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Tenuazonic acid from Stemphylium loti inhibits the plant plasma membrane H\(^+\)-ATPase by a mechanism involving the C-terminal regulatory domain

Peter K. Bjørk\(^1\), 0000-0002-5641-7791
Silas A. Rasmussen\(^2\), 0000-0001-7633-8743
Sisse K. Gjetting\(^1\), 0000-0002-7173-3880
Nanna W. Havshøi\(^1\),
Thomas Isbrandt Petersen\(^2\), 0000-0003-3938-0816
Johan Ø. Ipsen\(^1\), 0000-0001-5509-8496
Thomas O. Larsen\(^2\), 0000-0002-3362-5707
Anja T. Fuglsang\(^1\) 0000-0003-1153-8394

\(^1\)Department of Plant and Environmental Sciences, Faculty of Science, University of Copenhagen, Thorvaldsensvej 40, 1870 Frederiksberg C, Denmark;
\(^2\)Department of Biotechnology and Biomedicine, Technical University of Denmark Søltofts Plads, B. 221, 2800 Kongens Lyngby, Denmark

Author for correspondence:
Anja T. Fuglsang
Tel: +45 35 33 25 86
Email: atf@plen.ku.dk

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Summary

- Pathogenic fungi often target the plant plasma membrane (PM) H⁺-ATPase during infection. To identify pathogenic compounds targeting plant H⁺-ATPases, we screened extracts from 10 *Stemphylium* species for their effect on H⁺-ATPase activity.
- We identified *Stemphylium loti* extracts as potential H⁺-ATPase inhibitors, and through chemical separation and analysis, we identified tenuazonic acid (TeA) as a potent H⁺-ATPase inhibitor. By assaying ATP hydrolysis and H⁺ pumping, we confirmed TeA as a H⁺-ATPase inhibitor both in vitro and in vivo. To visualize in planta inhibition of the H⁺-ATPase, we treated pH-sensing *Arabidopsis thaliana* seedlings with TeA and quantified apoplastic alkalization.
- TeA affected both ATPase hydrolysis and H⁺ pumping, supporting a direct effect on the H⁺-ATPase. We demonstrated apoplastic alkalization of *Arabidopsis thaliana* seedlings after short-term TeA treatment, indicating that TeA effectively inhibits plant PM H⁺-ATPase in planta. TeA-induced inhibition was highly dependent on the regulatory C-terminal domain of the plant H⁺-ATPase.
- *Stemphylium loti* is a phytopathogenic fungus. Inhibiting the plant PM H⁺-ATPase results in membrane potential depolarization and eventually necrosis. The corresponding fungal H⁺-ATPase, PMA1, is less affected by TeA when comparing native preparations. Fungi are thus able to target an essential plant enzyme without causing self-toxicity.

**Key words:** fusicocecin, phytotoxin, plasma membrane H⁺-ATPase, *Stemphylium loti*, tenuazonic acid
Introduction

Plants have evolved an advanced, multilayered defense system that fends off pathogen invasion. In response, pathogens have adapted by developing increasingly advanced invasion strategies to overcome plant defense mechanisms. While plant defense relies on recognition of specific and conserved microbe structures, known as microbe-associated molecular patterns (MAMPs), pathogens must target vital plant functions or the proteins involved (Jones & Dangl, 2006; Boller & Felix, 2009). Many fungal species, both pathogenic and beneficial, target the plant plasma membrane (PM) H⁺-ATPase (Elmore & Coaker, 2011). Nutrient uptake in plant cells is highly reliant on the electrochemical gradient across the PM, which is established primarily by the PM H⁺-ATPase (Palmgren, 2001; Falhof et al., 2016). Protons are transported across the PM and energized by the hydrolysis of ATP into ADP and P. H⁺-ATPase is essential for plant growth and development, as well as specific functions, such as stomata dynamics (Kinoshita & Shimazaki, 1999; Inoue & Kinoshita, 2017). The H⁺-ATPase has 10 transmembrane helices, a large cytoplasmic domain, a C-terminal autoinhibitory domain, and it belongs to the family of P-type ATPases (Morth et al., 2011). The C-terminus comprises approximately 100 amino acids and does not have any defined structure. Rather, it is suggested to wind around the body of the H⁺-ATPase and physically constrain protein activity (Pedersen et al., 2007). H⁺-ATPase regulation often involves direct phosphorylation of specific sites in the C terminus. The well-studied Arabidopsis thaliana H⁺-ATPase (AHA2) is activated by phosphorylation of Thr⁸⁸¹ and Thr⁹⁴⁷, while it is inactivated by phosphorylation of Ser⁸⁸⁹ and Ser⁹³¹ (Jahn et al., 1997; Fuglsang et al., 2007; Rudashevskaya et al., 2012; Guzel Deger et al., 2015; Haruta et al., 2015). Deletion of the regulatory C-terminal domain or mutation of specific phosphorylation sites results in an activated pump that is insensitive to inhibition caused by phosphorylation.

Like plants, fungi are dependent on the PM H⁺-ATPase to generate an electrochemical gradient across the PM. The yeast (Saccharomyces cerevisiae) H⁺-ATPase (PMA1) shares structural similarity with its plant equivalent, but the C-terminally regulatory domain is much shorter (Portillo, 2000; Pedersen et al., 2007). PMA1 has only two known phosphorylation sites, both of which are responsive to glucose (Mazón et al., 2015). This difference in regulatory mechanisms between the H⁺-ATPases of plants and fungi creates the opportunity for fungal compounds to interact with plant H⁺-ATPases.
without affecting the fungus itself. Indeed, the fungal phytotoxin fusicoccin (FC), from the fungus *Fusicoccum amygdali*, specifically stabilizes binding between the regulatory C-terminal domain of the plant PM H\(^+\)-ATPase and its activating 14-3-3 protein, resulting in irreversible activation of the pump (Fuglsang *et al.*, 1999, 2003; Würtele *et al.*, 2003). FC is widely used to investigate H\(^+\)-ATPase activation in the laboratory; however, plants treated with FC lose control of stomata regulation and suffer from uncontrolled water loss and wilt (Marre, 1979).

Members of the fungal species *Stemphylium* are plant pathogens that cause leaf spots in crops such as asparagus (*Asparagus officinalus*), garlic (*Allium sativum*), parsley (*Petroselinum crispum*), pear (*Pyrus communis*), and sugar beet (*Beta vulgaris L.*)(Graf *et al.*, 2016; Gálvez *et al.*, 2016; Koike *et al.*, 2013; Köhl *et al.*, 2009; Tanahashi *et al.*, 2017; Hanse *et al.*, 2015). The taxonomy and metabolite production of *Stemphylium* spp. reveal a large family of both host-specific and non-host-specific pathogenic fungi, producing a vast number of diverse metabolites (Woudenberg *et al.*, 2017; Olsen *et al.*, 2018). However, the invasion strategy and plant pathogenicity of *Stemphylium* spp. remain elusive.

In this study, we screened a range of chemical extracts from different plant pathogenic fungi and identified Tenuazonic acid (TeA) from *Stemphylium loti* as specifically targeting the plant PM H\(^+\)-ATPase. TeA was previously shown to inhibit photosynthesis and the potential use of TeA as a herbicide targeting PSII was recently reviewed by Chen & Qiang (2017). Here we show that TeA inhibits plant PM H\(^+\)-ATPases at micromolar concentrations by a mechanism involving the C-terminal regulatory domain. Furthermore, we show that TeA targets the plant PM H\(^+\)-ATPase with a higher specificity compared to its *Saccharomyces cerevisiae* homolog, PMA1, when comparing native preparations of H\(^+\)-ATPase. These results suggest that *Stemphylium loti* targets the PM H\(^+\)-ATPase of the host cell upon infection as part of a mechanism that eventually leads to cell death.

