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Regulation of DJ-1 by glutaredoxin 1 *in vivo* – implications for Parkinson's disease

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Abstract

Parkinson's disease (PD) is the second most common neurodegenerative disease worldwide, caused by the degeneration of the dopaminergic neurons in the *substantia nigra*. Mutations in *PARK7* (DJ-1) result in early onset autosomal recessive PD, and oxidative modification of DJ-1 has been reported to regulate the protective activity of DJ-1 *in vitro*. Glutathionylation is a prevalent redox modification of proteins resulting from the disulfide adduction of the glutathione moiety to a reactive cysteine-SH; and glutathionylation of specific proteins has been implicated in regulation of cell viability. Glutaredoxin 1 (Grx1) is the principal deglutathionylating enzyme within cells, and it has been reported to mediate protection of dopaminergic neurons in *C. elegans*, however many of the functional downstream targets of Grx1 *in vivo* remain unknown. Previously, DJ-1 protein content was shown to decrease concomitantly with diminution of Grx1 protein content in cell culture of model neurons (SH-SY5Y and Neuro-2A lines). In the current study we aimed to investigate the regulation of DJ-1 by Grx1 *in vivo* and characterize its glutathionylation *in vitro*. Here, with *Grx1*^{-/-} mice we provide evidence that Grx1 regulates protein levels of DJ-1 *in vivo*. Furthermore, with model neuronal cells (SH-SY5Y) we observed decreased DJ-1 protein content in response to treatment with known glutathionylating agents; and with isolated DJ-1 we identified two distinct sites of glutathionylation. Finally, we found that overexpression of DJ-1 in

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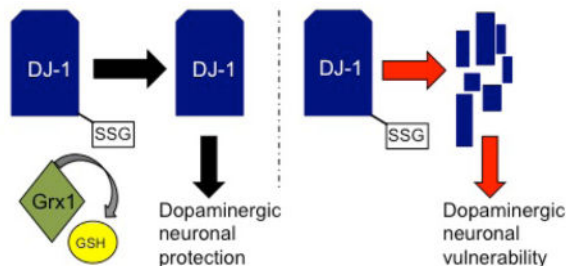
Conflict of Interest statement: None declared

Supporting Information Available

Supplemental items of data include western blot and qPCR analyses for Grx1^{-/-} and hGrx1-transgenic mice, western blot analyses of human samples, validation of purity and specificity of antibodies, HPLC-mass spectrometric chromatograms, fluorescence data for quantification of expression of exogenous DJ-1 in *C. elegans*, and cell culture data for individual components of the proteolytic inhibitor cocktail. This material is available free of charge via the Internet at <http://pubs.acs.org>

the dopaminergic neurons partly compensates for the loss of the Grx1 homolog in a *C. elegans in vivo* model of PD. Therefore; our results reveal a novel redox modification of DJ-1 and suggest a novel regulatory mechanism for DJ-1 content *in vivo*.

Graphical Abstract



Keywords

Redox; Parkinson's disease; glutaredoxin; DJ-1; glutathionylation; *C. elegans*

Introduction

Parkinson's disease (PD) is a debilitating neurodegenerative disease caused by the loss of dopaminergic neurons in the *substantia nigra* region of the midbrain [1, 2]. Although the etiology of PD is unclear, PD can be categorized into two groups: familial and sporadic. These classifications are based on whether PD is attributed to a genetic mutation or if the cause of the disease is undefined. In both cases oxidative stress and dysregulation of redox homeostasis have been implicated.

LRRK2 (Leucine-Rich Repeat Kinase 2) and DJ-1 (PARK7) are two proteins in which genetic mutations are linked to development of PD [3, 4]. Mutations in LRRK2 are shown to cause autosomal dominant PD in humans, and over-expression of mutant LRRK2 results in dysfunction of dopamine signaling or degeneration of dopaminergic neurons in animal models of PD [5, 6, 7, 8, 9]. Mutations in DJ-1 result in autosomal recessive, early onset PD in humans, through destabilization and loss of DJ-1 protein content [10]. In addition to mutations driving familial PD, post mortem brain samples from patients with sporadic PD have been reported to have decreased levels of DJ-1 protein within several areas of the brain including the nigral region [11]. Although studies have implicated DJ-1 as an anti-oxidant factor [12, 13], the molecular mechanism(s) by which DJ-1 protects cells from oxidative stress-induced death is currently unresolved [14]. Genetic manipulation has revealed a key residue on DJ-1, which is critical for the protection against oxidative stressors. Substitution of cysteine 106 with alanine (C106A) results in a "non-functional" DJ-1; over-expression of WT, but not C106A DJ-1, protects *Park7^{-/-}* mouse embryonic fibroblasts from H₂O₂-induced cell death [15]. In addition, it has been found that cysteine 106 can be oxidatively modified to the sulfinic acid. This modification has been reported to play a critical role in protecting M17 neuroblastoma cells from MPP⁺ toxicity [16]. These data suggest that

sufficient levels of DJ-1 protein and specific redox modification of DJ-1 play important roles in mediating neuronal protection.

Protein S-glutathionylation (Protein-SSG) is a prevalent oxidative modification involving the formation of a disulfide bond between glutathione (GSH) and a reactive protein-cysteine-SH [2]. Protein glutathionylation can result in changes in subcellular localization, protein function and/or protein levels [17, 18, 19]. Glutaredoxin 1 (Grx1) is the principal deglutathionylating enzyme in the cell, restoring the protein-SH and its function [2]. Grx1 has been shown to mediate neuronal protection both *in vitro* and *in vivo* [19, 20, 21, 22]. We recently reported evidence that PD patients have diminished levels of Grx1 specifically within the remaining dopaminergic neurons of the midbrain compared to controls [20]. Further we found that *Caenorhabditis elegans* (*C. elegans*) lacking the homolog of Grx1 displayed exacerbated losses of dopamine-dependent behavior and dopaminergic neurons in models of both familial and sporadic PD [20]. Thus, it appears that Grx1 mediates protection of dopaminergic neurons *in vitro* and *in vivo*, however the downstream molecular mechanism(s) that accomplish this protection remain to be elucidated. In this regard, knockdown of *Grx1* in model dopaminergic neurons was previously reported to increase cell death [21, 22], and to decrease DJ-1 protein content [22]. Over-expression of WT, but not C53S or C106S DJ-1, abolished the cell toxicity associated with diminished Grx1 function, suggesting that the content of DJ-1, its cysteine redox status, and its effect on neuronal cell viability are regulated *via* Grx1 [22]. Since this interpretation may also apply to our observations that loss of the Grx1 homolog in *C. elegans* exacerbates PD-like phenotypes [20], we set out in the current study to determine whether Grx1 regulates DJ-1 *in vivo*.

Here, we report evidence supporting a post-translational mechanism for regulation of DJ-1 protein content involving reversible glutathionylation and regulation by Grx1 *in vivo*. Accordingly, over-expression of DJ-1 partially compensates for neuronal toxicity due to the loss of Grx1 in models of PD.

Methods

Western blot analysis of mouse brain tissue

Mouse brain tissues from two independent cohorts obtained from Case Western Reserve University and The University of Vermont (a kind gift from Dr. Y. Janssen-Heininger), were dissected into three sections, midbrain, cortex and cerebellum. Midbrain portions of the brains were snap frozen in liquid nitrogen. Prior to use, midbrains were homogenized by physical disruption using a pestle in a 1.5 ml Eppendorf tube with Cell Lysis Buffer (Cell Signaling) containing protease inhibitors (Roche). Tubes were incubated on ice for 15 min followed by sonication three times for 5 sec each. Samples were clarified by centrifugation at 4°C for 30 min at 16,000 x g. Aliquots of the supernatant (70 µg of protein content) were run on a 15% SDS-PAGE gel, transferred to PVDF membrane using a semi-dry transfer apparatus (Bio-Rad) at 20 V for 1 h, blocked with 5% milk in Tris-buffered saline with 0.05% Tween-20 (TBST) for 1 h, and probed overnight with anti-actin (Sigma, diluted 1:5000), DJ-1 (Abcam, 1:5000) and Grx1 (Genscript, 1:1000) antibodies. Blots were washed three times for 10 min each in TBST and probed with secondary antibodies (Bio-Rad, 1:5000) for 1 h. Blots were developed using enhanced chemiluminescence (Thermo Fisher).

Rabbit polyclonal anti-Grx1 was generated by Genscript against recombinant human Grx1 produced in *E. coli* [20]. The western blots were scanned and band densities were quantified using ImageJ software (NIH).

Western blot analysis of human brain tissue

Postmortem brain tissue samples from histopathologically diagnosed PD and non-PD cases (Supplemental Table A), were used for this study according to approved IRB protocols. Frozen tissues of brainstem or pons were obtained from the NICHD Brain and Tissue Bank for Developmental Disorders at the University of Maryland (Baltimore, MD), and from the Case Western Reserve University/University Hospitals of Cleveland Brain Bank (Cleveland, OH). Areas of the frozen brainstem samples containing the substantia nigra were carefully dissected and were homogenized with Cell Lysis Buffer (Cell Signaling) plus 1 mM phenylmethanesulfonyl fluoride (Sigma) and Protease Inhibitor Cocktail (Sigma). Equal amounts of total protein extract were run on a 15% SDS-PAGE gel and transferred to PVDF (Millipore). Membranes were blocked in 5% milk in TBST for 1 h. Primary antibodies were incubated overnight in 5% milk in TBST, and washed three times in TBST prior to addition of the secondary antibody. Secondary antibodies were incubated for 1 h in 5% milk in TBST, washed three times in TBST and developed using ECL (Pierce). Antibodies used included anti-actin (Sigma), anti-DJ-1 (Abcam) and anti-Grx1 [20].

Western blot analysis of human blood cells

Human blood samples used for this study were collected according to approved IRB protocols (see patient information Supplemental Table B). PBMCs from 1.5 ml of cryogenically frozen blood were isolated by lysis of the red blood cells in 10 ml red blood cell lysis buffer (0.155 M ammonium chloride, 0.01 M potassium bicarbonate, and 0.00013 M EDTA) for 15 min on a rocker. The suspension was spun at 1,500g on a table-top centrifuge, the supernatant was removed, and 10 ml of red cell lysis buffer was added. This process was repeated two more times with incubation on the rocker shortened to 5 min each. The resulting PBMCs were lysed and were analyzed *via* western blotting in the same manner as described for the western blot analysis of mouse brain tissue except 35 μ g of lysate was loaded per sample.

Immunoprecipitation of DJ-1 from mouse brain lysate

Whole snap frozen C57BL/6 or FEB mouse brains were placed in 4 ml cell lysis buffer (Cell Signaling) plus 50 mM iodoacetamide (to block free cysteine-SH). Brains were mechanically homogenized *via* mortar and pestle, and then subjected to lysis *via* sonication on ice for 2 min, alternating 10 s on 10 s off. Lysate was then spun down at 16,000g at 4°C in a tabletop centrifuge for 1 hour. The resulting supernatant was divided into aliquots, with addition of either 50 μ l Protein G beads (Pierce) and 5 μ g anti-goat IgG (Santa Cruz); or 50 μ l Protein G beads and 5 μ g anti-DJ-1 (Goat, Abcam). Samples were rotated at 4°C for 48 h, washed three times in lysis buffer plus 0.25 M NaCl. Laemmli sample buffer (without reducing agents) was added and beads were boiled for 10 min. Samples were then run on a 15% non-reducing SDS-PAGE gel, transferred to a PVDF membrane, and probed for glutathionylation using an anti-glutathione antibody (anti-GSH, Millipore). Membranes were stripped by incubation in 0.1 M Tris-HCl pH 6.8, 2% SDS, and 0.11 M β -mercapto

ethanol at 50°C for 30 min followed by three washes in TBST for 15 min each. Blots were then probed with rabbit anti-DJ-1 (Cell Signaling) in the same manner as described above.

LC-MS analysis of proteins and peptides

Mass spectrometric (MS) analyses were performed using a LTQ Velos dual-pressure linear ion trap mass spectrometer equipped with an Accela 600 quaternary pump for in-line protein and peptide high performance liquid chromatography (HPLC) separation and an electrospray ionization source (Thermo Scientific, Waltham, MA). The electrospray ionization source was operated in the positive ion mode with the following parameters: normalized collision energy 35 kV, capillary temperature 300 °C, source voltage 5 kV, capillary voltage 43 V, and tube lens 105 V. Mass spectra were collected over a 200–2000 m/z range. To record mass spectra of intact proteins, 0.5 µg of DJ-1 or its glutathionylated form was loaded onto an XBridge BEH300 C4 column (3.5 µm, 2.1 × 50 mm) (Waters, Milford, MA) and eluted with a linear gradient (5–100%) of acetonitrile in water containing 0.1% formic acid. The gradient was developed over 20 min at a flow rate of 0.25 ml/min. Masses of the proteins were calculated by deconvolution of intact protein ions using ProMass for Xcalibur software (Thermo Scientific). The same instrumentation including column and solvents was employed to separate peptide products of tryptic digestion of DJ-1. However, to ensure adequate peptide separation the gradient of acetonitrile was developed over 40 min. The protein tryptic digestion was performed in 50 mM Tris-HCl, pH 7.5, at 37 °C overnight. Products of the reaction were directly analyzed by LC-MS. Peptides were identified based on a correlation of their tandem-mass spectra (MS/MS) with theoretically predicted ion fragmentation patterns.

The efficiency of protein glutathionylation was assessed by relative quantification of unmodified DJ-1 in the reaction sample. For this purpose, prior to the tryptic digestion, the glutathionylated DJ-1 sample was supplemented with an equal amount (12.5 µg) of ¹⁵N-labeled DJ-1. LC-MS analysis of the tryptic peptide mixtures revealed co-eluting native and ¹⁵N-labeled peptides. The relative ion intensities for pairs of native and ¹⁵N-labeled peptides at the same charge state were used to calculate the relative abundance of unmodified DJ-1 and thus the fraction of glutathionylated protein. Glutathionylation of cysteine residues were estimated using unmodified ¹⁵N DJ-1 as an internal standard. Quantification of modified tryptic peptides were estimated by subtraction of the peak areas of the non-modified peptides in the treated ¹⁴N-samples from those in the untreated ¹⁵N-samples.

