

Evidence both L-type and non-L-type voltage-dependent calcium channels contribute to cerebral artery vasospasm following loss of NO in the rat

Article

Accepted Version

McNeish, A., Altayo-Jimenez, F. and Garland, C. J. (2010) Evidence both L-type and non-L-type voltage-dependent calcium channels contribute to cerebral artery vasospasm following loss of NO in the rat. *Vascular Pharmacology*, 53 (3-4). pp. 151-159. ISSN 1537-1891 doi: <https://doi.org/10.1016/j.vph.2010.06.002> Available at <https://centaur.reading.ac.uk/25803/>

It is advisable to refer to the publisher's version if you intend to cite from the work. See [Guidance on citing](#).

To link to this article DOI: <http://dx.doi.org/10.1016/j.vph.2010.06.002>

Publisher: Elsevier

All outputs in CentAUR are protected by Intellectual Property Rights law, including copyright law. Copyright and IPR is retained by the creators or other copyright holders. Terms and conditions for use of this material are defined in the [End User Agreement](#).

www.reading.ac.uk/centaur

CentAUR

Central Archive at the University of Reading

Reading's research outputs online

Evidence both L-type and non-L-type voltage-dependent calcium channels contribute to cerebral artery vasospasm following loss of NO in the rat

A. J. McNeish,² Francesc Jimenez Altayo¹ and C. J. Garland^{1*}

¹Department of Pharmacology, University of Oxford, Mansfield Road Oxford OX1 3QT &

²Department of Pharmacy & Pharmacology, University of Bath, BA2 7AY

Running title: Calcium channels and cerebral vasospasm

* Author for correspondence:

Professor C J Garland,

Department of Pharmacology,

University of Oxford,

Mansfield Road,

Oxford OX1 3QT. UK

Phone: +44-1865-281119

Fax: +44-1865-271853

e-mail: christopher.garland@pharm.ox.ac.uk

Abstract

We recently found block of NO synthase in rat middle cerebral artery caused spasm, associated with depolarizing oscillations in membrane potential (E_m) similar in form but faster in frequency (*circa* 1Hz) to vasomotion. T-type voltage-gated Ca^{2+} channels contribute to cerebral myogenic tone and vasomotion, so we investigated the significance of T-type and other ion channels for membrane potential oscillations underlying arterial spasm. Smooth muscle cell membrane potential (E_m) and tension were measured simultaneously in rat middle cerebral artery. NO synthase blockade caused temporally coupled depolarizing oscillations in cerebrovascular E_m with associated vasoconstriction. Both events were accentuated by block of smooth muscle BK_{Ca} . Block of T-type channels or inhibition of Na^+/K^+ -ATPase abolished the oscillations in E_m and reduced vasoconstriction. Oscillations in E_m were either attenuated or accentuated by reducing $[Ca^{2+}]_o$ or block of K_V , respectively. TRAM-34 attenuated oscillations in both E_m and tone, apparently independent of effects against $K_{Ca3.1}$. Thus rapid depolarizing oscillations E_m and tone observed after endothelial function has been disrupted reflect input from T-type calcium channels in addition to L-type channels, other depolarizing currents appear to be unimportant. These data suggest that combined block of T and L-type channels may represent an effective approach to reverse cerebral vasospasm.

Key words: cerebral vasospasm; K-channels; T-type calcium channels; nitric oxide; endothelial cell dysfunction

1. Introduction

Cerebral arteries operate in a dynamic state of partial constriction (myogenic tone), providing the capacity to constrict or relax in response to changing levels of intraluminal pressure, shear stress and nerve activity. Myogenic tone is an intrinsic property of the smooth muscle, helping maintain constant total cerebral blood flow and adapting blood flow locally to meet metabolic demand.. Myogenic constriction is driven primarily by membrane depolarization leading to Ca^{2+} influx (Davis *et al.*, 1999; Hill *et al.*, 2001), possibly with a contribution via stretch-activated calcium sensitization (Schubert *et al.*, 2008). Myogenic tone is often superimposed by vasomotion in the form of synchronised oscillations in smooth muscle cell membrane potential (E_m), Ca^{2+} and tension. Although the physiological function of vasomotion in general is unclear, it may help to maintain a constant blood supply in many tissues, including the brain (Haddock *et al.*, 2005).

One key influence of basal NO release in the middle cerebral artery appears to be suppression of both myogenic tone (Golding *et al.*, 2001; Peng *et al.*, 1998; Zimmermann *et al.*, 1997) and vasomotion (Dirnagl *et al.*, 1993; Haddock *et al.*, 2002). This influence appears largely due to activation of BK_{Ca} (Mandala *et al.*, 2007; Yuill *et al.*, 2010). So block of NO generation and/or BK_{Ca} provides a means to mimic an aspect of endothelial dysfunction that is an early feature of cardiovascular disease, including disease conditions that predispose to vasospasm (Jewell *et al.*, 2004; Vanhoutte *et al.*, 2009). These conditions will also mimic the loss of NO observed after subarachnoid haemorrhage, where scavenging of NO by haemoglobin (Martin *et al.*, 1985) causes profound vasospasm (Toda *et al.*, 1991). Significantly, enhanced vasomotion (or vasospasm) can lead to a reduction of cerebral capillary blood flow and thus compromise of neuronal function (Biswal *et al.*, 1996; Pluta, 2005).

Several mechanisms, including the rho-kinase pathway, can contribute to the development and maintenance of constriction in smooth muscle, alongside calcium entry. However, in many vascular beds, including the cerebral vasculature changes in smooth muscle intracellular Ca^{2+} ($[\text{Ca}^{2+}]_i$) concentration are critical for myogenic tone and vasomotion (Haddock *et al.*, 2002; Haddock *et al.*, 2005; Yuill *et al.*, 2010). $[\text{Ca}^{2+}]_i$ increase involves release from intracellular stores and entry from the extracellular space via voltage-gated Ca^{2+} channels and non-selective cation channels, such as transient receptor potential channels (TRPC). Ca^{2+} influx through voltage-gated Ca^{2+} channels (VGCC) leads to global increases in smooth muscle cell $[\text{Ca}^{2+}]_i$ and constriction, and high voltage activated (L-type) Ca^{2+} channels appear central in this sequence (McCarron *et al.*, 1997; Moosmang *et al.*, 2003; Nelson *et al.*, 1990). These channels are expressed widely in vascular smooth muscle and their open probability increases over a physiologically relevant range (circa -50 to -30 mV) (Lacinova, 2005; Smirnov *et al.*, 1992). Low voltage activated or T-type Ca^{2+} channels are also expressed in vascular smooth muscle of resistance arteries (Braunstein *et al.*, 2008; Clozel *et al.*, 1997; Kuo *et al.*, 2008; Navarro-Gonzalez *et al.*, 2009; Perez-Reyes, 2003). But although they are normally active in the range *circa* -60 to -40 mV, the characteristic rapid inactivation of these channels argues against a significant role at physiologically relevant membrane potentials in the vasculature. Despite this, they have been implicated in the maintenance of vascular tone in a variety of arteries, including rat cremaster (VanBavel *et al.*, 2002), rat basilar (Navarro-Gonzalez *et al.*, 2009) and middle cerebral (Lam *et al.*, 1998) arteries, and direct measurements have shown high voltage-activated but nifedipine-insensitive Ca^{2+} currents, pharmacologically indistinguishable from T-type currents, in both guinea-pig and rat terminal mesenteric arteries (Morita *et al.*, 1999; Morita *et al.*, 2002).

