Kv4.3 channel downregulation mediates chronic post-lesional pacemaker acceleration in surviving dopamine substantia nigra neurons

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Abstract

Substantia nigra dopamine (SN DA) neurons are progressively lost in Parkinson disease (PD). While the molecular and cellular mechanisms of their differential vulnerability and degeneration have been extensively studied, we still know very little about potential functional adaptations of those SN DA neurons that – at least for some time – manage to survive during earlier stages of PD. We utilized a partial lesion 6-OHDA mouse model to characterize initial electrophysiological impairments and chronic adaptations of surviving identified SN DA neurons, both *in vivo* and *in vitro*. Early after lesion (3 weeks), we detected a selective loss of *in vivo* burst firing in surviving SN DA neurons, which was accompanied by *in vitro* pacemaker instability. In contrast, late after lesion (>2 months), *in vivo* firing properties of surviving SN DA neurons had recovered in the presence of 2-fold accelerated pacemaking *in vitro*. Finally, we show that this chronic cell-autonomous adaptation in surviving SN DA neurons was mediated by Kv4.3 channel downregulation. Our study demonstrates substantial homeostatic plasticity of surviving SN DA neurons after a single-hit non-progressive lesion, which might contribute to the phenotype of initially surviving SN DA neurons in PD.

Keywords

dopamine, 6-OHDA, Kv4.3 channel, substantia nigra, accelerated pacemaker

Introduction

One clinically relevant hallmark of Parkinson disease (PD) is the progressive degeneration of vulnerable neuronal populations, including substantia nigra dopamine (SN DA) neurons (Giguere et al., 2018; Kalia and Lang, 2015; Poewe et al., 2017). At the time of clinical diagnosis, more than 50% of SN DA neurons, in particular those in ventrolateral regions, are already lost (Gibb and Lees, 1991; Kordower et al., 2013). Moreover, Kordower and colleagues (Kordower *et al.*, 2013) demonstrated that most remaining SN DA neurons were rapidly lost (or lose their DA phenotype) within the first few years of clinically manifest PD. While neuropathological studies map the time course of degeneration with cellular resolution and also allow for a molecular characterization of the remaining neurons, they cannot investigate the functional properties of surviving neurons. *In vivo* recordings of putative human SN DA neurons in the context of DBS surgery provided a first glimpse of electrically active DA survivors (Dragicevic et al., 2015; Schiemann et al., 2012; Zaghloul et al., 2009), but a systematic investigation of these DA neurons is not feasible in a therapeutic setting. Thus, studying functional adaptations of surviving DA neurons in the context of PD is the domain of animal models, which themselves often capture only partial aspects of the disease. Among the various rodent PD models, 6-hydroxydopamine (6-OHDA) lesions have been extensively used to study the DA-depleted basal ganglia (Chen et al., 2021; Fieblinger et al., 2018; Kravitz et al., 2010; Parker et al., 2018; Parker et al., 2016; Rubi and Fritschy, 2020; Sharott et al., 2005; Sitzia et al., 2020; Ungerstedt, 1968; West et al., 2018). By comparison, a smaller number of studies applying only partial lesions focused on the properties of the remaining DA midbrain system, including its long-term adaptations and drug responses (Bez et al., 2016). These studies demonstrated that – in contrast to PD – 6-OHDA DA lesions are non-progressive (i.e. single-hit) and follow a stereotypical time course: an initial phase dominated by cell loss, inflammation and behavioral impairment for about 3 weeks, which is followed by a chronic phase (studied for up to 20 month in mice), where partial behavioral recovery, neurochemical and molecular adaptations and – to a certain degree – axonal sprouting of DA neurons occur (Bez *et al.*, 2016; Cenci and Bjorklund, 2020; Kirik et al., 1998; Schwarting and Huston, 1996; Winkler et al., 2002). Single cell

electrophysiological investigations of viable post-lesional DA neurons, however, have been exceedingly rare with only Hollerman & Grace providing a first data set (Hollerman and Grace, 1990). They found 6-OHDA-lesion-size-dependent changes in *in vivo* firing properties of putative DA neurons in rats, among them a decrease of *in vivo* burst firing after 50-60% (sized) lesions (Hollerman and Grace, 1990).

In contrast, cell-autonomous characteristics of surviving SN DA neurons after 6-OHDA lesion have not yet been studied. Here, we aim to close this gap by exploring the electrophysiological properties *in vivo* and *in vitro* of surviving SN DA neurons in a well-characterized, intra-striatal unilateral 6-OHDA lesion mouse model, where about 50% of SN DA neurons survive. We selected two time points of electrophysiological investigation based on our long-term behavioral characterization of motor phenotypes. In particular, we studied the dynamics of spontaneous locomotion and turning behavior in a drug-free setting. This identified a 3-week post-lesion time point (i.e. early phase), where motor impairment was prominent, and a stable late phase (>68 days post-lesion), where turning asymmetry had completely recovered. In contrast to the significant behavioral differences between early and late post-lesional time points, the extent of the striatal lesion and the degree of stereologically-determined TH-positive cell numbers were stable. This pointed to a relevant contribution of early functional impairment and late adaptation in a stable pool of surviving SN DA neurons as detailed below.

Results

Early behavioral impairments and long-term adaptations after partial lesion of substantia nigra dopamine neurons

In order to identify potential compensatory adaptations in surviving SN DA neurons, we utilized a C57Bl6N mouse model with a partial 6-OHDA lesion of the nigrostriatal pathway. The neurotoxin 6-OHDA was unilaterally infused into the right dorsal striatum, causing a partial degeneration of DA neurons in the ipsilateral substantia nigra, which was quantified by unbiased stereology (Fig. 1A). We continuously characterized motor behavior of the model (i.e. open field, cylinder test) until two different time points, either 21-days post-lesion (early phase) or more than 60 days post-lesion (late phase). Initially, animals were sacrificed for TH immunohistochemistry in midbrain and striatum. As shown in Fig. 1B, at the early phase, the number of TH-positive (i.e. dopaminergic) SN neurons was reduced post-6-OHDA (lower left panels) compared to control mice, where only vehicle (ACSF with ascorbic acid) was infused into the dorsal striatum (upper left panels). The right panel in Fig. 1B displays the stereological results of TH-positive SN neurons for N=9 vehicle-treated and N=9 6-OHDA-infused mice. The data were normalized to the stereological results of respective contralateral SN DA neurons. In contrast to vehicle, where no ipsilateral loss of SN DA neurons was detected, the number of ipsilateral surviving SN DA neurons in the early post-6-OHDA phase was on average reduced to about 40% (Fig. 1B; vehicle: 100 ± 2.644 %; 6-OHDA: 36.85 ± 3.463 %; $p < 0.0001$, Mann-Whitney test). We also carried out the unbiased stereology for TH-positive SN neurons in the late phase and detected a similar degree of loss (Suppl. Fig. 1C, vehicle: 102.6 ± 7.9 %; 6-OHDA: 39.96 ± 3.7 %; $p = 0.0262$, Mann-Whitney test). These results indicated that SN DA cell loss was stable throughout our observation period of more than 2 months. In addition, we analyzed the axonal compartment of DA midbrain neurons by determining striatal TH-optical densities, both in early and late phase (Suppl. Fig. 1). Similar to cell counts, we found an about 50% reduction of TH-immunosignal in the ipsilateral dorsal

striatum, both in early and late phase, whereas the ventral striatum was only mildly affected (ca. 20% reduction). In summary, our model induces a stable loss of SN DA neurons, associated with a stable reduction of striatal TH-immunoreactivity throughout the entire observation period. In other words, in accordance with previous chronic 6-OHDA studies (Bez *et al.*, 2016; Cenci and Bjorklund, 2020; Schwarting and Huston, 1996), this model showed no progression and is therefore suitable to study functional adaptations in a stable pool of surviving SN DA neurons.

