The cytoskeleton is disrupted by the bacterial effector HrpZ, but not by the bacterial PAMP flg22, in tobacco BY-2 cells

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Abstract

Plant innate immunity is composed of two layers. Basal immunity is triggered by pathogen-associated molecular patterns (PAMPs) such as the flagellin-peptide flg22 and is termed PAMP-triggered immunity (PTI). In addition, effector-triggered immunity (ETI) linked with programmed cell death and cytoskeletal reorganization can be induced by pathogen-derived factors, such as the Harpin proteins originating from phytopathogenic bacteria. To get insight into the link between cytoskeleton and PTI or ETI, this study followed the responses of actin filaments and microtubules to flg22 and HrpZ in vivo by spinning-disc confocal microscopy in GFP-tagged marker lines of tobacco BY-2. At a concentration that clearly impairs mitosis, flg22 can induce only subtle cytoskeletal responses. In contrast, HrpZ causes a rapid and massive bundling of actin microfilaments (completed in ~20 min, i.e. almost simultaneously with extracellular alkalinization), which is followed by progressive disintegration of actin cables and cytoplasmic microtubules, a loss of cytoplasmic structure, and vacuolar disintegration. Cytoskeletal disruption is proposed as an early event that discriminates HrpZ-triggered ETI-like defence from flg22-triggered PTI.

Key words: Actin, defence, flg22, Harpin Z, innate immunity, tobacco (Nicotiana tabacum L. cv. Bright Yellow 2).

Introduction

Plants lack the somatic adaptive immune system based on mobile defence cells characteristic for animal immunity. Plant defence, in contrast, is based upon an innate immunity of each individual cell (Jones and Dangl, 2006). This innate immunity comprises two distinct layers. The basal layer is evolutionarily ancient and triggered by conserved pathogen structures termed pathogen-associated molecular patterns (PAMPs). These PAMPs, such as flagellin, the subunit building the filament of bacterial flagellum, bind to specific receptors in the plasma membrane triggering so-called PAMP-triggered immunity (PTI). This basal layer of broad immunity is often accompanied by a more advanced and strain-specific immunity termed effector-triggered immunity (ETI), which is triggered by pathogen effectors that have to enter the cytoplasm of the host cell. The reason for this complexity is linked to coevolution between host and pathogen: PTI would be expected to select for pathogens, where the eliciting PAMPs are lost. However, since PAMPs are essential for the lifecycle of the pathogen, this evolutionary strategy does not work – a bacterial intruder lacking the PAMP flagellin would not elicit a PTI response, but it would also not be able to move. This dilemma stimulated, during a second round of host–pathogen warfare, the development of microbial effector proteins. These effectors are secreted into the cytoplasm of the host and suppress PTI (for review, see Tsuda and Katagiri, 2010). In response to these pathogen effectors, the host plant has evolved additional pathogen-specific receptors (encoded by so-called R genes) that specifically recognize the effectors in the cytoplasm and trigger the second layer of immunity, ETI (Boller and He, 2009). In many cases, ETI culminates in a plant-specific version of programmed cell death, the hypersensitive response.
often followed by systemic acquired resistance of the host. The conceptual dichotomy between PTI and ETI has been very valuable to interpret and classify the huge variety of plant defence responses, but this concept is presently on the move again. Recent studies show that the difference between PAMPs and effectors is more gradual than previously conceived (Thomma et al., 2011). Moreover, PTI and ETI share numerous common events (Tsuda and Katagiri, 2010). Thus, the apparent dichotomy might be a question of signal quantity rather than quality. In addition, plants can discriminate different pathogens and activate different responses that are appropriate for the respective pathogen. Therefore, at present, the PTI-ETI concept is extended towards a signature-based model (for review, see Aslam et al., 2009).

The archetypal elicitor of PTI is bacterial flagellin, which triggers defence responses in various plants (Gómez-Gómez and Boller, 2002). A synthetic 22-amino-acid peptide (flg22) from a conserved flagellin domain is sufficient to induce most of the cellular responses (Felix et al., 1999). A genetic screen in Arabidopsis thaliana using flg22 identified the Arabidopsis leucine-rich repeat receptor kinase FLS2, which binds flg22 (for review, see Chinchilla et al., 2006). Upon binding of the ligand, FLS2 is internalized by a receptor-mediated endocytic process that presumably has regulatory functions (Jones and Dangl, 2006).

