



"WE MAKE NGS BETTER"

HighPrep PCR

Efficient cleanup for post PCR and fragment size-selection for NGS library construction

Catalog Nos. AC-60001, AC-60005, AC-60050, AC-60250, AC-60500
Manual Revision 2.2
WI-72-08

- Magnetic bead-based chemistry
- No centrifugation or filtration
- Efficient cleanup
- Precise size-selection
- Efficient recovery of DNA fragments ≥ 100 bp

Protocol

Contents

Product Description and Process 1

Product Specifications 1

Storage and Preparation, Materials Supplied 2

Equipment and Reagents to Be Supplied by the User 2

PCR cleanup reaction using a 96-well plate 2

PCR cleanup reaction using a 384-well plate 3

Ordering and Related Product Information 5

For Research Use Only. Not for use in diagnostic procedures.

Information in this document is subject to change without notice.

MAGBIO GENOMICS, INC. DISCLAIMS ALL WARRANTIES WITH RESPECT TO THIS DOCUMENT, EXPRESSED OR IMPLIED, INCLUDING BUT NOT LIMITED TO THOSE OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. TO THE FULLEST EXTENT ALLOWED BY LAW, IN NO EVENT SHALL MAGBIO GENOMICS, INC. BE LIABLE, WHETHER IN CONTRACT, TORT, WARRANTY, OR UNDER ANY STATUTE OR ON ANY OTHER BASIS FOR SPECIAL, INCIDENTAL, INDIRECT, PUNITIVE, MULTIPLE OR CONSEQUENTIAL DAMAGES IN CONNECTION WITH OR ARISING FROM THIS DOCUMENT, INCLUDING BUT NOT LIMITED TO THE USE THEREOF, WHETHER OR NOT FORESEEABLE AND WHETHER OR NOT MAGBIO GENOMICS, INC. IS ADVISED OF THE POSSIBILITY OF SUCH DAMAGES.

TRADEMARKS

The trademarks mentioned herein are the property of MagBio Genomics, Inc. or their respective owners.

Product Description

HighPrep PCR is a paramagnetic bead-based reagent for amplicon purification and DNA size selection in NGS library preparation workflows. It selectively binds DNA fragments to magnetic beads while removing salts, primers, primer-dimers, and dNTPs. At a 1.8× bead ratio, the reagent provides reliable recovery of DNA fragments ≥100 bp. The purified DNA is then eluted in water or a low-salt buffer, ready for direct use in downstream applications.

This protocol can be used for manual procedures as well as a guideline for adapting the kit to automated liquid handling instruments. For availability of ready-to-run scripts, please contact MagBio Genomics.

Amplicons purified with the HighPrep PCR are ready to be used in the following applications:

- PCR
- Mutation detection and genotyping
- Next generation sequencing
- Microarrays
- Restriction enzyme digests
- Cloning

Process

HighPrep PCR uses a simple 3 steps procedure: Bind-Wash-Elute. HighPrep PCR is added to the PCR reaction sample. The protocol utilizes a magnet plate (magnet stand) for processing the PCR reaction sample. During the process, contaminants and salts are washed off and pure DNA is eluted, ready to be used in subsequent applications.

Product Specifications

Product Number	Description	Number of Reactions	Storage Conditions
AC-60001	HighPrep PCR (1 mL) (Sample size)	55	2-8°C DO NOT FREEZE
AC-60005	HighPrep PCR (5 mL)	277	
AC-60050	HighPrep PCR (50 mL)	2,777	
AC-60250	HighPrep PCR (250 mL)	13,888	
AC-60500	HighPrep PCR (500 mL)	27,777	

Number of reactions is based on typical 10 µL PCR reaction volume.
Volume of HighPrep PCR per reaction = 1.8 x (PCR Reaction Volume)

Storage and Preparation

- Store at 2-8°C. **DO NOT FREEZE.** HighPrep PCR is stable for 18 months when stored at 2-8°C.
- Keep at room temperature for 30 minutes prior to use.
- Thoroughly shake the HighPrep PCR to resuspend the beads before use.

Materials Supplied in the Kit

- HighPrep PCR

Equipment and Reagents to Be Supplied by the User:

- 80% Ethanol (Prepared from non denatured Ethanol)
- Reagent grade water, 10mM TRIS-HCL pH 8.0, or TE Buffer (Low TE, <0.1mM EDTA)
- Magnetic separation device compatible with 96 or 384 well PCR plate (see page 5)
- 96-well PCR plate or 384-well PCR plate
- Multichannel pipette

HighPrep PCR: PCR cleanup reaction using a 96-well plate

 Bring the **HighPrep PCR** to room temperature for at least 30 minutes before use.

1. Shake thoroughly the **HighPrep PCR** to fully resuspend the magnetic beads.
2. Transfer PCR reaction to appropriate 96-well plate. Follow the table in step 3 for the desired sample processing volume.
3. Add **HighPrep PCR** volume according to the PCR reaction.
See table below to determine appropriate volume.

PCR Reaction Volume (µL)	HighPrep PCR Volume at 1.8X (µL)*
10	18
20	36
50	90

* Formula used to calculate the volume of HighPrep PCR needed for PCR reaction:
HighPrep PCR volume per reaction = 1.8 X PCR reaction volume.

4. Mix the **HighPrep PCR** and PCR sample by pipetting up and down 6-8 times.
5. Incubate the mixture for 5 minutes at room temperature.

