

Rabbit Anti-GATAD1 antibody

SL11921R

Product Name	GATAD1
Chinese Name	眼发育相关基因蛋白 ODAG 抗体
Alias	ODAG; GATA zinc finger domain containing 1; Ocular development associated gene; GATA zinc finger domain-containing protein 1; ocular development-associated gene protein; GATAD1 protein; ocular development associated; ODAG; RG083M05.2; tcag7.279; GATD1_HUMAN.
Research Area	Cell biology Neurobiology Signal transduction Growth factors and hormones Epigenetics
Immunogen Species	Rabbit
Clonality	Polyclonal
React Species	(predicted: Human, Mouse, Rat, Chicken, Dog, Pig, Cow, Horse, Rabbit, Zebrafish, Sheep,) ELISA=1:5000-10000
Applications	not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Theoretical molecular weight	29kDa
Cellular localization	The nucleus
Form	Liquid
Concentration	1mg/ml
immunogen	KLH conjugated synthetic peptide derived from human GATAD1/ODAG: 61-160/269
Lsotype	IgG
Purification	affinity purified by Protein A
Buffer Solution	1M TBS(pH7.4) with 1% BSA, 3% Proclin300 and 50% Glycerol.
Storage	Shipped at 4°C. Store at -20 °C for one year. Avoid repeated freeze/thaw cycles.
Attention	This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

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The protein encoded by this gene contains a zinc finger at the N-terminus, and is thought to bind to a histone modification site that regulates gene expression. Mutations in this gene have been associated with autosomal recessive dilated cardiomyopathy. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Jun 2012]

Function:

ODAG (Ocular development-associated gene), a novel transcription factor located on chromosome 7, encodes a protein that may play a role in eye development. mRNA profiling in multiple human tissue indicates that ODAG is expressed in human CD56+ NK cells and thyroid tissue.

Subunit:

Component of a chromatin complex, at least composed of KDM5A, GATAD1 and EMSY.

Subcellular Location:

Nuclear

Tissue Specificity:

Ubiquitously expressed among various tissue types. Expressed in left ventricular myocytes.

Product Detail

DISEASE:

Defects in GATAD1 are the cause of cardiomyopathy, dilated type 2B (CMD2B) [MIM:614672]. CMD2B is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Patients are at risk of premature death.

Similarity:

Contains 1 GATA-type zinc finger.

SWISS:

Q8WUU5

Gene ID:

57798

Database links:

[Entrez Gene: 57798](#) Human



[Entrez Gene: 67210](#) Mouse

[SwissProt: Q8N5Y5](#) Human

[SwissProt: Q8WUU5](#) Human

[SwissProt: Q3TS14](#) Mouse

[SwissProt: Q8VCQ2](#) Mouse

[SwissProt: Q920S3](#) Mouse

[SwissProt: Q9CSG2](#) Mouse

[Unigene: 21145](#) Human