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## **LRRK2 mutations cause mitochondrial DNA damage in iPSC-derived neural cells from Parkinson's disease patients: Reversal by gene correction**

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### **Abstract**

Parkinson's disease associated mutations in *leucine rich repeat kinase 2 (LRRK2)* impair mitochondrial function and increase the vulnerability of induced pluripotent stem cell (iPSC)-derived neural cells from patients to oxidative stress. Since mitochondrial DNA (mtDNA) damage can compromise mitochondrial function, we examined whether *LRRK2* mutations can induce damage to the mitochondrial genome. We found greater levels of mtDNA damage in iPSC-derived neural cells from patients carrying homozygous or heterozygous *LRRK2* G2019S mutations, or at-risk individuals carrying the heterozygous *LRRK2* R1441C mutation, than in cells from unrelated healthy subjects who do not carry *LRRK2* mutations. After zinc finger nuclease-mediated repair of the *LRRK2* G2019S mutation in iPSCs, mtDNA damage was no longer detected in differentiated neuroprogenitor and neural cells. Our results unambiguously link *LRRK2* mutations to mtDNA damage and validate a new cellular phenotype that can be used for examining pathogenic mechanisms and screening therapeutic strategies.

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### **Conflict of interest**

J.W.L., B.S. and S.H.Z. filed a patent application about the genomic repair of *LRRK2* in fibroblasts and iPSCs. The remaining authors declare no conflicts of interest.

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## Keywords

Parkinson's disease; LRRK2; mitochondrial DNA damage; stem cells

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## Introduction

Mutations in *leucine-rich repeat kinase 2 (LRRK2)* are associated with sporadic and familial forms of Parkinson's disease (PD) (Paisan-Ruiz et al., 2004; Satake et al., 2009; Simon-Sanchez et al., 2009; Zimprich et al., 2004). Mitochondrial impairment is considered to be a critical factor in the pathogenesis of both sporadic and genetic forms of PD (Henchcliffe and Beal, 2008). Recently, using PD patient-specific induced-pluripotent stem cells (iPSCs) carrying *LRRK2* G2019S and R1441C mutations, both *LRRK2* mutations were linked to compromised oxidative phosphorylation and mitochondrial dynamics, rendering neural cells more vulnerable to mitochondria-associated stress (Cooper et al., 2012). However, the mechanisms by which *LRRK2* mutations lead to a loss of mitochondrial function are poorly understood.

In PD, reactive oxygen species (ROS) damage lipids and proteins (Sherer and Greenamyre, 2005), but less is known about damage to mtDNA (Sanders and Timothy Greenamyre, 2013). DNA damage, defined as any modification of DNA that can alter its coding properties or interfere with normal function in transcription or replication (Lindahl, 1993; Rao, 1993), is distinct from mutations, which are a change in the base sequence of the DNA. However, damage to mtDNA may lead to mtDNA mutations. The mitochondrial genome is particularly susceptible to oxidative damage, likely due to the proximity of mtDNA to ROS production at the inner mitochondrial membrane and the lack of protection afforded by histones (Yakes and Van Houten, 1997). Mitochondrial DNA damage can compromise metabolic functions, predispose to ROS generation and trigger cell death. Accumulation of mtDNA damage is a particular problem for the brain because neurons are post-mitotic and long-lived. In order to study these issues in a neuronal context, we applied cellular reprogramming technology to examine whether *LRRK2* mutations lead to mtDNA damage.

## Materials and Methods

### Induced pluripotent stem cell (iPSC) generation, propagation and differentiation and repair of the LRRK2 G2019S mutation

Institutional Review Boards approved the study. The subjects gave written, informed consent for participation in this study. Fibroblasts from a 4 mm skin punch biopsy were cultured using standard techniques (Byers et al., 2011; Cooper et al., 2012). The fibroblasts were reprogrammed using OCT4, SOX2, KLF4 and CMYC and standard methods (Chan et al., 2009). The additional iPSC lines are available from the Coriell Stem Cell Biobank (LRRK2 R1441C ND34394, ND34393, ND35884; LRRK2 G2019S ND35367) (Cooper et al., 2012). The iPSCs were propagated on irradiated/mitomycin-C inactivated mouse embryonic fibroblasts (Global Stem, Rockville, MD) and manually passaged (Cooper et al., 2012; Nguyen et al., 2011). The details of individual iPSC clones are listed in Supplemental Table 1. ZFN-mediated genomic repair of LRRK2 G2019S in iPSCs was performed.

