

Antifungal activity of the ribosome-inactivating protein BE27 from sugar beet (*Beta vulgaris* L.) against the green mould *Penicillium digitatum*

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SUMMARY

The ribosome-inactivating protein BE27 from sugar beet (*Beta vulgaris* L.) leaves is an apoplastic protein induced by signalling compounds, such as hydrogen peroxide and salicylic acid, which has been reported to be involved in defence against viruses. Here, we report that, at a concentration much lower than that present in the apoplast, BE27 displays antifungal activity against the green mould *Penicillium digitatum*, a necrotrophic fungus that colonizes wounds and grows in the inter- and intracellular spaces of the tissues of several edible plants. BE27 is able to enter into the cytosol and kill fungal cells, thus arresting the growth of the fungus. The mechanism of action seems to involve ribosomal RNA (rRNA) *N*-glycosylase activity on the sarcin–ricin loop of the major rRNA which inactivates irreversibly the fungal ribosomes, thus inhibiting protein synthesis. We compared the C-terminus of the BE27 structure with antifungal plant defensins and hypothesize that a structural motif composed of an α -helix and a β -hairpin, similar to the γ -core motif of defensins, might contribute to the specific interaction with the fungal plasma membranes, allowing the protein to enter into the cell.

Keywords: apoplast, defensin, green mould, plant defence, pokeweed antiviral protein (PAP), polynucleotide:adenosine glycosylase, rRNA *N*-glycosylase.

INTRODUCTION

Plants have developed protein-based defences against infections by fungi and other pathogens. Some of these proteins are common and have been classified into 17 families of pathogenesis-related (PR) proteins, and others occur specifically in some plant species (van Loon *et al.*, 2006). Most are induced by signalling compounds, such as salicylic acid, jasmonic acid, ethylene or hydrogen peroxide, and display antimicrobial activities *in vitro* through different enzymatic activities (van Loon *et al.*, 2006).

The plant apoplast plays a crucial role in plant defence as it contains proteins that trigger the plant defence response and inducible enzymes that are toxic to fungi, such as proteases, chitinases, glucanases, reactive oxygen species (ROS)-generating enzymes and defensins (Doehlemann and Hemetsberger, 2013).

The apoplast of sugar beet (*Beta vulgaris* L.) leaves also contains the ribosome-inactivating proteins (RIPs) BE27 and BE29 which represent different levels of glycosylation of the same protein (Iglesias *et al.*, 2005). These proteins are synthesized in response to virus infection (Girbes *et al.*, 1996) and are expressed in both infected (hypersensitive response) and non-infected (systemic acquired resistance, SAR) leaves (Iglesias *et al.*, 2005). Furthermore, the signalling compounds hydrogen peroxide and salicylic acid induce the expression of both proteins (Iglesias *et al.*, 2005, 2008). The hypothesis that BE27 is involved in plant defence against viruses is partially supported, although in non-physiological conditions, by the fact that the external application of BE27 to sugar beet leaves prevents infection by *Artichoke mottled crinkle virus* (AMCV) and because this protein displays polynucleotide:adenosine glycosylase activity against *Tobacco mosaic virus* (TMV) genomic RNA (gRNA) (Iglesias *et al.*, 2005).

RIPs are protein synthesis inhibitors that irreversibly inactivate mammalian ribosomes (Girbes *et al.*, 2004; Puri *et al.*, 2012). RIPs can also inhibit protein synthesis in other animals and yeasts (Girbes *et al.*, 2004), and some can inactivate plant and bacterial ribosomes, although usually these ribosomes are insensitive to the majority of RIPs (Girbes and Ferreras, 1998). These proteins are 28S ribosomal RNA (rRNA) *N*-glycosylases (EC 3.2.2.22) which cleave the *N*-glycosidic bond between adenine at position 4324 and its ribose in the 60S subunit of rat ribosomes. This adenine is located in the sarcin–ricin loop (SRL), which is crucial for anchoring the EFG or EF-2 elongation factors on the ribosome during messenger RNA–transfer RNA (mRNA–tRNA) translocation in prokaryotes and eukaryotes, respectively (Girbes *et al.*, 2004; Puri *et al.*, 2012). In addition, some RIPs release adenines from viral gRNAs (Barbieri *et al.*, 1997; Iglesias *et al.*, 2005) and herring sperm DNA (Barbieri *et al.*, 1997; Iglesias *et al.*, 2015). Thus, Barbieri *et al.* (1997) have proposed the name polynucleotide:adenosine glycosylases for these proteins and have suggested that extensive deadenylation of viral RNA and DNA might be responsible for the antiviral action of

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RIPs. From a structural point of view, RIPs have been classified into two types: type 1 [e.g. saporin, pokeweed antiviral protein (PAP), BE27 and ME] consisting of single-chain proteins, and type 2 (e.g. ricin, abrin and ebulin) consisting of an A (active) chain with RIP properties covalently linked to a B (binding) chain with lectin properties. The latter RIPs can enter cells more easily because the B chain allows binding to sugar-containing cell surface receptors and, for this reason, type 2 RIPs can be potent toxins, e.g. ricin, abrin or volkensin (Citores *et al.*, 2013; Ferreras *et al.*, 2011a). Type 1 RIPs from monocots lack the leader peptide (Di Maro *et al.*, 2014) and, for this reason, these proteins are synthesized and localized in the cytosol (Chaudhry *et al.*, 1994; Lanzanova *et al.*, 2011). By contrast, type 1 RIPs from dicots and type 2 RIPs contain a leader peptide (Di Maro *et al.*, 2014), thus being synthesized in the endoplasmic reticulum. Several studies have indicated that type 1 RIPs from dicots are localized exclusively (Di Cola *et al.*, 1999; Iglesias *et al.*, 2005; Park *et al.*, 2004; Yoshinari *et al.*, 1996) or mainly (Carzaniga *et al.*, 1994; Parente *et al.*, 2008; Yoshinari *et al.*, 1997) in the apoplast, whereas type 2 RIPs are localized in protein storage vacuoles (Lord and Spooner, 2011).

RIPs have been extensively researched because of their ability to inhibit protein synthesis and their utility as antiviral (Fang *et al.*, 2011) and antitumour (Pan *et al.*, 2014) agents. The most promising applications of RIPs in experimental medicine, especially in anticancer therapy, are related to their use as components of immunotoxins, conjugates or recombinant chimeras in which they are linked to tumour-targeting ligands or antibodies which mediate their binding and internalization by malignant cells (Becker and Benhar, 2012; Citores *et al.*, 2013; Ferreras *et al.*, 2011a; Polito *et al.*, 2011). However, some RIPs behave as defence proteins as they are induced by salt stress (Rippmann *et al.*, 1997), insect elicitors (Engelberth *et al.*, 2012), mechanical wounding (Engelberth *et al.*, 2012; Song *et al.*, 2000), pathogenic fungus (Tan *et al.*, 2013) and temperature and ultraviolet light (Qin *et al.*, 2009). In addition, some are induced by signalling compounds, such as jasmonic acid (Chaudhry *et al.*, 1994; Song *et al.*, 2000), salicylic acid (Qin *et al.*, 2009) or abscisic acid (Song *et al.*, 2000). Furthermore, several studies have indicated that RIPs protect plants from viral (Park *et al.*, 2004; Tumer *et al.*, 1997) and fungal (Park *et al.*, 2002) infection, and transgenic plants bearing RIP genes have been constructed that are resistant to viruses, fungi and insects (Corrado *et al.*, 2005; Dowd *et al.*, 2012; Huang *et al.*, 2008; Qian *et al.*, 2014).

