Inhibition of somatosensory mechanotransduction by Annexin A6

Ramin Raouf†1,2, Stéphane Lolignier†1,3, Jane E. Sexton†1, Queensta Millet1, Sonia Santana-Varela1, Anna Biller1, Alice M. Fuller1, Vanessa Pereira1, Jyoti S. Choudhary4, Mark O. Collins5, Stephen E. Moss6, Richard Lewis7, Julie Tordo8,9, Els Henckaerts8, Michael Linden8 and John N Wood*1

1 Molecular Nociception Group, Wolfson Institute for Biomedical Research, UCL, Gower Street, London, WC1E 6BT, UK

2 current address: Wolfson Centre for Age-Related Diseases, King’s College London, London, SE1 1UL, UK

3current address: NEURO-DOL, UMR1107 INSERM/Université Clermont Auvergne, 63000 Clermont-Ferrand, France

4Wellcome Genome Campus, Hinxton, Cambridge, CB10 1SA, UK

5 Department of Biomedical Science, The University of Sheffield, Western Bank, Sheffield S10 2TN UK

6Institute of Ophthalmology, UCL, 11-43 Bath Street, London, EC1V 9EL, UK

7 Institute for Molecular Bioscience, University of Queensland, Brisbane St Lucia, QLD 4072

8Department of Infectious Diseases, King’s College London School of Medicine, London, SE1 9RT, UK

9Current address: Cell and Gene Therapy Catapult, Guy’s Hospital, London, SE1 9RT, UK

*Correspondence: J.Wood@ucl.ac.uk

†These authors contributed equally to this work
Abstract

Mechanically-activated slowly adapting currents in sensory neurons have been linked to noxious mechanosensation. The conotoxin NMB-1 (noxious mechanosensation blocker-1) blocks such currents and inhibits mechanical pain. Using a biotinylated form of NMB-1 in mass spectrometry analysis, we identified 67 binding proteins in sensory neurons and a sensory neuron-derived cell line, of which the top candidate was annexin A6, a membrane associated calcium-binding protein. Annexin A6-deficient mice showed increased sensitivity to mechanical stimuli. Sensory neurons from these mice showed increased activity of the cation channel Piezo2, which mediates a rapidly adapting mechano-gated current linked to proprioception and touch, and a decrease in mechanically activated, slowly adapting currents. Conversely, overexpression of annexin A6 in sensory neurons inhibited rapidly adapting currents that were partially mediated by Piezo2. Furthermore, overexpression of annexin A6 in sensory neurons attenuated mechanical pain in a mouse model of osteoarthritis, a disease in which mechanically evoked pain is particularly problematic. These data suggest that annexin A6 can be exploited to inhibit chronic mechanical pain.

Introduction

Pain is the greatest clinical challenge of the age in numerical terms, and mechanically-evoked pain is a major issue in many pain states, particularly those linked to arthritis and musculoskeletal pain that occur more frequently in older people (1). A subset of primary nociceptive neurons are directly activated by mechanical stimuli (2). In these neurons, genes encoding mechanically-gated ion channels
are transcriptionally activated by nerve growth factor (NGF) and ion channel trafficking is potentiated by protein kinase C (PKC) (3). Mechanical pain is enhanced by NGF or phorbol esters, suggesting that the mechanically activated currents identified in sensory neurons have a major role in mechanosensation (3). Mechanically-gated currents in sensory neurons have been classified on the basis of their adaptation kinetics (4). Rapidly adapting (RA) currents have been linked to proprioception and touch through the activation of the mechanically-gated ion channel Piezo2 (5, 6). Deletion of Piezo2 with Advillin Cre-ER\textsuperscript{T2} abolishes >80% of RA currents in sensory neurons, as well as 50% of intermediately adapting (IA) currents (5). Slowly adapting (SA) and IA channels have also been linked to noxious mechanotransduction (7).

Although Piezo2 contributes to mechanical allodynia through EPAC mediated regulation (8), the principal mechanically-gated ion channels responsible for noxious mechanosensation have yet to be identified. Some TRP channels, for example TRPC3, seem to contribute to mechanosensory currents, although a contribution of endogenous Piezo channels is a potential confounding factor (9). A slowly adapting mechanosensitive channel, TMEM150c or TNN, expressed in muscle spindles and their afferents does not appear to have a role in mechanical pain (10). Potassium channels found in sensory neurons are also mechanosensors (11). Thus at least 4 distinct channel subtypes, (TRPs, TMEMs, Piezos and K channels) may contribute to somatosensory mechanotransduction.

A screen for toxins that block mechanosensitive channels has identified NMB-1 (Noxious Mechanosensation Blocker-1), a conotoxin peptide that selectively inhibits the persistent, non-adapting IA component and SA mechanically-activated currents in sensory neurons (7). NMB-1 binds to small diameter, peripherin-positive neurons (7), which are responsible for noxious mechanotransduction (12). NMB-1 also inhibits mechanotransduction currents in the cochlea and blocks behavioural responses to noxious mechanical stimuli. To identify the targets to which NMB-1
binds and therefore identify proteins that contribute to mechanotransduction, we developed a biotinylated form of NMB-1 that retained activity. Using sensory neurons and sensory neuron derived cell lines, we performed immunoblotting and pull-down assays using biotinylated NMB-1 followed by mass spectrometry analysis. Several candidates were identified including the Ca\(^{2+}\)-dependent, phospholipid-binding protein, annexin A6. Here we described a role for annexin A6 in mechanotransduction in sensory neurons and the implications of this role in blocking mechanical pain.

