


# Legumain Induces Oral Cancer Pain by Biased Agonism of Protease-Activated Receptor-2

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Oral squamous cell carcinoma (OSCC) is one of the most painful cancers, which interferes with orofacial function including talking and eating. We report that legumain (Lgmn) cleaves protease-activated receptor-2 (PAR<sub>2</sub>) in the acidic OSCC microenvironment to cause pain. Lgmn is a cysteine protease of late endosomes and lysosomes that can be secreted; it exhibits maximal activity in acidic environments. The role of Lgmn in PAR<sub>2</sub>-dependent cancer pain is unknown. We studied Lgmn activation in human oral cancers and oral cancer mouse models. Lgmn was activated in OSCC patient tumors, compared with matched normal oral tissue. After intraplantar, facial or lingual injection, Lgmn evoked nociception in wild-type (WT) female mice but not in female mice lacking PAR<sub>2</sub> in Nav1.8-positive neurons (*Par<sub>2</sub>Nav1.8*), nor in female mice treated with a Lgmn inhibitor, LI-1. Inoculation of an OSCC cell line caused mechanical and thermal hyperalgesia that was reversed by LI-1. *Par<sub>2</sub>Nav1.8* and *Lgmn* deletion attenuated mechanical allodynia in female mice with carcinogen-induced OSCC. Lgmn caused PAR<sub>2</sub>-dependent hyperexcitability of trigeminal neurons from WT female mice. *Par<sub>2</sub>* deletion, LI-1, and inhibitors of adenylyl cyclase or protein kinase A (PKA) prevented the effects of Lgmn. Under acidified conditions, Lgmn cleaved within the extracellular N terminus of PAR<sub>2</sub> at Asn<sup>30</sup>↓Arg<sup>31</sup>, proximal to the canonical trypsin activation site. Lgmn activated PAR<sub>2</sub> by biased mechanisms in HEK293 cells to induce Ca<sup>2+</sup> mobilization, cAMP formation, and PKA/protein kinase D (PKD) activation, but not β-arrestin recruitment or PAR<sub>2</sub> endocytosis. Thus, in the acidified OSCC microenvironment, Lgmn activates PAR<sub>2</sub> by biased mechanisms that evoke cancer pain.

**Key words:** asparaginyl endopeptidase; cancer pain; legumain; oral cancer; protease; protease-activated receptor-2

## Significance Statement

Oral squamous cell carcinoma (OSCC) is one of the most painful cancers. We report that legumain (Lgmn), which exhibits maximal activity in acidic environments, cleaves protease-activated receptor-2 (PAR<sub>2</sub>) on neurons to produce OSCC pain. Active Lgmn was elevated in OSCC patient tumors, compared with matched normal oral tissue. Lgmn evokes pain-like behavior through PAR<sub>2</sub>. Exposure of pain-sensing neurons to Lgmn decreased the current required to generate an action potential through PAR<sub>2</sub>. Inhibitors of adenylyl cyclase and protein kinase A (PKA) prevented the effects of Lgmn. Lgmn activated PAR<sub>2</sub> to induce calcium mobilization, cAMP formation, and activation of protein kinase D (PKD) and PKA, but not β-arrestin recruitment or PAR<sub>2</sub> endocytosis. Thus, Lgmn is a biased agonist of PAR<sub>2</sub> that evokes cancer pain.

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**Table 1. Patient profiles**

Patient #	Sex	Age	Ethnicity	Tumor location	Primary tumor stage	Nodal status
1	F	71	Hispanic	Mandibular gingiva	pT4a	pN0
2	M	57	Hispanic	Mandibular gingiva	pT2	pN2a
3	M	66	Hispanic	Floor of mouth, mandibular gingiva	pT4a	pN0
4	F	77	White/Non-Hispanic	Mandibular gingiva	pT4a	pN0
5	F	50	Asian	Tongue	pT1	pN0
6	M	93	Asian	Mandibular gingiva	pT2	pN0
7	F	81	White/Non-Hispanic	Maxillary gingiva	pT2	pN0

## Introduction

Up to 90% of cancer patients endure pain; oral cancer is one of the most painful (van den Beuken-van Everdingen et al., 2007). Pain often overwhelms oral cancer patients in the final months of life; these patients suffer most while speaking, drinking, or eating, and their quality of life plummets (Connelly and Schmidt, 2004; Kolokythas et al., 2007). Oral squamous cell carcinoma (OSCC) pain worsens with disease progression and responds poorly to opioids. Development of an alternative to opioids is stymied by our poor understanding of the mechanism driving cancer pain. While the etiology of oral cancer pain is not well understood, it is known that OSCC secretes mediators that sensitize and activate nociceptors within the cancer microenvironment and generate pain. These mediators include endothelin, ATP, nerve growth factor, and proteases; proteases produce pain by cleaving protease-activated receptor-2 (PAR<sub>2</sub>) on nociceptors (Pickering et al., 2008; Schmidt et al., 2007; Lam and Schmidt, 2010; Ye et al., 2011, 2014b; Lam et al., 2012). The proteases that activate PAR<sub>2</sub> in the OSCC microenvironment are unknown. Anaerobic metabolism in tumors and inflamed tissues acidifies extracellular fluid. Legumain (Lgmn; asparaginyl endopeptidase) is a cysteine protease of late endosomes and lysosomes with an acidic pH optimum. Although Lgmn has been implicated in tumor metastasis (Kembhavi et al., 1993), and patients with metastatic oral cancer report greater pain (Connelly and Schmidt, 2004), it is unknown whether Lgmn causes PAR<sub>2</sub>-dependent cancer pain.

PAR<sub>2</sub> is a G-protein-coupled receptor (GPCR) expressed by nociceptors that mediates neurogenic inflammation and pain (Steinhoff et al., 2000; Vergnolle et al., 2001). Proteases activate PAR<sub>2</sub> through distinct mechanisms. The canonical agonists trypsin, trypsinase, and kallikrein cleave within the extracellular N terminus of PAR<sub>2</sub>, which exposes a tethered ligand that binds to and activates the cleaved receptor (Nystedt et al., 1995; Böhm et al., 1996a; Corvera et al., 1997; Angelo et al., 2006). The biased agonists cathepsin S and elastase cleave at different sites within the PAR<sub>2</sub> N terminus, leading to distinct pathways of PAR<sub>2</sub> signaling and trafficking (Ramachandran et al., 2009; Zhao et al., 2014). PAR<sub>2</sub> couples to signaling pathways that sensitize and activate pain-related ion channels, including the transient receptor potential vanilloid (TRPV1, TRPV4) and ankyrin (TRPA1) channels, yielding sustained sensitization of nociceptors and chronic pain (Amadesi et al., 2006; Dai et al., 2007; Zhao et al., 2015). Since PAR<sub>2</sub> is upstream of these pronociceptive channels, blockade of proteases and of PAR<sub>2</sub> is advantageous for the management of cancer pain. Therefore, there is a need to identify proteases that remain active in the acidified extracellular milieu of tumors and to determine whether they activate PAR<sub>2</sub> to produce pain. Some proteases known to activate PAR<sub>2</sub> show diminished activity under acidic conditions (Hachem et al., 2003, 2005). Moreover, proteases are often profiled in diseased tissues by measurement of protein or mRNA, not activity.

Here, we report that Lgmn is a unique activator of PAR<sub>2</sub>. We found that Lgmn is reproducibly and robustly activated in OSCC patients and mice with OSCC. Under acidified conditions, Lgmn cleaved PAR<sub>2</sub> at a distinct site and activated PAR<sub>2</sub> by biased mechanisms, leading to hyperexcitability of nociceptors and nociceptive behavior in mice. A Lgmn inhibitor prevented OSCC pain in mice. Thus, we have identified Lgmn as a novel mediator and therapeutic target for OSCC pain.

## Materials and Methods

**Reagents.** We used recombinant human Lgmn (440,000 ng/ml, catalog #2199-CY-010, R & D System), Lgmn substrate Z-Ala-Ala-Asn-AMC (I-1865; Bachem), DMEM and HBSS (ThermoFisher), PAR<sub>2</sub> agonist 2-Furoyl-LIGRLO-NH<sub>2</sub> and PAR<sub>1</sub> agonist TFLLR-NH<sub>2</sub> (Tocris), Lgmn-generated PAR<sub>2</sub> activating peptide (RSSKGR; GL Biochem), and other reagents (Sigma) unless otherwise specified.

**Lgmn activation and inhibition.** Lgmn was activated per manufacturer protocol. Lgmn activation was confirmed by incubating Lgmn (1 ng/μl) with Z-Ala-Ala-Asn-AMC (200 μM) in dilution buffer [250 mM NaCl, 50 mM 2-(N-morpholino) ethanesulfonic acid (MES); pH 5.0] or MES-HBSS (HBSS, 50 mM MES pH 5.0, 5.5 or 6.0). Fluorescence (excitation 340 nm, emission 460 nm) was measured every 30 s for 10 min in a Flexstation three plate reader (Molecular Devices). Lgmn (1 ng/μl) was assayed for activity in the presence of the Lgmn inhibitors QDD100531 or QDD123427 (1 pM–10 μM) and PAR<sub>2</sub> antagonists I-343 (10 μM) or GB88 (10 μM; Farmer, 2013; Lieu et al., 2016; Jimenez-Vargas et al., 2018). Rich Williams provided the Lgmn inhibitors. The specificity of QDD100531 was demonstrated in Ness et al. 2015 (compound 9h in Supplementary Table 1); QDD100531 showed no reactivity to other proteases including cathepsin S, cathepsin B, caspase-3, caspase-8, or USP17 (Ness et al., 2015). QDD123427 showed similar specificity (R. Williams, personal communication). GB88 exhibited specific antagonist activity against four PAR<sub>2</sub> agonists differing in structure and mechanism; selectivity of GB88 for PAR<sub>2</sub> over PAR<sub>1</sub> and PAR<sub>4</sub> was also demonstrated (Suen et al., 2012). I-343, a member of the I-191 family of full PAR<sub>2</sub> antagonists, inhibited inositol phosphate-1 (IP<sub>1</sub>) generation induced by the PAR<sub>2</sub> agonists trypsin and 2-Furoyl-LIGRLO-NH<sub>2</sub>, but not IP<sub>1</sub> accumulation induced by ATP (Farmer, 2013; Jiang et al., 2018; Jimenez-Vargas et al., 2018). For studies in mice, activated Lgmn was diluted in 50 mM MES and 250 mM NaCl, at pH 5.0 (dilution buffer) to a concentration of 300 ng/20 μl. For the *in vivo* experiments with LI-1 (10 mM, 100 μl, diluted in DMSO; Lee and Bogyo, 2010) the inhibitor was injected into the tail vein 2 h before injection of Lgmn. LI-1 was a gift from Matthew Bogyo. It is a covalent Lgmn inhibitor that exhibits >20,000-fold selectivity for Lgmn over cathepsin B, cathepsin L, and caspase-3. It has previously been shown to inhibit all Lgmn activity *in vivo* within 1 h of administration (Edgington-Mitchell et al., 2016).

**OSCC patients.** Patients were screened and enrolled through New York University (NYU) Oral Cancer Center after consent. Detailed demographic information (age, sex, ethnicity, cancer location, primary tumor stage, and evidence of metastasis) was collected. During surgical resection, tumor and matched normal oral mucosa specimens were collected (normal was harvested at anatomically matched contralateral site). Specimens were frozen in liquid nitrogen and maintained at –80°C. The

Committee on Human Research at NYU Langone Medical Center approved human studies.

**Mice.** Female C57BL/6J (#000664) and NU/J *Foxn1*<sup>tmu</sup> athymic mice (#002019), four to eight weeks, were from The Jackson Laboratory. Female C57BL/6J and *F2rl1*<sup>-/-</sup> (B6. Cg-F2rl1<sup>tm1Msb/J</sup>) mice (#004993), four to eight weeks, from The Jackson Laboratory, were used for trigeminal ganglia (TG) dissociation. *F2rl1* conditional knock-out (KO) C57BL/6 mice were generated by genOway as described (Jimenez-Vargas et al., 2018). *Lgmn*<sup>-/-</sup> C57BL/6N mice were a gift from Thomas Reinheckel (Matthews et al., 2010). The NYU Institutional Animal Care and Use Committee approved mouse studies.

**Analysis of total and active Lgmn in tissues.** Snap frozen human and murine tissues were sonicated in 50 mM citrate pH 5.5, 0.5% CHAPS, 0.1% Triton X-100, and 4 mM DTT (10 μl/mg tissue). Solids were cleared by centrifugation and protein concentration was measured by BCA assay (Pierce). Protein was diluted in citrate buffer (50 μg/20 μl buffer), and LE28 was added from a 100× DMSO stock (1 μM final; Edgington et al., 2013). Samples were incubated at 37°C for 15 min, and the reaction was quenched with 5× sample buffer [200 mM Tris-Cl (pH 6.8), 8% SDS, 0.04% bromophenol blue, 5% β-mercaptoethanol, and 40% glycerol]. Protein was resolved on a 15% polyacrylamide gel under reducing conditions. LE28 binding was detected by scanning the gel for Cy5 fluorescence using a Typhoon 5 (GE Healthcare). Proteins were transferred to nitrocellulose membranes for immunoblotting with a goat anti-human Lgmn antibody (R & D AF2199, 1:1000 diluted in 50% Li-Cor blocking buffer and 50% PBS-T containing 0.05% Tween 20). Donkey-anti-goat-HRP (1:10,000; A15999; Invitrogen) was used for detection with Clarity Western ECL Substrate (Bio-Rad). Actin (Sigma A5060) and ponceau stain were controls.

**Tongue xenograft cancer model.** An orthotopic xenograft tongue cancer model was created by injecting HSC-3 into the tongue (Lam et al., 2012). NU/J *Foxn1*<sup>tmu</sup> athymic mice were injected in the left lateral tongue under anesthesia [1 × 10<sup>5</sup> HSC-3 human tongue OSCC cells suspended in 20 μl vehicle (1:1 mixture of DMEM and Matrigel; Corning, reference #354234), or vehicle alone]. After two weeks, the resulting xenografted tumors and vehicle-injected tongues were excised and snap frozen for protein analysis as above. Goat anti-mouse Lgmn (R & D AF2058) and donkey anti-goat IR-800 (Li-Cor) were used in the immunoblot.

**Paw xenograft cancer model.** The plantar surface of the right hind paw of NU/J *Foxn1*<sup>tmu</sup> athymic mice were inoculated with 1 × 10<sup>5</sup> HSC-3 in 20 μl of DMEM and Matrigel (Ye et al., 2011, 2014a). The paw xenograft model permits measurement of mechanical and heat hypersensitivity of the paw. By 14 d after inoculation, a visible tumor developed in the paw. After measuring baseline mechanical and thermal withdrawal thresholds, HSC-3 were inoculated into the hind paw. Mechanical and thermal withdrawal were measured at post inoculation days 3, 6, 10, and 13. On post inoculation day 14, LI-1 (10 mM, 100 μl) was injected into the tail vein. Mechanical and thermal withdrawal were measured at 1, 3, 6, 12, 24, and 48 h after injection of LI-1 into the paw cancer mouse model.

