Mouse LRRN1 Protein

Cat. No. LNR-MM101

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| Description | |
|-------------------------|--|
| Source | Recombinant Mouse LRRN1 Protein is expressed from HEK293 with His tag at the C-Terminus. |
| | It contains Ser26-Ala631. |
| Accession | Q61809 |
| Molecular Weight | The protein has a predicted MW of 69.28 kDa. Due to glycosylation, the protein migrates to 80-90 kDa based on Tris-Bis PAGE result. |
| Endotoxin | Less than 1EU per µg by the LAL method. |
| Purity | > 95% as determined by Tris-Bis PAGE |
| | > 95% as determined by HPLC |
| Formulation and Storage | |
| Formulation | Lyophilized from 0.22µm filtered solution in PBS (pH 7.4). Normally 8% trehalose is added as protectant before lyophilization. |
| Reconstitution | Centrifuge the tube before opening. Reconstituting to a concentration more than 100 µg/ml is recommended. Dissolve the lyophilized protein in distilled water. |
| Storage | -20 to -80°C for 12 months as supplied from date of receipt20 to -80°C for 3-6 months in unopened state after reconstitution.2-8°C for 2-7 days after reconstitution.Recommend to aliquot the protein into smaller quantities for optimal storage. Please minimize freeze-thaw cycles. |
| Background | |
| | Lrrn1 is required for the formation of MHBloss of function leads to a loss of the morphological constriction and loss of Fgf8. Cells overexpressing Lrrn1 violate the boundary and result in a loss of cell restriction between midbrain and hindbrain compartments. Lrrn1 also regulates the glycosyltransferase Lunatic Fringe, a modulator of Notch signalling, maintaining its expression in midbrain cells which is instrumental in MHB boundary formation. |

Assay Data

Tris-Bis PAGE



Mouse LRRN1 on Tris-Bis PAGE under reduced condition. The purity is greater than 95%.

SEC-HPLC

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Assay Data





The purity of Mouse LRRN1 is greater than 95% as determined by SEC-HPLC.