In Situ Mitochondrial Ca\textsuperscript{2+} Buffering Differences of Intact Neurons and Astrocytes from Cortex and Striatum*\textsuperscript{*1}

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The striatum is particularly vulnerable to neurological disorders, such as Huntington disease. Previous studies, with nonsynaptic mitochondria isolated from cortical and striatal homogenates, suggest that striatal mitochondria are highly vulnerable to Ca\textsuperscript{2+} loads, possibly influencing striatal vulnerability. However, whether and how neuronal and glial mitochondria from cortex and striatum differ in Ca\textsuperscript{2+} vulnerability remains unknown. We test this hypothesis using a novel strategy allowing comparisons of mitochondrial Ca\textsuperscript{2+} buffering capacity in cortical and striat lar neuron-astrocyte co-cultures. We provide original evidence that mitochondria not only in neurons but also in astrocytes from striatal origin exhibit a decreased Ca\textsuperscript{2+} buffering capacity when compared with cortical counterparts. The decreased mitochondrial Ca\textsuperscript{2+} buffering capacity in striatal versus cortical astrocytes does not stem from variation in mitochondrial concentration or in the rate of intracellular Ca\textsuperscript{2+} elevation, being mechanistically linked to an increased propensity to undergo cyclosporin A (CsA)-sensitive permeability transition. Indeed, 1 \textmu M CsA selectively increased the mitochondrial Ca\textsuperscript{2+} buffering capacity of striatal astrocytes, without modifying that of neurons or cortical astrocytes. Neither thapsigargin nor FKS06 modified mitochondrial Ca\textsuperscript{2+} buffering differences between cell types, excluding a predominant contribution of endoplasmic reticulum or calcineurin. These results provide additional insight into the mechanisms of striatal vulnerability, showing that the increased Ca\textsuperscript{2+} vulnerability of striatal versus cortical mitochondria resides in both intact neurons and astrocytes, thus positioning the striatum at greater risk for disturbed neuron-astrocyte interactions. Also, the selective effect of CsA over striatal astrocytes suggests that in vivo neuronal sheltering with this compound may indirectly result from astrocytic protection.

Brain regions and cell types vary in their susceptibility to neurological disorders. The striatum is particularly vulnerable to both acute (e.g. hypoglycemia, hypoxia, and global ischemia) and chronic (e.g. Huntington disease (HD)) disorders. Excitotoxicity and altered energy metabolism have been proposed as important determinants of striatal pathogenesis, leading to toxic increases in [Ca\textsuperscript{2+}]. (1). Studies with mitochondria isolated from brain homogenates raised the hypothesis that striatal mitochondria are more vulnerable to Ca\textsuperscript{2+} loads than cortical mitochondria, possibly contributing to selective striatal vulnerability in pathologies such as HD (2). However, because isolated mitochondria are deprived from their physiological cellular context, and brain homogenates contain mitochondria from both neurons and glia, two fundamental questions remain unanswered: first, whether the differences in Ca\textsuperscript{2+} vulnerability, observed with isolated mitochondria, also occur in situ (i.e. when mitochondria are in their physiological environment inside intact cells) and, second, whether mitochondrial vulnerability to Ca\textsuperscript{2+} loads differs among neurons and glia from cortex and striatum.

Striatal selectivity in HD is noteworthy, given that the disease-causing protein (mutant huntingtin) is widely expressed throughout the brain and not enriched in the striatum (3). Although HD may also affect nonstriatal regions, namely the cortex (4), typical striatal atrophy and neuronal loss greatly exceed those seen in nonstriatal regions, particularly when HD is not superimposed with Alzheimer’s disease or age-related volumetric loss (5). This suggests that vulnerability factors inherent to striatal neurons shape the selective pattern of HD neurodegeneration. One likely factor is the enrichment in N-methyl-D-aspartate (NMDA) receptors containing NR2B subunits, allowing prolonged Ca\textsuperscript{2+} influx following excitotoxic stimuli (6), but other vulnerability factors may assist. Mitochondria have an essential role in neuronal survival following excitotoxic [Ca\textsuperscript{2+}], elevations (7), and mitochondria-dependent Ca\textsuperscript{2+} handling is impaired in striatal neurons from HD mice (8). HD may in fact be aggravating inherent mitochondrial vulnerability in the striatum, which may reside in either neurons or glia. Indeed, glia may also contribute for non-cell-autonomous neurodegeneration in HD (9) or ischemic injuries (10, 11), thus stressing the importance of identifying vulnerability factors in both neurons and glia.

In the present study, we test the hypothesis that mitochondrial vulnerability to Ca\textsuperscript{2+} loads differs among intact neurons and astrocytes from cortex and striatum. We prepared primary neuron-astrocyte co-cultures from rat cortex and striatum.

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2. The abbreviations used are: HD, Huntington disease; CsA, cyclosporin A; CypD, cyclophilin D; DIV, days in vitro; ER, endoplasmic reticulum; FCCP, carboxyfluorescein-d-phthalocyanine-p-trifluoromethoxyphenyl) hydradoreze; FKS06, tacrolimus; GFAP, glial fibrillary acidic protein; MK-801, (S\textsuperscript{a},10R\textsuperscript{a}-methyl-10,11-dihydro[2,3]dihydro[1,4]oxepin-5,10-imine hydrogen maleate; NMDA, N-methyl-D-aspartate; NMDAR, NMDA receptor; PMPL, plasma membrane potential indicator; ROI, region of interest; TMRM\textsuperscript{*+}, tetramethylrhodamine methyl ester; ANOVA, analysis of variance.
whose cellular composition and functional differentiation were confirmed by immunolabeling and functional imaging studies, respectively. Here we report for the first time not only the development but also the application of an experimental strategy allowing comparisons of in situ mitochondrial Ca\(^{2+}\) buffering capacity in the preserved physiological context of intact brain cells. To allow balanced comparisons, similarly to standard procedures with isolated mitochondria experiments, we have introduced calibrations for differences in mitochondrial concentration and rate of intracellular Ca\(^{2+}\) elevation in intact cells.

Results in the present study provide additional insight into the mechanisms of striatal vulnerability, showing for the first time that the increased vulnerability of striatal mitochondria to Ca\(^{2+}\) loads is present in both intact neurons and astrocytes when compared with their cortical counterparts. Interestingly, our finding of a lower mitochondrial Ca\(^{2+}\) buffering capacity in intact striatal versus cortical astrocytes, mechanismically linked to increased cyclosporin A-dependent permeability transition, positions the striatum at higher risk for disturbed interactions between synaptic partners, neurons and astrocytes.