**4-Nitroquinoline 1-oxide (4NQO)-induced OSCC model.** An OSCC mouse model was generated by exposing mice to 4NQO (100 μg/ml) in drinking water for 16 weeks (Lam et al., 2012). Functional allodynia (gnaw-time) was measured with dolognawmeters (Dolan et al., 2010). Before administration of 4NQO, mice were examined to confirm the absence of oral abnormalities. 4NQO administration and dolognawmeter training over 15 sessions overlapped; baseline gnaw-time was calculated from the final 5 sessions. Functional allodynia was measured after 28 weeks. The tongue was harvested and a 1- to 2-mm coronal section was dissected from the most clinically suspicious region, fixed in 10% neutral buffer formalin, and processed for paraffin embedding and slide preparation. Four 5-μm hematoxylin and eosin (H & E)-stained tongue sections were evaluated for OSCC. Two pathologists blinded to group identity performed histopathologic analysis. Only mice with histologically confirmed OSCC were included in the analysis of nociception.

**Mechanical and thermal nociception in the hind paw.** To assess mechanical nociception, mice were placed on a platform with a metal mesh floor and acclimated for 1 h. Paw withdrawal threshold was measured

with von Frey filaments (Stoelting; Pickering et al., 2008). Withdrawal threshold was defined as the gram-force sufficient to elicit left hindpaw withdrawal. Withdrawal threshold for each animal was determined as the mean of three trials for each animal. Thermal hyperalgesia was measured with a paw thermal stimulator (IITC Life Sciences; Yamano et al., 2017). Mice were placed in a plastic chamber on a 25°C glass surface. A radiant heat source was focused on the left hind paw and withdrawal latency was measured as the mean of three trials taken at least 5 min apart in each mouse. The cutoff latency was established at 20 s. Lgmn (300 ng in 20 μl) or vehicle (control in 20 μl) was administered by intraplantar injection into the left hind paw under 1% isoflurane. Injections were made 1 h before the withdrawal tests (days 0, 1, and 4).

**Facial mechanical nociception.** Mice were placed individually in a transparent, mesh-floor, box and acclimated for 1 h every other day for two weeks. We measured withdrawal responses to mechanical stimulation of the left cheek with von Frey filaments ranging from 0.008 to 4 g-force (11 filaments in total) in ascending order (Deseure et al., 2003). We applied the von Frey filament to the cheek, defined by the area below the eye, between the nose and the ear. Each fiber was applied once; however, if the response to a von Frey filament was equivocal or the mouse was moving, the same von Frey filament was reapplied to the same area of the cheek 10 s after the first stimulus, or when the mouse stopped moving. The interval between applications of von Frey filaments of different intensities was 5 min. The facial nociception score was reported as a numerical average of the 11 responses in the following response categories: 0: no response; 1: detection, the mouse is aware of the filament that stimulates the face; the mouse turns its head slightly to the object; 2: reaction, the mouse turn its head away quickly, pulls it backward or reacts with a single face wipe; 3: escape/attack, the mouse quickly escapes from the object, attacks the object with its paw or mouth, or reacts with two facial swipes; 4: multiple facial grooming, the mouse responds to the filament stimulation with more than three facial wipes continuously. Hair on the left cheek was removed before subcutaneous injections of Lgmn. The whiskers were not trimmed. Lgmn (300 ng in 20 μl) was injected subcutaneously to the left cheek under 1% isoflurane. Injections were made 1 h before the facial mechanical withdrawal test at days 0, 1, and 4.

**Orofacial behavior.** The dolognawmeter quantifies a behavioral index of orofacial nociception (Dolan et al., 2010). The device measures the time taken to gnaw through a dowel and is a validated index of orofacial nociception in mice with OSCC. Mice were trained for 15 training sessions in the dolognawmeter or until the coefficient of variance of the time required to gnaw was below 0.2. A baseline gnaw-time (mean of the final five training sessions) was established for each mouse. After baseline gnaw-times were determined, treatment or drug injections were initiated and the mice underwent behavioral testing. Each response was analyzed relative to the mouse's baseline. Activated Lgmn, 300 ng in 20 μl, was injected into the tongue under isoflurane. The injection was performed at days 0, 1, and 4. One hour after injection, the mice were tested with a dolognawmeter.

**Quantification of Lgmn in OSCC cells.** Lgmn was measured in HSC-3 and DOK cells by ELISA. HSC-3 or DOK cells (~5000/well of a 12-well plate) were cultured for 72 h (~70% confluency; Lam et al., 2012). Medium was removed, cells were washed with 5 ml PBS without Ca<sup>2+</sup> and Mg<sup>2+</sup>, and DMEM (500 μl) was added to each well. After 48 h, medium was collected and centrifuged (1500 rpm, 4 min, 4°C). Cells were homogenized with 100 μl of RIPA buffer/well (Thermo Scientific, product #89901). Cell lysate was collected and centrifuged (1500 rpm, 10 min, 4°C). The pellet was discarded. A RayBio Human Lgmn ELISA kit (RayBiotech, code ELH-Lgmn-1) was used for Lgmn quantification. The standard curve was generated using the following concentrations: 7000, 2800, 1120, 448, 179, 72, 29, and 0 pg/ml of Lgmn provided with the kit. The optical densities of the standards and samples were read at 450 nm wavelength using a Promega GloMax luminometer (Promega BioSystems, Model E9032). Alternatively, cells were live-labeled with LE28 (1 μM, 0.1% DMSO) for 4 h, lysed on ice in PBS containing 0.1% Triton X-100, and cleared by centrifugation. Supernatants collected overnight in serum-free media were concentrated using an Amicon Filter with a 3-kDa cutoff. Total

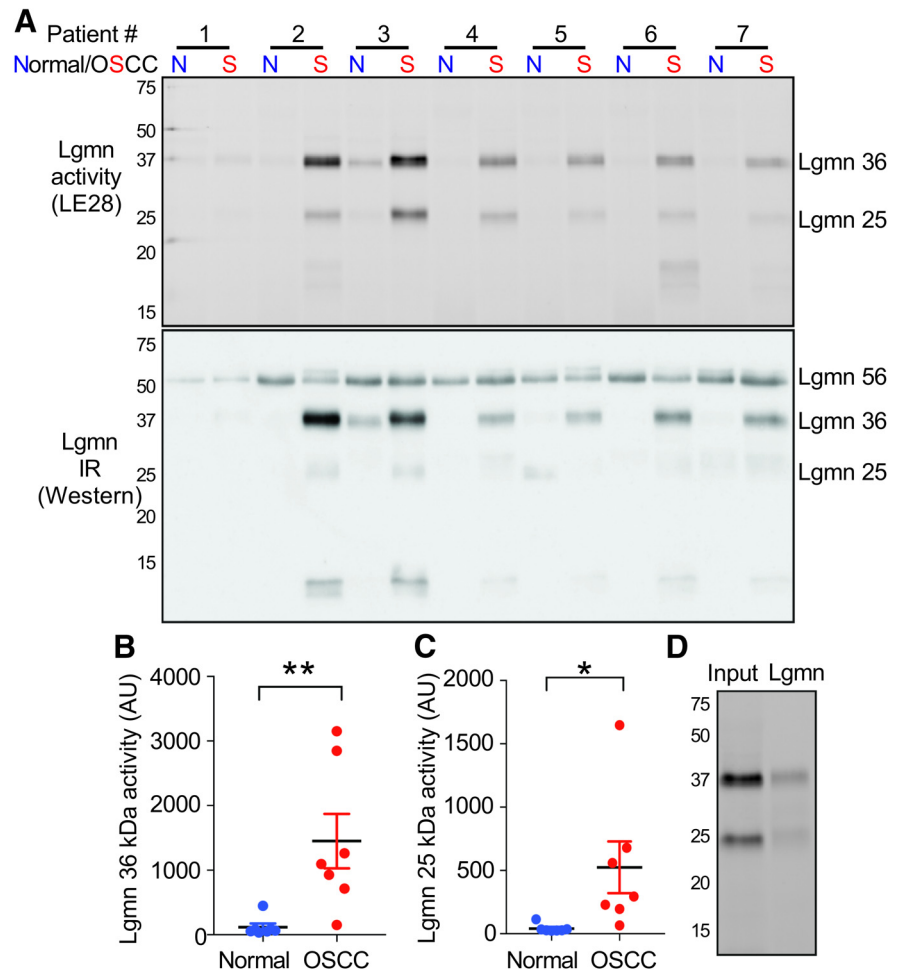


protein from whole-cell lysates or supernatants (~60 µg) were resolved by SDS-PAGE. Gels were scanned for Cy5 fluorescence and subject to Lgmn immunoblotting.

**Lgmn immunofluorescence in cancer cells.** HSC-3 and DOK cells were grown on cover slips at 37°C, 5% CO<sub>2</sub> for 48 h. Cells were washed with PBS and fixed in 4% paraformaldehyde at room temperature for 15 min. Cells were incubated with 3% bovine serum albumin (BSA) in PBS to block non-specific binding, then incubated with mouse monoclonal anti-Lgmn antibody (Santa Cruz Biotechnology, B-8: sc-133234, lot #A0610), 1:50, 4°C, overnight. Cells were washed in PBS and then incubated with goat anti-mouse secondary antibody conjugated to Alexa Fluor 488 (Life Technology, A11029), 1:300, room temperature for 3.5 h. Nuclei were stained with Hoechst (Thermo Scientific, product #62249, lot #RG2244203). The cover slips were mounted on slides in Fluoromount G (Electron Microscope Sciences). A laser scanning confocal microscope (LSM 700, Carl Zeiss) was used to obtain fluorescent images. The images were captured with a Zeiss Plan-Apochromat 63×/1.40 Oil DIC M27 objective lens for Lgmn signal quantification or 20× DIC objective lens for capturing the representative images. The fluorescent signal intensity of each cell was measured by a blinded researcher using NIH ImageJ. Controls included the following: (1) staining of the spleen from wild-type (WT) and *Lgmn*<sup>-/-</sup> mice; (2) preabsorption of the primary anti-Lgmn antibody with Lgmn; and (3) omission of the primary antibody. For the preabsorption negative control, the primary anti-Lgmn antibody (0.07 nM, equivalent to 1:50 dilution) was incubated in 10× higher concentration of activated Lgmn (0.7 nM) at 37°C for 48 h versus cells stained with the primary anti-Lgmn antibody, which was incubated in activated Lgmn vehicle. WT and *Lgmn*<sup>-/-</sup> mice were anesthetized with 100 mg/kg ketamine and 10 mg/kg xylazine (intraperitoneal) and transcardially perfused with 25 ml cold PBS, followed by 25 ml of 10% neutral formalin solution. The spleen was postfixed in 10% neutral formalin solution for 24 h, cryoprotected in 30% (v/v) sucrose in PBS for 2 d at 4°C, and embedded in Tissue-Tek<sup>R</sup> optimum cutting temperature (OCT) compound (Sakura Finetek). Sections (10 µm) were processed for immunofluorescence staining.

**Dissociation of TG neurons.** Mice were anesthetized with isoflurane, bilateral TG were removed and neurons were enzymatically dispersed as described (Ono et al., 2015). Ganglia were cut into 8–10 segments and incubated for 30 min in collagenase and dispase (12 mg collagenase, 14 mg dispase in 3 ml HBSS). Neurons were triturated and plated onto laminin/poly-L-ornithine-coated coverslips. Neurons were cultured in Leibovitz medium containing 10% fetal calf serum, with penicillin and streptomycin for 24 h (37°C, 5% CO<sub>2</sub>).

**Electrophysiological recording.** Hyperexcitability of small TG neurons (<20 µm, <30 pF) was quantified by measuring rheobase (Scheff et al., 2018). Whole-cell patch-clamp recordings were made using Axon patch 200B amplifier (Molecular Devices). Digidata 1440A (Molecular Devices) was used for data acquisition and pulse generation. Rheobase was measured using 250 ms square pulses starting from -10 pA with steps of 10 pA until the action potential threshold was reached. The resting membrane potentials were recorded at stable conditions without



**Figure 1.** Lgmn activation in human OSCC. **A**, Active Lgmn labeled by LE28 (upper panel) as shown by in-gel fluorescence and total Lgmn immunoreactivity (IR; lower panel) as shown by Western blotting in OSCC biopsies and patient-matched normal oral mucosa. The gel (upper panel) was transferred to nitrocellulose and immunoblotted for total Lgmn levels (lower panel). **B**, Densitometry of the 36-kDa species labeled by LE28, displayed as averages for all normal and OSCC samples ( $t_{(12)} = 3.124$ ,  $**p = 0.0088$ , when 36-kDa Lgmn activity in matched normal oral mucosa and SCC is compared, unpaired Student's *t* test). **C**, Densitometry of the 25-kDa species labeled by LE28, displayed as averages for all normal and OSCC samples ( $t_{(12)} = 2.367$ ,  $*p = 0.0356$ , when 25-kDa Lgmn activity in matched normal oral mucosa and SCC is compared, unpaired Student's *t* test). **D**, Immunoprecipitation of LE28-labeled tumor sample with a Lgmn-specific antibody to confirm the 36- and 25-kDa species ( $n = 7$ ).

current injection. Neurons with resting membrane potentials more positive than -40 mV were excluded from analysis. Input resistance was estimated from current-clamp recordings of the voltage response to 250 ms of 10 pA hyperpolarizing current. Pipette resistance was 4–5 mΩ in the following external solution: 140 mM NaCl, 5 mM KCl, 10 mM HEPES, 10 mM glucose, 1 mM MgCl<sub>2</sub>, and 2 mM CaCl<sub>2</sub>; pH was adjusted to 7.3–7.4 with NaOH. Pipette solution consisted of the following: 110 mM K-glucuronate, 30 mM KCl, 10 mM HEPES, 1 mM MgCl<sub>2</sub>, and 2 mM CaCl<sub>2</sub>; pH was adjusted to 7.25 with KOH. Rheobase was measured after 10 min incubation with Lgmn (20 ng/ml in external solution, pH 5.5), Lgmn (20 ng/ml) plus LI-1 (10 µM), or Lgmn vehicle (Lgmn activation buffer pH 5.5). To study the signaling pathways of Lgmn induced hyperexcitability, perforated patch-clamp was performed to avoid dialysis of cytoplasmic constituents. Amphotericin B (240 µg/ml) was used in the pipette solution. Neurons were preincubated with GF109203X (GFX, Tocris) or PKI-tide (both 1 µM, 30 min, 37°C) before challenge with Lgmn (20 ng/ml) or Lgmn vehicle. Rheobase was measured 10 min after incubation with Lgmn or vehicle.

