

A Novel Non-Neuronal *hSK3* Isoform with a Dominant-Negative Effect on *hSK3* Currents

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Key Words

Potassium channel • Alternative splicing • SK3 • KCNN3

Abstract

We have identified a *hSK3*-transcript, *hSK3_ex1c*, which is generated by alternative splicing. Isoform *hSK3_ex1c* lacks the cytosolic N-terminus and the first transmembrane helix and is exclusively expressed in non-neuronal tissues. *hSK3* transfected tsA cells showed a Ca²⁺-activated K⁺ current in patch-clamp experiments, whereas *hSK3_ex1c* transfected cells and cells co-transfected with both isoforms did not. We fused both isoforms to fluorescence proteins and observed *hSK3* localization predominantly in the plasma membrane. The co-expression of *hSK3* + *hSK3_ex1c* resulted in their cytoplasmic co-localization. Thus, *hSK3_ex1c* has a dominant-negative effect on *hSK3* by preventing its transport into the plasma membrane.

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Introduction

Small conductance calcium-activated potassium channels (SK channels) form a distinct subfamily of potassium channels, which comprises three members, SK1 to SK3 [1]. Their subunits have six transmembrane

helices, a P-loop region between the fifth and sixth transmembrane helix and cytosolic N- and C-terminal regions. All members are voltage-independent and their activation by elevated cytosolic Ca²⁺ concentrations is mediated via calmodulin [2-5]. They are involved in modulating neuronal firing [6, 7].

Alternative splicing is assumed to increase the functional diversity of potassium channels and might also be involved in the regulation of whole-cell current amplitude [8]. Alternative splicing within the SK channel subfamily has been shown only for SK1 [9, 10] and SK3 channels [11, 12]. In the case of SK3 channels two isoforms have been described. One isoform shows an altered outer pore region which changes its pharmacokinetic properties [12]. The other one lacks a cytosolic N-terminal region and the first transmembrane helix. This isoform is shown to have a dominant-negative regulatory effect on SK channels [11] and is shown to be expressed in neuronal tissues.

Here we describe an additional non-functional *hSK3*-isoform, *hSK3_ex1c*, which is generated by alternative splicing and is co-expressed with *hSK3* in lymphocytes, lung and skeletal muscle but contrary to the isoform previously reported by Tomita et al. [11], *hSK3_ex1c* is absent in neuronal tissues. Since this isoform is shown to have a dominant-negative effect on

hSK3, it possibly regulates SK3-channel activity in non-neuronal tissues.

Materials and Methods

If not assigned otherwise, all standard chemicals were obtained from Sigma (Germany). Scyllatoxin was obtained from Latoxan (France).

RT-PCR

Reverse transcription (RT) was carried out on 2 µg of total RNA from adult and fetal whole brain, lung, heart, skeletal muscle (Research Genetics) and a B-lymphocyte cell line (extracted using the RNeasy extraction kit, Qiagen) using superscript II (Invitrogen) according to manufacturer's protocol. Furthermore, as an additional template, a cDNA library from six to eight week old whole human embryos (gift from Dr. Francis Poulat, CNRS France) was used. *hSK3_ex1c* transcripts were amplified using the primers III-F1 (cctgccacgcgacatctg) and R10 (gacaccttccacagatg) and *hSK3* transcripts with the primers *hSK3*-F (gctctctggggtttg) and R10. Primers GAPDH-F (ggcgctggtcaccacgg) and GAPDH-R (ctgatgatcttgacgtgtg) were used as controls for the RT reaction and to test for any contaminating genomic DNA. PCR was carried out in a RT reaction mix without RNA, water and genomic DNA as a negative control.

cDNA construction

IMAGE clone 686525 was obtained from a cDNA library from human tonsillar cells, which were enriched for B cells and was distributed by the RZPD (AA255937, resource centre of the German human genomic project, RZPD: IMAGp998C061678). The entire coding region of isoform *hSK3_ex1c* was amplified from the IMAGE clone 686525 using the primers *hSK3_1c*-F1 (aaaactcgagccaccatggagagacataaaggac) and *hSK3*-R1 (aaatggatccccgcaactgctgaactgtgta) and cloned into pDsRED2-N1 (Clontech) via XhoI/BamHI sites. It encodes the fusion protein *hSK3_ex1c*-DsRED. The entire coding region of *hSK3* was amplified with the primers *hSK3*-F1 (aaatggatccccgcaactgctgaactgtgta) and *hSK3*-R1 and cloned via BamHI/XhoI into pEGFP-N1 (Clontech). It encodes the fusion protein *hSK3*-eGFP. In *hSK3_ex1c*-DsRED and *hSK3*-eGFP the fluorescent protein is fused to the C-terminal region of *hSK3* and *hSK3_ex1c*. The N-terminal coding part of *hSK3_ex1c* was isolated by HindIII/SfiI digestion from clone *hSK3_ex1c*-DsRED and was used to replace the HindIII/SfiI fragment in the clone pHygro-*hSK3*Q19 (a gift from Dr. Heike Jäger, University of Ulm) in order to generate an untagged *hSK3_ex1c* construct.

