Research article

Isolation and characterization of a Solanum tuberosum subtilisin-like protein with caspase-3 activity (StSBTc-3)

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A B S T R A C T

Plant proteases with caspase-like enzymatic activity have been widely studied during the last decade. Previously, we have reported the presence and induction of caspase-3 like activity in the apoplast of potato leaves during Solanum tuberosum- Phytophthora infestans interaction. In this work we have purified and identified a potato extracellular protease with caspase-3 like enzymatic activity from potato leaves infected with P. infestans. Results obtained from the size exclusion chromatography show that the isolated protease is a monomeric enzyme with an estimated molecular weight of 70 kDa approximately. Purified protease was analyzed by MALDI-TOF MS, showing a 100% of sequence identity with the deduced amino acid sequence of a putative subtilisin-like protease from S. tuberosum (Solenomics protein ID: PGSCO0033DMP400018521). For this reason the isolated protease was named as StSBTc-3. This report constitutes the first evidence of isolation and identification of a plant subtilisin-like protease with caspase-3 like enzymatic activity. In order to elucidate the possible function of StSBTc-3 during plant pathogen interaction, we demonstrate that like animal caspase-3, StSBTc-3 is able to produce in vitro cytoplasm shrinkage in plant cells and to induce plant cell death. This result suggest that, StSBTc-3 could exert a caspase executer function during potato- P. infestans interaction, resulting in the restriction of the pathogen spread during plant–pathogen interaction.

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1. Introduction

Plant immune system is broadly divided into two interconnected defensive layers. The first layer, called as pathogen associated molecular patterns (PAMPs)- triggered immunity (PTI) involves the recognition of conserved microbial elicitors (PAMPs) by plant pattern recognition receptors (PRRs) (Boller and Felix, 2009; Chinchilla et al., 2006; McLellan et al., 2013; Muthamilarasan and Prasad, 2013; Zipfel, 2008). The activation of PRRs results in active defense responses such as production of reactive oxygen species, callose deposition and synthesis of antimicrobial compounds (Clay et al., 2009; Reina-Pinto and Yephremov, 2009). However, there are some plant pathogen microorganisms able to attenuate PTI by the secretion of proteins (effectors) that manipulate host processes inducing the effector-triggered susceptibility (ETS) (McLellan et al., 2013; Muthamilarasan and Prasad, 2013; Hof et al., 2014; Jones and Dangl, 2006; Nimchuk et al., 2003). As a consequence of this, the second defense layer, called effector-triggered immunity (ETI), is activated. This mechanism comprises plant resistance (R) proteins which detect pathogen effectors, or their activity, often resulting in a localized cell death or hypersensitive response (HR) (McLellan et al., 2013; Muthamilarasan and Prasad, 2013; Hof et al., 2014; Jones and Dangl, 2006; Nimchuk et al., 2003).

Several extracellular proteases have been associated with host immunity and described during plant–pathogen interaction. Some examples are three papain-like proteases named as: 1) Rcr3 required for C. fulvum tomato resistance inducing hypersensitive reaction cell death; 2) NbCatB, for N. benthamiana Cathepsin B, required for the development of the hypersensitive response, and 3) StC14, a potato defense papain-like cysteine protease (Shabab et al., 2008). Additionally, a positive correlation has been reported between the potato field resistance to Phytophthora infestans and the expression pattern of STAP1 and STAP3, for Solanum tuberosum (effectors) that manipulate host processes inducing the effector-triggered susceptibility (ETS) (McLellan et al., 2013; Muthamilarasan and Prasad, 2013; Hof et al., 2014; Jones and Dangl, 2006; Nimchuk et al., 2003).

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Aspartic proteases 1 and 3 (Guevara et al., 2005, 2002). In *Arabidopsis thaliana* susceptibility to *Pseudomonas* bacterial infection is enhanced by the knockdown of the pepsin-like protease CDR1 (constitutive disease resistance 1) (Xia et al., 2004). On the other hand, in *A. thaliana*, the overexpression of an extracellular subtilisin-like protein named as *AtSBT3.3* produce an increase of the plant disease resistance response to oomycete attack (Ramirez et al., 2013). Also, *SlP69B* and *C*, two tomato subtilisin proteases, are induced in tomato after pathogen infection (Rawlings and Salvesen, 2013; Tornero et al., 1997).