Recently, we have tested other activities of BE27 that could play a defensive role against pathogens (Iglesias *et al.*, 2015), finding that BE27 displays rRNA *N*-glycosylase activity against yeast and *Agrobacterium tumefaciens* ribosomes, DNA polynucleotide:adenosine glycosylase activity against herring sperm DNA, and magnesium-dependent endonuclease activity against the supercoiled plasmid PUC19 (nicking activity). In addition, BE27 possesses superoxide dismutase (SOD) activity, thus

being able to produce the signal compound hydrogen peroxide. BE27 is also toxic to COLO 320 cells, inducing apoptosis in these cells. The combined effect of these biological activities could result in a broad action against several types of pathogen, such as viruses (polynucleotide:adenosine glycosylase activity), bacteria (rRNA *N*-glycosylase and nicking activities), fungi (rRNA *N*-glycosylase activity) and insects (via the induction of gut cell apoptosis) (Iglesias *et al.*, 2015).

In order to gain an insight into the protective properties of BE27 against pathogenic fungi, we have carried out experiments to test whether this protein is able to enter the fungal cytoplasm and arrest the growth of the fungus *Penicillium digitatum*, which infects a broad spectrum of edible plants, such as kiwifruit, sugarbeet, apple, plum, spinach and, mainly, citrus (<http://www.plantwise.org/KnowledgeBank/Datasheet.aspx?dsid=39570>) (Frisvad and Samson, 2004). At concentrations lower than those found in the apoplast, BE27 was able to enter the cytosol and kill fungal cells, thus arresting the growth of the fungus. The mechanism of action seemed to involve rRNA *N*-glycosylase activity on the SRL of the large-subunit rRNA which inactivates irreversibly the fungal ribosomes, thus inhibiting protein synthesis. In addition, we found that, close to its C-terminus, BE27 contains a structural motif composed of an α -helix and a β -hairpin similar to that present in antifungal defensins. Such a β -hairpin has been reported to be responsible for the interaction of defensins with the fungal plasma membrane, allowing these apoplastic proteins to enter into fungal cells (Lacerda *et al.*, 2014; Sagaram *et al.*, 2013).

RESULTS

Effect of BE27 on the growth of *P. digitatum*

To characterize the antifungal properties of BE27, we carried out experiments to test the effect of this protein on the growth of the fungus *P. digitatum*. As illustrated in Fig. 1, BE27 showed a strong growth-inhibiting effect on *P. digitatum* and reduced fungal growth in a concentration-dependent manner. Thus, 8, 4 and 0.6 $\mu\text{g/mL}$ of BE27 resulted in 70%, 57% and 24% growth inhibition, respectively, after 78 h of incubation with conidia as starting material. In addition, 8 $\mu\text{g/mL}$ of BE27 added to mycelia growing for 28 h inhibited fungal growth by 84%, suggesting that BE27 has a stronger effect on mycelial growth than on conidial germination. These concentrations are lower than those reached by BE27 and BE29 in the apoplast in *B. vulgaris* leaves. To estimate the concentration of BE27 and BE29 present in the apoplast, we studied the translation inhibitory activity of *B. vulgaris* crude extracts from 6-week-old plants untreated or treated with either hydrogen peroxide or salicylic acid, and 7-month-old plants collected in a crop field (Table 1). These data must be evaluated with care as they may be overestimated or underestimated because of the presence of other active substances in the crude extracts. The crude extract

obtained from 7-month-old plants was the least active in inhibiting protein synthesis in rabbit reticulocyte lysate. Such an extract contained 24 781 units of BE27 and BE29 per gram of tissue. From this preparation, we obtained 4.6 µg of BE27 and 7.3 µg of a mix containing approximately equal amounts of BE27 and BE29, for a total of 12 395 units/g of tissue, thus obtaining a purification yield of 50%. Considering 60 µL per gram of fresh weight as the volume of apoplastic fluid (Lopez-Millan *et al.*, 2000; Winter *et al.*, 1994), the concentration of BE27/29 must be at least 200 µg/mL, much higher than that required to stop the growth of *P. digitatum*. The concentration was even higher in 6-week-old plants grown in a chamber, and increased two- and seven-fold in hydrogen peroxide- and salicylic acid-treated plants, respectively (Table 1). It has been shown previously that beetin expression is greatly enhanced on treatment with the signalling compounds hydrogen peroxide and salicylic acid (Iglesias *et al.*, 2005). Thus, the apoplast, which is the

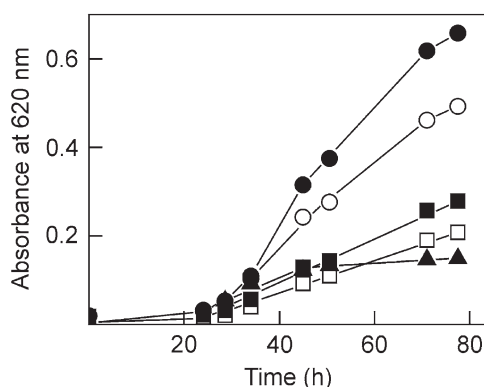


Fig. 1 Antifungal activity of BE27 against *Penicillium digitatum*. Antifungal activity of BE27 against *P. digitatum* was measured in a microtitre plate bioassay. Conidia of *P. digitatum* were grown in potato dextrose broth (PDB) medium in the presence of different concentrations of BE27. Fungal growth was measured as the increase in absorbance at 620 nm. The curves represent buffer control (filled circles), 0.6 µg/mL BE27 (open circles), 4 µg/mL BE27 (filled squares) and 8 µg/mL BE27 (open squares). Conidia of *P. digitatum* were grown in PDB for 28 h before exposure to 8 µg/mL BE27 (filled triangles).

Table 1 Quantification of BE27 and BE29 activity in crude extracts from *Beta vulgaris* leaves. Crude extracts were obtained as indicated in the Experimental procedures section. The values are referred to 1 g of fresh tissue.

Crops	Protein (µg/g of tissue)	IC ₅₀ * (ng/mL)	Specific activity† (units/mg of protein)	Total activity† (units/g of tissue)	Yield‡ (%)
Control	18500	310	3226	59677	—
Hydrogen peroxide-treated	22400	211	4739	106161	—
Salicylic acid-treated	18100	39.8	25125	454773	—
Crop field	22080	891	1122	24781	(100)
BE27	4.6	0.96	1041666	4791	50.0
BE27/29	7.3	0.96	1041666	7604	—

*IC₅₀ is the concentration giving 50% inhibition of protein synthesis in a rabbit reticulocyte lysate system.

