Single Channel Function of Inositol 1,4,5-trisphosphate Receptor
Type-1 and -2 Isoform Domain-Swap Chimeras

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ABSTRACT The InsP₃Rs proteins have three recognized domains, the InsP₃-binding, regulatory/coupling, and
channel domains (Mignery, G.A., and T.C. Südhof. 1990. EMBO J. 9:3893–3898). The InsP₃ binding domain
and the channel-forming domain are at opposite ends of the protein. Ligand regulation of the channel must involve
communication between these different regions of the protein. This communication likely involves the interced-
ing sequence (i.e., the regulatory/coupling domain). The single channel functional attributes of the full-length
recombinant type-1, -2, and -3 InsP₃R channels have been defined. Here, two type-1/type-2 InsP₃R regulatory/cou-
pling domain chimeras were created and their single channel function defined. One chimera (1-2-1) contained
the type-2 regulatory/coupling domain in a type-1 backbone. The other chimera (2-1-2) contained the type-1 reg-
ulatory/coupling domain in a type-2 backbone. These chimeric proteins were expressed in COS cells, isolated,
and then reconstituted in proteoliposomes. The proteoliposomes were incorporated into artificial planar lipid bi-
layers and the single-channel function of the chimeras defined. The chimeras had permeation properties like that
of wild-type channels. The ligand regulatory properties of the chimeras were altered. The InsP₃ and Ca²⁺ regula-
tion had some unique features but also had features in common with wild-type channels. These results suggest
that different independent structural determinants govern InsP₃R permeation and ligand regulation. It also sug-
gests that ligand regulation is a multideterminant process that involves several different regions of the protein.
This study also demonstrates that a chimera approach can be applied to define InsP₃R structure-function.

KEY WORDS: inositol trisphosphate receptor • Ca²⁺ release channel • intracellular Ca²⁺ signaling

INTRODUCTION

The inositol 1,4,5-trisphosphate receptor (InsP₃R)* genes encode intracellular Ca²⁺ release channels that are fundamental to many intracellular Ca²⁺-signaling processes in mammalian cells. The InsP₃Rs Ca²⁺-release channels are found in intracellular membranes, predominately the ER. These channels mobilize intracellular Ca²⁺ in response to InsP₃ generated by receptor-activated hydrolysis of phosphatidylinositol 4,5 bisphosphate in the inner leaflet of the surface membrane.

The InsP₃Rs are part of a family of three homologous proteins. The InsP₃Rs proteins have three recognized domains: the InsP₃-binding, regulatory/coupling, and channel domains (Mignery and Südhof, 1990, 1993; Südhof et al., 1991; Blondel et al., 1993). The ligand binding and channel domains are relatively well conserved between the different InsP₃R family members. The least conserved domain is the regulatory/coupling domain that contains several potential regulatory sites (including the putative Ca²⁺ binding region; Mignery et al., 1992; Sienaert et al., 1996, 1997). The regulatory/coupling domain physically links the amino-terminal InsP₃-binding domain to the carboxyl-terminal channel domain.

One consequence of this structural heterogeneity is thought to be the isoform-specific functional differences between the different InsP₃R family members observed in single channel studies (Watras et al., 1991; Perez et al., 1997; Hagar et al., 1998; Mak et al., 1998; Ramos-Franco et al., 1998a). A prominent isoform-specific functional attribute is how InsP₃ and cytosolic Ca²⁺ regulate these channels. For example, the Ca²⁺ sensitivity of the type-1 and -2 InsP₃R channels are quite different (Bezprozvanny et al., 1991; Ramos-Franco et al., 1998a). Under identical experimental conditions, one is sharply bell-shaped (type-1) while other is not (type-2). The type-1 and -2 channels are also regulated by InsP₃ differently. The type-2 InsP₃-R has the higher InsP₃ affinity (Südhof et al., 1991; Newton et al., 1994; Perez et al., 1997; Miyakawa et al., 2001) and higher InsP₃ efficacy (Ramos-Franco et al., 1998a).

The function of single recombinant type-1 and -2 InsP₃R channels has been defined and this function matches that of their native counterparts (Kaznacheeva et al., 1998; Ramos-Franco et al., 1998b, 2000; Mak et al., 2000; Boehning et al., 2001). The

*Abbreviation used in this paper: InsP₃R, inositol 1,4,5-trisphosphate receptor.
InsP$_3$ binding site and the channel pore are in distinct regions at opposite ends of the protein. The cytosolic region encompassing the InsP$_3$ binding domain region apparently undergoes a significant conformational change upon ligand binding (Mignery and Südhof, 1990). Despite the large linear separation of the InsP$_3$ binding and channel domains, Boehning and Joseph (2000) reported that there may be a direct association of these two domains. Thus, ligand regulation of the pore may involve interactions with distant regions of the protein (considering the linear protein sequence). Intuitively, these interactions may depend on properties of the intervening sequence (i.e., the regulatory/coupling domain). Single channel data suggest that these interactions may be different in the InsP$_3$R isoforms.

To test this possibility, the single channel function of type-1 and type-2 InsP$_3$R regulatory/coupling domain chimeras was defined here. Two domain-swap InsP$_3$R chimeras were tested. One chimera contained the type-2 regulatory/coupling domain in a type-1 backbone (i.e., the 1-2-1 chimera). The other chimera contained the type-1 regulatory/coupling domain in a type-2 backbone (i.e., the 2-1-2 chimera). The ligand regulatory properties of the chimeric channels were analyzed and compared with their full-length progenitors.

