

The TRPM7 channel kinase regulates store-operated calcium entry

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Key points

- Pharmacological and molecular inhibition of transient receptor potential melastatin 7 (TRPM7) reduces store-operated calcium entry (SOCE).
- Overexpression of TRPM7 in TRPM7^{-/-} cells restores SOCE.
- TRPM7 is not a store-operated calcium channel.
- TRPM7 kinase rather than channel modulates SOCE.
- TRPM7 channel activity contributes to the maintenance of store Ca²⁺ levels at rest.

Abstract The transient receptor potential melastatin 7 (TRPM7) is a protein that combines an ion channel with an intrinsic kinase domain, enabling it to modulate cellular functions either by conducting ions through the pore or by phosphorylating downstream proteins via its kinase domain. In the present study, we report store-operated calcium entry (SOCE) as a novel target of TRPM7 kinase activity. TRPM7-deficient chicken DT40 B lymphocytes exhibit a strongly impaired SOCE compared to wild-type cells as a result of reduced calcium release activated calcium currents, and independently of potassium channel regulation, membrane potential changes or changes in cell-cycle distribution. Pharmacological blockade of TRPM7 with NS8593 or waixenicin A in wild-type B lymphocytes results in a significant decrease in SOCE, confirming that TRPM7 activity is acutely linked to SOCE, without TRPM7 representing a store-operated channel itself. Using kinase-deficient mutants, we find that TRPM7 regulates SOCE through its kinase domain. Furthermore, Ca²⁺ influx through TRPM7 is essential for the maintenance of endoplasmic reticulum Ca²⁺ concentration in resting cells, and for the refilling of Ca²⁺ stores after a Ca²⁺ signalling event. We conclude that the channel kinase TRPM7 and SOCE are synergistic mechanisms regulating intracellular Ca²⁺ homeostasis.

(Resubmitted 3 January 2017; accepted after revision 20 January 2017; first published online 28 January 2017)

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Abbreviations CRAC, calcium release-activated calcium channel; CRACR2, CRAC-regulatory protein 2; cTRPM7, chicken TRPM7; cWT, chicken wild-type; ΔKinase, deleted kinase domain; ER, endoplasmic reticulum; FBS, fetal bovine serum; HEK, human embryonic kidney; hTRPM7, human TRPM7; IP₃, inositol trisphosphate; KAB, kinase activity buffer; KO, knockout; PI, propidium Iodide; PM, plasma membrane; SARAF, SOCE-associated regulatory factor; SOCE, store-operated calcium entry; STIM, stromal interaction molecule; Tg, thapsigargin; TRP, transient receptor potential; TRPM7, transient receptor potential melastatin 7; WT, wild-type; WT par, wild-type parental.

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Introduction

Changes in intracellular calcium concentration regulate a plethora of biological functions within a cell, including signal transduction, proliferation, motility, transcription, exocytosis and apoptosis. However, dysregulation of Ca^{2+} homeostasis may drive carcinogenesis by modulating cell migration and metastasis (Prevarskaya *et al.* 2014). Cells increase their intracellular Ca^{2+} concentration either by releasing Ca^{2+} from intracellular stores, such as the endoplasmic reticulum (ER) or sarcoplasmic reticulum or by opening Ca^{2+} conducting channels within the plasma membrane (PM). In many electrically non-excitabile cells, both processes are coupled, in that Ca^{2+} release from the ER engages calcium release-activated calcium channels (CRAC) in the PM, a process that is known as store-operated calcium entry (SOCE). This mechanism has two crucial functions: cellular signalling and store refilling.

Although SOCE and the corresponding CRAC channels have been described in the late 1980s and early 1990s, the molecular key players of SOCE remained elusive until extensive RNAi screens in *Drosophila* identified two proteins crucial for SOCE: the ER-resident Ca^{2+} sensor stromal interaction molecule STIM1 (Liou *et al.* 2005; Roos *et al.* 2005) and the PM-resident channel-forming protein Orai1 (Feske *et al.* 2006; Prakriya *et al.* 2006; Vig *et al.* 2006a,b; Zhang *et al.* 2006). STIM1 binds to and activates Orai1 channels upon store depletion, providing a Ca^{2+} influx pathway that is essential for proper cell function in almost all eukaryotic cells. Careful titration of the protein concentration of STIM1 and Orai1 thus determines the amount of Ca^{2+} influx (Hoover & Lewis, 2011; Kilch *et al.* 2013) and its subsequent downstream effects. Mammalian species contain three homologous CRAC channel proteins (Orai1, Orai2 and Orai3), as well as two homologous STIM molecules (STIM1 and STIM2). All Orai homologues exhibit distinct selectivity profiles for Ca^{2+} , Ba^{2+} and Na^{+} , differential pharmacological effects in response to 2-aminoethoxydiphenyl borate, and strikingly different feedback regulation by intracellular Ca^{2+} . STIM1 is thoroughly characterized in the literature, however knowledge about STIM2 awaits further study. Although STIM2 exhibits higher expression levels in primary human melanocytes (Stanisz *et al.* 2012) and is the prominent STIM protein in neurons (Berna-Erro *et al.* 2009) and dendritic cells (Bandyopadhyay *et al.* 2011), gene depletion studies of STIM2 exhibit minimal to no deleterious effects in most cell lines. A significant functional difference between STIM1 and STIM2 is the two-fold higher Ca^{2+} sensitivity of the EF-hand domain of STIM1 (Stathopoulos *et al.* 2009; Zheng *et al.* 2011), which allows STIM2 to activate Ca^{2+} influx at smaller decreases in Ca^{2+} store content. STIM2 has been identified as the feedback regulator for basal Ca^{2+} concentration

and ER store content (Brandman *et al.* 2007). STIM2 may be pre-coupled to Orai channels in the PM, enabling store independent activation of Ca^{2+} influx and regulation of basal cytosolic Ca^{2+} concentration. STIM2 can also remain in the ER, where it serves to activate CRAC channels after mild store depletion (Parvez *et al.* 2008; Thiel *et al.* 2013).

The regulation of SOCE/CRAC is subjected to multiple molecular mechanisms, including post-translational modifications such as phosphorylation (Parekh & Penner, 1995; Smyth *et al.* 2009, 2012; Kawasaki *et al.* 2010), ubiquitination (Keil *et al.* 2010; Eylonstein *et al.* 2011; Lang *et al.* 2012) and glycosylation (Kilch *et al.* 2013), or interaction partners such as CRAC-regulatory protein 2 (CRACR2)A and B (Srikanth *et al.* 2010) and Junctate (Srikanth *et al.* 2012), as well as inhibitory proteins such as Golli (Feng *et al.* 2006) and SOCE-associated regulatory factor (SARAF) (Palty *et al.* 2012).

Another Ca^{2+} -conducting channel of significant research interest in the field of cancer is transient receptor potential melastatin 7 (TRPM7), which contains an ion channel domain that transports Mg^{2+} , Co^{2+} , Mn^{2+} , Ni^{2+} and Zi^{2+} and an active protein kinase domain (Fleig & Chubanov, 2014). Knockout (KO) studies of TRPM7 have established it as a key player in cell growth and proliferation, as well as developmental processes (Nadler *et al.* 2001; Jin *et al.* 2008, 2012; Ryazanova *et al.* 2010; Lee *et al.* 2011; Sah *et al.* 2013a). Although the TRPM7 kinase domain is not essential for channel function, there is a functional coupling that modulates channel gating (Schmitz *et al.* 2003; Jansen *et al.* 2016). It is known that TRPM7 kinase activity is regulated by intracellular Mg^{2+} (Ryazanova *et al.* 2004) and caspases can cleave the kinase from the channel without affecting its phosphotransferase activity (Desai *et al.* 2012). Besides the autophosphorylation of TRPM7 itself, several targets of the kinase have been identified, including the assembly domain of myosin IIA, IIB, IIC (Clark *et al.* 2006), annexin I (Dorovkov & Ryazanov, 2004), phospholipase $\text{C}\gamma 2$ (Deason-Towne *et al.* 2012) and eEF2-k (Perraud *et al.* 2011), as well as components of chromatin-remodelling complexes (Krapivinsky *et al.* 2014).

As a result of the almost ubiquitous presence of these ion channels and their significance for cell physiology and the development of immune diseases and cancer, further research is needed to fully understand their regulatory mechanisms and downstream effects. In the present study, we investigated how TRPM7 contributes to and interacts with SOCE and how this affects the store refilling process and the store content in B lymphocytes. We found that, although TRPM7 itself is not a store-operated ion channel, its activity is acutely linked to SOCE and its facilitating modulatory effect is mediated by the kinase domain of the protein. In addition, TRPM7 contributes to Ca^{2+}

homeostasis by maintaining the filling state of intracellular Ca^{2+} stores in resting cells, and this effect is mediated by the ion channel domain of TRPM7. We hypothesize that TRPM7 channel activity is required to maintain the cellular Ca^{2+} store content and that TRPM7 kinase phosphorylates and thereby modulates STIM proteins and/or other SOCE components. This represents the first study showing a relationship between a TRPM channel, CRAC/SOCE and the ER store content and demonstrates a new facet of how SOCE can be regulated.

