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Title: Nitrooleic acid, an Endogenous Product of Nitrativ Stress, Activates Nociceptive Sensory Nerves via the Direct Activation of TRPA1.

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Running Title: OA-NO₂ activates TRPA1

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Abbreviations: OA, oleic acid; 9-OA-NO₂, 9-nitrooleic acid; ROS, reactive oxygen species; RNS, reactive nitrogen species; 4HNE, 4-hydroxynonenal; 4ONE, 4-oxononenal; PG, prostaglandin; SIN-1, 3-morpholino-sydnnonimine; carboxyPTIO, 2-(4-Carboxyphenyl)-4,4,5,5-tetramethylimidazole-1-oxyl-3-oxide; 15dPGJ₂; 15-deoxy- $\Delta^{12,14}$ -prostaglandin J₂; AITC, allyl isothiocyanate; DTT, dithiothrietol; Caps, capsaicin; HC-030031, 1,2,3,6-tetrahydro-1,3-dimethyl-N-[4-(1-methylethyl)phenyl]-2,6-; AP-18, 4-(4-chlorophenyl)-3-methylbut-3-en-2-oxime; hTRPA1-HEK, HEK cells stably transfected with human TRPA1 channels; hTRPV1-HEK, HEK cells stably transfected with human TRPV1 channels; nt-HEK, non-transfected HEK cells.

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Abstract

Transient Receptor Potential A1 (TRPA1) is a non-selective cation channel, preferentially expressed on a subset of nociceptive sensory neurons, that is activated by a variety of reactive irritants via the covalent modification of cysteine residues. Excessive nitric oxide during inflammation (nitrative stress), leads to the nitration of phospholipids resulting in the formation of highly reactive cysteine modifying agents, such as nitrooleic acid (9-OA-NO₂). Using calcium imaging and electrophysiology, we have shown that 9-OA-NO₂ activates human TRPA1 channels (EC₅₀, 1μM), whereas oleic acid had no effect on TRPA1. 9-OA-NO₂ failed to activate TRPA1 in which the cysteines at position C619, C639, C663 and the lysine at 708 had been mutated. TRPA1 activation by 9-OA-NO₂ was not inhibited by the NO scavenger carboxyPTIO. 9-OA-NO₂ had no effect on another nociceptive-specific ion channel, TRPV1. 9-OA-NO₂ activated a subset of mouse vagal and trigeminal sensory neurons, which also responded to the TRPA1 agonist allyl isothiocyanate and the TRPV1 agonist capsaicin. 9-OA-NO₂ failed to activate neurons derived from TRPA1^{-/-} mice. The action of 9-OA-NO₂ at nociceptive nerve terminals was investigated using an ex vivo extracellular recording preparation of individual bronchopulmonary C-fibers in the mouse. 9-OA-NO₂ evoked robust action potential discharge from capsaicin-sensitive fibers with slow conduction velocities (0.4 – 0.7 m/s), which was inhibited by the TRPA1 antagonist AP-18. These data demonstrate that nitrooleic acid, a product of nitrative stress, can induce substantial nociceptive nerve activation through the selective and direct activation of TRPA1 channels.

Introduction

Oxidative stress and nitrative stress have been implicated as contributing to acute and chronic inflammation (Radi, 2004; Szabo et al., 2007; Valko et al., 2007). Nitric oxide (NO) is an endogenous mediator with multiple cellular functions that is produced by many cell types including vascular endothelium, neutrophils, fibroblasts and nerves (Bian and Murad, 2003). NO, generated from L-arginine by NO synthases (NOS), reacts with the reactive oxygen species (ROS) superoxide - which is formed through multiple pathways in inflammation including NADPH oxidase, xanthine oxidase and perverted mitochondrial function - to produce the reactive nitrogen species (RNS) peroxynitrite (ONOO⁻) and nitrogen dioxide (*NO₂). RNS are potent inflammatory molecules that can react with lipids, proteins and DNA (Szabo et al., 2007). Within membranes, where the hydrophobic environment maximizes RNS production (Moller et al., 2007), RNS react with unsaturated fatty acids (e.g. oleic acid) causing the addition of a NO₂ group (nitration) (Freeman et al., 2008; Jain et al., 2008; Trostchansky and Rubbo, 2008). Nitrated fatty acids (e.g. nitrooleic acid) are highly reactive electrophilic compounds that can modulate a variety of cellular targets, including thiol residues and PPAR γ (Freeman et al., 2008; Trostchansky and Rubbo, 2008).

Nitrated fatty acids are detectable *in vitro* after exposure of fatty acids to RNS-donors (Jain et al., 2008; O'Donnell et al., 1999). Nitrated fatty acids have been measured in human plasma and red blood cells, with total nitrooleic acid (OA-NO₂) and total nitrolinoleic acid concentrations in plasma of 920nM and 630nM, respectively (Baker et al., 2005). Interestingly, oxidative stress-produced reactive unsaturated aldehydes such as 4-hydroxynonenal (4HNE) that tend to break off from the phospholipid during peroxidation (Gardner, 1989), 32% of OA-NO₂ in plasma and 72% in packed red blood cells is esterified (bound within phospholipid) (Baker et al., 2005). The intriguing possibility exists that esterified nitrated fatty acids represent a sink of bioactive

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mediators produced during nitrative stress that can induce subsequent cellular functions following liberation from the membrane by Phospholipase A₂ (Jain et al., 2008).

Inflammation elicits pain and reflexes due to the activation of somatosensory and visceral nociceptive sensory nerves. Recently a member of the Transient Receptor Potential (TRP) ion channel family termed TRPA1 has been demonstrated preferentially on nociceptive sensory nerves and is activated by irritants such as allyl isothiocyanate (AITC), cinnamaldehyde, bradykinin and phytocannabinoids (Bandell et al., 2004; Bautista et al., 2006; De Petrocellis et al., 2008; Jordt et al., 2004). ROS and reactive lipid peroxidation products have been shown to activate nociceptive neurons via TRPA1 (Andersson et al., 2008; Bautista et al., 2006; Macpherson et al., 2007b; Taylor-Clark et al., 2008b; Taylor-Clark et al., 2008c; Trevisani et al., 2007), likely via the direct covalent modification of key N-terminus cysteine groups (Hinman et al., 2006; Macpherson et al., 2007a; Trevisani et al., 2007). Given that the nitro group (NO₂) is one of the strongest electron-withdrawing groups known; and given that the rate constants for cysteine adduction by nitrated fatty acids exceed those of lipid peroxidation products (Baker et al., 2007); we predict that the reactive products of nitrative stress represent a group of highly potent endogenous TRPA1 activators, such that their production during inflammation may significantly enhance nociceptors activation.

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Methods

All experiments were approved by the Johns Hopkins Animal Care and Use Committee or conducted according to the requirements of the United Kingdom Animals (Scientific Procedures) Act (1986) and strictly conformed to the ethical standards of GlaxoSmithKline Pharmaceuticals as appropriate.

HEK293 cell culture: Wild-type HEK293 cells, cells stably expressing human TRPA1 (hTRPA1-HEK) or human TRPV1 (hTRPV1-HEK) were used in this study, as previously described (Taylor-Clark et al., 2008c). Cells were maintained in an incubator (37°C, 5% CO₂) in DMEM (containing 110 mg/L pyruvate) supplemented with 10% FBS and 500 mg/mL Geneticin as a selection agent. Cells were removed from their culture flasks by treatment with Accutase (Sigma), then plated onto poly-D-lysine-coated cover slips and incubated at 37°C for >1 h before experimentation.

hTRPA1-3C/K-Q mutant: cDNA (1.6µg/ml) for hTRPA1 and hTRPA1-3C/K-Q (C619, C639 and C663 mutated to serines; K708 mutated to glutamine) (Hinman et al., 2006) was expressed in HEK293 cells using GenJet In Vitro DNA Transfection Reagent (SignaGen Laboratories, Gaithersburg, MD).

