

# Product Datasheet

## LRRK2 Antibody - BSA Free

### NB110-58771SS

Unit Size: 0.025 ml

Store at 4C. Do not freeze.

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**NB110-58771SS**

LRRK2 Antibody - BSA Free

**Product Information**

|                                |                             |
|--------------------------------|-----------------------------|
| <b>Unit Size</b>               | 0.025 ml                    |
| <b>Concentration</b>           | 1 mg/ml                     |
| <b>Storage</b>                 | Store at 4C. Do not freeze. |
| <b>Clonality</b>               | Polyclonal                  |
| <b>Preservative</b>            | 0.02% Sodium Azide          |
| <b>Isotype</b>                 | IgG                         |
| <b>Purity</b>                  | Immunogen affinity purified |
| <b>Buffer</b>                  | PBS                         |
| <b>Target Molecular Weight</b> | 286 kDa                     |

**Product Description**

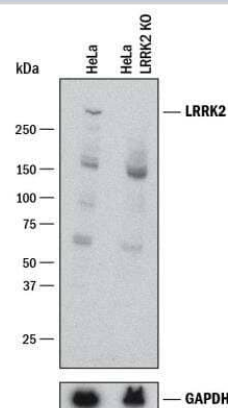
|                    |  |
|--------------------|--|
| <b>Host</b>        | Rabbit   |
| <b>Gene ID</b>     | 120892   |
| <b>Gene Symbol</b> | LRRK2  |
| <b>Species</b>     | Human, Mouse, Rat, Primate   |
| <b>Immunogen</b>   | A synthetic peptide made to a C-terminal region, within residues 2500-2527 of the human LRRK2 protein, conjugated to KLH. [Swiss-Prot# Q5S007] |

**Product Application Details**

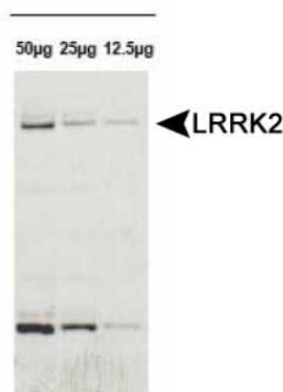
|                              |   |
|------------------------------|---|
| <b>Applications</b>          | Western Blot, Flow (Cell Surface), Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Frozen, Immunohistochemistry-Paraffin, Immunoprecipitation, Knockdown Validated, Knockout Validated  |
| <b>Recommended Dilutions</b> | Western Blot 1:1000, Immunohistochemistry 1:200-1:500, Immunocytochemistry/ Immunofluorescence 1:500-1:1000, Immunoprecipitation 1:10-1:500, Immunohistochemistry-Paraffin reported in scientific literature (PMID 24312256), Immunohistochemistry-Frozen, Flow (Cell Surface) reported in scientific literature (PMID 20483355), Knockout Validated, Knockdown Validated |

**Images**

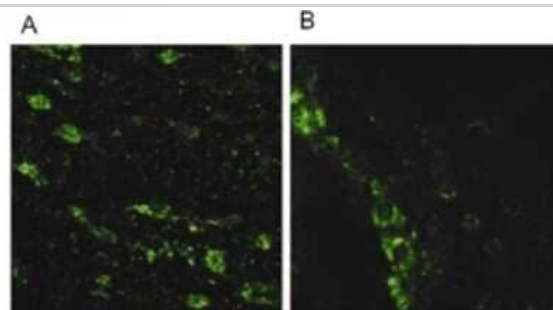
Western Blot: LRRK2 Antibody [NB110-58771] - Western blot shows lysates of HeLa human cervical epithelial carcinoma parental cell line and LRRK2 knockout (KO) HeLa cell line. PVDF membrane was probed with 1:1000 of Rabbit Anti-Human LRRK2 Polyclonal Antibody (Catalog # NB110-58771) followed by HRP-conjugated Anti-Rabbit IgG Secondary Antibody (Catalog #HAF008). Specific band was detected for LRRK2 at approximately 275 kDa (as indicated) in the parental HeLa cell line, but is not detectable in the knockout HeLa cell line. This experiment was conducted under reducing conditions.



