

## CaT1 and the Calcium Release-activated Calcium Channel Manifest Distinct Pore Properties\*

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**The calcium release-activated calcium channel (CRAC) is a highly Ca<sup>2+</sup>-selective ion channel that is activated on depletion of inositol triphosphate (IP<sub>3</sub>)-sensitive intracellular Ca<sup>2+</sup> stores. It was recently reported that CaT1, a member of the TRP family of cation channels, exhibits the unique biophysical properties of CRAC, which led to the conclusion that CaT1 comprises all or part of the CRAC pore (Yue, L., Peng, J. B., Hediiger, M. A., and Clapham, D. E. (2001) *Nature* 410, 705–709). Here, we directly compare endogenous CRAC with heterologously expressed CaT1 and show that they manifest several clearly distinct properties. CaT1 can be distinguished from CRAC in the following features: sensitivity to store-depleting agents; inward rectification in the absence of divalent cations; relative permeability to Na<sup>+</sup> and Cs<sup>+</sup>; effect of 2-aminoethoxydiphenyl borate (2-APB). Moreover, CaT1 displays a mode of voltage-dependent gating that is fully absent in CRAC and originates from the voltage-dependent binding/unbinding of Mg<sup>2+</sup> inside the channel pore. Our results imply that the pores of CaT1 and CRAC are not identical and indicate that CaT1 is a Mg<sup>2+</sup>-gated channel not directly related to CRAC.**

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In non-excitabile cells, calcium transients upon receptor stimulation generally consist of two phases: a rapid rise due to Ca<sup>2+</sup> release from IP<sub>3</sub>-sensitive<sup>1</sup> intracellular Ca<sup>2+</sup> stores, which is followed by sustained Ca<sup>2+</sup> entry through store-operated plasma membrane channels (1). Of all store-operated channels, CRAC is functionally the best described, but its molecular identity and the mechanism of its store-dependent activation remain elusive (2). Several members of the TRP family, a group of cation channels related to the *Drosophila* transient receptor potential (*trp*) gene product (3, 4), have been implicated in store-dependent Ca<sup>2+</sup> influx (5–9), but until recently none of them was found to exhibit the unique biophysical properties of CRAC (3).

In a recent study, Yue *et al.* (10) reported that CaT1 (11), a TRP family member homologous to *Caenorhabditis elegans* OSM-9, manifests the pore properties and store dependence of CRAC, and it has been proposed that CaT1 comprises all or part of the CRAC pore (10, 12, 13). To further investigate this intriguing possibility, we made a direct comparison of CaT1, heterologously expressed in human embryonic kidney (HEK-293) cells, and endogenous CRAC in rat basophilic leukemia (RBL) cells. RBL cells were chosen because they were the source of the CaT1 clone used by Yue *et al.* (10) and have been widely used as a model system to study CRAC (2). We confirm the resemblance of currents through CRAC and CaT1 but additionally present a number of novel features that are incompatible with the proposed equality of the CaT1 and CRAC pores.

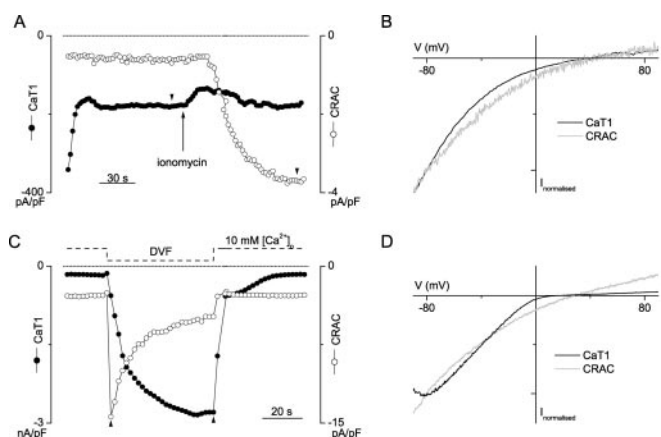
### EXPERIMENTAL PROCEDURES

**Cell Culture and Transfection**—HEK-293 cells and RBL-2H3 cells were grown in Dulbecco's modified Eagle's medium containing 10% (v/v) human serum, 2 mM L-glutamine, 2 units/ml penicillin, and 2 mg/ml streptomycin at 37 °C in a humidity-controlled incubator with 10% CO<sub>2</sub>. HEK-293 cells were transiently transfected with the pCINeo/IRES-GFP/mCaT1 vector using previously described methods (14), and electrophysiological recordings were performed between 8 and 24 h after transfection.

