Opioid-Mediated Modulation of Acid-Sensing Ion Channel Currents in Adult Rat Sensory Neurons

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ABSTRACT

Muscle ischemia, associated with peripheral artery disease (PAD), leads to the release of proinflammatory mediators that decrease extracellular pH and trigger the activation of proton-activated acid-sensing ion channels (ASIC). Claudication pain, linked with low blood flow, can be partially relieved by endogenous opioid peptide release. However, we previously reported that sustained ASIC currents in dorsal root ganglion (DRG) neurons were enhanced by naturally occurring endomorphin-1 and -2 opioid peptides, indicating a role of opioid involvement in hyperalgesia. The present study examined whether clinically employed synthetic (fentanyl, remifentanil) and the semisynthetic opioid (oxycodone) would also potentiate sustained ASIC currents, which arise from ASIC3 channel isoforms. Here, we show that exposure of each opioid to DRG neurons resulted in potentiation of the sustained ASIC currents. On the other hand, the potentiation was not observed in DRG neurons from ASIC3 knockout rats. Further, the enhancement of the ASIC currents was resistant to pertussis toxin treatment, suggesting that Gαi/Gαo G-proteins are not involved. Additionally, the potentiation of sustained ASIC currents was greater in DRG neurons isolated from rats with ligated femoral arteries (a model of PAD). The effect of all three opioids on the transient ASIC peak current was mixed (increase, decrease, no effect). The inhibitory action appears to be mediated by the presence of ASIC1 isoform expression. These findings reveal that, under certain conditions, these three opioids can increase ASIC channel activity, possibly giving rise to opioid-induced hyperalgesia.

Introduction

Acid-sensing ion channels (ASIC) belong to the epithelial Na⁺ channel/degenerin (ENaC/DEG) family and are primarily Na⁺-selective channels that open in response to a drop in extracellular pH. In mammals, there are six ASIC isoforms (ASIC1a and 1b, ASIC2a and 2b, ASIC3, and ASIC4) produced by four genes (Baron and Lingueglia, 2015; Kellenberger and Schild, 2015). A functional ASIC channel is made up of three subunits and is either homomeric or heteromeric (Kellenberger and Schild, 2015). ASIC1–3 are expressed primarily in the central and peripheral nervous systems (Deval and Lingueglia, 2015). In the peripheral nervous system, ASIC1 and ASIC3 are found in trigeminal and dorsal root ganglion (DRG) sensory neurons (Deval and Lingueglia, 2015).

Although DRG neurons express primarily ASIC1 and ASIC3 isoforms, their stoichiometry is not known. Moreover, under ischemic conditions the expression levels of both isoforms have been reported to be altered (Liu et al., 2010; Walder et al., 2010; Fagg et al., 2017). Both variants exhibit a high degree of H⁺ ion sensitivity, and their threshold for activation occurs when the extracellular pH reaches 7.2 (Gründer and Pusch, 2015; Kellenberger and Schild, 2015). ASIC3 channel stimulation results in a biphasic current that is unique to this subunit. There is an initial rapidly inactivating current that is followed by a sustained current that lasts as long as the extracellular environment remains acidic (Gründer and Pusch, 2015). The inactivation time constants (τ) for ASIC1a and ASIC3 are 1.2–3.4 and 0.3 seconds, respectively (Kellenberger and Schild, 2015). The difference in this biophysical parameter provides a convenient way to distinguish ASIC1 from ASIC3.

Recent studies have shown modulation of ASIC currents by signaling molecules that are released during chronic inflammatory conditions. For instance, lactic acid can enhance ASIC1 and ASIC3 peak current in sensory neurons (Immke and McCleskey, 2001; Molliver et al., 2005). Enhancement of sustained ASIC3 currents by arachidonic acid and lysophosphatidylcholine has been described elsewhere (Smith et al., 2007; Marra et al., 2016). Likewise, the endogenous opioid peptides, dynorphins, which stimulate kappa opioid receptors, have been reported to potentiate ASIC1 currents independent of G protein coupling (Sherwood and Askwith, 2009).