TRPA1^{-/-}/TRPV1^{-/-} mice: TRPA1^{-/-} mice (Taylor-Clark et al., 2008b) were successfully bred with TRPV1^{-/-} mice (Davis et al., 2000), yielding normal Mendelian numbers of offspring. Those mice that were homozygous deficient in both TRPA1 and TRPV1 were used in this study.

Dissociation of mouse sensory ganglia: Mouse vagal and trigeminal ganglia were isolated and enzymatically dissociated from wild-type C57BL/6J mice, TRPA1^{-/-} mice and TRPA1^{-/-}

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/TRPV1-/- mice using previously described methods (Taylor-Clark et al., 2008b; Taylor-Clark et al., 2008c). Isolated neurons were plated onto poly-D-lysine-coated and laminin-coated coverslips and used within 24 hours.

Calcium imaging: HEK293-covered coverslips were loaded with Fura 2 acetyoxymethyl ester (Fura-2 AM; 8 μ M) (Molecular Probes, Carlsbad, CA, USA) in DMEM (containing 110 mg/L pyruvate) supplemented with 10% FBS and incubated (40 min, 37°C, 5% CO₂). Neuron-covered coverslips were loaded with Fura-2 AM (8 μ M) in L-15 media containing 20% FBS and incubated (40 min, 37°C). For imaging, the coverslip was placed in a custom-built chamber (bath volume of 600 mL) and superfused at 4 mL/min with Locke solution (34°C; composition (mM): 136 NaCl, 5.6 KCl, 1.2 MgCl₂, 2.2 CaCl₂, 1.2 NaH₂PO₄, 14.3 NaHCO₃ and 10 dextrose (gassed with 95% O₂-5% CO₂, pH 7.3-7.4)) for 15 minutes before and throughout each experiment by an infusion pump. Changes in intracellular free calcium concentration (intracellular [Ca²⁺]_{free}) were measured by digital microscopy (Universal; Carl Zeiss, Inc, Thornwood, NY, USA) equipped with in-house equipment for ratiometric recording of single cells. The field of cells was monitored by sequential dual excitation, 352 and 380 nm, and the analysis of the image ratios used methods previously described to calculate changes in intracellular [Ca²⁺]_{free} (Taylor-Clark et al., 2008c). The ratio images were acquired every 6 seconds. Superfused buffer was stopped 30 seconds before each drug application, when 300 μ L buffer was removed from the bath and replaced by 300 μ L of 2X test agent solution added between image acquisitions. Following treatments, neurons were exposed to KCl (30 seconds, 75 mM) to confirm voltage sensitivity. At the end of experiments, both neurons and HEK cells were exposed to ionomycin (30 seconds, 1 μ M) to obtain a maximal response.

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For the analysis of Fura-2 AM loaded cells, the measurement software converted ratiometric information to intracellular $[Ca^{2+}]_{free}$ using Tsien parameters ($[Ca] = K_d ((R-R_{min})/(R_{max}-R)) (\beta)$) particular to this instrumentation and the HEK cells and dissociated mouse vagal neurones. Preliminary calibration studies yielded an R_{min} (352/380 ratio under calcium-free conditions) of 0.3 for both HEK cells and mouse sensory neurons and an R_{max} (352/380 ratio under calcium-saturating conditions) of 18 and 14 for HEK cells and neurons, respectively. β (380 in calcium-free conditions / 380 in calcium-saturating conditions) was estimated as being 10 and the K_d was estimated as being 224nM. In the following experimental studies we did not specifically calibrate the relationship between ratiometric data and absolute calcium concentration for each specific cell, choosing instead to use the parameters provided from the calibration studies and relate all measurements to the peak ionomycin response in each viable cell. This effectively provided the needed cell-to-cell calibration for enumerating individual cellular responses. Only cells that had a robust response to ionomycin were included in analyses. At each time point for each cell, data was presented as the percentage change in intracellular $[Ca^{2+}]_{free}$, normalized to ionomycin: $response_x = 100 \times ([Ca^{2+}]_x - [Ca^{2+}]_{bl}) / ([Ca^{2+}]_{max} - [Ca^{2+}]_{bl})$, where $[Ca^{2+}]_x$ was the apparent $[Ca^{2+}]_{free}$ of the cell at a given time point, $[Ca^{2+}]_{bl}$ was the cell's mean baseline apparent $[Ca^{2+}]_{free}$ measured over 120s, and $[Ca^{2+}]_{max}$ was the cell's peak apparent $[Ca^{2+}]_{free}$ during ionomycin treatment. For the neuronal experiments, neurons were defined as 'responders' to a given compound if the mean response was greater than the mean baseline plus 2 X the standard deviation. Only neurons that responded to KCl were included in analyses. Given that vagal and trigeminal ganglia likely comprise of heterogeneous neuronal populations it is important to emphasize the point that results are presented in two distinct ways. Firstly, the number of neurons responding (based on the criteria described above) to a given stimulus compared to the total number of neurons is reported. Secondly, the mean percentage change in intracellular $[Ca^{2+}]_{free}$ normalized to ionomycin of *those neurons that (based on the above criteria) were defined as 'responders'* is reported.

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Whole cell voltage clamp: Conventional whole cell patch-clamp recordings were performed at room temperature (21-24°C) using a Multiclamp 700B amplifier and pCLAMP 9 software (Molecular Devices, Sunnyvale, CA). Pipettes (3-4 M Ω) fabricated from borosilicate glass (Sutter Instruments, Novato) were filled with an internal solution composed of (mM): 140 CsCl, 4 MgCl₂, 10 HEPES, and 5 EGTA; pH was adjusted to 7.2 with CsOH. Cover slips were superfused continuously during recording with an external solution composed of (in mM): 140 NaCl, 2 MgCl₂, 5 CsCl, 10 HEPES, and 10 D-Glucose (pH adjusted to 7.4 with NaOH) and gassed with 95% O₂ – 5% CO₂. Only cells with <10 M Ω series resistances were used and compensated up to 80%. Currents were sampled at 500 Hz, and recordings were filtered at 10 kHz. The membrane potential was held at -60 mV through out the recording. Drugs were applied to the cell and record inward current. Data were analyzed using clampfit software and transferred to Excel spreadsheets or GraphPad Prism 4 for further analysis.

