Characterization of the channel-pores formed by *Bacillus thuringiensis* Cry46Ab toxin in planar lipid bilayers

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Abstract

Cry46Ab from *Bacillus thuringiensis* TK-E6 is a new mosquitocidal toxin with aerolysin-type architecture, and has been shown that co-administration of Cry46Ab with other mosquitocidal Cry toxins results in synergistic toxicity against *Culex pipiens* Coquillett (Diptera: Culicidae) mosquito larvae. Cry46Ab therefore is expected to find use in improving the insecticidal activity of *B. thuringiensis*-based bioinsecticides. In the present study, the mode of action of Cry46Ab was explored by single-channel measurements of Cry46Ab channel-pores. The single-channel conductances of channel-pores formed in planar lipid bilayers by Cry46Ab were determined to be 31.8 ± 2.7 pS in 150 mM NaCl and 24.2 ± 0.7 pS in 150 mM $CaCl_2$. Ion-selectivity measurements revealed that the channel-pores formed by Cry46Ab were cation selective. The permeability ratio of K^+ to Cl^- was approximately 4, and the preferences for cations were $K^+ > Na^+$, $K^+ > Ca^{2+}$, and $Ca^{2+} > Na^+$. A calcein release assay using liposomes suggested that Cry46Ab influences the integrity of membrane vesicles. Formation of cation-selective channel-pores has been observed with other insecticidal Cry toxins that have structures distinct from those of Cry46Ab; the capability of forming such pores may be a property required of insecticidal toxins.

Keywords
Bacillus thuringiensis· Cry46Ab toxin· planar lipid bilayer· single-channel analysis· calcein release assay

Introduction

Mosquito control is a central means by which mosquito-borne diseases such as malaria, viral hemorrhagic fever, and lymphatic filariasis can be prevented. Chemical insecticides have traditionally been used for mosquito control, but these agents have negative impacts on other organisms and the environment. These concerns therefore have prompted the development of alternative approaches such as biological control. The gram-positive spore-forming bacterium *Bacillus thuringiensis* subsp. *israelensis* (Bti) shows strong toxicity against *Anopheles*, *Aedes*, and *Culex* mosquito larvae, and is one of the most widespread and environmentally friendly components used in materials employed for mosquito control. The mosquitocidal activity of Bti resides in three major Cry toxins (Cry4Aa, Cry4Ba, and Cry11Aa) and a Cyt toxin (Cyt1Aa). The structures of Bti Cry toxins have been shown to share a similar three-domain architecture (domains I, II, and III) (Boonserm et al. 2005, 2006; de Maagd et al. 2001). On the other hand, Cyt1Aa has a single domain with a β-sheet in the center surrounded by two α-helical layers (Cohen et al. 2011). Cyt1Aa has a low mosquitocidal activity, but this Cyt toxin synergizes with the Cry toxins (Crickmore et al. 1995; Pérez et al. 2005; Wirth et al. 1997, 2005; Wu et al. 1994). Surprisingly, although Bti-based bioinsecticides have been used to control mosquitos for many years, resistance to the bioinsecticides has not yet

been reported in field populations of mosquitoes (Ben-Dov 2014). The lack of

resistance to Bti-based bioinsecticides has been attributed primarily to the different

modes of action of, and synergistic interactions among, Bti toxins (Crickmore et al.

1995; Poncet et al. 1995; Wirth et al. 2005). However, application of insecticides is

always accompanied by the risk of selecting insecticide resistance in larval mosquito

populations; thus, it is possible that mosquitoes may develop resistance to Bti-based

bioinsecticides in the future, as has been reported in field populations of lepidopteran

insects (Tabashnik et al. 1990). In fact, it has been shown in the laboratory that heavy

and continuous application of individual Bti-derived Cry toxins can induce resistance

(Georghiou and Wirth 1997; Wirth et al. 2012). It is therefore desirable to find toxins

with modes of action that differ from those of Bti toxins in order to prolong the useful

life of Bti-based bioinsecticides.