Methods

Cell culture

DT40 B lymphocytes, both wild-type (WT) parental (WT par) and genetically modified, were maintained in RPMI 1640 medium (Clontech, Palo Alto, CA, USA) with 10% fetal bovine serum (FBS), 1% chicken serum, 10 U ml⁻¹ penicillin/streptomycin and 2 mM glutamine. Next, 50 µg ml⁻¹ blasticidin was added to chicken TRPM7-KO (cTRPM7-KO) cells. Then, 50 µg ml⁻¹ blasticidin and 1 mg ml⁻¹ zeocin was added to all inducible overexpressing cell lines. DT40 cTRPM7-KO, chicken wild-type (cWT), human TRPM7 (hTRPM7)-deleted kinase domain (Δ Kinase) and hTRPM7-KR were maintained in culture medium supplemented with 15 mM MgCl₂ when uninduced (Schmitz *et al.* 2003). Overexpression in DT40 B cell lines was induced by 1 µg ml⁻¹ doxycycline for 48 h. KO of chicken TRPM7 in the DT40 V79.1 mutant was induced by 200 nM tamoxifen (Nadler *et al.* 2001).

Rat basophilic leukaemia-1 (obtained from the American Type Culture Collection, Manassas, Virginia) and tetracycline-inducible human embryonic kidney (HEK) hTRPM7 cells (Schmitz *et al.* 2003) were maintained in Dulbecco's modified Eagle's medium (Clontech) with 10% FBS and 10 U ml⁻¹ penicillin/streptomycin. The medium for HEK hTRPM7 cells was supplemented with 5 µg ml⁻¹ blasticidin and 200 µg ml⁻¹ zeocin. The overexpression in HEK cells was induced by the addition of 1 µg ml⁻¹ tetracycline for 24 h.

All cells were cultured under 5% CO₂ at 37°C and 95% humidity.

Constructs and transfection

hStim1 and *hStim2* constructs were subcloned into pCAGGS-IRES-GFP and pIRES-Neo, respectively. For transfection, 2 µg of DNA/10⁶ cells was electroporated into HEK-293 cells overexpressing TRPM7 with Nucleofactor II electroporator and kit L (Lonza, Basel, Switzerland). The cells were transfected in accordance with the manufacturer's instructions and cultured for 48 h before protein extraction.

Electrophysiology

Patch clamp recordings were performed at room temperature in the tight-seal whole-cell configuration. Recording electrodes with a resistance of 3–4 MΩ were used. Pipette and cell capacitance were electronically compensated before each voltage ramp with an EPC-10 patch clamp amplifier controlled using Patchmaster software (HEKA, Lambrecht, Germany). After establishing whole-cell configuration, voltage ramps from –100 to +120 mV (200 ms duration) for the measurement of TRPM7 currents and from –150 to +100 mV (50 ms duration) for the measurement of CRAC currents were applied every 2 s from a holding potential of 0 mV. Potassium currents were measured using voltage ramps from –100 to +100 mV with a holding potential of –80 mV. Membrane currents were sampled at 10 kHz and filtered at 2.9 kHz. Voltages were corrected for a liquid junction potential of 10 mV in standard bath solution. For leak current correction, the ramp current before current activation was subtracted and the currents were normalized to whole cell capacitance.

The internal pipette solution contained (in mM): 140 Cs-glutamate, 8 NaCl, 10 Cs-Hepes, 3 MgCl₂, 10 BAPTA and 0.02 inositol trisphosphate (IP₃) for recording CRAC currents and 140 Cs-glutamate, 8 NaCl, 10 Cs-Hepes and 5 mM EDTA for TRPM7 currents. For K⁺ currents, we used (in mM): 140 K-glutamate, 8 NaCl, 10 Hepes and 7.5 CaCl₂, buffered with 10 BAPTA to 1 µM free internal calcium.

Standard bath solution contained (in mM): 120 NaCl, 2.8 KCl, 2 MgCl₂, 10 CaCl₂, 10 CsCl, 10 Na-Hepes and 10 glucose for recording CRAC currents and 140 NaCl, 2.8 KCl, 2 MgCl₂, 1 CaCl₂, 10 Na-Hepes and 10 glucose for both TRPM7 and K⁺ currents.

Flow cytometric analysis of DNA content and cell cycle analysis

DNA content and cell cycle analyses were carried out after fixation of cells and staining with propidium iodide (PI). Briefly, 2–3 × 10⁶ cells of each condition were washed once with PBS, then fixed by adding 1 ml of ice-cold 70% ethanol and stored at 4°C overnight. Cells were washed twice with PBS to remove the EtOH, treated with 20 µg ml⁻¹ RNase A for 30 min and stained by adding 50 µg ml⁻¹ PI for another 30–45 min in the dark at 4°C. Cells were analysed using a BD FACSCalibur Flow Cytometer (Becton-Dickinson Biosciences, Franklin Lakes, NJ, USA) with a 488 laser and data were collected with a 585/42 filter. For cell cycle analysis, the singlets were separated by a gate and 25 000 events per experiment were counted. Analysis was performed using FlowJo software (FlowJo LLC, Ashland, OR, USA).

Fura-2AM based Ca²⁺ imaging

Cells were pre-plated for 30 min on glass coverslips and then loaded with 3.5 μM fura-2AM in RPMI for 15–20 min at 37°C and 5% CO₂ in a humidified incubator. For store refilling experiments, cells were loaded in suspension and washed afterwards with cell culture medium. To deplete the stores, cells were treated with 1 μM ionomycin for 5 min. Ionomycin was removed by washing the cells three times with cell culture medium, followed by plating the cells on glass coverslips for 20 min. Control cells were treated the same way, but without exposure to ionomycin. All experiments were performed in an open perfusion chamber with an upright microscope at room temperature. Images were analysed with TILLVision software (FEI Munich GmbH, Gräfelfing, Germany). The effect of high-K-induced depolarization on SOCE was assessed using a functional drug screening system (FDSS-7000EX; Hamamatsu Photonics KK, Japan), a 96/384-well fluorescent kinetic plate reader. Induced and uninduced DT40 V79.1 cells were loaded with 3.5 μM fura-2AM then plated in 96-well plate at a density of 60 000 cells well⁻¹ before measurement. Bath solutions contained (in mM): 140 NaCl or 140 KCl, 0 or 0.5 or 2 CaCl₂, 2 MgCl₂, 10 glucose and 10 Hepes (pH 7.2 with NaOH). Stock solutions of thapsigargin (Tg) were prepared in DMSO at a concentration of 1 or 5 mM.

Quantitative real-time PCR

Total RNA was isolated from 3×10^6 cells using a Qiagen RNeasy Kit in accordance with the manufacturer's instructions. RNA integrity was assessed using 1 μl of RNA on RNA Nano 6000 chips and a 2100 Agilent Bioanalyzer (Agilent Technologies Inc., Santa Clara, CA, USA). Next, 1 μg of RNA was converted to 20 μl of cDNA using a High Capacity cDNA RT Kit with RNase Inhibitor (Invitrogen, Carlsbad, CA, USA). The kit utilizes random priming. cDNA was further diluted to 8 ng μl^{-1} (based on the starting RNA concentration) with nuclease-free water. Quantitative real-time PCR was performed in a 10 μl reaction volume on a 384-well plate using 32 ng of cDNA in triplicate for the genes of interest and housekeeping genes.

The final concentration of the QuantiTect assay was 1 \times and for individual primers 625 nM for each F- and R-primer with a Quanti Q19 Tect SYBR green kit (Qiagen, Valencia, CA, USA). PCR conditions were: 15 min at 95°C, 15 s at 94°C, 30 s at 55°C (40 cycles) and 30 s at 72°C. Relative expression was calculated in accordance with the ΔCq method ($2^{-\Delta\text{Cq}}$). The primers used were: GAPDH forward: GGCACGCCACTACTATC; reverse: CCTGCATCTGCCCATTT; UB forward: GGG ATGCAGATCTTCGTGAAA; reverse: CTTGCCAGCA AAGATCAACCTT; TRPM7 (Qiagen).

Immunoblotting

For analysis of expression and phosphorylation levels, HEK293 hTRPM7 cells were induced with tetracycline. As shown in Fig. 6A, cells were treated with or without 30 μM NS8593 and lysed after 24 h. As shown in Fig. 7, cells were transfected with Stim1 or Stim2 constructs for 48 h before protein extraction (for details, see above). Lysis buffer contained (in mM): 10 Tris-HCl, 69 Na Cl, 5% glycerol, 0.5% Triton-X, 1 EDTA, 1 \times HALT phosphatase inhibitor cocktail mix (Thermo Scientific, Waltham, MA, USA) and protease inhibitor cocktail (Sigma-Aldrich, St Louis, MO, USA) in accordance with the manufacturer's instructions. For all western blot experiments, 300 μg of total proteins was used to immunoprecipitate TRPM7, Stim1 and Stim2 proteins using anti-HA, anti-FLAG and anti-Stim2 antibodies, respectively. The *in vitro* phosphorylation assay of TRPM7 was performed as described previously (Schmitz *et al.* 2003). Briefly, the immunoprecipitated TRPM7 proteins were treated alone with PBS or kinase activity buffer (KAB) or mixed with Stim1 or Stim2 then incubated with PBS or KAB. All samples were incubated for 30 min at 30°C on a rotating wheel. Samples were then denatured with β -mercaptoethanol for 5 min at 65°C and supernatants were transferred on a 3–8% gradient Tris-acetate gel, separated by electrophoresis, and electrotransferred on polyvinylidene fluoride membranes at a constant 15 V overnight. Immunoblots were probed with anti-HA (dilution 1:500; Roche, Basel, Switzerland), anti-Phospho-Mix (dilution 1:1000; Cell Signaling Technology, Beverly, MA, USA), anti-Flag (dilution 1:1000; Sigma-Aldrich) and anti-STIM2 (dilution 1:400; Anaspec, Inc., Fremont, CA, USA).