During the last decade, special attention of scientists has been focused on plant proteases with caspase-like activities. Although plants have no gene orthologous to caspases in their genomes, caspase-like activities had been associated with plant programmed cell death (PCD) by the activity based protein profiling (ABPP) technology (Chichkova et al., 2010; Coffen and Wolpert, 2004; Fernandez et al., 2012; Kolodziejek and van der Hoorn, 2010). In this way, two apoplastic serine dependent proteases (subtilisin-like proteases) with caspase-6 activities and related to plant PCD have been purified and described, they are named as: saspases (Coffen and Wolpert, 2004) and phytaspases (Chichkova et al., 2010). Additionally, two caspase-like activities have been related to destructive and non-destructive vacuole mediated PCD (Hara-Nishimura and Hatsugai, 2011). Destruction mechanism is initiated by the cysteine protease named as vacuolar processing enzyme (VPE) with caspase-1 activity (Hara-Nishimura et al., 2005; Hatsugai et al., 2004). The non-destructive PCD involves the 20S proteasome subunit PBA1 with caspase-3 activity (Hara-Nishimura and Hatsugai, 2011; Hatsugai et al., 2009). On the other hand, two recombinant expressed barley legumains named as *HvLeg-2* and -4 showed cysteine and caspase-like activities (Julian et al., 2013). A multifunctional role was assumed for these two cysteine peptidases, whereas *HvLeg-2* induces in leaves to biotic and abiotic stimuli, in seeds is induced by gibberellic acid, *HvLeg-4* respond in leaves to wounding and has an unknown role in the germinating seed (Julian et al., 2013). Recently, we have reported a positive correlation between apoplastic caspase-3 activity and potato field resistance to *P. infestans* infection, suggesting the induction and/or activation of apoplastic serine protease/s with caspase-3 activity during potato- *P. infestans* interaction (Fernandez et al., 2012).

In the present work, we describe the purification, identification and characterization of an apoplastic protease with caspase-3 activity from potato leaves infected with *P. infestans* (named as *StSBTc-3*). MALDI-TOF-MS identification of the isolated protein revealed that a *S. tuberosum* subtilisin like protein (Solgenomics protein ID: PGSC0003DMP400018521) is responsible of the caspase-3 activity. Additionally, we demonstrate that *StSBTc-3* is able to induce in vitro cytoplasmic shrinkage and cell death on tomato cells. These results provide new evidences about the type of proteases involved in the plant defense response mechanism during potato- *P. infestans* interaction.

![Fig. 1.](image-url) Purification steps of a caspase-3 like protease from IWF of 48 h *P. infestans* infected potato leaves. IWF of 48 h infected potato leaves was subjected to anion exchange chromatography and eluted with a linear gradient of 0–500 mM NaCl (A). Fractions with caspase-3 like activity from MonoQ eluate were subjected to size exclusion chromatography (B). Molecular weight of the isolated protease in its native state was estimated using a calibration curve for the size exclusion chromatography: (1) IgG (150 kDa); (2) BSA (67 kDa); (3) Lactoalbumin (35 kDa); Cytochrome C (12.7 kDa) and Vitamin B12 (1.355 kDa). The arrow indicates logMW from the isolated protease (C). Fractions from the size exclusion chromatography with caspase-3 like activity were pooled, desalted and analyzed by 15% SDS-PAGE (D).
2. Methods

2.1. Plant and fungal material: growth conditions

Potato plants of S. tuberosum L. cv. Pampeana INTA were grown in pots containing a 1:1 (v/v) mixture of soil/vermiculite and maintained at 18 °C for eight weeks with a 12 h photoperiod. Phytophthora infestans race R2 R3 R6 R7 R9, mating type A2, was grown on potato tuber slices of cv. Spunta at 18 °C and darkness. Seven days post-inoculation, mycelia was harvested in sterile water and stimulated to release zoospores by incubation for 2–3 h at 4 °C. After filtration through muslin, the resulting suspension was observed under light microscope to quantify spores and sporangia before being used as inoculum. Concentration was adjusted to 10^5 sporangia mL^{-1}.