In addition, limb ischemia in people with peripheral artery disease (PAD) can lead to intermittent claudication–leg pain

ABBREVIATIONS: ASIC, acid-sensing ion channels; DAMGO, [D-Ala²,N-MePhe⁴,Gly-ol]-enkephalin; Dil, DiICₙ(3)-tetramethylindocarbocyanine perchlorate; DRG, dorsal root ganglion; E-1/2, endomorphin 1/2; EGFP, enhanced green fluorescent protein; Fen, fentanyl; FP, freely profused; KO, knockout; Lig, ligated; MOR, mu opioid receptor; OIH, opioid-induced hyperalgesia; Oxy, oxycodone; PAD, peripheral artery disease; PTX, pertussis toxin; Rem, remifentanil; τ, inactivation time constant; WKY, Wistar-Kyoto.
induced by walking (Criqui and Aboyans, 2015). The pain associated with claudication partly results from tissue acido-
sis and inflammatory mediators, and is transduced by thin
muscle afferents. Prescription opioids make up one of the
treatment modalities for PAD patients with claudication, but
their use is limited by their tendency to cause tolerance,
physical dependence, and addiction resulting from the loss of
MOR at the surface level.

Moreover, the use of opioids for relief of either chronic pain or
postsurgical pain can, under certain conditions, lead to para-
doxical pain known as opioid-induced hyperalgesia (OIH). OIH,
a nociceptive sensitization state, occurs when the administra-
tion of opioids intended for pain relief leads to an increased
sensitivity to painful stimuli (Lee et al., 2011).

Endogenous opioid peptides are known to be released at
ischemic or inflammatory sites (Mousa et al., 2002). Previ-
ously, we found that the high-affinity mu opioid receptor
(MOR) peptides, endomorphin 1 (E-1) and endomorphin
2 (E-2), potentiated sustained ASIC3 currents in the absence
of MOR stimulation and G protein signaling (Farrag et al.,
2017). Thus, the present study expands on these findings and
determines whether the clinically employed opioids fentanyl
(Fen), remifentanil (Rem), and oxycodone (Oxy) would also
potentiate ASIC3 currents. The synthetic opioids Fen and Rem
(see Fig. 1A) have been reported to induce OIH when
administered after surgery (Fletcher and Martinez, 2014,
and references therein). Whether administration of the semi-
synthetic opioid Oxy (Fig. 1A) can lead to OIH is unknown.

**Materials and Methods**

**Animals.** All experiments were approved by the Penn State
College of Medicine or Medical College of Wisconsin Institutional
Animal Care and Use Committee (IACUC) and complied with the
National Institutes of Health guidelines. DRG neurons were isolated
from adult male Sprague-Dawley rats (125–175 g; Charles River, King
of Prussia, PA). For experiments shown in Fig. 3, Wistar-Kyoto (WKY)
and transgenic WKY ASIC3 knockout (KO) (ASIC3−/−) rats were
employed. Wild-type and mutant strain WKY rats were obtained from
the Gene Editing Rat Resource Center (Melinda Dwinell, Ph.D.,
PI/NHLBI; Medical College of Wisconsin). This ASIC3 KO strain
(WKY-Asic3−/−Mcwi; RGDBID: 12790599) was generated by clustered
regularly interspaced short palindromic repeats (CRISPR)/Cas9 tech-
nology, targeting the sequence GGCCCACACGCTCGGCGCA
GGG (protospacer adjacent motif underlined) into WKY/NCrl (Charles
River).
River Laboratories) rat embryos, resulting in 61 base pair deletions in exon 1 and predicted truncation of the normal ASIC3 open reading frame after only 22 amino acids.

**Femoral Artery Ligation and DRG Neuron Labeling.** Three days before the DRG neuron isolation, bilateral femoral artery ligations were performed with 5-0 silk sutures under anesthesia (3%-5% isoflurane), as described elsewhere (Copp et al., 2016). The wounds were closed with stainless steel wound clips. Under these conditions, there is a reduction of blood flow reserve capacity that nevertheless is sufficient to meet the metabolic demands at rest (Lash et al., 1995). Additionally, the triceps surae muscles were injected with the fluorescent neuronal tracer 3% DiI (3-tetramethylindocarbocyanine perchlorate; DiI; Thermo Fisher Scientific, Carlsbad, CA) prepared in dimethylsulfoxide. A total of 30 μl of DiI was injected per leg with a 30-gauge needle, which was left in the muscle for approximately 10 seconds to avoid tracer outflow.

