A Reliable Research Partner in Life Science and Medicine

DJ-1/PARK7 Polyclonal Antibody

catalog number: E-AB-92096

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

Description

Reactivity Human; Mouse; Rat

Immunogen Recombinant fusion protein of human DJ-1/PARK7

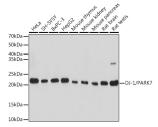
Host Rabbit Isotype IgG

Purification Affinity purification

Buffer Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

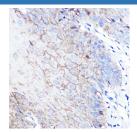
Applications	Recommended Dilution	
WB	1:500-1:2000	
IHC	1:50-1:200	
IF	1:50-1:200	

Data

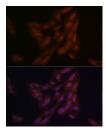


Western blot analysis of extracts of various cell lines using DJ-1/PARK7 Polyclonal Antibody at 1:1000 dilution.

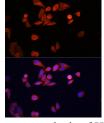
Observed-MW:22 kDa Calculated-MW:19 kDa



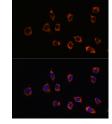
Immunohistochemistry of paraffin-embedded human esophageal using DJ-1/PARK7 Polyclonal Antibody at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.



Immunofluorescence analysis of C6 cells using DJ-1/PARK7 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.



Immunofluorescence analysis of HeLa cells using DJ-1/PARK7 Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.



For Research Use Only

Fax: 1-832-243-6017

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Immunofluorescence analysis of L929 cells using DJ-1/PARK7 Polyclonal Antibody at dilution of 1:100 (40x

lens). Blue: DAPI for nuclear staining.

Preparation & Storage

Storage Storage Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.

Shipping The product is shipped with ice pack, upon receipt, store it immediately at the

temperature recommended.

Background

The product of this gene belongs to the peptidase C56 family of proteins. It acts as a positive regulator of androgen receptor-dependent transcription. It may also function as a redox-sensitive chaperone, as a sensor for oxidative stress, and it apparently protects neurons against oxidative stress and cell death. Defects in this gene are the cause of autosomal recessive early-onset Parkinson disease 7. Two transcript variants encoding the same protein have been identified for this gene.

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