Detached leaves were placed on Petri dish capsules with wet filter paper and infected by spraying with sterile distilled water (control) or a suspension containing 10^5 mL^{-1} sporangia of P. infestans. Petri dishes were placed at 18 °C and 100% humidity until leaves were harvested at 48 h post-inoculation.

Tomato cell suspensions (Lycopersicon esculentum cv. Money Maker; line Msk8) (Laemmli, 1970) were grown at 25 °C in the dark at 125 r.p.m. in MS medium (Duchefa, Haarlem, the Netherlands) supplemented with 3% (w/v) sucrose, 5.4 mM 1-naphthylacetic acid, 1 mM 6-benzyladenine and vitamins (Duchefa) (Felix et al., 1993).

2.2. Isolation of intercellular washing fluid (IWF)

Intercellular washing fluid of potato infected leaves was obtained as previously described by a standard technique based in a vacuum infiltration-centrifugation procedure (Olivieri et al., 1998). Briefly, tissue was immersed in infiltration buffer (50 mM HCl-Tris (pH 8), 3.5% (w/v) NaCl, 0.1% (v/v) 2-mercaptoethanol and 20 μL/15 mL of Tween20) and subjected to three vacuum pulses of 10 s separated by 30 s intervals. Infiltrated leaves were recovered, dried on filter paper, placed in fritted glass filters, inserted in centrifuge tubes and centrifuged for 20 min at 400 × g at 4 °C. The recovered fluid was used immediately or conserved at −20 °C. As a marker enzyme for the cytosolic and vacuolar fraction Glucose-6-phosphate dehydrogenase (EC 1.1.1.49) and α-Mannosidase (EC 3.2.1.24) activity were measured as previously described (Guevara et al., 2002).

2.3. Protein concentration

Protein concentration was determined by the bicinchoninic acid method (Smith et al., 1985) using bovine serum albumin (BSA) as
Fig. 3. Potato subtilisin-like protein with caspase-3 activity (Sسبت-3) nucleotide and deduced amino acid sequences. A. Nucleotides are numbered from the first ATG codon of the transcript. The termination codon TGA is marked with an arrow. Signal peptide and protease conserved domains are indicated in shaded boxes. Tryptic peptides obtained by MALDI-TOF-MS are underlined in the mature protein. Amino acids corresponding to the catalytic triad are indicated with an asterisk. B. Domain organization of Sسبت-3. The structure of Sسبت-3 includes a signal peptide, a prosegment, two peptidase domains and a PA domain. Amino acids from the catalytic triad Asp144/His215/Ser542 are indicated with diamonds.
2.4. Assay for specific caspase-3 like activity

Caspase-3 like activity was determined using mammalian caspase-3 substrate N-Acetyl-Asp-Glu-Val-Asp-7-amido-4-trifluoromethylcoumarin (Ac-DEVD-AMC) as substrate (Fernandez et al., 2012). Briefly, DEVcDase activity of IWFs and eluates was measured in reaction buffer composed of 50 mM sodium acetate (pH 5.2), 2 mM DTT and 15 μM Ac-DEVD-AMC dissolved in dimethylsulfoxide. Release of fluorophore by cleavage was measured using a microplate fluorimeter (Fluoroskan asent, Thermo Electron Corporation) during a 2.5 h reaction time at 37 °C (excitation and emission wavelengths of 405 and 525 nm respectively). Results were corrected against a blank containing only buffer and substrate. All assays were performed in triplicate and DMSO concentration was less than 1.5%.

For protease inhibition study, isolated protein was preincubated with 2.5 mM phenylmethanesulfonyl fluoride (PMSF) for 1 h at 37 °C.

2.5. Purification of caspase-3 like activity from IWFs

A two-step chromatographic procedure was employed to purify caspase-3 like protease from IWF of 48 h infected potato leaves. Caspase-3 substrate (Ac-DEVD-AMC) was used to monitor specific caspase-3 like activity. The experiments were performed at room temperature.

