

Porcine anti-Interferon alpha IgG antibody ELISA Kit | IFNalpha IgG-AB elisa kit

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CATALOG #
AAA247711

REACTIVITY
Porcine

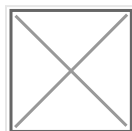
PRODUCT SYNONYMS

anti-Interferon alpha IgG antibody; Porcine anti-Interferon alpha IgG antibody ELISA Kit; IFNalpha IgG-AB elisa kit

PRODUCT SPECIFICATIONS

Reactivity	Porcine
Specificity	This assay has high sensitivity and excellent specificity for detection of IFN-alpha IgG-Ab. No significant cross-reactivity or interference between IFN- alpha IgG-Ab and analogues was observed. NOTE: Limited by current skills and knowledge, it is impossible for us to complete the cross-reactivity detection between IFN-alpha IgG-Ab and all the analogues, therefore, cross reaction may still exist in some cases.
Samples	Serum, plasma, cell culture supernatants, body fluid and tissue homogenate
Assay Type	Quantitative Competitive
Sensitivity	1.0 ng/mL
Preparation and Storage	Store all reagents at 2-8 degree C.

IMAGES



Standard Curve (Sample)

ADDITIONAL INFORMATION

Related Product Information for IFNalpha IgG-AB elisa kit

Intended Uses: This IFN-alpha IgG-Ab ELISA kit is a 1.5 hour solid-phase ELISA designed for the quantitative determination of Porcine IFN-alpha IgG-Ab. This ELISA kit for research use only, not for therapeutic or test applications!

Principle of the Assay: IFN-alpha IgG-Ab ELISA kit applies the competitive enzyme immunoassay technique utilizing a IFN-alpha IgG antigen and an IFN-alpha IgG-Ab-HRP conjugate. The assay sample and buffer are incubated together with IFN-alpha IgG-Ab-HRP conjugate in pre-coated plate for one hour. After the incubation period, the wells are decanted and washed five times. The wells are then incubated with a substrate for HRP enzyme. The product of the enzyme-substrate reaction forms a blue colored complex. Finally, a stop solution is added to stop the reaction, which will then turn the solution yellow. The intensity of color is measured spectrophotometrically at 450nm in a microplate reader. The intensity of the color is inversely proportional to the IFN-alpha IgG-Ab concentration since IFN-alpha IgG-Ab from samples and IFN-alpha IgG-Ab-HRP conjugate compete for the IFN-alpha IgG antigen binding site. Since the number of sites is limited, as more sites are occupied by IFN-alpha IgG-Ab from the sample, fewer sites are left to bind IFN-alpha IgG-Ab-HRP conjugate. A standard curve is plotted relating the intensity of the color (O.D.) to the concentration of standards. The IFN-alpha IgG-Ab concentration in each sample is interpolated from this standard curve.

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Small volumes of product may become entrapped in the seal of the vial during shipment and storage; if necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the cap.

