Plasticity-related gene 3 (LPPR1) and age at diagnosis of Parkinson disease

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Abstract

Objective

To identify modifiers of age at diagnosis of Parkinson disease (PD).

Methods

Genome-wide association study (GWAS) included 1,950 individuals with PD from the NeuroGenetics Research Consortium (NGRC) study. Replication was conducted in the Parkinson’s, Genes and Environment study, including 209 prevalent (PAGEP) and 517 incident (PAGEI) PD cases. Cox regression was used to test association with age at diagnosis. Individuals without neurologic disease were used to rule out confounding. Gene-level analysis and functional annotation were conducted using Functional Mapping and Annotation of GWAS platform (FUMA).

Results

The GWAS revealed 2 linked but seemingly independent association signals that mapped to LPPR1 on chromosome 9. LPPR1 was significant in gene-based analysis (p = 1E-8). The top signal (rs17763929, hazard ratio [HR] = 1.88, p = 5E-8) replicated in PAGEP (HR = 1.87, p = 0.01) but not in PAGEI. The second signal (rs73656147) was robust with no evidence of heterogeneity (HR = 1.95, p = 3E-6 in NGRC; HR = 2.14, p = 1E-3 in PAGEP + PAGEI and HR = 2.00, p = 9E-9 in meta-analysis of NGRC + PAGEP + PAGEI). The associations were with age at diagnosis, not confounded by age in patients or in the general population. The PD-associated regions included variants with Combined Annotation Dependent Depletion (CADD) scores = 10–19 (top 1%–10% most deleterious mutations in the genome), a missense with predicted destabilizing effect on LPPR1, an expression quantitative trait locus (eQTL) for GRIN3A (false discovery rate [FDR] = 4E-4), and variants that overlap with enhancers in LPPR1 and interact with promoters of LPPR1 and 9 other brain-expressed genes (Hi-C FDR < 1E-6).

Conclusions

Through association with age at diagnosis, we uncovered LPPR1 as a modifier gene for PD. LPPR1 expression promotes neuronal regeneration after injury in animal models. Present data provide a strong foundation for mechanistic studies to test LPPR1 as a driver of response to damage and a therapeutic target for enhancing neuroregeneration and slowing disease progression.
The underlying neurodegenerative process that causes Parkinson disease (PD) begins decades before the disease is diagnosed. The current view is that following an initial insult (e.g., toxicity, trauma, or genetic), the disease starts with an asymptomatic phase of unknown duration, followed by development of prodromal nonmotor symptoms such as constipation, anosmia, and sleep disorders. Years later, cardinal motor signs appear, at which point a diagnosis of PD is made. Age at onset of motor signs, and therefore the age at diagnosis of PD, is highly variable, ranging from teen ages to the 10th decade of life. The reason for this variation is unknown, and understanding it will likely shed light on factors that affect the rate of disease progression.

There is substantial evidence that genetic factors play a major role in age at onset of motor signs and age at diagnosis of PD. Genome-wide studies have identified numerous loci that associate with the risk of developing PD, but the risk factors do not explain the variation in age at onset. Three loci have been nominated as modifiers of age at onset in familial PD. The present study was aimed at identifying genetic modifiers for common idiopathic PD. We hypothesized that identification of the genetic basis to interindividual variability in age at diagnosis will provide insights into the intrinsic mechanisms that determine the rate of deterioration during preclinical disease.

Methods
This study was a case-control GWAS, followed by replication and functional annotation.

Standard protocol approvals, registrations, and patient consents
The study was approved by the institutional review boards at all participating institutions. Written informed consent was obtained from all patients and controls for participation in the study.

Participants
The study included 2 data sets. The NeuroGenetics Research Consortium (NGRC) data set used for the discovery GWAS, gene-based test, and functional annotations. The Parkinson’s, Genes and Environment (PAGE) study was used for replication. Participants’ characteristics are shown in table 1 and figure e-1 (links.lww.com/NXG/A66).