**MATERIALS AND METHODS**

**Materials**

Lipids (phosphatidylcholine, phosphatidylethanolamine, and phosphatidylserine) were obtained from Avanti Polar Lipids, Inc. D-mylo inositol 1,4,5-trisphosphate was purchased from LC Laboratories. Ryanodine was purchased from Calbiochem. All reagents and enzymes were of molecular biology grade.

**Construction and Transfection of InsP$_3$R Expression Vectors**

The chimeras were constructed using the strategy described below. The type-1 ligand binding, type-2 coupling, and type-1 channel domain chimera (1-2-1) were constructed exploiting a unique Bst BI restriction site located at the beginning of the channel domain. The type-2 ligand binding domain was produced using the strategy described below.

To test this possibility, the single channel function of type-1 and type-2 InsP$_3$R regulatory/coupling domain chimeras was defined here. Two domain-swap InsP$_3$R chimeras were tested. One chimera contained the type-2 regulatory/coupling domain in a type-1 backbone (i.e., the 1-2-1 chimera). The other chimera contained the type-1 regulatory/coupling domain in a type-2 backbone (i.e., the 2-1-2 chimera). The ligand regulatory properties of the chimeric channels were analyzed and compared with their full-length progenitors.

A third intermediate (pC) exploiting the unique Bst BI restriction site located at the beginning of the channel domain in the type 1 receptor was prepared. Specifically, the ligand-binding and regulatory/coupling type-2 sequence (residues 1–2,177) was inserted into the type-1 cDNA at residue 2225. This was accomplished by combining the ~6.4 kbp EcoRI-HindIII (nucleotides 1–6383) fragment from the full-length type-2 cDNA (pInsP$_3$R-T2) with a PCR product spanning the Hind III site at nucleotide 6383 and extending 3’ through nucleotide 6792 that had a BstBI site introduced at nucleotide 6779. The primers for this amplification were TCGGCTCCAAGGCTTCCGGCC and CGGATCTTGAATCTCTGGATTCGCC. This product was digested with HindIII-XbaI and the two fragments were ligated into EcoRI-XbaI cut pCMV5. Intermediate pC was digested with Bst BI-XbaI and the 2.47 kbp BstBI-XbaI fragment of the type-1 receptor (pInsP$_3$R-T1) was inserted to form a type-2 ligand binding/coupling/type-1 channel domain chimera (2-2-1, pD). The addition of the BstBI site at the 3’ end of the type-2 sequence resulted in the substitution of Y2178 to an F. The inter-domain linkages in the chimeras and native receptors are shown in Fig. 1.

The final assembly of the 1-2-1 chimera was accomplished by digesting pB with Bam HI and inserting the 2.85 kbp BamHI fragment of pB containing nucleotides 6530–6780 of the type-2 coupling domain and the type-1 channel domain (nucleotides 7000–9465).

The type-2 ligand binding and channel domains interspersed with the type 1 coupling/regulatory domain was constructed as follows. The type-1 InsP$_3$R (pInsP$_3$R-T1) was digested with BstBI-Nhe I and the resulting 11.33 kbp fragment was isolated. To this the Nhe I-BstBI fragment (nucleotides 7637–8535) of the type-2 receptor was inserted generating intermediate pE. The type-2 InsP$_3$R cDNA pH2-2 (Südhof et al., 1991) was linearized with BamHI and sequences spanning residues 6781–7637 were PCR amplified introducing a BstBI site at nucleotide 6781 (primers: CGCAGAATTCCTTGCACACAAACAGCACTTCGAG and GTGCGAGAGTCGGC). This fragment was digested with BstBI and inserted into BstBI-digested pE resulting in a 1-1-2 chimera (intermediate pH). Note, that in addition to introducing the BstBI site in this PCR product, V2180 was changed to an I. This allows the type-2 channel domain to be introduced into intermediate pH such that there are no spurious residues introduced and the actual type-1/-2 junction begins between I2226 of the type-1 receptor and F2181 of the type-2 receptor. The full-length type-1 cDNA (pInsP$_3$R-T1) EcoRI-XbaI fragment was inserted into similar sites of pGEM3Z in which the SphI site was abolished. Residues 236–4733 were excised by digestion with Eco RI-SphI and the resulting vector fragment was ligated with a PCR product spanning nucleotides 2142–4746 that introduces an XhoI site spanning residues 606 and 607 of the type-1 receptor to generate intermediate pG. Next, nucleotides 1–2081 of the type-2 InsP$_3$R cDNA were PCR amplified with the following oligonucleotides: GGAATTCCTCGGGACAGAGTCGGC and GGCTAGCTTTGGATTCGCC, which introduced a unique XhoI, site spanning residues 606 and 607. This PCR product was ligated into Eco RI-XhoI-digested intermediate pG at similar sites and then the entire EcoRI XbaI fragment was inserted into pCMV5 to generate a 2-1-1 chimera (intermediate pH). The final assembly of the 2-1-2 chimera was achieved by digesting the 1-1-2 plasmid (pF) with EcoRI-XhoI and ligating the EcoRI-XhoI fragment of the 2-1-1 chimera (pH).