†One unit of activity is defined as the amount of protein necessary to inhibit protein synthesis by 50% in 1 mL of rabbit reticulocyte lysate reaction mixture.

‡The yield is referred to the purification of BE27 and BE27/29 from 300 g of leaves from a crop field.

first cellular compartment to be invaded in the infection process (Doehlemann and Hemetsberger, 2013), contains sufficient BE27 to arrest fungal growth.

Microscopic analysis and BE27 uptake by *P. digitatum*

The effect of BE27 on the mycelial growth of *P. digitatum* was also visualized microscopically (Fig. 2). Corresponding to growth inhibition, microscopy studies revealed modifications of hyphal morphology after exposure to BE27. Compared with the untreated control, which presented regular and homogeneous hyphae, hyper-branching and aborted hyphal branches could be observed in the cultures treated with BE27 (Fig. 2a, left panels). Moreover, chitin staining with calcofluor white (CFW) revealed that the treatment with BE27 inhibited hyphal elongation, rendering cells significantly shorter than control cells (Fig. 2a, right panels).

It has been reported that *Trichosanthes kirilowii* Maxim plant cell cultures secrete three bifunctional plant enzymes with both rRNA *N*-glycosylase and chitinase activities (Shih *et al.*, 1997). Moreover, sugar beet leaves contain chitinases that can be induced by pathogen infection (Berglund *et al.*, 1995). To exclude the possibility that chitinase activity, either by contamination or as a consequence of an intrinsic activity of BE27, may be responsible for fungal growth inhibition, we performed experiments to assess whether our BE27 samples displayed such activity. We also tested samples with known chitinase activity, such as crude extracts from non-induced and signalling compound-induced sugar beet leaves, as well as the chitinase from *Streptomyces griseus*. The results indicated that no such activity was present in BE27 samples in standard conditions, in which the other mentioned samples gave values corresponding to the presence of chitinase activity as expected (data not shown).

Some antifungal apoplastic proteins, such as chitinases or glucanases, act outside the fungus cells (Doehlemann and Hemetsberger, 2013); however, in order to deplete fungal ribosomes, BE27 must enter into the cytosol. In order to study the interaction of BE27 with fungal cells, the location of fluorescent-labelled BE27 in treated hyphae was monitored by confocal microscopy. In these conditions, cyanine 3 (Cy3)-labelled BE27

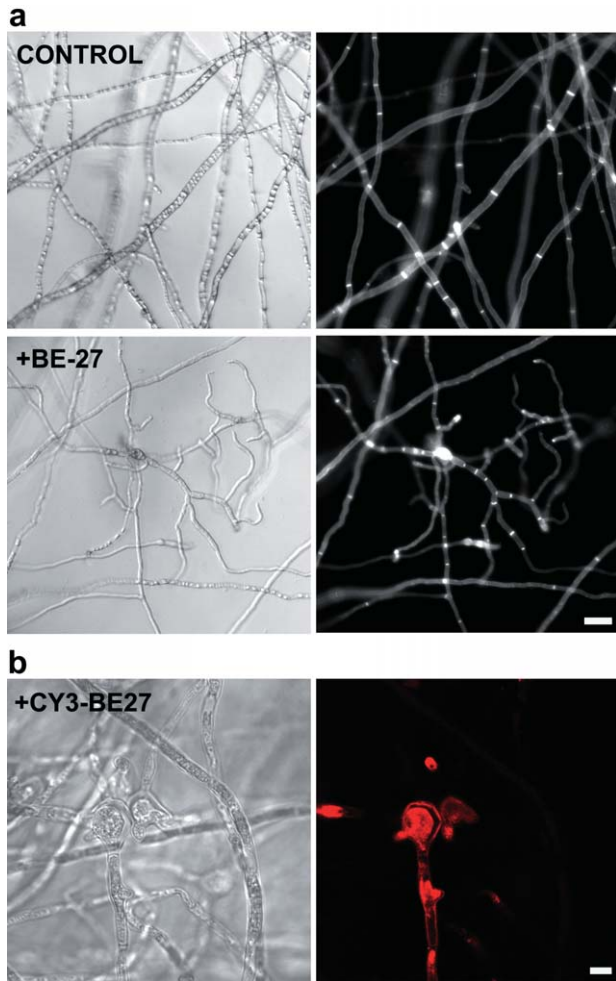


Fig. 2 (a) Morphological changes in *Penicillium digitatum* mycelium exposed to BE27. *Penicillium digitatum* mycelium was grown in the absence (control) or presence of 5 µg/mL of BE27. After 24 h of incubation, samples were stained with calcofluor white (CFW) and visualized under the microscope. Panels represent bright-field images (left) and fluorescence images indicative of CFW staining (right) for the same fields. Bar, 20 µm. (b) Interaction between BE27 and *P. digitatum*. To visualize the interaction of BE27 with fungal structures, conidia of *P. digitatum* were grown *in vitro* in potato dextrose broth (PDB) medium for 24 h before exposure to 5 µg/mL of CY3-BE27 for 24 h. After washing, the samples were visualized with a confocal laser microscope. The red channel image (right panel) and the corresponding interference contrast image (left panel) are shown. Bar, 5 µm.

retained full antifungal activity (data not shown). On incubation with 5 µg/mL of BE27 for 24 h, BE27 accumulated in the cytoplasmic space of terminal branches of *P. digitatum* hyphae (Fig. 2b).

rRNA glycosylase activity of BE27 on the *P. digitatum* ribosomes

Because the rRNA *N*-glycosylase activity of BE27 may play an antifungal role in sugar beet, we assayed the effect of BE27 on