Fig 1 C shows our continuous behavioral monitoring of unilateral 6-OHDA and vehicle-treated mice before and up to 68 days post-infusion. Based on previous studies (Cenci and Bjorklund, 2020; Schwarting and Huston, 1996), we focused on the dynamics of drug-free spontaneous turning behavior during open field locomotion (Fig. 1C, lefts panels). In contrast to vehicle-infused mice, which displayed a stable symmetric ratio (ca. 50%) of ipsi- to contralateral turning throughout the entire experiment, 6-OHDA-infused mice showed a dramatic shift to ipsilateral turning immediately after treatment (Figure 1C, right panel). Interestingly, ipsilateral turns occurred in long sequences of up to 40 individuals turns, a pattern not observed in controls (Suppl. Fig. 2). Importantly for this study, these turning sequences as well as the overall ipsilateral bias gradually but eventually completely recovered over 2-months post treatment (two-way-ANOVA, p-value across time $p < 0.0001$, p-value across groups $p < 0.0001$, significant difference between vehicle and 6-OHDA group till day 40, Šídák's multiple comparisons test). Analysis of turning features, such as diameter or velocity, revealed that the recovered contralateral turns were similar to turns performed by vehicle-treated mice (Suppl. Fig. 3). This could be indicative for a degree of functional recovery and/or functional compensation within the stable pool of surviving DA neurons. General locomotion, measured as performed track length per session, recovered in parallel to the turning bias (Suppl. Fig. 1). However, other motor behaviors, such as contralateral paw use during rearing in the cylinder test, did not show recovery throughout the experimental period (Fig. 1D) (21st post-infusion day: vehicle $38.5 \pm 5.24\%$, 6-OHDA $5.73 \pm 1.35\%$, $p < 0.0001$, Mann-Whitney test; 68th post-infusion day: vehicle $63.67 \pm 6.6\%$, 6-OHDA $3.167 \pm 1.8\%$, $p < 0.0001$, Mann-Whitney test), which is consistent with the partial but stable loss of SN DA neurons and reduction of dorsal striatal TH.

To investigate this model's homeostatic mechanisms from the perspective of the surviving SN DA neurons, we performed *in vivo* extracellular recordings and *in vitro* whole-cell patch-clamp recordings once within a period with strong turning bias, at the 21st day after lesion (early phase), and at a later time point – after reaching stable baseline levels (late phase).

Figure 1. Unilateral striatal 6-OHDA mouse model with stable loss of SN DA neurons resulted in delayed partial behavioral recovery.

(A) Experimental design, illustrating timeline of behavioral assays with termination points – groups ended either on the 21st, or later than the 64th post-infusion day. (B) Left panel: TH-DAB staining of the SN, 10x magnification and 60x magnification for vehicle and 6-OHDA-infused mouse. Scalebar left

200µm, scalebar right 25µm. Right panel: ratio of ipsilateral (infusion side) to contralateral side of surviving TH-positive neurons in SN at 21st post-infusion day. (C) Left: spontaneous locomotion of mice in open field arena for a 10 min session. Left, examples of an ACSF-infused mouse (vehicle), right example of a 6-OHDA-infused mouse at baseline (upper panels), 4th post-infusion day (middle panels) and 68th post-infusion day (lower panels). Right: ratio of ipsilateral to contralateral turning behavior for all experimental groups plotted against session days. Note recovery in the 6-OHDA-treated mice from day 4 to day 68 after initial strong turning bias (>90%). (D) Cylinder test quantified by ratio of contra- to ipsilateral forepaw use. Note significant loss of contralateral forepaw involvement, both at 21 and 64 days. All data are presented as mean ± standard error of mean (SEM).

Surviving SN DA neurons in the early post-6-OHDA phase exhibit impaired firing properties both *in vivo* and *in vitro*

In the early post-6-OHDA phase, we explored the *in vivo* spontaneous activity of surviving identified SN DA neurons in isoflurane-anesthetized mice, using single-unit extracellular recordings combined with juxtacellular labelling and post-hoc TH immunohistochemistry (see schema Fig. 2A). Figure 2B shows the *in vivo* extracellular electrical activity (top panel) and identification of a representative SN DA neuron (lower right panel) from a control mouse, 3 weeks after vehicle infusion. Note the typical combination of irregular single spike firing with occurrence of transient fast burst discharges (bursts were defined by the 80/160ms Grace criteria; see methods for details). The interspike interval (ISI) histogram, which captures ISIs from the entire recording time of 10 min for this cell, displays the typical features of an *in vivo* bursty DA neuron, i.e. a bimodal ISI distribution representing intra- and interburst intervals over the dynamic range between ~30-1000 ms (lower left panel). This recorded and labelled cell was localized in the medial SN and characterized as TH-immunopositive. In comparison, Fig. 2C

displays representative *in vivo* activity of a surviving identified SN DA neuron 3 weeks after 6-OHDA-infusion. Here, while irregular single-spike pacemaking was present similar to controls, fast burst events were almost completely absent (top panel). In accordance, the ISI histogram of this neuron, for the entire 10 min of recording, showed a unimodal, Gaussian-like distribution without fast intraburst and slow interburst intervals (Fig. 2C, lower left panel). This recorded and labelled cell was also localized in the medial SN and identified as TH-immunopositive.

Similar to these two example cells, all *in vivo* recorded DA neurons from the early phase were successfully juxtacellularly labeled with neurobiotin (NB), post-hoc immunohistochemically reconstructed and localized within the ventral midbrain (Fig 2D, vehicle: n =28, N =15; 6-OHDA: n =18, N =16). For best comparison, we restricted our analysis to those DA midbrain neurons localized in the medial half of the SN (mSN), where a sufficient number of surviving and electrically active DA neurons could be sampled after 6-OHDA lesion. In contrast, we only managed to detect a very small number of active DA neurons in the lateral SN (lSN), the most vulnerable part of the DA midbrain (Figure 2D). Consequently, Figure 2 E-H compares the group data of surviving mSN DA neurons at early phase between vehicle and 6-OHDA treatment. We detected only a small reduction (< 40 %) of overall firing rate for post-6-OHDA mSN DA neurons compared to vehicle-treated controls (Fig.2E; vehicle: mean frequency = 7.5 ± 0.92 Hz, 6-OHDA: mean frequency = 4.71 ± 0.6 Hz, $p = 0.0184$, Mann-Whitney test) and no differences in overall firing regularity expressed as CV (Fig. 2F; vehicle: CV = 53.96 ± 7.78 %, 6-OHDA: CV = 36.06 ± 3.7 %, $p = 0.1694$, Mann-Whitney test). In contrast, the reduction of bursting - expressed as the percentage of spikes fired in bursts (SFB) and burst rate - was dramatic (ca. 10-fold) and highly significant for mSN DA neurons in the 6-OHDA group compared to vehicle controls (Fig. 2G, vehicle: SFB = 31.22 ± 7.71 ; 6-OHDA: SFB = 2.24 ± 1.09 %; $p = 0.0032$, Welch's t test; Fig. 2H, vehicle: burst rate = 0.34 ± 0.09 Hz, 6-OHDA: burst rate = 0.04 ± 0.02 Hz; $p = 0.0066$, Welch's t test). All other analyzed firing parameters identified no additional differences between the two early groups (Suppl. Fig. 4). In summary, we detected a dramatic compression of the dynamic *in vivo* firing range in mSN DA neurons surviving the 6-OHDA lesion. In addition to the degree of DA cell loss, this functional

impairment of surviving DA neurons might be a novel contributing factor to the extensive motor impairment, observed during the early post-6-OHDA phase.

