Functional properties of a pore mutant in the *Drosophila melanogaster* inositol 1,4,5-trisphosphate receptor

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**Abstract** The inositol (1,4,5)-trisphosphate receptor (InsP\(_3\)R) is an intracellular calcium release channel that plays a crucial role in cell signaling. In *Drosophila melanogaster*, a single InsP\(_3\)R gene (itpr) encodes a protein (DmInsP\(_3\)R) that is ~60% conserved with mammalian InsP\(_3\)Rs. The functional properties of wild-type (WT) and mutant DmInsP\(_3\)Rs have recently been described [Srikanth et al., Biophys. J. 86 (2004) 3634–3646]. Here, we use the planar lipid bilayer reconstitution technique to describe single channel properties of a ka901 point mutant (G2630S) in the pore-forming region of DmInsP\(_3\)R. We find that homomeric ka901 channels are not functional, but the heteromeric WT:ka901 mutant channels display increased conductance, longer channel open time and altered ion selectivity properties when compared to WT DmInsP\(_3\)R. Obtained results are consistent with the gain of function phenotype observed in ka901/+ mutant flies.

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1. **Introduction**

The inositol (1,4,5)-trisphosphate receptor (InsP\(_3\)R) is an intracellular calcium (Ca\(^{2+}\)) release channel that plays a critical role in Ca\(^{2+}\) signaling [1]. The mammalian genome encodes three different isoforms of the InsP\(_3\)R-type I (InsP\(_3\)R1), type II (InsP\(_3\)R2) and type III (InsP\(_3\)R3) [2], while a single gene (itpr) codes for DmInsP\(_3\)R in the *Drosophila* genome [3,4]. All 3 mammalian isoforms share a common domain structure and 60–70% sequence identity [2]. InsP\(_3\)Rs are subjected to multiple levels of regulation [1,5,6]. The structural determinants responsible for InsP\(_3\)R1 conductance and gating properties [7,8] and modulation by Ca\(^{2+}\) [9–11], ATP [12] and phosphorylation [13] have been uncovered in recent functional experiments with InsP\(_3\)R1 wild-type and mutant channels.

The *Drosophila* InsP\(_3\)R protein (DmInsP\(_3\)R) shares the same domain structure and ~60% sequence identity with mammalian InsP\(_3\)R isoforms (reviewed in [14]), with the highest level of conservation in the amino-terminal ligand binding and the carboxy-terminal channel regions. Recently, single channel properties of the *Drosophila* wild-type isoforms and two single point mutants (wc703 and ug3) identified in a genetic screen have been described [15,16]. An additional mutant identified in genetic screen (ka901) contains a single point mutation in a putative pore-forming region of the DmInsP\(_3\)R. Expression of full-length ka901 mutant in S9F cells did not result in InsP\(_3\)R-gated channels in bilayers, indicating that ka901 homomeric channels are not functional [15]. To analyze ka901 functional properties, here we investigated functional properties of heteromeric WT:ka901 channels formed by co-infection of wild-type (WT) DmInsP\(_3\)R and ka901 viruses into S9F cells. The observed results provide a novel information about structural determinants of DmInsP\(_3\)R pore and explanation to the gain-of-function phenotype observed in ka901/+ mutant flies.

2. **Materials and methods**

2.1. **Expression of DmInsP\(_3\)R in S9F cells**

Generation of DmInsP\(_3\)R wild-type and ka901 recombinant baculovirus has been previously described [15]. Equal amounts of DmInsP\(_3\)R and ka901 baculovirus were used for co-infection in *Spodoptera frugiperda* (S9F) cells as previously described [15]. Expression of DmInsP\(_3\)R and ka901 isoforms in S9F cells was confirmed by Western blotting with the affinity purified anti-DmInsP\(_3\)R rabbit polyclonal antibody (IB-9075) that was previously described [15].

2.2. **Single channel recordings and analysis of DmInsP\(_3\)R activity**

Recombinant DmInsP\(_3\)R channels expressed in S9F cells were incorporated into the bilayer by microsomal vesicle fusion as previously described [10,12,15]. The cis chamber contained 250 mM HEPES-Tris, pH 7.35, and the trans chamber, which was held at virtual ground, contained 55 mM Ba(OH)\(_2\) dissolved in 250 mM HEPES, pH 7.35. Single-channel conductance values of heteromeric ka901 channels were determined from the slope of a linear fit to unitary current amplitude versus transmembrane voltage data in the range between +10 and ~30 mV.