Cell culture

tsA201 cells (tsA cells), a HEK cell line stably transfected with the SV40 T-antigen (formerly called s93/tsA1609neo) were used for all experiments [13, 14]. tsA cells were cultured in MEM medium containing glutamax-I and Earle's salts (GIBCO,

Germany) and 10 % fetal bovine serum. A mixture of 0.5 µg plasmid DNA encoding pEGFP-N1 (Clontech) and 2 µg of plasmid DNA (pHygro-*hSK3_ex1c* and pHygro-*hSK3*Q19) were transfected into tsA cells using FUGENE 6 (Roche) according to the manufacturer's protocol. Plasmids *hSK3_ex1c*-DsRED and *hSK3*-eGFP were transfected without any transfection marker. Three to four days after transfection, cells were suspended and transferred to poly-lysine coated glass slides prior to further usage.

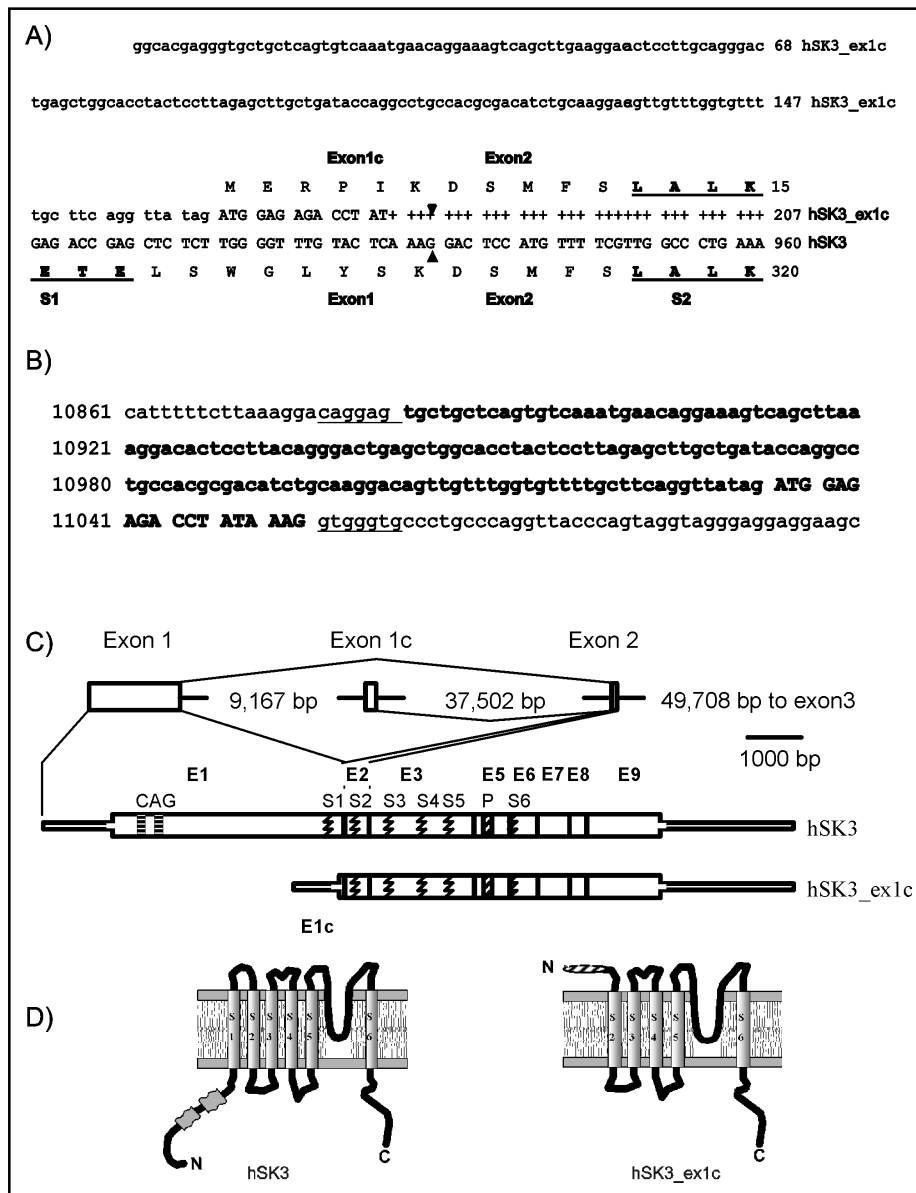
Patch-clamp experiments

Experiments were carried out using the whole-cell recording mode of the patch-clamp technique [15]. Electrodes were pulled from glass capillaries in two stages and fire-polished to resistances of 3.5 to 4 MΩ when filled with internal solution (in mM: 135 potassium aspartate, 2 MgCl₂, 10 HEPES, 10 EGTA and 8.7 CaCl₂ corresponding to [Ca²⁺]_{free} of 1 µM, pH 7.2). Membrane currents were measured with an EPC-9 amplifier (HEKA elektronik, Germany) using Pulse and Pulsefit (HEKA elektronik, Germany) as acquisition and analysis software. The cytosolic Ca²⁺-concentration was adjusted to [Ca²⁺]_{free} = 1 µM by whole-cell dialysis with internal solution. Na⁺-solution (in mM: 160 NaCl, 4.5 KCl, 2 CaCl₂, 1 MgCl₂, 5 HEPES, pH 7.4) and K⁺-solution (in mM: 164.5 KCl, 2 CaCl₂, 1 MgCl₂, 5 HEPES, pH 7.4) were used as external solutions. The membrane potential was held at -80 mV (in Na⁺-solution) or 0 mV (in K⁺-solution), clamped to -120 mV for 50 ms followed by 400 ms ramps from -120 to +60 mV. In between ramps the potential was clamped again to -80 mV (in Na⁺-solution) or 0 mV (in K⁺-solution). Whole-cell conductance was calculated from the slope of the ramp current measured between -90 mV to -70 mV. The whole-cell conductance observed in the presence of Na⁺-solution was presumed as background conductance and therefore subtracted from those conductances, found in the presence of K⁺-solution with and without scyllatoxin (ScTX). The fractional conductance was defined as the ratio between the conductance found in K⁺-solution and the conductance observed in K⁺-solution with and without ScTX. Data are given as mean ± SEM.