**Electrophysiology**—Patch clamp experiments were performed in the tight seal, whole-cell configuration at room temperature. Patch pipettes had DC resistances of 2–4 megaohms when filled with the standard intracellular solution containing (in mM): 145 cesium glutamate, 8 NaCl, 0, 3, or 10 MgCl<sub>2</sub>, 3.3 CaCl<sub>2</sub>, 10 EGTA, and 10 Hepes (pH 7.2). The free Ca<sup>2+</sup> concentration in this solution was calculated to be 100 nM. The standard extracellular solution contained (in mM): 145 NaCl, 2.8 KCl, 10 CsCl, 10 CaCl<sub>2</sub>, 2 MgCl<sub>2</sub>, 10 glucose, and 10 Hepes (pH 7.4). The divalent-free (DVF) solution contained (in mM): 150 NaCl, 10 EDTA, 10 glucose, and 10 Hepes (pH 7.4). To test the monovalent cation permeability sequence, DVF solutions were used in which NaCl was equimolarly replaced by LiCl, CsCl, KCl, or NMDG chloride.

Unless indicated differently, currents were measured in response to voltage ramps (–100 to +100 mV in 100 ms) applied at a frequency of 0.5 Hz from a holding potential of 0 mV. Series resistances were between 3 and 10 megaohms and were compensated for 50–80%. Voltage errors were generally <10 mV. Currents were sampled at 10 kHz and filtered at 2 kHz using an eight-pole Bessel filter. For the accurate determination of the currents through CRAC, background currents measured under identical ionic conditions but before store depletion were subtracted. Averaged data are expressed as the mean ± S.E. from

<sup>1</sup> The abbreviations used are: IP<sub>3</sub>, inositol triphosphate; CRAC, calcium release-activated calcium channel; HEK, human embryonic kidney; RBL, rat basophilic leukemia; DVF, divalent free; ECaC1, epithelial calcium channel 1; 2-APB, 2-aminoethoxydiphenyl borate; VOCC, voltage-operated calcium channel; NMDG, N-methyl-D-glucamine.



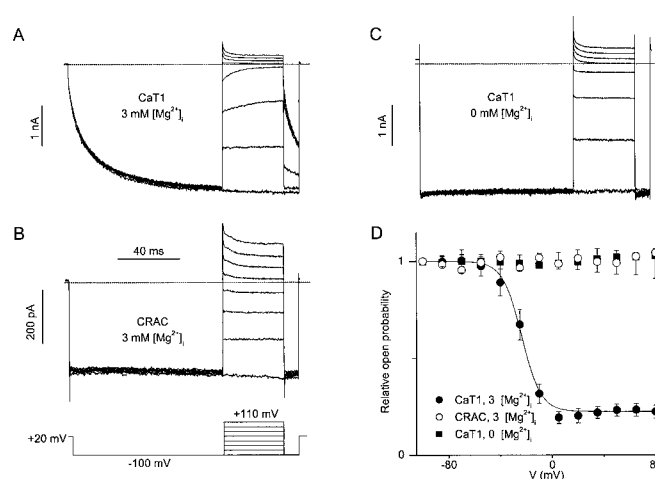
**FIG. 1. Comparison of the activation mechanism of CaT1 and CRAC and of their behavior in the absence of divalent cations.** A, time course of currents through CaT1 and CRAC at  $-80$  mV in standard extracellular solution ( $10$  mM  $\text{Ca}^{2+}$ ). CaT1 is instantaneously activated, whereas CRAC is activated by brief ( $5$  s) application of ionomycin ( $2$   $\mu\text{M}$ ). Currents were elicited by linear voltage ramps ( $2$  V/s) from  $-100$  to  $+100$  mV applied every  $2$  s from a holding potential of  $0$  mV. B, current-voltage relations for CaT1 and CRAC in  $10$  mM external  $\text{Ca}^{2+}$ . Traces were obtained in the experiments shown in A at the time points indicated by the arrowheads. For CRAC, the background current before application of ionomycin was subtracted. C, effect of switching from the standard extracellular solution to a DVF solution on currents through CaT1 and CRAC at  $-80$  mV. D, current-voltage relations for CaT1 and CRAC in DVF solution. Traces were obtained in the experiments shown in C at the time points indicated by the arrowheads. For CRAC, the background current in DVF before store depletion was subtracted. In panels B and D, traces were normalized to the current at  $-80$  mV.