**DRG Neuron Isolation and Dissociation.** Three days after artery ligation and tracer injection, the rats were anesthetized with CO₂ and sacrificed by decapitation. The lumbar (L₄–L₆) DRG were dissected and rapidly placed in ice-cold Hank’s balanced salt solution (Sigma-Aldrich, St. Louis, MO) and cleared of connective tissue. Thereafter, the ganglia were enzymatically dissociated in Earle’s balanced salt solution (Sigma-Aldrich) containing collagenase Type D (0.6 mg/ml; Roche, Mannheim, Germany), trypsin (0.35 mg/ml; Worthington, Lakewood, NJ), and DNase (0.1 mg/ml; Alfa Aesar, Ward Hill, MA) and shaken in a water bath for 40 minutes at 35°C. Afterward, the neurons were centrifuged twice for 6 minutes at 50 g. Thereafter, the DRG neurons were placed in minimum essential medium (Thermo Fisher Scientific) supplemented with 10% FBS, 1% glutamine, and 1% penicillin-streptomycin. The dissociated neurons were then plated onto poly-L-lysine-coated 35-mm polystyrene dishes and incubated in a humidified atmosphere at 37°C in 5% CO₂ and 95% air. In some experiments, the DRG neurons were pretreated overnight with pertussis toxin (PTX, 0.5 μg/μl; List Biologic Laboratories, Manassas, VA) to facilitate identification of transfected L-cells. After overnight incubation, electrophysiological recordings were obtained with Igor Pro (WaveMetrics, Lake Oswego, OR). The figures shown were obtained with Autodesk Graphic software, and the statistical analysis was performed with Prism (GraphPad Software, La Jolla, CA) and expressed as mean ± S.D. The sample sizes were prespecified before data collection. Data comparison between two groups was determined with the use of unpaired, two-tailed Student’s t test, while one-way ANOVA followed by Tukey’s test, while one-way ANOVA followed by Tukey’s (Fig. 3) or Bonferroni (Fig. 4) post hoc tests were employed for multiple comparisons. P < 0.05 was considered statistically significant. The intergroup comparisons were prespecified, and all results from the statistical tests are reported. The statistical comparisons performed with the groups shown in Fig. 3C (WKY vs. ASIC3 KO), Fig. 3D (WKY vs. ASIC1a-L cells vs. ASIC3-L cells), Fig. 4D (Oxy FP vs. Oxy LIG vs. Fen FP vs. Fen LIG vs. Rem FP vs. Rem LIG), and Fig. 4E (FP vs. LIG) were prespecified and not exploratory.

**Results**

**Enhancement of Sustained ASIC Currents by Clinical Opioids in Rat DRG Neurons.** In the first set of experiments, Dil-labeled neurons were exposed to opioids routinely employed for intraoperatively (Rem and Fen) or chronic pain (Oxy). Figure 2 shows superimposed ASIC current traces before (black traces) and during (red traces) application of Oxy (10 μM, Fig. 2Aii), Fen (10 μM, Fig. 2Bii), and Rem (10 μM, Fig. 2Cii). It can be observed that the amplitude of the sustained currents increased after opioid exposure when compared with pH 6.0 alone.

The summary dot plot shown in Fig. 2D illustrates the increase in sustained ASIC current for all three opioids. The concentration (10 μM) of clinical opioids chosen was based on our previous observations of the endorphin pharmacologic profiles that showed maximal ASIC3 potentiation between 3 and 10 μM (Farrag et al., 2017).