Statistical analysis

All data are reported as the mean \pm SEM. Currents were analysed with FitMaster, version 2.11 (HEKA) and IGOR Pro (Wavemetrics, Portland, OR, USA). CRAC current development was analysed at -130 mV and both potassium and TRPM7 currents were extracted at $+80$ mV for statistical analysis. Current amplitudes were normalized to cell size upon break-in as current density (pA/pF). The amplitude of the reported SOCE response represents the difference in fluorescence before and after calcium readdition $[(F_{\text{max}} - F_{\text{min}})/F_{\text{basal}}]$. The absolute intracellular Ca²⁺ concentration was estimated from the relationship $[\text{Ca}^{2+}]_i = K^*(R - R_{\text{min}})/(R_{\text{max}} - R)$ where the values of K , R_{min} and R_{max} were determined from an *in situ* calibration of fura-2-AM in DT40 cells as described by Grynkiewicz *et al.* (1985). Appropriate ANOVA or Student's *t* tests were performed to assess statistical significance. $P < 0.05$ was considered as statistically significant.

Results

TRPM7-deficient B lymphocytes show impaired SOCE

To investigate the role of the Ca²⁺ conducting-channels TRPM7 and CRAC in store-operated calcium entry, as

well as their potential interaction, we measured SOCE in different clones of DT40 Chicken B lymphocytes, either WT or TRPM7-deficient, using a fura-2-based Ca²⁺ imaging approach with a Ca²⁺ re-addition protocol. As shown in Fig. 1, SOCE was strongly reduced from a

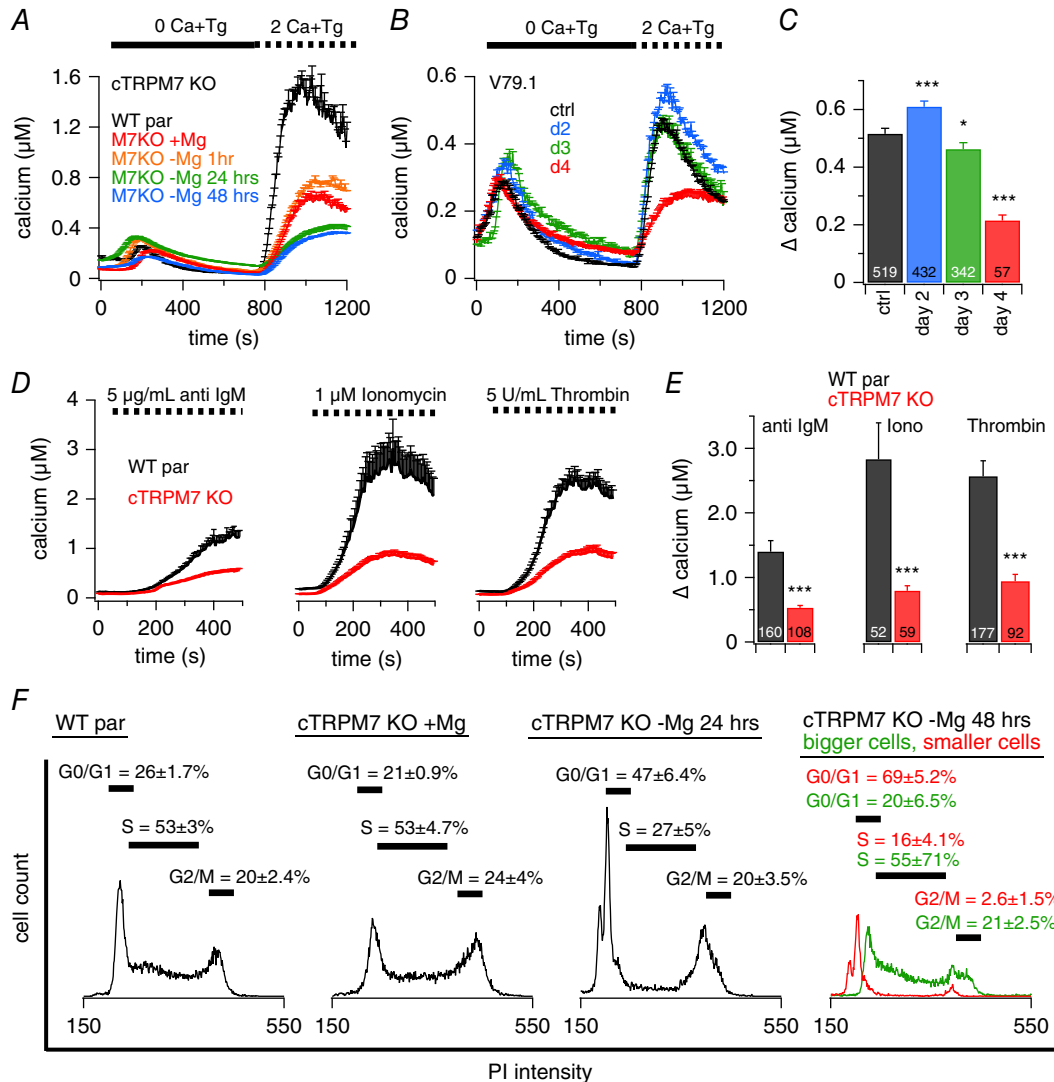


Figure 1. Stable or inducible TRPM7-KO decreases SOCE

Ca²⁺ imaging experiments of intact DT40 chicken B lymphocytes. A–D, at the beginning of the experiment, cells were kept in extracellular standard solution supplemented with 0.5 mM Ca²⁺. A and B, average [Ca²⁺]_i responses following store depletion by the addition of 5 μM Tg and readdition of 2 mM [Ca²⁺]_o (SOCE) in DT40 WT par cells (n = 469) or TRPM7-deficient cells (cTRPM7-KO) and inducible V79.1 TRPM7-KO DT40 cells (B). A, cTRPM7-KO cells were kept in medium with or without supplemented 15 mM Mg²⁺ for the indicated time (+Mg²⁺, n = 546; –Mg²⁺ 1 h, n = 304; –Mg²⁺ 24 h, n = 129; –Mg²⁺ 48 h, n = 207). Data showing 24 h of Mg²⁺ exposure (in green) are shifted by +50 nM Ca²⁺ to make them visible behind the overlapping 48 h of data (in blue). B, KO was induced with tamoxifen and cells were kept in Mg²⁺ supplemented medium. C, average of Ca²⁺ influx peaks assessed from baseline and obtained from SOCE measurements in cells in (B). D, average [Ca²⁺]_i responses following store depletion and Ca²⁺ influx by the addition of 2 mM [Ca²⁺]_o (SOCE) triggered by different stimuli as indicated, in DT40 WT par cells (anti-IgM n = 207, ionomycin n = 207, thrombin n = 177) or cTRPM7-KO cells (anti-IgM n = 108, ionomycin n = 59, thrombin n = 92). E, average Ca²⁺ influx peaks obtained from SOCE measurements in cells in (D). F, PI based cell cycle analysis of DT40 WT par and cTRPM7-KO cells kept in medium with or without supplemented 15 mM Mg²⁺ for the indicated time (representative traces of six independent experiments are shown). Stars indicate statistical significance: *P < 0.05; **P < 0.01; ***P < 0.001.

peak value of 1953 ± 134 nM Ca^{2+} in WT par cells to 756 ± 56 nM Ca^{2+} in stable DT40 cTRPM7-KO cultured in medium supplemented with 15 mM Mg^{2+} . Removal of the supplemented Mg^{2+} from the culture medium of cTRPM7-KO cells for the indicated period of time reduced SOCE even more, probably as a result of a quiescent metabolic transition (Sahni *et al.* 2010) (Fig. 1A). Similarly, the tamoxifen-inducible DT40 cTRPM7-KO cell line V79.1 also responded with reduced SOCE (Fig. 1B and C). The maximal increase in $[\text{Ca}^{2+}]_i$ after store depletion and external Ca^{2+} re-addition was reduced from 515 ± 19 nM in uninduced V79.1 to 214 ± 19 nM in cells 4 days after induction of the KO, although the cells were supplemented with 15 mM Mg^{2+} in the culture medium.

To test whether other stimuli of SOCE showed the same phenomenon, we used anti-IgM antibodies, ionomycin and thrombin to release Ca^{2+} from the stores. The impaired SOCE in cTRPM7-KO cells compared to WT par cells was also observed after induction of SOCE with these stimuli (Fig. 1D and E).