Cry46Ab is a crystal protein derived from *B. thuringiensis* strain TK-E6. Upon

activation by proteinase K, Cry46Ab shows high cytotoxicities to human leukemic T

cells (MOLT-4 and Jurkat), but not to human embryonal kidney cells (HEK293)

(Hayakawa et al. 2007). Cry46Ab is therefore named parasporin 2Ab exhibiting

preferential cytotoxicity against human cancer cells by Committee of Parasporin

Clasification and Nomenclature (http//parasporin.fitc.pref.fukuoka.jp/index.html). On

the other hand, it has recently been reported that Cry46Ab shows apparent toxicity

against *Culex pipiens* Coquillett (Diptera: Culicidae) mosquito larvae (Hayakawa et al.

2017). Toxicity spectra of Cry46Ab other than human leukemic T cells and *C. pipiens*

mosquito larvae remains to be elucidate.

Interestingly, co-administration of Cry46Ab with other mosquitocidal Cry

toxins, especially the combination of Cry46Ab with Cry4Aa from Bti, results in

significant synergistic toxicity against *C. pipiens* mosquito larvae (Hayakawa et al.

2017). Cry46Ab thus may be a good candidate to improve the activity of crystal toxin

insecticides currently in use, and may prolong the life of Bti-based bioinsecticides.

Cry46Ab shows highest homology (84 % identity at the amino acid sequence level) to

Cry46Aa, a toxin that also is referred to as parasporin-2Aa, from *B. thuringiensis* strain

A1547 (Ito et al. 2004). Cry46Ab also shows relatively high homology (approximately

40 % identity at the amino acid sequence level) to hydralysin, a toxin produced by the

green hydra *Chlorohydra viridissima* (Sher et al. 2005). Therefore, Cry46Ab is inferred

to be a member of the aerolysin-type β pore-forming toxins, a family that includes

parasporin-2Aa and hydralysin. In general, aerolysin-type toxins are produced by a very

diverse group of organisms, and this large family of toxins contains not only aerolysins

from *Aeromonas hydrophila* and related *Aeromonas* species, but also epsilon toxin from

Clostridium perfringens, alpha toxin from *C. septicum*, enterolobin from *Enterolobium contortisiliquum* (a Brazilian tree), and Mtx toxin from *Lysinibacillus sphaericus* (Knapp et al. 2010). Aerolysin-type toxins share structural similarities in their domains, and all act as cytolysins via pore formation. On the other hand, Cry toxins can be divided into several distinct homology groups (Schnepf et al. 1998), and most of them (nearly 90%) belongs to the three-domain Cry toxin group (see http://www.lifesci.sussex.ac.uk/home/Neil_Crickmore/Bt/). The three-domain Cry toxin is a member of the α pore-forming toxin which employ α-helices to insert into the membrane, and are believed to form cation-selective channel-pores (Xu et al. 2014).

Cry46Ab is a functional pore-forming toxin with the single-channel conductance of 103.3 ± 4.1 pS (mean $\pm$ standard deviation) in 150 mM KCl (Hayakawa et al. 2017), but other characteristics of Cry46Ab channel-pores remain unclear. In the present study, the mode of action of channel-pores formed by Cry46Ab was further analyzed using planar lipid bilayers and liposomes.

Materials and Methods

Preparation of recombinant Cry46Ab toxin

Recombinant Cry46Ab fused with glutathione *S*-transferase (GST-Cry46Ab)

was expressed in *Escherichia coli* BL21 as described previously (Hayakawa et al. 2017).

Briefly, *E. coli* cells harboring pGST-Cry46Ab-S1 (Hayakawa et al. 2017) were

cultured at 37 °C in TB medium containing ampicillin (100 μg/ml) until the optical

density at 600 nm (OD_{600}) reached 0.4-0.6, and then expression of recombinant

Cry46Ab was induced with 0.1 mM isopropyl β-D-1-thiogalactopyranoside at 30 °C for

4 h. The GST-Cry46Ab was purified using glutathione-sepharose 4B (GE Healthcare

Bio-Sciences AB, Uppsala, Sweden) according to the manufacturer's instructions.