We recently reported that middle cerebral arteries develop intense and sustained constriction, associated with a very rapid form of vasomotion, when NOS and/or BK_{Ca} channels were blocked, to mimic endothelial dysfunction. Furthermore, both constriction and vasomotion depended on

calcium entry via VGCCs and the oscillations in E_m were temporally linked to changes in smooth muscle $[Ca^{2+}]_i$ (Yuill *et al.*, 2010). The temporally linked oscillations in E_m , $[Ca^{2+}]_i$ and tension were similar to the widely described phenomenon of vasomotion, but displayed a much higher frequency (~ 1 Hz as opposed to $\sim 0.1-0.2$ Hz, Yuill *et al.*, 2010). The intense “vasospastic vasomotion” was reversed by inhibition L-type Ca^{2+} channels and clearly involved a complex action of NO that appeared to include stimulation of BK_{Ca} channels and a cGMP-independent closure of VGCCs (Yuill *et al.*, 2010). However, although a central role for VGCC, NO and BK_{Ca} was apparent, the importance of other ionic currents that might contribute to the rapid depolarizing oscillations was unclear.

Thus, the aim of the present study was to characterize the ionic mechanisms responsible for rhythmic oscillations in E_m and tension in rat isolated middle cerebral arteries following inhibition of BK_{Ca} channels and NOS. We probed channels that may lead to both depolarization (calcium, sodium and chloride channels) and repolarization (potassium channels). Our data suggest a novel role for both smooth muscle T-type Ca^{2+} channels and several potassium conductances in the both “vasospastic vasomotion” and the underlying maintenance of vasoconstriction.

2. Materials and Methods

2.1 Animals and tissue isolation

Male Wistar rats (200-250 g) were killed by cervical dislocation followed by decapitation, following institutional guidelines for animal welfare and schedule 1 of the Animals (scientific procedures) Act 1986. The brain was removed and immediately placed in ice-cold Krebs solution. Segments of the middle cerebral artery (~ 2 mm long) were dissected and stored in ice-cold Krebs for use within 30 min, with similar size vessels used in all experimental groups

2.2 Experimental protocols

Segments of middle cerebral artery (internal diameter ~150 μm) were mounted in a Mulvany-Halpern myograph (model 400A, Danish Myotechnology) in Krebs solution containing (mM): NaCl, 118.0, NaCO_3 , 24; KCl, 3.6; $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$, 1.2; glucose, 11.0; CaCl_2 , 2.5; gassed with 20% O_2 , 5% CO_2 and balance N_2 and maintained at 37°C. After equilibration for 20min, vessels were tensioned to 1-1.5mN (approximates wall tension at 60mmHg). Smooth muscle tension was recorded with an isometric force transducer and Powerlab software (ADI, Australia). Vessel viability was assessed by addition of exogenous K^+ (15-55 mM, total K^+ concentration); only vessels developing tension of ≥ 3 mN were used, following this Endothelial cell viability was assessed by the ability of the protease activated receptor 2 activating peptide; SLIGRL (20 μM) (Alexander *et al.*, 2008) to relax U46619 induced tone (100 nM) by ≥ 75 %, vessels with less relaxation were discarded. In some experiments, endothelial cells were removed by gently rubbing the luminal surface with a human hair; subsequent relaxation of < 15 % to SLIGRL (20 μM) was considered as successful removal and further abrasion often lead to damage of smooth muscle cells.

L-NAME (100 μM), indomethacin (10 μM) and iberiotoxin (100 nM) were added throughout the experiment (to block NO synthase (NOS), cyclooxygenase and BK_{Ca} channels, respectively), unless otherwise stated. In combination, these drugs gave a robust and sustained constriction (similar to vasospasm), and an associated rapid vasomotion. Similar responses were recorded in each case after inhibition of NOS alone, but vasomotion was more variable between preparations. Indomethacin had no effect on oscillations, but was included in the experimental cocktail to minimize any potential for confounding thromboxane signaling after NOS inhibition (Benyo *et al.*, 1998; McNeish *et al.*, 2007). Recordings were assessed in the presence of: the T-type (mibefradil 100 nM and NNC 55-0396 300 nM) and L-type (nifedipine 1 μM) Ca^{2+} channel blockers, the K_{Ca} channel blockers, apamin ($\text{K}_{\text{Ca}2.3}$ (SK_{Ca}), 50 nM), TRAM-34 ($\text{K}_{\text{Ca}3.1}$ (IK_{Ca}), 1 μM), iberiotoxin (BK_{Ca} , $\text{K}_{\text{Ca}1.1}$, 100nM) and charybdotoxin ($\text{K}_{\text{Ca}3.1}$, BK_{Ca} , 100 nM), the K_{IR} channel inhibitors BaCl_2 (30 μM) and CsCl_2 (10 mM), the Na^+/K^+ -ATPase inhibitor, ouabain (1 μM) and the voltage-gated

K⁺channel inhibitor, 4-aminopyridine (4-AP, 3 mM). Papaverine (150 μM) was added at the end of each experiment to assess overall tone. All blocking drugs were incubated for at least 20 min before data was recorded to ensure maximal effect. In most experiments smooth muscle membrane potential (E_m) and tension were measured simultaneously as previously described, using glass microelectrodes (filled with 2 M KCl; tip resistance, 80-120 MΩ) to measure E_m (Garland *et al.*, 1992).

2.3 Data analysis and statistical procedures

Results are expressed as the mean±s.e.mean of *n* animals. Tension values are given in mN (always per 2 mm segment) and E_m as mV. During the vasospastic vasomotion E_m is expressed as the mean E_m over a random 10s period of the rapid vasomotion where possible we have also reported the size of the depolarizing oscillations in mV. Vasodilatation is expressed as percentage reduction of the total vascular tone (myogenic tone plus vasoconstrictor response induced by either U46619 or the combination of L-NAME and iberiotoxin, as appropriate), quantified by relaxation with papaverine (150 μM). Graphs were drawn and comparisons made using either Student's t-test, or one-way ANOVA with Tukey's post-hoc test using Prism software (Graphpad, USA). $P \leq 0.05$ was considered significant.

