Product Name: NBPF1/9/10/12/14/15/16/20 Rabbit

Polyclonal Antibody Catalog #: APRab14424



Summary

Production Name NBPF1/9/10/12/14/15/16/20 Rabbit Polyclonal Antibody

Description Rabbit Polyclonal Antibody

Host Rabbit
Application WB,ELISA

Reactivity Human, Rat, Mouse

Performance

ConjugationUnconjugatedModificationUnmodified

Isotype IgG

Clonality Polyclonal Form Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw Storage

cycles.

Buffer Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.

Purification Affinity purification

Immunogen

Gene Name NBPF12

NBPF12; COAS1; KIAA1245; Neuroblastoma breakpoint family member 12; Chromosome 1 amplified sequence 1; NBPF10; Neuroblastoma breakpoint family

Alternative Names

member 10; NBPF16; Neuroblastoma breakpoint family member 16; NBPF1; KIAA1693;

Neuroblastoma brea

Gene ID 55672/400818/284565/25832

Q5TAG4/Q6P3W6/Q5SXJ2/Q3BBV0/Q3BBW0/Q3BBV1/Q8N660/Q5TI25.The

SwissProt ID antiserum was produced against synthesized peptide derived from human

NBPF1/9/10/12/14/15/16/20. AA range:121-150

Application

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838

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Dilution Ratio WB 1:500 - 1:2000. ELISA: 1:40000..

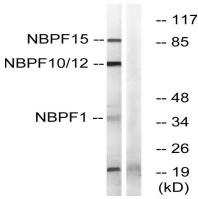
Molecular Weight 36kD

Background

This gene is a member of the neuroblastoma breakpoint family (NBPF) which consists of dozens of recently duplicated genes primarily located in segmental duplications on human chromosome 1. This gene family has experienced its greatest expansion within the human lineage and has expanded, to a lesser extent, among primates in general. Members of this gene family are characterized by tandemly repeated copies of DUF1220 protein domains. Gene copy number variations in the human chromosomal region 1q21.1, where most DUF1220 domains are located, have been implicated in a number of developmental and neurogenetic diseases such as microcephaly, macrocephaly, autism, schizophrenia, mental retardation, congenital heart disease, neuroblastoma, and congenital kidney and urinary tract anomalies. Altered expression of some gene family members is associated with several types of cancer. Thismiscellaneous:Encoded by one of the numerous copies of NBPF genes clustered in the p36, p12 and q21 region of the chromosome 1.,similarity:Belongs to the NBPF family.,similarity:Contains 10 NBPF domains.,similarity:Contains 2 NBPF domains.,similarity:Contains 3 NBPF domains.,similarity:Contains 6 NBPF domains.,similarity:Contains 7 NBPF domains.,similarity:Contains 8 NBPF domains.,tissue specificity:Expressed in a neuroblastoma cell line.,tissue specificity:Expressed in spinal cord.,tissue specificity:Expressed in spleen and fetal liver.,tissue specificity:Expressed in the mammary gland.,tissue specificity:Ubiquitously expressed with a higher expression observed in breast and liver. Also expressed in neuroblastoma cell line.,tissue specificity:Widely expressed. The only tissue which shows a weak expression is kidney.,

Research Area

Image Data



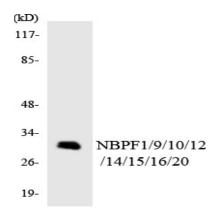
Western blot analysis of lysates from 293, HepG2, Jurkat, and COLO cells, using NBPF1/9/10/12/14/15/16/20 Antibody. The lane on the right is blocked with the synthesized peptide.

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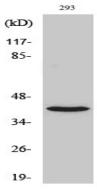
Product Name: NBPF1/9/10/12/14/15/16/20 Rabbit

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Western blot analysis of the lysates from HT-29 cells using NBPF1/9/10/12/14/15/16/20 antibody.



Western Blot analysis of various cells using NBPF1/9/10/12/14/15/16/20 Polyclonal Antibody

Note

For research use only.