**Cell lines.** Human embryonic kidney (HEK293) cells stably expressing the human (h)PAR<sub>2</sub> with extracellular N-terminal FLAG and intracellular C-terminal HA11 epitopes (HEK-FLAG-PAR<sub>2</sub>-HA cells) have been described (Böhm et al., 1996b). PAR<sub>1</sub> and PAR<sub>2</sub> were deleted from

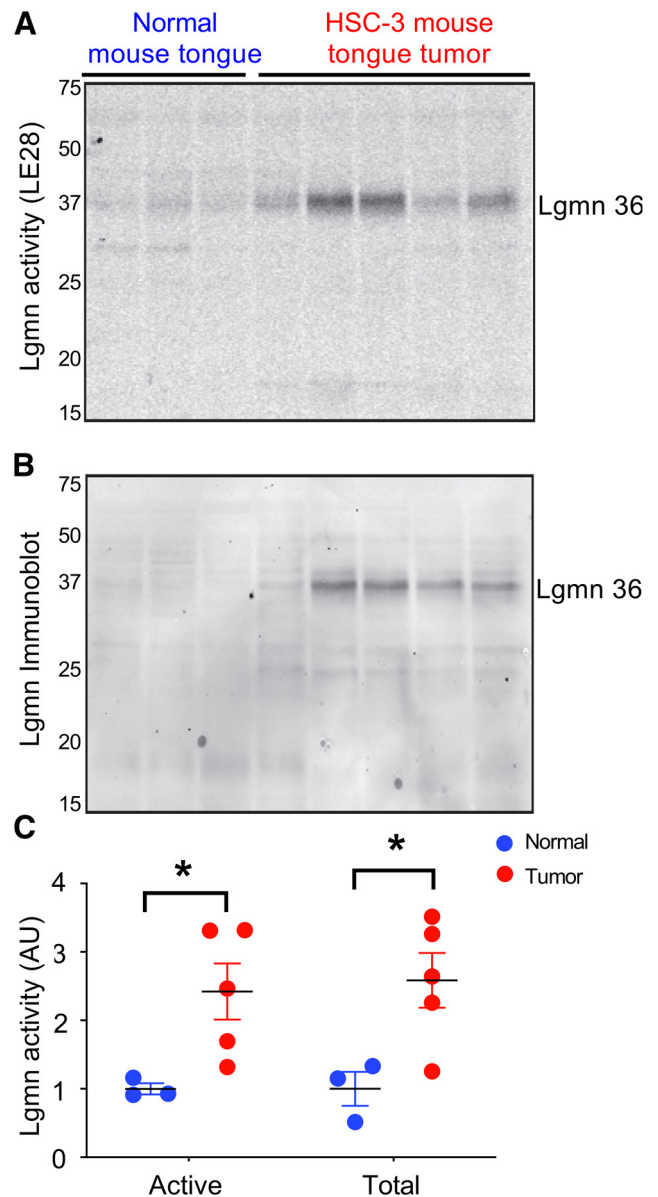
HEK293 cells using CRISPR/Cas9 (Ungefroren et al., 2017). Cells were maintained in DMEM with 10% fetal bovine serum and hygromycin (100 µg/ml, 5% CO<sub>2</sub>, 37°C). Dysplastic oral keratinocyte, DOK, cell number 94122104 from Sigma-Aldrich was cultured in DMEM/F12 (Invitrogen), 10% fetal bovine serum, 50 IU/ml penicillin/streptomycin and 5 µg/ml hydrocortisone. Human OSCC, HSC-3, cell number JCRB0623, was from Japanese Collection of Research Bioresources Cell Bank. HSC-3 was cultured in DMEM, 10% fetal bovine serum, and 50 IU/ml penicillin/streptomycin. DOK and HSC-3 cell lines were maintained at 37°C with 5% CO<sub>2</sub>.

**PAR<sub>2</sub> cleavage.** A peptide corresponding to hPAR<sub>2</sub> amino acids 21–50 was synthesized by American Peptide Company and dissolved in water at 300 µM. The peptide was diluted to 200 µM with Lgmn activation buffer (50 mM sodium acetate, 100 mM NaCl, pH 4.5) in the presence and absence of Lgmn (200 nM; final volume 25 µl). After overnight incubation at 37°C, the reactions were quenched with 25 µl of 50% acetonitrile containing 0.1% trifluoroacetic acid (TFA). Samples (2 µl) were subject to LCMS analysis with a Shimadzu LCMS 2020 fitted with a Phenomenex Luna 3 µm C8(2) column (100 Å, 100 × 2 mm). A gradient of 0% – 60% acetonitrile over 10 min with 0.05% TFA was used for separation.

**Immunofluorescence in HEK cells.** HEK-FLAG-PAR<sub>2</sub>-HA cells (45,000) were plated in eight-well ibiTreat µ-slides and incubated overnight. Cells were washed with HBSS and incubated with Lgmn (100 nM final in HBSS pH 5.5) or trypsin (10 nM final in HBSS pH 7.4; 1 h 37°C). Cells were fixed in 4% paraformaldehyde on ice for 10 min. Cells were incubated with blocking buffer [3% normal horse serum (NHS) and 0.1% saponin] for 30 min at RT. Primary antibodies in blocking buffer [rabbit anti-FLAG (1:250, Rockland) and mouse anti-HA (1:250; Ray Biotech)] were incubated overnight at 4°C. Cells were washed with PBS and secondary antibodies in blocking buffer (donkey anti-rabbit-Alexa Fluor 488 and donkey anti-mouse-Alexa Fluor 647; 1:500; ThermoFisher) were added for 1 h at RT. After washing with PBS, DAPI was added for 5 min followed by additional washing. Cells were imaged immediately on a Leica SP8 confocal microscope.

**On-cell Westerns.** HEK-FLAG-PAR<sub>2</sub>-HA cells were plated on poly-D-lysine (PDL)-coated 96-well plates (30,000 cells/well) and incubated overnight. Cells were washed two times in HBSS (pH 7.4) and placed in MES-HBSS (pH 5.0) for Lgmn assays or HBSS (pH 7.4) for trypsin assays. Cells were incubated with Lgmn (1 or 10 ng/µl final concentration) in MES-HBSS (pH 5.0), trypsin (10 nM final concentration) in HBSS (pH 7.4), or vehicle (buffer control) for 30 min at 37°C. Cells were washed with HBSS and fixed with 4% paraformaldehyde in PBS for 20 min on ice. Cells were washed three times in PBS and incubated with blocking buffer (PBS + 3% NHS) for 1 h at room temperature. Cells were incubated with mouse anti-FLAG antibody (1:500, Cell Signaling) in PBS + 1% NHS overnight at 4°C. Cells were washed three times in PBS, incubated with donkey anti-mouse Alexa Fluor 790 (1:1000, A11371, ThermoFisher) in PBS + 1% NHS for 1 h at room temperature. Cells were washed 1 time in PBS and incubated with the nuclear stain SYTO 82 Orange (1 µM, ThermoFisher) in saline for 30 min. Cells were washed three times with saline and then imaged on a GE HealthcareTyphoon imaging system (GE). FLAG immunofluorescence intensity was quantified using NIH ImageJ and was normalized to nuclear fluorescent intensity to correct possible cell loss.

**Measurement of intracellular Ca<sup>2+</sup>.** HEK-FLAG-PAR<sub>2</sub>-HA cells were plated on PDL-coated 96-well plates (25,000 cells/well) and incubated overnight. Cells were loaded with fura-2 AM (1 µM, Cayman Chemicals) in loading buffer (150 mM NaCl, 2.6 mM KCl, 0.1 mM CaCl<sub>2</sub>, 1.18 mM MgCl<sub>2</sub>, 10 mM D-glucose, 10 mM HEPES, 4 mM probenecid, 0.5% BSA, pH 7.4) for 1 h at 37°C. Cells were washed 2 times in HBSS and then placed in MES-HBSS (pH 5.0, 5.5, 6.0, or 7.4) for the Lgmn assays or HBSS (pH 7.4) for trypsin assays. Fluorescence was measured with 340 or 380 nm excitation and 530 nm emission with a Flexstation three plate reader. Baseline fluorescence was measured for 45 s (Zhao et al., 2015). Cells were challenged with Lgmn (1 or 10 ng/µl final concentration) in MES-HBSS (pH 5.0, 5.5, 6.0, or 7.4), trypsin (10 nM final concentration) in HBSS (pH 7.4), or vehicle (buffer control), and fluorescence was measured for an



**Figure 2.** Lgmn in xenograft model of OSCC. **A, B**, Active Lgmn labeled by LE28 (**A**), shown by in-gel fluorescence, and total Lgmn immunoreactivity (IR; **B**), shown by Western blotting of lysates from HSC-3 xenografts or control tongues. The gel from (**A**) was transferred to nitrocellulose and immunoblotted for total Lgmn levels (**B**). **C**, Densitometry of active and total Lgmn from **A, B**, respectively ( $n = 3-5$ ,  $t_{(6)} = 2.592$ ,  $*p = 0.0411$ , when active Lgmn in normal mouse tongue and xenograft cancer is compared, and  $t_{(6)} = 2.818$ ,  $*p = 0.0304$ , when total Lgmn in normal mouse tongue xenograft cancer is compared, unpaired Student's *t* tests).

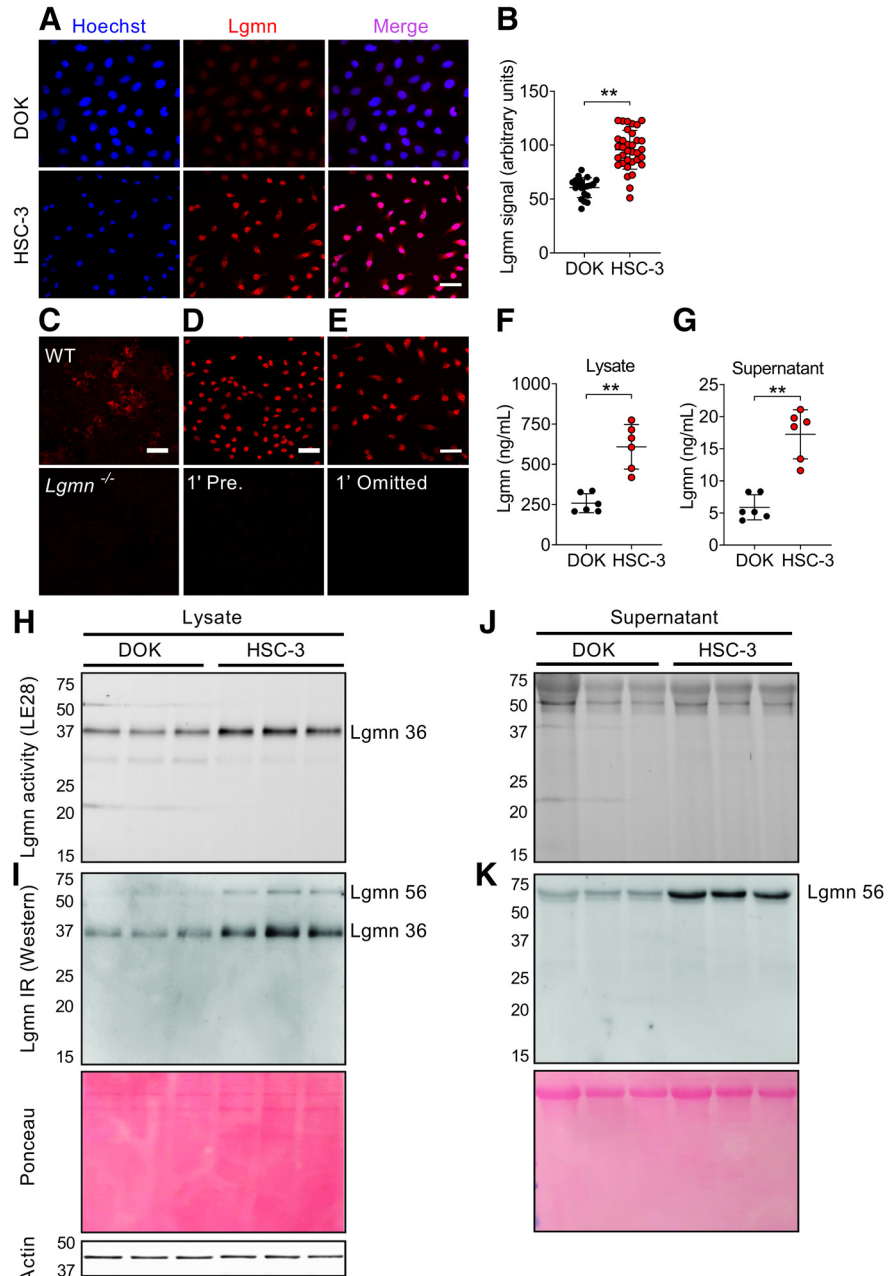
additional 180 s. To confirm that Lgmn-induced Ca<sup>2+</sup> responses were because of activation of PAR<sub>2</sub>, cells were incubated with the PAR<sub>2</sub> antagonists I-343 (10 µM) or GB88 (10 µM; Farmer, 2013; Lieu et al., 2016; Jimenez-Vargas et al., 2018), the PAR<sub>1</sub> antagonist SCH79797 (200 nM; Ahn et al., 2000), or vehicle in HBSS + 1% DMSO for 1 h at 37°C before the Ca<sup>2+</sup> assay. To confirm the requirement for protease activity, Lgmn was incubated with the Lgmn inhibitors QDD100531 (1 µM) or QDD123427 (100 nM; Ness et al., 2015) in HBSS + 1% DMSO for 1 h before the Ca<sup>2+</sup> assay. Cells were maintained with the inhibitors. To assess the requirement for hydrolysis of the Asn<sup>30</sup>↓Arg<sup>31</sup> site, a mutant PAR<sub>2</sub> in which Asn<sup>30</sup> was replaced with Ala (PAR<sub>2</sub>-ΔN30A) was generated (Twist Biosciences, San Francisco, CA). cDNA (5 µg) encoding PAR<sub>2</sub>-ΔN30A was expressed in HEK-PAR<sub>2</sub>-KO cells using polyethylenimine (ratio 1:6, DNA:PEI), and cells were studied after 48 h. To assess the capacity of Lgmn or trypsin to

desensitize PAR<sub>2</sub>-mediated Ca<sup>2+</sup> signaling, HEK-FLAG-PAR<sub>2</sub>-HA cells were preincubated with Lgmn (1 or 10 ng/μl final concentration) in MES-HBSS (pH 5.0), trypsin (10 nM final concentration) in HBSS (pH 7.4), or vehicle (buffer control) for 10 min at 37°C. Cells were washed and recovered in HBSS (pH 7.4) for 20 min at 37°C. Cells were then challenged with trypsin (10 nM). Desensitization of Ca<sup>2+</sup> signals to the second challenge with trypsin was calculated as a percentage of the responses in cells preincubated with the vehicle control.

**FRET assays of cAMP, protein kinase D (PKD), and ERK.** Genetically encoded FRET biosensors targeted to the cytosol were used to assess cAMP, PKD and ERK activation in living cells in real time (Zhao et al., 2015). HEK-FLAG-PAR<sub>2</sub>-HA cells were transfected with cDNA (5 μg) encoding the cAMP biosensor Cyto-EPAC (Jimenez-Vargas et al., 2018), the PKD biosensor Cyto-DKAR (Zhao et al., 2019), or the ERK biosensor Cyto-EKAR (Yarwood et al., 2017) using polyethylenimine (ratio 1:6, DNA:PEI). Cells were plated on PDL-coated 96-well plates (25,000 cells/well) and incubated overnight. Cells were washed 2× in HBSS and placed in MES-HBSS (pH 5.0) for the Lgmn assays. The cyan (470 nm) and yellow fluorescent protein (535 nm) emission ratios were measured with a CLARIOstar<sup>plus</sup> plate reader (BMG). Baseline fluorescent ratios were recorded for 5 min. Cells were challenged with Lgmn (1 or 10 ng/μl) in MES-HBSS (pH 5.0) or vehicle (buffer control), and FRET was measured for an additional 30 min. FRET ratios were normalized to the MES-HBSS vehicle control.