Burst discharge in DA neurons is orchestrated by the interplay of patterned synaptic inputs with their intrinsic excitability, giving rise to different types of bursting (Otomo et al., 2020). To identify a possible contribution of cell-autonomous changes in intrinsic excitability of surviving mSN DA neurons to their impaired *in vivo* bursting, we also studied these cells *in vitro* in synaptic isolation (see Schema 2I). Figure 2J shows representative stable and regular pacemaker activity of identified mSN DA neuron from a vehicle-infused control (lower right panel), as well as a subthreshold response to hyperpolarizing current injection (lower left panel). In contrast, pacemaker activity of mSN DA neurons post-6-OHDA infusion was unstable and characterized by intermittent periods of action potential failure (see Fig. 2K, upper panel). However, subthreshold responses were similar to controls (lower right panel). Analogous to the *in vivo* experiments, all *in vitro* recorded DA neurons were neurobiotin (NB) filled, identified, and mapped, resulting in an equivalent, medial clustering of surviving SN DA neurons (Fig. 2L). When *in vitro* pacemaker properties were compared between identified mSN DA neurons from the vehicle and the 6-OHDA groups (Fig. 2L), we detected no differences in mean firing rates (Fig. 2M, vehicle: firing rate = 1.699 ± 0.17 Hz, 6-OHDA: firing rate = 2.425 ± 0.25 Hz, $p = 0.1448$, Mann-Whitney test), rebound delays and sag-amplitudes (Fig. 2O, vehicle: rebound delay = 584.9 ± 95.8 ms, 6-OHDA: rebound delay = 429.6 ± 56.1 ms, $p = 0.1738$, Mann-Whitney test; Fig. 2P, vehicle: sag-amplitude = 15.3 ± 1.1 mV, 6-OHDA: sag-amplitude = 13.5 ± 0.96 mV, $p = 0.1395$, Mann-Whitney test), but a highly significant 5-fold increase of pacemaker irregularity expressed as CV (Fig. 2N, vehicle: CV = 6.531 ± 0.7147 %; 6-OHDA: CV = 33.99 ± 5.788 ; $p = 0.0007$, Mann-Whitney test). More detailed analysis revealed that the increased CV was mainly mediated by intermittent phases of firing failure, similar to the one shown in Fig. 2K (Suppl. Fig. 5). Further analysis, including a comparison of on-cell and whole-cell recordings, did not reveal additional differences. In short, we provided the first data set in the literature that surviving SN DA neurons 3-weeks after a 6-OHDA infusion are functionally impaired displaying a 10-fold reduced *in vivo* burst activity. This was accompanied by a 5-fold decrease of *in vitro* cell-autonomous pacemaker stability. In order to address the question whether these

altered properties were stable adaptations or reflected only transient impairment, we also studied *in vivo* and *in vitro* electrophysiology of surviving mSN DA neurons in the late phase, characterized by partial behavioral recovery.

Figure 2. Surviving SN DA neurons at early post-6-OHDA phase exhibited a 10-fold decrease in *in vivo* bursting and a 5-fold decrease in *in vitro* regularity

(A), (I) Experimental design, illustrating timeline of behavioral assays, followed by terminal *in vivo* juxtacellular recordings (A) or by terminal *in vitro* whole-cell recordings (I) at the 21st post-infusion day. (B), (C) Top: 10s original recording trace of spontaneous *in vivo* single-unit activity from SN DA neurons in vehicle (B) and 6-OHDA-infused mouse (C). Scalebars: 1s, 0.2mV. Below, left: corresponding ISI histograms. Below, right: corresponding confocal images of juxtacellularly labelled and immunohistochemically identified DA neuron. Note the sparse bursting of the surviving SN DA neuron from the 6-OHDA-infused mouse at early phase. (D), (L) Anatomical mapping of all extracellularly recorded and juxtacellularly labelled DA neurons (D), and of all *in vitro* recorded and filled DA neurons (L), projected to bregma -3.08 mm. Location of example SN DA neurons in (B), (C), (J), (K) are highlighted. Smaller symbols represent DA neurons that have been recorded and identified, but not included in the group data analysis in (E-H). (E-H) Scatter dot-plots, showing significant decrease of *in vivo* mean firing rate (E), percentage of SFB (G) and burst rate (H) and no significant differences in CV (F) between the vehicle and 6-OHDA-infused mice. Note the 10-fold decrease in SFB for 6-OHDA-infused mice. (J), (K) Top: 10s original recording trace of *in vitro* whole-cell recording of spontaneous SN DA neuron activity in a vehicle (J) and 6-OHDA-infused (K) mouse. Scalebars: 1s, 20mV. Note that the 6-OHDA DA neuron has a highly irregular pacemaking. Below, left: corresponding to hyperpolarizing current injection. Below, right: confocal images of NB filled and immunohistochemically identified DA neuron. (M-P) Scatter dot-plots, showing no difference in *in vitro* mean firing rate (M), rebound delay (O) or sag-component (P) and a 5-fold increase in CV (N). Immunohistochemical imaging for all four DA neurons are displayed in 10x and 60x magnifications (green, TH; red, NB), scalebars: 200µm, 20µm. All data are presented as mean ± standard error of mean (SEM).

Surviving mSN DA neurons in the late 6-OHDA phase display full recovery of *in vivo* burst firing associated with accelerated *in vitro* pacemaking

We studied and analyzed surviving mSN DA neurons in the late 6-OHDA phase in an identical fashion to the early phase (compare Fig. 3A/I and Fig. 2A/I). In contrast to the 10-fold difference in burst rate in early phase, we found no significant electrophysiological *in vivo* differences between the two groups in the late phase (compare Fig 3B with Fig. 3C). Among others (Suppl. Fig. 6), this implies that mean firing frequencies, CV, SFB and burst rates were similar in chronically surviving mSN DA neurons compared to controls (Fig. 3E, vehicle: mean firing rate = 6.95 ± 0.82 Hz; 6-OHDA: mean firing rate = 7.53 ± 1.12 Hz; $p = 0.6038$, Mann-Whitney test; Fig. 3F, vehicle: CV = $54.88 \pm 12.23\%$; 6-OHDA: CV = $53.33 \pm 12.65\%$, $p > 0.9999$, Mann-Whitney test; Fig. 3G, vehicle: SFB = $20.2 \pm 9.11\%$, 6-OHDA: SFB = $22.17 \pm 8.87\%$, $p = 0.8784$, Mann-Whitney test; Fig. 3H, vehicle: burst rate = 0.1728 ± 0.08 Hz; 6-OHDA: burst rate = 0.3071 ± 0.15 Hz, $p = 0.4308$, Mann-Whitney test). This suggests that surviving mSN DA neurons might have fully recovered more than 2-months after lesion. However, the *in vivo* results cannot distinguish between simple recovery (e.g. functional restoration) or adaptive homeostatic plasticity. Therefore, we also studied *in vitro* pacemaker properties of the DA neurons in the late post-6-OHDA phase. Here, we found clear evidence for chronic functional remodeling of surviving mSN DA neurons. When comparing intrinsic pacemaker frequencies, we noticed that late 6-OHDA survivors discharged almost 2-fold faster compared to vehicle-controls (compare Fig. 3J with 3K, Fig. 3M vehicle: firing rate = 1.474 ± 0.1653 Hz; 6-OHDA: firing rate = 2.713 ± 0.2057 Hz, $p = 0.0016$, Mann-Whitney test), while maintaining a stable and regular pattern (Fig. 3N, vehicle: CV = $9.515 \pm 1.7\%$, 6-OHDA: CV = $16.10 \pm 6.661\%$, $p = 0.175$, Mann-Whitney test). In line with this pacemaker acceleration, the sag component was increased, and the rebound delay shortened (Fig 3O, vehicle: rebound delay = 629.6 ± 96.38 ms, 6-OHDA: rebound delay = 364.2 ± 57.6 ms, $p = 0.0216$, Mann-Whitney test; Fig. 3P, vehicle: sag-amplitude = 15.06 ± 0.9487 mV, 6-OHDA: sag-amplitude = 19.11 ± 0.9118 mV, $p = 0.0072$, Mann-Whitney test). Other *in vitro* spike features were not significantly different between the two groups (Suppl. Fig. 7). Interestingly, breaking into whole-cell recording mode further accelerated firing

frequency, while only marginally increasing firing irregularity (Suppl. Fig. 7F). In summary, complete functional recovery of surviving mSN DA neurons was – at least in part – achieved by homeostatic plasticity of intrinsic excitability. Precisely, surviving mSN DA neurons not only recovered from instable pacemaking at the early phase, but managed to double their discharge rates in late phase. As this phenotype is reminiscent to the chronic adaptation of mSN DA neurons to mutant α -synuclein overexpression, (Subramaniam et al., 2014a), we aimed to identify the underlying biophysical and molecular changes.