C. elegans strains and maintenance

A list of *C. elegans* strains with their designation, corresponding genotype, and strain nomenclature is shown in Supplemental Table C. *C. elegans* strains were maintained on nematode growth medium (NGM) plates with *E. coli* OP50 as a bacterial food, as previously described [6].

Single-worm PCR

Individual worms were placed in PCR tubes containing 10 µl total volume consisting of 1x OneTaq buffer (New England Biolabs) and 0.5 mg/ml proteinase K (Sigma). Worms were

frozen on dry ice for 15 min, then lysed *via* incubation at 65°C for 90 min followed by 95°C for 30 min. PCR amplifications were performed using OneTaq polymerase following manufacturer's instructions (New England Biolabs). Identification of WT and mutant *djr-1.2* DNA was accomplished using the following primers (Integrated DNA Technologies), Forward: 5'-GCAAACCTCGAAAGCAGTTGC-3', Reverse WT: 5'-CGTATTGTTCCGGACGTAGC-3' Reverse knockout: 5'-TTTGACCGTGTTCCTTTGTCG-3'. Identification of *LRRK2* and *glrx-10* DNA was accomplished using the primers described previously [20].

Generation of *C. elegans* expressing Pdat-1::mCherry or Pdat-1::DJR1.2::mCherry in dopaminergic neurons

Plasmids containing Pdat-1::mCherry and Pdat-1::DJR-1.2::mCherry were constructed in the backbone of the pFxnRFP vector (a generous gift of Dr. S. Mitani, Tokyo Women's Hospital, Japan). The Pdat-1 insert was excised from the Pdat-1::pFxnEGFP plasmid [6] using the KpnI and HindIII sites. The DJR-1.2 open reading frame was PCR amplified from the DJR-1.2_pDONR201 plasmid (GE) with primers containing the BamHI and XhoI sites, Forward 5'-GATATAGGATCCATGGCTGCCCAAAGAG-3' and Reverse 5'-GTACTACTCGAGTACTTTGCCAAACACAG-3'. The mCherry insert was obtained by PCR amplification from the pGH8 plasmid (Addgene) with primers containing the NotI and BglII sites, Forward 5'-ATAATAGCGCCGCTTAATGGTCTCAAAG-3' and Reverse 5'-GCGCGCGAGATCTACTACTTATACAATTCATCC-3'. Restriction digestion was performed with appropriate selective enzymes according to the manufacturer's instructions (New England Biolabs). PCR products were purified with the QIAquick PCR purification kit (Qiagen). All DNA fragments were run on 1% agarose gels, followed by extraction with the QIAquick gel extraction kit (Qiagen). DNA inserts were ligated into pFxnRFP vector at room temperature for 2.5 h using T4 ligase (New England Biolabs) to generate the Pdat-1::mCherry and Pdat-1::DJR-1.2::mCherry plasmids. The plasmids were transformed into *E. coli* DH5 α cells (Life Technologies), propagated, purified on Qiagen miniprep columns, and verified by DNA sequencing. The DNA constructs were microinjected into *C. elegans* N2 strain by Knudra Transgenic (Murray, Utah). Stable worm lines with red fluorescent dopaminergic neurons expressing mCherry (Pdat-1::mCherry) or DJR-1.2 fused with mCherry (Pdat-1::DJR-1.2::mCherry) were selected, and their genotypes were verified by PCR.

Basal slowing assay

The basal slowing assay was performed as previously described [6]. Briefly, NGM assay plates were seeded with a ring of *E. coli* OP50. Age-synchronized worms were transferred to a non-seeded NGM plate and washed twice in S basal buffer (100 mM NaCl, 10 μ g/ml cholesterol, 50 mM potassium phosphate, pH 6.0). Worms were then transferred to either seeded or non-seeded assay plates, allowed to recover for 5 min, and their body bends were recorded in 20 s intervals.

Dopaminergic neuron counting

Approximately 30 age-synchronized adult worms were placed into 200 μ l of 10 mM tetramisole (Sigma) in raised wall four chamber slides (Lab-Tek). GFP neurons were

visualized with a Nikon microscope at 20x zoom. Dopaminergic CEP and ADE head neurons were assessed for degeneration in a blinded fashion, according to published protocols [20]. Neurons with detectable GFP signal were counted as positive, neurons lacking cell body GFP signal were counted as negative.

Treatment and analysis of SH-SY5Y cells

SH-SY5Y cells (ATCC) were grown in OPTI -MEM (Invitrogen) + 10% fetal bovine serum (Atlanta Biological) at 37 °C with 5% CO₂. For H₂O₂ treatment, cells were treated with increasing concentrations of H₂O₂ for 2 h as indicated in the figure legend. For diamide treatment experiments, cells were pretreated with either 10 μM MG-132 (Sigma) diluted in DMSO, 1 μM lactacystin (Santa Cruz) diluted in DMSO, 1:200 dilution of cell culture broad protease inhibitors (Sigma) diluted in DMSO, or DMSO (0.5%) for 7 h. Cells were then treated with 0.1 mM diamide or PBS for 2 h. All cells were lysed in Cell Lysis Buffer (Cell Signaling) containing protease inhibitors (Roche) on ice for 15 min with occasional mixing (Vortex). Lysates were spun down at 4°C for 30 min at 16,000 x g in a table-top centrifuge. The supernatant (35 μg of protein) was loaded onto a 15% SDS-PAGE gel and western blotting was completed in the same manner as the mouse midbrain tissue described above.

Validation of glutathionyl mixed disulfide specificity of the anti-GSH Antibody

Glutathionylation of proteins can be reversed by addition of disulfide reducing agents. In order to validate the interpretation of the reactivity of the anti-GSH antibody, anti-DJ-1 beads were used to immunoprecipitate DJ-1 from mouse brains, and the resultant effluent was processed on a non-reducing SDS-PAGE gel and transferred to a PVDF membrane. After transfer the blot was blocked in 5% milk for 1h and cut in half. Half of the blot was incubated with buffer and the other half was incubated with buffer plus 100 mM dithiothreitol (DTT) for 30 min. The blots were then washed and probed with the anti-GSH antibody overnight. The blot incubated in buffer alone revealed the prominent band at 45 kDa while the blot incubated in buffer plus DTT failed to show immunoreactivity at 45 kDa (Supplemental Figure S4). Blots were then stripped and re-probed for DJ-1 confirming the presence of DJ-1 on both halves of the membrane.

Generation of recombinant DJ-1

Recombinant DJ-1 was purified following the protocol described in [23]. Briefly 1.5 L of BL21(DE3) *E. coli* containing pET15b-DJ-1 was induced with 1 mM IPTG for 4 h. Cells were spun down and lysed via lysozyme treatment followed by sonication. Centrifugally cleared lysate containing 6x-His tagged DJ-1 was bound to Ni²⁺ His-select resin (Sigma), washed, eluted with 200 mM imidazole, cleaved with thrombin, and dialyzed overnight. Uncleaved DJ-1 was removed using a second pass over the Ni²⁺ column, thrombin was removed using benzamidine-sepharose resin, and co-purifying nucleic acids were removed from DJ-1 using a Q-sepharose column. DJ-1 (free of the 6x-His tag) was then concentrated and snap frozen in liquid nitrogen.

Generation of ^{15}N labeled DJ-1

Recombinant uniformly ^{15}N -labeled DJ-1 was generated as previously described [23]. Briefly, N-terminally 6x-His tagged DJ-1 was expressed from pET15b in BL21(DE3) *E. coli*. An inoculum of the cells was grown overnight in LB media containing 100 $\mu\text{g/ml}$ ampicillin at 37°C. 50 ml of this culture was spun down and washed twice in 10 ml of sterile $^{15}\text{NH}_4\text{Cl}$ M9 defined media [23] and transferred to 1 L of $^{15}\text{NH}_4\text{Cl}$ M9 media. These cells were grown at 37°C and DJ-1 expression was induced by the addition of 0.5 mM IPTG when the $A_{600} = 0.5$. Protein purification was performed as described above. ^{15}N labeling efficiency was determined using mass spectrometry as described [23] and the incorporation efficiency was >99%.

RNA isolation and qPCR Methods

Midbrain tissue from C57BL/6 and BALBc *Grx1*^{-/-} mice and age- and sex-matched controls (when available) was homogenized in 1 mL TRIzol (Life Technologies) using a Teflon and glass mortar and pestle. RNA was isolated according to manufacturer's instructions. Resulting RNA was converted to cDNA using SuperScriptII (Invitrogen) according to manufacturer's instructions. Park7 (DJ-1) expression (probe Mm00498538_m1) was analyzed using TaqMan Fast Universal PCR mix (Life Technologies) on the StepOnePlus machine (Life Technologies) using StepOne v2 software. Park7 expression was normalized to Gapdh (probe Mm99999915_g1) expression as an internal control.

RESULTS

Grx1 controls DJ-1 protein expression *in vivo*.

As reported previously, knockdown of Grx1 *via* shRNA results in decreased DJ-1 protein content in the SH-SY5Y [21] and Neuro2A cell lines [22], however it is unknown if Grx1 plays a role in mediating DJ-1 protein content *in vivo*. Midbrain lysates from *Grx1*^{-/-} and WT mice in the C57BL/6 background from two independent cohorts were subjected to western blot analysis, in order to investigate the potential control of DJ-1 by Grx1. Western blot analysis indicated that *Grx1*^{-/-} mice have diminished levels of DJ-1 protein compared to WT controls (Figure 1 A, B). Quantitative PCR was used to measure the mRNA levels of DJ-1 in the *Grx1*^{-/-} and WT mice in the C57BL/6 background. No statistical difference was found in the mRNA levels between the *Grx1*^{-/-} and WT mice (Supplemental Figure S1A). We also analyzed DJ-1 content in brain samples (kindly provided by Reiko Matsui, Boston University) from transgenic mice in which hGrx1 was overexpressed so that total Grx1 content (human plus mouse) was about 2-fold higher relative to WT mice [24]. In this case the DJ-1 content was not different (transgenic *vs.* WT), suggesting that the normal content of Grx1 is sufficient to maintain DJ-1 content, at least in the absence of increased oxidative stress (Supplemental Figure S1B). Notably, we also analyzed midbrain lysates from *Grx1*^{-/-} and WT mice in the BALB/c background and these mice did not show a difference in the DJ-1 protein or mRNA content (Supplemental Figure S1 C, D); thus indicating that differential genetic expression and/or posttranslational regulation in the BALB/c strain disconnects the regulatory effect of Grx1 on DJ-1.

In addition, to investigate whether DJ-1 and Grx1 protein levels correlate in human tissue, lysates from human peripheral blood mononuclear cells (PBMCs) and post-mortem midbrain tissue were subjected to western blot analysis. Protein levels of Grx1 and DJ-1 (normalized to actin) showed a significant positive correlation for protein isolated from both PBMCs (Figures 1 C) and midbrain tissue (Figures 1 D; blots are shown in Supplemental Figure 2S A, B). Taken together, these data provide *in vivo* evidence for the first time that Grx1 influences DJ-1 protein content in the midbrains of mice, and that Grx1 and DJ-1 protein contents are co-variant in human tissue.

Treatment of cells with glutathionylation-inducing agents results in loss of DJ-1 protein

As noted, knockdown of Grx1 in model dopaminergic neuronal (Neuro2A) cells *via* shRNA resulted in decreased DJ-1 protein, but not mRNA content [22]; however the mechanism by which this loss of DJ-1 protein occurred remains unclear. We hypothesized that loss of Grx1 results in accumulation of glutathionylated DJ-1 (DJ-1-SSG), which is then rapidly degraded. In order to test whether agents that promote DJ-1-SSG formation result in loss of DJ-1 content, SH-SY5Y cells were treated acutely with H₂O₂, which is known to result in protein-SSG formation [25, 26]. DJ-1 protein content was selectively diminished in a dose dependent manner (relative to actin, which is known to be glutathionylated without affecting its content) (Figure 2 A, B). Since H₂O₂ can modify cysteine residues to sulfinic and sulfonic acids in addition to protein-SSG, we wanted to limit the modification to glutathionylation of DJ-1. Accordingly, cells were treated separately with diamide, an oxidizing agent characterized to promote selective formation of protein-SSG [27, 28, 29]. Western blot analysis indicated that treatment with diamide also resulted in a significant loss of DJ-1 protein content compared to untreated controls. (Figure 2 C, D). Notably, the extent of diminution in DJ-1 protein content (~40%) was similar for both the H₂O₂- and diamide-treated cells. It was previously reported that loss of DJ-1 content concomitant with knockdown of Grx1 could be limited by treatment with a cocktail of inhibitors of proteolysis, consistent with the concept of enhanced degradation [22]. In order to learn whether loss of DJ-1 after diamide treatment of SH-SY5Y cells displayed the analogous pattern of non-proteasomal proteolysis as observed in Grx1-deficient Neuro2A cells [22], we pretreated the SH-SY5Y cells with the proteasome inhibitor MG-132 or with a broad protease inhibitor (PI) cocktail prior to diamide treatment [22]. Pretreatment with the broad protease inhibitors, prevented the diamide induced loss of DJ-1 (Figure 2 C, D). Notably, pretreatment of SH-SY5Y cells with either MG-132 or lactacystin inhibited the proteasome (as indicated by increased β -catenin content (Figure 2, E, G), but failed to blunt the diamide-induced loss of DJ-1 protein (Figure 2 E, F). These data suggest that treatment of SH-SY5Y cells with glutathionylation-inducing agents results in DJ-1 degradation in a proteasome-independent manner.

DJ-1 is glutathionylated *in vivo*

DJ-1 contains three cysteine residues located at positions 46, 53, and 106. Cysteines 53 and 106 have been reported previously to be oxidatively modified [23, 30], but there is no evidence for glutathionylation of DJ-1 in a physiological setting. In order to determine if DJ-1 undergoes glutathionylation *in vivo*, DJ-1 was immunoprecipitated from whole mouse brain homogenates and subjected to non-reducing SDS-PAGE and western blot analysis.

