The density of calretinin striatal interneurons is decreased in 6-OHDA-lesioned mice

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Abstract
Interneurons play a significant role in the functional organization of the striatum and some of them display marked plastic changes in dopamine-depleted conditions. Here, we applied immunohistochemistry on brain sections from 6-hydroxydopamine (6-OHDA) mouse model of Parkinson’s disease and sham animals to characterize the regional distribution and the morphological and neurochemical changes of striatal interneurons expressing the calcium-binding protein calretinin (CR).

Two morphological subtypes of calretinin-immunostained (CR +) interneurons referred, respectively, as small- and medium-sized CR + interneurons were detected in 6-OHDA- and sham-lesioned animals. The small cells (9–12 µm) prevail in the anterior and dorsal striatal regions; they stain intensely for CR and display a single slightly varicose and moderately arborized process. The medium-sized CR + interneurons (15–20 µm) are more numerous than the small CR + cells and rather uniformly distributed within the striatum; they stain weakly for CR and display 2–3 long, slightly varicose and poorly branched dendrites. The density of medium CR + interneurons is significantly decreased in the dopamine-depleted striatum (158 ± 15 neurons/mm³), when compared to sham animals (370 ± 41 neurons/mm³), whereas that of the small-sized CR + interneurons is unchanged (174 ± 46 neurons/mm³ in 6-OHDA-lesioned striatum and 164 ± 22 neurons/mm³ in sham-lesioned striatum). The nucleus accumbens is populated only by medium-sized CR + interneurons, which are distributed equally among the core and shell compartments and whose density is unaltered after dopamine denervation. Our results provide the first evidence that the medium-sized striatal interneurons expressing low level of CR are specifically targeted by dopamine denervation, while the small and intensely immunoreactive CR + cells remain unaffected. These findings suggest that high expression of the calcium-binding protein CR might protect striatal interneurons against an increase in intracellular calcium level that is believed to arise from altered glutamate corticostriatal transmission in Parkinson’s disease.

Keywords Basal ganglia · Parkinson’s disease · Dopamine · Calcium binding protein · Nucleus accumbens · Striatum

Introduction
The striatum is chiefly composed of γ-amino-butyric acid (GABAergic) spiny projection neurons, but it also comprises a small proportion of aspiny local circuit neurons (Graveland and DiFiglia 1985; Gerfen and Bolam 2010; Kawaguchi et al. 1995). Despite their relatively low number—it reportedly ranges from 2–3% of the total neuronal population in rodents (Rymar et al. 2004; Oorschot 2013) to 15–20% in non-human primates (Graveland and DiFiglia 1985)—the aspiny interneurons occupy a crucial position in the striatal microcircuitry (Kawaguchi et al. 1995; Tepper et al. 2018) and are involved in several motor and psychiatric disorders (Ding et al. 2011; Kataoka et al. 2010; Pisani et al. 2007). The role of interneurons in shaping and tuning striatal inputs and outputs is well recognized, but their exact function remains elusive, especially in pathological conditions.

In 6-OHDA-lesioned mice used as animal models of Parkinson’s disease, degeneration of nigrostriatal dopaminergic axons has been shown to lead to a significant reduction of dendritic arborization and spine density of striatal projection neurons (Gagnon et al. 2017; Fieblinger et al. 2014; Suarez et al. 2014; Gomez et al. 2019). The number and chemical makeup of various types of striatal interneurons have also been shown to be altered following degeneration of
dopamine afferent projections in humans (Huot et al. 2007), non-human primates (Petryszyn et al. 2016) and rodents (Unal et al. 2015; Ma et al. 2014; Mura et al. 2000). These changes are thought to result from altered striatal activity known to be regulated by glutamate and dopamine neurotransmission, as evidenced by the increase in corticostriatal excitatory input that follows striatal dopamine depletion in Parkinson’s disease, a condition that may lead to a neurotoxic raise of intracellular calcium levels (Blandini et al. 1996). In that context, calcium-binding proteins expressed by striatal cells, including calbindin, parvalbumin and calretinin (CR), are believed to play an important role by maintaining calcium homeostasis and exerting protective effect against increases in intracellular calcium level (Choi 2005; Mitchell et al. 1994; Lee et al. 2002; Figueredo-Cardenas et al. 1998). A similar phenomenon occurs at substantia nigra level where calbindin expressed by midbrain dopamine cells has been shown to be neuroprotective (German et al. 1992; Inoue et al. 2019).

The present study focuses on the fate of striatal interneurons that express CR, a calcium-binding protein of the “EF-hand” family, in mice that underwent a lesion of the nigrostriatal dopamine pathway. The striatum of mice harbors two morphological subtypes of aspiny calretinin-immunostained (CR+) interneurons (Petryszyn et al. 2014). Neurons of the first type stain intensely for CR, are endowed with a small cell body (9–12 µm) and a slightly varicose and moderately arborized process, and prevail in the anterior and dorsal sectors of the striatum. Neurons of the second type stain moderately for CR but outnumber those of the first type; they have a medium-sized cell body (15–20 µm) with 2–3 long, slightly varicose, poorly branched dendrites and are uniformly scattered throughout the striatum, but slightly more abundant in its ventral sector. It should be mentioned that, based on neurochemical makeup, the existence of three morphological subtypes of CR+ cells has recently been reported in the mouse striatum, each showing different size of cell bodies (Garas et al. 2018).

