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Modulation of iridovirus-induced apoptosis by endocytosis, early expression, JNK, and apical caspase

Nilesh S. Chitnis a,1, Susan M. D’Costa a,2, Eric R. Paul a,3, Shän L. Bilimoria a,b,*

a Department of Biological Sciences, Texas Tech University, Lubbock, TX 79409-3131, USA
b The Center for Biotechnology and Genomics, Texas Tech University, Lubbock, TX 79409-3131, USA

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Abstract

Chilo iridescent virus (CIV) is the type species for the family Iridoviridae, which are large, isometric, cytoplasmic dsDNA viruses. We examined the mechanism of apoptosis induction by CIV. High CIV doses (CIVXS; 400 μg/ml), UV-irradiated virus (CIVUV; 10 μg/ml) and CVPE (CIV protein extract; 10 μg/ml) induced apoptosis in 60% of treated Choristoneura fumiferana (IPRI-CF-124T) cells. Normal doses of infectious CIV (10 μg/ml) induced apoptosis in only 10% of C. fumiferana (CF) cells. Apoptosis was inhibited by Z-IETD-FMK, an apical caspase inhibitor, indicating that CIV-induced apoptosis requires caspase activity. The putative caspase in CF cells was designated Cf-caspase-i. CIVUV or CVPE enhanced Cf-caspase-i activity by 80% at 24 h relative to mock-treated cells. Since the MAP kinase pathway induces or inhibits apoptosis depending on the context, we used JNK inhibitor SP600125 and demonstrated drastic suppression of CVPE-induced apoptosis. Thus, the JNK signaling pathway is significant for apoptosis in this system. Virus interaction with the cell surface was not sufficient for apoptosis since CIVUV particles bound to polysterene beads failed to induce apoptosis. Endocytosis inhibitors (bafilomycin or ammonium chloride) negated apoptosis induction by CIVUV, CIVXS or CVPE indicating that entry through this mode is required. Given the weak apoptotic response to infectious CIV, we postulated that viral gene expression inhibited apoptosis. CIV infection of cells pretreated with cycloheximide induced apoptosis in 69% of the cells compared to 10% in normal infections. Furthermore, blocking viral DNA replication with aphidicolin or phosphonoacetic acid suppressed apoptosis and Cf-caspase-i activity, indicating that early viral expression is necessary for inhibition of apoptosis, and de novo synthesis of viral proteins is not required for induction. We show for the first time that, in a member of the family Iridoviridae, apoptosis: (i) requires entry and endocytosis of virions or virion proteins, (ii) is inhibited under conditions permitting early viral expression, and (iii) requires the JNK signaling pathway. This is the first report of JNK signal requirement during apoptosis induction by an insect virus.

Keywords: Chilo iridescent virus; Iridovirus; Apoptosis; Inhibition; Endocytosis; Viral early expression; JNK; Apical caspase; Insect; Lepidopteran cells

Introduction

Chilo iridescent virus (CIV) is the type species for the family Iridoviridae (Iyer et al., 2006). The Iridoviridae are large, cytoplasmic DNA viruses with an icosahedral outer capsid, an internal lipid membrane, and an electron-dense core containing a circularly permuted double-stranded DNA genome (209 kbp). CIV infects a wide range of insects (Williams et al., 2005), replicates productively in the cotton boll weevil Anthonomen grandis (Henderson et al., 2001), and induces metamorphic arrest and mortality in this host (McLaughlin et al., 1972). Earlier work showed that UV-treated CIV or viral “soluble” protein extract inhibited host protein synthesis in mosquito cells (Cerutti and Devauchelle, 1980). Recently, our group showed that a CIV virion protein extract (CVPE) induces mortality in neonate boll weevil larvae (Bilimoria, 2001) and apoptosis in spruce budworm Choristoneura fumiferana and boll weevil cell cultures (Paul et al., 2007).