Intracellular localization of *hSK3*-isoforms

Cells were washed with PBS buffer and were observed in PBS buffer without any fixation. For selecting cells, fluorescence was excited using a 100 W mercury lamp. GFP and DsRED fluorescence were assessed independently (GFP: Excitation 465 nm, band width 15 nm; emission 520 nm, long pass filtered; DsRED: Excitation 520 nm, band width 20 nm; emission 630 nm, band width 30 nm). Confocal assessment was carried out using the Laser Scanning System Radiance 2000 (Biorad) attached to an inverted microscope (Nikon Eclipse TE300, Nikon). A PlanApo 60X oil objective (Nikon) was used for all confocal measurements. Data recording and analyses were carried out using the Lasersharp 2000 software package (Biorad). GFP fluorescence was excited with the 488 nm line of an argon laser and emission was measured using an HQ510/30 filter (Nikon). DsRED fluorescence was excited with the 543 nm line of a green HeNe-laser and emitted light was long pass filtered using a E600/LP filter (Nikon).

Fig. 1. Isoform *hSK3_ex1c* is generated by alternative splicing. A) cDNA and protein sequence comparison of *hSK3* and *hSK3_ex1c*. Small letters represent untranslated and capital letters translated regions. Identical nucleotides are shown as + and exon boundaries with arrowheads. Exons 1, 2, and 1c are assigned. Protein sequences according to the shown cDNA regions are given. Amino acids within the transmembrane helices S1 and S2 are shown in bold letters and are underlined. B) Genomic sequence spanning exon 1c. The cDNA region from positions 10 to 180 (shown as bold letters) is flanked by typical splice acceptor and donor sites (underlined). C) Organization of the genomic region spanning exons 1, 1c, and 2 and a scheme for alternative splicing. Distances between the exons are given. Schemes for both cDNAs are shown below. Broad boxes represent the ORF and narrow boxes the UTRs. The Exons (E1, E1c, to E9), both CAG repeats and transmembrane helices S1 to S6 and the P-loop are shown. D) Presumed topology of *hSK3* and *hSK3_ex1c* subunits.



