NBN Antibody



PACO18273

Reactivity:

Product Information

Size: Protein Background:

50ul Mutations in this gene are associated with Nijmegen breakage syndrome, an autosomal recessive chromosomal instability syndrome characterized by microcephaly, growth

retardation, immunodeficiency, and cancer predisposition. The encoded protein is a member of the MRE11/RAD50 double-strand break repair complex which consists of 5

Human, Mouse, Rat member of the MRE11/RAD50 double-strand break repair complex which consists proteins. This gene product is thought to be involved in DNA double-strand break

Source: repair and DNA damage-induced checkpoint activation.

Rabbit Gene ID:

Isotype: NBN

lgG Uniprot

Applications: O60934

ELISA, IHC **Synonyms:**

Recommended dilutions: nibrin

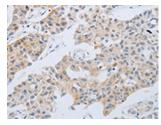
ELISA:1:2000-1:5000, IHC:1:25-1:100 **Immunogen:**

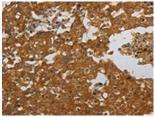
Synthetic peptide of human NBN.

Storage:

-20° C, pH7.4 PBS, 0.05% NaN3, 40% Glycerol

Product Images





The image on the left is immunohistochemistry of paraffin-embedded Human lung cancer tissue using PACO18273(NBN Antibody) at dilution 1/30, on the right is treated with synthetic peptide. (Original magnification: x—200).

The image on the left is immunohistochemistry of paraffin-embedded Human breast cancer tissue using PACO18273(NBN Antibody) at dilution 1/30, on the right is treated with synthetic peptide. (Original magnification: x—200).