Previous work has shown that the cell cycle state affects the amount of SOCE (Tani *et al.* 2007; Smyth *et al.* 2009). We therefore aimed to examine whether the reduced SOCE in TRPM7-deficient cells was a result of the effect of TRPM7 on cell-cycle distribution. Using PI-based flow cytometry analyses, we measured the cell cycle distribution of B lymphocytes under all the experimental conditions described above. cTRPM7-KO cells, cultured with an additional 15 mM Mg^{2+} in the cell culture medium, showed a similar cell cycle distribution as the WT par cells. The transition into the quiescent metabolic state occurs only upon removal of additional Mg^{2+} from the culture medium (Sahni *et al.* 2010) (Fig. 1F). The cells then undergo apoptosis, resulting in an increase of the amount of cells in G0/G1 and hardly any cells in M-phase. Forty-eight hours after removal of additional Mg^{2+} from the culture medium, there are predominantly two populations of cells that differ in cell size and in cell cycle distribution (Fig. 1F, right) and represent dying and living cells. Taken together, our results show that the SOCE phenotype in cTRPM7-KO cells is a result of the absence of TRPM7 and not the subsequent changes in cell cycle distribution and cell growth.

TRPM7-KO impairs SOCE by reducing I_{CRAC} rather than affecting K^+ conductances

To test whether TRPM7-KO affects potassium (K^+) conductances and thereby possibly changes membrane potential and calcium driving force, we used DT40 V79.1 cells cultured in the absence (uninduced) or presence of 200 nM tamoxifen (induced) to generate the cTRPM7-KO. We first investigated whether impaired SOCE persists under completely potassium-depolarized conditions. We performed fura-2-based Ca^{2+} imaging experiments under

physiological condition using standard Ringer solution containing 140 mM NaCl, as well as under a depolarized condition where NaCl was equimolarly replaced by KCl and collapsing the membrane potential to 0 mV. We found that TRPM7-KO reduced SOCE in a similar way under both physiological ($\sim 45\%$ inhibition) and depolarizing conditions ($\sim 40\%$ inhibition) (Fig. 2A and B).

To confirm these findings and more directly establish that K^+ conductances are not affected by TRPM7-KO, we next investigated whole-cell K^+ currents using experimental conditions that were designed to simultaneously reveal effects on both voltage- and calcium-activated K^+ currents, if present. Cells were held at -80 mV to keep voltage-dependent K^+ channels activatable and stimulated at 0.5 Hz with a 50 ms voltage ramp from -100 mV to $+100$ mV to reveal any voltage-dependent currents. We have previously used this voltage protocol successfully in Jurkat T cells to assess Kv1.3 (Ren *et al.* 2008). In parallel, we supplemented the internal solution with $1 \mu\text{M}$ Ca^{2+} to concomitantly activate any Ca^{2+} -gated K^+ channels. Under these conditions, we did not observe significant voltage-dependent currents in DT40 cells but, instead, observed a constitutive and linear conductance that reversed at around -55 mV, probably reflecting a resting K^+ conductance in both WT and cTRPM7-KO cells (Fig. 2C and D). Over time, we observed an increase in a slightly outwardly rectifying conductance that had a similar reversal potential and might have been a result of Ca^{2+} -dependent K^+ channels (Fig. 2C and D). We have not attempted to further characterize these currents because they were unaltered in WT and cTRPM7-KO cells. These data further support the conclusions derived from the Ca^{2+} measurements in intact cells under fully depolarized conditions (Fig. 2A and B), indicating that membrane potential effects are not responsible for the reduction in SOCE.

Next, we assessed the store-operated CRAC current (I_{CRAC}) in both WT and TRPM7-deficient cells using a voltage ramp protocol from -150 to $+100$ mV over 50 ms and delivered at 0.5 Hz intervals from a holding membrane potential of 0 mV. Figure 2E and F shows that I_{CRAC} was significantly reduced in cTRPM7-KO cells compared to cTRPM7 expressing cells. These results demonstrate that the mechanism by which TRPM7-KO impairs SOCE appears to result directly from reduced CRAC currents rather than indirectly through effects on K^+ currents and membrane potential changes.

Pharmacological blockade of TRPM7 shows that SOCE is acutely linked to TRPM7 function

To assess whether impaired SOCE in TRPM7-deficient B lymphocytes is a long-term adaptive effect as a result of the KO of TRPM7 protein or an acutely linked process, we performed fura-2-based Ca^{2+} imaging experiments

using different known TRPM7 or CRAC inhibitors. As expected, Gd^{3+} , which inhibits CRAC channels without affecting TRPM7 (Hermosura *et al.* 2002), decreased the maximal $[Ca^{2+}]_i$ influx peak after store depletion from 1259 ± 41 nM in control cells to 34 ± 4 nM $[Ca^{2+}]_i$ (Fig. 3A). The TRPM7 blockers NS8593 (Chubanov *et al.*

2012) and waixenicin A (Zierler *et al.* 2011) were also able to strongly inhibit the SOCE response to 243 ± 22 nM and 191 ± 30 nM $[Ca^{2+}]_i$, respectively. This effect was specific because, in control experiments in B lymphocytes deficient for TRPM7 expression, the TRPM7 blocker NS8593 failed to impair SOCE (Fig. 3B and D).

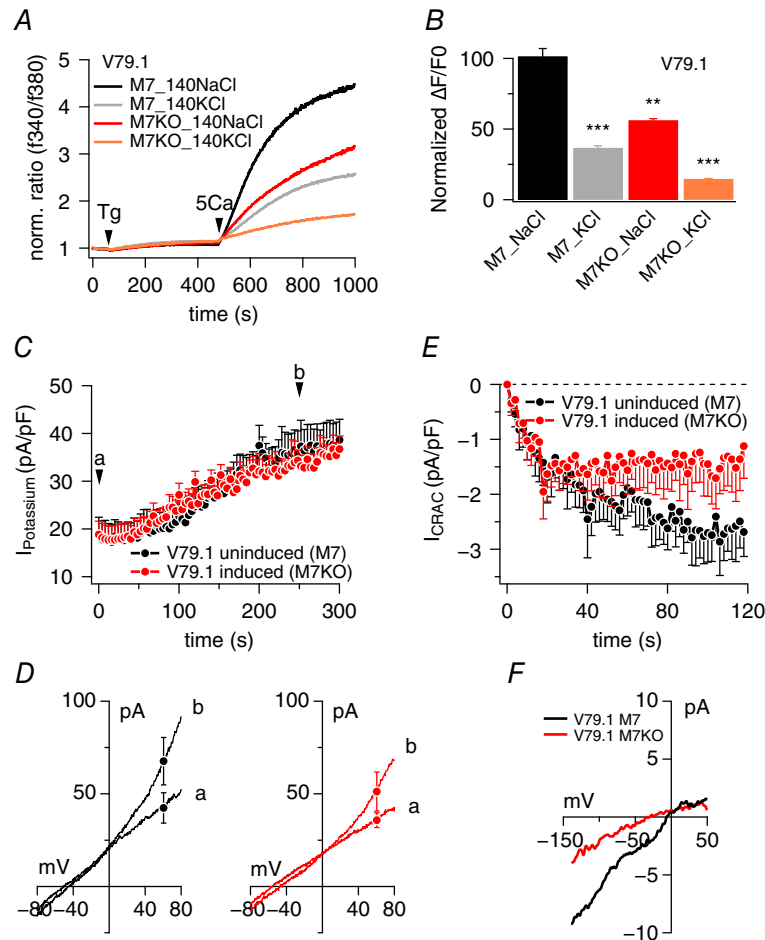


Figure 2. Impaired SOCE in TRPM7-deficient B lymphocytes is not due to altered K^+ currents

A, average $[Ca^{2+}]_i$ responses following store depletion by the addition of $5 \mu M$ Tg and readdition of 5 mM $[Ca^{2+}]_o$ (SOCE) in uninduced DT40 V79.1 or induced DT40 V79.1. Calcium signals were acquired on a 96-well fluorescence kinetic plate reader and measured in standard Ringer solution with 140 mM NaCl or in depolarized condition by replacing NaCl with KCl. Data are averages of triplicates. B, average peak Ca^{2+} influx responses assessed from baseline and obtained in (A) ($n = 3$ each). Asterisks indicate statistical significance: $*P < 0.05$; $**P < 0.01$; $***P < 0.001$. C, time course of average K^+ current development in uninduced DT40-V79.1 cells ($n = 9$) and in induced DT40 V79.1 ($n = 10$). Tamoxifen (200 nM) was used to induce TRPM7-KO and currents were recorded 4 days post-induction. Cells were held at -80 mV and stimulated with a 50 ms voltage ramp from -100 mV to $+100$ mV at 0.5 Hz to activate voltage-gated K^+ channels. Cells were concomitantly perfused with $1 \mu M$ Ca^{2+} to activate any additional Ca^{2+} -sensitive K currents. Currents were extracted at $+80$ mV, normalized to cell size and plotted as current density in pA/pF over the time of the experiment. D, average $I-V$ behaviour of K^+ currents extracted at the two time points labelled (a) and (b) in (C) from uninduced (black $I-V$ traces; $n = 9$) and induced-KO cells (red $I-V$ traces; $n = 10$). E, average CRAC currents (I_{CRAC}) evoked by $20 \mu M$ IP_3 in uninduced DT40 V79.1 ($n = 12$) or induced DT40 V79.1 ($n = 20$). Cells were held at 0 mV and stimulated with a 50 ms voltage ramp from -150 mV to $+100$ mV at 0.5 Hz. Cells were concomitantly perfused with $20 \mu M$ IP_3 to activate I_{CRAC} . Currents were extracted at -130 mV, normalized to cell size and plotted as current density in pA/pF over time of the experiment. F, average $I-V$ behaviour of CRAC currents extracted from 116 to 120 s. The black trace represents average current from uninduced cells ($n = 6$) and the red trace represents average current from M7KO cells ($n = 14$).