Protein concentration was estimated using a protein assay kit (Bio-Rad Laboratories,

Inc., Hercules, CA), with bovine serum albumin as the standard. The

mosquito-larvicidal activity of the GST-Cry46Ab was analyzed by bioassay using *C.*

pipiens larvae (3rd instar), as described previously (Hayakawa et al. 2017). Briefly, forty

micrograms of the GST-Cry46Ab was adsorbed onto 2 mg of latex beads (0.8-μm

diameter, Sigma-Aldrich Corp., St. Louis, MO), and then administered to the mosquito

larvae as a diet. In a preliminary experiment, latex beads adsorbed GST was not toxic

against *C. pipiens* larvae at a working concentration of 2 μg/ml (data not shown).

Purified GST-Cry46Ab was activated using a trypsin-immobilized column as

described previously (Hayakawa et al. 2017). The activation process was monitored by

analysis using sodium dodecyl sulfate - 15% polyacrylamide gel electrophoresis

(SDS-PAGE), and the activated Cry46Ab (a polypeptide of 29 kDa) was eluted as the

flow-through. Activated Cry46Ab was concentrated using Vivaspin 20 (10-kDa

molecular weight cut-off) centrifugal filter devices (GE Healthcare Bio-Sciences AB).

Lipid bilayer experiments

Pore formation by activated Cry46Ab in planar lipid bilayers was analyzed as

described previously (Hayakawa et al. 2017). Briefly, each instrument consisted of two

chambers (upper, *cis* chamber; lower, *trans* chamber), and the bottom of the *cis* chamber

was a thin sheet of polyvinyl chloride with a small circular hole (approximately φ200

μm). A lipid bilayer was prepared by painting an asolectin (phospholipids from soybean,

Sigma-Aldrich Corp.) solution (40 mg/ml n-decane) across the small hole. The

chambers were held at virtual ground, such that the voltage in the solution of the *cis*

chamber was connected to a patch-clamp amplifier by an Ag/AgCl electrode-defined

membrane potential. After injection of activated Cry46Ab (0.1 μg/ml) into the solution

in the *cis* chamber, the toxin was incorporated into the lipid bilayer while applying a

70-mV holding potential across the lipid bilayer. Data were analyzed using pClamp

software (ver 9.2, Axon Instruments, Foster City, CA).

Unless otherwise stated, the current amplitude of the resolvable steps was

recorded for each experiment, and the resulting data were plotted versus the

corresponding applied voltage to generate the current-voltage relationship. Channel

conductance was determined from the slope of the linear regressions on similar data

points. Corresponding values obtained in different experiments were subjected to

descriptive analysis by calculating (over the number of experiments in which the values

had been observed) the arithmetic mean and standard deviation.

To assess ion selectivity of the pores formed by Cry46Ab, membrane currents

through the channel-pore formed by Cry46Ab were recorded with a 4-fold gradient of

KCl across the lipid bilayer (600 mM KCl and 10 mM Tris-HCl (pH 8.0) in the *cis*

chamber, 150 mM KCl and 10 mM Tris-HCl (pH 8.0) in the *trans* chamber). The

zero-current reversal potential (V_R) was obtained as X-intercept of the linear showing

current-voltage relationship, and the P_K/P_{Cl} permeability ratio was calculated using the

Goldman-Hodgkin-Katz equation as follows (Benz et al. 1979).

$$V_R = \frac{RT}{F} \ln \frac{P_K[K^+]_{cis} + P_{Cl}[Cl^-]_{trans}}{P_K[K^+]_{trans} + P_{Cl}[Cl^-]_{cis}}$$

In the formula, R is molar gas constant, T is absolute temperature, and F is Faraday

constant.

Calcein release assay

Large unilamellar vesicles (LUVs) of liposome were prepared as described

previously (Shai et al. 1990) with some modifications. Briefly, 5 mg of asolectin

(Sigma-Aldrich Corp.) was dissolved in 500 µl of chloroform. The chloroform then was

removed using a circulating aspirator to generate a thin film of asolectin. The film was

suspended in 500 µl of calcein solution (100 mM calcein, 100 mM NaCl, and 50 mM

Tris-HCl, pH 8.3) by sonication for 10 min at intervals of 30 sec on, 30 sec off. The

resulting suspension was allowed to stand for 30 min at room temperature to permit the

formation of multilamellar vesicles, leading in turn to the subsequent formation of small

unilamellar vesicles of liposome. Liposomal LUVs were generated by subjecting the

small unilamellar vesicle suspension to repeated freeze-thaw cycles using liquid

nitrogen at room temperature. LUVs containing calcein were purified using

discontinuous sucrose density gradient centrifugation. The sucrose gradient was

generated by sequential layering of 35, 30, 25, 20, and 10 % (w/w) sucrose solutions.