Figure 3. Surviving SN DA neurons at late 6-OHDA phase recovered *in vivo* burst firing and doubled their intrinsic pacemaker frequency *in vitro*.

(A), (I) Experimental design, illustrating timeline of behavioral assays, followed by terminal *in vivo* juxtacellular recordings (A) or by terminal *in vitro* whole-cell recordings (I) after >64 post-infusion days. (B), (C) Top: 10s original recording trace of spontaneous *in vivo* single-unit activity from SN DA neurons in vehicle (B) and 6-OHDA-infused mouse (C). Scalebars: 1s, 0.2mV. Below, left: corresponding ISI histograms. Below, right: corresponding confocal images of juxtacellularly labelled and immunohistochemically identified DA neuron. (D), (L) Anatomical mapping of all extracellularly recorded and juxtacellularly labelled DA neurons (D), and of all *in vitro* recorded and filled DA neurons (L), projected to bregma -3.40 mm (D) and -3.08mm (L). Location of example SN DA neurons in (B), (C), (J), (K) are highlighted. Smaller symbols represent DA neurons that have been recorded and identified, but not included in the group data analysis in (E-H). (E-H) Scatter dot-plots, showing no significant difference in mean firing rate (E), CV (F), SFB (G) and burst rate (H). (J), (K) Top: 10s original recording trace of *in vitro* whole-cell recording of spontaneous activity from SN DA neurons in a vehicle (J) and 6-OHDA-infused (K) mouse. Scalebars: 1s, 20mV. Note that the 6-OHDA DA neuron has enhanced, but regular pacemaking. Below, left, corresponding to hyperpolarizing current injection. Below, right, confocal images of NB filled and immunohistochemically identified DA neuron. (M-P) Scatter dot-plots, showing doubling of the firing rate (M), no difference in CV (N), decrease in rebound delay (O) and increase of sag-component (P) for the 6-OHDA-treated mice in comparison to vehicle group. Immunohistochemical imaging for all four DA neurons are displayed in 10x and 60x magnifications (green, TH; red, NB), scalebars: 200µm, 20µm. All data are presented as mean ± standard error of mean (SEM).

Accelerated *in vitro* pacemaking of surviving mSN DA neurons is caused by downregulation of Kv4.3 channels

To test whether Kv4.3-mediated A-type channels are responsible for differences in late phase pacemaker rate between 6-OHDA and controls, we carried out a pharmacological occlusion experiment using 1 μ M AmmTx3, a selective Kv4.3 channel blocker (see scheme Fig. 4A). The presence of 1 μ M AmmTx3 not only accelerated discharge rates in both identified mSN DA neurons from vehicle- and 6-OHDA-infused mice (see Fig. 4BC for representative cells; Fig. 4D for mapping) but, importantly, also eliminated significant rate differences between the two treatment groups (Fig. 4E, vehicle: mean firing rate = 4.54 ± 0.18 Hz, 6-OHDA: mean firing rate = 4.84 ± 0.28 Hz, $p = 0.285$; Fig. 4 F, vehicle: CV = 7.84 ± 1.26 %, 6-OHDA: CV = 5.53 ± 0.44 %, $p = 0.2517$; both Mann-Whitney test). This result demonstrated that differences in Kv4.3 channel function were a main driver of the late post-6OHDA fast pacemaker phenotype. In addition, 1 μ M AmmTx3 also abolished more subtle differences in rebound delays and sag amplitudes (Fig. 4G, vehicle: rebound delay = 31.71 ± 2.4 ms, 6-OHDA: rebound delay = 28.4 ± 4.0 ms, $p = 0.0952$; Fig. 4H. vehicle: sag-amplitude = 15.61 ± 1.1 mV, 6-OHDA: sag-amplitude = 16.43 ± 0.95 mV, $p = 0.7434$; both Mann-Whitney test, Suppl. Fig. 8). In summary, these experiments demonstrated that reduced Kv4.3 function and/or expression are mediators of the post-6-OHDA late phase phenotype of accelerated *in vitro* pacing.

Next, we aimed to reveal whether this adaptive phenotype was associated with altered Kv4.3 protein expression. We performed Kv4.3 immunohistochemistry and confocal imaging in vehicle and 6-OHDA treated mice both on the ipsilateral and contralateral infusion side (see scheme in Fig. 4I). We defined DA-selective ROIs by using TH-immunopositive areas to measure the intensity of Kv4.3-immunosignals exclusively within DA neurons (i.e. TH-signal based mask) (Subramaniam *et al.*, 2014a). Figure 4J and 4K compare Kv4.3-immunoreactivities in the midbrain between/from contralateral control side and ipsilateral affected side in a 6-OHDA-treated mouse in the late post-6-OHDA phase. Note the reduction of TH-immunoreactivity on the ipsilateral side (upper panels). Higher resolution of Kv4.3-immunoreactivity in surviving TH-positive neurons showed lower Kv4.3-immunosignal on the treated side. Quantitative comparison of TH-ROIs of the entire SN between contralateral control side and 6-OHDA-treated side revealed a significant reduction of average Kv4.3-immunosignal on the 6-OHDA treated side (Figure 4L, $D = 0.2014$, $p < 0.0001$, two-sample Kolmogorov–Smirnov test). This reduction

was also detected in a selective analysis of Kv4.3-immunosignals in the medial SN (Figure 4M, contralateral side: 73.03 ± 0.389 , ipsilateral side: 54.8 ± 0.486 , $p < 0.0001$, unpaired t-test), the region where we carried out our functional characterization. In addition, Kv4.3 signal intensities showed weak correlation to the size of TH-positive ROIs ($R^2 = 0.0041$, $Sy.x = 1227$, $N = 4874$, $p < 0.0001$, simple linear regression, Suppl. Fig. 9). As Kv4.3 channel subunits are expected to be mostly located on the cell membrane, we also investigated whether Kv4.3-immunosignal was different across cell compartments. A decrease of Kv4.3-immunosignal was observed both in cell membrane and cytoplasm ROIs (Suppl. Fig. 9). In contrast, analogous quantification in vehicle-treated mice revealed no contra-ipsilateral differences in Kv4.3-immunoreactivity (Suppl. Fig. 9). In summary, these immunohistochemical experiments provided evidence for a downregulation of Kv4.3 channel subunits in mSN DA neurons, surviving a partial 6-OHDA lesion. This phenotype was therefore distinct from the overexpression of Kv4.3 channel protein, observed in mutant α -synuclein mice (Subramaniam *et al.*, 2014a). Thus, we provided first evidence for a selective functional adaptation of SN DA neurons in response to a single-hit non-progressive lesion.

Figure 4. Enhanced pacemaker frequency of SN DA neurons at the late post-6-OHDA phase was mediated by Kv4.3 channel downregulation

(A), (I) Experimental design, illustrating timeline of behavioral assays, followed by terminal *in vitro* whole-cell recordings under Kv4.3 channel blocker AmmTx3 (A) or by multi-labeling immunohistochemistry for exploring Kv4.3-channel expression differences (I) after >64 post-infusion days. (B), (C) Top: 10s original recording traces of *in vitro* whole-cell recording of spontaneous activity from SN DA neurons in AmmTx3-preincubated slices for a vehicle (B) and 6-OHDA-infused (C) mouse.

