

Jurkat Genomic DNA



1-800-632-7799
info@neb.com
www.neb.com



N4001S 005140716071

N4001S

15 µg Lot: 0051407 Exp: 7/16

100 µg/ml Store at -20°C

Description: Human male Jurkat (human acute T-cell leukemia) genomic DNA.

Source: Jurkat (acute T-cell leukemia) cells were grown to confluency in RPMI plus 10% fetal bovine serum. Genomic DNA was isolated by a standard genomic purification protocol (1), phenol extracted and equilibrated to 10 mM Tris-HCl (pH 7.5) and 1 mM EDTA.

Applications:

- A control for Methylation-Specific PCR (MSP) (2), Bisulfite sequencing, Methylation-sensitive Single-Nucleotide Primer Extension (Ms-SNuPE), Combined Bisulfite Restriction Analysis (COBRA), Bisulfite treatment and PCR-Single-Strand Conformation Polymorphism Analysis (Bisulfite-PCR-SSCP/BiPS).
- PCR, SNP Analysis, Southern Blotting
- Genomic DNA library construction

Supplied in: 10 mM Tris-HCl (pH 7.5) and 1 mM EDTA. Avoid multiple freeze/thaw cycles.

Quality Assurance: Purified free of contaminating proteins and RNA.

A_{260/280} Ratio: 1.92

Reference:

1. Sambrook, J. and Russell, D. (2001) *Molecular Cloning: A Laboratory Manual*, (3rd ed.), (pp. 6.4–6.12). Cold Spring Harbor: Cold Spring Harbor Laboratory Press.
2. Herman, J.G. and Baylin, S.B. (1996). U.S. Patent No. 5,786,146. John Hopkins University School of Medicine.

CERTIFICATE OF ANALYSIS

Jurkat Genomic DNA



1-800-632-7799
info@neb.com
www.neb.com



N4001S 005140716071

N4001S

15 µg Lot: 0051407 Exp: 7/16

100 µg/ml Store at -20°C

Description: Human male Jurkat (human acute T-cell leukemia) genomic DNA.

Source: Jurkat (acute T-cell leukemia) cells were grown to confluency in RPMI plus 10% fetal bovine serum. Genomic DNA was isolated by a standard genomic purification protocol (1), phenol extracted and equilibrated to 10 mM Tris-HCl (pH 7.5) and 1 mM EDTA.

Applications:

- A control for Methylation-Specific PCR (MSP) (2), Bisulfite sequencing, Methylation-sensitive Single-Nucleotide Primer Extension (Ms-SNuPE), Combined Bisulfite Restriction Analysis (COBRA), Bisulfite treatment and PCR-Single-Strand Conformation Polymorphism Analysis (Bisulfite-PCR-SSCP/BiPS).
- PCR, SNP Analysis, Southern Blotting
- Genomic DNA library construction

Supplied in: 10 mM Tris-HCl (pH 7.5) and 1 mM EDTA. Avoid multiple freeze/thaw cycles.

Quality Assurance: Purified free of contaminating proteins and RNA.

A_{260/280} Ratio: 1.92

Reference:

1. Sambrook, J. and Russell, D. (2001) *Molecular Cloning: A Laboratory Manual*, (3rd ed.), (pp. 6.4–6.12). Cold Spring Harbor: Cold Spring Harbor Laboratory Press.
2. Herman, J.G. and Baylin, S.B. (1996). U.S. Patent No. 5,786,146. John Hopkins University School of Medicine.

CERTIFICATE OF ANALYSIS