Impaired SOCE in TRPM7-KO B lymphocytes is restored by expression of TRPM7

Corroborating evidence linking TRPM7 functionality and SOCE is provided by the rescue experiments shown in Fig. 3E and J. Inducing the overexpression of hTRPM7 in the cTRPM7-KO DT40 cell line (cWT) (Fig. 3E and F) restores SOCE to almost the same degree seen in DT40 WT par cells (WT par: 1055 ± 43 nM; cTRPM7-KO: 515 ± 28 nM; cWT induced: 855 nM [Ca^{2+}]_i) (Fig. 3H and J). The overexpression was confirmed by quantitative RT-PCR (Fig. 3E), as well as by patch clamp experiments (Fig. 3F). Although the cTRPM7-KO cells did not exhibit any TRPM7 current, the average maximal current density was 284 pA/pF in the WT par cells and 766 pA/pF in the overexpressing cells (cWT). The correspondent I - V relationships are shown in Fig. 3G. These results demonstrate a clear interaction and a direct link between TRPM7 functionality and SOCE.

TRPM7 itself is not a store-operated channel

To test whether the observed decrease of SOCE in TRPM7-deficient cells stems from the contribution of Ca^{2+} entry through TRPM7 channels following store depletion, we performed fura-2-based imaging experiments where we inhibited CRAC currents either by the CRAC channel inhibitor Gd^{3+} or by using DT40 chicken B lymphocytes deficient in CRAC components (Wang *et al.* 2009). The addition of Gd^{3+} to the bath solution completely abolished SOCE in WT par, cTRPM7-KO and hTRPM7-overexpressing B lymphocytes (Fig. 4A and B).

We next examined SOCE in B lymphocyte clones deficient in one or more CRAC components. Although the lack of Orai1 had little effect on SOCE, the absence of Orai2 actually increased the signal (Fig. 4C and D), which is consistent with previous work (Holzmann *et al.* 2013). However, the double-KO of Orai1/Orai2 almost

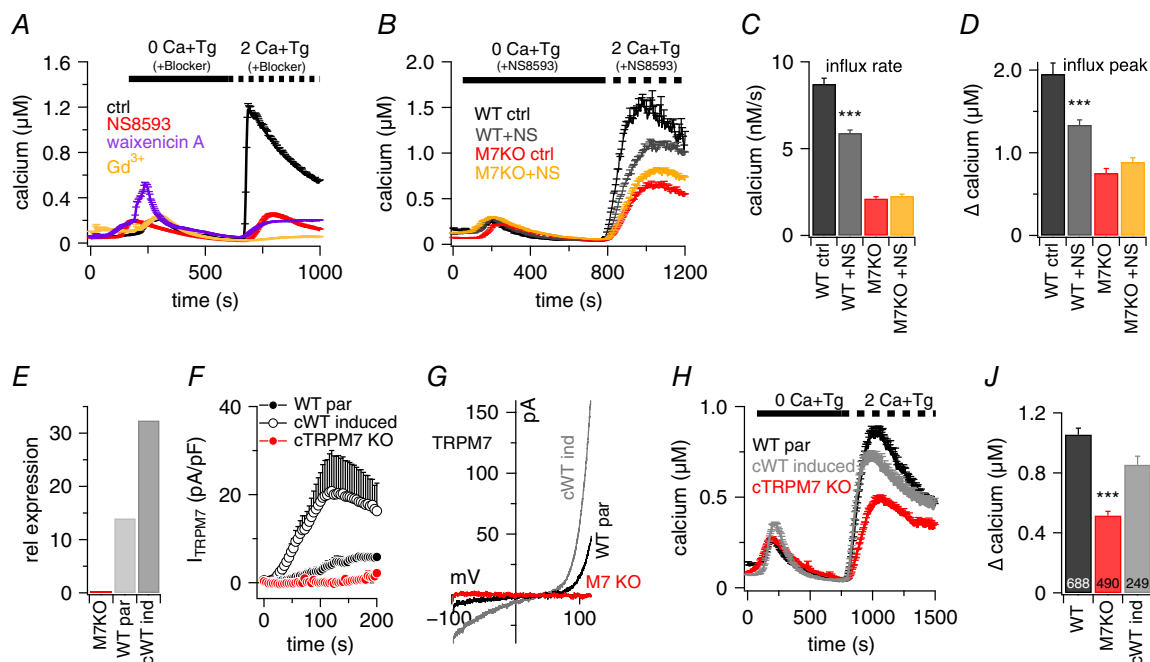


Figure 3. SOCE is acutely linked to TRPM7 function

A and B, average [Ca^{2+}]_i responses following store depletion by the addition of $5 \mu\text{M}$ Tg and readdition of 2 mM [Ca^{2+}]_o (SOCE) in rat basophilic leukaemia cells (A) or DT40 B cells (B). A, bath solution was non-supplemented (ctrl, $n = 79$) or supplemented with $1 \mu\text{M}$ Gd^{3+} ($n = 22$), $30 \mu\text{M}$ NS8593 ($n = 29$) or $10 \mu\text{M}$ waixenicin A ($n = 17$) at the time of application, as indicated by the black bar. B, bath solution was supplemented with $30 \mu\text{M}$ NS8593 or without (ctrl). Shown are traces for DT40 WT par cells with ($n = 732$) or without ($n = 469$) and cTRPM7-KO cells with ($n = 441$) or without ($n = 546$) blocker. C and D, average of Ca^{2+} influx rates (C) and maximum Ca^{2+} influx peaks assessed from baseline of each cell (D) and obtained from SOCE measurements in (B). E, quantitative RT-PCR analyses of TRPM7 expression [$2^{-\Delta\Delta\text{Cq}}$] in DT40 B cells cTRPM7-KO (M7KO), WT par and cWT induced (cWT ind) for hTRPM7 overexpression. The relative expression of TRPM7 was normalized to the average expression of GAPDH and UB house keeping genes. F, TRPM7 current densities in DT40 B cells cTRPM7-KO (red, $n = 5$), WT par ($n = 6$) and cWT after induction ($n = 6$) as assessed at $+80$ mV. G, I - V relationships of currents evoked by 200 ms voltage ramps extracted from cells measured in (F). H, average [Ca^{2+}]_i responses following store depletion by the addition of $5 \mu\text{M}$ Tg and readdition of 2 mM [Ca^{2+}]_o (SOCE) in DT40 B cells cTRPM7-KO ($n = 490$), WT par ($n = 688$) and cWT after induction ($n = 249$). I, average Ca^{2+} influx peaks assessed from baseline and quantified from SOCE measurements in (H). Asterisks indicate statistical significance: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

completely eliminated SOCE. Similarly, STIM1-KO cells exhibited no significant SOCE signal, whereas STIM2 reduced SOCE by 52%. Taken together, these results demonstrate that TRPM7 is not store-operated and does not contribute significantly to SOCE-mediated signals. Rather, it appears to play a regulatory role in support of SOCE.

The TRPM7 kinase domain modulates SOCE

Because the above experiments show that Ca^{2+} flux through TRPM7 does not contribute to SOCE, TRPM7 might shape SOCE through its kinase domain activity. To test this, we made use of two DT40 cell lines overexpressing kinase-dead hTRPM7 mutants after induction with doxycycline, namely hTRPM7- Δ -kinase and hTRPM7-K1648R (Schmitz *et al.* 2003; Yu *et al.* 2013). Although both TRPM7 variants lack kinase activity, they exhibit different degrees of ion channel function. As illustrated in Fig. 5A and B, the hTRPM7-K1648R mutant conducts currents comparable to the hTRPM7-WT current (cWT) (Fig. 3G) and the Δ -kinase mutation essentially eliminates conductivity through the pore of the channel. Fig. 5C and D depicts the results of the fura-2-based Ca^{2+} imaging experiments. Although overexpression of hTRPM7-WT (cWT) accomplishes the rescue of the cTRPM7-KO phenotype in SOCE, neither the hTRPM7- Δ -kinase, nor the hTRPM7-K1648R proteins, despite their differences in ion transport abilities, are able to do the same. These results demonstrate that the TRPM7 kinase domain rather than the TRPM7 channel domain regulates SOCE in DT40 B-lymphocytes.

TRPM7 currents are required to refill the internal Ca^{2+} stores after depletion

SOCE not only serves to translate signalling events from the PM into the cell, but also to refill the internal Ca^{2+} stores after signalling. Because we found that TRPM7 functionality affects SOCE, we next considered whether TRPM7 affects ER store content at rest and/or contributes to the refilling process after signalling. We examined the role of TRPM7 in regulating the resting Ca^{2+} store content (Fig. 5E and F), as well as the refilling of the stores after depletion (Fig. 5G and H), by performing fura-2-based intact-cell imaging experiments and using ionomycin to probe ER store contents in various DT40 cell lines.