Apoptosis is generally characterized by several criteria. These include caspase activation, cytoplasmic shrinkage, phosphatidylserine exposure, chromatin condensation, DNA fragmentation,
and cell blebbing (Hay and Kannourakis, 2002). Apoptosis proceeds through signal transduction pathways, usually involving activation of cysteine-aspartate-specific proteases, i.e., caspases. Various signals activate initiator (apical) caspases, which then trigger effector (executioner) caspases. In Drosophila, one mechanism includes interaction of Eiger (invertebrate tumor necrosis factor ligand) with the wengen membrane receptor. This leads to activation of JNK, which then triggers the central initiator caspase DRONC, instigating a cascade triggering the effector caspase Drice and ultimately apoptosis (Igaki et al., 2002). The Eiger-wengen mediated pathway appears to be the invertebrate equivalent of the “extrinsic” pathway found in mammalian systems. Alternate induction mechanisms in Drosophila include activation of DRONC through ER stress or through bcl-2-type stimulation and could represent invertebrate equivalents of “intrinsic” pathways. Induction of apoptosis in uninfected Spodoptera frugiperda (Lepidoptera: Noctuidae) cells appears to involve the equivalents of mammalian apical caspase and caspase-3 (Manji and Friesen, 2001) as initiator and effector caspases, respectively.

Recently, three iridoviruses from the genera Ranavirus and Lymphocystivirus were shown to induce apoptosis in cell culture (Essbauer and Ahne, 2002; Chinchar et al., 2003; Hu et al., 2004; Imajoh et al., 2004). Apoptosis induction by frog virus-3 required caspases, and induction of apoptosis by red sea bream iridovirus involved effector caspases-3 and 6 (Chinchar et al., 2003; Imajoh et al., 2004). However, the role of viral entry and endocytosis in the induction of apoptosis by the Iridoviridae is not established, and the significance of viral gene expression in apoptosis modulation has not been studied.

In this report, we show for the first time that in a member of the family Iridoviridae: (i) entry and endocytosis of virions or virion proteins are required for apoptosis induction, (ii) apoptosis is inhibited under conditions permitting early viral expression, (iii) apoptosis requires the JNK signaling pathway. This is the first report of JNK signal requirement during apoptosis induction by an insect virus.

Results

Normal and UV-treated Chilo iridescent virus induce apoptotic blebbing in spruce budworm cells

To determine if CIV replication was necessary for induction of apoptosis in C. fumiferana (CF) cells, we treated these cells with normal and UV-inactivated CIV (CIVUV) as described in Materials and methods. Normal, active CIV was used at two concentrations: high dosage (CIVXS; 400 μg/ml) and standard dosage (CIV; 10 μg/ml). Infected cells were incubated at 28 °C as well as 21 °C for 24 h and observed for cell blebbing. Fig. 1 shows that CIVXS induced 70% blebbing (A), whereas CIV induced 10% blebbing and 55% lysis in these cells (B). Lysis was indicated by the presence of cell debris. CIVUV (10 μg/ml) induced 62% blebbing and 5% lysis in CF cells (C). Mock-treated cells incubated at 21 °C or 28 °C did not undergo significant blebbing or lysis (D). The above results are consistent with apoptosis induction using CIV virion protein extract (CVPE) (Bilimoria, 2001; Paul et al., 2007).

Apoptosis induced by high-dosage CIV, CVPE and UV-irradiated CIV is negated by inhibitors of caspase activity

To determine whether apoptosis induced by high-dosage CIV, CVPE, or UV-irradiated CIV was caspase-dependent, CF

![Fig. 1. Induction of blebbing in Choristoneura fumiferana (CF124T) cells treated with Chilo iridescent virus and UV-treated CIV. (A) Cells treated with very large doses of CIV (400 μg/ml). (B) Cells infected with normal doses of CIV (10 μg/ml). (C) Cells treated with UV-irradiated CIV (CIVUV; 10 μg/ml). Incubation for panel A through panel C was at 21 °C. (D) Mock-treated cells were incubated at 21 °C or 28 °C. Observations were made by phase-contrast microscopy. Cells were photographed 24 h after infection or treatment. Magnification: 200×.](image-url)
cells were pretreated with pancaspase inhibitor (Z-VAD-FMK, Calbiochem, CA) and challenged with the above inducers. Pancaspase inhibitor reduced blebbing as follows: CIVXS-induced by 93% (Fig. 2A), CIVUV-induced by 97% (Fig. 2B), CVPE-induced by 99% (Fig. 2C). To establish whether the apoptotic pathway in this system requires an apical caspase, CF cells were pretreated with Z-IETD-FMK (a mammalian caspase-8 inhibitor shown to inhibit lepidopteran, Sf-caspase-X) (Manji and Friesen, 2001) prior to challenge with CIVXS, CIVUV, or CVPE. The results show that Z-IETD-FMK blocked apoptotic blebbing by at least 97% for CIVXS (Fig. 2A); 96% for CIVUV (Fig. 2B); 96% for CIVUV and 98% for CVPE (Fig. 2C). Mock treatment with RBSS yielded less than 2% blebbing (Fig 2D). These results suggest that apoptosis induction in CF cells by high-dosage CIV, CVPE, and UV-treated CIV is dependent on Sf-caspase-X-like activity, which we have designated *C. fumiferana* initiator caspase (Cf-caspase-i).