**EXPERIMENTAL PROCEDURES**

**Materials**—Fura-4F acetoxymethyl ester, tetramethylrhodamine methyl ester (TMRM\(^{+}\)), and 4-bromocamphotinic were from Invitrogen. Plasma membrane potential indicator (PMPI) (Component A from the R-8042 membrane potential assay kit) was from Molecular Devices Corp. (Sunnyvale, CA) (12). Culture media and supplements were from Invitrogen unless otherwise stated. NMDA, (S,R,10S)-(+)-5-methyl-10,11-dihydro[a,d]-cyclohepten-5,10-imine hydrogen maleate (MK-801), ifeneprol, myxothiazol, oligomycin B, rotenone, carboxyca
ide p-(trifluoromethoxyphenyl) hydrazone (FCCP), cyclosporin A, tacrolimus (FK506), thapsigargin, and other reagents were from Sigma.

**Cell Culture**—Primary cultures were generated from the offspring of Wistar rats (CRIFFA, Barcelona, Spain). Handling and care of animals were conducted according to the European Union guidelines for animal research (86/609/EEC; in agreement with the National Institutes of Health guidelines) and Portuguese law (Portarias 1005/92 and 1131/97). Cortical and striatal cultures were prepared and maintained as previously described for striatal cultures (8, 13) but without cytosine arabinoside to allow the growth of astrocytes. Briefly, hemcocitcrites (free from meninges, olfactory bulb, striatum, and hippocampus) and hemistriata were dissected from the same embryonic day 18 rats, pooled according to brain region (cortex or striatum), and processed in parallel (sister cultures). Dissociated cells (1 × 10\(^5\)) were plated in 12-mm round polyethyleneimine-coated glass coverslips and used in functional imaging experiments between 11 and 13 days in vitro (DIV).

**Immunocytochemistry**—Cells were fixed with 4% paraformaldehyde for 15 min at 37 °C and permeabilized with 0.4% Triton X-100 in phosphate-buffered saline for 15 min. Blocking was performed with 3% bovine serum albumin in phosphate-buffered saline for 30 min at room temperature. Cells were then incubated with primary antibodies for the neuronal marker (mouse anti-MAP-2 (microtubule-associated protein); Sigma; 1:500) and the astrocytic marker (rabbit anti-glial fibrillary acidic protein (GFAP); Dako; 1:250) in 3% bovine serum albumin phosphate-buffered saline for 1 h at room temperature. After washing, cells were incubated with appropriate secondary antibodies conjugated to either AlexaFluor 488 or 594 (Invitrogen; 1:200) for 1 h at room temperature and shielded from light. Nuclei labeling was performed with Hoechst 33342 (Invitrogen; 2 μg/ml; 10 min at room temperature). To test for antibody selectivity, primary antibodies were omitted in control coverslips processed in parallel. No immunolabeling was observed in control coverslips, which were used for shading correction of epifluorescent images, acquired with the same equipment settings.

**Monitoring of Dynamic Changes in [Ca\(^{2+}\)]\(_i\), Mitochondrial Membrane Potential (Δψ\(_m\)), and Plasma Membrane Potential (Δψ\(_p\)) in Single Cells**—Functional imaging at single-cell resolution was performed with a system composed by an inverted epifluorescence microscope (Eclipse TE300, Nikon, Tokyo, Japan) equipped with 20×, 40×, and 60× air objectives, a monochromator (Polychrome II; TILL Photonics, Martinsried, Germany), a CCD camera (C6790; Hamamatsu Photonics, Hamamatsu, Japan), and a computer with analysis software (AquaCosmos 2.5; Hamamatsu Photonics). Cells were loaded with the [Ca\(^{2+}\)]\(_i\) probe Fura-4F (5 μM) for 45 min at 37 °C. For simultaneous recordings of changes in [Ca\(^{2+}\)]\(_i\), and in TMRM\(^+\) fluorescence (indicative of changes in Δψ\(_m\)), 50 nM TMRM\(^+\) (quench mode) was present during Fura-4F loading and throughout the experiments. Simultaneous recordings of changes in [Ca\(^{2+}\)]\(_i\), and in Δψ\(_p\), were performed in Fura-4F-loaded cells with a constant PMPI concentration present throughout the experiment. The PMPI concentration was the same as previously described (12), corresponding to a 1:2,000 dilution of a PMPI stock solution prepared by reconstituting one R-8042 Component A vial in 1 ml of ultrapure water. Experiments were performed at 37 °C in buffer containing 133 mM NaCl, 5 mM KCl, 1 mM Na\(_2\)SO\(_4\), 0.4 mM KH\(_2\)PO\(_4\), 20 mM HEPES, 1.3 mM CaCl\(_2\), and 15 mM glucose, pH 7.4. Where indicated, glucose was replaced by 2 mM 2-deoxy-D-glucose, and mitochondria were energized with 10 mM pyruvate. Fura-4F was excited at 340 and 380 nm, TMRM\(^+\) at 500 nm. Dual Fura-4F and TMRM\(^+\) recordings were performed with a 520-nm long pass filter. There was no significant bleed-through between Fura-4F and TMRM\(^+\) or between Fura-4F and PMPI with these excitation and emission settings. Individual cells were identified as regions of interest (ROIs) for the determination of fluorescence time courses.

**In situ calibration of Ca\(^{2+}\) responses for each individual cell was performed at the end of every experiment, according to previously detailed procedures (8). Briefly, maximal 340/380 ratios (R\(_{max}\)) were determined with 20 μM 4-bromocamphotinic with 1.3 mM extracellular Ca\(^{2+}\), and minimal 340/380 ratios (R\(_{min}\)) were determined in Ca\(^{2+}\)-free buffer with 5 mM EGTA. This Ca\(^{2+}\)-free buffer contained neither TMRM\(^+\) nor PMPI, and thus the subsequent decays in their respective fluorescence should not be interpreted as changes in Δψ\(_m\) or Δψ\(_p\). In experiments addressing mitochondrial Ca\(^{2+}\) buffering capacity, maximal mitochondrial depolarization (Δψ\(_m\) collapse) was con-
firmed/evoked before \([\text{Ca}^{2+}]\), calibration by adding protonophore (1 \(\mu\text{M} \text{FCCP}\)), with ATP synthase reversal prevented by oligomycin (3 \(\mu\text{M}\)). Further details on the analysis of cellular responses are provided below.

**Image Processing and Data Analysis—Analysis of \([\text{Ca}^{2+}]_i\)** responses was performed as previously detailed (8). Briefly, background-corrected 340/380 Fura-4F ratios were normalized to the \(R_{\text{max}}\) (100 units) and \(R_{\text{min}}\) (0 units) determined for each single cell during in situ calibration. The left yy axes of \([\text{Ca}^{2+}]_i\) graphs are scaled with these normalized values, where 50 units should correspond to the \(\text{Ca}^{2+}\) probe \(K_p\), regardless of its absolute value inside the different cells. The right yy axis is an approximate log \([\text{Ca}^{2+}]_i\) scale adjusted based on the dynamic range of Fura-4F and \(K_p\) determinations (15, 16).