To determine whether the sustained current enhancement was mediated via stimulation of MOR, which couple to...
control solution (pH 7.4) for 3 minutes. Afterward, the cells pretreated with naloxone (30 μM) in neurons before (Ctrl [control], black trace) and after exposure to 10 μM Oxy, Fen, and Rem in control (A–Ci) and in neurons pretreated overnight in PTX (0.5 μg/μl; A–Ci). The solid bars indicate the 10-second exposure to either pH 6.0 alone or pH 6.0 + opioid. The cells were preexposed to the opioids (pH 7.4) for 3 minutes before application with pH 6.0 + opioid. The holding potential (Vh) for DRG neurons was −80 mV. (D) Summary dot plot with mean (± S.D.) of the Oxy-, Fen- and Rem-mediated enhancement of sustained ASIC currents in control and PTX-treated DRG neurons. The numbers in parenthesis indicate the number of recordings.

Gaα/Gαo G proteins, a group of Dil-labeled neurons was pretreated overnight with PTX. The results in Fig. 2 (right) show that all three MOR agonists enhanced the sustained currents in a similar magnitude as that observed in control neurons. The results for all PTX-treated groups are summarized in Fig. 2D.

In a separate set of experiments, we measured τ values of L-cells transfected with either ASIC1a or ASIC3 cDNA. The τ values for L-cells expressing ASIC1a are similar in magnitude to neurons isolated from ASIC3 KO rats (Fig. 3D). The ASIC3-expressing L-cells displayed τ values that were close to 0.3 seconds.

We next compared the effect of opioids on τ values in ASIC1a- and ASIC3-expressing L-cells. The τ values of the L-cell group expressing ASIC1a were 3.15 ± 0.38 and 2.47 ± 0.70 seconds (n = 4; P = 0.156, paired t test) before (pH 6.0) and during (pH 6.0 + 10 μM Oxy) exposure, respectively. Additionally, the τ values for ASIC3-expressing L-cells were 0.37 ± 0.08 and 0.45 ± 0.06 seconds (n = 19; P = 0.139, paired t test), before and during (pH 6.0 + 10 μM Oxy).

Effect of Muscle Ischemia on the Opioid-Mediated Enhancement of Sustained ASIC Currents in Rat DRG Neurons. Previous studies have shown that ASIC expression levels are altered under ischemic conditions such as that observed after femoral ligation (Liu et al., 2010; Xing et al., 2012). We have recently reported that ligation leads to both a decrease in ASIC1 and increase in ASIC3 expression levels in rat DRG neurons (Farrag et al., 2017). We thus compared the effect of the three clinical opioids on sustained ASIC currents in Dil-labeled neurons from both rats with freely perfused (FP, control) and ligated (LIG) femoral arteries.

The current traces shown in Fig. 4, Ai, Bi, and Ci are those obtained from control rats before (black traces) and during (red traces) opioid exposure. The traces shown to the right are those recorded in neurons isolated from rats with ligated arteries. The results summarized in Fig. 4D indicate that Oxy increases observed for Oxy alone and Oxy + naloxone were 83% ± 68% and 472% ± 386% (n = 5, P = 0.35), respectively. It should be noted that the higher increase observed with both Oxy + naloxone was a result of one cell exhibiting an enhancement of approximately 20-fold. Overall, these results indicate that the opioids modulate the sustained currents independent of Gaα/Gαo subunits or MOR activation, which is consistent with our previous report (Farrag et al., 2017).

To examine whether the potentiation of sustained ASIC currents by opioids was a result of ASIC3 activation alone, we next isolated DRG neurons from control (WKY) and ASIC3 KO rats. The ASIC currents shown in Fig. 3A show that exposure to Rem (10 μM) potentiated the sustained ASIC current almost 3-fold. On the other hand, the application of Rem (10 μM) to the neuron isolated from an ASIC3 KO rat failed to enhance the sustained ASIC currents (Fig. 3B). The summary dot plot in Fig. 3C indicates that Rem significantly (P < 0.001) enhanced ASIC currents when compared with DRG neurons isolated from ASIC3 KO rats. In 9 of 10 neurons, Rem exposure caused a slight inhibition of the sustained ASIC currents, and it nearly doubled the ASIC current in one neuron.