3. **Results**

The domain structure of the InsP\(_3\)R [2] is conserved between DmInsP\(_3\)R and mammalian InsP\(_3\)R isoforms. The ka901 mutation (G2630S) described in earlier studies [15,16] resides...
in a carboxy-terminal channel-forming domain of DmInsP$_3$R. G2630 residue mutated in ka901 is located within a putative pore-forming region and highly conserved within InsP$_3$R and RyanR gene families and in a region homologous to the selectivity filter of K$^+$ channel family (Fig. 1). Flies with one copy of ka901 mutation and a deficiency for the InsP$_3$R (itpr$^{50B0}$/itpr$^{50B0}$) die as second instar larvae [16], similar to the homozygous deficiency (itpr$^{50B0}$/itpr$^{50B0}$) [17], indicating that in homozygous state ka901 mutation behaves similar to the loss of function mutation. Consistent with this conclusion, ka901 mutants expressed in Sf9 cells by baculovirus infection failed to form functional channels in bilayers [15]. In contrast to these results, Ca$^{2+}$ release experiments with microsomal vesicles extracted from adult heads of ka901/+ heterozygotes showed over twofold higher calcium release as compared to wild-type [15], indicative of a gain-of-function phenotype.

To explain the discrepancy between ka901/ka901 and ka901/+ phenotypes, we reasoned that ka901 forms functional channels only when assembled with the wild-type (WT) DmInsP$_3$R subunits, but not in the homomeric state. To test this prediction and to examine the functional properties of WT: ka901 channels, we co-expressed ka901 and DmInsP$_3$R subunits in Sf9 cells at 1:1 ratio by baculovirus co-infection. The expression levels of WT:ka901 heteromers with microsomal vesicles extracted from adult heads of ka901/+ heterozygotes showed similar to wild-type DmInsP$_3$R (Fig. 2). In contrast to experiments with ka901 microsomes [15], InsP$_3$-gated channels were observed routinely (in 13 out of 18 experiments) in bilayer experiments with WT:ka901 microsomes. In three out of 18 experiments, the properties of observed channels were similar to the wild-type DmInsP$_3$R data not shown), and we reasoned that these channels result from WT homotetramers. However, in 10 out of 18 experiments the conductances and gating properties of the observed channels (Fig. 3B) were clearly distinct from the wild-type DmInsP$_3$R.

We reasoned that these channels are formed by WT:ka901 heteromers. Although exact stoichiometry of these heteromers in the bilayers is unknown, the frequency of these channels’ occurrence indicates that the presence of a single ka901 subunit in a tetramer is sufficient to change DmInsP$_3$R gating and conductance properties (see Section 4). Only heteromeric WT: ka901 channels were considered in the following analysis.

The mean current amplitude of WT:ka901 channels at 0 mV holding potential is equal to 1.87 ± 0.03 pA ($n = 4$) with a mean open dwell time of 6.2 ± 0.5 ms (Fig. 3B). In the same recording conditions, WT DmInsP$_3$R display unitary current amplitude of 1.8 pA and mean open dwell time of 4.3 ms [15]. Thus, unitary current is increased by 4% and mean open time is increased by 44% in the presence of ka901 subunit. In contrast to DmInsP$_3$R, WT:ka901 channels displayed a sub-conductance state with amplitude of 70% of the open state (Fig. 3B) in about 25% of the channel openings. This sub-conductance state was found to be a property of a single hybrid channel, as transitions between open and sub-conductance states were observed in membranes with only a single functional channel. The presence of a single functional tetramer in the bilayer was deduced by the amplitude of channel opening observed over 10–30 min. In cases where more than one functional channel is present in the bilayer, some openings have amplitudes which are greater than that of the single channel opening.

Single-channel conductances of 86.0 ± 0.2 pS and 39.0 ± 0.1 pS were estimated for the open and sub conductance states, respectively (Fig. 4), as compared with the 70.0 ± 0.1 pS conductance of the WT DmInsP$_3$R (Fig. 4) [15]. Extrapolation of the current–voltage curve to the intercept on the voltage axis indicated a −18 mV shift of approximated reversal potential for the $I/V$ curve of the full-open state of the WT:ka901.
channels when compared to wild-type channels (Fig. 4). Since 55 mM Ba\(^{2+}\) in the trans chamber is the sole current carrier in the system, this change in reversal potential in the mutant indicates an alteration in ion selectivity of the heteromeric WT:ka901 channel.

The observed effects on single-channel conductance, selectivity and channel gating are consistent with the position of the ka901 mutation (G2630S) in the pore region of DmInsP\(_3\)R (Fig. 1). In additional experiments, we determined the Ca\(^{2+}\)-sensitivity of WT:ka901 channels. Firstly, the presence of only heteromeric channels in the bilayer was confirmed by measuring current–voltage relationship as described above (Fig. 4). Subsequently, calcium dependence experiments were carried out with the same bilayer. We observed that the bell-shaped Ca\(^{2+}\)-dependence of the WT:ka901 channels was similar to that of the wild-type DmInsP\(_3\)R [15] (data not shown), suggesting that DmInsP\(_3\)R modulation by Ca\(^{2+}\) is not influenced by the ka901 mutation.