2.5.1. Ion-exchange chromatography on Mono-Q HR 5/5

Intercellular washing fluid from 48 h P. infestans infected leaves was dialyzed against 50 mM sodium acetate buffer, pH 5.2 at 4 °C using cellulose membranes (molecular weight cut off 12000Da, Sigma D9652-100FT). The resulting sample was loaded onto a Mono-Q HR 5/5 anion exchange column. The column was previously equilibrated in 50 mM sodium acetate (pH 5.2) and eluted with a linear gradient of 0-500 mM NaCl at 1 mL/min flow rate. Peaks were detected at λ = 280 nm and assayed for DEVcDase activity. Positive fractions were dialyzed against 0.05 M sodium acetate buffer pH 5.2 at 4 °C and dried in vacuum centrifuge (Savant AES 1010 Automatic Environmental Speed Vac®).

2.5.2. Size exclusion chromatography

Fractions with caspase-3 activity (corresponding to fractions: 16–18) eluted from the anion exchange chromatography were pooled and dried to 100 μL in vacuum centrifuge. The concentrated sample was loaded onto a Superose 12 10/300 size exclusion column (GE Healthcare GE, 17-5173-01) and eluted with 0.05 M sodium acetate buffer (pH 5.2). 0.3 M NaCl using a flow rate of 0.5 mL·min⁻¹. Peaks were detected at λ = 280 nm. All fractions were desalted by dialysis against 0.05 M sodium acetate buffer (pH 5.2) at 4 °C, dried in vacuum centrifuge and assayed for DEVcDase activity. Fractions with caspase-3 activity were submitted to SDS-PAGE (Laemmli, 1970).

2.6. SDS-PAGE

IWF and chromatographic fractions were analyzed by denaturing sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) using 15% (p/v) acrylamide gel (Laemmli, 1970). The samples were previously treated in denaturing buffer with β-mercaptoethanol before PAGE. Gels were stained with colloidal Coomassie Brilliant Blue G250 (Neuhooff et al., 1988).

2.7. MALDI-TOF mass spectrometry protein analysis

The protein bands obtained after several purification steps were cut and manually recovered, trypsin digested and mass spectrometric data were obtained using a MALDI-TOF-TOF spectrometer, Ultraflex II (Bruker), in the mass spectrometry facility CEQUIBIEM, Argentina (http://www.qb.fcen.uba.ar/cequibiem/). For calibration each spectrum, the following peaks of trypsin (m/z) were used: 842.51, 1045.56 and 2211.11. All the trypsin peptides were excluded as contaminants. Proteins were identified using the Mascot 1.9 search engine (Matrixscience, UK) on MSDB, NCBI, SwissProt and Solgenomics databases. A detailed analysis of peptide mass mapping data was performed using Mascot software, allowing the following parameters: specie Viridiplantae, ±50 ppm peptide mass tolerance, as well as cysteine carboxamidomethylation as possible modifications. The confidence in the peptide mass fingerprinting matches was based on the score level and confirmed by the accurate overlapping of the matched peptides with the major peaks of the mass spectrum. A threshold of 67 in Mascot score was statistically significant with p < 0.05.

2.8. Protease activity assay

Protease activity was determined with azocasein as substrate in 50 mM sodium acetate buffer (pH 5.2) and 0.5 μg of isolated protease. Samples were maintained at 37 °C for 2.5 h. Protease activity was monitored as an increase in the absorbance at 335 nm of the supernatant. The effect of the serine protease inhibitor PMSF was tested by previously incubation of the protease with 2.5 mM PMSF at 37 °C for 1 h.

2.9. Phylogenetic analysis of Solanaceae subtilisin-like proteins

Deduced amino acid sequences from StSBTc-3, tomato extracellular subtilisin-like proteins and tobacco phytaspase were retrieved by searching public databases at the Solgenomics (solgenomics.net/) and NCBI (http://www.ncbi.nlm.nih.gov/). Subcellular localization was predicted using either TargetP V1.0 (www.cbs.dtu.dk/services/TargetP®) and SignalP® (www.cbs.dtu.dk/services/SignalP®). Deduced amino acid sequences were aligned using the T-Coffee program (http://tcoffee.crg.cat/). Editing and sequence identity was calculated using the GeneDoc program (v2.6.003) (Nicholas and Nicholas, 1997). The accession numbers of the
2.10. Tomato cell suspension cell death assay

Tomato cell suspensions of 5 days old were incubated over night in 50 mM sodium acetate buffer (pH 5.2) with or without 0.1 ng/μl of SISBTc-3 or 0.1–1 ng/μl of trypsin (EC 3.4.21.4, SigmaTM) in the presence or not of the serine protease inhibitor PMSF (2.5 mM). Assays were performed on 96-well micro liter plates at 25 °C in darkness for 20 h. Subsequently, cells were washed with fresh culture medium, mounted on microscope slides and visualized using a fluorescence microscope (Nikon) with an excitation and emission filters of 480 nm and 525 nm respective. The production of green fluorescence is due to the penetration of the dye into cells with compromised plasma membrane and binding to nuclear DNA.