Changes in TMRM\(^+\) and PMPI fluorescence in single cells are expressed as arbitrary fluorescence units. Individual signals are corrected for epifluorescence shading, using images acquired in the same experimental xx-yy-zz plane, without TMRM\(^+\) (during \(R_{\text{min}}\) determination at the end of the experiment) and in the absence of PMPI (before starting the experiment, in PMPI-free buffer). Exposure times for image acquisition were kept constant in all experiments, with signal saturation and underexposure being avoided merely by adjusting camera gain before starting the experiments. Mathematical correction for different gains was introduced so that all fluorescent signals are compared under the same circumstances.

Conversions to mV of neuronal PMPI fluorescence (Figs. 1, B and D (ii), and 2, A (ii)) were performed in a manner similar to that described previously (12) but with in situ calibration for each individual neuron. Briefly, extracellular \(\text{Na}^+\) was replaced equimolarly by \(\text{K}^+\) in a stepwise fashion, up to 42 \(\mu\text{M}\) extracellular \(\text{K}^+\) in the assay buffer. \(\Delta\psi_p\) values (in mV) were derived from the Goldman-Hodgkin-Katz equation, assuming conductances for \(\text{K}^+\), \(\text{Na}^+\), and \(\text{Cl}^-\) of 100:1:100. PMPI fluorescence enhancements evoked by \(\text{K}^+\) in each neuron were plotted in a function of \(\Delta\psi_p\) and fit monoexponentially (correlation coefficients, \(r^2 > 0.990\)) to generate individual calibration curves to derive approximate \(\Delta\psi_p\) changes in mV.

Cell size was determined from the ROI area (\(\mu\text{m}^2\)) using micrometer-calibrated images. Neuronal areas reflect only the soma, due to impossibility of tracing the entire neuronal processes. Area conversion into approximate somatic diameter (e.g. Fig. 2B) was performed by a circular approximation using Equation 1,

\[
d = 2 \times \text{SQRT}(a/\pi) \quad \text{(Eq. 1)}
\]

where \(d\) represents the diameter, \(\text{SQRT}\) is the square root, \(a\) is the area, and \(\pi\) is the circular constant.

Mitochondrial concentration in single cells was estimated from the analysis of the average TMRM\(^+\) fluorescence in each ROI (for a representative measurement, see Fig. 3A (ii), solid line, \(\gamma - \delta\)). The difference between resting TMRM\(^+\) fluorescence (start of experiment) and nonmitochondrial fluorescence (TMRM\(^+\) signal after FCCP, prior to \(R_{\text{max}}\) determination; Figs. 3 and 4 (ii), solid lines) was assumed to reflect mitochondrial concentration. Our assumption is based on evidence that when the amount of TMRM\(^+\) stacked in the mitochondria surpasses the quench threshold, autoquenching occurs and fluorescence becomes invariant with matrix TMRM\(^+\) concentration (14); TMRM\(^+\) quenching in the present study is evidenced by the dequenching spikes in Fig. 4, A–D, ii, solid lines). Therefore, the resting cellular TMRM\(^+\) fluorescence in our experiments is primarily governed by the amount of mitochondria per cell volume (mitochondrial concentration) with superimposed nonmitochondrial fluorescence. This nonmitochondrial fluorescence reflects TMRM\(^+\) in extracellular buffer in the cytosol and non-specifically bound to cellular elements. Hence, subtracting the residual TMRM\(^+\) signal following full \(\Delta\psi_p\) collapse (as confirmed with FCCP in the presence of oligomycin) should eliminate the vast majority of nonmitochondrial fluorescence, thus yielding an estimate of mitochondrial concentration. We further substantiate this procedure by providing both medium and high magnification images (supplemental material) of TMRM\(^+\)-loaded neurons and astrocytes used in functional experiments. Collectively, the images show that the mitochondrial distribution is sparser in astrocytes that in neurons, visually showing a lower mitochondrial concentration. We also provide fluorescence intensity profiles across representative cells as well as time lapse experiments illustrating how \(\Delta\psi_p\) collapse with oligomycin plus FCCP influences TMRM\(^+\) distribution and fluorescence intensity values in mitochondria-rich and mitochondria-free cellular regions.

The rate of \([\text{Ca}^{2+}]_i\) elevation was estimated by curve-fitting (with nonlinear regression) the normalized Fura-4F 340/380 ratios in experiments such as in Fig. 4, where the combination of oligomycin and a respiratory chain inhibitor (myxothiazol) collapsed \(\Delta\psi_p\), preventing mitochondrial \(\text{Ca}^{2+}\) uptake (17). Best fit values, with \(r^2 > 0.990\), were obtained with Equation 2,

\[
y = I(\chi < x_o), \quad \text{THEN} \quad y = b, \quad \text{ELSE} \quad y = b + (t - b) \times (1 - e^{-(x-x_o)}) \quad \text{(Eq. 2)}
\]

where \(y\) represents normalized \([\text{Ca}^{2+}]_i\), at time \(x\), \(x_o\) is the first time point, \(b\) is basal \([\text{Ca}^{2+}]_i\), \(t\) is the top part of the curve (constrained to be below the 100 value achieved during \(R_{\text{max}}\)), \(e\) is Neper’s number, and \(K_{\exp}\) is the exponential constant used to compare the rate of \([\text{Ca}^{2+}]_i\) elevation among cells (e.g. Fig. 4A (i), \(K_{\exp}\text{ arrow}\)). It should be noted that myxothiazol, albeit at higher concentrations, was previously shown to inhibit \(\text{Ca}^{2+}\) influx through store-operated channels (18). However, 3 \(\mu\text{M}\) oligomycin (specifically oligomycin B, which has a 0.5 \(\mu\text{M} K_{\exp}\) for the inhibition of store-operated \(\text{Ca}^{2+}\) influx (19)) was present in our experiments either with or without myxothiazol, thus preventing differences in store-operated \(\text{Ca}^{2+}\) influx between these experiments.

