

RESEARCH PAPER

Endothelium-dependent hyperpolarization-related relaxations diminish with age in murine saphenous arteries of both sexes

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BACKGROUND AND PURPOSE

We investigated the effects of aging on the contributions of NO and endothelium-dependent hyperpolarization (EDH) to endothelium-dependent relaxation in saphenous arteries of male and female C57BL/6J mice aged 12, 34 and 64 weeks.

EXPERIMENTAL APPROACH

Vasomotor responses of saphenous arteries were analysed by wire myography in the absence and presence of stimuli of the endothelium, inhibitors of NOS, and inhibitors and stimulants of small ($K_{Ca2.3}$) and intermediate ($K_{Ca3.1}$) conductance calcium-activated potassium channels.

KEY RESULTS

Arterial relaxing responses to sodium nitroprusside and to ACh in the absence of pharmacological inhibitors (indomethacin and L-NAME), were similar in all age groups and sexes, but those mediated by endothelium-derived NO were slightly but significantly increased in 64-week-old male mice. In the presence of inhibitors, 12-week-old animals showed pronounced ACh-induced relaxation, which was significantly reduced in 34- and 64-week-old mice of both sexes. The EDH-related component of ACh-induced relaxations was abolished by TRAM-34 ($K_{Ca3.1}$ blocker) or UCL 1684 ($K_{Ca2.3}$ blocker). Although the maximal relaxation induced by NS309 (K_{Ca} activator) was not affected by aging, the sensitivity for NS309 significantly decreased with aging. The presence of SKA-31 (K_{Ca} modulator) potentiated relaxations induced by ACh in arteries of 12-week-old but not older mice.

CONCLUSION AND IMPLICATIONS

In a small muscular artery of mice of either sex, total endothelium-dependent relaxation is not affected by age. However, possibly due to changes in K_{Ca} channel function, the contribution of EDH to endothelium-dependent relaxations decreased with age. The contribution of endothelium-derived NO increases in old male mice.

Abbreviations

CRC, concentration response curve; DMSO, dimethyl sulfoxide; D_{opt} , optimal diameter; EDH, endothelium-dependent hyperpolarization; EDHF, endothelium-derived hyperpolarizing factor; EDNO, endothelium-derived NO; $K_{Ca2.3}/SK3$, small conductance calcium-activated potassium channel; $K_{Ca3.1}/IK1$, intermediate conductance calcium-activated

potassium channel; KRB, Krebs Ringer bicarbonate-buffered salt solution; INDO, indomethacin; L-NAME, N^ω-nitro-L-arginine methyl ester; NA, noradrenaline; NS309, 6, 7-dichloro-1H-indole-2, 3-dione 3-oxime; PHE, phenylephrine; SKA-31, naphtho[1,2-d]thiazol-2-ylamine; SNP, sodium nitroprusside; TRAM-34, 1-[(2-chlorophenyl)diphenylmethyl]-1H-pyrazole; UCL 1684, 6,12,19,20,25,26-hexahydro-5,27,13,18,21,24-trietheno-11,7-metheno-7H-dibenzo [b,n] [1,5,12,16] tetraazacyclotricosine-5,13-diium dibromide

Introduction

In humans, aging and male sex are associated with an increased risk for cardiovascular diseases (Castelli, 1984; Egashira *et al.*, 1993; Farhat *et al.*, 1996; Orshal and Khalil, 2004; Lloyd-Jones *et al.*, 2010). These pathologies are characterized by an altered balance between endothelium-derived relaxing and contracting factors. Most investigations of this endothelial dysfunction address the reduced endothelium-dependent vasodilatation of large conduit arteries as a result of reduced bioavailability of endothelium-derived NO (EDNO) (Tschudi *et al.*, 1996; Lesniewski *et al.*, 2009). However, in small muscular resistance arteries, marked endothelium-dependent vasodilatation can persist during pharmacological inhibition of NOS or deficiency of NOS3 (Huang *et al.*, 2000; Takaki *et al.*, 2008b). This has been attributed to endothelium-derived hyperpolarizing factor (EDHF) (Feletou and Vanhoutte, 1999; Busse *et al.*, 2002). This mechanism involves small and intermediate conductance calcium-activated potassium channels (K_{Ca}2.3 and K_{Ca}3.1, respectively) in the plasmalemma of the endothelial cells (Grgic *et al.*, 2009; Edwards *et al.*, 2010). Activation of these channels can cause endothelium-dependent vasodilatation by at least two mechanisms: (i) release of an alternative endothelium-derived relaxing factor such as K⁺ ions, H₂O₂ or epoxyeicosatrienoic acid; and (ii) conduction of the endothelial cell hyperpolarization to the underlying smooth muscle cells by heterocellular gap junctions (Feletou and Vanhoutte, 2009; de Wit and Griffith, 2010; Chadha *et al.*, 2011). In contrast to NO, little is known about the effects of risk factors on endothelium-dependent hyperpolarization (EDH)-related arterial responses. Both impairment (Sunano *et al.*, 1999; Wigg *et al.*, 2001; Bussemaker *et al.*, 2003; Haddock *et al.*, 2011) and a compensatory up-regulation (Taddei *et al.*, 2006; Goto *et al.*, 2012) have been reported. Aging alters vascular functions at the level of both endothelium and smooth muscle cell. Age-related changes in EDH have been documented in normal and pathological conditions in rat mesenteric arteries (Fujii *et al.*, 1993; Goto *et al.*, 2012). In the current study, we tested the hypotheses that endothelium-dependent relaxation is impaired by aging in mice as a result of reduced EDH-related responses and that this is more pronounced in male mice as compared with female. For this purpose we used saphenous arteries (small muscular arteries) of young, adult and aging male and female mice, a species that is increasingly used for experimental cardiovascular research. We recorded vasomotor responses in the absence and presence of stimuli of the endothelium, inhibitors of NOS and inhibitors and stimulants of K_{Ca}2.3 and K_{Ca}3.1 channels.

Methods

Solutions and drugs

Krebs Ringer bicarbonate-buffered salt solution (KRB) contained (in mM): 118.5 NaCl, 4.7 KCl, 2.5 CaCl₂, 1.2 MgSO₄, 1.2 KH₂PO₄, 25.0 NaHCO₃ and 5.5 glucose. The KRB solution was continuously aerated with 95% O₂/5% CO₂ and maintained at 37°C. In high K⁺-KRB solution was KRB in which all NaCl was replaced by KCl. Buffers containing intermediate K⁺ concentrations were prepared by mixing appropriate volumes of KRB and K⁺-KRB. Indomethacin (INDO) was purchased from Sigma Aldrich (Zwijndrecht, The Netherlands) and dissolved in ethanol. ACh, noradrenaline (NA), phenylephrine (PHE), N^ω-nitro-L-arginine methyl ester (L-NAME) and sodium nitroprusside (SNP) were purchased from Sigma Aldrich and dissolved in KRB solution. 6, 7-dichloro-1H-indole-2,3-dione 3-oxime (NS309), naphtho[1,2-d]thiazol-2-ylamine (SKA-31) and 1-[(2-chlorophenyl)diphenylmethyl]-1H-pyrazole (TRAM-34) were purchased from Sigma Aldrich and dissolved in dimethyl sulfoxide (DMSO). 6,12,19,20,25,26-hexahydro-5,27:13, 18:21,24-trietheno-11,7-metheno-7H-dibenzo [b,n] [1,5,12,16] tetraazacyclotricosine-5,13-diium dibromide (UCL 1684) was purchased from Tocris Bioscience (Bristol, UK) and dissolved in DMSO.

