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**HANDBOOK**

# **QuickGene-AutoS DNA Blood Kit AS-DB (IVD)**

**For extraction of genomic DNA from whole blood**

















Ver.CEIVD1.0

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All labels of Bottle, Reagent, Package and Manual including this HandBook use the symbols mentioned below.

Symbol	Description	Symbol	Description
	In Vitro Diagnostic Medical Device		Consult instruction for use
	Use by date		Manufacturer
	BATCH CODE		Caution
	CE MARKING		Catalogue Number
	Temperature limitation		Contains reagents sufficient for <n> number
	Date of Manufacture		Authorized representative in the European Community
	Contains		Adding

# 1. Introduction

QuickGene porous membrane to immobilize nucleic acid has large specific surface area and uniform & fine porousness. So QuickGene successfully extracts genomic DNA with high yield. QuickGene also uses pressured filtration technology, which enables producing new, compact and automatic instruments for rapid nucleic acid purification.

The specifications mentioned above will reduce reexamination and produce reliable observation. QuickGene-AutoS DNA blood kit / AS-DB (IVD) is intended to use for in vitro diagnostic medical device to sample DNA purifying from human whole blood.

## INDENDED USE

The combination of QuickGene-Auto12S/QuickGene-Auto24S system and QuickGene-AutoS DNA whole blood kit / AS-DB (IVD) is intended to isolate high quality genomic DNA automatically from human whole blood sample. Generally, DNA isolated by the system is useful for PCR based analysis like HLA typing or karyotyping to know patients' genotype before transplantation, and for Next Generation Sequencing (NGS) for selection of the molecular targeted agents. Such high quality genomic DNA is also suitable for the long term storage project like bio banking with less DNA degeneration/degradation. DNA isolated from the system can't be used for diagnosis, prevention, or treatment of a disease purpose directly. The system and the kit are intended for use by professional users adequately skilled in molecular biological techniques and trained to operate the system.



QuickGene DNA whole whole blood kit L / DB-L (IVD) does NOT intend to use the subject detection written in either List A or List B in Annex, II IVD directive 98/79/EC, or Self testing.

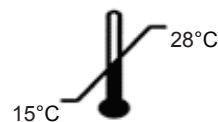
**Please be sure to read this handbook carefully before using the kit.**

**This Kit is only used with QuickGene-Auto12S or QuickGene-Auto24S as IVD marked product.**

## 2. Kit Components and Storage Conditions

### 2-1. Kit Components (48 Preps)

- |  |        |
|--|--------|
| <input type="checkbox"/> Protease(Lyophilized) / EDB | 1 vial |
| <input type="checkbox"/> Reagent strips              | 48 pcs |
| <input type="checkbox"/> 1 ml Long Tips              | 48 pcs |
| <input type="checkbox"/> Waste Tubes / (WT)          | 48 pcs |

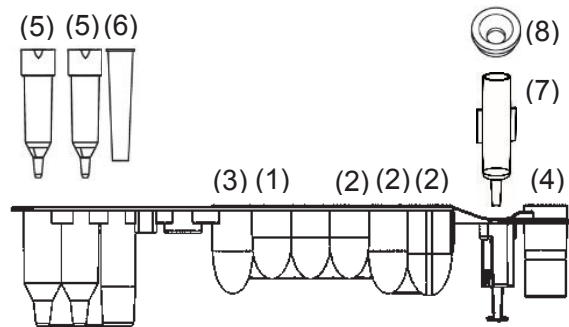


### 2-2. Storage Conditions

All reagents are stable at room temperature (15-28°C) until expiring date indicated at outer box. Reconstituted EDB is stable for 2 months when stored at 4°C. After dissolution, the EDB can be stored in a frozen state (-20°C) for at least 6 months. In such a case, do not repeat the freezing and thawing by dispensing 1.5 mL or 2 mL tubes.

