



Product Sheet

Geotrichum citri-aurantii (ATCC® 201505™)

Please read this **FIRST**



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: *Geotrichum citri-aurantii* (ATCC® 201505™)

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: Tm2

Deposited Name: *Geotrichum candidum* Link : Persoon, anamorph

Product Description:

An ampoule containing viable cells 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 307: Cornmeal agar

ATCC® Medium 336: Potato dextrose agar (PDA)

Growth Conditions

Temperature: 25°C to 30°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.

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1. To thaw a frozen ampoule, place in a **2530 °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 10 microliter (or any amount desired up to all) of the content onto a plate or broth with medium recommended.
3. Incubate the inoculum/strain at the temperature and conditions recommended.
4. Inspect for growth of the inoculum/strain regularly. The sign of viability is noticeable typically after 2-3 days of incubation. However, the time necessary for significant growth will vary from strain to strain.

Colony and Cell Morphology: After 5 days at 25°C colonies whitish and butyrous producing abundant sympodial and arthric conidia.

Notes

Deposited as *Geotrichum candidum*; citrus race; treatment with alcohols or citral inhibits arthrospore germination.

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 26S ribosomal RNA gene, partial sequence
AAATCAAAAACCTTTTAACAATGGATCTCTTGGTTCTCGTATCGATGAAGAACGCGAGCGAAAACGCGATAT
TTCTTGTGAATTGCAGAAGTGAATCATCAGTTTTTGAACGCACATTGCACTTTGGGGTATCCCCAAAGT
ATACCTGTTTGAGCGTTGTTTCTCTTGGAAATGCATTGCTTTTCTAAAATATCGAAACAAAATTGTTTG
TAACAAAAATCATTCAACCTCAGATCAAGTAGGACTACCCGCTGAACCTAAGCATATCAA
D1/D2 region of 26S ribosomal RNA gene
ATATCAATAAGCGGAGGAAAAGAAACCAACAGGGATTGCCTTAGTAACGGCGAGTGAAGCGGCAAA
AGCTCAAAATTTGAAATCGGCCAACAGGTCGAGTTGTAATTTGTAGATTGTATCTTGAGAGCGGATTAA
AGTCTGTTGGAATACAGCGCCTTAGAGGGTGACAGCCCGTAAAATCTATTCTCATTGTAAGATACTTT
CGAAGAGTCGAGTTGTTTGGGAATGCAGCTCTAAGTGGGAGGTAATTCCTTCAAAGCTAAATATTG
ACGAGAGACCGATAGCGAACAAGTACTGTGAAGGAAAGATGAAAAGCACCTTTGAAAAGAGAGTGA
AAAAGTACGTGAAATTGTTAAAAGGGAAGGGTATTGAATCAGACGTGGTGTGTTGTTTCAGCATTGTT
TCGGCAGTGTATTCAACAATACTAGGCCAAGGTGGGGTATTGGGAGTGAAAAGAAGTAGGAAT



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GTAGCTCCTCGGAGTGTATAACCTACTTTTCATAACTCCTTAGTCGCCTCAGGACTGCGCTTCGGCAAGG
ACTTTGGCATAATGATTCTATAACCGC



tomato field soil, Miyazaki, Japan



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



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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