Previous studies in rats in which the nigrostriatal dopaminergic pathway has been lesioned by means of the neurotoxin 6-OHDA led to contradictory findings about the fate of the CR+ striatal interneurons. Some studies reported a significant but transitory increase in the number of striatal CR+ neurons (Mura et al. 2000), but other investigations detected a permanent decrease of the same neuronal elements (Ma et al. 2014). Despite the fact that mice have become a widely used animal model to study various neurodegenerative conditions, including Parkinson’s disease, no data are currently available on the effect of striatal dopamine depletion on CR+ striatal interneurons in any mouse model of Parkinson’s disease. In the light of the paucity and conflicting nature of the data obtained in rats and because no information are currently available in mice, we undertook a detailed stereological study to determine the fate of the morphological subtypes of striatal CR+ interneurons in unilateral 6-OHDA-lesioned mice, a widely used animal model of Parkinson’s disease.

Materials and methods

Animals

Brains were obtained from 8 C57BL/6 male mice of 3 months old (Mus musculus; Charles River, QC, Canada). Animals were housed in a temperature-controlled room (21–25 °C) under a 12-h light/dark cycle and had free access to food and water. All protocols were approved by the Institutional Animal Care and Use Committee (Comité de Protection des Animaux de l’Université Laval, #2019-287/ VRR-18-107) and all procedures involving animals and their care were made in accordance with the Canadian Council on Animal Care’s Guide to the Care and Use of Experimental Animals (Ed2).

6-OHDA unilateral injection

Four mice received a unilateral 6-OHDA injection in the right medial forebrain bundle (mfb) to lesion the dopaminergic striatal afferent projections. Thirty minutes before 6-OHDA injections, mice received an i.p. injection of desipramine (25 mg/Kg) diluted in saline (0.9%) at a concentration of 2 mg/mL. Mice were then anaesthetized using 2% isoflurane and their heads were fixed in a stereotaxic apparatus. A hole was drilled and the following stereotaxic coordinates relative to bregma were aimed: anteroposterior = − 1.2 mm; mediolateral = 1.1 mm; dorsoventral = − 5.0 mm, corresponding to the mfb, according to the mouse brain atlas of Franklin and Paxinos (1997). A glass micropipette containing a freshly prepared solution of 6-OHDA diluted in ascorbic acid (0.02%) at a concentration of 6 µg/µL was slowly introduced in the mfb. The 6-OHDA was then pressure injected (0.125 µL/min, during 2 min). Following injection, the micropipette was left in place for 2 min before being slowly retracted. A total volume of 0.25 µL of 6-OHDA was injected into the mfb, corresponding to 1.5 µg of 6-OHDA. Four other mice served as the sham group: each animal received a unilateral injection of 0.02% ascorbic acid (vehicle), according to the procedure described above. After the surgery, the skin was sutured and the eight mice were allowed to recover.

Behavioral assessment

The severity of the dopamine lesion was assessed 20 days after 6-OHDA or vehicle unilateral injections. Mice from
the two experimental groups were introduced in a large glass cylinder and spontaneous motor behavior was recorded for 10 min, using a digital camera. The number of spontaneous rotations ipsilateral and contralateral to the injection side were then counted post hoc, by an experimenter blinded to the experimental treatment.

**Tissue preparation**

Forty days following 6-OHDA or vehicle stereotaxic injections, animals were deeply anesthetized with a mixture of ketamine and xylazine (100 mg/kg, 10 mg/kg, i.p.), and perfused transcardially with a wash of 0.9% saline (50 mL), followed by 2% acroleine diluted in phosphate buffer (PB, 100 mL) and by 4% paraformaldehyde (PFA; 100 mL, diluted in PB). Brains were dissected out, post-fixed for 1 h in a 4% PFA solution and cut with a vibratome (Leica VT1200S) into 50 µm thick coronal sections, which were serially collected in sodium phosphate buffer saline (PBS, 0.1 M, pH 7.4).

**Immunostaining of TH and DAT**

For each mouse, two coronal sections were selected through the striatum at 1.42 and 0.14 mm relative to the bregma, and one section through the substantia nigra at −3.10 mm from the bregma (Franklin and Paxinos 1997). These sections were immunostained for tyrosine hydroxylase (TH) to assess to the DA lesion caused by 6-OHDA injection. Briefly, sections were incubated for 30 min into a 0.5% solution of sodium borohydride (NaBH₄) diluted in PBS then rinsed thoroughly in PBS. Sections were then pre-incubated for 1 h into a blocking solution composed of 0.1% Triton X-100, 2% of normal donkey serum diluted in PBS. They were then incubated overnight (ON) with a polyclonal rabbit antibody against TH (Catalog #AB152, Millipore, 1:1000) and a monoclonal antibody against DAT (Catalog #MAB369, Millipore, 1:1000). Then, sections were rinsed in PBS and incubated for 2 h in the blocking solution to which a secondary antibody IRDye 680 anti-rabbit (Catalog #LIC-926-32223, Mandel, 1:1000) raised in donkey and an IRDye 800 antibody IRDye 680 anti-rabbit (Catalog #LIC-926-32223, Mandel, 1:1000) raised in goat were added. Sections were then rinsed and mounted on gelatine-coated slides, air-dried, and coverslipped with Dako fluorescence mounting medium (Catalog #S-3023, Mississauga, ON, Canada).