To trigger ETI-like programmed cell death, Harpin proteins have been used. These bacterial proteins, first discovered in Erwinia amylovora, a phytopathogenic bacterium causing the fire-blight disease of apple and pears (Wei et al., 1992), have acquired considerable interest as triggers for hypersensitive response-like cell death and systemic acquired responses. They act as components of a bacterial type-III secretion system and can induce host events characteristic for ETI, including production of reactive oxygen species, accumulation of defence-related transcripts, and cell death (for review, see Tampakaki et al., 2010). Harpin proteins comprise different types that fulfil different functions during type-III secretion: HrpN is translocated into the host cytoplasm and acts stably with liposomes and synthetic bilayer membranes (HrpZPsph), pv. phaseolicola (HrpZPbg), is localized in the apoplast and acts as helper protein supporting type-III secretion. Functional proof for a role in type-III secretion comes from experiments where HrpZ could be successfully integrated into the type III secretion model system of the mammalian pathogen Yersinia enterocolitica (Lee et al., 2001a). Furthermore, HrpZ was found to associate stably with liposomes and synthetic bilayer membranes and forms an ion-conducting pore in vitro. Whether HrpZ can trigger ETI of the strict sense, is not clear. The finding of pathovar-specific activities in A. thaliana (Haapalainen et al., 2012) indicates the existence of cognate R-gene products. Due to this conceptual uncertainty, the term ‘ETI-like’ response will be used throughout the current work.

In animal cells, the host cytoskeleton is a major target of type-III effectors, in particular actin microfilaments (Cossart and Sansonetti, 2004). Although various type-III effectors of plant pathogenic bacteria can suppress plant defence responses such as hypersensitive cell death and expression of defence genes (for review, see Takemoto and Hardham, 2004), it is unclear whether they target the cytoskeleton as observed in animal cells. So far, AvrBs3, an effector from Xanthomonas campestris, was reported to induce swelling of mesophyll cells, a response that could be indicative of disruption to the plant microtubule, and HopZ1a, a type-III secreted effector from P. syringae was claimed to interact with both tubulin heterodimers and polymerized microtubules (Marois et al., 2002; Lee et al., 2012). Since several plant pathogens produce anticytoskeletal compounds during invasion (reviewed by Kobayashi and Kobayashi, 2008), the cytoskeleton seems to be an important player in plant defence. In fact, actin filaments seem to act cooperatively with PEN2 and PEN3 in penetration resistance against a broad range of pathogenic fungi (Stein et al., 2006). The role of the cytoskeleton has been mainly discussed in the context of barrier responses to pathogen penetration, for example, by cell-wall papillae that can be observed at sites of penetration attempts. The formation of these papillae is preceded by a reorganization of the cytoskeleton, causing redistribution of vesicle traffic and cytoplasmic aggregation towards the penetration site (for reviews, see Takemoto and Hardham, 2004; Kobayashi and Kobayashi, 2008) and a somewhat slower migration of the nucleus (for review, see Schmelzer, 2002). In addition, the cytoskeleton participates in the execution of hypersensitive cell death that involves and requires massive remodelling of actin filaments and microtubules (for review, see Smertenko and Franklin-Tong, 2011).

This study group’s previous work addressed the role of the cytoskeleton in Harpin-triggered defence using two cell lines from Vitis that differ in their microtubular dynamics manifested by altered levels of tyrosinylated α-tubulin (Qiao et al., 2010). The line Vitis vinifera cv. Pinot Noir is susceptible to pathogens such as Plasmopara viticola and Erysiphe necator, whereas Vitis rupestris efficiently copes with infection by these pathogens (Jürges et al., 2009). Treatment with a commercial HrpN preparation could trigger various defence responses, including extracellular alkalization and induction of defence genes. However, these defence responses were weaker and less sensitive in the cell line, where microtubules were more dynamic. In response to the elicitor, the cortical microtubules disappeared, which was most pronounced in V. rupestris. This correlation indicated that stable microtubules act as negative regulators of defence signalling and are disrupted in response to the elicitor. If this is more than a correlation, pharmacological manipulation of microtubules should be able to activate defence genes. In fact, using resveratrol synthase and stilbene synthase as key genes of phytoalexin induction, it could be shown that pharmacological manipulation of microtubules could induce gene expression in the absence of Harpin (Qiao et al., 2010), suggesting that the cytoskeleton might play a third role in plant defence. In addition to its function in barrier responses to penetration attempts, and in the execution of hypersensitive cell death, it might also act more upstream in defence signalling. Using the
same cellular system, early defence responses were compared to flg22 (PTI, not leading to cell-death) and Harpin (ETI-like response, culminating in cell death) and it was shown that both responses shared some of early signal components, but differed in perception, the signature of oxidative burst, and the integration into a qualitatively different stilbene output, correlated with different cell-death responses (Chang and Nick, 2012).

