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Ion channel activity of the CSFV p7 viroporin in surrogates of the ER lipid bilayer

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1. Introduction

Modulation of the cellular ion balance in order to take over the cellular machinery seems to be a common feature for viruses. Several viruses encode at least one protein displaying ion channel (IC) activity. These proteins are known as viroporins, the term being viruses encode at least one protein displaying IC activity. The cellular machinery seems to be a common feature for viruses. Several

Viroporins comprise a family of non-structural proteins that play significant and diverse roles during the replication cycle of many animal viruses. Consequently, they have become promising targets for inhibitory drug and vaccine development. Structure-function traits common to all members of the family are their small size (ca. 60–120 aa), high hydrophobicity, and the presence of helical domains that transverse the membrane and assemble into oligomeric-permeating structures therein. The possibility that viroporins show in particular conditions any kind of specificity in the transport of ions and small solutes remains a point of contention in the field. Here we have approached this issue using the Classical Swine Fever Virus (CSFV) protein p7 viroporin as a model. We have previously reported that CSFV-p7 induces release of ANTS (MW: 427.33) from lipid vesicles that emulate the Endoplasmic Reticulum (ER) membrane, and that this process is dependent on pH, modulated by the lipid composition, and recreated by a C-terminal transmembrane helix. Here we have assayed CSFV-p7 for its capacity to form ion-conducting channels in ER-like planar lipid membranes, and established whether this activity is subject to regulation by the same factors. The analysis of electrophysiological recordings in ER membrane surrogates suggests that CSFV-p7 forms pores wide enough to allow ANTS release. Moreover, we were able to discriminate between two pore structures with slightly different sizes and opposite ion selectivities. The fact that the relative abundances of each pore type depend crucially on membrane composition strengthens the view that the physicochemical properties of the lipid bilayers present in the cell endomembrane system modulate viroporin activity.

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of the lipid bilayer the viroporin may have the potential either to establish nanometer-sized ion channels, or alternatively, participate in additional membrane permeabilization mechanisms allowing the transport of larger solutes exceeding the nanometer scale [35,36].

Interestingly, it has been observed that lipid charge influences the IC activity of some viroporins like the E protein of SARS-CoV [32], while vesicle permeabilization assays suggest that PV 2B pore-forming activity of some viroporins like the E protein of SARS-CoV [32], while

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2.6. Statistical processing

For comparison between vesicle leakage and channel formation activity, two kinds of assays were performed. For vesicle leakage results, at least four kinetic assays were performed and shown data correspond to arithmetic mean and standard error. In case of channel formation activity, 30 recordings comprising 10 traces with a duration of 20 s were analyzed. Each trace with at least one 2 pA event was considered as positive. The histograms of the current jump amplitude were made from at least 60 recordings and more than 550 events were analyzed for each histogram. The data were normalized to 1, using the number of total events in each case. The histograms were fitted to one or two Gaussian peaks, depending on the value of the adjusted R-squared. Each SD value is the square root variance of the corresponding Gaussian distribution. The histograms of permeability ratio contain data from 64 recordings.

3. Results and discussion

3.1. Ion channel activity of CSFV-p7

The CSFV-p7 protein possesses two hydrophobic, membrane-spanning regions (TM1 and TM2) separated by a short basic loop, an arrangement characteristic of class II viroporins (Fig. 1). Each hydrophobic domain is additionally intervened by a short loop with polar character, a feature also shared by integral membrane proteins that assemble into helical oligomers [22]. CSFV-p7 induces ANTS release from liposomes, a process that requires acid pH and evolves more efficiently in ER-like membranes [38,42]. The pore-forming activity and its regulation by pH and lipid composition can be recreated by the p7C peptide, representing the C-terminal domain including the TM1–TM2 connecting basic loop (Fig. 1). In order to improve the topological localization of the viroporin activity, p7-4, a TM2-based peptide lacking the basic loop, was designed. Interestingly, p7-4 also induces ANTS release from liposomes, but the process does not require acid pH and it is not inhibited by classic viroporin inhibitors [42]. These previous results could indicate that the TM1–TM2 connecting loop is essential for the CSFV-p7 activity regulation. Up to date, CSFV-p7 full protein and its constituent domains have been shown to participate in the permeabilization of lipid vesicles [38], but electrophysiological measurements investigating their pore-formation mechanisms at the single channel level are missing.

To this end, CSFV p7 viroporin was first assayed in planar lipid mixtures that mimicked the composition of ER membranes (Fig. 2). Although the ER is the main site of cholesterol and structural phospholipid synthesis, these lipids are rapidly transported to other organelles and, in fact, ER displays only low concentrations of sterols and complex sphingolipids [43]. Accordingly, we prepared planar bilayers based on the main constituent phospholipids of the ER membrane, namely, zwitterionic PC and PE plus the anionic PI mixed in a roughly 5:3:2 molar ratio [43].

