



Product Sheet

# *Malassezia yamatoensis* (ATCC® MYA-4956™)

Please read this **FIRST**



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

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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: *Malassezia yamatoensis* (ATCC® MYA-4956™)

American Type Culture Collection  
PO Box 1549  
Manassas, VA 20108 USA  
[www.atcc.org](http://www.atcc.org)

800.638.6597 or 703.365.2700  
Fax: 703.365.2750  
Email: [Tech@atcc.org](mailto:Tech@atcc.org)

Or contact your local distributor

## Description

**Strain Designation:** CBS 9725 [JCM 12262, M 9985]

**Deposited Name:** *Malassezia yamatoensis*

**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 2737: Modified Leeming & Notman Agar (MLNA)

ATCC® Medium 2693: modified Dixon (mDixon)

## Growth Conditions

**Temperature:** 30°C to 32°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 50 µL (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 6-7 days on Leeming & Notman modified (MLNA) agar at 32°C, colonies flat to convex, pale yellow-cream with undulating margin. Cells are ovoid to short cylindrical with monopolar budding.

## Notes

In general, the strain grows better on MLNA than mDixon.

Additional, updated information on this product may be available on the ATCC® web site at [www.atcc.org](http://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  
GTTTCTGTAGGTGAACCTGCAGAAGGATCATTAGTGAAAGCATGGGTCAGCCATACGGATGCGCAAG  
CGTCTCTGGCGACCTTTTTTCCATTTATCCAAACCCGTGTGCACTGTGTAGCGTGTCTTTTTGGATGCGC  
TACGAAATTTCTACACTCGTATGTTGTATGTACGTGAATTTGTTGGACCGTAAC TGCCCAACCAACTTTA  
CACAACTTTCGACAACGGATCTCTTGTTCTCCCATCGATGAAGAACGCGAGCAACGCGATAGGTAA  
TGTGAATTGCAGAATTCCGTGAATCATCGAATCTTTGAACGCACCTTGCGCTCCATGGTATTCCGTGGA  
GCATGCCTGTTGAGTGCCGTGAATCTCTCCCCCTTTGGGTTGCGAAAGCAGTCTAGCGCGCGGAG  
GTTGGATGGGCGATGCCTTGCAGGGCTGGCCGAAAAGCATTAGCGCCTCGCTAGAGCCTCTAAGGG  
AGAGGCCCAAGTGACTTGGCCTTGGTCGGCTATGCCAAACCAAGCAGGAGTGCAAGCGCAGCATGAT  
ACGTCAATTTGCTGTGTGCGCCGCCGCTTGGGGACGCCAAAGACAAGCGCTTGGAGCGAAGAGTGC  
GCTTCGAAGTAGCGTGAGACGATTGCTTGGAAACGCATTCCCTTTTTCTCTGCTCTCAATCAGGTAG  
GATCACCCGCCGAACCTAAG

D1D2 region of the 26S ribosomal RNA gene

CATATCATTAAGCGGAGGAAAAGAACTAACAAGGATCCCCTAGTAACGGCGAGTGAAGCGGGAA  
GAGCTCAACTTTGAAAGCTGGTACCTTTGGTGCCCGCTTGTAACTCGAGACGCGTTTTCCGTGCGGC  
GCTATGGACAAGTTCTTGGAAACAGGACATCGTAGAGGGTAAAATCCCGTACTTGCCATGGATGTAC



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CGTGCTTTGTGATACGCGCTCCAAGAGTCGAGTTGTTTGGGATTGCAGCTCAAATGGGTGGTAGACTC  
CATCTAAAGCTAAATATCGGGGAGAGACCGATAGCGAACAAGTACCCTGAGGGAAAGATGAAAAGC  
ACTTTGGAAAGAGAGTTAAAAGTACGTGAAATTGTCGAAAGGGAAGCGCTTGAAGTCAGCCATGCCG  
CTCAGGACTCAGCTTGGTTTTTCCGAGTGTATTTCTGGGTAGCAAGTCAGCATTGGTTTGGTGCCTCG  
GAGAAGCATCTGGGAATGTAGCGCCCTCGGGCGTGTATAGCCTAGGATTGGATACGACGTGCTAGA  
CCAAGGAACGACGCGCCCTTTTGGCGGGTCTTCGGACACCTTCGGCCTTAGGATGCTGGCGTAATGG  
CTTTAAGTGGC



Male patient with seborrheic dermatitis, Tokyo, Japan.



References and other information relating to this product are available online at [www.atcc.org](http://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](http://www.atcc.org).  
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