Single-cell genomic profiling of human dopamine neurons identifies a population that selectively degenerates in Parkinson’s disease

Tushar Kamath1,2,4, Abdulraouf Abdulraouf1,4, S. J. Burris1, Jonah Langlieb1, Vahid Gazestani1, Naeem M. Nadaf1, Karol Balderrama1, Charles Vanderburg1 and Evan Z. Macosko1,3✉

The loss of dopamine (DA) neurons within the substantia nigra pars compacta (SNpc) is a defining pathological hallmark of Parkinson’s disease (PD). Nevertheless, the molecular features associated with DA neuron vulnerability have not yet been fully identified. Here, we developed a protocol to enrich and transcriptionally profile DA neurons from patients with PD and matched controls, sampling a total of 387,483 nuclei, including 22,048 DA neuron profiles. We identified ten populations and spatially localized each within the SNpc using Slide-seq. A single subtype, marked by the expression of the gene AGTR1 and spatially confined to the ventral tier of SNpc, was highly susceptible to loss in PD and showed the strongest upregulation of targets of TP53 and NR2F2, nominating molecular processes associated with degeneration. This same vulnerable population was specifically enriched for the heritable risk associated with PD, highlighting the importance of cell-intrinsic processes in determining the differential vulnerability of DA neurons to PD-associated degeneration.

RESULTS

A molecular taxonomy of human SNpc DA neurons. To address the sampling challenge associated with DA neuron profiling, we developed a protocol based on fluorescence-activated nuclei sorting (FANS), to enrich midbrain DA neuron nuclei for use in snRNA-seq (Fig. 1a). In a scRNA-seq dataset of mouse midbrain22 (Extended Data Fig. 1a), we identified the gene Nr4a2 encoding a transcription factor (TF) as specific to mammalian midbrain DA neurons (area under the curve (AUC) = 0.76; Methods). To comprehensively profile all cell types in the SNpc, we isolated nuclei from eight neurotypical donors (Supplementary Table 1) and performed snRNA-seq on both NR4A2-positive and -negative nuclei (Extended Data Fig. 1b). In total, we generated 184,673 high-quality profiles (median number of unique molecular identifiers (UMIs) per individual, 8,810; median number of genes per individual, 3,590 (Extended Data Fig. 1c,d); median number of UMIs per cell, 8,086; median number of genes per cell, 3,462 (Extended Data Fig. 1e,f)). 43.6% of which were from the NR4A2-positive cytometry gate (Methods and Fig. 1b). We performed clustering analysis of each donor separately to assign profiles to one of seven main cell classes, including DA neurons (Methods and Extended Data Fig. 1g–j). The NR4A2-sorted profiles were 70-fold enriched for DA neurons (Fig. 1c), defined by a cluster with joint expression of TH, SLC6A3 and SLC18A2 (Extended Data Fig. 1i,j), genes whose products are essential for DA neurotransmission23.

To identify subtypes of DA neurons, we performed LIGER27 on 5,684 high-quality DA neuron nuclei, a 180-fold increase in absolute numbers over existing datasets of human midbrain DA neurons (Extended Data Fig. 2a). Based on our LIGER-derived, low-dimensional embedding (Extended Data Fig. 2b), we identified molecular mechanisms play important roles in the selective vulnerability of some DA neuron populations to PD degeneration.

1Broad Institute of Harvard and MIT, Stanley Center for Psychiatric Research, Cambridge, MA, USA. 2Harvard Graduate Program in Biophysics, Harvard University, Cambridge, MA, USA. 3Massachusetts General Hospital, Department of Psychiatry, Boston, MA, USA. 4These authors contributed equally: Tushar Kamath, Abdulraouf Abdulraouf. ✉e-mail: emacosko@broadinstitute.org

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9. Vahid Gazestani, 1, Evan Z. Macosko, 1, Karol Balderrama, 1, Charles Vanderburg, 1, and Tushar Kamath, 1,2,4

10. The loss of dopamine (DA) neurons within the substantia nigra pars compacta (SNpc) is a defining pathological hallmark of Parkinson’s disease (PD). Nevertheless, the molecular features associated with DA neuron vulnerability have not yet been fully identified. Here, we developed a protocol to enrich and transcriptionally profile DA neurons from patients with PD and matched controls, sampling a total of 387,483 nuclei, including 22,048 DA neuron profiles. We identified ten populations and spatially localized each within the SNpc using Slide-seq. A single subtype, marked by the expression of the gene AGTR1 and spatially confined to the ventral tier of SNpc, was highly susceptible to loss in PD and showed the strongest upregulation of targets of TP53 and NR2F2, nominating molecular processes associated with degeneration. This same vulnerable population was specifically enriched for the heritable risk associated with PD, highlighting the importance of cell-intrinsic processes in determining the differential vulnerability of DA neurons to PD-associated degeneration.

11. Recent advances in single-cell RNA-sequencing technology have begun to reveal cell-type-specific changes in several brain diseases. These enrichment results suggest that cell-intrinsic molecular mechanisms play important roles in the selective vulnerability of some DA neuron populations to PD degeneration.
Fig. 1 | A molecular census of DA neurons in the human substantia nigra. a, NR4A2 antibody-based enrichment strategy and snRNA-seq profiling workflow. b, UMAP representation of 104,097 NR4A2+ (left) and 80,576 NR4A2− (middle) nuclei from eight neurotypical donors. Profiles colored green are from clusters identified as DA neurons. c, Bar plot of proportions of DA neurons per replicate for NR4A2+ versus NR4A2− libraries (median fold enrichment, 70; n = 21 NR4A2+ and n = 16 NR4A2− libraries). d, UMAP representation of 15,684 DA neuron nuclei, colored by cell type. e, Dot plot showing expression of selected marker genes across DA clusters. f, UMAP representation of 104,097 DA neuron nuclei, colored by cell type. 

Ten transcriptionally distinct subpopulations (Methods, Fig. 1d,e and Extended Data Fig. 2c) with strong alignment across all donors (Methods; alignment score, 0.98; Extended Data Fig. 2d). Four DA clusters preferentially expressed SOX6 while the other six expressed CALB1, recapitulating a well-defined developmental axis of variation within midbrain DA neurons (Fig. 1e)24. All of our subtypes showed strong expression of genes that are essential for DA neurotransmission (Extended Data Fig. 2e)25. Previously defined dopaminergic markers20 also showed strong expression across our dataset, with mixed expression of subtype markers previously found that differentiate different rodent populations (Extended Data Fig. 2f). Further, the proportions of these broad subtypes generally matched stereotactic estimates of CALB1+ cells from previous immunohistochemical stains of postmortem human nigra21 (Extended Data Fig. 2f), suggesting no intrinsic bias in our tissue sampling method. The broad CALB1−/SOX6+ axis of variation was consistent across donors (Extended Data Fig. 2g) and clusters were consistently represented across integrative tools (Extended Data Fig. 2h–j and Methods), suggesting that our strategy consistently uncovered and preserved underlying biological states.
Our method for high-throughput profiling of DA subtypes enabled us to make robust comparisons amongst paras compacta DA neurons across species. Although recent profiling studies of primates have shown strong evolutionary conservation of cell types in mice, a few primate-specific specializations have been reported\(^{25,26}\). To investigate the evolutionary conservation of our human DA neuron subtypes, we sampled and integrated, with our human datasets, DA neuron profiles from four additional species spanning three phylogenetic orders: Primate, Scandentii and Rodentia (Fig. 1f). Cross-species analysis (Methods) identified eight clusters (Fig. 1g) that consistently followed the SOX6-CALB1 axis of variation (Extended Data Fig. 3a–c). While some profiles exhibited divergence in cellular proportions, integrative analysis largely maintained the cluster distinctions found in the human-only analysis (Extended Data Fig. 3d). Examining the contributions of each species to each cluster, we found that the DA8 population, composed primarily of the CALB1_GEM human population (Extended Data Fig. 3d), included profiles derived only from humans and macaque (Fig. 1g,h), and expressed a number of highly specific marker genes not found in any one other DA subtype in our integrative analysis (Extended Data Fig. 3e) and absent from a previous analysis of rodent DA subtypes (Extended Data Fig. 3f\(^{21,32}\)). Using in situ hybridization across the mouse midbrain (Methods), we found no murine DA neurons expressing the marker genes Fam83b and Gem for our DA8 population (Extended Data Fig. 3g–i), further confirming the lack of a cognate population in rodents.

