

#### FOXI1 Antibody (Center) Blocking peptide Synthetic peptide

Catalog # BP12180c

# Specification

# FOXI1 Antibody (Center) Blocking peptide - Product Information

Primary Accession

### <u>Q12951</u>

# FOXI1 Antibody (Center) Blocking peptide - Additional Information

Gene ID 2299

**Other Names** 

Forkhead box protein I1, Forkhead-related protein FKHL10, Forkhead-related transcription factor 6, FREAC-6, Hepatocyte nuclear factor 3 forkhead homolog 3, HFH-3, HNF-3/fork-head homolog 3, FOXI1, FKHL10, FREAC6

### Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage** Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions** 

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

# FOXI1 Antibody (Center) Blocking peptide - Protein Information

Name FOXI1

Synonyms FKHL10, FREAC6

#### Function

Transcriptional activator required for the development of normal hearing, sense of balance and kidney function. Required for the expression of SLC26A4/PDS, JAG1 and COCH in a subset of epithelial cells and the development of the endolymphatic system in the inner ear. Also required for the expression of SLC4A1/AE1, SLC4A9/AE4, ATP6V1B1 and the differentiation of intercalated cells in the epithelium of distal renal tubules (By similarity).

Cellular Location Nucleus.

**Tissue Location** Expressed in kidney.



# FOXI1 Antibody (Center) Blocking peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

### Blocking Peptides

### FOXI1 Antibody (Center) Blocking peptide - Images

### FOXI1 Antibody (Center) Blocking peptide - Background

This gene belongs to the forkhead family of transcriptionfactors which is characterized by a distinct forkhead domain. Thespecific function of this gene has not yet been determined; however, it is possible that this gene plays an important role in the development of the cochlea and vestibulum, as well asembryogenesis. Mutations in this gene may be associated with the common cavity phenotype. Two transcript variants encoding differentisoforms have been found for this gene.

#### FOXI1 Antibody (Center) Blocking peptide - References

Jonard, L., et al. Int. J. Pediatr. Otorhinolaryngol. 74(9):1049-1053(2010)Moreno-Estrada, A., et al. BMC Evol. Biol. 10, 267 (2010) :Vidarsson, H., et al. PLoS ONE 4 (2), E4471 (2009) :Ferrell, R.E., et al. Lymphat Res Biol 6(2):69-76(2008)Rodriguez-Antona, C., et al. Mol. Pharmacol. 63(5):1180-1189(2003)