### Assessment of iPSC pluripotency

Teratoma analyses of selected iPSC clones were performed in nonobese diabetic/severe combined immunodeficient mice (Applied StemCell). Karyotypes were analyzed by GTW banding at >400 band resolution (Cytogenetics Laboratory, Stanford University School of Medicine). Immunocytochemistry was performed as described (Mak et al., 2012) using primary antibodies raised against OCT4, SOX2, Tra1-60 and SSEA4 (All from Millipore).

Images of the immunofluorescence were taken using the Odyssey Infrared Imaging System (LI-COR Biosciences, USA). For RT-PCR, total RNA was extracted using the RNeasy Micro kit (Qiagen, Valencia, CA) and 150ng RNA was used for reverse-transcription into cDNA using the iScript cDNA Synthesis Kit (BioRad, Hercules, CA). Total reaction volume was 20  $\mu$ l; the resulting cDNA sample was diluted such that the final cDNA concentration was 18.75ug/uL. 4  $\mu$ l of the diluted cDNA sample was used as template for qPCR amplification. qPCR was performed using the Biorad CFX96 Real-Time system using Applied Biosystems TaqMan Gene Expression Assays for the following probes: EOMES (Hs00172872\_m1), DNMT3B (Hs00171876\_m1), FOXD3 (Hs00255287\_s1) and FOXA2 (Hs00232764\_m1).

### **iPSC differentiation into neuroprogenitor cells and neural cells**

Two differentiation protocols were used to generate cells for analysis of mtDNA damage. The analysis of mtDNA damage across neural cells from multiple patients and healthy subjects (Fig. 1) used a differentiation protocol that had previously been used to determine mitochondrial deficits in neural cells (Cooper et al., 2012). The differentiation of immature neuroprogenitor cells and more mature neural cells from repaired iPSCs for analyses of mtDNA damage was performed as described (Mak et al., 2012). MtDNA damage was increased in iPSC-derived cells carrying *LRRK2* mutations differentiated with either protocol.

### **Immunocytochemistry**

Cells were fixed with 4% PFA (EMS, RT15713) for 10 min, washed with PBS (Sigma, T8787), permeabilized with 0.3% Triton x-100 (Sigma, X-100) in PBS for 5 min, and then blocked with 5 % serum (Vector, S-10000) after washing. Cells were incubated with primary antibody overnight at 4°C then incubated with Alexa Fluor 488 (1:200, Invitrogen Molecular Probes, A11029) or Alexa Fluor 555 (1:200, Invitrogen Molecular Probes, A21429). Coverslips were mounted with Vectashield Mounting Medium for Fluorescence with DAPI (Vector Laboratories, H-1200) and imaged on a Nikon Eclipse Ti inverted fluorescence microscope. Antibodies that were used are summarized in (Supplemental Table 2).

### **DNA isolation and quantification for analysis of mtDNA damage**

Coded, pelleted samples of cells were received for blinded analysis. DNA was isolated using a high molecular weight genomic DNA purification kit (QIAGEN Genomic tip). DNA was quantified using the Picogreen dsDNA quantification assay (Molecular Probes). The fluorescence from the Picogreen was measured with a 485 nm emission filter and a 530 nm excitation filter using a microplate reader (SpectraMax Gemini EM). Lambda DNA was used to construct a standard curve in order to determine the concentration of unknown samples. Quality of the DNA prior to QPCR analysis was verified by running the DNA on a 0.6% ethidium bromide-stained agarose gel. Only DNA containing single bands of intact high molecular weight, and which showed negligible evidence of degradation, were used in the DNA damage assays. DNA samples were aliquoted and stored at  $-20^{\circ}\text{C}$ . Samples were thawed only once prior to downstream assays.

