Genomic DNA from *Candida parapsilosis* strain CBS 604

22019D-5™

Description

Genomic DNA isolated from *Candida parapsilosis* strain CBS 604 (ATCC 22019). This whole-genome sequenced product can be used in PCR and other molecular biology applications.

Organism: *Candida parapsilosis* (Ashford) Langeron et Talice
Derived from: *Candida parapsilosis* CBS 604 [CCRC 20515, DBVPG 6150, DMS 5784, IBL 2545, IFO 1396, IGC 2545, JCM 1785, NBRC 1396, NCYC 601, NRRL Y-12969, UCD 61-27] (ATCC 22019)
Genome sequenced strain: Yes
Type strain: No
Mass: 5 µg
Shipping information: Stored in 1X TE buffer

Storage Conditions

Product format: Freeze-dried
Storage conditions: 2°C to 8°C

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

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22019D-5

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. Resuspending the dried DNA in ≥ 250 µL may give better results.

Quality Control Specifications

Electrophoresis - RNA content: No RNA was detected by electrophoresis
Purity (A260/A280): 1.7 to 2.0
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.

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22019D-5

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 *Candida parapsilosis* strain CBS 604 (ATCC 22019D-5)

References

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

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22019D-5

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Revision

This information on this document was last updated on 2025-01-24

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