

**For Research Use Only.
Not for use in diagnostic procedures.**



Anti-*H. pylori* mAb

CODE No.	D369-3
CLONALITY	Monoclonal
CLONE	TMDU-D8
ISOTYPE	Mouse IgG3 κ
QUANTITY	100 μ L, 1 mg/mL
SOURCE	Purified IgG from hybridoma supernatant
IMMUNOGEN	Sonicated whole bacterial lysate of <i>H.pylori</i> (a combination of strains ATCC 43504, ATCC 43579, ATCC 43629)
FORMULATION	PBS containing 50% Glycerol (pH 7.2). No preservative is contained.
STORAGE	This antibody solution is stable for one year from the date of purchase when stored at -20°C.

APPLICATION-CONFIRMED

Immunohistochemistry 1 μ g/mL (paraffin section)

Heat treatment for paraffin embedded section: microwave oven, for 40 min. in 10 mM citrate buffer (pH 6.2)

SPECIES CROSS REACTIVITY on IHC

Species	Human	Mouse	Rat	Hamster
Sample	<i>H. pylori</i> -infected stomach	Not tested	Not tested	Not tested
Reactivity	+			

REFERENCES

- 1) Munari, M., *et al.*, *Blood*. **16**, 6612-6616 (2011)
- 2) Ito, T., *et al.*, *Lab. Invest.* **88**, 664-681 (2008)

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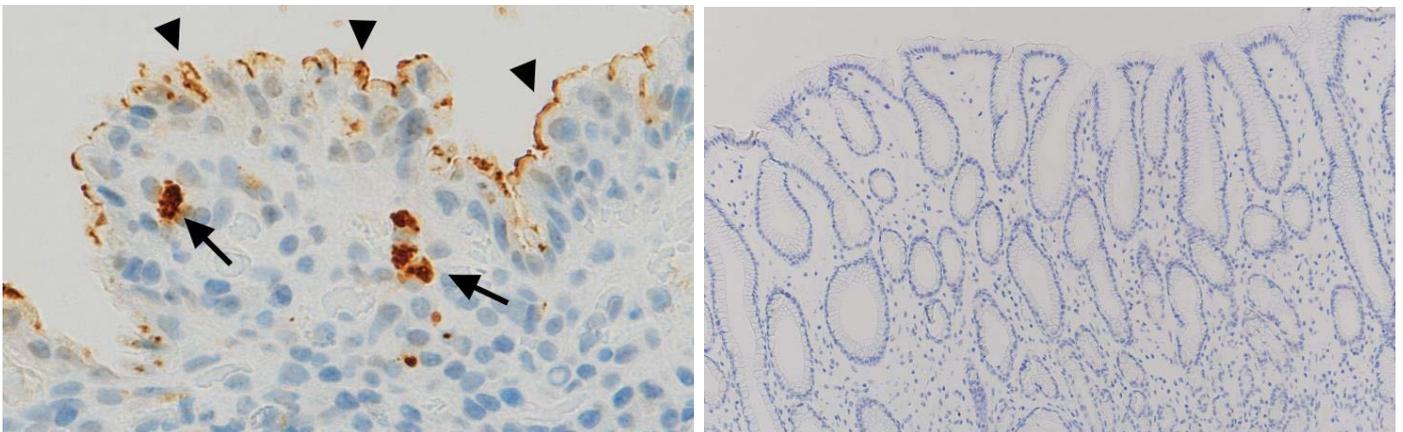
RELATED PRODUCTS

D369-3	Anti- <i>H. pylori</i> mAb
M078-3	Mouse IgG3 (isotype control)

Immunohistochemistry for formalin fixed paraffin-embedded section

- 1) Deparaffinize tissue sections in Xylene 3 times for 5 min. each.
- 2) Immerse the slides with Ethanol 3 times for 5 min. each.
- 3) Immerse the slides with PBS 3 times for 5 min.
- 4) Remove the slides from PBS and heat-treat with 10 mM Citrate buffer (pH 6.2) for 40 min. at 97°C using microwave oven.
- 5) Let the slide cool down until at room temperature in the Citrate buffer.
- 6) Remove the slides from the Citrate buffer and inactivate endogenous peroxidase with 3% H₂O₂ in Methanol for 10 min.
- 7) Wash the slides with PBS 2 times for 5 min each.
- 8) Remove the slides from PBS, and immerse the slides in Blocking buffer [20 mM HEPES, 1% BSA, 135 mM NaCl] for 5 min. at room temperature to block non-specific staining. Do not wash.
- 9) Tip off the blocking buffer, wipe gently around each section and cover tissues with primary antibody diluted with Blocking buffer as suggested in the **APPLICATION**. (The concentration of antibody will depend on the conditions.) Incubate the sections for 1 hr. at room temperature.
- 10) Wash the slides 2 times in PBS for 5 min. each.
- 11) Incubate the section with Histostar (Ms + Rb) (MBL; code no. 8460) for 30 min. at room temperature.
- 12) Wash the slides 3 times in PBS for 5 min. each.
- 13) Visualize by reacting for 10 min. with Histostar DAB Substrate Solution (MBL; code no. 8469). *DAB is a suspect carcinogen and must be handled with care. Always wear gloves.
- 14) Wash the slides 2 times in PBS for 5 min. each.
- 15) Counterstain in hematoxylin for 5 min., wash the slides 3 times in water for 5 min. each, and then immerse the slides in PBS for 5 min.
- 16) Dehydrate by immersing in Ethanol 3 times for 5 min. each, followed by immersing in Xylene 3 times for 3 min. each. Now ready for mounting.

(Positive control for Immunohistochemistry; *H. pylori*-infected stomach)



Immunohistochemical detection of H. pylori in human stomach

Left: *H. pylori* positive case (arrowhead: in mucous layer, arrow: in lamina propria)
Right: *H. pylori* negative case

Brown: Anti-*H. pylori* mAb (D369-3)
Blue: Hematoxylin

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