**EZgeneTM Whole Blood DNA Maxi kit**

**（BW-GD2314）**

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# Kit Contents

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| --- | --- | --- | --- |
| **Catalog #** | **GD2314-00** | **GD2314-01** | **GD2314-02** |
| Preps | 2 | 10 | 25 |
| ezBind DNA Maxi Columns | 2 | 10 | 25 |
| 50 mL Collection Tubes | 4 | 20 | 50 |
| Buffer BL | 21 mL | 102 mL | 255 mL |
| Buffer KB | 31 mL | 152 mL | 380 mL |
| 10 x Red Blood Cell Lysis Buffer | 20mL | 100 mL | 250 mL |
| DNA Wash Buffer | 15 mL | 3 x 24mL | 3 x 54mL |
| Elution Buffer | 18 mL | 90 mL | 210 mL |
| Protease K | 1.0 mL | 5.1mL | 13mL |
| RNase A (20 mg/mL) | 45 μL | 210 μL | 520 μL |
| Instruction Booklet | 1 | 1 | 1 |

# Introduction

The EZgeneTM Blood maxi gDNA kit provides a fast and easy method for isolating gDNA from blood. The system utilizes the reversible nucleic acid- binding properties of Biomiga’s ezBind membrane and the speed of spin column technology to yield high quality gDNA with the OD260/280 ratio of 1.8-

* 1. Up to 20 mL of fresh, frozen or anticoagulated whole blood treated with any common anticoagulant such as heparin, EDTA, or acid-citrate-dextrose can be readily processed at one time. This DNA Kit can also be used for the preparation of genomic DNA from buffy coat, serum, plasma, saliva, buccal swab and other body fluids. Purified DNA is ready for applications such as PCR, Southern Blotting, and Restriction Digestion.

The binding capacity per column is 1000 μg of gDNA. Use less than 20 mL of whole Blood or buffy coat is recommended.

# Storage and Stability

All EZgeneTM Blood maxi gDNA Kit components are guaranteed for at least 12 months from the date of production when stored as follows:

* + - Reconstituted Protease K store at -20°C.
		- All other materials at room temperature (15-25°C).

# Before Starting

The kit is designed to be simple, fast, and reliable provided that all steps are followed diligently. Please read the entire booklet and get all necessary supplies and equipments.

# Important

* **Dilute DNA Wash Buffer with absolute ethanol as follows: Add 60 mL (GD2314-00) or3 x 96mL (GD2314-01) or3 x 216mL (GD2314-**

**02) of absolute ethanol to each bottle**. The final concentration is 80%.

* Under cool ambient conditions, precipitates may form in Buffer BL. In case of such an event, heat the bottle at 37°C to dissolve before use.
* Preheat the Elution Buffer at 70°C **. Materials provided by user**
* Centrifuge capable of 5,000 x g
* Sterile 50 mL centrifuge tubes
* Water bath set to 55°C and 70°C
* Absolute ethanol

# Safety Information

Buffer BL contains chaotropic salts, which may form reactive compounds when combines with bleach, Do not add bleach or acidic solutions directly to the preparation waste, ware gloves and protective eyewear when handling.

# EZgeneTM Whole Blood DNA Maxi Isolation Protocols

Calculate the amount of 10 x Red Blood Cell Lysis Buffer to be used. Mix one part of 10 x Red Blood Cell Lysis Buffer with nine part of ddH2O before use.

Final concentration of Red Blood Lysis Buffer:

NH4Cl 155 mM

KHCO3 10 mM

Na2EDTA 0.1 mM

Adjust to pH7.4 with 1 M HCl or NaOH

1. Transfer **10 mL whole blood sample** (up to 20 mL) into a two 50 mL tube (10 mL each) and add **3 volumes** of **Red Blood Lysis Buffer**. For example, add 30 mL Red Blood Lysis Buffer to 10 mL whole blood. Mix by vortexing.

**Note:** Red Blood Cell Lysis Buffer is supplied as a 10 x concentrate and must be diluted with ddH2O according to bottle label before use.

1. Incubate for 15 to 20 min on ice, mixing by brief vortexing twice. The solution will become translucent.
2. Centrifuging at 450 g for 10 min at 4°C to precipitate the leukocytes. Discard the supernatant containing lysed red blood cells.
3. Wash the leukocytes with **2 volumes** of **Red Blood Lysis Buffer** of sample used in step 1. For example, add **20 mL Red Blood Lysis Buffer** if **10 mL** sample is used.
4. Centrifuging at 450 g for 10 min at 4°C again to precipitate the leukocytes. Discard the supernatant containing lysed red blood cells. Transfer the leukocytes together.
5. Add **0.5 mL** of **Protease K** .
6. If RNA-Free genomic DNA is required, add **20 μL** of **RNase A** (provided) to each sample and mix briefly by vortexing. **Add 10 mL** of **Buffer BL**. Mix well by vortexing at maximum speed for 30s.

Note: Precipitates may form in Buffer BL. Dissolve at 37°C before use.