**Regulatory element identification in dopaminergic neurons.** The identification of regulatory elements that drive the molecular identity of DA neurons can inform differentiation protocols for in vitro studies of DA neurons in PD, as well as drive the refinement of cellular replacement therapies for this disease. To understand the regulatory networks that may drive such transcriptional variation, we used single-cell regulatory network inference and clustering (SCENIC) to identify 84 regulons that were highly specific (P<0.05 and AUC>0.7, Wilcoxon rank-sum test; Methods) to the ten DA subtypes defined in our dataset. The top TFs ranked by AUC per DA subtype contained many TFs previously implicated in specifying DA identity, including those encoded by the genes SOX6, OTX2, SMAD1, PBX1, LMX1B, NEF2L1 and EN2 (Fig. 1i and Extended Data Fig. 4a)\(^{28}\). Even within the more homogenous SOX6 axis we identified several TFs with differential activity across subtypes, including the TF encoded by SMAD1, as well as some not previously implicated in DA neuron differentiation such as those encoded by CUX2 and ZNF91. The same SCENIC analysis on DA neuron data generated from the macaque midbrain (Extended Data Fig. 4b) identified highly overlapping sets of regulons, further corroborating the identification of selectively active TFs in each DA subtype.

**Localization of DA neurons in macaque midbrain by Slide-seq.** Dopaminergic neurons in the SNpc comprise the A9 group of catecholaminergic neurons and are divided into dorsal and ventral tiers, with the ventral tier showing greater vulnerability to PD-associated degeneration\(^{25,26-30}\). To spatially localize the ten DA populations defined in our human analysis (Fig. 1d), we performed Slide-seq\(^{31,32}\) on 27 arrays generated from nine coronal sections spanning the caudal 80% of the rostrocaudal axis of a *Macaca fascicularis* SNpc (Methods, Fig. 2a and Extended Data Fig. 5a,b). The A9 group was easily identifiable by visualizing the expression of DA neuron markers TH and SLC6A3 (Extended Data Fig. 5c). Using a recently described tool for decomposition of cell type mixtures in spatial transcriptomics data\(^{33}\) (Methods), we localized our ten DA subtypes to 3,206 Slide-seq-defined bead locations in the A9 group (Fig. 2d, Extended Data Fig. 5d and Methods) and positioned each relative to the demarcation between dorsal and ventral tiers (Fig. 2e, dotted line), defined from the expression of CALB1 and ALDHA1 (Extended Data Fig. 5c–e)\(^{34}\). The SOX6-AGTR1 population was the most strongly enriched in the ventral tier (Fig. 2d,e and Extended Data Fig. 5f–h; mean SOX6-AGTR1 relative distance, +490 μm), consistent with previous marker analyses performed on laser-capture microdissection subsets of A9 DA neurons\(^{34}\), while the CALB1_GEM and CALB1_TRHR populations were strongly enriched in the dorsal tier (Fig. 2d,e and Extended Data Fig. 5f–h; mean CALB1_GEM relative distance, −478 μm; mean CALB1_TRHR relative distance, −288 μm). Single-molecule fluorescence in situ hybridization (smFISH) (Extended Data Fig. 6a and Methods) of the human midbrain confirmed localization of the SOX6-AGTR1 subtype to the ventral tier (Fig. 2f,g and Extended Data Fig. 6b) and types CALB1_GEM and CALB1_TRHR to the dorsal tier (Fig. 2f,g and Extended Data Fig. 6c,d). Additionally, in situ analyses of two other SOX6 populations confirmed their lack of selective localization to either tier (Extended Data Fig. 6e–h), consistent with the Slide-seq results (Fig. 2e).

**Differentially vulnerable DA neurons in PD.** The strong ventral localization of the SOX6-AGTR1 population suggested that it may be especially susceptible to PD-associated degeneration. To identify cell-type-specific molecular alterations in PD, we profiled an additional 202,810 high-quality nuclei (Extended Data Fig. 7a–d); median number of UMIs per individual, 7,177; median number of genes per individual, 3,108; median number of UMIs and genes per cell, 6,939 and 3,076, respectively), including 6,364 DA neurons, from ten age-matched and postmortem-interval-matched (Extended Data Fig. 7e–g) individuals with documented pathological midbrain DA neuron loss, and a clinical diagnosis of either PD or LBD (Supplementary Table 2). Between PD/LBD and neurotypical control tissues there were no significant differences in tissue collection date (Extended Data Fig. 7h), and this covariate had no significant effect on the integrity of capture (P>0.05; Methods and Extended Data Fig. 7i). Finally, integrative analysis of these donors with our neurotypical controls (Methods) identified 68 transcriptionally defined subpopulations (Extended Data Fig. 7j–o) from our seven major cell classes, with minimal batch-dependent variation (alignment scores: 0.61–1.0; median alignment score across cell types, 0.76; Extended Data Fig. 7p).

We assessed the differential abundance between PD/LBD and aged control samples for both major cell classes and all molecularly defined subtypes. Among major cell classes, DA neurons showed the largest significant decline (P<0.05, Wilcoxon rank-sum test) as a fraction of cells per individual (Extended Data Fig. 8a). Among the 68 molecularly defined subpopulations, 11 showed significant proportional changes in association with PD/LBD (Fig. 3a; false discovery rate (FDR)-adjusted P<0.05). Among these, one proportionally increased population was a subset of microglia expressing GPNM (Extended Data Fig. 8b), which has been identified as a marker of disease-associated microglia in transcriptomic studies of Alzheimer's disease (AD)\(^{35,36}\) and was robust up to one-sixth of the original dataset size (Extended Data Fig. 8c). We also identified an increase in the VIM_LHX2 astrocyte subtype (Extended Data Fig. 8d), similarly robust to large losses in power due to sample size (Extended Data Fig. 8e). The VIM_LHX2 population expresses reactive markers, namely VIM and LHX2 (Extended Data Fig. 8d), suggesting that this population may play a role in responding to degenerative changes in PD/LBD SNpc.

The largest statistically significant decline was in the SOX6_AGTR1 DA population, while clusters CALB1_GEM and CALB1_TRHR were proportionally increased (Fig. 3b). These proportional changes were robust to differences in absolute numbers of DA neurons sampled per cluster (Supplementary Fig. 1a), clinical diagnosis (Supplementary Fig. 1b) and library quality (Supplementary Fig. 1c,d). We further developed a metric to visualize disease-associated enrichment...
In the first experiment we probed for cells expressing TH (Supplementary Table 3, Supplementary Fig. 2a–d and Methods). of PD (with concurrent documented midbrain DA neuron loss) divided among neurotypical donors and individuals who had died counted a total of 5,339 individual DA neurons (Supplementary of double- and triple-positive cells in each smFISH experiment and populations (Fig. 1e). We assayed the proportional representations enriched in the CALB1_TRHR, CALB1_GEM and CALB_RBP4, TH and formed on an additional 20 postmortem-frozen midbrains equally ers, we quantified DA subtype proportions using smFISH, per -alyzed cell profiles, identifying a gradient of susceptibility (Fig. 3c and or depletion within the low-dimensional embedding of jointly ana-lyzed cell profiles, identifying a gradient of susceptibility (Fig. 3c and Methods) that correlated with the expression of AGTR1 and ORs from mixed-effects association of single cells (MASC) (Fig. 3c).

Our flow cytometry procedure to isolate DA nuclei relies on Nissl staining of a 10-μm M. fascicularis midbrain slice adjacent to Slide-seq-assayed tissue. Circles indicate approximate location of the placement of the three Slide-seq arrays shown in b–d. RN, red nucleus; CP, cerebral peduncles; SNpcd, substantia nigra pars compacta dorsal; SNpcv, substantia nigra pars compacta ventral. Cartesian arrows indicate orientation of bead arrays in b–d; scale bar, 1 mm. Nissl staining was repeated nine times across macaque brain. b–d. Bead arrays colored by RCTD cell type definitions (Methods) corresponding to major cell type (b), CALB1+ or SOX6+ subtypes (c) and the three most spatially variable DA subtypes (d). e, Ridge plot for aggregated densities of CALB1 and SOX6 subtypes (top) and all ten DA subtypes (bottom) across 27 bead arrays (Methods, also includes definition of midline). f, Tiled image of one 10-μm human midbrain slice. White dotted line indicates the approximate A9 region; scale bar, 1 mm. Experiment was repeated once. g, Scatter plots showing relative location of triple- (yellow) and single-positive cells (Methods) from in situ hybridization of markers: CALB1+/GEM+ (left), CALB1+/TRHR+ (middle) and SOX6+/AGTR1+ DA neurons (right); scale bars, 1 mm. Experiment was repeated five times for SOX6+/AGTR1+ localization and once for CALB1+/GEM+ and CALB1+/TRHR+.

or depletion within the low-dimensional embedding of jointly ana-lyzed cell profiles, identifying a gradient of susceptibility (Fig. 3c and Methods) that correlated with the expression of AGTR1 and ORs from mixed-effects association of single cells (MASC) (Fig. 3c).