**Immunostaining of CR**

To assess the regional distribution of CR+ interneurons in the striatum, 6 equally spaced coronal sections (bregma 1.30 to −1.70 mm, interval of 600 µm) were randomly selected throughout the entire striatum from the 4 6-OHDA-lesioned mice and the 4 sham-lesioned animals. These sections were singly labeled for the calcium-binding protein CR. Briefly, the 6 striatal sections per animal were pre-incubated for 1 h in a blocking solution composed of 0.1% Triton X-100 and 2% of normal donkey serum diluted in PBS. They were then incubated ON with an antibody against CR (Catalog #7699/4, Swant) diluted 1:500 in the blocking solution. Sections were then rinsed in PBS and incubated for 2 h with an anti-rabbit biotinylated secondary antibody (Catalog #711-075-152, Vector Laboratories) raised in donkey and diluted 1:1000 in the blocking solution. Sections were rinsed and incubated for 1 h with the avidin–biotin-peroxidase complex (Catalog #PK4000, Vector Laboratories). Sections were then rinsed once in PBS and twice in Tris-saline buffer (TBS; 50 mM, pH 7.4) and incubated for 3 min in a solution containing 0.05% of 3,3’ dianimobenzidine (Catalog #D5637, Sigma) and 0.005% H₂O₂ diluted in TBS to reveal the bound peroxidase. Finally, sections were rinsed in TBS to stop the reaction and mounted on gelatine-coated slides, air-dried overnight, dehydrated in alcohol grade series, cleared in toluene and coverslipped with Permount.

To assess quantitatively the DA lesion, two coronal sections per mouse were selected through the striatum (bregma = 0.14 mm) and the nucleus accumbens (NAc, bregma = 1.42 mm) and doubly stained for TH and the dopamine membrane transporter (DAT) with infrared immunofluorescence. Sections were first incubated for 30 min with 0.5% NaBH₄, then rinsed thoroughly in PBS. They were then pre-incubated for 1 h in 0.5% Triton X-100 in PBS containing 2% of normal goat serum and normal donkey serum (blocking solution). Sections were incubated ON in the blocking solution containing the same TH antibody as above (Catalog #AB152, Millipore, 1:1000) and a monoclonal antibody against DAT (Catalog #MAB369, Millipore, 1:1000). Then, sections were rinsed in PBS and incubated for 2 h in the blocking solution to which a secondary antibody IRDye 680 anti-rabbit (Catalog #LIC-926-32223, Mandel, 1:1000) raised in donkey and an IRDye 800 antibody IRDye 680 anti-rabbit (Catalog #LIC-926-32219, Mandel, 1:1000) raised in goat were added. Sections were then rinsed and mounted on gelatine-coated slides, air-dried, and coverslipped with Dako fluorescence mounting medium (Catalog #S-3023, Mississauga, ON, Canada).

**Double immunostaining of CR and CB**

To determine the regional distribution of CR+ interneurons in the shell and core compartments of the NAc, 3 more sections (bregma 1.30, 1.10 and 0.90 mm) were selected throughout the NAc from the 4 6-OHDA-lesioned mice and the 4 sham-lesioned animals. These sections were doubly stained for CR and calbindin-D-28K (CB). Briefly,
the same protocol described above was used, except that normal goat serum was added to the blocking solution as well as a monoclonal primary antibody against CB (Catalog #C9848, Sigma, 1:500). In addition, a secondary antibody against mouse and coupled to Alexa Fluor 488 and raised in goat was used at a dilution 1:200 (Catalog #A11001, MolecularProbes).

**Data analysis**

**Assessment of the dopamine lesion**

In each mouse, the dopamine lesion induced by 6-OHDA stereotaxic injections was assessed from sections of the striatum and the NAc stained for TH and DAT using the IRDye 680, and the IRDye 800, respectively, as described above. Sections were scanned using the Odyssey imager (LiCor biotechnology). For each mouse, regions of interest were randomly selected by drawing 6 boxes in the striatum and 6 boxes in the NAc (3 in the core and 3 in the shell) of 150 µm² on both the intact (left) and lesioned (right) side. Mean TH and DAT immunoreactivity values obtained from the lesioned striatum or NAc, divided by immunoreactivity measured from the intact side × 100 was used to report the residual immunoreactivity.