### **Quantifying mtDNA damage using the quantitative polymerase chain reaction (QPCR) based assay**

Each cell line was independently differentiated three separate times. These three independently generated cell lines were analyzed blinded. The QPCR assay was performed in technical triplicate, which were then averaged, and the statistics performed on the means generated from the three independent experimental values. Cells were collected in lysis

buffer, centrifuged at  $10,000 \times g$  for 20 min at  $4^{\circ}\text{C}$ . The DNA was precipitated with isopropanol for 48 h at  $-80^{\circ}\text{C}$ . The QPCR assay was performed as described (Ayala-Torres et al., 2000; Santos et al., 2006). To confirm that QPCR assays were performed in the linear range, optimal number of cycles for each template was conducted to show that 50% reduction in the amount of template resulted in about 50% reduction in amplification. QPCR products that demonstrated 40–60% in the amplification of the target sequence when using 50% of the original template were considered acceptable.

The PCR amplification for the 16.2 kb fragment of human mtDNA had an annealing/extension time of 14 min for 28 cycles for NPCs and 33 cycles for iPSC-derived neural cells (Ayala-Torres et al., 2000). 10 ng was used to amplify the human large mtDNA fragment and the primer nucleotide sequences were 14841 and 15149 (Ayala-Torres et al., 2000). The final concentration of magnesium in the PCR for human mtDNA fragments was 1.1 mM. To ensure the quality and specificity, all PCR products were resolved on a 0.6% agarose gel and UV light used to visualize ethidium bromide-stained gels. Relative fluorescence of PCR products was quantified using Picogreen. Assuming a random distribution, the Poisson equation was used to calculate the number of DNA lesions. Based on this equation, the amplification is directly proportional to the fraction of undamaged DNA templates. As such, the average lesion frequency per strand is calculated as  $-\ln A_D/A_O$ , where  $A_D$  is the amplification of the damaged or experimental template, while  $A_O$  is the amplification of the undamaged or control template. Therefore, the results are shown as the number of lesions per strand normalized to 10kb. DNA damage can then be expressed mathematically as the number of DNA lesions per kilobase. All samples were analyzed in a blinded fashion.

### QPCR of a small mtDNA fragment

To ensure that the amplification of the large mtDNA fragment was not due to possible changes in mtDNA steady state levels, we amplified a human small mtDNA fragment (248 bp), since the amplification of this small fragment should be independent of damage. The profile for this PCR amplification was as follows: hot start for 10 min at  $75^{\circ}\text{C}$  when the DNA polymerase was added, an initial denaturation step for 1 min at  $94^{\circ}\text{C}$ , followed by 23 cycles for NPCs and iPSC-derived neural cells of denaturation for 1 min at  $94^{\circ}\text{C}$  and then annealing at  $60^{\circ}\text{C}$  for 45 seconds and extension at  $72^{\circ}\text{C}$  for 45 seconds. To complete the profile, a final extension for 10 min at  $72^{\circ}\text{C}$  was performed. The primer nucleotide sequences 14841 and 14620 were used to amplify the human 248-bp mtDNA fragment (Ayala-Torres et al., 2000). 10 ng of DNA template was used to amplify the human small mtDNA fragment. The PCR product was resolved on a 1.5 % agarose gel and UV light used to visualize ethidium bromide-stained gels. Relative fluorescence of PCR products were quantified using Picogreen.

### Statistical analysis

Data were analyzed by Student's t test or ANOVA using computer software (Prism, GraphPad, La Jolla, CA), and  $p < 0.05$  was considered statistically significant.

## Results and Discussion

Given the mitochondrial deficits of iPSC-derived neural cells from subjects carrying *LRRK2* mutations and the fact that mtDNA damage compromises mitochondrial and neuronal function, we carried out experiments to determine if *LRRK2* PD iPSC-derived neural cells accumulate mtDNA damage (Cooper et al., 2012). iPSCs were derived from three patients carrying the homozygous or heterozygous *LRRK2* G2019S mutation, two asymptomatic subjects carrying the heterozygous *LRRK2* R1441C mutation, and three age-matched healthy subjects without *LRRK2* mutations (Supplemental Table 1, Supplemental Fig 1) (Cooper et