PCR products and junctional sequences in the chimeric constructs were determined by DNA sequence analysis using either a Sequenase 2.0 sequencing kit (Amersham Biosciences) or outsourced to the Loyola University sequencing facility using an ABI Model 310 sequencer.
Equivalent numbers of COS-1 cells were transiently transfected with the chimeric cDNA using the DEAE-dextran method. Cells were incubated at 37°C (5% CO₂). Typical transfection efficiencies were generally ≥60% as determined by indirect immunofluorescence or via a green fluorescent reporter.

**CHAPS Solubilization and Gradient Sedimentation**

The transfected COS-1 cells were harvested 48–72 h posttransfection and microsomes were prepared following established procedures (Mignery et al., 1990; Ramos-Franco et al., 1998b, 2000). Briefly, cells were washed with PBS and scrapped into 50 mM Tris-HCl pH 8.3, 1 mM EDTA, 1 mM 2-mercaptoethanol, 1 mM PMSF, and lysed by 40 passages through a 27-gauge needle. Membranes were pelleted by a 20-min centrifugation (289,000 g), resuspended in buffer, and frozen at −80°C. Microsomal fractions were solubilized in 50 mM Tris-HCl pH 8.3, 1 mM EDTA, 1 mM 2-mercaptoethanol, 1 mM PMSF, 1.8% CHAPS on ice for 1 h. Insoluble fractions were eliminated by a 10-min centrifugation at 289,000 g. The supernatant containing solubilized channel complex were fractionated through 5–20% continuous sucrose (wt/vol) gradients or via a green fluorescent reporter.

**Expression and Isolation of Chimeric InsP₃R Channels**

The amino-terminal end of the receptor forms the ligand binding domain and the carboxyl terminus, which harbors the multiple membrane spanning sequences responsible for subunit assembly (Galvan et al., 1999; Galvan and Mignery, 2002), forms the permeation pathway (Ramos-Franco et al., 1999). Interspersed between these two domains is the coupling or modulatory domain (nearly 1,700 amino acids) that undergoes an apparent conformational change upon InsP₃ binding. This change may couple ligand occupancy to channel gating (Mignery and Südhof, 1990). In addition, this region contains several sequence motifs that may regulate the channel and include PKA-phosphorylation sites, ATP, calmodulin, and Ca²⁺ binding sites as well as alternative splicing events. Recently, electron microscopy of negatively stained InsP₃R revealed that the protein also undergoes a conformational change in response to elevated Ca²⁺ and that this change may also be important to regulation of InsP₃ channel activity (Hamada et al., 2002).

Two chimeric InsP₃Rs were used in this study. The 1-2-1 chimera contains the type-2 coupling domain inserted in the type-1 backbone (i.e., type-1 binding and channel domains). The 2-1-2 chimera contains the type-1 coupling domain inserted in the type-2 backbone (i.e., type-2 binding and channel domains). The chimeric receptors were constructed such that the junctional sequences between heterologous domains were relatively conserved (see Fig. 1). COS-1 cells were transiently transfected with the chimeric 1:2:1 and 2:1:2 expression constructs (Gorman, 1985). Cells were har-
vested 48–72 h posttransfection. Microsomal protein (10 µg) from the transfected cells was immunoblotted using type-1- and 2-specific amino- and carboxyl-terminal antibodies to confirm expression (Fig. 2). The antibodies directed against the type-1 isoform (T1NH and T1C) reacted with only the 1-2-1 chimera and type-1 control, pInsP3R-T1 (panels T1NH and T1C). The type-2 antibodies (T2NH and T2C) only reacted to the 2-1-2 chimera or the full-length type-2 control, pInsP3R-T2 (panels T2NH and T2C). Both chimeric proteins expressed at significant levels. There was, however, some weak cross-reactivity observed in all lanes apparently due to endogenous type-2 or perhaps type-3 isoforms in the COS cells using T2C (bottom). The type-2 and -3 receptor isoforms share some sequence homology over the carboxyl terminus. Additionally, the chimeras integrity was evaluated for the ability to bind InsP3. The two chimeras bound similar levels of 3H InsP3 as their wild type progenitors (unpublished data) and together with the recognition of the recombinant proteins by the appropriate carboxyl-terminal antibodies, we concluded that the constructs were faithfully expressed as full-length products.

The transfected COS-1 microsomal proteins were CHAPS solubilized and enriched on 5–20% sucrose linear gradients for incorporation into proteoliposomes. In all cases the chimeric proteins sedimented to positions on the sucrose gradients consistent with those of tetramers (unpublished data).

**Permeation Properties of the Chimeric InsP3R Channels**

Proteoliposomes containing the chimeric InsP3R channels were fused into artificial planar lipid bilayers. The method used was identical to that applied previously to define the single channel function of the native and recombinant InsP3R channels (Perez et al., 1997; Ramos-Franco et al., 1998a,b). The single channel function the full-length type-1 and -2 recombinant InsP3R channels were previously compared by Ramos-Franco et al. (2000). The permeation properties of the type-1 and -2 recombinant channels were indistinguishable. This is consistent with the highly conserved nature of the InsP3R channel domain.

