

## Human genomic DNA (HL-60)

CCL-240D™

## Description

High molecular weight genomic DNA isolated from the HL-60 human acute promyelocytic leukemia cell line (ATCC CCL-240). This cell line is positive for myc oncogene expression.

Organism: Homo sapiens, human Derived from: HL-60 (ATCC CCL-240)

Type strain: No Mass: 10 µg

### Storage Conditions

**Product format:** Frozen

Storage conditions: -20°C or colder

### Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic use, any human or animal consumption, or any diagnostic use.

### BSL<sub>1</sub>

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of Biosafety in Microbiological and Biomedical Laboratories (BMBL), U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies



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and procedures as well as any other applicable regulations as enforced by your local or national agencies.

## Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

## **Quality Control Specifications**

**Electrophoresis - RNA content:** No visible RNA detected in the agarose gel **Electrophoresis - digestion:** Tested and verified for PCR amplification and restriction digestion.

#### Notes

The HL-60 cell line (CCL-240) was established from a 36 year-old White female with acute promyelocytic leukemia. This cell line is positive for myc oncogene expression.

**Formulation:** 1X TE buffer (10 mM Tris-HCL, 1 mM EDTA, pH 8.0)

**Concentration:**See lot-specific information **Lot Number:**See lot-specific information

**Cell line description:**See lot-specific information

#### **Material Citation**

If use of this material results in a scientific publication, please cite the material in the following manner: Human genomic DNA (HL-60) (ATCC CCL-240D)

#### References



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References and other information relating to this material are available at www.atcc.org.

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#### Revision

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