Using an antibody that recognizes glutathionylated moieties (anti-GSH), we observed a prominent band at approximately 45 kDa (Figure 3 A, arrow). The blot was stripped and re-probed with an anti-DJ-1 antibody, which resulted in a band at the same molecular weight as recognized by the anti-GSH antibody. (Figure 3 B). The band located at approximately 45 kDa corresponds to the size of the dimeric DJ-1. This suggests that dimeric DJ-1-SSG can form an intermolecular bond between two DJ-1 monomers either in the cell or while running through the electric field in the gel. In order to validate that the approximately 45 kDa band is DJ-1 and not a non-specific band, mouse brain homogenates were immunoprecipitated with anti-DJ-1 beads to isolate DJ-1, and the sample was split into two aliquots. One aliquot was reduced with 100 mM dithiothreitol (DTT) and the other was left non-reduced; then the two samples were analyzed on western blots. The ~45 kDa band appeared in the non-reduced lane, but in the reduced lane a single band was present at approximately 22 kDa, corresponding to the molecular weight of monomeric DJ-1 (Supplemental Figure S3A). In addition, western blot analyses with the anti-DJ-1 antibody were performed with DTT-treated lysates of brain samples from WT and *DJ-1*^{-/-} mice (a kind gift from Vera Bonilha, Cleveland Clinic Foundation). An immunoreactive band appeared at approximately 22 kDa as expected for the WT samples; but no band was present for the samples from the *DJ-1*^{-/-} mice, documenting the specificity of the anti-DJ-1 antibody (Supplemental Figure 3S B).

Glutathionylation of proteins can be reversed by disulfide reducing agents. To confirm the interpretation of the reactivity of the anti-glutathione antibody, DJ-1 was immunoprecipitated from mouse brains, run on a non-reducing SDS-PAGE gel and transferred to a PVDF membrane. After transfer half of the blot was then incubated with 100 mM dithiothreitol (DTT) for 30 min to reduce glutathionylated cysteines. The blot incubated with DTT lost immunoreactivity with the anti-GSH antibody (Supplemental Figure 4S). Blots were then stripped and re-probed for DJ-1 confirming the presence of DJ-1 on both halves of the membrane. These data show that treatment with DTT results in ablation of reactivity with the anti-glutathione antibody, consistent with the conclusion that DJ-1 is glutathionylated within the midbrains of mice. It was also confirmed separately that actin which is known to be glutathionylated in intact cells [31] and has a similar molecular weight (42 kDa) to dimeric DJ-1 was not detected on the blot of the immunoprecipitated DJ-1 (data not shown).

Preferential glutathionylation of DJ-1 at cysteines 53 and 106

To further characterize the glutathionylation of DJ-1, 12.5 µg of pure recombinant DJ-1 was incubated with 5 mM GSH and 1 mM diamide for 30 min and subjected to liquid chromatography-mass spectrometry (LC-MS) analysis. MS spectra of the untreated (Figure 3 C) and treated (Figure 3 D) intact protein revealed two distinct sets of ions in the treated sample as compared to untreated DJ-1. The deconvoluted spectrum of untreated DJ-1 revealed the expected protein mass of 20036.3 Da, equal to the theoretical mass of the polypeptide based on the amino acid sequence. Reaction with GSH and diamide resulted in increases of 305 and 610 Da in the protein mass to 20341.5 and 20646.7 Da, corresponding to the additional masses of a single and two glutathionyl moieties covalently attached to DJ-1 (Figure 3 D).

To identify the sites of glutathione adduction, the glutathionylation reaction was repeated; the DJ-1 reaction mixture was dialyzed to remove GSH/diamide, digested with trypsin, and subjected to tandem MS/MS analysis. The trypsin digests clearly revealed the presence of two covalently modified peptides DVVIC⁵³PDASLEDAK ($m/z = 766.5 [M+2H]^+2$) and GLIAAIC¹⁰⁶AGPTALLAHEIGFGSK ($m/z = 756.4 [M+3H]^+3$) in the GSH/diamide treated sample (Figure 4 C–D) as compared to the unmodified DJ-1 (Figure 4 A–B). The identity of each of the peptides and sites of modification were confirmed by their MS/MS fragmentation patterns. A 305-Da shift in the b- and y-ion series observed between the modified and unmodified peptides indicated glutathionylation on the side chains of the Cys-53 (Figure 4 E) and Cys-106 (Figure 4 F) residues.

In order to estimate the relative extents of glutathionylation of the different cysteine residues, “light DJ-1” (¹⁴N, natural abundance) was treated with GSH and diamide. An equimolar amount of “heavy DJ-1” (¹⁵N-labeled) was untreated and used as an internal standard. Both ¹⁴N- and ¹⁵N-proteins were digested with trypsin, and subjected to LC-MS analysis (Supplemental Figure 5S, A–E). Quantification of the peak areas for the non-modified peptides indicated that major glutathionylation occurs at cysteine residues Cys53 and Cys106. The extent of glutathionylation was estimated by subtracting the peak area of the treated non-modified peptide (¹⁴N) from the peak area of the untreated standard peptide (¹⁵N), giving a value for the peak area of the glutathionylated peptide. Accordingly, the extent of glutathionylation of Cys53 and Cys106 were calculated to be 70 and 77%, respectively (Table 1). We did find a very minor fraction (~7%) of glutathionylated Cys46 (Table 1), but this was likely due to heterogeneity within the preparation of DJ-1 and/or unfolding of the protein in solution, as Cys46 is known to be buried in the hydrophobic core of the protein [32]. Taken together, our data show that isolated recombinant DJ-1 is preferentially glutathionylated at cysteine residues 53 and 106.

Loss of DJ-1 homolog exacerbates LRRK2-mediated dysfunction of dopamine-dependent movement in *C. elegans*

Overexpression of DJ-1 has been shown to mediate dopaminergic neuronal protection against LRRK2 toxicity *in vitro* [33]. In order to investigate the potential neuroprotective role of DJ-1 *in vivo*, we chose to use *C. elegans* as a model because of the simplicity of the dopaminergic neuronal system. *C. elegans* worms contain two distinct homologs to human DJ-1, DJR-1.1 and DJR-1.2. Two independent groups have reported that DJR-1.1 is expressed highly in the intestine and not within the head neurons [34, 35]. DJR-1.2, in contrast, has been reported to be expressed in the dopaminergic neurons of the worms [35]. Therefore, we used DJR-1.2-deficient animals in this study. Genetic crosses were made between the *djr-1.2*^{-/-} strain and transgenic animals expressing GFP in the dopaminergic neurons (used as WT control). In addition, the *djr-1.2*^{-/-} strain was crossed with lines that co-express GFP and pathogenic human LRRK2 mutants in the dopaminergic neurons (G2019S-LRRK2 or R1441C-LRRK2, hereafter referred to as G2019S and R1441C, respectively) [6]. Crosses were confirmed *via* PCR genotyping of *djr-1.2* and *LRRK2* (Figure 5A).

We examined the functional integrity of dopaminergic signaling in the *C. elegans* lines using a well-established assay of dopamine-dependent behavior, termed the basal slowing assay [36]. In agreement with our previous findings, transgenic expression of either G2019S or R1441C in the dopaminergic neurons of *C. elegans* resulted in age-dependent loss of dopamine-dependent movement behavior [6, 20]. Compared to WT, the mutant *djr-1.2*^{-/-} worms displayed a late onset loss of dopamine-dependent behavior on adult day 2 (Figure 5B). These data are in agreement with the previous finding that dauer-stage *C. elegans* lacking *djr-1.2* have a dopamine signaling defect [35]. Moreover, the hybrid lines, both deficient in DJR-1.2 and expressing G2019S or R1441C, displayed a more robust loss of dopamine-dependent behavior (Figure 5B). We also examined the morphological integrity of the GFP labeled-dopaminergic neurons *via* microscopy in young and aged worms. Expression of G2019S or R1441C resulted in an age dependent loss of dopaminergic neurons as previously documented [6, 20]. The *djr-1.2*^{-/-} mutant strain showed decreased survival of dopaminergic neurons compared to WT on adult day 8 (Figure 5C). Exacerbation of dopaminergic neuronal loss in the lines lacking DJR-1.2 and expressing LRRK2 mutants was limited and seen only on adult day 2 in the R1441C; *djr-1.2*^{-/-} line. These data indicate that loss of DJR-1.2 *in vivo* results in diminution of dopamine-dependent behavior, exacerbation of mutant LRRK2-induced diminution of dopamine-dependent behavior, but only incomplete dopaminergic degeneration and limited exacerbation of LRRK2-mediated dopaminergic degeneration.

Over-expression of DJ-1 homolog partially compensates for the loss of Grx1 in R1441C-mediated neurotoxicity

We previously found that loss of GLRX-10, the worm homolog of Grx1, exacerbates LRRK2-mediated dopaminergic dysfunction [20]. Our current data analyzing midbrain homogenate from *Grx1*^{-/-} mice indicates that loss of Grx1 results in a decreased protein content of DJ-1 *in vivo* (Figure 1 A, B), indicating that regulation of DJ-1 content is functionally downstream of Grx1. Synthesizing these observations, we hypothesized that the exacerbation of LRRK2 toxicity by deficiency of the Grx1 homolog in *C. elegans* [20] is due, at least in part, to increased glutathionylation and consequent loss of DJ-1. In order to test if DJ-1 overexpression could rescue the Grx1 deficiency, we generated R1441C; *glrx-10*^{-/-} worms specifically expressing DJR-1.2 fused to red fluorescent protein mCherry (DJR-1.2-mCherry) or mCherry (vector control) in the dopaminergic neurons. The compound worm lines were confirmed *via* PCR (Figure 6A) and fluorescence microscopy (Figure 6B). These worms were subjected to the basal slowing assay to assess their dopamine-dependent behavior.

Consistent with the previous report [20], R1441C; *glrx-10*^{-/-} worms displayed a robust loss in dopamine-dependent behavior. Expression of the mCherry control in the R1441C; *glrx-10*^{-/-} line resulted in no change in basal slowing response compared to its cognate strain (Figure 6C). Overexpression of DJR-1.2-mCherry in the dopaminergic neurons of the R1441C; *glrx-10*^{-/-} worms showed significant improvement in basal slowing response compared to the R1441C; *glrx-10*^{-/-} worms expressing the vector control on adult Day 0, and continued to show apparent but not statistically significant improvement on adult days 1 and 2. These data indicate that expression of DJR-1.2 can partially rescue the pathogenic

LRRK2 induced dysfunction of dopamine-dependent behavior in *C. elegans* lacking GLRX-10.

Discussion

DJ-1 plays an important role in mediating dopaminergic neuronal protection. De-stabilizing mutations in DJ-1 result in familial PD, and diminished protein content of DJ-1 has been reported in sporadic cases of PD [10, 11]. Furthermore, knockout or knockdown of DJ-1 results in increased susceptibility to toxin-induced dopaminergic cell death *in vivo* [37, 38]. These studies indicate that protein levels of DJ-1 play an important role in protecting against dopaminergic degeneration. Hence, deciphering the mechanisms of regulation of DJ-1 protein content represents an important focus of study in PD biology.

Reversible glutathionylation of proteins is recognized as an important regulatory mechanism for pathways that mediate cell death/viability [39]. For example, the FAS receptor is a signal transducing receptor that upon activation *via* Fas-ligand drives cell death through activation of the death domain. Fas-induced cell death has been shown to be regulated in part by glutathionylation at cysteine 294 [17]. Treatment of lung fibroblasts with Fas-ligand resulted in cell death when WT-Fas, but not C294A-Fas was expressed [17], suggesting that glutathionylation of Fas is a pre-requisite for cell death signal transduction. In addition, glutathionylation of the cell survival kinase Akt has been associated with decreased Akt activation and reduced cell viability in the human fetal retinal pigment epithelial cells when challenged with H₂O₂ [40]. Conversely, overexpression of Grx1 resulted in decreased levels of Akt glutathionylation in response to H₂O₂ and increased activation of Akt signaling [40]. These and other studies demonstrate the functional impact of selective protein glutathionylation on cell viability under stress conditions.

Our previous data revealed that Grx1 protein content was decreased in post-mortem samples of brain tissue from [20]. Our current data supports the interpretation that Grx1 regulates DJ-1 protein content *in vivo* and that DJ-1 is susceptible to glutathionylation *in vitro* and *in vivo*. Additional data supporting the idea that DJ-1 is regulated by glutathionylation have been reported. Unlike WT DJ-1, the C106A and C53A mutants of DJ-1 are resistant to degradation in Neuro2A cells in which Grx1 was knocked down [22]. Our current finding that cysteine residues 53 and 106 are more readily glutathionylated than cysteine 46 is consistent with data from the crystal structure of DJ-1 indicating that cysteine 46 is not solvent accessible and buried within the protein [32]. We hypothesize that glutathionylation of DJ-1 regulates its protein level in the cell. Consistent with this interpretation, treatment of SH-SY5Y cells with two known inducers of glutathionylation resulted in decreased DJ-1 protein content. This decrease in DJ-1 protein can be abrogated by pre-treatment with non-specific protease inhibitors suggesting that the loss of DJ-1 is caused by proteolytic degradation following glutathionylation. The result is in agreement with previously published data indicating that loss of Grx1 via shRNA knockdown in Neuro2A cells results in a loss of DJ-1 protein content [22]. In order to identify which component(s) of the proteolytic inhibitor cocktail blocked DJ-1 degradation and therefore which pathway(s) is dysregulated, we treated Neuro2A cells with the individual components and then knocked down Grx1 via shRNA. As expected, knockdown of Grx1 resulted in a loss of DJ-1 protein

content; however none of the individual components alone blunted the Grx1 knockdown-mediated loss of DJ-1 (Supplemental Figure 7S, A–F). These data suggest a complex mechanism of degradation likely involving multiple enzymes contributing to regulation of DJ-1 degradation.

