CaV1.2/CaV3.x channels mediate divergent vasomotor responses in human cerebral arteries

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The regulation of arterial tone is critical in the spatial and temporal control of cerebral blood flow. Voltage-gated Ca2+ (CaV) channels are key regulators of excitation–contraction coupling in arterial smooth muscle, and thereby of arterial tone. Although L- and T-type CaV channels have been identified in rodent smooth muscle, little is known about the expression and function of specific CaV subtypes in human arteries. Here, we determined which CaV subtypes are present in human cerebral arteries and defined their roles in determining arterial tone. Quantitative polymerase chain reaction and Western blot analysis, respectively, identified mRNA and protein for L- and T-type channels in smooth muscle of cerebral arteries harvested from patients undergoing resection surgery. Analogous to rodents, CaV1.2 (L-type) and CaV3.2 (T-type) α1 subunits were expressed in human cerebral arterial smooth muscle; intriguingly, the CaV3.1 (T-type) subtype present in rodents was replaced with a different T-type isofrom, CaV3.3, in humans. Using established pharmacological and electrophysiological tools, we separated and characterized the unique profiles of Ca2+ channel subtypes. Pressurized vessel myography identified a key role for CaV1.2 and CaV3.3 channels in mediating cerebral arterial constriction, with the former and latter predominating at higher and lower intraluminal pressures, respectively. In contrast, CaV3.2 antagonized arterial tone through downstream regulation of the large-conductance Ca2+-activated K+ channel. Computational analysis indicated that each Ca2+ channel subtype will uniquely contribute to the dynamic regulation of cerebral blood flow. In conclusion, this study documents the expression of three distinct Ca2+ channel subtypes in human cerebral arteries and further shows how they act together to orchestrate arterial tone.

INTRODUCTION

The spatial and temporal distribution of brain blood flow is set by networks of resistance arteries. Cerebral arteries are responsive to a range of humoral and mechanical stimuli (Furchgott and Zawadzki, 1980; Segal, 2000; Filosa et al., 2006), one of the key being arterial pressure (Knot and Nelson, 1998). Under dynamic conditions, elevated intraluminal pressure initiates arterial constriction by modulating ionic conductances that depolarize arterial smooth muscle membrane potential (Vm) (Knot and Nelson, 1998; Welsh et al., 2000, 2002). This electrical event in turn grades the opening of voltage-gated Ca2+ (CaV) channels, and the subsequent rise in cytosolic [Ca2+]i triggers the activation of myosin light chain kinase and actin–myosin cross-bridge cycling (Gallagher et al., 1997; Cole and Welsh, 2011). As CaV channels play a key role in setting arterial tone, these channels have long been a target of therapeutic interest, with blockers commonly used to moderate systemic hypertension and cerebral vasospasm (Godfraind, 2014).

The CaV channel family comprises 10 α1 pore-forming subtypes, among which the CaV1.2 channel predominates in the transduction of vascular smooth muscle contractility (Catterall, 2011). Given the robust vasodilatory response to L-type blockers (e.g., dihydropyridines), it is often assumed that this channel is the sole subtype expressed in rodent and human arteries (Knot and Nelson, 1998; Moosmang et al., 2003). Although a principal conductance, emerging animal studies indicate the presence of additional subtypes (Jensen et al., 2004; Kuo et al., 2010; Abd El-Rahman et al., 2013). Of particular note are T-type Ca2+ channels whose voltage profiles are leftward shifted, compared with L-type, positioning them to be more active at hyperpolarized Vm. Initial data argued that T-type channels were functionally similar to L-type in elevating cytosolic [Ca2+]i, albeit at
lower intraluminal pressures where arteries are hyperpolarized (Abd El-Rahman et al., 2013; Björling et al., 2013). Recent reports, however, indicate that T-type regulation of arterial tone is more sophisticated, with one subtype (CaV3.1) driving constrictor responses and the other (CaV3.2) mediating feedback dilation (Chen et al., 2003; Harraz et al., 2014; Thuesen et al., 2014). These animal observations are intriguing, but their translational significance has remained elusive as mechanistic animal observations are intriguing, but their translational significance has remained elusive as mechanistic studies have not extended to the human vasculature.

The goal of this study was to delineate Ca²⁺ channels in native human cerebral arterial smooth muscle and to determine how each conductance modulates arterial tone development. We hypothesized that both L- and T-type Ca²⁺ channels are present and that each conductance mediates distinctive vasomotor responses. Our investigation progressed from cells to tissues and involved the integrative use of pressurized myography, Western blot analysis, PCR, electrophysiology, and computational modeling. In human cerebral arterial smooth muscle cells, we documented for the first time the expression of L- (CaV1.2) and T-type (CaV3.2/CaV3.3) channels and characterized their unique physiological and pharmacological profiles. Subsequent observations showed that CaV1.2 and CaV3.3 augmented myogenic arterial tone in a manner consistent with their voltage dependence. In stark contrast, human CaV3.2 antagonized tone development and enabled indirect arterial vasodilation through its influence on large-conductance Ca²⁺-activated K⁺ (BKCa) channels. The divergent roles of Ca²⁺ channels are therapeutically important, as subtype-specific targeting could either suppress or enhance arterial tone. In summary, this human-based study is the first to illustrate the expression of three different CaV channels and to encode their distinctive influences on human cerebral arterial tone development.

**MATERIALS AND METHODS**

Human tissues and animal procedures

Excised human brain samples were obtained after institutional review board approval, written informed consent, and in accordance with the guidelines of the Declaration of Helsinki. Brain tissues were collected and placed in cold PBS, pH 7.4, containing (mM) 138 NaCl, 3 KCl, 10 Na₂HPO₄, 2 NaH₂PO₄, 5 glucose, 0.1 CaCl₂, and 0.1 MgSO₄, and transferred to the laboratory. Small superficial cerebral arteries (~150–250-µm diameter) were carefully dissected out of surrounding tissue and cut into segments. Animal procedures were approved by the Animal Care and Use Committee at the University of Calgary. In brief, female Sprague–Dawley rats (2–4-mo old) or C57BL/6 mice (2–4-mo old) were euthanized via CO₂ asphyxiation. The brain was carefully removed and placed in cold PBS. Middle and posterior cerebral arteries were carefully dissected out of surrounding tissue and cut into ~2-mm segments for enzymatic digestion.