al., 2012). Multiple iPSC clones were examined from each individual carrying the *LRRK2* R1441C mutation. We have previously shown that our protocol for iPSC-derived neural cells yields similar numbers of dopaminergic vs non-dopaminergic functional neurons (Cooper et al., 2012). Upon neuronal differentiation of the iPSC lines, cultures were harvested, pelleted and coded for blinded analysis. After receipt of the coded samples, DNA was purified and a quantitative polymerase chain reaction (QPCR)-based assay specific for the mitochondrial genome determined mtDNA damage. This method is based on the principle that various forms of DNA damage have the propensity to slow down or block DNA polymerase progression (Santos et al., 2006). Thus, if equal amounts of mtDNA from experimental and control specimens are amplified under identical conditions, the mtDNA sample with the least mtDNA damage will produce the greatest amount of PCR product (Santos et al., 2006). Using this approach, a significant increase in levels of mtDNA damage was found in neural cells derived from individuals carrying either the homozygous or heterozygous *LRRK2* G2019S (black bars,  $p < 0.0001$  ANOVA) or heterozygous R1441C (grey bars,  $p < 0.0001$  ANOVA) mutations compared to neural cells from healthy subjects (white bars, Fig 1a). Mitochondrial DNA copy number was similar across all clones (Fig 1b).

While increased levels of mtDNA damage in *LRRK2* neural cells were observed across the different pathogenic mutations, across multiple clones from single individuals and in siblings carrying the same mutation (Fig 1), we used two additional approaches to further strengthen our interpretation that *LRRK2* mutations induce mtDNA damage. *First*, we measured mtDNA damage in iPSC-derived neuroprogenitor cells (NPCs) from two brothers. NPCs from a PD patient carrying the heterozygous *LRRK2* G2019S mutation (iPSC clone L4a) showed greater levels of mtDNA damage compared to his healthy brother who did not carry the *LRRK2* mutation (iPSC clone C3; L4a; Fig 2,  $p = 0.0002$ ). *Second*, we addressed the issue of genetic and biological variability between patient and healthy subject control iPSCs. Ideally, isogenic cell lines that differ from the original culture lines only by a disease-causing mutation should be used for study. Without such isogenic lines, there may be difficulties in data interpretation, because personal genomic variation may cause functionally relevant differences between individuals. We therefore used zinc finger nucleases (ZFNs) to repair the *LRRK2* G2019S mutation (iPSC clone L4b<sup>WT/WT</sup>, Supplemental Table 1). In parallel, we used an iPSC clone from the same parental iPSC line (iPSC clone L4a) that was not modified during the ZFN process and had retained the *LRRK2* G2019S mutation (unmodified mutant iPSC clone L4e<sup>Unmod</sup>, Supplemental Table 1). Before assessing mtDNA damage, we differentiated repaired and control unmodified *LRRK2* G2019S iPSC clones into neuroprogenitor cells (NPCs) and neural cells (Mak et al., 2012). Immunocytochemistry revealed that while differentiated cultures expressed general neuronal markers (synapsin and alpha-synuclein), there was robust expression of markers for dopaminergic neurons (TH and VMAT2) in both mutant and gene-corrected cultures (Fig 3). Additionally, some neurons expressed GABA, but astroglial cells were not seen. The NPCs and neural cells differentiated from the repaired *LRRK2* G2019S iPSCs showed less mtDNA damage than otherwise isogenic cells from the control *LRRK2* G2019S unmodified iPSCs ( $p < 0.0001$  and  $p < 0.002$ , respectively; Fig 4a,b). The number of mtDNA genomes in neural cells and NPCs differentiated from repaired and control *LRRK2* G2019S iPSCs was similar ( $1.0 \pm 0.009$ ,  $1.0 \pm 0.002$  respectively). Mitochondrial DNA damage levels were also reversed to healthy control levels (Supplemental Fig 2), using a second independently generated ZFN corrected line (iPSC clone L4f<sup>WT/WT</sup> Supplemental Table 1).

Next, we examined whether cellular reprogramming or a neural phenotype was required for the mtDNA damage phenotype observed in differentiated NPCs and neural cells from *LRRK2* G2019S carriers. No differences in mtDNA damage levels between healthy subjects and *LRRK2* G2019S mutation carriers were detected in either fibroblasts (FBCs clones 1815

and 1828,  $-0.01 \pm 0.005$  lesions/10kb vs.  $-0.01 \pm 0.003$  lesions/10kb;  $p = 0.2$ ) or undifferentiated iPSCs (iPSC clones L4b<sup>WT/WT</sup> and L4e<sup>Unmod</sup> Supplemental Table 1,  $-0.01 \pm 0.02$  lesions/10kb vs.  $-0.01 \pm 0.02$  lesions/10kb;  $p = 0.99$ ). Mitochondrial DNA copy number was similar across all clones ( $1.02 \pm 0.007$ ,  $1.0 \pm 0.002$  FBCs, iPSCs respectively). Therefore, mtDNA damage is only observed in iPSC derived-NPCs and neural cells from individuals carrying the *LRRK2* G2019S mutation, and not in FBCs or undifferentiated iPSCs. Thus, the observed mtDNA damage is a neural specific phenotype.

