

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

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- Trockeneiszuschlag
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- Expressversand

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HSPA1L & TP53 Protein Protein Interaction Antibody Pair

Catalog #: DI0459 規格:[1 Set]

List All

Specification

Product Description:

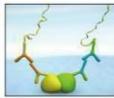
This protein protein interaction antibody pair set comes with two antibodies to detect the protein-protein interaction, one against the HSPA1L protein, and the other against the TP53 protein for use in *in* situ Proximity Ligation Assay. See Publication Reference below.

Application Image

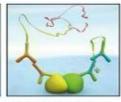
In situ Proximity Ligation Assay (Cell)



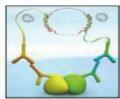
1. Incubate with target primary antibodies



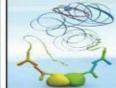
2. Add PLA probes PLUS and MINUS



3. Hybridize connector oligos



4. Ligation to form a complete DNA circle



5. Rolling circle amplification



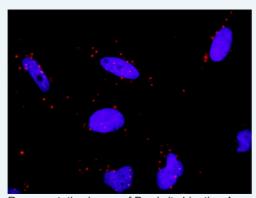
6. Add fluorescent probes to reveal interaction

Reactivity:

Human

Quality Control Protein protein interaction immunofluorescence result.

Testing:



Representative image of Proximity Ligation Assay of protein-protein interactions between HSPA1L and TP53. HeLa cells were stained with anti-HSPA1L rabbit purified polyclonal antibody 1:1200 and anti-TP53 mouse monoclonal antibody 1:50. Each red dot represents the detection of protein-protein interaction complex. The images were analyzed using an optimized freeware (BlobFinder) download from The Centre for Image Analysis at Uppsala University.

Supplied **Product:**

Antibody pair set content:

- 1. HSPA1L rabbit purified polyclonal antibody (20 ug)
- 2. TP53 mouse monoclonal antibody (40 ug)

*Reagents are sufficient for at least 30-50 assays using recommended protocols.

Storage Instruction:

Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -

2016/5/20

MSDS:

Download

Proteomics. 2013 Feb 8. [Epub ahead of print]

Publication Reference

 An analysis of protein-protein interactions in cross-talk pathways reveals CRKL as a novel prognostic marker in hepatocellular carcinoma.
Liu CH, Chen TC, Chau GY, Jan YH, Chen CH, Hsu CN, Lin KT, Juang YL, Lu PJ, Cheng HC, Chen MH, Chang CF, Ting YS, Kao CY, Hsiao M, Huang CY. Mol Cell

Applications

In situ Proximity Ligation Assay (Cell)

HSPA1L TP53

Gene Information

Entrez GeneID: 3305

Gene Name: HSPA1L

Gene Alias: HSP70-1L,HSP70-HOM,HSP70T,hum70t

Gene heat shock 70kDa protein 1-like

Description:

Omim ID: <u>140559</u>

Gene Ontology: Hyperlink

Gene Summary: This gene encodes a 70kDa heat shock protein. In conjunction with

other heat shock proteins, this protein stabilizes existing proteins against aggregation and mediates the folding of newly translated proteins in the cytosol and in organelles. The gene is located in the major histocompatibility complex class III region, in a cluster with two closely related genes which also encode isoforms of the 70kDa heat

shock protein. [provided by RefSeq

Other OTTHUMP00000029295,heat shock 10kDa protein 1-like,heat shock

Designations: 70kD protein-like 1

Gene Information

Entrez GeneID: 7157

Gene Name: TP53

Gene Alias: FLJ92943,LFS1,TRP53,p53

Gene tumor protein p53

Description:

Omim ID: 114480, 114500, 114550, 151623, 161550, 191170, 202300, 260350

Gene Ontology: Hyperlink

Gene Summary: This gene encodes tumor protein p53, which responds to diverse

cellular stresses to regulate target genes that induce cell cycle arrest, apoptosis, senescence, DNA repair, or changes in metabolism. p53 protein is expressed at low level in normal cells and at a high level in a variety of transformed cell lines, where it's believed to contribute to transformation and malignancy. p53 is a DNA-binding protein containing transcription activation, DNA-binding, and oligomerization domains. It is

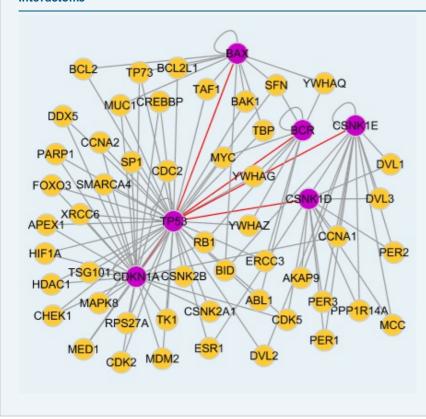
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postulated to bind to a p53-binding site and activate expression of downstream genes that inhibit growth and/or invasion, and thus function as a tumor suppressor. Mutants of p53 that frequently occur in a number of different human cancers fail to bind the consensus DNA binding site, and hence cause the loss of tumor suppressor activity. Alterations of this gene occur not only as somatic mutations in human malignancies, but also as germline mutations in some cancer-prone families with Li-Fraumeni syndrome. Multiple p53 variants due to alternative promoters and multiple alternative splicing have been found. These variants encode distinct isoforms, which can regulate p53 transcriptional activity. [provided by RefSeq

Other Designations:

p53 antigen,p53 transformation suppressor,p53 tumor suppressor,phosphoprotein p53,transformation-related protein 53

Interactome



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