Pressurized vessel myography

Human arterial segments were mounted in a customized arteriograph and superfused with warm (37°C) physiological saline solution containing: 119 mM NaCl, 4.7 mM KCl, 20 mM NaHCO₃, 1.1 mM KH₂PO₄, 1.2 mM MgSO₄, 1.6 mM CaCl₂, and 10 mM glucose, pH 7.4 (21% O₂, 5% CO₂, balance N₂). To limit the endothelial feedback influence on myogenic tone development, air bubbles were passed through the vessel lumen (~2–min). Arteries were equilibrated at 15 mmHg, and the contractile responsiveness was assessed by brief application of 60 mm KCl. After equilibration, intraluminal pressure was incrementally elevated from 20 to 100 mmHg and arterial external diameter was monitored under control conditions and in the presence of 200 nM nifedipine (CaV1.2 inhibitor), 1 µM NNC 55–0396 (T-type blocker), and/or 50 µM Ni²⁺ (CaV3.2 blocker). Maximal arterial diameter was subsequently assessed in Ca²⁺-free PSS (zero externally added Ca²⁺ plus 2 mM EGTA). Percentage of myogenic tone was used as a measure of myogenic responsiveness and was calculated at each intraluminal pressure as: % myogenic tone = 100* (D – D₀)/D₀, where D is external diameter under control conditions (Ca²⁺- PSS) or treated conditions (nifedipine, NNC 55–0396 or Ni²⁺), and D₀ is external diameter in Ca²⁺-free PSS. Percentage of maximal tone was used to ascertain the relative contribution of L- and T-type channels and was calculated at each intraluminal pressure as: % maximal tone = 100* (D₀ – D)/D₀, where D₀ is external diameter under control conditions (Ca²⁺- PSS) or treated conditions (nifedipine, NNC 55–0396 or Ni²⁺), and D₀ is external diameter in Ca²⁺-free PSS.

Isolation of cerebral arterial smooth muscle cells

Smooth muscle cells from human, rat, or mouse cerebral arteries were enzymatically isolated as described previously (Harraz and Welsh, 2013). In brief, arterial segments (1–2-mm long) were placed in an isolation medium (at 37°C for 10 min) containing (mM): 60 NaCl, 80 Na-glutamate, 5 KCl, 2 MgCl₂, 10 glucose, and 10 HEPES with 1 mg/ml BSA, pH 7.4. Vessels were then exposed to a two-step digestion process that involved: (1) 10–15-min incubation in isolation medium (37°C) containing 0.5 mg/ml papain and 1.5 mg/ml dithioerythritol; and (2) a 15-min incubation in isolation medium containing 100 µM Ca²⁺, 0.7 mg/ml of type F collagenase, and 0.3 mg/ml of type H collagenase. After incubation, tissues were washed repeatedly with ice-cold isolation medium and triturated with a fire-polished pipette. Isolated cells were kept in ice-cold isolation medium for use within ~6 h.

Quantitative PCR (qPCR)

Total RNA was isolated from human cerebral arteries or isolated smooth muscle cells using the RNeasy mini kit (QIAGEN) according to the manufacturer’s recommendations. Control total RNA preparations from human whole brain or skeletal muscle were obtained from Takara Bio Inc., whereas human retina total RNA was provided by P. Schnetkamp (University of Calgary, Calgary, Canada). Reverse transcription was performed using the QuantiTect reverse transcription kit (QIAGEN). For the negative control groups, all components except the reverse transcriptase were included in the reaction mixtures. Real-time PCR using intron-spanning primer sequences obtained from the qPrimer depot (Table S2; Cui et al., 2007) was performed using the Kapa SYBR Fast Universal qPCR kit (Kapa Biosystems). Human glyceraldehyde 3-phosphate dehydrogenase (GAPDH) gene was used as the reference gene. Control reactions and those containing cDNA from cerebral arteries were performed with 1 ng of template per reaction. Because of the limited quantities of RNA obtained from isolated smooth muscle cells (~200 cells), the entire cDNA yield from each preparation was used to assay the full set of test and housekeeping genes. The running protocol extended to 45 cycles consisting of 95°C for 5 s, 55°C for 10 s, and 72°C for 8 s using an Eppendorf Realplex 4 Mastercycler. PCR specificity was checked by dissociation curve analysis. Assay validation was confirmed by testing serial dilutions of pooled template cDNAs, suggesting a
linear dynamic range of 2.8–0.0028 ng of template and yielding percent efficiencies ranging from 80.4 to 108% (Table S3). No-template controls yielded no detectable fluorescence. Expression of the various Cav subtypes in human cerebral arteries or smooth muscle cells relative to whole brain (Cav1.2, Cav1.3, Cav2.1, Cav2.2, Cav2.3, Cav3.1, Cav3.2, and Cav3.3), skeletal muscle (Cav1.1), or retina (Cav1.4) was determined using the relative expression software tool (REST) version 2.0.13.

Western blot analysis

Whole human arteries were collected and frozen at −80°C in PBS, pH 7.4. The arteries were thawed on ice, centrifuged briefly to remove PBS, resuspended in low ionic strength buffer (10 mM Tris, pH 7.0, and 0.5 mM MgCl₂) containing Complete Ultra protease inhibitors (Roche), and incubated for 10 min on ice. The tissue was homogenized using a Polytron, followed by the addition of 1 volume solution A (0.5 M sucrose, 10 mM Tris, pH 7.0, 40 µM CaCl₂, 6 mM 2-mercaptoethanol, and 0.3 M KCl). After additional homogenization with the Polytron, the arteries were further subjected to 40 strokes in a Dounce homogenizer. The resulting crude lysate was centrifuged at 100,000 g for 1 h and resuspended in 50 µl of solution B (0.25 M sucrose, 10 mM Tris, pH 7.0, 20 µM CaCl₂, 3 mM 2-mercaptoethanol, and 0.15 M KCl). The protein concentration was determined with the Protein assay (Bio-Rad Laboratories) using BSA as the standard. The samples (4 µg per lane) were electrophoresed on a 5.6% polyacrylamide gel followed by transfer to PVDF membranes. The membranes were blocked for 1 h at room temperature using 0.2% Tropix I-Block (Applied Biosystems) in PBS followed by the addition of 1:200 diluted primary polyclonal antibodies raised against Cav1.2, Cav3.2, or Cav3.3 (Alomone Labs) overnight at 4°C. The membranes were washed three times for 10 min with PBS containing 0.1% Tween-20 followed by the addition of 1:10,000 goat anti-rabbit IRDye 800CW (Li-Cor Biosciences) for 1 h at room temperature, washed three times with PBS-Tween, and visualized using an infrared imaging system (Odyssey; Li-Cor Biosciences).