Our flow cytometry procedure to isolate DA nuclei relies on protein expression of NR4A2, which is downregulated in DA neurons in PD97. To address these and other potential confound-ers, we quantified DA subtype proportions using smFISH, per-formed on an additional 20 postmortem-frozen midbrains equally divided among neurotypical donors and individuals who had died of PD (with concurrent documented midbrain DA neuron loss) (Supplementary Table 3, Supplementary Fig. 2a–d and Methods). In the first experiment we probed for cells expressing TH, SOX6 and AGTR1 (Fig. 3d); in a second experiment, we identified cells in situ expressing TH, CALB1 and TMEM200A (Fig. 3d), a marker enriched in the CALB1_TRHR, CALB1_GEM and CALB_RBP4 populations (Fig. 1e). We assayed the proportional representations of double- and triple-positive cells in each smFISH experiment and counted a total of 5,339 individual DA neurons (Supplementary Fig. 2e) across 40 full SNpc sections. We confirmed that CALB1+/ TMEM200A+ DA cells were selectively enriched in PD (Fig. 3e; Wilcoxon rank-sum test P < 0.05), as were CALB1+/TMEM200A− DA cells, although the log fold change difference was lower (Fig. 3e; log2(fold change 1.32) for the TMEM200A− group versus log2(fold change 1.13) for the TMEM200A+ group). We also confirmed selective depletion of the SOX6_AGTR1 sub-population (Fig. 3e; Wilcoxon rank-sum test P < 0.05, log2(fold change −2.1); Supplementary Table 5).

Cellular localization of PD common variants. We next sought to better understand the origins of the selective vulnerability of SOX6_AGTR1 cells to neurodegeneration in PD. One possibility is that the human genetic risk for PD—which is established at birth—selectively acts within the vulnerable population. To test this, we examined the enrichment of expression of genes harboring either familial and common variants associated with PD. A total of 26 genes have been identified that harbor mutations, as ascertained by family studies, that confer substantial risk for PD98. We tested
the overlap between these genes and markers specifically expressed in eight major cell classes of the SNpc (Fig. 4a and Methods). The DA neuron gene set was the only one to show significant enrichment (Bonferroni-corrected P < 0.05, Fisher’s exact test; Methods) of genes that contained these familial variants (Fig. 4b), suggesting that a substantial fraction of these genes act within DA neurons to influence neurodegeneration.

We next examined the enrichment of common variant risk of sporadic PD within markers of our SNpc cell classes (Methods)9,40, as well as markers of cell classes defined by the additional profiling of 46,872 nuclei from postmortem dorsal striatum tissue of four neuropathologically normal individuals (Extended Data Fig. 9a–g and Supplementary Table 4). We observed strong, significant enrichment (Bonferroni-corrected P < 0.05) of heritable risk for sporadic PD (Methods) in markers of DA neurons (Fig. 4c, Extended Data Fig. 10a), also consistent with previous analyses18,39. Next, we tested the enrichment of PD common variant risk across our 68 transcriptionally defined subpopulations in human SNpc (Methods and Supplementary Table 7). Using two different statistical methods34,43, we found the largest and only statistically significant (Bonferroni-corrected P < 0.05; Methods and Supplementary Table 9) enrichment of PD genetic risk genes within the SOX6_AGTR1 cell subtype (Fig. 4d and Extended Data Fig. 10b). Relative enrichment by both methods was uninfluenced by the number of nuclei sampled per DA subtype (Extended Data Fig. 10c and Supplementary Table 7), and ordering of the significance of these enrichments was consistent across variations in gene set size (Extended Data Fig. 10d).

To further characterize the genetic loci enriched in SOX6_AGTR1 cells, we identified genes that were assigned high MAGMA z-scores and low P values for DA subtype differential expression, including the SOX6_AGTR1 population (Fig. 4e and Extended Data Fig. 10e). Some prioritized—and well-studied—genes were
expressed in multiple SOX6+ and CALB1+ subtypes, including SNC-A, MAPT and GAK. Other SOX6_AGTR1-specific genes included the GWA-prioritized genes WNT3 and IG5SF9B. Gene Ontology (GO) enrichment analysis (Methods) of SOX6+-specific genes identified relevant ontologies that included regulation of neuron death and Wnt signaling, reinforcing that the genetic loci identified as being expressed in this subtype probably act intrinsically to influence neurodegeneration (Fig. 4f).

Nomination of gene programs altered in dopaminergic neurons.

Finally, we sought to nominate gene regulatory programs significantly and specifically altered in the SOX6_AGTR1 population in our PD/LBD tissue donors. Using gene set enrichment analysis (GSEA) of known TF target sets (Methods), we identified a total of 13TFs whose targets were significantly (FDR significant P < 0.05) enriched in the SOX6_AGTR1 population but not in the other DA subtypes (Fig. 4g,h and Supplementary Fig. 3a). This analysis revealed a depletion of targets of the TF encoded by the gene LMX1A (Methods; normalized enrichment score <0), a TF involved in the developmental specification of midbrain DA neurons. Transcription factors whose targets were enriched in the SOX6_AGTR1 population included TP53, whose own transcript was also upregulated specifically in the SOX6_AGTR1 population (Supplementary Fig. 3b), as well as NR2F2; both TFs modulate the progression of midbrain DA neuron loss in mouse models of PD.

Discussion

In this study we generated a molecular taxonomy of human SNpc DA neurons, spatially localized them within the SNpc and identified one DA subpopulation, SOX6_AGTR1, that is highly susceptible to neurodegeneration in PD. Importantly, across both the substantia nigra and caudate, this population was the most strongly enriched for expression of genes associated with PD by previous GWAS, suggesting that the genetic risk of PD acts preferentially in the most vulnerable neurons to influence their survival. Transcriptional changes within SOX_AGTR1 cells in patients with PD implicated several canonical cell stress pathways—including those regulated by TFs encoded by the genes TP53 and NR2F2—as important to the process of PD-associated neuronal death.

Our snRNA-seq analysis of SNpc DA neurons provides a comprehensive taxonomy of these critically important cells in humans. Our map will help guide bulk transcriptomic studies of PD in localization of disease-associated signals to specific human DA subtypes. Further, DA subtype definitions will allow the refinement of in vitro DA neuron differentiation protocols, which could prove useful in genetic screens of neuronal susceptibility and the testing of candidate therapeutic molecules. Interestingly, although nine of our ten populations showed homology to rodent DA populations, one cluster of cells, CALB1_GEM, was found only in our snRNA-seq data from macaque and human and not from mouse, rat or tree shrew. We localized CALB1_GEM cells exclusively to the dorsal tier.
The partitioning of heritable disease risk preferentially to the most vulnerable DA population provides evidence that the genetic influences of PD-associated degeneration are preferentially cell intrinsic. This result—which is consistent with previous efforts to partition heritable risk of PD24,26—contrasts with similar analyses performed on late-onset AD genetic risk that particularly implicate microglia and other populations of myeloid origin in nonautonomous neuroimmune mechanisms. Thus, despite overlapping pathologies and the clinical phenomenon of these two diseases, there exist substantial differences in how genetic risk may manifest in producing the disease, an insight that could prove useful in the advancement of specific therapeutic opportunities and biomarkers.

The heritability enrichment within PD-vulnerable neurons themselves also provides a crucial opportunity to address a long-standing question in PD about the primacy of midbrain DA neuron death relative to neurodegeneration in other regions. For example, one hypothesis of PD pathogenesis posits that PD pathology ascends through the brain (for example, via alpha-synuclein aggregates) from outside the central nervous system, through medullary brainstem structures. The extension of profiling efforts to these other structures—and comparison of heritability enrichment in vulnerable cell types—could help to clarify whether these other, vulnerable structures are degenerating due to primary influences of the disease or are secondary to midbrain DA loss.

Our identification of TFs whose activity is up- or downregulated, as nominated by our enrichment of targets in our differential expression, specifically within the vulnerable SOX6_AGTRI population implicates specific cellular pathways in the process of DA neuron death in PD. The TF encoded by NR2FP2, for example, has previously been shown to promote mitochondrial dysfunction in several disease models, including those of heart failure and PD27. Upregulation of the TF encoded by TP53 provides a link to other neurodegenerative diseases, such as amyotrophic lateral sclerosis, in which TP53 has been implicated in motor neuron death28,29. Cross-disease integrative analyses may reveal conserved molecular processes that are prime candidates for therapeutic intervention.

Online content
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Animal husbandry of *Mus musculus*. Animals were group housed with a 12:12 h light/dark schedule and allowed to acclimate to their housing environment for 2 weeks post arrival. For *M. musculus* housing, ambient temperature was strictly maintained between 68 and 72 °Fahrenheit and humidity strictly maintained between 30 and 30%. All procedures involving animals at Massachusetts Institute of Technology (MIT) were conducted in accordance with the US National Institutes of Health Guide for the Care and Use of Laboratory Animals, and were approved by the MIT Committee on Animal Care. All procedures involving animals at the Broad Institute were conducted in accordance with the US National Institutes of Health Guide for the Care and Use of Laboratory Animals, under protocol no. 0120-09-16.