As illustrated in Fig. 5E and F, cTRPM7-KO cells had significantly less Ca^{2+} in their stores than WT par cells, as indicated by reduced Ca^{2+} release responses to ionomycin challenge ($184 \pm 4 \text{ nM}$ vs. $341 \pm 7 \text{ nM}$ Ca^{2+}). Interestingly, the phenotype could be rescued by overexpressing either hTRPM7-WT (cWT) or hTRMP7-K1648R ($392 \pm 10 \text{ nM}$ and $476 \pm 15 \text{ nM}$ Ca^{2+}) but not by the non-conductive hTRPM7- Δ -kinase protein (Fig. 5E and F). Thus, overexpression of hTRMP7-K1648R can rescue the store content phenotype of cTRPM7-KO cells, even though it is unable to rescue the SOCE phenotype in those cells. This indicates that the constitutive activity of TRPM7 contributes to the filling state of resting cells. We further demonstrate that CRAC channels also appear to participate in this process because ionomycin responses in resting DT40 B cells are reduced in cells deficient in Orai1 and Orai2 ($264 \pm 6 \text{ nM}$) or STIM1 ($107 \pm 4 \text{ nM}$), whereas STIM2-deficient cells have a similar store content

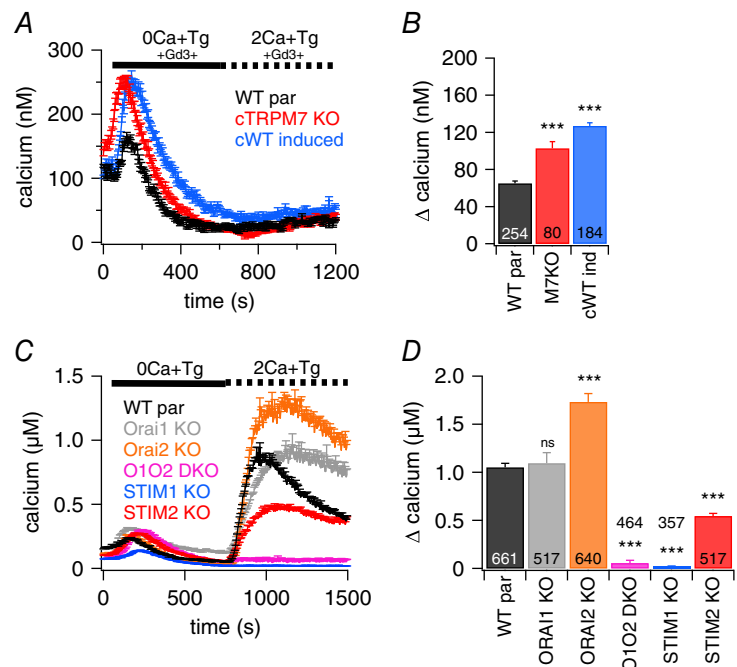


Figure 4. TRPM7 itself is not a store-operated channel
A and C, average $[\text{Ca}^{2+}]_i$ responses following store depletion by the addition of $5 \mu\text{M}$ Tg and readdition of 2 mM $[\text{Ca}^{2+}]_o$ (SOCE) in intact DT40 B cells. A, traces are shown for cTRPM7-KO ($n = 80$), WT par ($n = 254$) and cWT after induction ($n = 184$). Bath solution was supplemented with $1 \mu\text{M}$ Gd^{3+} . C, traces are shown for WT par ($n = 661$), Orai1-KO ($n = 517$), Orai2-KO ($n = 640$), Orai1/Orai2 double KO (O1O2 DKO; $n = 464$), STIM1-KO ($n = 357$) and STIM2-KO ($n = 517$). B and D, average of Ca^{2+} influx peaks obtained from SOCE measurements in cells in (A) or (C), respectively. The maximum Ca^{2+} release was measured for each individual cell from the resting baseline, averaged and plotted. Asterisks indicate statistical significance: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

as WT par cells, yielding similar ionomycin responses (354 ± 13 nM) (Fig. 5E and F). These results show that TRPM7 and CRAC currents control the intracellular Ca^{2+} store content at rest. Surprisingly, however, and in contrast to other cell systems, where STIM2 is considered to regulate basal cytosolic and ER levels (Brandman *et al.*

2007; Várnai *et al.* 2009; Gruszczynska-Biegala *et al.* 2011), it appears that, in the DT40 B cells, STIM1 may be more important for this resting activity of CRAC channels than STIM2.

In addition to contributing to Ca^{2+} store content under resting conditions, TRPM7 appears to also participate in

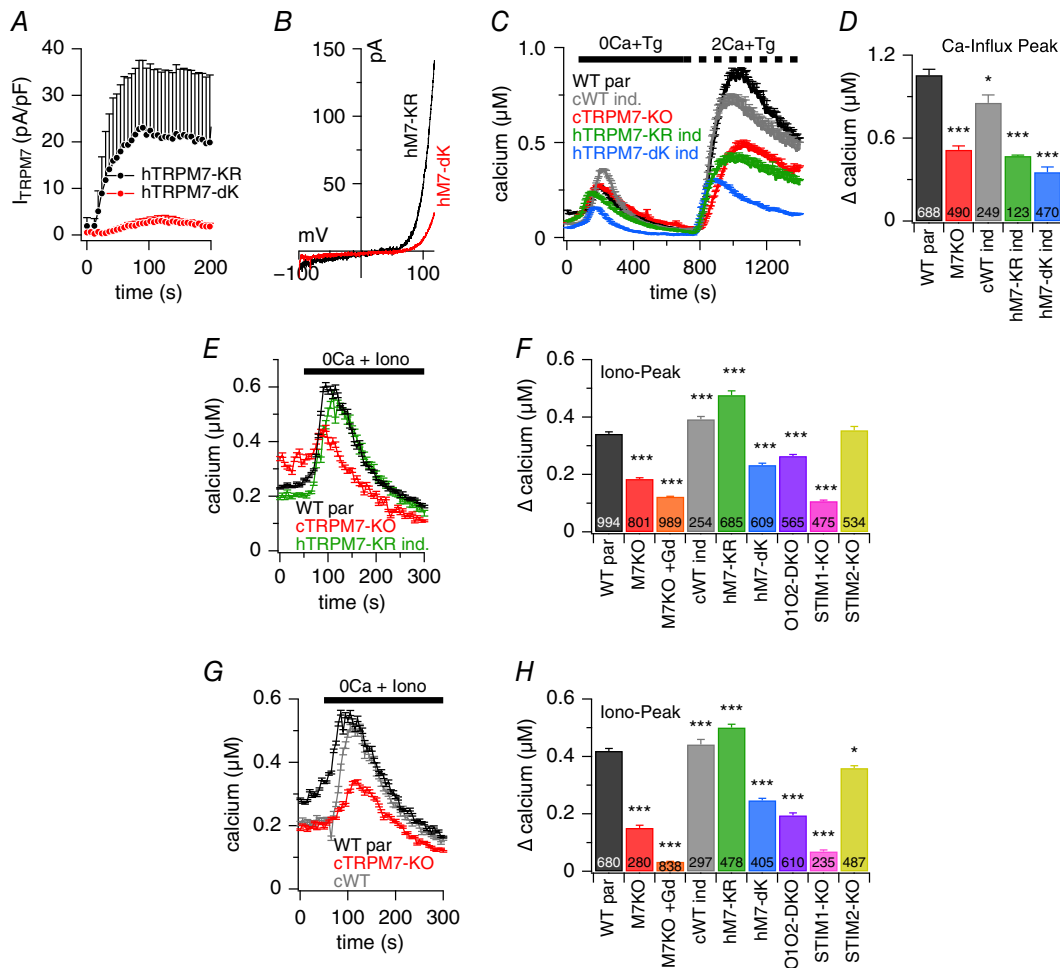


Figure 5. TRPM7 kinase domain affects SOCE and TRPM7 conductivity is needed to refill Ca^{2+} stores

A, TRPM7 current densities in DT40 B cells overexpressing either hTRPM7- Δ Kinase ($n = 6$) or hTRPM7-KR mutants ($n = 6$), 48 h after induction. B, I - V behaviour of currents evoked by 50 ms voltage ramps obtained from cells measured in (A). Holding potential was 0 mV. C, average $[\text{Ca}^{2+}]_i$ responses following store depletion by the addition of $5 \mu\text{M}$ Tg and readdition of indicated $[\text{Ca}^{2+}]_o$ (SOCE) in DT40 B cells WT par ($n = 688$) and cWT ($n = 249$), cTRPM7-KO ($n = 490$), hTRPM7- Δ Kinase ($n = 470$) and hTRPM7-KR mutant ($n = 123$) after induction. D, average of maximal Ca^{2+} influx assessed from baseline and obtained from SOCE measurements in (C). E, average $[\text{Ca}^{2+}]_i$ responses following store depletion by application of $1 \mu\text{M}$ ionomycin (Iono) in DT40 B cells WT par and cTRPM7-KO, and cTRPM7-KR mutant after induction. Ionomycin was applied in the absence of extracellular Ca^{2+} in the solution. In one experiment, cTRPM7-KO cells were analysed in the presence of $1 \mu\text{M}$ Gd^{3+} in the bath solution. F, average of Ca^{2+} release peaks assessed from baseline and obtained from measurements in (E), and additional $[\text{Ca}^{2+}]_i$ responses not depicted in (E) but using the same experimental approach for cTRPM7-KO with $1 \mu\text{M}$ Gd^{3+} in the bath, induced hTRPM7 KR mutant, induced hTRPM7- Δ Kinase mutant, Orai1/Orai2 double KO (O1O2 DKO), STIM1-KO and STIM2-KO. Numbers in black indicate number of cells averaged. G, for the refilling experiments, the experimental procedure was the same as shown for control cells in (E) but cells were treated with $1 \mu\text{M}$ ionomycin 20 min before the measurement (see Methods). Average $[\text{Ca}^{2+}]_i$ responses following store depletion by the addition of $1 \mu\text{M}$ ionomycin (Iono) in DT40 B cells WT par, cTRPM7-KO and cWT after induction. H, average of Ca^{2+} release peaks from baseline and obtained from measurements in (G) and additional DT40 mutant cell lines as investigated in (F). Asterisks indicate statistical significance: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

