PRODUCT INFORMATION

Product name: PAX8 antibody

Product type: Primary antibodies

Description: Rabbit polyclonal to PAX8

Immunogen: 3 synthetic peptides (human) conjugated to KLH

Reacts with: Hu, Ms

Tested applications: ELISA, WB and IF

GENE INFORMATION

Gene Symbol: PAX8

Gene Name: paired box 8

Ensembl ID: ENSG00000125618

Entrez GeneID: 7849

GenBank Accession number: X69699.1

Omim ID: 167415 Swiss-Prot: Q06710

Molecular weight of PAX8: 29.083kDa

Function: Transcription factor for the thyroid-specific expression of the genes exclusively expressed in the thyroid cell type, maintaining the functional differentiation of such cells.

Expected subcellular localization: nucleus.

Expected tissue specificity: Expressed in the excretory system, thyroid gland and Wilms tumors.

Involvement in disease: Defects in PAX8 are the cause of congenital hypothyroidism non-goitrous type 2 (CHNG2) [MIM:218700]. CHNG2 is a disease characterized by thyroid dysgenesis, the most frequent cause of congenital hypothyroidism, accounting for 85% of case. The thyroid gland can be completely absent (athyreosis), ectopically located and/or severely hypoplastic. Ectopic thyroid gland is the most frequent malformation, with thyroid tissue being found most often at the base of the tongue.

Summary: This gene encodes a member of the paired box (PAX) family of transcription factors. Members of thes gene family typically encode proteins that contain a paired box domain, an octapeptide, and a paired-type homeodomain. This nuclear protein is involved in thyroid follicular cell development and expression of thyroid-specific genes. Mutations in this gene have been associated with thyroid dysgenesis, thyroid follicular carcinomas and atypical follicular thyroid

adenomas. Alternateively spliced transcript variants encoding different isoforms have been described. [provided by RefSeq]

APPLICATION NOTE

Recommended dilution:

- ELISA: Antibody specificity was verified by direct ELISA against the 3 immunogen peptides. A titer of 1/60000 has been determined. Appropriate specificity controls were run.
- WB: 1/5000.IF: 1/1000.

Optimal dilutions/concentration should be determined by the end user.

Raised in: Rabbit

Clonality: Polyclonal

Isotype: IgG

Purity: Purified Antibody

Storage buffer: 0.5 X PBS, containing a final concentration of 50% Glycerol, 0.1% BSA and

0.01% Thimerosal.

Form: Liquid

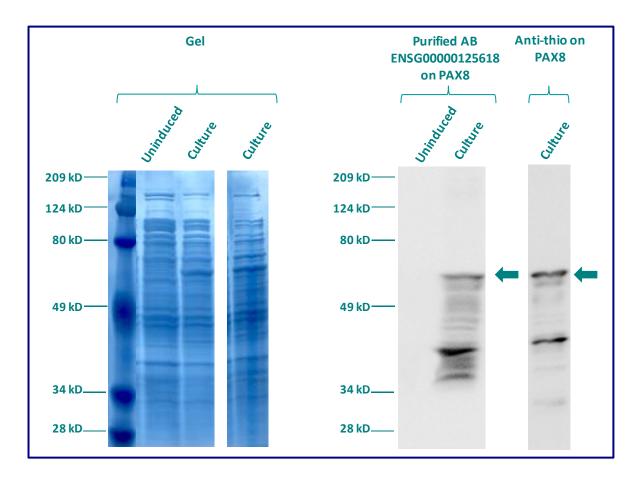
Storage instruction: Store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.

The purified antibody ENSG00000125618 has been tested at 1/5000 on uninduced (negative control) and induced culture of E.coli (one shot Top10 competent cells).

An anti-Thio (positive control) has been tested at 1/5000 on uninduced (negative control) and induced culture of E.coli (one shot Top10 competent cells) as a positive control.

Plasmid name: pBAD-DEST49.

Molecular weight of PAX8: 62.2kDa (48.2kDa + another 14kDa for the tag).



Gel concentration: 10%

Blocking: in 5% non-fat milk-PBST solution

1st Antibody: The antibodies are diluted in blocking buffer.

- Dilute the purified antibody ENSG00000125618 at 1:5000
- Dilute the anti-thio at 1:5000

60 minutes of incubation

2nd Antibody: The antibody is diluted in blocking buffer.

• Dilute the anti-Rabbit IgG HRP conjugated at 1/10000

60 minutes of incubation

IMMUNOFLUORESCENCE ANALYSIS

Immunofluorescence analysis of Paired box protein Pax-8 (PAX8) expression in 6 cells lines (HELA, 293T/17, Capan-2, SAOS2, SH-SY5Y, Skin 3,44). The purified Antibody ENSG00000125618 has been tested at 1/5000.

Red staining : cytoskeleton (microtubules/ α -tubuline)

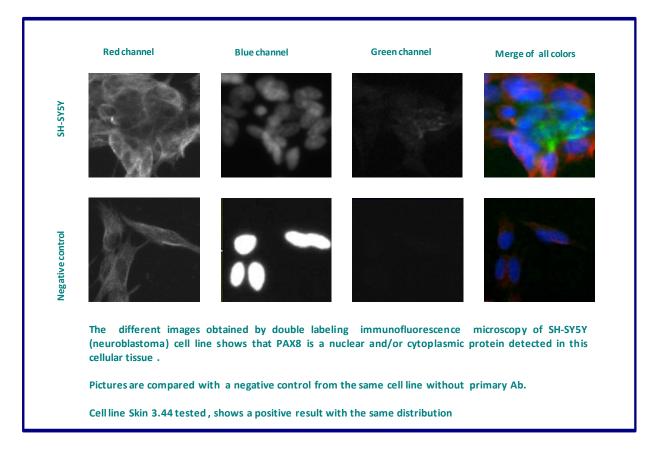
Blue staining: nucleus (Hoechst)

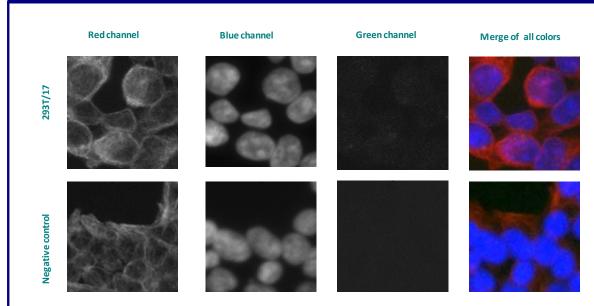
Green staining: anti-PAX8 antibody (purified)

Expected subcellular location: Nucleus

Expected tissue specificity: Expressed in the excretory system, thyroid gland and Wilms

tumors





The different images obtained by double labeling immunofluorescence microscopy of 293T/17 (kidney embrionic) cell line shows that PAX8 is not detected in this cellular tissue under these conditions.

Pictures are compared with a negative control from the same cell line without primary Ab.

Remaining cell lines tested gave a negative result under these conditions