Electrophysiological recordings

Whole-cell currents were recorded using a patch-clamp amplifier (Axopatch 200B; Molecular Devices), filtered at 1 kHz, digitized at 5 kHz, and stored on a computer for offline analysis with Clampfit 10.3 software. Whole-cell capacitance was measured with the cancellation circuitry in the voltage-clamp amplifier. A 1-M NaCl–agar salt bridge between the reference electrode and the pipette puller (Narishige PP-830), and backfilled with pipette solution contained (mM): 110 K aspartate, 30 KCl, 10 NaCl, 2 MgCl₂, 2 CaCl₂, 10 glucose, and 10 HEPES, pH 7.4. The pipette solution contained (mM): 110 K aspartate, 30 KCl, 10 NaCl, 2 MgCl₂, 10 HEPES, and 0.05 EGTA, pH 7.2, with 200 µg/ml amphotericin B. Membrane currents were gap-free recorded while the cells were voltage clamped at −40 mV. As reported previously (Harraz et al., 2014) and to analyze the frequency and amplitude of spontaneous transient outward K⁺ currents (STOCs), the cutoff value was set to be three times the single-channel conductance.

BKV channels. Perforated patch-clamp electrophysiology was used to measure whole-cell K⁺ currents in isolated human smooth muscle cells. The bath solution contained (mM): 134 NaCl, 4 KCl, 2 MgCl₂, 2 CaCl₂, 10 glucose, and 10 HEPES, pH 7.4. The pipette solution contained (mM): 110 K aspartate, 30 KCl, 10 NaCl, 2 MgCl₂, 10 HEPES, and 0.05 EGTA, pH 7.2, with 200 µg/ml amphotericin B. Membrane currents were gap-free recorded while the cells were voltage clamped at −40 mV. As reported previously (Harraz et al., 2014) and to analyze the frequency and amplitude of spontaneous transient outward K⁺ currents (STOCs), the cutoff value was set to be three times the single-channel conductance.

Computational modeling

Network structure. A single complete arterial network of the cerebral vasculature was acquired using a micrograph of a resected brain biopsy. The micrograph was then manually segmented to obtain a network containing six orders (first–sixth) of branching arteries. This network was supplied by a single inflow vessel and contained 20 outflow vessels. To obtain a mathematical representation suitable for blood flow simulations, the network was discretized into 135 cylindrical vessel segments (213–1,700-mm long). Arterial diameter values were initially traced and set constant for each order according to average measured values (in μm): 600, 396, 261, 173, 114, and 75 for first, second, third, fourth, fifth, and sixth order vessels, respectively.

Blood flow. A previously developed theoretical model (Pries et al., 1990, 1994; Goldman and Popel, 2000) was used to calculate pressures and two-phase (erythrocyte and plasma) steady-state flows in the cerebral arterial network considered. This hemodynamic model implements conservation of blood volume flow (Q) and erythrocyte volume flow (QHb) at each node joining two or three vessel segments, where Hb is the discharge (flowaveraged) hematocrit. These conservation equations are given below, where the sums are over all segments i connected to node j:

$$\sum_i Q_{ij} = 0 \quad \text{and} \quad \sum_i H_i Q_{ij} = 0.$$  

These equations are solved using the Poiseuille pressure–flow relation $AP = \eta LQ/(R^4)$, where R and L are segment radius and length, respectively, and $\eta$ is the blood viscosity. The model includes the following empirical properties of blood rheology:

(a) the Fahraeus effect, which determines how tube (volume-averaged) hematocrit varies with vessel diameter; (b) the Fahraeus–Lindqvist effect, which determines how viscosity varies with diameter and hematocrit; and (c) the phase separation (or plasma skimming) effect, which determines how erythrocytes are distributed at diverging bifurcations. The model requires specification of hematocrit at the inlet segment and specification of pressures at the upstream ends of the inlet segments and downstream ends of the outlet segments. All flow simulations used an inlet hematocrit of 0.42, and inlet and outlet pressures of 75 and 40 mmHg, respectively, in accordance with earlier studies that monitored the arterial pressure drop in cerebral vascular networks (Faraci and Heistad, 1990).

Myogenic tone and diameters. The pressure–diameter relationship measured in isolated human vessels under control conditions (see Fig. 1) was fit with a fourth-order polynomial, and this was used to adjust diameters from the initially constant values within each order given above based on the calculated pressures in each segment. Solutions were obtained iteratively, by alternating updates of pressures and flows with updates of diameters, until pressures, diameters, and flows became constant (steady state) throughout the model network. The same method was used for simulations in the presence of nifedipine, NNC, or Ni²⁺ (see Figs. 6 and 7), by using fourth-order polynomial fits to the measured variations in the pressure–diameter relationship.

Statistical analysis
Data are expressed as means ± SEM, and n = a/b indicates the number of vessels or cells (a) per the number of human subjects, rats, or mice (b). Where appropriate, paired or unpaired t tests, or one-way ANOVA was performed using SigmaPlot 12.5 to compare the effects of a given condition/treatment on arterial diameter, or whole-cell current. P values ≤0.05 were considered statistically significant. Averaged I–V relationships were fit to the following Peak Gaussian third Parameter function:

\[ I(\text{V}) = \text{I}_{\text{max}}*\exp\left(-0.5\left(\text{V} - \text{V}_{\text{max}}\right)/\text{b}\right)^2 \],

where \( \text{I}_{\text{max}} \) is peak current (I), \( \text{V}_{\text{max}} \) is \( \text{V} \) at \( \text{I}_{\text{max}} \), and \( \text{b} \) is the slope of the distribution. The voltage dependence of activation and steady-state inactivation were described with single Boltzmann distributions of the following forms:

Activation: \( I(\text{V}) = \text{I}_{\text{max}}/(1 + \exp[(-\text{V} - \text{V}_{50})/\text{k}]) \);

Inactivation: \( I(\text{V}) = \text{I}_{\text{max}}/(1 + \exp[\text{V} - \text{V}_{50}/\text{k}]) \),

where \( \text{I}_{\text{max}} \) is the maximal activatable current, \( \text{V}_{50} \) is the voltage at which the current is 50% activated or inactivated, and \( \text{k} \) is the slope of the distribution.