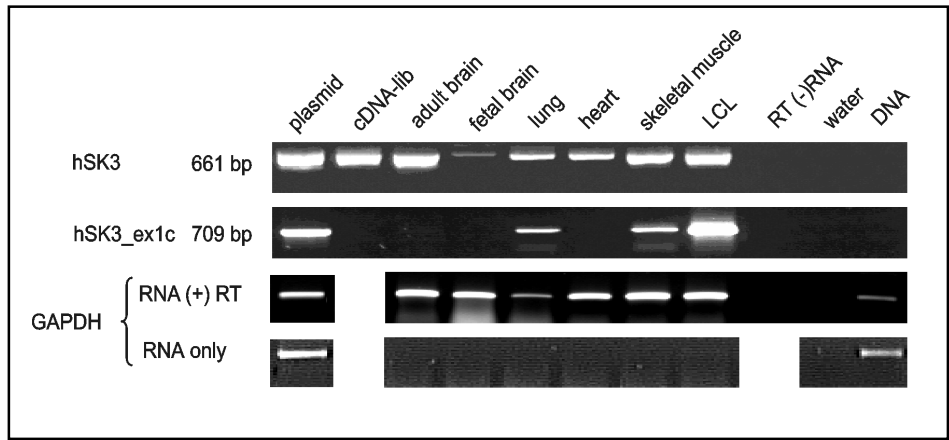
Results

The insert of IMAGE clone 686525 spanned 1984 bp. Its sequence is identical to the original known *hSK3* sequence (Accession no.: AJ251016) 3' from position 928 of the *hSK3* ORF but it differs 5' from this position (Figure 1A). Except for a 10 bp region from the 5' end of the IMAGE clone, the differing cDNA sequence overlaps with the genomic sequence (Accession No.: AF336797) from positions 10882 to 11052 and is flanked by typical splice acceptor and donor sites (Figure 1B). The remaining 10 bp 5' region of the cDNA could not be detected in the genomic sequence, neither by blast search, nor by sequence comparison with the sequence of the *hSK3/KCNN3* gene (AF336797). This might indicate that there

may be an additional, still unknown exon within the *hSK3/KCNN3* gene. The detected transcript represented by IMAGE clone 686525 was generated by alternative splicing and we have named it *hSK3_ex1c* (Figure 1C). The longest ORF in the newly identified cDNA starts at position 163 (Figure 1A) and encodes an *hSK3* isoform that lacks the first transmembrane helix and the cytosolic N-terminal region of *hSK3* (Figure 1D). Its extracytosolic region is 7 amino acids longer than that of *hSK3-1B* [11].

The expression of *hSK3* and *hSK3_ex1c* was investigated by RT-PCR (Figure 2). Transcripts encoding *hSK3* were observed in all tissues tested as well as in the human embryonic cDNA library. Contrary to this, *hSK3_ex1c* transcripts were only observed in lung, skeletal muscle and the B-lymphocyte cell line, but were

Fig. 2. Expression of *hSK3* and *hSK3_ex1c* isoforms using RT-PCR. The expected fragment length of the PCR products and the transcripts' names are given on the left hand side. RNA sources are given above each lane (cDNA: cDNA library form six- to eight-week old human embryos, LCL: lymphocyte cell line). Plasmid DNA (plasmid) was used as positive control. PCR was carried out on the reverse transcriptase reaction mix without RNA (RT(-)RNA), water, and genomic DNA (DNA) were negative controls.



In order to confirm the reverse transcriptase reaction, PCRs were carried out using glycerin-aldehyde-dehydrogenase specific primers (GAPDH; RNA(+))RT). Since these primers also amplify genomic DNA, they were used to test for contaminating genomic DNA (GAPDH; RNA only) using the tested RNAs without prior reverse transcriptase reaction as a template.

absent in the cDNA library derived from 6 to 8-week old human embryos, heart, fetal and adult brain. Therefore we were interested in the functional properties of the newly identified isoform.

Since both isoforms were shown to be co-expressed in several tissues we also investigated the effect of *hSK3_ex1c* expression on *hSK3*, by co-expressing them in tsA cells (Figure 3). The data represent results from three independent transfection experiments. tsA cells expressing isoform *hSK3* alone showed a whole-cell conductance of 1.6 ± 0.3 nS ($n=7$) with Na^+ -solution as bath solution, which was increased to 5.4 ± 1.2 nS ($n=7$) when the external solution was exchanged to K^+ -solution. Current amplitudes corresponding to much higher whole-cell conductances were described in a previous report for rSK3 expressing HEK293 cells [16]. Even in this study the published current amplitudes varied between different experiments. Therefore, the comparable low current amplitudes observed here are most likely due to the applied expression system. In the presence of 10 nM ScTX in K^+ -solution, a mean whole-cell conductance of 1.9 ± 0.4 nS ($n=7$) was observed.

Whole-cell conductances found with Na^+ -solution as external solution were assumed to be background conductances and were therefore subtracted from those observed in K^+ -solution with and without ScTX. After subtracting the background conductance, the fraction of remaining conductance in the presence of 10 nM ScTX in K^+ -solution was calculated as the fraction of whole-cell conductance with K^+ -solution as bath solution to be 0.22 ± 0.04 nS ($n=7$). *hSK3_ex1c* expressing cells showed a whole-cell conductance of 0.6 ± 0.1 nS ($n=12$) in Na^+ -solution which did not significantly increase in K^+ -