C-fiber extracellular recordings: Mice were killed by CO₂ asphyxiation followed by exsanguination. The innervated isolated trachea/bronchus preparation was prepared as previously described (Nassenstein et al., 2008). Briefly, the airways and lungs with their intact extrinsic innervation (vagus nerve including vagal ganglia) were taken and placed in a dissecting dish containing Krebs bicarbonate buffer solution composed of (mM): 118 NaCl, 5.4 KCl, 1.0 NaH₂PO₄, 1.2 MgSO₄, 1.9 CaCl₂, 25.0 NaHCO₃ and 11.1 dextrose, and equilibrated with 95% O₂ and 5% CO₂ (pH 7.2–7.4) (also containing indomethacin (3 μ M)). Connective tissue was trimmed away leaving the trachea and lungs with their intact nerves. The airways were then pinned to the larger compartment of a custom-built two-compartment recording chamber which was lined with silicone elastomer (Sylgard). A vagal ganglion was gently pulled into the adjacent compartment of the chamber through a small hole and pinned. Both compartments were separately superfused with Krebs bicarbonate buffer (37°C). A sharp glass electrode was pulled by a Flaming Brown

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micropipette puller (P-87; Sutter Instruments, Novato, CA, USA) and filled with 3 M NaCl solution. The electrode was gently inserted into the vagal ganglion so as to be placed near the cell bodies. The recorded action potentials were amplified (Microelectrode AC amplifier 1800; A-M Systems, Everett, WA, USA), filtered (0.3 kHz of low cut-off and 1 kHz of high cut-off), and monitored on an oscilloscope (TDS340; Tektronix, Beaverton, OR, USA) and a chart recorder (TA240; Gould, Valley View, OH, USA). The scaled output from the amplifier was captured and analyzed by a Macintosh computer using NerveOfIt software (Phocis, Baltimore, MD, USA). To measure conduction velocity, an electrical stimulation (S44; Grass Instruments, Quincy, MA, USA) was applied to the center of the receptive field. NerveOfIt software analysis was also able to discriminate individual nerve fiber responses on the rare occasion that more than one bronchopulmonary afferent was recorded from during stimulation (electrical, mechanical or chemical) of the lung tissue. The conduction velocity of the individual bronchopulmonary afferents was calculated by dividing the distance along the nerve pathway by the time delay between the shock artifact and the action potential evoked by electrical stimulation. Drugs were intratracheally applied as a 1ml bolus over 10s.

In the extracellular recording studies, the action potential discharge was quantified off-line and recorded in 1s bins. A response was considered positive if the number of action potentials in any 1s bin was >2 times the average background response. The background activity was usually either absent or less than 2 Hz. The peak frequency evoked by a stimulus was quantified as the maximum number of action potentials that occurred within any 1s bin.

Chemicals: Stock solutions (200X+) of all agonists were dissolved in 100% ethanol (final concentration of 0.5% ethanol or less). 9-OA-NO₂ and oleic acid were purchased from Cayman Chemicals (Ann Arbor, MI). Fura 2AM was purchased from Molecular Probes (Carlsbad, CA). AP-18 was purchased from Biomol (Plymouth Meeting, PA). HC030031 was purchased from

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Tocris Bioscience (Ellisville, MO). All other chemicals were purchased from Sigma-Aldrich (St. Louis, MO). Due to the reported instability of 9-OA-NO₂ in aqueous solutions (half-life_{aq} of approximately 2 hours (Gorczyński et al., 2007)), 9-OA-NO₂ was dissolved into the appropriate buffer within 5 minutes of experimental use.

Results

9-OA-NO₂ activates TRPA1 expressed on HEK293 cells

We have previously shown that HEK293 cells stably transfected with human TRPA1 (hTRPA1-HEK) responded, in calcium imaging assays, to the reactive electrophilic products of oxidative stress, such as 4HNE and 8-iso Prostaglandin A₂ (Taylor-Clark et al., 2008b; Taylor-Clark et al., 2008c). Using the same hTRPA1-HEK cells we found that 9-OA-NO₂ (30nM-30μM, for structure see Fig 1A) activated TRPA1 channels (maximum response of 47±1.5% of ionomycin, n=156) with an approximate EC₅₀ of 1μM (Figs 1B and 1C). The potency of 9-OA-NO₂ at TRPA1 channels is 10 fold greater than the canonical selective TRPA1 agonist allyl isothiocyanate (AITC), which activated the hTRPA1-HEK cells (maximum response of 50±1.3% of ionomycin, n=269) with an approximate EC₅₀ of 10μM (Fig 1C). 9-OA-NO₂ (100nM-100μM) failed to activate ntHEK cells (maximum response of 3.2±0.13% of ionomycin, n=369, Fig 1B), suggesting that the increase in cytosolic calcium in the HEK cells caused by 9-OA-NO₂ required TRPA1 channels. We investigated the effect of oleic acid on TRPA1 channels. Oleic acid (100nM-100μM) failed to activate hTRPA1-HEK cells (maximum response of 0.73±0.12% of ionomycin, n=168, Fig 1C), suggesting that the addition of the NO₂ group onto the fatty acid was crucial to TRPA1 channel activation.

We have previously demonstrated that the highly reactive electrophile 4-oxononanal (4ONE), which activates hTRPA1-HEK cells with an approximate EC₅₀ of 2 μM, is also an agonist for TRPV1 channel, although only at 100μM (Taylor-Clark et al., 2008b). We investigated whether

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or not 9-OA-NO₂ was also capable of activating TRPV1 channels. Using calcium imaging of hTRPV1-HEK cells we found that 9-OA-NO₂ (100nM-100μM) failed to activate TRPV1 channels (n=168, Fig 1C). As expected, the hTRPV1-HEK cells responded robustly to the canonical TRPV1 agonist capsaicin (300nM, maximum response of 34±1.1% of ionomycin, n=168, data not shown).

We next investigated the mechanism by which 9-OA-NO₂ activates TRPA1 channels. It has been reported that nitrated fatty acids may mediate some of their biological actions via the release of nitric oxide (NO) (Schopfer et al., 2005). It is conceivable that NO then directly nitrosylates the channel causing activation, as has been shown for other TRP channels (Yoshida et al., 2006). Indeed, NO donors have been shown to have a weak agonist effect on TRPA1 channels (Sawada et al., 2008; Takahashi et al., 2008). We confirmed the weak agonist effect of two different NO donors, sodium nitroprusside and SIN-1, on TRPA1 activity: sodium nitroprusside (300μM) and SIN-1 (300μM) caused an increase in calcium in hTRPA1-HEK cells (n=75) of 8.7±0.56% of ionomycin and 5.0±0.31% of ionomycin, respectively (data not shown). We then investigated the contribution of NO to 9-OA-NO₂-induced TRPA1 activation using the NO scavenger carboxy-PTIO. Incubating hTRPA1-HEK cells with 1mM carboxy-PTIO for 10 min had no effect on the activation of TRPA1 by 9-OA-NO₂ (3μM): maximum response of 49±2.3% of ionomycin (n=80) and 46±1.7% of ionomycin (n=212) for vehicle and carboxy-PTIO treatments, respectively (Fig 2A). Overall the data suggests that NO is unlikely to play a major role in the activation of TRPA1 channels by 9-OA-NO₂.

