

DESCRIPTION

Species Reactivity	Human
Specificity	Detects pro/active forms of human MMP-2 in Western blots. In Western blots, no cross-reactivity with recombinant human (rh) MMP-9, rhMMP-1, or rhMMP-3 is observed.
Source	Monoclonal Mouse IgG ₁ Clone # 36006
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Chinese hamster ovary cell line CHO-derived recombinant human MMP-2
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

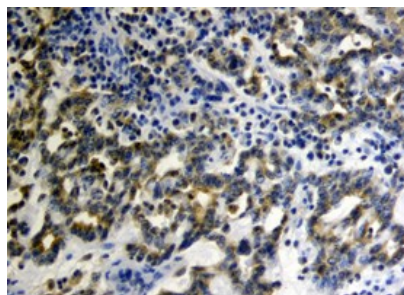
APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. [General Protocols](#) are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Dual RNAscope ISH-IHC Compatible	3-25 µg/mL	Immersion fixed paraffin-embedded sections of human stomach
Western Blot	1 µg/mL	Recombinant Human MMP-2 Western Blot Standard (Catalog # WBC025) under non-reducing conditions only
Immunohistochemistry	50-100 µg/mL	See Below

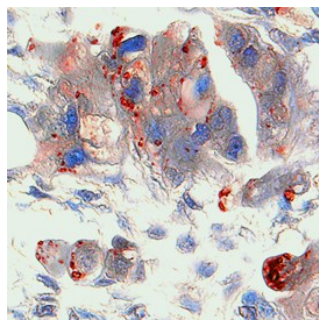
DATA

Immunohistochemistry



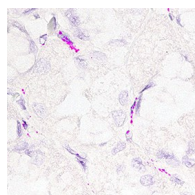
MMP-2 in Human Ovarian Cancer Tissue. MMP-2 was detected in immersion fixed paraffin-embedded sections of human ovarian cancer tissue using Mouse Anti-Human MMP-2 Monoclonal Antibody (Catalog # MAB902) at 10 µg/mL overnight at 4 °C. Tissue was stained using the Anti-Mouse HRP-DAB Cell & Tissue Staining Kit (brown; Catalog # CTS002) and counterstained with hematoxylin (blue). View our protocol for [Chromogenic IHC Staining of Paraffin-embedded Tissue Sections](#).

Immunohistochemistry

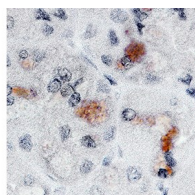


MMP-2 in Human Ovarian Cancer Tissue. MMP-2 was detected in immersion fixed paraffin-embedded sections of human ovarian cancer tissue using Mouse Anti-Human MMP-2 Monoclonal Antibody (Catalog # MAB902) at 15 µg/mL overnight at 4 °C. Tissue was stained (red) and counterstained with hematoxylin (blue). View our protocol for [Chromogenic IHC Staining of Paraffin-embedded Tissue Sections](#).

In-situ Hybridization



In Situ Hybridization (ISH)



Immunohistochemistry (IHC)

Detection of MMP-2 in Human Stomach. Formalin-fixed paraffin-embedded tissue sections of human stomach were probed for MMP2 mRNA (ACD RNAScope Probe, catalog #311751; Fast Red chromogen, ACD catalog # 322750). Adjacent tissue section was processed for immunohistochemistry using mouse anti-human MMP2 monoclonal antibody (R&D Systems catalog # Catalog # MAB902) at 5µg/mL with overnight incubation at 4 degrees Celsius followed by incubation with anti-mouse IgG VisUCyte HRP Polymer Antibody (Catalog # Catalog # VC001) and DAB chromogen (yellow-brown). Tissue was counterstained with hematoxylin (blue). Specific staining was localized to fibroblasts.

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS. For liquid material, refer to CoA for concentration.
Shipping	Lyophilized product is shipped at ambient temperature. Liquid small pack size (-SP) is shipped with polar packs. Upon receipt, store immediately at the temperature recommended below.
Stability & Storage	<p>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</p> <ul style="list-style-type: none"> • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

Matrix metalloproteinases are a family of zinc and calcium dependent endopeptidases with the combined ability to degrade all the components of the extracellular matrix. MMP-2 (gelatinase A), a type IV collagenase, can degrade a broad range of substrates including type IV, V, VII and X collagens as well as elastin and fibronectin. It is believed to act synergistically with interstitial collagenase (MMP-1) in the degradation of fibrillar collagens as it degrades their denatured gelatin forms. MMP-2 has been shown to be associated with many connective tissue cells as well as neutrophils, macrophages and monocytes. Structurally, MMP-2 may be divided into several distinct domains: a pro-domain which is cleaved upon activation; a catalytic domain containing the zinc binding site; a fibronectin-like domain thought to play a role in substrate targeting; and a carboxyl terminal (hemopexin-like) domain containing 2 N-linked glycosylation sites.