WizPrep™ gDNA Mini Kit (Blood)

IVD in vitro Diagnostic Use Only



DESCRIPTION

The WizPrep™ gDNA Mini Kit (Blood) provides a fast and simple method to isolate genomic DNA from various blood sample including fresh whole blood, frozen blood, blood dried spot, serum and plasma etc.

The WizPrep™ gDNA Mini Kit (Blood) uses silica-membrane technology to eliminate the cumbersome steps associated with loose resins or slurries. The kit is ready for use and can purify the genomic DNA from a wide variety of blood source samples, and the whole process is completed in less than 20 minutes.

Purified DNA is suitable for PCR, restriction endonuclease digestion and Southern Hybridization.

KIT CONTENTS

Contents	100 prep	250 prep	Storage
GB Buffer	25 ml	65 ml	R/T
W1 Buffer (concentrate) ⁽¹⁾	30 ml	75 ml	R/T
W2 Buffer (concentrate) ⁽¹⁾	12 ml	30 ml	R/T
Elution Buffer	10 ml	25 ml	R/T
Proteinase K (lyophilized)*	22mg x 2	22mg X 5	4℃
Spin Columns**	100 pcs	250 pcs	R/T
Instruction Manual	1 pcs	1 pcs	

- 1): Add absolute ethanol to the W1 and W2 Buffer prior to initial use (see the bottle label for volume).
- * After receiving the Proteinase K, please store at 4°C. After dissolved in distilled water, store the Proteinase K solution at 4°C for up to 3 months. For longer storage (up to 1 year), the Proteinase K solution should be divided into small aliquot and stored at -20°C.
- ** All Spin Columns are sterilized by electron beam.

REAGENTS AND EQUIPMENT TO BE SUPPLIED BY USER

- 96-100% ethanol (to prepare W2 Buffer)
- 1.5ml microcentrifuge tubes
- Sterile RNase-free pipette tips and Manual pipettors
- Centrifuge for microcentrifuge tubes
- Equipment for sample disruption and homogenization
- Personal protection equipment (lab coat, gloves, goggles)

KIT SPECIFICATIONS

Parameter	Characteristics	
Format	Silca-membrane spin column	
Sample materials	< 100 mg tissue	
Fragment size	> 200 nt	
Typical yield	3-12 μg from 200 μl whole blood	
	20-30 μg from 200 μl buffy coat	
	15-25 µg from 5 X 106 cultured cells	
A260/280	1.7 - 1.9	
Elution volume	50 μΙ	
Preparation time	< 20 minutes	
Binding capacity	200 ug	

QUALITY CONTROL ANALYSIS

In accordance with Wizbiosolutions Inc. ISO 13485-certified Quality Management System, each lot WizPrep™ gDNA Mini Kit (Blood) is tested against predetermined specifications to ensure consistent product quality.

PROTOCOL

To check before start

- 1) Add absolute ethanol to the W1 and W2 Buffer prior to initial use (see the bottle label for volume).
- 2) Dissolve each vial of Proteinase K (22mg) in 1,100µl of D.W
- 3) If a precipitate has formed in GB Buffer, dissolve by incubating at 56°C before use.

1. Lysis

- Add 200µl of whole blood to 1.5ml microcentrifuge tube. (Use up to 200 µl plasma, serum, buffy coat, body fluids, or up to 5 x 106 lymphocytes in 200µl of PBS).
- Add $\ 200\mu l$ of GB Buffer and $\ 20\mu l$ of Proteinase K and mix by vortexing.
- Incubate at 56 $^{\circ}\text{C}\,$ for 10 minutes. During incubation, invert the tube every 5 minutes.

2. Binding

- Add 200µl of 100% ethanol to the sample lysates and mix by vortexing briefly.
- Connect Spin Column to 2.0ml Collection tube.
- Apply the mixture to the Spin Column and centrifuge for 1 min. at 13,000 rpm.
- Transfer the Spin Column to the new Collection tube (providede) and discard the flow-through with Spin Column.

3. Wash

- Add 500µl of W1 Buffer to the Spin Column and centrifuge for 1 min. at 13,000 rpm.
- Discard the flow-through and re-connect with Spin Column.
- Add 500µl of W2 Buffer (ethanol added) in the Spin Column and centrifuge for 1 min. at 13,000 rpm.
- Discard the flow-through and re-connect the Spin Column and centrifuge for 2 min. at 13,000 rpm.

4. Elution

- Connect the Spin Column and new 1.5 ml tube.
- Add 50-100µl of Elution Buffer and incubate at R/T for 1 min.
- Centrifuge for 1 min. at 13,000 rpm.
- Discard the Spin Column and eluted purified DNA for use next step.

5. DNA Storage

• The purified DNA sample may be stored at -20°C for a few days. It is recommended that samples be placed at -70°C for long term storage.



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TROUBLESHOOTING

Problem	Possiblity	Suggestions
Column clogged	Sample overloading	Reduce sample volume or separate into multiple tubes.
	Precipitate was formed at DNA Binding Step	Reduce the sample material. Before loading the column, break up precipitate in ethanol-added lysate
Low yield	Incorrect DNA Elution Step Incomplete DNA Elution	Ensure that Elution Buffer was added and absorbed to the center of the Column matrix. Elute twice to increase yield
Eluted DNA does not perform well in downstream applications	Residual ethanol contamination RNA contamination Genomic DNA was degraded	 Following the washing step, dry the Column with additional centrifugation at full speed for 3 minutes. Perform Optional RNA Degradation Step.
	30 dog. 200	• Use fresh sample, long storage may result in fragmentation of genomic DNA.

SYMBOL GLOSSARY

REF	Catalogue number	•••	Manufacturer
LOT	Batch code	IVD	In-vitro diagnostic use only
1	Temperature limit	[]i	Instructions for use
Ω	Use-by date		

ORDERING INFORMATION

Product	Cat No.	Package
WizPrep™ gDNA Mini Kit (Blood)	W71050-100	100 prep
WIZI TEP GENTA WITH THE (BIOOU)	W71050-250	250 Prep



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