**CVPE and UV-irradiated CIV induce apical caspase activity**

To confirm whether an apical enzyme is indeed activated by CVPE, UV-irradiated CIV, and CIV as suggested by the inhibitor data (Fig. 2), we tested for apical caspase activity using Caspase-Glo Assay kit (Promega, WI) as described in Materials and methods. In this assay the LETD sequence, which is specific for apical caspasps, was used as substrate (Lien et al., 2004). Fig. 3 shows that CVPE increased apical caspase activity by 89% as compared to mock treatment by 24 h post-treatment. UV-irradiated CIV increased apical caspase activity by 88%, and enzyme activity in CIV-infected cells increased by 65% compared to mock treatments. These results are consistent with the caspase inhibitor data (Fig. 2) and further support a major role for an apical caspase during apoptosis induction in the CIV system.

**JNK inhibitor negates apoptosis induction by CVPE**

JNK plays an important role in regulation of *Drosophila* apoptosis through several pathways (Mizutani et al., 2003; Kanda and Miura, 2004). JNK signaling can augment or inhibit apoptosis depending on the context. Therefore, we examined the effect of JNK inhibitor in the CIV-CF cell system. Fig. 4 shows that pretreatment with JNK inhibitor SP600125 (25 nM) reduced CVPE-induced apoptosis by 87%, suggesting a major role for JNK signaling in the regulation of CIV-induced apoptosis. Treatment with JNK inhibitor alone or mock treatment with Tris-NaCl buffer induced negligible blebbing.

**Inhibition of endocytosis negates apoptosis induction by CVPE, UV-irradiated CIV and high-dose CIV**

It was important to determine whether interaction of viral proteins with the cell surface in the CIV-CF cell system is
sufficient for inducing apoptosis or whether there is a requirement for cellular entry. The use of ammonium chloride to inhibit acidification of endosomes in insect cells is well documented (Hacker and Hardy, 1997; IJkel et al., 2000). Fig. 5A shows that 10 mM ammonium chloride reduced apoptosis induction by CVPE (83%), CIVUV (77%), and high-dose CIV (69%). Fig. 5B shows that bafilomycin A1 (1 μM), an inhibitor of endocytosis (Mizutani et al., 2003; Long et al., 2006), also reduced apoptosis induction by CVPE (93%), CIVUV (93%), and high-dose CIV (91%) in CF cells. These results suggest that induction of apoptosis by CVPE, UV-irradiated CIV, or high-dose CIV is endocytosis-dependent.

**Virus–cell surface interaction is not sufficient for induction of apoptosis**

To confirm that induction of apoptosis requires viral entry rather than surface interaction of virions with cells, we attached CIVUV to polystyrene beads as described in Materials and methods (Paran et al., 2001). UV-treated CIV (100 μg/ml) was attached to 12 mg polystyrene beads (average diameter 10 μm) per manufacturer’s protocol (Polysciences, Inc., PA). Beads were extensively washed after conjugation with virus, and attachment of virus to beads was confirmed using a virion-associated kinase assay. Kinase activity has been demonstrated in CIV particles (Monnier and Devachelle, 1980), and our group recently showed kinase activity in CVPE (Paul et al., 2007). Therefore, interaction of bead-bound CIV with cells was confirmed with kinase assays on whole cells. CIVUV to polystyrene beads as described in Materials and methods (Paran et al., 2001). This would allow interaction of virions with the cell surface but preclude viral entry.

