MBead Buffy Coat Genomic DNA Kit

Cat No. SN008-0100 Size: 100 Reactions
Cat No. SN008-0004 Size: 4 Reactions
Sample: Up to 300 µl of the buffy coat
Format: Magnetic Bead System
Operation time: 10-15 minutes
Release volume: 200 µl

Description

This magnetic bead genomic DNA purification kit was specifically designed to isolate the genomic DNA from the Buffy Coat. Its unique buffer system will efficiently lyse cells and degrade proteins, allowing for the DNA to be easily bound by the surface of the magnetic beads. The RNA and other non-specific binding particles are removed with a wash buffer, and the genomic DNA is then released into the Release Buffer. The genomic DNA can be purified manually within 10-15 minutes (using most magnetic separators) or the kit can be easily adapted to satisfy most automated nucleic acid purification systems.

Features

- Fast, reproducible, and easy processing using a magnetic bead system.
- Isolate high quality genomic DNA.
- Recovered genomic DNA is compatible with various downstream applications.

Applications

- Restriction Enzyme Digestion.
- PCR amplification.
- Southern Blotting.
- Real-Time assay.

Kit Contents

Contents	SN008-0100	SN008-0004
Magnetic Bead	2 ml X 1 vial	80 µl X 1 vial
Lysis Buffer	30 ml X 1 bottle	1.5 ml X 1 vial
Wash Buffer	80 ml X 1 bottle	2 ml X 2 vial
Release Buffer	20 ml X 1 bottle	1 ml X 1 vial

Quality Control

The quality of the MBead Buffy Coat Genomic DNA Kit is tested on a lot-to-lot basis to ensure consistent product quality.

Required Materials

- Absolute ethanol
- 1.5 ml microcentrifuge tubes
- Magnetic separator
- Water bath / Dry bath

MBead Buffy Coat Genomic DNA Kit Protocol

Step 1 Lysis

1. Transfer the Buffy Coat (up to 300 µl) into a 1.5 ml microcentrifuge tube and add 300 µl of the Lysis Buffer.
2. Mix well and incubate at 65°C for 5 minutes. During this time, pre-heat the Release Buffer to 65°C for the Step 4.
3. Add 300 µl of the absolute ethanol to the lysate and mix well.

Step 2 DNA Binding

1. Add 20 µl of Magnetic Beads. Mix well by gently shaking for 3 minutes.
2. Place the tube in a magnetic separator for 30 seconds.
3. Remove the solution (If the mixture becomes viscous, increase magnetic bead separation time)

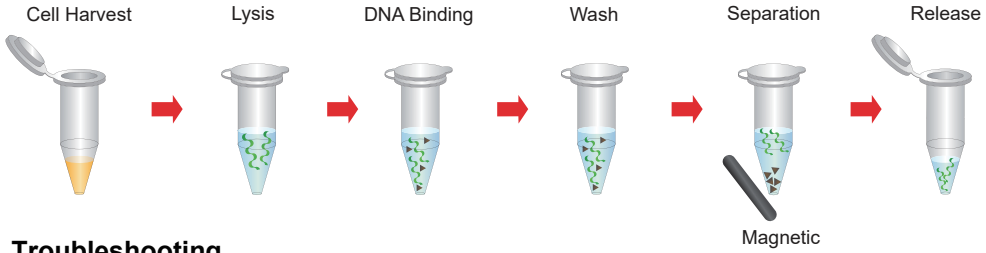
Step 3 Wash

1. Add 800 µl of the Wash Buffer and mix well (Following the wash, the mixture will no longer be viscous).
2. Place the tube in a magnetic separator for 30 seconds. Remove the solution.

Step 4 Release

1. Add 200 µl of the Release Buffer (pre-heated to 65°C) and mix well.
2. Incubate for 3 minutes at 65°C (during the incubation, shake the tube vigorously every minute).
3. Place the tube in a magnetic separator for 1 minute.
4. Carefully transfer ONLY the clean portion of the solution to a clean tube.

NOTE: Be sure and allow the magnetic beads to disperse completely during the binding, wash and elution steps.



Troubleshooting

Refer to the table below to troubleshoot problems that you may encounter when you did genomic DNA isolation with the kit.

Problem	Cause	Solution
DNA is sheared or degraded	Lysate mixed too vigorously	Use the appropriate pipette tip set for the required volume, lower it under the reading line of the solution to mix the sample, pipet up and down gently to mix.
	DNases contamination	Maintain a sterile environment while working (e.g. wear gloves and use DNase-free reagents).
RNA containment	Incomplete removal of the RNase	RNase A treatment.
Low yields of genomic DNA	Incomplete lysis and homogenization	Complete lysis. Use the appropriate method for the lysate preparation based on the amount of starting materials.
	Incorrect handling of Magnetic Beads	Vortex the tube containing the Magnetic Beads to fully resuspend the beads before adding them to your sample.
	Incorrect elution conditions	Add the Release Buffer (50-100µl) and incubate for 3 min at 65°C.
	The quality of the starting material may not be optimal	Use fresh sample and process immediately after collection, or freeze the sample at -80°C or in the liquid nitrogen.
High background on UV measurement	Residual beads released	Repeat the magnetic separation and transfer the eluate to a clean tube.

Related Ordering Information

Cat. No.	Description	Size
SM101-0500	*Taq* DNA polymerase	500 U
SM200-0100	PCR SUPERMIX	100 Reactions
SM201-0100	Hot Start SUPERMIX	100 Reactions
SM255-0100	Ultrapure Proteinase K	100 mg
SA001-0500	AGAROSE Tablet, 0.5g	100 Tablets
SL001-1000	Novel Juice Supplied in 6X Loading Buffer	1 ml
SD003-R600	100 bp DNA Ladder H3 RTU	600 µl
SD010-R600	1 Kb DNA Ladder RTU	600 µl
SD013-R600	XLarge DNA Ladder RTU	600 µl
ST040-4000	100 mM dNTP Set	4 x 1 ml
ST046-1000	100 mM dNTP Set	4 x 250 µl
ST025-1000	2.5 mM dNTP Mix	1 ml
ST010-1000	10 mM dNTP Mix	1 ml

Caution

- Check buffers before use for salt precipitation. Re-dissolve any precipitate by warming up to 37°C.
- During operation, always wear a lab coat, disposable gloves, and protective equipment.
- All products are for research use only.