Genomic DNA from *Aspergillus clavatus* strain Kral

1007D-2™

Description

Genomic DNA isolated from *Aspergillus clavatus* QM 1276. This fungal strain is also available as ATCC® Catalog No.: 1007.

Organism: *Aspergillus clavatus* Desmazieres

Derived from: *Aspergillus clavatus* QM 1276 [ATCC 9598, ATCC 9602, CBS 513.65, CCC-T-191b, IMI 15949, LSHB Ac86, LSHB Ac95, MIT 213, NCTC 3887, NCTC 978, NRRL 1, NRRL 1656, QM 7404, Thom 107, WB 1] (ATCC 1007)

Genome sequenced strain: Yes

Type strain: No

Mass: 2 µg

Shipping information: Stored in 1X TE buffer

Storage Conditions

Product format: Freeze-dried

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 1

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 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.

Handling Procedures

Centrifuge tube prior to opening to prevent loss of pelleted material

1. Rehydrate contents of vial with molecular grade H_2O.
DNA is dried in 1X Tris buffer.
2. Place vial at 37°C for 1 hour or at +2°C to 8°C overnight.
3. For more complete rehydration and to fully recover DNA incubate the sample overnight at 4°C while rocking, then incubate for 1 hour at 65°C.

Quality Control Specifications

Electrophoresis - RNA content: No RNA was detected by electrophoresis
Integrity: Integrity of DNA was determined by electrophoresis on a 1% agarose gel stained with SYBR Safe™, and was found to be of high molecular weight.
Functional tests: Functional activity was confirmed by PCR amplification of approximately 1500 base pairs fragment of rRNA gene cluster including ITS1-5.8S-ITS2 region.
Identity: Identity confirmed by sequencing of ITS1, 5.8S gene and ITS2 regions of ribosomal RNA (~ 500 base pairs).

Notes

Genomic DNA is appropriate for PCR and other molecular biology applications.

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: Genomic DNA from *Aspergillus clavatus* strain Kral (ATCC 1007D-2)

References

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

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Revision

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This information on this document was last updated on 2024-10-24

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