UV-treated CIV (100 μg/ml) was attached to 12 mg polystyrene beads (average diameter 10 μm) per manufacturer’s protocol (Polysciences, Inc., PA). Beads were extensively washed after conjugation with virus, and attachment of virus to beads was confirmed using a virion-associated kinase assay. Kinase activity has been demonstrated in CIV particles (Monnier and Devachelle, 1980), and our group recently showed kinase activity in CVPE (Paul et al., 2007). Therefore, interaction of bead-bound CIV with cells was confirmed with kinase assays on whole cells. CIVUV–bead conjugate and gamma32P-ATP were added to cells and incubated at 21 °C for 24 h. Staurosporine and genistein were added to stop all kinase activity after incubation. The cell lysate was analyzed on SDS–PAGE and visualized by phosphorimaging. Interaction of CIVUV–bead conjugate with the cell surface was confirmed by the presence of at least one phosphorylated cellular protein. CIVUV–beads in supplemented media devoid of...
Low-pH-dependent endocytosis is required for CIV infection

Since CVPE and CIVUV required low-pH-dependent endocytosis for induction of apoptosis, we hypothesized that CIV infection should also require endocytosis. Therefore, we tested for virus replication in the presence and absence of ammonium chloride (10 mM) and bafilomycin A1 (1 μM) by observing for viral cytopathic effect and DNA replication. Fig. 6 shows that CIV-induced cytopathic effect (consisting of hypertrophy, syncytia formation, granularity, or lysis) was drastically reduced in the presence of AC (85% reduction) and bafilomycin A1 (95% decrease). Fig. 6 also shows that viral DNA levels (determined by dot blot analysis; see Materials and methods) in CIV infections containing AC or Bf were significantly lower (more than 95% after correction for mock infections) than those in normal CIV infections and barely above those in mock-infected cells. These results indicate that low-pH-dependent endocytosis is required for CIV infection.

Pretreatment with cycloheximide augments apoptosis by CIV and CVPE

In order to determine whether de novo protein synthesis is required for induction of apoptosis after CIV infection, we added cycloheximide (200 μg/ml) to CF cells 1 h prior to infection. In cells with cycloheximide, CIV infection resulted in 70% apoptotic cells as determined by blebbing assay (Fig. 7) and 60% apoptosis as determined by TUNEL assay (Fig. 8). Similar levels of blebbing (75%) were observed when CF cells were exposed to CIVUV after prior treatment with CHX (data not shown). CIV infections without inhibitor resulted in less than 10% apoptosis (Figs. 7 and 8B). Cycloheximide alone or mock infections resulted in less than 2% blebbing. These results indicate that de novo protein synthesis is not required for apoptosis induction upon infection of CF cells with CIV.

Viral early expression inhibits CIV-induced apoptosis

CIV infection induces a much lower level of apoptosis in CF cells than CIVXS, CVPE, or CIVUV (Figs. 1, 2A, 7, 8B). Moreover, inhibition of viral protein synthesis enhances CIV-induced apoptosis, suggesting that CIV gene expression is required for suppression of apoptosis. To determine if viral early gene expression blocks apoptosis induction upon CIV infection, we pretreated CF cells with aphidicolin (5 μg/ml) (D’Costa et al., 2001) or with phosphonoacetic acid (PAA; 200 μg/ml) prior to infection. Fig. 9 shows that CIV infection induced only 2% blebbing in the presence of aphidicolin and 1% blebbing with phosphonoacetic acid pretreatment compared to 10% during CIV infections of untreated cells. Thus, blebbing levels are significantly lower under conditions permitting only early gene expression compared to those allowing both early and late gene expression. Late gene expression down regulates many early genes in CIV (D’Costa et al., 2004), which probably leads to the observed 10% enhancement of blebbing. Fig. 9 also shows that inhibition of viral DNA replication in CIV infections where CIVUV or CVPE had been added as apoptosis inducers resulted in similarly drastic reductions in apoptotic blebbing. Cells challenged with CIV and CIVUV together (data not shown) manifested blebbing in 65% of the cell population, probably due to an overwhelming dose of viral proteins. Reciprocal experiments involving CVPE treatment for 1 h followed by CIV and PAA resulted in 40% blebbing and 35% cell lysis, probably due to excessive dose of viral proteins (data not shown). These
results suggest that one or more viral early gene products inhibited apoptosis induction by CIV, CVPE, or CIV_UV.