Online supplemental material
Fig. S1 reports low and high voltage-activated currents in human cerebral arterial smooth muscle, and that high millimolar Ba²⁺ masks the separation of the two current components. Fig. S2 illustrates the effects of high concentrations of Ba²⁺ as the charge carrier on the I–V plots recorded in human smooth muscle. Fig. S3 depicts the biophysical properties of the nifedipine-sensitive and -insensitive currents in human cerebral arterial smooth muscle. Fig. S4 shows kinetic analysis of inward currents before and after the application of nifedipine. Fig. S5 compares representative traces and reversal potentials of Ba²⁺ currents recorded in human, mouse, and rat cerebral arterial smooth muscle cells. Table S1 reports the demographic characteristics of the origin of tissues and cells used. Tables S2 and S3 show the primer

Figure 1. Human cerebral arteries exhibited myogenic activity. (A) Photomicrographs and timeline highlighting the general methodological procedures. (B and C) Representative traces and summary data illustrate that pressurized human cerebral arteries displayed myogenic activity whereby they constricted as intraluminal pressure sequentially increased from 20 to 100 mmHg in PSS. In Ca²⁺-free PSS, cerebral arteries passively dilated (n = 7/6; three females; paired t test; *, P < 0.05; italic values denote probability). (D) Percentage of myogenic tone (see Materials and methods) was directly proportional to the applied intraluminal pressure (n = 6/6; three females; *, P < 0.05). Error bars denote SEM.
sequences and validation parameters for the qPCR analysis. The online supplemental material is available at http://www.jgp.org/cgi/content/full/jgp.201511361/DC1.

RESULTS

Human cerebral arteries exhibit myogenic activity
Pial cerebral arteries, isolated from patients undergoing brain resection surgery (Fig. 1 A and Table S1), were cannulated and mounted in a myograph and pressurized from 20 to 100 mmHg to assess myogenic reactivity. In physiological Ca^2+ (1.6 mM), human cerebral arteries gradually constricted to increases in intraluminal pressure (Fig. 1, B and C); this active response was abolished by removing extracellular Ca^2+ (0 mM Ca^2+ / 2 mM EGTA). Fig. 1 D illustrates that the percentage of myogenic tone increased progressively from 10% at 20 mmHg to 33% at 100 mmHg.

Human cerebral arterial smooth muscle expresses L- and T-type Ca^2+ channels
Given the preceding Ca^2+-dependent observations, we screened for the α1 pore-forming subunits of CaV channel subtypes (CaVx.x). Beginning with qPCR, analysis of human cerebral arteries and isolated smooth muscle cells revealed that all CaV subunits were expressed in the former, whereas only CaV1.1, CaV1.2, CaV1.4, CaV3.2, and CaV3.3 were expressed in the latter (Fig. 2 A). When standardized to reference tissue (whole brain: CaV1.2, CaV1.3, CaV2.1, CaV2.2, CaV2.3, CaV3.1, CaV3.2, and CaV3.3; skeletal muscle: CaV1.1; retina: CaV1.4), CaV1.2 and CaV3.2 were enriched in whole arteries (Fig. 2 B),

Figure 2. Human cerebral arteries express L- and T-type CaV channels. (A) Screening mRNA for CaV channel subtypes (CaVx.x) in reference tissues, cerebral arteries (n = 3 PCR reactions on three groups of arteries from three patients; two females), or isolated smooth muscle cells (n = 5 PCR reactions on five digests of smooth muscle cells obtained from arteries of three patients; two females). qPCR was used to calculate the expression levels relative to the housekeeping gene human glyceraldehyde 3-phosphate dehydrogenase (GAPDH). (B and C) Expression of CaVx.x mRNA in human cerebral arteries (B) or smooth muscle cells (C) was assessed relative to levels in reference tissues (whole brain: CaV1.1, CaV1.3, CaV2.1, CaV2.2, CaV2.3, CaV3.1, CaV3.2, and CaV3.3; skeletal muscle: CaV1.1; retina: CaV1.4). (D) Protein expression of the α1 subunits of CaV1.2, CaV3.2, and CaV3.3 in human cerebral arteries was determined by Western blotting. Blots are representative of three independent experiments. Error bars denote SEM.

Figure 3. CaV channel currents in human cerebral arterial smooth muscle cells. (A) Representative traces of inward currents from one isolated cell under conditions in which 1.8 mM Ca^2+ and 1.8 or 10 mM Ba^2+ were alternatively used as the extracellular charge carrier. (B) Averaged effect (bar graphs and I-V plots) of the extracellular charge carrier on the amplitude of the inward current (n = 5/2; one female; *, P < 0.05). (C) Normalized I-V plots displayed a depolarizing shift upon changing from 1.8 mM Ca^2+ to 1.8 mM Ba^2+ and then 10 mM Ba^2+. (D) Representative trace and I-V plot of 10 mM Ba^2+ currents (n = 40/11; five females) using the step protocol displayed. (E) Averaged I-V plots obtained using the ramp protocol (inset; n = 26/7; three females; gray denotes SEM). (F) Scatter plots of the correlation between current amplitude (pA/pF) and the age of the subjects from which cells were obtained. Left plot was obtained from 45 cells from 13 subjects (six females). Middle and right plots represent females (n = 19/6) and males (n = 26/7), respectively. Error bars denote SEM.
and CaV1.2, CaV3.2, and CaV3.3 displayed enriched expression in isolated smooth muscle cells (Fig. 2 C). Endothelial contamination of isolated smooth muscle samples was minimal as Von Willebrand factor mRNA levels decreased dramatically (~96%) compared with whole arteries. Focusing on the enriched targets, Western blot analysis confirmed the protein expression of CaV1.2, CaV3.2, and CaV3.3 in human cerebral arteries (Fig. 2 D).

Inward currents through human CaV channels
Using patch-clamp electrophysiology, we then monitored inward currents in cerebral arterial smooth muscle. Freshly isolated human cells were bathed in 1.8 mM Ca2+ and 1.8 or 10 mM Ba2+; the fist mimics physiological extracellular conditions, whereas the latter two accentuate charge flow through open Ca2+ channels. Stepping from −90 mV to voltage steps ranging from −50 to 40 mV, we observed a voltage-dependent inward current that was approximately fivefold less in 1.8 mM Ca2+ than in high [Ba2+] (Fig. 3 A and B). Fig. 3 C illustrates the voltage dependence of the whole-cell currents, with normalized I-V plots showing that Ba2+ evoked a depolarizing shift compared with Ca2+. Notably, normalized Ca2+ current (Fig. 3 C) exhibited a hyperpolarized threshold and an additional inward hump peaking at −30 mV, suggestive of a low voltage-activated current component (Fig. S1). The rightward shift associated with Ca2+ replacement could be further advanced by working with solutions containing 50 or 110 mM Ba2+ (Fig. S2). Given the small amplitude of native Ca2+ currents, subsequent recordings were performed in 10 mM Ba2+; step and ramp voltage protocols elicited robust inward Ba2+ currents (Fig. 3, D and E). Linear regression analysis (Fig. 3 F) indicated an inverse correlation between peak current amplitude and age of subjects (r2 = 0.531; P < 0.0001; age range 19–70 yr; n = 45/13; six females). This correlation was particularly evident in female (r2 = 0.6057; P < 0.0001; 19–58 yr; n = 19/6) rather than male (r2 = 0.0001; P = 0.9566; 33–70 yr; n = 26/7) subjects.

