

# PAX6 (Stem Cell Marker) Antibody - With BSA and Azide

Mouse Monoclonal Antibody [Clone PAX6/1166 ]

Catalog # AH12033

## Product Information

---

|                   |   |
|-------------------|---|
| Application       | IHC, IF, FC                                   |
| Primary Accession | <a href="#">P26367</a>                        |
| Other Accession   | <a href="#">5080</a> , <a href="#">611376</a> |
| Reactivity        | Human   |
| Host              | Mouse   |
| Clonality         | Monoclonal                                    |
| Isotype           | Mouse / IgG1, kappa                           |
| Clone Names       | PAX6/1166                                     |
| Calculated MW     | 46683   |

## Additional Information

---

|                  |   |
|------------------|---|
| Gene ID          | 5080  |
| Other Names      | Paired box protein Pax-6, Aniridia type II protein, Oculorhombin, PAX6, AN2   |
| Application Note | IHC~~1:100~500 IF~~1:50~200 FC~~1:10~50   |
| Storage          | Store at 2 to 8°C.Antibody is stable for 24 months.   |
| Precautions      | PAX6 (Stem Cell Marker) Antibody - With BSA and Azide is for research use only and not for use in diagnostic or therapeutic procedures. |

## Protein Information

---

|                   |  |
|-------------------|--|
| Name              | PAX6   |
| Synonyms          | AN2  |
| Function          | Transcription factor with important functions in the development of the eye, nose, central nervous system and pancreas. Required for the differentiation of pancreatic islet alpha cells (By similarity). Competes with PAX4 in binding to a common element in the glucagon, insulin and somatostatin promoters. Regulates specification of the ventral neuron subtypes by establishing the correct progenitor domains (By similarity). Acts as a transcriptional repressor of NFATC1- mediated gene expression (By similarity). |
| Cellular Location | Nucleus {ECO:0000250 UniProtKB:P63015}. [Isoform 5a]: Nucleus {ECO:0000250 UniProtKB:P63016}   |
| Tissue Location   | [Isoform 1]: Expressed in lymphoblasts.  |

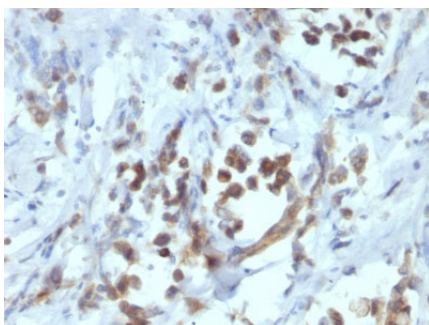
## Background

---

Pax genes contain paired domains with strong homology to genes in *Drosophila*, which are involved in programming early development. Lesions in the Pax-6 gene account for most cases of aniridia, a congenital malformation of the eye, chiefly characterized by iris hypoplasia, which can cause blindness. Pax-6 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. The Pax-6 gene encodes a transcriptional regulator that recognizes target genes through its paired-type DNA-binding domain. The paired domain is composed of two distinct DNA-binding subdomains, the amino-terminal subdomain and the carboxy-terminal subdomain, which bind respective consensus DNA sequences. The human Pax-6 gene produces two alternatively spliced isoforms that have the distinct structure of the paired domain.

## Images

---



Formalin-fixed, paraffin-embedded human Gastric Carcinoma stained with PAX6 Monoclonal Antibody (PAX6/1166).

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