The liposome suspension was mixed with 2 volumes of 64 % (w/w) sucrose to yield a

suspension in a solution of approximately 40 % (w/w) sucrose, and this mixture was

placed at the bottom of sucrose gradient. After centrifugation at 200,000 x g for 1.5 hr at

4 °C, the thin orange-colored layer of the liposome fraction was collected; the orange

color indicated the presence of self-quenched calcein. The liposomes then were

subjected to a second round of purification using sucrose density gradient centrifugation (for two purifications total).

The ability of Cry46Ab to disrupt liposomes was evaluated by a calcein release assay. Calcein is self-quenched at high concentration (100 mM), as in the liposome, but emits fluorescence when calcein is diluted by disruption of the liposome. The fluorescence intensity of calcein released from the liposomes was measured using a spectrofluorophotometer (RF-5300pc, Shimadzu, Osaka, Japan) with excitation at 490 nm and emission at 520 nm. Fluorescence intensity was monitored every 20 sec for 5 min after addition of the activated Cry46Ab to the liposome suspension. The amount of fluorescence released was expressed as a percentage of the maximum fluorescence intensity achieved by the addition of Triton X-100 at a final concentration of 1% which disrupt liposome completely (fluorescence recovery).

Results

Recombinant Cry46Ab

Recombinant Cry46Ab was successfully expressed as a GST fusion in *E. coli*, and the molecular mass of purified GST-Cry46Ab was approximately 60 kDa, similar to the expected mass (59.309 kDa) (Fig. 1A). After treatment using a trypsin-immobilized

column, GST-Cry46Ab was processed into a polypeptide of 29 kDa (Fig. 1 A), similar

in size to that of the activated Cry46Ab that was shown to be toxic against human

leukemic T cells (Hayakawa et al. 2007) and against *C. pipiens* larvae (Hayakawa et al.

2017). GST-Cry46Ab exhibited apparent toxicity against *C. pipiens* larvae, with an

LC_{50} value (95% confidence limits) of 1.01 (0.98 – 1.05) µg/ml (Fig. 1B).

Single-channel measurement

Pore formation by activated Cry46Ab in the planar lipid bilayer was assessed

previously, when the single-channel conductance was determined to be 103.3 ± 4.1 pS

in 150 mM KCl (Hayakawa et al. 2017). In the present study, to characterize more

precisely the channel-pores formed by Cry46Ab in a planar lipid bilayer, the ability of

Cry46Ab to permeabilize an artificial lipid bilayer was analyzed in salt solutions

containing NaCl or $CaCl_2$, salts distinct from the KCl used in the previous experiments.

In a symmetrical salt solution containing 150 mM NaCl and 10 mM Tris-HCl

(pH 8.0), apparent channel currents were detected between −100 and +100 mV at

approximately 30 min after the addition of activated Cry46Ab to the solution in the *cis*

chamber (Fig. 2A). These channel currents were observed only in the presence of toxin.

The recordings of membrane currents were similar to those observed previously in 150

mM KCl (Hayakawa et al. 2017), and were characterized as long periods of closure

interrupted by bursts of current jumps and rapid flickering between open and closed

states (Fig. 2A). The current-voltage relationship was linear at several conductance

levels. Most frequently observed was a line indicating a channel conductance of $31.8 \pm$

2.7 pS, followed (successively less frequently) by lines indicating conductances of 59.7

$\pm$ 1.1, 89.8 $\pm$ 0.7, 119.4 $\pm$ 1.3 and 150.6 $\pm$ 2.8 pS (Fig. 2B and C). In fact, 639 single

events were recorded in total, consisting of 259, 127, 82, 76, and 34 events at the

respective channel conductances (Fig. 2B and C). As the channel conductances

observed in this experiment occurred almost universally in multiples of approximately

30 pS, channel conductance higher than 30 pS was considered to be the result of the

synchronized opening and closing of multiple channels. The conductance of single

channels formed by Cry46Ab on the planar lipid bilayer thus was determined to be 31.8

$\pm$ 2.7 pS in 150 mM NaCl.

