





Product Sheet

Trichophyton tonsurans (ATCC® 10270™)

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
2

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: *Trichophyton tonsurans* (ATCC® 10270™)

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: IOC 2415 [Leao 2415]

Deposited Name: *Trichophyton mentagrophytes* (Robin) Blanchard

Product Description: An ampoule containing viable cells (yeast cells, spores, or agar cubes with 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 28: Emmons' modification of Sabouraud's agar

ATCC® Medium 200: YM agar or YM broth

ATCC® Medium 1245: YEPD

Growth Conditions

Temperature: 24°C to 26°C

Atmosphere: Typical aerobic

Recommended Procedure

Frozen ampoules packed in dry ice should either be thawed immediately or stored in liquid nitrogen. If liquid nitrogen storage facilities are not available, frozen ampoules may be stored at or below -70°C for approximately one week. **Do not under any circumstance store frozen ampoules at refrigerator freezer temperatures (generally -20°C).** Storage of frozen material at this temperature will result in the death of the culture.

1. To thaw a frozen ampoule, place in a **25°C to 30°C** water bath, until just thawed (**approximately 5 minutes**). Immerse the ampoule just sufficient to cover the frozen material. Do not agitate the ampoule.
2. Immediately after thawing, wipe down ampoule with 70% ethanol and aseptically transfer at least 50 µL (or 2-3 agar cubes) of the content onto a plate or broth with medium recommended.
3. Incubate the inoculum/strain at the temperature and conditions recommended. Inspect for growth of the inoculum/strain regularly. The sign of viability is noticeable typically after 2-4 days of incubation. However, the time necessary for significant growth will vary from strain to strain.

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. The sign of viability is noticeable typically after 1-2 days of incubation. However, the time necessary for significant growth will vary from strain to strain.

Colony and Cell Morphology: Mycelium white, dense, velutinous. Hyphae hyaline, guttulate

Notes

No special notes.

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

DNA Sequence


18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S ribosomal RNA gene, partial sequence
GGTTTCCTAGGTGAACCTGCGGAAGGATCATTAAACGCGCAGGCCGGAGCCGGCCCCCAGGATAG
GGCCAAACGTCCTCAGGGGTGAGCAGATGTGCGCCGCGCTACCGCCCCATTCTTGTCTACCTTACTC
GGTTGCCTCGGCGGGCCGCGCTCTCCAGGAGAGCCGTTCCGGCAGCCTCTCTTTATAGCGGCTCAACG
CTGGACCGCGCCCGCGGAGACAGACGCAAAAAAAAAATTCTTTCAGAAGAGCTGTCTAGTCTGAGCG




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TTAGCAAGCAAAAATCAGTTAAAACCTTCAACAACGGATCTCTTGGTTCCGGCATCGATGAAGAACGC
AGCGAAATGCGATAAGTAATGTGAATTGCAGAATTCGGTGAATCATCGAATCTTTGAACGCACATTGC
GCCCTTGGCATTCCGGGGGGCATGCCTGTTTCGAGCGTCATTTAGCCCTCAAGCCCGGCTTGTGTGA
TGGACGACCGTCCGGCGCCCCGTCTCTGGGGTGCGGGACGCGCCGAAAGCAGTGGCCAGGCCG
CGATTCCGGCTTTCTAGGCGAATGGCAACAAACCAGCGCTCCAGGACCGGCCCTGGCCTCAAA
ATCTGTTTTATACCTATCAGTTGACCTCGGATCAGGTAGGGATAC

D1D2 region of the 28S ribosomal RNA gene

ATATCAATAAGCGGAGAAAAGAAACCAACAGGGATTGCCAGTAACGGCGAGTGAAGCGGCAA
GAGCTCAAATTTGAAATCTGGCCTCCCCGGGGGCCGAGTTGTAAATTTGCAGAGGATGTTTCGGGTGC
GGCCGCGCTAAGTTCTTGAACAGGACGTCAGAGAGGGTGAGAATCCCGTCTTGGCGGGCGGT
CCGCGCCCGTGTGAAGTCTCTGACGAGTCTGAGTTGTTTGGGAATGCAGCTCTAAGCGGGTGGTAAA
TTTCATCTAAAGCTAAATATTGGCCGGAGACCGATAGCGCACAAAGTAGAGTATCGAAAGGTTAAAA
GCACCTTGAAGGGAGTTAAACAGCACGTTGAAATTTGAAAGGGAAGCGCTTCCGGCCAGACTCG
GGGGCGGGTTCAGCGGGTCTCGTCCGCCGTATTCTCGTTTTCCCGGCCAGCATGTTTCGACG
GCCGTCAAAGGCCCCCGAATGTGTCTCTCGGGACGCTTATAGCCGGGGTGAATGCGGCC
GTCGGGACTGAGGAACCGCTCCGGCTCGGATGCTGGCGTAATGGCCGTAAGCGGC



Isolation

Tinea capitis



References

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



Biosafety Level: 2

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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Disclaimers

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