**CIV early gene expression inhibits apical caspase activity**

Since CIV early gene expression blocked apoptosis, it was important to determine the effect of this temporal phase on apical caspase activity (putative C_f-caspase-i) vital to apoptosis induction in this system (Figs. 2, 3). Therefore, we assayed C_f-caspase-i activity in CF cells at 24 h post-infection (or treatment) under conditions allowing early but not late viral expression. Fig. 10 shows that significant levels of apical caspase activity were detected in CIV infections with or without inhibition of early gene expression with cycloheximide (CHX-CIV, CIV) and in cells treated with CIV virion protein extract (CVPE). On the other hand, when phosphonoacetic acid blocked viral expression to early genes, CIV infection suppressed C_f-caspase-i activity to mock infection levels. These data strongly suggest that CIV UV-induced apoptosis requires de novo viral protein synthesis, and is dependent on apical caspase and JNK-like factors. These are the first such findings for a member of the family *Iridoviridae*.

Induction of profuse apoptotic blebbing with very high doses of CIV (400 μg/ml) probably results from virion components initiating rapid apoptosis, which preempts viral gene expression and consequent viral replication. These results are consistent with our observation that UV-irradiated CIV is capable of inducing apoptosis in CF cells. The ability of UV-irradiated cytoplasmic DNA viruses to induce apoptosis varies with virus and cell type (Hay and Kannourakis, 2002).

Low-pH-dependent endocytosis of CVPE and of UV-irradiated CIV was necessary for induction of apoptosis, implicating a requirement for cellular entry. This was further supported by preclusion of cellular entry when CIV_UV was attached to polystyrene beads and concurrent abolition of apoptotic activity (Fig. 5). Binding of CVPE and CIV_UV to polystyrene beads did not present steric hindrance problems as determined by the ability of virion-associated kinase activity to phosphorylate at least one cellular polypeptide. The requirement for cellular entry in induction of apoptosis has been shown for other cytoplasmic DNA viruses such as African swine fever virus (ASFV) (Carrascosa et al., 2002) and vaccinia virus (Ramsey-Ewing and Moss, 1998). Several viruses outside this group also require cellular entry as a prerequisite for apoptosis induction. These include *Autographa californica* nucleopolyhedrovirus (AcMNPV) (LaCount and Friesen, 1997), Sindbis virus (Jan and Griffin, 1999), human reovirus (Connolly and Dermody, 2002), and murine coronavirus (Liu et al., 2003).

The JNK pathway plays a critical role in insect immunity. JNK (coded by basket in *Drosophila*) (Riesgo-Escovar et al., 1996; Sluss et al., 1996) is activated upon infection of lepidopteran (Wojda et al., 2004) and *Drosophila* species (Kanda and Miura, 2004). In *Drosophila*, Eiger (invertebrate tumor necrosis factor; TNF) functions as a cell death ligand and binds to the TNF receptor-like protein, wengen. Eiger requires the JNK pathway and the mammalian caspase-9 homologue DRONC, which it activates (Igaki et al., 2002). JNK-mediated proapoptotic action does not always require gene expression (Lei and Davis, 2003).

Our data show that JNK and apical caspase inhibitors each block CIV-induced apoptosis in *C. fumiferana* cells, and this is
consistent with the general scheme outlined for apoptosis in *Drosophila*. Therefore, as summarized in Fig. 11, we postulate that Z-IETD-FMK inhibits a DRONC-like caspase (\(Cf\)-caspase-

i) to suppress apoptosis (Figs. 2 and 3) in CF cells. The JNK inhibitor, SP600125, blocks JNK activity and impedes DRONC-mediated apoptosis in CF cells (Fig. 4). Preliminary data (not shown) indicate that JNK inhibitor also inhibits caspase activity, suggesting that JNK is upstream of apical caspase activation. Thus, the suppression of apoptosis by inhibitors of apical caspase and JNK underscores the importance of the JNK signaling pathway in iridovirus-induced apoptosis.