Western Blot: LRRK2 Antibody [NB110-58771] - NB110-58771 in increasing amounts of human lymphoblast



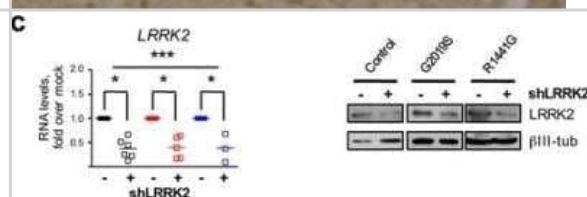
Immunocytochemistry/Immunofluorescence: LRRK2 Antibody [NB110-58771] - Immunofluorescence of Lrrk2 with NB110-58771 within (A) Substantia nigra neurons (B) Chains of migrating neuroblasts.



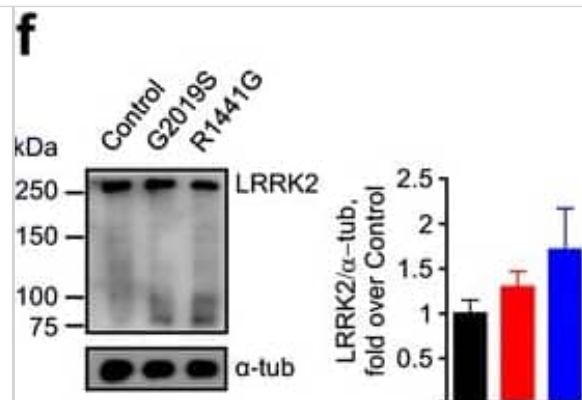
Immunohistochemistry-Frozen: LRRK2 Antibody [NB110-58771] - DAB Immunostaining in frozen sections of mouse brainstem.



Western Blot: LRRK2 Antibody [NB110-58771] - LRRK2 RNA levels 5 days after shRNA lentiviral transduction. The empty vector was used as the mock control. Lines represent the mean  $\pm$  SEM of three to four independent silencing experiments. A representative immunoblot analysis of LRRK2 protein levels 5 days after transduction is also shown. Beta-III-tubulin was the loading control. Image collected and cropped by CiteAb from the following publication (<https://jneuroinflammation.biomedcentral.com/articles/10.1186/s12974-016-0761-x>) licensed under a CC-BY license.



**Western Blot: LRRK2 Antibody [NB110-58771] - Characterization of iPSC-derived DA neurons with LRRK2 mutations.** a Diagram showing the DA differentiation protocol used for neural induction of human iPSC lines. b Temporal gene expression analyzed by qRT-PCR at three time points: induction (3 weeks), expansion (4–5 weeks), & maturation (>6 weeks). Each point represents the mean  $\pm$  SEM of at least two independent differentiation experiments. c Representative images of mature neuronal cultures showing expression of neuronal ( $\beta$ III-tubulin, Tau, &  $\alpha$ -synuclein) & dopaminergic (TH, NURR1) markers. Nuclei were counterstained with Hoechst. Scale bars: 50  $\mu$ m. d Quantification of immunostainings. Data are represented as mean  $\pm$  SEM of counts from at least two different lines for each genotype. e Representative western blot analyses of TH, Tau, & GFAP with  $\beta$ III-tubulin as loading control in iPSC-derived mature neurons. f Representative immunoblots & quantification of LRRK2 expression in mature neuronal cultures.  $\alpha$ -tubulin was the loading control & data were normalized to control WT neurons. Bars represent the mean  $\pm$  SEM of at least two different lines per genotype. DIV days in vitro, GEL gelatin, POL poly-ornithine, FBN fibronectin, LMN laminin, N2 N2 supplement, bFGF basic fibroblast growth factor, SAG smoothened agonist, LDN LDN-193189, CHIR CHIR99021, SB SB431542, BDNF brain-derived neurotrophic factor, AA ascorbic acid, B27 B27 supplement, dbcAMP dibutyryl cyclic adenosine monophosphate, TGF $\beta$ III transforming growth factor  $\beta$ III, GDNF glial derived neurotrophic factor. See Additional file 2 for uncropped blots Image collected & cropped by CiteAb from the following publication (<http://jneuroinflammation.biomedcentral.com/articles/10.1186/s12974-016-0761-x>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