Delineating L- and T-type channels in human cerebral arterial smooth muscle
The whole-cell inward Ba2+ current depicted in Fig. 3 reflected simultaneous influx through L-type (CaV1.2) and T-type (CaV3.2/CaV3.3) Ca2+ channels. To effectively delineate between the two types, nifedipine was applied at a concentration (200 nM) that fully abolishes...
exhibited rapid activation (T-type: \( \tau_{\text{Act}} \), 2.1 ms; L-type: \( \tau_{\text{Act}} \), 7.5 ms), and deactivated slower (T-type: \( \tau_{\text{Deact}} \), 8.9 ms; L-type: \( \tau_{\text{Deact}} \), 3.2 ms) compared with subtracted nifedipine-sensitive current (Fig. S3) or total control current (Fig. S4). Linear regression analysis (Fig. 4 G) indicated a negative correlation between age and the amplitude of the L-type (\( r^2 = 0.7396; P < 0.0001; 36–70 \) yr; \( n = 19/6 \)) but not for T-type current (\( r^2 = 0.0753; P = 0.2164; 33–70 \) yr; \( n = 22/7 \)).

Cross-species comparison of CaV channel activity
Past studies have shown that rat and mouse arterial smooth muscle express CaV1.2 (L), CaV3.1 (T), and CaV3.2 (T) subtypes (Kuo et al., 2010; Abd El-Rahman et al., 2013; Björling et al., 2013; Thuesen et al., 2014). Human cells displayed a similar expression profile, with the exception being that CaV3.1 was substituted by CaV3.3 (Fig. 2). Using patch-clamp electrophysiology, we compared the electrical properties of cerebral arterial \( \text{Ca}^{2+} \) channels between species using 10 mM Ba\(^{2+} \) as the charge carrier. Values of cell capacitance (pF), indicative of cell size, varied across species (Fig. 5 A). Inward current densities (pA/pF) were lower in human smooth muscle cells than in mouse or rat; however, L- but not T-type currents (Liao et al., 2007; Harraz and Welsh, 2013); \( \sim 75\% \) block was observed (Fig. 4 A). Note that the use of different prepulse voltages was not successful in the separation of low and high voltage-activated Ca\(^{2+} \) channel currents in rodent cerebral arterial smooth muscle (Kuo et al., 2010; Abd El-Rahman et al., 2013). To test whether the residual nifedipine-insensitive current was dominated by a T-type conductance, we applied 1 \( \mu M \) NNC 55–0396. This T-type blocker abolished the remaining current, and I-V plots showed that nifedipine- and NNC-sensitive current components peaked (\( V_{\text{max}} \)) at 19.7 ± 0.4 mV and 2.3 ± 0.9 mV, respectively (Fig. 4, B and C).

Caution is warranted when using pharmacological T-type blockers given their reported off-target effects on L-type channels (Kuo et al., 2010; Abd El-Rahman et al., 2013; Harraz and Welsh, 2013). Experiments in Fig. 4 (D–F) confirmed the successful delineation as nifedipine evoked hyperpolarized shifts in the voltage dependence of: (a) steady-state inactivation (\( V_{50\text{Inact}} \)) at \(-35.4 \pm 2.3 \) mV vs. control, \(-12.7 \pm 0.9 \) mV; (b) activation (\( V_{50\text{Act}} \)) at \(-20.3 \pm 1.3 \) mV vs. control, \(-5.2 \pm 0.8 \) mV; and (c) peak window current (\( -35.4 \) mV vs. control \(-14.1 \) mV). Kinetic analysis additionally noted that the nifedipine-insensitive T-type component inactivated faster (Fig. 4, E and F).

Figure 5. Electrophysiological comparison of Ca\(_V\) channels in cerebral arterial smooth muscle cells of humans, mice, and rats. (A) Scatter plot of cell capacitance (pF) obtained from human (H, black), mouse (M, red), or rat (R, green) cerebral arterial smooth muscle cells. (B) Absolute (pA/pF) and normalized I-V plots of inward currents (\( n = 40/12 \) human; six females; 25/15 mouse; 37/20 rat). (D) Bar graphs of current amplitude before (control) and after nifedipine (200 nM) treatment. Normalized plot fits (bottom) display the hyperpolarized shifts of T-type nifedipine-insensitive currents. *, \( P < 0.05 \), significant from respective control. (E) I-V plot fits of residual T-type currents. Bottom inset displays \( V_{\text{max}} \) values (\( n = 21/10 \) human; six females; 25/15 mouse; 25/14 rat). (F) Percentage of inactivation at different time points after a voltage step from \(-90 \) to \(10 \) mV in the absence and presence of nifedipine. (G) Averaged effect of 200 nM nifedipine on voltage dependence of steady-state inactivation (left) and activation (right); average normalized data (% \( I/I_{\text{max}} \)) were fit with single Boltzmann distributions. Error bars denote SEM.
normalized currents displayed comparable dependence on voltage (Fig. 5, B and C). Whole-cell currents inactivated faster in mouse smooth muscle cells, whereas the reversal potential of rat currents was depolarized relative to human or mouse (Fig. S5). 200 nM nifedipine elicited comparable blockade across species, and the residual insensitive currents displayed 10–20-mV hyperpolarized shifts consistent with T-type activity (Fig. 5, D and E). In all three species, L-type channel inhibition quickened inactivation and leftward shifted the activation/inactivation profiles (Fig. 5, F and G).

L- and T-type Ca$^{2+}$ channels contribute to human myogenic tone development

In rodent cerebral arteries, studies have shown that L- and T-type channels (CaV1.2 and CaV3.1) facilitate myogenic tone development at high and low intraluminal pressures, respectively (Abd El-Rahman et al., 2013; Björling et al., 2013). Both conductance types (L-type: CaV1.2; T-type: CaV3.3 substituting for CaV3.1) are present in human cerebral arteries and, as such, comparable experiments were conducted on pressurized human vessels to elucidate potential roles of Ca$^{2+}$ channels. Similar to electrophysiology, L-type Ca$^{2+}$ channel blockade preceded T-type suppression to minimize off-target effects. 200 nM nifedipine induced a vasodilator response in human vessels that was limited at low pressure and maximal at 80–100 mmHg. The additional application of a T-type blocker (1 µM NNC 55–0396) modestly dilated arteries in a pressure-dependent manner, with peak responses at 40–60 mmHg (Fig. 6, A–C). Data in Fig. 6 (B and C) displayed distinctive L- and T-type vasomotor trends, although statistical significance was not achieved because of limited observations ($n = 3/3; P = 0.087–0.157$).