As mentioned previously, τ values for ASIC1 and ASIC3 are quite distinct. ASIC1 exhibit slow τ, while τ for ASIC3 is fast (Kellenberger and Schild, 2015). Therefore, we next measured τ for both groups of DRG neurons. The plot in Fig. 3D indicates that τ values observed in WKY DRG neurons range from 0.1 to 4.4 seconds, indicative of ASIC current heterogeneity. However, the τ values of DRG neurons from ASIC3 KO were significantly (P < 0.001) greater when compared with control neurons. The lowest τ value for DRG neurons from KO rats was 1.6 seconds, which suggests that ASIC currents do not exhibit ASIC3-like τ values.

In a separate set of experiments, we measured τ values of L-cells transfected with either ASIC1a or ASIC3 cDNA. The τ values for L-cells expressing ASIC1a are similar in magnitude to neurons isolated from ASIC3 KO rats (Fig. 3D). The ASIC3-expressing L-cells displayed τ values that were close to 0.3 seconds.

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exposure led to a significantly ($P < 0.05$) greater enhancement of sustained ASIC currents. Although both Fen and Rem application exhibited greater sustained ASIC currents when compared with neurons from freely perfused muscle, no statistical significance was reached (Fig. 4D).

Figure 4E depicts the summary of the measured $\tau$ for both group of neurons. It can be observed that the mean $\tau$ of DRG neurons from ligated rats was significantly ($P < 0.01$) lower than that from control rats (2.41 vs. 0.88 seconds). The lower $\tau$, characteristic of ASIC3 channels, observed in neurons from the LIG rat group was likely a result of greater expression of this channel subunit. The mean $\tau$ for WKY KO rats (2.70 seconds; Fig. 3D) was comparable to that observed in Sprague-Dawley rats (Fig. 4E).

**Effect of Opioids on Transient Peak Currents in Rat DRG Neurons and in ASIC-Transfected L-Cells.** In the current study and previously (Farrag et al., 2017), we observed that whereas the opioids consistently enhanced the sustained ASIC currents, their effect on transient peak currents was mixed. That is, application of the opioids would increase, decrease, or have a minimal effect on the peak currents (see Figs. 2–4). Given that functional ASIC are trimeric, the actual ASIC1 and ASIC3 contribution to the total ASIC peak currents is difficult to ascertain. We reasoned that the varied responses may result from isoform heterogeneity in DRG neurons. Thus, we next used $\tau$ values to sort the DRG neurons isolated from FP and LIG rats into “ASIC1-like” and “ASIC3-like.” That is, neurons with $\tau$ greater than 0.8 seconds were grouped into the former, and the rest were grouped into the latter.

Figure 5A is a summary dot plot of the change in transient peak currents after exposure to Oxy (10 $\mu$M) in neurons grouped based on this criterion. The plot indicates that in ASIC1- and ASIC3-like neurons isolated from FP rats, Oxy application, for the most part, led to inhibition of the transient currents while in approximately 30% of cells in each group peak current enhancement was observed. We then compared the effect of Oxy on DRG neurons isolated from rats with ligated femoral arteries. In this group of neurons, Oxy application primarily blocked peak currents of ASIC1-like cells (6/7) but remained mixed in the ASIC3-like group of cells.

We next employed this criterion in L-cells expressing ASIC1 or ASIC3 alone (i.e., homomeric) or ASIC3 cotransfected with ASIC1 at a 1:1 and 1:3 ASIC3:ASIC1 ratio. The dot plot in Fig. 5B summarizes the effect of Oxy (10 $\mu$M) on the transient peak currents. The plot shows that Oxy exerted mixed effects on homomeric ASIC1 and ASIC3 peak currents. However, the coexpression of ASIC3 and ASIC1a led to mainly inhibitory effects with either a 1:1 or 1:3 ASIC3:ASIC1a ratio. This suggests that the opioid-mediated inhibitory of peak currents are more likely to occur under conditions where both ASIC1a and ASIC3 isoforms are coexpressed.

