Wash silica membrane

1st wash

Add **2 mL Buffer BQ2**. Centrifuge **2 min** at **4,500 x g**.

It is not necessary to discard the flow-through after the first washing step.

2nd wash

Add **2 mL Buffer BQ2**. Centrifuge **10 min** at **4,500 x g**. Remove the column carefully from the rotor in order to avoid that the flow-through comes in contact with the column outlet.

By prolonged centrifugation during this second washing step, residual ethanolic washing Buffer BQ2 is removed from the silica membrane of the NucleoSpin® Blood L Column.

+ 2 mL BQ2

4,500 x g
2 min



+ 2 mL BQ2

4,500 x g
10 min



5 Dry silica membrane

The drying of the NucleoSpin® Blood L Column is performed by prolonged centrifugation time (10 min) in the 2nd wash step.

6 Elute highly pure DNA

Insert the column into a new Collection Tube (15 mL) and apply **200 µL preheated Buffer BE (70 °C)** directly to the center of the silica membrane. Incubate at **room temperature** for **2 min**. Centrifuge at **4,500 x g** for **2 min**.

For alternative elution procedures see section 2.4.

+ 200 µL BE
(70 °C)

RT
1 min



4,500 x g
2 min



5.3 Genomic DNA purification with NucleoSpin® Blood XL

Before starting the preparation:

- Check if Buffer BQ2 and Proteinase K were prepared according to section 3.
- Set an incubator or water bath to 56 °C.
- Preheat Elution Buffer BE to 70 °C.
- For centrifugation, a centrifuge with a **swing-out rotor** and appropriate buckets capable of reaching 4,000–4,500 x g is required.

1 Lyse blood sample

Pipette up to **10 mL blood** (or body fluid) sample (equilibrated to room temperature) and **500 µL Proteinase K** into a 50 mL tube (not provided).

If you process ≤ 5 mL blood, sample loading with a single centrifugation step is possible (step 3).

If processing buffy coat, do not use more than 2 mL and add PBS to adjust the volume to 10 mL.

If cultured cells are used, resuspend up to 1×10^8 cells in a final volume of 10 mL PBS.

Add **10 mL Buffer BQ1** (if processing less than 10 mL blood, add one volume of Buffer BQ1) to the samples and vortex the mixture vigorously for 10 s.

Note: Vigorous mixing is important to obtain high yield and purity of DNA.

Incubate samples at **56 °C for 15 min**.

Let the lysate cool down to room temperature before proceeding with addition of ethanol.

The lysate should become brownish during incubation with Buffer BQ1. Increase incubation time with Proteinase K (up to 20 min) and vortex once or twice during incubation if processing older or clotted blood samples.



10 mL blood

**+ 500 µL
Proteinase K**

+ 10 mL BQ1

Mix

**56 °C
15 min**

2 Adjust DNA binding conditions

Add **10 mL ethanol (96–100%)** (if processing less than 10 mL blood, add one volume of ethanol) to each sample and mix by inverting the tube 10 times.

**+ 10 mL
ethanol**
Mix

Note: High local ethanol concentration must be avoided by immediate mixing after addition.

Be sure that the lysate has cooled down to room temperature (about 5 min) before loading it onto the columns. Loading of hot lysate may lead to diminished yields.

3 Bind DNA

For each preparation, take one **NucleoSpin® Blood XL Column** placed in a Collection Tube and load **15 mL of lysate**. Do not moisten the rim of the column. Close the tubes with screw caps and centrifuge **3 min** at **4,000 x g**. Discard flow-through.

Discarding the flow-through may be omitted. Be careful after the second loading step during removal of the tube from the centrifuge and the removal of the column from the tube: keep tube with column upright to avoid contact of flow-through with the column outlet.

Usually the lysate will start to flow through the column even before centrifugation. This will not adversely affect DNA yield or purity. Keep NucleoSpin® Blood XL Column in an upright position as liquid may pass through the ventilation slots on the rim of the column even if the caps are closed.

Load **15 mL of the remaining lysate** to the respective NucleoSpin® Blood XL Column. Again, avoid moistening the rim. Centrifuge **3 min** at **4,000 x g**. Discard the flow-through and place the column back into the Collection Tube.

Remove the Collection Tube with the column carefully from the rotor and avoid that the flow-through comes in contact with the column outlet.

If you process ≤ 5 mL blood no loading of remaining lysate is necessary.



