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Minimum conditions for the induction of cortical spreading depression in brain slices


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Submitted 14 March 2014; accepted in final form 12 August 2014

CORTICAL SPREADING DEPRESSION (CSD) is a fundamental response to brain injury. A propagating wave of massive gray matter depolarization, CSD has been incontrovertibly shown to occur in human brain insults, including stroke, subarachnoid hemorrhage, and trauma (Lauritzen et al. 2011). CSD is also highly conserved in phylogeny, occurring in all mammals tested, as well as birds, reptiles, fish, and amphibians (Marshall 1959; Young 1980). CSD-like events even occur in insects in situations of systemic stress, where they appear to convey an adaptive advantage (Rodgers et al. 2007; Armstrong et al. 2009).

Yet, there is very strong circumstantial evidence that CSD also occurs in migraine, a condition in which no apparent brain injury or massive systemic stress occurs (Olesen et al. 1981; Woods et al. 1994; Cao et al. 1999; Bowyer et al. 2001; Hadjikhani et al. 2001). Moreover, unlike in stroke, subarachnoid hemorrhage, or trauma, humans undergoing migraine aura are conscious and able to report their experience (Russell and Olesen 1996). This implies that CSD must be inducible under conditions of relative physiological normalcy.

Previous models of CSD, however, have largely been limited to massive ionic, metabolic, or traumatic insults that would not be compatible with the awake, behaving, and apparently uninjured state of migraine (Somjen 2001; Charles and Brennan 2009). It is important to know whether there are conditions that would allow induction of CSD without overt injury.

A first step in understanding the “naturalistic” induction of CSD is to systematically study conditions that will set off this all-or-none event. The pioneering work of Matsuura and Bureš (1971) estimated that a surprisingly large volume of tissue, ~1 mm³, might be required to initiate CSD in rat brains in vivo under high-potassium solution stimulation. However, their approach necessarily involved unintended damage to superficial cortical layers and lack of explicit control of stimulation areas. This may have led to significant overestimation of “minimum” tissue volumes required for CSD induction.

We previously reported the development of a microfluidic device that allows precise focal control of chemical environments of a brain slice (Tang et al. 2011). By locally applying solutions of different concentrations of potassium ions over different areas of the cortical layers of mouse brain slices, we study conditions necessary for CSD induction. We find that CSD is inducible under a wide range of physiologically plausible parameters, with smaller stimulation areas necessary during exposure to extracellular potassium levels ([K⁺]ₑ) expected in trauma and significantly larger stimulation areas necessary under [K⁺]ₑ expected without overt injury. Our data thus define a set of conditions upon which the physiologically realistic modeling of CSD (in silico, in vitro, and in vivo) may proceed.

MATERIALS AND METHODS

Ethical approval. Experimental protocols were approved by the Animal Research Committees of University of California, Los Angeles, California and the University of Utah.

Brain slice preparation and maintenance. Male C57Bl/6 mice (age: 1–3 mo; n = 22) were deeply anesthetized with isoflurane and decapitated. The brain was rapidly removed and immersed in an ice-cold high-sucrose low-sodium artificial cerebrospinal fluid (ACSF) solution of the following composition (in mM): 216 sucrose, 2 KCl, 2 MgSO₄, 1.25 NaH₂PO₄, 2 CaCl₂, 1 MgCl₂, 26 NaHCO₃, and 10 glucose bubbled with 95% O₂-5% CO₂. The brain was then glued to a stage, and coronal slices of 400-μm thickness were cut in an ice-cold
oxygenated sucrose solution using a vibratome (MA752 Motorised Advance Vibroslice; Campden Instruments). The slices were then incubated in an oxygenated ACSF solution (in mM: 125 NaCl, 3 KCl, 1.25 NaH₂PO₄, 2 CaCl₂, 1 MgCl₂, 25 NaHCO₃, and 10 glucose) at room temperature for at least 1 h before the experiments.

**Microfluidic brain slice chamber.** The microfluidic device and experimental setup is shown in Fig. 1. The device, described in detail in our prior work (Tang et al. 2011), includes embedded microchannels, microposts, and fluidic ports made of polydimethylsiloxane (PDMS). These microstructures are designed to mechanically support a brain slice and focially deliver a specific chemical stream. The outer chamber walls allow a continuous bath perfusion of ACSF to maintain slice viability.