**Results**

**Identification of NMB-1 binding proteins shows that NMB-1 interacts with annexin A6**

We have shown that biotinylated NMB-1 (bNMB-1) can specifically bind peripherin-positive dorsal root ganglion (DRG) neurons rather than large diameter sensory neurons. bNMB-1 retains its activity in blocking slowly adapting mechanically evoked currents in these neurons (7). We used this modified form of the toxin to identify proteins that interact with NMB-1 in rat DRG neurons. We developed an immunoblotting protocol to stain interacting proteins on Western Blots of DRG or cerebellum membrane protein extracts and found that the bNMB-1 can bind specific bands from DRG protein extracts whereas the same pattern of staining was not present in protein extracts from cerebellum (Fig. S1a). Due to the low yield of protein recovery from PVDF membrane strips, we identified binding proteins by juxtaposing alongside the blot an identically processed polyacrylamide gel stained with metallic silver to visualize all the proteins (Fig. S1a). We used LC-MS/MS analysis to identify peptides present in bands from the gel that corresponded to positions of bNMB-1 reactive bands on the blot (Data file S1). To restrict this list of potential NMB-1 interacting partners to proteins associated with the mechanotransduction complex, we used a pull-down approach using the differentiated nociceptor cell line ND-C, a DRG neuron-derived cell line that shows discrete mechanically activated currents akin to DRG neurons in culture (13, 14). We performed a pull-down assay for NMB-1 from differentiated
ND-C cell protein extracts (Fig. S1b) and identified proteins present in the pull-downs using mass spectrometry (Data file S2). We reasoned that the list of proteins from the intersection of the two tables would be the smallest set containing bNMB-1 interacting proteins and found that annexin A6 was the most abundant protein shared between the two lists (Table 1).

annexin A6 is a calcium-dependent membrane binding protein that is expressed by most sensory neurons. It has been suggested to modulate Ca$^{2+}$ conductance in DRG neurons (15) and to mediate mechanotransduction currents in lipid bilayers. Incubation of WT and Anxa6 KO DRG tissue with bNMB-1 and fluorescently tagged Avidin (Fig. S2a-f) revealed a substantial decrease in binding of bNMB-1 to Anxa6 KO tissue (Fig. S2g). The residual binding observed in Anxa6 KO tissue also confirmed that NMB-1 bound not only annexin A6 complexes but also other membrane proteins in sensory neurons. We also confirmed that an immunoreactive Annexin A6 band ran at the appropriate molecular weight to one of the bands selected for sequencing (protein band labelled “5” in Fig. S1a), was present in the preparation that produced an annexin A6 signal on mass spectrometry analysis (Fig. S2h). Based on this evidence for an interaction between NMB-1 and annexin A6, we next examined the effects of annexin A6 on mechanotransduction in DRG neurons.

**Annexin A6 deletion potentiates mechanical pain in mice and mechanosensory currents of sensory neurons.**

Using global Anxa6 KO mice, we investigated the role of annexin A6 in mechanosensation in vivo. WT and Anxa6 KO mice showed similar tactile thresholds, as assessed with the von Frey test (Fig. 1a). However, in response to noxious mechanical pressure, Anxa6 KO mice showed a 17.5% decrease in the force required to elicit a response (Fig. 1b). These findings suggest that annexin A6 is involved in noxious, but not innocuous mechanosensation.
We then investigated mechanotransduction at the cellular level in cultured sensory neurons from WT and Anxa6 KO mice dorsal root ganglia by recording mechanically-activated (MA) currents in whole cell patch clamp configuration \((2, 16)\). Currents were evoked by applying focal pressure to the cell membrane with a polished glass pipette in cells were held at -70 mV. Sensory neurons were either mechanically insensitive (non-responding), or mechanically sensitive with an MA current defined by its adaptation kinetics as rapidly adapting (RA), intermediately adapting (IA), or slowly adapting (SA). Cell capacitance did not differ between WT and Anxa6 KO neurons, whatever their response to mechanical stimulation (Fig. 2a). We observed a 1.8 fold decrease in the percentage of neurons displaying a SA current, mostly in favour of non-responding neurons which were increased 1.9 fold in the KO (Fig. 2b). The minimum stimulation required to elicit a > 40 pA MA current was similar in WT and KO neurons whatever the current kinetics (Fig. 2c). However, the maximum density of transient RA and IA currents when combined was 3.5 fold higher in Anxa6 KO neurons compared to WT neurons. There was a 4.1 fold decrease in the maximum density of SA currents in Anxa6 KO compared to WT neurons (Fig. 2d). When plotted against the displacement produced by the stimulation probe, these observations showed an substantial increase in RA and IA current density and a decrease in SA current density in Anxa6 KO neurons compared to WT neurons (Fig. 2e-h and Fig. S3a-b). These data support a potential contribution of annexin A6 to the SA current.

We next assessed whether overexpressing annexin A6 in DRG neurons had the opposite effect on mechanotransduction to that induced by knocking out Anxa6. Cultured DRG neurons were transfected with a CMV-driven expression vector containing variant 1 of human ANXA6 together with the fluorescent protein tdTomato, or the corresponding empty vector. MA currents were recorded in tdTomato-positive neurons 2 days after transfection. Cell capacitance did not differ between cells transfected with ANXA6 or empty vector, whatever their response to mechanical stimulation (Fig. 3a). Transfection did not alter the percentage of cells with the different types of MA current (Fig. 3b) or...
MA current thresholds (Fig. 3c), whatever the current kinetics. However, we observed a 2.63 fold decrease in the maximum RA and IA current density (Fig. 3d) in neurons transfected with the ANXA6-containing construct compared to those transfected with empty vector. An increase in the maximum SA MA current density recorded in neurons transfected with the ANXA6-containing construct was also observed, although this was not significant (Fig. 3d). The significant decrease in RA and IA current density can be seen in current density / displacement curves whereas for SA currents, neurons transfected with the EV or with the ANXA6 expression vector showed similar currents (Fig. 3e-h and Fig. S3c-d).

