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Amplicon clean-up using SPRI beads

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1 Works for me dx.doi.org/10.17504/protocols.io.7nxhmfn

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STEPS MATERIALS

NAME	CATALOG #	VENDOR
Agencourt AMPure XP	A63880	Beckman Coulter
Elution Buffer (EB)	19086	Qiagen
QuantiFluor(R) ONE dsDNA System, 100rxn	E4871	Promega

1 ⚠

Vortex SPRI beads thoroughly to ensure they are well resuspended, the solution should be a homogenous brown colour.

Agencourt AMPure XP
by Beckman Coulter
Catalog #: A63880

2 Add an equal volume (1:1) of SPRI beads to the sample tube and mix gently by either flicking or pipetting. For example add **50 µl** SPRI beads to a **50 µl** reaction.

3 Pulse centrifuge to collect all liquid at the bottom of the tube.

4 Incubate for **00:05:00** at room temperature.

5 Place on magnetic rack and incubate for **00:02:00** or until the beads have pelleted and the supernatant is completely clear.

6 Carefully remove and discard the supernatant, being careful not to touch the bead pellet.

7 Add **200 µl** of room-temperature [M]**70 % volume** ethanol to the pellet.

8 Carefully remove and discard ethanol, being careful not to touch the bead pellet.

9 **go to step #7** and repeat ethanol wash.

10 Pulse centrifuge to collect all liquid at the bottom of the tube and carefully remove as much residual ethanol as possible using a P10 pipette.

11 With the tube lid open incubate for **00:01:00** or until the pellet loses it's shine (if the pellet dries completely it will crack and become difficult to resuspend).

12 Resuspend pellet in **30 µl** Elution Buffer (EB), mix gently by either flicking or pipetting and incubate for **00:02:00** .

Elution Buffer (EB)
by Qiagen
Catalog #: 19086

13 Place on magnet and transfer sample to a clean 1.5mL Eppendorf tube ensuring no beads are transferred into this tube.

14 Quantify **1 µl** product using the Quantus Fluorometer using the ONE dsDNA assay.

QuantiFluor(R) ONE dsDNA System, 100rxn
by Promega
Catalog #: E4871

Quantus
Fluorometer
Promega E6150

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