The slice rests on a network of microposts that allow bath perfusion of the underside as well as the top surface of the brain slice. At select locations, the posts are replaced by fluidic port assembly. For each assembly, one application port is surrounded by several suction ports to create a confined chemical plume, which delivers chemicals into a brain slice by diffusion.

Underneath the microposts and ports are microchannels, which connect to an injection pump and a suction pump for fluidic delivery and removal, respectively. Sections of silicone tubing (ID: 1/8") are bonded to the inlets and outlets of the microchannels. A syringe pump (sp100i Pump; WPI, Sarasota, FL) is used to deliver an ACSF solution with a set potassium concentration ([K⁺]) into each application port. A vacuum pump (BUCHI V-700, New Castle, DE) with a feedback pressure control module is used to adjust suction flows (mean velocities of the order of 0.01 m/s) and achieve a controlled confined plume of a potassium-rich ACSF solution.

**Plume control and size determination.** Figure 1C shows images of the plume of an ACSF solution labeled with fluorescein over a fluidic port with and without suction, demonstrating that we can successfully confine the plume by a purely fluidic control without requiring any physical seals. The images are shown without a brain slice placed over the fluidic port for clarity, but the creation of a confined plume in the presence of a brain slice was also demonstrated as discussed in previous work (Tang et al. 2011; Fig. 2C).

To determine conditions for CSD induction in terms of the area of de polarization, we used fluidic ports of different radii. The radius of each port was defined as the distance from the port center to the inner boundary of the suction ports (Fig. 2A, top inset). The port radii used in the present work were 0.1, 0.15, 0.2, 0.25, 0.36, and 0.5 mm.

Because of the complexity of light diffraction and scattering within a brain tissue, we empirically determined a correlation between the plume radius under a brain slice and fluorescence images using partially exposed fluidic ports as illustrated in Fig. 2A. The plume radius correlated well with the physical port radius, falling within the range defined by the inner and outer boundaries of the suction ports shown as the dashed lines.

We report normalized plume radius throughout. Plume area was calculated by measuring the area above 50% pixel intensity and using the circle area equation to derive an equivalent or normalized radius to allow comparison. Normalized radius was used because in seven experiments (1 with 0.15-mm port size, 1 with 0.2-mm port size, 1 with 0.25-mm port size, 2 with 0.36-mm port size, and 2 with 0.5-mm port size), port size and positioning were such that the plume boundary was outside the slice boundary. In these cases, the area of the plume over the slice was not circular.

**Induction and optical recording of CSD in brain slices.** Brain slices were transferred from a tissue culture incubator to the microfluidic device maintained at 32 ± 1°C, with a continuous ACSF bath perfusion bubbled with 95% O₂-5% CO₂. This bath solution was delivered to the chamber using gravity at 2 ml/min. A piece of filter paper was placed at the bath outlet to minimize flow fluctuations. The perfusion rate was verified using an analytic balance, and the chamber temperature was monitored and kept within 32 ± 1°C throughout the experiment.

CSD was induced by applying a high-[K⁺] oxygenated ACSF solution containing different molarities of KCl substituted for equimolar NaCl. A small amount of fluorescein (< 0.1%) in the ACSF solution allowed visualization of the plume (Fig. 2C). If no CSD was observed for the first set value of the potassium ion concentration, the concentration was increased in steps of 2–10 mM until we reached a threshold concentration. The molarities were 10, 12, 15, 25, 30, 40, 60, 70, 80, 120, 130, and 140 mM above normal ACSF [K⁺]; thus total [K⁺] was 13, 15, 18, 28, 33, 43, 63, 73, 83, 123, 133, and 143 mM. Once a CSD event was observed in these ascending thresholding experiments, we conducted a confirmative experiment by returning to the lower concentration from a prior step and verifying that no CSD would be induced below the detected threshold. To ensure that there was not a systematic bias inherent to our ascending thresholding procedure, we also performed reverse experiments where we began at high values of [K⁺] applied on the slice surface and then decreased the applied potassium concentration step by step until a minimum threshold level was reached. We observed no difference in the thresholds obtained using the two procedures.

