ALDOA Antibody, Biotin conjugated



PACO24619

Reactivity:

Human

Source:

Product Information

Size: Protein Background:

50ug Plays a key role in glycolysis and gluconeogenesis. In addition, may also function as

scaffolding protein. Defects in ALDOA are the cause of glycogen storage disease type 12 (GSD12) [MIM:611881]; also known as red cell aldolase deficiency. A metabolic disorder

associated with increased hepatic glycogen and hemolytic anemia. It may lead to

myopathy with exercise intolerance and rhabdomyolysis. Belongs to the class I fructose-

bisphosphate aldolase family.

Rabbit Gene ID:

Isotype: ALDOA

lgG Uniprot

Applications: P04075

ELISA Synonyms:

Recommended dilutions: Fructose-bisphosphate aldolase A (EC 4.1.2.13) (Lung cancer antigen NY-LU-1) (Muscle-

type aldolase), ALDOA, ALDA

Immunogen:

Recombinant Human Fructose-bisphosphate aldolase A protein (2-364AA).

Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, PH 7.4

Product	Images
---------	---------------

N/A N/A