2.4 Drugs, chemicals, reagents and other materials

Exogenous K⁺ was added as an isotonic physiological salt solution in which all the NaCl was replaced with an equivalent amount of KCl. Concentrations of K⁺ used are expressed as final bath concentration. L-NAME (N^G-nitro-L-arginine methyl ester), indomethacin, mibefradil, NNC 55-0396, nifedipine, barium chloride, cesium chloride, ouabain, 4-aminopyridine and papaverine were all obtained from Sigma (Poole, U.K.). U46619 (9,11-Dideoxy-11 α ,9 α -epoxymethanoprostaglandin F2 α) was from Calbiochem (UK). Apamin, charybdotoxin, iberiotoxin and tetrodotoxin from Latoxan (Valence, France). SLIGRL (serine-leucine-isoleucine-glycine-arginine-leucine-NH₂)

from Auspep (Parkville, Australia). TRAM-34 was a generous gift from Dr H. Wulff (University of California, Davis). All stock solutions were dissolved in distilled water except SLIGRL and charybdotoxin, dissolved in 0.9% NaCl, U46619 and TRAM-34, dissolved in dimethylsulfoxide (DMSO), nifedipine, dissolved in ethanol, and indomethacin which was dissolved in Na₂CO₃ (2%); vehicle controls were performed for drugs dissolved in DMSO ethanol and Na₂CO₃. All nomenclature conforms to the BJP guide to receptors and ion channels (Alexander *et al.*, 2008)

3. Results

3.1. Effect of inhibiting Nitric oxide synthase and BK_{Ca} on E_m and tension.

Rat middle cerebral arteries developed spontaneous myogenic tone equivalent to 1.3±0.1 mN (≈15 % of maximum tension, with 55 mmol/L KCl, n=40) with a resting membrane potential (E_m) of -42.1±0.9 mV (n=40). Addition of the NOS inhibitor, L-NAME (100 μM), and the cyclooxygenase inhibitor, indomethacin (10 μM) tended to evoke smooth-muscle cell depolarization (E_m -39.2±1.0 mV, n=38) and constriction (4.6±0.2 mN, n=40; *P*<0.05). In all vessels, the depolarization developed into ongoing oscillations (amplitude 7±0.9 mV) followed by equivalent changes in tone (amplitude 0.12±0.01 mN). We have previously reported similar observations in the rat middle cerebral artery following NOS inhibition alone (Yuill *et al.*, 2010). With L-NAME present, stimulation of the endothelium with 20μM SLIGRL evoked hyperpolarization of 20.9±1.7 mV (n=31) associated with 77.6±3.6 % relaxation (n=37).

In the presence of L-NAME and indomethacin, the BK_{Ca} channel inhibitor, iberiotoxin (100 nM), evoked further depolarization (to E_m -35.7±1.1 mV, n=40; *P*<0.05) and constriction (to 5.7±0.2 mN, n=41; *P*<0.05), associated with a marked increase in the amplitude of oscillations in E_m temporally linked to tension (Figure 1B). The oscillations in E_m and tension had a frequency of 0.84±0.02 Hz and 0.80±0.05 Hz and amplitude of 22.6±1.3 mV and 0.19±0.02 mN, respectively (n=41). All

subsequent experiments were performed in the presence of L-NAME, indomethacin and iberiotoxin unless stated. Removal of the endothelium abolished SLIGRL-mediated relaxation (20 μ M), but failed to affect oscillations in E_m (frequency of 0.76 ± 0.10 Hz; amplitude of 19.4 ± 4.3 mV) and tension (frequency of 0.66 ± 0.10 Hz; amplitude of 0.16 ± 0.05 mN, $n=3$).

3.2 $[Ca^{2+}]_o$ but not Na^+ or Cl^- currents modify oscillations in membrane potential

The voltage-dependent Na^+ channel blocker, tetrodotoxin (1 μ M) did not modify either the frequency or amplitude of oscillations in E_m (control: frequency of 0.84 ± 0.03 Hz, amplitude of 16.1 ± 1.4 mV; tetrodotoxin: frequency of 0.90 ± 0.02 Hz, amplitude of 15.4 ± 2.7 mV, $n=3$) and tension (control: frequency of 0.73 ± 0.11 Hz, amplitude of 0.13 ± 0.01 mN; tetrodotoxin: frequency of 0.67 ± 0.14 Hz, amplitude of 0.12 ± 0.01 mN, $n=3$).

To investigate if the oscillations were dependent upon Ca^{2+} influx, we decreased extracellular Ca^{2+} in steps from 2.5 to 0 mM. Simultaneous measurements of changes in E_m and tension showed that decreasing Ca^{2+} evoked depolarization (from -37.0 ± 2.1 to -26.8 ± 2.4 mV, $n=7$; $P<0.05$) and relaxation (from 5.4 ± 0.6 to 0.3 ± 0.1 mN, $n=7$; $P<0.05$). Under these conditions, the amplitude of oscillations in E_m and tension was diminished (Figure 1C and D). The calcium-dependent Cl^- channel inhibitor, DIDS (150 μ M), did not alter spontaneous oscillations in E_m or tension (data not shown) nor did it modify E_m (control: -44.4 ± 3.0 mV; DIDS: -49.2 ± 3.9 mV, $n=5$) or tension (control: 4.7 ± 0.6 mN; DIDS: 3.6 ± 0.5 mN, $n=5$).

3.4. Effect of Ca^{2+} channel blockers on oscillations in E_m and tension

The T-type selective Ca^{2+} channel blocker, mibefradil (100 nM) abolished oscillations in E_m (Figure 2A), significantly reduced oscillations in tension (control: frequency of 0.79 ± 0.07 Hz, amplitude of 0.16 ± 0.04 mN; mibefradil: frequency of 0.09 ± 0.05 Hz, amplitude of 0.04 ± 0.02 mN, $n=5$; $P<0.05$) and evoked relaxation (control: 5.1 ± 0.7 mN; mibefradil: 3.5 ± 0.6 mN, $n=5$) and depolarization (E_m

control: -36.1 ± 1.3 mV; E_m mibefradil: -30.4 ± 2.5 mV, $n=5$). The more selective T-type Ca^{2+} channel antagonist, NNC 55-0396 (300 nM) also significantly reduced oscillations in E_m (Figure 2B) and in tension (control: frequency of 0.79 ± 0.09 Hz, amplitude of 0.17 ± 0.03 mN; NNC 55-0396: frequency of 0.20 ± 0.07 Hz, amplitude of 0.07 ± 0.02 mN, $n=6$; $P < 0.05$) and evoked relaxation (control: 5.5 ± 0.5 mN; NNC 55-0396: 3.0 ± 0.5 mN, $n=6$; $P < 0.05$). However, NNC 55-0396 did not significantly modify mean E_m (E_m control: -36.2 ± 3.2 mV; E_m NNC 55-0396: -31.9 ± 3.2 mV, $n=6$). Further addition of the L-type voltage-gated Ca^{2+} channel inhibitor, nifedipine (1 μ M), relaxed (control: 5.8 ± 0.7 mN; NNC 55-0396 3.5 ± 0.6 mN; NNC 55-0396+nifedipine: 0.5 ± 0.1 mN, $n=4$; $P < 0.05$) and hyperpolarized (E_m control: -33.0 ± 3.7 mV; E_m NNC 55-0396: -30.0 ± 4.6 ; E_m NNC 55-0396+nifedipine: -36.7 ± 3.4 mV, $n=4$; Figure 2C). We also assessed the effect of NNC 55-0396 against middle cerebral artery basal myogenic tone in the absence of inhibitors. NNC 55-0396 (300 nM) did not significantly affect E_m (E_m control: -51.6 ± 2.47 ; E_m NNC 55-0396: -48.9 ± 2.0 mV, $n=5$) or relax myogenic tone (control: 1.04 ± 0.18 ; NNC 55-0396: 0.69 ± 0.09 mN) under these conditions. The additional presence of nifedipine (1 μ M) also failed to evoke hyperpolarization (E_m NNC 55-0396+ nifedipine: -51.5 ± 2.8 mV) but did cause significant relaxation (NNC 55-0396+ nifedipine: 0.44 ± 0.04 mN, $n=4$, $P < 0.05$). We have previously demonstrated similar relaxation of basal myogenic tone with nifedipine alone (Yuill *et al.*, 2010).