At least three experiments were conducted for each combination of port size and applied potassium ion concentration. To consider a limiting case of an infinitely large application port, additional exper-

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**Fig. 1.** Microfluidic slice chamber allowing controlled focal depolarization. A: brain slice rests on posts in a modified submerged slice chamber. Posts allow bath perfusion both above and below the slice. Selected posts are replaced by microfluidic ports, consisting of 1 application port and 4 suction ports. *Inset* shows a microfluidic fluidic port, and dashed arrows represent application flow, solid arrows represent suction flow. Use of the suction ports allows a plume of high-K⁺ concentration ([K⁺]) solution to be precisely confined. Ports are made of varying sizes, and concentration of perfusion solution is varied. B: schematic of the experimental setup showing the chamber integrated with epifluorescence imaging. C: schematic view of 1 application/suction port (left) and fluorescence images of the applied flows with suction (right, top) and without (right, bottom). Scale bar = 0.1 mm. PDMS, polydimethylsiloxane.
imments were conducted where a potassium-rich ACSF solution was applied over an entire slice via bath perfusion rather than through a microfluidic port.

During each experiment, the applied potassium concentration was maintained at a set value for 2 min or until CSD was observed, whichever was shorter. The fluidic port was then flushed with a solution of oxygenated ACSF at a normal potassium ion concentration, and the slice was allowed to rest for 10 min before retesting. At the end of each experiment, the viability of the slice was confirmed by local injection and suction rates were adjusted so that the resulting slice bottom plume area matched with those measured in thresholding experiments. Cellular uptake or other active transport mechanisms were not considered.

Indirect confirmation of the mass transport model using two-photon microscopy. We performed two-photon microscopy experiments to determine the concentration profile of fluorescein isothiocyanate-dextran molecules (FITC-dextran, 3,000–5,000 molecular weight; Sigma; dissolved in ACSF) applied to a brain slice using one microfluidic port size (0.25-μm radius) and compared the experimental results with our model prediction.

Experiments were performed using a Sutter movable objective microscope coupled to a Ti Sapphire laser (MaiTai HP, Spectra Physics, CA) tuned to 810 nm for FITC-dextran and sulfrodhamine 101 excitation. A high numerical aperture water-immersion objective (Carl Zeiss W Plan-Apochromat 101/1.40 DIC) was used. When necessary, two channels were simultaneously acquired using Chroma filters [535/50 nm (green) and 610/75 nm (red)]. Scanimage (Pologruto et al. 2003) was used to control image acquisition.

Three-hundred-micrometer-thick brain slices were incubated in an ACSF solution with 1 μM sulfrodhamine 101 (SR101; Invitrogen, Carlsbad, CA) added at 32°C for at least 30 min. SR101 labels astrocytes throughout the thickness of a slice (Nimmerjahn et al. 2004) and thereby allows anatomical registration of two-photon images. Before each optical scan, the location of the slice top surface was first determined by lowering the microscope focus until the first image of where SR101 stained astrocytes (red) became visible. Images were acquired with a z-axis step size of 10 μm from the first image where astrocytes were identified down to the bottom of the slice.

Fluorescence intensity decays with depth z due to light scattering and absorption by the slice (Helmchen and Denk 2005), which can be modeled using the Beer-Lambert law:

$$ F_{2PLSM} \propto P_0 \times e^{-z/s_l} $$

Here $F_{2PLSM}$ is the fluorescent intensity, $P_0$ is the laser power applied, $z$ is the depth, and $s_l$ is the mean free path describing the scattering properties of the tissue (Helmchen and Denk 2005). The fluorescence
RESULTS

Reliability of CSD induction and recording. CSD waves were successfully induced by locally applying a solution of potassium rich ACSF onto cortical layers of mice brain slices. CSD was observed as an approximately concentric propagating wave of decreased, followed by increased, light reflectance (Fig. 2C), propagating out from the port site at a speed of \( \sim 3 \) mm/min (median: 3.0, 25, 75 percentile 2.3, 3.7 mm/min).

Approximately three to four CSD waves could be induced in a given slice at each test location over periods >1 h. In preliminary experiments we also used direct current field stimulation condition with balanced suction and application flows of potassium ions to verify that the optical signal is a strong correlate of the surface area \( (r^2) \) of the brain slice that is exposed to high [K\(^+\)].

The minimum concentration required to evoke CSD was \( \sim 15 \) mM, which was achieved with bath perfusion. An extrapolation of Eq. 2 yields the minimum plume radius corresponding to this concentration of \( \sim 0.57 \) mm at this concentration.