NGRC is a case-control study of genetically unrelated participants, including 2000 PD cases and 1986 controls. Patients were enrolled sequentially from movement disorder clinics in Portland (OR), Seattle (WA), Albany (NY), and Atlanta (GA). Controls were spouses of patients or community volunteers, self-reported as being free of neurologic disease. The eligibility criterion for cases was diagnosis of PD by a movement disorder specialist according to the UK Brain Bank criteria. The eligibility criteria for controls were no neurologic disease and genetically unrelated to patients. Age was defined as age at study entry. Age at diagnosis was extracted from medical records or ascertained by self-report. Age at onset of the first motor sign was obtained using a self-administered questionnaire. Age at onset and age at diagnosis were highly correlated in the NGRC (r² = 0.91, p < 2E-16). All participants were whites of European descent.

PAGE is a cross-sectional study nested in the longitudinal NIH-American Association of Retired Persons Diet and Health Study. Participants were enrolled in 1995–1997 (irrespective of PD) via a food frequency questionnaire mailing, and in the 2004–2006 follow-up visit were asked if they had been diagnosed with a major chronic disease including PD. Participants who had been diagnosed with PD before enrollment (before 1998) were designated as prevalent PD (PAGEp, N = 209), participants who were diagnosed during follow-up (1998–2006) were designated as incident PD (PAGEi, N = 517), and participants who did not have PD were designated as controls (N = 1,549). All participants in this study were non-Hispanic whites.

Genotyping
NGRC participants were genotyped on Illumina HumanOmn1-Quad v1-0 B array and Immunochip array. Genotypes and samples were filtered by call rate, minor allele frequency (MAF) < 0.01, Hardy-Weinberg, and cryptic relatedness, as described before. Imputation was performed using IMPUTE v2.3.0, with the 1000G Phase3 integrated variant set (October 2014) as reference. Imputed single nucleotide polymorphisms (SNPs) with info score < 0.9 or MAF < 0.01 were excluded. A total of 8.5 million SNPs (900,000 genotyped and 7.6 million imputed) were used in the analysis.

PAGE participants were genotyped for rs73656147 (block 1) and rs17763929 (block 2). SNPs were chosen based on statistical significance and availability of predesigned validated TaqMan assay from Thermo Fisher (rs73656147 assay number = C__97534229_10; rs17763929 assay number = C__34297681_10).

Population structure
Principal component (PC) analysis is used to infer population-specific genetic differences, which arise from ancestry differences.
in allele frequencies and can obscure genetic association studies if not accounted for. NGRC PC analysis was conducted using a pruned subset of 100K SNPs from the GWAS as previously described.15 The top 3 PCs (effect sizes PC1 = 0.2%, PC2 = 0.06%, and PC3 = 0.06%) were included in the GWAS and adjusted for in all downstream analyses involving the NGRC. The PAGE data sets used for replication did not have ancestry informative markers (AIMs); however, a subset of the participants (396 of 726 PD cases) was previously genotyped with the Immunochip array. We conducted PC analysis using a pruned subset of 100K SNPs from the GWAS as previously described.16 The PGESW data set without PC adjustment and using the subset that had ancestry informative markers (AIMs) and adjusting for PC1-3. PAGEI and PAGEP were classified as prevalent PD (PAGEP), participants who were diagnosed with PD during follow-up (between 1998 and 2006) were classified as incident PD (PAGEI), and participants who did not have PD were designated as controls. Because PAGE participants were of similar age at entry, the method of classifying the participants into prevalent vs incident cases inevitably assigned earlier ages at diagnosis to the prevalent group and later diagnoses to the incident group.

### Statistics

**Discovery**

GWAS was conducted using PD cases only (1,950 NGRC participants with known age at diagnosis). Association between 8.5M SNPs and age at diagnosis was tested using Cox regression in ProbABEL v0.5.0,17 specifying an additive genetic model, treating age at diagnosis as a quantitative trait, and adjusting for PC1-3. The statistical outcome of Cox regression was hazard ratios (HRs) and corresponding p values. Statistical significance was set at p < 5E-8. Manhattan plots and quantile-quantile (QQ) plots were generated using FUMA v1.3.0.20 Genomic inflation factor (λ) was calculated using the estlambdfunction in GenABEL v1.8 in R.21 LocusZoom22 was used to visualize the chr9:103,865,000–104,055,000 region (GWAS peak). Haploview v4.223 was used to generate linkage disequilibrium (LD) plots of D’ and r² for SNPs in the chr9:103,865,000–104,055,000 region with GWAS p < 1E-4. LD between 2 SNPs was calculated using 1000G Phase3 v5 in LDlink.24 Linear regression was used to estimate and test differences in mean age at diagnosis (β). Conditional analysis was performed using coxph function in the survival v2.41 R package. Moving average plots (MAPs) were generated using the freqMAP v0.2 R package.25

Gene-based analysis was conducted using summary statistics from the GWAS and LD from the 1000G Phase3 EUR to map the GWAS SNPs to 18,985 protein-coding genes (hg19 build) and to calculate gene-based p values, using MAGMA v1.06,26 as implemented in FUMA v1.3.0.20 Statistical significance was set at Bonferroni-corrected p < 2.6E-6 (0.05/18,985).