$n$  cells. The significance of differences between means was evaluated using Student's unpaired  $t$  test.

## RESULTS

Immediately after obtaining the whole-cell configuration, CaT1-transfected HEK-293 cells exposed to  $10$  mM extracellular  $\text{Ca}^{2+}$  and with intracellular  $\text{Ca}^{2+}$  buffered at  $100$  nM (to maintain full  $\text{Ca}^{2+}$  stores) displayed a large membrane current (Fig. 1, A and B). This current subsequently decayed to reach a lower, but still robust, steady state level (Fig. 1A). The CaT1-mediated current was strongly inwardly rectifying (Fig. 1B) and reversed at positive potentials ( $39 \pm 7$  mV;  $n = 18$ ) in agreement with previous work (10). Depletion of intracellular  $\text{Ca}^{2+}$  stores with the  $\text{Ca}^{2+}$  ionophore ionomycin ( $2$   $\mu\text{M}$ ) did not induce any store-dependent activation of CaT1 but instead caused a transient inhibition of the CaT1-current by  $26 \pm 4\%$  ( $n = 8$ ) (Fig. 1A). A similar inhibition by  $30 \pm 6\%$  ( $n = 7$ ) was observed when  $\text{Ca}^{2+}$  stores were passively depleted by application of the sarcoendoplasmic reticulum  $\text{Ca}^{2+}$ -ATPase inhibitor, 2,5-di-*t*-butyl-1,4-benzohydroquinone ( $20$   $\mu\text{M}$ ). We attribute the inhibitory effect of store-depleting agents to an increase of intracellular  $\text{Ca}^{2+}$  as has been previously described for epithelial calcium channel (ECaC1; see Refs. 15 and 16), a close homologue of CaT1. Under identical intra- and extracellular ionic conditions, RBL-2H3 cells displayed only a very small background current that reversed at negative potentials ( $-41 \pm 2$  mV;  $n = 26$ ). Depletion of  $\text{Ca}^{2+}$  stores by extracellular application of  $2$   $\mu\text{M}$  ionomycin (Fig. 1A) or intracellular perfusion with  $\text{IP}_3$  ( $20$   $\mu\text{M}$ ; data not shown) activated CRAC, as indicated by the appearance of a  $\text{Ca}^{2+}$ -selective, inwardly rectifying current that reversed at positive potentials ( $42 \pm 7$  mV;  $n = 26$ ) (Fig. 1B).

The permeability of both CaT1 and CRAC in the presence of extracellular  $\text{Ca}^{2+}$  is at least 100 times higher for  $\text{Ca}^{2+}$  than for  $\text{Na}^+$ , but removal of all divalent cations from the extracellular medium renders both channel pores highly permeable for



**FIG. 2. Comparison of the voltage dependence of CaT1 and CRAC.** A and B, current traces for CaT1 and CRAC in DVF solution in response to the voltage step protocol shown below. C, current traces for CaT1 when  $\text{MgCl}_2$  was omitted from the intracellular solution. D, voltage dependence of the channel open probability for CaT1 and CRAC with  $3$  mM intracellular  $\text{Mg}^{2+}$  and for CaT1 in the absence of intracellular  $\text{Mg}^{2+}$  ( $n = 6-12$ ). Currents were measured during the final step to  $-100$  mV and normalized to the steady state current at this potential.