**Materials and Methods**

**Chemical materials**

Tenuazonic acid (TeA) (cat #610-88-8) was purchased from Santa Cruz Biotechnology (Dallas, TX, USA). Fusicoccin (FC) (cat #F0537) was purchased from Sigma-Aldrich (St. Louis, MO, USA).
Purification of spinach plasma membranes

PM-enriched vesicles from *Spinacia oleracea* (baby spinach) were isolated using two-phase partitioning as described by Lund and Fuglsang (2012). Fresh *Spinacia oleracea* leaves (30 g) were homogenized in buffer (50 mM MOPS, 5 mM EDTA, 50 mM Na₄P₂O₇, 0.33 M sucrose, and 1 mM Na₂MoO₄, pH 7.5) and centrifuged for 15 min at 10 000 g. The supernatant was collected and centrifuged for 30 min at 50 000 g, and the pellet was resuspended in 330/5 buffer (0.33 M sucrose and 5 mM K₂HPO₄). Following PM were enriched by two-phase partitioning (in 20% Dextran T500 solution, 0.33 M sucrose, 5 mM K₂HPO₄, and 4 mM KCl). PM fractions were frozen in liquid N₂ and stored at −80°C.

In cases where plant material was pretreated, *Spinacia oleracea* leaves were incubated with 5 μM TeA or an equal volume of 1% DMSO (control) for 15 minutes at room temperature prior to homogenization.

Plant material for bioimaging and growth tests

For perfusion assays, *Arabidopsis thaliana* (ecotype Col-0) seeds stably expressing the pH sensor apo-pHusion (Gjetting et al. 2012) were surface sterilized and grown under long-day light conditions (16 h light/8 h dark) on agar plates containing ½ × Murashige and Skoog (MS) (Murashige & Skoog, 1962) + 1% (w/v) sucrose for four to five days before perfusion imaging experiments.

In order to test the effect of TeA on root growth, *Arabidopsis thaliana* (Col-0) seeds were surface sterilized using 1-5% w/w sodium hypochlorite and 0.73% w/w HCl. Seeds were saturated over night at 4°C on ½ x MS including vitamins (1% sucrose, 0.7% plant agar). Germinated and grown for 6 days under long-day light conditions (16h light/8h dark, at 20°C) before transferring to ½ x MS agar containing 0, 2.5, 5, 10 or 20 μM TeA. Seedlings were grown for another 6 days, and growth were measured every second day. Image analysis was done using ImageJ 1.47v.

Perfusion assays

Roots of four- to five-day-old seedlings were immobilized with agar on Teflon-coated slides, covered with a droplet of bath solution (0.1 mM CaCl₂, 0.5 mM KCl, and 10 μM MES, pH 5.5) and left to
stabilize for 5–10 min before mounting on a Leica SP5-X confocal laser scanning microscope (Leica Microsystems, Mannheim, Germany). Using a 20x dipping objective and a perfusion setup as described by Gjetting et al. (2012), either bath solution or 10 μM TeA was added. Imaging data for apo-pHusion fluorescence in the root elongation zone were acquired in xyt-mode using a white light laser with line-by-line sequential scanning (line average 2) of the fluorescent proteins EGFP (excitation 488 nm; emission 500–530 nm) and mRFP1 (excitation 558 nm; emission 600–630 nm). The pinhole was set to an airy disc of 2.

Perfusion experiments showing no focal shift or unstable baseline before initial changing of buffer were selected for data analysis. Imaging data were analyzed using the open-source software ImageJ (https://imagej.nih.gov/ij/index.html). Background values were subtracted based on average intensity in areas without cells. Ratio calculations were created using pixel-by-pixel division of EGFP with mRFP1 generating floating 32-bit images. Regions of interests (ROIs) were chosen for calculating average pixel intensities to represent the largest possible area with cells in the same optical plane. Calibration curves were created using 10 mM MES/10 mM MOPS and 10 mM citrate. pH was adjusted to between 5 and 7 with KOH.

Yeast strain growth and plasmids
Saccharomyces cerevisiae strain RS-72 (MATa ade1-100 his4-519 leu2-3, 112) was transformed with plasmids containing cDNA of the Saccharomyces cerevisiae H+-ATPase, PMA1 (pMP 400); AHA2 (pMP 1625); or C-terminally truncated A. thaliana H+-ATPases, aha2Δ40 (pMP 810), aha2Δ77 (pMP210) or aha2Δ92 (pMP 132), under control of the PMA1 promoter (Regenberg et al., 1995). In the RS-72 strain, the promoter of the endogenous PM H+-ATPase PMA1 gene is replaced with a galactose-dependent promoter, GAL1. When cells are grown on galactose, the endogenous H+-ATPase PMA1 is expressed, whereas changing to glucose-based media requires complementation with a functional PM H+-ATPase. Transformed yeast cells were precultured in 5 ml of SGAH medium (galactose, yeast nitrogen base, adenine, histidine), transferred to 100 ml SGAH and cultured overnight. The cell culture was transferred to 500 ml sterile YPD medium for 20 h before membrane harvest. All cultures were grown at 30°C, 200 rpm.
Purification of yeast microsomes, plasma membranes and ER enriched membranes

Transformed *Saccharomyces cerevisiae* cells were harvested essentially as described by Villalba *et al.* (1992). To obtain PM and ER enriched fractions, microsomal fractions (MFs) were resuspended in GTED$_{20}$ (20% glycerol (v/v), 100 mM Tris-HCl, pH 7.5, 1 mM EDTA, pH 8, and 1 mM DTT) or STED$_{10}$ (as GTED but with 10% sucrose w/v), placed on top of an 8-ml sucrose step gradient consisting of increasing concentrations of sucrose in STED buffer (29, 33, 43 and 53 % STED), and centrifuged for 16-20 h at 154 000 $g$ at 4°C. PM fractions were collected at the 43/53% interface, ER fractions at the 29/33% interface, resuspended in ice-cold MilliQ water, and centrifuged for 1 hr at 250 000 $g$ at 4°C. The pellet was resuspended in GTED$_{20}$, frozen in liquid N$_2$, and stored at -80°C.

Protein determination

Protein concentration was determined as described by Bradford (1976) with γ-globulin as standard.

SDS-PAGE and western blotting.

SDS-gel electrophoresis (SDS-PAGE) was performed according to standard techniques. SDS-PAGE gels were stained using Coomassie Brilliant Blue (CBB) to visualize separated proteins. For immunoblotting PM H$^+$/ATPase was detected by an antibody directed against the central part of AHA2. Phosphothreonine residues were detected by an anti-P-Thr antibody from Zymed (#71-8200; dilution1:600).