Image processing was performed with Aquacosmos 2.5 (Hamamatsu Photonics) and with ImageJ (National Institutes of Health; available on the World Wide Web). Calculations on numerical data derived from ROIs were automated in Excel spreadsheets (Microsoft Corp., Redmond, WA). Values throughout are presented as mean ± S.E. unless otherwise stated. Nonlinear regressions, curve fitting, and t tests were performed using GraphPad Prism version 4.0 (San Diego, CA). Other statistical analyses were performed with Statistical Package for the Social Sciences version 15.0 for Windows (SPSS Inc., Chicago, IL). Type I error probability was set at 0.05 for all analyses.
Characterization of Neurons and Astrocytes in Cortical and Striatal Cultures—Immunolabeling revealed that cortical and striatal cultures were predominantly composed of neurons and astrocytes (over 95% of all nuclei stained with Hoechst 33342 were also positive for either the neuronal marker, MAP-2, or the astrocytic marker, GFAP) (supplemental Fig. 1). At 11 DIV, cells were functionally differentiated. Neurons responded to K⁺ or NMDA receptor (NMDAR) activation with \([\text{Ca}^{2+}]_{i}\) elevation and plasma membrane depolarization in a concentration-dependent manner (Fig. 1, A and C, and solid lines in B and D). In contrast, astrocytes did not respond to NMDA (up to 100 \(\mu\text{M}\)), despite exhibiting changes in plasma membrane polarization upon K⁺ challenging. Also, no changes in astrocytic \([\text{Ca}^{2+}]_{i}\) were detected during either K⁺ or NMDA challenging, despite clear responses to \(\text{Ca}^{2+}\)-ionophore during \([\text{Ca}^{2+}]_{i}\) calibration (\(R_{\text{max}}\); Fig. 1, A and C, and dashed lines in B and D). These clear functional differences ensured cellular identification, particularly in experiments with PMPI (e.g. Fig. 1, A and C (ii), bottom left; Fig. 3, A and B (ii), dashed lines).

To appraise differences between cortical and striatal neurons, we integrated three parameters (somatic diameter, NMDA pEC\(_{50}\), and proportion of functional NMDAR-2B) in a cluster analysis with \(n = 23\) random neurons of each type. The average somatic diameter was larger for cortical neurons (17.4 ± 0.4 \(\mu\text{m}\) versus 12.9 ± 0.2 \(\mu\text{m}\); \(p < 0.001\)). Despite similar...
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NMDA pEC$_{50}$ (5.48 ± 0.05 versus 5.36 ± 0.05, cortical and striatal neurons, respectively; $p = 0.087$), the contribution of the NMDAR-2B subtype for [Ca$$^{2+}$$], responses was higher in striatal (37.0 ± 2.1%) versus cortical neurons (24.9 ± 1.9%; $p < 0.001$). This latter parameter was derived from the degree of attenuation with ifenprodil (2 μM; selective NMDAR-2B antagonist; IFEN in Fig. 2A) of plateau responses to 10 μM NMDA (a concentration close to the NMDA EC$_{50}$) and is expressed as a percentage of maximal attenuation with MK-801 (5 μM; non-subtype-selective NMDAR antagonist). These three parameters were plotted three-dimensionally, and a “two-step cluster analysis” (Statistical Package for the Social Sciences version 15.0; blind to both the cortical or striatal origin of the data points as well as the expected number of clusters) separated them into two clusters (Fig. 2B, red dashed line), nearly completely (40:46) matching their brain origin and thus showing strong differences among the two neuronal populations. Meaningfully, these differences are reminiscent of the in vivo condition, namely the smaller somatic size of striatal neurons and particularly their higher proportion of functional NMDAR-2B, which is considered a relevant determinant for their selective vulnerability (6, 20).

Interestingly, three-dimensional cluster analysis with morphofunctional parameters also separated neurons and astrocytes with minor overlap (supplemental Fig. 2), showing the potential application of this strategy to automated image analysis. Most importantly, evidence of morphological and functional differences among cells underlines the significance of comparing their mitochondrial properties in situ.

**In Situ Mitochondrial Ca$$^{2+}$$ Buffering Capacity in Intact Neurons and Astrocytes**—To monitor mitochondrial Ca$$^{2+}$$ buffering in situ, cells were loaded with Fura-4F ($K_d$ ∼ 1 μM), a probe we have previously shown adequate to identify the mitochondrial set point for Ca$$^{2+}$$ accumulation (8). Because we aimed as studying intact cells, we avoided standard strategies such as ionophores or proteolytic enzymes as a means to expose mitochondria to Ca$$^{2+}$$. Instead, we devised a strategy using a controlled ATP depletion to elevate [Ca$$^{2+}$$]$_i$, which was buffered at the expense of $\Delta \psi_m$ by mitochondria energized with the cell-permeable substrate pyruvate (10 mM).

To initiate the Ca$$^{2+}$$ challenging of in situ mitochondria, glycolytic and mitochondrial ATP generation were prevented by replacing glucose with 2-deoxy-D-glucose (2 mM) and by adding the ATP synthase inhibitor oligomycin (3 μM). This procedure initiated an elevation of [Ca$$^{2+}$$], in both neurons and astrocytes (Fig. 3, A–D, i, O), reaching a “plateau” (Fig. 3A (i), α point), after which [Ca$$^{2+}$$]$_i$ was buffered for a variable period until the onset of a massive increase in [Ca$$^{2+}$$]$_i$ (Fig. 3A (i), β point), suggesting widespread Ca$$^{2+}$$ overload in the in situ mitochondrial population of each single cell, which invariably led to cell death.

Evidence that the above described [Ca$$^{2+}$$]$_i$ plateau results from mitochondrial Ca$$^{2+}$$ buffering is as follows: (i) the plateau occurs at [Ca$$^{2+}$$]$_i$, close to the mitochondrial set point for Ca$$^{2+}$$ accumulation (0.3–1 μM) (7) (Fig. 3, A–D (i)); (ii) no plateau is observed if mitochondria are prevented from uptaking Ca$$^{2+}$$ (Fig. 4, A–D (i)); (iii) the plateau is immediately disrupted, and mitochondrial Ca$$^{2+}$$ is released to the cytosol, upon $\Delta \psi_m$ collapse with protonophore (1 μM FCCP) (supplemental Fig. 3B); (iv) mitochondrial depolarization, as judged from the analysis of TMRM$^+$ redistribution (Fig. 3, A–D (ii), solid lines), is ongoing throughout the [Ca$$^{2+}$$]$_i$ plateau and near completion at the time where the plateau can no longer be sustained (Fig. 3, β point).