3.5. Involvement of $K_{Ca3.1}(IK_{Ca})$ and $K_{Ca2.3}(SK_{Ca})$ channels

Addition of the $K_{Ca3.1}$ channel blocker, TRAM-34 (1 μ M) markedly decreased the amplitude of oscillations in E_m and tension (Figure 3B) followed by small relaxation (control: 6.5 ± 0.4 mN; TRAM-34: 5.3 ± 0.2 mN, $n=5$; $P < 0.05$). TRAM-34 did not significantly reduce E_m , (E_m control: -33.9 ± 2.6 mV; E_m TRAM-34: -28.7 ± 2.3 mV, $n=4$). The effect of TRAM-34 on the amplitude of oscillations in E_m (control: 19.4 ± 4.3 mV; TRAM-34: 3.0 ± 1.6 mV, $n=3$; $P < 0.05$) and tension

(control: 0.16 ± 0.05 mN; TRAM-34: 0.04 ± 0.02 mN, $n=3$) was similar after endothelium removal. The oscillations in E_m and tension were not modified by the $K_{Ca2.3}$ blocker apamin (50 nM), either alone (Figure 3C) or in the additional presence of TRAM-34 (Figure 3D). Likewise, charybdotoxin (100 nM) alone (Figure 4B) or with apamin (Figure 4C) did not affect either oscillations in E_m and tension (Figure 4D and E), or mean tension and E_m .

3.6. Involvement of K_{IR} channels, voltage-gated K^+ channels and the Na^+/K^+ -ATPase

Inhibition of K_{IR} channels with $BaCl_2$ (30 μ M) did not affect the amplitude of oscillations in E_m (control: 31.1 ± 4.6 mV; $BaCl_2$: 30.9 ± 6.1 mV, $n=5$) or tension (control: 0.20 ± 0.03 mN; Ba^{2+} : 0.19 ± 0.03 mV, $n=5$). However, another inhibitor of K_{IR} , CsCl (10 mM) increased the amplitude of oscillations in E_m (control: 30.8 ± 2.0 mV; CsCl: 40.4 ± 3.4 mV, $n=5$; $P < 0.05$), but not in tension (control: 0.20 ± 0.02 mN; CsCl: 0.19 ± 0.02 mV, $n=5$), while 4-aminopyridine (4-AP; 3 mM to block K_v) increased the amplitude of oscillations in both E_m (control: 17.75 ± 2.07 ; 4-AP: 30.62 ± 7.33 mV, $n=3$) and tension (control: 0.17 ± 0.01 ; 4-AP: 0.25 ± 0.07 mN; $n=3$, Figure 5A). The Na^+/K^+ -ATPase inhibitor, ouabain (1 μ M) evoked relaxation (control: 5.8 ± 0.3 mN; ouabain: 4.6 ± 0.2 mN, $n=3$; $P < 0.05$) but without significantly reducing E_m (control: -42.2 ± 1.2 mV; ouabain: -38.4 ± 1.6 mV, $n=3$). However, ouabain did reduce both the amplitude (Figure 5B and C) and frequency (control: 0.97 ± 0.11 Hz; ouabain: 0.31 ± 0.16 Hz, $n=3$; $P < 0.05$) of oscillations in E_m , without altering oscillations in tension (Figure 5C and frequency control: 1.30 ± 0.22 Hz; plus ouabain: 0.92 ± 0.22 Hz, $n=3$). The subsequent addition of 4-AP in the presence of ouabain caused depolarization (ouabain: -38.4 ± 1.6 mV; ouabain+4-AP: -34.6 ± 3.3 mV, $n=3$; $P < 0.05$), and increased oscillations in E_m (Figure 5C) and increased tension overall (ouabain: 4.6 ± 0.2 mN; ouabain+4-AP: 5.5 ± 0.2 mN, $n=3$; $P < 0.05$).

4. Discussion

This study provides the first demonstration, that rhythmic oscillations in membrane potential and tension as well as the associated spasm in rat isolated middle cerebral arteries following inhibition of BK_{Ca} channels and/or NOS reflect Ca²⁺ influx via T-type Ca²⁺ channels, in addition to L-type Ca²⁺ channels. These data extend our previous observation that NOS inhibition leads to L-type Ca²⁺ channel opening and arterial spasm, characterized by sustained constriction and superimposed by rapid vasomotion (Yuill *et al.*, 2010). Vasomotion was of much higher frequency than previously recorded in other vascular beds, and as such we refer to it as ‘vasospastic’ vasomotion. We also provide evidence for modulation of both the Ca²⁺ dependent vasomotion and constriction through Na/K ATPase and a K⁺ conductance.

Consistent with previous work, our data suggest that calcium influx underlies rhythmic oscillations in the constricted rat middle cerebral artery, and that oscillations in membrane potential and tension are linked to oscillations in intracellular [Ca²⁺]_i as well as spasm in both middle cerebral and basilar arteries (Haddock *et al.*, 2002; Navarro-Gonzalez *et al.*, 2009; Yuill *et al.*, 2010). Reducing extracellular calcium diminished the amplitude of oscillations, led to relaxation and paradoxically depolarized the membrane. The relaxation presumably reflected the reduction in peak E_m associated with the reduced amplitude of oscillations in E_m. Some small oscillations in tension did persist and may reflect vasomotion-independent intracellular calcium release, as reported in some arterial beds: for review see Haddock *et al.*, 2005 (Haddock *et al.*, 2005). As oscillations were insensitive to the calcium-dependent Cl⁻ channel inhibitor, DIDS and the voltage-dependent Na⁺ channel blocking agent, tetrodotoxin they certainly appeared to be mediated exclusively by calcium conductance. These data contrast with the basilar artery, where inhibition of Cl⁻ channels in this larger artery abolished calcium-dependent oscillations, leading to hyperpolarization and relaxation (Haddock *et al.*, 2002), and parenchymal arterioles, where calcium-dependent oscillations were blocked with tetrodotoxin (Filosa *et al.*, 2004). However, our data are consistent with human pial arteries, where tetrodotoxin was also without effect against oscillations in diameter (Gokina *et al.*, 1996). In the

present study, oscillations in muscle membrane potential were also resistant to direct damage of the endothelium, suggesting this monolayer may not influence rapid vasomotion associated with arterial spasm.

