

NIH 3T3 Mouse Genomic DNA



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



N4004S 005140516051

N4004S

15 µg **Lot: 0051405** **Exp: 5/16**
100 µg/ml **Store at -20°C**

Description: NIH 3T3 (mouse embryonic fibroblast cell line) genomic DNA.

Source: NIH 3T3 (mouse embryonic fibroblast) cells are grown to confluency in DMEM 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: 1.91

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

NIH 3T3 Mouse Genomic DNA



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



N4004S 005140516051

N4004S

15 µg **Lot: 0051405** **Exp: 5/16**
100 µg/ml **Store at -20°C**

Description: NIH 3T3 (mouse embryonic fibroblast cell line) genomic DNA.

Source: NIH 3T3 (mouse embryonic fibroblast) cells are grown to confluency in DMEM 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: 1.91

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