the refilling process after depletion of Ca^{2+} stores. This is illustrated in Fig. 5G and H, where experiments are depicted in which the transient application of ionomycin was used to fully deplete Ca^{2+} stores, followed by a 20 min waiting period that allowed cells to refill their stores. Some 20 min after depletion, the store content in cTRPM7-KO cells was reduced by ~64% compared to the WT par cells. In the additional presence of Gd^{3+} for blocking SOCE, there was no refilling of ER stores in cTRPM7-KO cells at all. Cells overexpressing hTRPM7-WT (cWT) or hTRMP7-K1648R exhibited normal refilling levels comparable to WT par cells. The overexpression of hTRPM7- Δ -kinase refilled the stores 41% less than the hTRPM7-WT protein. The deficiency in SOCE components also affected the store refilling process because the Orai1/Orai2 double KO, which did not show any SOCE in the Ca^{2+} re-addition measurements (Fig. 4), reduced the signal by ~54%. Remarkably, the STIM1-KO was more effective than the Orai1/Orai2 KO in that it reduced ionomycin-induced Ca^{2+} responses by ~84%, whereas STIM2-KO achieved merely a 14% reduction. Taken together, these experiments demonstrate a major

role of TRPM7 Ca^{2+} current in the refilling of intracellular Ca^{2+} stores. It appears that TRPM7 and SOCE work hand in hand to maintain the intracellular Ca^{2+} balance under resting conditions, as well as after Ca^{2+} signalling.

The TRPM7 kinase domain may be targeting STIM proteins

The results of the present study demonstrate that acute SOCE regulation by TRPM7 requires a functional kinase domain, irrespective of the remaining ion channel conductivity. To examine the target of the TRPM7 kinase domain, we made use of the TRPM7 channel blocker NS8593 to investigate whether the channel activity of TRPM7 is needed for efficient kinase activity. As a read out of kinase activity, we assessed the auto-phosphorylation of TRPM7 protein in immunoblot experiments. Tetracycline-inducible HEK293 cells overexpressing hTRPM7 (Schmitz *et al.* 2003) and treated with NS8593 showed a 42% reduction in autophosphorylation of hTRPM7, illustrating the modulatory role of the TRPM7 current for kinase-activity (Fig. 6A and B). This

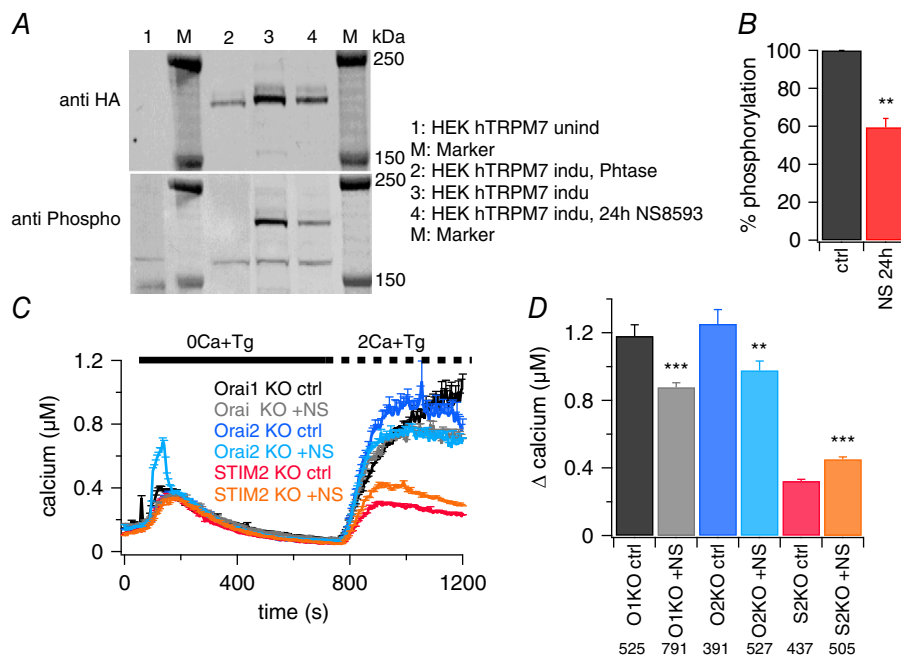


Figure 6. TRPM7 kinase domain may be targeting STIM proteins

A, western blot analyses of TRPM7 expression and phosphorylation in tetracycline-inducible HEK hTRPM7 cells. (1) uninduced cells (M, marker); (2) cells induced with tetracycline (24 h) and cell lysate treated with λ phosphatase; (3) cells induced with tetracycline (24 h); and (4) cells induced with tetracycline and treated with $30 \mu\text{M}$ NS8593 (24 h) (a representative blot for four independent experiments is shown) (M, marker). B, percentage of hTRPM7 phosphorylation from cells treated with or without NS8593; lane 3 vs. lane 4 in (A). Phosphorylation was normalized to total protein level (HA-signal) and phosphorylation in untreated cells was set to 100%. C, average $[\text{Ca}^{2+}]_i$ responses following store depletion by the addition of $5 \mu\text{M}$ Tg and readdition of 2 mM $[\text{Ca}^{2+}]_o$ (SOCE) in DT40 B cells Orai1-KO, Orai2-KO and STIM2-KO with $30 \mu\text{M}$ NS8593 (+NS) or without (ctrl) in the bath solution. Number of cells are shown in the corresponding bar graphs in (D). D, average of Ca^{2+} influx peaks assessed from baseline and obtained from SOCE measurements in (C). Asterisks indicate statistical significance: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

result shows that NS8593 not only blocks the current through the TRPM7 pore, but also decreases the TRPM7 kinase activity.

Next, we used this pharmacological tool to examine which of the molecular components of SOCE might be the targets of the TRPM7 kinase. The rationale behind this approach was that blocking the ion flux through TRPM7 should affect the kinase activity, and removing the target of phosphorylation in B lymphocytes would render the SOCE of these cells resistant to NS8593 inhibition. Accordingly, we performed fura-2-based intact-cell imaging experiments in which we examined the Ca^{2+} signals in the presence of the TRPM7 blocker NS8593 in different B lymphocyte clones deficient in one of the CRAC components. As depicted in Fig. 6C and D, the TRPM7 blocker was able to reduce SOCE in Orai1-KO (26% reduction) and Orai2-KO (21% reduction) cells but not in STIM2-KO cells. Because the STIM1-KO cells did not show any SOCE, they could not be used for this type of experiment. These results suggest an effect of the TRPM7 kinase domain on STIM2 proteins, although we cannot rule out a possible effect on STIM1 as well.

Given our results indicating a possible mechanism involving STIM proteins, we assessed whether STIM1 and STIM2 are the direct targets of TRPM7 kinase. We transfected tetracycline-inducible HEK-293 overexpressing hTRPM7 with either STIM1 plasmid (Fig. 7A) or STIM2 plasmid (Fig. 7B). Although the anti-phosphomix detected the *in vitro* phosphorylation of hTRPM7 (Fig. 7B), we were unable to detect any phosphorylation of STIM1 and STIM2. The absence of STIM phosphorylation may be a result of the potential limiting factor of the

sensitivity of the technique (detection limit) or the antibodies, or possibly because TRPM7 may affect STIM indirectly by engaging another protein partner of STIM or Orai proteins.

Discussion

In the present study, we report a new mechanistic target of the TRPM7 channel kinase: SOCE. The present study confirms that TRPM7 channel activity, although not directly contributing to SOCE, influences TRPM7 kinase function, thereby modifying the efficacy of Ca^{2+} entry via SOCE. In addition, constitutive TRPM7 ion channel activity serves to maintain Ca^{2+} store contents in resting unstimulated cells. Thus, our results suggest that TRPM7 and CRAC currents work hand in hand to regulate cytosolic Ca^{2+} signals as well as refill intracellular Ca^{2+} stores.