Measurements of in situ mitochondrial Ca$$^{2+}$$ buffering were thus derived from the time difference between the aforementioned α and β points (β − α = duration of the [Ca$$^{2+}$$]$_i$ plateau), a procedure that follows the same reasoning applied to isolated
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**FIGURE 3.** In situ monitoring of mitochondrial Ca\(^{2+}\) buffering capacity in neurons and astrocytes from cortex and striatum. A–D, changes in \([\text{Ca}^{2+}]\), \(i\) and in TMRM\(^{+}\) (\(ii\); solid lines) or PMPI (\(ii\); dashed lines) fluorescence in arbitrary units (A.F.U.). The solid or dashed line in \(i\) and \(ii\) depicts measurements performed simultaneously in the same single cell. Representative cells with identical \([\text{Ca}^{2+}]\), kinetics were chosen (\(i\)) in order to allow comparison of changes in TMRM\(^{+}\) and PMPI fluorescence (\(ii\)). Assay buffer contained 10 mM pyruvate, and glucose was replaced by 2 mM 2-deoxy-o-glucose before the addition of oligomycin (3 \(\mu\text{M}\); O). FCCP addition (1 \(\mu\text{M}\); F) confirms full \(\Delta\psi_{m}\) collapse. \(R\) and \(C\), \(R_{\text{max}}\) and \(R_{\text{min}}\), determinations, respectively. A (\(i\)), \(\beta - \alpha\), duration of the \([\text{Ca}^{2+}]\), plateau (i.e. apparent mitochondrial Ca\(^{2+}\) buffering). A (\(ii\)), \(\gamma - \delta\), mitochondrial concentration. Up to 50 random single cells of each type were analyzed per experiment; only a few are shown for clarity, being representative of \(n = 7\) independent experiments.

Mitochondria experiments with steady Ca\(^{2+}\) infusion (13, 21). Results for the duration of the \([\text{Ca}^{2+}]\), plateau are shown in Fig. 5A, where a two-factor (region \(\times\) cell type) ANOVA revealed a significant region \(\times\) cell type interaction (\(F_{(1,24)} = 4.63; p = 0.042\)), a significant main effect of the cell type (neurons \(\text{versus}\) astrocytes; \(F_{(1,24)} = 23.7; p < 0.001\)), and no significant effect of the brain region (cortical \(\text{versus}\) striatal; \(F_{(1,24)} = 1.23; p = 0.279\)).

These results, per se, suggest that in situ mitochondrial Ca\(^{2+}\) buffering capacity is higher in neurons than in astrocytes, with no differences between cortical and striatal neurons, and with mitochondria in striatal astrocytes exhibiting the lowest Ca\(^{2+}\) buffering capacity, thus affording the lowest protection and precipitating loss of Ca\(^{2+}\) homeostasis and cell death. Still, following a similar reasoning to standard isolated mitochondria experiments, we investigated whether differences in mitochondrial concentrations and/or rate of Ca\(^{2+}\) elevation among cell types might be influencing these results.

**Mitochondrial Concentration and Rate of [Ca\(^{2+}\)], Elevation in Intact Neurons and Astrocytes**—Because the cellular mitochondrial concentration, and the rate of [Ca\(^{2+}\)], elevation may affect the time during which intact cells can buffer [Ca\(^{2+}\)], (more mitochondria or less Ca\(^{2+}\) = more time; less mitochondria or more Ca\(^{2+}\) = less time), we have assessed whether such parameters differ among neurons and astrocytes from cortical and striatal origin. Mitochondrial concentrations were measured as detailed under “Experimental Procedures” (e.g. \(\gamma - \delta\) in Fig. 3A (\(ii\), solid line; see also supplemental material). Analysis of TMRM\(^{+}\) fluorescence under matrix quenching conditions (50 nM; as demonstrated by the dequenching spikes in Fig. 4, A–D (\(ii\), solid lines) revealed a higher mitochondrial concentration in neurons than in astrocytes, expressed by a larger difference between the TMRM\(^{+}\) fluorescence before oligomycin and that remaining after FCCP, which reflects nonmitochondrial fluorescence (Figs. 3 and 4, A and C (\(ii\) versus B and D (\(ii\), solid lines)). Two-factor (region \(\times\) cell type) ANOVA of the mitochondrial concentration (Fig. 5B) revealed a significant main effect of the cell type (neurons \(\text{versus}\) astrocytes; \(F_{(1,16)} = 24.5; p < 0.001\)) and no significant effect of brain region (cortical \(\text{versus}\) striatal; \(F_{(1,16)} = 1.11; p = 0.308\)) nor of region \(\times\) cell type interaction (\(F_{(1,16)} = 0.253; p = 0.622\)). Hence, these results suggest that the lower “apparent” mitochondrial Ca\(^{2+}\) buffering capacity for astrocytes (Fig. 5A) is partly due to their lower mitochondrial concentration, when compared with neurons (Fig. 5B). Interestingly, despite the differences in the average area of cortical and striatal neuronal somas (\(\sim 1.2 \times 220 \pm 4 \text{ versus } 177 \pm 4 \mu\text{m}^2\), \(n = 526\) and 477 neurons, respectively) and especially between cortical and striatal astrocytes (\(\sim 2.5 \times 324 \pm 8 \text{ versus } 796 \pm 30 \mu\text{m}^2\), \(n = 472\) and 267 astrocytes, respectively), the mitochondrial concentrations in both types of...
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**A**

| Cortical neurons | Cortical astrocytes | Striatal neurons | Striatal astrocytes |
|------------------|---------------------|------------------|--------------------|
| ![Image](image1.png) | ![Image](image2.png) | ![Image](image3.png) | ![Image](image4.png) |

**B**

| Cortical neurons | Cortical astrocytes | Striatal neurons | Striatal astrocytes |
|------------------|---------------------|------------------|--------------------|
| ![Image](image5.png) | ![Image](image6.png) | ![Image](image7.png) | ![Image](image8.png) |

**C**

| Cortical neurons | Cortical astrocytes | Striatal neurons | Striatal astrocytes |
|------------------|---------------------|------------------|--------------------|
| ![Image](image9.png) | ![Image](image10.png) | ![Image](image11.png) | ![Image](image12.png) |

**D**

| Cortical neurons | Cortical astrocytes | Striatal neurons | Striatal astrocytes |
|------------------|---------------------|------------------|--------------------|
| ![Image](image13.png) | ![Image](image14.png) | ![Image](image15.png) | ![Image](image16.png) |

**FIGURE 4.** Intracellular Ca\textsuperscript{2+} dynamics in neurons and astrocytes from cortex and striatum under conditions where mitochondria are prevented from buffering [Ca\textsuperscript{2+}]. A–D, changes in [Ca\textsuperscript{2+}], (i) and in TMRM\textsuperscript{+} (ii) solid or dashed lines, respectively) fluorescence in arbitrary units (A.F.U.). The solid or dashed line in i and ii depicts measurements performed simultaneously in the same single cell. i, representative cells with identical [Ca\textsuperscript{2+}]\textsubscript{i} kinetics to allow comparisons in ii. Oligomycin (O; 3 \textmu M) and myxothiazol (M; 2 \textmu M) were added where indicated. FCCP (F; 1 \textmu M) addition and in situ determination of R\textsubscript{max} and R\textsubscript{min} (R and r, respectively) were performed at the indicated times for cells depicted by solid lines and slightly earlier for those shown by dashed lines (A and B, 10 min earlier; C and D, 5 min earlier). This time variation results from the need to anticipate and avoid excessive Fura-4F leakage to allow accurate [Ca\textsuperscript{2+}]\textsubscript{i} calibration. Assay buffer contained 10 mm pyruvate, and glucose was replaced by 2 mm 2-deoxy-D-glucose before the addition of oligomycin (3 \textmu M). A(i), K\textsubscript{exp} the rate of [Ca\textsuperscript{2+}]\textsubscript{i} elevation. Up to 50 random single cells of each type were analyzed per experiment; only a few are shown for clarity, being representative of n = 5 independent experiments.

neurons as well as in both types of astrocytes is remarkably similar (Fig. 5B). These results show that mitochondrial concentration is primarily governed by the cell type (neurons versus astrocytes) rather than by its region of origin (cortical versus striatal).