4. Discussion

The absence of functional channels from ka901 homomers in bilayer experiments [15] agrees with the genetic finding that the allelic strength of ka901 is equivalent to that of the null allele *itp*\(^{90b0}\) [16]. However, when co-expressed with the wild-type DmInsP\(_3\)R, ka901 mutant behaves as a “gain of function” mutant in the vesicle flux assay [15]. As shown here, these results can be explained by increasing single-channel conductance (23% increase) and open dwell time (44% increase) of heteromeric WT:ka901 DmInsP\(_3\)R channels. These effects on conductance and gating properties of DmInsP\(_3\)R are consistent with the position of ka901 mutation (G2630S) within the predicted pore-forming region of DmInsP\(_3\)R (Fig. 1). The pore-forming regions of InsP\(_3\)R and RyanR families are highly conserved and contain a putative pore-forming motif GXRXGGGI/VGD (the G2630 residue of DmInsP\(_3\)R mutated in ka901 is underlined). The G4826 residue in RyanR2 is homologous to the G2630 residue mutated in ka901. Mutation studies of this residue revealed conductance properties that support our observations in ka901 [18,19]. The G4826C mutant does not form functional homomeric channels in bilayer experiments [19], similar to what we observe with ka901 homomeric channels. Co-transfection studies of G4826C with wild-type RyanR2 showed the presence of a single type of hybrid channel and not four types of hybrid channels, as seen in the case of the G4824A mutant and RyanR2 co-transfections [18,19]. Interestingly, similar to our observation of increase in conductance of WT:ka901 hybrid channels, the conductance of the full open state of WT:G4826C hybrid channels is also increased by ~25% when compared with RyanR2. The exact subunit composition of heteromeric channels is not known for either WT:ka901 DmInsP\(_3\)R (Figs. 3 and 4) or WT:G4826 RyanR2 [19]. The observation of a single sub-conductance state and a single full-open state indirectly suggests that some hybrid channels may not be functional. Thus, properties of WT:ka901 DmInsP\(_3\)R channels mirror those of WT:G4826C RyanR2 channels, presumably due to the similarity between the cysteine (CH2-SH) and serine (CH2-OH) side chains found in G4826C and ka901 mutants, respectively. Most likely, the presence of a larger side-chain in
these mutants in narrow selectivity filters of DmInsP3R or RyanR2 channels puts a limit on a number of mutant subunits in a functional channel. The presence of a single sub-conductance state indicates the presence of only one functional heteromer. Multiple sub-conductance states were observed in case of the G4824A mutant of RyanR2, which indicated the stoichiometry of wild-type and mutant channels in the tetramer [18]. Similarly, we reasoned that the presence of different sub-conductance states would indicate the presence of multiple heteromers in different stoichiometries. Instead, we observed only a single sub-conductance state. Since a tetramer consisting of four ka901 subunits does not form functional channel [15], we reason that the most likely functional tetramer would comprise 3WT and 1 ka901 subunits. An alternate stoichiometry is also possible.

Increase in conductance of the single functional state in WT:ka901 DmInsP3R (Fig. 4) or WT;G4826 RyanR2 [19] may arise from an interaction of the side chain (either -OH or -SH) with the back bone. The idea of an interaction between the side chain and the backbone is supported by another study, of the I4898T mutation of RyanR1, which has been linked to central core disease [20]. The I4898 position in RyanR1 corresponds to the isoleucine residue in the GGGI/VGD putative pore-forming motif of RyanR and InsP3R, and thus lies adjacent to the G residue mutated in ka901. It has been shown that RyanR channels are leaky in cells co-transfected with WT RyanR1 and I4898T mutants [20]. Similar to our findings with ka901, homomeric tetramers of the I4898T mutant of RyanR1 are non-functional as judged by Ca2+ photometry experiments [20]. Consistent with our observations of ka901 channels, it seems likely that the leaky channel arises from an interaction of the –OH (in the threonine side chain) with the backbone, while in the mutant homotetramer the pore is blocked. Overall, these results highlight the importance of the putative pore-forming motif for ion conductance by InsP3R and RyanR channels. These results also agree with previous analysis of InsP3R1 [7,8] pore forming mutants and with the known structure of the potassium channel selectivity filter, which is composed of a similar sequence [21]. In summary, our results demonstrate the conserved function of the G2630 residue in InsP3R and RyanR Ca2+ channel families.

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