An “extrinsic” pathway has been postulated for lepidopteran, *S. frugiperda* cells (IPLB-SF-21), based on the induction of apoptosis by mammalian FADD and TNFα, which utilize caspase-8 in mammalian systems (Vucic et al., 1997; Maguire et al., 2000). A putative apical caspase designated \(Sf\)-caspase-X is inhibited by Z-IETD-FMK in *S. frugiperda* cells (Manji and Friesen, 2001). Our data show that Z-IETD-FMK also blocks CVPE-induced apoptosis (Fig. 3) and suggest that CVPE-induced apoptosis in *C. fumiferana* cells is mediated through a putative apical caspase, designated *C. fumiferana* initiator caspase (\(Cf\)-caspase-i). There is some evidence that caspase-8 homologue (DREDD) and FADD homologue (DFADD) detected in *Drosophila* may be contributing to the apoptotic process in insects (Chen et al., 1998; Hu and Yang, 2000); however, this is not firmly established. On the other hand, it is well established that DRONC plays a central role in *Drosophila* apoptosis as well as metamorphosis and could well be the common conduit for all caspase-dependent apoptosis in insect systems (Walldhuber et al., 2005; Hay and Guo, 2006). It should also be noted that CVPE as well as CIV induce metamorphic arrest and mortality in the cotton boll weevil (Bilimoria, 2001).

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**Fig. 9.** Effect of early viral gene expression on apoptosis induction by CVPE in CF124T cells. Late gene expression in CIV-infected CF124T cells was blocked by inhibiting viral DNA replication with 5 \(\mu\)g/ml aphidicolin (APH) or 200 \(\mu\)g/ml phosphonoacetic acid (PAA). Viral early gene expression was permitted for 6 h after infection with CIV (10 \(\mu\)g/ml), and cells were challenged with CVPE (10 \(\mu\)g/ml) or with CVPE (10 \(\mu\)g/ml). Cells were then incubated at 21 °C and observed for apoptotic blebbing 24 h after initial treatment or infection. MOCK: untreated cells; APH/PAA: cells treated with aphidicolin or phosphonoacetic acid, respectively; CVPE: cells treated with CVPE only (A1, B1) or with CVPE only (A2, B2); CIV-APH/PAA and CVPE: cells pretreated with APH or PAA, infected with CIV, then challenged with CVPE (A1, B1) or CVPE (A2, B2) at 6 hpi. Results presented are means of 3 separate experiments in which 100 cells from 5 random fields were counted. Error bars represent standard deviations.

**Fig. 10.** Effect of CIV early gene expression on activation of apical caspase. CF124T cells were assayed for caspase-8 activity after treatment with CVPE. Incubation was at 21 °C for 24 h except where noted. Reagents and inhibitors were in RBSS at the following concentrations: CIV (10 \(\mu\)g/ml), phosphonoacetic acid (PAA; 200 \(\mu\)g/ml), cycloheximide (CHX; 200 \(\mu\)g/ml), and CVPE (10 \(\mu\)g/ml). CHX: treatment with cycloheximide alone. CHX-CIV: cells pretreated with CHX and infected with CIV compared to treatment with CHX only. PAA: phosphonoacetic acid-treated cells. CIV-PAA: cells pretreated with PAA and infected with CIV. CIV-PAA-CVPE: cells pretreated with PAA, infected with CIV, and then challenged with CVPE after 6 hpi. CIV: cells infected with CIV (10 \(\mu\)g/ml) at 24 hpi. CIV 3HPT: cells treated with CVPE and assayed at 3 h post treatment. CIV 3HPI: apical caspase assay carried out 3 h after infection with CIV compared to CVPE 3HPT. CVPE: cells treated with CVPE alone. Caspase-8 assay was carried out as described in Materials and methods. Values indicated are relative luminosity units. MOCK treatment was with RBSS only (bar and horizontal gray line). Values for all treatments at \(t=0\) were not significantly different from mock treatments. Error bars represent standard deviations for 3 independent experiments.
Given this and the relationship between the JNK pathway and DRONC in *Drosophila*, we postulate that Z-IETD-FMK suppresses apoptosis in the CIV-CF cell system by inhibiting putative DRONC-like *Cf*-caspase-i (Fig. 11).

We have provided strong evidence suggesting inhibition of apoptosis by one or more viral early gene products. CIV virion protein extract (CVPE) and UV-irradiated CIV induced strong apoptosis as observed by blebbing and TUNEL assays, but protein extract (CVPE) and UV-irradiated CIV induced strong apoptosis by one or more viral early gene products. CIV virion putative DRONC-like promotes apoptosis in the CIV-CF cell system by inhibiting gray. During CIV infection, early gene expression inhibits infection. Note: processes compromised by addition of inhibitor are indicated in gray.

During CIV infection, early gene expression inhibits infection. Fig. 11 depicts a working model incorporating our data. This is the first report showing that endocytosis and JNK-like activity are essential components of the apoptotic pathway in a member of the family Iridoviridae and that apoptosis in this virus group is suppressed by early viral expression. This is also the first report of JNK signal requirement during apoptosis induction by an insect virus.

