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R1075PS Polyclonal Antibody to Cholesterol Oxidase - Ig Fraction

Alternate names: CHOD, Cholesterol oxidase, EC 1.1.3.6

Quantity: 0.1 mg

Concentration: 1.0 mg/ml (by UV absorbance at 280 nm)

Background: Cholesterol oxidases exist as both type I and type II oxidases and are implicated in

bacterial pathogenesis. In addition, they are important as clinical reagents, potential

larvicides, and tools in cell biology.

Host: Goat

Immunogen:Cholesterol oxidase from StreptomycesFormat:State: Lyophilized purified Ig fraction

Purification: Delipidation, salt fractionation and ion exchange chromatography

followed by extensive dialysis against the buffer

Buffer System: 0.02 M Potassium phosphate, 0.15 M Sodium chloride, pH 7.2

Preservatives: 0.01% (w/v) Sodium azide

Reconstitution: Restore with 0.1 ml of deionized water (or equivalent).

Applications: Western blot: 1/500-1/5,000.

Immunoprecipitation: 1/100. ELISA: 1/5,000-1/20,000.

This product has been assayed against 1.0 μ g of Cholesterol oxidase [microorganism] in a standard sandwich ELISA using peroxidase conjugated affinity purified anti-Goat IgG and ABTS as a substrate for 30 minutes at room temperature. A working dilution of

1/2,000 to 1/8,000 of the reconstitution concentration is suggested.

Other applications not tested. Optimal dilutions are dependent on conditions and

should be determined by the user.

Specificity: This product detects Cholesterol oxidase (microorganism). Cross reactivity against

Cholesterol oxidase from other sources is unknown.

Immunoelectrophoresis give a single precipitin arc against anti-goat serum as well as

purified and partially purified Cholesterol oxidase [microorganism].

Storage: Store lyophilized at 2-8°C for 6 months or at -20°C long term.

After reconstitution store the antibody undiluted at 2-8°C for one month or (in

aliquots) at -20°C long term.

Avoid repeated freezing and thawing. Shelf life: one year from despatch.



Pictures:

Goat anti Cholesterol oxidase antibody Cat.-No. R1075P (Lot 6534) was used to detect purified Cholesterol oxidase under reducing (R) and non-reducing (NR) conditions. Reduced samples of purified Cholesterol oxidase contained 4% BME and were boiled for 5 minutes. Samples of ~1µg of protein per lane were run by SDS-PAGE. Protein was transferred to nitrocellulose and probed with a 1/3000 dilution of primary antibody (4° C inblocking buffer). Detection shown was using Dylight 488 conjugated donkey anti goat. Images were collected using the BioRad VersaDoc System.

