Wild-type and Brachyolmia-causing Mutant TRPV4 Channels Respond Directly to Stretch Force*1

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Whether animal ion channels functioning as mechanosensors are directly activated by stretch force or indirectly by ligands produced by the stretch is a crucial question. TRPV4, a key molecular model, can be activated by hypotonicity, but the mechanism of activation is unclear. One model has this channel being activated by a downstream product of phospholipase A2, relegating mechanosensitivity to the enzymes or their regulators. We expressed rat TRPV4 in Xenopus oocytes and repeatedly examined >200 excised patches bathed in a simple buffer. We found that TRPV4 can be activated by tens of mm Hg pipette suction with open probability rising with suction even in the presence of relevant enzyme inhibitors. Mechanosensitivity of TRPV4 provides the simplest explanation of its various force-related physiological roles, one of which is in the sensing of weight load during bone development. Gain-of-function mutants cause heritable skeletal dysplasias in human. We therefore examined the brachyolmia-causing R616Q gain-of-function channel and found increased whole-cell current densities compared with wild-type channels. Single-channel analysis revealed that R616Q channels maintain mechanosensitivity but have greater constitutive activity and no change in unitary conductance or rectification.

In stark contrast with vision, smell, and most tastes, which are based on G-protein-coupled receptors, the mechanical senses (hearing, touch, balance, monitoring blood pressure or systemic osmolarity etc.) are poorly understood in molecular terms. Although prokaryotic mechanosensitive channels are clearly activated directly by membrane stretch force (6), how eukaryotic mechanosensitive channels are activated is still under debate. Models range from direct activation by stretches from membrane and/or cytoskeleton to indirect activation through stretch-produced ligands (7, 8). The mammalian two-pore K+ channels TREK1 and TAASK are activated by both membrane stretch and polyunsaturated fatty acids (PUFAs)2 (9). There, cytoskeletal disruption actually increases mechanosensitivity, suggesting that the cytoskeleton does not transmit stretch force to these channels (10).

Among the Ca2+-permeable transient receptor potential potential channels, yeast TRPY1 (11) and animal TRPV4 (transient receptor potential channel subtype V4) (12) have been studied extensively for their response to osmotic or mechanical stimuli. TRPY1 can be activated directly by suction applied to excised membrane patches (11, 13). However, results from such a test limited preliminary studies found in the TRPV4 literature are inconsistent and even contradictory (1, 2, 14), likely reflecting mechanical complexities of patches excised from animal cells and/or molecular heterogeneity.

Rodent TRPV4 channels were first cloned repeatedly by following hypotonicity-induced Ca2+-signals (1, 2). Rat trpv4 complements the mechano- and osmosensitivity defects of the osm9 mutant worm (15). How osmotic force activates TRPV4 is unclear, however. Besides hypotonicity, this polymodal channel is also activated by PUFAs (16) among other stimuli (12). One study used enzyme inhibitors to block the hypotonicity response and concluded that swelling activates TRPV4 through a specific PUFAs, namely 5,6-epoxyeicosatrienoic acid, produced by an enzyme pathway comprising phospholipase A2 (PLA2) and cytochrome P450 epoxygenase (EPO) (3, 16). This model therefore has the PUFA-producing enzyme(s) or their upstream regulator(s) as the force sensor and TRPV4, a ligand-gated and not a mechanically gated channel. Rat TRPV4 expressed in budding yeast can also be activated by hypotonicity. This finding questions this model because yeast has no PLA2 or PUFAs (17). Because stretching TRPV4-expressing patches should provide a direct test and because TRPV4 expresses strongly in Xenopus oocyte (18), we have undertaken a systematic examination of the possible direct mechanosensitivity of TRPV4 so expressed. Here, we report that TRPV4 is in fact directly mechanosensitive.

Among its many functions, TRPV4 appears to gauge forces sustained by bones. Unloading-induced bone loss is suppressed in trpv4−/− mice (19). Recently, TRPV4 gain-of-function (GOF) mutations were found to cause autosomal dominant blockage of bone development in human. In heterozygotes, clinical manifestations range from post-natal brachyolmia (4), spondylometaphyseal dysplasias, and metatropic dysplasia (5, 20) to infantile and neonatal lethality (5). Expressed in HEK cells, some, although not all, of these GOF alleles present larger whole-cell currents than the wild type (4, 20). No patch-clamp examination of these mutant channels has been reported to date. We conducted an extensive analysis of the brachyolmia-causing R616Q allele and found it to have an increased constitutive activity but also to display direct force activation.