ATPase assay

H$^+$/ATPase activity was assayed according to the modified protocol of Baginski *et al.* (1967) described by Wielandt *et al.* (2015). For each well, 0.5–3 μg of protein was added in a 60 μl volume. For initial screening, the assay was performed at 30°C for 30 min, with 2.5 mM ATP, pH 7, in ATPase assay buffer (20 mM MOPS, 8 mM MgSO$_4$, 50 mM KNO$_3$ [inhibitor of V-ATPases], 5 mM NaN$_3$ [inhibitor of mitochondrial ATPases], and 0.25 mM Na$_2$MoO$_4$ [inhibitor of acid phosphatases]). For kinetics studies, an ATP regenerating system (5 mM phosphoenolpyruvate and 0.05 μg μl$^{-1}$ pyruvate kinase) was added. For competition assays between TeA and FC/14-3-3, PM fractions were incubated with either 20 μM TeA and 1μg 14-3-3 or 20 μM FC and 1μg 14-3-3 protein for 10 min at
room temperature prior to assay initiation. PM from *S. oleracea* was used for control experiments and incubated with 10 μM FC, 1μg 14-3-3 protein or both for 10 min prior to assay start (total assay volume of 60 μl). All ATPase assays were performed using three technical replicates and two biological replicates.

Proton pumping assay

*Spinacia oleracea* PM vesicles were turned inside out by the detergent Brij 58 (Johansson et al., 1995). Proton transport across the membrane was assayed using a modified version of the method described by Lund and Fuglsang (2012). The fluorescent probe 9-amino-6-chloro-2-methoxyacridine (ACMA) can move freely across membranes in the unprotonated state. As inside-out vesicles become loaded with H⁺ upon activation of the H⁺-ATPase, ACMA becomes protonated and, thus, trapped inside vesicles, resulting in fluorescence quenching (excitation 412 nm; emission 480 nm). The assay was performed using 0.03 μg μl⁻¹ protein in reaction buffer (11.4 mM MOPS, 56.8 mM K₂SO₄, 22.7 mM glycerol, 2.3 mM ATP, 1 μM ACMA, 0.05% Brij 58, and 60 nM valinomycin, pH 7.0) and started by addition of MgSO₄ to a final concentration of 3 mM. The assay was stopped by addition of 10 μM of the H⁺ ionophore nigericin.

Dose-response and enzyme kinetics analysis

The dose response of ATP hydrolysis of plant PM to inhibitors, i.e. fungal extracts and TeA, or agonists, i.e. FC/14-3-3, was determined by monitoring ATP hydrolysis in response to increasing inhibitor/agonist concentration. Kinetic data were obtained by monitoring ATP hydrolysis in response to increasing ATP concentration. All ATP hydrolysis data were obtained as described in the Materials and Methods section “ATPase assay”. The concentration of inhibitor needed to reach half-inhibition, IC₅₀, was calculated by nonlinear regression fitted to log[inhibitor] versus the normalized response (variable slope, bottom constraint =0.0) using GraphPad Prism 5.0a for Mac OS X, GraphPad Software (www.graphpad.com). The kinetic parameters *V*ₘₐₓ and *K*ₘ were estimated by nonlinear regression using GraphPad Prism 5.0a.

Statistics
All experiments were performed as technical triplicates (minimum) with two or more biological replicates. Values are presented as mean ± SEM; \( P \) values were calculated by one- or two-way analysis of variance (ANOVA) followed by Dunnett’s posttest analysis or Bonferroni posttest using GraphPad Prism 5.0a. \( P < 0.05 \) was considered statistically significant (*, \( P < 0.05 \); **, \( P < 0.01 \); ***, \( P < 0.001 \)).

Sequence alignment
The PMA1 amino acid sequence was obtained from the Saccharomyces Genome Database (www.yeastgenome.org). BLASTp was employed to identify sequences of H\(^+\)-ATPases from *Stemphylium lycopersici*, *Alternaria alternata*, and *Fusarium oxysporum* with close similarity to that of PMA1 using the ExPASy Bioinformatics Resource Portal (www.expasy.org). BLASTp was employed to identify sequences of H\(^+\)-ATPases from *Solanum lycopersicum*, *Brassica rapa*, and *Glycine max* with close similarity to AHA2 using ExPASy. Sequences were aligned using MUSCLE alignment software (Edgar, 2004), and pairwise distances were calculated using MEGA6 software (www.megasoftware.net). C-terminal domains were predicted using TMHMM v. 2.0 software (www.cbs.dtu.dk/services/TMHMM-2.0).

Fungal cultivation and fractionation
Several *Stemphylium* species were grown on potato dextrose agar (PDA), malt extract agar (MEA), and yeast extract with supplements (YES) plates for 10–14 days at 25°C in the dark. Fungal growth was then harvested and extracted with ethyl acetate containing 0.5% (v/v) formic acid. The extracts were concentrated *in vacuo*. *Stemphylium loti* FIP 217 was identified as the most active in the H\(^+\)-ATPase assay and was selected for further examination. A large-scale extract from *Stemphylium loti* FIP 217 was generated by cultivation of *Stemphylium loti* FIP 217 on an equal number of PDA and YES plates (100 plates in total). Fungal growth was extracted with ethyl acetate containing 1% (v/v) formic acid, and the pooled extract was concentrated *in vacuo*.

The large-scale *Stemphylium loti* FIP 217 extract was initially fractionated on diol-functionalized silica (Biotage, Uppsala, Sweden) using an Isolera autoflasher (Biotage, Uppsala, Sweden). The column was eluted stepwise using heptane, dichloromethane, ethyl acetate, and methanol in 50%
increments. The active fractions were pooled and taken through another round of fractionation using a C_{18} packed column. Fractions were eluted using a stepwise gradient of A) H_{2}O + 50 ppm trifluoroacetic acid and B) methanol + 50 ppm trifluoroacetic acid, from 10% methanol to 100% methanol in increments of 10% methanol, using an Isolera autoflasher (Biotage, Uppsala, Sweden), with each fraction being three column volumes (45 ml).

**UHPLC-DAD-HRMS**

Extracts and fractions were analyzed by ultra high-performance liquid chromatography-diode array detection-high-resolution mass spectrometry (UHPLC-DAD-HRMS) using a 1290 Infinity UHPLC machine (Agilent, Santa Clara, CA, USA). Separation was achieved on a 2.0 × 150-mm Poroshell phenyl-hexyl column (2.7 Å; Agilent Technologies, Santa Clara, CA, USA). The column was eluted with a linear gradient consisting of A) H_{2}O + 20 mM formic acid and B) acetonitrile + 20 mM formic acid starting at 10% B and increasing to 100% B over 15 min and held at this composition for 2 min before returning to initial conditions. MS detection was performed by a 6540 series quadropole time-of-flight (QTOF) mass spectrometer equipped with a dual jet stream electrospray ionization (ESI) source operating in positive mode. Data-dependent auto MS/MS acquisition was performed using fixed collision induced dissociation (CID) collision energies of 10, 20, and 40 eV. The data generated were dereplicated employing an in-house MS/HRMS natural product database (Kildgaard et al., 2014).

**Results**

**Screening for inhibitors**

We established a bioactivity-based screening procedure to identify fungal compounds targeting the plant PM H^{+}-ATPase (Fig. 1). Crude extracts from 10 Stemphylium species (here named sp1–sp10) were initially tested for their effects on PM H^{+}-ATPase activity in vitro. All metabolites were extracted using ethyl acetate, dissolved in DMSO, and screened for effects on the activity of Spinacia oleracea PM H^{+}-ATPase by quantifying ATP hydrolysis (Fig. 2). Most extracts had an inhibitory effect on H^{+}-ATPase activity, except sp8, which did not affect ATP hydrolysis. We chose the extract from sp7, Stemphylium loti, for further analysis.
To identify the inhibitory compound(s), extracts of *Stemphylium loti* were further fractionated using normal-phase (NP) and reversed-phase liquid flash. The initial extract from *Stemphylium loti* was separated into 10 fractions (fractions 1.1–1.10), each of which was then tested individually (Fig. 2b). We used three different concentrations of extract to assess whether the inhibitory effect was dose-dependent (Fig. 2b). Fractions 1.9 and 1.10 were the best candidates and were further tested for their dose dependency. These two fractions inhibited PM H\(^+\)-ATPase activity in a dose-dependent manner with IC\(_{50}\) values of 0.82 and 0.64 µg µg\(^{-1}\) protein, respectively (Fig. 3a, Table 1). Fractions 1.9 and 1.10 were further fractionated into 10 fractions (2.1–2.10) by reversed-phase HPLC. Again, we tested each fraction individually for its effect on ATP hydrolysis by the plant H\(^+\)-ATPase (Fig. 2c). The inhibitory molecules were clearly separated into fractions 2.6 and 2.7. The IC\(_{50}\) values were 0.77 and 0.33 µg µg\(^{-1}\) protein, respectively (Fig. 3b, Table 1).