Load 15 mL
4,000 x g
3 min

Load residue
4,000 x g
3 min

4 Wash silica membrane

1st wash

Add **7.5 mL Buffer BQ2** to the NucleoSpin® Blood XL Column. Centrifuge **2 min** at **4,000 x g**.

It is not necessary to discard the flow-through after the first washing step.

2nd wash

Add **7.5 mL Buffer BQ2**. Centrifuge **10 min** at **4,000 x g**. Remove the column carefully from the rotor to avoid that flow-through gets in contact with the column outlet.

By prolonged centrifugation during this second washing step, residual ethanolic Buffer BQ2 is removed from the silica membrane of the NucleoSpin® Blood XL Column.



+ 7.5 mL BQ2

4,000 x g
2 min



+ 7.5 mL BQ2

4,000 x g
10 min

5 Dry silica membrane

The drying of the NucleoSpin® Blood XL Column is performed by prolonged centrifugation time (10 min) in the 2nd wash step.

6 Elute highly pure DNA

Insert the column into a new Collection Tube (50 mL) and apply **1000 µL of preheated Buffer BE (70 °C)** directly to the center of the silica membrane. Incubate at **room temperature** for **2 min**. Centrifuge at **4,000 x g** for **2 min**.

For alternative elution procedures see section 2.4.



+ 1000 µL BE
(70 °C)

RT
2 min



4,000 x g
2 min

5.4 Genomic DNA purification with NucleoSpin® Blood QuickPure

Before starting the preparation:

- Check if Buffer BQ2 and Proteinase K were prepared according to section 3.
- Set an incubator or water bath to 70 °C.
- Preheat Elution Buffer BE to 70 °C.

1 Lyse blood sample

Pipette **25 µL Proteinase K** and up to **200 µL blood**, buffy coat or body fluid sample (equilibrated to room temperature) into 1.5 mL microcentrifuge tubes (not provided).

For sample volumes less than 200 µL, add PBS to adjust the volume to 200 µL. If cultured cells are used, resuspend up to 5×10^6 cells in a final volume of 200 µL PBS.

Add **200 µL Lysis Buffer BQ1** to the samples and vortex the mixture vigorously (10–20 s).

Note: Vigorous mixing is important to obtain high yield and purity of DNA.

Incubate samples at **70 °C** for **10–15 min**.

The lysate should become brownish during incubation with Buffer BQ1. Increase incubation time with Proteinase K (up to 30 min) and vortex once or twice vigorously during incubation if processing older or clotted blood samples.

200 µL blood

**+ 25 µL
Proteinase K**

+ 200 µL BQ1

Mix

**70 °C
10–15 min**



2 Adjust DNA binding conditions

Add **200 µL ethanol (96–100 %)** to each sample and vortex again.

**+ 200 µL
ethanol**

Mix



3 Bind DNA

Apply the samples to the **NucleoSpin® Blood QuickPure Columns** placed in a Collection Tube and centrifuge **1 min** at **11,000 x g**. If the samples are not drawn through the matrix completely, repeat the centrifugation at higher *g*-force (up to 15,000 x *g*). Discard Collection Tube with flow-through.

Load lysate

**11,000 x g
1 min**



4 Wash silica membrane

Place the NucleoSpin® Blood QuickPure Column into a new Collection Tube (2 mL) and add **350 µL Buffer BQ2**. Centrifuge **3 min** at **11,000 x g**. Discard Collection Tube with flow-through.

Optional: Place the NucleoSpin® Blood QuickPure Column into a new Collection Tube (2 mL; not provided) and add 200 µL Buffer BQ2. Centrifuge 1 min at 11,000 x g. Discard flow-through and Collection Tube and proceed to step 6.

This additional washing step is only recommended if the DNA is intended for use as a template in especially critical PCRs. In the vast majority of cases, you can save time by this step.



+ 350 µL BQ2



**11,000 x g
3 min**

5 Dry silica membrane

The drying of the NucleoSpin® Blood QuickPure Column is performed by the 3 min centrifugation in step 4.

6 Elute highly pure DNA

Place the NucleoSpin® Blood QuickPure Column in a 1.5 mL microcentrifuge tube (not provided) and add **50 µL prewarmed Buffer BE (70 °C)**. Dispense buffer directly onto the silica membrane. Incubate at **room temperature** for **1 min**. Centrifuge **1 min** at **11,000 x g**.

For alternative elution procedures see section 2.4.