**BRET assays of β-arrestin recruitment.** HEK293 cells were transfected with cDNA encoding the PAR<sub>2</sub>-RLuc8 (1 μg) and β-arrestin-1-YFP (4 μg) with polyethylenimine (ratio 1:6, DNA:PEI; Jensen et al., 2013). Cells were plated on PDL-coated 96-well white walled plates (30,000 cells/well) and incubated overnight. Cells were washed 2× in HBSS and placed in MES-HBSS (pH 5.0) for the Lgmn assays or HBSS (pH 7.4) for trypsin assays. Coelenterazine-h (5 μM, Nanolight, Pinetop AZ) was added to the cells and the cells were challenged with Lgmn (1 or 10 ng/μl) or trypsin (10 nM). RLuc8 luminescence (480 nm) and YFP fluorescence (530 nm) emission were measured using a CLARIOstar<sup>plus</sup> plate reader. Baseline fluorescence ratios were recorded for 2.5 min. The BRET ratio was normalized to vehicle control and baseline.

**Experimental design and statistical analysis.** We used GraphPad Prism 7 and 8 (GraphPad Prism, GraphPad Software) for the statistical analysis. Results are expressed as mean ± SEM. For cell-based assays, triplicate measurements were made from four to five experiments; differences were evaluated by one-way or two-way ANOVA and Dunnett's or Tukey's multiple comparisons test. One-way ANOVA and Tukey's or Sidak's multiple comparisons and Student's *t* test were used for *in vivo* behavioral experiments and comparison of rheobase. Unpaired Student's *t* test was used to compare values between two groups.



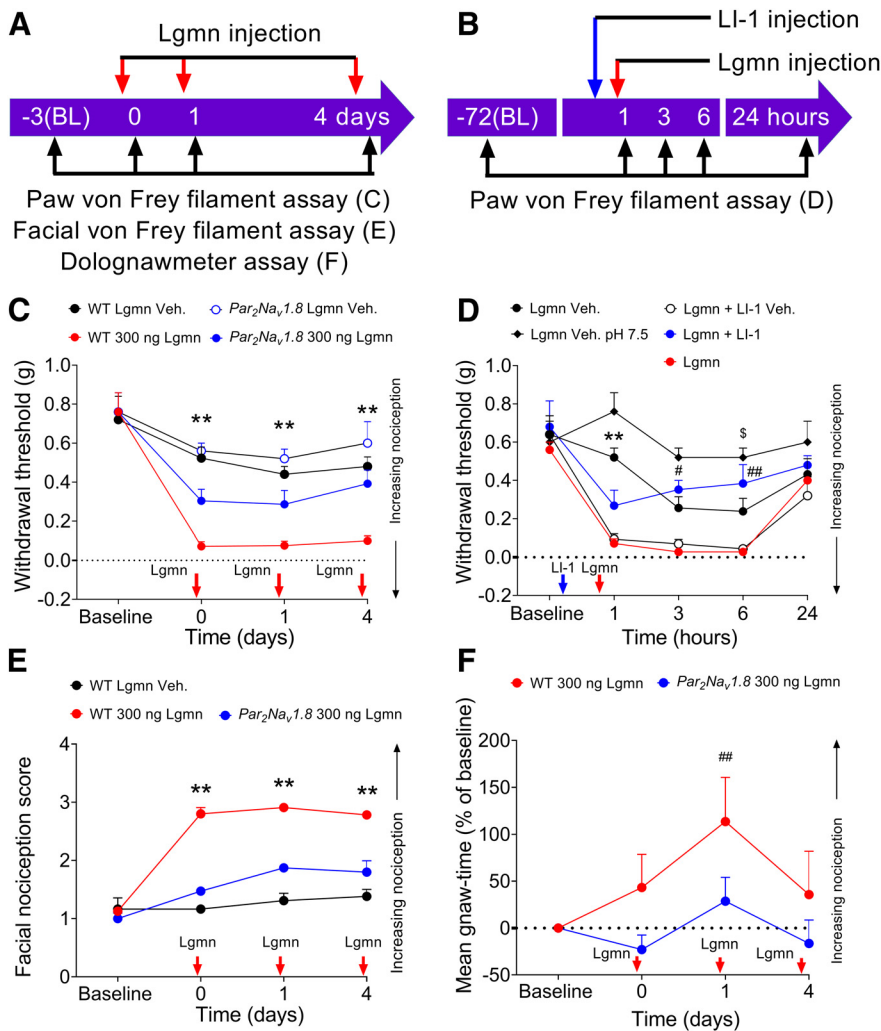
**Figure 3.** Expression of Lgmn in OSCC cells. **A**, Localization of immunoreactive Lgmn (red) in DOK and HSC-3. **B**, Lgmn signal intensity in DOK and HSC-3 was quantified in individual cells by NIH ImageJ ( $t_{(52)} = 8.11$ ,  $**p = 8.53E-11$ , when Lgmn signal of DOK is compared with HSC-3,  $n = 20$  and 34 cells in DOK and HSC-3, respectively, unpaired Student's *t* test). **C**, Localization of immunoreactive Lgmn in the spleen of WT and *Lgmn*<sup>-/-</sup> mice. **D**, Preabsorption of Lgmn antibody with Lgmn eliminated HSC-3 staining. **E**, Omission of the anti-Lgmn antibody resulted in lack of HSC-3 staining. **F**, **G**, Quantification of Lgmn protein by ELISA in DOK and HSC-3 cell lysate and supernatant. The concentration of Lgmn in HSC-3 cell lysate and supernatant was three times higher than that of DOK.  $N = 6$  experiments in each group (in **F**,  $t_{(10)} = 5.70$ ,  $**p = 0.0002$ , when the Lgmn concentration in supernatant from HSC-3 is compared with the Lgmn protein concentration in supernatant from DOK, unpaired Student's *t* test. In **G**,  $t_{(10)} = 6.49$ ,  $**p = 0.000069$ , when the Lgmn concentration in lysate from HSC-3 is compared with the Lgmn concentration in lysate from DOK, unpaired Student's *t* test). Scale bar in **A**, **C**, **D**, **E**: 50 μm. **H**, Labeling of active Lgmn with LE28 in DOK and HSC-3 cell lysate, as shown by in-gel fluorescence. **I**, Lgmn immunoblot in DOK and HSC-3 cell lysate in **H**, with ponceau stain and actin immunoblot to verify equal loading. **J**, Labeling of Lgmn with LE28 in DOK and HSC-3 cell supernatant, as shown by in-gel fluorescence. **K**, Immunoblot of Lgmn from DOK and HSC-3 cell supernatant.

## Results

### Lgmn is activated in human and murine OSCC

To determine whether Lgmn is activated in OSCCs, we collected OSCC specimens and matched normal oral mucosa from seven





**Figure 4.** Lgmn-evoked nociception. **A**, Experiment timeline to test the effect of Lgmn (red arrow) on paw, facial mechanical nociception, and oral function on WT and *Par2Nav1.8* mice. **B**, Experiment timeline to test the effect of Lgmn (red arrow) and LI-1 (blue arrow) on paw mechanical nociception on WT mice. **C**, Effects of Lgmn or vehicle (Veh) on paw withdrawal in WT and *Par2Nav1.8* mice. Arrows indicate time of Lgmn or vehicle injection. Relative to WT Veh. mice, withdrawal thresholds were significantly lower in WT Lgmn mice, but not *Par2Nav1.8* mice, at days 0, 1, and 4 (interaction  $F_{(9,64)} = 2.47, p = 0.02$ ;  $**p = 0.0002$  at d0,  $**p = 0.0033$  at d1,  $**p = 0.002$  at d4, respectively, when WT mice treated with 300 ng Lgmn are compared with WT mice treated with Lgmn vehicle,  $n = 5$  in each group, two-way ANOVA, Tukey's multiple comparisons). **D**, Effects of LI-1 on Lgmn-evoked mechanical allodynia in WT mice. Blue arrow indicates time of LI-1 or vehicle administration. Red arrow indicates time of Lgmn or Lgmn vehicle injection. Withdrawal thresholds were measured at 1, 3, 6, and 24 h after Lgmn injection. The effect of Lgmn varied with time (interaction  $F_{(16,100)} = 2.98, p = 0.005$ , two-way ANOVA,  $n = 5$  in each group, Tukey's multiple comparisons). One hour after injection of LI-1, mean withdrawal threshold was lower in Lgmn versus Lgmn vehicle ( $**p = 0.0002$ ). LI-1 prevented the nociceptive effect of Lgmn at 3 and 6 h after Lgmn injection ( $#p = 0.04$  and  $##p = 0.009$ , respectively, when Lgmn plus LI-1 is compared with Lgmn plus LI-1 vehicle), but not 1 h after LI-1 injection ( $p = 0.30$ ). Lgmn Veh. at pH 5 reduced the withdrawal threshold more than Lgmn Veh. at pH 7.5 at 6 h after paw injection ( $F_{(4,100)} = 24.5, ^5p = 0.04$ , when Lgmn Veh. at pH 5.0 is compared with Lgmn Veh. at pH 7.5, two-way ANOVA, Tukey's multiple comparisons). **E**, Lgmn (300 ng) was injected into the cheek at days 0, 1, and 4 following baseline facial withdrawal measurements. Arrows indicate Lgmn injection into the cheek. Lgmn induced facial mechanical allodynia in WT but not *Par2Nav1.8* mice. The means of the facial nociception score were significantly increased in WT mice versus *Par2Nav1.8* mice at days 0, 1, and 4 ( $F_{(3,36)} = 71.69, **p = 2.98E-15$  at d0, d1, and d4, when WT 300 ng Lgmn is compared with *Par2Nav1.8* 300 ng Lgmn,  $n = 5$  in each group, two-way ANOVA, Sidak's multiple comparisons). **F**, Lgmn (300 ng) significantly increased gnaw-time at 1 d after injection in WT mice versus baseline, but not in *Par2Nav1.8* mice ( $F_{(3,33)} = 3.26, ##p = 0.0045$  at d1, when WT 300 ng Lgmn is compared with baseline,  $n = 8$  and 6 in WT and *Par2Nav1.8* mice, respectively, two-way ANOVA, Tukey's multiple comparisons).

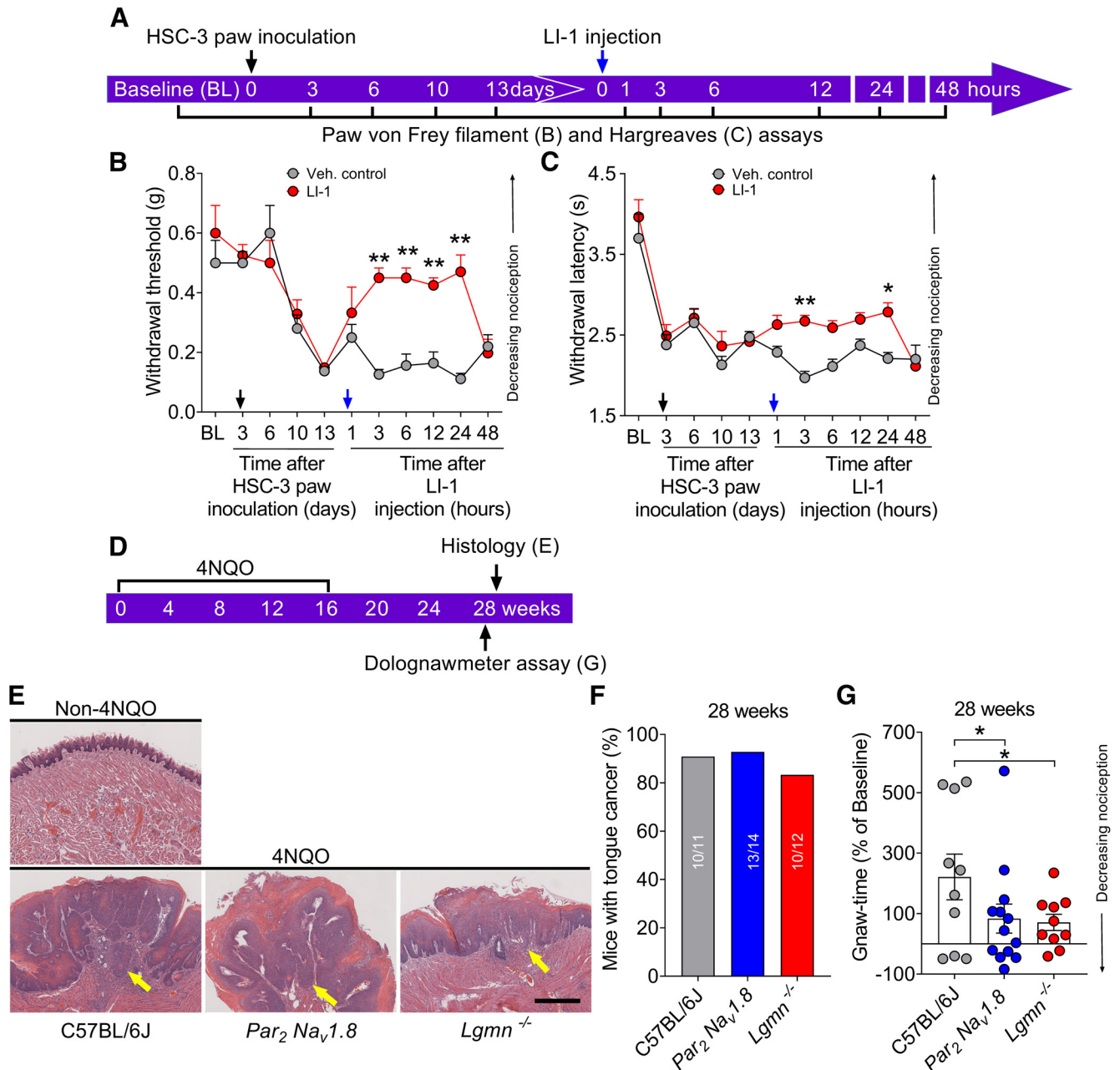
patients (Table 1). Specimens were incubated with a fluorescently quenched activity-based probe (LE28) selective for Lgmn (Edgington et al., 2013). Two LE28-labeled species of 36 kDa and 25 kDa were activated in OSCC versus normal tissue (Fig. 1A–D). Immunoprecipitation with a Lgmn-specific antibody confirmed

the identity of mature forms of Lgmn (Fig. 1D). Immunoblotting revealed total levels of mature Lgmn 36 kDa increased in all OSCC versus normal tissue (Fig. 1A). The 56 kDa inactive Lgmn zymogen (pro-Lgmn) was detected in all specimens. Total Lgmn and Lgmn activity of 36 kDa were up-regulated in a murine OSCC xenograft model (human OSCC cells (HSC-3) inoculated in nude mouse tongues) versus normal (Fig. 2A–C).

We examined Lgmn expression in HSC-3 and dysplastic oral keratinocytes (DOK, non-cancer cell line) by immunofluorescence and ELISA. Immunoreactive Lgmn was detected in cytosolic granules of HSC-3 and DOK cells, and expression was confirmed by ELISA of cell lysate and conditioned medium (supernatant; Fig. 3A,F,G). The intensity of Lgmn staining was higher in HSC-3 (Fig. 3B). Controls for the selectivity of the Lgmn antibody included absence of staining of spleen from *Lgmn*<sup>-/-</sup> mice (Fig. 3C), abolition of Lgmn staining by Lgmn antibody preabsorption with Lgmn (Fig. 3D), and lack of staining when the Lgmn antibody was omitted (Fig. 3E). The Lgmn protein concentration in lysate and supernatant from HSC-3 was higher than the concentration from DOK (Fig. 3F,G). The levels of Lgmn in lysate (Fig. 3H,I) and supernatant (Fig. 3J,K) were higher in HSC-3 compared with DOK. Immunoblotting revealed upregulation of Lgmn 56-kDa zymogen and Lgmn 36-kDa mature form in HSC-3 versus DOK (Fig. 3J). HSC-3 cells also secreted more pro-Lgmn than DOKs (Fig. 3J,K). These results suggest that OSCC cells express and release more Lgmn than dysplastic keratinocytes.