In experiments performed in 150 mM KCl and pH 5.0 the complete protein (p7Full) exhibited a high IC activity with a variety of current levels and lifetimes (Fig. 2). In those experiments different kinds of events appear. On the one hand, traces showing well-defined “opening” and “closing” rapid events, and on the other hand, larger current levels with longer lifetimes without almost any small flickering (Fig. 2A). This effect could be due to the simultaneous opening of several channels or could correspond to the formation of different channel structures. In order to discriminate between single and multiple channel insertions, the analysis of all recordings allowed resolving the stepwise changes (ΔI) in each trace. The current vs. time traces were analyzed with the software Clampfit 10.1, so that average current values for each level were obtained once the corresponding zero-voltage baseline was subtracted. Histograms of the current jump amplitudes of the recorded
peptides were synthesized and analyzed in planar bilayers resembling abilization [38,42]. Thus, to map IC activity, CSFV p7 sequence-based dependencies on sequences different to those sustaining vesicle permeability are necessary. For instance, similar results were obtained in the absence of cholesterol using a PC:PE (7:3 molar ratio) planar bilayers (data not shown). In contrast, the short interhelical loop presents a regulatory function and determines the sensitivity to pH changes. Previous work demonstrated that the lack of anionic phospholipids in the lipid composition, abated p7 permeabilizing activity observed in planar membranes is consistent with a large leakage value. Furthermore, when the pH is increased to 7.4, the remarkable decrease in recordings displaying ion channel activity is in agreement with the almost inextensible leakage found in LUVs. Of note, single channel currents of 6 pA or larger were occasionally found in the recordings at neutral pH (less than 25% of the total events, see right panels).

Interestingly, the pH is not the only critical parameter controlling ion channel activity but the membrane composition is also crucial. Previous work demonstrated that the lack of anionic phospholipids in the lipid composition, abated p7 permeabilizing activity measured in vesicles [38,42]. Control experiments performed in a 3-lipid PC:PE:Chol (7:3:10 molar ratio) mixture at pH 5.0 showed only tiny currents, which is compatible with the small leakage found in LUVs of similar composition. By selecting this control mixture, it was not our intention to analyze the effects of cholesterol on channel activity (this would require thorough analysis, which was beyond the scope of the present work), but rather to test the absence of the anionic phospholipid PI. In fact, similar results were obtained in the absence of cholesterol using PC:PE (7:3 molar ratio) planar bilayers (data not shown).

3.2. Sequences involved in CSFV-p7 ion channel activity

The CSFV p7 IC activity observed in the previous experiments might depend on sequences different to those sustaining vesicle permeabilization [38,42]. Thus, to map IC activity, CSFV p7 sequence-based peptides were synthesized and analyzed in planar bilayers resembling ER composition. As displayed in the diagram of Fig. 1, p7N and p7C sequences spanned the potential N- and C-terminal TM1 and TM2 regions of CSFV p7, respectively, and overlapped along the conserved polar loop sequence [38]. In the case of p7-4, the sequence was selected to span the hydrophobic C-terminal TM2 of CSFV p7. The resulting sequence was long enough to transverse the membrane, although interrupted by the turn-promoting 49MTNPPVK55 polar sequence. The pore-forming activities of these peptides were assessed in artificial lipid bilayers, as shown in Fig. 4A. Both p7C and p7-4 peptides exhibited substantial IC activity, in agreement with the high values of leakage induced by these peptides in LUVs. In contrast, p7N showed almost null currents both in electrophysiological recordings and in LUVs. These data confirm a similar dependence of vesicle permeabilization and IC activities on the C-terminal helix of CSFV-p7.

However, the histograms of the current jump amplitudes of the recorded traces denoted certain differences between the IC activities displayed by p7C and p7-4 (Fig. 4B). Whereas p7C recreated with remarkable accuracy p7full behavior (i.e., a prevalent single channel current of ca. 6 pA and a shoulder at 10–12 pA), p7-4 favored lower intensity currents of ca. 2–3 pA and no evidence for a bias towards a larger structure could be found in the histogram. Thus, the addition of the 33MRDEPI38 turn sequence seems to be required for precisely recapitulating p7full IC activity by the TM2-based peptides. This observation was further supported by the next set of experiments.