Given that 9-OA-NO₂ activates TRPA1 channels in a NO-independent manner and that oleic acid has no effect on the channel, it is likely that the electrophilic C=C-NO₂ moiety is responsible for the TRPA1 channel activation. This would be consistent with previous reports that direct covalent

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modification of TRPA1 channel cysteines induces activation (Hinman et al., 2006; Macpherson et al., 2007a). 9-OA-NO₂ is a reactive electrophilic molecule that has been shown to readily adduct amino acid residues such as cysteines (Michael reaction) (Baker et al., 2007). Indeed when the TRPA1 activation potency of 9-OA-NO₂ is compared with other endogenous TRPA1 'covalently-modifying' agonists investigated in our heterologous system (Fig 2B, and see (Taylor-Clark et al., 2008b; Taylor-Clark et al., 2008c)), the rank order of -LogEC₅₀ (9-OA-NO₂ (6) ≥ 4ONE (5.8) > 4HNE (5) > 8-iso PGA₂ (4.5)) is almost identical to the rank order of second-order rate constants for the reaction of glutathione's cysteine residue in model systems (9-OA-NO₂ (183M⁻¹s⁻¹) > 4ONE (145M⁻¹s⁻¹) > 4HNE (1.3M⁻¹s⁻¹) > 8-iso PGA₂ (0.7M⁻¹s⁻¹)) (Baker et al., 2007). This suggests that modification of cysteines may play a role in 9-OA-NO₂'s activation of TRPA1. For human TRPA1 channels, covalent modification (Michael reaction)-induced TRPA1 activation is dependent on the presence of three crucial cysteine residues (C619, C639 and C663) and one lysine residue (K708) on the channel's intracellular N-terminus (Hinman et al., 2006). We hypothesized that 9-OA-NO₂ would fail to activate TRPA1 channels with mutations at these four residues. Using plasmid cDNA encoding the mutant hTRPA1-3C/K-Q and wild-type hTRPA1 channels (see methods), we found that, as expected, 9-OA-NO₂ (3μM) activated wild-type hTRPA1 channels (maximum response of 50±2.1% of ionomycin, n=166) but failed to activate the mutant channel (maximum response of 3.0±1.0% of ionomycin, n=18), whereas menthol (300μM), which activates TRPA1 channels independently of C619, C639 and C663 (Xiao et al., 2008), did activate the mutant (maximum response of 33±7.0% of ionomycin, Fig 2C). Neither 9-OA-NO₂ nor menthol activated ntHEK cells (maximum response of 0.45±0.12% of ionomycin and 1.7±0.16% of ionomycin (n=88), respectively). The data clearly suggest that 9-OA-NO₂ activates TRPA1 via covalent modification.

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Interestingly, unlike products of oxidative stress such as 4ONE and 4HNE, 9-OA-NO₂ has been shown to form Michael adducts with cysteine residues in a manner that can be reversed by the thiol-containing reducing agents dithiothriitol (DTT) and glutathione (GSH) (Batthyany et al., 2006). Thiol-reducing agents are present *in vivo* in both extracellular and intracellular compartments and their levels are sensitive to the redox state of the environment (Szabo et al., 2007; Valko et al., 2007). DTT (1-5mM) has previously been shown to reverse the activation of TRPA1 channels by H₂O₂ but not by 15dPGJ₂ or 4HNE (Andersson et al., 2008; Takahashi et al., 2008). In calcium imaging assays of hTRPA1-HEK cells, we found that 7 min treatment with 10mM DTT failed to reverse the activation of TRPA1 by 9-OA-NO₂ (3μM): maximum response of 51±1.1% of ionomycin (n=204) and 49±1.2% of ionomycin (n=259) for vehicle and DTT treatments, respectively (Fig 2D). We found a similar lack of reversibility when using 250μM glutathione (GSH) ethyl ester (membrane permeable form of GSH) (data not shown).

9-OA-NO₂ activates TRPA1-expressing nociceptive sensory neurons

We and others have previously identified TRPA1 channel responses in subpopulations of native somatosensory and visceral sensory neurons from trigeminal, vagal and dorsal root ganglia (Bandell et al., 2004; Jordt et al., 2004; Nassenstein et al., 2008). These TRPA1-expressing neurons almost always also respond to capsaicin, the TRPV1 agonist. Here we used calcium imaging to address the hypothesis that 9-OA-NO₂ would activate sensory neurons that also respond to AITC, the canonical TRPA1 channel agonist, and to capsaicin. 9-OA-NO₂ (10μM) activated approximately 60% of trigeminal dissociated sensory neurons and 40% of vagal dissociated neurons (Fig 3A). AITC (100μM) and capsaicin (1μM) activated similar proportions of the dissociated neurons, although there appeared to be a greater percentage of capsaicin-sensitive neurons in the vagal ganglia compared to the trigeminal ganglia. When only those

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neurons that responded to 9-OA-NO₂ were combined, the mean responses demonstrated that 9-OA-NO₂ (10μM) caused a robust activation of sensory neurons (maximum response of 35±3.4% of ionomycin and 36±3.3% of ionomycin for vagal and trigeminal neurons, respectively) that also responded strongly to AITC and capsaicin (Fig 3B and 3C), suggesting that 9-OA-NO₂'s actions were restricted to nociceptive neurons that were activated by TRPA1 and TRPV1 agonists. Next, we assessed the potency of 9-OA-NO₂ elicited responses in vagal sensory neurons. As with the hTRPA1-HEK cells, 0.1μM 9-OA-NO₂ evoked only minor responses, which increased dramatically at 1μM and at 10μM began to approach a maximum, suggesting a similar order of magnitude between 9-OA-NO₂'s responses at heterologously expressed TRPA1 channels and native sensory neurons (Fig 3D).

TRPA1^{-/-} neurons are insensitive to 9-OA-NO₂

To confirm the molecular identity of the 9-OA-NO₂-activated channels in native nociceptors, we compared responses of wild-type vagal neurons with vagal neurons derived from TRPA1^{-/-} mice. In calcium imaging assays, 80 out of 195 wild-type vagal neurons responded to 9-OA-NO₂ (10μM) with a maximum response of 35±3.4% of ionomycin. However, only 34 out of 124 TRPA1^{-/-} neurons responded to 9-OA-NO₂, with a dramatically reduced maximum response of 6.8±0.6% of ionomycin, indicating that TRPA1 channels were responsible for the great majority of the 9-OA-NO₂ response (Fig 4A). As expected, the TRPA1^{-/-} neurons also failed to respond to AITC (100μM) but responded robustly to capsaicin (1μM). Although 9-OA-NO₂ had no observable effect on our hTRPV1-HEK cells, it was possible that the minor residual 9-OA-NO₂-induced response in TRPA1^{-/-} neurons was due to electrophile-dependent TRPV1 activation (Salazar et al., 2008; Taylor-Clark et al., 2008b). We addressed this hypothesis using mice with genetic deletion of both TRPA1 and TRPV1 channels. Consistent with our hTRPV1-HEK cell

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data, the neuronal responses of TRPA1^{-/-}/TRPV1^{-/-} mice were indistinguishable from TRPA1^{-/-} mice, with 9-OA-NO₂ (10μM) activating 20/101 neurons with a maximum response of 5.3±0.9% of ionomycin (Fig 4A). As expected TRPA1^{-/-}/TRPV1^{-/-} neurons also failed to respond to both AITC and capsaicin.

We further investigated the responses of native neurons to 9-OA-NO₂ in whole cell voltage clamp of vagal neurons. In the nominal absence of Ca²⁺, 9-OA-NO₂ (10μM) induced an inward current in 7 out of 10 wild type neurons (mean current density 12.5±8.3 pA/pF), which was reversed by HC030031 (20μM)(Fig 4B and 4D), the selective TRPA1 antagonist with an IC₅₀ of approximately 1μM (McNamara et al., 2007). However, 9-OA-NO₂ (10μM) had virtually no effect on TRPA1^{-/-} neurons, with only 1 of 9 responding (current density 0.19 pA/pF) (Fig 4C and 4D). As expected, Capsaicin (1μM) responses were no different between neurons from wild-type and TRPA1^{-/-} mice: 4 out of 8 responded with mean current density of 170±52 pA/pF and 7 out of 9 responded with mean current density of 158±59 pA/pF, respectively. Taking the calcium imaging and voltage clamp data together, we conclude that the activation of native neurons by 9-OA-NO₂ is overwhelmingly dependent on TRPA1 channels.