We analyzed fractions 2.6 and 2.7 by HPLC-DAD-HRMS and dereplicated the data using a MS/HRMS library (Kildgaard et al., 2014). This revealed TeA to be a prevalent compound and possible candidate for PM H\(^+\)-ATPase inhibition. Commercially available TeA was purchased for further experiments. TeA inhibited PM H\(^+\)-ATPase-mediated ATP hydrolysis with an IC\(_{50}\) value of 8 ng TeA µg\(^{-1}\) protein, equivalent to 0.7 µM under these assay conditions. This corresponds to a 103-, 81-, 96-, and 41-fold increase in inhibitory effect compared to fractions 1.9, 1.10, 2.6, and 2.7, respectively (Table 1).

To confirm the inhibitory effect of TeA on the PM H\(^+\)-ATPase, we employed another assay measuring H\(^+\) pumping rate as opposed to the previously used assay measuring ATP hydrolysis. Accumulation of protons in inside-out vesicles was measured using the fluorescent probe ACMA. Upon protonation, ACMA is trapped inside the vesicles, leading to a decrease in fluorescence. We treated inside-out vesicles with TeA in the range of 0.1 to 12 µM (Fig. 4a). A clear reduction in fluorescence was observed upon TeA addition. The initial rate of proton transport into vesicles was estimated by linear regression of the decrease in fluorescence from time 0 to 120 s. The resulting slope was plotted as a function of TeA concentration (Fig. 4b). TeA concentrations of 3, 6, and 12 µM significantly inhibited proton pumping compared to the control. Taken together, these data indicate
that TeA inhibits both ATP hydrolysis and proton pumping, confirming that TeA is an inhibitor of the plant PM H⁺-ATPase \textit{in vitro}.

TeA mode of action
To characterize the mechanism of TeA inhibition, we investigated the effect of TeA on H⁺-ATPase kinetics by combining increasing concentrations of the substrate, ATP, with different concentrations of TeA (Fig. 5a). A preliminary Hanes–Woolf plot revealed similar $K_m$ values independent of TeA, while $V_{\text{max}}$ was affected in the presence of both 1 and 2.5 µM TeA. The data were fitted as a noncompetitive inhibitor using nonlinear regression. This showed that $V_{\text{max}}$ decreased from 332.4 nmol P i mg⁻¹ protein min⁻¹ to 219.3 nmol P i mg⁻¹ protein min⁻¹ and to 83.8 nmol P i mg⁻¹ protein min⁻¹ after treatment with 1 and 2.5 µM TeA, respectively (Table 2). These results indicate that TeA inhibits H⁺-ATPase activity by acting as a noncompetitive inhibitor of ATP.

To further investigate the inhibitory effect of TeA on the H⁺-ATPase, we treated \textit{Spinacia oleracea} leaves \textit{in vivo} with 5 µM TeA, or DMSO as a control, for 15 min prior to purification of PMs. \textit{In vivo} treatment with TeA did not affect $K_m$, while $V_{\text{max}}$ was decreased in the subsequent ATPase activity test (Fig. 5b), thus confirming the data obtained from the \textit{in vitro} treatment experiment (Table 3). As the observed decrease in $V_{\text{max}}$ could be due to degradation of H⁺-ATPase, PM fractions were separated on an SDS gel, transferred to a nitrocellulose membrane and the H⁺-ATPase detected by immunoblotting. The bands corresponding to the PM H⁺-ATPase were detected around 100 kDa. No degradation was observed in the TeA-treated sample compared to the control (Fig. 5c). Also, the degree of Threonine phosphorylation was tested. Also no difference was observed between the two samples. These results indicate that the observed decrease in $V_{\text{max}}$ is not caused by protein degradation, nor by posttranslational modification of the Thr-residue in the very C-terminal end, but rather by a TeA-mediated inhibition of the H⁺-ATPase.

Effect of TeA on \textit{Arabidopsis thaliana} root growth
In order to investigate the effect of TeA on plant growth, a growth experiment was set up in which 6-days old \textit{Arabidopsis thaliana} seedlings were moved to substrates containing 0-20 µM TeA for another six days of growth. TeA treatment resulted in reduced root growth on all concentrations, with
20 μM being the most effectful (Fig 6a, b). Root growth was significantly different already at day 2 on all concentrations employed (Fig. 6a). Seedlings treated with TeA exhibited fewer side roots and the abaxial side of leaves were purple due to stress, compared to controls (Fig. 6b). A close look at the epidermis revealed an increased number of root hairs at 5μM TeA, but at 20 μM the cells in the elongation zone looked small and collapsed, compared to the normally elongated form of the cells.

Live imaging of TeA effects on root cells
As the next step, we wanted to investigate the inhibitory effect of TeA on the cellular level. Seedlings expressing the ratiometric fluorescent pH biosensor apo-pHusion, which is targeted to the apoplast (Gjetting et al. 2012), were used in a perfusion assay for live imaging of four- to five-day-old Arabidopsis thaliana seedlings. Apoplastic pH was measured in elongating root cells as changes in the ratio of EGFP/mRFP1 intensity in response to treatment with either 10 μM TeA or bath solution (control). TeA treatment resulted in an apoplastic alkalinization starting after ~90 s of treatment, peaking after ~ 3–4 min, and lasting for the entire 10 min of the experiment (Fig. 7a). The apoplastic pH increased with 0.1 ratiometric unit indicating a pH change larger than 1 pH unit (see insert). In addition, root cells perfused with TeA appeared to accumulate fluorescent protein in fragmented endoplasmic reticulum compartments after ~ 5 min, and this accumulation increased in intensity over the course of the experiment (Fig. 7b, t = 10:00).

TeA treatment of plant material interferes with fusicoccin activation of PM H⁺-ATPases
The fungal toxin FC is commonly used to study H⁺-ATPase activation as it stabilizes binding of activating 14-3-3 protein to the C-terminus of the plant H⁺-ATPase. In vivo treatment with TeA suggests that H⁺-ATPases inactivated by TeA are not degraded, but rather permanently inactivated. To investigate the mechanism of binding between TeA and the H⁺-ATPase, we initially confirmed that FC and 14-3-3 together activate the PM H⁺-ATPase (Fig. 8a). We then incubated PM fractions with 20 μM FC + 14-3-3 (1 μg) for 10 min and measured ATP hydrolysis at increasing TeA concentrations (Fig. 8b). A clear effect of TeA was observed, as TeA was able to reverse the activated conformation induced by FC. The data were analyzed using nonlinear regression, revealing an IC₅₀ value of 5.4 μM, significantly higher than 0.7 as obtained in the original material.
Similarly, we pre-incubated PM fractions with 20 µM TeA and 14-3-3 protein for 10 min and measured ATP hydrolysis at increasing FC concentrations (Fig. 8c). Previous experiments suggested significant TeA-induced inhibition of H⁺-ATPase activity after 10 min of treatment, and without added FC, we observed little to no Pi in the ATP hydrolysis assay (10 nmol Pi mg⁻¹ min⁻¹). However, increasing the concentration of FC reduced the inhibition, although specific activity was very low compared to that in FC-treated PM fractions in the absence of TeA (Fig. 8a,b,c).