Below, left: corresponding hyperpolarizing current injection. Scalebars: 1s, 20mV. Below, right: confocal images of NB filled and immunohistochemically identified DA neuron (green, TH; red, NB). 10x , 60x magnification, scalebars: 200μm, 20μm. (D) Anatomical mapping of all *in vitro* recorded and filled DA neurons, projected to bregma -2.92 mm. (E-H) Scatter dot-plots, showing loss of differences in *in vitro* electrophysiological parameters. (J), (K) Top: 4x magnification of midbrain of a 6-OHDA-infused mouse, >64 days post-lesion – contralateral side (J), and corresponding ipsilateral side (K). Middle: 60x magnification in the highlighted area from 4x image (green, TH; red, Kv4.3). Bottom, left: zoom-in on an example ROI (highlighted in 60x image). Bottom, right: color-coded Kv4.3-channel immunohistochemical signal intensity in the example ROI. Note Kv4.3-channel signal decrease in surviving DA neurons on the ipsilateral to the injection side. (L) Histogram showing intensity of Kv4.3 immunosignals for all TH-positive ROIs, from ipsilateral, lesioned, side (in red) and from contralateral side (in grey). Inset, same data shown as a cumulative distribution. Note a clear right-shift to lower intensities for the ipsilateral side. (M) Comparison of mean TH-colocalized Kv4.3 immunosignals from medial SN from ipsilateral, lesioned, side, and contralateral, as control side. All data are presented as mean ± standard error of mean (SEM).

Discussion

Our study provides the first combined *in vitro* & *in vivo* electrophysiological characterization of identified DA neurons in the substantia nigra, surviving a partial 6-OHDA lesion. Studying these surviving SN DA neurons at two time points, we discovered time-dependent post-6-OHDA selective differences of firing properties both *in vivo* and *in vitro*. Early after lesion and coinciding with prominent behavioral impairments in our model, we detected a selective and dramatic (about 10-fold) reduction of *in vivo* burst firing in identified surviving SN DA neurons. This is reminiscent of an early study by Hollermann & Grace (Hollerman and Grace, 1990), who reported reduced bursting in putative SN DA neurons one week after partial lesion. At the early time point, we also found unstable *in vitro* pacemaker activity in these cells, while the mean firing rates were not different compared to those from vehicle-infused controls. In contrast, late (>2 months) after lesion, where substantial behavioral recovery had occurred, we detected no differences in *in vivo* firing rates and patterns – including burst discharges – of surviving SN DA neurons between 6-OHDA- and vehicle-infused mice. Surprisingly, the *in vitro* pacemaker rate was almost twice as fast in post-6-OHDA DA survivors compared to controls. Finally, we identified Kv4.3 channel downregulation as the cause for this chronic post-lesional pacemaker acceleration. In essence, we revealed for the first time the homeostatic adaptations of electrophysiological properties in surviving SN DA neurons.

Regarding the functional phenotype of surviving SN DA neurons, we identified a sequence of early but transient impairment, followed by chronic functional adaptation. While we did not investigate the mechanisms underlying the early impairment of *in vivo* burst firing, we focused our in-depth analysis on the chronic phase. Here, we identified a slowly-developing chronic Kv4.3 channel-mediated mechanism of pacemaker adaptation. Previously, we had already identified Kv4.3 channels in SN DA neurons as a pathophysiological target in a transgenic mutant (A53T-SNCA) α -synuclein mouse model (without cell loss) (Subramaniam *et al.*, 2014a). In this model, we found a pacemaker acceleration caused by oxidative impairment of Kv4.3 channels. In contrast to the post-lesional SN DA neurons in

this study, the activity of SN DA neurons in the α -synuclein model was accelerated also *in vivo*. Thus, α -synuclein pathology causes an allostatic shift of the *in vivo* firing rate setpoint of SN DA neurons, while post-lesional SN DA neurons recover their *in vivo* firing setpoint by homeostatic pacemaker plasticity. Interestingly, both models converge on the pacemaker potassium channel Kv4.3 but in a different fashion. While Kv4.3 subunits were downregulated on protein level in post-lesional SN DA neurons, mutant α -synuclein induced oxidative Kv4.3 dysfunction as well as protein upregulation (Subramaniam *et al.*, 2014a). We are aware that neither of the mouse models fully capture the pathophysiology of PD, where α -synuclein pathology, cell loss, inflammation and compensatory mechanisms are expected to occur side by side (Surmeier *et al.*, 2017b). At least at the transcriptional level, we know that Kv4.3 mRNA is upregulated in human SN DA neurons from PD patients (Dragicevic *et al.*, 2015; Schiemann *et al.*, 2012). This is similar to α -synuclein models and opposite to post-lesional Kv4.3 downregulation, shown here, suggesting that Kv4.3 channel mediated compensatory mechanisms are finally overwhelmed during PD progression. Moreover, the results of this study imply that α -synuclein pathology interferes with the homeostatic capacity of surviving SN DA neurons, in particular the regulation of Kv4.3 channels. Indeed, we recently showed a Kv4.3 gain-of-function phenotype for vagal motoneurons in response to mutant SNCA expression (Chiu *et al.*, 2021). Thus, Kv4.3 channels – a major “brake” of the pacemaker rate (Khaliq and Bean, 2008; Liss *et al.*, 2001; Tarfa *et al.*, 2017) – is an emerging downstream target across several PD models. It would be interesting to further investigate functional and molecular regulation of this potassium channel in chronic and progressive PD models, where pathology and cell loss go hand in hand.

In addition to cell-autonomous adaptations in surviving SN DA neurons, we assume that their chronic *in vivo* electrophysiological phenotype is also shaped by network level plasticity in response to dopamine-depletion. Our finding that the mean *in vivo* discharge rates were not different to controls in the presence of an accelerated intrinsic pacemaker strongly suggests a shift of the synaptic excitation-inhibition (E-I) balance towards more inhibition. Numerous studies have found altered synaptic inhibition in the dopamine-depleted basal ganglia (recently reviewed in (Zhang *et al.*, 2021)). In particular, Heo and colleagues recently demonstrated a chronic E-I balance shift toward more

inhibition across several PD models including 6-OHDA (Heo et al., 2020). They identified a substantial contribution of additional GABA synthesis and release from reactive astrocytes in the midbrain. It has also been well-established that basal ganglia GABA neurons fire in more synchronized hence more effective fashion after dopamine depletion (Cagnan et al., 2015; Evans et al., 2020; Milosevic et al., 2018; Phillips et al., 2020; Tinkhauser et al., 2020; Wichmann et al., 2018). This has recently been confirmed by elegant *in vivo* single cell resolved studies using either gCAMP-based calcium monitoring or *in vivo* patch-clamp approaches (Ketzeff et al., 2017; Kravitz et al., 2010; Parker et al., 2018; Parker et al., 2016; Sitzia et al., 2020). The chronic acceleration of pacemaker rate in response to enhanced inhibition would also be expected within the framework of homeostatic plasticity (Turrigiano, 2012). However, the mechanisms of firing rate set-point control, recently elucidated for hippocampal neurons (Styr et al., 2019), is still unknown for SN DA neurons.

Our study has several limitations. As already discussed, we explored a single-hit partial lesion model, while PD is a chronic progressive neurodegenerative process. While we demonstrate that electrophysiological phenotypes per se are flexible and change in response to lesion, we also have to consider distinct DA subpopulations within the SN. We recently showed that lateral SN DA neurons projecting to DLS, the most vulnerable DA subpopulation (corresponding to the ventro-lateral SN DA neurons in humans (Duzel et al., 2009; McCutcheon et al., 2019)) possess a distinct *in vivo* firing phenotype in control mice (Farassat et al., 2019). Therefore, recording from this population in PD models would be highly relevant. However, in our current partial 6-OHDA lesion model, these lateral SN DA neurons were too severely lost to allow for more than anecdotal post-lesion electrophysiological analysis (see Fig. 3). Here, a milder version of the current model would be needed to identify their homeostatic plasticity. Regarding the surviving DA neuron in the medial SN, we previously showed that they have distinct projection targets including DLS, DMS and lateral shell of nucleus accumbens (Farassat et al., 2019). In the current study, we have not aimed for identification of axonal projections to avoid potentially confounding effects of additional brain surgery and to not further complicate already challenging experiments. Nevertheless, this important issue should be

addressed in follow-up studies by e.g. molecular subtyping approaches (Heymann et al., 2020; Poulin et al., 2018; Poulin et al., 2020; Saunders et al., 2018).