9-OA-NO₂ induces action potential discharge from visceral C-fibers via TRPA1

The effect of 9-OA-NO₂ on nociceptive nerve endings was analyzed using extracellular recording techniques in an *ex vivo* vagal innervated mouse lung preparation (Kollarik et al., 2003).

Nociceptive vagal C-fibers were considered those nerves fibers that responded with action potential discharge to capsaicin and α,β methylene ATP. We have previously shown that TRPA1 agonists only activate this bronchopulmonary nerve population (Nassenstein et al., 2008; Taylor-Clark et al., 2008b). In 7 experiments the C-fiber under study (conduction velocities ranged from

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0.4 – 0.7 m/s) responded strongly with action potential discharge to capsaicin (0.3 μ M) and α , β methylene ATP (10 μ M). In 7 of 7 of these capsaicin sensitive nerve fibers, 9-OA-NO₂ (30 μ M) evoked action potential discharge (Fig 5A and 5B). The action potential discharge in response to a 1 ml infusion of 9-OA-NO₂ delivered over a 20 second period had an onset within the 20 sec delivery period, and generally persisted for only about 2-3 min. The total number of action potentials averaged 227 \pm 51 (Fig 5A). The peak frequency of discharge induced by 9-OA-NO₂ averaged 12 \pm 2Hz, which was about 50% of that observed with capsaicin (24 \pm 4Hz, added at the end of the experiment). The response to 9-OA-NO₂ was reproducible within a given nerve fiber. Treating the tissue a second time 20 minutes after the cessation of action potential discharge, resulted in a response not significantly different than the first response (Fig 5A). AP-18 (30 μ M), the selective TRPA1 antagonist with an IC₅₀ of approximately 3 μ M (Petrus et al., 2007), nearly abolished the 9-OA-NO₂-induced action potential discharge in the lung C-fibers in 5 out of 5 fibers tested (Fig 5A and 5B). We have previously noted that there is a subpopulation of α , β methylene ATP-sensitive fibers in the mouse lung that are insensitive to capsaicin (Kollarik et al., 2003). We evaluated two of these capsaicin-insensitive fibers, and both were found to be also insensitive 9-OA-NO₂.

Discussion

Our findings demonstrate that the nitrated fatty acid nitrooleic acid (9-OA-NO₂) is a stimulator of somatosensory and visceral nociceptors via the selective and direct activation of TRPA1 channels. Based on our concentration-response analysis in both neurons and hTRPA1-HEK cells we can conclude that this compound is the most potent endogenous TRPA1 agonist thus far described.

9-OA-NO₂ activated hTRPA1 channels in a heterologous system at concentrations just above those levels found in plasma samples from healthy humans (Baker et al., 2005) and well within the range of concentrations that OA-NO₂ induces other (non-neuronal) biological effects (Freeman et al., 2008; Trostchansky and Rubbo, 2008). The parent compound of 9-OA-NO₂, oleic acid, had no effect on hTRPA1, suggesting that the addition of the highly electrophilic nitro group was responsible for 9-OA-NO₂'s actions on TRPA1, rather than the hydrocarbon chain or the carboxylic acid group. Nitrated fatty acids have been shown to be stable in lipophilic environments but in aqueous solutions they have been shown to release NO (Freeman et al., 2008). However, compared to other nitrated fatty acids OA-NO₂ is relatively stable (half-life_{aq} of approximately 2 hours) (Gorczynski et al., 2007) and so it was not surprising that the NO scavenger carboxyPTIO had no effect on 9-OA-NO₂ TRPA1 activation. The capacity of NO itself to activate TRPA1 is not yet clear. Although we confirmed, using SIN-1 and sodium nitroprusside, the mild activation of TRPA1 channels by the NO donors demonstrated by other groups (Sawada et al., 2008; Takahashi et al., 2008), these studies are confounded by reports that NO donors, including sodium nitroprusside and SNAP which are typically thought to have no oxidative effect, also produce superoxide (Pieper et al., 1994). Superoxide (and its downstream ROS H₂O₂) can independently induce TRPA1 activation (Sawada et al., 2008). Further studies are needed to confirm a direct interaction of NO with the TRPA1 channel. Nevertheless, based on the

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potency and efficacy of 9-OA-NO₂'s activation of TRPA1 it is likely that, regardless of any direct (minor) effect of NO on the channel, the primary mechanism of nitrative stress-induced nociceptors activation is via the nitration of fatty acids.

Using a mutated TRPA1 channel that has mutations at C619, C639, C663 and K708, which renders it insensitive to electrophilic activation (Hinman et al., 2006), we were able to show that 9-OA-NO₂ activated the channel through interactions with these nucleophilic residues. This was unsurprising given the evidence that the activation was NO-independent and the activation potency ratio of 9-OA-NO₂ and other endogenous electrophiles was similar to their described Michael addition to cysteines (Baker et al., 2007).

Using the reducing agent DTT (and GSH) we found that the activation of hTRPA1 channels by 9-OA-NO₂ was not reversible by these agents. DTT has previously been shown to reverse TRPA1 activation by H₂O₂ but not by the reactive unsaturated aldehyde 4HNE (Andersson et al., 2008; Sawada et al., 2008), which activates TRPA1 via Michael addition of C619, C639, C663 and K708 (Trevisani et al., 2007). Our finding of a non-reversible Michael addition by 9-OA-NO₂ was unexpected, as OA-NO₂ and nitrolinoleic acid have both been shown to form reversible Michael adducts with cysteine residues on GSH and other peptides/proteins (Batthyany et al., 2006), although OA-NO₂ does irreversibly react with xanthine oxidoreductase (Kelley et al., 2008). It would be prudent not to overextend any conclusions based on these *in vitro* heterologous system studies, but there remains the possibility that for OA-NO₂ there is a mismatch between its addition of cysteines on TRPA1 (pro-nociception) and its addition of cysteines on antioxidants (anti-nociception), which would augment the efficacy of neuronal activation *in vivo*.

TRPA1 channels are preferentially expressed on small diameter nociceptive sensory neurons in trigeminal, vagal and dorsal root ganglia (Bandell et al., 2004; Jordt et al., 2004; Nassenstein et

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al., 2008). In our experiments we found that 9-OA-NO₂ activated a population of small diameter neurons that also responded to AITC, another TRPA1 agonist, and capsaicin, the TRPV1 agonist. Similar results were seen for both trigeminal and vagal neurons, suggesting that somatosensory and visceral nociceptors are activated by the nitrated fatty acid. Using TRPA1^{-/-} vagal neurons we confirmed in calcium imaging and voltage clamp studies the molecular identity of the target of 9-OA-NO₂ in sensory neurons as being TRPA1, which is consistent with TRPA1 being the sole target of acrolein and 4HNE (products of lipid peroxidation) (Bautista et al., 2006; Macpherson et al., 2007b; Trevisani et al., 2007). Previous studies had shown that the lipid peroxidation product with the greatest electrophilicity, 4ONE (Baker et al., 2007; Doorn and Petersen, 2002), was not only able to activate TRPA1 channels but also gated TRPV1 channels (Taylor-Clark et al., 2008b). Given that 9-OA-NO₂ is more electrophilic than 4ONE, as determined by cysteine adduction (Baker et al., 2007), we would have predicted that 9-OA-NO₂ would have activated hTRPV1-HEK cells. In addition, another oleic acid derivative, N-oleylethanolamine, has been shown to gate TRPV1 channels (Movahed et al., 2005). However, 9-OA-NO₂ failed to activate TRPV1 channels and there was no difference between the response to 9-OA-NO₂ in vagal neurons from TRPA1^{-/-} mice and TRPA1^{-/-}/TRPV1^{-/-} mice. This lack of effect suggests that TRPV1 activation by reactive molecules is not solely dependent on the degree of electrophilicity.