Both flg22 and Harpin affected cortical microtubules and actin filaments, which was more pronounced in the V. ruprestris line and for treatment with Harpin. However, since the cytoskeleton had to be visualized by either immunofluorescence (microtubules) or by fluorescent phalloidin (actin filaments), both requiring fixation of the cells, only the bulk changes of the cytoskeleton occurring at progressive stages of the response became detectable. To get clearer insight into the timing of the response, the current work investigated the tobacco BY-2 system. In this system, GFP-tagged marker lines for the cytoskeleton have been established, which allows the following the cytoskeletal response over time in living cells and thus also detection of the earlier stages of cytoskeletal remodelling. The cellular responses of BY-2 to flg22 (PTI) and HrpZ (ETI-like response) are compared with focus on the cytoskeleton. Consistent with the results from grapevine cells, strong and rapid cytoskeletal responses to HrpZ were observed, contrasting with very mild changes triggered by flg22. However, extending previous results by spinning-disc confocal microscopy and life-cell imaging, it is now shown that these responses initiate early and proceed in parallel with extracellular alkalization. The response of actin filaments (using the GFP-AtFABD2 marker) to flg22 or HrpZ was followed over time in individual cells by spinning-disc confocal microscopy. Confocal images were recorded with an AxioObserver Z1 (Zeiss, Jena, Germany) using a 63 × LCI-Neofluar Imm Corr DIC objective (NA 1.3), the 488 nm emission line of an Ar-Kr laser, and a spinning-disc device (YOKOGAWA CSU-X1 5000). The cells on the slide were maintained at 27 °C by in microscope temperature control stage and time-lapse series recorded by capturing Z-stack every 2 min over a period of 20 min (for flg22), or every 20 min over a period of 120 min (for HrpZ). The responses were elicited by 10 μM (flg22) or 57.6 μM (HrpZ). The long-term response was assessed from Z-stack collected from single recordings at lower concentrations at 3 day (for flg22, 100 nM) and 2 day (for HrpZ, 2.59 μM) after treatment. Mock controls were done with the respective solvents (water for flg22 and 5 mM MES for HrpZ) and treated in exactly the same manner as the samples.

Materials and methods

Cell lines and cultivation

BY-2 (Nicotiana tabacum L. cv. Bright Yellow-2) suspension cell lines (Nagata et al., 1992) were cultivated in liquid medium containing 4.3 g l⁻¹ Murashige and Skoog salts (Duchefa Biochemie; containing 1 l⁻¹ sucrose, 200 mg KH₂PO₄, 100 mg myo-inositol, 1 mg thiamine, and 0.2 mg 2,4-dichlorophenoxycetic acid, final pH 5.8). In addition to the non-transformed BY-2 wild type, transgenic lines were used in this study that expressed the actin-binding domain 2 of plant fimbrin in fusion with GFP under control of the constitutive CaMV 35S promoter (AtFABD2, Sano et al., 2005) and a transgenic line expressing the β-tubulin AtTuB6 from A. thaliana in fusion with GFP driven by the CaMV 35S promoter (Hohenberger et al., 2011). The cells were subcultivated weekly by inoculating 1.0–1.5 ml stationary cells into 30 ml fresh medium in 100 ml Erlenmeyer flasks. The cells were incubated in darkness at 27 °C under constant shaking on a KS260 basic orbital shaker (IKA Labortechnik) at 150 rpm. The media for the transgenic cell lines were complemented with either 30 mg l⁻¹ hygromycin (GFP-AtFABD2) or with 50 mg l⁻¹ kanamycin (GFP-AtTuB6), respectively.

Elicitors

The flg22 peptide QRLSTGSRINSAKDDAQLQIA (Felix et al., 1999) was purchased from a commercial source (Laboratoire de Biotechnologie du Luxembourg). The gene for the expression of Harpin from P. syringae p.v. phaseolicola (HrpZpH) was cloned into the vector pET21a (Novagen, Darmstadt, Germany) and transferred into Escherichia coli BL21 (DE3) RIL (Agilent Technologies, USA) by electroporation (Li et al., 2005). The recombinant protein was expressed and purified as follows. The transformed cells were grown at 30 °C in LB with 100 mg l⁻¹ ampicillin to an optical density at 600 nm approximately 0.6–0.8. Then, expression was induced with 1 mM isopropyl β-D-1-thiogalactopyranoside for about 6 h at 30 °C. Cells were spun down and the sediment resuspended in extraction buffer (50 mM Tris-HCl pH 8.0, 100 mM NaCl, 1 mM EDTA, 1 mM phenylmethanesulphonyl fluoride, freshly added). Cells were lysed by sonication and proteins denatured by boiling for 10 min. Cell debris and denatured proteins were removed by centrifugation (4 °C, 21,000 g, 10 min) and the supernatant transferred to a new beaker. Then, solid ammonium sulphate was slowly added to 45% (w/v) of a saturated solution while stirring the mixture on ice. The precipitated protein was resuspended in 5 mM MES (pH 5.5) and desalted by dialysis using regenerated cellulose tubular membrane (size exclusion 6 kDa, wall thickness 30 μM; ZelluTrans/Roth dialis- sis membranes T2, Roth, Karlsruhe, Germany). The concentration of purified Harpin was quantified by the amido black protein dye assay (Popov et al., 1975) against bovine serum albumin as calibration reference.