To test whether in situ mitochondria of the different cell types were being exposed to different rates of [Ca\textsuperscript{2+}]\textsubscript{i} elevation, we performed experiments similar to those depicted in Fig. 3 but with the further addition of myxothiazol (2 \textmu M) after oligomycin (Fig. 4). The underlying reasoning was that although oligomycin allows mitochondrial Ca\textsuperscript{2+} uptake, the combination of oligomycin and a respiratory chain inhibitor (myxothiazol) prevents it by collapsing \Delta\psi\textsubscript{m} (17), thus allowing a measurement of the rate at which [Ca\textsuperscript{2+}]\textsubscript{i} is rising inside cells, in close analogy with removing mitochondria from the “test tube” and measuring the rate at which Ca\textsuperscript{2+} is being infused in isolated mitochondria experiments (13). Indeed, with myxothiazol present, \Delta\psi\textsubscript{m} in both neurons and astrocytes rapidly collapsed (Fig. 4, A–D (ii), solid lines; TMRM\textsuperscript{+} dequenching), and the [Ca\textsuperscript{2+}]\textsubscript{i} elevation initiated with oligomycin progressed without interruption until near Fura-4F saturation, showing the absence of mitochondrial Ca\textsuperscript{2+} uptake (Fig. 4, A–D (i)). Compare with Fig. 3, A–D (i)). By means of nonlinear regression with Equation 2, as detailed under “Experimental Procedures,” we calculated the rate constant, K\textsubscript{exp} allowing comparisons of the rate of [Ca\textsuperscript{2+}]\textsubscript{i} elevation among cell types. Two-factor (region \times cell type) ANOVA of this parameter (Fig. 5C) revealed a significant region \times cell type interaction (F\textsubscript{(1,16)} = 14.5; p = 0.002) as well as a significant main effect of the cell type (neurons versus astrocytes; F\textsubscript{(1,16)} = 8.45; p = 0.010) and of the brain region (cortical versus striatal; F\textsubscript{(1,16)} = 11.2; p = 0.004). These results show that mitochondria in cortical neurons are exposed to a significantly higher rate of [Ca\textsuperscript{2+}]\textsubscript{i} elevation when compared with the other cell types (cortical astrocytes, striatal neurons, or striatal astrocytes) for which similar rates were observed. Hence, the absence of differences in mitochondrial Ca\textsuperscript{2+} buffering between cortical and striatal neurons depicted in Fig. 5A is only “apparent.” Indeed, in order to buffer for similar periods a faster rate of Ca\textsuperscript{2+} elevation with identical mitochondrial concentrations, cortical neurons should possess mitochondria with higher Ca\textsuperscript{2+} buffering capacities than striatal neurons.

To allow comparisons of the present in situ mitochondrial Ca\textsuperscript{2+} buffering data with what should be expected if the mito-
Mitochondria were isolated, we calibrated the “apparent values” for each cell type (Fig. 5A), dividing them by the respective average mitochondrial concentrations (Fig. 5B) and multiplying by the respective average rates of \([\text{Ca}^{2+}]_i\) elevation (Fig. 5C).

Two-factor (region × cell type) ANOVA of the calibrated values (Fig. 5D) revealed a significant main effect of the cell type (neurons versus astrocytes; \(F_{1,24} = 4.63; p = 0.042\); observed power = 54.2%) and of brain region (cortical versus striatal; \(F_{1,24} = 23.7; p < 0.001\); observed power = 99.7%), with no significant region × cell type interaction (\(F_{1,24} = 2.35; p = 0.138\)). The observed power for the main effect of cell type and of brain region was 87.1 and 98.4%, respectively. Meaningfully, these results provide original evidence that mitochondria not only in neurons but also in astrocytes from striatal origin exhibit a decreased \([\text{Ca}^{2+}]_i\) buffering capacity when compared with their cortical counterparts.

**Mechanisms Underlying in Situ Mitochondrial \([\text{Ca}^{2+}]_i\) Buffering Differences**—Mitochondria have been shown to interact with the endoplasmic reticulum (ER) in the regulation of intracellular \([\text{Ca}^{2+}]_i\) signaling (22). Thus, a possible mechanism underlying differences in in situ mitochondrial \([\text{Ca}^{2+}]_i\) buffering between cell types might involve heterogeneity in their ER \([\text{Ca}^{2+}]_i\) levels and/or ER-mitochondria \([\text{Ca}^{2+}]_i\) signaling. To test this hypothesis, we challenged cortical and striatal cells with 1 \(\mu\)M thapsigargin prior to the determination of mitochondrial \([\text{Ca}^{2+}]_i\) buffering capacity. Thapsigargin acutely and selectively arrests ER \([\text{Ca}^{2+}]_i\) pumps, depleting and preventing ER \([\text{Ca}^{2+}]_i\) storage (23, 24). We consistently observed transient \([\text{Ca}^{2+}]_i\) elevations following acute exposure to thapsigargin in astrocytes but less frequently in neurons (see supplemental Fig. 3 for representative tracings). Previous studies evidenced typically small \([\text{Ca}^{2+}]_i\) increases in neurons challenged with thapsigargin, detected with the high affinity Fura-2 probe (25), which may escape detection with the intermediate affinity Fura-4F in our experiments. Although this suggests that ER stores may differ in neurons and astrocytes, thapsigargin treatment did not significantly (\(p > 0.05\)) modify the mitochondrial \([\text{Ca}^{2+}]_i\) buffering capacity of each cell type (as a percentage of respective control: cortical neurons = 97.3 ± 5.8%, cortical astrocytes = 92.4 ± 2.3%, striatal neurons = 95.1 ± 2.2%, striatal astrocytes = 96.7 ± 5.8%; n = 3 independent experiments), arguing against a
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**DISCUSSION**

The present study is the first to examine whether mitochondria in neurons and astrocytes from cortex and striatum differ in Ca\(^{2+}\) buffering capacity. Here we describe a novel strategy allowing *in situ* measurements of mitochondrial Ca\(^{2+}\) buffering in intact cells. In addition, we present evidence that mitochondria in striatal neurons and astrocytes exhibit a decreased Ca\(^{2+}\) buffering capacity when compared with their cortical counterparts. Also, we show that differences between neuronal and astrocytic mitochondria are region-specific and that mitochondrial Ca\(^{2+}\) buffering differences between cortical and striatal astrocytes are abolished with 1 \(\mu M\) CsA. These results provide additional insight into the mechanisms of selective striatal neurodegeneration and into the relative Ca\(^{2+}\) vulnerability of neuronal and astrocytic mitochondria, as discussed below.

