

CYP21A2 rabbit pAb

Cat No.: ES4954

For research use only

Overview

Product Name CYP21A2 rabbit pAb

Host species Rabbit
Applications WB;ELISA

Species Cross-Reactivity Human;Rat;Mouse;

Recommended dilutions Western Blot: 1/500 - 1/2000. ELISA: 1/40000. Not

yet tested in other applications.

Immunogen Synthesized peptide derived from the Internal

region of human CYP21A2.

Specificity CYP21A2 Polyclonal Antibody detects endogenous

levels of CYP21A2 protein.

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name Steroid 21-hydroxylase

Gene Name CYP21A2

Cellular localization Endoplasmic reticulum membrane; Peripheral

membrane protein. Microsome membrane;

Peripheral membrane protein.

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal
Concentration 1 mg/ml
Observed band 55kD
Human Gene ID 1589
Human Swiss-Prot Number P08686

Alternative Names CYP21A2; CYP21; CYP21B; Steroid 21-hydroxylase;

21-OHase; Cytochrome P-450c21; Cytochrome P450 21; Cytochrome P450 XXI; Cytochrome P450-C21;

Cytochrome P450-C21B

Background cytochrome P450 family 21 subfamily A member

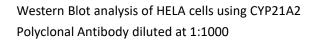
2(CYP21A2) Homo sapiens This gene encodes a member of the cytochrome P450 superfamily of

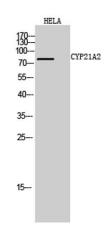


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enzymes. The cytochrome P450 proteins are monooxygenases which catalyze many reactions involved in drug metabolism and synthesis of cholesterol, steroids and other lipids. This protein localizes to the endoplasmic reticulum and hydroxylates steroids at the 21 position. Its activity is required for the synthesis of steroid hormones including cortisol and aldosterone. Mutations in this gene cause congenital adrenal hyperplasia. A related pseudogene is located near this gene; gene conversion events involving the functional gene and the pseudogene are thought to account for many cases of steroid 21-hydroxylase deficiency. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008],





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