Next we tested the dependence of p7C and p7-4 IC activities on solution pH and membrane composition, as previously done in the case of the full protein (Fig. 3). Interestingly, each peptide’s IC activity seems to display a particular behavior, which was mirrored by their effects on vesicle leakage. On the one side, p7C performs like the full CSFV p7 protein both in LUVs and in electrophysiological recordings (Fig. 5A). On the other side p7-4 shows a distinctive feature in ER membranes. Increasing pH does not inhibit the ability of p7-4 to induce membrane permeabilization (Fig. 5B). This seems a particular characteristic of ER membranes, because PI lacking control membranes show in both peptide samples very low currents at pH 5. The ANTS leakage induced by p7-4 also shows a significant reduction under these conditions as compared to PI-containing samples. Although the reduction in leakage is not so marked, once again, the correlation between planar electrophysiology and LUVs is maintained supporting our finding that in the absence of the 33MRDEPI38 turn, pores could be efficiently formed, but their activity was not pH dependent. Thus, the short interhelical loop presents a regulatory function and determines the sensibility to pH changes.

Overall, the electrophysiological recordings of peptides inserted in ER-like bilayers are consistent with the formation of a CSFV p7 channel, in which the C-terminal hydrophobic region is a pore lining transmembrane helix and the basic loop regulates the ion permeation process, whereas the N-terminal hydrophobic region functional role remains unknown. In this manner, experiments in planar membranes correlate with previously obtained vesicle leakage results and support the hypothesis.
that the same p7 structure could be responsible for both, ion conductance in planar bilayers and ANTS release from vesicles.

### 3.3. Effects of lipid composition on CSFV-p7 ion channel activity

Results in Figs. 3 and 5 demonstrate that lipid composition has critical effects on the permeabilization induced by CSFV p7 protein and its derived peptides. In the ER-like ternary mixture two conspicuous components, PE and PI, may impart a negative curvature or increase the negative surface charge density of the membrane monolayers, respectively.

Thus, to test intrinsic curvature and surface charge effects on p7full IC activity, we probed a variety of lipid compositions that differed slightly from the canonical ratio 5:3:2 (PC:PE:PI) of ER membranes. Fig. 6 compiles the histograms of current jumps recorded for lipid bilayers made by first keeping constant the amount of charged lipids (PI fixed) and changing the intrinsic curvature (decreasing PC:PE ratio) and then the other way round (i.e., PE fixed and decreasing PC:PI ratio).

**Fig. 4.** Pore-forming activity of p7N, p7C and p7-4 peptides in ER-like planar bilayers and vesicles at pH 5.0. A) Left: Comparison of the ability to form channels in planar bilayers (light gray) with leakage induced in LUVs by the addition of p7 peptides to a peptide-to-lipid mole ratio of 1:250 (dark gray). Right: Representative traces of IC activity. B) Histograms of the current jump amplitude measured for the active p7C and p7-4 peptides. The histograms have been fitted to two Gaussian peaks (left panel) and single Gaussian peak (right panel), respectively.

**Fig. 5.** Comparison between vesicle leakage and channel formation induced by p7C and p7-4 peptides. Effect of pH and lipid composition on IC activity and LUV permeabilization induced by p7C (A) or p7-4 (B). Conditions otherwise as in the previous Fig. 4.
As previously discussed (Fig. 2B), the main peak at 6 pA and the shoulder at ca. 10 pA observed in the 5:3:2 ER-like composition might denote the prevalence of a single channel, with slight contribution of simultaneous double insertions. As could be expected,[46,47] the effect of the lipid charge in the channel function is remarkable: a decrease in the PI content (6:3:1 and 5:4:1 ratios) enhances a defined second peak in the histogram centered at 15 pA, which comprises current values that triplicate those of the main peak centered at 5 pA. In contrast, the excess of PI (3:3:4) yields a huge dispersion with no clearly defined maximum. On the other hand, altering the intrinsic curvature of the monolayer by including an excess of PE (3:6:1) leads to a single well-defined maximum at 5 pA. This is probably related to the fact that lipids with intrinsic negative curvature, like PE, restrict the formation of protein–lipid combined structures because of the energy penalty involved in the bending of the bilayer leaflet [48]. Notably, the intermediate composition (5:4:1) containing low PI and almost equimolar PC:PE ratio leads to the best definition of the 5 and 15 peaks in the histogram.

The overall understanding of Fig. 6 becomes extremely complex due to the combined action of charge and curvature effects. The only straightforward conclusion that can be drawn is that slight changes in any of these factors yield dramatic changes in the current histograms. In view of that, we see no evident reason to claim that there is a predominant channel structure that depending on the membrane conditions appears only in the form of single channels (5:3:2 or 3:6:1 mixtures) or also in the form of simultaneous multiple insertions (6:3:1 or 5:4:1). Quite the contrary, it seems more likely to suggest that at least two different pore structures can be formed, being the relative abundance and characteristics of each one crucially modulated by the lipid composition in the membrane.