Mitochondria are high capacity Ca\(^{2+}\) buffering systems that play a key role in protecting cells from Ca\(^{2+}\)-induced damage (17, 30). Hence, differences in mitochondrial Ca\(^{2+}\) buffering are hypothesized to assist the selective vulnerability of cellular populations in the brain. A previous study, using mitochondria isolated from cortical and striatal homogenates, raised the hypothesis that striatal mitochondria might be “supersensitive to Ca\(^{2+}\),” thus contributing for selective striatal vulnerability in HD (2). However, data reflected a mixed nonsynaptic mitochondrial population in which the predominance of neuronal (26) or glial (31) mitochondria is debatable. The ensuing question of whether striatal mitochondrial vulnerability involves neuronal or glial mitochondria or both has so far remained unanswered.

Results in this study support the last possibility by showing that mitochondrial Ca\(^{2+}\) buffering capacity is decreased not only in striatal *versus* cortical neurons but also in striatal *versus* cortical astrocytes. This implies that striatal neurons, in addition to the vulnerability present in their own mitochondria, are paired with more vulnerable astrocytic partners than cortical neurons. Also, our finding that 1 \(\mu M\) CsA selectively increased mitochondrial Ca\(^{2+}\) buffering capacity in striatal astrocytes, rendering it identical to that of cortical astrocytes, without modifying neuronal mitochondrial Ca\(^{2+}\) buffering, has important implications. Indeed, it suggests that the ability of CsA to protect striatal neurons in *in vivo* HD models (32) could be the indirect consequence of reducing the mitochondrial vulnerability of their astrocytic neighbors.

Vulnerable astrocytes may aggravate the susceptibility of neuronal neighbors by failing to provide adequate support under pathophysiological conditions (9, 11, 33). Mitochondrial impairment in astrocytes may compromise protective astrocyte-neuron lactate trafficking and/or cause them to produce diffusible reactive oxygen species that damage nearby neurons (10, 34). Meaningfully, neurons rely on astrocytes for an efficient clearance of extracellular glutamate, an activity shown to be defective in HD astrocytes (35). We propose that deficits in this and other energy-dependent astrocytic activities will be aggravated in striatal astrocytes, since their Ca\(^{2+}\)-vulnerable mitochondria should more frequently arrest ATP synthase or even reverse its activity, thus decreasing ATP availability.

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**FIGURE 6.** Effect of cyclosporin A on the mitochondrial Ca\(^{2+}\) buffering capacity of intact neurons and astrocytes from cortex and striatum. Bars, mean and S.E. of “apparent mitochondrial Ca\(^{2+}\) buffering capacity,” assessed in experiments identical to those described in Figs. 3 and 5A, and performed in the absence (control; white bars) or presence of cyclosporin A (1 \(\mu M\) CsA, black bars). Data are from three independent experiments and were analyzed by *t* test comparison between CsA-treated cells and their respective controls, evidencing a significant (*p < 0.05*) effect in striatal astrocytes but not in other cell types (n.s., nonsignificant).

Major contribution of the ER in explaining differences in mitochondrial Ca\(^{2+}\) buffering between cortical and striatal neurons and astrocytes.

Differences in Ca\(^{2+}\) buffering between isolated brain mitochondria mixtures have been previously associated with the occurrence of permeability transition and/or the levels of mitochondrial cyclophilin D (26, 27). Thus, we tested whether cyclosporin A (CsA; an inhibitor of mitochondrial permeability transition that binds to cyclophilin D) influenced the differences in mitochondrial Ca\(^{2+}\) buffering exhibited by intact neurons and astrocytes. We performed experiments identical to those described in the legend to Fig. 3, both in the absence (control) and in the continuous presence of 1 \(\mu M\) CsA, including a 15-min preincubation period. Interestingly, results depicted in Fig. 6 showed that this CsA treatment significantly increased mitochondrial Ca\(^{2+}\) buffering capacity in striatal but not cortical astrocytes and not in neurons from either brain regions. Still, because CsA is also known to inhibit calcineurin, and thereby regulate mitochondrial phenomena such as fragmentation (28), we also tested the effect of FK506, which shares the property of calcineurin inhibition with CsA but has no effect on mitochondrial permeability transition (29). Cells were preincubated for 15 min with FK506, which was kept throughout the experiments at the previously used concentration of 0.6 \(\mu M\) (28). We found no significant differences (*p > 0.05*) in mitochondrial Ca\(^{2+}\) buffering between control cells and those where calcineurin was inhibited (as a percentage of respective control: cortical neurons = 96.4 ± 3.2%, cortical astrocytes = 95.8 ± 2.4%, striatal neurons = 103.9 ± 3.5%, striatal astrocytes = 93.6 ± 4.2%; *n* = 3 independent experiments). Taken together, these findings suggest that the selective effect of CsA on striatal astrocytes is unrelated to calcineurin inhibition but rather reflects inhibition of permeability transition. Thus, an increased propensity to undergo CsA-sensitive permeability transition in striatal astrocytes is a likely mechanistic explanation for their decreased mitochondrial Ca\(^{2+}\) buffering capacity when compared with cortical astrocytes.
ATP synthase reversal allows mitochondria to delay $\Delta\psi_m$ collapse, provided that cells have sufficient glycolytic ATP. Thus, mitochondria may appear to be more resistant to injury in cells with abundant glycogen stores, as we have previously shown for HD cellular models (8). Given the limited glycogen availability in neurons as opposed to astrocytes (36), the phenomenon of earlier mitochondrial damage in astrocytes than in neurons, thus acting as a major confounding factor when comparing mitochondrial function in these cell types. To avoid this, we inhibited the use of glycogen stores and ATP synthase reversal in our experiments with intact neurons and astrocytes. For the purpose of data comparison, it should be noted that glycogen stores are typically unavailable in isolated mitochondria and when comparing synaptic with nonsynaptic fractions (26, 27).