Animals

Male and female C57BL/6J mice (Charles River, Wilmington, MA, USA) aged 12, 34 and 64 weeks were housed in standard cages (constant room temperature and humidity, 12 h light/12 h dark cycles) and had free access to standard chow diet (pellet) and tap water. All procedures were performed in accordance with the Committee for Animal Care and Use of Maastricht University.

Wire myography

Tissue preparation. Animals were killed by CO₂/O₂ inhalation. Saphenous arteries were dissected free from surrounding fat and connective tissue and directly mounted in a wire myograph (Danish Myo Technology, Aarhus, Denmark). The anatomical location of these vessels is illustrated in the supplement (Supporting Information Fig. S1). Arterial segments (2 mm) were distended to the diameter at which maximal contractile responses to 10 μM NA could be obtained (Hilgers *et al.*, 2010). Optimal diameters (D_{opt}) and maximal contractile responses to NA are summarized in Supporting Information Table S1. The maximal relaxing response to ACh (10 μM) was recorded during contraction induced by 10 μM NA and arterial segments which showed less than 85% relaxation were discarded from the experiments. We performed several pharmacological protocols with the aim of discriminating

between different pathways of endothelium-dependent relaxation. These were not intended to document normal physiological control of arterial function.

Contributions of NO, EDH and COX products to endothelium-dependent relaxation. Initially, a concentration response curve (CRC) for PHE (0.01–10 μ M) was recorded. During the contraction induced by 10 μ M PHE, an ACh CRC (0.01–10 μ M) was generated. After 20 min recovery in drug-free solution, arteries were contracted using K^+ (40 mM), and an ACh CRC (0.01–10 μ M) was recorded during contraction induced by K^+ . These experiments were repeated in the presence of the COX inhibitor INDO (10 μ M) and in the presence of both INDO (10 μ M) and the NOS inhibitor L-NAME (100 μ M).

Contribution of $K_{Ca2.3}$ and $K_{Ca3.1}$ to EDH-related relaxation. Arterial segments were exposed to INDO (10 μ M) and L-NAME (100 μ M). A PHE CRC (0.01–10 μ M) was constructed and the effects of the K_{Ca} channel activator NS309 and of the positive allosteric modulator of K_{Ca} channels SKA 31 (1 μ M) on this CRC were recorded. EDH-related relaxation in response to ACh (0.01–10 μ M) was studied in the presence of INDO (10 μ M) and L-NAME (100 μ M) to rule out interference of NO and prostaglandins with K_{Ca} channels (Bolotina *et al.*, 1994). To study the contribution of K_{Ca} channel subtypes, 1 μ M UCL 1684 (Rosa *et al.*, 1998) was used to block small conductance K_{Ca} channels, while 10 μ M TRAM-34 was used to block intermediate conductance K_{Ca} channels (Hilgers *et al.*, 2010; Hilgers and Webb, 2007).

Sensitivity of vascular smooth muscle to NO. Arteries were contracted with PHE (10 μ M) in the presence of INDO (10 μ M) and L-NAME (100 μ M), and the relaxing effects of the NO donor SNP (0.01–10 μ M) were recorded.

Statistical analysis

All CRCs for contractile stimuli were expressed as a percentage of the maximal response to 10 μ M NA prior to the administration of any pharmacological inhibitor. Relaxing responses were expressed as percentage of the level of pre-contraction. Individual CRCs were fitted to a non-linear sigmoid regression curve (Graph Pad Prism 5.0, Graph Pad Software, La Jolla, CA, USA). Sensitivity (pEC_{50}) and maximal effect (E_{max}) are shown as mean $\pm$ SEM. pEC_{50} and E_{max} were compared by unpaired *t*-test. Two-way analysis of variance followed by a Bonferroni *post hoc* test was used to compare multiple groups. A *P* value <0.05 was considered statistically significant and <0.1 as indicating a trend.

Results

Contractile reactivity

We characterized contractile responses in the saphenous arteries of different age groups of both sexes. The optimal diameters and the maximal contractile responses to 10 μ M NA were comparable in all age groups of both sexes (Supporting Information Table S1). Furthermore, the sensitivity (pEC_{50} ; Supporting Information Table S1) and maximal con-

traction (E_{max}) to PHE (0.01–10 μ M) or K^+ (40 mM) in the absence or presence of NOS and COX inhibitors were comparable in all age groups of both sexes (Supporting Information Table S1).

Relaxing responses to ACh during PHE-induced contraction

In the absence of inhibitors, the sensitivity of PHE-contracted saphenous arteries to ACh (0.01–10 μ M)-induced relaxation and the maximal response were similar in 12-, 34- and 64-week-old male (Figure 1A) and female (Figure 1D) mice. The presence of INDO (10 μ M) had no effect on the sensitivity or the maximal response to ACh in any age group or sex (Figure 1B, E; Table 1), demonstrating that, in our setting, COX products did not contribute to the relaxing responses.

EDNO responses

To evaluate the contribution of EDNO to arterial relaxation, we inhibited EDH-related relaxation by depolarizing the vessels with high potassium buffer [K^+ 40 mM] and COXs by INDO. In saphenous arteries of male mice, the sensitivity to ACh was not changed under these conditions in the three age groups (Figure 2A; Table 1), but the maximal relaxation to ACh was significantly greater in 64-week-old animals ($70 \pm 3\%$) than in 12-week-old mice ($58 \pm 3\%$, $P = 0.022$) and a trend was observed for 34-week-old animals ($56 \pm 6\%$, $P = 0.0956$). In contrast, arteries of female mice, showed similar sensitivity (Figure 2C; Table 1) and maximal relaxation to ACh in all age groups (60 ± 3 , 48 ± 6 and $57 \pm 1\%$ in 12-, 34- and 64-week-old mice, respectively).