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3 The abbreviations used are: PUFA, polyunsaturated fatty acid; PLA2, phospholipase A2; EPG, cytochrome P450 epoxygenase; GOF, gain-of-function; 4α-PDD, 4α-phorbol 12,13-didecanoate; BPB, bromophenacyl bromide.
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EXPERIMENTAL PROCEDURES

Oocyte Expression—Wild-type or mutant alleles were PCR-amplified using high-fidelity PfuUltra polymerase (Stratagene) and integrated into pGH19, an oocyte-specific cRNA expression plasmid (21). cRNA was synthesized from XhoI-linearized templates using an mMESSAGE mMACHINE T7 kit (Ambion). Stage V and VI Xenopus laevis oocytes were injected with 40 nl of diluted RNA solution. Because TRPV4 expression, particularly of GOF alleles, is toxic to oocytes, 1 μM ruthenium red (Sigma) was added to the ND96 incubation buffer (22). Channel activities increased over several days. Currents were measured after 1–5 days depending on the experimental requirement.

Electrophysiological Techniques—Two-electrode voltage-clamp recording was carried out with HS2A head stages and a VG-2A×100 virtual bath clamp connected to a GeneClamp 500 amplifier interfaced through a Digidata 1440A digitizer acquired using pClamp10 software (all from Axon Instruments). The base bath solution contained 66 mM KCl, 100 mM sorbitol, 1.8 mM BaCl₂, and 5 mM K⁺-HEPES, pH 7.2 (all from Sigma). 4ucharbol 12,13-didecanoate (4α-PDD; 3 μM; Sigma) was added directly to the bath. Sorbitol was omitted from the base solution to form the hypotonic solution. Data were analyzed using both pClamp10 and SigmaPlot 2000 (SPSS) software. Patch clamp was performed with a borosilicate glass pipette with an ~1-μm diameter opening at the tip, recorded in excised inside-out mode with symmetric 98 mM KCl, 1 mM MgCl₂, and 10 mM K⁺-HEPES, pH 7.2 (all from Sigma), unless stated otherwise. Data were acquired at 10 kHz through an eight-pole Bessel filter at 1 kHz, played back at >5 kHz for analysis. Suctions were applied through a syringe and gauged with a PX140 pressure sensor (Omega).

RESULTS

Wild-type and R616Q Macroscopic Currents—We examined whole-oocyte currents under two-electrode voltage clamp. Xenopus oocytes 3 days after injection of 4 ng of wild-type TRPV4 cRNA presented small currents that could be greatly amplified by the addition of the synthetic phorbol ester activator 4α-PDD at 3 μM (n > 30) (Fig. 1a, upper) (23). As expected of TRPV4, these currents show prominent outward rectification (24). Uninjected oocytes (n > 10) or those injected with TRPV4 cRNA with a M680K mutation engineered at the presumed filter (15) showed no such currents (n > 10) (Fig. 1a, lower). Perfusion of a hypotonic solution also activated these currents over minutes, which subsided upon return to isotonicity (n > 10). This cycle could be repeated with the same oocyte except when blocked by the TRP blocker ruthenium red (Fig. 1b). Although a native stretch-activated channel is observed in excised patches from oocytes, hypotonicity fails to activate currents in the majority of whole oocytes examined by two-electrode voltage clamp (25).

Oocytes injected with 4 ng of cRNA of the brachyolm-causing R616Q mutant allele presented very large currents even without 4α-PDD or hypotonicity, the magnitudes of which are greater compared with oocytes injected with 10-fold more wild-type cRNA (Fig. 1c). Despite variations, expressed currents clearly increased with the amount of cRNA injected. Assuming the same production of channels per unit injected RNA, comparisons show that the R616Q channel had ~30 times the steady-state activities of the wild-type channel (Fig. 1d). R616Q-expressing oocytes exhibited macroscopic currents that showed outward rectification, ruthenium red blockage, and stimulation by 4α-PDD and hypotonicity like the wild type (data not shown). Oocytes rarely survived injection of >40 ng of wild-type TRPV4 cRNA or >4 ng of R616Q cRNA. The toxicity of stronger TRPV4 GOF mutants selected through yeast genetics has also been observed (23).