Here we demonstrate that *LRRK2* mutations are associated with mtDNA damage, even in neural cells derived from presymptomatic mutation carriers. Thus, mtDNA damage could be an early event in the pathogenesis of PD. The use of ZFNs to repair the genetic mutation in otherwise isogenic cells abrogated the mitochondrial phenotype, thereby providing strong evidence that *LRRK2* mutations *cause* mtDNA damage in neural cells. The mechanisms that result in mitochondrial dysfunction in PD include intrinsically high levels of reactive oxygen species (ROS) (Guzman et al., 2010) and environmental factors that can cause significant oxidative damage and neurodegeneration (Betarbet et al., 2000; Langston et al., 1983). Whether the robust and reproducible mtDNA damage phenotype causes *LRRK2* associated mitochondrial dysfunction or is a consequence thereof requires further investigation; however, *LRRK2*-induced mtDNA damage is a new cellular phenotype and molecular marker of disease-causing *LRRK2* mutations. Having an easily quantifiable phenotype will now allow us to dissect cause-effect relationships, underlying mechanisms of *LRRK2* pathogenesis and potential responses to therapeutic interventions.

To date, a few reports suggest a modest increase in ROS as a result of pathogenic *LRRK2* mutations (Angeles et al., 2011; Heo et al., 2010). In addition, pathogenic *LRRK2* mutations can increase susceptibility to oxidative stress (Cooper et al., 2012; Ng et al., 2009; Nguyen et al., 2011). While *LRRK2* mutations were not associated with problems regulating  $O_2^-$  levels by iPSC-derived neural cells (Cooper et al., 2012), measurement of ROS levels at different stages of iPSC differentiation would elucidate whether levels of ROS are correlated with levels of mtDNA damage observed in the current study.

Because *LRRK2* mutations are a common cause of both sporadic and autosomal dominant PD, examination of the molecular targets of *LRRK2* and their potential roles in mtDNA damage is likely to provide critical mechanistic and therapeutic insights into both *LRRK2*-related and idiopathic PD. The QPCR-based assay we used simultaneously detects a wide variety of types of mtDNA damage, including strand breaks, apurinic/apyrimidinic sites, modified purines and pyridines and DNA repair intermediates. Future studies to identify the specific types of mtDNA damage in mutant *LRRK2* neural cells will help identify the critical DNA repair pathways involved and may suggest additional therapeutic targets.

In conclusion, our data demonstrate that mtDNA damage is induced in neural cells by PD-associated mutations in *LRRK2*, and this phenotype can be functionally reversed or prevented by ZFN-mediated genome editing in iPSCs. These results indicate that mtDNA damage is likely a critical early event in the neuronal dysfunction that leads ultimately to *LRRK2*-related PD. Moreover, the cellular phenotype described here suggests new avenues for mechanistic exploration and provides a novel platform for screening therapeutics.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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## References