3. Results and discussion

3.1. Purification of a protease with caspase-3 like activity from IWF of potato leaves infected with P. infestans

We have previously reported that caspase-3 activity found in potato leaves during P. infestans infection was mainly attributable to apoplastic proteases (Fernandez et al., 2012). In order to isolate and identify the protein/s with this caspase activity, intercellular washing fluid (IWF) from potato leaves after 48 h of infection with P. infestans was analyzed by chromatographic methods.

First, IWF was subjected to size protein anion exchange liquid chromatography (MonoQ HR 5/5) and eluted with a linear gradient of NaCl (0–500 mM) as described in Materials and methods. One peak with caspase-3 activity (Pool I) was eluted at 175–200 mM NaCl (Fig. 1A).

With the aim to purify protein/s with caspase-3 activity, Pool I was subjected to size exclusion chromatography (Superose 12, GE Healthcare) with a 17–5173-01. Protein elution profile obtained from this purification step is shown in Fig. 1B. Only one protein peak with caspase-3 like activity (Pool II) was detected. Estimated molecular weight of proteins from Pool II by size exclusion chromatography was 70 kDa approximately (Fig. 1C). Subsequently, proteins pattern of Pool II were analyzed by 15% SDS-PAGE (Fig. 1D). As shown in Fig. 1D, only one protein band with an estimated molecular weight of 75 kDa approximately, was visualized. Results obtained from estimated molecular weight by size exclusion chromatography and SDS-PAGE analysis, demonstrate that, a monomeric and extracellular protease is being responsible of the caspase-3 activity determined and is induced in the potato leaves apoplast after P. infestans infection (Fernandez et al., 2012). In contrast with these results, data reported by Han et al. (Han et al., 2012) show that, caspase-3 activity detected during xylem development in Populus tomentosa is attributed to the 20S proteasome, an intracellular large protein complex. This antecedent and the results obtained in this work, demonstrate that, in plants, caspase-3 activity is involved during cell death process, either in the cell death induced after tissue differentiation (Han et al., 2012) or in the cell death induced during potato- P. infestans interaction to circumscribe pathogen dispersion (Fernandez et al., 2012). However, several differences have been found between the enzymes able to exert caspase-3 activity involved in these types of cell death. In the first case (Han et al., 2012) proteins with caspase-3 activity are heteromeric and with intracellular localization, whereas in the second case (Fernandez et al., 2012), protein with caspase-3 activity is monomeric and extracellularly located.

3.2. Identification of a subtilisin-like protease with caspase-3 activity

With the aim to identify the potato monomeric extracellular protease with caspase-3 activity, protein band obtained from the SDS-PAGE analysis of Pool II (Fig. 1D), was digested in gel by trypsin and followed by MALDI-TOF mass spectrometry analysis and subsequently MS/MS analysis. The resulting spectra were used to search for matching proteins in the NCBI database (http://www.ncbi.nlm.nih.gov) using the Mascot search program (http://www.matrixscience.com, Matrix Science, London, England) and the potato genome database Solgenomics (solgenomics.net/).

Fig. 2 shows MALDI-TOF mass spectra and the sequences of the three tryptic peptides obtained. BLASTp and tBLASTn searches against the potato genome database revealed 100% of identity with a subtilisin-like protease from S. tuberosum (Transcript and protein ID: >PGSC0003DM4000018521) respectively with an E-value of 8.5 e–04 (Fig. 3). Minor identities with higher E-values were found to the tryptic peptides with other putative proteins corresponding to subtilisin-like proteases of S. tuberosum in this database. In this way, tryptic peptides matched with a subtilisin-like protease (PGSC0003DM400012119) with an identity of 93.33% and an E-value of 5e-05, a subtilase (PGSC0003DM40001990) with an identity of 73.33% and an E-value of 0.0029, a subtilisin-like protease preproenzyme (PGSC0003DM400027005) with an identity of 68.42% and an E-value of 0.10 and a subtilisin-like protease (PGSC0003DM40064549) with an identity of 58.06% and an E-value of 0.006. Fig. 3 shows the analysis of the nucleotide and deduced protein sequences of this subtilisin-like protease from S. tuberosum.