**Stereological counting of CR + cells**

In each mouse, the 6 equally spaced sections selected across the entire striatum (600 µm interval) were examined with a confocal Laser Scanning Microscope (LSM700, Zeiss) equipped with a camera (AxioCam), a motorized stage (X and Y axes) and a Z-axis indicator (Leica Z-axis control) and controlled by a computer running StereoInvestigator software. A detailed scanning of the regional distribution of CR + interneurons throughout the striatum was achieved by dividing the structure into 8 distinct sectors: antero-dorso-lateral (ADL), antero-dorso-median (ADM), antero-ventro-lateral (AVL), antero-ventro-median (AVM), postero-dorso-lateral (PDL), postero-dorso-median (PDM), postero-ventro-lateral (PVL) and postero-ventro-median (PVM). The contour of the striatum was first outlined on each CR-immunostained coronal section using a 4X/0.10 objective, according to the stereotaxic atlas of Franklin and Paxinos (1997). A vertical line parallel to the midline and passing by the center of the striatum was first traced, dividing the structure into medial and lateral sectors (Fig. 1A). A horizontal line, perpendicular to and centered on the vertical line, was added to delineate four sectors on each brain section. The anteroposterior axis was then divided in two by considering the first 3 coronal sections as representative of anterior (pre-commissural) sectors and the last 3 of posterior (post-commissural) sectors. The 3 sections (200 µm interval) taken through the NAc of each mouse and doubly stained for CR and CB were also analyzed stereologically, the CB immunostaining being used to delineate the core from the shell. The number of doubly stained cells encountered was counted.

The sampling process leading to the estimation of the total number of CR + interneurons in each of the striatal sectors and in the core and the shell of the NAc began by using a grid formed by 157 × 157 µm squares placed over each section. At each intersection of the grid that fell into the sector, a counting frame of the same size was drawn and examined with a 40X/1.4 objective leading to the examination of the entire area of each sector (Fig. 1B). On each selected section, a mean number of 48.9 ± 4.7 sampling sites in each striatal sector, 42.1 ± 4.2 in the shell and 39.4 ± 4.0 in the core, were...
examined. Neurons that fell inside the counting frame and did not contact the exclusion lines were counted whenever their nucleus came into focus within a 12 µm thick optical disector centered in the section (Fig. 1C). An average of 202 ± 29 CR + cells were counted in each striatum ipsilateral to the sham or 6-OHDA lesion whereas 181 ± 24 were counted in each striatum contralateral to the lesion, yielding coefficients of error (Gundersen, \(m = 1\) and 2nd Schmitz-Hof) ranging from 0.06 to 0.12 and from 0.06 to 0.14, respectively. In the NAc, an average of 103 ± 21 CR interneurons were counted ipsilateral to the sham or 6-OHDA lesion, while 99 ± 19 were counted contralateral to the lesion, yielding coefficients of error (Gundersen, \(m = 1\) and 2nd Schmitz-Hof) between 0.07 and 0.22, respectively. These coefficients of error are used to assess the precision of the estimation by taking into account the shape of the region of interest, the distribution of neurons within this region and the sampling criteria, predicting the accuracy of stereological procedure (Gundersen et al. 1999; Schmitz and Hof 2000). For each striatal and NAc region, the density of CR + cells was expressed in number of CR + neurons per mm³ of tissue, using the total number estimated by the optical disector and the volume of each sector, as estimated by Cavalieri’s method (García-Fiñana et al. 2003).

Quantification of mouse spontaneous motor behavior following surgery revealed a marked preferential rotational behavior towards the 6-OHDA-lesioned side (right). In 6-OHDA-lesioned mice, we observed that 97 ± 2% of spontaneous rotations were directed towards the lesioned side, compared to a normal value of 54 ± 12% for sham-lesioned animals. Our quantitative immunohistochemical analysis revealed a significant decrease of TH immunoreactivity in the 6-OHDA-lesioned striatum, compared to the intact counterpart and to sham-lesioned animals (Fig. 2). The 6-OHDA group showed a TH residual content of only 9 ± 1% in the lesioned striatum compared to 105 ± 4% for the sham group. In the core and shell compartments of the NAc, the 6-OHDA group of mice presented a similar TH residual content of 20 ± 11% and 25 ± 14%, that were both significantly lower than what was observed in sham animals (110 ± 13% and 102 ± 3%). The efficacy of the 6-OHDA lesion of the dopamine striatal and NAc afferent axons was also supported by a significant decrease of DAT immunoreactivity, the striatum of 6-OHDA-lesioned mice showing a DAT residual content of only 7 ± 2% compared to 103 ± 10% for the sham group. Similarly, the NAc of 6-OHDA-lesioned animals was characterized by a DAT residual content of 14 ± 6% and 21 ± 9% in the core and the shell, compared to 110 ± 18% and 96 ± 4% for the sham-lesioned experimental group. As expected, no significant differences were observed in the striatum and the NAc between the intact side of 6-OHDA and the sham-lesioned or intact side of sham animals, for both TH and DAT staining.