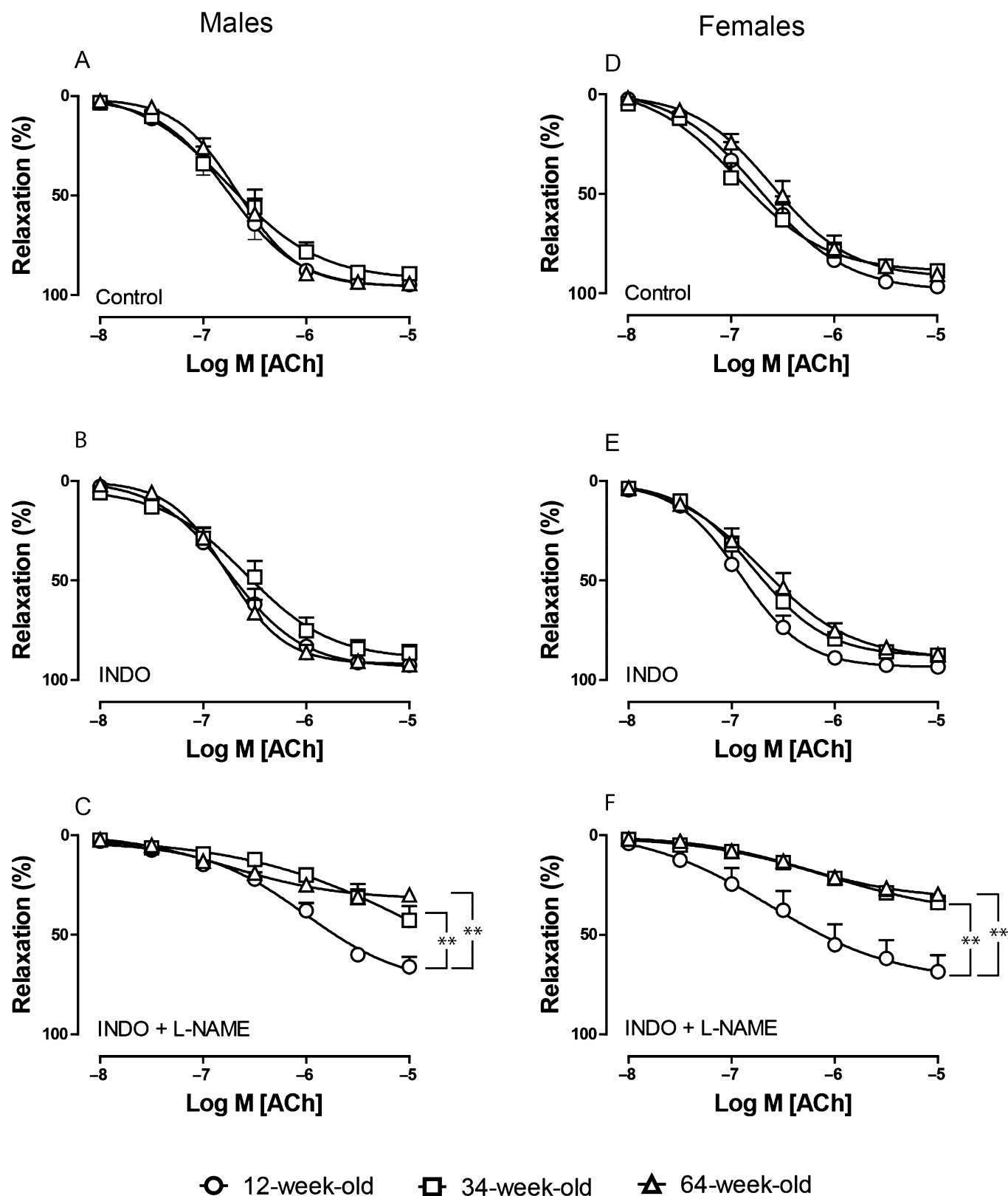
In both male and female mice, K^+ -contracted arteries, when treated with INDO (10 μ M) plus L-NAME (100 μ M), did not relax in response to ACh (Figure 2B, D). We, therefore, conclude that both NO and EDH (as will be described later) contribute to the ACh-induced vasorelaxation in murine saphenous arteries. In addition, we demonstrated that the contribution of NO to maximal relaxation increases in old male but not female mice.

Relaxing responses to SNP

To determine the endothelium-independent response to NO, PHE-contracted arteries were treated with INDO (10 μ M) and L-NAME (100 μ M) to block COXs and NOS, respectively. Subsequently, the relaxing response to the NO donor SNP (0.01–10 μ M) was measured. Sensitivity and maximal relaxation to SNP were similar in all three age groups and in both sexes (Figure 3A, B; Table 1). Relaxing responses to the endothelium-independent NO donor SNP were not affected by aging and sex, indicating that the sensitivity of the vascular smooth muscle cells to NO is unchanged.

ACh-induced EDH-related relaxations

To characterize the contribution of EDH to vascular relaxation, arteries were treated with L-NAME (100 μ M) plus INDO (10 μ M) and were contracted with 10 μ M PHE. In the presence of NOS and COX inhibitors, the sensitivity to ACh was reduced, but did not differ between age groups (Figure 1C, F; Table 1). The maximal effect of ACh, however, did decrease with age under these conditions. In 12-week-old male and female mice, ACh induced a pronounced maximal relaxation

**Figure 1**

Effect of aging on relaxing responses to ACh (0.01–10 μ M) during PHE-induced (10 μ M) contraction in saphenous arteries of male (A–C) and female mice (D–F). Circles, 12-week; squares, 34-week; triangles, 64-week-old mice. (A, D) In the absence of pharmacological inhibitors. (B, E) In the presence of INDO (10 μ M). (C, F) In the presence of both INDO (10 μ M) and L-NAME (100 μ M). Values are shown as means $\pm$ SEM ($n = 7$ –14). ** $P < 0.001$.

Table 1

Effect of aging on endothelium-dependent and -independent relaxations

Age (weeks)	12			34			64		
	pEC ₅₀	E _{max} (%)	n	pEC ₅₀	E _{max} (%)	n	pEC ₅₀	E _{max} (%)	n
Males									
Relaxation responses to ACh									
Control	6.6 ± 0.1	93 ± 1	9	6.7 ± 0.1	90 ± 3	6	6.6 ± 0.1	94 ± 1	8
INDO	6.7 ± 0.1	92 ± 1	9	6.6 ± 0.1	88 ± 3	6	6.7 ± 0.1	92 ± 2	8
INDO + L-NAME	6.1 ± 0.1	67 ± 4	8	NA	43 ± 7*	6	NA	31 ± 2*	7
INDO + L-NAME + SKA-31	6.6 ± 0.1 [†]	76 ± 2	5	NA	32 ± 3*	4	NA	34 ± 3*	5
INDO + L-NAME + TRAM-34	5.8 ± 0.3	6 ± 1 [†]	3	ND	ND		ND	ND	
INDO + L-NAME + UCL 1684	5.1 ± 2.8	9 ± 2 [†]	3	ND	ND		ND	ND	
INDO + L-NAME + TRAM-34 + UCL 1684	5.2 ± 1.2	7 ± 2 [†]	4	NA	12 ± 3	3	NA	12 ± 3	3
Relaxation responses to SNP									
INDO + L-NAME	7.2 ± 0.1	97 ± 1	8	7.2 ± 0.1	99 ± 1	5	7.1 ± 0.1	97 ± 1	7
EDNO responses									
INDO	6.1 ± 0.1	58 ± 3	8	6.1 ± 0.1	56 ± 6	7	6.2 ± 0.1	70 ± 3*	5
INDO + L-NAME	NA	6 ± 2	3	NA	15 ± 1	3	NA	22 ± 2	8
Relaxation responses to NS309									
INDO + L-NAME	5.8 ± 0.1	87 ± 2	6	5.3 ± 0.1*	85 ± 2	3	5.2 ± 0.1*	78 ± 2	5
Females									
Relaxation responses to ACh									
Control	6.8 ± 0.1	98 ± 1	11	6.8 ± 0.1	88 ± 3	10	6.6 ± 0.1	90 ± 5	9
INDO	6.9 ± 0.1	94 ± 1	9	6.8 ± 0.1	88 ± 3	14	6.7 ± 0.1	87 ± 2	8
INDO + L-NAME	5.9 ± 0.1	65 ± 7	12	6.0 ± 0.2	34 ± 4*	12	6.2 ± 0.2	28 ± 3*	11
INDO + L-NAME + SKA-31	6.4 ± 0.1 [†]	66 ± 6	2	ND	24 ± 3	4	ND	32 ± 8	4
INDO + L-NAME + TRAM-34 + UCL 1684	NA	19 ± 6 [†]	2	NA	8 ± 3	4	NA	18 ± 3	4
Relaxation responses to SNP									
INDO + L-NAME	7.6 ± 0.2	98 ± 1	7	7.5 ± 0.1	96 ± 1	8	7.4 ± 0.1	98 ± 1	11
EDNO related responses									
INDO	6.3 ± 0.1	60 ± 3	8	6.2 ± 0.1	48 ± 6	7	6.3 ± 0.2	57 ± 1	5
INDO + L-NAME	NA	16 ± 2	4	NA	18 ± 3	3	NA	19 ± 5	3
Relaxation responses to NS309									
INDO + L-NAME	5.7 ± 0.1	82 ± 1	6	5.3 ± 0.1*	77 ± 4	9	5.4 ± 0.1*	86 ± 4	5

E_{max} expressed as % reduction of the maximal contractile response to 10 μM PHE except for EDNO responses (% reduction of maximal contractile response to 40 mM K⁺). All values are shown as mean ± SEM.

*P < 0.05 compared with arteries of 12-week-old animals of the same sex under the same condition.