The permeation properties of the two chimeric channels were defined in solutions containing 220 mM CsHEPES cis (20 mM trans), pH 7.4, and 1 mM EGTA ([Ca2+]FREE = 200 nM). The cis-chamber solution also contained 1 µM InsP3. The channel was spontaneously active exhibiting frequent and rapid opening events. Sample single channel recordings obtained from the 1-2-1 chimeric channel at three different steady-state membrane potentials are presented in Fig. 3 (left). Open events at −70 mV are shown as downward deflec-
tions (zero current level marked). Open events at −20 and 0 mV are shown as upward deflections. Averaged unit current values obtained from both the 1-2-1 chimera (filled symbols; n = 9) and 2-1-2 chimera (open symbols; n = 8) are plotted as a function of membrane potential (Fig. 3). The slope conductances of the chimeric channels were nearly identical (220 pS). Unitary current reversed near −45 mV, indicating the chimeric channels were cation selective (i.e., permeating Cs⁺). These data are very similar to those obtained in similar studies on the recombinant full-length type-1 and -2 channels (Ramos-Franco et al., 1998b, 2000). The data is also similar to that obtained from native full-length type-1 and -2 channels (Ramos-Franco et al., 1998a). Thus, the permeation pathway of the chimeric channels functions quite normal retaining features common to their native and full-length recombinant progenitors.

*InsP₃ Sensitivity of the Chimeric InsP₃R Channels*

The InsP₃ sensitivity of the chimeric channels was defined. Sample single channel recordings of both chimeras in the absence of InsP₃ and three different InsP₃ concentrations are shown in Fig. 4 A. These recordings were made at 0 mV and open events are shown as upward deflections. The current carrier was Cs⁺ and solutions contained 200 nM free Ca²⁺. The addition of InsP₃ (cis-chamber) activated both chimeric channels. Application 1 µM InsP₃ robustly activated the 1-2-1 chimera to high Po levels but activated the 2-1-2 chimera to a lesser extent. The relatively short single channel recordings shown (Fig. 4 A) were selected to illustrate relative changes in open probability (Po).

Mean steady-state Po of the two chimeric channels at different InsP₃ concentrations is plotted in Fig. 4 B. These data represent the behavior of 11 different 2-1-2 channels and 8 different 1-2-1 channels. The dose of InsP₃ required for half maximal activation (EC₅₀) of the 2-1-2 chimeric channel was 13.9 ± 38 nM (Hill coefficient = 1.27). The EC₅₀ of the 1-2-1 chimeric channel was 261 ± 114 nM (Hill coefficient = 1.43). These values are statistically different (P < 0.05). Previously published data collected on full-length recombinant type-1 (Ramos-Franco et al., 1998b) and type-2 (Ramos-Franco et al., 2000) InsP₃R channels are presented as dashed lines. The EC₅₀ of InsP₃ activation of the full-length recombinant type-1 channel was 97 ± 66 nM. The EC₅₀ full-length type-2 recombinant channel was 122 ± 100 nM. The InsP₃ EC₅₀ for the 1-2-1 chimera was in a range similar to that of the full-length recombinant channels. The InsP₃ EC₅₀ for the 2-1-2 chimera was shifted substantially to lower InsP₃ concentrations. Different efficacies of InsP₃ in the activation of the chimeric channels were also noted. At very low InsP₃ concentration (i.e., 1 nM, the lowest dose applied here), the 2-1-2 chimera had a Po of near 0.18. This was similar to the Po of the 2-1-2 chimera channel when no InsP₃ was added. In contrast, the Po of the 1-2-1 chimera was near 0 at 1 nM InsP₃ or when no InsP₃ was added. The 1-2-1 chimera had a substantially higher maximal Po ~2–4 fold higher than either of the full-length channels. Thus, InsP₃ regulation of the pores of the chimeric channels is not identical to that of their full-length counterparts. The fact that InsP₃ regulates their pores and that this regulation has certain similarities with wild-type InsP₃ regulation implies that, (a) the ligand-binding domains in the chimeric channels modulate the interaction of InsP₃ binding and channel domains and, (b) this interaction depends on the identity of the interceding coupling or modulatory sequence.
The gating kinetics (open and closed dwell times) of the two chimeric channels in the presence of 1 μM InsP₃ are shown in Fig. 5. Under these conditions, the 1-2-1 chimera was characterized by frequent long open events and short closed events. This was not the case for the 2-1-2 chimera. The gating kinetics of the two chimeras in the absence of InsP₃ and at different InsP₃ concentrations are summarized in Table I.

The Ca²⁺ Sensitivity of the Chimeric InsP₃R Channels
The Ca²⁺ sensitivity of the 2-1-2 chimeric channel is shown in Fig. 6. These data were collected in the presence of 1 μM InsP₃ (cis-chamber). Sample single channel recordings at two different free Ca²⁺ concentrations are presented (Fig. 6, left). The free Ca²⁺ concentration was adjusted by altering the Ca²⁺-EGTA mixture and verified using a Ca²⁺ electrode. Channel activity (Po) of the 2-1-2 chimeric channel was similar at 10 nM and 10 μM free Ca²⁺ (cis-chamber). Mean Po (± SE, n = 4) of the 2-1-2 chimeric channel at different free Ca²⁺ concentrations is plotted in Fig. 6 (right). These data show that the activity of the 2-1-2 chimera was not Ca²⁺ dependent over a wide free Ca²⁺ range (10 nM to 100 μM). This implies that the interceding type-1 regulatory/coupling domain (i.e., containing the putative Ca²⁺ regulatory sites) does not confer Ca²⁺ binding information to the surrounding type-2 backbone.