In the mouse striatum, two types of CR + interneurons are observed with opposite dorsoventral distribution, whereas only one type is present in the NAc

Based on their soma size and somatodendritic morphological features, we detected two types of striatal CR + interneurons in the murine specimens used in the present study. Neurons of the first type were intensely stained for CR and endowed with a small-sized (9–12 µm) cell body displaying slightly varicose and beaded processes (Fig. 3A, C). The vast majority of them were unipolar, but a small number
Fig. 2 Assessment of the dopamine lesion following unilateral 6-OHDA injection in the medial forebrain bundle. A Coronal sections taken through the striatum (STR, bregma 1.42 and 0.14), the nucleus accumbens (NAc, bregma 1.42) and the substantia nigra pars compacta (SNc, bregma 3.1) were immunostained for tyrosine hydroxylase (TH) in sham- (upper row) and 6-OHDA- (lower row) lesioned mice. Histograms show TH (B) and dopamine transporter (DAT, C) residual content in the sham- (light green) and 6-OHDA- (dark green) lesioned striatum as well as in the core and shell compartments of the NAc. Dots represent individual data obtained for each mouse. *P < 0.05 for sham- vs 6-OHDA-lesioned mice using Mann–Whitney statistical test.

Fig. 3 Confocal images of calretinin-immunostained (CR+) interneurons in the sham- (A, B) and 6-OHDA- (C, D) lesioned striatum. In the striatum of both experimental groups, small-sized (A, C) and medium-sized (B, D) CR+ cells were observed.
of bipolar cells were also observed. Some of the small CR+ interneurons formed clusters along the subventricular zone and the subcallosal streak, especially in the anterior part of the striatum. Interestingly, this type of CR+ interneurons was completely absent from the NAc. In the striatum itself, the small aspiny CR+ interneurons displayed an anteroposterior and dorsoventral decreasing gradient. Indeed, the quantification of the small-sized CR+ interneurons in the sham-lesioned striatum showed that they prevailed in the pre-commissural striatal sectors (222 ± 32 cells/mm$^3$) and are less abundant in the post-commissural striatum (49 ± 14 cells/mm$^3$, Fig. 4A). The density of these interneurons was also significantly higher in the dorsal half of the sham-lesioned striatum (252 ± 41 cells/mm$^3$) compared to its ventral counterpart (65 ± 23 cells/mm$^3$). A similar density was observed in the medial (159 ± 31 cells/mm$^3$) and lateral (169 ± 28 cells/mm$^3$) striatal sectors. Such neurons are totally absent from the NAc.

Fig. 4 Histograms depicting the density of the small (A, C) and medium (B, D) CR+ cells in different sectors of the sham- (light green bars) and 6-OHDA- (dark green bars) lesioned striatum and nucleus accumbens. In C and D, histograms show the density of CR+ interneurons in the whole lesioned (empty bars) and intact (hatched bars) striatum of sham- and 6-OHDA-lesioned mice. In the sham- and 6-OHDA-lesioned striatum, a significant anteroposterior and dorsoventral decreasing gradient was observed for the small-sized CR+ cells (A) whereas a dorsoventral increasing gradient characterized the medium-sized CR+ neurons (B). In contrast to the small-sized CR+ cells, the density of the medium-sized is significantly decreased in all sectors of 6-OHDA-lesioned striatum, but unaffected in the NAc (B). A similar decreasing trend is observed in the intact striatum (D). *$P<0.05$, †$P=0.057$ using two-way ANOVA followed by Tukey multiple comparison test.
Neurons of the second type are weakly labeled for CR and are endowed with a medium-sized (15–20 µm) cell body displaying smooth aspiny dendrites (Fig. 3B, D). The density of medium-sized CR+ interneurons in the entire striatum was twice as high than that of the small-sized interneurons (370 ± 41 cells/mm³ vs 164 ± 22). Our stereological quantification in the sham-lesioned striatum indicated that medium-sized CR+ interneurons are distributed according to a dorsoventral increasing gradient (Fig. 4B). In contrast to the small-sized CR+ interneurons, medium-sized CR+ cells prevailed in the ventral half of the striatum (466 ± 54 cells/mm³ vs 276 ± 43 for the dorsal counterpart), whereas they appeared rather homogeneously distributed along the anteroposterior (396 ± 46 cells/mm³ vs 312 ± 68) and mediolateral (325 ± 43 cells/mm³ vs 423 ± 57) axes.

In contrast to the small-sized CR+ interneurons observed exclusively in the striatum, the medium-sized interneurons were also found in the NAc, where they were significantly more numerous than in the striatum (1237 ± 335 cells/mm³ vs 370 ± 41). Morphologically, the medium-sized CR+ interneurons observed in the NAc were similar to those found in the striatum (Fig. 5).