solution (0.8 ± 0.1 nS; $n=12$). A similar situation was observed, when tsA cells co-expressed both isoforms (whole-cell conductance of 0.4 ± 0.1 nS; $n=12$ for Na^+ -solution and 0.6 ± 0.1 nS; $n=12$ for K^+ -solution). This indicates, that the isoform *hSK3_ex1c* is able to reduce SK3 currents, if it is co-expressed with the functional *hSK3* isoform. A similar effect was described for the *hSK3-1B* isoform [11]. In that case it was shown that *hSK3-1B* was able to trap *hSK3* subunits in the cytoplasm, which could explain the reduction of SK3 whole-cell currents. We assumed a similar mechanism for the isoform *hSK3_ex1c* described here. Therefore we investigated the intracellular localization of *hSK3* and *hSK3_ex1c*. We fused both isoforms at their C-termini to different fluorescent proteins and expressed them in tsA cells (Figure 4). The data represent the results from two independent transfection experiments. Cells expressing isoform *hSK3* fused to eGFP (*hSK3-eGFP*) showed a whole-cell conductance of 21.2 ± 8.2 nS ($n=13$) with Na^+ -solution and 63.0 ± 16.4 nS ($n=13$) with K^+ -solution as external solutions. This conductance in K^+ -solution was reduced by 10 nM ScTX (27.1 ± 8.8 nS; $n=13$), that corresponds to a fractional conductance of 0.16 ± 0.04 ($n=13$) of the remaining current, when the background conductance is subtracted. Interestingly, the whole-cell conductance of cells expressing *hSK3-eGFP* was strongly increased compared to those cells expressing the wild type *hSK3* without the fused GFP. Cells expressing isoform *hSK3_ex1c*-DsRED (*hSK3_ex1c* fused with its C-term to DsRED) did not show any increased whole-cell conductance after changing the bath solution from Na^+ -solution (0.3 ± 0.1 nS; $n=9$) to K^+ -solution (0.7 ± 0.1 nS, $n=9$). This, however, was the case

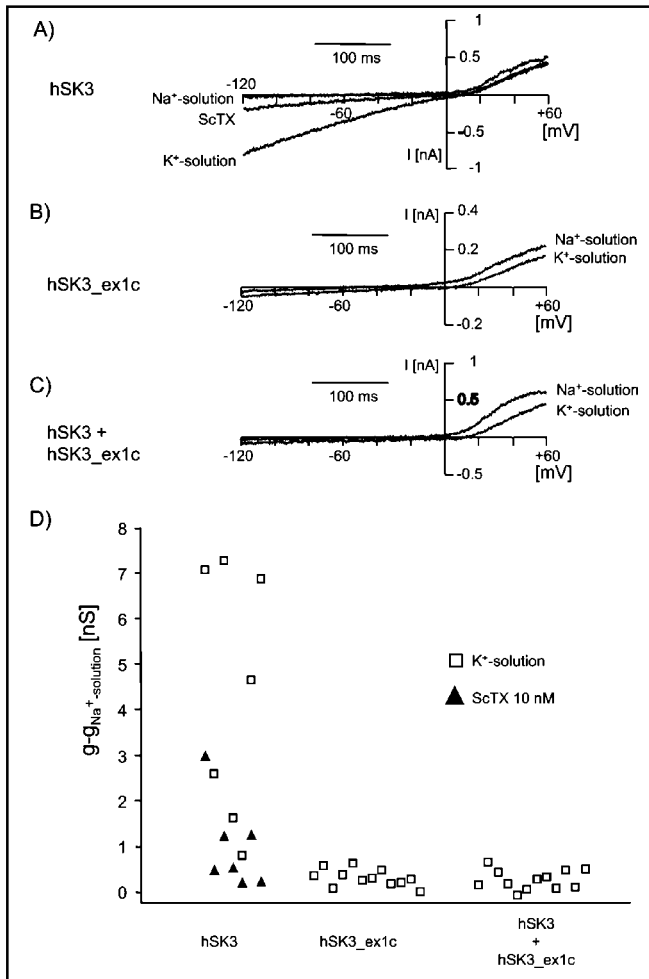


Fig 3. Expression of isoforms *hSK3* and *hSK3_ex1c* in tsA cells. Isoforms *hSK3* and *hSK3_ex1c* were transiently expressed in tsA cells and currents were measured in the whole-cell mode of the patch-clamp technique with 1 μ M free internal Ca^{2+} and Na^+ -solution and K^+ -solution as external solutions. SK3 currents were blocked by adding 10 nM ScTX to K^+ -solution. Representative current traces for cells expressing *hSK3* (A), *hSK3_ex1c* (B) and for cells co-expressing both isoforms (*hSK3* + *hSK3_ex1c*; C) are shown above. Whole-cell conductances found for each experiment are shown in diagram D. Whole-cell conductance found in Na^+ -solution was assumed as background conductance and was therefore subtracted from currents measured in K^+ -solution with and without 10 nM ScTX. 10 nM ScTX blocks SK3 currents. Cells expressing *hSK3_ex1c* and those co-expressing both isoforms did not show conductance above 1 nS and therefore ScTX was not tested on these cells.

when *hSK3-eGFP* and *hSK3_ex1c-DsRED* were co-expressed (1.9 ± 0.4 nS in Na^+ -solution and 10 ± 2.7 nS in K^+ -solution; $n=12$). The remaining whole-cell conductance in K^+ -solution in the presence of 10 nM ScTX was found to be 5.5 ± 1.7 nS ($n=8$), which