We previously found that E-2 caused significant potentiation of sustained ASIC currents. Thus, in the next set of experiments we examined whether coapplication of E-2 and Oxy would exert primarily potentiating effects on ASIC peak currents. DRG neuron from LIG rats, with presumably greater ASIC3 expression levels, and L-cells transfected with ASIC1a and/or ASIC3 were preincubated in Oxy (pH 7.4) for 3 minutes. Thereafter, both E-2 and Oxy (both at 10 $\mu$M) were applied to the cells. The results in Fig. 5A indicate that in ASIC1-like cells, E-2 + Oxy primarily (6/8) blocked peak currents but enhanced the currents in most of ASIC3-like cells (8/10). Similarly, in L-cells coexpressing both ASIC channel subunits resulted in block of the peak currents by both agonists. Yet both E-2 and Oxy potentiated all ($n = 9$) ASIC3-expressing L-cells, though in one cell the enhancement was relatively small.

**Discussion**

The results of our present study demonstrate that the synthetic and semisynthetic opioids Oxy, Fen, and Rem, like endogenous opioid peptides, can potentiate sustained ASIC currents in acutely isolated DRG neurons innervating triceps surae muscle. The opioid-mediated potentiation was also significantly greater in DRG neurons isolated from rats with ligated femoral arteries, a model of PAD. Our findings further suggest...
that the enhancement of the sustained currents does not result from stimulation of G protein subunits (Gαi/Gαo), known to couple to MOR in rat DRG neurons (Hassan et al., 2014), because PTX pretreatment did not prevent current potentiation. Additionally, enhancement of the sustained ASIC3 currents occurred in the presence of the MOR blocker naloxone, suggesting that MOR stimulation is not involved. Rather, it is likely that the opioids interact directly with ASIC3 channel subunits. Moreover, the three opioids exerted varying effects on the peak transient currents: enhanced, blocked, or no effect. Thus, the overall modulation of ASIC currents by opioids appears to be multifactorial.

Most studies that have reported the effect of inflammatory mediators on ASIC currents have focused on peak transient currents. Dynorphins, for example, are thought to interact with ASIC1 channel subunits that potentiate ASIC currents (Sherwood et al., 2012). Similar observations have been reported with FFRM amides (Askwith et al., 2000). In cases where it is shown that there is crosstalk between the receptor and ASIC channels, this has also reported the effect on peak transient currents. Unlike these studies, however, our results show that the clinical opioids can exert variable effects, as has also been observed with endogenous opioid peptides (Farrag et al., 2017).

In the present study, we employed the desensitization time constant as an indicator of ASIC isoform influence on whole-cell ASIC current response to opioids. Sorting τ values into “ASIC1-” and “ASIC3-like” showed that exposure of DRG neurons to opioids will exert primarily an inhibitory effect on transient peak currents. Under conditions where ASIC3 expression dominates (i.e., DRG from LIG group), a greater fraction of peak currents will be potentiated by opioids. The varied responses demonstrate that ASIC stoichiometry exhibits a complex pharmacologic profile resulting from the sum of two independent types of actions that comprise whole-cell ASIC currents. These observations, which have been previously reported by others (Babinski et al., 2000; Hesselager et al., 2004; Hattori et al., 2009; Kusama et al., 2013), are thought to serve as a regulatory mechanism in response to slight changes in extracellular pH.
One study that examined the effect of either morphine or [D-Ala²,N-MePhe⁴,Gly-ol]-enkephalin (DAMGO) in rat DRG found that either agonist blocked the peak transient currents (Cai et al., 2014). However, the effect on sustained ASIC currents was not reported. Unlike a previous report (Farrag et al., 2017) and our present study, the investigators showed that the inhibition of the peak transient currents was due to crosstalk between MOR and ASIC isoforms that employed Ca²⁺ signaling events. Although morphine and Oxy share similar structures, we observed sustained ASIC current enhancement in the presence of PTX and in L-cells that do not express MOR. Our results also show that Oxy exerted variable effects on peak currents particularly in DRG neurons from FP rats, though most were inhibitory.

Ischemic pain, often observed with claudication and angina, is associated with severe muscle ischemia. The sustained component of ASIC currents is thought to play a major role in pain and pressor responses associated with tissue ischemia as well as inflammation (Naves and McCleskey, 2005; Salinas et al., 2009; Sluka and Gregory, 2015). The sustained current appears to be responsible for the long-lasting pain observed under chronic ischemic conditions. Previous studies have reported that ASIC isoform expression levels in sensory neurons are altered under inflammatory or ischemic conditions. For instance, the carrageenan-induced muscle inflammation in mice has been reported to increase ASIC2 and ASIC3 expression levels while ASIC1 is unchanged (Walder et al., 2010). Femoral ligation in rats has been shown to significantly increase ASIC3 expression while decreasing ASIC1 levels (Liu et al., 2010; Xing et al., 2012; Farrag et al., 2017).