The density of the small-sized CR+ interneurons in the striatum is unaffected by 6-OHDA lesion

The small-sized CR+ interneurons had similar morphological features and CR immunoreactivity in the 6-OHDA and the sham-lesioned striatum (Fig. 3A, C). They were also regionally distributed according to the same anteroposterior (248 ± 71 cells/mm³ vs 38 ± 16, \(P = 0.029\)) and dorsoventral (308 ± 80 cells/mm³ vs 31 ± 12, \(P = 0.029\)) decreasing gradient (Fig. 4A). Overall, we found a similar density of the small-sized CR+ interneurons in the sham-lesioned striatum (164 ± 22 cells/mm³) and the 6-OHDA-lesioned striatum (174 ± 46 cells/mm³), indicating that dopamine striatal deafferentation had no significant effect on this specific subtype of CR+ interneurons. This finding was supported by the absence of statistical differences in the density of the small-sized CR+ interneurons between the intact (177 ± 44 cells/mm³) and the 6-OHDA-lesioned striatum (174 ± 46 cells/mm³) of 6-OHDA mice. No significant differences were observed between the intact (120 ± 10 cells/mm³) and the sham-lesioned striatum (164 ± 22 cells/mm³), nor between the intact striatum (upper panels) and the shell (lower panels) compartment of the NAc (Fig. 5B, C).
of the sham group \(120 \pm 10 \text{ cells/mm}^3\) and the intact striatum of 6-OHDA mice \(177 \pm 44 \text{ cells/mm}^3\), Fig. 4C).

**In 6-OHDA-lesioned mice, the density of the medium CR + interneurons is significantly decreased in the striatum but unaltered in the NAc.**

The medium-sized CR + interneurons displayed similar morphological features in the 6-OHDA and the sham-lesioned striatum, with no signs of degeneration in the 6-OHDA-lesioned animals. In 6-OHDA-lesioned mice, they also displayed a similar level of CR immunoreactivity and regional distribution than in controls, their density being higher in the ventral striatal sectors (233 ± 17 cells/mm³) than in dorsal counterparts (82 ± 15 cells/mm³, Fig. 4B). Comparison of the density of medium-sized CR + interneurons in the 6-OHDA lesioned striatum (158 ± 15 cells/mm³) revealed a significant decrease compared to what was observed in the sham-lesioned striatum (370 ± 41 cells/mm³). Although the decrease appeared rather homogeneous throughout the whole striatum (40–72% decrease), we found the highest decrease in the dorsal (70%) and lateral (72%) striatal sectors and the lowest in the ventral (50%) and medial (40%) counterparts. Interestingly, the density of the medium-sized CR + cells was also found to be almost similarly decreased in the intact striatum of 6-OHDA mice (143 ± 53 cells/mm³), when compared to either the sham-lesioned (370 ± 41 cells/mm³) or the intact striatum (316 ± 29 cells/mm³) of sham animals (Fig. 4D). Although non-statistically significant, these results suggest an indirect consequence of the 6-OHDA lesion on the density of the medium-sized CR + striatal cells of the striatum.

In the NAc, the density of the medium-sized CR + neurons was similar between 6-OHDA (1074 ± 204 cells/mm³) and sham-lesioned mice (1237 ± 335), with no significant change in their regional distribution between the core (1146 ± 215 cells/mm³) and the shell (1104 ± 227), when compared to sham animals (1220 ± 345 and 1340 ± 357, Fig. 4B). As expected, similar densities were observed between the 6-OHDA-lesioned NAc (1074 ± 204 cells/mm³) and intact NAc (1193 ± 169) of 6-OHDA mice, as well as between sham-lesioned (1237 ± 335 cells/mm³) and intact NAc (1258 ± 167 cells/mm³) of sham animals. No changes in the proportion of the CR + /CB + doubly stained neurons were noted between 6-OHDA and sham-lesioned NAc, either in the core (8 ± 4% vs 7 ± 4) or in the shell (26 ± 3% vs 26 ± 3).

**Discussion**

The present study has provided the first detailed quantitative description of changes that affect CR + striatal interneurons in a mouse model of Parkinson’s disease. Two morphologically distinct types of CR + aspiny striatal interneurons were detected in mice: (a) small, unipolar cells that stained intensely for CR, displayed beaded dendrites and were scattered according to a dorsoventral decreasing gradient; (b) medium-sized bipolar neurons that stain weakly for CR, displayed smooth dendrites and were scattered according to a dorsoventral increasing gradient (see also Petryszyn et al. 2014; Garas et al. 2018). This study also provides a detailed description of the distribution of CR + cells in the NAc of mice. In contrast to the striatum, where both small and medium-sized CR + interneurons occurred, the mouse NAc contains only medium-sized CR + cells that are rather evenly distributed between its core and shell compartments. The medium-sized CR + cells in NAc were morphologically similar to the CR + neurons of the same size that occurred in the striatum. These results are in accordance with previous observations made in the rat (Hussain et al. 1996) and mouse (Trouche et al. 2019) NAc.