Recordings of membrane currents in the solution containing 150 mM $CaCl_2$

and 10 mM Tris-HCl (pH 8.0) yielded a profile similar to those observed in KCl and

NaCl solutions (Fig. 3A). In the current-voltage relationship, the most frequently

observed line was one indicating a channel conductance of 24.2 $\pm$ 0.7 pS (Fig. 3B and

C). Although the frequency was low, a line indicating a conductance of 52.6 $\pm$ 3.9 pS

also was observed (Fig. 3B and C). Three hundred and sixty-four single events were recorded in total, and 230 and 71 of the events occurred on lines indicating channel conductances of 24.2 ± 0.7 and 52.6 ± 3.9 pS, respectively (Fig. 3B and C). Thus, the conductance of single channels formed by Cry46Ab on the planar lipid bilayer was determined to be 24.2 ± 0.7 pS in 150 mM $CaCl_2$.

Ion selectivity of the pores formed by Cry46Ab

Membrane currents through the channel-pore formed by Cry46Ab also were recorded with a 4-fold gradient of KCl across the lipid bilayer. As a result, various channel currents with different levels were detected between -50 and $+50$ mV upon addition of activated Cry46Ab to the solution in the *cis* chamber (data not shown). In this assay, the current amplitudes of the resolvable steps were recorded, pooled for 8 independent experiments, and averaged over the number of steps at each applied voltage (Fig. 4A). Considering the results of single-channel measurements in this study and in the previous report (Hayakawa et al. 2017), Cry46Ab is thought to form channel-pores with very similar conductances in planar lipid bilayers, and higher channel currents are thought to be caused by synchronized opening and closing of multiple channels. It was therefore reasonable to speculate that all of the linear showing current-voltage

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relationship observed in the present experiment share the same X-intercept. In fact, the

current-voltage relationship constructed for this measurement approached linearity, and

the V_R was determined to be -18.38 mV (Fig. 4A). The P_K/P_{Cl} permeability ratio

calculated from this V_R value (using the Goldman-Hodgkin-Katz equation (Benz et al.

1979)) was 3.67, demonstrating a higher permeability for K^+ than for Cl^-.

In addition, the effect of pH on the channel-pore formed by Cry46Ab was

assessed. The membrane currents were recorded under the same experimental

conditions as above, but 10 mM Tris-HCl (pH 6.0) was used as the buffer instead of 10

mM Tris-HCl (pH 8.0). As a result, various channel currents were detected at voltages

between −50 and +50 mV, with these currents occurring at levels distinct from those

observed in the measurement using a buffer of pH 8.0 (data not shown). The current

amplitudes of the resolvable steps were recorded, pooled for 7 independent experiments,

and averaged over the number of steps at each applied voltage. The current-voltage

relationship approached linearity, and the V_R was determined to be -20.04 mV; the

P_K/P_{Cl} permeability ratio calculated from this V_R value was 4.25 (Fig. 4B). Thus, the

channel-pores formed in the planar lipid bilayer by Cry46Ab appear to be cation

selective.

Preferred cations

To determine the preferred cations (K^+, Na^+, or Ca^{2+}) for the channel-pores

formed by Cry46Ab, membrane currents were measured across the lipid bilayer when

different salt solutions were present on the two sides of the chamber. In this experiment,

KCl and NaCl were used at a concentration of 150 mM, and $CaCl_2$ was used at a

concentration of 75 mM to provide Cl^- at a concentration equivalent to those in the KCl

and NaCl solutions.