BLASTp search of the subtilisin-like protease from S. tuberosum (ID: PGSC0003DM400018522) obtained against the MEROPS peptidase database (Rawlings et al., 2012), allowed us to classify the subtilisin-like protease as a protease belonging to the S8A subfamily (unassigned peptidase) of serine proteases. In accordance with this result, we show in Fig. 4 the inhibition by PMSF, a specific inhibitor of serine proteases, of protease and caspase-3 activities of the purified potato subtilisin-like protease.

So far, potato database contains several subtilisin-like predicted protein sequences deduced annotated as a result of the potato genome sequencing, however, there are not reports about subtilisin proteases transcripts and/or proteomics in potato. In this

Fig. 5. Alignment of SISBTc-3 deduced amino acid sequence with other Solanaceae subtilisin-like proteases. Multiple sequence alignment of the aminoacid sequence from SISBTc-3, tobacco pytaspase and tomato subtilisin like proteases was performed using the T-Coffee program. The accession numbers of the aligned protease sequences are as follows: SISBTc-3 (PGSC0003DM400018521); SISBT1 (CAA674251); SISBT2 (CAA674301); SISBT3 (NP_0012347471); SISBT4a (NP_0012347801); SISBT4b (CAA705951); SISBT4c (CAA706061); SISBT4e (CAA706211); SPl69A (CAA677241); SPl69B (CAA677251); SPl69C (CAA64121); SPl69D (CAA677271); SPl69E (CAA604131); SPl69F (CAA604141); SITmp (AAB387431) and NPhytaspase (ACT347641). Conserved residues which are common to all sequences are shadowed in black; less identity is shadowed in gray scale.
report we show the purification and identification of a *S. tuberosum* subtilisin-like protease named as *StSBTc-3* (subtilisin-like protein with caspase-3 activity) involved in plant cell death during potato-*P. infestans* interaction. Other plant subtilisin-like proteases with caspase-6 activities have been purified and identified, namely, saspas (Coffen and Wolpert, 2004) and phytaspases (Chichkova et al., 2010), however neither of them were able to cleave DEVD, a specific substrate of caspase-3.

### 3.3. Sequence analysis of *StSBTc-3*

Most SBTs from plants and other organisms show the typical pre-pro-protein structure of secretory proteins, comprising a signal peptide at the N-terminus, a cleavable prodomain, a subtilisin-like mature peptide, a protease associated domain (PA) (Schaller, 2004). According with the data above described, the *StSBTc-3* predicted amino acid sequence contains a signal peptide, a propeptide and a mature peptide of 659 aminoacids (Fig. 3). Consistent with the estimated molecular weight of 70 kDa obtained by size exclusion chromatography and SDS-PAGE (Fig. 1C and D), molecular weight prediction of SDS-PAGE (Fig. 3C and D), molecular weight prediction of *StSBTc-3* deduced amino acid sequence was of 70.5097 kDa for the mature protein (Solgenomics ID: PGSC0003DMP400018521).

SignalP and TargetP program predictors of subcellular location of eukaryotic proteins predicted a cleavage site between positions 22-23 of the full length pre-pro-protein. This result suggests the presence of a secretory signal peptide of 22 aminoacids (Met¹-Ala²²) consistent with *StSBTc-3* extracellular localization previously reported (Fernandez et al., 2012). Bioinformatic analysis performed with CD-search (NCBI) and Pfam (Sanger Institute) programs, suggested the presence of four domains highly conserved in precursors of subtilisin-like serine proteases: a peptidase inhibitor I9 domain (Thr¹⁰⁵-His¹¹², E-value: 5.68e-15), a peptidase S8 family domain (Val¹⁰⁹-Ser¹³⁵, E-value: 7.60e-100), a protease associated (PA) superfamily domain (Thr¹⁴⁷-Ile¹⁷⁰, E-value: 1.37e-06) and a peptidase S8_S53 superfamily domain (Lys¹⁵²-Ser¹⁷⁰, E-value: 4.77e-29) (Fig. 3B).