### PAR<sub>2</sub> expression on Nav1.8-positive nociceptors is necessary for Lgmn-induced mechanical allodynia

To investigate whether Lgmn causes allodynia by activating PAR<sub>2</sub> on nociceptors, we administered Lgmn (300 ng intraplantar injection) for 3 d (0, 1, 4 d) to WT (C57BL/6J) and *Par2Nav1.8* mice, which lack PAR<sub>2</sub> in Nav1.8-positive neurons. We measured paw withdrawal responses to stimulation of the plantar surface with von Frey filaments at baseline and 1 h after each Lgmn injection (Fig. 4A). Lgmn induced mechanical allodynia on all 3 d in WT mice; however, mechanical allodynia was attenuated in *Par2Nav1.8* mice by 51% on day 0

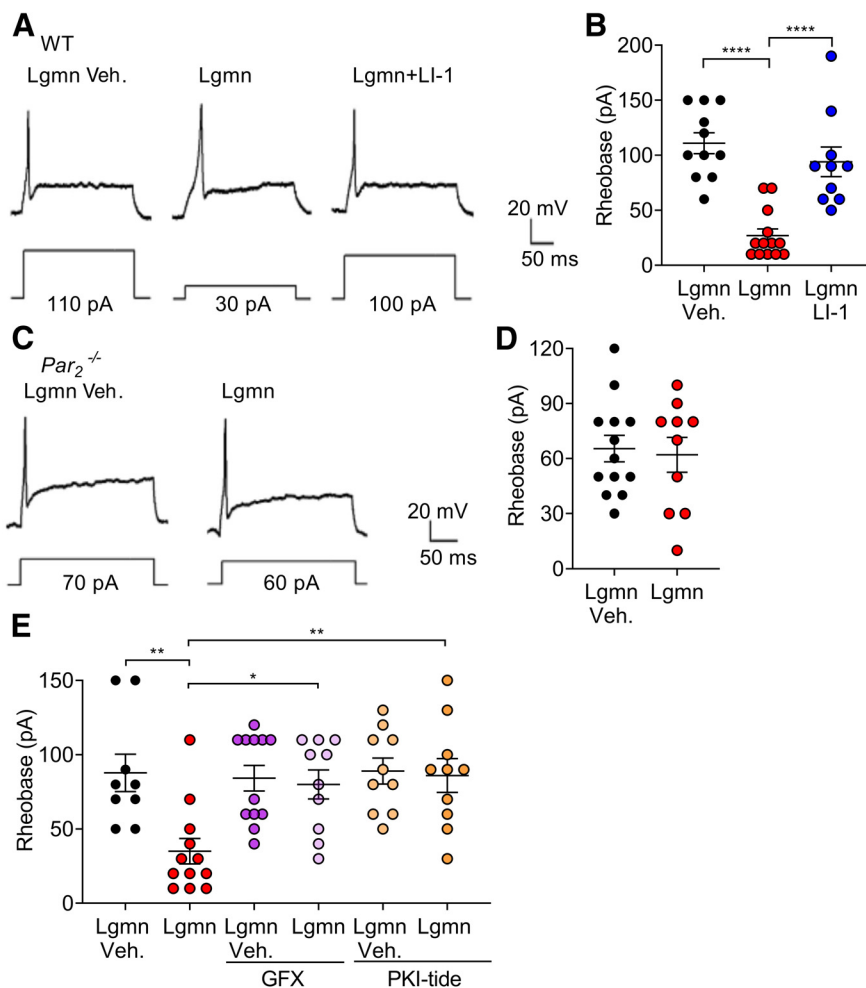


**Figure 5.** Contributions of Lgmn and PAR<sub>2</sub> to OSCC pain. **A**, The experimental protocol included baseline measurements of withdrawal threshold to mechanical stimulation and withdrawal latency to thermal stimulation, HSC-3 inoculation, injection of LI-1, and measurements of nociception. **B**, Mechanical allodynia. HSC-3 inoculation (black arrow) produced mechanical allodynia after 10 d. LI-1 (blue arrow) reversed cancer-induced mechanical nociception versus vehicle control after 3, 6, 12, and 24 h following injection but not after 48 h ( $F_{(1,77)} = 32.42$ ,  $**p = 0.0003$  at 3 h,  $**p = 0.0015$  at 6 h,  $**p = 0.0068$  at 12 h,  $**p = 2.15E-07$  at 24 h, when Veh. control is compared with LI-1,  $n = 8$  in each group, two-way ANOVA, Sidak's multiple comparisons). **C**, Thermal hyperalgesia. HSC-3 inoculation (black arrow) produced thermal hyperalgesia beginning at 3 d. LI-1 (blue arrow), versus vehicle control, reduced thermal hyperalgesia after 3 and 24 h but not after 48 h ( $F_{(10,140)} = 24.45$ ,  $**p = 0.0029$  at 3 h,  $*p = 0.0278$  at 24 h, when Veh. control is compared with LI-1,  $n = 8$  in each group, two-way ANOVA, Sidak's multiple comparisons). **D**, The experimental protocol included administration of 4NQO to the mice for 16 weeks, measurement of OSCC nociception (gnaw-time) with the dolognawmeter, and histologic analysis of the tongue to confirm cancer. **E**, Representative histologic images of the tongue from C57BL/6J mice that did not receive 4NQO, and of the tongues from C57BL/6J, *Par<sub>2</sub>Na<sub>v</sub>1.8*, and *Lgmn*<sup>-/-</sup> mice at 28 weeks following 4NQO administration. Arrows indicate tongue carcinoma. Scale bar: 100 μm. **F**, The percentage of C57BL/6J, *Par<sub>2</sub>Na<sub>v</sub>1.8*, and *Lgmn*<sup>-/-</sup> mice that developed tongue cancer at 28 weeks after 4NQO administration. **G**, Change of gnaw-time versus baseline (percentage change of gnaw-time at baseline was set as 0%; data not shown) of C57BL/6J ( $n = 10$ ), *Par<sub>2</sub>Na<sub>v</sub>1.8* ( $n = 13$ ), and *Lgmn*<sup>-/-</sup> ( $n = 10$ ) mice with 4NQO-induced tongue cancer ( $F_{(1,30)} = 16.28$ ,  $*p = 0.029$ , when C57BL/6J is compared with *Par<sub>2</sub>Na<sub>v</sub>1.8*,  $*p = 0.024$ , when C57BL/6J is compared with *Lgmn*<sup>-/-</sup>, two-way ANOVA with Tukey's multiple comparisons).

(Fig. 4C). To confirm that the nociceptive action of Lgmn required enzymatic activity and to test the analgesic potential of a Lgmn inhibitor, we administered the Lgmn-selective inhibitor, LI-1 (10 mM, 100 μl, i.v.; Edgington-Mitchell, 2016) to WT mice 120 min before intraplantar injection of Lgmn and measured paw

withdrawal at 1, 3, 6, and 24 h after Lgmn injection (Fig. 4B). LI-1 attenuated Lgmn-induced mechanical allodynia by 44% 1 h after Lgmn injection (Fig. 4D). Thus, Lgmn-induced mechanical allodynia in mice requires PAR<sub>2</sub> expression on Na<sub>v</sub>1.8-expressing nociceptors and Lgmn enzymatic activity.





**Figure 6.** PAR<sub>2</sub> mediated Lgmn-induced hyperexcitability in TG neurons. **A**, Representative raw traces of whole-cell patch-clamp recordings showing membrane potential response at rheobase of TG neurons from WT mice treated with Lgmn vehicle, Lgmn, and Lgmn + LI-1. **B**, Rheobase of TG neurons in different treatment groups. Lgmn vehicle:  $110.0 \pm 31.4$  pA,  $n = 11$ ; Lgmn:  $26.9 \pm 22.1$  pA,  $n = 13$ ; Lgmn + LI-1:  $94.0 \pm 42.4$  pA,  $n = 10$  ( $F_{(2,31)} = 23.14$ ,  $****p = 1.23E-6$ , when Lgmn and Lgmn vehicle are compared,  $****p = 6.93E-5$ , when Lgmn and Lgmn + LI-1 are compared, one-way ANOVA with Tukey's multiple comparisons). **C**, Representative raw traces of whole-cell patch-clamp recordings showing membrane potential response at rheobase of TG neurons from *Par2*<sup>-/-</sup> mice treated with Lgmn and Lgmn vehicle. **D**, Rheobase of TG neurons from *Par2*<sup>-/-</sup> mice. Lgmn vehicle:  $65.3 \pm 26.0$  pA,  $n = 13$ ; Lgmn:  $62.0 \pm 30.1$  pA,  $n = 10$ . **E**, Lgmn induced hyperexcitability and PKC-dependent or PKA-dependent pathways. Perforated patch-clamp recordings were used to measure rheobase of TG neurons. Neurons were preincubated with GFX  $1 \mu\text{M}$  and PKI-tide  $1 \mu\text{M}$  before Lgmn or Lgmn vehicle treatments. Rheobase was measured after neurons were challenged with Lgmn and Lgmn vehicle. Lgmn vehicle,  $87.7 \pm 12.5$  pA,  $n = 9$ ; Lgmn,  $35.6 \pm 8.5$  pA,  $n = 12$ ; GFX + Lgmn vehicle,  $84.1 \pm 8.5$  pA,  $n = 12$ ; GFX + Lgmn,  $80.0 \pm 9.7$  pA,  $n = 10$ ; PKI-tide + Lgmn vehicle,  $89.0 \pm 8.7$  pA,  $n = 10$ ; PKI-tide + Lgmn,  $86.0 \pm 11.3$  pA,  $n = 10$  ( $F_{(5,57)} = 4.93$ ,  $***p = 0.0052$ , when Lgmn and Lgmn vehicle are compared,  $*p = 0.0198$  when the Lgmn and Lgmn + GFX are compared,  $**p = 0.0198$ , when Lgmn and Lgmn + PKI-tide are compared, one-way ANOVA with Tukey's multiple comparisons).

### PAR<sub>2</sub> expression on Na<sub>v</sub>1.8-positive nociceptors is necessary for Lgmn-induced orofacial nociception

We used reflexive and operant assays to test whether Lgmn induces nociception in the orofacial region. We injected Lgmn into the cheek and measured facial withdrawal to stimulation with von Frey filaments. Lgmn (300 ng) was injected subcutaneously for 3 d (0, 1, 4 d) to WT and *Par2*<sup>-/-</sup> mice. Withdrawal was measured at baseline and 1 h after each injection (Fig. 4A). Lgmn induced facial mechanical allodynia on all 3 d in WT mice, but the nociceptive effect was attenuated in *Par2*<sup>-/-</sup> mice by 81% on day 0 (Fig. 4E). For operant behavioral testing, Lgmn (300 ng) was injected into the tongue. Dolognawmeters quantified a behavioral index of nociception 1 h after injection (Fig. 4A). Lgmn

induced orofacial dysfunction in WT but not *Par2*<sup>-/-</sup> mice (Fig. 4F). Thus, expression of PAR<sub>2</sub> in Na<sub>v</sub>1.8-expressing nociceptors is necessary for Lgmn-induced pain.

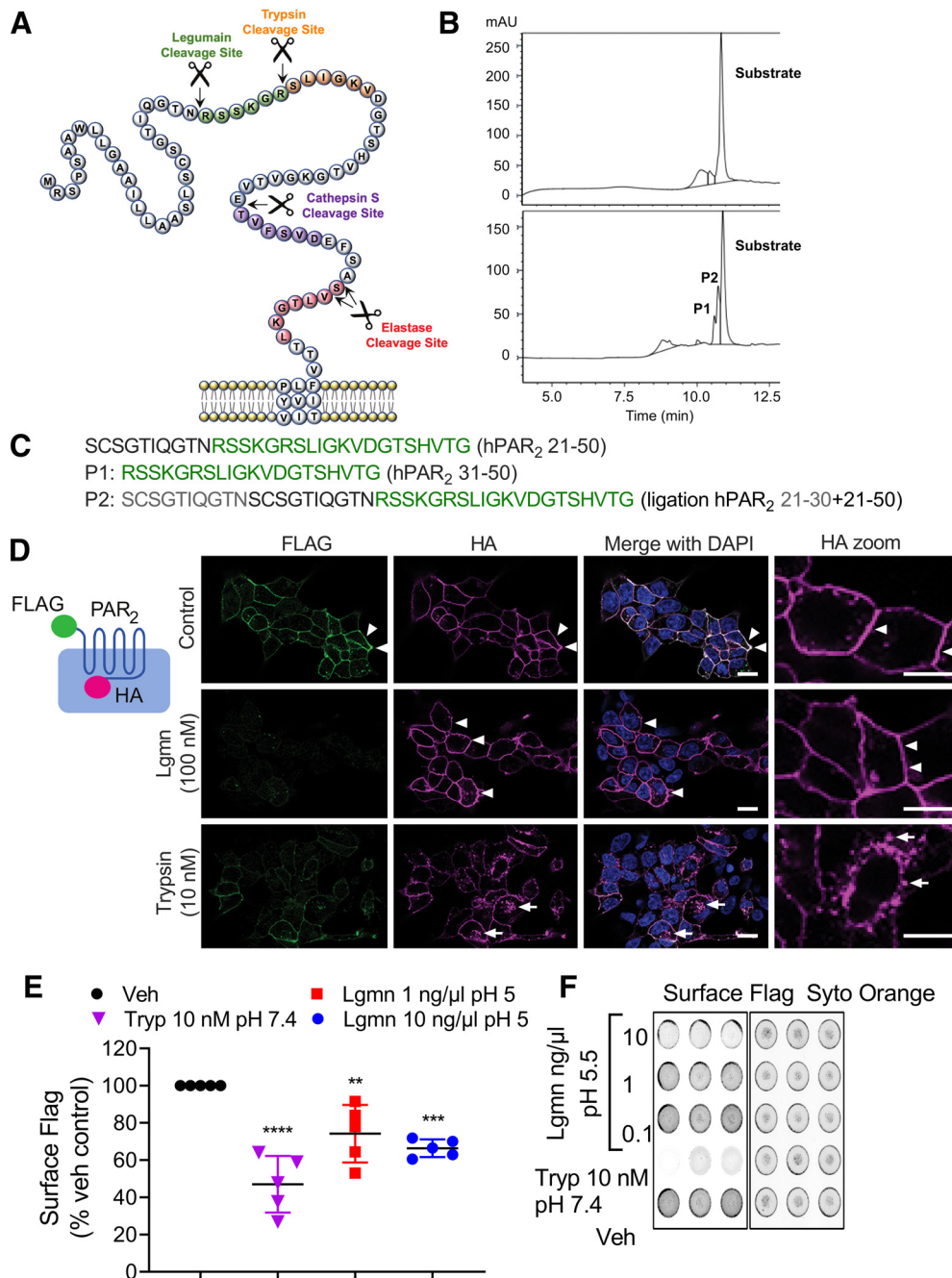
### Lgmn mediates OSCC nociception

We used LI-1 to study whether Lgmn secreted from HSC-3 contributes to nociception in the xenograft paw cancer model. After measuring baseline mechanical withdrawal and thermal latency in the paws of NU/J *Foxn1*<sup>tm</sup> athymic mice, we inoculated HSC-3 cells, which highly express Lgmn (Fig. 5A). Post-inoculation withdrawal measurements verified cancer-generated mechanical allodynia or thermal hyperalgesia. Mice were then treated with LI-1, and mechanical and thermal nociception were assessed (Fig. 5A). LI-1 reversed OSCC-induced mechanical withdrawal at 3, 6, 12, and 24 h after injection (Fig. 5B), and reversed thermal hyperalgesia at 3 and 24 h postinjection (Fig. 5C). LI-1 had no effect 48 h postinjection. An OSCC mouse model was generated with 4NQO administered over 16 weeks in *Lgmn*<sup>-/-</sup>, *Par2*<sup>-/-</sup> and WT mice. Oral mechanical allodynia was measured with dolognawmeters at week 28. Tongues were removed, sectioned, stained with H & E, and reviewed independently by two pathologists (Fig. 5D). OSCC was confirmed in all groups (Fig. 5E). OSCC incidence was >80% at 28 weeks after the administration of 4NQO in all groups (Fig. 5F). *Lgmn*<sup>-/-</sup> and *Par2*<sup>-/-</sup> mice showed significantly less mechanical allodynia than the WT mice (Fig. 5G).