Unitary Conductance—Inside-out patches excised from TRPV4-expressing oocytes exhibited a large unitary conduct-
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a. WT R616Q

+100mV 200A 1s 100ms

R616Q

b. WT R616Q

-100mV -5 0 1 1.5 nPo

100mV

c. nPo

-100mV -100mV

FIGURE 2. Spontaneous microscopic TRPV4 currents. a, spontaneous single-channel activities from a patch excised from an oocyte highly expressing the wild type (left) and a typical one expressing R616Q (right) examined at +100 mV (upper) or −100 mV (lower). Segments of the recordings are displayed at a faster time base showing discrete opening and closing events. C marks the closed levels. b, I-V plots showing the unitary conductances of the wild type (●) and R616Q (○). c, nPo-V plot from a typical wild-type TRPV4-expressing patch showing activation by positive voltages.

ance, not observed in un.injected oocytes or those injected with filter mutant M680K RNA (data not shown). The spontaneous activities of wild-type or R616Q channels tended to run down immediately after patch excision. Channel activity was allowed to relax to a more stable level before quantification (Fig. 2a). I-V plots of both the wild-type and R616Q channels show unitary conductance rectification, being 98 picosiemens outward and 45 picosiemens inward (Fig. 2b), easily distinguishable from the ~30-picoseimien conductance native to the oocyte (supplemental Fig. S1) (26), which sometimes appeared in the same patches. The activity of TRPV4 could also be distinguished from that of this native channel by its 4α-PDD activation and time-dependent blockage by ruthenium red back-filled in the patch pipette (data not shown). Ensemble open probabilities (nPo) of wild-type and mutant channels were small at negative voltages and increased steeply at positive voltages (Fig. 2, a and c). Matching the amount of cRNA injected and incubation period, the R616Q patches invariably had higher basal activities than the wild-type patches. Activation of multiple R616Q channels at positive voltage gave a noisy impression (Fig. 2a); however, the unitary conductance of R616Q could nonetheless be measured from the basal activities of oocytes injected with less RNA (see Fig. 5) or at negative potential (Fig. 2a).

Suction Activates TRPV4—Tens of mm Hg suction pulses applied to membrane patches excised from expressing oocytes directly activated TRPV4 channels (from >200 patches of >50 oocytes). Such activations are shown in the sample traces in Fig. 3a for the wild-type channel and in Fig. 5 (a, b, f, and g) for the R616Q mutant channel. From the same excised patches, after the rundown of spontaneous activities, nPo increased with the pipette suction (Fig. 3, b and c). This increased nPo was maintained over the course of 10–20 s of pressure exposure (e.g. Fig. 5, a, b, f, and g), and in the cases in which pressure sensitivities were assessed before patch excision, no significant changes were observed after excision (data not shown). In no instances could the TRPV4 nPo reach saturation before the applied suction ruptured the patch or the seal at ~60–200 mm Hg. Starting with sizable spontaneous activities, the nPo of R616Q TRPV4 increased 3–4-fold, with a 10–15% S.E. between 0- and 60-mm Hg suction (Fig. 3c, open symbols; and Fig. 5, b and j). A similar rise in the wild-type nPo was seen between 30- and 60-mm Hg suction (Fig. 3c, closed symbols). The R616Q basal activities appeared to cause a left shift of the nPo versus force response curve, but variability and other technical limitations precluded slope comparison (Fig. 3c). The basal activities of the wild type among different patches were low and variable. Estimation of mechanosensitivity as the -fold increase in nPo was therefore carried out with R616Q instead of the wild type.

A PLA2 Inhibitor Blocks General Activation of TRPV4 in Intact Oocytes—It has been reported that hypotonic but not 4α-PDD activation of TRPV4 expressed in cultured cells can be blocked by preincubating them for 20 min in the presence of the PLA2 inhibitor bromophenacyl bromide (BPB) at 100 μM (3). Similar treatment had no effect on the ability of hypotonicity or 4α-PDD to activate TRPV4 in whole oocytes. Because of the possibility that the access of BPB was restricted by the large size and complex geometry of the oocyte, we extended the incubation time for several hours. Indeed, we found that the hypotonic response was greatly inhibited by extended incubation (Fig. 4, a and c). However, we found that the 4α-PDD response was also reduced (Fig. 4, b and d) in fact to a greater extent (Fig. 4e). It thus appears that BPB can greatly inhibit the ability of either 4α-PDD or hypotonicity (but not depolarization) to activate TRPV4 in intact oocytes. In previous work, the apparent specificity of BPB inhibition of the hypotonic response could have reflected a partial inhibition of PLA2 due to the shorter incubation, resulting in obvious inhibition of only the weaker hypotonic response. It is notable that depolarization activation was not substantially affected by BPB in the absence of additional stimulation, underscoring the independence of voltage- from chemical- and mechanical-dependent gating (23). Anthranilic acid, another PLA2 inhibitor, at 20 μM for 15 min (27) also had no effect on the TRPV4 hypotonic response. 1 h of treatment with 50 μM anthranilic acid was toxic to oocytes.

