## **Quick Start Guide**

# GenElute™ Mammalian Genomic DNA Kit

## G1N10, G1N70, G1N350

## Reagents to Prepare

- Dilute the Wash Solution Concentrate with 10 mL (10 preps), 80 mL (70 preps) or 360 mL (350 preps) of 95–100% ethanol. After each use, tightly cap the diluted Wash Solution to prevent ethanol evaporation.
- Dissolve one bottle of Proteinase K in water to obtain a 20 mg/mL stock solution. This solution can be stored for several days at 2–8 °C or for long term storage, aliquots may be stored at –20 °C until needed.

#### **Protocol**

All spins at  $\geq$  6,500 x g unless otherwise noted.

#### **Release DNA from Cells**

- 1. Cultured Cells
  - a. Pellet up to  $5 \times 10^6$  cells by spinning at 300 x g. Discard medium.
  - b. Resuspend cells in 200  $\mu$ L Resuspension Solution. **Optional:** Add 20  $\mu$ L RNase A. Mix & incubate at room temperature for 2 min.
  - c. Add 20 µL Proteinase K, followed by 200 µL Lysis Solution C. Vortex throughly. Incubate at 70°C for 10 min.
  - d. Continue to Bind DNA to Column.

#### **Release DNA from Tissues**

- 1. Mammalian Tissues
  - a. Mince up 25 mg tissue on ice. Transfer to microcentrifuge tube.
  - b. Add 180  $\mu$ L Lysis Solution T to tissue & 20  $\mu$ L Proteinase K. Vortex or invert to mix. Incubate at 55°C until fully digested (2-4 h). **Optional:** After digestion add 20  $\mu$ L RNase A. Mix and incubate at room temperature for 2 min.
  - c. Continue to Bind DNA to Column.

#### **Bind DNA to Column**

- 2. Column Preparation: Add 500 uL Column Preparation Solution to assembled column in collection tube and spin at  $12,000 \times g$  for 1 minute. Discard flowthrough.
- 3. Add 200  $\mu$ L ethanol to cell lysate. Vortex or invert to mix.
- 4. Transfer to column and spin for 1 min. Discard collection tube with flowthrough.

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#### Wash to remove contaminants

- 5. Transfer column to new collection tube. Add 500  $\mu L$  diluted Wash Solution to column. Spin 1 min and discard flowthrough. Repeat once.
- 6. Spin 3 min to completely dry the column.

### **Elute purified DNA**

- 7. Transfer column to new collection tube.
- 8. Add 200  $\mu L$  Elution Solution to column. Spin 1 min. Repeat elution in the same or new tube if desired.

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