## 2-3. Reagent strip components

<input type="checkbox"/>	Lysis Buffer	LDB	270 µl	(1)
<input type="checkbox"/>	Wash Buffer	WDB	750 µl, 3 positions	(2)
<input type="checkbox"/>	(>99%) Ethanol		270 µl	(3)
<input type="checkbox"/>	Elution Buffer	CDB	250 µl	(4)
<input type="checkbox"/>	Short Tip		2	(5)
<input type="checkbox"/>	Tip pack		1	(6)
<input type="checkbox"/>	Cartridge		1	(7)
<input type="checkbox"/>	Pressure Adapter		1	(8)



## 3. Other Required Materials, Not Supplied in This Kit

### [1] Reagents

- Nuclease-free water (for dissolving EDB)

### [2] Equipment

- QuickGene-Auto12S or QuickGene-Auto24S
- Micropipettes and tips
- Tube stand
- 2 ml microtubes for samples

Recommendation product: BM EQUIPMENT Cat. BM4020

SARSTEDT Cat.72.695.700, Cat.72.695.500S



When using a tube other than the recommended product, check the compatibility with the strip and equipment heater part beforehand.

- 1.5 ml or 2 ml microtubes for elution of DNA

Recommendation product: BM EQUIPMENT Cat. BM4015,

SARSTEDT Cat.72.706.700



When using a tube other than the recommended product, check the compatibility with the Collection holder beforehand.

## 4. Safety Warnings



All reagents and items should be considered chemically and biologically hazardous. Wearing a laboratory coat, disposable gloves and safety goggles during the experiments are highly recommended. In case of contact between the reagents and the eyes, skin, or clothing, wash immediately with water. (See the Safety Data Sheet for specific recommendations, <http://www.kurabo.co.jp/bio/English/>)

### ◆ Protease EDB



CONT

Proteinase, Bacillus neutral



Danger! Wear protective gloves/protective clothing/eye protection/face protection. IF ON SKIN (or hair): take off immediately all contaminated clothing. Rinse skin with water/shower. IF INHALED: Remove person to fresh air and keep comfortable for breathing. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. Call a POISON CENTER/doctor/.../if you feel unwell.

### ◆ LDB (Lysis Buffer)



CONT

Guanadine hydrochloride

Warning! Wear protective gloves/protective clothing/eye protection/face protection. IF exposed or concerned: Get medical advice/attention. Supplemental hazard information: 2,4,7,9-tetramethyldec-5-yne-4,7-diol May produce an allergic reaction.

### ◆ EtOH (Ethanol)



Warning! Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking. Keep container tightly closed. Ground/bond container and receiving equipment. Use explosion-proof electrical/ventilating/lighting/equipment. Use only non-sparking tools. Take precautionary measures against static discharge. Do not breathe dust/fumes/gas/mist/vapours/spray. Avoid breathing dust/fumes/gas/mist/vapours/spray. Wash contaminated parts thoroughly after handling. Do not eat, drink or smoke when using this product.

### ◆ WDB (Wash Buffer)



Warning! Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking. Keep container tightly closed. Ground/bond container and receiving equipment. Use explosion-proof electrical/ventilating/lighting/equipment. Use only non-sparking tools. Take precautionary measures against static discharge. Do not breathe dust/fumes/gas/mist/vapours/spray. Avoid breathing dust/fumes/gas/mist/vapours/spray. Wash contaminated parts thoroughly after handling. Do not eat, drink or smoke when using this product.

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◆ **CDB (Elution Buffer)**

- Do not drink or ingest. Avoid contact with eyes.
- If contact with eyes, skin, or clothing occurs, rinse thoroughly with water. Consult a physician if necessary.

◆ Use or storage of Reagent strips at the specified temperature (15°C – 28°C).

◆ Any solution and waste fluid containing LDB should not be mixed with bleach.

◆ **In the case of using potentially infectious samples:**

Wear a suitable laboratory coat, disposable gloves and safety goggles during the experiments.

◆ **Disposal of waste fluid and consumables when using potentially infectious samples:**

After use, dispose of potentially infectious samples and consumables by incineration, high-temperature decontamination, sterilization, or disinfection in accordance with applicable laws. When entrusting waste disposal to licensed hazardous waste disposal contractors, use specially controlled waste management forms (manifest), if applicable.

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## 5. Precautions

### ◆ Handling of Starting Material

- Small amount of samples should be adjusted to 200 µl with PBS (sterilized) before loading.
- Use a whole blood sample treated with EDTA-2Na, EDTA-2K, or heparin.
- Use a whole blood sample within 3 days after collection. The yield of DNA might decrease, or degradation of DNA might be caused when a blood sample preserved for a long time is used.
- The yield of DNA might decrease when the number of leucocytes exceeds  $2 \times 10^6$  cells/200 µl. In such cases, adjust the number of leucocytes by diluting the sample with PBS (sterilized) to below  $2 \times 10^6$  cells/200 µl.
- The Cartridge (CA) might clog when the number of leucocytes exceeds  $5 \times 10^6$  cells/200 µl. We recommend that you dilute the sample with PBS (sterilized) and then perform extraction.