A previous study observed a quiescent metabolic transition of TRPM7-deficient B lymphocytes (TRPM7-KO) after removal of additional Mg^{2+} from the cell culture medium, causing a shift in the cell cycle distribution of the cells towards G_0/G_1 phase and resulting in a reduction in SOCE (Sahni *et al.* 2010). In the present study, we demonstrate that TRPM7-KO cells, cultured with additional Mg^{2+} , show a normal cell cycle distribution comparable to WT DT40 lymphocytes and nevertheless have impaired SOCE (Fig. 1). Furthermore, TRPM7 blockers such as NS8593 and waixenicin A also decrease SOCE (Fig. 3), indicating a direct and acute correlation between the two mechanisms. The TRPM7-KO-mediated phenotype can be rescued by overexpressing hTRPM7-WT but not by the kinase

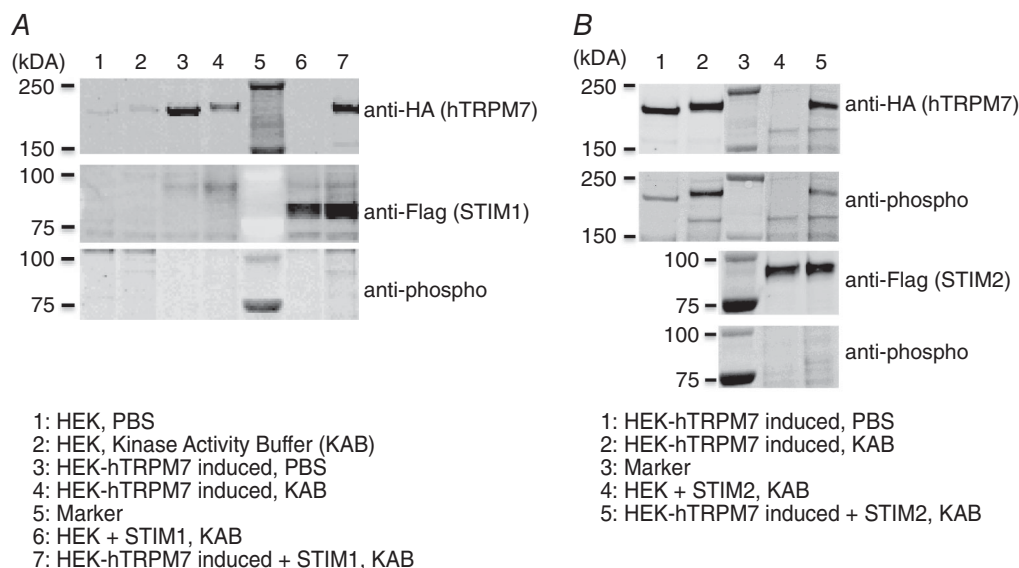


Figure 7. No detectable phosphorylation of STIM proteins was observed by Western blot

A, representative western blot of STIM1 co-expression with TRPM7 and its phosphorylation in HEK hTRPM7 cells. B, representative western blot of STIM2 co-expression with TRPM7 and its phosphorylation in HEK hTRPM7 cells.

dead mutant hTRPM7-K1648R or hTRPM7- Δ Kinase, suggesting that kinase activity of the protein rather than its ion channel function imparts a regulatory effect on the molecular CRAC components Orai and/or STIM (Fig. 5).

Indeed, the literature reports several ways of regulating SOCE by phosphorylation. Previous work identified two serine residues in the STIM1 protein that are phosphorylated in mitotic cells and are capable of suppressing SOCE (Smyth *et al.* 2009). Another study showed that Orai1 is a direct target of the Ca^{2+} -dependent kinase PKC β , and the phosphorylation of S27 and S30 abolishes SOCE (Kawasaki *et al.* 2010). However, phosphorylation can also augment SOCE. We have previously reported that Orai1 expression levels are regulated by serum- and glucocorticoid-inducible kinase 1 (SGK1), a growth factor-regulated kinase (Eylenstein *et al.* 2011). An enhancing effect has also been shown when STIM1 is phosphorylated by extracellular-signal-regulated kinases 1 or 2 on serine residues (Poza-Guisado *et al.* 2013). Our present data indicate that STIM2 might be the phosphorylation target of the endogenous kinase of TRPM7 rather than Orai (Fig. 6). Although one study has reported that Stim2 protein can be phosphorylated (Williams *et al.* 2001), we were unable to detect any phosphorylation in our current efforts (Fig. 7). The absence of STIM phosphorylation may be a result of the sensitivity of the technique (detection limit), the quality and sensitivity of antibodies, or possibly because TRPM7 may affect SOCE indirectly by engaging regulatory protein partners of STIM or ORAI. Possible candidates for such indirect modulatory effects include Septin, junctate, Golli, SARAF and CRACR2A, (Walsh *et al.* 2010; Srikanth *et al.* 2012; Hooper & Soboloff, 2015), warranting further investigation to determine the direct target of TRPM7 phosphorylation and its connection to SOCE.

There are numerous ways in which TRP channels and SOCE can interact in different cellular systems to generate a precisely adapted spatio-temporal Ca^{2+} signalling (Saul *et al.* 2014). Our results depict for the first time an acute functional link between TRPM7 and store-operated calcium entry. I_{CRAC} is certainly the best described store-operated calcium entry mechanism; however, other channels, especially members of the 'canonical' TRP sub-family, have also been reported to be store-operated (Parekh & Putney, 2005; Pedersen *et al.* 2005; Ramsey *et al.* 2006). Taken together, these Ca^{2+} entry mechanisms are essential for signalling and store refilling. TRPM7 is constitutively active and can be modulated by internal Mg^{2+} or ATP concentrations (Nadler *et al.* 2001; Demeuse *et al.* 2006). We found that TRPM7 itself is not a store-operated channel, at least not in DT40 B lymphocytes. However, the Ca^{2+} current through TRPM7, and thus the channel domain of the protein, contributes significantly in maintaining the filling state of internal

Ca^{2+} stores under unstimulated resting conditions without modulating SOCE itself. To our knowledge, this is the first demonstration that the Ca^{2+} current through TRPM7 is linked to store refilling and maintenance of ER Ca^{2+} store contents.

Several studies discuss the effects of the TRPM7 knockdown or KO *in vitro* or *in vivo*, including inhibitory effects on T cell migration (Kuras *et al.* 2012), the requirement of TRPM7 expression to obtain cardiac automaticity (Sah *et al.* 2013b) or the propensity of TRPM7 expression levels to enhance the metastatic potential of invasive human breast cancer cells *in vivo* (Middelbeek *et al.* 2012). Furthermore, the TRPM7 kinase domain has been shown to regulate the Ca^{2+} -sensitivity of G-protein induced histamine release in primary mouse mast cells (Zierler *et al.* 2016). Ca^{2+} signalling in B cells is fundamentally important for the development of adaptive immunity (Baba & Kurosaki, 2016). The apparent dependence of SOCE on functional TRPM7 kinase activity may represent a feedback loop for B-cell receptor induced Ca^{2+} signalling that is in relation to the availability of intracellular Mg-ATP for the cell.

The connection between TRPM7 and SOCE should be considered when discussing TRPM7 effects because the possibility exists that these effects may be secondary to regulatory effects on SOCE. This may pertain to previous studies that have proposed a strong role for TRPM7 in mediating Ca^{2+} influx or Ca^{2+} oscillations and secondary physiological effects such as fibroblast proliferation (Du *et al.* 2010) or oocyte fertilization, egg activation and embryo development (Carvacho *et al.* 2016). Our data raise the distinct possibility, if not likelihood, that much of the TRPM7-dependent Ca^{2+} entry attributed to TRPM7 might result from the additional secondary, indirect support of SOCE via the kinase activity of TRPM7 and the regulation of I_{CRAC} . The dysregulation of Ca^{2+} homeostasis through either mechanism may drive the above-mentioned cellular responses, as well as the expression of malignant phenotypes, such as proliferation, migration, invasion and metastasis. Therefore, it might not be coincidental that both channels, CRAC and TRPM7, have been linked to various types of cancer (Chen *et al.* 2013). For this reason, both CRAC and TRPM7 channels may be promising targets for developing therapeutic strategies for diseases in which these mechanisms are involved.

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Additional information

Competing interests

The authors declare that they have no competing interests.

Author contributions

RP conceived and co-ordinated the study. TK and MF designed, performed and analysed the experiments and prepared the illustrations. AF performed and analysed experiments. FDH isolated and provided waixenicin A. RP, TK, MF and AF wrote and edited the paper. All authors reviewed the results and approved the final version of the manuscript submitted for publication.

Funding

This work was supported by DFG to TK (Ki 1821/1-1) and partially supported by NIH/NCI U54CA143727 (RP) and NIH/NIGMS P20 GM103466 (FDH).

Acknowledgements

DT40 B lymphocyte cell lines WT par, cTRPM7-KO, cWT, hTRPM7 Δ Kinase and inducible hTRPM7-KO V79.1, hTRPM7 KR were kindly provided by Andrew Scharenberg (Seattle) and Carsten Schmitz (Denver), respectively. DT40 B lymphocyte cell lines Orai1 KO, Orai2 KO, Orai1Orai2 DKO, STIM1-KO and STIM2-KO were kindly provided by Tomohiro Kurosaki (Osaka) and Donald Gill (Philadelphia). The authors thank the University of Hawaii Cancer Centre Genomics Shared Resource (supported by NIH/NCI CA071789-13) for performing the quantitative RT-PCR assays for the present study. Some of the services for this research were provided with the help of the University of Hawaii John A. Burns School of Medicine Molecular and Cellular Immunology Core, which is supported in part by NIH/NIGMS P20GM103516 from the Centres of Biomedical Research Excellence (COBRE) program.