**Cryosectioning and brain preparation.** Postmortem human midbrain and dorsal striatum (caudate nucleus) tissue were flash-frozen at −80 °C and cryosectioned at ∼−20 °C into 60-µm sections. For midbrain preparation, pigmented regions of human midbrain were microdissected and between five and ten 60-µm sections prepared. Following microdissection, samples were placed on dry ice. Following microdissection, samples were placed on dry ice.

**Generation of single-nucleus suspensions from frozen human midbrain and caudate samples.** To each cryosectioned sample, 1 ml of extraction buffer (ExB) was added to an Eppendorf tube. Samples were triturated before being placed in a six-well plate. Samples were then triturated 20 times with ExB, every 2 min, until the entire volume was aspirated and diluted with 45 ml of wash buffer (WB) in a 50-ml Falcon tube and then split into four or five rounds of trituration and rest were performed. The entire volume of the well was then passed twice through a 26-gauge needle into the same well. The pellets were then resuspended in 250 µl of WB and mixed thoroughly by trituration. This 250-µl solution was then pooled with four other pellets from the same original midbrain and resuspended into approximately 1 ml of WB in a 1.5-ml Eppendorf tube.

**Immunolabeling and blocking of human nuclei.** Approximately 100 µl of 10% bovine serum albumin in WB (final concentration, 1%) was added to the concentrated nuclei. The antibody NRR42-A647 (Santa Cruz, no. sc-376984 AF647) was then added at a concentration of 1:300. Additionally, one caudate nucleus was stained and sorted with NeuN (see metadata at Broad SCP link in Data Availability), which was performed by the addition of an anti-NeuN–PE antibody (EMD Millipore Corp., clone A60; no. FCMA8317) at a concentration of 1:1,500. Samples were then covered in aluminum foil and incubated on a rotator at 4 °C for 1 h. Following incubation, Eppendorf tubes were spun at 150 g for 10 min in a swing-bucket benchtop centrifuge. The supernatant was then gently aspirated, then WB was added to achieve a total sample volume of 1 ml. Samples were then stained with DAPI (ThermoFisher, no. D2264) at 1:1,000 dilution and filtered with a 70-µm filter.

**FANS for enrichment of dopaminergic nuclei.** Flow-sorting parameters for DAPI gating are described in Martin et al. (https://www.protocols.io/view/frozen-five-nucleus-extraction-for-10xv3-sneeq-b62khge; see protocols.io link for all buffers and solution concentrations). All steps were performed on ice or cold blocks and all tubes, tips and plates were precooled for >20 min before starting isolation. Briefly, 60-µm sections of midbrain (∼500 µl) were placed in a single well of a 12-well plate and 2 ml of ExB was added to each well. Mechanical dissociation was performed by trituration using a P1000 pipette, pipetting 1 ml of solution slowly up and down with a 1-ml Rainin tip (no. 30389212), without creation of froth or bubbles, a total of 20 times. Tissue was allowed to rest in the buffer for 2 min and trituration was repeated. In total, four or five rounds of triturations and rest were performed. The entire volume of the well was then passed twice through a 26-gauge needle into the same well. Approximately 2 ml of solution was transferred into a 50-ml Falcon tube and filled with WB for a total of 30 ml of solution, which was then split across two 50-ml Falcon tubes (∼15 ml of solution in each tube). The tubes were then spun in a swinging-bucket centrifuge for 10 min at 600 g and 4 °C. Following spinning, the majority of supernatant was discarded (∼500 µl remaining with the pellet). Solution from two Falcon tubes were then pooled into a single tube of ∼1,000 µl of concentrated nuclear tissue solution. DAPI was then added to the solution at the manufacturer’s (Thermo Fisher Scientific, no. 62248) recommended concentration (1:1,000).

Following sorting, nuclei concentration was counted using a hemocytometer before loading into a 10X Genomics 3′ V3 Chip.

**Generation of single-nuclei suspensions from *M. fascicularis,* **Tupai a belangeri** and *Rattus norvegicus.* Fresh frozen *R. norvegicus* and *T. belangeri* brains were mounted in a cryostat. *T. belangeri* and *R. norvegicus* nuclear isolation was adapted from a previously published protocol (https://www.protocols.io/view/frozen-tissue-nuclei-extraction-for-10xv3-sneeq-b62khge). Following nuclear isolation, NRR42 staining was performed on isolated nuclei (no. sc-376984, Santa Cruz). The staining and flow-sorting parameters followed was similar to human midbrain and macaque nuclei sorting, as above. Both NRR42- and DAPI-labeled nuclei were flow sorted into one sample. Following nuclear isolation, nuclei were loaded into a 10X Genomics 3′ V3 Chip.

**Generation of Slide-seqV2 libraries from *M. fascicularis,* to generate Slide-seqV2 data.** *M. fascicularis* flash-frozen brain tissue samples were equilibrated in a cryostat at −20 °C and sectioned to the region of interest into 10-µm sections. Samples were transferred to the Slide-seqV2 array puck by placing that on top of the tissue transfer to an Eppendorf tube with 1 ml of wash buffer (50 mM sodium chloride sodium citrate (SSC)+1.20 RNase Inhibitor, Lucigen). Libraries were generated according to the published Slide-seqV2 protocol (https://www.protocols.io/view/library-generation-using-slide-seqv2-bxipkcn). Reagents and solutions, along with oligonucleotide sequences, are listed under the Materials section in protocols.io. All steps were followed except for step 10: following removing of beads from two washes with TE–W (10 mM Tris–HCl, pH 7.5, 1 mM EDTA, Ethidiumiodinametacetica acid, +0.05% Tween-20), the bead pellet was immediately resuspended in 200 µl of 0.1 N NaOH.

Libraries were sequenced based on the standard illumina protocol. Samples were pooled at a concentration of 4 nM and read structure was specified according to protocols.io. All samples were sequenced on either NovaSeq 6000 SP or S2 flowcells.

**snRNA-seq and library preparation.** For all single-nuclei experiments, the 10X Genomics (v3) kit was used according to the manufacturer’s protocol recommendations. Library preparation was performed according to the manufacturer’s recommendations. Libraries were pooled and sequenced on either a NextSeq or NovaSeq4000.

**Hybridization chain reaction on tissue sections.** Postmortem human and mouse midbrain tissues flash-frozen at −80 °C were cryosectioned at −15 to −20 °C to create 12-µm sections on SuperFrost Plus slides, which were frozen at −80 °C until staining. Slides were allowed to warm up to room temperature (RT) before being placed in 4% paraformaldehyde for 15 min at RT. Slides were then washed three times with 70% ethanol for 5 min before incubation in 70% ethanol for 2 h. After incubation, slides were then incubated at 37 °C in Probe Hybridization buffer (Molecular Instruments) for 10 min in a humidified chamber. The probe solution was then freshly prepared by the addition of 0.4 pmol of each probe set (Molecular Instruments) per 100 µl of Probe Hybridization buffer. Slides were then incubated overnight at 37 °C in a humidified chamber. After 18–24 h, sections were sequentially washed in the following solutions: (1) 75% probe wash buffer and 25% 5xSSCT (SSC+10% Tween-20), (2) 50% probe wash buffer and 50% 5xSSCT, (3) 25% probe wash buffer and 75% 5xSSCT and (4) 100% 5xSSCT. Slides were then washed for 5 min at RT in 5xSSCT and allowed to prefly by Probe Amplification buffer (Molecular Instruments) for ≥30 min at RT. Hairpins were then freshly prepared by the addition of 1 µl of hairpin per 100 µl of Amplification Buffer (Molecular Instruments), and snap-frozen using a thermocycler with the following settings: 95° for 90 s then cooling to RT (20 °C) at a rate of 3 °C min−1. Sections to hairpins were added to the desired volume of amplified hairpin buffer.

Following overnight incubation at RT in a humidified chamber, slides were washed twice for 30 min at RT with 5xSSCT. An appropriate amount of Fluoromount Gold plus NucBlue was added, and slides were then covered slip and sealed with clear nail polish and stored at 4 °C until imaging. Probe names and their associated accession numbers are listed as follows: TH (NM_001366795), CALR (NM_001366795), DDC (NM_001366795), SCN2A (NM_001366795), SLC25A9 (NM_001366795), FXN (NM_001366795), TRIM2 (NM_001366795), TRHR (NM_001366795), GABA (NM_001366795), and T (NM_001366795). The sections were imaged using a microscope with a 20x objective. Mounting and imaging of macroslide midbrain sections. Fresh macaque midbrain sections (10 µm) were equilibrated to RT and excess condensate wiped off. Sections were fixed in 70% ethanol for 2 min, followed by rehydration in ultrapure water for 30 s. Excess water was wiped off and slides were stained with Arcturus Histogen Solution (ThermoFisher, no. 12241-05) for 4 min. Excess
Slides were sequentially fixed in 70, 90 and 100% ethanol for 30 s, 10 s and 1 min, dye was tapped off and slides were rehydrated in water for 10 s for destaining. Cell type clustering and annotation of human datasets.