Therefore, annexin A6 inhibited transient MA currents in DRG neurons, as shown by the decrease in these currents in neurons overexpressing the human clone, and by their enhancement in Anxa6 KO DRG neurons. Anxa6 deletion was linked to a loss of SA currents, suggesting a possible contribution of annexin A6 to SA mechanotransduction complexes. Gene deletion studies have shown that >80% of rapidly adapting currents and 50% of intermediate adapting currents are lost in the absence of Piezo2 (5). Thus, annexin A6 has a functional link with this channel in sensory neurons. We therefore studied the effects of annexin A6 on MA currents produced by Piezo2 in a heterologous expression system to further explore this interaction.

Expression vectors containing the human ANXA6 or PIEZO2 clones (together with a tdTomato or GFP fluorescent protein, respectively), or their corresponding empty vectors, were co-transfected into ND-C cells (14). Membrane capacitance of recorded cells was similar between the groups (Fig. 4a). Annexin A6 transfection alone did not affect the endogenous MA current threshold, maximum density or current density / probe displacement relationship (Fig. 4b-d). Piezo2 alone produced a low-threshold RA current that is modulated by annexin A6. Indeed, the mechanical threshold of this current increased 2.5 fold in cells transfected with PIEZO2 alone compared to cells transfected with both
**PIEZ02 and ANXA6** (Fig. 4b). The maximum density of Piezo2 current was also reduced by 2.8 fold with **ANXA6** co-expression (Fig. 4c), a decrease that was also significant on current density / displacement curves (Fig. 4d and Fig. S3e). These data confirmed that the effects of annexin A6 on RA in DRG neurons are largely due to inhibition of Piezo2 mediated currents.

In addition to annexin A6, we tested other identified peptide binding partners of NMB-1 for a role in mechanosensation. ATP6V1B2 is the brain isoform of a vacuolar ATP synthase subunit which is expressed in cochlear hair cells and has been linked to a role in hearing (17). However, expression of ATP6V1B2 in HEK293T or ND-C cells did not produce a mechanically activated current (Fig. S4a-d) although co-expression of the subunit with **PIEZ02** partially reduced its peak amplitude similarly to annexin A6 (Fig. S4a-b).

**Overexpression of ANXA6 reduces mechanical pain in a mouse model of osteoarthritis.**

We extended our over-expression studies to a behavioural analysis of mechanosensation in a mouse model of osteoarthritis pain where mechanically evoked pain is particularly problematic. We generated AAV viruses (serotype TT) carrying the human **ANXA6** variant 1 clone upstream of an IRES-GFP construct and under the control of a CMV promoter. Although a sensory neuron-specific promoter would have been preferable for these studies, the size limitations on AAV inserts precluded this approach. AAV-TT carrying **ANXA6** or the empty control virus was delivered intrathecally to infect sensory neuron cell bodies, an approach that has been experimentally validated (18). We used GFP downstream of an IRES in the constructs to show that DRG neurons were effectively transduced by the virus.

Pain thresholds were followed for 12 weeks before a single dose of monosodium iodoacetate (MIA) was injected into the left knee, and behavioural assessment continued for another 3 weeks
whereupon mice were sacrificed to confirm successful transduction of DRG neurons (Fig. S5). AAV-mediated overexpression of ANXA6 decreased noxious mechanical sensitivity without affecting thermal sensitivity (Fig. 5a-b). The 50% paw withdrawal thresholds in response to von Frey filaments (19), a test that characterizes mechanical sensitivity, was similar in the two groups at week 12 (Baseline before MIA injection, Fig. 5c). One week after injection of MIA, all treatment groups showed a significant decrease in 50% threshold compared to the threshold recorded at baseline (Fig. 5d). Control (AAV-TT-IRES-GFP treated) mice showed a further 2.2 fold55% drop in 50% threshold from one week to three weeks post MIA while the 50% threshold of AAV-TT-ANXA6-IRES-GFP treated mice dropped only by 1.13 fold12% during this time period.

Mechanical hypersensitivity in AAV-TT-ANXA6-IRES-GFP treated mice was also attenuated in the weight bearing test, which evaluates mechanical pain by comparing the body weight applied on the MIA-injected paw over the total weight applied on both hind paws. In control treated mice, weight on the ipsilateral paw dropped by 1.15 fold one week after MIA injection and by 1.08 fold at three weeks. This asymmetry was significant compared to baseline at both time points whereas ANXA6-treated mice did not develop a significant weight bearing asymmetry at any time point. These results indicate that annexin A6 overexpression can prevent the onset of the weight bearing asymmetry induced by MIA and suggest that gene therapy delivery of annexin A6 has potential utility in treating human osteoarthritis pain by inhibiting mechanosensation in vivo.

**Discussion**

The present studies demonstrated the ability of annexin A6 to modulate mechanotransduction in sensory neurons. Annexins are a superfamily of Ca\(^{2+}\)-dependent, phospholipid-binding proteins which are involved in diverse functional processes. They are highly evolutionarily conserved; annexin A6, previously known as p68, calelectrin and protein III (20), is a compound gene which appears to derive
from the evolutionarily early splicing of annexin A5 and annexin A10. Annexin A6 links the cellular cytoskeleton to the internal lipid bilayer of the cell membrane through calcium-dependent interactions with negatively charged phospholipids. We identified annexin A6 as a binding target of the biotinylated Conotoxin NMB-1 which blocked behavioural responses to noxious mechanical stimuli and attenuated the persistent component of IA and SA MA currents. We hypothesised that annexin A6 would therefore be a potential candidate for a role in mechanotransduction. Annexin A6 has been suggested to form a cation-selective ion channel in lipid bilayers that is modulated by cGMP (21), or that low pH potentiates the ability of annexin A6 to form cation channels (22).