### Replication

Cox regression (coxph function in the survival v2.41 R package) was used to replicate the association of 2 SNPs with age at diagnosis. We used the same model as the NGRC (additive genetic model, treating age at diagnosis as a quantitative trait). Because of the availability of PCs only in a subset of PAGE, analyses were conducted twice: using the full PAGE data set without PC adjustment and using the subset that had AIMs and adjusting for PC1-3. PAGEI and PAGEP were treated separately and were combined using meta-analysis after testing for heterogeneity. If p of heterogeneity was <0.1, the fixed-effect model was used. Meta-analysis was performed using the metagen function in the meta v4.8 R package.

### Table 1 Data sets and participants’ characteristics

|                      | Discovery (NGRC) | Replication (PAGE) |
|----------------------|------------------|--------------------|
|                      | PD               | Controls           | PAGEp   | PAGEi   | Controls |
| N                    | 2,000            | 1,986              | 209     | 517     | 1,549    |
| Male/Female          | 1,346/654        | 769/1,217          | 164/45  | 396/121 | 1,213/336|
| Age at enrollment mean ± SD | 67.3 ± 10.7       | 70.3 ± 14.1        | 62.6 ± 4.9 | 63.2 ± 4.9 | 63.4 ± 4.9 |
| Age at follow-up mean ± SD | NR              | NR                 | 73.9 ± 4.9 | 74.5 ± 4.9 | 74.0 ± 4.9 |
| N with age at onset data | 1,999            | NR                 | 0       | 0       | NR       |
| Age at onset mean ± SD | 58.3 ± 11.9      | NR                 | NA      | NA      | NR       |
| N with age at diagnosis data | 1,950            | NR                 | 209     | 517     | NR       |
| Age at diagnosis range | 25–90            | NR                 | 42–72   | 53–81   | NR       |
| Age at diagnosis mean ± SD | 60.4 ± 11.4      | NR                 | 59.9 ± 6.6 | 69.4 ± 5.4 | NR       |

**Abbreviations:** NA = not available; NGRC = NeuroGenetics Research Consortium; NR = not relevant; PAGE = Parkinson’s Genes and Environment.

Participants were non–Hispanic whites and genetically unrelated. Data on the NGRC participants were collected at enrollment: patients already had the diagnosis of PD and controls were free of neurologic disease. NGRC participants were enrolled at 4 sites: Oregon, Washington, New York, and Georgia. Age at onset mean ± SD were as follows: Oregon = 56.5 ± 12.8, Washington = 58.7 ± 11.8, New York = 59.4 ± 11.5, and Georgia = 58.7 ± 11.1. Age at diagnosis mean ± SD were as follows: Oregon = 59.6 ± 11.7, Washington = 60.7 ± 11.6, New York = 60.9 ± 11.1, and Georgia = 60.3 ± 10.6. PAGE participants were originally enrolled in the longitudinal NIH-AARP diet study in 1995–1997. Their PD status was investigated in 2004–2006. Participants who had the diagnosis of PD before 1998 were classified as prevalent PD (PAGEP), participants who were diagnosed with PD during follow-up (between 1998 and 2006) were classified as incident PD (PAGEI), and participants who did not have PD were designated as controls. Because PAGE participants were of similar age at entry, the method of classifying the participants into prevalent vs incident cases inevitably assigned earlier ages at diagnosis to the prevalent group and later diagnoses to the incident group.
Functional annotation

Functional annotation was conducted in FUMA v1.3.0, using SNPs with GWAS p < 1E-6 and all variants in r² ≥ 0.6 with them, and included CADD analysis, eQTL mapping, 3D chromatin interaction mapping (Hi-C), annotation of enhancers, tissue-specific expression of genes identified via Hi-C and eQTL mapping, and their age-specific expression in the brain (BrainSpan.org). The false discovery rate (FDR) was used to correct for multiple testing. STRUM was used to predict the effect of a missense on the structural stability of a protein.