Microscopy

The response of actin filaments (using the GFP-AtFABD2 marker) and microtubules (using the GFP-AtTuB6 marker) to flg22 or HrpZ was followed over time in individual cells by spinning-disc confocal microscopy. Confocal images were recorded with an AxioObserver Z1 (Zeiss, Jena, Germany) using a 63 × LCI-Neofluar Imm Corr DIC objective (NA 1.3), the 488 nm emission line of an Ar-Kr laser, and a spinning-disc device (YOKOGAWA CSU-X1 5000). The cells on the slide were maintained at 27 °C by in microscope temperature control stage and time-lapse series recorded by capturing Z-stack every 2 min over a period of 20 min (for flg22), or every 20 min over a period of 120 min (for HrpZ). The responses were elicited by 10 μM (flg22) or 57.6 μM (HrpZ). The long-term response was assessed from Z-stack collected from single recordings at lower concentrations at 3 day (for flg22, 100 nM) and 2 day (for HrpZ, 2.59 μM) after treatment. Mock controls were done with the respective solvents (water for flg22 and 5 mM MES for HrpZ) and treated in exactly the same manner as the samples.

Measurement of extracellular alkalization

Extracellular alkalization was measured by combining a pH meter (pH 12, Schott Handylab) with a pH electrode (Lot 403-M8-57/120, Mettler Toledo). BY-2 cells were pre-equilibrated on an orbital shaker for around 30 min and then treated with different concentrations of flg22 (1–500 nM) or HrpZ (0.26–14.4 μM). Values for ΔpH were calculated as differentials of treatment versus mock control. Peak values were used as estimate for ΔpHmax and were reached at around 600 s (10 min) for flg22 and around 900–1200 s (15–20 min) for HrpZ. As positive controls for the activity of HrpZ, suspension cell lines from V. ruprestris and V. vinifera cv. Pinot Noir (Qiao et al., 2010; Chang and Nick, 2012) were added to the study.

Phenotyping of cellular responses

Packed cell volume as indicator for growth was determined at day 3 after subcultivation/treatment. Aliquots (2 ml) were sampled from the culture flask with a sterile, scaled pipette (Greiner) while shaking to ensure a homogenous distribution of cells, the pipettes sealed with Nescofilm and positioned vertically at 4 °C till full sedimentation of the cells. The packed cell volume was determined from the scale of the pipettes. Mitotic index was determined in aliquots of 500 μl at day 3 after subcultivation/treatment after staining for 2 min with 10 ng ml⁻¹
Hoechst 33258 (Sigma-Aldrich, Neu-Ulm, Germany) and addition of one drop of 10% (v/v) Triton X-100 (Roth). Cells were immediately scored under a AxioImager Z.1 using a DAPI filter set (excitation at 365 nm, beam splitter at 395 nm, and emission at 445 nm) by means of a Fuchs-Rosenthal haematocytometer (Thoma, Freiburg, Germany). For each data point, 2000 cells obtained from three independent experimental series were scored. To monitor changes in cell shape, differential interference contrast images of central sections were acquired by the AxioImager Z.1 in the ApoTome mode at day 3 after subcultivation/treatment using the mosaic and length measurement tools of the AxioVision software. As measure of cell shape, the ratio of cell width over cell length was measured and clustered into 11 classes increasing in steps of 0.2 and one class with values >2.0. Frequency distributions were constructed for three independent experimental series of 300–500 individual cells measured for each repeat. Mortality was assessed according to the method by Gaff and Okong’O-Ogola (1971) using 2.5% (w/v) Evans Blue (Sigma-Aldrich) in aliquots of 200 μl using custom-made staining chambers to remove the medium. The frequency of the dead cells (stained in blue) was scored using a Fuchs-Rosenthal haematocytometer under bright-field illumination with a AxioImager Z.1/ApoTome microscope. Mortality values were determined from three independent experiments with 1500 cells scored for each data point. To evaluate division synchrony (Campanoni et al., 2003), frequency distributions over the number of cells per individual file were constructed from approximately 2000 cell files from three independent experimental series.