The Ca²⁺ sensitivity of the 1-2-1 chimeric channel is shown in Fig. 7. These data were also collected in the presence of 1 μM InsP₃. Sample single channel recordings at three different free Ca²⁺ concentrations are presented (Fig. 7, left). Channel activity (Po) of the 1-2-1 chimera at a low free Ca²⁺ concentration (20 nM) was relatively high (~0.6). Compared with the either of the
recombinant full-length channels (type-1 or -2; Ramos-Franco et al., 1998b, 2000), this is a usually high activity level under these conditions. From this apparently basal activity level (at 1 μM InsP₃), the Po of the 1-2-1 chimera increased to ~0.85 at intermediate free Ca²⁺ concentrations (150 nM) and then decreased again to ~0.6 at higher Ca²⁺ concentrations (e.g., 10 μM). Mean Po (± SE, n = 7) of the 1-2-1 chimeric channel is plotted (solid symbols) as a function of free Ca²⁺ concentrations in Fig. 7 B. For comparison, the Ca²⁺ dependencies of the full-length recombinant type-1 and -2 InsP₃R channels are also shown (Fig. 7 B, dashed lines). The full-length recombinant type-1 channel has the classical bell-shaped Ca²⁺ dependency (Ramos-Franco et al., 1998b). The full-length type-2 recombinant channel has a more sigmoidal Ca²⁺ dependency (Ramos-Franco et al., 2000). The Ca²⁺ dependency of the 1-2-1 chimera has a bell-shaped component, like that of the full-length type-1 channel, but this bell-shaped Ca²⁺ dependency arises from a substantially elevated basal Po level. Further, the peak of the bell-shaped component is shifted to lower Ca²⁺ concentrations.

**TABLE I**

Dwell Times of the 2-1-2 and 1-2-1 Chimeras

| [InsP₃] | τ₁₁ | τ₂₁ | τ₁₂ | τ₂₂ |
|---------|-----|-----|-----|-----|
|         | ms  | ms  | ms  | ms  |
| 0       | 1.01 ± 0.04 (0.89) | 16.31 ± 7.24 (0.11) | 2.02 ± 0.09 (0.70) | 50.6 ± 19.51 (0.30) |
| 50 nM   | 1.43 ± 0.21 (0.86) | 21.49 ± 1.51 (0.14) | 3.05 ± 0.48 (0.71) | 57.78 ± 10.65 (0.29) |
| 1 μM    | 1.69 ± 0.50 (0.79) | 22.13 ± 11.79 (0.21) | 1.63 ± 0.56 (0.87) | 19.98 ± 7.75 (0.13) |

1-2-1 Chimera data

| [InsP₃] | τ₁₁ | τ₂₁ | τ₁₂ | τ₂₂ |
|---------|-----|-----|-----|-----|
|         | ms  | ms  | ms  | ms  |
| 0       | 1.49 ± 0.31 (0.62) | 22.55 ± 14.03 (0.38) | 2.99 ± 1.22 (0.70) | 52.25 ± 14.62 (0.30) |
| 50 nM   | 1.33 ± 0.34 (0.64) | 11.45 ± 2.8 (0.36) | 2.74 ± 0.82 (0.43) | 47.95 ± 13.24 (0.57) |
| 1 μM    | 4.70 ± 1.25 (0.60) | 16.14 ± 3.05 (0.40) | 1.19 ± 0.22 (0.47) | 42.29 ± 41.51 (0.55) |

*Values represent means and standard deviations of three to five different channels.
Interdependency of InsP₃ and Ca²⁺ Sensitivity of the 1-2-1 Chimeric Channel

The pore of the 1-2-1 chimera is regulated by both InsP₃ and Ca²⁺. This implies that the InsP₃ and Ca²⁺ binding regions of the protein (i.e., the binding and regulatory/coupling domain, respectively) both interact with the channel domain. It has been suggested that InsP₃ and Ca²⁺ may interact to regulate the InsP₃R pore (Kafstan et al., 1997; Mak et al., 1998). Here, possible interdependency of InsP₃ and Ca²⁺ in regulation of the 1-2-1 chimera pore was evaluated by comparing its Ca²⁺ sensitivity at two InsP₃ concentrations.