**Materials and methods**

**Virus rearing and purification**

Chilo iridescent virus was raised in larvae of the greater wax moth *Galleria mellonella* and purified by sucrose gradient centrifugation as described previously (Henderson et al., 2001).

**Cell culture**

IPRI-CF-124T (CF) cells (Bilimoria and Sohi, 1977) from the spruce budworm *C. fumiferana* and BRL-AG-3A (AG) cells (Stiles et al., 1992) from the boll weevil *A. grandis* were cultured in Corning 25-cm² flasks using Hink’s TNM-FH medium supplemented with 10% fetal bovine serum (HyClone laboratories) and incubated at 28 °C. CF and AG cells were typically subcultured at 6-day intervals at a ratio of 1:10 (Henderson et al., 2001).

**Virus infections**

IPRI-CF-124T cells (7.5 × 10⁵ cells/ml) in 24-well plates (Corning) were inoculated with suspensions of purified CIV at 10 μg/ml in unsupplemented medium. TCID₅₀ assays developed in our laboratory indicated that 10 μg/ml of purified CIV represents 10⁹ IU/ml (D’Costa, S. M., Perales, M., and Costa et al., 2004). The first three of these ORFs are similar to *iap* genes in *Cydia pomonella* granulosis virus (CpGV; Crook et al., 1993) and have the following motifs, respectively: ring finger, BIR repeat profile, and bipartite nuclear localization signal. ORF 284R is similar to an ASV gene coding for a polypeptide (p27) with an IAP domain (Chacon et al., 1995) and is expressed exclusively at late times in CIV infections (D’Costa et al., 2004). The temporal status of ORFs 157L, 193R, and 332L are not known, but published data do not preclude an immediate-early status for ORF 193R or ORF 284R (D’Costa et al., 2004). IAP genes, dIAP 1–3, are described in *Drosophila* where they inhibit caspase-9 homologue, DRONC, and *Drosophila* tumor necrosis factor receptor-associated factor 1, dTraf1. Our data do not preclude early expression of a CIV gene product other than *iap* that inhibits apical caspase activity and either blocks apoptosis or recruits cellular factors for this purpose (Saito et al., 2006).

Taken together, our results show that, in response to entry of CIV or virion proteins, *C. fumiferana* cells undergo apoptosis through direct or indirect activation of JNK (and therefore the MAP kinase pathway) and caspases, possibly involving the putative apical caspase, *Cf*-caspase-i. However, upon infection with CIV, this response is rapidly and significantly inhibited by viral early gene expression. Excessive viral load overwhelms control mechanisms, resulting in profuse, immediate apoptosis.

**Preparation of CIV protein extract**

CHAPS [3-(3-chloroaminopropyl) dimethylamino-1-propane-sulfonate] (Sigma-Aldrich, St. Louis, MO) were used to prepare CIV protein extract (CVPE) from purified virions by a procedure modified from Cerutti and Devauchelle (1980) and described by Paul et al. (2007).
Blebbing assay

Blebbing assays were carried out as described by Paul et al. (2007). Actinomycin D (4 μg/ml) was used as positive control and heat-inactivated CVPE (65 °C for 30 min) was the negative control. Experiments were performed in triplicate and percentages of cells showing blebs were calculated.

UV-irradiated virus

CIV (2 ml at 1.6 mg/ml) was added to 60-mm² plates (Corning) and exposed to UV radiation (254 nm) using a 60-watt germicidal UV lamp (Contamination Control Inc.) at a distance of 10 cm for 45 min. UV inactivation of CIV was confirmed by testing CF cells infected with normal and UV-irradiated CIV (CIVUV) for viral DNA replication by dot blot analysis (D’Costa et al., 2001) and quantification of blots by scintillation counting. CF cells were infected with CIVUV and CIV at normal (10 μg/ml) and excess (400 μg/ml; CIVXS) doses. To determine the effect of infection with CIVUV vs. CIV, 10 μg/ml of each virus preparation was added to 24-well plates seeded with CF cells, incubated at 21 °C for 18 h and observed by phase-contrast microscopy for cytopathic effect (hypertrophy, syncytia formation, granularity or lysis) or apoptotic blebbing.