[†]P < 0.05 compared with arteries of 12-week-old animals of the same sex treated with INDO plus L-NAME.

ND, not determined; NA, not applicable.

(E_{max} 67 ± 4% and 65 ± 7% in males and females, respectively) that was significantly decreased in 34- and 64-week-old mice (E_{max} 43 ± 7 and 31 ± 2% in males, 34 ± 4 and 28 ± 3% in females, respectively; Figure 1C, F and Table 1). We, therefore, conclude that EDH-related relaxations are attenuated with age in both sexes.

Contribution of K_{Ca}2.3 and K_{Ca}3.1 to EDH-related relaxation

To further characterize EDH-related relaxation and specifically to study the roles of K_{Ca}3.1 and K_{Ca}2.3 channels, we used

TRAM-34 and UCL-1684 to block K_{Ca}3.1 and K_{Ca}2.3 channels, respectively. Saphenous arteries were incubated with a combination of INDO (10 μM), L-NAME (100 μM), TRAM-34 (10 μM) and UCL-1684 (1 μM), and were made to contract with PHE. In all age groups of both sexes, the relaxation response to ACh was completely blocked in the presence of the combination of these inhibitors [Figure 4A–C (males), 4D–F (females); Table 1]. This demonstrated that K_{Ca}3.1 and K_{Ca}2.3 channels account for the observed EDH-related effects. To evaluate the role of the individual channel subtypes in EDH-related relaxation in 12-week-old mice, saphenous arter-

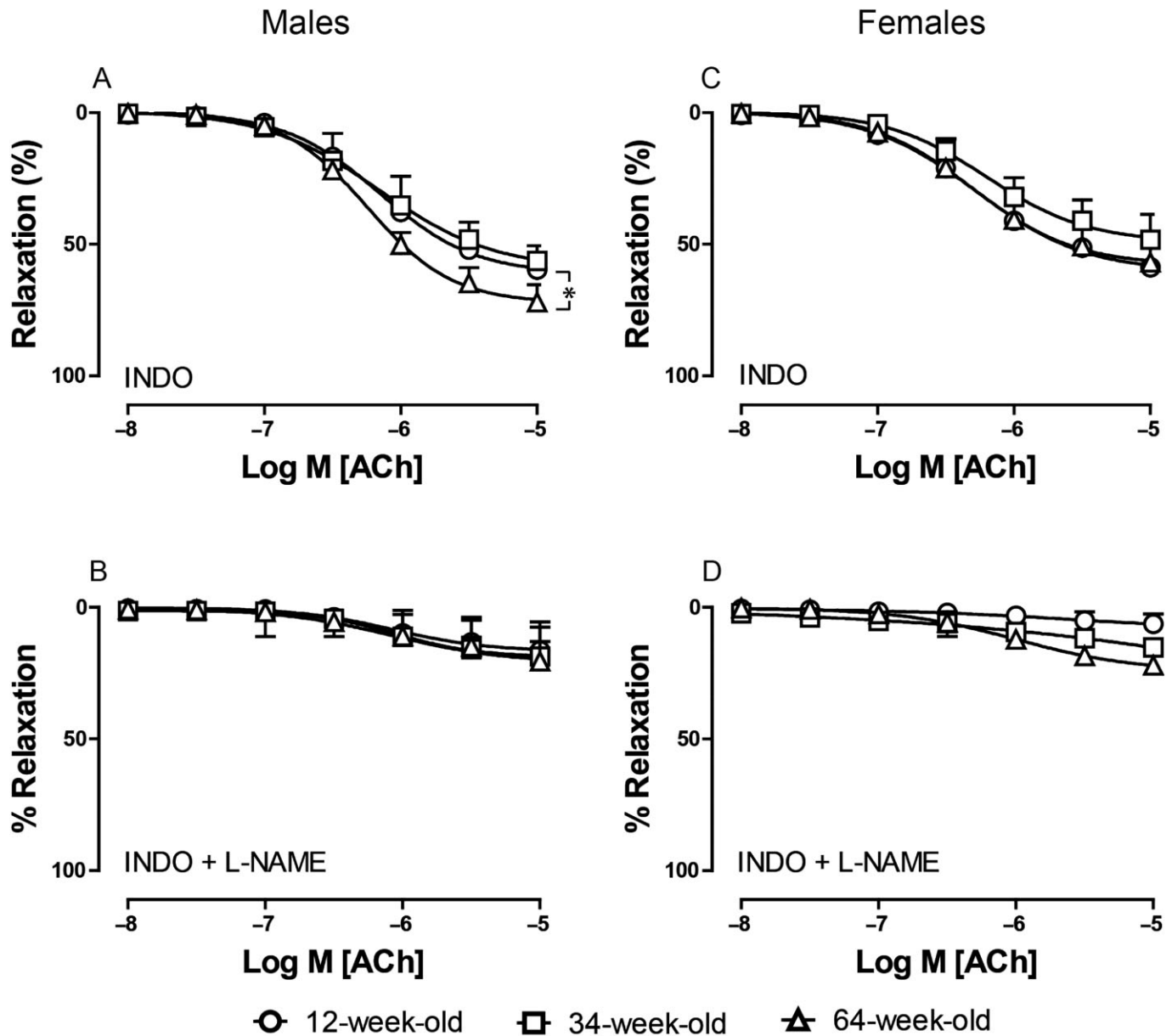


Figure 2

Effect of aging on relaxing responses to ACh (0.01–10 μ M) during K^+ -induced (40 mM) contraction in saphenous arteries of male (A–B) and female (C–D) mice in the presence of INDO (A, C), INDO (10 μ M) and L-NAME (100 μ M) (B, D). Circles, 12-week; squares, 34-week; triangles, 64-week-old mice. Values shown as means $\pm$ SEM ($n = 3$ –8). * = $P < 0.05$.

ies were treated with either TRAM-34 (10 μ M) or UCL-1684 (1 μ M) in combination with INDO and L-NAME. In young male mice, neither in the presence of TRAM-34 nor in the presence of UCL-1684 the arteries showed any residual relaxation (Figure 4A, Table 1), indicating that $K_{Ca3.1}$ and $K_{Ca2.3}$ can each mediate full and complete EDH-related relaxations in saphenous arteries of the mouse. Similar findings were obtained with 1 μ M instead of 10 μ M (Supporting Information Fig. S3).

K_{Ca} channel activation by NS309

Furthermore, we characterized the relaxation response to NS309, a channel activator that does not discriminate

between $K_{Ca3.1}$ and $K_{Ca2.3}$ channels (Strobaek *et al.*, 2004) during PHE-induced contraction in the presence of INDO (10 μ M) plus L-NAME (100 μ M). In 34- and 64-week-old male and female mice, sensitivity to the relaxing effect of NS309 was significantly reduced in males (pEC_{50} 5.3 ± 0.1 , 5.2 ± 0.1) and females (pEC_{50} 5.3 ± 0.1 , 5.4 ± 0.1) compared with 12-week-old animals (pEC_{50} 5.8 ± 0.1 in males and 5.7 ± 0.1 in females), but E_{max} was unchanged (Figure 5A, B; Table 1). This finding demonstrates that the sensitivity to K_{Ca} channel activation decreases with age. The relaxation responses induced by NS309 are largely endothelium dependent at lower concentrations (Supporting Information Fig. S4). Pharmacological inhibitors TRAM-34 and UCL-1684 alone or in

