

🕜 Tel: +1-301-363-4651 🛛 Email: cusabio@cusabio.com 🙆 Website: www.cusabio.com 🌘

MIF Antibody

Product Code	CSB-RA146975A0HU
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P14174
Immunogen	A synthesized peptide derived from human MIF
Species Reactivity	Human
Tested Applications	ELISA, WB; Recommended dilution: WB:1:500-1:5000
Relevance	Pro-inflammatory cytokine. Involved in the innate immune response to bacterial pathogens. The expression of MIF at sites of inflammation suggests a role as mediator in regulating the function of macrophages in host defense. Counteracts the anti-inflammatory activity of glucocorticoids. Has phenylpyruvate tautomerase and dopachrome tautomerase activity (in vitro), but the physiological substrate is not known. It is not clear whether the tautomerase activity has any physiological relevance, and whether it is important for cytokine activity.
Form	Liquid
Form Conjugate	Liquid Non-conjugated
Form Conjugate Storage Buffer	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
Form Conjugate Storage Buffer Purification Method	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography
Form Conjugate Storage Buffer Purification Method Isotype	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG
Form Conjugate Storage Buffer Purification Method Isotype Clonality	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal
Form Conjugate Storage Buffer Purification Method Isotype Clonality Product Type	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal Recombinant Antibody
Form Conjugate Storage Buffer Purification Method Isotype Clonality Product Type Immunogen Species	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal Recombinant Antibody Homo sapiens (Human)
Form Conjugate Storage Buffer Purification Method Isotype Clonality Product Type Immunogen Species Research Area	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Rabbit IgG Monoclonal Recombinant Antibody Homo sapiens (Human)
Form Conjugate Storage Buffer Purification Method Isotype Clonality Product Type Immunogen Species Research Area Gene Names	Liquid Non-conjugated Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Affinity-chromatography Affinity-chromatography Rabbit IgG Monoclonal Recombinant Antibody Homo sapiens (Human) Immunology

Image





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Description

MIF is an inflammatory cytokine and a crucial innate immune system regulator. MIF overexpression has been associated with increased inflammation and immunopathology. Anti-MIF antibodies have been shown to effectively inhibit tumor development and tumor-associated angiogenesis, implying that MIF is implicated not only in inflammatory and immunological responses but also in tumor cell proliferation. MIF is overexpressed in a wide range of cancers. When binding to the CD74 receptor, MIF stimulates a number of cell signaling pathways, including the PI3K/AKT and MAPK signaling pathways, which increase tumor cell proliferation and survival.

The MIF antibody genes were cloned from B cells that were derived from immunized animals with A synthesized peptide derived from human MIF and then introduced into the plasma vectors, which were transfected into mammalian cell lines for up-scaling expression. The product was purified by A synthesized peptide derived from human MIF to obtain the recombinant antibody against MIF. This recombinant MIF antibody is reactive with the MIF protein from Human. It is recommended for use in the ELISA, WB.