### ◆ Use of Reagent

- After addition of nuclease-free water to EDB leave it for 30 min or more at room temperature with occasionally stirring. Use it after confirming the powder is completely dissolved. The yield of DNA might decrease or the Cartridge (CA) might clog when dissolution of EDB is insufficient.

### ◆ Procedure of Extraction

- Before starting operation, please make sure the following things:
  - Waste Tubes and 1.5 ml or 2 ml microtubes (for elution) are set in the Collection holder.
  - Reagent strips are set correctly in the Reagent holder.
  - 1 ml Long tips and 2 ml microtubes (Whole blood and EDB included) are set in the Reagent strip.
  - The lid of Reagent holder is completely closed.
  - Reagent holder and Collection holder are properly set in the holder guide.
- All operations should be performed at room temperature (15 to 28 °C). In case of using at lower or higher temperature, it may affect the extraction performance.
- Except for unavoidable circumstances, please do not turn off the QG-Auto12S or QG-Auto24S device during operation. You cannot resume operation from the same process.
- Refer to the Operation Manual of QuickGene-Auto12S / QuickGene-Auto24S for details.



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## 6. Quality Control

- As part of the stringent quality assurance program in KURABO INDUSTRIES LTD., the performance of QuickGene-AutoS DNA Blood Kit / AS-DB (IVD) is evaluated routinely on a lot-to-lot uniformity.
- Yield and quality of extracted genomic DNA are checked by measuring the absorbance at 260 nm, ratio of absorbance (260 nm/280 nm).

## 7. Product Description

Table 2 shows the average of yield and purity (A260/280) of genomic DNA extracted from 200 µl of whole blood samples.

The yield varies depending upon sample conditions.

Table 2

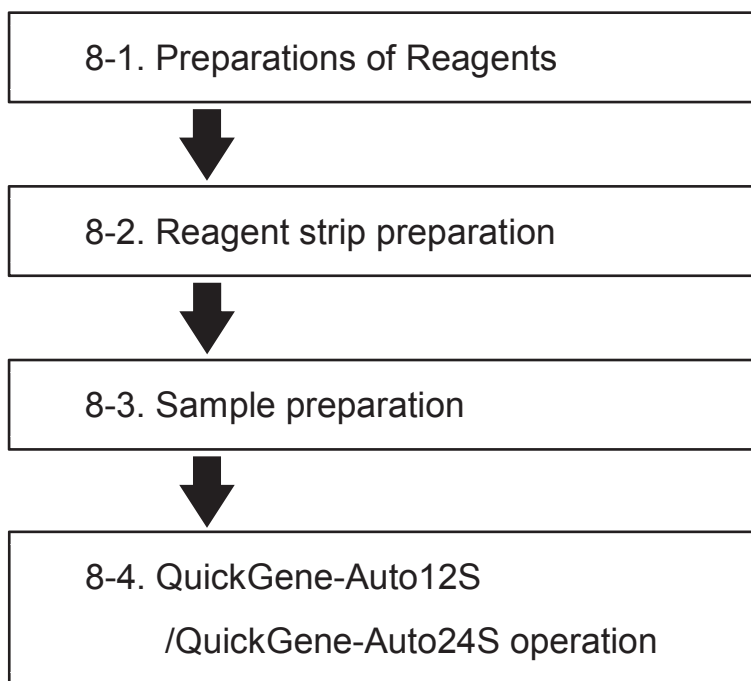
Sample	Amount of genomic DNA (µg)	A260/280
Whole blood (200 µl)	5.8	1.95

\*DNA and RNA are included in eluate extracted with this kit.

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## 8. Protocol

### [Overview Flow Chart]



### 8-1. Preparations of Reagents

#### ◆ EDB (Lyophilized)

**ADD** 3.3 ml water

When using EDB, pipette 3.3 ml of nuclease-free water into the vial containing lyophilized Protease. Dissolve it completely. Reconstituted EDB is stable for 2 months when stored at 4°C. Storage at –20°C will prolong the life of EDB, but repeated freezing and thawing should be avoided. Dividing the solution into aliquots and storage at –20°C is recommended.

#### Notices

Dissolve EDB completely by the following method, and then use the solution.