The importance of 9-OA-NO₂'s actions on TRPA1 channels was confirmed at the level of the sensory nerve terminals in *ex vivo* extracellular bronchopulmonary C-fiber recordings. We have previously shown that mouse vagal afferent capsaicin-sensitive C-fibers innervating the lungs can be activated by TRPA1 agonists and that these responses are abolished by TRPA1 antagonists and by genetic deletion of TRPA1 channels (Nassenstein et al., 2008; Taylor-Clark et al., 2008b). As predicted from the *in vitro* studies, 9-OA-NO₂ induced robust action potential discharge from bronchopulmonary capsaicin-sensitive C-fiber terminals in a manner that was inhibited by the

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selective TRPA1 antagonist AP-18. This result is consistent with previous reports that lipid peroxidation products and AITC and other isothiocyanates evoke pain, local reflexes and central reflexes through the activation of TRPA1 channels on nociceptive sensory nerves (Andersson et al., 2008; Bessac et al., 2008; Taylor-Clark et al., 2008a; Taylor-Clark et al., 2008b; Trevisani et al., 2007).

Many of NO's pathophysiological and cytotoxic effects are thought to be mediated by the NO-derived RNS (Radi, 2004), which have been shown to contribute *in vivo* to inflammatory models (Salvemini et al., 2006) and to multiple disease states including asthma, COPD, viral-induced pneumonia, cystic fibrosis, ischemic-reperfusion injury, circulatory shock, arthritis, colitis and pain (Radi, 2004; Ricciardolo et al., 2006; Szabo et al., 2007). In addition, there are exogenous stimuli that could contribute to the formation of RNS, for example nitrogen oxides (NO_x) in polluted air and cigarette smoke. The contribution of nitrated fatty acids, including OA-NO₂, to the activation of nociceptive sensory nerves in these conditions is as yet unknown. High nanomolar levels of nitrated fatty acids have been detected in plasma and cell membranes (Baker et al., 2005), and evidence suggests that these levels increase during inflammation (Balazy and Poff, 2004). In addition, much of the detected nitrated fatty acid is esterified (Baker et al., 2005), which suggests that subsequent activation of Phospholipase A₂ may release more nitrated fatty acids long after conditions of nitrative stress have diminished (Jain et al., 2008). Finally, it is likely that nociceptive nerves are themselves capable of producing RNS (and presumably nitrated fatty acids) as they express proteins capable of synthesizing superoxide (NADPH oxidase (Dvorakova et al., 1999)) and NO (nNOS (Mazzone and McGovern, 2008)). Taken together, our findings that OA-NO₂ is a potent endogenous activator of TRPA1 suggests a novel relevant mechanism by which excessive NO production and nitrative stress can contribute to nociception in inflammation.

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Footnotes

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Figure legends

Figure 1: Activation of hTRPA1-HEK cells by the nitrated fatty acid 9-OA-NO₂.

A, Structural formulae of oleic acid and 9-OA-NO₂. B, Mean \pm S.E.M. Ca²⁺ responses of hTRPA1-HEK cells to putative endogenous TRPA1 agonist 9-OA-NO₂ (0.1 to 10 μ M). All drugs were applied for 60s (blocked line). Black line denotes responses of hTRPA1-HEK cells (n=156), broken line denotes responses of nt-HEK cells (n=369). C, Dose-response relationships of Ca²⁺ responses of hTRPA1-HEK cells for 9-OA-NO₂, AITC and oleic acid, and responses of hTRPV1-HEK cells for 9-OA-NO₂ (0.03 to 300 μ M) (data comprise >156 cells). Data represent the maximum response during the 60s agonist treatment taken from mean cell response versus time curves (note that the S.E.M. is contained within symbol).

Figure 2: 9-OA-NO₂ activates TRPA1 via covalent modification.

A, Mean \pm S.E.M. Ca²⁺ responses of hTRPA1-HEK cells to 9-OA-NO₂ (3 μ M, 60s), with (grey line, n=212) and without (black line, n=80) pretreatment with NO scavenger carboxyPTIO (1mM). B, Dose-response relationships of Ca²⁺ responses of hTRPA1-HEK cells for 9-OA-NO₂, 4HNE, 4ONE and 8-iso PGA₂ (data comprise >156 cells)(some data taken from (Taylor-Clark et al., 2008b; Taylor-Clark et al., 2008c)). Data represent the maximum response during the 60s agonist treatment taken from mean cell response versus time curves (note that the S.E.M. is contained within symbol). C, Mean \pm S.E.M. Ca²⁺ responses to 9-OA-NO₂ (3 μ M, 60s) and menthol (300 μ M, 180s) of ntHEK cells (broken line, n=88), HEK cells transiently transfected with hTRPA1 (black line, n=166) and HEK cells transiently transfected with hTRPA1-3C/K-Q

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(grey line, n=18). D, Effect of DTT (10mM, grey line, n=259) and vehicle (black line, n=204) on the mean \pm S.E.M. Ca^{2+} responses of hTRPA1-HEK cells to 9-OA- NO_2 (3 μM , 60s).

Figure 3: 9-OA- NO_2 activates a subset of trigeminal and vagal neurons.

A, Representative Fura 2AM ratiometric image of Ca^{2+} responses of trigeminal (top) and vagal (bottom) neurons to 9-OA- NO_2 (10 μM), AITC (100 μM), capsaicin (Caps, 1 μM) and KCl (75mM). Percentage of KCl-sensitive neurons responding to the agonists is illustrated below each image. B and C, Mean \pm S.E.M. Ca^{2+} responses of trigeminal (B) and vagal (C) neurons responding to 9-OA- NO_2 (10 μM). Response to AITC (100 μM) and capsaicin (Caps, 1 μM) also shown. Blocked line denotes the 30s application of agonist. D, Dose-response relationship of Ca^{2+} responses of vagal neurons to 9-OA- NO_2 (0.1 to 10 μM). Data represent the maximum response during the 30s agonist treatment taken from mean cell response versus time curves. All neurons responded to KCl (75 mM) applied immediately prior to ionomycin.