monovalent cations (2, 10). This loss of selectivity is presumably because of the removal of divalents from negatively charged residues (aspartates or glutamates) lining the  $\text{Ca}^{2+}$ -selective pore (17), as has been previously shown for voltage-gated  $\text{Ca}^{2+}$  channels and for ECaC1 (18, 19). When switching from the normal extracellular solution containing  $10$  mM  $\text{Ca}^{2+}$  to a DVF solution, the CaT1-mediated current gradually increased following an exponential time constant of  $9 \pm 4$  s ( $n = 14$ ) (Fig. 1C). We interpret this time-dependent increase of the current as the slow removal of the  $\text{Ca}^{2+}$ -induced inactivation of CaT1, as has been previously demonstrated for its close homologue, ECaC1 (15, 16). After reaching its maximal amplitude, the monovalent CaT1 current remained stable for more than 15 min (Fig. 1C and data not shown). In contrast, the current through CRAC, when preactivated by ionomycin in the normal extracellular solution containing  $10$  mM  $\text{Ca}^{2+}$ , reached its maximal amplitude immediately after switching to DVF solution and decayed subsequently with a time constant of  $12 \pm 4$  s ( $n = 17$ ) (Fig. 1C).

Current-voltage relationships obtained during linear voltage ramps in DVF solutions revealed several significant differences between CaT1 and CRAC (Fig. 1D). First, the reversal potential for CaT1 ( $18 \pm 2$  mV;  $n = 8$ ) was significantly less positive than for CRAC ( $45 \pm 6$  mV;  $n = 8$ ). We examined whether this difference reflected different monovalent cation permeability profiles of the CaT1 and CRAC channel pores and found that, although both channels display a  $\text{Na}^+ \approx \text{Li}^+ > \text{K}^+ > \text{Cs}^+ > \text{NMDG}$  permeability sequence, CRAC has a much lower relative  $\text{Cs}^+$  permeability ( $P_{\text{Cs}}/P_{\text{Na}} = 0.48 \pm 0.03$  for CaT1 versus  $0.12 \pm 0.04$  for CRAC;  $p < 0.001$ ). Second, inward rectification was much more pronounced for CaT1 than for CRAC. The rectification score (defined as the ratio of the current amplitudes at potentials  $40$  mV negative and positive to the reversal potential) was  $9.9 \pm 1.1$  ( $n = 15$ ) for CaT1 compared with  $1.5 \pm 0.1$  for CRAC ( $n = 19$ ;  $p < 0.001$ ). Third, the current-voltage relationship for CaT1 currents measured using the linear voltage ramp protocol displayed a negative slope at potentials below  $-80$  mV, which was not observed for CRAC. As shown below, this negative slope originates from the time-dependent removal of intracellular  $\text{Mg}^{2+}$  block.

Monovalent cation currents through CaT1 activated during membrane hyperpolarization from  $+20$  mV to  $-100$  mV and deactivated during subsequent steps to less hyperpolarized