Finally, we would like to speculate about the possible implications of our findings for PD. In principle, homeostatic responses, as identified here, could have a dual nature. On one hand, they might render the neuron more robust, thereby slowing the rate of cell loss. On the other hand, the homeostatic adaptations may amplify innate vulnerability and accelerate disease progression. The latter would be a variant of the “stressful pacemaker” hypothesis of DA vulnerability (Chan et al., 2007; Surmeier, 2007; Surmeier et al., 2017a) and its potential clinical implications (Guzman et al., 2018; Liss and Striessnig, 2019; Ortner, 2021; Ortner et al., 2017; Parkinson Study Group, 2020). In summary, we provide the first steps towards defining the homeostatic repertoire of surviving DA neurons, but deeper insights are necessary to rationally design neuroprotective interventions in PD.

Methods

Animals

Male C57Bl6/N mice (Charles River Laboratories) were used for the study. Mice were 8 weeks old, group housed and maintained on a 12-hour light-dark cycle. All experiments and procedures involving mice were approved by the German Regierungspräsidium Darmstadt (V54-19c20/15-F40/30). In total, 115 mice were used for this study (see table below).

	Early phase		Late phase	
	Mice (#)	Cells (#)	Mice (#)	Cells (#)
<i>In vivo</i> juxta & behavior – vehicle	15	12 – medial/ 28 – all	12	10 – medial/ 26 – all
<i>In vivo</i> juxta & behavior – 6-OHDA	16	9 – medial/ 16 – all	10	9 – medial/ 11 – all
<i>In vitro</i> – vehicle	3	16	2	9
<i>In vitro</i> – 6-OHDA	6	37	5	25
<i>In vitro</i> & AmmTx3 – vehicle	-	-	3	15
<i>In vitro</i> & AmmTx3 – 6-OHDA	-	-	3	14
	Mice (#)		Mice (#)	
Kv4.3 Immunocytochemistry	8		8	
SN quantification – vehicle	9		3	
SN quantification – 6-OHDA	9		3	
Sum = 115				

Stereotactic 6-OHDA infusion

All surgeries were performed under general anesthesia in areflexic state. Prior to the induction of anesthesia, a premedication of 0.2 mg/kg atropine (atropine-sulfate, Braun Melsungen AG, Melsungen) was given as an intraperitoneal (i.p.) injection to stabilize circulation. Anesthesia was induced in a plastic chamber, which was flooded with 5% Isoflurane (Florene®, AbbVie Deutschland GmbH & Co. KG, Ludwigshafen, Germany) in pure oxygen (0.4 l/min). For maintenance of anesthesia, isoflurane was delivered through a breathing mask with a flow rate of 0.35 l/min and its concentration was regulated to 1.5-2.2% using an adjustable vaporizer (Uno, Zevenaar, Netherlands). The depth of anesthesia was controlled by testing the toe pinch reflex and the breathing rate (1-2Hz). Body temperature (36°C) and respiration were constantly monitored. Lidocaine/prilocaine ointment (25

mg/g, Emla® creme, AstraZeneca GmbH, 22876 Wedel) was applied prior to surgery and after suturing of the wound as local anesthetics. Additional analgesia was provided by subcutaneous injection of carprofen (4 mg/kg in NaCl, 1:100, Rimadyl®, Pflizer GmbH, Berlin, Germany) after infusion. Eye lubricant (Visidic, Bausch and Lomb, Berlin, Germany) was used to protect eyes from desiccation.

Desipramine hydrochloride (20 mg/kg, Sigma Aldrich) was injected i.p. 20-40 min before intracranial infusions to prevent 6-OHDA uptake by noradrenergic neurons. The desipramine solution was prepared in sterile, isotonic NaCl solution (B. Braun Melsungen AG, Germany) at the day of surgery. The infusion solutions are based on sterile artificial cerebrospinal fluid (ACSF, Harvard Apparatus, Holliston, MA, USA) with 0.2% L-ascorbic acid (used also as a vehicle solution). The 6-OHDA solution (0.02% 6-hydroxydopamine hydrochloride in ACSF with 0.2% L-ascorbic acid) was prepared at the day of infusion, stored on ice, and shielded from light.

During surgery, the animals were placed on a heating pad and were fixed in a stereotactic frame (Model 1900, Kopf Instruments, Tujunga, USA) with a stereotactic arm and a connected three-way digital positioning display. The scalp was opened by a longitudinal cut to expose the skull with bregma and lambda on display. With a centering scope (Model 1915, Kopf Instruments, Tujunga, USA), the bregma-lambda distance was measured and examined for suitable anatomy (4.4±0.2 mm distance). Afterwards, the skull was aligned to a reference frame with a stereotaxic alignment indicator (Model 1905, Kopf Instruments, Tujunga, USA) and the manipulator system was referenced to bregma.

Using a stereotaxic drill (Model 1911, Kopf Instruments, Tujunga, USA) with a 500 µm diameter drill bit, a hole above the right striatum was drilled (coordinates: ML: +1.9 mm, AP: +0.5 mm to bregma). ACSF or 6-OHDA solution were loaded to a 10 µl NanoFil syringe (World Precision Instruments Inc., Sarasota, FL, USA) with a 35G blunt needle, which was mounted on a MicroSyringe Pump (UMP3-1, World Precision Instruments) and controlled by a SYSMicro4 Controller (World Precision Instruments). Using the stereotactic arm, the needle was slowly lowered (about 750 µm/min) to a position of -2.2 mm below the brain surface (infusion site coordinates: ML: +1.9 mm, AP: +0.5 mm, DV: -2.2 mm to

bregma). Anatomical references are based on Franklin and Paxinos (2008). A volume of 6 μ l was infused with a flow rate of 250 nl/min. Once the volume was infused, the needle rested for 5 minutes in that position before it was slowly moved out of the brain. Directly before and after infusion, proper functioning of the syringe system and the needle was checked. Finally, after suture the animal was placed on a heating pad for full recovery. Oats, wet food pellets and water were placed inside the cage to ease consumption.

Behavioral testing

Open field

Spontaneous locomotion (track length, wall distance, time in center and number of rearings) and rotations of all mice were monitored in open field (50 $\times$ 50 cm, center 30 $\times$ 30 cm; red illumination, 3 lx) for 10 min in 3 baseline sessions and every 4th or 7th day post-infusion of ACSF/6-OHDA till the day of *in vivo* or *in vitro* experiment (e.g. 21st or >68th post-operative day). The open field was cleaned before and after each mouse with 0,1% acetic acid in distilled water. Using a video tracking system (Viewer II/III, Biobserve) spontaneous behavior was recorded and analyzed both online and offline. Data was extracted from Viewer as Excel-tables and the final analysis was made with custom made Matlab-scripts.

Cylinder test

Forelimb use during explorative activity was explored with cylinder test. The test was performed at corresponding termination time point (20-21st and 64th post-infusion day). Mice were placed individually in a glass beaker (9 cm diameter, 19 cm height) at room light and were video recorded with a camera (Logitech HD Webcam C615) for about 5 min. No habituation was allowed before video recording. The glass cylinder was cleaned before and after every mouse with 0,1% acetic acid in distilled water. Only weight-bearing wall contacts made by each and both forelimb on the cylinder

wall were scored. Wall exploration was expressed in terms of the percentage of contralateral to the infusion side (in the 6-OHDA-infused mice also impaired forepaw) to all forelimb wall contacts.

