

ANCHOR

SpeedPURE® Multi Kit (96) (RUO)



1. Turn on the ANCHOR BEAD 32 and log in
2. Invert each Multi Plate 3 times to resuspend the beads
3. Gently tap each Multi Plate 3 times on desk to collect the reagents at the bottom
4. Remove the seal of each Multi Plate
5. Add following reagents to the same well (lysis buffer, blue) (Fig. 1)
 - 8 μ L Internal Control (IC, if ANCHOR PCR Kits are used)
 - 20 μ L Proteinase K (PK)
6. Add 200 μ L (or 400 μ L, depending on protocol) of sample to first well (lysis buffer, blue)
7. Open the ANCHOR BEAD 32 door and place Multi Plates into the instrument (Fig. 2)
Ensure the Multi Plates are securely positioned with the plate label facing the front of the instrument
8. Insert the SpeedPURE® Magnetic Rod Covers – always use two rod covers per plate (two or four in total) to ensure proper protection of the magnetic rods (Fig. 3)
9. Ensure all magnetic rod covers click into place and are inserted firmly

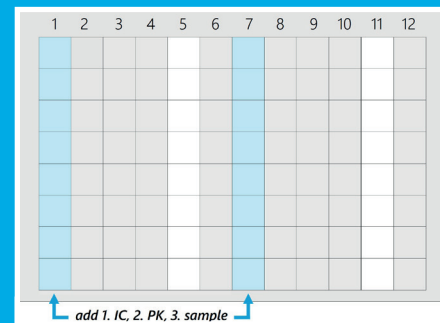


Fig. 1





Fig. 2



Fig. 3

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- On the display select  (upper left corner)
- Choose the desired protocol (Fig. 4)
 - SpeedPURE_200 for 200 µL sample input
 - SpeedPURE_400 for 400 µL sample input
- Select  (lower right corner) to start the sample preparation
- After the run is complete:
 - Remove and discard magnetic rod covers
 - Remove Multi Plates from instrument and collect eluates from columns 6 and 12 (Fig. 5)
 - Discard used Multi Plates

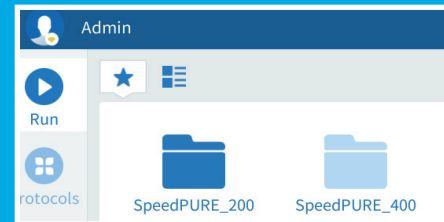
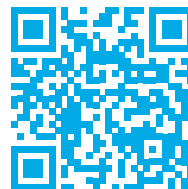


Fig. 4

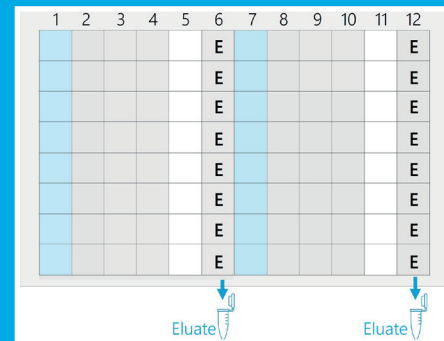


Fig. 5

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