Results

Actin responses to flg22 are subtle; those to HrpZ are drastic

To gain insight into the earlier stages of cytoskeleton remodelling in plant defence, this study followed the response of actin filaments to flg22 and HrpZ in individual BY-2 cells expressing the GFP-AtFABD2 marker by spinning-disc confocal microscopy. To ensure saturation of possible responses even for early time points, 10 μM flg22 was used, based on dose–response studies (to be discussed). Z-stack were collected through the entire cell every 2 min after addition of flg22 through the first 20 min (Fig. 1A). The transvacuolar network of actin cables emanating from the nucleus did not reveal any significant changes (Fig. 1A, upper row). However, a response of the fine meshwork of the cortical actin filaments underneath the cell membrane became detectable from ~5–10 min after elicitation (Fig. 1A, middle and lower rows). Here, the finer filaments disappeared and cortical actin cables changed orientation and often became more prominent (for instance, compare the images for 20 min

Fig. 1. Response of actin filaments in BY-2 to flg22 in vivo visualized by the GFP-AtFABD2 marker and spinning-disc confocal microscopy. (A) Time series after treatment with 10 μM flg22; upper row, merged Z-stack of confocal sections showing the layers in cell centre; middle and lower rows, details from two cortical regions of the same cell. (B) Solvent control (water). (C) Long-term response (3 days) after elicitation with 100 nM flg22. Observations are representative of at least five independent experimental series with a population of 100 individual cells for each treatment. Bars, 50, 10, and 5 μm (A), 10 and 5 μm (B), and 25 μm (C).
The cytoskeleton is disrupted by HrpZ, but not by flg22 in tobacco BY-2 cells

versus 4 min in the middle row of Fig. 1A). A responsiveness of the cortical actin meshwork was also observed in long-term studies using lower concentrations of flg22 such as 100 nM (Fig. 1C), where actin filaments in many (but not all) cells disintegrated into punctuate arrays, whereas the perinuclear network still persisted. None of these responses was observed in the mock control (Fig. 1B).

The rather subtle actin response to flg22 was in contrast to the drastic changes induced by HrpZ (Fig. 2). Merged Z-stack of confocal sections showing the layers in cell centre (upper row) and the cortical regions of the same cell (middle row) demonstrated that actin filaments had already undergone disruption at 20 min after elicitation with 57.6 µM HrpZ. Differential-interference contrast images of these cells showed that this disruption was accompanied by a dramatic breakdown of cytoplasmic architecture manifest at 60 min after elicitation and a disintegration of the vacuole (white arrows) visible at 90 min after elicitation. This was not observed in the solvent control (5 mM MES buffer). An inspection of earlier time points showed that actin filaments condensed into cortical aggregations that were clearly detectable at 10 min after elicitation (Supplementary Fig. S1A, available at JXB online) followed by contraction of actin towards the nucleus. A long-term experiment conducted at a ~20-fold reduced concentration (2.59 µM HrpZ), where most cells survived, as deduced from mortality scores (to be discussed), produced cortical actin filaments that were bundled and disorganized (Supplementary Fig. S1B).

Microtubules respond to HrpZ, but not to flg22

Parallel to actin, this study followed the response of microtubules using the GFP-AtTuB6 marker (Fig. 3). In the vast majority of cells, a treatment with 10 µM flg22 did not cause any changes, neither in the radial microtubules emanating from the nucleus (Fig. 3A, upper row) nor in the cortical microtubules underneath the membrane (Fig. 3A, lower row). Even for a long-term treatment, these microtubules did not respond (data not shown). However, in rare cases, aberrant microtubule structures could be observed (Supplementary Fig. S2). In these cells, plaque-like aggregations of signals appeared in the vicinity of the nuclear envelope either as a transient stage in response to a high concentration of flg22 (10 µM), or as long-term response to lower concentration (100 nM). As already found for actin, treatment with HrpZ (57.6 µM) caused drastic changes, mainly in the radial microtubules which were found to be eliminated (Fig. 3B, upper row), whereas cortical microtubules became thinner as compared to the control, but still maintained their integrity (Fig. 3B, lower row).

Fig. 2. Response of actin filaments in BY-2 to HrpZ in vivo visualized by the GFP-AtFABD2 marker and spinning-disc confocal microscopy; time series after treatment with a saturating concentration of HrpZ (57.6 µM); upper row, merged Z-stack of confocal sections showing the layers in cell centre; middle and lower rows, confocal sections and differential-interference contrast (DIC) images of cortical regions of the same cell, showing the breakdown of cytoplasmic architecture manifest at 60 min after elicitation and the disintegration of the vacuole (white arrows) at 90 min after elicitation. The solvent control consisted in 5 mM MES buffer. Observations are representative of at least five independent experimental series with a population of 100 individual cells for each treatment. Bars, 25 µm.
Fig. 3. Response of microtubules in BY-2 to 10 μM flg22 (A) or 57.6 μM HrpZ (B) in vivo visualized by the GFP-AtTuB6 marker and spinning-disc confocal microscopy, showing merged Z-stack of confocal sections showing the layers in cell centre and the cortical regions of the same cell. Controls (con) show the response to the solvent (A water, B 5 mM MES buffer). Observations are representative of at least five independent experimental series with a population of 100 individual cells for each treatment. Bars, 25 μm (A) and Bars, 50 μm (B).
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Apoptotic alkalization as rapid defence readout is more sensitive to flg22 than HrpZ