As such, the relative contribution of the two conductances to developed myogenic tone was dynamic, with T-type channels (CaV3.3) prevailing at lower pressures when the $V_m$ was relatively hyperpolarized and L-type channels (CaV1.2) dominating at higher pressures when smooth muscle is strongly depolarized (Fig. 6 D).

**CaV3.2 and BK<sub>Ca</sub>-mediated STOC activity**

To discriminate between the contributions of CaV3.2 and CaV3.3 to myogenic constriction, we examined the effects of a specific CaV3.2 blocker (50 µM Ni$^{2+}$) on isolated human smooth muscle cells and arteries. At this concentration, Ni$^{2+}$ was verified to selectively suppress CaV3.2 currents, but not other L- or T-type channels expressed in arterial smooth muscle or expression systems (Lee et al., 1999; Harraz and Welsh, 2013; Harraz et al., 2014). Starting with electrophysiology, we confirmed that 50 µM Ni$^{2+}$ partially suppressed the nifedipine-insensitive CaV3.x current (Fig. 7, A and B). A kinetic analysis followed, and time constant measurements revealed that the Ni$^{2+}$-sensitive current activated and inactivated faster (CaV3.2: $\tau_{\text{act}}$, 0.9 ± 0.1 ms; $\tau_{\text{inact}}$, 17.1 ± 3.3 ms) than the insensitive current (CaV3.3: $\tau_{\text{act}}$, 4.3 ± 0.3 ms; $\tau_{\text{inact}}$, 48.7 ± 9.5 ms), consistent with Ni$^{2+}$ selectively blocking CaV3.2 (Fig. 7, C and D) (Lee et al., 1999; Perez-Reyes, 2003). We next applied 50 µM Ni$^{2+}$ onto pressurized human arteries, and contrary to the vasodilation observed with a broad spectrum T-type blocker (NNC 55–0396; Fig. 6), Ni$^{2+}$ enhanced myogenic constriction at pressures ≤60 mmHg (Fig. 7, E and F). Interestingly, Ni$^{2+}$- and NNC 55–0396-induced vasomotor changes were pressure dependent but modestly different from one another, a finding that aligns with the voltage profiles of CaV3.2 and CaV3.3 (Fig. 7 G). The maximal arterial tone was directly proportional to intraluminal pressure. In contrast, the T-type contribution (red) was inversely proportional and significantly different from that of L-type at 80 and 100 mmHg. Percentage of the maximal tone was mathematically calculated from three independent experiments ($n = 3/3$; one female; $\ast$, $P < 0.05$). Right inset represents fits of L- versus T-type components of maximal tone within a range of pressure steps (mmHg). Error bars denote SEM.
preceding Ni²⁺ observations suggest, in accordance with rat studies (Harraz et al., 2014), that Ca₉₃.₂ mediates a negative feedback response by downstream triggering of BKCa channels to initiate STOCs. Perforated patch-clamp electrophysiology indicated the presence of STOCs in human smooth muscle cells and showed that 50 µM Ni²⁺ significantly reduced the frequency (control, 1.30 ± 0.35 Hz; Ni²⁺, 0.69 ± 0.17 Hz) but not the amplitude (control, 6.87 ± 0.09 pA; Ni²⁺, 6.56 ± 0.25 pA) of these events (Fig. 7 H).

**Predicted changes in cerebral blood flow by Ca²⁺ channel modulators**

Myography experiments (Figs. 6 and 7) showed that human cerebral arterial Ca²⁺ channel subtypes elicited divergent vasomotor effects whereby Ca₉₁.₂/CaV₃.₃ augmented but CaV₃.₂ counterbalanced myogenic constriction. To conceptually ascertain how each channel subtype could influence cerebral blood flow, a computational model was developed using an arterial network map (lengths and basal diameters) generated from a photomicrograph of excised human brain tissue (see Materials and methods; Fig. 8, A–C). Intraluminal pressure within the virtual model varied, with inlet and outlet pressures set to 75 and 40 mmHg, respectively (Faraci and Heistad, 1990). The pressure–diameter relationship measured in isolated human vessels in the absence and presence of nifedipine, NNC 55–0936, or Ni²⁺ was fit with a fourth-order polynomial, and these percentage values were used to adjust the initially constant diameters within each segment (Fig. 8 D). This integration enabled the calculation of steady-state distributions of pressure, diameter, and blood flow for each arterial segment (135 in total; Fig. 8 E). Two-dimensional plots and color-coded virtual networks indicated that the suppression of Ca₉₃.₂ and to a lesser extent CaV₃.₃ increased network blood flow in a pressure/diameter-dependent manner. In contrast, Ca₉₃.₂ inhibition constricted arterial segments and decreased cerebral blood flow (Fig. 8 F).

**DISCUSSION**

This study delineated CaV channels in human cerebral arteries and determined subtype-specific contributions to arterial tone development. Experiments progressed from cells to tissues and used pressurized myography,
PCR, Western blotting, electrophysiology, and computational modeling. Messenger RNA and protein analyses revealed the expression of L- and T-type CaV pore–forming subunits in cerebral arterial smooth muscle. Patch-clamp recordings subsequently confirmed the presence of inward current divisible into L- and T-type components in isolated human smooth muscle cells. In pressurized human arteries, L- (CaV1.2) and T-type (CaV3.3) channel suppression dilated arteries in a pressure-dependent manner at higher and lower intraluminal pressures, respectively. Subtype-specific targeting of CaV3.2 paradoxically implicated this conductance in restraining arterial constriction via a mechanism involving BKCa channels. In a final set of experiments, computational modeling conceptually illustrated how each CaV subtype plays a distinctive role in setting cerebral blood flow. To summarize, this study reports for the first time the expression and divergent functions of three CaV channel subtypes in human cerebral arteries.

