abcam

Product datasheet

Anti-Sall4 antibody [EPR10642] ab181087

Recombinant RabMAb

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Overview

Product name Anti-Sall4 antibody [EPR10642]

Description Rabbit monoclonal [EPR10642] to Sall4

Host species Rabbit

Tested applications Suitable for: IHC-P Species reactivity Reacts with: Human

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control Human yolk sac carcinoma tissue

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity - Long-term security of supply - Animal-free production For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Properties

Form

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.5% BSA

Purity Protein A purified

Monoclonal Clonality Clone number EPR10642

Isotype lgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab181087 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P	★ 南 南 南 (1)	1/50 - 1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Target

Function Probable transcription factor.

Tissue specificity Expressed in testis.

Involvement in disease

Defects in SALL4 are the cause of Duane-radial ray syndrome (DRRS) [MIM:607323]; also known as Okihiro syndrome. DRRS is a disorder characterized by the association of forearm malformations with Duane retraction syndrome.

Defects in SALL4 are the cause of oculootoradial syndrome (OORS) [MIM:147750]. Oculootoradial syndrome is an autosomal dominant condition characterized by upper limbs anomalies (radial ray defects, carpal bones fusion), extraocular motor disturbances, congenital bilateral non-progressive mixed hearing loss. Other less consistent malformations include heart involvement, mild thrombocytopenia and leukocytosis (before age 50), shoulder girdle hypoplasia, imperforate anus, kidney malrotation or rectovaginal fistula. The IVIC syndrome is an allelic

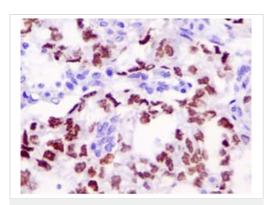
disorder of Duane-radial ray syndrome (DRRS) with a similar phenotype.

Sequence similaritiesBelongs to the sal C2H2-type zinc-finger protein family.

Contains 7 C2H2-type zinc fingers.

Cellular localization Nucleus.

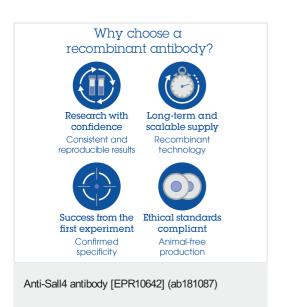
Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sall4 antibody
[EPR10642] (ab181087)

Immunohistochemical analysis of paraffin-embedded Human yolk sac carcinoma tissue labeling Sall4 with ab181087 at a 1/100 dilution.

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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