One of the earliest responses detected is a modification of plasma membrane permeability, in particular Ca^{2+}, H^+, and K^+ and anion fluxes that can be conveniently followed as changes of extracellular pH (Jabs et al., 1997; reviewed in Felix et al., 1999; Nürnberger and Scheel, 2001). To test, whether the differential cytoskeletal response to flg22 versus HrpZ was caused by a lack of sensitivity of BY-2 cells to flg22, this study therefore used extracellular alkalization as a fast cellular response monitoring plant defence. Already at a low concentrations (1 nM), flg22 could trigger a pH reaching a maximum at ~600 s after elicitation (Fig. 4A). This peak increased in amplitude, but was not advanced in time, when the concentration of flg22 was increased. A dose–response of maximal alkalization over the concentration of flg22 showed a saturation of the response from 200 nM at 0.27 pH units (Fig. 4B). HrpZ in a concentration of 1.73 μM could trigger an alkalization that was comparable in amplitude (Fig. 4C). To verify that this recombinantly produced elicitor was biologically active, the same concentration of HrpZ was tested in the two grapevine cell lines. For the more responsive V. rupestris, up to almost 0.7 pH units were obtained, for the less responsive V. vinifera cv. Pinot Noir the response was close to that found in BY-2. This pattern shows that the activity of HrpZ used in the current experiments was comparable to that of the commercial Harpin elicitor used in the previous studies (Qiao et al., 2010; Chang and Nick, 2012). A dose–response of maximal alkalization over the concentration of HrpZ in BY-2 (Supplementary Fig. S3) showed that stronger alkalization can be induced in BY-2 as well, but that this requires higher concentrations of the elicitor as compared to V. rupestris indicating differences in sensitivity between these cell lines. It should also be mentioned that the time points when these maxima were reached were between 900 and 1500 s, depending on the concentration, which is significantly later than the timing found for flg22 (Fig. 4A).

Cell division and cellular morphogenesis are altered by HrpZ

The morphogenesis of plant cells depends on the cytoskeleton. The cytoskeletal responses induced by HrpZ should therefore, as a consequence, alter cellular morphogenesis. Therefore, this study phenotyped growth, cell shape, and division synchrony as sensitive targets of cytoskeletal remodeling. A dose–response curve of packed cell volume as reporter for culture growth over the concentration of elicitor (Fig. 5A) showed a significant inhibition with increasing concentration of flg22 or HrpZ, respectively. This inhibition became more prominent with time (compare the curves for days 3 and 4 in Fig. 5A). To test, whether the decreased growth was caused by inhibition of cell division, the dose–responses of mitotic index were monitored at day 3 (at the peak of mitotic activity) after inoculation. For flg22, even for a high concentration of 1 μM flg22, mitotic index was only partially reduced (by some third as compared to the control). In contrast,
HrpZ reduced the mitotic index drastically to almost zero (Fig. 5B). The effects of flg22 and HrpZ on cell shape were investigated, since the axially of cell expansion depends on the organization of cortical microtubules. Frequency distributions for the ratio between cell width over cell length were constructed (Fig. 5C). For high concentrations of flg22 such as 1 μM, the frequency of cells with ratios of width/length 0.4–0.6 decreased, whereas cells with ratios >1 increased. This means that elongated cells became rare; whereas broader cells became more frequent (Fig. 5D left). This trend became progressively evident from flg22 concentrations exceeding 10 nM. The situation was inverse for treatments with HrpZ. Here the frequency of cells with a ratio of width/length 0.2–0.4 increased for as the elicitor concentration reached 1.73 μM, which means that cells became progressively elongated (Fig. 5D right). Lower concentrations of HrpZ were not effective in affecting cell shape (data not shown).

Since the effects of flg22 on cellular morphogenesis were very subtle in comparison with those of HrpZ, division synchrony was used as the most sensitive morphological marker. Cell divisions in tobacco suspension cells are synchronized by signals that depend on polar auxin transport which can be monitored as frequency peak of cells composed of six cells (Campanoni et al., 2003). Since polar auxin transport is strongly dependent on actin organization (reviewed in Nick, 2010), the division synchrony can be used as highly sensitive marker for actin remodelling and had been used successfully to monitor cell-death-related actin bundling (Chang et al., 2011). In response to flg22, the amplitude of this peak increased progressively up to 3-fold as compared to the control (Supplementary Fig. S4), suggesting that flg22 can modulate the actin cytoskeleton although this is not accompanied by bulk changes. Conversely, as to be expected from its actin bundling effect, a permissive concentration of HrpZ (2.59 μM) elevates the frequency of hexacellular files (Supplementary Fig. S4).