A century ago, Bayliss (1902) first described the inherent ability of resistance arteries to constrict to elevated arterial pressure, a phenomenon known as the myogenic response. In the cerebral circulation, this myogenic response is essential to maintain brain perfusion, and a global rise in smooth muscle \([\text{Ca}^{2+}]\) is critical to the transduction process. Like most stimuli, elevated intraluminal pressure drives a rise in cytosolic \([\text{Ca}^{2+}]\), through the induction of arterial depolarization and subsequent activation of voltage-gated L-type CaV1.2 channels (Knot and Nelson, 1998). Although a dominant player, animal studies have noted other Ca\(^{2+}\) channels (T-type, CaV3.1/CaV3.2) in cerebral arteries (Kuo et al., 2010; Abd El-Rahman et al., 2013). Distinct from the “long-lasting” kinetics and high voltage activation profile of the “L”-type conductance, “T”-type channels are characteristically “transient” and activate at more hyperpolarized voltages (Perez-Reyes, 2003; Catterall, 2011). Although Ca\(^{2+}\) entry events through L-type channels (CaV1.2 sparklets) have been characterized in vascular smooth muscle (Navedo et al., 2005), CaV3.x sparklets remain to be investigated. They would likely display spatiotemporal characteristics distinct from CaV1.2 and which according to their single channel properties would be more transient in nature and smaller in magnitude (Perez-Reyes, 2003). Recent studies in rodents have implied that T-type channels modulate myogenic tone, albeit in a manner unique and sometimes paradoxical to L-type channels (Chen et al., 2003; Kuo et al., 2010; Abd El-Rahman et al., 2013; Björling et al., 2013; Harraz et al., 2014). Despite the growing interest and therapeutic potential, human studies to date have failed to establish whether T-type channels are present in key vascular beds and if so, how they influence arterial tone.

Figure 8. Predicted influence of L- and T-type Ca\(^{2+}\) channels on blood flow in human cerebral arteries. (A and B) A photomicrograph of excised human brain tissue was used to generate an arterial network of defined dimensions. (C) The segmented network was skeletonized into vessel segments producing a mathematical representation useful for blood flow simulations; 1 marks the first inflow segment, and the other numbered boundary segments are outflows. Inlet and outlet pressures were set to 75 and 40 mmHg, respectively. (D and E) Using myography data, we modeled the vasomotor behavior of virtual arterial segments across a range of pressures (40–75 mmHg), in the absence and presence of nifedipine, NNC 55-0936, or Ni\(^{2+}\). Responses were incorporated into the model and steady-state distributions of diameter; pressure and blood flow were calculated for each arterial segment. (F) Color-coded maps of changes in blood flow illustrate the effects of CaV1.2, CaV3.3, or CaV3.2 channels on network perfusion.
Within this context, we began to isolate human cerebral arteries from patients undergoing resection surgery. Using vessel myography, we observed for the first time that human cerebral arteries constrict robustly to an increase in intraluminal pressure, with percentages of myogenic responses aligning with previous animal studies (Knot and Nelson, 1998; Cole and Welsh, 2011). The removal of extracellular Ca\textsuperscript{2+} abolished myogenic constriction, an observation consistent with elevated pressure evoking Ca\textsuperscript{2+} flux through Ca\textsubscript{V} channels (Knot and Nelson, 1998). 10 genes encode for the \( \alpha \) pore-forming subunits of Ca\textsuperscript{2+} channels (Catterall, 2011), and molecular analyses showed robust mRNA expression of three subtypes (Fig. 2). Ca\textsubscript{V}1.2 (L-type) and Ca\textsubscript{V}3.2 (T-type) were present in isolated human smooth muscle cells prescreened for endothelial contamination. Messenger RNA of Ca\textsubscript{V}3.3 was also observed while Ca\textsubscript{V}3.1 was minimally expressed, a finding distinct from rodents (Kuo et al., 2010; Abd El-Rahman et al., 2013; Björling et al., 2013). Compared with Ca\textsubscript{V}3.1, Ca\textsubscript{V}3.3 channel displays slower activation/inactivation kinetics and activates at relatively depolarized potentials (Monteil et al., 2000; Frazier et al., 2001; Chemin et al., 2002). One could rationalize, in this regard, that Ca\textsubscript{V}3.3 may be better suited than Ca\textsubscript{V}3.1 to drive cellular functions requiring sustained Ca\textsuperscript{2+} flux (Chemin et al., 2002), such as smooth muscle contraction and arterial tone development. Western blot analysis confirmed protein expression of all three subtypes in whole arteries.

Having detected their message and protein expression, electrophysiological recordings followed to separate L- and T-type conductances in human cerebral arterial smooth muscle. Human cells demonstrated a small inward current in physiological Ca\textsuperscript{2+} solutions that displayed a voltage-dependent IV profile consistent with Ca\textsuperscript{2+} channel activation (Fig. 3). Exchange of physiological Ca\textsuperscript{2+} for higher millimolar Ba\textsuperscript{2+} dramatically enhanced inward current, masked the low voltage–activated Ca\textsuperscript{2+} current component, and induced a depolarized shift in the voltage profile without destabilizing the cell. The latter shift is presumably responsible for masking the hyperpolarized current component and is reflective of a high divalent cation screening effect and barium’s preferential permeability through L-type channels (Perez-Reyes, 2003; Li et al., 2010). Using nifedipine at concentrations that fully block the smooth muscle splice variant of L-type Ca\textsubscript{V}1.2 channels (Liao et al., 2007; Harraz and Welsh, 2013), a partial suppression of Ba\textsuperscript{2+} currents occurred. Note that nifedipine has been shown to lack an effect on Ca\textsubscript{V}3.x in vascular smooth muscle or expression systems when applied in the submicromolar ranges (Harraz and Welsh, 2013; Harraz et al., 2014). Three lines of evidence subsequently confirmed that L- and T-type channels, respectively, carried the nifedipine-sensitive and -insensitive components. First, the nifedipine-insensitive current was actively blocked by T-type inhibitors. Second, the presumed L- and T-type currents displayed depolarized and hyperpolarized profiles, respectively, in accordance with conductances being high and low voltage activated. Third, the residual insensitive conductance activated and inactivated faster, in keeping with the transient nature of T-type channels (Catterall, 2011). Ca\textsubscript{V}3.x currents recorded in human arterial smooth muscle display kinetics similar to those in heterologous expression systems. Note, however, they do display a depolarized voltage-dependent profile because of the use of high millimolar Ba\textsuperscript{2+} as the permeant recording ion (Perez-Reyes, 2003; Harraz and Welsh, 2013). Overall, these findings are unique in the vascular literature and are the first to show that multiple Ca\textsubscript{V} channels are not only present but also electrically active in native human arterial smooth muscle.