Other cell types using a Wilcoxon rank-sum test from the presto package (https://github.com/MacoskoLab/PuckCaller/), with default settings. (treeshrew_2.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macaque (M_fascicularis_5.0.fasta) and tree shrew (Rnor_6.0.fasta), mouse (mm10), macro...
SCENIC analysis to identify differentially regulated regulons. To identify differentially regulated regulons associated with specific dopaminergic subtypes, we employed SCENIC with user-recommended settings from the SCENIC vignette (https://github.com/gordon-lab/SCENIC_R/inst/doc/SCENIC_Running.html). Briefly, we averaged the log-normalized gene values of single-nucleus profiles from our dataset of non-PD control individuals per dopaminergic subtype. We filtered on genes that had at least one count per bulk profile across all ten subtypes. We ran a correlation analysis using GENIE3 and then ran SCENIC to determine TF modules within these correlations. Using AUCell, we scored cells based on regulon activity and plotted these scaled regulon scores on a per-dopaminergic subtype basis. To determine statistically significant differences in regulon activity, we ran a Wilcoxon rank-sum test (using the presto package https://github.com/immunogenomics/presto) on regulon scores between major dopaminergic subtypes and ranked cells based on their AUC values and used AUCell, we scored cells based on regulon activity and plotted these scaled regulon scores on a per-dopaminergic subtype basis. To identify differentially regulated regulons associated with specific dopaminergic subtypes, we used SCENIC to determine TF modules within these correlations. Using AUCell, we scored cells based on regulon activity and plotted these scaled regulon scores.

Robust cell type decomposition analysis of Slide-seqV2 data. To assign cell types from the single-nucleus reference of M. fascicularis to Slide-seqV2 pucks, we used the robust cell type decomposition (RCTD) package. RCTD allows for spatial pixels to be assigned sparse mixtures of the reference cell types, which is congruous to Slide-seqV2’s near single-cell resolution. The macaque reference annotations used are detailed below (integration of dopaminergic neurons across species). We combined high-level annotation with the more detailed subtype definition of DA subtypes, based on an integrative analysis with our reference human control dataset (Extended Data Fig. 5d; see below). The gene set used in RCTD integration was created by combining a reference-wide set of highly variable genes (first 5,000 genes under a variance-stabilizing transformation) with those selected in neuronal cell type identification. Further, Slide-seqV2 pucks were filtered and retained with at least 50 UMIs and 20 genes.

To aid RCTD in identification of beads with artificial signatures, we added a ‘cell type’ to the reference that corresponded to cells that did not pass standard quality control metrics and that expressed high levels of URR, and oxidative phosphorylation genes heavily enriched in rough ER, which may be differentially retained in nuclei dissociation preparations. Beads assigned by RCTD solely to this artifact classification were discarded from later analysis.

The mapping was performed with RCTD’s doublet-aware mode, which is designed to most accurately decompose cells captured by Slide-seqV2’s ten micron beads. RCTD classifies each bead as a mixture of reference cell types; additionally, it categorizes these classifications as ‘simply’, ‘doublet uncertain’, ‘doublet certain’ or ‘reject’, depending on the number of cell types assigned and assignment confidence. We filtered out ‘reject’ beads and used classifications from the other three categories for later analysis.

To delineate each puck’s medial–lateral midline (for plotting the distances of subtypes CALB1 and SOX6), TH and CALB1 gene expression were plotted and positioned manually on the boundary between each gene’s expression to calculate a line of best fit. Integration of dopaminergic neurons across species. To jointly define dopaminergic neurons across species, we used LIGER’s projection method. Briefly, we ran LIGER’s projection method on the reduced dimensionality of the integrated macaque midbrain data. We defined a major dopaminergic neuron across species if it was represented in our single-cell dataset. We identified DA subtypes across all nonhuman species. Gene-by-cell DGE matrices were normalized and centered. Variable gene selection was performed, followed by non-negative matrix factorization (NMF), Louvain clustering and uniform manifold approximation and projection (UMAP) projection based on low-dimensional embedding from NMF. Low-quality nuclei were defined as those with >10% of their reads mapping to mitochondrial transcripts, and putative doublets were defined by coexpression of one or more major cell type markers identifiable from a recently published study of mouse midbrain. DA neurons from all species were subsetted based on expression of clusters with high levels of TH and/or SLC18A2 for tree shrew, which did not have the gene TH annotated as such in the reference genome. We produce the multispecies integrative analysis shown in Fig. 1a, where we first integrated mouse and human DA neurons using LIGER with the following parameters: (1) number of variable genes, 1,080; (2) kernel width = 10 (number of nearest neighbors); and clusters were determined using the Louvain clustering algorithm in LIGER with a resolution of 1.5. Joint low-dimensional embedding was visualized using the UMAP algorithm.

To define a reference scRNA-seq dataset for our RCTD analysis of Slide-seq data from the tree shrew, we projected DA neurons from M. fascicularis onto our neurotypical human control dataset (Extended Data Fig. 5d). To accomplish this, we subsetted our macaque midbrain data to DA neurons only (defined by expression of TH, SLC6A3 and SLC18A2). Macaque DA neuron profiles were then projected onto the human neurotypical control reference map using the online INMF branch of LIGER, with the setting ‘projection = TRUE’. After quantile normalization, datasets were jointly clustered via a nearest-neighbor graph (knn_k = 45, number of nearest neighbors), and clusters were determined using the Louvain clustering algorithm in LIGER with a resolution of 1.2. Joint low-dimensional embedding was visualized using the UMAP algorithm. The macaque cluster with high levels of mitochondrial and oxidative phosphorylation genes was flagged for removal, but was used for RCTD mapping analysis (see above). Putative doublets (defined by the coexpression of one or more major cell type markers identifiable from a recently published study of mouse midbrain) were also removed from integrative analysis. DA subtypes were assigned to macaque nuclei based on the nearest-neighbor graph and coclustering with human DA cell types.

Differential abundance assessment of cell types in association with PD and other covariates. To identify differentially abundant cell subpopulations in association with PD, we employed MASC. MASC is a generalized, mixed-effect model that models a binomial distribution whose mean is a linear function of covariates in our model: sex, status (control or disease) and SNC (whether nuclei were captured in a NRR4A2-positive or -negative library). To jointly define differential cell abundance across major cell types and subtypes while accounting for both disease and control individuals and, for each bin, we determined the total number of cells per individual within that bin. We then averaged those values across individuals per bin for both case and control, then scaled these scores across all bins to provide a normalized estimate of the relative abundance of certain DA subtypes across the UMAP space.

Differential expression analysis. To identify differentially expressed genes across all major cell types, we employed model-based analysis of single-cell transcriptomes (MAST). MAST is a mixed-effect hurdle model that models droplet-based, single-nucleus/cell expression data as a mixture of a binomial and normal distribution (for log-normalized nonzero expression values) while systematically accounting for predefined covariates. We included the following fixed-effect covariates in our model: log(number of UMIs), sex, percentage of reads that map to mitochondrial genes per nucleus (percent.mito) and status (control or disease) to test the effect of the disease on each cell type. We additionally included the library from which the nuclei were sampled as a random-effect covariate in the model, to account for intracellular correlation of expression data. To evaluate the effect of disease on expression, we performed differential expression analysis across all major cell types. For all non-neuronal cell types we used only cells from NRR4A2-negative libraries; for neurons, we used only NRR4A2-positive libraries. Genes were defined as differentially expressed if they reached statistical significance in at least 10% of samples and FDR-adjusted P < 0.05. We ran MAGMA on the set annotation setting to test the significance of this result across major cell types and subtypes while accounting for batch variation. We included all the same covariates as above (except for disease status), and included additional covariates of cell type identity and a covariate for whether the cell derived from a NRR4A2-positive or -negative library. To create marker gene sets, we selected genes with coefficient >0 and FDR-adjusted P < 0.05.

