

GALE rabbit pAb

Cat No.: ES16281

For research use only

Overview

Product Name GALE rabbit pAb

Host species Rabbit
Applications WB

Species Cross-Reactivity Human; Mouse; Rat Recommended dilutions WB 1: 500-2000

Immunogen Synthesized peptide derived from human GALE AA

range: 104-154

Specificity This antibody detects endogenous levels of GALE at

Human/Mouse/Rat

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 GALE Gene Name GALE

Cellular localization cytosol, extracellular exosome,

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal Concentration 1 mg/ml

Observed band

Human Gene ID 2582 Human Swiss-Prot Number Q14376

Alternative Names

Background This gene encodes UDP-galactose-4-epimerase

which catalyzes two distinct but analogous reactions: the epimerization of UDP-glucose to UDP-galactose, and the epimerization of

UDP-N-acetylglucosamine to

UDP-N-acetylgalactosamine. The bifunctional nature

of the enzyme has the important metabolic

consequence that mutant cells (or individuals) are dependent not only on exogenous galactose, but



+86-27-59760950 ELKbio@ELKbiotech.com

www.elkbiotech.com



referred to as galactosemia type 3, a disease characterized by liver damage, early-onset cataracts, deafness and mental retardation, with symptoms ranging from mild ('peripheral' form) to severe ('generalized' form). Multiple alternatively spliced transcripts encoding the same protein have been identified. [provided by RefSeq, Jul 2008],

Western blot analysis of lysates from PC-12 cells, primary antibody was diluted at 1:1000, 4° over night

also on exogenous N-acetylgalactosamine as a

glycoproteins and glycolipids. Mutations in this gene result in epimerase-deficiency galactosemia, also

necessary precursor for the synthesis of



