

A Reliable Research Partner in Life Science and Medicine

## Recombinant p63(alpha) Monoclonal Antibody

catalog number: AN300136P

Note: Centrifuge before opening to ensure complete recovery of vial contents.

#### **Description**

Reactivity Human

**Immunogen** A synthetic peptide corresponding to the C-terminus of the human p63(alpha p63).

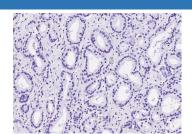
HostRabbitIsotypeIgGCloneA1250PurificationProtein A

Buffer 0.2 µm filtered solution in PBS

Applications Recommended Dilution

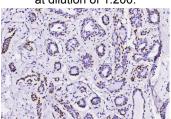
**IHC-P** 1:100-1:500

#### Data



Immunohistochemistry of paraffin-embedded human
prostatic carcinoma using p63(alpha) Monoclonal Antibody prostate using p63(alpha) Monoclonal Antibody at dilution of at dilution of 1:200.

Immunohistochemistry of paraffin-embedded human
prostatic carcinoma using p63(alpha) Monoclonal Antibody at dilution of 1:200.



Immunohistochemistry of paraffin-embedded human breast carcinoma using p63(alpha) Monoclonal Antibody at dilution of 1:200.

## **Preparation & Storage**

**Background** 

**Storage** This antibody can be stored at 2°C-8°C for one month without detectable loss of

activity. Antibody products are stable for twelve months from date of receipt when stored at -20°C to -80°C. Preservative-Free. Avoid repeated freeze-thaw cycles.

Shipping lce bag

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## For Research Use Only

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 Rev. V1.2

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This gene encodes a member of the p53 family of transcription factors. The functional domains of p53 family proteins include an N-terminal transactivation domain, a central DNA-binding domain and an oligomerization domain. Alternative splicing of this gene and the use of alternative promoters results in multiple transcript variants encoding different isoforms that vary in their functional properties. These isoforms function during skin development and maintenance, adult stem/progenitor cell regulation, heart development and premature aging. Some isoforms have been found to protect the germline by eliminating oocytes or testicular germ cells that have suffered DNA damage. Mutations in this gene are associated with ectodermal dysplasia, and cleft lip/palate syndrome 3 (EEC3); split-hand/foot malformation 4 (SHFM4); ankyloblepharon-ectodermal defects-cleft lip/palate; ADULT syndrome (acro-dermato-ungual-lacrimal-tooth); limb-mammary syndrome; Rap-Hodgkin syndrome (RHS); and orofacial cleft 5.

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