Heritability enrichment analyses. To determine which cell types and subtypes are enriched for heritable risk of traits, we used MAGMA40 (https://ctg.cncr.nl/software/magma) and stratified LD score regression (s-LDSC). For AD and PD, we downloaded publicly available summary statistics from the most recently available studies. We then performed a single-nucleotide polymorphism to gene set enrichment analysis using the equivalent webserver (FUMA: https://fuma.ctglab.nl/) using default parameter settings. For all major cell types/subtypes, we took the top 3,500 marker genes ranked by z-score as determined from MAST (see above) for each cell type and subtype as a defined gene set. For 17 subtypes we were unable to reach sufficient nuclei to acquire 3,500 marker genes and, instead, used the maximum number of genes identified that reached statistical significance (Supplementary Table 10). Importantly, despite setting upper bounds on the maximum number of differentially expressed genes for these smaller cell types, there was no correlation with the number of nuclei sampled and P values from MAGMA for our DA subtypes (Extended Data Fig. 10c). We ran MAGMA on the set annotation setting to test the significance of whether each marker gene set was enriched for a given risk of the trait. The resulting P values were Bonferroni corrected for multiple hypothesis testing across major cell types and subtypes. To run a stratified LD score for partitioning of heritability across cell type gene sets, we followed the standard procedure defined by the s-LDSC wiki (https://github.com/bulk/lidsc/wiki/Cell-type-specific-analyses). Briefly, we first munged

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all summary statistics with the munge_sumstats.py script provided with the s-LDSC package. We next defined a reference gene set based on all genes expressed in our substantia nigra dataset. We specified baseline weights based on the model defined for cell-type-specific analyses. Finally, we ran partitioned heritability on the gene sets across all cell types and subtypes. Resulting P-values were Bonferroni corrected for multiple hypothesis testing across major cell types and subtypes.

To create the ‘pseudo-Manhattan’ plots in Fig. 4c, we ordered the nominated MAGMA genes by their z-score and the −log(P-value) from our MAST analysis. GO signatures (Fig. 4f) for top genes that drive the association of the SOX6 signature with the common variants of PD were determined by taking all genes higher than MAST z > 2.4568. We used enrichR on these genes to determine enriched GO biological processes using the GO_Biological_Process_2017 library.

Familial PD variant enrichment analysis. To test whether there exists any enrichment of genes previously nominated as containing variants that cause familial PD, we gathered a list of known familial PD genes from a recent whole-exome sequencing study. We ran Fisher’s exact test using the geneOverlap package between genes that were considered specifically expressed in a major cell type, as determined by the top 10% of genes ordered by the AUC metric with the Wilcoxon rank-sum test, and these 26 genes. The resulting P-values were then corrected for multiple hypothesis testing using the Bonferroni method (eight tests for major cell types).

GSEA of TF gene sets across dopamine subtypes. To identify TFs and their downstream target genes as enriched, we used GSEA from the GSEA package. The choice of GSEA on the list of differentially expressed genes over a SCENIC-based approach was motivated by recent studies suggesting the use of random-effect models (as we have carried out via MAST) to control for false discovery in single-cell expression data with multiple replicates. We first gathered all TFs from three TF regulon libraries in the enrichR database: TRRUST, ENCODE and ARCHS4. We then ran GSEA with the following parameters—minsize = 1, maxsize = 500, nperm = 1,000—to test the enrichment of a TF targets list in an ordered list of differentially expressed genes per subtype. All P-values were FDR corrected for hypothesis testing across the ten subtypes tested.

Regression analyses. To determine the lines of best fit for Extended Data Fig. 10c (number of nuclei sampled versus heritability enrichment significance), we performed an ordinary least-squares regression on −log(P-transformed P-values against the number of nuclei sampled per cluster for P-values generated by both MAGMA and s-LDSC. A Wald test was performed to assess whether the absolute value of the coefficient of the slope of the line was significantly > 0. To determine any bias associated with case-control status on the median number of UMIs per library, we used a linear mixed-effect model and included covariates of case-control status as a fixed effect and the library as a random effect (Extended Data Fig. 7h). To determine the influence of year of collection on the quality of sampling, we regressed the median number of UMIs acquired per library against the year of collection for each tissue sample (Extended Data Fig. 7i).
Extended Data Fig. 1 | See next page for caption.
Extended Data Fig. 1 | High-throughput snRNA-seq profiling of SNpc. a, Expression of tyrosine hydroxylase (Th) and Nr4a2 in a published scRNA-seq study of mouse midbrain(21). b, Representative FANS plot for enriching midbrain dopaminergic neurons. APC-A (x-axis) represents NR4A2 antibody channel and Brilliant Violet (y-axis) represents DAPI channel. The NR4A2 gate was thresholded to select the top 2.5–4% of all nuclei (red box). c,d, Number of genes (C) and UMIs (unique molecular indicators) (D) per nucleus grouped by donor (n = 11,577 nuclei for subject 3298; 49,759 nuclei from 3322; 29,230 nuclei for 3345; 23,747 nuclei for subject 3346; 24,856 nuclei for subject 3482; 25,243 nuclei for subject 4956; 6,967 for subject 5610; and 13,294 nuclei for subject 6173). e,f, log10 number of genes (nGene) (E) and log10 number of UMIs (nUMI) (F) per cell type (n = 15,684 DA neuronal nuclei; 46,860 non-DA neurons; 5,866 OPC nuclei; 14,579 astrocyte nuclei; 76,837 oligodendrocyte nuclei; 16,755 microglia/macrophage nuclei; and 8,092 endothelial cell/fibroblast nuclei). For all box plots, center bar indicates median value and lower and upper hinges correspond to first and third quartiles respectively. Whisker distance from upper and lower hinges represents ≤1.5xIQR. DA, dopaminergic; non-DA, non-dopaminergic, OPC, oligodendrocyte precursor cell; Olig, oligodendrocyte; Endo/pericyte, endothelial cells/pericytes; MG, microglia/macrophage. g,h, UMAP representations of the NR4A2+ (G) and NR4A2− (H) gated nuclei, colored by cluster. i,j, Violin plots of marker genes for the seven major cell classes in the NR4A2+ (I) and NR4A2− (J) gated nuclei. DA, dopaminergic; non-DA, non-dopaminergic; Astro, astrocyte; MG, microglia/macrophages; OPC, oligodendrocyte precursor cell; Olig, oligodendrocyte; Endo, endothelial cells.
Extended Data Fig. 2 | See next page for caption.
Extended Data Fig. 2 | High-throughput profiling and characterization of human midbrain DA neuron diversity. a, Bar plot (log scale) of total number of midbrain DA neurons profiled in this study and all previous single-cell studies of the human midbrain. b, Factor loadings obtained from LIGER integrative analysis projected onto UMAP embedding of DA neuron analysis. Title of each plot indicates highest loading DA subtype and top five highest loading genes (see Methods) for each factor. c, Upset plot of gene sets specific to each DA subtype determined by MAST differential expression analysis (Methods). d, SNpc UMAP representation of 15,684 DA profiles colored by individual. e, Violin plots showing scaled expression of genes previously defined by La Manno et al. as specific to midbrain DA neurons (left) and specific to subtypes identified in the mouse brain (right). f, Box plot of proportion of SOX6+ and CALB1+ DA neurons per individual (n = 8 neurotypical controls). Center bar indicates median value and lower and upper hinges correspond to first and third quartiles respectively. Whisker distance from upper and lower hinges represent ≤1.5×IQR. g, Expression of (in descending order) SLC6A3, SLC18A2, TH, SOX6, and CALB1 per individual. h, UMAP embedding of DA neurons colored by annotations determined via Harmony (see Methods). i, Density plots depicting −log10-transformed p-values of a resampling test (see Methods) to determine concordance of LIGER annotations in the Harmony-derived UMAP embedding. j, Scatter plot depicting average expression per DA subtype for the LIGER-derived annotations (X-axis, Fig. 1d) and Harmony-derived annotations (Y-axis, Extended Data Fig. 2h).
Extended Data Fig. 3 | See next page for caption.
Extended Data Fig. 3 | Cross-species analysis of DA subtypes. a–c, Expression of SLC18A2 (DA neuron marker) (A), SOX6 (B), and CALB1 (C) across all five species sampled. d, Confusion matrix showing overlap of human cells within clusters defined by the cross-species integrative analysis and the human-only analysis (Fig. 1d). The clusters colored in red correspond to the primate-specific population (DA8) and cognate population in the human-only analysis (CALB1_GEM). e, Dot plot of additional marker genes for the primate-specific population, DA8. f, Feature plots of genes defined by Poulin et al as specific to subtypes of DA neurons in the mouse midbrain. g, Single-molecule fluorescence in situ hybridization (smFISH) images depicting absence of Fam83b + mouse nigral DA neurons. Top row = scanned and tiled image. Middle = white inset from top row indicating midline DA neurons. Bottom row = black inset from top row showing high-resolution images of mouse pars compacta dorsal tier. Th (green), Calb1 (red), and Fam83b. Top row scale bar = 100 microns. Middle/bottom row scale bar = 10 microns. h, Single-molecule FISH images depicting absence of Gem + mouse nigral DA neurons. Top = scanned and tiled view of in situ hybridization stains of mouse midbrain. Middle and bottom rows indicate high resolution example images of Th +/ Calb1+/Gem- cells in the dorsal tier (middle) and midline A9 region (bottom). Top row scale bar = 100um. Middle/bottom row scale bar = 10 microns. i, High-resolution view of in situ hybridization stains of Acta2 and Gem showing detection of Gem in smooth muscle vasculature but not DA neurons. Scale bar = 20 microns. Experiment was repeated twice for Fam83b and Gem in situ analyses and once for the Acta2 co-localization in mural cells.
Extended Data Fig. 4 | Additional regulon analyses across all DA subtypes. a, Top ten differentially enriched (FDR-adjusted p-value < 0.05, Wilcoxon rank-sum two-sided test) regulons per DA subtype. Scaled regulon score is the column-wise average scaled regulon activities per subtype as determined from SCENIC analysis (see Methods). b, Scatter plot indicating correlation of between scaled significant regulons from SCENIC analysis of macaque (X-axis) and human (Y-axis) DA subtype profiles.
Extended Data Fig. 5 | See next page for caption.
Extended Data Fig. 5 | Slide-seq mapping of DA neurons. 

