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Datasheet for ABIN1701447

**anti-GALE antibody (AA 21-120) (Biotin)**

## Overview

Quantity:	100 µL
Target:	GALE
Binding Specificity:	AA 21-120
Reactivity:	Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GALE antibody is conjugated to Biotin
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Immunohistochemistry (Frozen Sections) (IHC (fro))

## Product Details

Immunogen:	KLH conjugated synthetic peptide derived from human GALE/Galactowaldenase
Isotype:	IgG
Cross-Reactivity:	Mouse
Predicted Reactivity:	Human,Rat,Pig,Horse
Purification:	Purified by Protein A.

## Target Details

Target:	GALE
Alternative Name:	GALE/Galactowaldenase ( <a href="#">GALE Products</a> )

## Target Details

Background:	<p>Synonyms: FLJ95174, FLJ97302, Galactose 4 epimerase UDP, Galactowaldenase, galE, GALE_HUMAN, OTTHUMP00000002991, OTTHUMP00000002994, OTTHUMP000000037931, OTTHUMP000000044857, SDR1E1, short chain dehydrogenase/reductase family 1E member 1, UDP galactose 4 epimerase, UDP glucose 4 epimerase, UDP-galactose 4-epimerase, UDP-glucose 4-epimerase.</p> <p>Background: GALE is a 348 amino acid protein that functions as the third enzyme in the Leloir pathway of galactose metabolism. A member of the sugar epimerase family, GALE exists as a homodimer, binds FAD as a cofactor and catalyzes the epimerization of UDP-N-acetylglucosamine to UDP-N-acetylgalactosamine and UDP-glucose to UDP-galactose. The gene encoding GALE maps to human chromosome 1p36.11 and mutations in this gene lead to the development of complex disorder known as epimerase-deficiency galactosemia (EDG) or galactosemia type 3, which is characterized by mental retardation, liver damage, cataracts and deafness.</p>
Gene ID:	2582
Pathways:	<a href="#">Response to Water Deprivation</a> , <a href="#">Cellular Glucan Metabolic Process</a>

## Application Details

Application Notes:	WB 1:300-5000 IHC-P 1:200-400 IHC-F 1:100-500
Restrictions:	For Research Use only

## Handling

Format:	Liquid
Concentration:	1 µg/µL
Buffer:	Aqueous buffered solution containing 0.01M TBS ( pH 7.4) with 1 % BSA, 0.03 % Proclin300 and 50 % Glycerol.
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20°C for 12 months.

Handling

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Expiry Date: 12 months