We found that annexin A6 KO mice had heightened sensitivity to noxious mechanical stimuli and sensory neurons from these mice showed increased RA and IA mechanically-evoked currents. Conversely, overexpressing annexin A6 reduced the RA current produced by the mechanotransduction channel Piezo2 in sensory neurons and heterologous expression systems. Intriguingly, loss of annexin A6 led to reduced SA currents. However, annexin A6 alone does not encode a mechanosensitive current (Fig. 4d). It is thus nonetheless possible that annexin contributes to the complex that forms the SA channel in sensory neurons.

Annexin A6 inhibited RA and IA mechanotransduction currents, suggesting a possible role for annexin A6 in cutaneous mechanosensation. Annexin A6 has been linked to roles in mechanosensory functions in other physiological systems. For example, in mice overexpressing annexin A6 in cardiomyocytes, there is a decrease in their contractility which is presumed to be the result of reduced free intracellular Ca\(^{2+}\) and reduced Ca\(^{2+}\) release upon depolarisation (23). Similarly, in skeletal muscle, annexin A6 regulates the gating properties of the sarcoplasmic reticulum Ca\(^{2+}\) release channel, potentially linking it to a role in excitation-contraction coupling that mediates muscle contraction (24).
Current understanding of the functions of annexin A6 provides insight into the potential mechanisms by which it could regulate mechanotransduction. When Ca$^{2+}$ is low, annexins are soluble monomeric proteins although when intracellular Ca$^{2+}$ levels rise, annexins can bind to phospholipid membranes (15). Membrane binding for annexins has been proposed to require the formation of trimers that aggregate into a hexagonal array surrounding a target protein (15, 25, 26). This aggregation affects membrane properties, for example altering Ca$^{2+}$ mediated membrane fluidity (27-30) and potentially interfering with protein-protein or protein-phospholipid interactions and affecting Ca$^{2+}$ dependent second messenger pathways (31). If annexin A6 can regulate membrane fluidity, increasing membrane rigidity in a similar way as annexin A5 for example (30), it is possible that this could increase the force required to elicit a mechanically activated response. In this situation, it is possible that RA and IA currents, which are typically evoked by a lower intensity of mechanical displacement, would be most sensitive to changes produced by overexpressing annexin A6 in cells. This notion is consistent with the attenuation of RA and IA current amplitude in DRG neurons overexpressing annexin A6. Whether annexin A6 expression is regulated in sensory neurons in physiological conditions is currently unknown but evidence from cardiomyocytes demonstrates the potential for its expression to be dynamically regulated in cells (32). Our gene therapy data suggests that annexin A6-mediated regulation of mechanosensation also has the potential to be clinically useful. Gene therapy offers a target-specific alternative to existing pharmacological interventions and is a viable treatment method for analgesia in the clinic (33). Similarly, in preclinical models, gene therapy for pain relief is effective in multiple pain conditions (18, 34-37) including rheumatoid arthritis (38). We used an AAV serotype with a high tropism for sensory neurons, AAV-TT, to overexpress human ANXA6 in vivo and found that overexpressing ANXA6 attenuated mechanical hypersensitivity in the MIA model of osteoarthritis pain.
Other studies have identified proteins such as TRPC3, TRPC6 and TRPA1 as mechanosensors that function in a context dependent manner (39, 40). Other proteins identified by NMB-1 peptide binding are now being examined for a possible role in mechanotransduction. The data presented suggest that annexin A6 may participate in the channel complex responsible for the SA mechanotransduction current that remains to be characterised. The present study also demonstrates an interesting role for annexin A6 in regulating mechanotransduction that may be clinically useful in terms of gene therapy.

Materials and methods

Ligand blotting assay using DRG and cerebellum membrane protein extracts

Membrane proteins were isolated using a previous published protocol (41). Briefly, DRG (70 ganglia) and cerebellum were excised from P0 rats and homogenized using a glass Teflon homogenizer in homogenization buffer (0.32 m sucrose, 10 mm HEPES, pH 7.4, and 2 mm EDTA plus protease inhibitors leupeptin, pepstatin, phenylmethylsulfonyl fluoride, benzamidine, aprotinin). The homogenates was centrifuged for 15 min at 1000 x g and the supernatant centrifuged at 200,000 x g for 15 min and extracted with lysis buffer 50 mm HEPES, pH 7.4, 2 mm EDTA, protease inhibitors, 4% SDS and 2% Triton X-100. The samples were resolved using SDS-PAGE on 8 - 12% gels and transferred to PVDF or nitrocellulose filters (R&D systems). The membrane was blocked with 0.1% Tween-20 in PBS, pH7.4 (wash buffer) and incubated with biotinylated NMB-1 (2 or 5 µM) for 15 min. The membranes were washed and treated with chilled 4% paraformaldehyde. This crosslinking step enhances the binding of NMB-1(7). The membranes were incubated with alkaline phosphatase-conjugated streptavidin (Chemicon) for 2 hours at room temperature. bNMB-1 was detected using alkaline phosphatase colour reaction using BCIP/NBT kit from Vector Laboratories according to the manufacturer’s instructions. Levamisole Solution (Vector Laboratories) was added to inhibit endogenous alkaline phosphatase activity. The appearance of blue precipitate was monitored on both
bNMB-1 treated and control blots. Control experiments were also carried out where NMB-1 was substituted with goat anti biotin IgG to ensure specificity of NMB-1 blotting assay. Reactive bands on the PVDF or nitrocellulose filters were cut and sent to MRC Clinical Sciences Centre (Imperial College, London UK) for analysis. However, attempts to extract the proteins from the filter and process for LC-MS/MS or perform in situ digestion did not produce sufficient quantities of material required for mass spectroscopy. As an alternative approach, we split into two a mini-gel after resolving proteins samples. One half was electrotransfered onto nitrocellulose filter and processed for bNMB-1 binding; and the other half was silver stained (Silver Stain Kit, Pierce, ThermoFisher). The silver stained half was aligned with the bNMB-1 blot based on protein markers’ position and other landmarks. Gel bands corresponding to the position of the reactive bands on the blots were excised from the gel. The samples were analyzed using LC-MS/MS on an Agilent Q-TOF LC/MS.