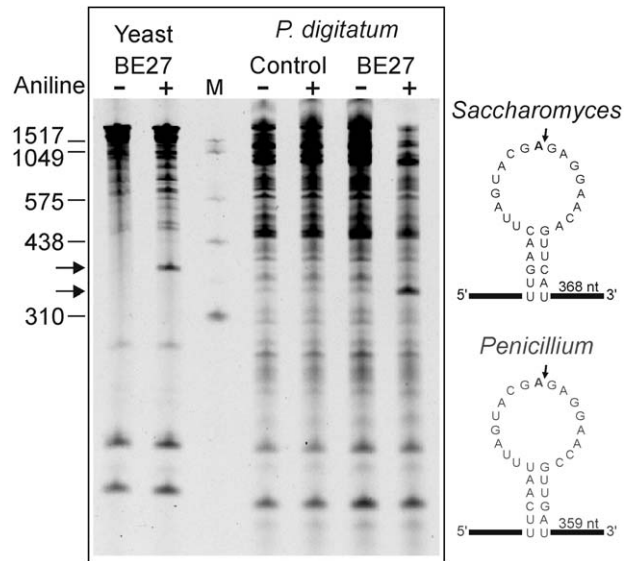


Fig. 3 rRNA *N*-glycosylase activity of BE27 on yeast and *Penicillium digitatum* ribosomes. Left: rRNA *N*-glycosylase activity was assayed as indicated in Experimental procedures. Each lane contained 3 µg of RNA isolated from either untreated (control) or BE27-treated ribosomes from yeast or *P. digitatum*. The arrows indicate the RNA fragments released as a consequence of ribosome-inactivating protein (RIP) action on acid aniline treatment (+). Numbers indicate the size of the standards (M) in nucleotides. Right: sarcin-ricin loop of the large rRNA from yeast and *Penicillium*. The sequences from *Saccharomyces cerevisiae* (accession number J01355) and *Penicillium solitum* (JN642222) were downloaded from the National Center for Biotechnology Information (NCBI) sequence database (<http://www.ncbi.nlm.nih.gov/nucleotide/>). The large rRNA 3' end from *Penicillium* was determined by alignment with the large rRNA from *Saccharomyces*. The adenine released by RIP action (bold type), the site of splitting by the acid aniline (arrow) and the size of the generated fragment are also indicated.

ribosomes from *P. digitatum*. As shown in Fig. 3, BE27 displayed rRNA *N*-glycosylase activity on these ribosomes, as indicated by the release of the diagnostic fragment (Iglesias *et al.*, 2015) on treatment with acid aniline. For comparative purposes, Fig. 3 also shows the rRNA *N*-glycosylase assay on yeast. The released fragments displayed sizes around 380 and 350 nucleotides for yeast and *P. digitatum*, respectively, in accordance with that expected for SRL deglycosylation (368 and 359 nucleotides for yeast and *Penicillium*, respectively; Fig 3). Therefore, this RIP might enter into the fungal cells and inactivate their ribosomes, avoiding the propagation of the pathogen, as has been reported for ME, a type 1 RIP from *Mirabilis expansa* (Ruiz & Pav.) Standl. (Park *et al.*, 2002).

Enzymatic activity of BE27 in *P. digitatum* cultures

BE27, at concentrations of 4 µg/mL, showed a strong growth inhibitory effect on *P. digitatum* cultures in liquid medium (Figs 1

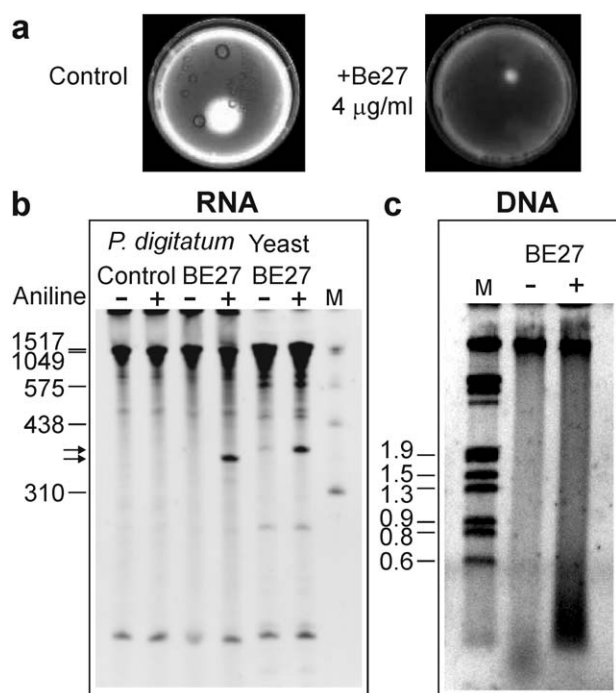


Fig. 4 Antifungal and rRNA *N*-glycosylase activity of BE27 against *Penicillium digitatum*. (a) *Penicillium digitatum* was grown in potato dextrose broth (PDB) in the absence (control) or presence of 4 µg/mL BE27 for 5 days. The mycelium was extensively washed with sterile water and harvested to extract the RNA and DNA. Representative photographs of two plates are shown. (b) rRNA *N*-glycosylase activity was assayed as indicated in Experimental procedures. Each lane contained 3 µg of RNA isolated from either untreated (control) or BE27-treated cultures from *P. digitatum*. For comparative purposes, yeast RNA depurinated by BE27 is also included in the assay. The arrows indicate the RNA fragments released as a consequence of ribosome-inactivating protein (RIP) action on acid aniline treatment (+). Numbers indicate the size of the standards (M) in nucleotides. (c) The DNA was isolated as indicated in Experimental procedures and 3 µg was electrophoresed. The numbers indicate the corresponding size of the standards (λ DNA *HindIII/EcoRI*) in Kb.

and 4a). The RNAs from these cultures were isolated and analysed to detect the presence of the RNA fragment diagnostic of RIP rRNA *N*-glycosylase (Fig. 4b). The diagnostic fragment was absent in the RNA from control cultures and present in that from cultures grown in the presence of BE27. This indicates that BE27, at a concentration of 4 µg/mL, is able to enter into fungal cells in a manner that allows it to reach the cytosolic ribosomes.

Initially, the toxicity of RIPs to animal cells was attributed to their ability to inhibit protein synthesis leading to cell death, but there is an increasing body of evidence indicating that the main cause of RIP toxicity in these cells is their ability to induce apoptosis (Das *et al.*, 2012). COLO 320 cells treated with BE27 exhibited the morphological features characteristic of apoptosis, strong dose-dependent activation of the effector caspase 3/7 and the breakdown of the nuclear DNA into oligonucleosomal frag-

ments (Iglesias *et al.*, 2015). DNA from *P. digitatum* was isolated from the cultures shown in Fig. 4a and analysed to detect the presence of the oligonucleosomal fragments (Fig. 4c). Although a smear of degraded DNA was observed in BE27-treated fungi compared with the untreated control, no internucleosomal cleavage was visible, suggesting that BE27 intoxication can occur by non-apoptotic mechanisms.