Due to a finite pumping speed, it took (means ± SD) 17 ± 8 s for the plumes to reach 90% of the final stimulation areas. CSD was induced relatively quickly thereafter, on average after 12 s (mean CSD initiation time 29 ± 8 s). There was no explicit dependence of time-to-CSD-induction on the plume size or potassium ion concentration.

Minimum tissue volume for CSD induction. CSD both in vitro and in vivo is a three-dimensional phenomenon; its initiation thus occurs in a volume of tissue, whose parameters constrain the physiological mechanisms of induction. We used our plume area/concentration data and numerical simulation (Tang et al. 2011), validated with two-photon microscopy plume imaging, to estimate the minimum depolarized tissue volumes necessary for CSD initiation (Fig. 4).

To account for the hindered diffusion due to ion movements in tortuous pathways within the brain slice, the diffusivity of K\(^+\) in free medium \( (2 \times 10^{-9} \text{ m}^2/\text{s}) \) is reduced to represent the apparent diffusion coefficient (ADC),

\[
D_{\text{Apparent}} = \frac{D}{\lambda^2}
\]

Here \( \lambda \) is the tortuosity of the brain extracellular space. The value of \( \lambda \) employed in our simulation was either 1.8 or 2.2, which are the upper and lower limits of ADC reported in independent previous studies (Nicholson 1993; Nicholson and Tao 1993; Binder et al. 2004; Syková and Nicholson 2008; Zador et al. 2008).

Figure 4A uses these parameters to generate an expected plume profile through a 400-µm brain slice over time; \( y \) axis is the distance from slice bottom; \( x \) axis is the shows extracellular [K\(^+\)], relative to the slice surface. Traces are at 10-s intervals. There is a clear time dependence to plume concentration that needs to be incorporated for precise plume modeling. Note that the depth of the affected cortical volume is determined primarily by the diffusion of potassium ions into a brain slice, which is in turn a function of the time. Since the onset of CSD
Minimum cortical volumes required for CSD induction. A: time-dependent K⁺ plume modeling. Predicted extracellular potassium concentration is plotted across the slice thickness at the port center location as a function of time. The concentration at each location increases as a function of time due to continuous supply of high-[K⁺] solution at the slice bottom surface. A schematic of the slice model is shown in the inset. Concentration is plotted along the dotted line above the port center. B: validation of model with two-photon microscopy. Left: predicted concentration profiles superimposed with isoconcentration contours from 10 to 90% of the applied solution concentration. Right: experimentally determined fluorescent intensities across the thickness of a slice. The experimental profile is truncated by the presence of dead cell layers on both surfaces of the slice that did not result in fluorescence signals from SR101 staining. FEM, finite element method; 2PLSM, 2-photon laser scanning microscopy. Scale bar = 0.1 mm. C: minimum tissue volumes for CSD induction. Contour lines show expected volumes of K⁺ plume at 50 and 10% of the slice surface concentration. Contours are plotted for minimum (D₀₉₅, dots) and maximum (D₀₉₀, dashes) diffusion constants reported in the literature. Expectation is that CSD volumes would fall within these contours; closer to 50% line. Experimentally determined plume radii at different applied [K⁺] were used to predict the volume of tissue expected to induce CSD. We used experimentally determined plume area at each experimentally applied [K⁺] to predict the volume of tissue >15 mM [K⁺]. Our reasoning for using a 15-mM cutoff is that this was the minimum concentration able to induce CSD; we would not expect the volumes enclosed by lower concentrations to be relevant for induction. For each diffusion coefficient (D₀₉₅, black; D₀₉₀, white), the mean and standard deviation tissue volumes and applied [K⁺], for volumes of tissue predicted to be >15 mM, are plotted. The area bounded by the error bars (shaded) represents the expected range of CSD ignition volumes from our experiments. The range of predicted volume of tissue necessary to induce CSD at 15 mM is 0.03–0.06 mm³. At a cutoff of 30 mM (the lowest value achieved by a circular focal plume rather than bath perfusion), the range of predicted volume is 0.005–0.012 mm³.

DISCUSSION

CSD is suggested as the physiological mechanism behind the migraine aura. However, it is surprising that such a massive ionic and metabolic event, incontrovertibly seen in conditions of brain injury, could be generated in the awake uninjured brain. CSD does not possess a single threshold; like other phenomena [e.g., the action potential (FitzHugh 1955)] the threshold for all-or-none activity is a multidimensional surface (manifold) (Dahlem and Isele 2013).
to sample this surface as completely as possible in a brain slice preparation.

