

PureHelix™ Genomic DNA Prep Kit [Rice] (Ver. 3.0) Column type

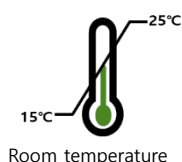
Kit contents

PureHelix™ Genomic DNA Prep Kit [Rice] (Ver.3.0)			
Cat. No.	GCTR50 (50preps)	GCTR100 (100preps)	GCTR200 (200preps)
NGD1	15ml	30ml	60ml
NPS2	15ml	30ml	60ml
EB	5ml	10ml	20ml
WB	11ml (Add ethanol 44 ml)	22ml (Add ethanol 88 ml)	44ml (Add ethanol 176 ml)
MaxBinder™ Solution	5ml	10ml	20ml
Nanozyme Mix	2ea (Dry)	4ea (Dry)	8ea (Dry)
Cell Resuspension Solution	3ml	6ml	12ml
Proteinase K (10mg/ml)	0.5ml (Dry)	0.5ml (Dry) x 2ea	0.5ml (Dry) x 4ea
Column Set (with cap, 50ea/ Blue Box)	1box	2box	4box
Instructions for Use	1ea	1ea	1ea

Description

PureHelix™ Genomic DNA Prep Kit [Rice] is designed for rapid and pure isolation of total DNA from rice. The spin column based method completely removes PCR inhibitors such as divalent cations and proteins resulting in a high purity preparation of genomic DNA. There is no use of phenol or chloroform, handling is safe and does not produce any harmful waste. DNA purified with this kit is suitable for a variety of applications, including PCR amplification, digestion with restriction endonucleases and membrane hybridizations.

Storage



Shelf life



Quality Control

Each lot of **PureHelix™ Genomic DNA Prep Kit [Rice]** was tested against predetermined specifications to ensure consistent product quality.

Protocol

Important things to do before starting

- Before using **WB**, add **absolute ethanol** according to the bottle label to obtain a working solution. You may use 80% ethanol, instead of WB. Ethanol does not supplied in this kit.
- Add **0.5 ml** of distilled water to the provided **Proteinase K** tube for making **10mg/ml** concentration, and then store at -20°C.
- **Add 1.5 ml** of **Cell Resuspension Solution** to provided **NanoZyme mix tube**, and then Store at -20°C.

1. Cell Lysis

- 1) Add **300 µl of NGD1** and **50 µl of Nanozyme Mix** to ground single grain of rice in a 1.5ml microcentrifuge tube. Vortex vigorously for 30-60 sec.
- 2) Add **8 µl of Proteinase K (10mg/ml)** and mix by pipetting.
Incubate at 60°C for 1 hour, and then cool the tube on ice for 5 min.
- 3) Add **300 µl of NPS2**. Vortex briefly.
- 4) Place the tube on ice for 5 min, and centrifuge for 5 min at 12,000 rpm.
※ **The precipitate will be a tight pellet. If the pellet is not tight, repeat this step.**
- 5) Transfer **600 µl of the supernatant** into a clean 1.5ml tube.
- 6) Add **100 µl of absolute ethanol** and vortex vigorously.

2. Column Activation [Optional]

※ **These steps are required for the best yield.**

- 1) Place a Spin Column into a 2 ml collection tube.
- 2) Add **100 µl of MaxBinder™ Solution** into the Spin Column.
- 3) Centrifuge at 12,000 rpm for 30 sec and immediately proceed to next step.
You need not discard the flow-through from the collection tube.

3. Loading

- 1) Pipet **400 µl of the mixture** from step 1 (Cell Lysis) into a spin column sitting in a 2 ml collection tube.
- 2) Centrifuge for 30 sec at 12,000 rpm. Discard the flow-through.
- 3) Pipet **the remains of the mixture** from step 1 into the spin column.
- 4) Centrifuge for 30 sec at 12,000 rpm. Discard the flow-through.

4. Washing

- 1) Add **500 µl of WB (80% ethanol)** into the spin column.
- 2) Centrifuge for 30 sec at 12,000 rpm. Discard the flow-through.
※ Repeat these steps for high-purity DNA preparation.
- 3) Centrifuge at 12,000 rpm for 2 min to remove residual ethanol.

5. Elution

- 1) Discard the 2 ml collection tube and place the spin column in a clean 1.5 ml tube.
Add **40-50 µl of EB solution or distilled water** into the center of the column.
- 2) Centrifuge at 12,000 rpm for 2 min. Discard the spin column.
Store the eluted DNA at 4°C or -20°C.

Products

Cat. No.	Products	Size
GCTR50	PureHelix™ Genomic DNA Prep Kit [Rice] (Ver. 3.0) Column type	50preps
GCTR100	PureHelix™ Genomic DNA Prep Kit [Rice] (Ver. 3.0) Column type	100preps
GCTR200	PureHelix™ Genomic DNA Prep Kit [Rice] (Ver. 3.0) Column type	200preps