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





MONOCLONAL ANTIBODY

Anti-monomeric Keima-Red mAb

Code No. M126-3M Clone 2F7

Subclass Mouse IgG2a Quantity 100 µL

Concentration 1 mg/mL

BACKGROUND: The fluorescent protein "Keima-Red," which was cloned from the stony coral whose Japanese name is "Komon-Sango." *CoralHue*™ monomeric Keima-Red (mKeima-Red) absorbs light maximally at 440 nm and emits red light at 620 nm. Thus mKeima-Red exhibits an extremely large Stokes shift (180 nm). The red fluorescence is stable under usual aerobic conditions.

SOURCE: This antibody was purified from hybridoma (clone 2F7) supernatant using protein A column. This hybridoma was established by fusion of mouse myeloma cell P3U1 with Balb/c mouse lymphocyte immunized with recombinant *CoralHue*™ monomeric Keima-Red.

FORMULATION: 100 μg IgG in 100 μL volume of 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.

REACTIVITY: This antibody reacts with *CoralHue*[™] monomeric Keima-Red on Western blotting.

APPLICATIONS:

Western blotting; 1 µg/mL

Immunoprecipitation; Not recommended

*Clone 3C9 is suitable for this application. Please refer to

the data sheet (MBL, code no. M127-3M).

<u>Immunohistochemistry</u>; Not tested <u>Immunocytochemistry</u>; Not tested <u>Flow cytometry</u>; Not tested

Detailed procedure is provided in the following **PROTOCOL**.

INTENDED USE:

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

REFERENCES:

- 1) Lee, J., et al., J. Cell Biol. 217, 1613-1622 (2018) [WB]
- 2) Yamashita, SI., et al., J. Cell Biol. 215, 649-665 (2016) [WB]
- 3) Kogure, T., et al., Nat. Biotechnol. 24, 577-581 (2006)

Related Products:

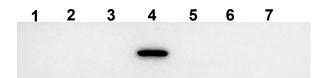
Please visit our website at https://ruo.mbl.co.jp/.

The descriptions of the following protocols are examples. Each user should determine the appropriate condition.

PROTOCOL:

SDS-PAGE & Western blotting

- 1) Mix the sample with equal volume of Laemmli's sample buffer.
- 2) Boil the samples for 3 minutes and centrifuge. Load 10 μ L of sample per lane on a 1-mm-thick SDS-polyacrylamide gel and carry out electrophoresis.
- 3) Blot the protein to a polyvinylidene difluoride (PVDF) membrane at 1 mA/cm² for 1 hour in a semi-dry transfer system (Transfer Buffer: 25 mM Tris, 190 mM glycine, 20% methanol). See the manufacturer's manual for precise transfer procedure.
- 4) To reduce nonspecific binding, soak the membrane in 10% skimmed milk (in PBS, pH 7.2) for 1 hour at room temperature, or overnight at 4°C.
- 5) Incubate the membrane with primary antibody diluted with 1% skimmed milk (in PBS, pH 7.2) as suggested in the **APPLICATIONS** for 1 hour at room temperature. (The concentration of antibody will depend on the conditions.)
- 6) Wash the membrane with PBS (5 minutes x 6 times).
- 7) Incubate the membrane with 1:10,000 Anti-IgG (Mouse) pAb-HRP (MBL; code no. 330) diluted with 1% skimmed milk (in PBS, pH 7.2) for 1 hour at room temperature.
- 8) Wash the membrane with PBS (5 minutes x 6 times).
- 9) Wipe excess buffer off the membrane, then incubate it with appropriate chemiluminescence reagent for 1 minute. Remove extra reagent from the membrane by dabbing with paper towel, and seal it in plastic wrap.
- 10) Expose to an X-ray film in a dark room for 20 seconds. Develop the film as usual. The condition for exposure and development may vary.



Western blotting analysis of Azami-Green (1), Dronpa-Green (2), Kaede (3), Keima-Red (4), Kikume Green-Red (5), Kusabira-Orange (6) and Midoriishi-Cyan (7) from E. coli using M126-3M.

CoralHue™ Keima-Red is a product of co-development with Dr. Atsushi Miyawaki at the Laboratory for Cell Function and Dynamics, the Brain Science Institute, and the Institute of Physical and Chemical Research (RIKEN).

Use of *CoralHue*[™] Keima-Red requires a license from MBL Co., Ltd. MBL grants non-profit research organizations the right to use the product for non-commercial research purposes. For commercial entities a commercial license is required. For more information, please contact support@mbl.co.jp.