

## Product Information

**Anti-Collagen IV antibody, Mouse monoclonal**  
clone J3-2, purified from hybridoma cell culture

Catalog Number **SAB4200500**

### Product Description

Anti-Collagen IV (mouse IgM isotype) is derived from the hybridoma J3-2 produced by the fusion of mouse myeloma cells and splenocytes from BALB/c mice immunized with a placenta preparation rich in basement membrane collagen.<sup>1</sup> The isotype is determined by ELISA using Mouse Monoclonal Antibody Isotyping Reagents, Catalog Number ISO2. The antibody is purified from culture supernatant of hybridoma cells grown in a bioreactor.

Anti-Collagen IV recognizes human, monkey, bovine, dog, rat and mouse collagen IV. The product may be used in several immunochemical techniques including immunoblotting (~170kDa), immunocytochemistry, immunohistochemistry, immunoprecipitation and ELISA.<sup>1</sup>

The composition of the extracellular framework of all vertebrates is dominated by a class of molecules known as collagens, each with a unique feature suited to its function and location.<sup>2</sup> A member of this family, collagen IV, is the primary collagen found in the extracellular basement membranes separating a variety of epithelial and endothelial cells. It is a major component of the dermal-epidermal junction, where it is mostly found in the lamina densa. Collagen IV is a heterotrimeric molecule containing two  $\alpha 1$ -like and one  $\alpha 2$ -like chains. Multiple diseases have been associated with the molecule including Alport and Goodpasture's syndrome, as well as several rheumatological and dermatological diseases including acquired epidermolysis bullosa.<sup>3</sup> Many malignancies have been associated with increased degradation of type IV collagen. Moreover, types I and IV collagens can induce chemoresistance by directly interacting with integrins on cancer cells. Nevertheless, non-collageneous domains of collagen IV released by proteolysis can also inhibit tumor angiogenesis.<sup>2</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 Material Safety Data Sheet for information regarding hazards and safe handling practices.

### Storage/Stability

For extended storage, freeze at  $-20^{\circ}\text{C}$  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 concentration of 1.0-2.0  $\mu\text{g/mL}$  is recommended using human placenta extracts.

Immunohistochemistry: a working concentration of 2.0-4.0  $\mu\text{g/mL}$  is recommended using human liver tissue sections.

**Note**: In order to obtain the best results using various techniques and preparations, we recommend determining optimal working dilutions by titration.

### References

1. Sundarraj, N., and Willson, J., *Immunology*, **10**, 133-140 (1982).
2. Egeblad, M., et al., *Curr. Opin. Cell Biol.*, **22**, 697-706 (2010).
3. Abreu-Velez, A.M., and Howard, M.S., *N. Am. J. Med. Sci.*, **4**, 1-8 (2012).

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