### Lgmn induces PAR<sub>2</sub>-dependent hyperexcitability of trigeminal neurons

To determine whether Lgmn causes PAR<sub>2</sub>-dependent hyperexcitability of TG neurons, we measured rheobase in WT mice using whole-cell patch-clamp. Neurons were studied in acidic buffer (external solution, pH 5.5). Lgmn

(20 ng/ml, 10 min) decreased rheobase versus vehicle. There was no significant difference between the resting membrane potentials of TG neurons from WT mice pretreated with Lgmn versus Lgmn vehicle (Lgmn:  $-55.7 \pm 2.2$  mV,  $n = 13$ ; Lgmn vehicle:  $-56.6 \pm 3.9$  mV;  $n = 11$ ,  $t_{(22)} = 0.207$ ,  $p = 0.8378$ , unpaired Student's *t* test). The mean input resistance was increased in TG neurons pretreated with Lgmn compared with Lgmn vehicle, but there was no statistically significant difference (Lgmn:  $659.3 \pm 93.9$  M $\Omega$ ,  $n = 13$ ; Lgmn vehicle:  $526.9 \pm 79.9$  M $\Omega$ ;  $n = 11$ ,  $t_{(22)} = 1.052$ ,  $p = 0.3443$ , unpaired Student's *t* test). To investigate requirement for activity, we preincubated Lgmn with LI-1 (10  $\mu\text{M}$ , 10 min) or vehicle. LI-1 prevented the effect of Lgmn on



**Figure 7.** Lgmn cleavage of PAR<sub>2</sub>. **A**, Lgmn PAR<sub>2</sub> cleavage sites and PAR<sub>2</sub> N terminus indicating known cleavage sites. **B**, **C**, HPLC chromatograms (**B**) and product identification by mass spectrometry (**C**) of degradation of a fragment of PAR<sub>2</sub> (hPAR<sub>2</sub><sup>21–50</sup>) by Lgmn, showing that Lgmn cleaves PAR<sub>2</sub> at Asn<sup>30</sup>↓Arg<sup>31</sup>. **D**, Localization of PAR<sub>2</sub> using antibodies to extracellular N-terminal FLAG and intracellular C-terminal HA epitopes (inset) in HEK-FLAG-PAR<sub>2</sub>-HA cells incubated with vehicle (control), Lgmn, or trypsin. Arrows denote plasma membrane localization; arrowheads denote endosomal localization. Scale bar: 20 μm. **E**, **F**, On-cell Western showing that trypsin and Lgmn remove the extracellular N-terminal FLAG epitope from HEK-FLAG-PAR<sub>2</sub>-HA cells. **E**, Quantification of triplicate observations from  $n = 5$  individual experiments ( $F_{(5,22)} = 16.03$ ,  $**p = 0.0034$ , Lgmn 1 ng/μl compared with vehicle,  $***p = 0.0002$ , Lgmn 100 ng/μg compared with vehicle,  $****p = 2.5E-7$  trypsin compared with vehicle, one-way ANOVA with Dunnett's test,  $n = 5$ ). **F**, Representative images of FLAG immunoreactivity and Syto Orange stain.

rheobase (Fig. 6A,B). To investigate the contribution of PAR<sub>2</sub>, we analyzed Lgmn-induced hyperexcitability in TG neurons from *Par<sub>2</sub><sup>-/-</sup>* mice. Lgmn (20 ng/ml, 10 min) did not affect rheobase of *Par<sub>2</sub><sup>-/-</sup>* neurons (Fig. 6C,D).

To evaluate the signaling pathway that mediates effects of Lgmn on excitability, perforated patch-clamp recordings were made from TG neurons pretreated with inhibitors of protein kinase C (PKC; GFX, 1 μM; Coultrap et al., 1999) or protein kinase A (PKA; PKI-tide, 1 μM; Ohlstein et al., 1990) for 30 min at 37°C

before treatment with Lgmn (20 ng/ml) or vehicle. GFX and PKI-tide prevented Lgmn-induced hyperexcitability (Fig. 6E). Thus, Lgmn causes hyperexcitability of TG nociceptors through Lgmn enzymatic activity, expression of PAR<sub>2</sub>, PKC activity, and PKA activity.

#### Lgmn cleaves PAR<sub>2</sub>

To determine whether Lgmn can cleave PAR<sub>2</sub> and identify the cleavage site, Lgmn (200 nM in acetate buffer, pH 4.5)

**Table 2. Masses of hPAR<sub>2</sub> N-terminal peptide cleavage products identified by mass spectrometry**

Peptide	Expected	Found
Substrate:	M: 2991.268	
SCSGTIQGTNRSSKGRSLIGKV <sub>0</sub> GTSHVTG	(M + 2)/2: 1496.64	1496.45
	(M + 3)/2: 998.10	998.10
	(M + 4)/4: 748.83	748.80
Product 1:	M: 2042.263	
RSSKGRSLIGKV <sub>0</sub> GTSHVTG	(M + 2)/2: 1022.14	1022.45
	(M + 3)/3: 681.76	681.95
	(M + 4)/4: 511.57	
Product 2:	M: 3958.274	
SCSGTIQGTNSCSGTIQTNRSSKGRSLIGKV <sub>0</sub> GTSHVTG•H <sub>2</sub> O	(M + 2)/2: 1980.15	
	(M + 3)/3: 1320.43	1320.10
	(M + 4)/4: 990.58	990.40

was incubated with a peptide corresponding to residues 21–50 (S<sup>21</sup>CSGTIQTNRSSKGRSLIGKVDGTSHVTG<sup>50</sup>) of the extracellular N terminus of hPAR<sub>2</sub> (200 μM). Digest was analyzed by HPLC and mass spectrometry. Cleavage products were detected corresponding to PAR<sub>2</sub><sup>31–50</sup> and PAR<sub>2</sub><sup>21–30</sup> fused to PAR<sub>2</sub><sup>21–50</sup> (Fig. 7A–C; Table 2). Thus, Lgmn cleaves the N terminus of hPAR<sub>2</sub> at Asn<sup>30</sup>↓Arg<sup>31</sup> consistent with its preference for asparagine residues and ability to ligate peptides with C-terminal asparagine residues to free N termini (Mikula et al., 2017).

To determine whether Lgmn cleaves intact PAR<sub>2</sub> at the plasma membrane, hPAR<sub>2</sub> with an extracellular FLAG epitope and intracellular HA epitope was expressed in HEK293 cells (Fig. 7D). HEK-FLAG-PAR<sub>2</sub>-HA cells were incubated with Lgmn (100 nM, HBSS pH 5.0), trypsin (10 nM, HBSS pH 7.4), or vehicle (buffer control). FLAG and HA were localized by immunofluorescence and confocal microscopy. In vehicle-treated cells, FLAG and HA colocalized at the plasma membrane (Fig. 7D). After incubation with Lgmn, FLAG was depleted from the plasma membrane whereas HA was retained at the plasma membrane, consistent with PAR<sub>2</sub> cleavage and removal of the extracellular FLAG epitope. After incubation with trypsin, FLAG was depleted from the plasma membrane, and HA was detected within endosomes, consistent with PAR<sub>2</sub> cleavage and endocytosis (Böhm et al., 1996a).

On-cell Western was used to quantify removal of the FLAG epitope. In Lgmn incubated cells (1 or 10 ng/μl, MES-HBSS pH 5.0, 30 min, 37°C), FLAG immunoreactivity was reduced 26 ± 7% (1 ng/μl) or 34 ± 2% (10 ng/μl) versus vehicle-treated cells (Fig. 7E,F). After trypsin incubation (10 nM, HBSS pH 7.4, 30 min, 37°C), FLAG immunoreactivity was reduced 53 ± 7% versus vehicle-treated cells. Nuclear stain (Syto Orange) confirmed that proteases did not remove cells from the plate (Fig. 7F).

Thus, Lgmn can cleave intact PAR<sub>2</sub> at the surface of HEK cells and remove the extracellular FLAG epitope. Lgmn cleaves PAR<sub>2</sub> at Asn<sup>30</sup>↓Arg<sup>31</sup>, proximal to the trypsin cleave site (Arg<sup>36</sup>↓Ser<sup>37</sup>). Lgmn does not evoke endocytosis of PAR<sub>2</sub>.

### Lgmn activity and Lgmn-induced Ca<sup>2+</sup> signaling are pH dependent

Trypsin, tryptase, and kallikreins cleave PAR<sub>2</sub> at Arg<sup>36</sup>↓Ser<sup>37</sup> and induce coupling to G<sub>αq</sub> and mobilization of intracellular Ca<sup>2+</sup> (Böhm et al., 1996a; Corvera et al., 1999; Oikonomopoulou et al., 2006). To examine whether Lgmn can mobilize Ca<sup>2+</sup>, HEK-

FLAG-PAR<sub>2</sub>-HA cells were challenged with Lgmn (1 or 10 ng/μl, MES-HBSS pH 5.0, 5.5, 6.0 or HBSS pH 7.4), trypsin (10 nM, HBSS pH 7.4), or vehicle (buffer control). Change in intracellular Ca<sup>2+</sup> was measured using fura-2 AM. Trypsin increased Ca<sup>2+</sup>, reflected by increased F340/380 nm emission, which rapidly declined toward basal (Fig. 8A). At pH 7.4, 10 ng/μl Lgmn induced a small increase in Ca<sup>2+</sup>, whereas 1 ng/μl Lgmn did not elicit a Ca<sup>2+</sup> response (Fig. 8A). At a lower pH of 5.0, 1 and 10 ng/μl Lgmn caused sustained and concentration-dependent increases in Ca<sup>2+</sup> (Fig. 8B,E). At pH 5.0 and 5.5, 10 ng/μl Lgmn increased Ca<sup>2+</sup>, compared with pH 6.0 and 7.4 (Fig. 8C,D); however, at pH 5, 1 ng/μl Lgmn increased Ca<sup>2+</sup> compared with pH 5.5 and 6.0 (Fig. 8E,F). Lgmn activity assays confirmed the acidic pH optimum of Lgmn, which was active at pH 5.0 and 5.5 but not pH >6.0 (Fig. 8G). Lgmn increased Ca<sup>2+</sup>, in the absence of extracellular Ca<sup>2+</sup>, indicating intracellular mobilization (Fig. 8H). The Lgmn inhibitors, QDD100531 (1 μM) and QD123427 (100 nM), prevented Lgmn-evoked (1 ng/μl) Ca<sup>2+</sup> signals (Fig. 8I,J), and caused concentration-dependent inhibition of activity (Fig. 8K; Ness et al., 2015).

### Lgmn induces Ca<sup>2+</sup> signaling through PAR<sub>2</sub>

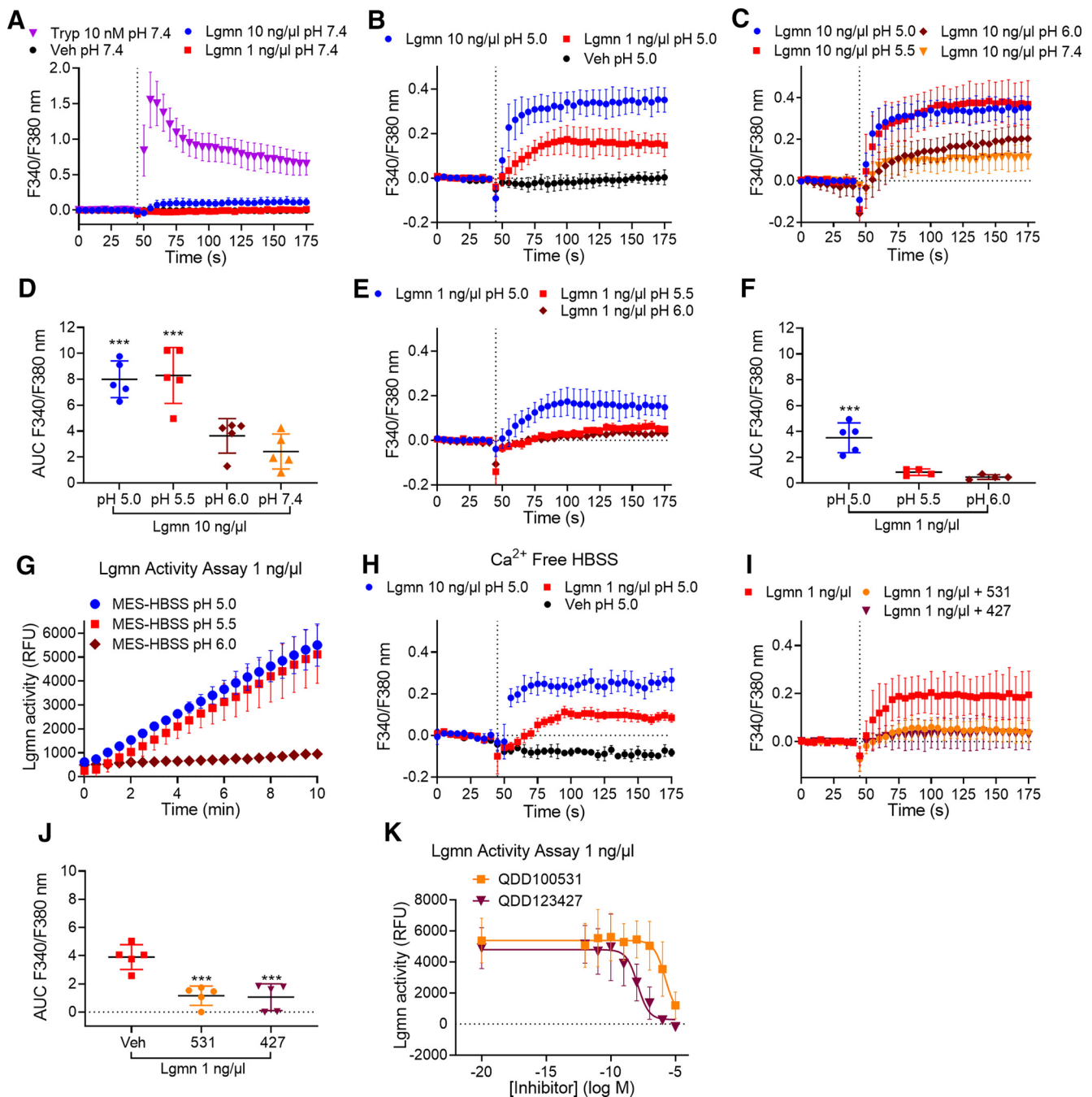
To determine whether Lgmn induces Ca<sup>2+</sup> signaling through PAR<sub>2</sub>, we used specific antagonists for PAR<sub>2</sub> and cells genetically deleted for PAR<sub>2</sub>. PAR<sub>2</sub> antagonists, I-343 and GB88 (10 μM; Farmer, 2013; Lieu et al., 2016; Jimenez-Vargas et al., 2018) abolished Lgmn-stimulated (1 and 10 ng/μl) Ca<sup>2+</sup> responses (Fig. 9A–C). I-343 and GB88 did not inhibit the enzymatic activity of Lgmn (1 ng/μl), which was slightly increased in the presence of these antagonists (Fig. 9D). To determine whether Lgmn induced a Ca<sup>2+</sup> increase through PAR<sub>1</sub>, we used the specific PAR<sub>1</sub> antagonist SCH79797 (Ahn et al., 2000). SCH79797 (200 nM) did not alter Lgmn-induced Ca<sup>2+</sup> responses (Fig. 9E,F). To confirm the pharmacologic evidence that PAR<sub>2</sub>, and not PAR<sub>1</sub>, mediated Lgmn-evoked Ca<sup>2+</sup> signals, PAR<sub>2</sub> or PAR<sub>1</sub> was deleted from HEK293 cells using CRISPR/Cas9 (Ungefroren et al., 2017). In PAR<sub>2</sub>-KO cells, the PAR<sub>2</sub> selective agonist 2-Furoyl-LIGRLO-NH<sub>2</sub> did not increase Ca<sup>2+</sup> (Fig. 9G); however, the PAR<sub>1</sub> selective agonist TFLLR-NH<sub>2</sub> increased Ca<sup>2+</sup> in PAR<sub>2</sub>-KO cells (Fig. 9H). In PAR<sub>1</sub>-KO cells, TFLLR-NH<sub>2</sub> did not increase Ca<sup>2+</sup> except at a high concentration (10 μM; Fig. 9I), whereas 2-Furoyl-LIGRLO-NH<sub>2</sub> increased Ca<sup>2+</sup> in PAR<sub>1</sub>-KO cells (Fig. 9J). Lgmn (10 ng/μl) mobilized Ca<sup>2+</sup> in PAR<sub>1</sub>-KO but not PAR<sub>2</sub>-KO cells (Fig. 9K).