When 150 mM KCl was provided in the *cis* chamber and 150 mM NaCl was

provided in the *trans* chamber, incorporation of Cry46Ab into the lipid bilayer yielded

positive membrane current without holding potential (Fig. 5A). Similar preferential

permeability was observed when the salt solutions were exchanged between the *cis* and

trans chambers (150 mM NaCl in the *cis* chamber and 150 mM KCl in the *trans*

chamber) (Fig. 5B). This result demonstrated that the Cry46Ab channel-pore has a

higher permeability for K^+ than for Na^+, and that the movement of K^+ across the lipid

bilayer occurs in both directions through this channel-pore. On the other hand, when the

measurement was performed with 150 mM KCl in the *cis* chamber and 75 mM $CaCl_2$ in

the *trans* chamber, positive membrane current was observed (Fig. 5C), indicating

preferential permeability for K^+ compared to Ca^{2+}. When the measurement was

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performed with 150 mM NaCl in the *cis* chamber and 75 mM CaCl$_2$ in the *trans*

chamber, negative membrane current was observed (Fig. 5D), indicating preferential

permeability for Ca^{2+} compared to Na$^+$. Therefore, the cation preferences of the

channel-pores formed by Cry46Ab were K$^+$ > Na$^+$, K$^+$ > Ca^{2+}, and Ca^{2+} > Na$^+$.

Calcein release assay

LUVs of asolectin were created with the self-quenched fluorescent dye calcein

entrapped in their interior cavities, and membrane perturbation was monitored by the

increase in fluorescence due to the dilution of calcein in the surrounding solution. After

addition of activated Cry46Ab to the LUV suspension, the fluorescence intensity

increased in a rapid and dose-dependent manner (Fig 6). The estimated fluorescence

recoveries at 5 min after addition of Cry46Ab at 10.0, 2.5, and 0.6 μg/ml (final

concentration) were 59.2, 46.2, and 15.9 %, respectively (Fig 6). These observations

suggested that Cry46Ab (and presumably, the channel-pore formed by Cry46Ab)

influences the integrity of membrane vesicles.

Discussion

We previously demonstrated that Cry46Ab is a functional pore-forming toxin

with a single-channel conductance of 103.3 ± 4.1 pS in 150 mM KCl (Hayakawa et al.

2017), but other characteristics of these channel-pores remained unclear. Therefore, to

characterize the channel-pore more precisely, single-channel analyses were performed

using NaCl or CaCl$_2$ solution, that is, employing salts distinct from the

previously-tested KCl. The measurements of membrane currents in both 150 mM NaCl

and 150 mM CaCl$_2$ solutions revealed typical current transitions between open and

closed states, and a result similar to that previously observed in the 150 mM KCl

solution. Single-channel conductances of the channel-pores in 150 mM NaCl and in 150

mM CaCl$_2$ were 31.8 ± 2.7 and 24.2 ± 0.7 pS, respectively. However, the single-channel

conductance of the channel-pore in 150 mM NaCl was apparently lower than that in 150

mM KCl; this observation may suggest some difference in characteristics of the

Cry46Ab channel-pores compared to those formed by aerolysin. Notably, aerolysin

from *S. sobria* has been shown to form channel-pores with similar single-channel

conductance in both KCl and NaCl solutions (650 pS in 1 M KCl; 600 pS in 1 M NaCl)

(Chakraborty et al. 1990).

Measurement of membrane currents in a 4-fold KCl gradient (600 mM in the

cis chamber, 150 mM in the *trans* chamber) demonstrated that the channel-pores formed

by Cry46Ab permeabilized K$^+$ in preference to Cl$^-$; the P_K/P_{Cl} permeability ratios

calculated at pH 6.0 and 8.0 were 3.67 and 4.25, respectively. The cation preference of

the Cry46Ab channel-pores were determined to be $K^+ > Na^+$, $K^+ > Ca^{2+}$, and $Ca^{2+} > Na^+$.