Previous investigations have revealed that GABAergic cells expressing CR form the most abundant class of interneurons in the striatum of primates (Cicchetti et al. 2000; Wu and Parent 2000) and there is evidence that newborn CR + interneurons continue to be added to the adult striatum in both normal (Ernst et al. 2014) and pathological conditions (Wei et al. 2011). In most species investigated thus far, the striatum was described as containing either small-, medium- or large-sized CR + interneurons or a combination thereof: rats (Mura et al. 2000; Rymar et al. 2004; Wu and Parent 2000; Garas et al. 2018), human (Cicchetti et al. 2000; Parent et al. 1995; Petryszyn et al. 2014) and non-human primates (Petryszyn et al. 2014, 2016; Wu and Parent 2000). The primate striatum harbors a unique population of giant CR + interneurons, which express choline acetyltransferase (ChAT), a faithful marker of cholinergic neurons (Bernacer et al. 2012; Kataoka et al. 2010; Petryszyn et al. 2014, 2016). These large-sized CR + / ChAT+ interneurons are significantly decreased in number in the striatum of parkinsonian monkeys, a change that was shown to reflect a higher level of CR expression by ChAT+ interneurons rather than an increase in their absolute number (Petryszyn et al. 2016).

Such giant CR + interneurons that express ChAT do not occur in the striatum of mice. The latter contains a significant number of medium-sized CR + interneurons, which are morphologically and neurochemically similar to striatal CR + interneurons of the same size detected in other species, and a smaller number of small-sized, unipolar CR + cells that prevail in the anterodorsal sector of the striatum, near the subventricular zone, an area that retains its neurogenic capacity throughout adult life (Ming and Song 2011). The immature appearance of these small CR + interneurons and their presence near the neurogenic zone of the striatum suggest that they might be issued from postnatal neurogenesis.
as it appears to be the case in normal adult rats (Dayer et al. 2005; Garas et al. 2018), rabbits (Luzzati et al. 2006) and human (Ernst et al. 2014). Evidence for small CR + neurons generated during early postnatal development has also been obtained in mice (Revishchin et al. 2010a, 2010b). These immature looking cells were described as having a perikaryon and processes coated with polymorphous spines; they, thus, differ markedly from the typical GABAergic CR + aspiny striatal interneurons as well as from the small-sized CR + cells described in the present study.

In addition to the typical small-, medium- and large-sized striatal CR + interneurons known to be present in rats (Rymar et al. 2004; Wu and Parent 2000; Garas et al. 2018), two additional types of CR + cells were described following a study of rats with a 6-OHDA-lesion of the nigrostriatal pathway (Mura et al. 2000). In that study, neurons of the first type were detected in both dopamine-depleted and intact striata and consisted of small unipolar CR + cells endowed with an irregular dendritic tree that branches very near to the cell body in a bush-like pattern. Neurons of the second type, present only in the 6-OHDA-lesioned striatum, were less numerous and composed of uni- or bipolar CR + cells with dendrites displaying tiny and widely interspaced spine-like structures. The morphological features of these small CR + cells in the rat striatum do not match with those observed in the present study in mice. Furthermore, in that single study, an overall increase of CR + cells was noted in dopamine-lesioned striatum 3 weeks following 6-OHDA injection, but this augmentation tended to normalized at 6, 10, and 18 weeks post-lesion (Mura et al. 2000). Whether this increase could solely be explained by a postnatal production of these two new types of CR + cells or by a transient increase in the expression of CR is unclear. In this regard, it should be reminded that the expression of CR appears to be transitory in many neuronal types and could vary according to the maturation state of the neurons (Schwaller 2014).

The present investigation of the dopamine-depleted striatum in mice has revealed that the two types of striatal CR + cells detected previously in normal animals (Petryszyn et al. 2014) occur in the 6-OHDA-lesioned striatum and its sham counterpart, their regional distribution, morphological characteristics and neurochemical makesups being totally unaltered by dopamine depletion. However, the density of the medium-sized CR + interneurons remained unaffected. Such a 6-OHDA-induced decrease in the number of medium-sized CR + striatal cells with no alteration of the small CR + cells observed here in mice is at variance with the transient increase in the number of striatal CR + cells reported in rats under the same pathological condition (Mura et al. 2000). Such a discrepancy could be due either to species differences or variations in experimental designs. It should be noted that the present observations were made almost 6 weeks following 6-OHDA lesion, a period during which the increase noted in rats was shown to be back to normal (Mura et al. 2000). It is, thus, tempting to hypothesize that the increase in the number of CR + cells noted 3 weeks following 6-OHDA injection in rats is the result of a higher level of CR expression, as part of an early compensatory mechanism designed to cope with increased intracellular calcium level that might occur following altered glutamatergic corticostriatal transmission that is believed to characterize Parkinson’s disease (Calabresi et al. 1996; Cepeda et al. 1998). In regard to the preservation of the small striatal CR + cells observed in the present study, it is worth noting that these neurons abound in the anterior sector of the striatum, which is principally innervated by cortices of the associative type. Such a specific topographical and connectional relationship suggests that the small CR + interneurons are chiefly involved in the integrative role of the striatum, whereas the medium-sized CR + interneurons, which are affected by dopamine depletion, are principally concerned with the sensorimotor or limbic function of the striatum, which are dealt with in more posterior and ventral aspects of the structure. The topographical distribution and high level of CR expression displayed by the small-sized CR + striatal cells might render them less vulnerable than the medium-sized CR + interneurons to increased calcium levels derived from enhanced striatal excitatory input arising from the cerebral cortex (Mitchell et al. 1994; Blandini et al. 1996; Calabresi et al. 1996; Calabresi et al. 1996; Meshul et al. 1999).