In vivo electrophysiology

Extracellular recording

In vivo extracellular single-unit activities of SN and VTA neurons were recorded in ACSF-infused (vehicle) and 6-OHDA-infused mice, similar procedures were used in other studies from our lab (Farassat *et al.*, 2019; Schiemann *et al.*, 2012; Subramaniam *et al.*, 2014a). Briefly, mice were anesthetized (isoflurane; induction 4.5-5%, maintenance 1–2% in 0.4 l/min O₂) and placed into a stereotactic frame. The craniotomies were performed as described above to target the lateral SN (AP: -3.08 mm, ML: 1.4 mm) and medial SN (AP: -3.08 mm, ML: 0.9 mm). Borosilicate glass electrodes (10–25 MΩ; Harvard Apparatus, Holliston, MA, USA) were made using a horizontal puller (DMZ-Universal Puller, Zeitz, Germany) and filled with 0.5 M NaCl, 10 mM HEPES (pH 7.4) and 1.5% neurobiotin (Vector Laboratories, Burlingame, CA, USA). A micromanipulator (SM-6; Luigs & Neumann, Ratingen, Germany) was used to lower the electrodes to the recording site. The single-unit activity of each neuron was recorded for at least 10 minutes at a sampling rate of 12.5 kHz (for firing pattern analyses), and then for another one minute at a sampling rate of 20 kHz (for the fine analysis of AP waveforms). Signals were amplified 1000x (ELC-03M; NPI electronics, Tamm, Germany), notch- and bandpass-filtered 0.3–5000 Hz (single-pole, 6 dB/octave, DPA-2FS, NPI electronics) and recorded on a computer with an EPC-10 A/D converter (Heka, Lambrecht, Germany). Simultaneously, the signals were displayed on an analog oscilloscope and an audio monitor (HAMEG Instruments CombiScope HM1508; AUDIS-03/12M NPI electronics). Midbrain DA neurons were initially identified by their broad biphasic AP (> 1.2 ms duration) and slow frequency (1–8 Hz) (Grace and Bunney, 1984a; b; Ungless and Grace, 2012). AP duration was determined as the interval between the start of initial upward component and the minimum of following downward component.

568

569 *Juxtacellular labeling of single neurons*

570 In order to identify the anatomical location and neurochemical identity of the recorded neurons, they
571 were labeled post-recording with neurobiotin using the juxtacellular *in vivo* labeling technique
572 (Pinault, 1996). Microiontophoretic currents were applied (1–10 nA positive current, 200ms on/off
573 pulse, ELC-03M, NPI Electronics) via the recording electrode in parallel to the monitoring of single-unit
574 activity. Labeling was considered successful, when: the firing pattern of the neuron was modulated
575 during current injection (i.e., increased activity during on-pulse and absence of activity in the off-
576 pulse), the process was stable for at least 20s, and was followed by the recovery of spontaneous
577 activity. This procedure allowed for the exact mapping of the recorded DA neuron within the SN and
578 VTA subnuclei (Franklin and Paxinos, 2008) using custom written scripts in Matlab (MathWorks,
579 Natick, MA, USA), combined with neurochemical identification using TH-immunostaining.

580

581 *In vitro* electrophysiology

582 *Slice preparation*

583 Animals were anesthetized by intraperitoneal injection of ketamine (250 mg/kg, Ketaset, Zoetis) and
584 medetomidine-hydrochloride (2.5 mg/kg, Domitor, OrionPharma) prior to intracardial perfusion using
585 ice-cold ACSF consisting of the following (in mM): 50 sucrose, 125 NaCl, 2.5 KCl, 25 NaHCO₃, 1.25
586 NaH₂PO₄, 2.5 glucose, 6 MgCl₂, 0.1 CaCl₂ and 2.96 kynurenic acid (Sigma-Aldrich), oxygenated with
587 95% O₂ and 5% CO₂. Rostral coronal midbrain slices (bregma: -2.92 mm to -3.16 mm) were sectioned
588 at 250 µm using a vibrating blade microtome (VT1200s, Leica). Slices were incubated for 1 h before
589 recordings in a 37°C bath with oxygenated extracellular solution with extra 1µM AmmTx3, containing
590 the following (in mM): 22.5 sucrose, 125 NaCl, 3.5 KCl, 25 NaHCO₃, 1.25 NaH₂PO₄, 2.5 glucose, 1.2
591 MgCl₂ and 1.2 CaCl₂.

592

In vitro patch-clamp recordings

Slices were placed in a heated recording chamber (37°C) that was perfused with oxygenated extracellular solution at 2-4 ml min⁻¹. CNQX (20 µM; Biotrend), gabazine (SR95531, 4 µM; Biotrend) and DL-AP5 (10 µM; Cayman Chemical) were added to inhibit excitatory and inhibitory synaptic transmission. Neurons were visualized using infrared differential interference contrast videomicroscopy with a digital camera (VX55, Till Photonics) connected to an upright microscope (Axioskop 2, FSplus, Zeiss). Patch pipettes were pulled from borosilicate glass (GC150TF-10; Harvard Apparatus LTD) using a temperature-controlled, horizontal pipette puller (DMZ-Universal Puller, Zeitz). Patch pipettes (4-6 MΩ) were filled with a solution containing the following (in mM): 135 KClu, 5 KCl, 10 HEPES, 0.1 EGTA, 5 MgCl₂, 0.075 CaCl₂, 5 NaATP, 1 LiGTP, 0.1% neurobiotin, adjusted to a pH of 7.35 with KOH. Recordings were performed using an EPC-10 patch-clamp amplifier (Heka electronics) with a sampling rate of 20 kHz and a low-pass filter (Bessel, 5 kHz). For analysis, action potential thresholds (mV) were determined at $dV_m/dt > 10$ mV/ms.

Immunohistochemistry

Following *in vivo* recordings, animals were transcardially perfused, as described previously (Farassat *et al.*, 2019; Schiemann *et al.*, 2012; Subramaniam *et al.*, 2014a). Fixed brains were sectioned into 60µm (midbrain) or 100µm (forebrain) coronal sections using a vibrating microtome (VT1000S, Leica). *In vitro* slices were fixed in paraformaldehyde after finishing the experiment. Sections were rinsed in PBS and then incubated (in blocking solution (0.2 M PBS with 10% horse serum, 0.5% Triton X-100, 0.2% BSA). Afterwards, sections were incubated in carrier solution (room temperature, overnight) with the following primary antibodies: polyclonal guinea pig anti-tyrosine hydroxylase (TH; 1:1000; Synaptic Systems), monoclonal mouse anti-TH (1:1000; Synaptic Systems) or polyclonal rabbit anti-TH (1:1000; Synaptic Systems); mouse anti-Kv4.3 (1:1000, Alomone Labs). In sequence, sections were again washed in PBS and incubated (room temperature, overnight) with the following secondary antibodies: goat anti-guinea pig 488, goat anti-rabbit 488, goat anti-mouse 488, goat anti-mouse 568

(all 1:750; Thermofisher). Streptavidin AlexaFluor-568 and Streptavidin AlexaFluor-488 (both 1:1000; Invitrogen) were used for identifying neurobiotin-filled cells. Sections were then rinsed in PBS and mounted on slides with fluorescent mounting medium (Vectashield, Vector Laboratories).