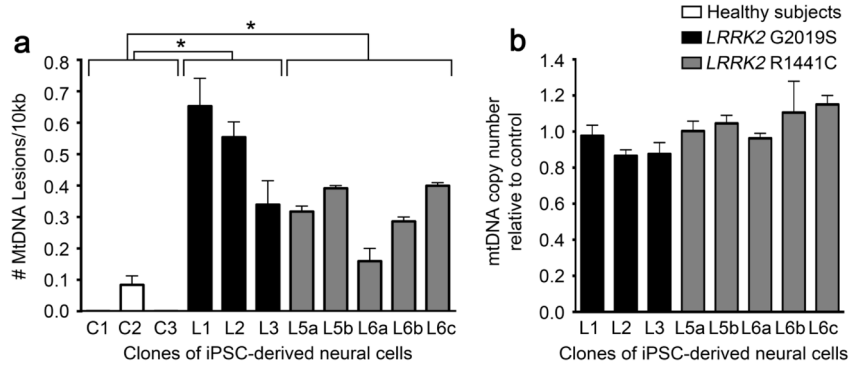
- Angeles DC, et al. Mutations in LRRK2 increase phosphorylation of peroxiredoxin 3 exacerbating oxidative stress-induced neuronal death. *Hum Mutat.* 2011; 32:1390–7. [PubMed: 21850687]
- Ayala-Torres S, et al. Analysis of gene-specific DNA damage and repair using quantitative polymerase chain reaction. *Methods.* 2000; 22:135–47. [PubMed: 11020328]
- Betarbet R, et al. Chronic systemic pesticide exposure reproduces features of Parkinson's disease. *Nat Neurosci.* 2000; 3:1301–6. [PubMed: 11100151]
- Byers B, et al. SNCA triplication Parkinson's patient's iPSC-derived DA neurons accumulate alpha-synuclein and are susceptible to oxidative stress. *PLoS One.* 2011; 6:e26159. [PubMed: 22110584]
- Chan EM, et al. Live cell imaging distinguishes bona fide human iPS cells from partially reprogrammed cells. *Nat Biotechnol.* 2009; 27:1033–7. [PubMed: 19826408]
- Cooper O, et al. Pharmacological rescue of mitochondrial deficits in iPSC-derived neural cells from patients with familial Parkinson's disease. *Sci Transl Med.* 2012; 4:141ra90.
- Guzman JN, et al. Oxidant stress evoked by pacemaking in dopaminergic neurons is attenuated by DJ-1. *Nature.* 2010; 468:696–700. [PubMed: 21068725]
- Henchcliffe C, Beal MF. Mitochondrial biology and oxidative stress in Parkinson disease pathogenesis. *Nat Clin Pract Neurol.* 2008; 4:600–9. [PubMed: 18978800]
- Heo HY, et al. LRRK2 enhances oxidative stress-induced neurotoxicity via its kinase activity. *Exp Cell Res.* 2010; 316:649–56. [PubMed: 19769964]
- Langston JW, et al. Chronic parkinsonism in humans due to a product of meperidine analog synthesis. *Science.* 1983; 219:979–980. [PubMed: 6823561]
- Lindahl T. Instability and decay of the primary structure of DNA. *Nature.* 1993; 362:709–15. [PubMed: 8469282]
- Mak SK, et al. Small molecules greatly improve conversion of human-induced pluripotent stem cells to the neuronal lineage. *Stem Cells Int.* 2012; 2012:140427. [PubMed: 22567022]
- Ng CH, et al. Parkin protects against LRRK2 G2019S mutant-induced dopaminergic neurodegeneration in *Drosophila*. *J Neurosci.* 2009; 29:11257–62. [PubMed: 19741132]
- Nguyen HN, et al. LRRK2 mutant iPSC-derived DA neurons demonstrate increased susceptibility to oxidative stress. *Cell Stem Cell.* 2011; 8:267–80. [PubMed: 21362567]
- Paisan-Ruiz C, et al. Cloning of the gene containing mutations that cause PARK8-linked Parkinson's disease. *Neuron.* 2004; 44:595–600. [PubMed: 15541308]
- Rao KS. Genomic damage and its repair in young and aging brain. *Mol Neurobiol.* 1993; 7:23–48. [PubMed: 8318166]
- Sanders LH, Timothy Greenamyre J. Oxidative damage to macromolecules in human Parkinson disease and the rotenone model. *Free Radic Biol Med.* 2013; 62:111–20. [PubMed: 23328732]
- Santos JH, et al. Quantitative PCR-based measurement of nuclear and mitochondrial DNA damage and repair in mammalian cells. *Methods Mol Biol.* 2006; 314:183–99. [PubMed: 16673882]
- Satake W, et al. Genome-wide association study identifies common variants at four loci as genetic risk factors for Parkinson's disease. *Nat Genet.* 2009; 41:1303–7. [PubMed: 19915576]
- Sherer TB, Greenamyre JT. Oxidative damage in Parkinson's disease. *Antioxid Redox Signal.* 2005; 7:627–9. [PubMed: 15890006]
- Simon-Sanchez J, et al. Genome-wide association study reveals genetic risk underlying Parkinson's disease. *Nat Genet.* 2009; 41:1308–12. [PubMed: 19915575]