In Fig. 8A, sample single channel recordings illustrate the Ca²⁺ sensitivity of the 1-2-1 chimera in the presence of 0.2 μM InsP₃ (cis-chamber). The Po of the 1-2-1 chimera at 40 nM free Ca²⁺ was relatively high (≈0.5). The Po increased at higher Ca²⁺ concentrations (150 nM and 20 μM). The corresponding total amplitude histograms (Fig. 8A, right) also illustrate the Ca²⁺-dependent changes in Po (unit current remained constant). The mean Po of the 1-2-1 chimera (n = 9) in the presence of 0.2 μM InsP₃ is plotted (open circles) as a function of free Ca²⁺ concentrations in Fig. 8B. The Ca²⁺ dependency of this channel in the presence of 1 μM InsP₃ is represented by the thin bell-shaped line (reproduced here from Fig. 7B). To compare the two curves, the Po at 10 nM was subtracted from each. These offset-subtracted curves are superimposed in the inset (Fig. 8B). The general shape of the 1-2-1 chimera’s Ca²⁺ dependency was different at the two InsP₃ concentrations. At one InsP₃ level (1.0 μM), it was bell-shaped (EC₅₀ = 62.8 ± 13.1; IC₅₀ = 159.5 ± 35.9). The Hill coefficients associated with these values (EC₅₀ and IC₅₀ at 1 μM InsP₃) were about three. At the other InsP₃ level (0.2 μM), the channel’s Ca²⁺ dependency was sigmoidal (EC₅₀ = 216.6 ± 43.9; Hill coefficient = 3.2). The EC₅₀ values of Ca²⁺ activation at the two different InsP₃ concentrations were statistically different (P < 0.05). These data suggest that the Ca²⁺ regulation of the 1-2-1 chimera pore is modulated by InsP₃. Thus, the interceding type-2 regulatory/coupling domain (i.e., containing putative Ca²⁺ regulatory sites) appears to exchange Ca²⁺ binding information in both directions (i.e., to the amino-terminal InsP₃-binding domain and the carboxyl-terminal channel domain).

DISCUSSION

Several labs have defined the single-channel function of InsP₃,R channels by reconstituting these channels into planar lipid bilayers (Bezprozvanny et al., 1991; Watras et al., 1991; Ramos-Franco et al., 1998a). There has been particular attention lately on defining the isoform-specific attributes of the InsP₃,R channels. Some studies have focused on the type-3 InsP₃,R channel (Hagar et al., 1998; Mak et al., 2000). Others have focused on the type-1 InsP₃,R channel (Bezprozvanny et al., 1991; Kaznacheyeva et al., 1998). Our group has focused on the type-2 InsP₃,R channel (Perez et al., 1997; Ramos-Franco et al., 1998a, 2000).

All three InsP₃,R channels have similar permeation properties. However, their regulation by Ca²⁺ and InsP₃,
is isoform specific. All three InsP₃R channel have been cloned and are routinely expressed in heterologous systems (Kaznacheyeva et al., 1998; Ramos-Franco et al., 1998b, 2000; Mak et al., 2000). Thus, the stage is set to use molecular genetic manipulations to explore the molecular determinants of InsP₃R channel structure and function.

The InsP₃R Permeation Pathway

The single-channel properties of the two chimeras were similar to each other and to their full-length wild-type counterparts (Ramos-Franco et al., 1998a,b, 2000). Specifically, their conductance (~220 pS in a 220/20 Cs⁺ gradient) and selectivity (reversal potential of ~45 mV in Cs-HEPES salts) were preserved. This indicates that the genetic manipulations required to create these domain-swap chimeras did not interrupt the structural integrity of the permeation pathway. However, the Ca²⁺ and InsP₃ regulation of the pores formed by the chimeric proteins was not identical to that of the wild-type channels. The pores were ligand regulated (in most cases) but the molecular manipulations clearly altered the nature of this regulation. These results are consistent with a previous study involving gross TMR-deletion mutants.

Figure 7. Ca²⁺ sensitivity of the 1-2-1 chimeric channel. (A) Sample single channel recordings from the 1-2-1 chimera are shown at three different Ca²⁺ concentrations in the presence of 1 μM InsP₃. The holding potential was 0 mV and open events are upward deflections. The current carrier was Cs⁺. Corresponding total amplitude histograms (3–5 min of recording) are shown for each Ca²⁺ concentrations. (B) Mean Po of the 1-2-1 chimeric channel is plotted as a function of Ca²⁺ concentration (filled circles). Data points represent means (± SE). Data represent seven different 1-2-1 chimeric channels. The Ca²⁺ dose response curves of the full-length type-1 and -2 InsP₃R channels are also shown (dashed lines; Ramos-Franco et al., 2000).
Ramos-Franco et al., 1999). There a gross deletion of the four amino-terminal TMRs generated mutant proteins that formed channels when reconstituted in bilayers. The permeation pathway of these channels had the well-defined permeation properties of the InsP$_3$R channel but was not ligand regulated. Deletion of the four TMRs effectively disrupted the information flow from the binding and the regulatory/coupling domain to the channel domain. Thus, it appears that the InsP$_3$R pore structure is a robust structure that is relatively resistant to manipulations around it. This is not true for the regulation of the InsP$_3$R pore (i.e., how often it opens and closes). The ligand regulation of the pore is clearly impacted by the presence and identity of the structures surrounding it. The salient point here is that the permeation and ligand regulatory properties of the InsP$_3$R channel appear to be governed independently by different structural determinants of the protein.