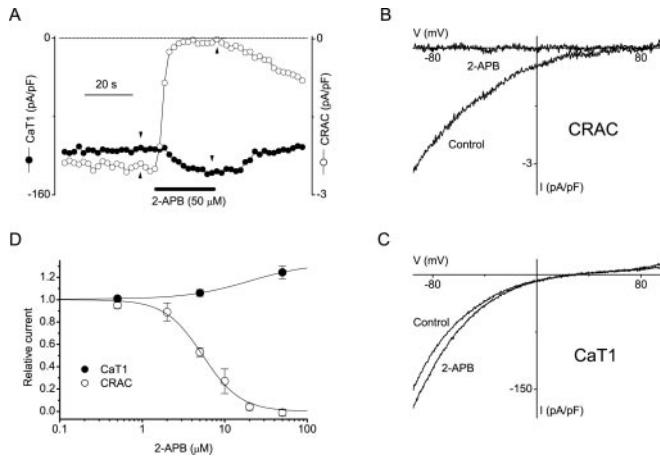
potentials (Fig. 2A). The fraction of open channels at the end of each voltage step was assessed from the initial current amplitude during a final step to  $-100$  mV and exhibited a Boltzmann dependence on membrane potential (20), with a slope factor of 8 mV and half-maximal deactivation at  $-24$  mV (Fig. 2D). This voltage dependence was abolished when  $Mg^{2+}$  was omitted from the intracellular solution (Fig. 2, C and D), indicating that the gating mechanism of CaT1 is a voltage-dependent block/unblock of the channel pore by intracellular  $Mg^{2+}$ . Using pipette solutions containing different concentrations of  $Mg^{2+}$ , we found that half-maximal block at 0 mV was achieved with  $\sim 200$   $\mu M$  intracellular  $Mg^{2+}$ . In contrast, monovalent cation currents through CRAC did not exhibit such voltage dependence (Fig. 2, B and D), not even with 10 mM intracellular  $Mg^{2+}$  ( $n = 5$ ; data not shown).

It should be noted that CaT1 currents remained strongly inwardly rectifying in symmetric monovalent cation solutions and in the absence of intracellular  $Mg^{2+}$ , indicating that rectification represents an intrinsic pore property of CaT1. Moreover, the strong inward rectification and voltage-dependent  $Mg^{2+}$  block of CaT1 were preserved in cell-free inside-out patches (data not shown).

2-aminoethoxydiphenyl borate (2-APB), an inhibitor of  $IP_3$  receptors, is known to inhibit store-dependent  $Ca^{2+}$  influx

pathways in a variety of cells (21–25). We confirmed that 50  $\mu M$  2-APB caused a rapid and complete block of CRAC in RBL cells (Fig. 3A), which was voltage-independent (Fig. 3B and data not shown) and slowly reversible upon washout (Fig. 3A). The inhibitory effect of 2-APB on CRAC was dose-dependent with half-maximal inhibition at 5.5  $\mu M$  (Fig. 3D). In contrast to a recent study (25), we did not observe a consistent initial increase in CRAC activity upon application of 2-APB. However, we cannot exclude the possibility that some 2-APB-dependent potentiation was masked by the rapid blocking effect of the drug. Although 2-APB has been shown to be membrane-permeable, two observations indicate that the target for 2-APB block of CRAC is exposed to the extracellular medium. First, the onset of block occurred with a time constant of  $\sim 2$  s (Fig. 3A), which is almost as fast as block of CRAC by extracellular  $Zn^{2+}$ ,  $Cd^{2+}$ , or  $La^{3+}$  (data not shown) and much faster than what has been reported for  $IP_3$  receptor-mediated effects of 2-APB in intact cells (22). Second, dialysis of the cytoplasm with a pipette solution supplemented with 50  $\mu M$  2-APB did not markedly prevent activation of CRAC by ionomycin (data not shown; see also Ref. 23). From this we conclude that the interaction site for 2-APB is either on the extracellular part of the CRAC channel itself or at least closely associated therewith. In remarkable contrast with its inhibitory effects on CRAC, 50  $\mu M$  2-APB potentiated CaT1-mediated currents by  $\sim 25\%$  (Fig. 3, A and D). The potentiating effect of 2-APB did not show marked voltage dependence (Fig. 3C), was fully reversible upon washout (Fig. 3A), and could be repeated several times in the same cell. A similar but less pronounced potentiation was obtained with 5  $\mu M$  2-APB (Fig. 3D). Application of 250  $\mu M$  2-APB also caused a potentiation of CaT1, which was mostly followed by a slow and moderate inhibition of the currents by  $\sim 20\%$  and/or activation of linear leak currents. Thus, CRAC is at least 100 times more sensitive for 2-APB block than CaT1.