We designed our experiments to address the feasibility of CSD under physiologically realistic conditions. In contrast to most CSD models (Aitken et al. 1998; Ayata et al. 2006; Brennan et al. 2007) we focused on \([K^+]_e\) that could be expected in vivo, in a range whose maximum was approximate intracellular \([K^+]\). This maximum concentration is relevant to situations such as brain trauma or stroke, where cellular lysis occurs. The lower limits of the range were designed to test the “K⁺ ceiling,” levels of extracellular K⁺ ranging from 10 to 20 mM, beyond which both neuronal and vascular behavior changes dramatically (Somjen et al. 1976; Heinemann and Lux 1977; Somjen 1979; Knot et al. 1996). These concentrations are achievable in vivo during pathological states, such as seizure (Heinemann et al. 1977; Heinemann 1986), that are recoverable without apparent cellular injury. Intermediate levels (30–80 mM) are seen during passage of the CSD wave (Somjen 2001) and are also likely relevant to tissue in the vicinity of brain injury such as a contused or infarcted region.

Matsuura and Bureš (1971) estimated the minimum radius of depolarized tissue necessary for CSD induction to be approximately 600–960 μm and the minimum \([K^+]_e\) between 12 and 45 mM. No direct correlation between the two quantities was reported.

The range of minimum concentrations reported in the previous study is consistent with our results. However, a minimum affected tissue volume calculated from the minimum radii reported by Matsuura and Bureš is 0.9–3.7 mm³. This is much larger than our estimated values of ~0.03–0.06 mm³.

Such discrepancy may in part be due to differences in preparation (rat vs. mouse brain, in vivo vs. in vitro). Perhaps more importantly, the insertion of a 200-μm diameter capillary in the previous study was likely to have caused damage to superficial cortical layers, which are believed to be most sensitive to depolarization and therefore CSD induction (Marshall et al. 1950; Leão 1951; Világi et al. 2001).

In contrast to these in vivo methods, an advantage of our preparation is that we were able to directly control both the concentration of the potassium ions applied to a cortical surface and the focal area exposed to the depolarizing stimuli. We were also able to directly observe the induction of CSD and precisely measure the surface area of tissue exposed to the depolarizing stimulus. By directly observing and modeling fluid transport in slices, we were also able to predict the minimum volume depolarized with greater confidence.

Our results show that CSD can be induced in small volumes of tissue exposed to high physiological-range \([K^+]_e\) as well as larger volumes exposed to lower concentrations. Our combined data show that the relationship between \([K^+]_e\) and plume area is approximately constant. This implies that susceptibility to CSD might be determined by the total mass of K⁺ in the extracellular space. The question then arises whether there is a true equivalence between relatively high-[K⁺]/small-radius experiments and low-[K⁺]/large-radius experiments: i.e., whether high-[K⁺]/small-radius stimuli elicit CSD because they are actually generating low-[K⁺]/large-radius area of stimulation due to diffusion. Our data suggest this is not the case because high-[K⁺]/small-radius experiments did not differ in time to CSD induction from low-[K⁺] large-radius experiments. If a diffusion-generated large low concentration plume were being generated, high-[K⁺]/small-radius experiments would have longer induction times.

At \([K^+]_e\) below ~30 mM, the relationship between \([K^+]_e\) and plume radius flattened. For any \([K^+]_e\) in this range, a larger area of exposed cortex was required to generate CSD. This change in behavior may reflect a \([K^+]_e\) buffering capacity that is operant below, and exceeded above ~30 mM. It could also reflect the fact that beyond 0.18-mm port radius, our circular port assemblies overlapped the boundaries of the slice, potentially increasing variability. Finally at \([K^+]_e\) <15 mM, CSD was not elicitable at any area of stimulation including whole slice superfusion. At 15 mM the relationship between \([K^+]_e\) and area became completely flat, revealing an absolute minimum threshold for CSD induction. Presumably below this \([K^+]_e\) buffering mechanisms can counter any increases, at least for the duration of \([K^+]_e\) elevation in our experimental paradigms.

To summarize, our data suggest a \([K^+]_e\) buffering capacity that at concentrations <15 mM is intact, that at concentrations <30 mM is challenged but still operant, and that at concentrations >30 mM is essentially saturated. It is likely that different buffering mechanisms are at play.