**HrpZ, but not flg22, can induce cell death**

A characteristic feature of ETI is programmed cell death, leading to a hypersensitive response occurring at infection sites. Therefore, this study followed mortality in response to flg22 and HrpZ over time. For flg22, elevated mortality was not observed for a range of concentrations (Fig. 6 left). Even for treatment with 200 nM (data not shown), mortality was only 3.4% at day 3. On the contrary, HrpZ induced cell death from day 2, reaching nearly 70% at day 3 after treatment with 8.64 μM HrpZ (Fig. 6 right).

**Discussion**

This work addressed the response of the cytoskeleton to flg22 (a canonical trigger for PTI) and HrpZ (a bacterial elicitor triggering an ETI-like response) by spinning-disc confocal microscopy and life-cell imaging in the BY-2 tobacco cell line. The motivation was the observation that a commercial Harpin elicitor induced cytoskeletal responses in two grapevine cell lines that differ in their microtubular dynamics (Qiao et al., 2010). A limitation of the grapevine system was the need to visualize microtubules by immunofluorescence, and actin by fluorescent phalloidin, both protocols requiring chemical fixation. Since the cytoskeletal response could thus not be followed over time in individual cells, only bulk changes of cytoskeletal organization became evident, which means that the early stages of these responses were not detected. This limitation was circumvented in the current work by using tobacco BY-2 cells, where transgenic fluorescent marker lines are available. A second drawback of the previous experimental system was the fact that the commercial Harpin elicitor is a HrpN species, which requires translocation into the host cytoplasm. Therefore, this study used HrpZ, which acts as helper protein supporting for type-III secretion in the apoplastic face of the membrane. Based on these two changes, it is now shown that the cytoskeletal responses differ depending on the nature of the elicitor and the nature of the cytoskeletal element: whereas both actin and microtubules responded drastically and rapidly to HrpZ, the cytoskeletal responses to flg22 remained very subtle. Both responses initiated early and could be detected from ~5–10 min after elicitation. The time course for the actin response either accompanied (for flg22) or even preceded (for HrpZ) extracellular alkalization, a very early response for defence, preceding calcium influx, mitogen activated protein (MAP) kinases, reactive oxygen species and plant hormones (salicylic acid, jasmonate, ethylene), and induction of defence-related genes. Generally, actin responded more sensitively as compared to microtubules, although the microtubules in the cell centre disintegrated in response to HrpZ (parallel with actin filaments).

To understand the differential early cytoskeletal response to flg22 versus HrpZ, it is relevant to compare the perceptive mechanisms for these elicitors. The PAMP flg22 is a ligand of the leucine-rich repeat receptor kinase FLS2 (Gómez-Gómez and Boller, 2000; Zipfel et al., 2004; Chinchilla et al., 2006). A dose–response curve using apoplastic alkalization as readout shows that around 100 nM were required to elicit a significant response (Fig. 4B), which places BY-2 into the less responsive systems. For comparison, cell lines of V. rupestris produce a half-maximal response at <5 nM flg22, whereas, on the other hand, V. vinifera cv. Pinot Noir needs >800 nM flg22 for half-activation (Chang and Nick, 2012). Although BY-2 does not classify for being highly flg22 sensitive, flg22 is clearly sensed at much lower concentrations as compared to HrpZ, where >5 μM are required to get half-maximal alkalization (Supplementary Fig. S3), consistent with findings in grapevine cells (Chang and Nick, 2012). Both elicitors also differ in the timing of apoplastic alkalization, HrpZ occurs about 5–10 min later as compared to flg22 leading to a model, where the link between flg22 and alkalization is more direct, whereas the link between Harpin and alkalization is indirect. Again, this difference in timing is not confined to BY-2, but has also been observed in the grapevine system (Chang and Nick, 2012).

