

## Product Information

### Monoclonal Anti-EDEM3, Clone EDEM3-1

produced in mouse, purified immunoglobulin

Product Number **E0409**

#### Product Description

Monoclonal Anti-EDEM3 (mouse IgG1 isotype) is derived from the hybridoma EDEM3-1 produced by the fusion of mouse myeloma cells and splenocytes from BALB/c mice immunized with a synthetic peptide corresponding to a fragment of human EDEM3 (GeneID: 80267), conjugated to KLH. The corresponding sequence is identical in mouse and rat EDEM3. The isotype is determined using a double diffusion immunoassay using Mouse Monoclonal Antibody Isotyping Reagents, Product Number ISO2.

Monoclonal Anti-EDEM3 reacts with human, mouse, and rat EDEM3. The antibody may be used in various immunochemical techniques including immunoblotting (~120 kDa).

EDEM3 (ER degradation enhancer, mannosidase alpha-like 3), a soluble EDEM homolog, enhances glycoprotein endoplasmic reticulum-associated degradation (ERAD) and mannosyl trimming.<sup>1</sup> Proteins that fail to fold in the ER are transferred from the ER to the cytosol, where they are destroyed by the ubiquitin-proteasome system.<sup>2,3</sup> Quality control in the ER is regulated by productive folding and ERAD mechanisms. Accelerated refolding and degradation of unfolded proteins are induced in response to ER stress by a transcriptional program termed the unfolded protein response (UPR).<sup>4</sup> Three EDEM homologs, EDEM1, EDEM2 and EDEM3 have been identified, which are transcriptionally upregulated upon ER stress by the activated IRE1/Xbp-1 branch.<sup>5</sup> In mammalian cells, EDEM1 is localized to the ER, mainly as a soluble glycoprotein, interacts with the COOH-terminus of calnexin and lacks mannosidase activity.<sup>6</sup> Over-expression of EDEM1 accelerates ERAD by promoting the release of terminally misfolded glycoproteins from calnexin.<sup>6-8</sup> EDEM3 accelerates ERAD of misfolded glycoproteins as well, but in contrast to EDEM1, it greatly stimulates mannosidase trimming *in vivo*.<sup>1</sup>

#### Reagent

Supplied as a solution in 0.01 M phosphate buffered saline, pH 7.4, containing 15 mM sodium azide as a preservative.

Antibody concentration: ~1.0 mg/mL

#### Precautions and Disclaimer

For R&D use only. Not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

#### Storage/Stability

For continuous use, store at 2–8 °C for up to one month. For extended storage, freeze at –20 °C in working aliquots. Repeated freezing and thawing, or storage in “frost-free” freezers, is not recommended. If slight turbidity occurs upon prolonged storage, clarify the solution by centrifugation before use. Working dilution samples should be discarded if not used within 12 hours.

#### Product Profile

**Immunoblotting:** A working antibody concentration of 1-2 µg/mL is recommended using a whole extract of mouse 3T3 or rat NRK cells.

**Note:** In order to obtain best results in various techniques and preparations, it is recommended to determine optimal working dilutions by titration.

#### References

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2. Hosokawa, N. et al., *EMBO Rep.*, **2**, 415-422 (2001).
3. Kostova, Z., and Wolf, D.H., *EMBO J.*, **22**, 2309-2317 (2003).
4. Oda, Y. et al., *J. Cell Biol.*, **172**, 383-393 (2006).
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6. Oda, Y. et al., *Science*, **299**, 1394-1397 (2003).
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VS,ST,TD,KAA,PHC,MAM 03/19-1