



DNAPure Reagent (Genomic DNA Isolation Reagent)

For research use only

DNAPure Reagent 20ml	Cat no: YT9091
DNAPure Reagent 50ml	Cat no: YT9092
DNAPure Reagent 100ml	Cat no: YT9093

DNAPure Reagent is a complete and ready-to-use reagent for the isolation of genomic DNA from tissue and liquid samples. The DNAPure Reagent is a novel guanidine-detergent lysing solution which permits selective precipitation of DNA from a cell lysate. The procedure can be completed in 30-40 minutes with a genomic DNA recovery of 70 - 100%. The isolated DNA can be used, without additional purification, for Southern analysis, dot blot hybridization, molecular cloning, PCR and other molecular biology and biotechnology applications.

Reagents required, but not supplied: 100% ethanol – 75% ethanol – Nuclease free water

Cell Lysis :

- 1- Cells grown in monolayer: Add 0.75-1.0 ml of DNAPure Reagent per 10 cm² culture plate area.
- 2- Add 1 ml of DNAPure Reagent to $1-3 \times 10^7$ cells, either in pellet or in suspension (volume < 0.1 ml)
- 3- Homogenize 10-40 mg tissue in 1 ml of DNAPure Reagent

PROTOCOL

- 1- **LYSIS \ HOMOGENIZATION:** add 1 ml DNAPure reagent to 10-40 mg tissue, 10^7 cells or 0.1 ml liquid sample.
- 2- **INCUBATION:**
 - I. Incubate for 10 minutes at room temperature
 - II. Incubate on ice for 10 minutes
- 3- **CENTRIFUGATION:** Sediment the homogenate for 3 min at $10,000 \times g$ at room temperature. transfer the resulting viscous supernatant to a fresh tube.
- 4- **DNA Precipitation:** add 1ml of cold 100% ethanol of the same volume (add 1ml ethanol per 1 ml of DNAPure Reagent used for the isolation). Invert several times to mix well, DNA should quickly become visible as a cloudy precipitate.
- 5- **CENTRIFUGATION:** centrifuge at $15000 \times g$ for 2-3 min at room temperature to pellet the DNA and discard the supernatant.
- 6- **DNA WASH:**
 - I. Wash the DNA precipitate in 1ml ethanol 70% .Centrifuge at 15,000g for 1-2 minute and Discard the supernatant
 - II. Wash the DNA precipitate in 0.5ml ethanol 70%. Centrifuge at 15,000g for 1-2 minute and Discard the supernatant
- 7- **DNA SOLUBILIZATION:** Air dry the DNA by storing in an open tube for 1-15 minutes after removing the ethanol. (If the DNA pellet overdried, it will be much more difficult to dissolve.) dissolve the pellet in 50-100µl nuclease free water or 10mM Tris-HCL .