



Product Sheet

Candida metapsilosis (ATCC® 14054™)

Please read this **FIRST**



Storage Temp.
Frozen: -80°C or colder
Freeze-Dried: 2°C to 8°C
Live Culture: See Propagation Section



Biosafety Level
1

Intended Use

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

Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: *Candida metapsilosis* (ATCC® 14054™)

American Type Culture Collection
PO Box 1549
Manassas, VA 20108 USA
www.atcc.org

800.638.6597 or 703.365.2700
Fax: 703.365.2750
Email: Tech@atcc.org

Or contact your local distributor

Description

Strain Designation: 3175

Deposited Name: *Candida parapsilosis* (Ashford) Langeron et Talice

Product Description: An ampoule containing viable cells (may include spores and mycelia) suspended in cryoprotectant.

Propagation

The information recommended in this section is to assist users in obtaining living culture(s) for their studies. The recommendation does not imply that the conditions or procedures provided below are optimum. Experienced researchers may initiate the growth of a culture in their own way.

ATCC® Medium 200: YM agar or YM broth

ATCC® Medium 1245: YEPD

ATCC® Medium 28: Emmons' modification of Sabouraud's agar

Growth Conditions

Temperature: 24°C to 26°C

Atmosphere: Typical aerobic

Recommended Procedure

For **freeze-dry (lyophilized) ampoules:**

1. Open an ampoule according to enclosed instructions.
2. From a single test tube of **sterile distilled water** (5 to 6 mL), withdraw approximately 0.5 to 1.0 mL with a sterile pipette and apply directly to the pellet. Stir to form a suspension.
3. Aseptically transfer the suspension back into the test tube of sterile distilled water.
4. Let the test tube sit at room temperature (25°C) undisturbed **for at least 2 hours**; longer (e.g., overnight) rehydration might increase viability of some fungi.
5. Mix the suspension well. Use several drops (or make dilutions if desired) to inoculate recommended solid or liquid medium. Include a control that receives no inoculum.
6. Incubate the inoculum at the propagation conditions recommended.
7. Inspect for growth of the inoculum/strain regularly. Viability is typically noticeable after 3-5 days of incubation. However, the time necessary for significant growth will vary from strain to strain.

Colony and Cell Morphology: After 4 day at 25°C colonies cream-coloured, semi-dull to glistening and smooth. Yeast cells globose to oval 2,0-3,5 x 3,0-4,5 µm, single or budding. Long, elongated cells sometimes produced, pseudomycelium produced forming very straight, regular chains of elongated cells after 2-3 days.

Notes

Deposited as *Candida parapsilosis*

Additional, updated information on this product may be available on the ATCC® web site at www.atcc.org.

DNA Sequence

D1/D2 region of 26S rRNA gene

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TTCCAATTCGACCTCAAATCAGGTAGGACTACCCGCT
GAACTTAAGCATATCAATAAGCGGAGGAAAAGAAACCAACAGGGATTGCCTTAGTAGCGGCGAGTG
AAGCGGCAAAAGCTCAAATTTGAAATCTGGCACTTTCAGTGTCCGAGTTGTAATTTGAAGAAGGTATC
TTTGGGTCTGGCTCTTGTCTATGTTTCTTGGAACAGAACGTCACAGAGGGTGAGAATCCCCTGCGATGA
GATGACCCAGACCTATGTAAGTTCCTTCGAAGAGTCGAGTTGTTGGGAATGCAGCTCTAAGTGGGT
GGTAAATCCATCTAAAGCTAAATATTGGCGAGAGACCGATAGCGAACAAGTACAGTGATGGAAAGA
TGAAAAGAACTTTGAAGAGAGAGTGAAGAAAGTACGTGAAATTTGAAAGGGAAGGGCTTGAGATC
AGACTTGGTATTTGTATGTTACTCTTTCGGGGGTGGCCTCTACAGTTTACCGGGCCAGCATCAGTTTGG
GCGGTAGGAGAATTGCAAGAAATGTGGCACTGCTTCGGTAGTGTGTTATAGTCTTTGTGATACTGCC
AGCCTAGACTGAGGACTGCGGCTTCGGCTTAGGATGTTGGCATAATGATCTTAAGTCGC
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Isolation

Human, New York, USA

References



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Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the current publication of the *Biosafety in Microbiological and Biomedical Laboratories* from the U.S. Department of Health and Human Services Centers for Disease Control and Prevention and National Institutes for Health.

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Additional information on this culture is available on the ATCC web site at www.atcc.org.
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