Formation of cation-selective channel-pores in planar lipid bilayers has been observed

with other insecticidal Cry toxins, including Cry1Aa (Grochulski et al. 1995), Cry1Ac

(Slatin et al. 1990), Cry1C (Schwartz et al. 1993), Cry3Aa (Slatin et al. 1990), Cry3B2

(Von Tersch et al. 1994), and Cry4Ba (Puntheeranurak et al. 2004). Like Cry46Ab, these

molecules are insecticidal Cry toxins, but unlike Cry46Ab, these other toxins are

members of the α pore-forming toxin family and do not share structural similarity with

aerolysin-type toxins. It is believed that, after sequential interaction with specific

receptors facilitating formation of the toxin oligomer, insecticidal Cry toxins insert into

the cell membrane of columnar cells in the midgut tissue and create pores that allow cell

uptake of cations. The uptake of cations causes the influx of excess water into the

midgut cells, leading to swelling and bursting, a cell lysis process referred to as

colloid-osmotic lysis (Knowles and Ellar 1987). Considering this previous work and the

observations in the present work, the formation of cation-selective channel-pores by

crystal proteins may be a property required of insecticidal toxins. In the present study,

the effect of channel-pores formed by Cry46Ab on the integrity of membrane vesicles

also was assessed by the calcein release assay. After addition of activated Cry46Ab to

LUVs containing self-quenched calcein, fluorescence intensity increased quickly in a

dose-dependent manner. This observation suggested that Cry46Ab, and presumably the

channel-pore formed by Cry46Ab, influences the integrity of membrane vesicles.

On the other hand, among aerolysin-type toxins, the enterotoxin from *C.*

perfringens forms highly cation-selective channel-pores in planar lipid bilayers (Benz

and Popoff 2018), but other aerolysin-type toxins (including aerolysin from *A. sobria*

(Chakraborty et al. 1990)), alpha toxin from *C. septicum* (Ballard et al. 1993), and

epsilon toxin from *C. perfringens* (Petit et al. 2001)) form slightly anion-selective

channel-pores in planar lipid bilayers. All of these aerolysin-type toxins are classified as

β pore-forming toxins that, upon binding to the receptor of the target cell, form toxin

oligomers that can insert into the cell membrane and form channel-pores. Formation of

channel-pores induces ion fluxes leading to membrane depolarization and ultimately

cell death (Abrami et al. 2000). The amphipathic β-hairpin in the middle domain of

these toxin molecules is thought to constitute the transmembrane β-barrel cylinder

(Melton et al. 2004). The β-hairpin is composed of a striking pattern of alternating

charged and non-charged amino acids; the charged amino acids are thought to line the

lumen of the β-barrel cylinder (Melton et al. 2004). It is believed that the composition

and/or sequence of charged amino acids in the β-hairpin region is responsible for the ion

selectivity of aerolysin-type toxins. The corresponding β-hairpin can be identified in the structure of Cry46Ab based on its sequence similarity with Cry46Aa (parasporin-2Aa, Akiba et al. 2009). Therefore, it will be of interest to determine the amino acid residues in the β-hairpin of Cry46Ab that affect the ion selectivity, as well as the effect of this domain on to Cry46Ab's insecticidal activity. Currently, the residues that determine the ion selectivity of channel-pores remain unclear; Cry46Ab may be useful as a model for elucidating the mechanism of ions permeation through such channel-pores.

Co-administration of Cry46Ab with other mosquitocidal Cry toxins (including Cry4Aa, Cry11Aa, and Cry11Ba) has been shown to result in synergistic toxicity against *C. pipiens* mosquito larvae (Hayakawa et al. 2017). Cry4Aa (Boonserm et al. 2006), Cry11Aa (Fernández et al. 2005), and Cry11Ba (Likitvivatanavong et al. 2009) are three-domain Cry toxin, and are therefore hypothesized to form cation-selective channel-pores in a manner similar to that of other insecticidal Cry toxins. On the other hand, it is unclear whether this synergistic toxicity is generated in association with the cation-selective channel-pore formation. Specifically, it is possible that the mechanism of this synergistic toxicity depends on multiple other factors in addition to pore formation.

