

FOXG1 Antibody

Catalog # ASC11891

Product Information

Application	WB, IF, E
Primary Accession	P55316
Other Accession	NP_005240 , 32307177
Reactivity	Human, Mouse, Rat
Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Calculated MW	52352
Concentration (mg/ml)	1 mg/mL
Conjugate	Unconjugated
Application Notes	FOXG1 antibody can be used for detection of FOXG1 by Western blot at 1 - 2 μ g/ml. For immunofluorescence start at 20 μ g/mL.

Additional Information

Gene ID	2290
Other Names	Forkhead box protein G1, Brain factor 1, BF-1, BF1, Brain factor 2, BF-2, BF2, hBF-2, Forkhead box protein G1A, Forkhead box protein G1B, Forkhead box protein G1C, Forkhead-related protein FKHL1, HFK1, Forkhead-related protein FKHL2, HFK2, Forkhead-related protein FKHL3, HFK3, FOXG1
Target/Specificity	FOXG1; FOXG1 antibody is human, mouse and rat reactive.
Reconstitution & Storage	FOXG1 antibody can be stored at 4°C for three months and -20°C, stable for up to one year.
Precautions	FOXG1 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name	FOXG1
Function	Transcription repression factor which plays an important role in the establishment of the regional subdivision of the developing brain and in the development of the telencephalon.
Cellular Location	Nucleus {ECO:0000255 PROSITE-ProRule:PRU00089, ECO:0000269 PubMed:21280142}
Tissue Location	Expression is restricted to the neurons of the developing telencephalon.

Background

FOXG1 belongs to the forkhead family of transcription factors which is characterized by a distinct forkhead domain and plays an important role in the development of the brain and telencephalon (1). It is essential for the proliferation of progenitor cells in the cerebral cortex and influences regional patterning in the mammalian telencephalon (2,3). Expression is restricted to the neurons of the developing telencephalon (3). The loss of FOXG1 results in an accelerated rate of neuronal differentiation and the shortening of the neurogenetic period in the embryonic cerebral cortex (2,4).

References

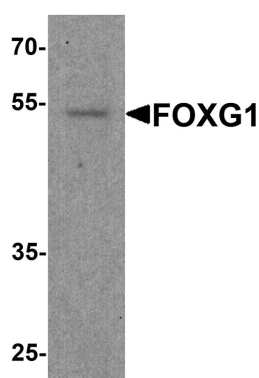
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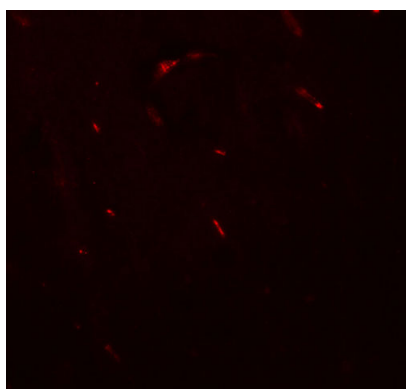
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Mencarelli MA, Spanhol-Rosseto A, Artuso R, et al. Novel FOXG1 mutations associated with the congenital variant of Rett syndrome. *J. Med. Genet.* 2010; 47:49-53.

Images



Western blot analysis of FOXG1 in rat brain tissue lysate with FOXG1 antibody at 1 µg/ml.



Immunofluorescence of FOXG1 in mouse brain tissue with FOXG1 antibody at 20 µg/ml.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.