Figure 4: 9-OA- NO_2 fails to activate TRPA1-/- vagal neurons

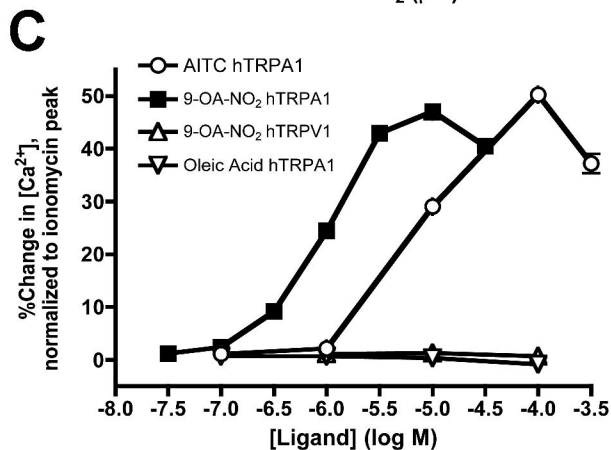
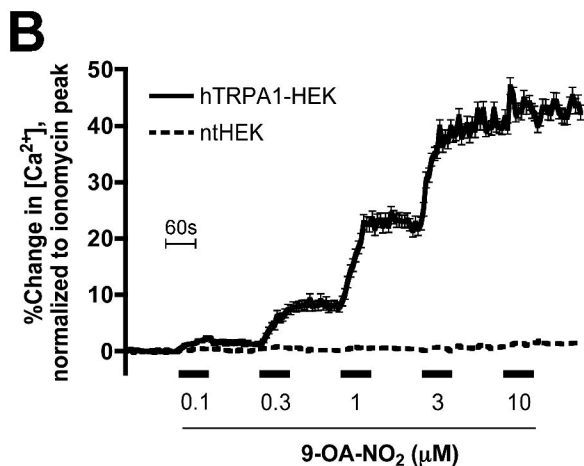
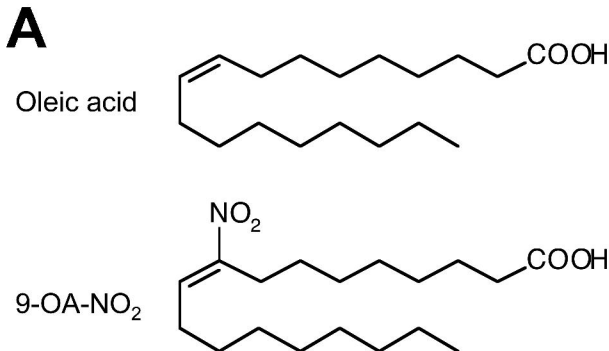
A, Mean \pm S.E.M. Ca^{2+} responses of vagal neurons responding to 9-OA- NO_2 (10 μM). Response to AITC (100 μM) and capsaicin (Caps, 1 μM) also shown. Data comprise of neurons from wild-type mice (black squares, 80 out of 196 neurons responding), neurons from TRPA1-/- mice (red squares, 34/124), and neurons from TRPA1-/-TRPV1-/- mice (blue squares, 20/101). Blocked line denotes the 30s application of agonist. All neurons responded to KCl (75 mM) applied immediately prior to ionomycin. B, Representative trace of the inward current evoked in a wild-type vagal neuron (held at -60mV) by 9-OA- NO_2 (10 μM) (and reversed by HC030031 (20 μM)) and capsaicin (1 μM). C, Representative trace of the inward current evoked in a TRPA1-/- vagal

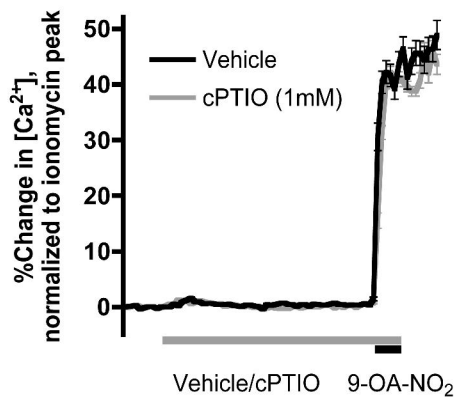
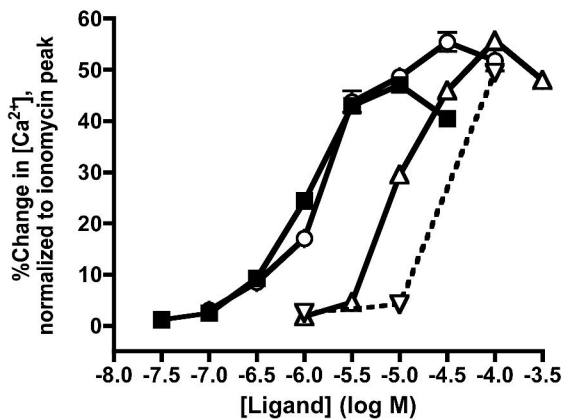
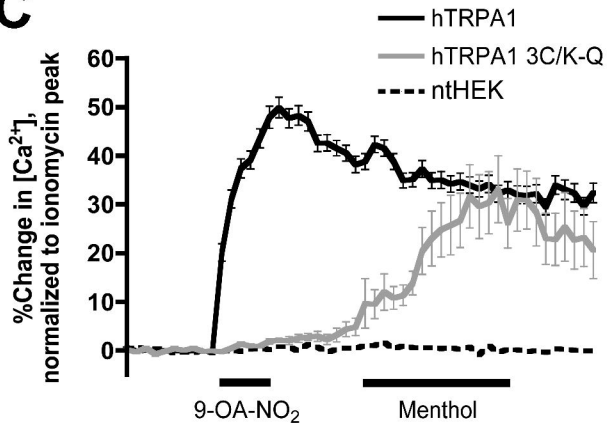
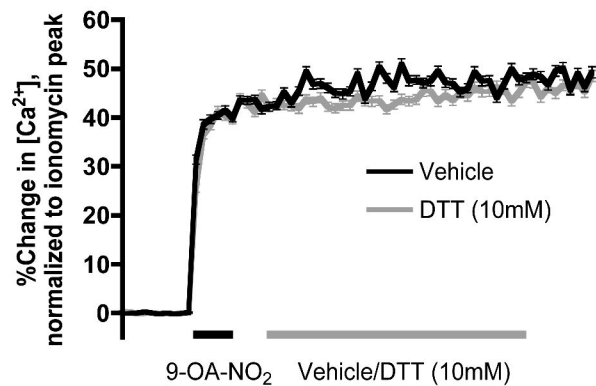
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neuron (held at -60mV) by 9-OA-NO₂ (10 μM) and capsaicin (1 μM). D, Mean ± S.E.M. Inward current density (pA/pF) of vagal neurons responding to 9-OA-NO₂ (10 μM). Data comprise of neurons from wild-type mice (black column) and neurons from TRPA1^{-/-} mice (white column).

Figure 5: 9-OA-NO₂ activation of C-fiber terminals.

A, Mean ± S.E.M. Total action potential discharge from individual identified bronchopulmonary C-fibers to 9-OA-NO₂ (30 μM) in paired experiments: control and vehicle-treated fibers (n=7) and control and AP-18 (30μM)-treated fibers (n=5). All C-fibers responded to capsaicin (300nM) at the end of the experiment. Only one fiber was assessed in each preparation. B, Representative trace of action potential discharge from a single bronchopulmonary C-fiber evoked by 9-OA-NO₂ (30 μM) in the absence and presence of AP-18 (30μM), followed by the response to capsaicin (300nM). Inset, action potential waveform of the individual bronchopulmonary C-fiber.



A**B****C****D**

A

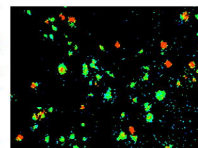
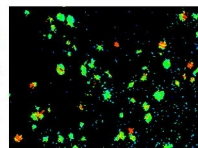
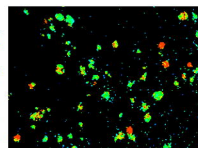
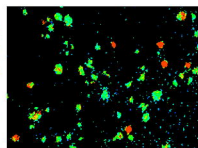
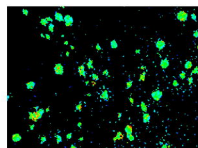
Background

9-OA-NO₂

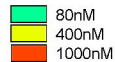
AITC

Caps

KCl



Trigeminal

[Ca²⁺]_{free}

58%

70%

58%

100%

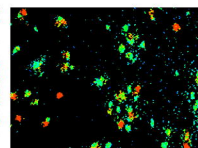
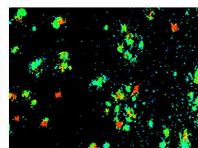
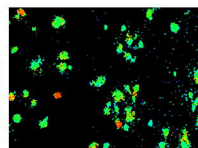
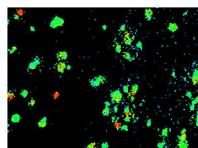
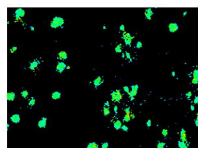
Background