Data availability

NGRC genotype and phenotype data are available at dbGaP ncbi.nlm.nih.gov/gap accession number phs000196.v3.p1.

Results

GWAS

In SNP-based GWAS, the most significant signal for association, at p = 5E-8, mapped to LPPR1 on chromosome 9q31.1 (figure 1, A and B). In the gene-based test, LPPR1 achieved p = 1E-8, surpassing the genome-wide statistical significance threshold of p < 2.6E-6 (figure 1, C and D). The p values were not inflated (λ = 1.007 SNP based, λ = 1.04 gene based). Analysis of LD in the region revealed 2 haplotype blocks with seemingly independent signals for association (figure 1, E and F). There was strong LD among SNPs in each block, but weak LD between the blocks (r² < 0.2) because of a recombination hot spot between them (figure 1F). The 2 blocks were in a ~200 Kb region inside LPPR1. Block 1 consisted of 51 SNPs with MAF > 0.01, which yielded HR = 2.02–1.88, with p = 9E-7 to 2E-5 for association with age at diagnosis. Block 2 consisted of 39 SNPs with MAF = 0.02, which yielded HR = 1.88–1.85, with p = 5E-8 to 7E-7. We chose 1 SNP to represent each block for replication: rs73656147 for block 1 (MAF = 0.01, HR = 1.95, p = 5E-8) and rs17763929 for block 2 (MAF = 0.02, HR = 1.88, p = 5E-8), both in Hardy-Weinberg (p > 0.3), with little correlation between them (r² = 0.2). Conditional analysis conducted to determine whether the 2 blocks were tagging the same or different disease-associated variants was inconclusive because although the signals were weakened when adjusted for each other, neither was abolished when conditioned on the other (table e-1, links.lww.com/NXG/A70). The 8 categories of stratification were family history, sex, cigarette smoking, caffeine intake, nonsteroidal anti-inflammatory drugs use, recruitment site, Jewish heritage, and the European country of ancestral origin. The association signal for rs73656147 (block 1) was robust across all strata. rs17763929 (block 2) showed evidence of heterogeneity as a function of recruitment site and the European country of ancestral origin. Given these results, we tested the association of the 2 SNPs with PCs. rs17763929 was associated with PC1 (p = 7E-6) and PC3 (p = 8E-3), and rs73656147 was not (p > 0.05 for PC1-3), indicating the presence of population structure in block 2 but not in block 1.

Replication

In comparison to NGRC, which had a 65-year range for age at diagnosis, the PAGE data sets had a narrower range of less than 30 years. Because PAGE participants were of similar age at study entry, the method of classifying the participants into prevalent PD (diagnosis before entry) vs incident PD (diagnosis after entry) inevitably assigned earlier ages at diagnosis to the prevalent group (PAGEP) and later diagnoses to the incident group (PAGEI). Mean age at diagnosis in PAGEP was 59.9 ± 6.6 years, which was similar to the NGRC (60.4 ± 11.4). PAGEI participants were on average 10 years older at diagnosis (69.4 ± 5.4, range 53–81 years). Given the disparity in the range and mean ages at diagnosis, we analyzed PAGEp and PAGEI separately.

Association of rs73656147 (block 1) with age at diagnosis replicated robustly (table 3). There was no evidence of heterogeneity between PAGEp and PAGEI in the association of rs73656147 with age at diagnosis, although the signal was stronger in PAGEp than in PAGEI, which is not surprising, given that the former is enriched in cases with earlier age at diagnosis. To gauge robustness of the associa
Figure 1 Results of genome-wide association study for age at diagnosis of PD