**SDS-PAGE and Western blot using antibody recognizing annexin A6**

The samples were resolved using SDS-PAGE on a 7.5% precast gel (Bio-rad) and transferred to a PVDF membrane (Merck). The membrane was blocked with 5% Marvel in 1 x PBST for 1 hr at room temperature then incubated with Anti-annexin VI antibody (Abcam ab31026) at a concentration of 1:500 at 4°C overnight. The membrane was washed in 1 x PBST then incubated with HRP-conjugated goat anti-rabbit IgG (Thermo Scientific) at a concentration of 1:1600 for 1 h 30 min at room temp. After further washing the membrane was immersed in Pierce ECL Western Blotting substrate (Thermo Scientific), and then developed with an exposure time of 1 minute.

**NMB-1 pull-down from ND-C cell extracts**

Seven 10 cm plates of sub-confluent, actively dividing ND-C cells were treated for three days with D-MEM containing 1 mM dibromo-cAMP and 0.2% FBS. The cultures were incubated with 5 µM biotinylated NMB-1 for 3.5 minutes and fixed with chilled 4% PFA for 10 minutes. Extraction buffer
containing protease inhibitors, 0.1% SDS and 1% Triton X-100 was added directly to the plate. The cells were scraped and extracted for three hours on ice. The protein extract was vortexed, spun down and incubated with prepared Neuroavidin gel matrix (ThermoFisher Scientific) overnight at 4 C. The beads were washed three times and the bound proteins eluted according to the manufacturer’s instructions by heating to 95 C for 10 minutes in 2% or 5% β-mercaptoethanol. Control samples were processed as above omitting biotinylated NMB-1.

**LC-MS/MS analysis of NMB-1 binding proteins**

Eluates from NMB-1 pull-down experiments were separated by SDS-PAGE electrophoresis using a NuPAGE 4-12% Bis-Tris gel (1.5 mm x 10 well, Invitrogen) using MOPS buffer. The gel was stained overnight with colloidal Coomassie blue (Sigma). Each lane was excised into 16 bands that were cut into 1-2 mm cubes and in-gel digested overnight using trypsin (sequencing grade; Roche). Peptides were extracted from gel bands twice with 50% acetonitrile/0.5% formic acid and dried in a SpeedVac (Thermo). Peptides were resuspended using 0.5% formic acid were analysed online using an Ultimate 3000 Nano/Capillary LC System (Dionex) coupled to an LTQ FT hybrid mass spectrometer (Thermo Electron) equipped with a nanospray ion source. Peptides were desalted on-line using a micro-Precolumn cartridge (C18 Pepmap 100, LC Packings) and then separated using a 30 min RP gradient (4-32% acetonitrile/0.1% formic acid) on a BEH C18 analytical column (235 μm id x 10 cm) (Waters).

The mass spectrometer was operated in standard data dependent acquisition mode controlled by Xcalibur 2.0. The instrument was operated with a cycle of one MS (in the FTICR cell) acquired at a resolution of 100,000 at m/z 400, with the top five most abundant multiply-charged ions in a given chromatographic window subjected to MS/MS fragmentation in the linear ion trap. All data were processed using BioWorks V3.3 (Thermo Electron) and searched using Mascot server 2.2 (Matrix Science) against a mouse IPI sequence database (June, 2007) using following search parameters:
trypsin with a maximum of 2 mis-cleavages, 20 ppm for MS mass tolerance, 0.5 Da for MS/MS mass tolerance, with 3 variable modifications of Acetyl (Protein N-term), Carbamidomethyl (C), and Oxidation (M). False discovery rates determined by reverse database searches and empirical analyses of the distributions of mass deviation and Mascot Ion Scores were used to establish score and mass accuracy filters. Application of these filters to this dataset resulted in a < 1% false discovery rate as assessed by reverse database searching. Protein hits from all datasets were BLAST-clustered using a threshold of 95% sequence homology over at least 50% of sequence length.

**Proteomics data processing**

All data were processed using BioWorks V3.3 (Thermo Electron) and searched using Mascot server 2.2 (Matrix Science) against a mouse IPI sequence database (June, 2007) using following search parameters: trypsin with a maximum of 2 mis-cleavages, 20 ppm for MS mass tolerance, 0.5 Da for MS/MS mass tolerance, with 3 variable modifications of Acetyl (Protein N-term), Carbamidomethyl (C), and Oxidation (M). False discovery rates determined by reverse database searches and empirical analyses of the distributions of mass deviation and Mascot Ion Scores were used to establish score and mass accuracy filters. Application of these filters to this dataset resulted in a < 1% false discovery rate as assessed by reverse database searching. Protein hits from all datasets were BLAST-clustered using a threshold of 95% sequence homology over at least 50% of sequence length.

**Animals**

All experiments were approved by the UK Home Office and performed under a Home Office project licence (PPL 70/7382). 7-8 weeks old male C57Bl6/J mice, Anxa6 knock-outs or wild-type littermates were used and kept under 12/12 light cycle with food and water ad libitum. Anxa6 global knock-outs were previously generated by disruption of exon 3 with insertion of a neo cassette(42). Experimenters were blind to the genotype or treatment of the animals. AAV viruses were injected intrathecally (10
µl at \(2.4 \times 10^{12}\) VP/ml) 20 min following intravenous injection of mannitol (200 µl at 25%) to potentiate sensory neurons transduction (43). Osteoarthritis was induced by injection of 0.5 mg MIA (monosodium iodoacetate) (Sigma, UK) into the right knee joint cavity under light isoflurane anaesthesia(44).