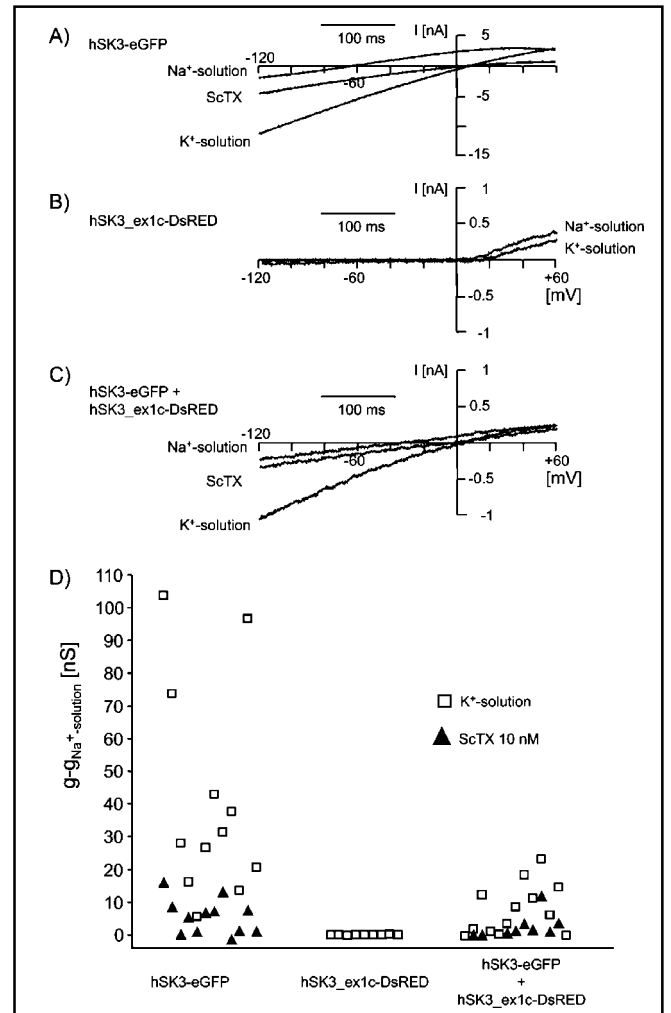
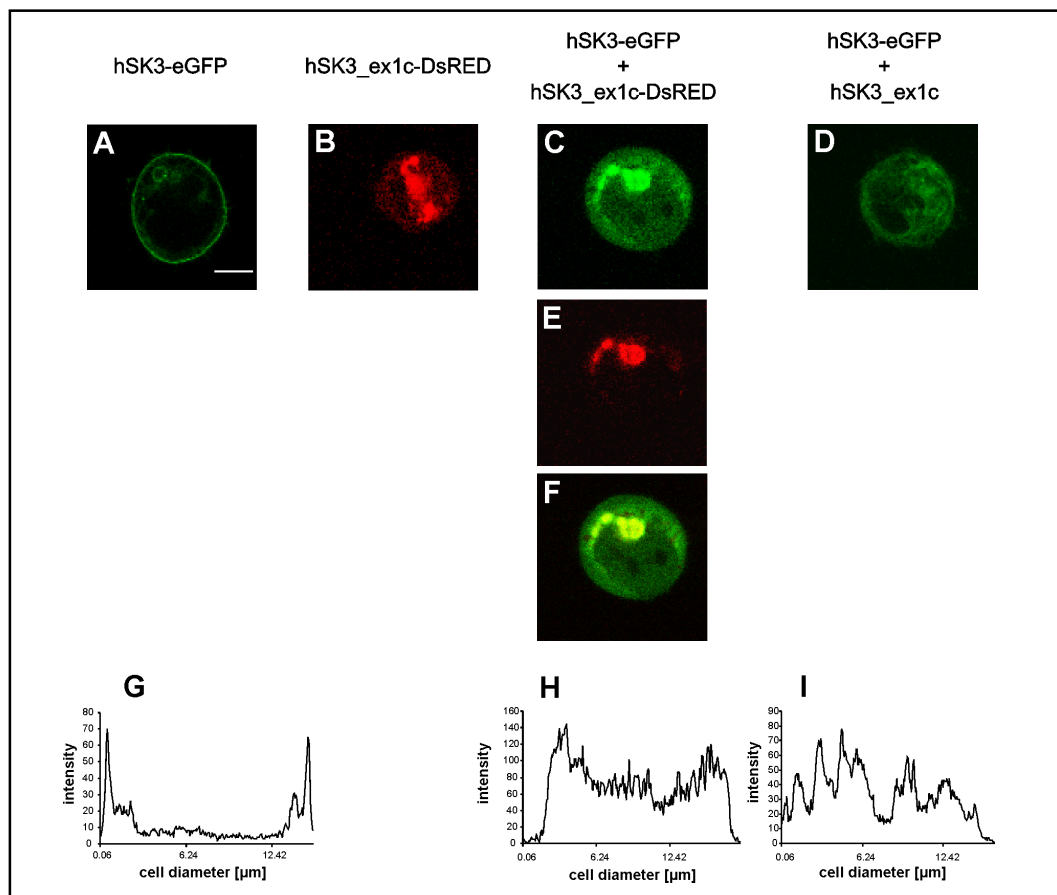