DAB immunocytochemistry

For DAB (3,3'-diaminobenzidine) staining procedures, a Vectastain ABC Staining Kit (Vector Laboratories) was used. Coronal sections of midbrain (30 μ m) areas were cut and rinsed in PBS (3x10 min). Similar to previous immunolabeling procedures, unspecific antigen binding sites were blocked by incubation of the sections with blocking solution (60 min, room temperature). Subsequently, sections were incubated with primary antibody against TH (rabbit anti TH) overnight, rinsed in PBS (3x10 min), and were incubated with biotinylated secondary antibodies (biotinylated anti-rabbit) for two hours at RT. In parallel, an avidin-biotin complex (ABC) was formed by pre-incubation of avidin (1:1000) with biotinylated HRP (1:1000) in PBS for two hours at room temperature. Sections were rinsed in PBS (3x10 min) prior to incubation with ABC solution (60 min, room temperature). Next, sections were rinsed in PBS (2x10 min) and Tris-buffer (1x10 min). Finally, DAB oxidation was developed by application of 2 % H₂O₂, 2 % NiCl₂ and 4 % DAB in Tris-buffer using a DAB Substrate Kit (Vector Laboratories, Burlingame, USA). NiCl₂ enhances sensitivity and intensity of DAB precipitation product. DAB oxidation was developed for 2 to 5 minutes and was stopped with Tris-buffer once a specific high-contrast signal was detectable. Sections were rinsed in Tris-buffer (3x10 min) and transferred onto gelatin-covered slides, air-dried overnight, and dehydrated in consecutive ascending alcohol concentrations (50 %, 70 %, 90 % and 2x 100 %; 10 min each) followed by dehydration in xylol (2x 100 %; 10 min each). Finally, sections were mounted under glass coverslips with hardening mounting medium (Vectamount, Vector Laboratories, Burlingame, USA).

Unbiased stereology measurements

For quantification of total cell loss, TH-DAB labeled SN DA neurons were counted using unbiased stereology based on optical dissection (Gundersen, 1986). In coronal sections (30 μ m), the region of interest was selected based on anatomical landmarks including the medial lemniscus, which separates SN and adjacent VTA. Stereological counting provides unbiased data based on random, systematic sampling using an optical fractionator. This method involves counting of neurons with an optical dissector, a three-dimensional probe placed through a reference space (Gundersen, 1986). The optical dissector forms a focal plane with a thin depth of field through the selected sections. Objects in focus of this focal plane are located within the reference section and are counted, while objects outside of the focal plane are not counted. On top of the optical dissector, a counting frame is applied. Counting frames ensure that all neurons have equal probabilities of being selected, regardless of shape, size, orientation, and distribution. To avoid counting of capped neurons at the border of a section, an additional guard zone was deployed at the upper and lower borders of each section. DA neurons within the counting frame as well as those crossing the green line (acceptance line) were counted, while DA neurons crossing the red line (rejection line) excluded. Moreover, only neurons with a detectable nucleus in focus within the optical dissector were counted. For quantification of total cell loss, StereoInvestigator software (V5, MicroBrightField, Colchester, USA) was used in combination with BX61 microscope (Olympus, Hamburg, Germany). The region of interest was selected and marked using a low magnification objective lens (2x, NA 0.25, Olympus) and 12-30 serial sections of 30 μ m thickness were counted, covering the entire caudo-rostral extent of the SN. To count the number of DA neurons in the area of the SN pars compacta, a high magnification oil-immersion objective lens (100x, NA 1.30, Olympus) was used.

Optical density measurements

Optical density measurements of TH-DAB labeled striatal sections were performed using ImageJ software (<http://rsbweb.nih.gov/ij/>). Following TH-DAB labeling and TH-immunohistochemistry,

images of five coronal sections (100 μ m) covering the rostrocaudal axis of the striatum were captured using laser-scanning microscope (Nikon Eclipse90i, Nikon GmbH). Images were gray scale converted and mean gray values of desired striatal areas were encircled and measured. Unspecific mean gray values were measured in a defined cortical area (100x100 pixels) that displayed no specific TH signal due to the absence of DA innervation and were subtracted. The ventral edge of lateral ventricles served as an anatomical landmark to separate dorsal and ventral areas. For all animals, the measurement from the ipsilateral to the infusion side were divided by the contralateral side to calculate the relative optical density of the striatum.

Immunohistochemical Kv4.3 channel signal quantification

A Nikon Eclipse 90i microscope was used for fluorescent signal detection, excitation wavelength of 488 nm for TH-signal and 568 nm for Kv4.3-channel signal. From each animal, 4 midbrain slices covering the caudal, intermedial and rostral regions, were selected and imaged for overview with 4x magnification. Then 60x magnification was used to acquire data from 4 areas within each SN (4 images on the ipsilateral and 4 images on the contralateral to infusion side). All images were acquired using the same laser and camera settings. Images were exported from Nikon NIS-Elements Advanced Research (Version 4.20.03) software as 8-bit TIFF files for quantification. Data was analyzed using custom made Python 3.0 scripts with matplotlib, numpy, scimage and scipy modules. First, TH immunosignals were converted to a binary image via Otsu-thresholding algorithm. Then, the resulting binary image was used as a mask for Kv4.3 channel immunosignal detection. For all ROIs surface areas and mean Kv.4.3 channel signal intensity was measured. By applying erosion and dilatation algorithms on the ROIs, membrane and cytoplasm areas were segregated, allowing isolation of Kv4.3 channel immunosignal intensity for these cell compartments. Background Kv4.3 channel immunosignal was quantified in TH-immunosignal areas below the Otsu-threshold. All data were then grouped according to medio-lateral and ipsi-/contralateral position for both vehicle and 6-OHDA group. Graphs and statistical analysis for this data was performed using Python custom made scripts.

695

696 Statistical Analysis

697 *Spike train analyses*

698 Spike time-stamps were extracted by thresholding above noise levels with IgorPro 6.02 (WaveMetrics,
699 Lake Oswego, OR, USA). Firing pattern properties such as mean frequency, coefficient of variation (CV)
700 and bursting measures were analyzed using custom scripts in Matlab. In order to estimate burstiness
701 and intra-burst properties, we used the burst detection methods described in Grace & Bunney (Grace
702 and Bunney, 1984a; b). All non-burst related ISIs (excluding all ISIs that followed the Grace and Bunney
703 criteria, as well as all pre- and post-burst ISIs) were used to calculate the single spike firing frequency
704 and single spike coefficient of variation.

705 For analysis of general firing patterns, autocorrelation histograms (ACH) were plotted using custom
706 Matlab scripts. We used established criteria for classification of *in vivo* firing patterns based on visual
707 inspection of autocorrelograms (Farassat *et al.*, 2019; Schiemann *et al.*, 2012; Subramaniam *et al.*,
708 2014a): single spike-oscillatory (≥ 3 equidistant peaks with decreasing amplitudes), single spike-
709 irregular (< 3 peaks, increasing from zero approximating a steady state), bursty-irregular (narrow peak
710 with steep increase at short ISIs) and bursty-oscillatory (narrow peak reflecting fast intraburst ISIs
711 followed by clear trough and repetitive broader peaks).

712 Statistics

713 Categorical data is represented as stacked bar graphs. To investigate the assumption of normal
714 distribution, we performed the single-sample Kolmogorov-Smirnov test. The Mann-Whitney-Test,
715 one-/two-way ANOVA were performed in non-parametric data to determine statistical significance.
716 Categorical parameters, such as ACH-based firing pattern, were analyzed with the Chi-squared test.
717 Statistical significance level was set to $p < 0.05$. All data values are presented as means $\pm$ SEM.
718 Statistical tests were performed using GraphPad Prism 9 (GraphPad Software, San Diego, CA, USA),
719 Matlab and Python. The scatter plots are represented with median or mean $\pm$ SEM. The resulting p

values were compared with Bonferroni-corrected α -level or Tukey *post hoc* comparison. A value of $p \leq 0.05$ was considered to be statistically significant; $p \leq 0.05 = *p \leq 0.005 = **p \leq 0.0005 = ***$. Graphs were plotted using GraphPad Prism software (9.0c), Matlab and Python.

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Additional information

Author contributions

LK & JR designed the study. LK performed the lesions, behavioral & *in vivo* electrophysiology, JS performed the *in vitro* experiments. Analysis was carried out jointly by LK, JS, NF and JR. NF taught LK *in vivo* electrophysiology and juxtacellular labelling. LK & JR wrote the manuscript.

Declaration of Interests

The authors declare no conflict of interests.

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Ethics

Animal experimentation: All experiments and procedures involving mice were approved by the German Regierungspräsidium Darmstadt (V54-19c20/15-F40/30).

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