Blockers of 5,6′-Epoxyeicosatrienoic Acid Synthesis Have No Effect on the Direct Mechanosensitivity of TRPV4 in Excised Patches—We quantified the effects of PLA2 or EPG inhibitors on the mechanosensitivities of R616Q and wild-type alleles of oocyte-expressed TRPV4 in excised patches as well. Ten R616Q membrane patches were sampled, and their responses to 60-mm Hg pipette suction were registered as controls. This procedure was repeated on oocytes preincubated in BPB (Fig. 5, upper panels) or 17-octadecynoic acid, an inhibitor of EPG (3,
16) (Fig. 5, lower panels) for 1–3 h. The inhibitors were also present in the recording chamber, where they had direct access to the excised membrane patches. No significant changes in channel responses to suctions were observed upon incubation of these inhibitors. Qualitatively similar results were also obtained with the wild-type channels, with no difference in the mechanosensitivity observed after inhibitor exposure (data not shown). Thus, the direct mechanosensitivity of TRPV4 in excised patches seems to be independent of the production of arachidonic acid or its metabolite 5,6-epoxyeicosatrienoic acid.

DISCUSSION

We found rat TRPV4 to respond directly to pipette suctions that stretch the excised membrane patches. This mechanosensitivity was robust and consistent, observed in wild-type or R616Q mutant patches (Figs. 3 and 5). The response of TRPV4 to stretch force in excised patches bathed in a simple buffer rules out any covalent chemistry dependent on diffusible cytoplasmic factors being involved in transduction.

Inhibition of either PLA2 by BPB or EPG by 17-octadecynoic acid had no effect on the direct mechanosensitivity of TRPV4 in excised patches (Fig. 5). However, prolonged exposure to BPB
strongly inhibited both the hypotonic and 4α-PDD responses of intact oocytes (Fig. 4). This could reflect a Ca$^{2+}$-dependent feedback mechanism that could facilitate an all-or-none response of TRPV4 to stimulation. PLA2 requires Ca$^{2+}$ binding to its C2 domain for activity (28). Threshold initial Ca$^{2+}$ entry through the force-gated TRPV4 could subsequently act to activate the Ca$^{2+}$-dependent PLC, releasing membrane-bound IP$_3$ and DAG, which would subsequently lead to IP$_3$ and DAG-sensitive Ca$^{2+}$ influx. BPB would block such a feedback. Such a Ca$^{2+}$-induced Ca$^{2+}$ release feedback is fairly common and has been documented for the mechanosensitive TRPY1 (11, 13).

Beyond the control of systemic osmolarity (1, 29), TRPV4 has been implicated in other force-related transductions in vivo, including flow-mediated dilation of arteries (30), strain-induced endothelial cell reorientation (31), viscosity-coupled epithelial ciliary activity (32, 33), and osteoclast response to weight load on bones (19). In these instances, a direct mechanosensitivity of the TRPV4 channel itself provides the simplest explanation.

We encountered patch-to-patch heterogeneity, which may reflect differences in patch elasticity or other geometric and mechanical complexities of the membrane (34) with its subtending cytoskeleton (35, 36) and/or heterogeneity of the channel populations (e.g. degree of phosphorylation?). These intricacies are beyond the ability of the present technical resolution to sort out and may underlie the statistical variances reported within studies (3, 16) and contradictory claims between studies (1, 2, 14) found in the TRPV4 literature.

Recent discoveries on heritable human diseases due to TRPV4 mutations illustrate the medical importance of TRPV4 (4, 27). We found the brachyolmia-causing R616Q allele to retain wild-type unitary conductance and rectification but to have constitutive activities in the absence of applied stimuli (Figs. 1–4). R616 is predicted to be at the base of the fifth transmembrane helix (S5) in the TRPV4 subunit. Recent forward genetic screening for TRPV4 mutations that are toxic to yeast yielded GOF alleles nearby (L619P, L624P). These yeast-selected mutations have similar but much stronger GOF phenotypes than R616Q (23). These findings support a general Shaker-like topology for TRPV4, with S5 being part of the main gate. Because the mutant channel has direct mechanosensitivity and constitutive activities, it likely behaves in osteoclasts or other relevant cells as a wild-type channel under mechanical stress, leading to the GOF phenotypes in vivo (4, 5). High-resolution analyses of other disease-causing alleles under patch clamps should further our understanding of the disease process.

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