To confirm that Lgmn activates PAR<sub>2</sub> by cleavage at the Asn<sup>30</sup>↓Arg<sup>31</sup> site, we generated a mutant receptor in which the Asn<sup>30</sup> residue was mutated to Ala, which would not be recognized by Lgmn. PAR<sub>2</sub>-ΔN30A was transfected into PAR<sub>2</sub>-KO HEK cells. 2-Furoyl-LIGRLO-NH<sub>2</sub> (10 μM) robustly increased Ca<sup>2+</sup> in HEK-PAR<sub>2</sub>ΔN30A cells, whereas Lgmn (10 ng/μl) had no effect (Fig. 9L). To determine whether Lgmn activates PAR<sub>2</sub> by exposure of a tethered ligand, we synthesized a hexapeptide, R<sup>31</sup>SSKGR<sup>36</sup>, corresponding to a potential tethered ligand revealed by Lgmn cleavage of PAR<sub>2</sub> at the Asn<sup>30</sup>↓Arg<sup>31</sup> site. However, R<sup>31</sup>SSKGR<sup>36</sup> (10 μM to 0.01 nM) did not alter Ca<sup>2+</sup> in HEK-PAR<sub>2</sub> cells (Fig. 9M). These data confirm that the Lgmn-driven Ca<sup>2+</sup> response is PAR<sub>2</sub> dependent but does not involve exposure of a tethered ligand domain.

### Lgmn desensitizes PAR<sub>2</sub>, but does not induce an association between PAR<sub>2</sub> and β-arrestin-1

Processes that terminate PAR<sub>2</sub> signaling at the plasma membrane include β-arrestin-mediated desensitization of PAR<sub>2</sub>,

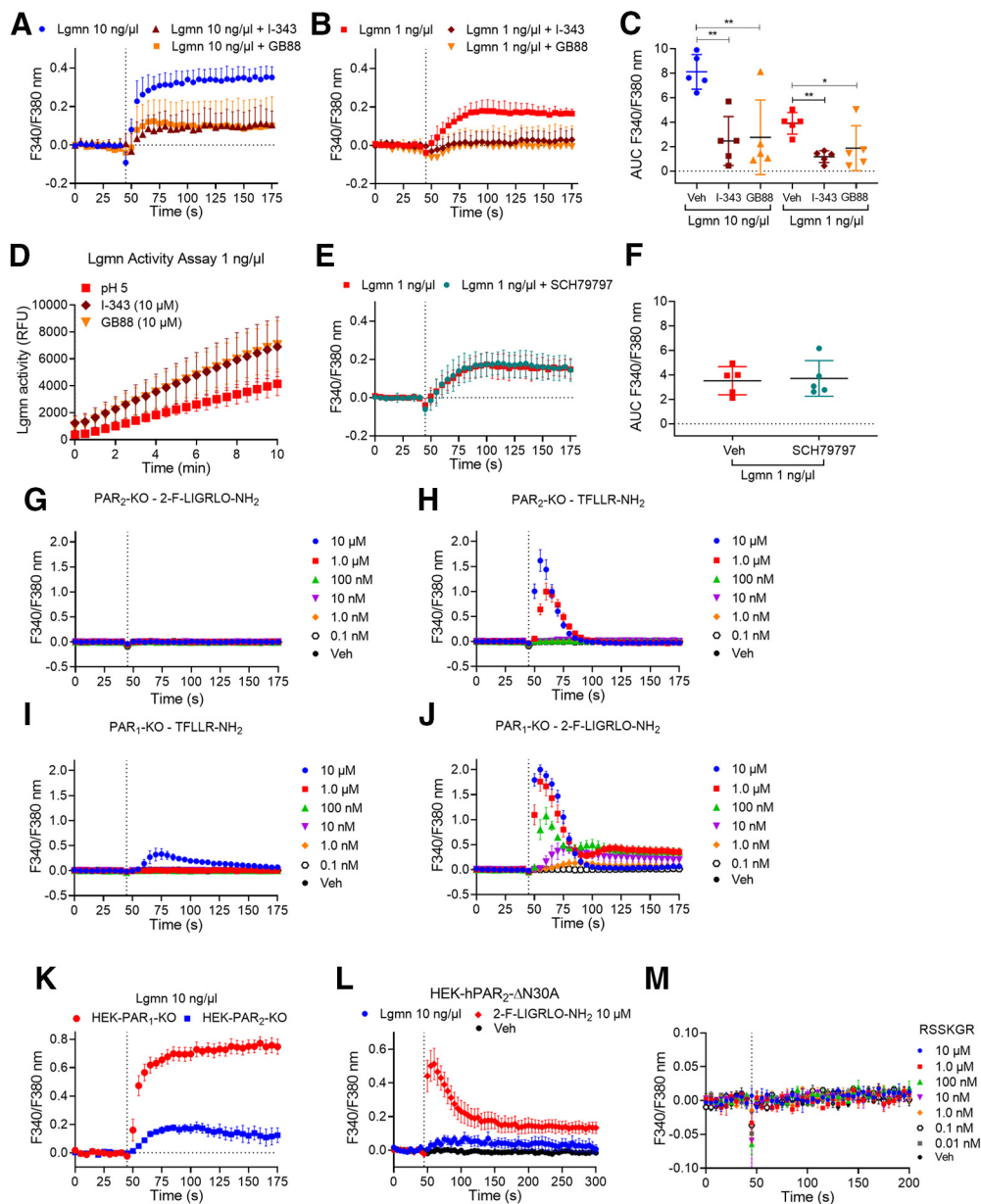




**Figure 8.** pH dependence of Lgmn activity and stimulation of Ca<sup>2+</sup> signaling. Time course (**A–C**) and area under curve (AUC; **D**) of trypsin-evoked and Lgmn-evoked Ca<sup>2+</sup> signaling in HEK-FLAG-PAR<sub>2</sub>-HA cells at pH 7.4, 6.0, 5.5, and 5.0 ( $F_{(3,16)} = 17.5$ ,  $***p = 0.003$ , pH 5.0 compared with pH 7.4,  $***p = 0.0001$ , pH 5.5 compared with pH 7.4, one-way ANOVA with Tukey's test,  $n = 5$ ). **E, F**, Time course (**E**) and AUC (**F**) of Lgmn (1 ng/μl)-evoked Ca<sup>2+</sup> signaling in HEK-FLAG-PAR<sub>2</sub>-HA cells at pH 6.0, 5.5, and 5.0 ( $F_{(2,10)} = 22.68$ ,  $***p = 0.0009$  for pH 5.0 compared with pH 5.5 and  $***p = 0.0003$  for pH 5.0 compared with pH 6.0, one-way ANOVA with Tukey's test,  $n = 5$ ). **G**, Lgmn activity assays at pH 5.0, 5.5, and 6.0. **H**, Effects of depletion of extracellular Ca<sup>2+</sup> on Lgmn responses. **I, J**, Time course (**I**) and AUC (**J**) of the effects of the Lgmn inhibitors QDD100531 (531) and QD123427 (427) on Lgmn Ca<sup>2+</sup> signals ( $F_{(2,12)} = 22.32$ ,  $***p = 0.0002$  for 531 and  $***p = 0.0002$  for 427 compared with vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ). **K**, Lgmn activity assay in the presence of graded concentrations of Lgmn inhibitors QDD100531 and QDD123427 (pH 5.0).

PAR<sub>2</sub> cleavage and removal of activation sites and tethered ligand domains, and PAR<sub>2</sub> endocytosis (Böhm et al., 1996a; Déry et al., 1999; DeWire et al., 2007). Since Lgmn cleaves PAR<sub>2</sub> proximal to the trypsin site, subsequent inhibition of trypsin signaling would likely reflect PAR<sub>2</sub> desensitization. To examine desensitization, HEK-FLAG-PAR<sub>2</sub>-HA cells were incubated with Lgmn (1 or 10 ng/μl, MES-HBSS pH 5.0), trypsin (10 nM, HBSS pH 7.4), or vehicle (buffer control; Fig. 10A). Cells were washed and

recovered in HBSS pH 7.4 for 20 min and then challenged with trypsin (10 nM). In cells preincubated with vehicle, trypsin challenge at 30 min robustly increased Ca<sup>2+</sup> (Fig. 10B,C). Initial challenge with trypsin also increased Ca<sup>2+</sup>, but response to a second challenge at 30 min was reduced by  $53.7 \pm 6.3\%$  versus the response in vehicle-treated cells, consistent with desensitization and endocytosis of PAR<sub>2</sub>. Initial challenge with Lgmn (10 ng/μl) slightly increased Ca<sup>2+</sup>, but the response to a second challenge at



**Figure 9.** Lgmn signals through PAR<sub>2</sub>. **A–C**, Time course (**A**, **B**) and area under curve (**C**) of the effects of the PAR<sub>2</sub> antagonists I-343 and GB88 on Lgmn Ca<sup>2+</sup> signals ( $F_{(2,12)} = 9.89$  for Lgmn 10 ng/μl with  $**p = 0.0036$  for I-343 and  $**p = 0.0052$  for GB88 compared with vehicle,  $F_{(2,12)} = 6.84$  for Lgmn 1 ng/μl with  $**p = 0.0073$  for I-343 and  $*p = 0.0384$  for GB88 compared with vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ). **D**, Lgmn activity assays in the presence of the PAR<sub>2</sub> antagonists I-343 and GB88 (pH 5.0). Triplicate observations from  $n = 5$  individual experiments. **E**, **F**, Time course (**E**) and area under curve (**F**) of the effects of the PAR<sub>1</sub> antagonist SCH79797 on Lgmn Ca<sup>2+</sup> signals. **G–J**, Effects of the PAR<sub>2</sub> agonist 2-Furoyl-LIGRLO-NH<sub>2</sub> and the PAR<sub>1</sub> agonist TFLLR-NH<sub>2</sub> on Ca<sup>2+</sup> signals in PAR<sub>2</sub>-KO (**G**, **H**) and PAR<sub>1</sub>-KO HEK293 (**I**, **J**) cells. **K**, Time course of Lgmn Ca<sup>2+</sup> signals in HEK-PAR<sub>1</sub>-KO and HEK-PAR<sub>2</sub>-KO cells. **L**, Time course of Lgmn Ca<sup>2+</sup> signals in HEK-PAR<sub>2</sub>ΔN30A cells. **M**, Effects of potential PAR<sub>2</sub> activating peptide R<sup>31</sup>SSKGR<sup>36</sup> Ca<sup>2+</sup> signaling in HEK-FLAG-PAR<sub>2</sub>-HA cells.

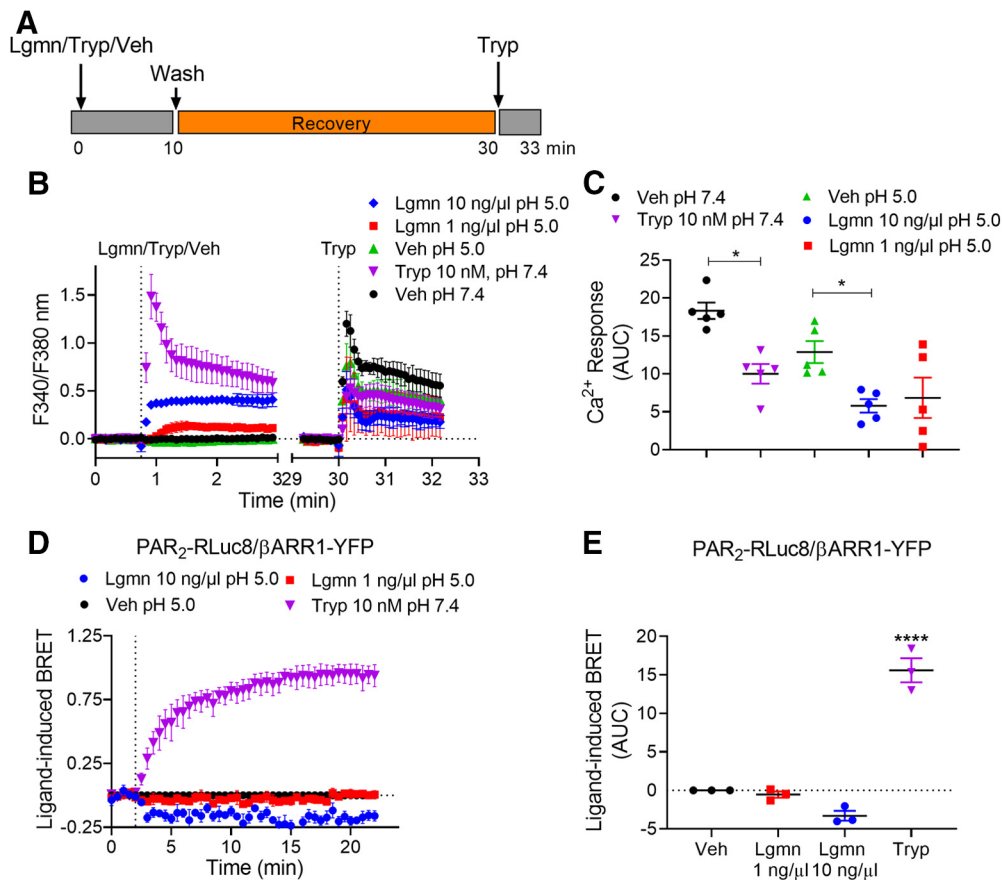
30 min was reduced by  $49.9 \pm 6.9\%$  versus response in vehicle-treated cells, consistent with desensitization of PAR<sub>2</sub>.