Taken together, these results indicate that TeA effectively inhibits the plant PM H⁺-ATPase even when the regulatory domain is displaced by association with 14-3-3 protein.

The plant PM H⁺-ATPase regulatory C-terminus is required for TeA inhibition.

TeA inhibition of the plant PM H⁺-ATPase was dose-dependent with an IC₅₀ value of 8 ng (Fig. 3c, f), equivalent to 0.7 µM (Table 4, Fig. 8a). We previously demonstrated that TeA-induced inhibition is noncompetitive with respect to ATP both in vitro and in vivo (Fig. 5). The C-terminus of plant H⁺-ATPases is a well-described autoregulatory domain, where both activation and inhibition are determined by phosphorylation of specific residues (Fuglsang et al., 2007), leading to binding of regulatory proteins (Fuglsang et al., 2003, 2007; Yang et al., 2010). Consequently, we sought to determine the importance of the C terminus in TeA-induced inhibition of the H⁺-ATPase. Wild-type and C-terminally truncated mutants of the well-studied PM H⁺-ATPase from Arabidopsis, AHA2, were heterologously expressed in Saccharomyces cerevisiae. We purified membranes from Saccharomyces cerevisiae containing aha2Δ92, aha2Δ77, aha2Δ40 and wild type AHA2. All membranes were assayed for ATP hydrolysis in response to increasing TeA concentration (Fig. 9). TeA inhibited AHA2 activity (Fig. 9a) in a comparable manner to what we observed in plant derived material (Fig. 9b), but with a much higher IC₅₀ value. In contrast, TeA was not able to inhibit ATP hydrolysis of recombinant aha2Δ92 expressed in Saccharomyces cerevisiae (Table 4), even at concentrations of 50 µM TeA. The same tendency was observed for aha2Δ77, indicating that TeA inhibition is dependent on the presence of the regulatory C-terminus. TeA inhibited both AHA2 and aha2Δ40, but with a much higher IC₅₀ value (14.3 and 3.7 µM (Table 4)) than observed in plant tissue. This could indicate that binding of TeA to the H⁺-ATPase is sensitive to the protein conformation which might change due to the different lipid environment in the two organisms.
Another interesting observation is that at low concentrations of TeA, a slight activation is observed for some of the H\(^+\)-ATPases (Fig 9e and f; aha2\(\Delta\)40 and PMA1). The PM H\(^+\)-ATPase of *Saccharomyces cerevisiae*, PMA1, and the plant PM H\(^+\)-ATPase are structurally comparable (Morth *et al.*, 2011), although the C-terminal regulatory domains are markedly different. The C-termini of PMA1 and AHA2 were predicted to contain 43 and 111 amino acid residues, respectively. The PMA1 C-terminal has two phosphorylation sites, Ser911 and Thr912 (Lecchi *et al.*, 2007), whereas the C-terminus of the *Arabidopsis thaliana* H\(^+\)-ATPase, AHA2, has seven reported phosphorylation sites (Rudashevskaya *et al.*, 2012), (Haruta & Constabel, 2003; Fuglsang *et al.*, 2014). We purified PM fractions from *Saccharomyces cerevisiae* expressing PMA1 and analyzed ATP hydrolysis at increasing TeA concentrations. TeA inhibition of PMA1 was in the same range as when AHA2/aha2\(\Delta\)40 was expressed in yeast. For PMA1 IC\(_{50}\) was found to be 7.8 \(\mu\)M TeA compared to 14.33 and 3.7\(\mu\)M for AHA2 and aha2\(\Delta\)40 respectively. Again this is higher than what is found in plant material (Fig. 9b, f, Table 4).

Taken together, these results indicate that TeA interacts with the regulatory C-terminus of the plant H\(^+\)-ATPase to inhibit activity and that the length of the C-terminus is important for this interaction. Furthermore, the data also indicate that the lipid composition of the membrane is of importance for the effect of TeA.

Sequence differences between H\(^+\)-ATPases from plants and fungi

The present data highlight the importance of the C-terminus of the plant H\(^+\)-ATPase in TeA-induced inhibition. We speculate that PMA1 is less sensitive to TeA because its regulatory domain differs from plants. We analyzed the amino acid sequence of AHA2 to identify highly similar sequences from *Solanum lycopersicum*, *Brassica rapa*, and *Glycine max* for alignment. Likewise, we used the amino acid sequence of PMA1 to identify highly similar sequences from *Stemphylium lycopersici*, *Alternaria alternata*, and *Fusarium oxysporum* for alignment (Fig 10). This revealed that the C-termini of all plant H\(^+\)-ATPases are highly conserved, whereas the fungal C-termini are less conserved. Besides the differences in length, few similarities were observed among fungal H\(^+\)-ATPases, and we expect that posttranslational regulation is essential for both plant and fungal H\(^+\)-ATPases.
Pairwise distance, a measure of the number of substitutions per site between sequences, was calculated using MEGA6 software to assess differences in amino acid composition between pairs of sequences. The pairwise distance between AHA2 and corresponding sequences from Solanum lycopersicum, Brassica rapa, and Glycine max was 0.104, 0.118, and 0.121, respectively, confirming that very few amino acids differ between these sequences. Conservation of the regulatory domain in an essential enzyme highlights its importance in normal cell function, and we speculate that activity regulation through posttranslational modulation of the H^+-ATPase C-termini is shared among the plants examined.

In order to test if TeA affects fungal growth, Saccharomyces cerevisiae expressing either PMA1 or AHA2 were incubated with increasing concentrations of TeA and the growth was monitored for 48 hrs. No effect was found (data not shown). This result further supports that the ability of TeA to reach the fungal H^+-ATPase is limited compared to the possibility of reaching the plant H^+-ATPase when inserted in native membranes.

Discussion

Stemphylium spp. invade numerous plant species, and cause leaf spots, ultimately leading to decreased yield and accumulation of mycotoxins in harvested crops (Hanse et al., 2015). In this study, we screened extracts from 10 different Stemphylium species for their effect on plant PM H^+-ATPase activity, to investigate whether this essential enzyme was targeted by any of the fungi. PM fractions were purified from Spinacia oleracea leaves using the aqueous two-phase partitioning. Extracts from nearly all screened Stemphylium species inhibited ATP hydrolysis to various extents, suggesting that several species modulate host H^+-ATPase activity upon pathogen attack.