According to Yue *et al.* (10), the properties of CaT1, notably its activation by active store depletion, were dependent on the expression level of the channel protein. Therefore, we wanted to make sure that the differences we observed between CaT1 and CRAC were independent on the number of CaT1 channels per cell. Like Yue *et al.* (10), we measured CaT1 currents within 12 h after transfection and selected for cells that had current densities below 100 pA/pF at  $-100$  mV with 10 mM extracellular  $Ca^{2+}$  (mean peak amplitude,  $78 \pm 9$  pA/pF;  $n = 8$ ). Despite the fact that the average CaT1 current density in this group of cells was  $\sim 5$  times lower than the overall average ( $356 \pm 44$  pA/pF;  $n = 25$ ), they displayed all of the above



**FIG. 3. Comparison of the effects of 2-APB on CaT1 and CRAC.** A, time course of currents through CaT1 and CRAC measured at  $-80$  mV in standard extracellular solution showing the effect of 50  $\mu M$  2-APB. B and C, current voltage relations for CRAC (B) and CaT1 (C) obtained during the experiments shown in A at the time points indicated by the arrowheads. D, dose-response relation for the effect of 2-APB on CaT1 and CRAC. Data points ( $n = 4-9$ ) were normalized to the current measured in the absence of 2-APB.

**TABLE I**  
Summary of similarities and differences between CaT1 and CRAC

Properties of CaT1 and CRAC were determined in the present study unless indicated otherwise. pS, picosiemens.	CaT1	CRAC (RBL cells)
<b>Similarities</b>		
Calcium selectivity	$P_{Ca}/P_{Na} > 100^a$	$P_{Ca}/P_{Na} > 100^b$
Anomalous mole fraction behavior	Yes <sup>a</sup>	Yes
Divalent permeability	$Ca^{2+} > Ba^{2+} \approx Sr^{2+} > Mg^{2+}$ <sup>a</sup>	$Ca^{2+} > Ba^{2+} \approx Sr^{2+} > Mg^{2+}$ <sup>b</sup>
Monovalent permeability	$Na^+ \approx Li^+ > K^+ > Cs^+$	$Na^+ \approx Li^+ > K^+ > Cs$
Effect of $La^{3+}$ , $Zn^{2+}$ , $Cd^{2+}$	Block	Block <sup>c</sup>
Single-channel conductance (150 $Na^+$ )	42–58 pS <sup>a</sup>	55 pS <sup>d</sup>
<b>Differences</b>		
Effect of ionomycin	Inhibition	Activation
Current in DVF	Stable	Transient
Inward rectification in DVF	Strong	Weak
$P_{Cs}/P_{Na}$	$0.48 \pm 0.03$	$0.12 \pm 0.04$
Effect of intracellular $Mg^{2+}$	Voltage-dependent block	No effect
Effect of extracellular 2-APB	Potentiation	Inhibition

<sup>a</sup> From Ref. 10.  
<sup>b</sup> From Ref. 32.  
<sup>c</sup> From Ref. 33.  
<sup>d</sup> From Ref. 34.



described CaT1 features: inhibition of 24% by 2  $\mu\text{M}$  ionomycin ( $n = 2$ ); strong inward rectification in DVF solution (rectification score,  $12.7 \pm 3.2$ ;  $n = 4$ ); relative  $\text{Cs}^+$  permeability ( $P_{\text{Cs}}/P_{\text{Na}}$ ) of  $0.52 \pm 0.06$  ( $n = 5$ ); voltage-dependent gating in the presence of intracellular  $\text{Mg}^{2+}$  with a slope factor of 8 mV and half-maximal deactivation at  $-25$  mV ( $n = 4$ ); potentiation to 121% of control by 50  $\mu\text{M}$  2-APB ( $n = 2$ ). Thus, the biophysical and pharmacological properties that distinguish heterologously expressed CaT1 from endogenous CRAC in RBL cells are not influenced by the expression level of the CaT1 channel protein. Moreover, virtually identical results were obtained in CaT1-expressing Chinese hamster ovary cells (data not shown), indicating that the expression system does not significantly affect the properties of the CaT1 channel.