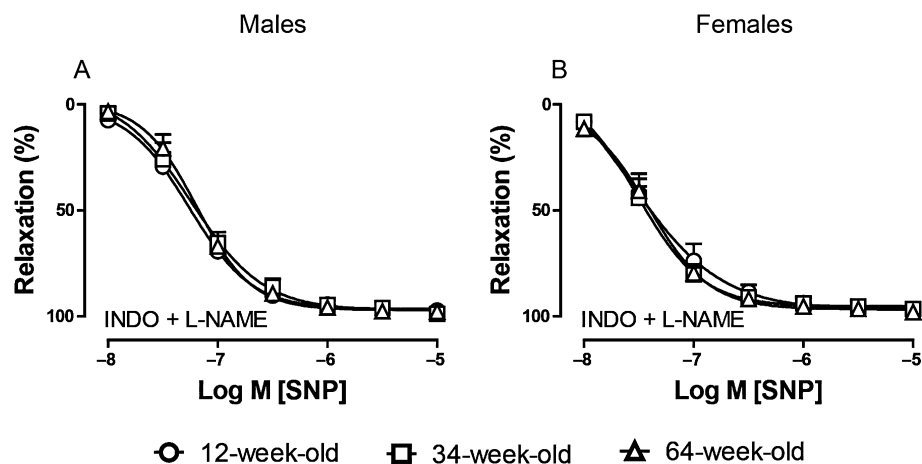


Figure 3

Effect of aging on relaxing responses to SNP (0.01–10 μ M) during PHE-induced (10 μ M) contraction in saphenous arteries of male (A) and female (B) mice in the presence of INDO (10 μ M) and L-NAME (100 μ M). Circles, 12-week; squares, 34-week; triangles, 64-week-old mice. Values are shown as means $\pm$ SEM; ($n = 5$ –11).

combination completely block NS309-induced endothelium-dependent relaxation in young male mice (Supporting Information Fig. S4).

K_{Ca} channel activation by SKA-31

Further we studied the potentiating effect of SKA-31, a newly developed allosteric modulator of *K_{Ca}*2.3 and *K_{Ca}*3.1 (Sankaranarayanan *et al.*, 2009; Hasenau *et al.*, 2011). The direct relaxing effect of SKA-31 was minor, and only observed at concentrations exceeding 3 μ M (Supporting Information Fig. S2). The potentiating effects of this compound were measured in PHE-contracted arteries in the presence of INDO (10 μ M) plus L-NAME (100 μ M). In 12-week-old mice of both sexes, SKA-31 potentiated ACh-induced relaxation at a concentration of 1 μ M [Figure 6A (males), 6D (females), Table 1], but this agent did not potentiate ACh-induced relaxations in 34- and 64-week-old males (Figure 6B, C) and females (Figure 6E, F). This underscores our hypothesis that *K_{Ca}*3.1 and *K_{Ca}*2.3 channel functions decrease with age in both sexes.

Discussion and conclusion

In the present study, the relative contribution of NO, EDH and prostaglandins to endothelium-dependent relaxation was studied in young, adult and aged mice of both sexes. We initially studied contractile responses induced by either PHE or *K⁺*. These contractile responses were not affected by age or sex. Interestingly, the contractile responses were also not affected by the presence of INDO (to block contributions of COX products) or the presence of both INDO and L-NAME.

Thereafter, we evaluated the responses mediated by EDNO in the presence of INDO in arteries that were depolarized with *K⁺*. In addition to activating arterial smooth muscle and peri-arterial nerves by depolarization, high *K⁺* inhibits EDH. Responses mediated by EDNO were not affected by age in arteries isolated from female mice. Surprisingly, in arteries

isolated from old male mice, EDNO-mediated responses were increased. This was not due to changes in the sensitivity of vascular smooth muscle for NO as these, in line with earlier findings in other labs (DeSouza *et al.*, 2000; Taddei *et al.*, 2001), were similar in all age groups. Our findings regarding EDNO are intriguing, since it is generally accepted that NO bioavailability decreases with age (Tschudi *et al.*, 1996; Barton *et al.*, 1997; Taddei *et al.*, 2001). However, these studies on EDNO were performed on larger conduit arteries. In contrast, our study focused on the contribution of different vasoactive factors to endothelial function in saphenous arteries, which are muscular resistance arteries. The discrepancy between the previous studies and our findings may thus lie in the well-known and very profound regional vascular and interspecies heterogeneity.

The contribution of prostaglandins and EDH to endothelium-dependent relaxation was studied in PHE-contracted arteries. Total endothelium-dependent relaxation did not differ between age groups or sexes. Endothelium-dependent contractions, prominent in other types of murine arteries (e.g. mouse aorta; Zhou *et al.*, 2005) were not observed and ACh-induced relaxations were not affected by the presence of INDO. Thus, in contrast to other murine arteries (aorta, carotid and femoral arteries) (Zhou *et al.*, 2005; Liu *et al.*, 2012), COX products do not contribute to endothelium-dependent relaxation in murine saphenous arteries.

In line with other studies in rats (Fujii *et al.*, 1993; Goto *et al.*, 2000; 2012), EDH-related relaxations decreased with age in murine saphenous arteries. Because the maximal ACh-induced relaxations did not change with age, we assume that the residual EDH-related responses together with the response mediated by EDNO are sufficient for full relaxation even in arteries isolated from older mice. It has been demonstrated that endothelial *K_{Ca}* channels are critically involved in the EDH-related phenomenon in various arteries *in vitro* and *in vivo* (Brahler *et al.*, 2009; Grgic *et al.*, 2009; Kohler and Ruth, 2010). Suppression of *K_{Ca}*2.3 expression by doxycycline