**Plasmids**

*hANXA6* transcript variant 1 clone (NP_001146) was purchased from Origene (cat. #RC202086) and sub-cloned into a pIRES2-tdTomato vector, originally made from a pIRES2-AcGFP1 backbone. *hPiezo2* (NP_071351 with SNPs rs7234309 and rs3748428) was cloned into an IRES-eGFP vector as previously described (8).

**AAV-TT virus production and purification**

HEK293T cells were co-transfected with a pAAV-CMV-hANXA6-IRES-GFP plasmid containing the *hANXA6* transcript variant 1 clone (NP_001146) upstream a GFP, flanked by the ITRs of AAV2, together with a helper plasmid providing the *rep* and AAV-TT *cap* genes in trans(45). 72 h after co-transfection with PEI-max (Polysciences, 3.5 ml per mg DNA), cells were harvested by centrifugation and lysed by 4 freeze-thaw cycles (from -80°C to 37°C) in lysis buffer (150 mM NaCl, 50 mM Tris pH 8.5) before benzonase treatment (5000 U / cell factory 10 chamber, corresponding to 6320 cm² growth area, 37°C for 30 min). Lysates were cleared by centrifugation and filtered at 0.22 µm. Cell culture supernatant was also harvested and precipitated by addition of ammonium sulfate to 31.3%. The resulting pellet was also resuspended in lysis buffer, treated with benzonase, clarified by centrifugation and filtered.

Recombinant AAV-TT virions were purified by fast protein liquid chromatography (ÄKTApurifier system, GE Healthcare) and an AVB sepharose affinity column (buffer A: PBS, pH 8; buffer B: 0.5 M glycine, pH 2.7). The collected fractions were then dialysed against PBS overnight and viruses were
concentrated with Vivaspin 20 centrifugal concentrators (100 kDa MWCO, Sartorius). Viral genome and capsid titers were determined by real time quantitative PCR and SDS-PAGE, respectively. Primers used for PCR quantification were the following: GFP forward 5’ GAC GGC AAC ATC CTG GGGCAC AAG 3’ and GFP reverse 5’ CGG CGG CGG TCA CGA ACT C 3’. A standard curve was generated by loading known quantities of the linearized pAAV-CMV-ANXA6-IRES-GFP vector plasmid. The vectors were finally diluted to 2.4 x 10^{12} VP/ml in saline before injection.

**Behaviour**

Tactile sensitivity of the plantar surface of animals was measured using the up-down method for obtaining 50% threshold using von Frey hairs (19). Animals were habituated to the von Frey chambers for 2 h before the experiment. Noxious mechanical sensitivity was assessed using Randall Selitto apparatus. Briefly, a blunt probe was applied to the tail with increasing force and the force at which the animal withdrew was recorded(46). The test was performed 3 times for each animal.

Animals' weight distribution was assessed using an incapacitance meter (Linton Instrumentation). Mice were placed in the rearing position, with the upper body in a cylindrical chamber. The average weight applied by each hindpaw on the two sensors was measured over a 5s period of steady behaviour with both paws in contact with the sensors. Ipsilateral/contralateral weight ratios were calculated from 3 averaged consecutive measurements.

**Immunohistochemistry**

DRG and spinal cord were taken from mice following intracardiac perfusion with 4% paraformaldehyde (PFA), post-fixed for 2 h in 4% PFA and cryoprotected overnight in 30% sucrose (all solutions made from phosphate buffer saline). Tissues were embedded in OCT compound and stored at -20°C until 10 and 30 μm cryosections of DRGs and spinal cord, respectively, were performed and mounted onto Superfrost Plus slides (Thermo Scientific). Slides were dried for 2 h, blocked in 10% goat
serum, and incubated overnight with a primary antibody anti-GFP (Abcam #ab290, 1:2000), anti-
ANXA6 (Abcam #ab31026, 1/400), and/or with biotin-conjugated IB4 (Sigma #L2140, 1:500). Slides
were then incubated 2 h with Alexa Fluor 488-conjugated goat anti-rabbit IgG (Abcam #ab150077,
1:1000) and Alexa Fluor 568-streptavidin conjugate (Life technologies, #S-11226, 1:1000). For NMB-
1/annexin A6 interaction studies, lumbar DRG tissue was extracted from 3 mice per genotype and
snap frozen on dry ice in OCT before being stored at -80°C. Sections were prepared as described above
before being incubated in ice cold 4% PFA for 8 mins. Samples were blocked with protein block (Abcam
#64226) for 2 hours at RT before being incubated with 2µM bNMB-1 (or no NMB-1 as control) in
protein block overnight at 4°C. Samples were then treated with FITC-conjugated Avidin (Invitrogen
#434411; 1:1000) at RT for 2 hours. Images were taken on a Leica SP8 confocal microscope.
Fluorescence was quantified in 6-10 sections per mouse using ImageJ software and normalized to
background fluorescence.

Culture and transfection of DRG neurons
Immediately after CO₂ euthanasia, thoracic and lumbar dorsal root ganglions were removed from 6-
12 weeks old mice and subsequently digested for 35 min at 37°C in Ca²⁺- and Mg²⁺-free Hanks’
balanced salt solution (HBSS) containing 5 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid
(HEPES), 10 mM glucose, 5 mg/ml collagenase type XI and 10 mg/ml dispase. After digestion, ganglia
were gently triturated in Dulbecco’s modified Eagle’s medium (DMEM) containing 10% qualified foetal
bovine serum (FBS) using fire-polished glass Pasteur pipettes. When required, neurons were
electroporated using the Neon Transfection System (Life Technologies) according to the supplier’s
recommendations, in 10 µl tips and with 0.6 µg plasmid per reaction, applying 2 x 20 ms pulses of 1100
V. Cells were plated on poly-L-lysine and laminin coated dishes, in DMEM containing 10% FBS and 125 ng/ml 7S nerve growth factor (NGF). Neurons were kept at 37°C in 5% CO₂ and used for patch clamp studies the following two days, or at 48 ± 4 h for electroporated neurons.