It is widely accepted that Cry toxin recognizes the specific receptors on midgut

epithelial cells of susceptible insect larva. The specific receptors promote

toxin-membrane interaction, and is believed to play a crucial role in the insecticidal

spectra. Since Cry46Ab also exhibits specificity, a specific receptor should be present

on the target cell membrane. On the other hand, formation of channel-pores is probably

a fundamental property of Cry toxins, it is believed that Cry toxins at high concentration

is able to form channel-pores in a membrane even without specific receptors. In fact, it

has been demonstrated that several Cry toxins form channel-pores in the planar lipid

bilayer without specific receptors (Grochulski et al. 1995; Puntheeranurak et al. 2004;

Schwartz et al. 1993; Slatin et al. 1990; Von Tersch et al. 1994). In the present study, we

used 0.1 µg/ml of Cry46Ab for single-channel measurements, and the concentration was

demonstrated to be high enough to form channel-pores in planar lipid bilayer. On the

other hand, calcein release assay revealed that much higher concentration of Cry46Ab

(>0.6 µg/ml) required to influence the integrity of membrane vesicles. It is speculated

that if the membrane with specific receptors is available, amount of Cry46Ab required

for measurements is reduced. It is plausible to conclude that the channel-pores observed

in the present study is central to the toxicity of Cry46Ab.

Acknowledgments

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Figure legends

Fig. 1 The recombinant Cry46Ab toxin. (A) The recombinant Cry46Ab expressed as a

GST fusion was purified using glutathione beads, and was activated using a

trypsin-immobilized column. Protein was analyzed by 15% SDS-PAGE. A total of 1 µg

of protein was applied in each lane. (B) Mosquitocidal activities of GST-Cry46Ab. The

experiments were repeated independently six times. Mortality rates, observed at 48 h after administration, are presented as mean (standard deviation).

Fig. 2 Single-channel analysis of activated Cry46Ab in symmetrical 150 mM NaCl solutions. (A) Representative segments of typical current traces. The current levels corresponding to the open state of the channels are indicated by dashed lines. (B) Representative current-voltage relationship obtained from the Cry46Ab current steps. The experiment was repeated three times independently, and the mean and standard deviation are shown. (C) Frequency of conductance observed on the Cry46Ab current steps. Six hundred and thirty-nine single events were recorded in total.

Fig. 3 Single-channel analysis of activated Cry46Ab in symmetrical 150 mM CaCl$_2$ solutions. (A) Representative segments of typical current traces. The current levels corresponding to the open state of the channels are indicated by dashed lines.   (B) Representative current-voltage relationship obtained from the Cry46Ab current steps. The experiment was repeated more than three times independently, and the mean and standard deviation are shown. (C) Frequency of conductance observed on the Cry46Ab current steps. Three hundred and sixty-four single events were recorded in total.

Fig. 4 Ion selectivity of the channel-pore formed by activated Cry46Ab. Anion-cation selectivity of the channel-pore was determined using 4-fold gradients of KCl across the lipid bilayer. Current amplitudes of the resolvable steps were recorded, pooled for 8 independent experiments, and averaged over the number of steps at each applied voltage. The current-voltage relationship was constructed and the zero-current reversal potential (V_R) was determined. The P_K/P_{Cl} permeability ratios were calculated from this V_R value using the Goldman-Hodgkin-Katz equation. (A) Measurement in pH 8.0 solution. (B) Measurement in pH 6.0 solution.

Fig. 5 Selectivity order of cations for the channel-pore formed by activated Cry46Ab. Selectivity order of cations for the channel-pore formed by activated Cry46Ab was determined by membrane current measurement under asymmetrical buffer conditions using (A) 150 mM KCl / 150 mM NaCl, (B) 150 mM NaCl / 150 mM KCl, (C) 150 mM KCl / 75 mM $CaCl_2$, (D) 150 mM NaCl / 75 mM $CaCl_2$ (*cis* chamber / *trans* chamber), respectively. Membrane current was measured without holding potential upon incorporation of Cry46Ab into the lipid bilayer.

Fig. 6 Calcein release assay. LUVs of asolectin were created with the self-quenched

fluorescent dye calcein entrapped in their interior cavities. After addition of activated

Cry46Ab to the LUV suspension, the increase in fluorescence intensity was monitored

every 20 sec for 5 min. Fluorescence recovery was expressed as a percentage of the

maximum fluorescence intensity, achieved by the addition of Triton X-100. Open circles

(○), filled circles (●) and squares indicate final concentration of Cry46Ab at 10.0, 2.5,

and 0.6 μg/ml, respectively.