In the rat middle cerebral artery, opening L-type Ca^{2+} channels is essential for vasoconstriction and vasomotion to develop, because inhibition of these channels abolishes vasomotion and fully reverses tone (Yuill *et al.*, 2010). Surprisingly, in the present study under similar vasospastic conditions, oscillations in E_m , and the associated oscillations in tension were abolished and followed by relaxation after calcium influx through T-type Ca^{2+} channels was blocked with mibefradil (at a concentration selective for block of T-type Ca^{2+} channels; 100 nM) or with a T-type Ca^{2+} channel selective, non-hydrolysable analogue of mibefradil, NNC 55-0396, (Huang *et al.*, 2004). Block of oscillations with the putative T-type Ca^{2+} channel blockers was not associated with a net hyperpolarization and complete relaxation, contrasting with the L-Type Ca^{2+} channel blocker nifedipine. So the effect of mibefradil or NNC 55-0396 is unlikely to reflect a non-specific effect against L-type channels. Furthermore, although mibefradil apparently reduced blood pressure and myogenic tone by an action on L-type calcium channels (Moosmang *et al.*, 2006), the lower concentration of mibefradil used in the present study is relatively specific against T-type Ca^{2+} channels. In fact, mibefradil seems only to inhibit the L-type Ca^{2+} channels after tissue metabolism (Wu *et al.*, 2000). The non-hydrolysable analogue of mibefradil NNC 55-0396 is selective for T-type Ca^{2+} channels, with no reported block of L-type Ca^{2+} channels even in concentrations as high as 100 μM (Huang *et al.*, 2004). Mibefradil has also been reported to block both Cl^- (Bernd *et al.*, 1997) and Na^+ (Eller *et al.*, 2000; Guatimosim *et al.*, 2001) channels, but again with much higher concentrations (μM) than employed in the current study. Furthermore, the fact that blockers such as DIDs and TTX had no effect against vasospastic vasomotion makes an action of mibefradil against these channels extremely unlikely. Therefore, the differential effect of mibefradil and NNC-0396 compared to the selective blocker L-type Ca^{2+} channel blocker nifedipine indicate a critical role for

T-type Ca^{2+} channels in vasospastic vasomotion. Further, these channels contribute significantly to overall constriction in the middle cerebral artery.

Vasospastic vasomotion only developed once the smooth muscle cells depolarized to *circa* -40 mV, so it may be that T-type channels involved in vasomotion have gating properties similar to high voltage activated Ca^{2+} channels. This is surprising, as by definition T-type Ca^{2+} channels activate at low potentials and then quickly inactivate (Lacinova, 2005). However, our data are consistent with studies reporting T-type Ca^{2+} channels that influence vascular tone and have properties similar to high-voltage activated Ca^{2+} channels, (Navarro-Gonzalez *et al.*, 2009). Both T- and L-type Ca^{2+} channels are expressed in rat basilar (Navarro-Gonzalez *et al.*, 2009) and middle cerebral arteries (Kuo *et al.*, 2008), and in each artery the $\text{CaV}_{3.2}$ (or T-type) is the most abundant VGCC alpha subunit expressed. Human recombinant T-type Ca^{2+} channels (Kaku *et al.*, 2003) and T-type Ca^{2+} channels co-expressed with auxiliary subunits (Wyatt *et al.*, 1998) do have gating properties similar to high voltage activated channels, so channels in the middle cerebral artery may be similar. Peripheral arteries also contain VGCCs with similar biophysical properties to high voltage activated channels (-50 to -20 mV), and are pharmacologically indistinguishable from T-type Ca^{2+} channels in both the guinea-pig and rat (Morita *et al.*, 1999; Morita *et al.*, 2002). In these small mesenteric arteries, T-type Ca^{2+} channels are the predominant voltage-gated subtype (Gustafsson *et al.*, 2001; Jensen *et al.*, 2004), and show increased window current due to non-inactivation at physiological E_m (Jensen *et al.*, 2009).

As nifedipine abolished vasomotion, we propose that L-type channels are key for the initial depolarization and constriction (Yuill *et al.*, 2010); whereas T-type are activated subsequently and as such are critical for the vasospastic vasomotion. Interestingly, input from T-type Ca^{2+} channels seem to be important for the initial constriction in the basilar artery, whereas L-type channels are critical for vasomotion (Navarro-Gonzalez *et al.*, 2009). But taken together, these results all suggest

a functional coupling between L- and T-type Ca^{2+} channels, as previously suggested in renal (Hansen *et al.*, 2001) and mesenteric arterioles (Braunstein *et al.*, 2008). As such, this might explain why neuroprotection in ischemic stroke is more effective in patients given blockers for more than just L-type VGCCs (Kobayashi *et al.*, 1998).

As VGCC and hence vasospastic vasomotion are inhibited by a complex interaction between NO and smooth muscle cell BK_{Ca} channels in middle cerebral arteries, we attempted to characterise further the vasospastic vasomotion. Endothelial cell damage did not affect the vasomotion, so we inhibited a variety of K^{+} currents. Both inwardly rectifying and voltage-gated K^{+} channels participate in maintenance of resting membrane potential and vascular tone (Ko *et al.*, 2008; Nelson *et al.*, 1995; Sobey, 2001). However, inhibition of K_{IR} channels with CsCl or barium did not affect vascular tone, although CsCl did slightly increase E_{m} oscillation amplitude. As Ba^{2+} was without effect, this small change most likely reflected a non-selective action of CsCl. Inhibition of K_{v} channels with 4-AP also had little effect, causing only a small increase in the amplitude of oscillations in E_{m} and tension. This is consistent with the reported role of these channels in rat mesenteric artery where inhibition of K_{v} increased rhythmic contractions (Gustafsson *et al.*, 1994). So voltage-gated K^{+} channels did not appear to play any major role in vasospastic vasomotion.

Our data do suggest that $\text{Na}^{+}/\text{K}^{+}$ -ATPase might contribute to vasomotion, as ouabain caused relaxation and reduced the amplitude and frequency of oscillations in E_{m} , although surprisingly without affecting oscillations in tension. Ouabain can attenuate intercellular communication in smooth muscle (Harris *et al.*, 2000; Martin *et al.*, 2003; Matchkov *et al.*, 2007) and the synchronized changes in vascular $[\text{Ca}^{2+}]_{\text{i}}$ (Koenigsberger *et al.*, 2004) that lead to vasomotion (Chaytor *et al.*, 1997; Matchkov *et al.*, 2004; Peng *et al.*, 2001). So in part, ouabain may alter membrane potential oscillations by modifying cell-cell communication. Interestingly, ouabain

effects were reversed by 4-AP, again indicating that K_V might contribute under some conditions to influence vasomotion.

The ability of the $K_{Ca3.1}$ channel inhibitor TRAM-34 to reduce rather than enhance the amplitude of oscillations in E_m and tension was also unexpected. This effect was on the smooth muscle, as it was not altered by removal of the endothelium, and in contrast to TRAM-34, charybdotoxin, a mixed BK_{Ca} and $K_{Ca3.1}$ inhibitor failed to modify the oscillations. One explanation, is that TRAM-34 inhibits non-selective cation channels in the smooth muscle, similar to its action in isolated immune cells (Schilling *et al.*, 2007). Non-selective cation channels are present in rat middle cerebral artery smooth muscle cells and appear to contribute to the calcium entry and vascular tone (Marrelli *et al.*, 2007; Welsh *et al.*, 2002). So data with TRAM-34 suggest that non-selective cation channels may play an important role in the calcium entry events underpinning depolarization and vasomotion after NOS inhibition in the middle cerebral artery.