For computational simplicity and limitation in real-time concentration measurement capabilities, our model considered diffusion as the only mechanism for K⁺ ion distribution in the extracellular space. Other physiological K⁺ regulation mechanisms need to be considered to fully predict the minimum condition for CSD induction: 1) K⁺ release from neuronal and glial activity; 2) active K⁺ uptake by neurons and glial cells; 3) K⁺ spatial buffer by glial cells; and 4) passive KCl uptake by glial cells (Kraig and Nicholson 1978; Largo et al. 1996; Holthoff and Witte 2000; Walz 2002; MacAulay and Zeuthen 2012). The relative importance of the mechanisms varies based on specific experiment conditions and different brain regions (Kofuja and Newman 2004). Moreover, the relative importance of different mechanisms likely differs at different \([K^+]_e\). When \([K^+]_e\) is below ~12–15 mM, active potassium uptake by neurons and glial cells likely plays a significant role in clearing potassium from the extracellular space (Cordingley and Somjen 1978; Xiong and Stringer 2000). Under higher \([K^+]_e\), the active uptake of potassium likely saturates (Baker 1968; Walz and Hertz 1982); findings consistent with our results. Finally, experimental conditions could also have affected our results and simulations: 1) slice thickness; 2) vertical distance between plume origin and slice; 3) spatial location of the center of the K⁺ plume (as different cortical layers have different CSD susceptibility; Marshall et al. 1950; Leão 1951; Világi et al. 2001); 4) correspondence of fluorescent marker concentrations to \([K^+]_e\); and 5) microfluidic device design.

The control we achieve over the ionic and fluidic environment was not possible without a brain slice preparation. However, an obvious limitation of this approach is that the vascular changes that accompany and likely influence CSD were not sampled. Vascular activity is important because of its homeostatic role in vivo and because cerebral arterioles constrict rather than dilate at \([K^+]_e\) that induce CSD (Knot et al. 1996). This means that vascular delivery and clearance of metabolites are compromised at precisely the levels of \([K^+]_e\) where neurons and glia are entering a critical transition to depolarization. Implications will need to be tested in vivo. One might predict
that minimum volumes of depolarization could actually be lower in the presence of a functional vasculature. On the other hand, vascular clearance of potassium ions could slow the approach of $[K^+]_e$ to threshold values and thus have opposite effects. Our use of a mouse model may also not completely mimic conditions expected in humans. Mouse neural and cellular density is greater than that of humans, and the neuron/glia ratio is higher (Tsai et al. 2009). The implication is that mouse tissue might support CSD at smaller depolarized volumes than human tissue. The fact that mice are lissencephalic while humans are gyrencephalic is likely not relevant to our slice experiments but has implications for CSD induction and propagation (Dahlem and Hadjikhani 2009).

If CSD occurs in awake behaving humans, as is proposed for the migraine aura, it must occur under conditions that do not result in either incapacitation or lasting injury. The volume of depolarized tissue determined from our experiments is small enough that transient, focal alterations in perfusion or metabolism might be sufficient to set off CSD. In this context it is interesting to note that our predicted minimum volumes of depolarized tissue are similar to the predicted volume of a cortical column (Mountcastle 1997) or penetrating artery territory (Woolsey et al. 1996). One could speculate that a discrete sensory column exposed to excessive stimulation, or a single penetrating artery undergoing reversible spasm, might be sufficient to cause CSD without causing lasting tissue damage. Conclusions. We have used a microfluidic device in mouse brain slice preparation to evaluate the minimum conditions necessary to induce CSD. We find that CSD is inducible under a range of conditions including those likely to be encountered in brain injury and those likely during awake behaving states such as migraine.

ACKNOWLEDGMENTS

We thank Dr. Andrew Charles for use of equipment. Present address of H. E. López-Valdés: Universidad Nacional Autonoma de Mexico, Av Universidad 3000, Ciudad Universitaria, 04510 Ciudad de México, Distrito Federal, Mexico. GRANTS

This work was supported by the National Institute of Neurological Disorders and Stroke Grants NS-059072 and NS-070084 (to K. C. Brennan) and Department of Defense Grant CDMRP PR 100060 (to K. C. Brennan). DISCLOSURES

No conflicts of interest, financial or otherwise, are declared by the author(s).

AUTHOR CONTRIBUTIONS


REFERENCES


