

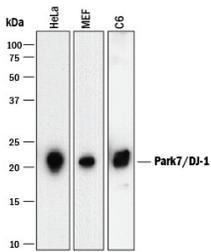
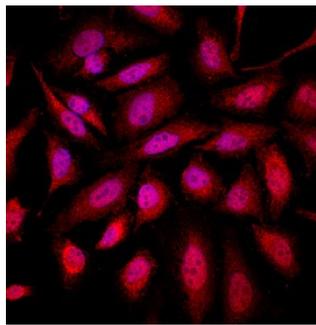
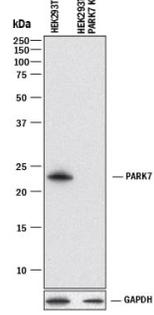
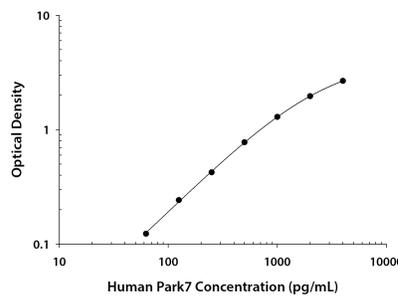
DESCRIPTION	
<b>Species Reactivity</b>	Human/Mouse/Rat
<b>Specificity</b>	Detects human, mouse, and rat PARK-7 in Western blots. In sandwich immunoassays, it is specific for human PARK-7.
<b>Source</b>	Recombinant Monoclonal Mouse IgG <sub>2A</sub> Clone # 925805R
<b>Purification</b>	Protein A or G purified from cell culture supernatant
<b>Immunogen</b>	<i>E. coli</i> -derived recombinant human PARK-7 Ala2-Asp189 Accession # Q99497
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

## APPLICATIONS

**Please Note:** Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
<b>Western Blot</b>	1 µg/mL	See Below
<b>Immunocytochemistry</b>	8-25 µg/mL	See Below
<b>Knockout Validated</b>	Park7/DJ-1 is specifically detected in HEK293T human embryonic kidney parental cell line but is not detectable in Park7/DJ-1 knockout HEK293T cell line.	
<b>ELISA</b>	This antibody functions as an ELISA detection antibody for Human Park7//DJ-1 when paired with Mouse Anti-Human Park7/DJ-1 Monoclonal Antibody (Catalog # MAB39952).	
	<i>This product is intended for assay development on various assay platforms requiring antibody pairs. We recommend the Human Park7/DJ-1 DuoSet ELISA Kit (Catalog # DY3995) for convenient development of a sandwich ELISA.</i>	

## DATA

<p><b>Western Blot</b></p>  <p><b>Detection of Human, Mouse, and Rat Park7/DJ-1 by Western Blot.</b> Western blot shows lysates of HeLa human cervical epithelial carcinoma cell line, MEF mouse embryonic feeder cells, and C6 rat glioma cell line. PVDF membrane was probed with 1 µg/mL of Mouse Anti-Human/Mouse/Rat Park7/DJ-1 Monoclonal Antibody (Catalog # MAB39951) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # HAF018). A specific band was detected for Park7/DJ-1 at approximately 22 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.</p>	<p><b>Immunocytochemistry</b></p>  <p><b>Park7/DJ-1 in HeLa Human Cell Line.</b> Park7/DJ-1 was detected in immersion fixed HeLa human cervical epithelial carcinoma cell line using Mouse Anti-Human/Mouse/Rat Park7/DJ-1 Monoclonal Antibody (Catalog # MAB39951) at 10 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Mouse IgG Secondary Antibody (red; Catalog # NL007) and counterstained with DAPI (blue). Specific staining was localized to cytoplasm and nuclei. View our protocol for <a href="#">Fluorescent ICC Staining of Cells on Coverslips</a>.</p>
<p><b>Knockout Validated</b></p>  <p><b>Western Blot Shows Human Park7/DJ-1 Specificity by Using Knockout Cell Line.</b> Western blot shows lysates of HEK293T human embryonic kidney parental cell line and Park7 knockout HEK293T cell line (KO). PVDF membrane was probed with 1 µg/mL of Mouse Anti-Human/Mouse/Rat Park7/DJ-1 Monoclonal Antibody (Catalog # MAB39951) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # HAF018). A specific band was detected for Park7/DJ-1 at approximately 23 kDa (as indicated) in the parental HEK293T cell line, but is not detectable in knockout HEK293T cell line. GAPDH (Catalog # MAB5718) is shown as a loading control. This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.</p>	<p><b>ELISA</b></p>  <p><b>Human Park7/DJ-1 ELISA Standard Curve.</b> Recombinant Human Park7/DJ-1 protein was serially diluted 2-fold and captured by Mouse Anti-Human Park7/DJ-1 Monoclonal Antibody (Catalog # MAB39952) coated on a Clear Polystyrene Microplate (Catalog # DY990). Mouse Anti-Human/Mouse/Rat Park7/DJ-1 Monoclonal Antibody (Catalog # MAB39951) was biotinylated and incubated with the protein captured on the plate. Detection of the standard curve was achieved by incubating Streptavidin-HRP (Catalog # DY998) followed by Substrate Solution (Catalog # DY999) and stopping the enzymatic reaction with Stop Solution (Catalog # DY994).</p>

## PREPARATION AND STORAGE

**Reconstitution** Reconstitute at 0.5 mg/mL in sterile PBS.

**Shipping** The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.  
\*Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C

**Stability & Storage** Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.

## BACKGROUND

Park7, also known as DJ-1, is a cytoplasmic protein that belongs to the ThiJ/Pfp1/DJ-1 superfamily of highly conserved proteins that function as protein chaperones, catalases, proteases and kinases. Park7 is widely expressed in the brain as well as in peripheral tissues. It exists as a homodimer that can be localized in the cytoplasm, nucleus and mitochondria. Park7 is a redox-sensitive protein that has been ascribed various functions including that as a redox sensor and antioxidant protein. Mutations in Park7 are associated with a small percentage of hereditary early onset Parkinson's disease. Human and mouse Park7 share 92% amino acid sequence identity.