Despite their diverse demographic origins, Ca\textsuperscript{2+} channel currents were resolved in all tested cells. With that said, an intriguing inverse relationship between current amplitude and donor age existed (Fig. 3), similar to reports in animal models (Georgeon-Chartier et al., 2013; Fukuda et al., 2014). This relationship was sexually dimorphic and manifested exclusively in females. These observations suggest that Ca\textsuperscript{2+} channel expression and/or activity may be under the regulation of sex hormones (Bowles et al., 2004; Ulrich et al., 2007); however, caution is warranted given the limited sample size and the pathological state of the human subjects. Further analysis showed that the preceding age–current correlation stemmed primarily from variations in L-type activity (Fig. 4), a finding that indicates that the efficacy of calcium channel blockers in the treatment of constrictive disorders may be altered or compromised in the elderly (Kumar and Hall, 2003; Fukuda et al., 2014). Recent rodent studies have suggested that both L- and T-type channels play a direct role in elevating cytosolic [Ca\textsuperscript{2+}], and myogenic tone, albeit at specific \( V_{\text{Ms}} \) (Abd El-Rahman et al., 2013; Björling et al., 2013). In particular, the L-type Ca\textsubscript{V}1.2 channels appear to dominate at elevated pressures when vessels are depolarized. In comparison, the low voltage–activated T-type Ca\textsubscript{V}3.1 channels are thought to facilitate tone development at lower pressures when smooth muscle resides in a hyperpolarized state (Abd El-Rahman et al., 2013; Björling et al., 2013). Human L- and T-type channels could operate similarly, and to pursue this hypothesis, a two-step approach was instituted. Starting with a comparative electrical analysis, findings in Fig. 5 illustrate that human, rat, and mouse smooth muscle cells all exhibit L- and T-type current components, with human cells displaying smaller amplitudes. No substantive difference was observed in activation/steady-state inactivation voltage profiles, although inactivation kinetics were faster in mouse smooth muscle. Given these similarities, we next pressurized human cerebral arteries from 20 to...
100 mmHg and performed vessel myography under control conditions and in the presence of L-type (nifedipine) and T-type (NNC 55–0396) channel blockers. In a pressure-dependent manner, L- and T-type blockers evoked vasodilation in myogenically responsive arteries (Fig. 6). L- and T-type channels dominated myogenic tone at higher and lower pressures, respectively, in accordance with the physiological window currents of CaV1.2 and CaV3.x channels. Reduced Ca\textsuperscript{2+} flux through CaV3.3 channels is most likely responsible for the vasodilator response to NNC 55–0396, as findings below note that selective CaV3.2 blockade induces arterial constriction. Our observations are the first to invoke a role for T-type channels in human cerebral arteries, and arguably this conductance may dominate Ca\textsuperscript{2+} entry in distal arteries where intraluminal pressures are \(\sim30–45\%\) less than systemic (Faraci and Heistad, 1990). If true, it follows that constrictive disorders in smaller arteries may be more effectively treated by simultaneous L- and T-type channel suppression, a view consistent with clinical findings (Beltrame et al., 2004; Ogura et al., 2012; Sugano et al., 2013).

Studying T-type channels in native cells has been consistently limited by the lack of subtype-specific modulators. One exception is Ni\textsuperscript{2+}, a divalent cation that preferentially blocks CaV3.2 channels when applied in the low micromolar range (Lee et al., 1999; Harraz et al., 2014). In rat cerebral arterial smooth muscle and HEK cells, Ni\textsuperscript{2+} abolished CaV3.2, but not other high or low voltage-activated Ca\textsuperscript{2+} currents. Previous experiments have additionally shown that Ni\textsuperscript{2+} lacked any effects on cells and arteries where CaV3.2 channels had been genetically ablated (Harraz and Welsh, 2013; Harraz et al., 2014). Exploiting this tool, electrophysiological experiments followed by kinetic analysis confirmed that 50 \(\mu\text{M}\) Ni\textsuperscript{2+} inhibited CaV3.2 but not CaV3.3 channels in human cerebral arterial smooth muscle cells. In paradoxical contrast to the vasodilation induced by a broad-spectrum T-type blocker, Ni\textsuperscript{2+} enhanced myogenic constriction at pressures \(\leq60\) mmHg (Fig. 7). Such results suggest a distinctive role of CaV3.2 channels, perhaps involving a vasodilator conductance sensitive to Ca\textsuperscript{2+} and/or voltage. In this regard, recent rodent literature has argued that CaV3.2 can trigger Ca\textsuperscript{2+} release from the sarcoplasmic reticulum in the form of Ca\textsuperscript{2+} sparks, and that the subsequent generation of Ca\textsuperscript{2+} sparks elicits hyperpolarization via BK\textsubscript{Ca} channels (Chen et al., 2003; Harraz et al., 2014). To address this possibility, we monitored BK\textsubscript{Ca}-mediated STOC activity in human cells, and in support of our hypothesis, Ni\textsuperscript{2+} inhibited these transient hyperpolarizing currents. It is worth noting that previous whole-cell and whole arterial recordings argued against a direct modulatory role of Ni\textsuperscript{2+} on K\textsuperscript{+} channels when applied at low micromolar concentrations (Harraz et al., 2014). Our findings challenge the stereotypic view that Ca\textsuperscript{2+} channels only facilitate myogenic constriction. They further highlight the view that if T-type modulators are to be used therapeutically to treat cardiovascular disorders (Giordanetto et al., 2011), subtype specificity should be thoughtfully considered.

With CaV1.2, CaV3.2, and CaV3.3 channels distinctively influencing arterial diameter, logic dictates that unique blood flow responses should ensue. Although it is impractical to monitor cerebral blood flow in small human arteries, such responses can be predicted in a virtual arterial network (Pries et al., 1990, 1994; Goldman and Popel, 2000). Using architectural and vasomotor data from human arteries, we consequently designed a computational model that mimics a native arterial network. Findings in Fig. 8 highlight that CaV1.2 or CaV3.3 inhibition dilated virtual cerebral arteries and increased network blood flow. CaV1.2 blockade elicited a particularly profound response in larger arteries, vessels that harbor higher intraluminal pressures. Subtype-specific suppression of CaV3.2 alternatively constricted arterial segments, diminishing network cerebral blood flow. Such predictions conceptually emphasize the differential significance of Ca\textsuperscript{2+} channels in matching blood flow with brain metabolic activity.

This study identified Ca\textsubscript{V} channels in human cerebral arteries and ascertained how each conductance influences myogenic tone. Molecular and electrophysiological analyses demonstrated that \(\alpha_1\) subunits of L- (CaV1.2) and T-type (CaV3.2 and CaV3.3) channels are robustly expressed in human cerebral arterial smooth muscle. In myogenically active human arteries, CaV1.2 inhibition induced vasodilation at high intraluminal pressures, whereas CaV3.3 blockade induced modest dilation at lower pressures when vessels resided in a hyperpolarized state. Paradoxically, cerebral arterial CaV3.2 channels appear to attenuate myogenic tone through the downstream activation of BK\textsubscript{Ca} channels.

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