abcam

Product datasheet

Alexa Fluor® 488 Anti-SOX9 antibody [EPR12755] ab202686



2 Images

Overview

Product name Alexa Fluor® 488 Anti-SOX9 antibody [EPR12755]

Description Alexa Fluor® 488 Rabbit monoclonal [EPR12755] to SOX9

Host species Rabbit

Conjugation Alexa Fluor® 488, Ex: 495nm, Em: 519nm

Tested applications Suitable for: ICC/IF Species reactivity Reacts with: Human

Immunogen Recombinant fragment corresponding to Human SOX9 aa 400 to the C-terminus.

Database link: P48436

Run BLAST with Run BLAST with

Positive control ICC/IF: HepG2 cells.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit **General notes**

monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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Properties

Form Liquid

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

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: PBS, 1% BSA, 30% Glycerol (glycerin, glycerine)

Purity Protein A purified

Clonality Monoclonal
Clone number EPR12755

Isotype IgG

Applications

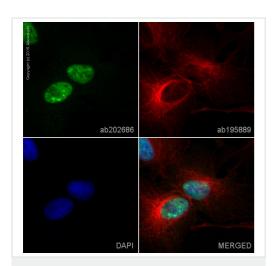
The Abpromise guarantee Our Abpromise guarantee covers the use of ab202686 in the following tested applications.

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

Application	Abreviews	Notes
ICC/IF		1/100. This product gave a positive signal in HepG2 cells fixed with 4% formaldehyde (10 min).

Target		
Function	Plays an important role in the normal skeletal development. May regulate the expression of other genes involved in chondrogenesis by acting as a transcription factor for these genes.	
Involvement in disease	Defects in SOX9 are the cause of campomelic dysplasia (CMD1) [MIM:114290]. CMD1 is a rare often lethal, dominantly inherited, congenital osteochondrodysplasia, associated with male-to-female autosomal sex reversal in two-thirds of the affected karyotypic males. A disease of the newborn characterized by congenital bowing and angulation of long bones, unusually small scapulae, deformed pelvis and spine and a missing pair of ribs. Craniofacial defects such as cle palate, micrognatia, flat face and hypertelorism are common. Various defects of the ear are ofter evident, affecting the cochlea, malleus incus, stapes and tympanum. Most patients die soon after birth due to respiratory distress which has been attributed to hypoplasia of the tracheobronchial cartilage and small thoracic cage.	
Sequence similarities	Contains 1 HMG box DNA-binding domain.	
Cellular localization	Nucleus.	

Images



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-SOX9 antibody [EPR12755] (ab202686)

ab202686 staining SOX9 in HepG2 cells. The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab202686 at 1/100 dilution (shown in green) and <u>ab195889</u>, Mouse monoclonal to alpha Tubulin (Alexa Fluor[®] 594), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



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