In silico structural analysis of BE27

It has been suggested that a β -hairpin present in domain 3 of the dicot type 1 RIPs may be necessary for the toxicity of such proteins (Di Maro *et al.*, 2014). To ascertain the main structural characteristics that may be involved in the antifungal behaviour of BE27, a three-dimensional structure was predicted by comparative modelling using several RIP crystal structures as templates. The selected best model was found to have a confidence score (C-score) of 1.24, template modeling (Tm) score of 0.89 ± 0.07 and root-mean-square deviation (RMSD) of 3.3 ± 2.3 Å, which satisfied the range of parameters for molecular modelling. Figure 5a shows such a model that displays the characteristic fold of type 1 RIPs and the A chain of type 2 RIPs, in which the confluence of three domains creates the active site: N-terminal domain 1, consisting of β -strands and α -helices; domain 2, characterized by α -helices; and C-terminal domain 3, formed by two α -helices and two β -strands (Di Maro *et al.*, 2014). Although the models obtained by comparative modelling have a limited value and should be confirmed by X-ray diffraction studies, we used such a model to compare the structure of BE27 with some of the well-known antifungal peptide defensins (Fig. 5b–f). Plant defensins are small proteins of 45–54 amino acid residues which, despite their low level of amino acid sequence identity, present a conserved tertiary structure that consists of a triple-stranded antiparallel β -sheet and one α -helix that are stabilized into a compact shape by three to five disulfide bridges (Lacerda *et al.*, 2014). A β -hairpin (the γ -core motif) has been reported to be essential for membrane interactions allowing the entry of defensins into fungal cells (Lacerda *et al.*, 2014; Sagaram *et al.*, 2013). Such a β -hairpin is preceded by the α -helix. As shown in Fig. 5b–e, BE27 displayed the same motif as defensins next to its carboxy terminus, consisting of a helix of 10 amino acid residues linked to an opposing β -hairpin by a loop of five residues. The β -hairpin is composed of two antiparallel β -strands of eight and six residues linked by a loop of five residues. This structure is almost identical to that of PAP-S, a RIP isolated from *Phytolacca americana* L. (Fig. 5c). Figure 5b and 5c also show the helix and γ -core motif from two representative antifungal defensins: SPE10 from *Pschyrhizus erosus* (L.) Urb. (Song *et al.*, 2011) and MtDf4 from *Medicago truncatula* Gaertn. (Sagaram *et al.*, 2013). Both defensins contain a helix of 11 residues and an opposing β -hairpin containing two β -strands (five and seven residues). These structures are characterized by the presence of several positive charges

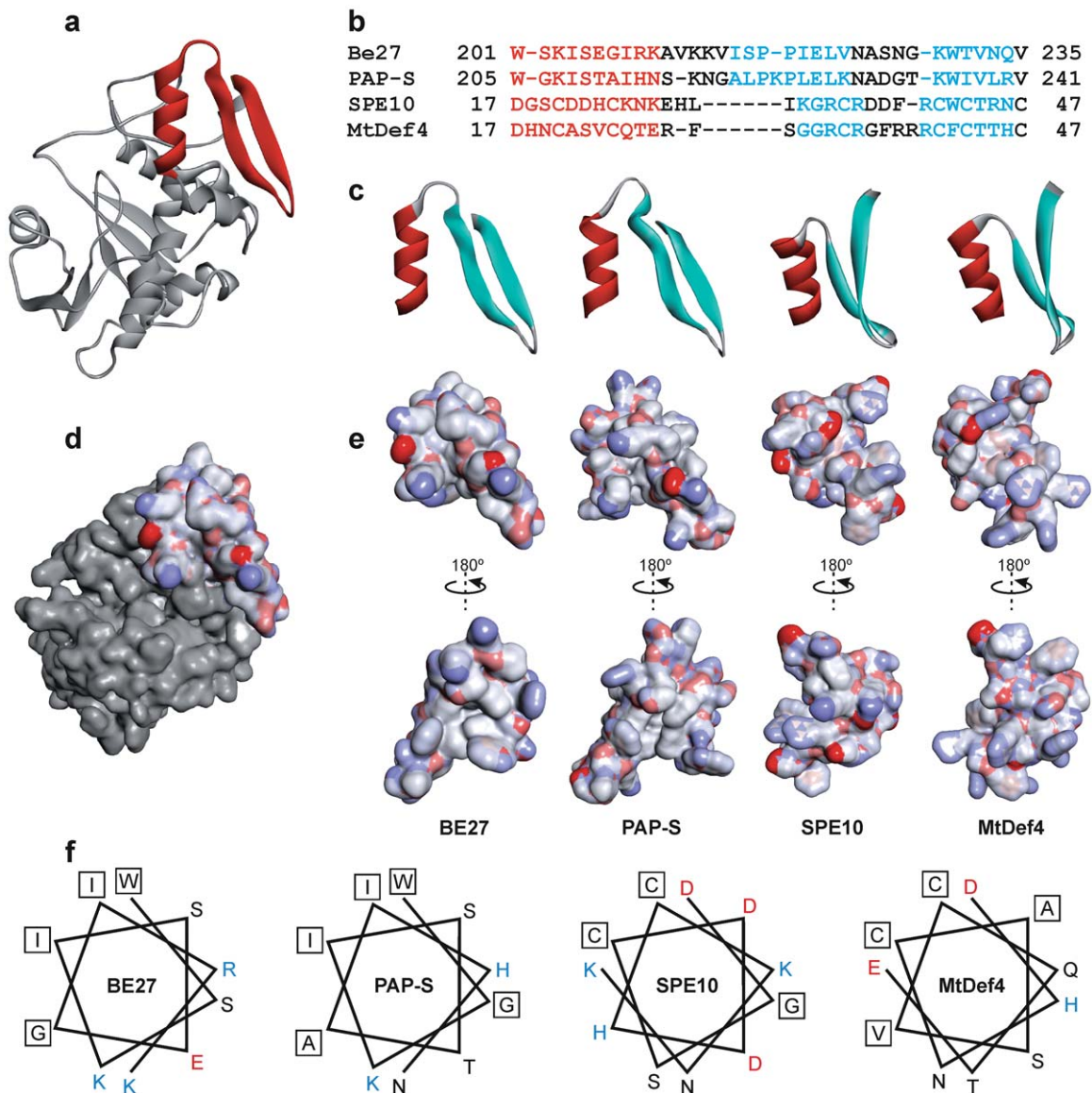


Fig. 5 Structure of the α -helix/ γ -core motif of BE27 compared with PAP-S (pokeweed antiviral protein) and the defensins SPE10 and MtDef4. (a) Structure of BE27 indicating the position of the α -helix/ γ -core motif from amino acid 201 to amino acid 235 (red ribbon). (b) Sequence alignments based on the three-dimensional alignments using the program Expresso (Di Tommaso *et al.*, 2011). Colouring of the alignments indicates the presence of α -helix (red) or β -sheet (cyan) secondary structures. (c) Ribbon representation of the α -helix/ γ -core motif from BE27 expanding from amino acid 201 to amino acid 235 and their homologues from PAP-S, SPE10 and MtDef4. The α -helix and β -strands are coloured in red and cyan, respectively. (d, e) Electrostatic surface of the α -helix/ γ -core motif from BE27 localized in the whole protein (d) or compared with the same motif in PAP-S, SPE10 and MtDef4 (e). Electrostatic potential is indicated in red (negative charge), white (neutral) and blue (positive charge). (f) Helical wheel drawing of the helices of BE27, PAP-S, SPE10 and MtDef4. The amino acid charge is indicated in red (negative), black (neutral) and blue (positive). Hydrophobic amino acids are boxed.