a, Box plots showing, for each bead array, the log number of genes captured per bead (n = 27 bead arrays). Each bead array is a part of a triplet set serially created during the dissection (Methods). Center bar indicates median number of genes and lower and upper hinges correspond to first and third quartiles of data, respectively. Whisker distance from upper and lower hinges represent no more than 1.5× interquartile range. 

b, Anatomical plates of Macaca fascicularis brain (71) in sagittal view with coronal sectioning. Highlighted red region is the substantia nigra. Black arrows indicate approximate locations of start (green highlighted plate) and end (blue highlighted plate) of sets of bead arrays sampled via Slide-seq. 

c, Spatial plot for the pucks in Set 1 showing each bead’s expression of TH, ALDH1A1, CALB1, and SLC6A3. 

d, UMAP showing integrative analysis (Methods) of control DA neuron snRNA-seq profiles and Macaca fascicularis DA neuron snRNA-seq profiles. Left — colored by DA types defined by the human integrative analysis (Fig. 1d). Right — colored by species. 

e, Expression of CALB1 across all 27 bead arrays showing putative location of the substantia nigra pars compacta dorsal tier.

f, Bead arrays showing localization of all 10 DA subtypes from Set 1 (Fig. 2b-d) as determined by RCTD (Methods).

g, Location of aggregated SOX6+ and CALB1+ subtypes across all 27 bead arrays.

h, Location of three highly spatially variable subtypes (SOX6_AGTR1, CALB1_TRHR, CALB1_GEM) across all 27 bead arrays.
Extended Data Fig. 6 | Single-molecule fluorescence in situ hybridization of DA neurons in postmortem human midbrain. 

a, Top row - representative images of TH in situ hybridization in postmortem midbrain (Methods, scale bar = 10um). Bottom row - masks obtained from FIJI denoting location and size of DA neuron (Methods). 
b-f, Representative images for: TH+/SOX6+/AGTR1+ (B), TH+/CALB1+/TRHR+ (C), TH+/CALB1+/GEM+ (D), TH+/SOX6+/GFRA2+ (E), TH+/SOX6+/PART1+ (F). White arrows indicates lipofuscin-associated autofluorescence and * indicates neuromelanin-associated autofluorescence defined by the absolute co-localization of pixel values across all channels. Scale bar = 10um. 
g, Location of SOX6+/GFRA2+ (G) and SOX6+/PART1+ (H) across human midbrain, yellow = triple-positive cells, green = single-positive (TH+) cells. Sections for (G) and (H) were obtained from 1.8 mm from the most rostral/anterior aspect of the pars compacta. Scale bar = 1mm for (G) and (H). Experiment was repeated once for each subpopulation.
Extended Data Fig. 7 | See next page for caption.
Extended Data Fig. 7 | Case-control scRNA-seq integrative analysis of SNpc. a,b, Box plots showing, per individual, the median number of UMIs (A) and genes (B) per profile (n = 8 neurotypical controls and 10 PD/LBD samples). ns, not significant Wilcoxon rank-sum two-sided test p > 0.05. Center bar indicates median value and lower and upper hinges correspond to first and third quartiles respectively. Whisker distance from upper and lower hinges represent no more than 1.5*IQR* range. c,d, Violin plots of number of UMIs (C) and genes (D) per profile across eight major cell classes grouped by disease (PD/LBD, blue) and control (Ctrl, red). e-f, Box plots showing, per individual, the age at death (E) and postmortem interval (F) stratified by case (PD/LBD) and control (Ctrl) (n = 8 neurotypical controls and 10 PD/LBD samples). Center bar indicates median value and lower and upper hinges correspond to first and third quartiles respectively. Whisker distance from upper and lower hinges represent no more than 1.5*IQR* range (ns = not significant, Welch two sample t-test, p = 0.61 for (E) and p = 0.18 for (F)). g, Stacked bar plot of sex stratified by disease status. h, Histogram of the year of collection of postmortem midbrain samples colored by case-control status (see Supplementary Tables 1 and 2 for raw values). i, Scatter plot of the collection year for each sample (x-axis) and the median number of UMIs (y-axis). Blue line indicates line of best fit as determined by lm.fit in R. Grey shading indicates standard error of fit. P-value corresponds to significance of slope from a mixed-effect model (see Methods). j-o, UMAP representations of Non-DA neurons (n = 91,479 nuclei) (J), Astrocytes (n = 33,506 nuclei) (K), Oligodendrocyte precursor cells (OPCs) (n = 13,691 nuclei) (L), Oligodendrocytes (n = 178,815 nuclei) (M), Endothelial cells/pericytes (n = 14,903 nuclei) (N), and Microglia/macrophages (n = 33,041 nuclei) (O) colored by cell type (left) and individual (right). P, Alignment scores (see Methods) for each of the seven non-DA cell classes.
Extended Data Fig. 8 | Analyses and robustness testing of cell type proportional changes in PD/LBD. a, Box plot of log2 ratio of cell types normalized to median of control ratios. Pink and blue dots denote control and PD/LBD individuals, respectively (* = p < 0.05, p = 0.034, Wilcoxon rank-sum two-sided test, n = 8 neurotypical controls and 10 PD/LBD postmortem samples). Center bar indicates median value and lower and upper hinges correspond to first and third quartiles respectively. Whisker distance from upper and lower hinges represent no more than 1.5*interquartile range. b, Left = UMAP of 33,041 microglia/macrophage cells. Middle and right = expression of GPNMB and SULT1C2 marking specific expression in the annotated microglia cluster. c, Dot plot of downsampling analysis for two microglia subtypes. Red dotted line is FDR-adjusted p-value = 0.05. d, Left = UMAP of 33,506 astrocytes. Middle and right = expression of VIM and LHX2 respectively marking specific expression in the annotated astrocyte cluster. e, Dot plot of downsampling analysis for three astrocyte subtypes. Red dotted line is FDR-adjusted p-value = 0.05.
Extended Data Fig. 9 | Analysis of human dorsal striatum by snRNA-seq. a,b, Median number of UMIs (A) and genes (B) per donor. c,d Violin plots showing the number of UMIs (C) and (D) across eight major cell classes defined by clustering. (n = 9,096 iSPNs; 10,503 dSPNs; 8,433 astrocytes; 1,418 endothelial cells/fibroblasts; 2,115 inhibitory neurons; 2,445 OPCs; 10,667 oligodendrocytes), direct spiny projection neurons; iSPN, indirect spiny projection neuron; OPC, oligodendrocyte precursor cells. For box plots in (C) and (D), center bars indicates median value and lower and upper hinges correspond to first and third quartiles respectively. Whisker distance from upper and lower hinges represent no more than 1.5*interquartile range. e, UMAP representation of 46,872 single nuclei from the dorsal striatum colored by major cell class. Astro, astrocyte; Olig, oligodendrocyte; Endo cells, endothelial cell/pericyte. f, Dot plot showing specific expression of selected marker genes across the eight major cell classes. g, Percent contribution of four tissue donors to the eight major cell classes.
Extended Data Fig. 10 | Genetic enrichment analyses of the substantia nigra cell types and subtypes. a, Bar plot of -log10-transformed p-values from s-LDSC (stratified LD score) enrichment of Alzheimer’s (left) and Parkinson’s disease (right) across 16 cell types from dorsal striatum (caudate) and SNpc. Bars are colored for significantly (Bonferroni-corrected p-value < 0.05) enriched cell types. b, Dot plot of -log10-transformed p-values for s-LDSC analysis of PD genetic risk in the 68 transcriptionally defined SNpc clusters. Clusters are grouped on the y-axis by major cell class. Red dotted line indicates the Bonferroni significance threshold (p-adjusted < 0.05). c, Scatter plot of number of nuclei sampled per DA subtype (x-axis) and -log10-transformed p-values from MAGMA (top) and s-LDSC (bottom) analyses. Blue line indicates the line of best fit as determined by lm.fit package (Methods). P-value determined from a two-sided significance testing of the slope coefficient from the best fit line (n = 9 DA subtypes, see Methods). d, Dot plot of -log10-transformed p-values for MAGMA analysis of DA subtype markers across gene set sizes (Methods) for PD common variant study(38). e, Pseudo-Manhattan of genes specific to each DA subtype (Extended Data Fig. 2c) ordered by scaled -log10-transformed p-values from MAST (Methods). Genes are those prioritized by MAST (Z-score > 4.568).
Reporting Summary