**+ 50 µL BE
(70 °C)**

**RT
1 min**



**11,000 x g
1 min**

6 Appendix

6.1 Troubleshooting

Problem	Possible cause and suggestions
No or poor DNA yield	<p data-bbox="309 344 736 367"><i>Low concentration of leukocytes in sample</i></p> <ul data-bbox="309 379 981 480" style="list-style-type: none"> • Prepare buffy coat from the blood sample: Centrifuge whole blood at room temperature (3,300 x g; 10 min). Three different layers will be visible after centrifugation. Leukocytes are concentrated in the intermediate layer (= buffy coat).
	<p data-bbox="309 523 512 545"><i>Incomplete cell lysis</i></p> <ul data-bbox="309 558 981 703" style="list-style-type: none"> • Sample not thoroughly mixed with lysis buffer/Proteinase K. The mixture has to be vortexed vigorously immediately after addition of lysis buffer. • Proteinase K digestion is not optimal. Never add Proteinase K directly to lysis buffer. Incubate for 15–20 min at 70 °C /56 °C.
	<p data-bbox="309 746 613 769"><i>Reagents not applied properly</i></p> <ul data-bbox="309 782 981 855" style="list-style-type: none"> • Prepare buffers and Proteinase K solution according to instructions (section 3). Add ethanol to lysates before loading them on columns.
Poor DNA quality	<p data-bbox="309 898 745 920"><i>Suboptimal elution of DNA from the column</i></p> <ul data-bbox="309 933 981 1145" style="list-style-type: none"> • Preheat Buffer BE to 70 °C before elution. Apply Buffer BE directly onto the center of the silica membrane. • Elution efficiencies decrease dramatically if elution is performed with buffers of pH < 7.0. Use slightly alkaline elution buffer like Buffer BE (pH 8.5). • Mix vigorously once during the 70 °C /56 °C incubation step especially when working with older or clotted blood samples.
	<p data-bbox="309 1189 613 1211"><i>Reagents not applied properly</i></p> <ul data-bbox="309 1224 981 1297" style="list-style-type: none"> • Prepare buffers and Proteinase K solution according to instructions (section 3). Add ethanol to lysates and mix before loading them on columns.
	<p data-bbox="309 1340 512 1362"><i>Incomplete cell lysis</i></p> <ul data-bbox="309 1375 981 1449" style="list-style-type: none"> • Sample not thoroughly mixed with lysis buffer/Proteinase K. The mixture has to be vortexed vigorously immediately after addition of lysis buffer.

- Proteinase K digestion is not optimal. Do not add Proteinase K directly to lysis buffer. Incubate for at least 15–20 min at 56 °C / 70 °C.

RNA in sample

- If RNA-free DNA is desired, add 20 µL RNase A solution (20 mg/mL) before addition of lysis buffer.

Old or clotted blood samples processed

- For isolation of DNA from older or clotted blood samples, we recommend prolonging Proteinase K incubation to 30 min and vortexing several times during this step. Especially for NucleoSpin® Blood L/XL with troublesome blood samples performance can be improved by the following steps:

First incubate the lysate for 10–15 min at room temperature. Incubate for 15 min at the recommended 56 °C afterwards. Clear lysate before addition of ethanol. It is recommended performing a short centrifugation step of about 30–60 s after the lysis of the sample material (before addition of ethanol) in order to pellet non-lysed clumps.

In case of difficult blood samples it might happen that the washing steps with ethanolic Buffer BQ2 are not sufficient to remove all contamination. An additional wash step with a buffer including chaotropic salt is recommended, for example water / BQ1 / ethanol mix (1:1:1). Afterwards, the washing step with ethanolic Buffer BQ2 should be performed to completely remove the chaotropic salt of the wash buffer.

Poor DNA
quality
(continued)

Suboptimal
performance
of genomic
DNA in
enzymatic
reactions

Carry-over of ethanol

- Be sure to remove all of ethanolic Buffer B5 / BQ2 before eluting the DNA. If the level of B5 / BQ2 after the second wash has reached the column outlet for any reason, discard flow-through, place the column back into the Collection Tube, and centrifuge again.
-

Suboptimal
performance
of genomic
DNA in
enzymatic
reactions
(continued)

Contamination of DNA with inhibitory substances

- If DNA has been eluted with Tris/EDTA-buffer (TE), make sure that EDTA does not interfere with downstream applications or repurify DNA and elute in Buffer BE.
- If preparing DNA from older or clotted blood samples, extend Proteinase K incubation to 30 min and vortex once or twice during this step.
- If the A_{260}/A_{280} ratio of the eluate is below 1.6, repeat the purification procedure:

For **NucleoSpin® Blood**: Add 1 volume of Buffer B3 plus 1 volume ethanol to the eluate, load on NucleoSpin® Blood Column, and proceed with step 3 of the corresponding protocol.