which, in the case of BE27, are supplied by the amino acids lysine (Lys)-203, arginine (Arg)-209, Lys-210, Lys-213, Lys-214 and Lys-229 (Fig. 5b). These amino acids confer to these structures a highly positively charged electrostatic surface (Fig. 5e), which is almost fully exposed in the BE27 structure (Fig. 5d). Such types of amino acids are required to interact, through electrostatic interactions, with the negatively charged sphingolipids of the fungal plasma

membrane (Lacerda *et al.*, 2014). BE27 also displays tryptophan (Trp)-230, a hydrophobic residue conserved in antifungal defensins (Fig. 5b). This residue [phenylalanine (Phe) in other defensins] has been reported to be a key factor for antifungal action in SPE10 (Song *et al.*, 2011). As shown in Fig. 5f, the α -helix of BE27 assumes an amphipathic conformation with opposing nonpolar [Trp-201, isoleucine (Ile)-204, glycine (Gly)-207, Ile-208] and polar [serine

(Ser)-202, Lys-203, Ser-205, glutamic acid (Glu)-206, Arg-209, Lys-210] faces oriented along the long axis of the helix. In addition, the polar face displays a high cationicity because of the presence of one Arg and two Lys. This type of conformation has been reported to permit an efficient interaction with biological membranes in antimicrobial peptides (Zelezetsky and Tossi, 2006). A similar conformation is present in the PAP α -helix, whereas such a conformation is not evident in SPE10 or MtDef4 (Fig. 5f).

DISCUSSION

The type 1 RIP BE27 has been reported as a defence protein synthesized in response to virus infection whose mechanism of action lies in its RNA polynucleotide:adenosine glycosylase activity, thus conferring it with the ability to depurinate and destroy viral RNAs (Iglesias *et al.*, 2005). However, other activities have been reported recently which might favour the resistance of beet against a broad variety of pathogens (Iglesias *et al.*, 2015). The combined effect of these biological activities could result in a broad action against several types of pathogen, such as viruses, bacteria, fungi or insects. Because several studies have indicated that some RIPs protect plants from fungal infection (Park *et al.*, 2002) and transgenic plants bearing RIP genes have been constructed that are resistant to fungi (Corrado *et al.*, 2005; Huang *et al.*, 2008; Qian *et al.*, 2014), we have tested the ability of BE27 to arrest the growth of the fungus *P. digitatum*. The *Penicillium* genus includes necrotrophic fungi that colonize the wounds and grow in the inter- and intracellular spaces of plant tissues (de Capdeville *et al.*, 2008). Crucial regulatory processes and protein–protein interactions take place in the apoplast during fungal infection (Doehlemann and Hemetsberger, 2013). Among the PR proteins which are secreted into the apoplast during basal resistance responses are chitinases, glucanases, thionins, osmotins, proteases and defensins (Doehlemann and Hemetsberger, 2013; van Loon *et al.*, 2006). Here, we report that BE27 is another apoplastic protein able to enter and kill fungal cells, thus arresting the growth of the fungus. The mechanism of action seems to involve rRNA *N*-glycosylase activity on the *Penicillium* major rRNA which inactivates irreversibly its ribosomes, thus inhibiting protein synthesis. By contrast, beet ribosomes are not sensitive to BE27 (Iglesias *et al.*, 2008). This contrasts with data reported for PAP from *Phytolacca americana* L. leaves, which inactivates their 'conspecific' ribosomes (Prestle *et al.*, 1992). PAP accumulates in the apoplastic space and has been postulated to be part of a general suicide strategy of some higher plants, rendering wounded tissue inefficient for the establishment of an infection by viruses or parasitic fungi (Park *et al.*, 2004; Prestle *et al.*, 1992). The different RIPs seem to display different antifungal capacities. It has been reported that ME, a type 1 RIP from *Mirabilis expansa* (Ruiz & Pav.) Standl., shows substantial inhibitory activity against *Rhizoctonia solani* growth, whereas the type 1 RIP saporin-S6

(from *Saponaria officinalis* L.) does not display such activity, despite the fact that saporin-S6 is more active against *R. solani* ribosomes (Park *et al.*, 2002). ME was targeted to the surface of fungal cells and transferred into the cells; thus, ME caused ribosome depurination and subsequent fungal mortality. In contrast, saporin did not interact with fungal cells, correlating with its lack of antifungal activity (Park *et al.*, 2002). It is worth mentioning that ME did not inhibit the growth of another fungus, *Trichoderma reesei* (Park *et al.*, 2002), suggesting that RIP antifungal activity could be specific.

The DNA of *P. digitatum* cultures appeared to be degraded, but did not show internucleosomal cleavage on exposure to BE27, thus suggesting that the mechanism of fungal growth inhibition is ribosome inactivation, followed by necrotic cell death. This contrasts with the effect of some RIPs on insects, in which apoptosis in their gut cells seems to be the mechanism of toxicity (Shahidi-Noghabi *et al.*, 2011). However, we cannot exclude the possibility that the SOD activity reported for BE27 (Iglesias *et al.*, 2015) may contribute to the toxicity towards fungal cells. SOD activity generates hydrogen peroxide, which can be toxic to fungi or may directly or indirectly stimulate plant defence responses (Doehlemann and Hemetsberger, 2013; van Loon *et al.*, 2006). In this respect, it has been reported that SOD and other antioxidant enzymes from rice coordinately participate in the resistance to rice stripe virus (Hao *et al.*, 2011), and that SOD transgenes confer resistance to the fungus *Cercospora beticola* in sugar beet (Tertivanidis *et al.*, 2004).

Fluorescence microscopy of *P. digitatum* after incubation with BE27 conjugated with the fluorescent dye Cy3 indicates that BE27 is able to internalize and reach the cytoplasm of fungal cells. This is worth mentioning because type 1 RIPs internalize poorly and high concentrations are necessary to become toxic to intact cells. In fact, BE27 displays IC₅₀ values of 2.2×10^{-6} and 4×10^{-7} M to COLO 320 cells (the most sensitive cellular culture) after 48 and 72 h of exposure, respectively (Iglesias *et al.*, 2015). These concentrations are equivalent to 59 and 11 μ g/mL, respectively, higher than that required to arrest *P. digitatum* growth. In order to study the structural peculiarities that may be involved in the antifungal behaviour of BE27, a three-dimensional structure was predicted by comparative modelling and compared with PAP and the defensins SPE10 from *Pschyrhizus erosus* (Song *et al.*, 2011) and MtDef4 from *M. truncatula* (Sagaram *et al.*, 2013). Defensins are small proteins with a conserved signature of cysteines, which can form three to four disulfide bridges (Lacerda *et al.*, 2014). An amino acid sequence alignment of antifungal defensins from plants shows that they do not present conservative amino acid sequences, except for the cysteine residues and a Gly residue (Lacerda *et al.*, 2014). However, defensins present a well-conserved three-dimensional structure which forms one α -helix and three antiparallel β -sheets. These proteins can interact specifically with some negatively charged sphingolipids present at the