In summary, inhibition of either BK_{Ca} channels and/or NOS evokes vasospasm and fast, rhythmic oscillations in E_m and tension that are mediated by Ca^{2+} influx via both T-type and L-type Ca^{2+} channels. Our data suggest the T-type channels are active at physiologically relevant membrane potentials and can therefore make an important contribution to the control of cerebrovascular blood flow during vasospasm associated with disease states where NO synthesis or action is impaired, such as cerebral ischemia or subarachnoid haemorrhage.

Acknowledgments: This work was supported by the British Heart Foundation (FS/06/076/21988) and the Wellcome Trust. Francesc Jiminez-Altayo and Alister McNeish contributed equally to this manuscript

Figure Legends

Figure 1. Original traces showing simultaneous recordings of membrane potential (upper panels) and tension (lower panels) in rat middle cerebral arteries under control resting conditions (A) or in the presence of L-NAME (100 μM), indomethacin (10 μM) and iberiotoxin (100 nM; B). Under control conditions, membrane potential and tension are relatively stable. In the presence of L-NAME, indomethacin and iberiotoxin, the smooth muscle cells depolarized and developed regular depolarizing oscillations, which were associated with constriction and oscillations in tension; the peaks in membrane potential immediately preceded peaks in tension. Decreasing extracellular calcium (from 2.5 to 0 mM Ca^{2+}) caused depolarization and relaxation. Average data are shown in (C and D) showing the oscillation amplitude in membrane potential (C) and tension (D) in control vessels (2.5 mM Ca^{2+}) and in the presence of 1, 0.5, 0.25 and 0 mM Ca^{2+} . Data expressed as means \pm S.E.M. * $P \leq 0.05$ indicates a significant difference from control by one-way ANOVA with Tukey's post-hoc test, $n=3-7$.

Figure 2. Original traces showing the effect of either (A) the T-type Ca^{2+} channel blocker, mibefradil (100 nM), (B) the more selective T-type Ca^{2+} channel antagonist, NNC 55-0396 (300 nM) or (C) the combined application of NNC 55-0396 (300 nM) and the L-type Ca^{2+} channel blocker, nifedipine (1 μM), on simultaneous recordings of membrane potential (upper panels) and tension (lower panels). Both mibefradil and NNC 55-0396 caused relaxation and abolished oscillations in membrane potential and tension; combined application of NNC 55-0396 and nifedipine caused greater hyperpolarization and relaxation than NNC-550396 alone, $n=4-6$. Parallel lines (//) indicate a time break between same recordings from a single vessel the first time break shows the response at approximately 5 min post addition of drug the second time break corresponds to the maximum response which is approximately 15 min following addition .

Figure 3. Original traces showing control conditions (A), the effect of the $K_{Ca}3.1$ channel blocker, TRAM-34 (1 μ M; B), the $K_{Ca}2.3$ channel blocker, apamin (50 nM; C) or the combined application of TRAM-34 and apamin (D) on simultaneous recordings of membrane potential (upper panels) and tension (lower panels). Average data for oscillation amplitude in membrane potential (left panel) and tension (right panel) in control vessels and in the presence of TRAM-34, apamin and TRAM-34 + apamin are also shown (E). Data are expressed as means \pm S.E.M. * $P \leq 0.05$ indicates a significant difference from control by one-way ANOVA with Tukey's post-hoc test, $n=3-9$.

Figure 4. Original traces showing control conditions (A) and the effect of either the BK_{Ca} and $K_{Ca}3.1$ channel blocker, charybdotoxin (100 nM; B) or the combined application of charybdotoxin (100 nM) and the $K_{Ca}2.3$ channel blocker, apamin (50 nM; C) on simultaneous recordings of membrane potential (upper panels) and tension (lower panels). Average data are shown for the oscillation amplitude in membrane potential (D) and tension (E) in control vessels and in the presence of charybdotoxin and charybdotoxin + apamin. Data expressed as means \pm S.E.M. $n=3-4$.

Figure 5. Original traces showing (A) the effect of the voltage-gated K^+ channel inhibitor, 4-AP (3 mM) and (B) the effect of the Na^+/K^+ -ATPase inhibitor, ouabain (1 μ M) on simultaneous recordings of membrane potential (upper panels) and tension (lower panels). Average data are shown in (C) showing the oscillation amplitude in membrane potential (left panel) and tension (right panel) in control vessels and in the presence of ouabain and ouabain + 4-AP. Data expressed as means \pm S.E.M. */ $P \leq 0.05$ indicates a significant difference from either control or from ouabain alone, respectively, using one-way ANOVA with Tukeys post-hoc test, $n=3$.

References:

Alexander, SPH, Mathie, A, Peters, JA (2008) Guide to Receptors and Channels (GRAC), 3rd edition. *Brit J Pharmacol* **153**(S2): S1-S1.

- Benyo, Z, Gorlach, C, Wahl, M (1998) Involvement of Thromboxane A2 in the Mediation of the Contractile Effect Induced by Inhibition of Nitric Oxide Synthesis in Isolated Rat Middle Cerebral Arteries. *J Cereb Blood Flow Metab* **18**(6): 616-618.
- Bernd, N, Jean, P, Masahiro, K, Felix, V, Thomas, V, Guy, D (1997) Inhibition by mibefradil, a novel calcium channel antagonist, of Ca²⁺- and volume-activated Cl⁻ channels in macrovascular endothelial cells. *Brit J Pharmacol* **121**(3): 547-555.
- Biswal, BB, Hudetz, AG (1996) Synchronous oscillations in cerebrocortical capillary red blood cell velocity after nitric oxide synthase inhibition. *Microvasc Res* **52**(1): 1-12.
- Braunstein, TH, Inoue, R, Cribbs, L, Oike, M, Ito, Y, Holstein-Rathlou, NH, Jensen, LJ (2008) The Role of L- and T-Type Calcium Channels in Local and Remote Calcium Responses in Rat Mesenteric Terminal Arterioles. *J Vasc Res* **46**(2): 138-151.
- Brayden, JE, Nelson, MT (1992) Regulation of arterial tone by activation of calcium-dependent potassium channels. *Science* **256**(5056): 532-535.
- Chaytor, AT, Evans, WH, Griffith, TM (1997) Peptides homologous to extracellular loop motifs of connexin 43 reversibly abolish rhythmic contractile activity in rabbit arteries. *J Physiol* **503** (Pt 1): 99-110.
- Clozel, JP, Ertel, EA, Ertel, SI (1997) Discovery and main pharmacological properties of mibefradil (Ro 40-5967), the first selective T-type calcium channel blocker. *J Hypertens Suppl* **15**(5): S17-25.
- Davis, MJ, Hill, MA (1999) Signaling Mechanisms Underlying the Vascular Myogenic Response. *Physiol Rev* **79**(2): 387-423.
- Dirnagl, U, Lindauer, U, Villringer, A (1993) Nitric oxide synthase blockade enhances vasomotion in the cerebral microcirculation of anesthetized rats. *Microvasc Res* **45**(3): 318-323.
- Eller, P, Berjukov, S, Wanner, S, Huber, I, Hering, S, Knaus, H-G, Toth, G, Kimball, SD, Striessnig, J (2000) High affinity interaction of mibefradil with voltage-gated calcium and sodium channels. *Brit J of Pharmacol* **130**(3): 669-677.
- Filosa, JA, Bonev, AD, Nelson, MT (2004) Calcium Dynamics in Cortical Astrocytes and Arterioles During Neurovascular Coupling. *Circ Res* **95**(10): e73-81.
- Garland, CJ, McPherson, GA (1992) Evidence that nitric oxide does not mediate the hyperpolarisation and relaxation to acetylcholine in the rat small mesenteric artery. *Br J Pharmacol* **105**: 429-435.
- Gokina, NI, Bevan, RD, Walters, CL, Bevan, JA (1996) Electrical activity underlying rhythmic contraction in human pial arteries. *Circ Res* **78**(1): 148-153.
- Golding, EM, Steenberg, ML, Johnson, TD, Bryan, RM (2001) Nitric oxide in the potassium-induced response of the rat middle cerebral artery: a possible permissive role. *Brain Res* **889**(1-2): 98-104.
- Guatimosim, S, Sobie, EA, dos Santos Cruz, J, Martin, LA, Lederer, WJ (2001) Molecular identification of a TTX-sensitive Ca²⁺ current. *Am J Physiol Cell Physiol* **280**(5): C1327-1339.