1. Incubate the sample at 50°C for 20 minutes. Briefly vortex the tube once during incubation.
2. Add **6 mL absolute ethanol** to the lysate. Vortex at maximum speed for 10s. Briefly centrifuge the tube to collect any drops from the lid.
3. Place the column into the sample 50 mL tube. Transfer up to **20 mL** sample from step 9 to the DNA maxi column, and centrifuge at 4,000 g for 5 min to bind DNA. Discard flow-through liquid and reuse the collection tube.
4. Place the column into the collection tube. Add **15 mL** of **Buffer KB,** and centrifuge at 4,000 g for 5 min. Discard flow through liquid .
5. Add **15 ml of DNA Wash Buffer** *(Add ethanol before use)****.*** Centrifuge at 4,000 g for 5 min. Discard flow through liquid. Repeat step 12 to wash the column again.
6. Put the empty column, **with the lid open**, into the same collection tube and centrifuge at 5,000 g for 15 min to dry the column. **Note：**Residual ethanol

will be removed more efficiently with the lid of the column open. It is critical to remove the ethanol before elution. **Note:** Please dry the column in a vacuum oven at 50oC for 20 min if ethanol did not remove clearly.

1. Place the column into a sterile new 50 mL microtube, add **5 mL Elution Buffer**. Incubate at 65oC for 10 min . Centrifuge at 10,000 g for 5 min to elute the DNA. Add another **3 mL** of preheated Elution buffer to elute. If maxium yield is required, please repeat wash for serval times with elution buffer. Combine the DNA eluant.

# Determination of Yield and Quality

The total DNA yield can be determined by a spectrophotometer using deionized water, Tris-HCl Buffer, or Elution Buffer as a blank. Dilute the DNA in TE buffer and calculate the concentration using the following equation:

[DNA] = (Absorbance 260) x (0.05 μg/μL) x (Dilution Factor)

The quality of DNA can be assessed by measuring absorbance at both 260 nm and at 280 nm. A 260/ 280 ratio of 1.7-1.9 corresponds to 85%-95% purity. Expected yields will range from 4 μg - 12 μg of DNA per 250 μL of whole blood, depending on the source of the sample, its age, and the method of storage. Yields are generally 5-fold higher with buffy coat samples.

# Trouble Shooting Guide

|  |  |  |
| --- | --- | --- |
| **Problem** | **Cause** | **Possible Solution** |
| **Clogged Column** | Incomplete Lysis | Add the correct volume of Buffer BL and incubate for specified time at 65°C . It may be necessary to extendincubation time by 10 min. |
|  | Sample is too Large | If using more than 10 mL of Blood, increase volumes of Protease K, Buffer BL,. Pass aliquots of lysate through onecolumn successively. |
|  | Sample is too viscous | Divide sample into multiple tubes |
| **Low DNA Yield** | Clogged Column | See above |
| Poor elution | Repeat elution or increase elution volume (see notes on elution on page 6) Incubation of column at 70°C for 5 minwith Elution Buffer may increase yields. |
|  | Improper washing | Wash Buffer Concentrate must be diluted with absolute (100%) ethanol as specified on page 3 before use. |
| Buffy Coat Used | With Buffy Coat samples, use absolute ethanol, rather thanisopropanol . |
| **260****/280Low A****/A Ratio** | Extended centrifugation during elution. | Resin from the column may be present in elute and affect the OD absorbance. Avoid centrifugation at speed higherthan 15,000 x g. The trace resin in the eluted DNA will not interfere with PCR or restriction digests. |
|  | Poor cell lysis due to incomplete mixingwith Buffer BL. | Repeat the procedure, this time making sure to vortex the sample with Buffer BL immediately and completely. |
|  | Hemoglobin Remainson column | After application of sample to the column, wash once with2 mL of Buffer BL. |
| **No DNA****Eluted** | Poor Cell Lysis due toimproper mixing with Buffer BL | Mix thoroughly with Buffer BL prior to loading the column. |
|  | Absolute ethanol notadded to Buffer BL. | Before applying the sample to the column and aliquot ofBuffer BL/ethanol solution must be added. |
|  | No Ethanol added to Wash BufferConcentrate | Dilute Wash Buffer with the indicated volume of absolute ethanol before use (page 3). |
| **Washing Leaves Colored Residue in Column** | Incomplete Lysis due to improper mixingwith Buffer BL. | Buffer BL is viscous and the sample must be mixed thoroughly |
| No Ethanol added toWash Buffer Concentrate | Dilute Wash Buffer with the indicated volume of absolute ethanol before use. |
| **Eluted Material has Red or****Brown Color** | Sample Volume is tooLarge | Reduce sample volume, and proceed with protocol. |
| Hemoglobin remains in column | After applying sample, wash column once with 3 mL of Buffer BL. |

# Limited Use and Warranty

This product is intended for *in vitro* research use only. Not for use in human.

This product is warranted to perform as described in its labeling and in BEIWO’s literature when used in accordance with instructions. No other warranties of any kind expressed or implied, including, without limitation, implied warranties of merchantability or fitness for a particular purpose, are provided by BEIWO. BEIWO’s sole obligation and purchaser’s exclusive remedy for breach of this warranty shall be, at the option of BEIWO, to replace the products, BEIWO shall have no liability for any direct, indirect, consequential, or incidental damage arising out of the use, the results of use, or the inability to use it product.

For technical support or learn more product information, please contact us or visit our website.



[**Contact Us**](https://www.sigmaaldrich.cn/CN/en/collections/offices)**:** 400-115-2855

[www.beiwobiomedical.com](http://www.beiwobiomedical.com)

[**Customer Support**](https://www.sigmaaldrich.cn/CN/en/support/customer-support)**:**

market@beiwobiomedical.com

**Technical Support****:**

tech@beiwobiomedical.com