In order to test whether DJ-1 plays a functional role in regulating dopaminergic viability in concert with Grx1 *in vivo* we used *C. elegans* as a model system. We previously found that *C. elegans* lacking GLRX-10, the Grx1 homolog, displayed an exacerbation of the neurodegenerative phenotype in both familial and sporadic models of PD [20]. Our observation of decreased levels of DJ-1 in *Grx1*^{-/-} mice led us to hypothesize that decreased DJ-1 could be at least partly responsible for the more toxic PD-like phenotype in the *C. elegans* lines lacking GLRX-10. Protein content of DJ-1 within the *glrx-10*^{-/-} worms was not determined because the antibodies for human DJ-1 failed to specifically recognize *C. elegans* DJR-1.2 (data not shown). As an alternative approach to test if an elevated content of DJ-1 could compensate for the loss of GLRX-10 in *C. elegans*, DJR-1.2 was specifically expressed in the dopaminergic neurons of R1441C-LRRK2; *glrx10*^{-/-} worms. Using the well-established basal slowing assay to determine integrity of dopamine-dependent behavior, we found that expression of DJR-1.2 could partially compensate for the loss of GLRX-10. Indeed, the protective effect of DJ-1 appears to occur early in the life of the worm and then fades in adulthood (Figure 6C). Fluorescence microscopic analysis was conducted on adult days 0 and 2 to assess levels of mCherry in control worms or in worms expressing exogenous mCherry-tagged DJ-1 protein. Quantification of mCherry/GFP intensity showed no difference in the expression of exogenous mCherry in the control lines on adult days 0 and 2 as expected (Supplemental Figure S6, left panel). Notably there was also no change in the mCherry/GFP ratio in the DJ-1:mCherry worms (Supplemental Figure S6, right), indicating that diminution of DJ-1 mediated protection with age is not due to a loss of the exogenous DJ-1 protein. These data suggest several alternative hypotheses for why the effect of the overexpressed DJ-1 fades with time, including (a) the levels of exogenous DJ-1 fail to sustain protection due to an increased burden of ROS over time in these LRRK2-expressing worms; (b) DJ-1 is functionally inactivated in the worms due to increased ROS; and (c) the DJ-1 mediated dopaminergic neuronal protection involves co-factors that may be deactivated by glutathionylation in the absence of the Grx1 homolog. One or more of these hypotheses may be supported by previous findings. For example, it has been reported that pathogenic LRRK2 drives increased ROS expression *in vitro* [41]; DJ-1 has been reported to exist in a (possibly inactive) sulfonic acid state after oxidative challenge [42]; the glutathionylated protein database [43] reports that the DJ-1 binding partner thioredoxin-1 can be glutathionylated. Taken together these observations provide a plausible explanation for the transient protective effect of elevation of DJR-1.2 content, contributing some compensation for loss of the Grx1 homolog and some protection against mutant LRRK2-mediated toxicity *in vivo*.

Although DJ-1 provides some protection against exacerbation of LRRK2 toxicity by glutaredoxin deficiency, the lack of complete rescue indicates that other protein(s) also likely play a role in mediating Grx1 neuronal protection. For example, Akt1 is a well-established survival kinase that has been shown to play neuroprotective roles in a variety of contexts including PD [44]. Notably, the G2019S- and R1441C-LRRK2 mutants showed a decreased

ability to phosphorylate and activate Akt1 *in vitro*, suggesting that pathogenic mutations of LRRK2 drive toxicity at least in part through loss of a critical cell survival signaling mechanisms involving Akt [45]. Proper activation/regulation of Akt is critical for Akt mediated protection. Considering the context of Grx1 deficiency, it is important to note that Akt1 has been reported in other contexts to be regulated both directly [40] and likely indirectly [46] through reversible protein glutathionylation. Thus, treatment of human retinal pigment epithelial cells with H₂O₂ resulted in Akt1 glutathionylation, impaired phosphorylation of Akt1, and cell death [40]. Over-expression of Grx1 resulted in decreased levels of glutathionylated Akt1, normalized phosphorylation/activation of Akt1, and increased cell viability when challenged with H₂O₂ [40]. These data indicate that glutathionylation of Akt itself or of regulators of Akt may interfere with its cell-survival activity. In particular, Akt1 is positively regulated through phosphorylation by the upstream kinase PKC [47], and PKC itself appears to be negatively regulated by glutathionylation [43]. Thus, purified recombinant PKC treated with diamide and ³⁵S-GSH was found to be glutathionylated and inactivated. Furthermore, glutathionylation of intracellular PKC was confirmed by documenting DTT-reversible incorporation of ³⁵S-radioactivity into PKC that was immunoprecipitated from diamide-treated NIH3T3 cells in which GSH had been radiolabeled [46]. These data suggest that glutathionylation and concomitant inactivation of PKC would decrease Akt1 phosphorylation and result in an impaired ability to respond to stress. Taken all together, loss of Grx1 within dopaminergic neurons besides affecting cell viability through diminution of DJ-1 may also result in an impaired stress response due to hyper-glutathionylation of PKC and Akt. The increased glutathionylation and consequent inactivation of DJ-1 and the Akt pathway may combine to explain the exacerbated toxicity seen when the Grx1 homolog is genetically ablated in models of familial and sporadic PD. [20]. Hence, further studies are warranted to examine the synergistic impact of dysregulation of glutathionylation on neuronal cell survival mechanisms involving DJ-1 and Akt.

Therapeutic advancement in the general PD population has not changed greatly since the introduction of levodopa as the primary therapy for PD in the 1960's. Currently there are no therapeutic approaches available to slow the progression of dopaminergic degeneration. The ideal intervention would slow or even reverse the degeneration of dopaminergic neurons by targeting a dysregulated pathway common to both familial and sporadic PD. Although familial mutations represent only approximately 10 percent of all PD cases, post-translational modifications mimicking the functional consequences of the genetic mutations may play a role in the development of sporadic PD. In our previous study post-mortem PD brain tissue was found to have diminished levels of Grx1 [20], supporting the possibility that glutathionylation of key proteins involved in PD development/progression may contribute to PD pathogenicity. Our current data are consistent with the idea that glutathionylation of DJ-1 results in its degradation leading to decreased protein levels. This loss of DJ-1 protein mimics the outcome of the pathogenic L166P mutation that results in reduced steady-state levels of DJ-1 [48]. A search of the database dbGSH reveals that several other proteins implicated in the etiology of PD are susceptible to glutathionylation [43]. These proteins include VPS35, UCH-L1, and EIF4G1. Although these proteins can be glutathionylated, it is unclear if glutathionylation affects the function of these proteins in an analogous manner to

pathogenic mutations, so further study is warranted. Nevertheless, the potential impact of deficient Grx1 on glutathionylation status and function of proteins implicated in PD suggests that pharmacologically increasing Grx1 activity in neurons may decrease dopaminergic toxicity by regulating not only DJ-1 protein content, but also other downstream targets that are altered *via* glutathionylation.

Statistical analysis

Statistical analyses of the reported data, as noted in the figure legends, were performed using ANOVA followed by Tukey's post hoc test, Student's t-Test, or one-way linear regression. All statistical analyses were run using Microsoft Excel or GraphPad Prism.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations

ATCC	American Type Culture Collection
<i>C. elegans</i>	<i>Caenorhabditis elegans</i>
Grx1	Glutaredoxin 1
GSH	Glutathione
GSSG	Glutathione disulfide
Protein-SSG	Glutathionylated protein
HPLC	High performance liquid chromatography

PBMCs	Human peripheral blood mononuclear cells
DJ-1-SSG	Glutathionylated DJ-1
LRRK2	Leucine Rich Repeat Kinase 2
LC-MS	Liquid chromatography-mass spectrometry
m/z	Mass/charge ratio
MS	Mass spectrometry (mass spectrometric)
NGM	Nematode growth media
PD	Parkinson's disease
TBST	Tris-buffered saline with 0.05% Tween-20

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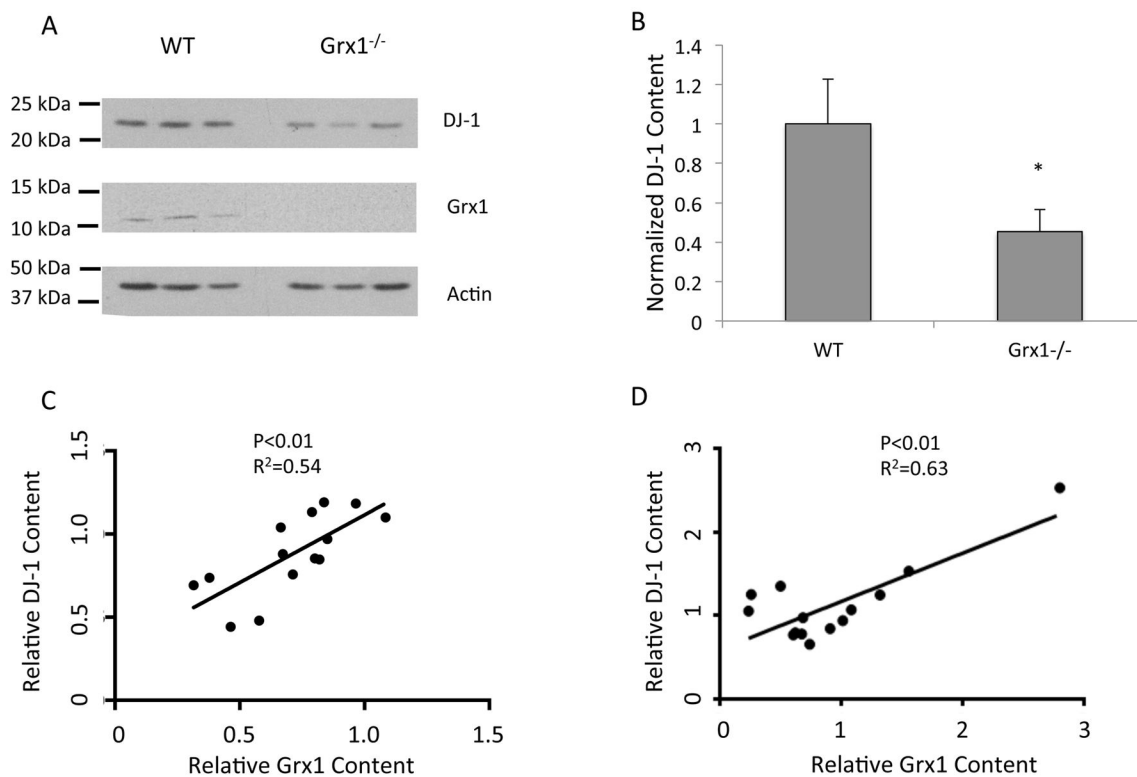
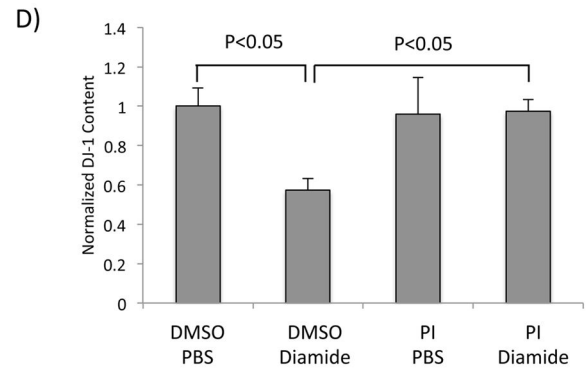
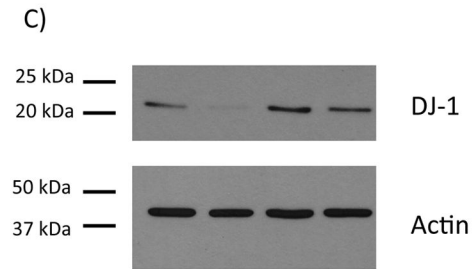
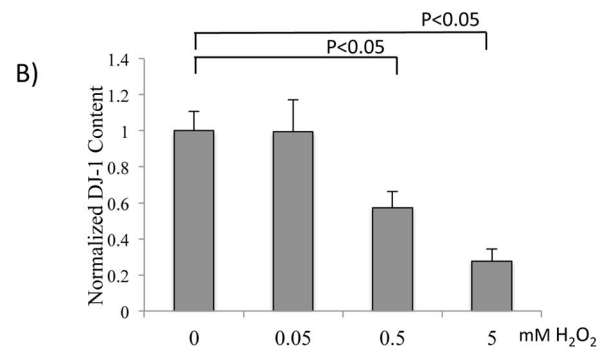
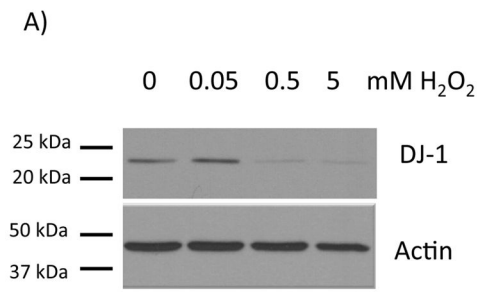


Figure 1. DJ-1 protein content is regulated by Grx1 *in vivo*

(A) Western blot analyses of midbrain homogenates from WT and *Grx1*^{-/-} mice probed with anti-Grx1, anti-DJ-1, and anti-actin antibodies, respectively. (B) Quantification of DJ-1 protein content (normalized to actin) in the lysates of midbrain samples from WT and *Grx1*^{-/-} mice; n = 6 independent WT and *Grx1*^{-/-} mice/**p*<0.05. Statistical analysis was completed using Student's T-test. Error bars represent SEM. (C) Positive correlation between the protein contents of DJ-1 and Grx1 (normalized to actin) in human PBMCs; n = 14 independent biological samples/*p*<0.01, *R*²=0.54. Statistical analysis was completed using a one-way linear regression test. (D) Positive correlation between the contents of the DJ-1 and Grx1 proteins (normalized to actin) in lysates of human post mortem midbrain tissue; n = 14 independent biological samples/*p*<0.01, *R*²=0.63. Statistical analysis was completed using a one-way linear regression test.