- Gustafsson, F, Andreasen, D, Salomonsson, M, Jensen, BL, Holstein-Rathlou, N (2001) Conducted vasoconstriction in rat mesenteric arterioles: role for dihydropyridine-insensitive Ca²⁺ channels. *Am J Physiol Heart Circ Physiol* **280**(2): H582-590.
- Gustafsson, H (1993) Vasomotion and underlying mechanisms in small arteries. An in vitro study of rat blood vessels. *Acta Physiol Scand Suppl* **614**: 1-44.
- Gustafsson, H, Nilsson, H (1994) Rhythmic contractions in isolated small arteries of rat: role of K⁺ channels and the Na⁺,K⁽⁺⁾-pump. *Acta Physiol Scand* **150**(2): 161-170.
- Haddock, RE, Hill, CE (2002) Differential activation of ion channels by inositol 1,4,5-trisphosphate (IP₃)- and ryanodine-sensitive calcium stores in rat basilar artery vasomotion. *J Physiol* **545**(Pt 2): 615-627.
- Haddock, RE, Hill, CE (2005) Rhythmicity in arterial smooth muscle. *J Physiol* **566**(Pt 3): 645-656.
- Hansen, PB, Jensen, BL, Andreasen, D, Skott, O (2001) Differential expression of T- and L-type voltage-dependent calcium channels in renal resistance vessels. *Circ Res* **89**(7): 630-638.
- Harris, D, Martin, PEM, Evans, WH, Kendall, DA, Griffith, TM, Randall, MD (2000) Role of gap junctions in endothelium-derived hyperpolarizing factor responses and mechanisms of K⁺-relaxation. *European Journal of Pharmacology* **402**(1-2): 119-128.
- Hill, MA, Zou, H, Potocnik, SJ, Meininger, GA, Davis, MJ (2001) Signal Transduction in Smooth Muscle: Invited Review: Arteriolar smooth muscle mechanotransduction: Ca²⁺ signaling pathways underlying myogenic reactivity. *J Appl Physiol* **91**(2): 973-983.
- Huang, L, Keyser, BM, Tagmose, TM, Hansen, JB, Taylor, JT, Zhuang, H, Zhang, M, Ragsdale, DS, Li, M (2004) NNC 55-0396 [(1S,2S)-2-(2-(N-[(3-benzimidazol-2-yl)propyl]-N-methylamino)ethyl)-6-fluoro-1,2,3,4-tetrahydro-1-isopropyl-2-naphthyl cyclopropanecarboxylate dihydrochloride]: a new selective inhibitor of T-type calcium channels. *J Pharmacol Exp Ther* **309**(1): 193-199.
- Jaggard, JH, Porter, VA, Lederer, WJ, Nelson, MT (2000) Calcium sparks in smooth muscle. *Am J Physiol Cell Physiol* **278**(2): C235-256.
- Jensen, LJ, Holstein-Rathlou, NH (2009) Is there a role for T-type Ca²⁺ channels in regulation of vasomotor tone in mesenteric arterioles? *Can J Physiol Pharmacol* **87**(1): 8-20.
- Jensen, LJ, Salomonsson, M, Jensen, BL, Holstein-Rathlou, N-H (2004) Depolarization-induced calcium influx in rat mesenteric small arterioles is mediated exclusively via mibefradil-sensitive calcium channels. *Br J Pharmacol* **142**(4): 709-718.
- Jewell, RP, Saundry, CM, Bonev, AD, Tranmer, BI, Wellman, GC (2004) Inhibition of Ca⁺⁺ sparks by oxyhemoglobin in rabbit cerebral arteries. *J Neurosurg* **100**(2): 295-302.
- Kaku, T, Lee, TS, Arita, M, Hadama, T, Ono, K (2003) The gating and conductance properties of Cav3.2 low-voltage-activated T-type calcium channels. *Jpn J Physiol* **53**(3): 165-172.
- Knot, HJ, Nelson, MT (1998) Regulation of arterial diameter and wall [Ca²⁺] in cerebral arteries of rat by membrane potential and intravascular pressure. *J Physiol* **508** (Pt 1): 199-209.

- Ko, EA, Han, J, Jung, ID, Park, WS (2008) Physiological roles of K⁺ channels in vascular smooth muscle cells. *J Smooth Muscle Res* **44**(2): 65-81.
- Kobayashi, T, Mori, Y (1998) Ca²⁺ channel antagonists and neuroprotection from cerebral ischemia. *Eur J Pharmacol* **363**(1): 1-15.
- Koenigsberger, M, Sauser, R, Lamboley, M, Beny, JL, Meister, JJ (2004) Ca²⁺ dynamics in a population of smooth muscle cells: modeling the recruitment and synchronization. *Biophys J* **87**(1): 92-104.
- Kuo, IK, Cribbs, LL, Grayson, TH (2008) Expression of multiple voltage dependent calcium channels in the middle cerebral artery of adult rats. In: *9th International Symposium on Resistance Arteries (ISRA)*. Hamilton Island, Australia.
- Lacinova, L (2005) Voltage-dependent calcium channels. *Gen Physiol Biophys* **24 Suppl 1**: 1-78.
- Lam, E, Skarsgard, P, Laher, I (1998) Inhibition of myogenic tone by mibefradil in rat cerebral arteries. *Eur J Pharmacol* **358**(2): 165-168.
- Mandala, M, Heppner, TJ, Bonev, AD, Nelson, MT (2007) Effect of endogenous and exogenous nitric oxide on calcium sparks as targets for vasodilation in rat cerebral artery. *Nitric Oxide* **16**(1): 104-109.
- Marrelli, SP, O'Neil, RG, Brown, RC, Bryan, RM, Jr. (2007) PLA2 and TRPV4 channels regulate endothelial calcium in cerebral arteries *Am J Physiol* Vol. 292, pp H1390-1397.
- Martin, PEM, Hill, NS, Kristensen, B, Errington, RJ, Griffith, TM (2003) Ouabain exerts biphasic effects on connexin functionality and expression in vascular smooth muscle cells. *Br J Pharmacol* **140**(7): 1261-1271.
- Martin, W, Villani, GM, Jothianandan, D, Furchgott, RF (1985) Selective blockade of endothelium-dependent and glyceryl trinitrate- induced relaxation by hemoglobin and by methylene blue in the rabbit aorta. *J Pharmacol Exp Ther* **232**(3): 708-716.
- Matchkov, VV, Gustafsson, H, Rahman, A, Briggs Boedtkjer, DM, Gorintin, S, Hansen, AK, Bouzinova, EV, Praetorius, HA, Aalkjaer, C, Nilsson, H (2007) Interaction between Na⁺/K⁺-pump and Na⁺/Ca²⁺-exchanger modulates intercellular communication. *Circ Res* **100**(7): 1026-1035.
- Matchkov, VV, Rahman, A, Peng, H, Nilsson, H, Aalkjaer, C (2004) Junctional and nonjunctional effects of heptanol and glycyrrhetic acid derivatives in rat mesenteric small arteries. *Br J Pharmacol* **142**(6): 961-972.
- McCarron, JG, Crichton, CA, Langton, PD, MacKenzie, A, Smith, GL (1997) Myogenic contraction by modulation of voltage-dependent calcium currents in isolated rat cerebral arteries. *J Physiol* **498** (Pt 2): 371-379.
- McNeish, AJ, Dora, K, Garland, C (2006) NO can suppress myogenic tone in rat middle cerebral arteries by activating BKc. *Faseb Journal* **20**(5): A1113-A1113.
- McNeish, AJ, Garland, CJ (2007) Thromboxane A2 inhibition of SKCa after NO synthase block in rat middle cerebral artery. *Br J Pharmacol* **151**(4): 441-449.