Caspase inhibitor treatments

Pancaspase inhibitor Z-VD-FMK or caspase-8 inhibitor (Z-IETD-FMK; Calbiochem, CA) was used as appropriate to inhibit caspase activities. CF cells were seeded in either 24-well or 60-well plates (Corning). Cells were incubated with pancaspase inhibitor (50 μM), caspase-8 inhibitor (50 μM), or JNK inhibitor SP600125 (25 nM) for 1 h at 28 °C. These cells were then challenged with either CVPE (10 μg/ml), CIV (400 μg/ml), or CIVUV (10 μg/ml) and incubated for an additional 24 h. The percent of cell population undergoing blebbing was monitored as described above.

Apical caspase enzyme assay

IPRI-CF-124T cells (7.5 × 10⁵ cells/ml) in 24-well plates (Corning) were inoculated with CVPE (10 μg/ml), CIV (10 μg/ml), or CIVUV (10 μg/ml) for 24 h at 21 °C. Caspase-8 assay was carried out using the Caspase-Glo Caspase-8 Assay Kit (Promega, WI) as per manufacturer’s protocol. Relative luminescence was measured with a Reporter microplate luminometer (Turner Designs, CA).

TUNEL assay

The TUNEL assay (terminal deoxynucleotidyl transferase [TdT]-mediated dUTP nick-end labeling) was carried out using the TdT-FragEL DNA fragmentation detection kit, Calbiochem (Cambridge, MA) as described by Paul et al. (2007). Five randomly selected fields were counted per slide, and percentages of TUNEL-positive cells (brown nuclei) were calculated from a sample size of at least 100 cells.

Ammonium chloride (AC) and bafilomycin A1 (Bf)-mediated inhibition of endocytosis

CF cells (6.3 × 10⁵ cells/ml complete TNM-FH) were seeded at 400 μl per well in Corning 24-well trays. Cells were pretreated with ammonium chloride (10 mM) or bafilomycin A1 (1 μM) for 30 min prior to addition of CVPE (10 μg/ml) and UV-treated CIV (10 μg/ml; see above) then incubated at 28 °C for 24 h. The concentrations of AC and Bf were maintained at 10 mM and 1 μM, respectively, through the treatment phase.

CIV replication

CF cells (6.3 × 10⁵ cells/ml complete TNM-FH) were seeded at 400 μl per well in Corning 24-well trays. Parallel sets of cells were pretreated with ammonium chloride (10 mM) or bafilomycin A (1 μM) for 30 min prior to CIV infection or not treated. Cells were incubated at 21 °C for 48 h. Hypertrophy, syncytia formation, granularity, or lysis indicated CPE. Cell monolayers were lysed with 0.5 M NaOH. Suspensions were neutralized with 10 M ammonium acetate and blotted on nitrocellulose membrane using a dot-blot apparatus (Bio-Rad). Dot blotting was carried out as described previously (D’Costa et al., 2001).

Preparation of polystyrene-linked UV-treated CIV

UV-treated CIV (100 μg/ml) was attached to 12 mg polystyrene beads (average diameter 10 μ) per manufacturer’s protocol (Polysciences, Inc., PA) except that sodium azide was omitted to prevent cell toxicity. Beads were extensively washed after conjugation with virus. Viral attachment to beads was confirmed using the assay for a kinase demonstrated in CIV particles (Monnier and Devauchelle, 1980) and recently detected in CVPE (Paul et al., 2007). The amount of CIVUV conjugated with beads was determined using the Bradford assay. Interaction of bead-bound CIV was confirmed with kinase assays on whole cells. CIVUV–bead conjugate (10 μg protein/ml) and gamma32P-ATP (10 μM) were added to cells. (ATP cannot traverse the cell membrane.) Cells were incubated at 28 °C for 1 h and washed thoroughly. Staurosporine (1 μM) and genistein (50 μM) were added to stop all kinase activity after incubation. The cell lystate was run on SDS–PAGE and analyzed by phosphorimaging. Interaction of CIVUV–bead conjugate with the cell surface was confirmed by the presence of at least one phosphorylated cellular protein. CIVUV (10 μg/ml) and CIVUV–bead conjugate (10 μg protein/ml) were added to CF cells, incubated at 28 °C for 24 h, and assayed for cell blebbing by staining for cells with Gram’s safranin for 2 min.

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