Add 3.3 ml of nuclease-free water, close the cap and mix with inversion the bottle.

Leave it for 30 min or more at room temperature with occasionally stirring it. Use it after confirming the powder is completely dissolved. The yield of DNA might decrease or the Cartridge (CA) might clog when dissolution of EDB is insufficient.

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## 8-2. Reagent strip preparation



To avoid contamination of nuclease, wear disposable gloves during preparation of Reagent strips and microtubes.



Refer to the Operation Manual of QuickGene-Auto12S / QuickGene-Auto24S for details.

- <1> Prepare the Collection holder and Reagent holder on the workbench.
- <2> Load the waste tube and 1.5 ml or 2 ml microtube into the Collection holder.
- <3> Remove the Reagent strips from the kit box, place it in the Reagent holder, and insert 1 ml Long Tip in the specified position.

## 8-3. Sample preparation

### Notices

Follow the protocol of <1> to <3> exactly.

In case the procedure is changed, the yield of DNA may not be obtained.



Wear a suitable laboratory coat, disposable gloves and safety goggles during the experiments.

To avoid contamination of nuclease, wear disposable gloves during preparation of Reagent strips and microtubes.



Refer to the Operation Manual of QuickGene-Auto12S / QuickGene-Auto24S for details.

- <1> Add 30  $\mu$ l of EDB (previously dissolved in nuclease-free water) to the bottom of a 2 ml microtube.

Recommendation product of 2 ml microtube: M&S Instruments Cat. 4020

SARUSTEDT Cat.72.695.700

Cat.72.695.500S

- <2> Add 200  $\mu$ l of a whole blood sample.
- <3> Set the 2 ml microtube containing whole blood and EDB in the specified position on the Reagent strip.

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## 8-4. QuickGene-Auto12S/QuickGene-Auto24S operation



To avoid contamination of nuclease, wear disposable gloves during preparation of Reagent strips and microtubes.



Refer the Operation Manual of QuickGene-Auto12S / QuickGene-Auto24S for the details before using the device.

- <1> Open the front door and put the Collection holder and Reagent holder to the specified positions on the machine.
- <2> Turn on the device.  
The device proceeds through a self-check and moves to the home position about all moving parts.
- <3> At the Home screen, select the "DNA WHOLE BLOOD".
- <4> Chose the elution volume
- <5> Make sure all the accessories has been putted in the system. Tick the check list then the "Next" button will show up.
- <6> Press the "Next" button.
- <7> Check the protocol information is correct, then press the "Start" button to proceed the isolation. Then processing will be started.
  - During the running step, the touch panel show the processing and remaining time.
  - Operation status can be confirmed by blinking process name (LYSIS, BINDING, WASH, ELUTE, FINISH).
  - Do not open the front door of the device while running. If you open the front door, please read the Operation manual of QG-Auto12S / QG-Auto24S and resume operation.
  - To pause, touch the "Pause" button on the operation panel. The end confirmation screen will be displayed, please press "Yes" to finish.
- <8> After finishing the protocol, the beeper will call and the process name "FINISH" flashes on the operation panel.  
After confirming that the device is completely stopped open the front door, take out the Reagent holder and the Collection holder.  
Take out the elution tube from the Collection holder.
  - The standard yield is 4 to 8 µg from 200 µl of whole blood samples.
  - If you do not use DNA immediately, please close the tube lid tightly and store at 4°C or -20°C.
  - In case of storing genomic DNA for a long time, it is recommended to preserve them at -20°C.

## 9. Troubleshooting

Review the information below to troubleshoot the experiments with QuickGene-AutoS DNA Blood Kit / AS-DB (IVD). For system related problems, see the QuickGene-Auto240L Operation manual.