We further separated and screened the Stemphylium loti extract for its effect on ATP hydrolysis and identified TeA as the main compound responsible for H^+-ATPase inhibition (Fig. 3). TeA also inhibited proton pumping, confirming TeA as a plant H^+-ATPase inhibitor (Fig. 4). Identification of TeA in Stemphylium loti extracts contrasts with a recent publication by Olsen et al. (2018) claiming that Stemphylium loti does not produce TeA; however, revisiting their data revealed TeA to be a highly abundant component in all four extracts of Stemphylium loti screened (Olsen, KJK, pers. comm., 2018).
TeA blocking the electron transport chain of PSII is proposed as the main mechanism of TeA action in phytotoxicity (Chen et al., 2007, 2010, 2015). TeA inhibits the O$_2$ evolution rate of PSII in isolated thylakoid membranes when treated with as little as 1 μM TeA; however, even 2 mM TeA is not able to inhibit O$_2$ evolution completely (Chen et al., 2007), suggesting that the effect does not result from a high-affinity interaction. Mesophyll cells in epidermal segments treated with 250 and 500 μM TeA for 30 min to 6 h show clear accumulation of reactive oxygen species (ROS) leading to cell death after prolonged exposure (Chen et al., 2010). This requirement for long exposure and high concentrations contrasts with our results. We observed an effect on the plant PM H$^+$-ATPase after in vivo treatment with 5 μM TeA for 15 min and determined an IC$_{50}$ value of only 0.7 μM TeA in our ATP hydrolysis assay. Prolonged exposure to TeA will inhibit transfer of electrons from QA to QB; however, our present study suggests that an earlier response to TeA is the inhibition of the PM H$^+$-ATPase. Considering the low IC$_{50}$ value of TeA on plant PM H$^+$-ATPases (0.7 μM), we expect that most H$^+$-ATPases will be effectively inhibited in plant tissue treated with 250 μM TeA. Effective inhibition of proton transfer across the PM leads to depolarization and eventually necrosis (Golstein & Kroemer, 2007). When one cell suffers from necrosis, cell debris will spread to neighboring cells causing increased ROS (Apel & Hirt, 2004). The ROS accumulation observed in response to TeA by Chen and coworkers (2010) may therefore be explained by necrosis due to severe membrane depolarization, rather than blocking of the electron transport chain.

A decrease in ATP hydrolysis and proton pumping could be caused by protein degradation, and TeA is reported to inhibit the release of newly synthesized proteins (Shigeura & Gordon, 1963). To rule out protein degradation as the cause for decreased H$^+$-ATPase activity, we pretreated leaves of Spinacia oleracea with 5 μM TeA, for 15 min prior to PM purification. Immunobloting analysis revealed that TeA pretreatment did not alter H$^+$-ATPase levels in PM fractions nor their phosphorylation status (Fig. 5c). We also investigated the effect of TeA on the kinetic constants $V_{\text{max}}$ and $K_m$. Pretreatment of Spinacia oleracea with TeA prior to PM purification resulted in a decrease in $V_{\text{max}}$ from 223.1 to 165.8 ng Pi mg$^{-1}$ protein min$^{-1}$ compared to the control. The ATP dose necessary to reach half of $V_{\text{max}}$ and $K_m$ remained largely unaffected, suggesting that TeA does not compete for the ATP-binding site of the PM H$^+$-ATPase.
In order to visualize the effect of TeA on plant growth, the root length of *A. thaliana* seedlings grown on TeA was measured. The effect of TeA was apparent already on day 2 of growth with TeA. After day 6, root length of seedlings treated with 2.5 μM TeA was on average 1.6 cm shorter than non-treated seedlings. Treatment with 20 μM TeA resulted in little to no growth, with average growth 0.1 cm after 6 days. This suggests that TeA is a potent toxin, which can be used to inhibit plant growth.

To confirm that TeA is a H⁺-ATPase inhibitor *in planta*, we employed a live-imaging method using the apoplastic pH sensor apo-pHusion (Gjetting et al., 2012). Apoplastic pH increased within 2 to 4 min after treatment with 10 µM TeA (Fig. 8). By calibrating the change in fluorescence to known pH values, we estimated the increase in apoplastic pH to be from ~5 to 7 after TeA treatment. Intracellular pH is approximately pH 7 in healthy plant cells, indicating that TeA treatment effectively eliminates the electrical gradient across the PM. Apoplastic pH did not return to initial pH for the duration of these experiments, suggesting an irreversible inactivation of H⁺-ATPases. TeA is therefore a potent inhibitor of H⁺-ATPase *in planta*.

PM H⁺-ATPase activation by 14-3-3 includes dislocation of the C-terminal regulatory domain and is dependent on phosphorylation of the penultimate threonine residue (Fuglsang et al., 1999). The fungal toxin FC stabilizes complex formation between the C-terminal domain and the 14-3-3 protein (Fuglsang et al., 2003; Würtele et al., 2003). By pretreating PM fractions for 10 min with FC and 14-3-3 protein, before TeA, we demonstrated that TeA can” overcome” the stable binding complex between the C-terminus, 14-3-3 and FC. The IC₅₀ value of TeA on FC-activated PM H⁺-ATPase fractions was 5.4 μM. For untreated PM, the IC₅₀ value of TeA was 0.7 μM, indicating that FC/14-3-3 protein treatment increased the amount of TeA required to reach half-inhibition by 7.8-fold. Conversely, FC/14-3-3 protein was not able to reciprocally reactivate PM fractions incubated with 20 μM TeA for 10 min prior to assay start. In the absence of FC and 14-3-3 protein, we observed a specific activity of 10.1 P₁ mg⁻¹ protein min⁻¹, which was increased to 44.1 P₁ mg⁻¹ protein min⁻¹ when treated with 50 μM FC and 14-3-3 protein (1 μg μl⁻¹). This demonstrates that PM H⁺-ATPases inhibited by TeA to only a small degree can be reactivated by FC/14-3-3 protein, although this
requires an elevated dose. These data suggest that the regulatory domain is not only involved in FC binding, but also is important for TeA-induced inhibition. The *Arabidopsis thaliana* H⁺-ATPase, AHA2, is permanently activated by removal of 77 or 92 amino acid residues of the C-terminal regulatory domain (Palmgren et al., 1990; Regenberg et al., 1995). Interestingly, we did not observe TeA-mediated inhibition of *aha2Δ92* or *aha2Δ77*, confirming the importance of the regulatory domain in TeA-mediated H⁺-ATPase inhibition. The H⁺-ATPase native to *Saccharomyces cerevisiae*, PMA1, responded to TeA treatment but required a higher dosage compared to the H⁺-ATPase protein purified from *Spinacia oleracea*. Interestingly, expression and purification of AHA2 in *Saccharomyces cerevisiae* led to an increase in the IC₅₀ value, reaching a value comparable to the one found for PMA1. Similarly, activity of *aha2Δ40*, where only 40 C-terminal amino acid residues were removed, was not affected by TeA to the same degree as the *Spinacia oleracea* H⁺-ATPase. This indicates that not only is the C-terminal, but also the chemical environment important for TeA inhibition. When AHA2 is inserted into yeast membranes the accessibility of TeA seems to be significantly reduced. One can imagine that TeA binds to a site generated between the body of the protein and is stabilized by the C-terminus. We have previously demonstrated that lysophospholipids can activate the plant PM H⁺-ATPase, whereas they do not have any effects on the fungal Pm H⁺-ATPase (Wielandt et al., 2015). Additionally, this regulation involved the regulatory C-terminal domain. The regulatory domains of PMA1 and *aha2Δ40* are of comparable length but do not show any sequence similarity, which might explain the difference in observed IC₅₀ values. Based on these data, we suggest that inhibition by TeA requires the full regulatory domain of the plant H⁺-ATPase.