cell membrane of pathogens, causing an increase in its permeabilization, leading to cell leakage and death by necrosis, or even to the transportation of these proteins to the intracellular environment, activating programmed cell death (Lacerda *et al.*, 2014; Thevissen *et al.*, 2003). The defensin specificity of binding to different sphingolipids or rafts containing sphingolipids and ergosterol is responsible for the disparity in toxicity of the different defensins against different fungi and the lack of toxicity towards plant cells (Thevissen *et al.*, 2003). It has been reported that the carboxy-terminal amino acid sequence is a major determinant of the antifungal activity of defensins (Sagaram *et al.*, 2013). This sequence is characterized by the presence of two antiparallel β -strands with an interposed loop and is preceded by an α -helix. Such a structural motif is characterized by a highly positively charged electrostatic surface (Lacerda *et al.*, 2014). In this study, we have identified a similar structural motif in the carboxy-terminal amino acid sequence of BE27 and PAP. This motif is shared by all apoplastic type 1 RIPs from dicots, and absent in the cytoplasmic type 1 RIPs from monocots (Di Maro *et al.*, 2014). In BE27, this motif also contains many positively charged amino acids required for the interaction with the negatively charged sphingolipid heads in the fungal membranes (Lacerda *et al.*, 2014; Thevissen *et al.*, 2003). However, the cysteines and disulfide bonds, a characteristic of all antifungal defensins, are absent in BE27. Because plant defensins lack a distinct hydrophobic core, the protein fold is stabilized primarily by disulfide bonds. This stabilization is achieved in type 1 RIPs by the presence of surrounding chains from the rest of the protein, particularly by the carboxy-terminal extension that forms an interface with the rest of the protein. The amphipathic α -helix of BE27 and PAP, characterized by a high cationicity of the polar face, might allow an efficient interaction with the fungal membrane, as has been reported for some antimicrobial peptides (Zelezetsky and Tossi, 2006). This might be necessary because PAP and, especially, BE27 present less cationic amino acids in their β -hairpins than SPE10 or MtDef4. Such amino acids have been found to be relevant for interaction with the negatively charged sphingolipids of fungal plasma membranes (Lacerda *et al.*, 2014; Sagaram *et al.*, 2013; Thevissen *et al.*, 2003). The insertion of BE27 into the lipid bilayer might be achieved by the presence of the β -hairpin, because a well-defined β -hairpin conformation has been reported to be necessary for disruption of the lamellar structure of the lipid membrane and insertion into the bilayer of β -hairpin antimicrobial peptides (Mani *et al.*, 2005).

The involvement of the above-mentioned structure in the entry and toxicity of BE27 and PAP in fungal cells is in accordance with studies reported on recombinant PAP mutants lacking several amino acids in their C-terminus (Baykal and Tumer, 2007; Hudak *et al.*, 2004; Hur *et al.*, 1995; Tumer *et al.*, 1997). Recombinant PAP is toxic to both transgenic tobacco (Tumer *et al.*, 1997) and yeast (Baykal and Tumer, 2007; Hudak *et al.*, 2004; Hur *et al.*,

1995) bearing the PAP gene, because this protein inactivates the ribosomes of the host. By contrast, C-terminal deletion mutants lacking a 25-amino-acid sequence (thus including part of the β -hairpin) and displaying rRNA *N*-glycosylase activity were not toxic to either host. This was a result of both a diminished stability of the mutant protein (Tumer *et al.*, 1997) and a failure in their sorting from the endoplasmic reticulum to the cytosol through the reticulum membrane (Baykal and Tumer, 2007; Hudak *et al.*, 2004; Hur *et al.*, 1995; Tumer *et al.*, 1997). However, this failure in the sorting to the cytosol was maintained by mutants having deletions just downstream of the β -hairpin. Although the authors of these articles sympathized with the hypothesis that the sequence of the last 10 amino acids was responsible for targeting to the membranes, such a sequence is not present in BE27 because it possesses a six amino acid shorter C-terminal sequence. A possible explanation is that the lack of these amino acids placed in the interface between the α -helix/ γ -core motif and the rest of the protein might diminish the stability of the β -hairpin structure in RIPs which, unlike defensins, lack the stabilizing disulfide bonds.

Further work will be conducted to study the *in vitro* and *in vivo* antifungal potential of BE27 and other RIPs against different fungi. The study of the toxicity of RIP mutants against fungal pathogens will clarify the role played by the C-terminus in the interaction with the fungal plasma membrane.

EXPERIMENTAL PROCEDURES

Materials

The sources of the chemicals used in this work have been indicated previously (Ferrerias *et al.*, 2011b) and most were obtained from Sigma-Aldrich (St Louis, MO, USA). The strain of *P. digitatum* was isolated in our laboratory and typified by the Spanish Type Culture Collection (CECT), Valencia, Spain. Potato dextrose agar (PDA) medium (purchased from Sigma-Aldrich) contains the following (per litre): 300 g potato infusion, 20 g dextrose, 15 g agar and 0.1 g chloramphenicol. Potato dextrose broth (PDB) medium (purchased from Sigma-Aldrich) contains the following (per litre): 6 g potato extract and 20 g dextrose. BE27 was isolated from 300 g of sugar beet leaves collected in September from a crop field at Cobos de Cerrato (Palencia, Spain) following a procedure described previously (Iglesias *et al.*, 2005).

Preparation of crude extracts from sugar beet leaves

All plants were grown in a growth chamber under conditions of 12 h light and 12 h dark at 20 °C. Treatment of laboratory-grown sugar beets with either hydrogen peroxide or salicylic acid was carried out by spraying dilute solutions (5 mM) every 24 h over 3 days. The crude leaf extracts were obtained from leaves that were ground in a ceramic mortar with liquid nitrogen; 30–50 mg were transferred to an Eppendorf tube and extracted for 10 min with 10 volumes of 5 mM sodium phosphate (pH 7.5) buffer containing 140 mM NaCl. After centrifugation at 10 000 g at 5 °C for 10 min, the supernatant was removed and stored at –20 °C.

Production of conidia and mycelium

The fungus was cultured on PDA slants for 8–10 days in a 26 °C incubator. A conidial suspension of *P. digitatum* was prepared by gently scraping the culture surface using a sterile glass rod after the addition of 2–3 mL of 0.1% aqueous solution of Tween-20. The spore suspension obtained was concentrated by centrifugation and dissolved in 0.1% aqueous solution of Tween-20. The concentration of the conidial suspension was quantified with a haemocytometer and the suspension was stored at –20 °C. Mycelium was obtained by inoculation of 250-mL portions of PDB in 1-L sterile Erlenmeyer flasks with 0.5 mL of spore suspension. The flasks were incubated at 26 °C on a horizontal shaker (250 rpm). After growing *P. digitatum* for 4–5 days, the mycelium was harvested by filtration through filter paper under vacuum, washed with distilled water and stored at –20 °C.