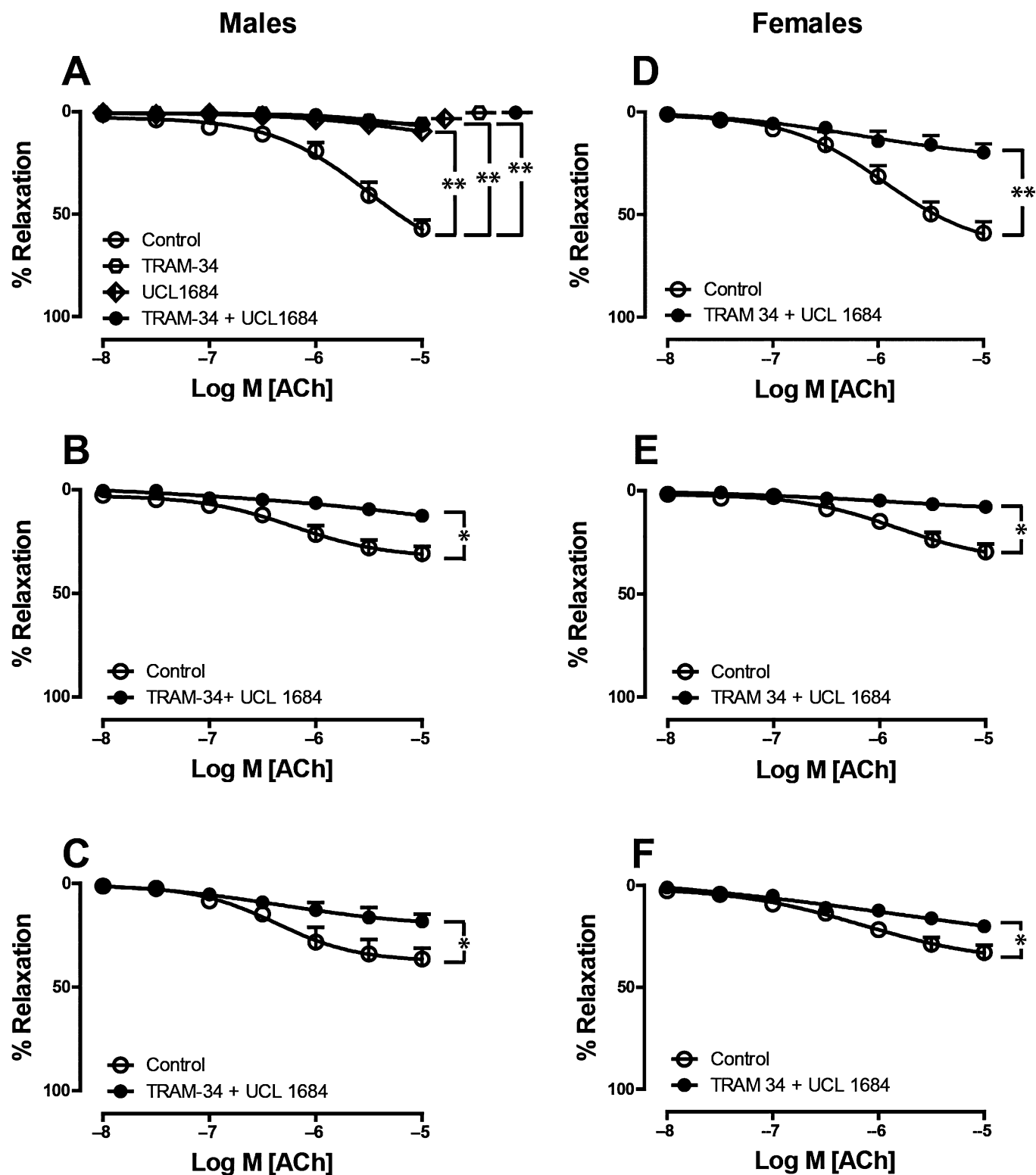


Figure 4

Effects of $K_{Ca}3.1$, $K_{Ca}2.3$ channel inhibitors on EDH-related relaxations during PHE-induced ($10 \mu\text{M}$) contraction in saphenous arteries of male (A–C) and female (D–F) mice aged 12 (A, D), 34 (B, E) and 64 (C, F) weeks. All experiments were performed in the presence of INDO ($10 \mu\text{M}$) and L-NAME ($100 \mu\text{M}$). In addition, arteries were treated with either TRAM-34 ($10 \mu\text{M}$) or UCL 1684 ($1 \mu\text{M}$) (12-week-old male mice only, A), or the combination of both (all age groups). Values are shown as means $\pm$ SEM ($n = 3$ –4). $^{**}P < 0.001$. $^{*}P < 0.05$.

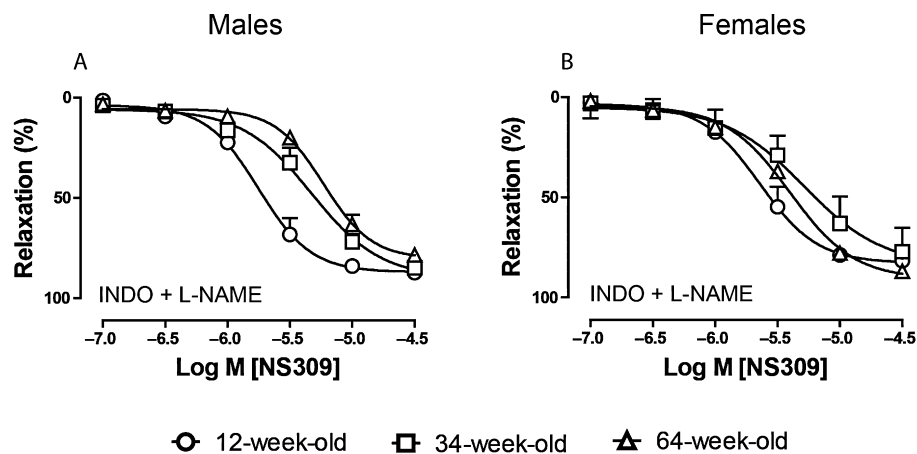


Figure 5

Effect of aging on relaxing responses to the $K_{Ca2.3}$ and $K_{Ca3.1}$ channel opener NS309 during PHE (10 μ M)-induced contraction in saphenous arteries of male (A) and female (B) mice. All experiments were performed in the presence of INDO (10 μ M) and L-NAME (100 μ M). Circles, 12-week; squares, 34-week; triangles, 64-week-old mice. Values are shown as means $\pm$ SEM ($n = 3-9$).

administration in SK3^{T/T} mice resulted in a pronounced elevation of blood pressure (~30 mmHg) (Taylor *et al.*, 2003). Deletion of $K_{Ca3.1}$ ($K_{Ca3.1}^{-/-}$ mice) decreased endothelial and smooth muscle hyperpolarizations in response to ACh and significantly increased arterial blood pressure (~20 mmHg) (Si *et al.*, 2006). Mice that were both $K_{Ca3.1}$ -deficient and $K_{Ca2.3}$ (SK3^{T/T})-depleted showed impaired EDH-dependent relaxation as well as increased arterial blood pressure (Brahler *et al.*, 2009). To establish whether $K_{Ca3.1}$ and $K_{Ca2.3}$ are also involved in the EDH-related responses of murine saphenous arteries, we evaluated their contribution to EDH-related relaxations in arteries of young and old female and male mice. Blocking both channels simultaneously fully inhibited EDH-related responses in both sexes and all age groups. Similar results have been reported for murine mesenteric arteries in the presence of TRAM-34 and apamin (Harrington *et al.*, 2007). Surprisingly, blocking only one of the two channel types, $K_{Ca2.3}$ or $K_{Ca3.1}$, had a similar effect as the blockade of both channels simultaneously. This is unlikely due to a lack of specificity of the inhibitors for either channel type (Rosa *et al.*, 1998; Wulff *et al.*, 2000). Interaction between $K_{Ca2.3}$ and $K_{Ca3.1}$ may, therefore, be crucial for EDH-related relaxation in murine saphenous arteries. In the future, co-immunoprecipitation experiments or *in situ* proximity ligation assays could establish the nature of the interaction between these channels. These experiments, however, were beyond the scope of our studies described here.

We relied on a classical pharmacological approach to study differences in channel expression and channel properties. In theory, decreased channel expression should result in a decreased E_{max} , but unchanged pEC_{50} when relaxations are induced by channel openers. In contrast, one would expect similar E_{max} , but altered pEC_{50} when the channel open probability is changed. Accordingly, we analysed relaxations induced by NS309 (an opener of K_{Ca} channels) to characterize receptor properties. We found that the E_{max} of NS309-induced relaxation was similar in all age groups and both sexes. However, arteries isolated from older mice were significantly

less sensitive for NS309. It thus seems that aging alters the function of the K_{Ca} channels. To further address whether aging affects the characteristics of K_{Ca} channels in the endothelial cells of saphenous arteries, we used SKA-31, a recently developed positive allosteric modulator of $K_{Ca3.1}$ and $K_{Ca2.3}$ channels (Sankaranarayanan *et al.*, 2009; Hasenau *et al.*, 2011). In saphenous arteries of young mice, SKA-31 potentiated EDH-related relaxations, but not in those of 34- and 64-week-old mice. This, together with our findings with NS309, suggests that the functional properties of K_{Ca} channels are critically affected by age. Alternatively, the density of heterocellular gap junctions which have been shown to conduct endothelial hyperpolarization resulting from K_{Ca} channel activation to the underlying smooth muscle may decrease with age as recently documented (Sandow *et al.*, 2004). Also, the mechanism by which smooth muscle cell hyperpolarization ultimately leads to relaxation and the role of intracellular calcium stores and plasmalemmal calcium channels merits further investigation. In addition to this, previous studies showed that NOS-derived H_2O_2 can be involved in EDH-related relaxations of both large and small arteries of mice and rats (Fujiki *et al.*, 2005; Drouin *et al.*, 2007; Takaki *et al.*, 2008a). This has to be further addressed in the future studies.