Aligning the C-terminal regulatory domains from H⁺-ATPases of the plants *Arabidopsis thaliana*, *Solanum lycopersicum*, *Brassica rapa*, and *Glycine max* confirmed the very high degree of conservation of the PM H⁺-ATPase between the four species (Fig.10). This allows phytotoxins to be functional against many host plant species by targeting the regulatory domain to modulate plant H⁺-ATPase activity. *Stemphylium* species are reported to target many different plant species, and targeting this highly conserved domain provides an opportunity to cause necrosis in many plants using the same metabolite.
To further illustrate that *Stemphylium loti* can target plant H⁺-ATPases without affecting the activity of the endogenous H⁺-ATPase, we aligned the amino acid sequences of H⁺-ATPases from four plant species with H⁺-ATPase sequences from four fungi. The C-termini of the plant PM H⁺-ATPases were predicted to be 111–114 amino acids long, while the C-termini from *Saccharomyces cerevisiae*, *Stemphylium lycopersici*, *Alternaria alternata*, and *Fusarium oxysporum* H⁺-ATPases were predicted to contain only 13–46 amino acids.

In addition to the theory that the plant C-terminus is important for the binding, also the conformational change between E1 to E2 might play a role for binding of TeA. The binding site might be more exposed in one conformation as compared to another. Here we can speculate that a compound only binds when the binding site is exposed, which could be when the protein is in the E2 formation.

In conclusion, the fungus *Stemphylium loti* produces the phytotoxin TeA that specifically targets the plant PM H⁺-ATPase. We demonstrated through extensive biochemical evaluation and direct visualization using apo-pHusion plants that TeA inhibits the plant PM H⁺-ATPase. Inhibition of the PM H⁺-ATPase results in depolarization of the membrane potential and eventually necrosis. The corresponding fungal H⁺-ATPase, PMA1, is much less affected, and we propose that this is due to structural differences in the C-terminal regulatory domains, but, just as importantly, also due to the different lipid composition of the plasma membrane of fungi compared to plants. By utilizing these differences, fungi are able to target an essential plant enzyme without causing self-toxicity.

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**Author contributions**

S.R. and T.I.: fungal growth and compound purification. P.B. and J.I.: biochemical screening. P.B. and N.H.: biochemical characterization of TeA. S.G: bioimaging of TeA effect *in planta*. N.H: Tea
effect on plant growth. P.B.: sequence alignment. T.L. and A.F. conceived and coordinated experiments. P.B.: first draft of the manuscript with input from S.R., T.L. and A.F.

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Figure legends:

Fig 1. Screening of fungal extracts for their effect on ATPase enzyme activity. Fungi were grown on plates, harvested and metabolites extracted. The extracts were added to plant H⁺-ATPase assays in order to identify effectors of the PM H⁺-ATPase. Fractions exhibiting a strong effect on the H⁺-ATPase were further separated and re-tested. This procedure was repeated until a relatively clean target were identified.
Fig. 2 H⁺-ATPase activity of Spinacia oleracea plasma membranes in the presence of metabolites from Stemphylium spp. (a) Addition of total metabolites (30 µg) from 10 different Stemphylium spp., (b) Stemphylium loti extracts (from panel a) fractionated into 10 fractions (1.1–1.10), and (c) fractionation of fractions 1.9 and 1.10 (from panel b) into 10 new fractions, 2.1–2.10. Fractions 1.1-1.10 was added as 0.02 (white bars), 0.16 (blue bars), 2.5 (black bars), and 20 µg µg⁻¹ protein (red bars). Fractions 2.1-2.10 was added as 0.01 (white bars), 0.2 (blue bars), 1.5 (black bars), and 11.5 µg µg⁻¹ protein (red bars). Values are means ± SEM. Data were analyzed by two-way analysis of variance (ANOVA), and H⁺-ATPase activity was compared to H⁺-ATPase activity of the control (no addition of extracts, not shown) with a Bonferroni posttest. *, P < 0.05; **, P < 0.01; ***, P < 0.001; n = 3.

Fig. 3 Dose-dependent inhibition of H⁺-ATPase activity measured in Spinacia oleracea plasma membranes (PMs) in response to tenuazonic acid (TeA) treatment. Values are means ± SEM. Experiments with Stemphylium loti fractions 1.9 and 1.10 (a), and 2.6 and 2.7 (b) were carried out in triplicate (n = 3), while experiments with TeA (c) were carried out in triplicate and on three independent PM fractions (n = 9). (d, e, f) Data were analyzed using nonlinear regression and fitted to log[inhibitor] (µg) versus the normalized response for determination of IC₅₀ values.

Fig. 4 Tenuazonic acid (TeA) effect on H⁺ pumping of Spinacia oleracea plasma membrane (PM) vesicles. (a) Accumulation of protons in inside-out PM vesicles of Spinacia oleracea treated with increasing concentrations of TeA visualized and quantified using the fluorescent probe ACMA. Addition of MgSO₄ initiates ATP-stimulated proton pumping into vesicles, while addition of nigericin releases the trapped protons by acting as a proton ionophore. Values are means ± SEM based on two technical replicates (n = 2) and are representative of two independent PM purifications. (b) The initial slope of each curve (from t = 0 s to t = 120 s) was estimated by linear regression and plotted as a function of TeA concentration (µM). Values are means ± SEM based on two technical replicates and are representative of two independent PM fractions. Data were analyzed using one-way analysis of variance (ANOVA), and Dunnett’s multiple comparisons test was used to calculate the difference in
slopes compared to the control: ***, $P < 0.001$.

**Fig. 5** Kinetic analysis of *Spinacia oleracea* plasma membrane (PM) H$^+$-ATPase inhibition by tenuazonic acid (TeA). Specific activity was plotted as a function of ATP concentration (mM) at the indicated concentrations of TeA. (a) *In vitro* treatment. (b) *In vivo* treatment of *Spinacia oleracea* leaves, 5 µM TeA or DMSO (control) for 15 min before purification of plasma membranes. Values are means ± SEM of four technical replicates of two independent PM samples ($n = 8$). (c) Immunodetection of PM H$^+$-ATPase and Phospho-Threonine in the plasma membrane samples (20 µg per lane). +/- indicates pretreatment with TeA or not.

**Fig. 6** TeA effect on *Arabidopsis thaliana* root growth. Seeds (Col-0) were germinated and grown for 6 days on ½MS, before transfer to ½MS agar containing 0, 2.5, 5, 10, 20 µM TeA respectively and further grown for 6 days. (a) Root length of 12-day old seedlings was measured on day 0, 2, 4, 6 after transfer to plates with TeA. Values are given in means ±SEM, $n = 32-48$. Root lengths of treated plants were highly significant (***, $P < 0.001$) for all treatments compared to control after 2 days growth with TeA. (b) Images of 12-day old seedlings.

**Fig. 7** *In vivo* tenuazonic acid (TeA)-induced alkalization of the apoplast in elongating root cells of four- to five-day-old *Arabidopsis thaliana* roots expressing apo-pHusion. (a) Seedlings were mounted with agar on a Teflon-coated microscope slide and covered with a droplet of bath solution. Root cells in the elongation zone were viewed on an SP5-X confocal laser scanning microscope, using a ×20 dipping objective and a perfusion setup for addition of TeA. After c. 2 min, 500 µl of 10 µM TeA (black curve) or bath solution (control, red curve) was added (black arrow) and the experiment was run for 10 min in total. Normalized ratio measurements of EGFP/mRFP1 are given as means ± SEM of $n = 4$ replicates. pH changes in response to TeA treatment were highly significant (***, $P < 0.001$) in comparison with controls (Student’s t-test). Inset shows *in planta* pH calibration curve using 10 mM MES, 10 mM MOPS, and 10 mM citrate buffer with pH adjusted to pH 5–7. (b) Confocal overlay images of EGFP (green) and mRFP1 (red) channels at three timepoints during the experiment: $t = 0$ before addition of TeA, $t = 4:30$ at the peak of the response, and $t = 10:00$ after 10 min. Scale
bar: 50 µm.