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For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

- The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
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- The statistical test(s) used AND whether they are one- or two-sided
  
  *Only common tests should be described solely by name; describe more complex techniques in the Methods section.*

- A description of all covariates tested
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- A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
- For null hypothesis testing, the test statistic (e.g. $F$, $t$, $r$) with confidence intervals, effect sizes, degrees of freedom and $P$ value noted
  
  *Give $P$ values as exact values whenever suitable.*

- For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
- For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
- Estimates of effect sizes (e.g. Cohen’s $d$, Pearson’s $r$), indicating how they were calculated

Our web collection on statistics for biologists contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection

- Human, mouse, tree shrew, and rat raw single-nuclei sequencing data were processed, aligned, and converted to a digital gene expression matrix by DropSeqTools v2.4.0 with default settings. Macaque single-nuclei raw sequencing data were processed, aligned, and converted to a digital gene expression matrix by CellRanger v5. The human caudate nuclei experiment sequencing data were processed, aligned, and converted to a digital gene expression matrix by CellRanger v3. Slide-seq raw sequencing data were processed, aligned, and converted to a digital gene expression and x-y coordinate matrices using SlideSeqTools v0.2. In situ hybridization data was acquired with Nikon NIS Elements AR software.

Data analysis

- All data was analyzed using Seurat v2.3.4, LIGER v0.4.2, SCENIC v1.2.4, RCTDv1.0, MASTv1.8.2, MASC v0.1.0-alpha, MAGMAv1.8, LDSCv1.0.1, Harmony v1.0, FGSEAv3.14, and SLMv1.3.0 on R v3.6.3. In addition, all scripts and custom code to reproduce the main figures are available at the following github repo: https://github.com/tkamath1/Kamathetal2022. All in situ hybridization images were analyzed with ImageJ v2.1.0/1.53c and Nikon NIS Elements AR software. The Sony SH800S software was used to analyze, visualize, and process flow cytometry data.

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.
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Policy information about availability of data
All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:
- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our policy

All processed data, UMAP coordinates, and annotations have been made freely available to download and visualize at the Broad Institute Single Cell Portal (note two links, one for the single-nuclei data and the other for the Slide-seq data): https://singlecell.broadinstitute.org/single_cell/study/SCP1768/ and https://singlecell.broadinstitute.org/single_cell/study/SCP1769/. Raw and processed data to support the findings of this study have been deposited in GEO under accession number GSE178265. For transcription factor analysis, the TRRUST 2019, Encode and CHEA Consensus, and ARCHS4 TF-coexpression public datasets were used. All are available for download via the enrichR website: https://maayanlab.cloud/Enrichr/#libraries

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Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size

No statistical methods were used to pre-determine sample sizes but our sample sizes are similar to those reported in previous publications using single-cell analyses to identify vulnerable and resistant cell populations (PMID: 33432193)

Data exclusions

For transcriptomic data, a UMI cutoff of 650 was set to remove low-quality nuclei from the analysis.

Replication

For differential abundance assessment, we performed in situ hybridization to validate the selective loss of the SOX6+/AGTR1+ DA neurons. The identification of a loss of this cell population was nominated by our differential abundance analysis and validated by an orthogonal assay using single-molecule FISH repeated once across an external set of 10 PD and 10 neurotypical control postmortem samples and performed once. Experiments to generate single-nuclei RNA-sequencing enriched DA neurons were validated across postmortem human and macaque tissue, having been repeated 19 times, 18 times using postmortem human samples, and a single time with a macaque postmortem sample by a wholly different experimentalist.

Randomization

No randomization occurred during the study. Given differences in age and sex across two arms of the study, those covariates were included in all differential abundance and expression calculations.

Blinding

For in situ validation of selective neuronal susceptibility and resistance and the localization/quantification of DA subtypes, all samples were blinded before staining, imaging, and quantification. Blinding was not performed for any other experiment as the single-nuc. RNA-sequencing data were generated agnostic to the hypothesis identified in the study.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

| n/a | Involved in the study |
|-----|-----------------------|
| ☐   | Antibodies            |
| ☒   | Eukaryotic cell lines |
| ☐   | Palaeontology and archaeology |
| ☐   | Animals and other organisms |
| ☐   | Human research participants |
| ☒   | Clinical data         |
| ☐   | Dual use research of concern |

Methods

| n/a | Involved in the study |
|-----|-----------------------|
| ☒   | ChIP-seq              |
| ☒   | Flow cytometry        |
| ☒   | MRI-based neuroimaging |
Antibodies

Antibodies used
1 - anti-NR4A2-A647 antibody (Santa Cruz, sc-376984, A647)
2 - anti-NeuN-PE; clone A60 (EMD Millipore Corp., FCMAB317PE)

Validation
anti-NR4A2-647 validation from Santa Cruz website (https://www.scbt.com/p/nurr1-antibody-f-5)
"[anti-NR4A2] available conjugated to...Alexa Fluor® 647 for WB (RGB), IF, IHC(P) and FCM"

Anti-Neun-PE validation from EMD website (https://www.emdmillipore.com/US/en/product/Milli-Mark-Anti-NeuN-PE-Antibody-clone-A60,MM_NF-FCMAB317PE?RefererURL=https%3A%2F%2Fwww.google.com%2F#overview)
"This Milli-Mark Anti-NeuN-PE Antibody, clone A60 is validated for use in FC (flow cytometry) for the detection of NeuN."

Animals and other organisms

Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research

Laboratory animals
Animals used:
Mus musculus - C57BL/6J, 3 months, male/female
Rattus norvegicus - 0.2 years, male
Tupaia belangeri - 3 years, male
Macaca fascicularis - 8 years, male
Mice were group housed with a 12-hour light-dark schedule and allowed to acclimate to their housing environment for two weeks post arrival. For mus musculus housing, ambient temperature was strictly maintained between 68 and 72 degrees Fahrenheit and humidity was strictly maintained between 30 and 50%. All procedures involving animals at MIT were conducted in accordance with the US National Institutes of Health Guide for the Care and Use of Laboratory Animals approved by the Massachusetts Institute of Technology Committee on Animal Care. All procedures involving animals at the Broad Institute were conducted in accordance with the US National Institutes of Health Guide for the Care and Use of Laboratory Animals under protocol number 0120-09-16.

Wild animals
No wild animals were used in this study.

Field-collected samples
No field-collected samples were used in this study.

Ethics oversight
All housing and procedures involving rodents were conducted in accordance with the US National Institutes of Health Guide for the Care and Use of Laboratory Animals under protocol number 0129-09-16 and approved by the Broad Institute Committee on Animal Care (IACUC). All non-human primate (NHP) tissue was processed in compliance with the Broad Institute IBC (IBC#: 2016-00127). All human tissue falls under a "Not Engage" designation determined by the Broad Institute IACUC (Federal-wide assurance #: FWA00014055).

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Human research participants

Policy information about studies involving human research participants

Population characteristics
This study involved the generation of RNA-sequencing data from postmortem human midbrain samples which fall under a "Not Engage" policy (NE5200) approved by the Broad IACUC (Federal Wide Assurance number: FWA00014055). Human midbrain samples age and biological sex are listed in Extended Data Tables 1-4. No genotypic information was obtained on these samples. Past diagnoses are listed in Extended Data Tables 1 and 2.

Recruitment
No recruitment was performed for this study.

Ethics oversight
This study falls under a "Not Engaged" determination as approved by the Broad Institute IACUC (FWA00014055).

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Flow Cytometry

Plots
Confirm that:
- The axis labels state the marker and fluorochrome used (e.g. CD4-FITC).
- The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).
- All plots are contour plots with outliers or pseudocolor plots.
- A numerical value for number of cells or percentage (with statistics) is provided.

Methodology
Sample preparation
All samples were prepared according to the Methods section of this paper. A detailed protocol for single-nuclei isolation is available via Protocols.io at: https://www.protocols.io/view/frozen-tissue-nuclei-extraction-for-10xv3-sseq-bi62khge. Nuclei
isolates were obtained from cryosections of postmortem mammalian tissue.

| Instrument          | Samples were flow-sorted using a Sony SH800 Cell Sorter |
|---------------------|---------------------------------------------------------|
| Software            | The standard Sony SH800S software was used for flow cytometry analysis |
| Cell population abundance | Abundances of the NR4A2+ neurons were determined post-sorting using single-nucleus RNA-sequencing |
| Gating strategy     | No preliminary FSC/SSC gating was performed. All gating is performed as listed in the Methods of the paper and the associated protocols.io link for the DAPI gating: https://www.protocols.io/view/frozen-tissue-nuclei-extraction-for-10xv3-snseq-bi62khge?step=19. |

Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary Information.