9-OA-NO₂

AITC

Caps

KCl



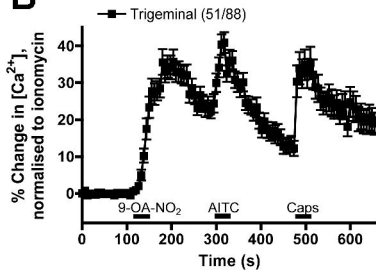
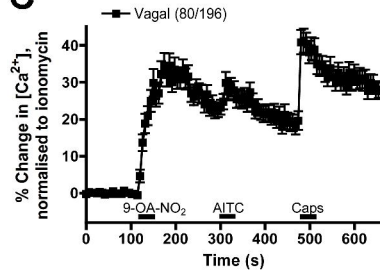
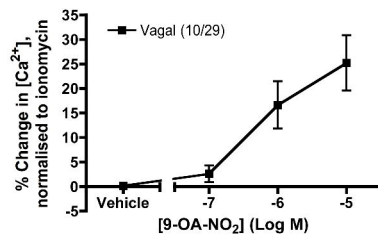
Vagal

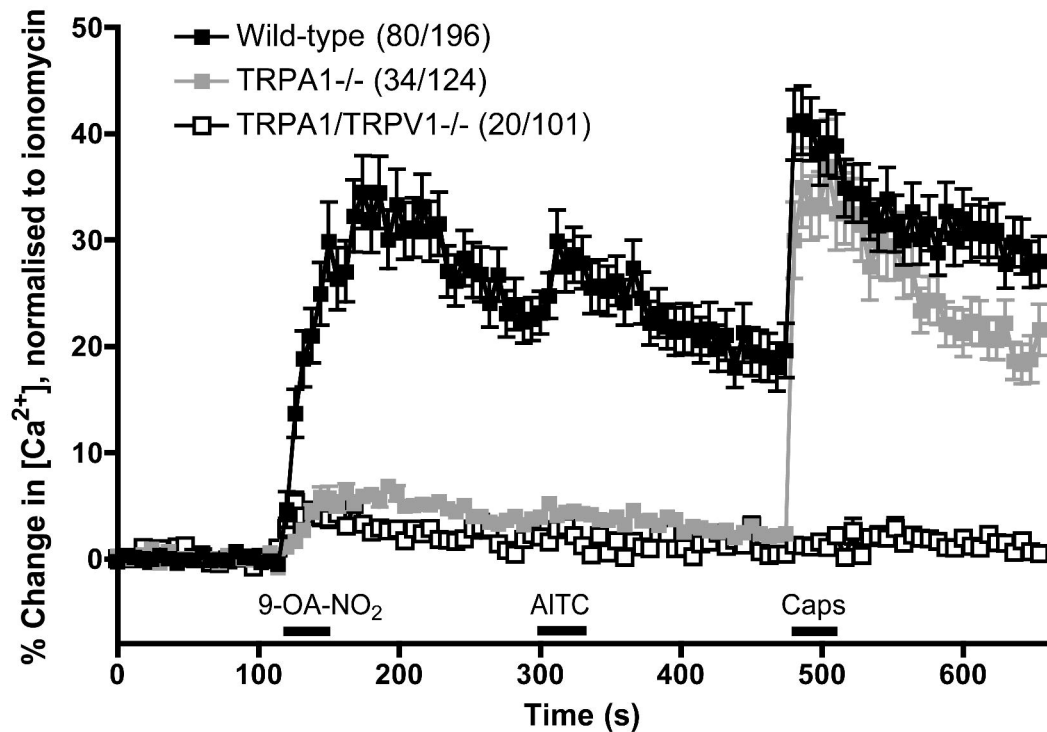
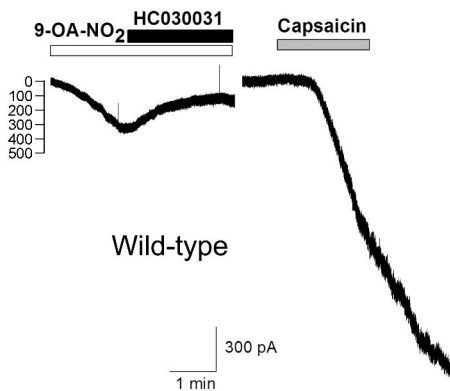
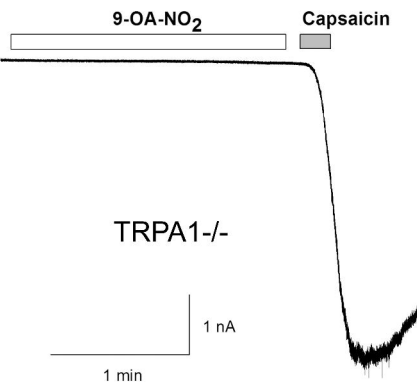
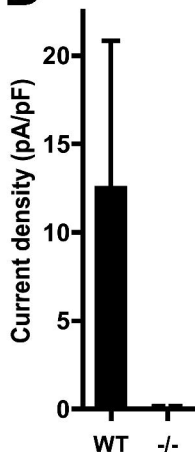
41%

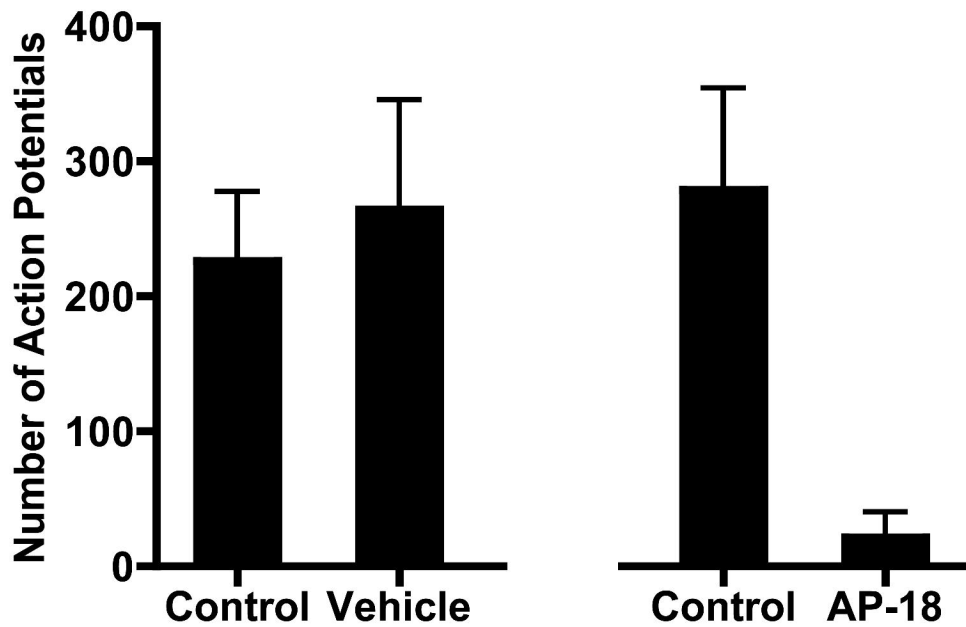
36%

62%

100%

B**C****D**

A**B****C****D**

A**B**