- Yakes FM, Van Houten B. Mitochondrial DNA damage is more extensive and persists longer than nuclear DNA damage in human cells following oxidative stress. *Proc Natl Acad Sci U S A.* 1997; 94:514–9. [PubMed: 9012815]
- Zimprich A, et al. Mutations in LRRK2 cause autosomal-dominant parkinsonism with pleomorphic pathology. *Neuron.* 2004; 44:601–7. [PubMed: 15541309]

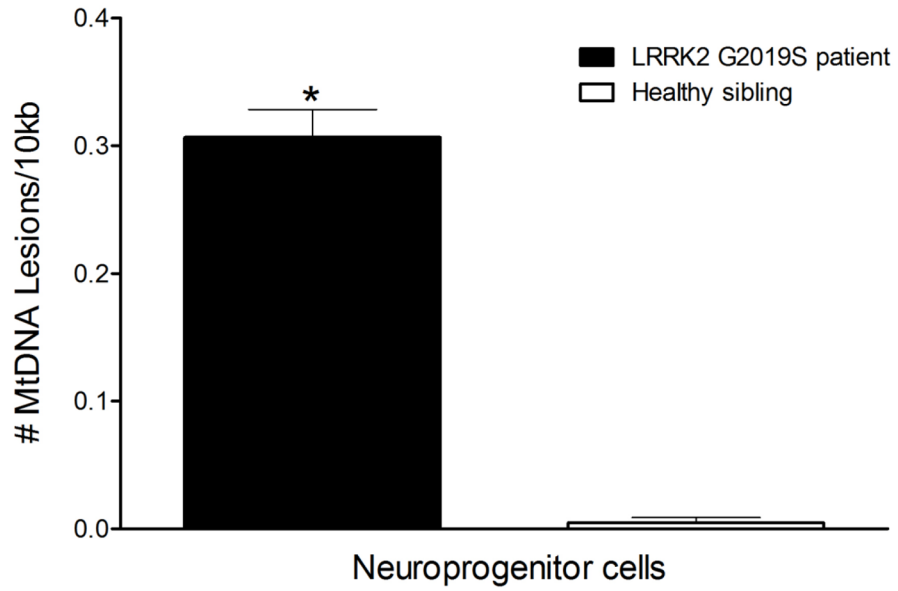


**Highlights**

1. LRRK2 PD-associated mutations induce mtDNA damage in iPSC-derived neural cells
2. MtDNA damage is functionally reversed by ZFN-mediated genome editing
3. MtDNA damage is an early biomarker of LRRK2-related PD neuronal dysfunction