We have attempted to quantify $K_{Ca3.1}$ and $K_{Ca2.3}$ expression using semi-quantitative immunohistochemistry. However, none of the presently available antibodies meets the criteria of specificity in paraffin sections of murine tissue (for details see online Supporting Information Figs. S5–S9). This suggests that at least some published data using such antibodies should be treated with scepticism. It should, however, be pointed out that dedicated immunofluorescence analyses of whole mount preparations (Senadheera *et al.*, 2012) would have been more suited.

The current study addressed age and sex differences in the vasomotor responses of murine saphenous arteries. We were able to demonstrate clear effects of aging on EDH-related relaxations. However, we did not observe major differences

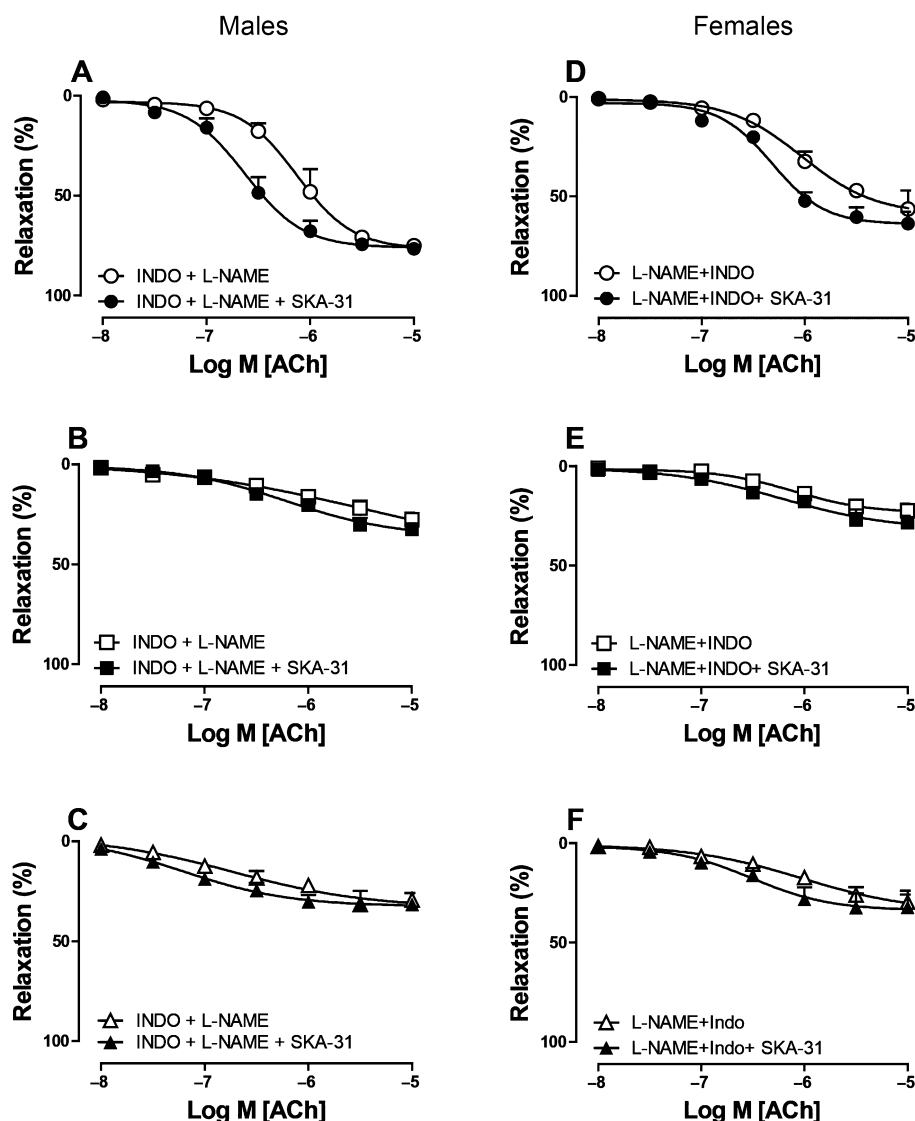


Figure 6

Potentiating effect of SKA-31 on relaxing responses in saphenous arteries of male (A–C) and female (D–F) mice of 12 (A, D), 34 (B, E) and 64 (C, F) weeks of age. Arteries were treated with either INDO (10 μ M) and L-NAME (100 μ M), or the combination of L-NAME (100 μ M), INDO (10 μ M) and SKA-31 (1 μ M). Relaxing responses to ACh were recorded (0.01–10 μ M) during PHE-induced (10 μ M) contraction. Values are shown as means $\pm$ SEM ($n = 4$ –5).

between the sexes, although a trend for increased sensitivity to exogenous NO was seen in female arteries and a modest increase in EDNO was observed in old male arteries. These mild sex differences regarding vasomotor function are surprising in view of the increased risk for cardiovascular disease associated with male gender. In addition, others have shown such gender differences previously, for example, the relative contribution of NO and EDH to endothelium-dependent relaxation differs in mesenteric arteries of male and female rats (McCulloch and Randall, 1998). In addition, studies on resistance arteries of endothelial NOS/COX-1 double-knockout mice (eNOS^{-/-}/COX-1^{-/-}) suggested that EDH is the main endothelium-dependent relaxing mechanism in female mice, whereas NO and PGI₂ are the predominant mediators in male mice (Scotland *et al.*, 2005). It is not easy to pinpoint

the factors responsible for the discrepancy between our findings and earlier studies but clearly regional vascular heterogeneity and differences between species regarding the contribution of endothelium-derived relaxing factors to endothelium-dependent relaxation play a role (Hecker, 2000; Campbell and Gauthier, 2002).

Clinical implications

In vivo animal studies showed that treatment with SKA-31 lowers blood pressure (Sankaranarayanan *et al.*, 2009; Damkjaer *et al.*, 2012). In addition, K_{Ca}3.1- and K_{Ca}2.3-deficient mice are hypertensive (Taylor *et al.*, 2003; Si *et al.*, 2006). Thus, sensitizing K_{Ca} channels could be considered for the treatment of hypertension. However, in our current *ex vivo* study, arteries isolated from young mice were more sen-

sitive to NS309 than those of older animals. In addition, SKA-31 was unable to potentiate EDH-related relaxations of saphenous arteries isolated from 34- or 64-week-old mice of both sexes. This implies that therapeutic interventions aimed at lowering blood pressure via K_{Ca} may be less efficacious in old than in young patients.

Summary

The present study shows that endothelium-dependent relaxations of murine saphenous arteries are maintained in aging mice of both sexes. However, the relative contribution of EDH to these relaxations diminishes with increasing age. The present study also shows that in murine saphenous arteries, both $K_{Ca3.1}$ and $K_{Ca2.3}$ are of crucial importance for EDH-related relaxation. Older mice display reduced sensitivity for K_{Ca} channel activators, possibly explaining the decreased contribution of EDH to endothelium-dependent relaxation in these mice. Only minor sex differences regarding endothelium-mediated relaxations were observed.

Acknowledgements

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Conflicts of interest

The authors declare that there are no conflicts of interest.

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Supporting information

Additional Supporting Information may be found in the online version of this article at the publisher's web-site:

Figure S1 The blue line shows the anatomical location of saphenous artery used in the present study. The black line corresponds to the femoral artery.

Figure S2 Direct relaxing responses to SKA-31 (0.001–10 μ M) during PHE (10 μ M) – induced contraction in saphenous arteries of 12-week-old male mice ($n = 4$).

Figure S3 Relaxing responses to ACh (0.001–10 μ M) in PHE (10 μ M)-contracted saphenous arteries in the presence of L-NAME (100 μ M), INDO (10 μ M) (+L + I; diamonds), and the selective K_{Ca} 3.1 channel blocker TRAM-34 at either 10 μ M (+L + I + T; squares) or 1 μ M (+L + I + T; squares) concentration. Values are expressed as mean $\pm$ SEM ($n = 4$ –6).