- Moosmang, S, Haider, N, Bruderl, B, Welling, A, Hofmann, F (2006) Antihypertensive Effects of the Putative T-Type Calcium Channel Antagonist Mibefradil Are Mediated by the L-Type Calcium Channel Cav1.2. *Circ Res* **98**(1): 105-110.
- Moosmang, S, Schulla, V, Welling, A, Feil, R, Feil, S, Wegener, JW, Hofmann, F, Klugbauer, N (2003) Dominant role of smooth muscle L-type calcium channel Cav1.2 for blood pressure regulation. *EMBO J* **22**(22): 6027-6034.
- Morita, H, Cousins, H, Onoue, H, Ito, Y, Inoue, R (1999) Predominant distribution of nifedipine-insensitive, high voltage-activated Ca²⁺ channels in the terminal mesenteric artery of guinea pig. *Circ Res* **85**(7): 596-605.
- Morita, H, Shi, J, Ito, Y, Inoue, R (2002) T-channel-like pharmacological properties of high voltage-activated, nifedipine-insensitive Ca²⁺ currents in the rat terminal mesenteric artery. *Br J Pharmacol* **137**(4): 467-476.
- Navarro-Gonzalez, MF, Grayson, TH, Meaney, KR, Cribbs, LL, Hill, CE (2009) Non-L-Type Voltage-Dependent Calcium Channels Control Vascular Tone of the Rat Basilar Artery. *Clin Exp Pharmacol Physiol* **36**(1): 55-66.
- Nelson, MT, Patlak, JB, Worley, JF, Standen, NB (1990) Calcium channels, potassium channels, and voltage dependence of arterial smooth muscle tone. *Am J Physiol Cell Physiol* **259**(1): C3-18.
- Nelson, MT, Quayle, JM (1995) Physiological roles and properties of potassium channels in arterial smooth muscle. *Am J Physiol* **268**(4 Pt 1): C799-C822.
- Peng, H-L, Jensen, PE, Nilsson, H, Aalkjar, C (1998) Effect of acidosis on tension and [Ca²⁺]_i in rat cerebral arteries: is there a role for membrane potential? *Am J Physiol Heart Circ Physiol* **274**(2): H655-662.
- Peng, H, Matchkov, V, Ivarsen, A, Aalkjaer, C, Nilsson, H (2001) Hypothesis for the initiation of vasomotion. *Circ Res* **88**(8): 810-815.
- Perez-Reyes, E (2003) Molecular physiology of low-voltage-activated t-type calcium channels. *Physiol Rev* **83**(1): 117-161.
- Pluta, RM (2005) Delayed cerebral vasospasm and nitric oxide: review, new hypothesis, and proposed treatment. *Pharmacol Ther* **105**(1): 23-56.
- Schilling, T, Eder, C (2007) TRAM-34 inhibits nonselective cation channels. *Pflugers Arch* **454**(4): 559-563.
- Schubert, R, Lidington, D, Bolz, S-S (2008) The emerging role of Ca²⁺ sensitivity regulation in promoting myogenic vasoconstriction. *Cardiovasc Res* **77**(1): 8-18.
- Smirnov, SV, Aaronson, PI (1992) Ca²⁺ currents in single myocytes from human mesenteric arteries: evidence for a physiological role of L-type channels. *J Physiol* **457**(1): 455-475.
- Sobey, CG (2001) Potassium channel function in vascular disease. *Arterioscler Thromb Vasc Biol* **21**(1): 28-38.

- Toda, N, Kawakami, M, Yoshida, K (1991) Constrictor action of oxyhemoglobin in monkey and dog basilar arteries in vivo and in vitro. *Am J Physiol Heart Circ Physiol* **260**(2): H420-425.
- VanBavel, E, Sorop, O, Andreasen, D, Pfaffendorf, M, Jensen, BL (2002) Role of T-type calcium channels in myogenic tone of skeletal muscle resistance arteries. *Am J Physiol Heart Circ Physiol* **283**(6): H2239-2243.
- Vanhoutte, PM, Shimokawa, H, Tang, EH, Feletou, M (2009) Endothelial dysfunction and vascular disease. *Acta Physiol (Oxf)* **196**(2): 193-222.
- Welsh, DG, Morielli, AD, Nelson, MT, Brayden, JE (2002) Transient Receptor Potential Channels Regulate Myogenic Tone of Resistance Arteries. *Circ Res* **90**(3): 248-250.
- Wu, S, Zhang, M, Vest, PA, Bhattacharjee, A, Liu, L, Li, M (2000) A Mibefradil Metabolite Is a Potent Intracellular Blocker of L-Type Ca²⁺ Currents in Pancreatic beta -Cells. *J Pharmacol Exp Ther* **292**(3): 939-943.
- Wyatt, CN, Page, KM, Berrow, NS, Brice, NL, Dolphin, AC (1998) The effect of overexpression of auxiliary Ca²⁺ channel subunits on native Ca²⁺ channel currents in undifferentiated mammalian NG108-15 cells. *J Physiol* **510** (Pt 2): 347-360.
- Yuill, KH, McNeish, AJ, Kansui, Y, Garland, CJ, Dora, KA (2010) Nitric Oxide Suppresses Cerebral Vasomotion by sGC-Independent Effects on Ryanodine Receptors and Voltage-Gated Calcium Channels. *J Vasc Res* **47**(2): 93-107.
- Zimmermann, PA, Knot, HJ, Stevenson, AS, Nelson, MT (1997) Increased Myogenic Tone and Diminished Responsiveness to ATP-Sensitive K⁺ Channel Openers in Cerebral Arteries From Diabetic Rats. *Circ Res* **81**(6): 996-1004.