The decreased in the density of the medium-sized CR + interneurons reported here in the striatum of 6-OHDA-lesioned mice is congruent with the results of another study in 6-OHDA rats, in which a significant decrease of CR + interneurons was observed in the ipsilateral striatum, 5 weeks following unilateral 6-OHDA injection (Ma et al. 2014). In the latter study, however, the existence of various morphological subtypes of CR + cells has not been taken into account in the quantitative analysis so that it is not possible to know if some types of CR + striatal interneurons might have been spared in the 6-OHDA-lesioned rats, as it is the case in 6-OHDA-lesioned mice.

Weather the decrease of the medium-sized CR + interneurons reported in the present study is caused by neuronal death or result from a decreased in CR expression following dopamine striatal denervation is unknown. However, it should be noted that the same level of CR immunoreactivity was observed in the sham- and 6-OHDA-lesioned striatum and that no morphological signs of neuronal degeneration were observed for these medium CR + striatal cells in the 6-OHDA-lesioned striatum. Through its capacity of maintaining intracellular calcium homeostasis, CR is believed confer to neurons a certain protection against neurodegenerative processes, particularly glutamatergic excitotoxicity, which is considered a major pathogenic component of
Parkinson’s disease (Choi 2005). This could explain why the medium-sized CR+ were selectively affected in the 6-OHDA-lesioned striatum, being less immunoreactive for CR than the small-sized CR+ cells for which the density appears unchanged.

Also of interest is the fact that the intact striatum of 6-OHDA-lesioned mice also showed a trend towards decreased density of CR+ medium interneurons, a tendency not observed in the sham-lesioned mice. Although not statistically significant, such a finding suggests an indirect effect of the dopamine lesion on striatal CR+ interneurons. A possible influence of dopamine on CR+ striatal neurons has been alluded to previously (Kawaguchi et al. 1995), but the mechanism whereby such a putative dopaminergic action might be exerted has remained elusive. Immunohistochemical studies in rats have shown that CR+ interneurons express a low level of the dopaminergic D5 receptors (Rivera et al. 2002), but such neurons appear to be completely devoid of the D1 and D2 receptors (Petryszyn et al. 2014). Thus, dopamine is likely to exert its influence upon CR+ striatal interneurons indirectly, probably by modulating the glutamatergic excitatory striatal projections of cortical or thalamic origin. The fact that the most significant decrease of the medium-sized CRcells was observed in the dorsolateral sectors of 6-OHDA-lesioned striatum and that no changes were observed in the NAc indicates that the medium-sized striatal interneurons expressing low level of CR and located in the sensorimotor striatal territory might be particularly vulnerable to an increase in intracellular calcium. This possibility is further strengthened by data indicating that perturbations of glutamatergic corticostriatal transmission induced by striatal dopamine depletion are particularly conspicuous in the sensorimotor striatal territory (Meshul et al. 1999; Blandini et al. 1996).

The intensity of the immunostaining for TH and DAT in the intact striatum of 6-OHDA-lesioned striatum was similar to that in sham animals. Hence, despite the fact that 5–10% of nigrostriatal axons have been reported to cross the midline in normal condition (Fass and Butler 1981), it is difficult to attribute the bilateral decrease of medium CR+ striatal cells to a lower dopamine concentration in the intact striatum of 6-OHDA-lesioned striatum compared to sham animals. Instead, the involvement of the corticostriatal projections could be a more likely explanation. Corticostriatal neurons are able to reach their target by either an ipsilateral, contralateral or bilateral projection (Hooks et al. 2018; Kiritani et al. 2012; Parent and Parent 2006). Hence, a bilaterally altered corticostriatal transmission could indeed explain, at least in part, the bilateral reduction of CR+ medium cells in 6-OHDA-lesioned mice. It is worth noting that various bilateral effects have often been reported following unilateral 6-OHDA lesion (Avila-Costa et al. 2005; Jedrzejewska et al. 1990; Nikolaus et al. 2003), including a transient increase in the number of striatal CR+ neurons following unilateral 6-OHDA-lesion in rats (Mura et al. 2000). However, a potential systemic effect of the 6-OHDA injection cannot be ruled out, particularly because increased level of pro-inflammatory cytokines and reduced concentrations of neurotrophic factors have previously been reported in the 6-OHDA-lesioned mice (Antunes et al. 2020).

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Author contributions SP was in charge of running experiments. SP and LS were in charge of data and statistical analyses. DG provided guidance for stereology. MP designed experiments and wrote the manuscript. AP revised the manuscript.

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Declarations

Conflict of interest The authors have no conflict of interest to declare.

Ethical approval All protocols were approved by the Institutional Animal Care and Use Committee (Comité de Protection des Animaux de l’Université Laval, #2019-287/VRR-18-107) and all procedures involving animals and their care were made in accordance with the Canadian Council on Animal Care’s Guide to the Care and Use of Experimental Animals (Ed2).

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