Figure 8. (A) Ca$^{2+}$-dependent gating of 1-2-1 chimeric channels at low InsP$_3$ concentrations. Sample single channel recordings from the 1-2-1 chimera are shown at three different Ca$^{2+}$ concentrations in the presence of 0.2 $\mu$M InsP$_3$. The holding potential was 0 mV and open events are upward deflections. The current carrier was Cs$^+$. Corresponding total amplitude histograms (3-5 min of recording) are shown for each Ca$^{2+}$ concentrations. (B) InsP$_3$ sensitivity of 1-2-1 chimeric channel Ca$^{2+}$ dependence. Mean Po of the 1-2-1 chimeric channel is plotted as a function of Ca$^{2+}$ concentration (open circles) in the presence of 0.2 $\mu$M InsP$_3$. Data points represent means ($\pm$ SE). Data represent 3-11 single channels. The Ca$^{2+}$ dose response curve of the 1-2-1 chimeric channel in the presence of 1 $\mu$M InsP$_3$ is shown as the bell-shaped curve. The Po at 1 nM free Ca$^{2+}$ was subtracted from both curve fits and the resulting curves are plotted on an expanded scale in the inset.
**InsP₃ Regulation**

The type-1 and -2 InsP₃R proteins contain InsP₃ binding sites with different InsP₃ affinities (Südhof et al., 1991; Newton et al., 1994; Perez et al., 1997). The InsP₃ binding affinity of the soluble binding domain of the type-2 InsP₃R is ~3 times higher than that of the type-1 receptor (Südhof et al., 1991). A similar relationship is evident in the InsP₃ sensitivity of the native and recombinant InsP₃R channels (Ramos-Franco et al., 1998a,b, 2000). Together, these data imply that the relative InsP₃ affinity of the type-1 and -2 channels is defined by structural determinants that lie within the binding domain of the protein.

Here, the InsP₃ sensitivity of two chimeric InsP₃R channels with different binding domains was defined. The InsP₃ modulation of the chimeric channels was different than that of either of the full-length progenitors. The type-2 binding domain did impart high InsP₃ sensitivity. The type-1 domain did impart lower InsP₃ sensitivity. This confirms that the identity of the binding domain does indeed play a role in establishing InsP₃ affinity. Efficacy of InsP₃ activation of the 2-1-2 chimera was similar to that of the full-length recombinant type-2 channel (Po’s 0.42 vs. 0.40, respectively). Efficacy of InsP₃ activation of the 1-2-1 chimera was about fourfold higher than that of the full-length recombinant type-1 channel (Po’s 0.70 vs. 0.17, respectively). Interestingly, there was also substantial activity of the 2-1-2 chimera (Po ~0.2) at very low InsP₃ concentrations (or when no InsP₃ was added). Such sustained channel activity is not observed in the full-length progenitor channels. Thus, the transduction of InsP₃ binding to channel opening is altered in these chimeric channels. These alterations could be due to the identity of the introduced regulatory/coupling domains or abnormal interdomain crosstalk. In either case, the implication is that structural determinants that define InsP₃ binding affinity in the binding domain are not the only factors that define InsP₃ regulation of the channel.

**Ca²⁺ Regulation**

Several Ca²⁺ binding sites and residues key to the calcium sensor (E2100 of the rat type-1 isoform) that regulate the InsP₃R channel have been shown to reside in the regulatory/coupling domain of the receptor (Mignery and Südhof, 1990; Sienaert et al., 1996, 1997; Miyakawa et al., 2001; Nadif-Kasari et al., 2002). Additionally, all three isoforms contain Ca²⁺ binding sites in the ligand binding as well as channel domains (Sienaert et al., 1996, 1997, 2002; Sipma et al., 1999). The details of calcium’s role in activation and inhibition are beginning to emerge, yet much more information is required to elucidate which critical residues are involved in the complex modulation of the receptor by this ion.

The 2-1-2 chimera has the type-1 regulatory/coupling domain and the 1-2-1 chimera the type-2 regulatory/coupling domain. The activity of the 2-1-2 chimera was not Ca²⁺ sensitive. One possible interpretation is that the 2-1-2 chimera does not bind Ca²⁺. Considerable care was taken to craft smooth junctional transitions between the different receptor domains and these transitions are relatively distant from the putative Ca²⁺ binding sites. However, a failure to bind Ca²⁺ may involve coordination of different regions of the protein and this coordination may not be available in the 2-1-2 chimera structure. Another possibility, and perhaps more likely, is a process that transduces Ca²⁺ binding into channel regulation is disrupted in the 2-1-2 chimera. In this case, the data would suggest that the type-1 regulatory/coupling domain does not communicate well with the type-2 channel (and/or binding) domain. This result is not entirely surprising in that the type-1 and -2 coupling/regulatory domains are the least conserved (having only ~60% identity). These domains contain large interspersed stretches of heterologous sequence. This provides multiple putative modulatory binding sites that may contribute to the observed isoform-specific Ca²⁺ signaling of these channels. The implication is that the type-1 and -2 coupling domains are not interchangeable.

The function of the 1-2-1 chimera was Ca²⁺ dependent. Its Ca²⁺ dependence was bell shaped (like that of the full-length type-1), but this bell shape was substantially offset from 0 Po. This Po offset may be related to the unusually high InsP₃ efficacy of this chimera. Its bell-shaped Ca²⁺ dependency was centered at lower Ca²⁺ concentrations compared with that of the full-length type-1 receptor. Activation at such low Ca²⁺ levels is typical of the full-length type-2. Thus, chimera has features common to its progenitors. It is interesting that a channel containing the type-2 regulatory/coupling domain has a bell-shaped component of Ca²⁺ sensitivity reminiscent of that generally associated with the full-length type-1 channel. This could imply that some of the structural determinants of the bell-shaped Ca²⁺ modulation may lie outside the regulatory/coupling domain. Alternatively, it could mean that structural determinants outside the regulatory/coupling domains alter the bell-shaped Ca²⁺ modulation in the native full-length type-2 context. Numerous Ca²⁺ binding motifs have been detected in other regions of these receptors, including their ligand binding and channel domains. Further, it has been suggested that InsP₃R Ca²⁺ inactivation may be modulated (or “tuned”) by InsP₃ binding (Hagar et al., 1998; Mak et al., 1998). In any event, these data support the notion that InsP₃R ligand (InsP₃ and Ca²⁺) regulation is a multideterminant and isoform-specific process.
Potential InsP₃ and Ca²⁺ Interaction