In our current study, the synthetic opioids, particularly Oxy, significantly enhanced the sustained ASIC currents in DRG neurons from rats with ligated femoral arteries. This group of neurons also exhibited a significantly lower desensitization constant in the same group of neurons, which is indicative of greater ASIC3 contribution to the total ASIC currents. In addition, the effect of Oxy on ASIC3 channel currents increased \( \tau \) values by approximately 20%, while \( \tau \) for ASIC1 currents decreased 20%. These slight changes suggest Oxy exposure may increase the response of ASIC3 channels to acid under acidic conditions, whereas the opposite would occur with ASIC1. More recently, E-1 was shown to also slow desensitization of ASIC3 currents recorded in Xenopus oocytes (Vyvers et al., 2018).

Although the amplitude of native sustained ASIC currents is relatively smaller than the peak current, it persists as long as the extracellular pH remains acidic. The sustained ASIC3 currents have been shown to make up of the overlap of both activation and inactivation curves, also referred to as window currents (Yagi et al., 2006). Signaling molecules, such as arachidonic acid, lysophosphatidylcholine, and endomorphins, enhance the window currents by both shifting the pH activation curve leftward to more basic pH values and the inactivation curve rightward to more acidic values (Deval et al., 2008; Marra et al., 2016; Farrag et al., 2017). The end result is an increase in the probability that closed nonactivated ASIC3 channels will open within this pH range when exposed these signaling elements and increase nociceptor excitability.

Our observations of the opioid-mediated sustained ASIC current potentiation bear physiologic relevance to the pathologic conditions such as PAD. Figure 6 illustrates our hypothesis by which OIH can occur from muscle ischemia and long-term opioid use. Under conditions where exercising muscle is freely perfused (Fig. 6A), lactic acid release will decrease extracellular pH. As a result, E-2 is released by DRG neurons in the muscle and dorsal horn.

Note that Ca²⁺ ions are chelated by lactic acid, which potentiates ASIC activity. At the presynapse, the afferent-mediated release of E-2 causes MOR stimulation such that the ascending pain signaling pathway is blocked. Figure 6B depicts an ischemic muscle with low blood flow that is accompanied by the following: 1) an increase in local inflammatory mediators, 2) an increase in ASIC3 isoform expression, 3) a decrease in ASIC1 expression, 4) an increase in E-2 release, and 5) an increase in Ca²⁺ ion chelation by lactic acid. The end result is sensory neuron hyperexcitability and pain.

Claudication pain can be alleviated with prescription opioids such as Oxy (Samolsky Dekel et al., 2010). However, the pain relief comes at the cost of developing opioid tolerance and eventual addiction. In some cases, it is likely that opioid tolerance can also lead to OIH. That is, with the continued use of opioid medication and possibly the increased E-2 release at the ischemic site, patients exhibit sensitization to painful stimuli. This has been documented with Rem and Fen (Lee et al., 2011; Kim et al., 2014; Santonocito et al., 2018).

In summary, our results show that three clinically employed, high-affinity MOR agonists potentiate ASIC currents in DRG neurons, with a more pronounced effect in neurons isolated...
from animals with ligated arteries. Like other inflammatory mediators, such as lactic acid, arachidonic acid, dynorphin peptides, and endomorphins, these opioids can effectively modulate ASIC currents independent of receptor activation. The results identify a novel, direct interaction of ASIC3 channel isoforms with opioids. This interaction may help to explain and understand the molecular mechanism associated with OIH that occurs without MOR stimulation.

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Authorship Contributions

Participated in research design: Zaremba, Ruiz-Velasco.
Conducted experiments: Zaremba, Ruiz-Velasco.
Performed data analysis: Zaremba, Ruiz-Velasco.
Wrote or contributed to writing of the manuscript: Zaremba, Ruiz-Velasco.

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