Antifungal activity measurements

Growth inhibitory assays of BE27 against *P. digitatum* were performed in 96-well microtitre plates. Conidia of *P. digitatum* (1000 spores/well) were incubated at 26 °C in 150 µL of PDB medium in the presence of different concentrations of BE27. Fungal growth was monitored spectrophotometrically using a microtitre plate Multiskan EX reader (Thermo Scientific, Waltham, MA, USA) and microscopically after 24, 28, 34, 45, 50, 71 and 78 h of incubation. Each experiment was performed in quadruplicate.

Mycelium for RNA and DNA extraction was prepared from cultures grown in 25-mm plates containing 2 mL of PDB medium inoculated with 1333 spores in the absence or presence of 4 µg/mL BE27. The plates were incubated at 26 °C on a horizontal shaker (40 rpm). After the growth of *P. digitatum* for 5 days, the mycelium was harvested by filtration through filter paper under vacuum, extensively washed with sterile water, weighed and stored at –80 °C. The experiments were carried out with six plates and performed in duplicate.

Microscopy of *P. digitatum* mycelium

Conidia of *P. digitatum* were grown *in vitro* in PDB medium for 24 h before exposure to BE27 (5 µg/mL) for 24 h. After incubation, samples were stained with 0.1% (w/v) of the chitin dye CFW for 5 min in the dark. Finally, mycelium was washed, resuspended in 20% glycerol and visualized under a Zeiss Axiovert 200M fluorescence microscope (Zeiss, Jena, Germany).

Cy3-BE27 labelling and confocal laser scanning microscopy of *P. digitatum* samples

BE27 fluorescence labelling with Cy3 was performed using the Cy3 Mono-Reactive Dye Pack from Amersham/GE-Healthcare (Barcelona, Spain) following the manufacturer's instructions. Conidia of *P. digitatum* were grown *in vitro* in PDB medium for 24 h before exposure to Cy3-BE27 (5 µg/mL) for 24 h. After washing, the samples were visualized with an LSM 510 META confocal microscope (Carl Zeiss, Jena, Germany).

rRNA *N*-glycosylase activity of BE27 on yeast and *P. digitatum* ribosomes

Preparation of the 30 000 g (S30) supernatants from yeast and *P. digitatum* was performed as described elsewhere (Iglesias *et al.*, 2015). Fresh baker's yeast (7.5 g) was ground in an unvitrified mortar precooled at –20 °C, with 15 g of alumina for 30 min at 4 °C, extracted with two volumes of buffer containing 10 mM Tris-HCl pH 7.6, 10 mM magnesium acetate, 10 mM KCl and 6 mM 2-mercaptoethanol, and centrifuged at 30 000 g for 20 min. This supernatant was designated S30 and was stored at –80 °C. The *P. digitatum* S30 supernatant was obtained from 15 g of mycelium as described for yeast. The rRNA *N*-glycosylase activity of beetin was assayed in 100-µL samples of S30 supernatants from yeast or *P. digitatum*, which were incubated with BE27 for 1 h at 30 °C. After treatment, the RNA was extracted with phenol and treated with aniline for 15 min at 23 °C. RNA samples were separated on a 5% urea–polyacrylamide gel, and stained with ethidium bromide (Iglesias *et al.*, 2015).

rRNA *N*-glycosylase activity in *P. digitatum* cultures

RNA from *P. digitatum* grown in the presence of BE27 was obtained from 15 mg of mycelium ground in a ceramic mortar with liquid nitrogen, using the RNA Plant Minikit (Qiagen, Madrid, Spain), according to the manufacturer's procedure. RNA samples were treated with aniline for 15 min at 23 °C, separated on a 5% urea–polyacrylamide gel and stained with ethidium bromide (Iglesias *et al.*, 2015).

DNA fragmentation analysis in *P. digitatum* cultures

The DNA from *P. digitatum* grown in the presence of BE27 was obtained from 20 mg of mycelium ground in a ceramic mortar with liquid nitrogen; 15 mg were transferred to an Eppendorf tube and suspended in 100 µL of 1 M sorbitol containing 0.1 M ethylenediaminetetraacetic acid (EDTA), pH 7.4, 0.1% 2-mercaptoethanol and 100 units of lyticase. After incubation at 30 °C for 1 h, the DNA was isolated following the instructions of the Genomic Prep Cells and Tissue DNA Isolation Kit (GE-Healthcare, Barcelona, Spain). DNA (3 µg) electrophoresis was carried out in 1% agarose gels using TBE buffer (0.089 M Tris, 0.089 M boric acid, 2 mM EDTA, pH 8.0) at 50 V for 4 h. DNA was stained for 20 min with Gel Red (Biotium, Inc., Hayward, CA, USA) and visualized with an ultraviolet lamp.

Molecular modelling of BE27 and structural comparison

The complete amino acid sequence of BE27 (accession number AAS67266.1) (Iglesias *et al.*, 2005) was downloaded from the National Center for Biotechnology Information (NCBI) sequence database (<http://www.ncbi.nlm.nih.gov/protein/>). Three-dimensional structural modelling was carried out on the I-TASSER server (Roy *et al.*, 2010; <http://zhanglab.ccmb.med.umich.edu/I-TASSER>). Protein structures 2q8w (PAP-S1aci), 3ctk (bouganin), 1qci (PAP), 1wuc (bouganin), 2z4u (PD-L4), 1ift (ricin A-chain) and 1apa (abrin A-chain) were chosen by I-TASSER as templates in the modelling. The three-dimensional structures of PAP-S

(accession code 1GIK), SPE10 (3PSM) and MtDef4 (2LR3) are available in the Protein Data Bank site (<http://www.rcsb.org/pdb/home/home.do>). Three-dimensional alignments of BE27, Psd1 and MtDef4 with PAP-5 were performed using the program Expresso (Di Tommaso *et al.*, 2011; <http://tcoffee.org.cat/apps/tcoffee/do:expresso>). Study and graph representations of protein structures were performed with the aid of the Discovery Studio 3.5 suite (<http://accelrys.com/>). The helical wheels were generated using WHAT 2.0 (Saier *et al.*, 2014; <http://www.tcd.org>).

Other procedures

Chitinase activity was evaluated as described previously (Naher *et al.*, 2012) using chitin as substrate and measuring the absorbance at 585 nm on addition of Ehrlich reagent (*p*-dimethylaminobenzaldehyde dissolved in glacial acetic acid that contained HCl). Protein synthesis was performed with a coupled transcription–translation *in vitro* assay using a rabbit reticulocyte lysate system as described elsewhere (Ferrerias *et al.*, 2011b). Protein concentrations were determined using the spectrophotometric method of Kalb and Bernlohr (1977). The RIP content in the crude extracts was calculated as indicated by Barbieri *et al.* (2006).

ACKNOWLEDGEMENTS

This work was supported by grant BIO/VA39/14 (Consejería de Sanidad, Junta de Castilla y León) to L.C. C.G. was supported by an Eurotango II fellowship (Erasmus Mundus Program). We thank Judy Callaghan (Monash University, Melbourne, Australia) for correcting the manuscript.

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