**Culture and transfection of ND-C cells**

ND-C cells, a hybridoma between neonatal rat dorsal root ganglia neurons and mouse neuroblastoma (13), were cultured in DMEM containing 10% FBS and kept at 37°C in 5% CO₂. Two days prior to patch clamp recordings, the cells were electroporated using the Neon Transfection System according to the supplier’s recommendations, in 10 µl tips and with 0.6 µg plasmid per reaction, applying 3 x 20 ms pulses of 1100 V.

**Electrophysiological recordings**

Small (< 30 pF) or large (> 30 pF) DRG neurons whose somata were not in contact with those of other neurons, and who were displaying clear fluorescent signal when transfection was performed, were selected for electrophysiological recordings. Patch pipettes were pulled from borosilicate glass capillaries using a PC-10 puller (Narishige Group) and had a resistance of 1.5-3.5 MΩ. The pipette solution contained (in mM): 125 CsCl, 4.8 CaCl₂, 1 MgCl₂, 4 MgATP, 0.4 Na₂GTP, 10 ethylene glycol tetra-acetic acid (EGTA) and 10 HEPES (pH: 7.4 adjusted with CsOH; osmolarity: 310 mOsm adjusted with sucrose). The bath solution contained (in mM): 132 NaCl, 3 KCl, 2.5 CaCl₂, 1 MgCl₂, 10 glucose, 10 HEPES (pH: 7.4 adjusted with NaOH; osmolarity: 310 mOsm adjusted with sucrose). Voltage and current clamp recordings were performed using a MultiClamp 200B amplifier and an Axon DigiData 1440A digitizer (Molecular Devices). Data were recorded and stored using Clampex 10 (Molecular Devices). Recordings were low-pass filtered at 5 kHz and sampled at 20 kHz. Capacity transients were cancelled; however, series resistances were not compensated. Voltages were not corrected for junction potentials and recordings were performed at room temperature. Holding command was set
at -70 mV before and during mechanical stimulation. When required, Vm measurements were performed immediately after achieving whole cell configuration. Off-line analysis and fits were performed using Clampfit 10 (Axon Instruments, Molecular Devices Inc.).

**Mechanosensory current analysis**

Mechanical stimulation of cell bodies was achieved using a heat-polished glass pipette (tip diameter of approximately 2 µm) controlled by a piezo-electric crystal drive (Burleigh LSS-3000) and placed above the centre of the cell soma at an angle of 70° to the surface of the dish. The probe was positioned so that a 10 µm movement did not visibly contact the cell but that an 11 µm movement, considered as a 1 µm stimulus, produced an observable membrane deflection. The probe was moved at a speed of 1 µm/ms. 250 ms mechanical steps were applied every 10 s in 1 µm increments. Currents were characterized as a function of the signal given by the highest stimulation intensity applied before the seal was lost. Cells were considered as non-responding when no current > 20 pA was observed in response to a 12 µm stimulus. Responding cells were then classified regarding to the adaptation kinetics of their currents. Rapidly adapting MA currents had a decay kinetic that was best described by a bi-exponential fit. Intermediately adapting MA currents had a decay kinetic that was best described by a mono-exponential fit. Slowly adapting currents showed minimal decay (<20%) during the 250ms stimulation.

**Data analysis**

Raw data were analysed for statistical significance using either unpaired Student's t-test, one-way or two-way ANOVA with repeated measures, as appropriate, followed by Dunnett’s multiple comparison test, Fisher’s LSD multiple comparisons test, or Chi-Square contingency test as indicated. Analysis were performed using GraphPad Prism 6 software. All values are shown as mean ± SEM.
Supplementary Materials

Fig. S1. Isolation of NMB-1 peptide binding partners.

Fig. S2. Interaction between NMB-1 and annexin A6.

Fig. S3. Representative traces of mechanically activated currents in vitro.

Fig. S4. ATP6V1B2 does not confer mechanosensitivity to HEK or ND-C cells but inhibits Piezo2 current.

Fig. S5. Transduction of DRG neurons using AAV-TT serotype.

Table S1. Top NMB-1 binding candidates in ND-C cells.

Data File S1. NMB-1 binding candidates in DRG neurons

Data File S2. NMB-1 binding candidates from differentiated ND-C cells

References and Notes


34. J. Braz, C. Beaufour, A. Coutaux, A. L. Epstein, F. Cesselin, M. Hamon, M. Pohl, Therapeutic efficacy in experimental polyarthritis of viral-driven enkephalin overproduction in sensory


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Figures legends

**Figure 1. Sensitivity of Anxa6 KO mice to noxious mechanical stimuli.** (a) Response to stimulation of the paw with von Frey hairs in WT and Anxa6 KO mice. (b) Tail withdrawal threshold of WT and Anxa6 KO mice in response to the application of a blunt probe. n = 9 WT and 8 KO mice. **p<0.01 Student’s unpaired t-test.