Fig 4. Expression of isoforms tagged with fluorescence proteins. Isoforms fused with their C-terms to fluorescence proteins behave the same as the wild type isoforms, when expressed in tsA cells. Currents were measured as described. Na^+ - and K^+ -solution with and without 10 nM ScTX were used as external solutions. Representative current traces for cells expressing *hSK3-eGFP* (A), *hSK3_ex1c-DsRED* (B) and for cells co-expressing both isoforms (*hSK3-eGFP* + *hSK3_ex1c-DsRED*; C) are shown. Whole-cell conductances for each experiment are shown in the diagram below (D). Whole-cell conductance found in Na^+ -solution was subtracted from the one in K^+ -solution with and without 10 nM ScTX. Cells expressing the GFP tagged isoform *hSK3-eGFP* showed conductances up to 110 nS. These currents were blocked by 10 nM ScTX. Contrary to that, cells expressing *hSK3_ex1c-DsRED* did not show any significantly increased whole-cell conductance. Cells co-expressing *hSK3-eGFP* + *hSK3_ex1c-DsRED* showed a significant whole-cell conductance in K^+ -solution. These currents were blocked by 10 nM ScTX

corresponds to a remaining fractional conductance of 0.14 ± 0.05 ($n=8$) (after subtraction of background conductance).

Fig 5. Intracellular localization of *hSK3* isoforms. *hSK3*-eGFP is shown as green fluorescence in A,C and D. *hSK3_ex1c*-DsRED is shown as red fluorescence in B and E. Merged images for GFP and DsRED fluorescence of co-expressed *hSK3*-eGFP and *hSK3_ex1c*-DsRED are shown in F. The averaged intensity histograms for GFP fluorescence are shown in G-I. Thus the isoform *hSK3_ex1c* prevents *hSK3* translocation to the plasma membrane. Scale bar 5 μm .



The mean whole-cell conductance observed for cells expressing the GFP-tagged *hSK3* isoform, *hSK3*-eGFP, is approximately 12 fold higher compared to cells expressing wild type *hSK3* channels. Even when *hSK3*-eGFP is co-expressed with *hSK3_ex1c*-DsRED, significant whole-cell currents were observed, which were carried by SK3 channels. This may indicate an incomplete inhibition of *hSK3* channels by the *hSK3_ex1c*-DsRED construct. However, since the SK3 current varies strongly between different cells co-expressing the tagged isoforms, it is much more likely, that the incomplete inhibition is due to the higher expression level of functional *hSK3*-eGFP rather than to the fact that the fused fluorescence proteins interfere with subunit assembly.

The intracellular localization of *hSK3*-eGFP and *hSK3_ex1c*-DsRED were studied using confocal laser scanning microscopy (Figure 5). The fluorescent pattern of cells expressing *hSK3*-eGFP alone (Figure 5A) was strikingly different from those co-expressing *hSK3*-eGFP and *hSK3_ex1c*-DsRED (Figure 5C), or with the untagged isoform *hSK3_ex1c* (Figure 5D). Whereas *hSK3*-eGFP-expressing cells showed a sharp annular distribution of fluorescence, the co-expressing cells show

a more uniform distribution. This observation is confirmed by the average intensity histograms (Figure 5G-I), which represent the average intensity along five axes drawn through the cell. The yellow colored regions in the merged images of cells co-expressing both fluorescence labeled isoforms indicated their co-localization (Figure 5F). Cells expressing the *hSK3_ex1c*-DsRED construct alone, showed an intracellular speckling fluorescence pattern that aggregates in some regions (Figure 5B). This indicates that isoform *hSK3_ex1c* is not incorporated into the plasma membrane. In contrast, such an incorporation has been shown for the functional isoform *hSK3* (Figure 5A). In addition, *hSK3_ex1c* seems to be able to prevent the incorporation of *hSK3* subunits into the plasma membrane, which could explain the reduction of Ca^{2+} -activated K^{+} current, as has been shown for *hSK3*-1B [11].

Discussion

The recently reported isoform *hSK3*-1B was described as an isoform having a dominant-negative regulatory effect on SK currents [11]. This isoform was shown to be generated by alternative splicing and was

presumed to encode a shortened SK3 subunit, which is lacking the first transmembrane-spanning S1 helix and the cytosolic N-terminal region [11]. Here we describe a similar *hSK3* isoform, *hSK3_ex1c*, which is also generated by alternative splicing. As in the case of *hSK3-1B*, *hSK3_ex1c* lacks a cytosolic N-terminus and the S1 helix, but contrary to *hSK3-1B*, the longest predicted ORF of *hSK3_ex1c* transcripts encodes a protein, which is extended towards its extracytosolic N-terminus by seven amino acids. Like *hSK3-1B*, *hSK3_ex1c* is able to suppress SK3 currents by trapping functional isoforms in the cytoplasm.