Figure S4 Relaxing responses to the K_{Ca} 2.3 and K_{Ca} 3.1 channel opener NS309 in phenylephrine (10 μ M)-contracted saphenous arteries with (E) or without (-E; circles) endothelium. In the intact arteries, experiments were performed in the presence of LNAME (100 μ M), INDO (10 μ M), (+E + L + I; diamonds), the selective K_{Ca} 3.1 channel blocker TRAM-34 (+E + L + I + T; 10 μ M; squares), the selective K_{Ca} 2.3 channel blocker UCL-1684 (+E + L + I + U; 1 μ M; inverted triangles) and the combined incubation of TRAM-34 and UCL-1684 (+E + L + I + T + U; triangle). Values are expressed as mean $\pm$ SEM ($n = 5$ –6).

Figure S5 Expression of K_{Ca} 3.1 channels in saphenous arteries of 12- (A, E), 34- (B, F) and 64-week-old (C, G) male mice. All tissues were fixed in 4% formaldehyde and embedded in paraffin. Four millimetre sections were rehydrated and exposed for 20 min to 0.3% H_2O_2 at RT to block endogenous peroxidases. Subsequently, sections were incubated in a humidified chamber (overnight, 4°C) with successively sheep antibodies against IK1 [anti- K_{Ca} 3.1, 1:600 in normal goat serum (NGS); Alomone Labs (Jerusalem, Israel); product number: APC-064; Lot number: AN-02] and horseradish peroxidase-coupled rabbit antibodies against sheep IgG (1:400 in NGS, DAKO, Glostrup, Denmark). The localization of HRP was visualized with 3, 3-diaminobenzidine (Sigma Aldrich). All sections were counterstained with hematoxylin. Negative controls (A, B, C, D) were incubated with secondary antibody only. Mouse brain (12-week-old male) was used as positive control in absence (D) of and in the presence (H) of primary antibody. Corresponding magnified insets are marked with asterisk. Staining of endothelium (arrow) was observed in 12-week-old, but not in 34- and 64-week-old mice. However, we also observed staining in smooth muscle cells, which may be due to incomplete specificity of the antibodies. We also tried lower concentrations of the primary antibody, which only decreased the staining in the endothelium relative to the surrounding structures. Nevertheless, these observations are in line with reduced K_{Ca} 3.1 protein expression in endothelium with aging.

Figure S6 Expression of K_{Ca} 2.3 channels in saphenous arteries of 12- (A, E), 34- (B, F) and 64-week-old (C, G) male mice. Tissues were fixed and sections processed as described in the legend of Supporting Information Fig. S5, except that sections were incubated with sheep antibodies directed against SK3 (anti- K_{Ca} 2.3 N-terminal, 1:200 in NGS; Alomone Labs; Product number: APC-025; Lot number: AN-04) and horseradish peroxidase-coupled rabbit antibodies against sheep IgG (1:400 in NGS, DAKO, Glostrup, Denmark). The localization of HRP was visualized with 3, 3-diaminobenzidine (Sigma Aldrich). All sections were counterstained with hema-

toxylin. Negative controls (A, B, C, D) were incubated with secondary antibody only. Mouse brain (12-week-old male) was used as positive control in absence (D) of and in the presence (H) of primary antibody. Corresponding magnified insets are marked with asterisk. No staining was observed, indicating that the SK3 antiserum did not recognize K_{Ca} 2.3 channels in the sections. Usage of whole mount arterial preparation merits further investigation.

Figure S7 Expression of K_{Ca} 3.1 channels in saphenous arteries of 12- (A, E), 34- (B, F) and 64-week-old (C, G) male mice. All tissues were fixed in 4% formaldehyde and embedded in paraffin. Four millimetre sections were rehydrated and exposed for 20 min to 0.3% H_2O_2 at RT to block endogenous peroxidases. Subsequently, sections were incubated in a humidified chamber (overnight, 4°C) with successively sheep antibodies against IK1 (anti- K_{Ca} 3.1, 1:600 in normal goat serum (NGS); SIGMA; Product number: P 4997) and horseradish peroxidase-coupled rabbit antibodies against sheep IgG (1:400 in NGS, DAKO, Glostrup, Denmark). The localization of HRP was visualized with 3, 3-diaminobenzidine (Sigma Aldrich). All sections were counterstained with hematoxylin. Negative controls (A, B, C, D) were incubated with secondary antibody only. Mouse brain (12-week-old male) was used as positive control in absence (D) of and in the presence (H) of primary antibody. Corresponding magnified insets are marked with asterisk. Staining of endothelium (arrow) was observed in 12-week-old, but not in 34- and 64-week-old mice. However, we also observed staining in smooth muscle cells, which may be due to incomplete specificity of the antibodies. We also tried lower concentrations of the primary antibody, which only decreased the staining in the endothelium relative to the surrounding structures. Nevertheless, these observations are in line with reduced K_{Ca} 3.1 protein expression in endothelium with aging.

Figure S8 Expression of K_{Ca} 2.3 channels in saphenous arteries of 12- (A, E), 34- (B, F) and 64-week-old (C, G) male mice. Tissues were fixed and sections processed as described in the legend of Supporting Information Fig. S7, except that sections were incubated with sheep antibodies directed against SK3 (anti- K_{Ca} 2.3 N-terminal, 1:200 in NGS; SIGMA; Product number: P 0608) and horseradish peroxidase-coupled rabbit antibodies against sheep IgG (1:400 in NGS, DAKO, Glostrup, Denmark). The localization of HRP was visualized with 3, 3-diaminobenzidine (Sigma Aldrich). All sections were counterstained with hematoxylin. Negative controls (A, B, C, D) were incubated with secondary antibody only. Mouse brain (12-week-old male) was used as positive control in absence (D) of and in the presence (H) of primary antibody. Corresponding magnified insets are marked with asterisk. No staining was observed, indicating that the SK3 antiserum did not recognize K_{Ca} 2.3 channels in the sections.

Figure S9 Expression of K_{Ca} 3.1 (A, B, C) and K_{Ca} 2.3 (D) channels in gut of WT (A, B, D) and knockout K_{Ca} 3.1 (C) mice. All tissues were fixed in 4% formaldehyde and embedded in paraffin. Four millimetre sections were rehydrated and exposed for 20 min to 0.3% H_2O_2 at RT to block endogenous peroxidases. Subsequently, sections were incubated in a humidified chamber (overnight, 4°C) with sheep antibodies against IK1 (anti- K_{Ca} 3.1 1:600 in normal goat serum (NGS); SIGMA; Product number: P 4997) or SK3 (anti- K_{Ca} 2.3 1:200 in normal goat serum (NGS); SIGMA; Product number:

P 0608) and horseradish peroxidase-coupled rabbit antibodies against sheep IgG (1:400 in NGS, DAKO, Glostrup, Denmark). The localization of HRP was visualized with 3,3'-diaminobenzidine (Sigma Aldrich). All sections were counterstained with hematoxylin. Negative controls (A) were incubated with secondary antibody only. With SK3 antiserum no staining was observed in the gut (D). However, IK1 antiserum resulted in intense staining in the gut of WT (B)

compared with $K_{Ca}2.3$ knockout (C) mice. But $K_{Ca}3.1$ KO mice also showed some amount of staining in the gut (C, arrow), which may be due to incomplete specificity of the antibodies. Unfortunately due to the limited availability of tissues from $K_{Ca}2.3$ KO mice further investigation was not feasible in our lab.

Table S1 Effect of aging on optimal diameter and contractile responses.