Genome-wide association was tested between 8.5 million SNPs and age at diagnosis in 1,950 PD cases from the NGRC, using the Cox hazard ratio regression method and adjusting for principal components (PC1–3). (A) Manhattan plot of SNP-based GWAS. Tallest peak, at \( p = 5 \times 10^{-8} \), was on chromosome 9q31.1. (B) QQ plot of SNP-based GWAS. The observed \( p \) values were not inflated (\( \lambda = 1.007 \)). (C) Manhattan plot of gene-based GWAS. \( LPPR1 \) was at \( p = 1 \times 10^{-8} \). Statistical significance threshold was \( p < 2.6 \times 10^{-6} \), which is Bonferroni corrected for the 18,985 protein-coding genes tested. (D) QQ plot of gene-based GWAS. The observed \( p \) values were not inflated (\( \lambda = 1.04 \)). (E) \( r^2 \) (top panel) and \( D' \) (bottom panel). Linkage disequilibrium (LD) across the SNPs that gave \( p < 1 \times 10^{-4} \) for association with age at diagnosis reveals 2 blocks represented by rs73656147 (left triangle) and rs17763929 (right triangle). (F) Magnified map of the associated region (chr9:103,865,000–104,055,000), showing that PD-associated SNPs map to \( LPPR1 \) and form 2 haplotype blocks separated by recombination hot spots (blue spikes). (G) Chromatin state of \( LPPR1 \) (Roadmap 111 Epigenomes), showing that active enhancers (yellow), transcription start site (red), and transcripts (green) of \( LPPR1 \) are seen only in stem cells and the brain and that the GWAS SNPs align with regulatory elements. ESC = embryonic stem cell; iPSC = induced pluripotent stem cell; TssA = active transcription start site (TSS); TssAFlnk = flanking active TSS; TxFlnk = transcription at gene 5' and 3'; Tx = strong transcription; TxWk = weak transcription; EnhG = genic enhancers; Enh = enhancers; ZNF/Rpts = zinc-finger genes and repeats; Het = heterochromatin; TssBiv = bivalent/poised TSS; BivFlnk = flanking bivalent TSS/ enhancer; EnhBiv = bivalent enhancer; ReprPC = repressed polycomb; ReprPCWk = weak repressed polycomb; Quies = quiescent.
Table 2 Association of LPPR1 variants with age and age at diagnosis is driven by age at diagnosis

| Block 1 | Block 2 |
|---------|---------|
| rs73656147 | rs17763929 |
| **N** | 1,950 | 1,950 |
| **Cox HR** | 1.95 | 1.95 |
| **p Value** | 3E-6 | 3E-6 |
| **LR β [95% CI]** | -6.00 [-9.18 to -2.83] | -5.98 [-9.16 to -2.81] |

Abbreviations: CI = confidence interval; HR = hazard ratio; LR = linear regression; β = effect size on age at diagnosis (in years) per copy of minor allele. The associations were tested in the NGRC data set using Cox regression, and the effect sizes were estimated using linear regression (LR). HR is the age-for-age increase in the odds of event per copy of the minor allele, as estimated using Cox regression. β is the difference in years in age at diagnosis between carriers of 1 minor allele vs no minor allele, as estimated using linear regression. Age at diagnosis was the primary outcome of the study. Minor alleles of rs73656147 and rs17763929 were associated with higher HR and younger age at diagnosis (Ia). The association was not influenced by sex (Ib), which was expected because, unlike PD risk, which is significantly associated with sex (OR = 3.26, p < 2E-16), age at diagnosis is not associated with sex (HR = 0.99, p = 0.83). Minor alleles were also associated with younger ages in cases (II), but not in controls (III). Because age and age at diagnosis were correlated (r² = 0.74, p < 2E-16), an association with one will show as an association with both. In conditional analysis, the association with age at diagnosis persisted when adjusted for age (IV), but the association with age was abolished when adjusted for age at diagnosis (V), suggesting that age at diagnosis was the driving force and association with age was a by-product of the correlation.

Association of rs17763929 (block 2) with age at diagnosis showed significant heterogeneity between PAGE and NGRC for the association of rs73656147 with age at diagnosis. Meta-analysis yielded HR = 2.14, p = 1E-3 for replication and HR = 2.00, p = 9E-9 for replication and discovery. Mean difference in age at diagnosis per copy of rs73656147 minor allele was -6.0 (95% confidence interval: -9.18 to -2.83) years in the NGRC, -5.53 (-9.72 to -1.34) in PAGEP, -0.84 (-4.22 to 2.55) in PAGEH, and -4.08 (-7.45 to -0.70) in the meta-analysis of the 3 data sets.