There is a consensus that the function of the single InsP₃R channels is isoform specific (Hagar et al., 1998; Mak et al., 1998; Ramos-Franco et al., 1998a, 2000). Studies of type-1 and -3 InsP₃R channel regulation indicate that InsP₃ binding regulates Ca²⁺ inactivation of the channel (Hagar et al., 1998; Mak et al., 1998). The underlying mechanism of the Ca²⁺-InsP₃ interaction is debated. Kaftan et al. (1997) has proposed that two InsP₃ binding sites regulate InsP₃R channels. A high affinity InsP₃ binding site activates the channel while a lower affinity InsP₃ binding site modulates the Ca²⁺ inactivation of the channel. Occupancy of the low affinity site reduces the degree of Ca²⁺ inactivation. In contrast, Mak et al. (1998) has proposed that a single high affinity InsP₃ binding site “tunes” the Ca²⁺ sensitivity of the InsP₃R channel. Increased InsP₃ occupancy of this site reduces the affinity of the Ca²⁺ inactivation process (i.e., inactivation occurs at higher Ca²⁺ levels).

There are certain considerations that must be taken into account if our results are to be interpreted in the type-1/type-3 regulatory context outlined above. First, the InsP₃ binding affinities of the type-2 and -3 isoforms are very different relative to the type-1 channel (type-2 < type-1 << type-3; Newton et al., 1994). Second, all three isoforms share at least one potential Ca²⁺ regulatory site (E2100; Miyakawa et al., 2001) but likely have others. Third, it has been suggested that calmodulin binding may be critical in InsP₃ and Ca²⁺ regulatory interaction (Michikawa et al., 1999) and the calmodulin–InsP₃R interaction may be isoform-specific (for review see Nadif-Kasari et al., 2002). For example, the type-3 does not contain the calmodulin-binding motif as the type-1 or -2 receptors. Recently, the complex role of calmodulin in the modulation of the type-1 receptor has been examined by Nosyreva et al. (2002), in which a mutant (W1577A) abolished Ca²⁺-dependent CaM binding of the rat type-1 receptor. Despite the absence of CaM binding, the channel activity showed a similar biphasic response to Ca²⁺ as the wild-type receptor and similar sensitivity to exogenously applied CaM. At the present, the role of CaM in the regulation of the receptor family is a confounding and elusive property of the receptor that remains to be elucidated. Fourth, the same InsP₃ and Ca²⁺ regulatory interaction scheme may simply not apply to all three InsP₃R channel isoforms. It may be misleading to interpret the ligand regulation of the three channels in a single conceptual framework.

The possibility that InsP₃ alters the Ca²⁺ dependence of the 1-2-1 chimera channel was examined here. In the presence of 1 μM InsP₃, the Ca²⁺ dependence of the 1-2-1 chimera had a bell-shaped component. In the presence of a lower InsP₃ concentration (0.2 μM), the Ca²⁺ dependence of the 1-2-1 chimera was sigmoidal (i.e., lost its bell-shaped component). This data is difficult to explain in the context of either the Kaftan et al. (1997) or Mak et al. (1998) Ca²⁺/InsP₃ regulatory schemes. For example, the apparent Ca²⁺ inactivation was lost at lower InsP₃ concentrations in the chimeric channels. Also, the EC₅₀ of Ca²⁺ activation was InsP₃ dependent (i.e., it was more sensitive to Ca²⁺ activation at high InsP₃ concentrations). This is not observed in the previously published Ca²⁺/InsP₃ regulatory results on full-length channels (Kaftan et al., 1997; Hagar et al., 1998; Mak et al., 1998, 2001). In any event, chimeric channel behavior was defined over a wide range of [Ca²⁺]s (10⁻⁶ to 2 × 10⁻³ M) at two [InsP₃]s (1 and 0.2 μM; see Fig. 8 B). The Ca²⁺ sensitivity of the 1-2-1 chimeras was different at the two [InsP₃]s tested (i.e., its Ca²⁺ sensitivity was InsP₃ dependent). This suggests that this channel does not behave as a solely Ca²⁺-gated or InsP₃-gated channel. Thus, the chimera results clearly show that there is an InsP₃/Ca²⁺ regulatory interaction. This interaction has three components. One involves InsP₃ modulation of the EC₅₀ of Ca²⁺ activation. A second involves InsP₃ modulation of high Ca²⁺ inactivation. The third involves InsP₃ modulation of the Po offset at low Ca²⁺ concentrations.

These initial chimera studies demonstrate that it is possible to use a chimera strategy to define InsP₃R structure-function. The large domain-swap chimeras created here formed functional channels, but these structural manipulations introduced unique ligand regulatory attributes. Application of a more conservative chimera strategy that exchanges certain nonconserved motifs and/or regions of heterologous sequence, in tandem with these bulk domain exchanges, will likely lead to important information regarding isoform-specific structure and function.

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