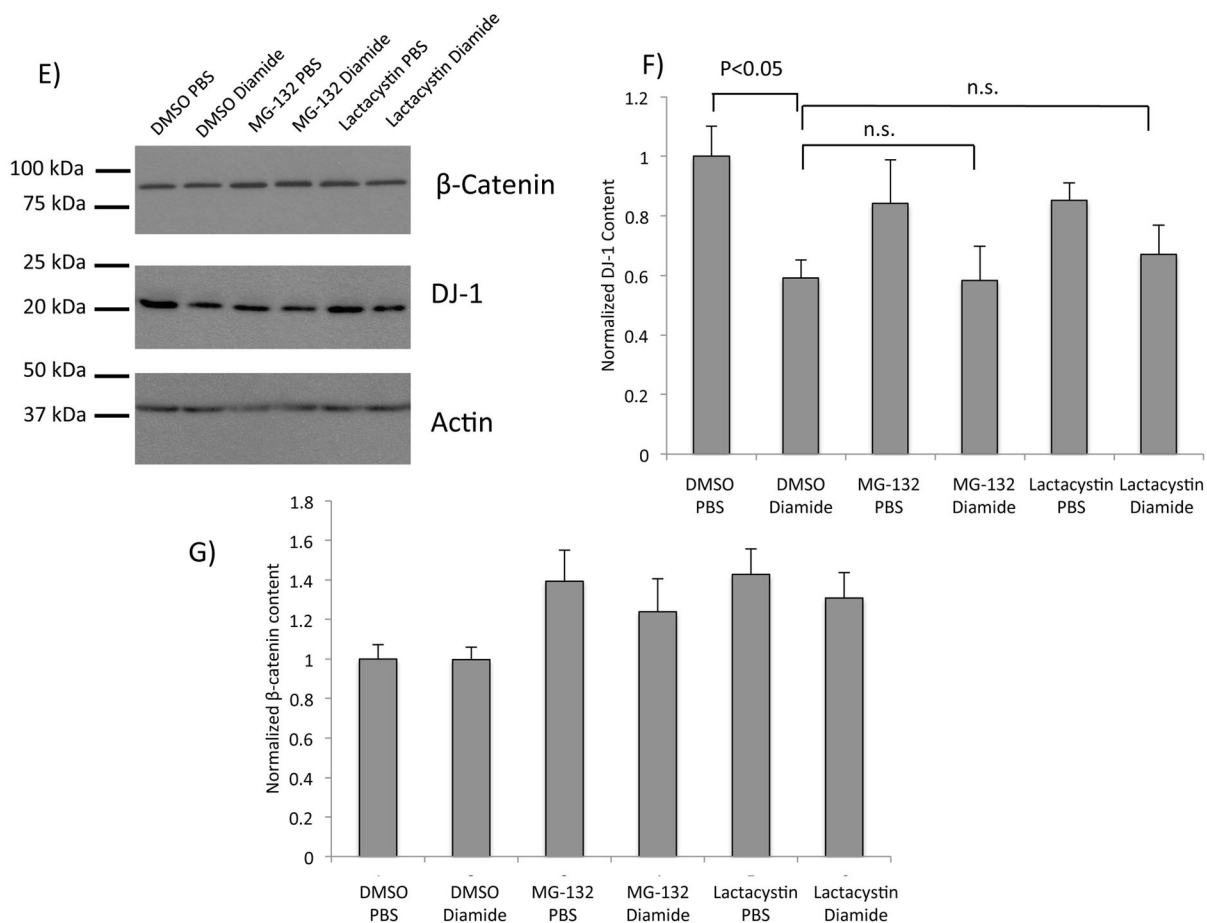


Figure 2. DJ-1 protein content is diminished after treatment with glutathionylation-inducing agents

(A) Representative western blot analysis of lysates from SH-SY5Y cells treated with increasing concentrations of H₂O₂. The blot was partitioned and probed with anti-DJ-1 or anti-actin, respectively. (B) Quantification of normalized DJ-1 content after H₂O₂ treatment; n = 8 independent biological replicates. (C) Representative western blot analysis of lysates from SH-SY5Y cells pretreated with DMSO, or a broad protease inhibitor cocktail (PI, *see Methods*); then treated with PBS or 0.1 mM diamide. The blot was partitioned and probed with anti-DJ-1 or anti-actin, respectively. Lane 1, DMSO + PBS; lane 2, DMSO + diamide; lane 3, PI + PBS; lane 4, PI + diamide. (D) Quantification of normalized DJ-1 content after diamide and PI treatments; n=8 independent biological replicates. (E) Representative western blot analysis of lysates from SH-SY5Y cells pretreated with DMSO, MG-132, or lactacystin; then treated with PBS or 0.1 mM diamide. The blot was partitioned and probed with anti-β-catenin, anti-DJ-1, and anti-actin, respectively. (F) Quantification of DJ-1 protein content (normalized to actin); n = 8 independent biological replicates. Statistical analyses for quantifications shown in (B), (D), and (F) were completed using ANOVA followed by Tukey's post hoc test. Error bars on quantification for (B), (D), and (F) represent SEM.

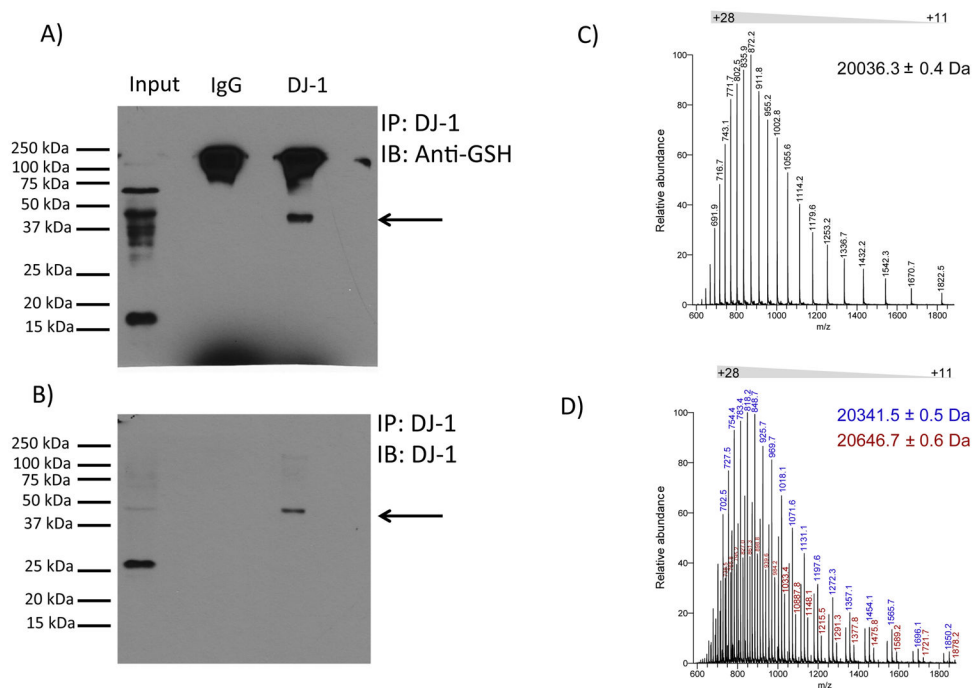


Figure 3. DJ-1 is glutathionylated *in vivo* and *in vitro*

(A) Representative western blot analysis of immunoprecipitated DJ-1 from mouse brain homogenate probed with anti-glutathione (GSH) antibody. Replicate analyses of four independent biological samples gave similar results. Arrow denotes a band of immunoreactivity at approximately 45 kDa. (B) Immunoprecipitated DJ-1 recognized by anti-DJ-1 antibody. The blot shown in (A) was stripped and re-probed with anti-DJ-1 antibody. Again, this was replicated with the four different samples. Arrow denotes a band of immunoreactivity at approximately 45 kDa. Overlay of the films shown in panels A and B documented coincidence of the 45 kDa band, representing glutathionylated DJ-1. (C) Electrospray ionization mass spectrum of intact recombinant DJ-1. The mass spectrum shows a distribution of charged states (+11 to +28) for intact DJ-1 protein. Deconvolution of the spectrum reveals a protein mass of 20036.3 Da, which is identical to the theoretical mass of recombinant DJ-1. (D) Electrospray ionization mass spectrum of glutathionylated DJ-1. Incubation of DJ-1 protein with 5 mM GSH and 1 mM diamide led to an increase in protein mass of 305-Da (blue ion series, deconvoluted mass = 20341.5 Da) or 610-Da (red ions series, deconvoluted mass = 20646.7 Da), corresponding to covalent modification of DJ-1 with one or two glutathione moieties, respectively.

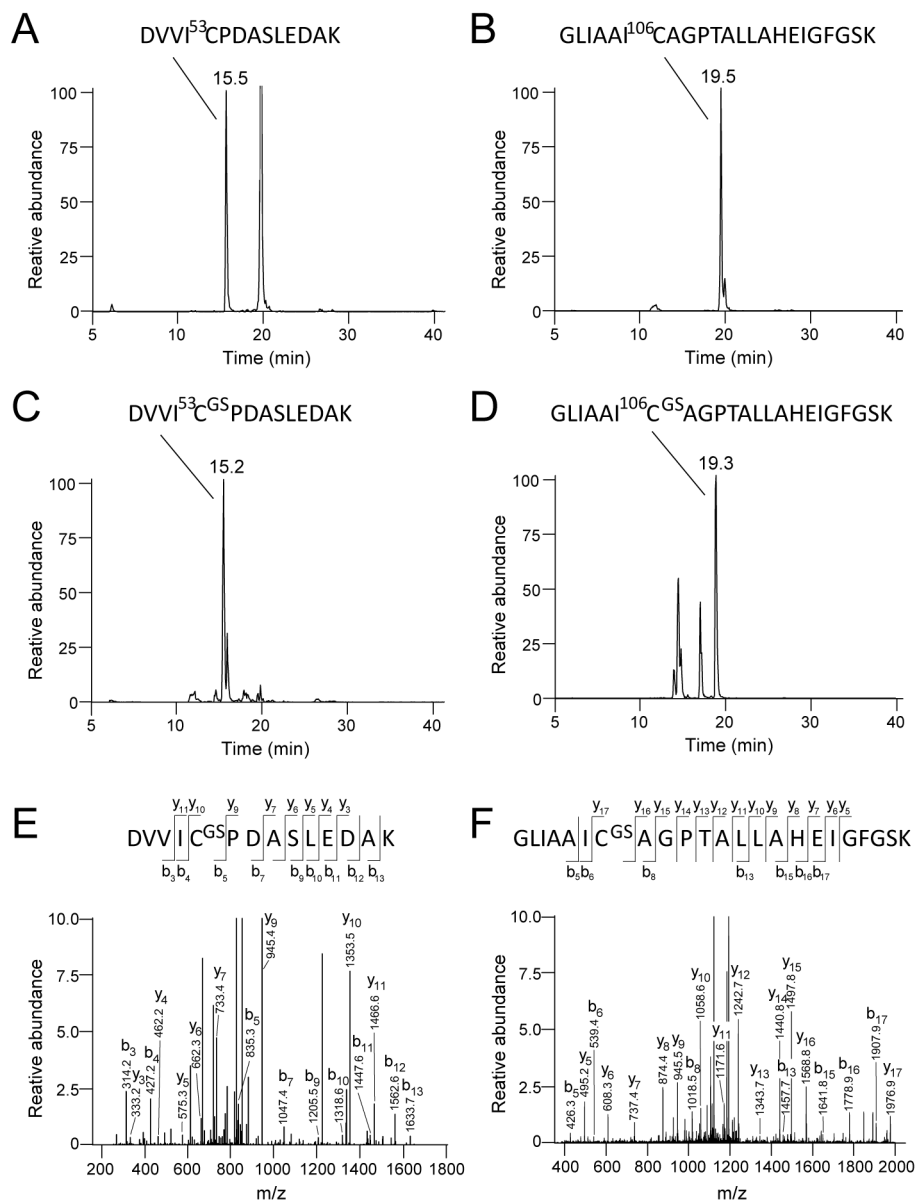


Figure 4. Identification of the glutathionylation sites of DJ-1 by tandem mass spectrometry. Selective Ion chromatographs of tryptic digest of DJ-1 treated with 5 mM GSH and 1 mM diamide revealed two Cys containing peptides (C) DVVIC⁵³PDASLEDAK ($m/z = 766.5 [M+2H]^{+2}$) and (D) GLIAAIC¹⁰⁶AGPTALLAHEIGFGSK ($m/z = 1258.2 [M+2H]^{+2}$), with corresponding masses increased by 305-Da as compared to the untreated sample shown in (A) and (B). (E) and (F): MS/MS fragmentation pattern of these peptides identifies Cys53 and Cys106 residues as the sites of glutathionylation. According to Biemann nomenclature, the series of b and y ions are identified in the mass fragmentograms and in the diagrams of the Cys-containing peptides at the top of panels E and F. For example, the site of glutathionylation at Cys-53 is identified by the loss of 408 mass units for the y₉ ion versus the y₁₀ ion, corresponding to the loss of the cysteinylglutathione moiety.

