



Product Sheet

# *Mycoplasma hyorhinis* (ATCC® 29052™)

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: *Mycoplasma hyorhinis* (ATCC® 29052™)

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

**Designation:** DBS 1050 [3T-6]

**Deposited Name:** *Mycoplasma hyorhinis* Switzer

**Product Description:** ATCC® CCL-10™ (BHK-21) cell culture or a specially formulated cell-free medium\* is suitable for propagation and maintenance. \*The following publication proposes a medium for the cultivation of this strain, but it has not yet been tested in our lab. Applied and Environmental Microbiology, Vol. 61(5), May 1995, p.1976-1979.

## Propagation

### Growth Conditions

**Temperature:** 37°C

**Atmosphere:** 5% CO<sub>2</sub>

**Recommended Host:** BHK-21 (ATCC® CCL-10™)

### Propagation Procedure

1. BHK-21 cells (ATCC® CCL-10™) are grown in Eagles MEM with non-essential amino acids and Earle's BSS (90%), plus 10% Fetal Bovine Serum under 5% CO<sub>2</sub> (9 mL of the medium and 1 mL of FBS in 25 cm<sup>2</sup> plastic tissue culture T-flasks) for approximately three days.
2. When the cells appear to be at optimal condition, change to fresh medium.
3. For scaling up: once the monolayer of cell line is formed, remove the entire medium; add 5 mL of PBS, then incubate for 5 minutes in atmosphere of 5% CO<sub>2</sub>. After 5 minutes, remove PBS and add 1 mL of trypsin, then incubate for 2 minutes at recommended atmosphere. After two minutes, add 5 mL of fresh medium to the flask, the monolayer begins to detach. Take the entire content out of the flask and place in a centrifuge tube and spin for 5 minutes at 2000 rpm. Remove supernatant and resuspend the pellet with 1 mL of the fresh medium. Inoculate the pellet into two 75 cm<sup>2</sup> plastic tissue culture T-flasks containing 22.5 mL of fresh medium and 10% FBS (2.5 mL) and incubate at recommended atmosphere for approximately three days.
4. Reconstitute a vial of ATCC® 29052™ with 2.0 mL of the culture medium, and add approximately 1.0 mL to each of two T-flasks of BHK-21 cells.
5. Incubate at 37°C under 5% CO<sub>2</sub> until the monolayers begin to detach. This will normally take three to five days. Additional time may be required when initially growing from the freeze-dried vial.
6. Pool the monolayers and medium from the flasks and centrifuge at 9000 rpm for 35 minutes.
7. Resuspend the pellet in a small amount of complete tissue culture medium. Additional passages may be made or cells prepared for storage. Either freeze the suspension by adding an equal amount of 20% glycerol, or freeze-dry using an appropriate cryoprotectant.

## Notes

This strain is not maintained readily on conventional agar-broth media.

Additional information on this culture is available on the ATCC® web site at [www.atcc.org](http://www.atcc.org).

## References

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

### Disclaimers

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Please see the enclosed Material Transfer Agreement (MTA) for further details regarding the use of this product. The MTA is also available on our Web site at [www.atcc.org](http://www.atcc.org)

Additional information on this culture is available on the ATCC web site at [www.atcc.org](http://www.atcc.org).

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