**Figure 2. Mechanotransduction in Anxa6 KO sensory neurons.** Mechanotransduction was assessed in cultured sensory neurons from WT and Anxa6 KO DRGs. (a) Membrane capacitances of recorded WT and Anxa6 KO neurons with RA, IA or SA mechanogated current and in non-responsive (NR) neurons. (b) Distribution of the recorded neurons based on the MA current displayed. (c) Minimal stimulation intensity required to elicit a > 40 pA MA current in WT and Anxa6 KO neurons. (d) Maximum current density (Imax/Cm) recorded in WT and Anxa6 KO neurons. (e-h) Current density plotted against the displacement of the mechanical probe for RA (e), IA (f) and SA (g) currents and the merged transient RA/IA currents (h). (a, c, d) * p < 0.05, ** p < 0.01, *** p < 0.001 WT compared to Anxa6 by two-way ANOVA followed by Fisher’s LSD multiple comparisons test. (b) * p < 0.05 by Chi-Square test. (c) two-way ANOVA Followed by Sidak p=0.13, 0.80, 0.99 for RA, IA, and SA respectively. (d) two-way ANOVA Followed by Sidak p=0.046, 0.45, and 0.059 for RA, IA, and SA respectively (e, f, g, h) two-way ANOVA p= 0.002, 0.0007, 0.0007, and <0.0001 for RA, IA, SA and RA/IA respectively. n values are given in (b) and represent the number of neurons.

**Figure 3. Mechanotransduction in sensory neurons overexpressing ANXA6.** Sensory neurons from WT mice were cultured and transfected with a human ANXA6-pIRES2-tdTomato construct or the corresponding pIRES2-tdTomato empty vector (EV). Only neurons expressing tdTomato were selected for mechano-clamp experiments. (a) Membrane capacitance of control and ANXA6 overexpressing neurons. (b) Distribution of the recorded neurons based on the MA current displayed. (c) Minimal stimulation intensity required to elicit a > 40 pA MA current in control and ANXA6-overexpressing neurons. (d) Maximum current density (Imax/Cm) recorded in control and Anxa6-overexpressing neurons. (e-h) Current density plotted against the displacement of the mechanical probe for RA (e), IA (f) and SA (g) currents and merged transient RA/IA currents (h). (a, c, d) * p < 0.05, ** p < 0.01, EV compared to ANXA6 by two-way ANOVA followed by Fisher’s LSD multiple comparisons test. (b) Chi-Square contingency test p=0.7452. (e, f, g, h) two-way ANOVA p=0.97, 0.0004, <0.0001, and <0.0001 for SA, RA, IA and RA/IA respectively. n values are given in (b) and represent the number of neurons.
Figure 4. Piezo2 MA current modulation by ANXA6 co-expression. ND-C cells were co-transfected with hPIEZO2-IRES-eGFP and/or hANXA6-pIRES2-tdTomato or their respective empty vectors (EV) IRES-eGFP and pIRES2-tdTomato according to the indicated combinations, and mechanotransduction was assessed by the mechano-clamp technique. (a) Membrane capacitance of cells transfected as indicated. (b) Minimal stimulation intensity required to elicit a > 40 pA MA current in cells transfected as indicated. (c) Maximum recorded MA current density in cells transfected as indicated. (d) Current density plotted against the displacement of the mechanical probe in cells transfected as indicated. (a-d) * p < 0.05, ** p < 0.01, *** p < 0.001 compared to PIEZO2+ EV by one-way ANOVA followed by Fisher’s LSD test (a-c) or two-way ANOVA followed by Fisher’s LSD multiple comparisons test (d). n = 4 EV + EV, 6 EV + ANXA6, 10 PIEZO2+ EV, and 5 PIEZO2 + ANXA6 cells.

Figure 5. Effect of virally-mediated overexpression of ANXA6 in sensory neurons on noxious mechanical sensitivity and mechanical hypersensitivity in the MIA model of osteoarthritis pain. WT mice received one intrathecal injection of AAV-TT viruses containing either a hANXA6-IRES-GFP construct or the IRES-GFP control. Pain thresholds were then monitored for 12 weeks after which osteoarthritis was induced by a unilateral injection of monosodium iodoacetate (MIA) into the knee joint and mechanical hypersensitivity was assessed for another 3 weeks. (a) Noxious mechanical sensitivity of mice injected with the indicated viruses and submitted to the paw pressure test. (b) Sensitivity to noxious heat of mice injected with the indicated viruses. (c) Mechanical hypersensitivity following unilateral OA induction in mice injected with the indicated viruses. (d) Weight bearing asymmetry as evaluated by the weight ratio between ipsilateral and contralateral rear paws in mice injected with the indicated viruses. Data are shown as Mean±SEM; (c-f) two-way repeated measures ANOVA with Dunnett’s multiple comparison test; ** p < 0.01, *** p < 0.001 compared to 0 within AAV-TT- IRES-GFP; + p < 0.05, ++ p < 0.01 compared to 0 within AAV-TT- ANXA6-IRES-GFP; n = 12 mice per group.
Table 1. NMB-1 interacting candidates shared between DRG and ND-C cells. Mouse Uniprot numbers are given. Total matches refer to combined numbers of positive peptide matches for each candidate.

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Figure 1

(a) 50% Paw withdrawal threshold (g) for WT and Anxa6 KO mice.

(b) Force (g) comparison between WT and Anxa6 KO mice, with a significant difference indicated by "**".
Figure 2

(a) Membrane capacitance

(b) MA currents distribution

(c) MA current threshold

(d) Maximum current density

(e) RA current density / displacement

(f) IA current density / displacement

(g) SA current density / displacement

(h) RA/IA current density / displacement
Figure 3

(a) Membrane capacitance ($C_m$)

(b) MA currents distribution

(c) MA current threshold

(d) Maximum current density

(e) RA current intensity / displacement

(f) IA current intensity / displacement

(g) SA current density / displacement

(h) RA/IA current density / displacement
Figure 4

(a) Membrane capacitance

(b) MA current threshold

(c) Maximum current density

(d) Current density / displacement

Legend:
- EV + EV
- EV + ANXA6
- PIEZO2 + EV
- PIEZO2 + ANXA6

Significance levels:
- * p < 0.05
- ** p < 0.01
- *** p < 0.001
Figure 5