6



Jul 07, 2020

Ampure Bead Cleanup

Alex Shaw¹, Manasi Majumdar², Catherine Troman¹, Javier Martin², Nick Grassly¹

¹Imperial College London; ²National Institute for Biological Standards and Control

1 Works for me This protocol is published without a DOI.

Poliovirus Sequencing Consortium

Alex Shaw

PROTOCOL CITATION

Alex Shaw, Manasi Majumdar, Catherine Troman, Javier Martin, Nick Grassly 2020. Ampure Bead Cleanup. **protocols.io**

https://protocols.io/view/ampure-bead-cleanup-bf85jry6

LICENSE

This is an open access protocol distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

CREATED

May 11, 2020

LAST MODIFIED

Jul 07, 2020

PROTOCOL INTEGER ID

36861

- 1 Prepare the AMPure XP beads for use; resuspend by vortexing.
- 2 Add required volume of resuspended AMPure XP beads to the reaction and mix by pipetting.
- 3 Incubate on a rotator for 5 minutes at room temperature.
- 4 Spin down the sample and pellet on a magnet. Keep the tube on the magnet, and pipette off the supernatant.
- 5 Keep on magnet, wash beads with 100 μL of freshly prepared 70% ethanol without disturbing the pellet.
- 6 Remove the 70% ethanol using a pipette and discard.

Repeat steps 5 and 6.

7	
8	Spin down and place the tube back on the magnet.
9	Pipette off any residual 70% ethanol.
10	Briefly allow to dry.
11	Remove the tube from the magnetic rack and resuspend pellet in elutant.
12	Incubate for 2 minutes at RT.
13	Pellet beads on magnet until the eluate is clear and colourless.
14	Remove desired volume of eluate and store in a clean tube. Avoid disturbing the pelleted beads.