

Goat Anti-Human MIP-1α Affinity Purified pAb

Catalog No. PA0952 **Quantity**: 100 μg

Description: MIP-- α belongs to the family of chemotactic cytokines known as chemokines. In humans

there are two major forms, MIP-1α and MIP-1β, also known as CCL3 and CCL4,

respectively. They are produced by many cells, particularly macrophages, dendritic cells, and lymphocytes. MIP-1 are best known for their chemotactic and proinflammatory

effects but can also promote homoeostasis.

Specificity: Human

Host:

Immunogen: Recombinant Human MIP-1α

Goat

Formulation: Lyophilized from sterile filtered PBS, pH 7.2.

Purification: Affinity chromatography on immoblized hMIP-1α matrix

Reconstitution: Centrifuge vial prior to opening. Reconstitute in sterile water to a concentration of 0.1 -

1.0 mg/mL.

Applications: ELISA, Western Blot, Immunohostochemistry

Application Notes: ELISA

To detect human MIP-1 α by direct ELISA (using 100 μ L/well antibody solution) a concentration of at least 0.5 μ g/mL of this antibody is required. This antigen affinity purified antibody, in conjunction with compatible secondary reagents, allows the

detection of 0.2-0.4 ng/well of recombinant human MIP-1a.

Western Blot:

To detect human MIP-1 α by Western Blot analysis this antibody can be used at a concentration of 0.1-0.2 μ g/mL. Used in conjunction with compatible secondary reagents the detection limit for recombinant human MIP-1 α is 1.5-3.0 μ g/lane, under either

reducing or non-reducing conditions.

Immunohistochemistry

To detect human MIP-1 α by immunohistochemistry, the recommended concentration is 5.0 μ g/mL with an overnight incubation at 4 $^{\circ}$ C. Heat-induced antigen retrieval at pH 9.0

is recommended.

The optimal concentration should be determined by the user for each specific application.

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Storage & Stability: Lyophilized product is stable for two years when kept at -20 °C. Reconstituted product is

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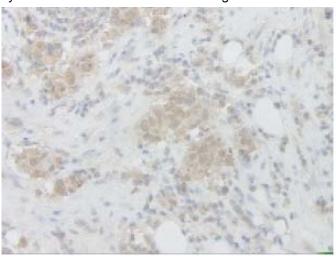
stable for two weeks at 2-8 °C. Avoid repeated freeze-thaw cycles.

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Western blot detection of human MIP-1 alpha using PA0952 at 0.1-0.2 $\mu g/mL$.

1 2 3 4 5 4 7 8 9 39 11 12

Immunohistochemical detection of MIP-1 alpha in formalin-fixed, paraffin-embedded sections of human breast carcinoma. An HRP-labeled polymer detection system was used with a DAB chromagen.



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