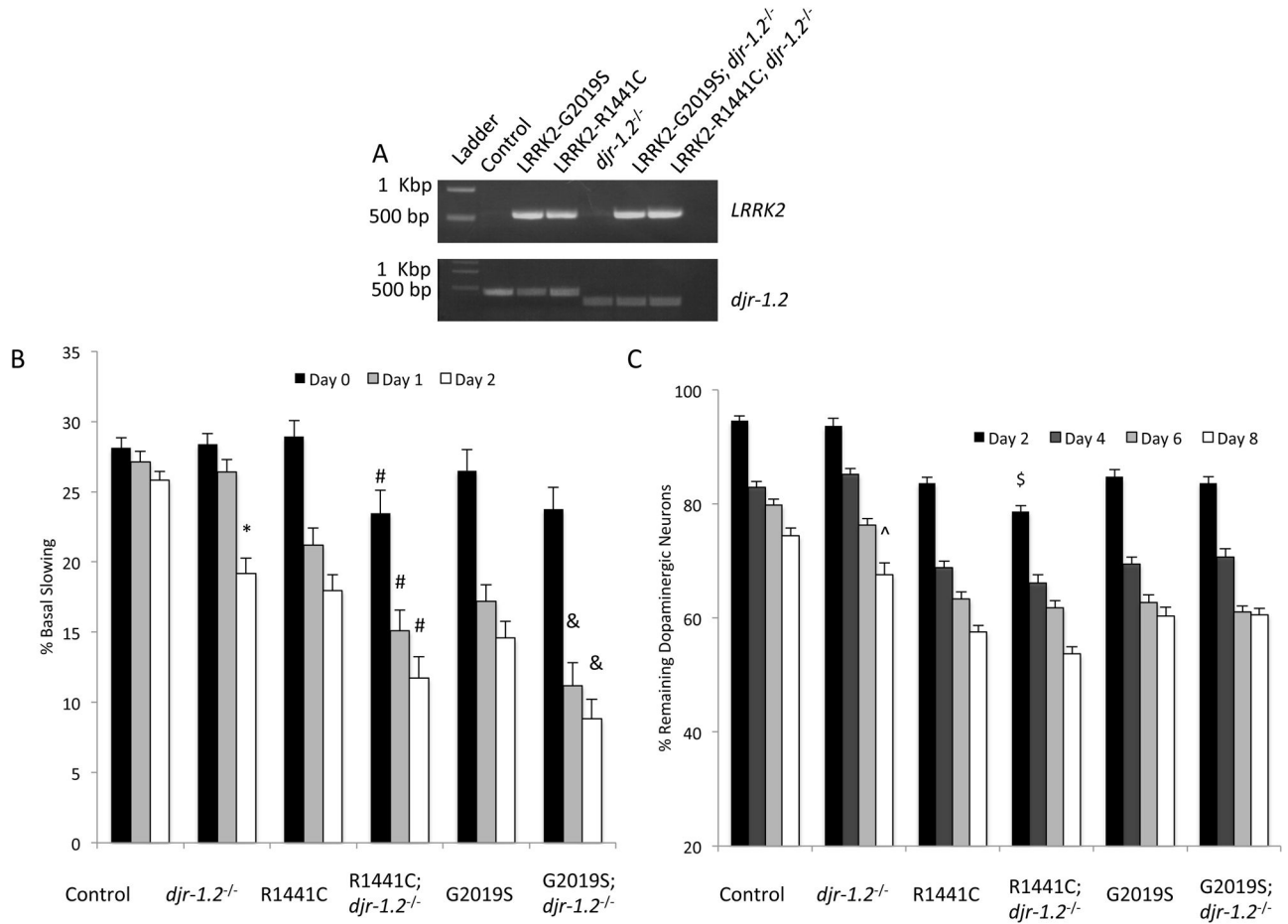


Figure 5. Loss of DJR-1.2 induces PD phenotypes in *C. elegans*, exacerbates mutant LRRK2-mediated impairment of dopamine dependent behavior, but does not consistently exacerbate mutant LRRK2-mediated dopaminergic neurodegeneration

(A) Representative PCR analysis of *LRRK2* (upper panel) and WT *djr-1.2* or mutant *djr-1.2* DNA (lower panel) in *C. elegans* lines. Primers of *djr-1.2* were designed to generate two distinct PCR products depending on the inclusion (upper band) or exclusion (lower band) of the first exon of *djr-1.2* in the *C. elegans* lines. (B) Basal slowing data demonstrating that loss of DJR-1.2 results in dysfunction of dopamine-dependent movement, and exacerbates R1441C- and G2019S-mediated dysfunction of dopamine-dependent movement; * $p < 0.05$, comparing control to *djr-1.2*^{-/-}; # $p < 0.05$, comparing R1441C to R1441C; *djr-1.2*^{-/-}; & $p < 0.05$, comparing G2019S to G2019S; *djr-1.2*^{-/-}. (C) Dopaminergic degeneration data demonstrating that loss of DJR-1.2 results in late onset degeneration, but does not exacerbate mutant LRRK2-mediated degeneration consistently. ^ $p < 0.05$ comparing control to *djr-1.2*^{-/-}, \$ $p < 0.05$ comparing R1441C to R1441C; *djr-1.2*^{-/-}. For (B) and (C) error bars represent SEM. n = 3 independent groups of worms for all days, ~10 worms (B) or ~30 worms (C) per genotype per experiment. Statistical analysis was completed using one-way ANOVA followed by Tukey’s post-hoc test. Statistical comparisons between WT and LRRK2 expressing lines recapitulated previously published experimental data [20], and were not included to simplify the figure.

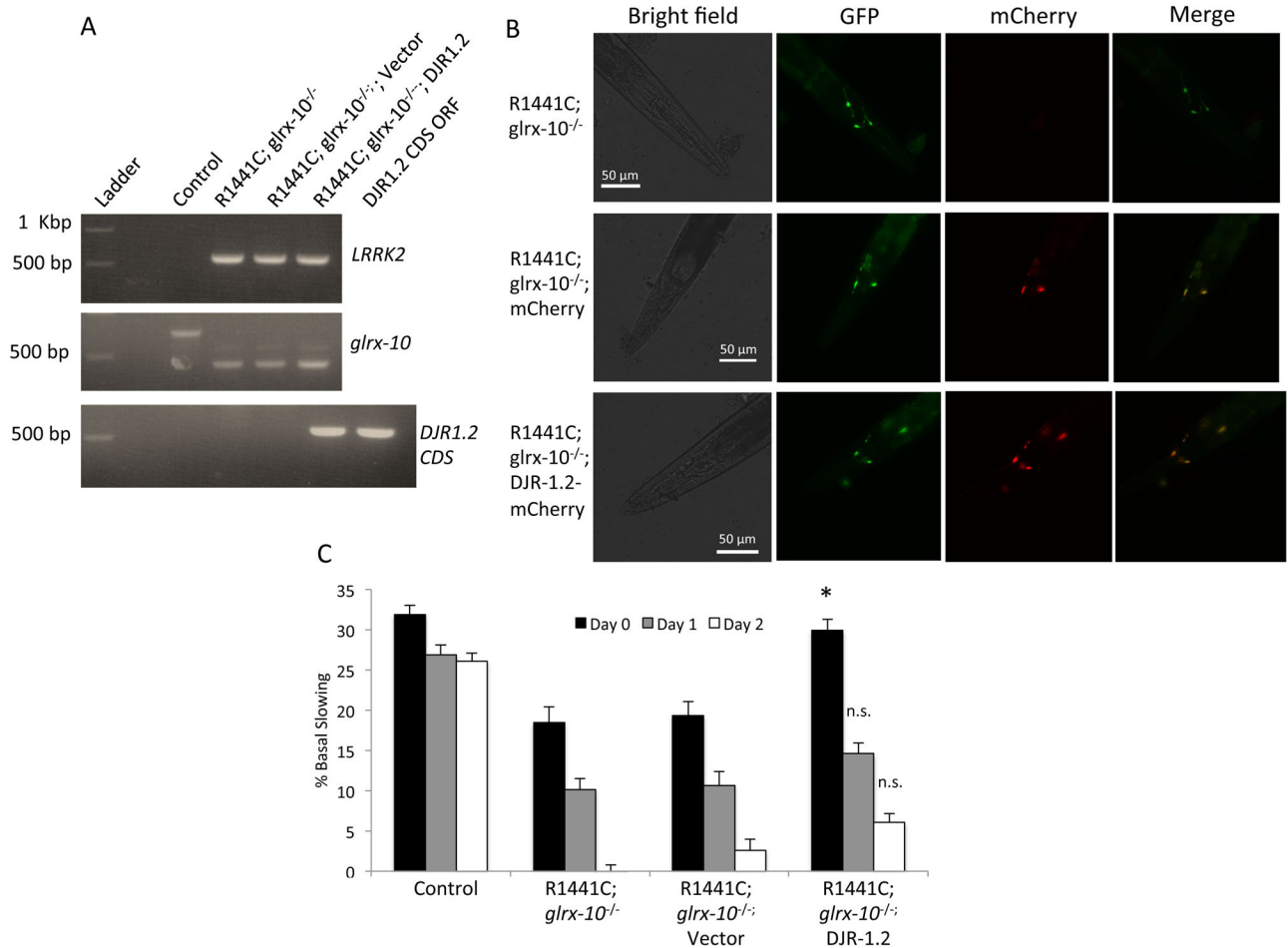


Figure 6. Over-expression of DJR-1.2 partially compensates for the loss of GLRX-10
 (A) PCR analysis of *LRRK2* (upper panel), WT or mutant *glrx-10* DNA (middle panel), and coding DNA sequence (CDS) of *djr-1.2* (lower panel) indicating the generation of R1441C; *glrx-10*^{-/-}; vector (mCherry) and R1441C; *glrx-10*^{-/-}; DJR-1.2-mCherry expressing *C. elegans* lines. (B) Fluorescent images of the head region of the newly generated *C. elegans* depicting the expression of Pdat-1::mCherry and Pdat-1::DJR-1.2:mCherry in the dopaminergic neurons. Expression of GFP tagged dopaminergic neurons co-localize with expression of both Pdat-1::mCherry and Pdat-1::DJR-1.2::mCherry indicating dopaminergic expression of mCherry and DJR-1.2-mCherry, respectively. (C) Basal slowing data demonstrating that over-expression of DJR-1.2 partially rescues the LRRK2-R1441C mediated dysfunction of dopamine-dependent movement in the *glrx-10*^{-/-} background. * $p < 0.05$ compared to R1441C; *glrx-10*^{-/-}; vector. Error bars represent SEM. $n = 3$ independent groups of worms for all days, 10 worms per genotype per experiment. Statistical analysis was completed using one way ANOVA followed by Tukey's post-hoc test.

Table 1

Site	Peptide	Native (Area)	¹⁵ N-labeled(Area)	Modified (%)
C46	DPVQCSR	14,214,288	15,332,719	7.3
C53	DVVICPDASLEDAK	69,277,937	229,809,312	70.0
C106	GLIAAICAGPTALLAHEIGFGSK	268,610,760	1,179,067,215	77.2 ^A

Quantitation of estimated glutathionylation of cysteines of DJ-1. Glutathionylation of cysteine residues 46, 53, and 106 were estimated using unmodified ¹⁵N DJ-1 as an internal standard. Quantification of modified tryptic peptides were estimated by subtraction of the peak areas of the non-modified peptides in the treated ¹⁴N-samples from those in the untreated ¹⁵N-samples (Supplemental Figure 3).

^AExample of calculation of extent of glutathionylation of Cys106: $1,179,067,215$ (¹⁵N non-modified area) - $268,610,760$ (¹⁴N non-modified area) = $910,456,455$ (estimated area of the glutathionylated peptide). $910,456,455$ (estimated area of the glutathionylated peptide) / $1,179,067,215$ (¹⁵N non-modified area) $\times 100 = 77.2\%$ (Estimated percentage of glutathionylated peptide as compared to total internal standard).

Supplementary Material

Regulation of DJ-1 by glutaredoxin 1 *in vivo* – implications for Parkinson's disease

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Supplemental Figure Legends

S1. QPCR and Western blot data for transgenic Grx1 and WT mice. (A) QPCR analysis of DJ-1 mRNA from WT and *Grx1*^{-/-} mice in the C57BL/6 background. N=3 mice per group, no statistical difference *via* Student's T-Test. (B) Western blot analysis of transgenic mice over-expressing hGrx1; tissue probed with anti-Grx1, anti-DJ-1, and anti-actin antibodies, respectively. (C) Western blot analysis of WT and *Grx1*^{-/-} mice in the BALB/c background; tissue probed with anti-Grx1, anti-DJ-1, and anti-actin antibodies, respectively. N=3 mice per group. (D) QPCR analysis of DJ-1 mRNA from WT and *Grx1*^{-/-} mice in the BALB/c background. N=3 mice per group, no statistical difference *via* Student's T-Test.

S2. Western blot data for human samples. (A) Western blot analysis of samples extracted from human PBMCs probed with anti-Grx1, anti-DJ-1, and anti-actin antibodies, respectively; n= 14 independent biological samples. (B) Western blot analysis of samples extracted from human midbrain tissue probed with anti-Grx1, anti-DJ-1, and anti-actin antibodies, respectively; n = 14 independent biological samples.

S3. Validation of the DJ-1 antibody. (A) Western blot analysis of immunoprecipitated DJ-1. Left lane, reduced (+ DTT); right lane, non-reduced (- DTT). (B) Western blot analysis of brain tissue homogenates from DJ-1^{-/-} and WT mice treated with DTT and probed with anti-DJ-1, and anti-actin antibodies, respectively. N=2 mice per genotype.

S4. Validation of the DJ-1 glutathionylation. DJ-1 was immunoprecipitated from mouse brain lysate, run under non-reducing conditions and transferred to PVDF membrane. Half of the blot was soaked in TBST buffer without DTT, and the other half was soaked in TBST buffer containing 100 mM DTT. The blots then were probed with the anti-GSH antibody, stripped, and re-probed with the anti-DJ-1 antibody for confirmation of DJ-1 protein. Loss of anti-GSH signal in the presence of DTT indicates that the disulfide reducing agent ablates the anti-GSH signal consistent with loss of the glutathionyl mixed disulfide adduct of DJ-1 (A) Western blot analysis of the membrane incubated with TBST (B) Western blot analysis of membrane incubated in TBST and DTT. Arrows indicate the ~45 kDa immunoreactive band.

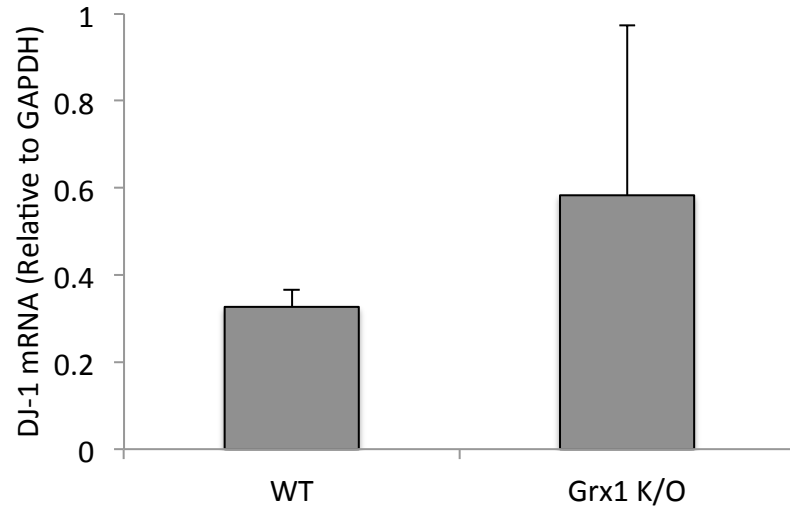
S5. Quantification of glutathionylated DJ-1 peptides. (A) and (B) represent selective ion chromatograms for two cysteine-containing DJ-1 peptides unmodified by glutathionylation; (C) and (D) correspond to the same peptides obtained from ¹⁵N-labeled DJ-1, serving as internal standards. Areas under the ion intensity peaks, colored in gray were used to calculate ratios between ¹⁵N-enriched and natural abundance peptides to determine the fraction of each peptide covalently modified by glutathione. (E) and (F) represent MS spectra averaged at the peak of ion intensities for DVVIC⁵³PDASLEDAK and GLIAAIC¹⁰⁶AGPTALLAHEIGFGSK peptides, respectively. Ions and their isotopic distribution for both ¹⁵N-enriched (blue) and natural abundance (black) peptides are shown.