We performed a phylogenetic analysis in order to identify putative subtilisin-like proteases of Solanaceae with caspase-like activity (Chichkova et al., 2010; Meichtry et al., 1999). So, we compared the amino acid deduced sequences of *StSBTc-3* with fifteen amino acid sequences of subtilisin-like proteases previously reported to Solanaceae species, two of them, *StSBTc-3* and tobacco phytaspase, with caspase-like activity.

Comparison of the *StSBTc-3* deduced amino acid sequence with GenBank and EMBL databases reveals a high degree of conservation with other subtilisin-like proteins from other Solanaceae species (Fig. 5). *StSBTc-3* has the highest sequence identity with tomato SBT4b (62%) (CAA07059.1), tomato SBTc-3 (CAA07060.1) and tobacco phytaspase (ACT34764.1) (46%), tomato SBTc-3 and tobacco phytaspase (ACT34764.1) (46%), tomato SBTc-3 and tobacco phytaspase (ACT34764.1) (46%), tomato SBTc-3 and tomato meiotic proteinase Tmp (AAAB38743.1) (28%) (Fig. 3).

![Fig. 6. Phylogenetic relationship of Solanaceae subtilisin-like proteases. The unrooted phylogenetic tree was created from the SBTc-3, tomato subtilisin-like proteases and tobacco phytaspase amino acid sequences annotated in the Solgenomics (www.solgenomics.net) database. Aminoacid multiple sequence alignment of the proteases: SSBTc-3 (PGSC0003DMP400018521), SSBT1 (CAA67429.1), SSBT2 (CAA67430.1), SSBT3 (NP_001234774.1), SSBT4a (NP_001234780.1), SSBT4b (AAAT059.1), SSBT4c (AAAT060.1), SSBTc-3 (CAA67429.1) and SSBTc-3 (CAA67430.1) (60%). Lower identities were detected with tobacco phytaspase (ACT34764.1), tomato SBT1 (CAA67429.1) and 2 (CAA67430.1) (35%), the tomato P69 group (SIP69A (CAA67724.1); SIP69B (CAA67725.1); SIP69C (CAA64121.1); SIP69D (AAAS767271.1); SIP69E (AAAS64131.1); SIP69F (AAAS64141.1)) (35%) and the tomato meiotic proteinase Tmp (AAAS64141.1) (28%) (Fig. 5).](image-url)
Data obtained from the phylogenetic analysis of StSBTc-3 (Fig. 6) show that despite this protein has higher similarity with SSBT3 and lower with SSBT1 (Figs. 5 and 6), it shares more structural properties with SSBT3 than with SSBT3. While SSBT3 has a fibronectin III-like domain as a C-terminal extension, SSBT1 and StSBTc-3 do not (Figs. 3 and 5). Also, tomato SSBT3 has been described to form homodimers through its PA domain. However, results obtained here showed that, as tomato SSBT1, StSBTc-3 is a monomeric enzyme in its active and mature form (Fig. 1C and D). Although SSBTc-3 is closer related to tomato SSBT3 than to tobacco phytophastase it shares more structural and activity specificities with the tobacco phytophastase (Fig. 6). As well as phytophastase (Chichkova et al., 2010, 2004), potato StSBTc-3 shows caspase activity and is related to the PCD that occurs in the hypersensitive response during plant–pathogen interactions. Data presented here suggest that there are both, structural similarities and identity in the amino acid sequences between subtilisin-like proteases of Solanaceae species. This fact suggests that will be necessary to perform assays to test the potential caspase-like activity of these proteases (SSBT1 and SSBT3) one by one.