Given that Lgmn desensitizes PAR<sub>2</sub>, we sought to determine whether Lgmn recruits  $\beta$ -arrestin-1 to PAR<sub>2</sub>. After trypsin cleavage, PAR<sub>2</sub> becomes phosphorylated by GPCR kinases and interacts with  $\beta$ -arrestins, which mediate desensitization and endocytosis (Corvera et al., 1999). However, after cathepsin S or elastase cleavage, PAR<sub>2</sub> neither recruits  $\beta$ -arrestins nor induces endocytosis (Zhao et al., 2014, 2015). We showed that Lgmn does not induce PAR<sub>2</sub> endocytosis (Fig. 7D). It is unknown whether PAR<sub>2</sub> associates with  $\beta$ -arrestin-1 following Lgmn cleavage. Thus, we examined BRET between PAR<sub>2</sub>-Rluc8 and  $\beta$ -arrestin-1-YFP following treatment with Lgmn. Trypsin

(10 nM), but not Lgmn (1 or 10 ng/μl), stimulated PAR<sub>2</sub>-Rluc8/ $\beta$ -arrestin-1-YFP BRET (Fig. 10D,E). These results accord with the inability of Lgmn to evoke PAR<sub>2</sub> endocytosis.

#### Lgmn activates PAR<sub>2</sub>-mediated cAMP formation and activation of PKD and ERK in HEK293 cells

After activation by trypsin, PAR<sub>2</sub> couples to G $\alpha_q$ , leading to mobilization of intracellular Ca<sup>2+</sup>, generation of cAMP, and activation of ERK and PKD (DeFea et al., 2000; Amadesi et al., 2009). ERK contributes to sensitization of nociceptors (Ji et al., 1999), and PKD promotes mobilization of PAR<sub>2</sub> from Golgi and recovery of responses to extracellular proteases (Amadesi et al., 2009; Zhao et al., 2019). To examine whether Lgmn-activated PAR<sub>2</sub>



**Figure 10.** Lgmn desensitization of PAR<sub>2</sub> Ca<sup>2+</sup> signaling and recruitment of  $\beta$ -arrestin-1. **A**, HEK-FLAG-PAR<sub>2</sub>-HA cells were exposed to trypsin or Lgmn for 10 min, washed, and then challenged with trypsin 20 min after washing. **B**, Time course of Ca<sup>2+</sup> signaling. **C**, Recovery of trypsin responses [area under curve (AUC) from **B**] in cells pretreated with vehicle, trypsin, or Lgmn ( $F_{(4,20)} = 1.61$ ,  $*p = 0.0116$  for trypsin, and  $*p = 0.0372$  for Lgmn 10 ng/ $\mu$ l compared with vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ). **D**, **E**, BRET assays of recruitment of  $\beta$ -arrestin-1 to PAR<sub>2</sub>. **D**, Time courses. **E**, AUC ( $F_{(3,8)} = 96.95$ ,  $****p = 1.25E-6$  for trypsin versus vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ). All assays were done in triplicate.

couple to a similar range of downstream effectors, we expressed FRET biosensors of cytosolic cAMP (Cyto-Epac), cytosolic PKD (Cyto-DKAR), and cytosolic ERK (Cyto-EKAR) in HEK-FLAG-PAR<sub>2</sub>-HA cells. Lgmn caused concentration-dependent activation of cAMP (Fig. 11A,B), PKD (Fig. 11C,D), and ERK (Fig. 11E,F) within the cytosol. cAMP and PKD responses were robust; however, the ERK response was small and detected only after treatment with a higher Lgmn concentration.

## Discussion

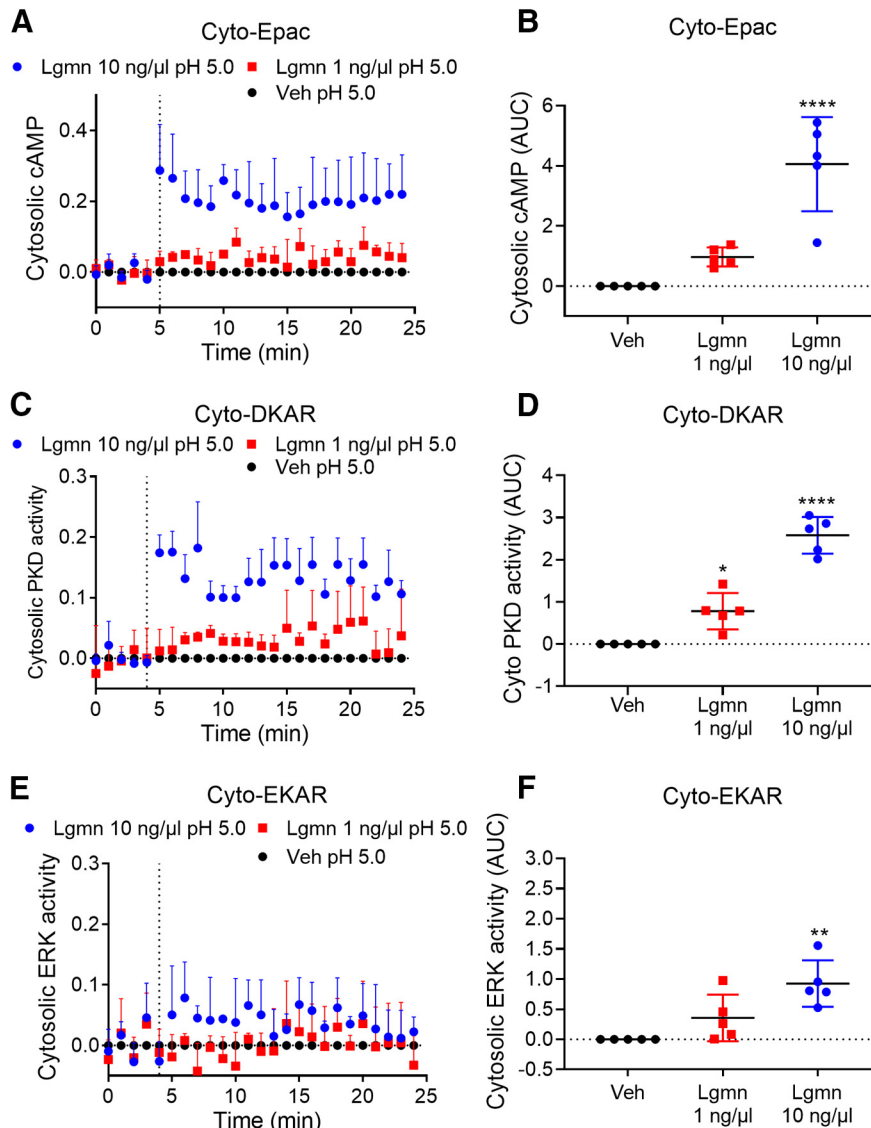
We report that Lgmn is secreted from OSCC cells and is robustly and reproducibly activated in human and mouse OSCCs compared with normal mucosa. Under acidic conditions, Lgmn cleaves and activates PAR<sub>2</sub> by biased mechanisms to evoke sustained hyperexcitability of nociceptors. We confirmed that PAR<sub>2</sub> and Lgmn contribute to OSCC pain in OSCC mouse models that recapitulate the progression of OSCC observed in humans; genes for Lgmn and PAR<sub>2</sub> on nociceptors were deleted in these mice. Lgmn contributes to cancer hallmarks including proliferation, invasion, and metastasis (Murthy et al., 2005; Vasiljeva et al., 2006; Li et al., 2013; Ohno et al., 2013; Edgington-Mitchell et al., 2015). While Lgmn has been reported to produce bone cancer pain through neurotrophin receptors (Yao et al., 2017), the role of the Lgmn/PAR<sub>2</sub> axis has not been described and could be therapeutically exploited.

The mechanism responsible for Lgmn activation in oral cancer is unresolved. Lgmn is synthesized as pro-Lgmn and traffics

through the endoplasmic reticulum and Golgi (Dall and Brandstetter, 2016). Lgmn is packaged and activated in the acidic environment of lysosomes (Dall and Brandstetter, 2016). At pH >6.0, acidic residues unfold and lose proteolytic activity. pH in cancers varies (5.4–6.7; Meyer et al., 1948; Vaupel et al., 1981; Newell et al., 1993; Gillies et al., 1994); the Lgmn activation mechanism remains obscure (Dall and Brandstetter, 2012). Exosomes released from OSCC might exhibit a pH low enough to activate Lgmn. A single report reveals acidic exosomes in cancer patients (Logozzi et al., 2019). Cells from human OSCCs, including the cell line used in this study (HSC-3), secrete exosomes (Dayan et al., 2012; Li et al., 2019). Stabilization between a RGD motif in the catalytic domain and the integrin  $\alpha_v\beta_3$  might also activate Lgmn at a higher pH (Liu et al., 2012).

We demonstrated that Lgmn causes cancer-associated nociception through PAR<sub>2</sub> activation on Na<sub>v</sub>1.8-expressing neurons. While all nociceptors express Na<sub>v</sub>1.8, some non-nociceptors, including low-threshold mechanoreceptors that mediate touch sensation, also express Na<sub>v</sub>1.8 (Shields et al., 2012). Since Lgmn activates cathepsins (Edgington-Mitchell et al., 2016), which can also activate PAR<sub>2</sub> (Zhao et al., 2014), it is possible that Lgmn activates PAR<sub>2</sub> directly or indirectly (Edgington-Mitchell et al., 2016). However, we found that Lgmn directly cleaves a fragment of hPAR<sub>2</sub> at a unique Asn<sup>30</sup>↓Arg<sup>31</sup> site, consistent with known Lgmn selectivity. Site mutation prevented Lgmn-evoked signaling, confirming this mechanism of proteolytic activation. Lgmn evoked hypersensitivity of TG neurons from WT mice





**Figure 11.** Lgmn activation of cAMP, PKD and ERK signaling and recruitment of  $\beta$ -arrestin-1. **A–F**, FRET assays of cytosolic cAMP (**A, B**), cytosolic PKD (**C, D**), and cytosolic ERK (**E, F**) in HEK-FLAG-PAR<sub>2</sub>-HA cells. **A, C, E**, Time courses. **B**, Area under curve (AUC) for cytosolic cAMP ( $F_{(2,12)} = 26.43$ , \*\*\*\* $p = 0.00,004$  for Lgmn 10 ng/μl compared with vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ). **D**, AUC for cytosolic PKD ( $F_{(2,12)} = 69.77$ , \*\*\*\* $p = 2.5E-7$  for Lgmn 10 ng/μl and \* $p = 0.0117$  for Lgmn 1 ng/μl compared with vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ). **F**, AUC for cytosolic ERK ( $F_{(2,12)} = 10.99$ , \*\* $p = 0.0015$  for Lgmn 10 ng/μl compared with vehicle, one-way ANOVA with Tukey's test,  $n = 5$ ).

(determined by patch clamp). These effects of Lgmn were prevented by a Lgmn inhibitor and absent in neurons from mice lacking PAR<sub>2</sub>, confirming necessity of PAR<sub>2</sub> activation. The nociceptive behavior we measured accords with Lgmn-induced neuronal hypersensitivity. Lgmn induced nociceptive responses in anatomic regions innervated by DRG (paw) and TG (craniofacial) neurons; a Lgmn inhibitor and selective deletion of PAR<sub>2</sub> in Na<sub>v</sub>1.8 neurons attenuated nociceptive responses. A Lgmn inhibitor eliminated chronic mechanical and thermal nociception in mice inoculated with OSCC cells.

OSCC patients complain of mechanical-induced and function-induced pain and not spontaneous pain (Connelly and Schmidt, 2004; Kolokythas et al., 2007). Our operant orofacial pain assay and automated device to perform the assay (dolog-nawmeter) quantifies a behavioral index of mechanical allodynia during gnawing (comparable to chewing in humans). Lgmn-induced mechanical allodynia and heat hyperalgesia in OSCC patients might involve PAR<sub>2</sub> sensitization of TRPV4 and TRPV1

ion channels, respectively (Grant et al., 2007; Sipe et al., 2008). TRPV4 mediates mechanosensation, while TRPV1 responds to heat and acids (Caterina et al., 1999; Liedtke and Friedman, 2003; Liedtke et al., 2003; Suzuki et al., 2003a,b). TRPV4 and TRPV1 are sensitized by adenylyl cyclase-dependent, PKA-dependent, and PKC $\epsilon$ -dependent mechanisms, which yield ion channel phosphorylation (Numazaki et al., 2002; Amadesi et al., 2006; Zhao et al., 2019). We showed that Lgmn cleavage of PAR<sub>2</sub> activates adenylyl cyclase and cAMP formation; cAMP unleashes catalytic subunits of PKA, which subsequently phosphorylate TRPV channels. We also showed that Lgmn alters rheobase through PKC. Lgmn robustly activates PKD, which likely contributes to PAR<sub>2</sub> trafficking from Golgi to plasma membrane (Zhao et al., 2019).

HEK293 cell experiments revealed that Lgmn mobilizes intracellular calcium, stimulates formation of cAMP, and activates PKD and ERK. Selective inhibitors of Lgmn abolished the

calcium responses; we therefore infer that protease activity is necessary. Lgmn-evoked signals were detected only under mildly acidic conditions, consistent with the acidic pH optimum of Lgmn. We infer that Lgmn-evoked calcium signaling required cleavage/activation of PAR<sub>2</sub> because PAR<sub>2</sub> antagonism or deletion and mutation of the cleavage site abolished signals. PAR<sub>1</sub> antagonism or deletion had no effect. Further studies are needed to reveal mechanisms by which Lgmn activated PAR<sub>2</sub> signals to regulate channel activity and nociception. Trypsin activation of PAR<sub>2</sub> involves exposure of a tethered ligand domain; peptides mimicking the tethered ligand activate the receptor (Hollenberg et al., 1996). Lgmn activation does not involve a tethered ligand; a synthetic peptide corresponding to the revealed N terminus was inactive. Trypsin-activated PAR<sub>2</sub> recruits  $\beta$ -arrestins and then internalizes; Lgmn did not promote  $\beta$ -arrestin recruitment or receptor endocytosis. Thus, like cathepsin S and elastase (Zhao et al., 2014, 2015), Lgmn activates PAR<sub>2</sub> by biased mechanisms to evoke pain.

Our findings are relevant for OSCC patients with pain. While the role of PAR<sub>2</sub> in OSCC pain is clear, antagonism of PAR<sub>2</sub> as a pain therapy approach is challenging. Access to the PAR<sub>2</sub> binding pocket frustrates development of a clinically viable PAR<sub>2</sub> antagonist (Goh et al., 2009; Suen et al., 2014; Boitano et al., 2015). Moreover, PAR<sub>2</sub> continues to signal following cleavage and endocytosis (Jimenez-Vargas et al., 2018); however, we showed that Lgmn-cleaved PAR<sub>2</sub> was not endocytosed. Furthermore, blockade of the Lgmn/PAR<sub>2</sub> axis with a Lgmn inhibitor abrogates OSCC pain in mice. Accordingly, a pain therapy strategy that utilizes blockade of Lgmn is physiologically expedient and holds great clinical potential.

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