S6. Quantification of mCherry and DJR-1.2:mCherry protein content in the dopaminergic neurons of *C. elegans* by fluorescence microscopy. Quantification indicates that content of mCherry and DJ-1:mCherry protein (relative to GFP expression specifically in the dopaminergic neurons) are equivalent and do not change over time.

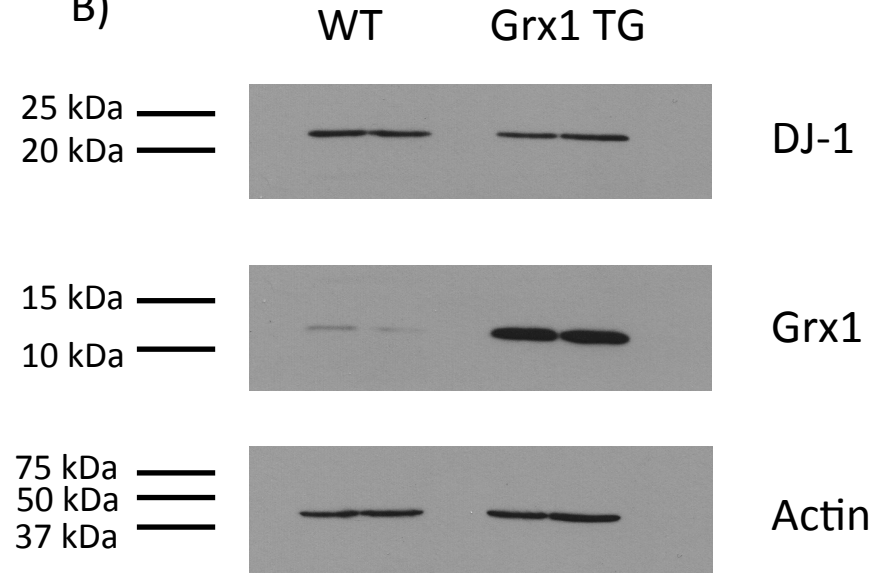
S7. Quantification of relative DJ-1 protein content in Neuro-2A cells transfected with scRNA/Grx1 shRNA and treated with individual protease inhibitors from the protease inhibitor cocktail. Neuro-2A cells transfected with scRNA/Grx1 shRNA were treated with individual protease inhibitors or with DMSO (**D**; vehicle) for 48 hr with fresh medium containing inhibitor or vehicle replaced every 12 hr. Panels show representative Western blots of DJ-1 along with respective loading control β -tubulin (Tub) and densitometric analyses. The inhibitors added with scRNA/shRNA are: (A) 52 μ M 4-(2-aminoethyl) benzenesulfonyl fluoride, (**Ae**), (B) 40 nM aprotinin, (**Ap**), (C) 2 μ M bestatin (**Be**), (D) 700 nM E-64 (**E64**), (E) 1 μ M leupeptin (**Le**) and (F) 750 nM pepstatin A (**Pe**). All data were analyzed using one-way ANOVA followed by Student-Newman-Keuls test to determine significant differences with * representing p < 0.05. n = 3-6 independent samples in each experimental group. None of the individual protease inhibitors tested alone afforded protection of DJ-1 from degradation, unlike the complete cocktail.

Supplemental Figure S1

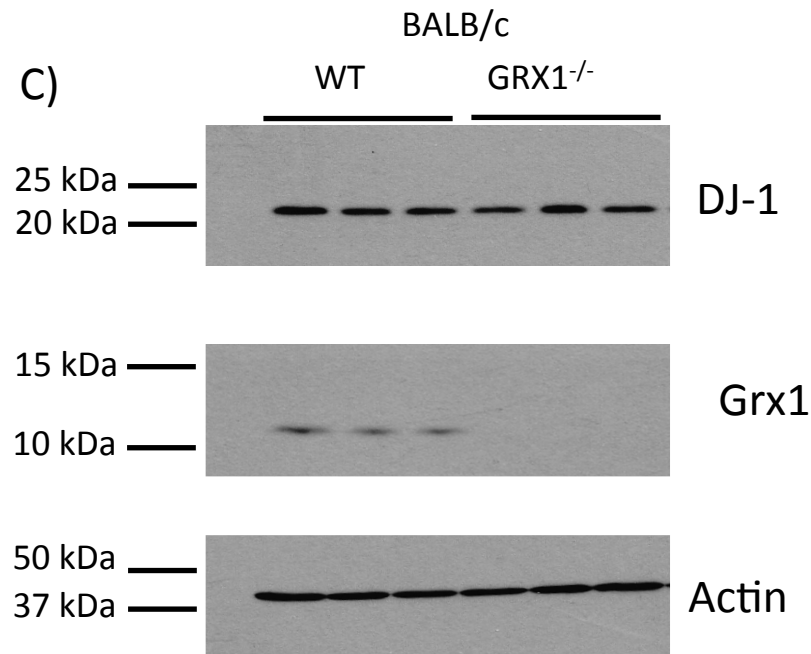
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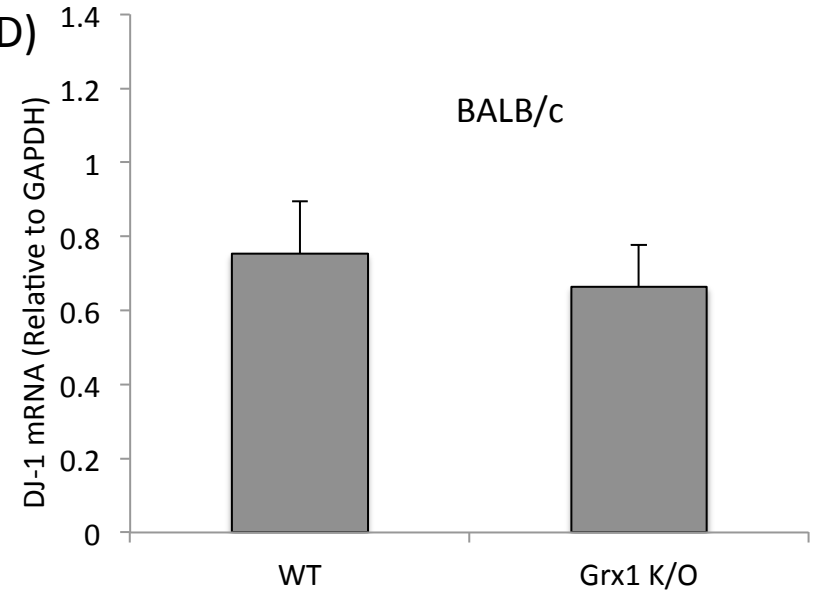
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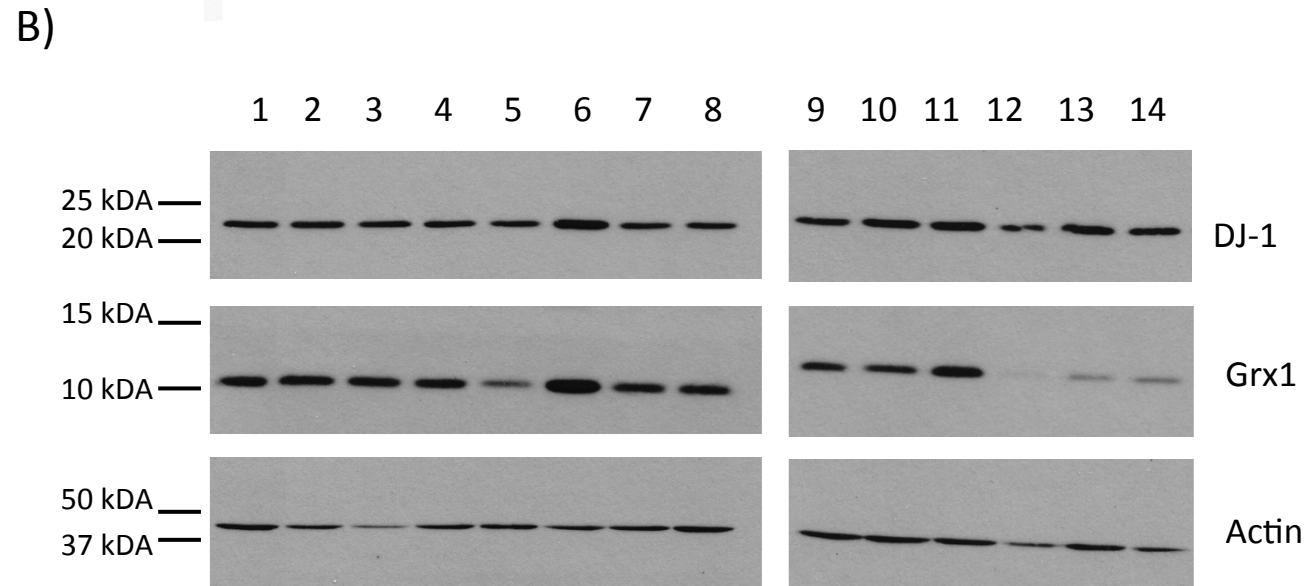
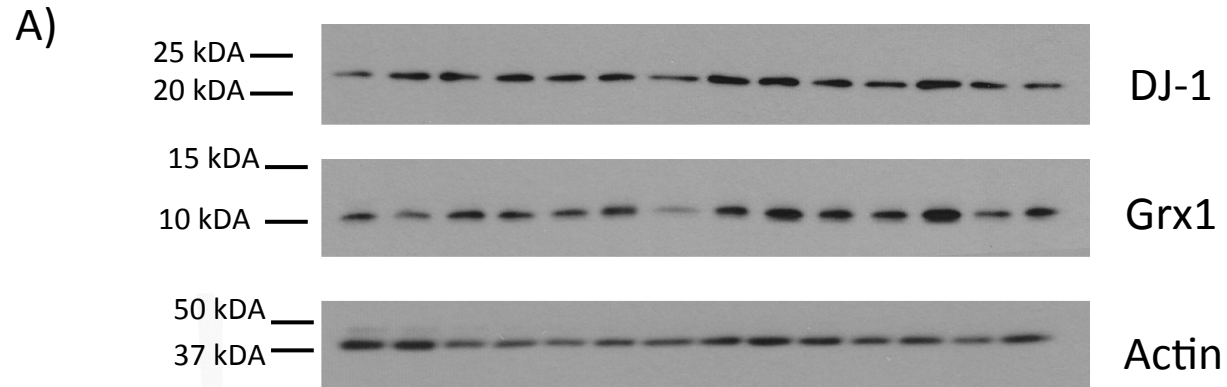
C)



D)

