

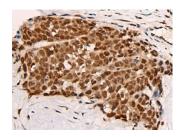
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HGH1 Polyclonal Antibody

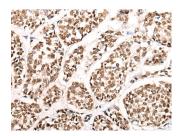
Catalog No.E-AB-17755ReactivityHStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsIHC,ELISAIsotypeIgG

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

Images



Immunohistochemistry of paraffinembedded Human thyroid cancer tissue using HGH1 Polyclonal Antibody at dilution of 1:45(×200)



Immunohistochemistry of paraffinembedded Human esophagus cancer tissue using HGH1 Polyclonal Antibody at dilution of 1:45(×200)

Immunogen Information

Immunogen Synthetic peptide of human HGH1

Gene Accession NP057542 **Swissprot** Q9BTY7

Synonyms Brain protein 16,BRP16,C8orf30A,D15Ertd741e,F20

3A,Fam203a,FLJ40907,LOC51236,MGC94185

Product Information

Buffer PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

Purify Antigen affinity purification

Dilution IHC 1:30-1:150, ELISA 1:5000-1:10000

Background

HGH1, is also known as BRP16, Brp16 is a 256 amino acid protein encoded by a gene on human chromosome 8. Made up of nearly 146 million bases, chromosome 8 encodes about 800 genes. Translocation of portions of chromosome 8 with amplifications of the c-Myc gene are found in some leukemias and lymphomas, and typically associated with a poor prognosis. Portions of chromosome 8 have been linked to schizophrenia and bipolar disorder. Trisomy 8, also known as Warkany syndrome 2, most often results in early miscarriage but is occasionally seen in a mosaic form in surviving patients who suffer to a varying degree from a number of symptoms including retarded mental and motor development, and certain facial and developmental defects. WRN is a DNA helicase encoded by chromosome 8 and shown defective in those with the early aging disorder Werner syndrome. Chromosome 8 is also associated with Pfeiffer syndrome, congenital hypothyroidism and Waardenburg syndrome.