### 3.4. Identification of two distinct CSFV-p7 ion channel structures

The global analysis of Fig. 6 provides interesting clues but it does not strictly discriminate whether the current jump histograms showing several peaks reveal different pore conformations or alternatively, single and simultaneous multiple insertions of the same pore type. To obtain additional insights on this issue, we analyzed in detail opening and closure events in current traces focusing on the membrane composition of 5:4:1 (PC:PE:PI) that displays the best definition of two different peaks in the corresponding histogram. In Fig. 7 we show representative traces of the two different current states that could be tentatively associated to two distinct oligomerization states or pore structures. In both traces, well-defined “opening” and “closing” current jumps become apparent.

Selectivity experiments carried out in the same lipid mixture 5:4:1 (PC:PE:PI) further supported the existence of two distinct ion-conducting p7 structures (Fig. 8). In the presence of a concentration gradient between both sides of the membrane, there is a net flux of ions through membrane pores and hence an electric current is measured. The applied voltage that is needed to make zero the electric current (the so-called reversal potential, $E_{rev}$) reveals the preferential passage of either positive or negative ions. In the experimental protocol used here (see the Materials and methods section), a positive applied potential means that the pore is selective to cations and a negative one is associated to an anionic selectivity. We performed reversal potential experiments at pH 5.0 under a concentration ratio of 5 ($C_{cis} = 150$ mM KCl, $C_{trans} = 750$ mM KCl) or 10 ($C_{cis} = 150$ mM KCl, $C_{trans} = 1.5$ M KCl). Fig. 8 shows the permeability ratio ($P_+/P_-$) distribution from performed reversal potential experiments. $P_+/P_-$ > 1 values are tied to a cationic selectivity whereas lower than 1 values indicate a preference for anions. We see two different groups of values, one connected to a very weak anionic selectivity and other to a cationic one. Note that a cluster of identical small structures displaying a higher

![Fig. 6. Histograms of the current jump amplitude measured for the CSFV-p7 protein in different lipid mixtures PC:PE:PI. 5:3:2 (A), 3:3:4 (B), 6:3:1 (C), 5:4:1 (D), 3:6:1 (E). The histograms have been fitted to two Gaussian peaks in all panels but B, where no clear peak is visible.](image)

![Fig. 7. CSFV-p7 channel current recordings in 150 mM KCl at pH 5 on the membrane composition of 5:4:1 (PC:PE:PI). Traces of two different current state jumps, representatives of two peaks observed in histogram of current jump of 5:4:1 composition shown in previous figure.](image)
conduction would yield the same reversal potential as any of its constituents, not a different value with opposite sign. However, the reason for opposite selectivity in each type of pores remains unclear. This lipid dependence of CSFV p7 protein channel activity resembles that reported for the HCV p7 protein [49] although there are some differences regarding the lipid compositions explored, the sub-conductance states and the cationic selectivity found in that case.

4. Conclusions

Given the great interest in viroporins because of their impact on health of humans and animals, many studies have focused on the ability of viroporins to permeabilize cell membranes by using lipid vesicles as model systems. However, approaches probing their pore-formation mechanisms at the single channel level are still scarce [50]. To this end, we have carried out detailed functional studies contrasting IC activity and leakage experiments using same lipid compositions and subject to the same regulatory factors.

For the particular case of CSFV-p7 viroporin, vesicle assays and electrophysiology in planar bilayers seem different sides of the same coin. Under a variety of conditions we observe a good correspondence between the relative number of channel insertions and the leakage observed in LUVs. In both techniques either the pH or membrane composition are equally critical to regulate the permeabilization of the membrane system. Remarkably, electrophysiological recordings show reproducible nanometer-sized pores that in accordance with the solved structure for HCV p7 should allow the transport of solvated ions, water and small molecules like ANTS. The pores display mild ion selectivity, compatible with the documented non-specificity of viroporins.

The analysis of CSFV p7 based peptides in planar bilayers is particularly interesting to identify the sequences involved in CSFV-p7 ion channel activity. Thus, the conclusion that the C-terminal hydrophobic region would remain unknown in the CSFV protein. The analysis of CSFV p7 based peptides in planar bilayers is particularly interesting to identify the sequences involved in CSFV-p7 ion channel activity. Thus, the conclusion that the C-terminal hydrophobic region would remain unknown in the CSFV protein.

Interestingly, lipid modulation of IC activity emerges as a distinctive feature of some viroporins. In the case of CSFV p7, minimal changes in either the lipid charge or the intrinsic lipid curvature originates dramatic changes in the current histograms. The lack of structural information about CSFV-p7 viroporin makes difficult even a qualitative explanation of the lipid regulation the IC activity. The combined analysis of current traces and selectivity experiments indicates that different current states would probably correspond to, at least, two different independent pore structures.

Author declaration

We wish to confirm that there are no known conflicts of interest associated with this publication and there has been no significant financial support for this work that could have influenced its outcome.

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