## Publications

- Labib M, Wang Z, Kim Y et al. Identification of druggable regulators of cell secretion via a kinome-wide screen and high-throughput immunomagnetic cell sorting Nature biomedical engineering 2023-11-27 [PMID: 38012306]
- Lopez de Maturana R, Lang V, Zubiarrain A et al. Mutations in LRRK2 impair NF- $\kappa$ B pathway in iPSC-derived neurons. J Neuroinflammation. 2016-11-18 [PMID: 27863501] (WB, Human)
- Skibinski G, Nakamura K, Cookson MR, Finkbeiner S. Mutant LRRK2 toxicity in neurons depends on LRRK2 levels and synuclein but not kinase activity or inclusion bodies. J Neurosci. 2014-01-08 [PMID: 24403142] (ICC/IF)
- Nakamori M, Takahashi T, Nishikawa T et al. Molecular Markers for Granulovacuolar Degeneration Are Present in Rimmed Vacuoles. PLoS One. 2013-11-28 [PMID: 24312256] (IHC-P, ICC/IF, Human)
- Melrose H. Update on the functional biology of Lrrk2. Future Neurol. 2008-01-01 [PMID: 19225574] (WB, Mouse)
- Melrose HL, Dachsel JC, Behrouz B et al. Impaired dopaminergic neurotransmission and microtubule-associated protein tau alterations in human LRRK2 transgenic mice. Neurobiol Dis 2010-12-01 [PMID: 20659558] (WB, Mouse)
- Melrose, HL et al. A comparative analysis of leucine-rich repeat kinase 2 (Lrrk2) expression in mouse brain Lewy Body disease. Neurosci 147: 1047-1058. 2007-01-01 [PMID: 17611037] (IF/IHC, WB, Mouse)
- Lee H, Melrose HL, Yue M et al. Lrrk2 localization in the primate basal ganglia and thalamus: a light and electron microscopic analysis in monkeys. Exp Neurol;224(2):438-47. 2010-08-01 [PMID: 20483355] (IF/IHC, WB, ICC/IF, Primate)

## Procedures

### Serum protocol for LRRK2 Antibody (NB110-58771)

LRRK2 Antibody:

Immunostaining of frozen sections using streptavidin or Exvtravidin peroxidase and DAB

1. Make 10 micron frozen sections.
2. Fix tissue sections on slides in cold acetone, 10% buffered folrmalin or 4% paraformaldehyde for 8 to 10 min.
3. Rinse slides with PBS by immersion for 2 min. Repeat.
4. Block all slides with 5% milk in PBST for 30 min. at RT.
5. Remove blocking solution and add NB 110-58771 LRRK2 primary antibody 1:100 to 1:200 in PBS with 0.1% triton X-100 with 5% milk.
6. Incubate slides at 4degrees Celcius, overnight.
7. Rinse slides with PBST by immersion for 2 min. Repeat.
8. Add biotinylated donkey anti-rabbit secondary antibody (1:500 in PBST with 5% milk). Incubate 1 hour at RT.
9. Rinse with PBST by immersion 2 min. Repeat.
10. Add streptavidin-HRP or Extravidin-HRP. Incubate at RT for 1 hr.
11. Rinse with PBS by immersion for 1 min. Repeat.
12. Make DAB solution.
13. Add DAB to slides. Incubate 5-7 minutes.
14. Stop DAB reaction by immersion in water.
15. Counterstain sections, if desired.
16. Dehydrate,defat and coverslip slides.





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### **Limitations**

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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