

HELINI™
Purefast
Soil Nucleic acid
Purification Buffers

Cat. No: 2012 - 25/50/100 Purifications

Handbook

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Kit components

No. of reactions	25	50
Buffer-A	10ml	20ml
Buffer-B	5ml	10ml
Buffer-C	5ml	10ml
Buffer-D	12.5	25ml
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Storage : in room temperature.

Technical Assistance

For technical assistance and more information, please contact;
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Safety Information

When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. Discard sample and assay waste according to your local safety regulations.

Introduction

HELINI Purefast Soil buffers are designed for rapid and cost-effective small-scale preparation of high quality viral nucleic acid from Soil. It effectively removes humic substance and inhibitors from the soil. The supernatant can be directly used with spin column purification kit. Isolated Viral nucleic acid can be used directly in RT-PCR/PCR.

Material Required

- Micro Pipettes Variable Volume 0.5-10 μ l, 10-100 μ l, and 100-1000 μ l
- Sterile pipette tips with aerosol barrier 2-20 μ l, 10-100 μ l, and 100-1000 μ l
- Disposable powder-free gloves
- Vortex mixer / Water bath
- Centrifuge with rotor for 1.5 ml reaction tubes
- 1.5ml/2ml centrifuge tubes

Important Notes:

All purification steps should be carried out at room temperature.

All centrifugations should be carried out in a table-top micro-centrifuge at $>12000 \times g$ (12000-14000 rpm, depending on the rotor type).

Procedure:

1. Transfer 100 to 200mg of dry soil sample into 2ml centrifuge tube.
2. Add 300 μ l of Buffer-A and mix well by vortexing for 2 min.
3. Add 200 μ l of Buffer-B and mix well by vortexing for 2 min.
4. Add 150 μ l of Buffer-C and mix well by vortexing for 2 min.
5. Add 500 μ l of Buffer-D and mix well by vortexing for 2 min.
6. Centrifuge at 10000rpm for 2min.
7. Transfer 200 μ l of clear supernatant into fresh 1.5ml centrifuge tube.
8. Add 200 μ l of Shrimp viral nucleic acid Binding buffer and continue the protocol.

Recommendation for Real-time PCR:

Use 5 - 20 μ l of elute

Manufactured and Marketed by

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