6. Place the sample plate on the 96 magnetic separation device for 3 minutes, or until the solution clears. Beads will pull to the side of the well.
7. With the sample plate still on the magnet, remove and discard the supernatant by pipetting.
 ⚠ *Do not disturb the attracted beads while aspirating the supernatant.*
8. With the sample plate on the magnet, add 200 μL of 80% Ethanol to each well and incubate for 30 seconds at room temperature.
9. With the plate still on the magnet, remove and discard the supernatant by pipetting.
10. Repeat steps 8-9 for a total of two 80% Ethanol washes.
11. Dry the beads by incubating the plate for ~ 5 minutes at room temperature with the plate still on the magnetic separation device.
 ⚠ *It is critical to completely remove all traces of alcohol but take caution in not over drying the beads as this will reduce the yield.*
12. Remove the sample plate from the magnetic separation device. Add 40 μL of elution buffer (reagent grade water, TRIS-HCl pH 8.0 or TE buffer) to each well and pipette up and down 5 times to mix. Prewarming the elution buffer to 55°C can increase the yield.
13. Incubate for 2 minutes at room temperature.
14. Place the sample plate back on the magnetic separation device and wait 3 minutes, or until the magnetic beads clear from the solution.
15. Transfer the eluate (cleared supernatant) to a new plate for subsequent applications or store at -20°C for later use

HighPrep PCR: PCR cleanup reaction using a 384-well plate

⚠ Bring the **HighPrep PCR** to room temperature for at least 30 minutes before use.

1. Shake thoroughly the **HighPrep PCR** to fully resuspend the magnetic beads.
2. Transfer PCR reaction to appropriate 384-well plate. Follow the table in step 3 for the desired sample processing volume.
3. Add **HighPrep PCR** volume according to the PCR reaction. See table below to determine appropriate volume.

PCR Reaction Volume (μL)	HighPrep PCR Volume at 1.8X (μL)*
5	9
7	12.6

* Formula used to calculate the volume of HighPrep PCR needed for PCR reaction:
 HighPrep PCR volume per reaction = 1.8 X PCR reaction volume.

4. Mix the **HighPrep PCR** and PCR sample by pipetting up and down 6-8 times.
5. Incubate the mixture for 5 minutes at room temperature.
6. Place the sample plate on the 384 magnetic separation device for 2 minutes, or until the solution clears. Beads will pull to the side of the well.
7. With the sample plate still on the magnet, remove and discard the supernatant by pipetting.
⚠ Do not disturb the attracted beads while aspirating the supernatant.
8. With the sample plate on the magnet, add 30 μL of 80% Ethanol to each well and incubate for 30 seconds at room temperature.
9. With the plate still on the magnet, remove and discard the supernatant by pipetting.
10. Repeat steps 8-9 for a total of two 80% Ethanol washes.
11. Dry the beads by incubating the plate for 3-5 minutes at room temperature with the plate still on the magnetic separation device.
⚠ It is critical to completely remove all traces of alcohol but take caution in not over drying the beads as this will reduce the yield.
12. Remove the sample plate from the magnetic separation device. Add 30 μL of elution buffer (reagent grade water, TRIS-HCl pH 8.0 or TE buffer) to each well and pipette up and down 5 times to mix. Prewarming the elution buffer to 55°C can increase the yield.
13. Incubate for 2 minutes at room temperature.
14. Place the sample plate back on the magnetic separation device and wait 2 minutes, or until the magnetic beads clear from the solution.
15. Transfer the eluate (cleared supernatant) to a new plate for subsequent applications or store at -20°C for later use.

Custom DNA Size-selection Protocols for NGS Applications

To obtain a custom protocol for DNA size-selection of a specific fragment size, contact:

US/Canada/Worldwide: support@magbiogenomics.com

Europe: info.europe@magbiogenomics.com

Ordering

HighPrep PCR

Catalog No.	Product
AC-60005	HighPrep PCR (5 mL)
AC-60050	HighPrep PCR (50 mL)
AC-60250	HighPrep PCR (250 mL)
AC-60500	HighPrep PCR (500 mL)

Related Products

MagQuant Plus DNA Kit V2

Catalog No.	Product	Description	Preps
MQP-50096	MagQuant Plus DNA Kit V2 (96 Preps)	Magnetic bead-based kit for normalization of DNA concentration, and quantitation of DNA for NGS and other applications.	96
MQP-50384	MagQuant Plus DNA Kit V2 (384 Preps)		384
MQP-51920	MagQuant Plus DNA Kit V2 (1,920 Preps)		1,920

HighPrep Blood & Tissue DNA Kit

Catalog No.	Product	Description	Preps
HPBTS-D96	HighPrep Blood & Tissue DNA Kit (96 preps)	Magnetic bead-based kit for Genomic DNA isolation from 20-250 μ L of blood, lysate of tissues, mouse tails, cultured cells, or buccal swabs	96
HPBTS-D96x4	HighPrep Blood & Tissue DNA Kit (384 preps)		384

Magnetic Separation Devices

Catalog No.	Description
MYMAG-96X	Magnetic Separation Device (96 well ring format)
MBMS-12	MagStrip Magnet Stand (1.5 mL x 12)
MBMS-31550	15 mL and 50 mL Magnetic Stand Combo (3 x 15 mL and 3 x 50 mL)



www.magbiogenomics.com