Association of rs17763929 (block 2) with age at diagnosis showed significant heterogeneity between PAGE1 and PAGEP (table 3), as it had within the NGRC (table e-2, links.lww.com/NXG/A70). The association with rs17763929 replicated in PAGEP but not in PAGE1. There was significant heterogeneity between PAGE1 and NGRC, but not between PAGEP and NGRC. Meta-analysis of PAGEP and NGRC yielded HR = 1.88, p = 4E-9 for full PAGE data and HR = 1.95, p = 3E-9 for the PAGE subsample adjusted for PC1-3. Including PAGE1 with PAGEP and NGRC in a random-effects meta-analysis diluted the effect size to HR = 1.53, p = 0.04. Mean difference in age at diagnosis per copy of rs17763929 minor allele was -5.65 (-8.20 to -3.11) years in the NGRC, -3.62 (-7.23 to -0.02) in PAGEP, and 0.62 (-1.34 to 2.58) in PAGE1.

Functional annotation

Hi-C analysis showed significant (FDR < 1E-6) chromatin interaction between the PD-associated LPPR1 SNPs and promoters of LPPR1 and several genes on chromosome 9 (figure 2, A). Some of the SNPs that were significant in Hi-C mapped to enhancers in the brain (table 4 and figure 1, G). Eleven of the genes identified through Hi-C are expressed in the brain: LPPR1, SEC61B, MSANTD3-TMEFF1, TMEFF1, GALNT12, MURC, GRIN3A, NR4A3, ALG2, MRPL50, and ZNF189 (figure 2, B and C). The expression of LPPR1 in the brain is the strongest in early prenatal stage and decreases with developmental stage and increasing age (figure 2, C).

CADD analysis, a scoring system for deleteriousness of genetic variants, identified 5 SNPs in block 1 and 3 in block 2, with CADD = 10–19 (table 4), which places them among the top 10% (CADD > 10) to 1% (CADD > 20) of most deleterious mutations in the genome.27 rs41296085 (CADD = 18, in block 1) is a missense (p.Ser12Ala) in exon 2, predicted to structurally destabilize the LPPR1 protein (ΔΔG = -1.2). The remainder of the variants with high CADD scores are in introns. eQTL analysis revealed an association between rs117451395 (block 1) with expression levels of GRIN3A (FDR = 4E-4).

Discussion

There has been intense research on PD risk factors, which so far has resulted in identification of numerous causative genes, 40 susceptibility loci, several environmental factors, and a few genes that interact with the environmental factors to increase or reduce the risk of developing PD. In contrast, we know little about factors that affect the rate of disease progression. In this study, we attempted to identify genetic modifiers of age at diagnosis, a reflection of rate of progression, using an
unbiased genome-wide approach, followed by independent replication, and functional annotation.

We uncovered evidence for association of genetic variants in neuronal plasticity-related gene 3 (LPPR1) with age at diagnosis of PD. Two signals of association were detected, each representing a haplotype block of SNPs. The variants that were associated with earlier age at diagnosis had low allele frequencies (MAF = 0.01–0.02), as were the variants that were previously found for age at onset of familial PD. The low allele frequencies may be one reason why modifier genes have been more difficult to detect than common variants that associate with risk.

Table 3 Replication

| Data sets            | N PD cases | Age at diagnosis | Block 1 rs73656147 | Block 2 rs17763929 |
|----------------------|------------|------------------|---------------------|--------------------|
|                      |            | Mean ± SD        | HR      | p Value | HR      | p Value |
| NGRC (discovery)     | 1,950      | 60.4 ± 11.4      | 1.95     | 3E-6    | 1.88    | 5E-8    |
| PAGEp (replication)  | 209        | 59.9 ± 6.6       | 2.88     | 7E-4    | 1.87    | 0.01    |
| PAGEp with PC1-3     | 113        | 59.9 ± 6.8       | 2.17     | 0.05    | 3.03    | 4E-3    |
| PAGEp (replication)  | 517        | 69.4 ± 5.4       | 1.62     | 0.07    | 1.04    | 0.41    |
| PAGEp with PC1-3     | 283        | 69.2 ± 5.3       | 1.48     | 0.16    | 1.03    | 0.45    |
| Meta-analysis A      | Heterogeneity rs73656147 | Heterogeneity rs17763929 |
| PAGEp and PAGEi      | ns         | 0.08             | 2.14     | 1E-3    | 1.34    | 0.31    |
| NGRC and PAGEp       | ns         | ns               | 2.08     | 2E-8    | 1.88    | 4E-9    |
| NGRC and PAGEi       | ns         | 0.01             | 1.90     | 9E-7    | 1.42    | 0.23    |
| NGRC and PAGEp and PAGEi | ns         | 0.02             | 2.00     | 9E-9    | 1.53    | 0.04    |
| Meta-analysis B      |            |                  |          |         |         |         |
| PAGEp and PAGEi      | ns         | 0.02             | 1.73     | 0.07    | 1.67    | 0.34    |
| NGRC and PAGEp       | ns         | ns               | 1.97     | 6E-7    | 1.95    | 3E-9    |
| NGRC and PAGEi       | ns         | 0.02             | 1.89     | 2E-6    | 1.43    | 0.23    |
| NGRC and PAGEp & PAGEi | ns         | 0.02             | 1.91     | 5E-7    | 1.68    | 0.05    |