For **NucleoSpin® Blood QuickPure**: Add 1 volume of Buffer BQ1 plus 1 volume ethanol to the eluate, load on NucleoSpin® Blood QuickPure Column, and proceed with step 3 of the corresponding protocol

For **NucleoSpin® Blood L / XL**: Add 1 volume of Buffer BQ1 plus 1 volume ethanol to the eluate, load on NucleoSpin® Blood L / XL Column, and proceed with step 3 of the corresponding protocol.

6.2 Ordering information

Product	REF	Pack of
NucleoSpin® Blood	740951.10/.50/.250	10/50/250
NucleoSpin® Blood L	740954.20	20
NucleoSpin® Blood XL	740950.10/.50	10/50
NucleoSpin® Blood QuickPure	740569.10/.50/.250	10/50/250
Buffer BQ1	740923	125 mL
Buffer B3	740920	100 mL
Buffer B5 Concentrate (for 100 mL Buffer B5)	740921	20 mL
Buffer BW	740922	100 mL
Proteinase K	740506	100 mg
RNase A	740505.50 740505	50 mg 100 mg
Collection Tubes (2 mL)	740600	1000

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

6.3 Reference

Vogelstein B., and D. Gillespie. 1979. Preparative and analytical purification of DNA from agarose. Proc. Natl. Acad. Sci. USA 76: 615-619.

6.4 Product use restriction/warranty

NucleoSpin® Blood kit components are intended, developed, designed, and sold FOR RESEARCH PURPOSES ONLY, except, however, any other function of the product being expressly described in original MACHEREY-NAGEL product leaflets.

MACHEREY-NAGEL products are intended for GENERAL LABORATORY USE ONLY! MACHEREY-NAGEL products are suited for QUALIFIED PERSONNEL ONLY! MACHEREY-NAGEL products shall in any event only be used wearing adequate PROTECTIVE CLOTHING. For detailed information please refer to the respective Material Safety Data Sheet of the product! MACHEREY-NAGEL products shall exclusively be used in an ADEQUATE TEST ENVIRONMENT. MACHEREY-NAGEL does not assume any responsibility for damages due to improper application of our products in other fields of application. Application on the human body is STRICTLY FORBIDDEN. The respective user is liable for any and all damages resulting from such application.

DNA/RNA/PROTEIN purification products of MACHEREY-NAGEL are suitable for *IN-VITRO*-USES ONLY!

ONLY MACHEREY-NAGEL products specially labeled as IVD are also suitable for *IN-VITRO*-diagnostic use. Please pay attention to the package of the product. *IN-VITRO*-diagnostic products are expressly marked as IVD on the packaging.

IF THERE IS NO IVD SIGN, THE PRODUCT SHALL NOT BE SUITABLE FOR *IN-VITRO*-DIAGNOSTIC USE!

ALL OTHER PRODUCTS NOT LABELED AS IVD ARE NOT SUITED FOR ANY CLINICAL USE (INCLUDING, BUT NOT LIMITED TO DIAGNOSTIC, THERAPEUTIC AND/OR PROGNOSTIC USE).

No claim or representations is intended for its use to identify any specific organism or for clinical use (included, but not limited to diagnostic, prognostic, therapeutic, or blood banking). It is rather in the responsibility of the user or - in any case of resale of the products - in the responsibility of the reseller to inspect and assure the use of the DNA/RNA/protein purification products of MACHEREY-NAGEL for a well-defined and specific application.

MACHEREY-NAGEL shall only be responsible for the product specifications and the performance range of MN products according to the specifications of in-house quality control, product documentation and marketing material.

This MACHEREY-NAGEL product is shipped with documentation stating specifications and other technical information. MACHEREY-NAGEL warrants to meet the stated specifications. MACHEREY-NAGEL's sole obligation and the customer's sole remedy is limited to replacement of products free of charge in the event products fail to perform as warranted. Supplementary reference is made to the general business terms and conditions of MACHEREY-NAGEL, which are printed on the price list. Please contact us if you wish to get an extra copy.

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