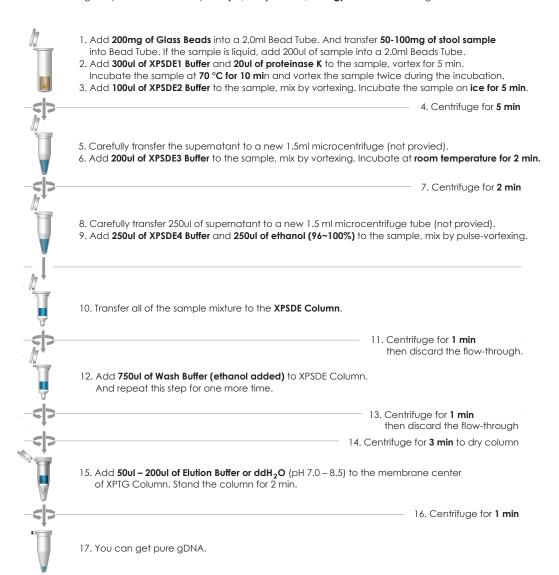
eCube Stool DNA Mini Kit

* Things to do before starting

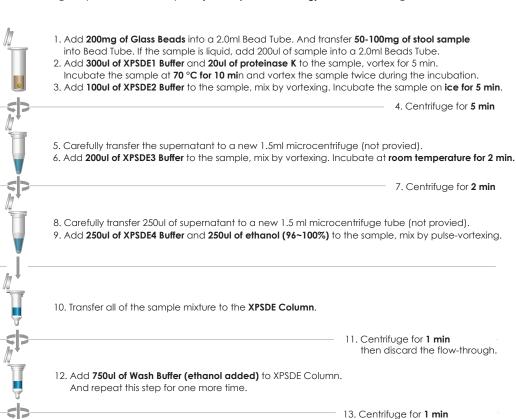
- 1. Add 1.1 ml ddH₂O to each Proteinase K tube to make a 10mg/ml stock solution. Store the stock solution at 2-8°C.
- 2. For 50preps, add 80ml ethanol (96-100%) to Wash Buffer when first open. For 200preps, add 320ml ethanol (96-100%) to Wash Buffer when first open.
- 3. Prepare a water baths to 70 °C before the operation.
- 4. All centrifuge steps are done at full speed (14,000 rpm or 10,000 x g) in a microcentrifuge.



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15. Add **50ul – 200ul of Elution Buffer or ddH_2O** (pH 7.0 – 8.5) to the membrane center of XPTG Column. Stand the column for 2 min.

16. Centrifuge for 1 min

17. You can get pure gDNA.



then discard the flow-through

14. Centrifuge for 3 min to dry column