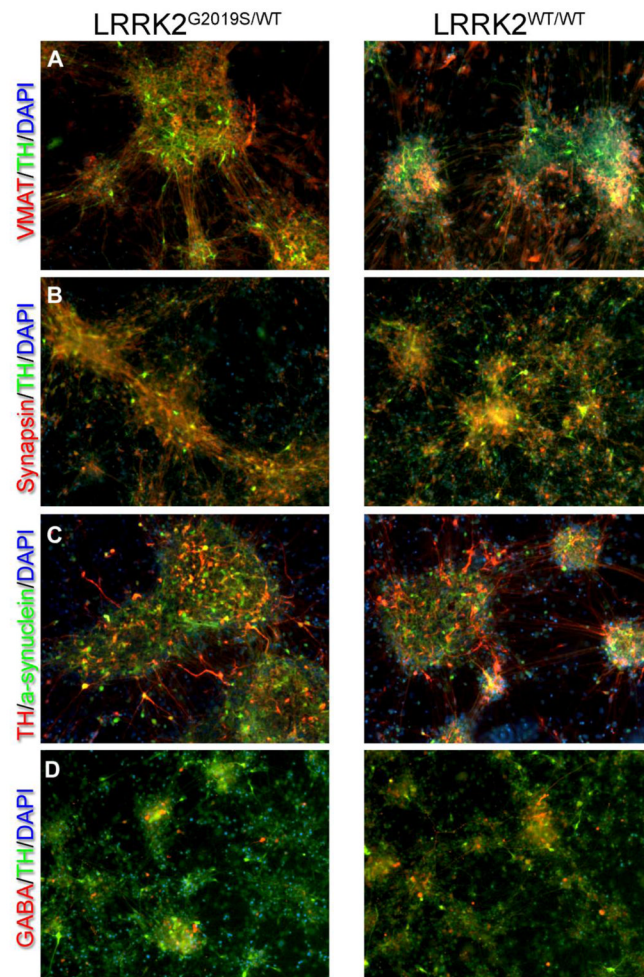


**Fig. 1.** *LRRK2* neural cells exhibit greater levels of mtDNA damage than neural cells from healthy subjects. **(A)** Neural cells were differentiated from iPSCs derived from *LRRK2* mutation carriers with the G2019S (black bars) and R1441C (gray bars) mutations and from healthy subjects (white bars). Mitochondrial DNA lesions were increased in neural cells from individual iPSC clones carrying *LRRK2* mutations (clones L1–3 and L5–6, Supplemental Table 1) relative to neural cells from healthy subjects’ iPSCs (clones C1–3, Supplemental Table 1). **(B)** In parallel, neural cells from individuals carrying *LRRK2* mutations (black and grey bars) contained a similar number of mtDNA copies as neural cells from healthy subjects. QPCR-based assay was performed in triplicate with three experimental replicates. Data are presented as mean ± SEM.

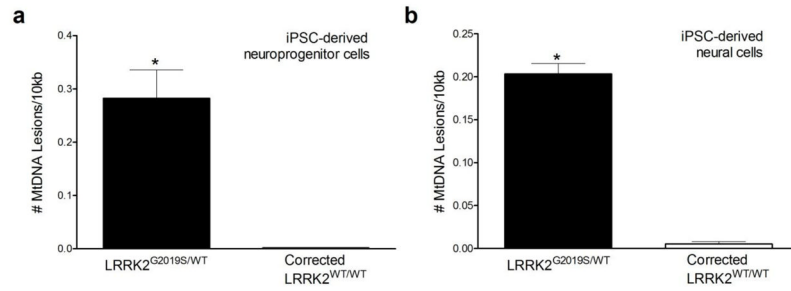


**Fig. 2.**

Neuroprogenitor cells carrying the *LRRK2* G2019S mutation show more mtDNA damage than neuroprogenitor cells from iPSCs of a healthy sibling without the *LRRK2* G2019S mutation. (A) Neuroprogenitor cells were differentiated from iPSCs carrying the *LRRK2* G2019S mutation (black bar) and from a healthy sibling (white bar). mtDNA lesions were increased in neuroprogenitor cells carrying the *LRRK2* G2019S mutation (clone L4a, Supplemental Table 1) relative to mtDNA from healthy sibling (clone C3, Supplemental Table 1). QPCR-based assay was performed in triplicate with three experimental replicates. Data are presented as mean  $\pm$  SEM, \* $p = 0.0002$ .



**Fig. 3.** (A–D) Representative images of immunocytochemistry show that differentiated neural cells derived from both mutant  $LRRK2^{G2019S/WT}$  and gene-corrected  $LRRK2^{WT/WT}$  iPSCs show equivalent expression of general neuronal markers (synapsin and alpha-synuclein) as well as robust expression of dopaminergic markers (TH & VMAT2). While GABA-containing neurons were also seen, astroglia (GFAP) were not (data not shown). Also note that our previous work provided compelling electrophysiological evidence that our protocol produces a true neuronal phenotype (Cooper et al., 2012).



**Fig. 4.** Genomic repair of the *LRRK2* G2019S mutation reduced mtDNA damage in neuroprogenitor cells and neural cells. NPCs (A) and neural cells (B) differentiated from iPSCs that retained *LRRK2* G2019S mutation after ZFN transfection (clone L4e<sup>Unmod</sup>, black bar, Supplemental Table 1) exhibited greater levels of mtDNA damage than cells differentiated from ZFN-corrected iPSCs (clone L4b<sup>WT/WT</sup>, white bar, Supplemental Table 1, \* $p < 0.002$ ). QPCR-based assay was performed in triplicate with three experimental replicates. Data are presented as mean  $\pm$  SEM.