The physiological role of these isoforms is not yet known. An inactive isoform of a cardiac potassium channel K_vLQT_1 has been described [17]. It is assumed that this isoform could play a role as a regulator for whole-cell K_vLQT_1 -current amplitude. This assumption was confirmed, since mutations in this isoform were shown to correlate with inherited cardiac arrhythmias [18]. The dominant-negative effect of the inactive K_vLQT_1 isoform is abolished in the presence of the β -subunit IsK [18]. By analogy, similar roles could be assumed for the inactive *hSK3* isoforms, *hSK3-1B* and *hSK3_ex1c*. In addition the existence of an as yet unknown β -subunit could also be assumed, which influences the dominant-negative effect of *hSK3-1B* and *hSK3_ex1c*.

In principle, the current reduction in the experiments with *hSK3* channels co-expressed with *hSK3_ex1c* could be due to the fact that *hSK3_ex1c* subunits, although non-functional, could still bind calmodulin. SK channel activation has been shown to be mediated by calmodulin, which constitutively binds to the C-terminus of SK channel subunits [2-4]. This binding of calmodulin to *hSK3_ex1c* subunits would compete with calmodulin-*hSK3* channel interaction resulting in less functional *hSK3* channels that could be activated by Ca^{2+} binding to calmodulin. However, in the experiments with the *hSK3-eGFP*, we observed an approximately 12-fold increase in functional *hSK3* channels. This suggests that the amount of calmodulin in the cell is sufficient to mediate the activation of such a high number of *hSK3* channels. The *hSK3_ex1c* subunits would have to bind at least this amount of calmodulin to account for the reduction in functional *hSK3* channels via that mechanism. Therefore it seems unlikely that the mechanism of complete current reduction in the co-expression experiments is due to a lack of calmodulin being able to act on *hSK3* subunits.

An interesting observation is the strongly increased current amplitude in cells expressing *hSK3-eGFP*. We have no explanation for this phenomenon. This effect is

due to a higher channel expression level of functional *hSK3-eGFP* channels compared to the wild type channel. One might speculate that the increase in functional channels is due to an increase in protein synthesis or protein stability. Alternatively, the eGFP-tag might stabilize SK3 subunits in the membrane or may increase their transport into the plasma membrane. Contrary to *hSK3-1B*, *hSK3_ex1c* is not expressed in brain tissues, but is expressed in skeletal muscle. Therefore, it could play a role in modulating SK3 whole-cell current amplitude in, for example, skeletal muscle. So far nothing is known about the regulation of the expression of SK3 transcripts. However, the fact that *hSK3-1B* and *hSK3_ex1c* show different tissue distributions implies that their expression might be regulated independently. This could provide a differential modulation of *hSK3* whole-cell current amplitude. The expression of *hSK3_ex1c* may be the reason why there are no apamin binding sites [19, 20] nor SK3 currents found in adult skeletal muscle [21]. Interestingly, functional SK3 channels are observed after denervation of skeletal muscle and in skeletal muscles of patients with myotonic dystrophy [21-23]. Such an up-regulation of SK3 channel activity might be explained at least in part by an alteration of *hSK3_ex1c* expression and not solely via an up-regulation of *hSK3* transcripts [22]. The inducible up-regulation of SK3 channels in a transgenic mouse is shown to induce abnormal respiratory response to hypoxic conditions and compromised parturition [24]. A suppression of SK3 channels in the same mouse model leads to an increase of the frequency of phasic urinary bladder smooth muscle contraction [25]. This report is in line with the observation, that the selective blockage of IK1 and SK3 channels suppresses the vasodilation, which is mediated by the endothelium-derived hyperpolarizing factor EDHF [26]. These findings underline the importance of SK3 channel activity as a determinant for smooth muscle contractility. Furthermore, all SK channels are thought to be involved in regulating the glucose-dependent insulin release of pancreatic islet cells [27]. All these findings emphasize the importance of SK3 channel regulation for many physiological events. An additional possibility of SK3 channel regulation beside SK3 channel expression could be important for a more robust fine tuning of cellular function than it is possible solely by an expressional regulation of SK3 channels. Therefore, isoform *hSK3_ex1c* could play a vital role in many homeostatic processes like smooth muscle contractility and control of blood pressure, in respiratory rhythmogenesis and the control of glucose serum level.

Note added in proof: After acceptance of this manuscript another paper was published describing the identical isoform hSK3_ex1c as hSK3-1C (Kolski-Andreaco AA, Tomita H, Shakkottai VV, Gutman GA,

Cahalan MD, Gargus JJ, Chandy KG. *Journal of Biological Chemistry* 2003 Nov24 [Epub ahead of print]) confirming and extending our findings.

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