Isolated brain mitochondria are mixtures of neuronal and glial mitochondria. These mixtures are heterogeneous in response to $Ca^{2+}$, as shown within nonsynaptic fractions (37), and when comparing synaptic with nonsynaptic fractions (26, 27), strengthening the hypothesis that neurons and glia differ in $Ca^{2+}$ buffering capacity. Presently, it is still unfeasible to isolate functional neuronal and glial mitochondria from the adult brain, narrowing the pursuit of this hypothesis to studies with primary cultures of brain cells. A previous study addressed this hypothesis using suspensions of permeabilized cells and evidenced that mitochondria in cerebellar neurons (1 DIV) buffered less $Ca^{2+}$ than in cortical astrocytes (10 DIV) (38). However, given the different regions under study for neurons (cerebellum) and astrocytes (cortex), it cannot be generalized that neuronal mitochondria buffer less $Ca^{2+}$ than astrocytic mitochondria. Indeed, within the same region (cortex or striatum), we saw the opposite, namely that neuronal mitochondria buffered more $Ca^{2+}$ than astrocytic mitochondria. Still, this did not apply across regions, since mitochondria in striatal neurons did not buffer more $Ca^{2+}$ than in cortical astrocytes, highlighting the cellular and regional specificity of mitochondrial $Ca^{2+}$ buffering differences.

Addressing the mechanisms underlying mitochondrial $Ca^{2+}$ buffering differences and considering possible mitochondria-ER interactions in $Ca^{2+}$ signaling (22), we tested whether selectively depleting and arresting ER $Ca^{2+}$ pumps with thapsigargin modified in situ mitochondrial $Ca^{2+}$ buffering. Because thapsigargin did not significantly affect mitochondrial $Ca^{2+}$ buffering in any of the four cell types, it is unlikely that ER heterogeneity explains their differences in mitochondrial $Ca^{2+}$ buffering capacity. This finding by no means undermines the physiological relevance of ER-mitochondria $Ca^{2+}$ signaling, particularly at low $[Ca^{2+}]_i$, being probably explained by the wide disparity in the amounts of maximal $Ca^{2+}$ accumulation by the two organelles and by ER saturation occurring before $[Ca^{2+}]_i$ reaches the set point for mitochondria net uptake. Although intact cells may differ in $Ca^{2+}$ handling mechanisms other than the mitochondria (e.g. pumps, buffers, and exchangers), in our experimental conditions, we have introduced calibrations for the rate of $Ca^{2+}$ elevation ($K_{exp}$) in the different cell types when mitochondria $Ca^{2+}$ buffering was selectively inhibited, which yielded a measurement of net $[Ca^{2+}]_i$ elevation considering $Ca^{2+}$ handling mechanisms (entry and extrusion pathways) other than the mitochondria.

Alternative mechanisms may involve a differential mitochondrial sensitivity to permeability transition. We tested whether mitochondrial $Ca^{2+}$ buffering was modified by CsA (inhibitor of permeability transition and calcineurin), and by FK506 (calcineurin inhibitor). The absence of effect of FK506 suggests that calcineurin activity does not modify mitochondrial $Ca^{2+}$ buffering, possibly because calcineurin activation is subsequent to mitochondrial depolarization (28), when the driving force for $Ca^{2+}$ buffering is exhausted. Moreover, it highlights the inhibition of permeability transition rather than calcineurin as the prevailing effect of CsA in our experiments. Therefore, given that 1 $\mu M$ CsA abolished the differences in in situ mitochondrial $Ca^{2+}$ buffering between cortical and striatal astrocytes, our results show that mitochondria in striatal astrocytes exhibit an increased propensity to undergo a CsA-sensitive form of permeability transition. The lack of a CsA (1 $\mu M$) effect over the other cell types suggests the occurrence of CsA-insensitive forms of permeability transition (39, 40). However, it is also possible that higher concentrations of CsA might be required for an effect to be observable, as suggested from recent data with isolated synaptic mitochondria, where 5 $\mu M$ but not 1 $\mu M$ CsA increased mitochondrial $Ca^{2+}$ buffering capacity (27). Unfortunately, testing of this possibility in intact neurons is hampered by their vulnerability to greater than 1 $\mu M$ CsA concentrations (39, 41). The need for high CsA concentrations in order to inhibit mitochondrial permeability transition may be associated with high levels of the CsA target, cyclophilin D (CypD) (27). Indeed, CypD levels were shown to be inversely correlated with the mitochondrial resistance to permeability transition, namely throughout brain development (42). Meaningfully, the highest levels of brain CypD protein were found in 1-day-old rats and progressively decreased, so that at 10 days of age, CypD expression became nearly identical to that of mature 3-month-old animals (Fig. 4d in Ref. 42). These findings suggest that the relative differences in mitochondrial $Ca^{2+}$ vulnerability between 1 DIV cerebellar neurons and 10 DIV cortical astrocytes (38) might change considerably when both cell types have 10 DIV. Moreover, the stabilization of CypD expression after 10 days of age (42) supports the modeling of brain mitochondrial $Ca^{2+}$ buffering with mature cultures of brain cells (11–13 DIV in the present study). Indeed, sufficient time in culture is essential for neurons to develop a network of processes containing mitochondria that is probably heterogeneous with respect to that in the cell body (43–45).

The decreased mitochondrial $Ca^{2+}$ buffering of striatal versus cortical neurons in the present study reflects neuronal mitochondria as a whole. In light of recent findings with synaptic versus nonsynaptic cortical mitochondria (26), it would be interesting to compare the $Ca^{2+}$ vulnerability of synaptic mitochondria in striatal and cortical neurons, since it may contribute to differences in the present study and influence pathological synaptic dysfunction in HD (46) as well as the greater susceptibility of striatal neurons to toxic insults (1, 6, 47). We recently compared the distribution and trafficking of in situ mitochondria within the processes of cortical and striatal neurons and found similar fractional occupations but faster mito-
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Mitochondria movement in cortical neurons. Although similar mitochondrial fractional occupation in processes is in agreement with the similar overall mitochondrial concentrations reported in the present study, the differences in mitochondrial movement speed raise the interesting hypothesis that mitochondria in cortical neurons may more rapidly and efficiently handle $Ca^{2+}$ buffering needs in distant processes by means of faster trafficking.

In summary, the present study shows for the first time that the increased vulnerability of striatal versus cortical mitochondria to $Ca^{2+}$ loads resides in both intact neurons and astrocytes. Mitochondria in striatal astrocytes have an increased propensity to undergo Ca-sensitive permeability transition, positioning the striatum at greater risk for disturbed neuron-astrocyte interactions and probably contributing to selective striatal demise in HD and other neurological disorders. The selective effect of $1\,\mu M$ Ca$^{2+}$ over striatal astrocytes suggests that in vivo neuronal shielding with this compound may indirectly result from astrocytic protection.

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