The low sensitivity to HrpZ indicates that this elicitor is not perceived through a canonical protein receptor, consistent with findings from other groups: For HrpZPsp, from
The cytoskeleton is disrupted by HrpZ, but not by flg22 in tobacco BY-2 cells

**Fig. 5.** Cellular responses to flg22 (left) and HrpZ (right). (A) Dose–response of packed cell volume as indicator of culture growth at day 3 (peak of mitotic activity) and day 4 (onset of cell expansion) after inoculation, respectively. (B) Dose-response of mitotic index at day 3 after inoculation (scored from a population of 2000 individual cells for each data point). (C) Effect on cell shape; frequency distributions for the ratio between cell width over cell length were constructed from 2500 individual cells for each experiment. (D) Differential-interference contrast images of representative cells either raised under control conditions (centre) or after treatment with either 1 μM flg22 or 1.73 μM HrpZ. Data in (A) and (B) are mean ± standard errors from three biological replicates.
P. syringae, cation-permeable pores have been reported (Lee et al., 2001a). HrpZ is highly conserved in P. syringae strains and, unlike the harpin HrpN of E. amylovora which is an essential pathogenicity factor (Wei et al. 1992), seems to act as helper protein supporting type-III secretion. Instead of being translocated into the host cytoplasm, HrpZ has been localized in the apoplast (Brown et al., 2001). HrpZ can bind to phosphatidic acid and, upon insertion into vesicles prepared from plant plasma membranes, it can cause vesicle disruption (Lee et al., 2001b; Haapalainen et al., 2011). Despite this obviously different mode of perception for HrpZ, there seems to be some sort of specificity: when HrpZ proteins originating from two different pathovars of P. syringae were administered to A. thaliana, both HrpZ types caused cell death but the modulation of anionic currents was specific, and in a phage display screen for peptide binding, different motif preferences were observed (Haapalainen et al., 2012).

Thus, whereas the flg22 signal is transduced by a classical receptor kinase activity, the HrpZ signal acts by a localized loss of membrane integrity, which will directly impinge upon cortical actin. Actin is known to stabilize membrane integrity in a great number of systems (Papakonstanti et al., 2000; for review, see Koivusalo et al., 2009) and seems to be directly linked with the plasma membrane of plant cells as demonstrated by TIRF-microscopy in BY-2 protoplasts expressing the GFP-FABD2 marker (Hohenberger et al., 2011). This membrane-associated actin population in the one hand stabilizes membrane integrity (Hohenberger et al., 2011) and sensitively responds to perturbations of membrane integrity by rapid detachment from the plasma membrane, which is then followed by bundling of actin cables, contraction towards the nucleus, and programmed cell death (Berghöfer et al., 2009). The current observations integrate well into the accumulating body of evidence linking actin remodelling with programmed cell death across eukaryotic cells in general (for review, see Gourlay and Ayscough, 2005; Franklin-Tong and Gourlay, 2008) and plant cells in particular (Smertenko and Franklin-Tong, 2011). For instance, during self-incompatibility in poppy, actin remodelling is necessary and sufficient to activate programmed cell death in the male gametophyte (Thomas et al., 2006). Conversely, the programmed cell death of embryonic suspensors during the somatic embryogenesis of conifers requires actin remodelling which is necessary for the embryo proper to become committed for embryogenesis (Smertenko et al., 2003).

The link between impaired membrane integrity by HrpZ-induced pore formation and actin remodelling will be the topic of future investigations. A molecular candidate for this link might be reactive oxygen species, since they not only participate in the signalling culminating in programmed cell death (for review, see Gechev et al., 2006), but also trigger actin reorganization in plant programmed cell death (Wilkins et al., 2011). The remodelling of actin in response to changes of redox potential is observed across all eukaryotic kingdoms pointing to ancient origins (Franklin-Tong and Gourlay, 2008). In fact, when in Harpin-triggered grapevine cells the NADPH oxidase as major source of apoplastic oxidative burst was blocked by diphenylene iodonium chloride, or when apoplastic hydrogen peroxide was scavenged by addition of catalase, this impaired the induction of stilbene synthase, a key player for grapevine phytoalexin synthesis (Chang et al., 2011).

Compared to actin, the response of microtubules seems to be more indirect. Even at the time when the actin cytoskeleton is breaking down in response to HrpZ, cortical microtubules, although disturbed in their orientation, still maintain a certain degree of integrity (Fig. 3B). Since plant microtubules have been recently shown to respond to oxidative imbalance (Livanos et al., 2012), they might simply respond to the actin-dependent oxidative burst caused by pore formation. Microtubules modulate deformation-sensitive calcium channels (Ding and Pickard, 1993; Mazars et al., 1997) and therefore modulate apoplastic alkalinization (Chang and Nick, 2012). Alkalinization in response to Harpin elicitors occurs later as compared to that triggered by flg22 in both BY-2 (this work) and grapevine (Chang and Nick, 2012). This might be caused by the time span required for the reactive oxygen species to disassemble the microtubules modulating the calcium channel. Oryzalin and taxol could activate defence genes in the absence of Harpin elicitors in grapevine, indicating a function of microtubules upstream of gene induction (Qiao et al., 2010). Interestingly, latrunculin B and

Fig. 6. Mortality in response to flg22 (left) and HrpZ (right). Data are mean ±standard errors from three independent experimental series with a population of 2000 individual cells scored after staining with 2.5% Evans Blue.
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