Preparing SPRI beads

This document describes a method for preparing SPRI beads, also known as Ampure (Beckman). There is some great info related to this protocol to be found here.

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Reagents and materials

- 1M Tris-Cl, pH 7.5 (Thermo Scientific, CAT#15567027)
- carboxylate-modified beads (Thermo Scientific, CAT#65152105050250, CAT#45152105050250)
- PEG-8000 (Promega, CAT#V3011)
- 0.5M EDTA (Thermo Scientific, CAT#15575020)
- 5M NaCl (Thermo Scientific, CAT#AM9760G)
- Tween 20 (Sigma, CAT#P9416)
- RNA clean water (Thermo Fisher, CAT#10977023)
- 1.5mL or 2.0mL Safe-Lock tubes (Fisher Scientific, CAT#05-402-25 or CAT#05-402-7)
- 50mL tubes (VWR, CAT#89093-190)

Solution recipes

- 50% (w/v) PEG-8000 12.5g in 25mL of water (see **Note 1**)
- 10% (v/v) Tween 20 1mL in 10mL of water
- TE solution (make 50mL)
 - 10mM Tris-Cl, pH 7.5 (500uL from 1M stock)
 - 1mM EDTA (100uL from 0.5M stock)

Protocol

- 1. Vortex mix SeraMag beads and transfer 1mL of each to a 2mL microtube.
- 2. Place tube on a magnetic rack and discard the supernatant after the beads settle.
- 3. Resuspend beads in 1mL of TE solution and pipette mix. Place back on the magnetic rack and discard the supernatant.
- 4. Repeat the previous step two additional times for a total of 3 rinses.
- 5. Resuspend the beads in 1mL TE solution of TE and set aside.
- 6. In a fresh 50mL tube, combine:
 - 1. 25mL of 5M NaCl

- $2.\,$ 500uL of 1M Tris-Cl, pH $7.5\,$
- $3.\ 500\mathrm{uL}$ of $0.5\mathrm{M}$ EDTA
- 4. 2.75 mL of water
- 7. Add the beads in TE solution to the prepared 50mL tube. Vortex mix.
- 8. Slowly add 20mL of 50% (w/v) PEG-8000 solution.
- 9. Add 250uL of 10% (v/v) Tween 20 solution to the 50mL tube and mix gently by inversion until the solution appears homogeneous.
- 10. Wrap the tube in tinfoil and store at +4C.

Notes

Note 1 - This PEG solution will be very viscous and make take a long time to dissolve completely. I suggest leaving it on a rotary mixer and walking away for a couple of hours.