### (1) Low yield or no DNA obtained

Cause	Action
Inappropriate storage conditions for whole blood sample	Use a whole blood sample within 3 days after collection as much as possible. The yield of DNA might decrease, or degradation of DNA might be caused when a blood sample preserved for a long time is used.
Inadequate dissolution of EDB	After addition of nuclease-free water to EDB leave it for 30 min or more at room temperature with occasionally stirring it. Use it after confirming the powder is completely dissolved.
Insufficient enzymatic activity of EDB	Reconstituted EDB is stable for 2 months when stored at 4°C. Do not use EDB preserved for a longer period than 2 months. Storage at -20°C will prolong the life of EDB, but repeated freezing and thawing should be avoided. Dividing the solution into aliquots and storage at -20°C is recommended.
Inappropriate addition order of reagents and whole blood sample	When preparing lysates, perform the additions to a 2 ml microtube in the following order : EDB (previously dissolved in 3.3 ml of nuclease-free water) --> Whole blood sample
Inappropriate volume of whole blood sample	If the volume of a whole blood sample is too much, reduce it to the prescribed volume (200 µl). Small amount of sample should be adjusted to 200 µl with PBS (sterilized) before loading.
Use of too much amount of leucocytes	The yield of DNA might decrease when the number of leucocytes exceeds $2 \times 10^6$ cells/200 µl. In such cases, adjust the number of leucocytes by diluting the sample with PBS (sterilized) to below $2 \times 10^6$ cells/200 µl.
DNA degradation	Refer to (3) "DNA degradation".

### (2) Clogging of Cartridge (CA) occurs

Cause	Action
Use of too much amount of a whole blood sample	Reduce it to the prescribed volume (200 µl).
Use of too much amount of leucocytes	The Cartridge (CA) might clog when the number of leucocytes exceeds $5 \times 10^6$ cells/200 µl. The yield of DNA might decrease when the number of leukocytes exceeds $2 \times 10^6$ cells/200 µl. In such case, we recommend that you dilute the sample with PBS (sterilized) to below $2 \times 10^6$ cells/200 µl, and then perform extraction.

### (3) DNA degradation

Cause	Action
Inappropriate storage conditions for whole blood sample	Use a whole blood sample within 3 days after collection as much as possible. The yield of DNA might decrease, or degradation of DNA might be caused when a blood sample preserved for a long time is used.

### (4) Purity of DNA is low

Cause	Action
Inappropriate storage conditions for whole blood sample	Use a whole blood sample within 3 days after collection as much as possible. The yield of DNA might decrease, or degradation of DNA might be caused when a blood sample preserved for a long time is used.
Insufficient enzymatic activity of EDB	Reconstituted EDB is stable for 2 months when stored at 4°C. Do not use EDB preserved for a longer period than 2 months. Storage at -20°C will prolong the life of EDB, but repeated freezing and thawing should be avoided. Dividing the solution into aliquots and storage at -20°C is recommended.

### (5) Subsequent experiments such as PCR etc. do not proceed well

Cause	Action
Inappropriate amount of DNA is used	Determine the DNA concentration based on the absorbance at 260 nm.
Low purity of DNA	Refer to (4) "Purity of DNA is low".
DNA degradation	Refer to (3) "DNA degradation".

### (6) A precipitate is formed in reagents

Cause	Action
Stored at low temperature	Store this kits at room temperature (15-28°C). If a precipitate is formed, dissolve the precipitate by incubation at 37°C. Cool down it to room temperature before use.

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## 10. Attention for Administrator and Operator

To avoid any accident and unfortunate risk, administrator and operator adhere closely to the rule mentioned below.

### 10-1. Administrator should

- a. Set specific space to apart working area from standard area for using this kit and instrument.
- b. Make operation learn the risk of using QuickGene to operator including potential operator.
- c. Approve blood sample without any infection from donor before use.

### 10-2. The education for operator consists at least below

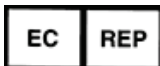
#### Operator should keep

- a. To be out of danger in the process of DNA isolation using this kit and instrument.
- b. To take steps to prevent any infection from unknown virus.
- c. To be conscious the important of double blind methods to secure the personal information from the sequence information of DNA, which is isolated from whole blood with this kit and instrument.
- d. To protect environment by avoiding the exposure of disposal plastic ware in this kit and the waste materials after operation.
- e. Not to be conducted this kit and instrument by the third party, who did not learn the education.
- f. To work with the kit and instrument in the specific directive area.
- g. The best physical condition and to check own healthy situation before the operation.
- h. Not to sterilize this kit and instrument by high pressure and high temperature.

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\* Trademark and exclusion item

Right to registered name etc. used in this handbook is protected by law especially even in the case of no denotation.



**MedNetGmbH**

Borkstrasse 10,48163 Muenster, Germany



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## **KURABO INDUSTRIES LTD.**

Advanced Technology Division, Bio-Medical Department

14-30, Shimokida-Cho, Neyagawa,

Osaka 572-0823, Japan

TEL +81-72-820-3079 FAX +81-72-820-3095

URL; <http://www.kurabo.co.jp/bio/English/>