

KitVANCE DURA DNA EXTRACTION FROM Animal Tissue

Designed for isolating high-quality genomic DNA from Animal Tissue samples

KIT COMPONENTS AND DESCRIPTION

The kit comes in a compact box with labeled vials/bottles. Each kit contains enough reagent for ~50 extractions. Components are divided into consumables (used per extraction) and reusable items. Protocol can be carried out at room temperature.

EQUIPMENT AND REAGENTS SUPPLIED

- L Buffer (10 ml vial)
- Proteinase K (20mg/ml)
- B Buffer (10 mL vial)
- WB1 (25 mL vial)
- WB2 (25 mL vial)
- EB (10 ml)
- Spin Column 1.5 ml (50 unit)
- Collection tubes 1.5 ml (50 units)

PROCEDURE

Total Time: 45 Minutes | Format: Spin Column

- 1- Grind 50 mg sample then Add 200 μ L of L Buffer the tube, add 20 μ L Proteinase K Incubate at 56°C for 20 minutes. Mixing by vortex for 5 seconds
- 2- Add 200 μ L Universal Binding Buffer and 200 μ L Ethanol (96–100%) or **Isopropanol**. Vortex for 5 seconds.
- 3- Transfer the entire mixture (~650 μ L) into the Silica Spin Column. Centrifuge at 10,000 rpm for 1 minute. Discard the flow-through.
- 4- Add 500 μ L Wash Buffer 1. Centrifuge at 10,000 rpm for 1 minute. Discard the flow-through.
- 5- Add 500 μ L Wash Buffer 2. Centrifuge at 10,000 rpm for 1 minute. Discard the flow-through.
- 6- Place the column back into the collection tube. Centrifuge at Max Speed for 1 minutes.

 **Critical:** This step removes residual ethanol. Do not skip, or your DNA may fail in PCR!

- 7- Transfer: Place the dry column into a new 1.5 mL tube (provided).
- 8- Add ~25 μ L Elution Buffer directly to the center of the column membrane.
- 9- For higher yield, use Elution Buffer pre-warmed to 70°C.

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10- Centrifuge at Maximum Speed for 1 minute.

Storage: The purified DNA is now ready for use. Store at 4°C for immediate use or -20°C for long-term storage.



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