

Mouse Anti-Bovine IgA GMA-3071 (IL-A71)

Description

IgA is a 320 kDa antibody dimer secreted by plasma cells found in the mammary gland, intestine, respiratory tract and skin. At these surfaces IgA neutralizes viruses and protects against pathogen adherence to the body.

Technical Information

Antibody: Mouse monoclonal, IgG_{2a}

Specificity: Bovine IgA¹
Cross-reactivity: Not tested
Immunogen: Bovine Ig

Formulation and Storage

Purity: IgG purified by protein G affinity

chromatography from serum-free

cell culture supernatant.

Product Formulation: Lyophilized from $a \ge 1 \text{ mg/ml}$

solution in 20 mM NaH₂PO₄ 0.15 M NaCl, 1.0% (w/v) mannitol, pH 7.4. Concentration determined by absorbance at 280 nm using an extinction coefficient of 1.4 ($\varepsilon_{0.1\%}$).

Reconstitution: Reconstitute with deionized water.

Storage: Aliquot and store at -20°C for

prolonged periods. Avoid freezethaw cycles. Alternatively add 0.02% (w/v) sodium azide and

store at 4°C.

Country of Origin: Hybridoma country of origin-

Kenya.

Subcloned and produced- USA.

Available Formats: 0.1 mg and 0.5 mg

References

¹ Williams, D.J.L., Newson, J., Naessens, J. 1990. *Vet. Immunol. Immunopathol.* 24:267-283.

Applications

For research use only.

ELISA: Recommended concentration for

use in a solid-phase ELISA is

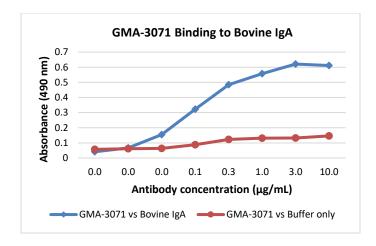
 $0.10 \,\mu g/mL$

Investigator should titrate for

specific application.

ELISA Data

Antibody specificity was confirmed by solid-phase ELISA.



Bovine IgA (Cedar Lane #CLFA07) was coated onto an ELISA plate at a concentration of 14 μ g/mL, for a final coating concentration of 40 nM, in coating buffer, 0.2M carbonate-bicarbonate. Serial dilutions of GMA-3071 were incubated with the antigen.

A goat anti-mouse Ig horseradish peroxidase (HRP) conjugated secondary antibody was used to detect GMA-3071 bound to IgA. O-phenylenediamine dihydrochloride (OPD) was used as a substrate.

Reaction was read on a plate reader at an absorbance of 490 nm after a 7.5-minute development time.