#### DISCUSSION

To evaluate the hypothesis that CaT1, a member of the TRP family of cation channels, constitutes all or part of CRAC, the calcium release-activated  $\text{Ca}^{2+}$  channel, we made a direct comparison of the biophysical and pharmacological properties of both channels under identical experimental conditions. At first sight, the similarities between CRAC and CaT1 (Table I) are indeed striking. However, it should be noted that many of the common characteristics of the CRAC and CaT1 pores (high  $\text{Ca}^{2+}$  selectivity; anomalous mole fraction effect; divalent and monovalent cation permeability sequence; block by inorganic cations; loss of selectivity in the absence of divalents; single-channel conductance to  $\text{Na}^+$  in the range of 10–100 picosiemens) are not only shared by ECaC1 (15, 26, 27), which has a pore region virtually identical to the pore region of CaT1, but also shared to a large extent by voltage-operated  $\text{Ca}^{2+}$  channels (VOCCs) (17, 28–30), which have pore regions that are only weakly homologous to CaT1/ECaC1. In VOCCs and ECaC1, high  $\text{Ca}^{2+}$  selectivity depends on negatively charged amino acids (glutamates and/or aspartates) in the pore region (18, 19). It is believed that high affinity binding of  $\text{Ca}^{2+}$  ions to these sites prevents, by electrostatic repulsion, monovalent cations from passing through the pore while simultaneously allowing high rates of  $\text{Ca}^{2+}$  flux (17). Such a configuration implies that the pore becomes readily permeable to monovalent cations when the extracellular concentration of  $\text{Ca}^{2+}$  and other divalent cations is so low that the  $\text{Ca}^{2+}$ -binding sites are no longer occupied (20). Intermediate  $\text{Ca}^{2+}$  concentrations block monovalent permeation but yield only small  $\text{Ca}^{2+}$  fluxes. Thus, the conductance of these channels as a function of the extracellular  $\text{Ca}^{2+}$  concentration goes through a minimum, a phenomenon termed anomalous mole fraction behavior, which has been observed in VOCCs (17), ECaC1/CaT1 (10, 27), and CRAC (31). Moreover, because negatively charged  $\text{Ca}^{2+}$ -binding sites generally display a high affinity for transition metals such as  $\text{Cd}^{2+}$ ,  $\text{Zn}^{2+}$ , or  $\text{La}^{3+}$  (20), it is not unexpected that these cations exert an inhibitory effect on the different  $\text{Ca}^{2+}$  selective channels. Thus, it appears that most of the pore properties common for CRAC and CaT1 are features inherent to all highly  $\text{Ca}^{2+}$ -selective pores in biological membranes.

As summarized in Table I, CaT1 differs from CRAC in RBL cells in the following salient features: effect of ionomycin-induced store depletion; current decay in DVF solution; relative permeability for  $\text{Cs}^+$  versus  $\text{Na}^+$ ; inward rectification; voltage-dependent block by intracellular  $\text{Mg}^{2+}$ ; and effect of extracellular 2-APB. Obviously, some of these differences might be explained by cell-specific modulation of channel function. However, monovalent permeability, channel rectification, and voltage-dependent block represent biophysical properties that are

directly related to the channel pore (20). Thus, our data demonstrate that the pore properties of CRAC and CaT1 are clearly distinct, thereby ruling out the possibility that CaT1 encompasses the full CRAC pore. We cannot definitely exclude the possibility that the CRAC pore consists of CaT1 plus an additional subunit, but this additional channel subunit would then have to abolish the intracellular  $\text{Mg}^{2+}$ -sensitivity of CaT1, provide for the 2-APB inhibition, and significantly alter the ion selectivity and inward rectification of the putative heteromultimeric channel. An alternative and presumably more likely explanation is that CRAC and CaT1 are two distinct channels with a number of overlapping properties but nevertheless encoded by different genes. In that case, we have provided a number of useful tools to discriminate between both channels in native tissue.

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## **CaT1 and the Calcium Release-activated Calcium Channel Manifest Distinct Pore Properties**

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