Abbreviations: HR = hazard ratio; NGRC = NeuroGenetics Research Consortium; ns = not statistically significant; PAGE = Parkinson’s, Genes and Environment; PC = principal component; PD = Parkinson disease.

Two SNPs with signals for association with age at diagnosis of PD in the NGRC data set (discovery) were genotyped and tested for association with age at diagnosis of PD in the PAGE data set (replication). PAGE participants were designated as PAGEp if they were diagnosed before study entry or PAGEi if they were diagnosed during the study. Cox regression was used to test the association of SNP (additive model) with age at diagnosis (quantitative trait) and to calculate hazard ratios (HRs) and the corresponding significance (p). NGRC was adjusted for PC1-3 in GWAS and meta-analyses. Only a subset of PAGE had ancestry informative markers (AIMs) for which PC could be calculated; thus, results are shown for the full PAGE data set without PC adjustment and for the PAGE subsample with PC adjustment. p values are 2-sided for NGRC and one-sided for PAGE because of the directionality of the hypothesis being replicated. Meta-analysis A: NGRC (PC1-3 adjusted) and PAGE (all data without PC adjustment), Meta-analysis B: NGRC (PC1-3 adjusted) and PAGE (subset of data adjusted for PC1-3). rs73656147 replicated robustly with no evidence of heterogeneity across data sets. rs17763929 replicated in PAGEp and showed significant heterogeneity between PAGEi and PAGEp or NGRC.

Meta-analysis was conducted using the fixed-effects model if there was no evidence for heterogeneity (p ≥ 0.1) and the random effects model if there was heterogeneity (p < 0.1).

unbiased genome-wide approach, followed by independent replication, and functional annotation.

We uncovered evidence for association of genetic variants in neuronal plasticity-related gene 3 (LPPR1) with age at diagnosis of PD. Two signals of association were detected, each representing a haplotype block of SNPs. The variants that were associated with earlier age at diagnosis had low allele frequencies (MAF = 0.01–0.02), as were the variants that were previously found for age at onset of familial PD. The low allele frequencies may be one reason why modifier genes have been more difficult to detect than common variants that associate with risk.

The association with block 1 replicated robustly in both PAGEp and PAGEi. Block 2 signal replicated in PAGEp but not in PAGEi. Block 2 has a complex LD structure, with evidence of population substructure, which limits generalizability of results. Failure to capture a signal for block 2 in PAGEi may be because we had genotype on only 1 SNP in block 2 for PAGE, which did not fully capture the complexity of block 2. PAGEi participants being significantly older than NGRC and PAGEp participants may also be a factor. LPPR1 promotes neuroregeneration, but its expression diminishes with age to nearly undetectable level by age 40 years (figure 2C). One can speculate that some detrimental variants may not have an effect after a certain age when the gene is no longer expressed.

Functional annotation of the PD-associated variants in LPPR1 revealed the presence of several variants with predicted deleterious effects, including a missense that destabilizes the structure of LPPR1, a regulatory element that associates with expression levels of GRIN3A, and enhancers that interact with promoters of LPPR1 and several other genes in the brain. Some of the candidate genes that were identified via interaction with LPPR1 play key roles in pathways that are implicated in PD, including GRIN3A (which encodes a subunit of NMDA receptor involved in the glutamate-regulated ion channels in the brain), SEC61B (protein transport apparatus of the endoplasmic reticulum membrane), MURC (Rho...
kinase signaling), and MRPL50 (mitochondrial ribosomal protein).

