

# Pax2 polyclonal antibody

Catalog: BS91021

Host: Rabbit

Reactivity: Human

## **BackGround:**

Pax genes contain paired domains with strong homology to genes in Drosophila which are involved in programming early development. The PAX2 gene is expressed in primitive cells of the kidney, ureter, eye, ear, and central nervous system. More specifically, in human embryo sections, PAX2 is expressed in the optic vesicle and later in the retina, in the otic vesicle and later in the semicircular canals of the inner ear, and in mesonephros, metanephros, adrenals, spinal cord, and hindbrain. PAX2 mutations can be responsible for renal hypoplasia, either isolated or associated with various ophthalmologic manifestations ranging from retinal coloboma to microphthalmia. Lesions in the PAX6 gene accounts for most cases of aniridia, a congenital malformation of the eye, chiefly characterized by iris hypoplasia, which can cause blindness. PAX6 is involved in other anterior segment malformations besides aniridia, such as Peters anomaly, a major error in the embryonic development of the eye with corneal clouding with variable iridolenticulocorneal adhesions.

#### **Product:**

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

**Molecular Weight:** 

## 45 kDa

**Swiss-Prot:** 

Q02962(Human)

## **Purification&Purity:**

ProA affinity purified

**Applications:** 

WB:1:1,000

IHC:1:50-1:100

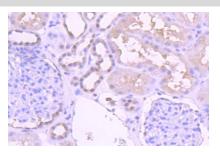
## **Storage&Stability:**

Store at +4  $^{\circ}$ C after thawing. Aliquot store at -20  $^{\circ}$ C or -80  $^{\circ}$ C. Avoid repeated freeze / thaw cycles.

## **Specificity:**

Pax2 polyclonal antibody detects endogenous levels of Pax2 protein.

## **DATA:**



Immunohistochemical analysis of paraffin-embedded human kidney

tissue using anti-Pax2 antibody. Counter stained with hematoxylin.

## Note:

For research use only, not for use in diagnostic procedure.

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