

DESCRIPTION

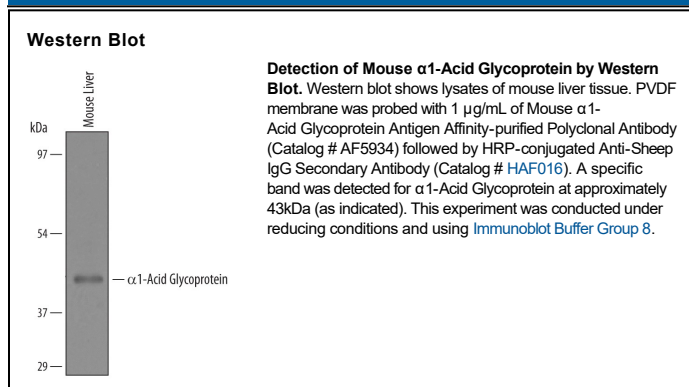
Species Reactivity	Mouse
Specificity	Detects mouse α 1-Acid Glycoprotein in direct ELISAs and Western blots. In direct ELISAs, less than 1% cross-reactivity with human α 1-Acid Glycoprotein is observed.
Source	Polyclonal Sheep IgG
Purification	Antigen Affinity-purified
Immunogen	Mouse plasma-derived α 1-Acid Glycoprotein Accession # P02763
Formulation	Lyophilized from a 0.2 μ m 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 μ m 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
Western Blot	1 μ g/mL	See Below

DATA



PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <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

α 1-Acid Glycoprotein (AGP; also OMD/Orosomucoid) is a 40-46 kDa member of the immunocalin subfamily, lipocalin family of molecules. In mouse, circulating AGP is principally the product of hepatocytes that originates from multiple related genes (AGP-1, -2 & -3 in *Mus musculus*). Circulating AGP-1 and -2 are both 189 amino acids (aa) in length, the principal sources of protein, and show 83% aa identity; AGP-3 contributes little to the AGP pool. In mouse blood, AGP is normally 200-400 μ g/mL. In response to inflammatory mediators (IL-6; IL-1), its concentration will rise 2 to 10 fold. More importantly, a complex glycosylation pattern will also change, transitioning from modestly branched to highly branched oligosaccharides. This change is reflected in its bioactivity, which has been shown to be a function of carbohydrate branching. AGP is generally considered to be a suppressor of inflammation. Rat and human AGP share only 70% and 47% aa identity with mouse AGP, respectively.