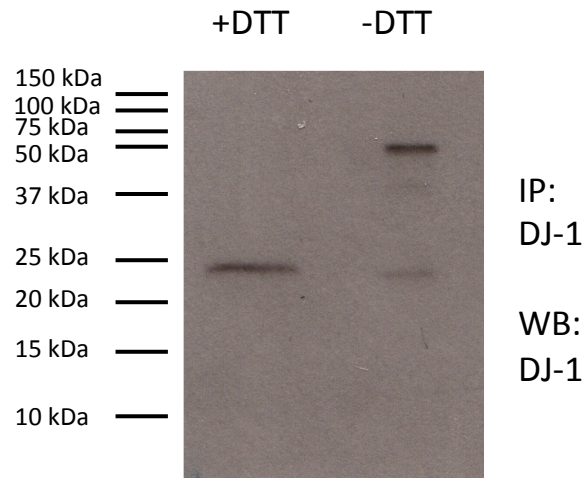


Supplemental Figure S2

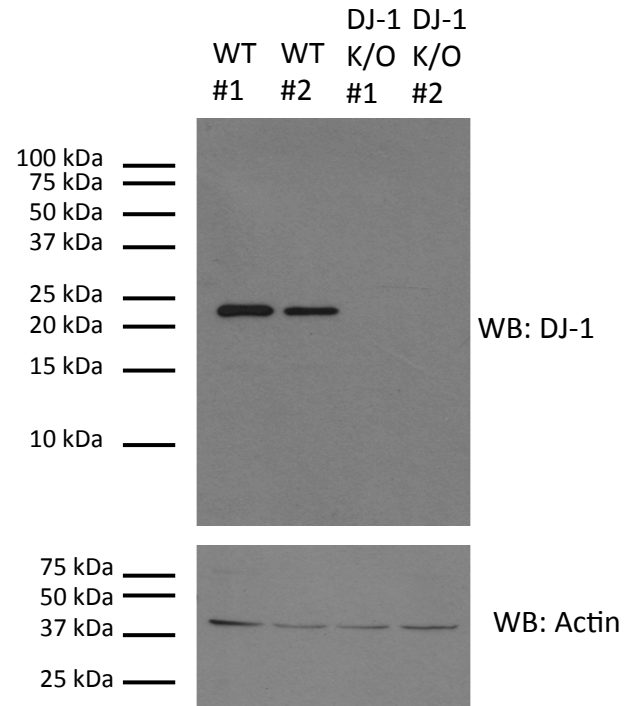


Supplemental Figure S3

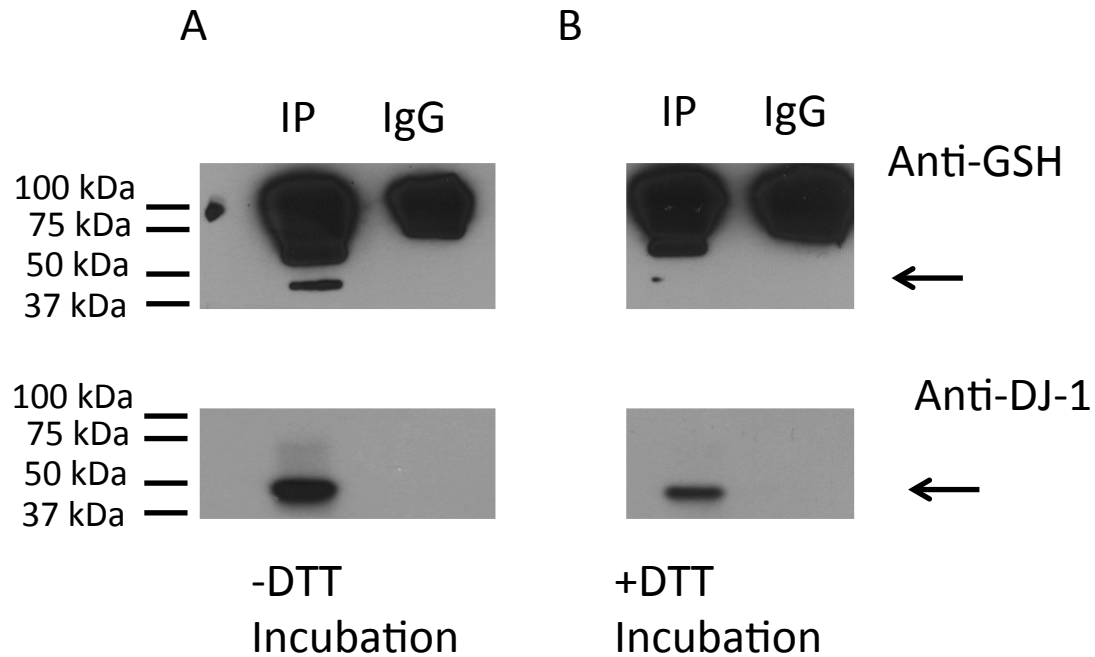
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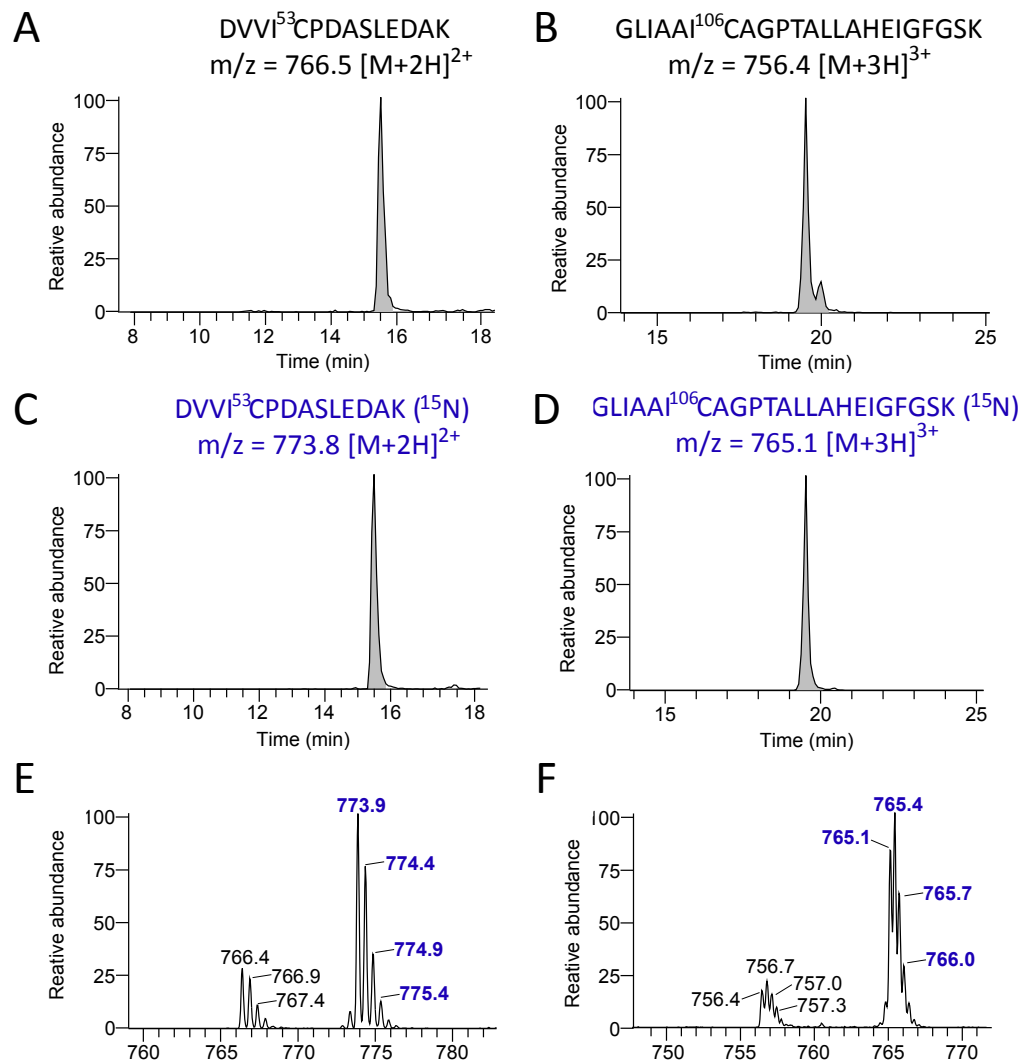
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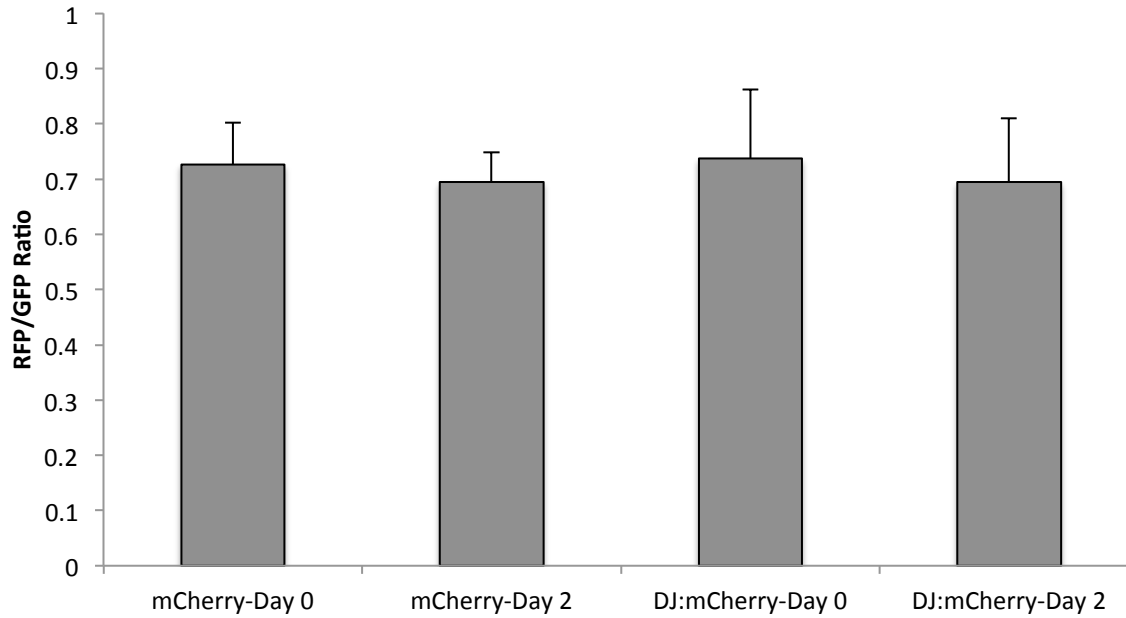
Supplemental Figure S4



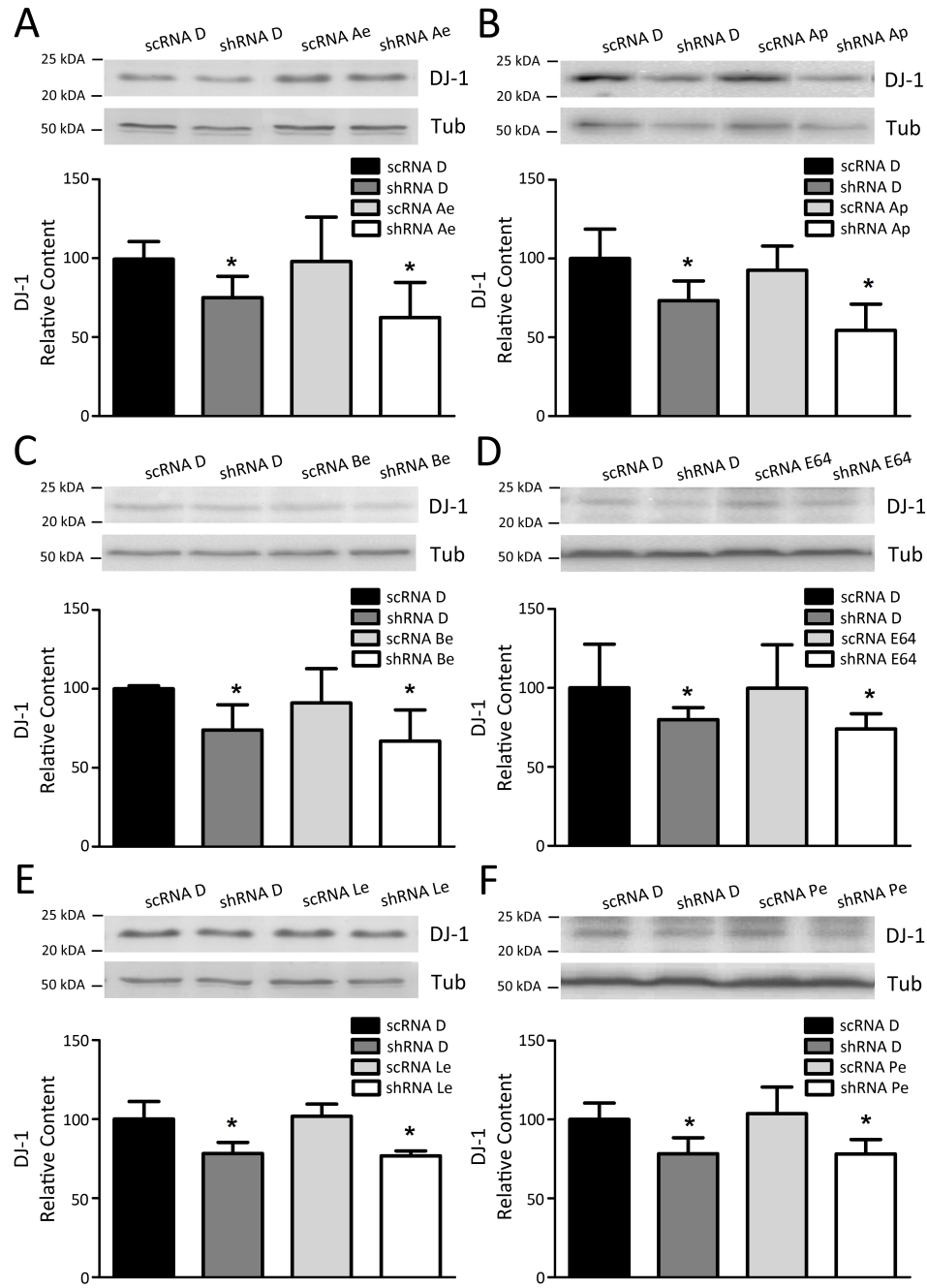
Supplemental Figure S5



Supplemental Figure S6



Supplemental Figure S7



Supplemental Tables

A Patient information for PBMC analysis

Patient Reference #	Age	Sex	Diagnosis
137	62	F	PD
219	67	M	PD
228	77	M	PD
236	67	M	PD
254	65	M	PD
256	74	M	PD
257	80	M	PD
260	46	M	PD
262	73	M	PD
264	57	M	PD
266	70	M	PD
268	64	M	PD
270	75	M	PD
271	58	F	PD

Cases used for Western blot analysis from PBMC lysate.

B Patient information for brain lysate analysis

Patient Reference #	Sex	Age	Diagnosis
P08-40	F	86	PD
P08-41	F	83	PD
P08-42	M	75	PD
P08-43	M	71	PD
P08-34	M	51	Control
P08-35	M	89	Control
P08-36	F	86	Control
P08-37	M	73	Control
P08-38	M	69	Control
PD1	M	77	PD
PD2	F	49	PD
PD3	F	61	PD
PD4	M	73	PD

Cases used for Western blot analysis from post mortem midbrain tissue.

C

Designation	Genotype	Strain	Reference
Control (GFP)	<i>lin-15(n765ts)X; cwrIs730[Pdat-1::GFP, lin-15(+)]</i>	SGC730	[6]
R1441C	<i>lin-15(n765ts)X; cwrIs851[Pdat-1::GFP, Pdat-1::LRRK2(R1441C), lin-15(+)]</i>	SGC851	[6]
G2019S	<i>lin-15(n765ts)X; cwrIs856[Pdat-1::GFP, Pdat-1::LRRK2(G2019S), lin-15(+)]</i>	SGC856	[6]
R1441C; <i>glrx-10^{-/-}</i>	<i>glrx-10(tm4634)I; cwrIs851</i>	SGC302	[20]
<i>djr-1.2^{-/-}</i>	<i>djr-1.2(tm1348)V</i>	tm1346	NBP
R1441C; <i>djr-1.2^{-/-}</i>	<i>djr-1.2(tm1348)V; cwrIs851</i>	SGC10	This study
G2019S; <i>djr-1.2^{-/-}</i>	<i>djr-1.2(tm1348)V; cwrIs856</i>	SGC11	This study
R1441C; <i>glrx-10^{-/-}</i> ; DJR-1.2	<i>glrx-10(tm4634)I; cwrIs851; cwrEx20[Pdat-1::DJR-1.2::mCherry]</i>	SGC20	This study
R1441C; <i>glrx-10^{-/-}</i> ; Vector	<i>glrx-10(tm4634)I; cwrIs851; cwrEx22[Pdat-1::mCherry]</i>	SGC22	This Study

NBP: National Bioresource Project (Japan)

Caenorhabditis elegans strains used in this study