3.4. StSBTc-3 ability to induce cell death in tomato cell culture

Caspases in mammalian PCD play a role as initiators and effectors (Boatright and Salvesen, 2003). Caspases –2, –8, –9 and –10 have been classified as initiator caspases, and they cleave inactive pro-forms of effector caspases, thereby activating them. On the other hand, caspases –3, –6 and –7 have been classified as effector caspases as they cleave other protein substrates within the cell, to trigger the apoptotic process and ultimately leading to cell death (Green and Reed, 1998; Los et al., 2001). According with the previously demonstrated in several reports (Rotari et al., 2004; Thornberry and Lazebnik, 1998), we have demonstrated that caspase-3 activity is required for completion of cell death induced in potato leaves to restrict the pathogen spread, during potato-P. infestans interaction in resistant potato cultivars (Fernandez et al., 2012). In order to determine if StSBTc-3 is able to induce cell death in plant cells, 0.1 ng/μl of the subtilisin-like protease or buffer (control) were incubated with tomato cell cultures for 20 h in the previously described conditions prior to SYTOX Green treatment. Results obtained (Fig. 7) show that StSBTc-3 was able to decrease tomato cell viability in a 75% during 20 h of incubation. Additionally, we observed cytoplasm shrinkage as an effect of the cells incubation with active StSBTc-3 (Fig. 7A). We have previously reported that StSBTc-3 caspase-3 activity is inhibited by PMSF (Fernandez et al., 2012). According with this antecedent, no changes in the tomato cell culture viability were observed when StSBTc-3 was previously incubated with 2.5 mM PMSF (Fig. 7B) compared to control. On the other hand, when tomato cells were incubated with 0.1 and 1 ng/μl of trypsin, no effect on the tomato cell viability was observed. These results suggest that StSBTc-3 cell death induction ability is related with the caspase-3 activity exerted by this protease. This observation is in accordance with the effects produced by executor caspases in animals, where caspase-3 is believed to be the major executioner to induce the cleavage of the PARP (Poly ADP-ribose polymerase), DNA fragmentation, cytoplasmic shrinkage, chromatin condensation, and final programmed death in animal cells (Thornberry and Lazebnik, 1998; Budihardjo et al., 1999; Cohen, 1997).

Fig. 7. In vitro effect of StSBTc-3 on tomato cell suspensions. Tomato cell suspensions were incubated in 50 mM sodium acetate buffer (control) with or without 0.1 ng/μl of protease or 0.1 ng/μl of trypsin, in the presence or not of the serine protease inhibitor PMSF (2.5 mM) for 20 h at 25 ℃. Subsequently, cells were incubated with the vital probe SYTOX Green for 10 min and examined using a fluorescence microscope. A. Bright-field (left panels) and fluorescence images (right panels) are shown. Bar: 25 μm. B. Dead cell quantification. Values were normalized as a percentage of the mean of protease treated cells. The data represents the mean ± SD of three replicates. Different letters indicate significant differences with p < 0.05 for one way ANOVA and Tukey tests. Abbreviations: pm, plasma membrane; cw, cell wall; and n, nuclei. The arrow indicates cytoplasm shrinkage.

4. Conclusions

In this work, we described the purification of a S. tuberosum subtilisin-like protease with caspase 3 activity (StSBTc-3) induced in potato leaves after P. infestans infection. Results obtained from size exclusion chromatography demonstrate that StSBTc-3 mature enzyme is monomeric with a molecular weight of 70 kDa. Analysis
of deduced amino acid sequence shows that the potato subtilisin like protein here purified presents the typical pro-pro-protein structure of secretory proteins, as well as most subtilisin like pro-
teases from plants previously reported. SSBT3c-3 present a highest
amino acid sequence identity with subtilisin like proteins identified in other Solanaceae. However, whereas SSBT3c-3 is phylogenetic
conserved related with tomato SSBT3, it shares more structural char-
teristics with tomato SSBT1. Therefore, this report constitutes the first evidence of isolation of a potato apoplastic subtilisin like protein with caspase-3 like activity which may help to complete the family of “caspase-like” proteases, previously considered as “incomplete”. Additionally, we demonstrate that caspase-3 enzym-
atic activity of SSBT3c-3 is able to induce plant cell death. Results presented here support the hypothesis that, SSBT3c-3 is an executor caspase induced in the apoplast of potato leaves of resistant culti-
vars after infection with P. infestans, causing cell death and as consequence of this, restriction of pathogen spread.

Contributions
FMB performed the assays, drew the figures and analyzed the data. GMG designed the research and the manuscript, and analyzed the data. GMG and FMB wrote the manuscript. GRD revised the manuscript and checked English language.

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