LPPR1 is one of the 5 members of a brain-specific gene family that modulates neuronal plasticity during development, aging, and after brain injury. LPPR1 is the strongest driver of axonal outgrowth in the gene family. Studies in mice have shown that after neuronal injury, overexpression of LPPR1 enhances axonal growth, improves motor behavior, and promotes functional recovery. Extrapolating to our findings, we propose that LPPR1 is involved, not necessarily in the cause of PD, rather in response to damage, and influences the efficacy of regeneration and the subsequent rate of deterioration in preclinical PD. The actual cause of injury and neuronal death is not stipulated in this hypothesis; it could be head trauma, environmental toxins or genetic, but once the initial damage is incurred, it is the efficacy of intrinsic mechanisms of repair that determine the rate of disease progression. Present findings provide a strong foundation for mechanistic studies to investigate the role of LPPR1 in PD and
Table 4 Functionally significant variants

| Block | GWAS SNP | position:alleles | GWAS p | $r^2$ | eQTL | CADD | Hi-C/EnhBrain |
|-------|----------|------------------|--------|------|------|------|---------------|
| 1     | rs77351585 | 9:103874925:C:T  | 2E-06  | —    | —    | 18   | Hi-C/EnhBrain |
| 1     | rs73495940 | 9:103875807:G:C  | 9E-07  | Lead | —    | —    | Hi-C |
| 1     | rs150164200| 9:103875896:A:C  | 2E-06  | 1    | —    | 10.4 | —            |
| 1     | rs117583993$^a$ | 9:103876647:G:A | 3E-06  | 1    | —    | —    | Hi-C/EnhBrain |
| 1     | rs148874623 | 9:103939117:A:C  | 9E-06  | 1    | —    | 12.1 | —            |
| 1     | rs117451395 | 9:103941039:C:T  | 1E-05  | 1    | GRIN3A | —    | Hi-C |
| 1     | rs41296085 | 9:103947810:T:G  | 2E-05  | 1    | —    | 18 (missense)$^b$ | Hi-C |
| 1     | rs117900237 | 9:103959240:G:A  | 2E-05  | 1    | —    | 10.5 | Hi-C |
| 2     | rs17763929 | 9:103984900:A:G  | 5E-08  | Lead | —    | —    | Hi-C |
| 2     | rs61188842 | 9:103988006:C:T  | 8E-05  | 0.6  | —    | —    | Hi-C/EnhBrain |
| 2     | rs117058418 | 9:104011717:T:C  | 2E-07  | 1    | —    | 10.4 | Hi-C |
| 2     | rs117314512 | 9:104014244:G:A  | 2E-07  | 1    | —    | 12.4 | Hi-C |
| 2     | rs149155028 | 9:104032402:TTC:T | 1E-05  | 0.7  | —    | 18.6 | Hi-C |

Functional annotation was conducted on SNPs with GWAS $p < 1E-6$ and SNPs that were in high LD with them ($r^2 > 0.6$). Variants are shown if they are the lead SNPs (most significant) for the block, or an eQTL (FDR = 4E-4), or had a CADD score >10, or had both significant evidence for 3D chromatin interaction (Hi-C, FDR < 1E-6) and overlapped with an enhancer in the brain. Block 1 is a single block of SNPs in high LD. Block 2 has a complex LD structure with at least 3 subhaplotypes (figure e1-C, links.lww.com/NXG/A66). Variants are shown with their rs accession number, chromosome position and the 2 alleles (major: minor), GWAS $p$ value for association with age at diagnosis of PD, and their correlation ($r^2$) with the lead SNP of the block. eQTL: an SNP that is associated with gene expression, in this case, $\Delta \Delta G= −1.2$, which predicts a destabilizing effect on the protein structure of LPPR1.

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**Author contributions**

Z.D. Wallen: statistical analysis, review, and critique of the manuscript. H. Chen: creation of the PAGE data set, acquisition of data, review, and critique of the manuscript. E.M. Hill-Burns: assembly and QC of phenotype and genotype data, imputation, review, and critique of the manuscript. S.A. Factor and C.P. Zabetian: creation of the NGRC data set, review, and critique of the manuscript. H. Payami: study concept, design and execution, creation of the NGRC data set, wrote the manuscript, and obtained funding.

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