**Fig. 8** Competition between Fusicoccin (FC) and tenuazonic acid. (a) Activation of *Spinacia oleracea* PM H⁺-ATPase preincubated with either FC, 14-3-3 or FC/14-3-3 prior to assay start. (b) Dose-dependent TeA inhibition of *Spinacia oleracea* PM H⁺-ATPase preincubated with FC and 14-3-3 protein prior to assay start. (c) Dose-dependent FC/14-3-3 activation of *Spinacia oleracea* PM H⁺-ATPase preincubated with 20 µM TeA prior to assay start. (b, c) Data were analyzed using nonlinear regression and fitted to log[agonist] (µM) versus the normalized response (variable slope) to determine the half maximal effective concentration (EC₅₀) value. FC (10 µM) and 14-3-3 (1 µg) were used in all treatments. Treatments with 14-3-3 alone or FC/14-3-3 were significant different from non-treated (**, P < 0.01;***, P < 0.001). Experiments were performed in triplicates of two independent PM fractions, and values represent means ± SEM (n = 6).

**Fig. 9** The effect of tenuazonic acid (TeA) on plasma membrane (PM) H⁺-ATPases with C-terminal deletions. TeA-induced inhibition of H⁺-ATPase activity measured on (a) AHA2 expressed in *Saccharomyces cerevisiae*, (b) PMs from *Spinacia oleracea*, (c, d, e) C-terminally truncated versions of AHA2; *aha2Δ92* (c), *aha2Δ77*(d) and *aha2Δ40* (e), all expressed in *Saccharomyces cerevisiae*, (f) endogenous *Saccharomyces cerevisiae* H⁺-ATPase PMA1. ATP hydrolytic activity experiments were performed in triplicates of three independent fractions (n = 9), except for AHA2 and *aha2Δ77* (two independent fractions, n = 6). Values represent means ± SEM. Data were analyzed using nonlinear regression and fitted to log[TeA] versus the response (variable slope) (constrains bottom =0.0) to determine the IC₅₀ values.

**Fig. 10** Alignments of amino acid sequences of the C termini of H⁺-ATPases from the plants *Arabidopsis thaliana, Solanum lycopersicum, Brassica rapa*, and *Glycine max*, and the fungi *Saccharomyces cerevisiae, Stemphylium lycopersici, Alternaria alternata*, and *Fusarium oxysporum*. Numbers indicate length of the C terminus, and colors indicate conserved amino acids.
Table 1 IC$_{50}$ values of *Stemphylium loti* fractions 1.9, 1.10, 2.6, and 2.7 and tenuazonic acid (TeA) in ATP hydrolysis assays using *Spinacia oleracea* plasma membrane H$^+$-ATPases

| *Stemphylium loti* extract | IC$_{50}$ (µg µg$^{-1}$ protein) | Ratio: IC$_{50}$ of extract/IC$_{50}$ of TeA |
|---------------------------|----------------------------------|------------------------------------------|
| Fraction 1.9              | 0.82 (0.69–0.97)                 | 102.7                                    |
| Fraction 1.10             | 0.65 (0.55–0.76)                 | 80.7                                     |
| Fraction 2.6              | 0.77 (0.68–0.87)                 | 96.3                                     |
| Fraction 2.7              | 0.33 (0.28–0.38)                 | 40.9                                     |
| TeA (commercial)          | 0.008 (0.007–0.009)              | 1                                        |

ATP hydrolysis was analyzed using nonlinear regression and fitted to log[inhibitor] versus the normalized response to determine IC$_{50}$ values. For *Spinacia oleracea* PMs treated with fractions 1.9, 1.10, 2.6, and 2.7, $n = 3$. For *Spinacia oleracea* PMs treated with TeA, $n = 9$. 

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Table 2 Effect of tenuazonic acid (TeA) on $K_m$ and $V_{\text{max}}$ of the *Spinacia oleracea* plasma membrane (PM) H⁺-ATPase

| Tenuazonic acid (μM) | $K_m$ (mM ATP) | $V_{\text{max}}$ (nmol P₁ mg⁻¹ min⁻¹) |
|---------------------|---------------|-------------------------------------|
| Control             | 1.15 (0.59 to 1.71) | 332.4 (262.9 to 401.8) |
| 1.0                 | 2.56 (0.74 to 4.38)  | 219.3 (132.5 to 306.1) |
| 2.5                 | 10.96 (0.0 to 37.0)  | 83.8 (−76.6 to 244.2)  |

ATP hydrolysis of *Spinacia oleracea* PM fractions was measured at different ATP concentrations in response to various TeA doses, and enzyme kinetic constants, $K_m$ and $V_{\text{max}}$, were determined from nonlinear regression of the Michaelis–Menten equation. Values are means ± SEM of four technical replicates of two independent PM fractions ($n = 8$).
Table 3 Kinetic parameters, $K_m$ and $V_{max}$, for *Spinacia oleracea* plasma membrane (PM) H$^+$-ATPases in response to tenuazonic acid (TeA) pretreatment

|          | $K_m$          | $V_{max}$         |
|----------|----------------|-------------------|
| Control  | 0.53 (0.21–0.86) | 223.1 (178.0–268.2) |
| TeA 5 μM | 0.48 (0.02–0.93) | 165.8 (115.1–216.5) |

ATP hydrolysis in *Spinacia oleracea* PM fractions purified from leaves that had been pretreated with 5 μM TeA was measured at various ATP concentrations. The enzyme kinetic constants, $K_m$ and $V_{max}$, were determined from nonlinear regression of the Michaelis–Menten equation. Values are means ± SEM of four technical replicates of two independent PM fractions ($n = 8$).
Table 4 IC₅₀ values of tenuazonic acid (TeA) for *Spinacia oleracea* H⁺-ATPase, *Saccharomyces cerevisiae* PMA1, and truncated *Arabidopsis thaliana* H⁺-ATPases aha2Δ92 and aha2Δ40

|                       | IC₅₀ (µM)         |
|-----------------------|------------------|
| *Spinacia oleracea* H⁺-ATPase | 0.68 (0.59–0.78) |
| AHA2 (expressed in yeast) | 14.33 (12.0–17.1) |
| PMA1                  | 7.8 (5.8–10.4)   |
| aha2Δ92               | N.D              |
| aha2Δ77               | N.D              |
| aha2Δ40               | 3.7 (2.7–5.0)    |

ATP hydrolysis of plasma membrane (PM) fractions from *Spinacia oleracea* and *Saccharomyces cerevisiae* expressing PMA1, aha2Δ92, aha2Δ77 and aha2Δ40 or ER enriched fraction with AHA2, all treated with increasing concentrations of TeA. Data were analyzed using nonlinear regression and fitted to log[TeA] versus the response (variable to slope) (bottom constrain =0.0) to determine the IC₅₀ values. N.D. not determined as data could not be fitted to the model.
Screening of Stemphylium spp. PM H^+\text{\textsuperscript{\text{ATPase activity (Percentage of control)}}}

Fractionation of S. loti extract

Fractionation of 1.9 and 1.10
(a) Relative fluorescence (%)

- Control
- 0.4 μM
- 6 μM
- 12 μM

Time (s)

(b) Slope

TeA (μM)

MgSO₄
Nigericin

TeA (μM)

***

-0.5
-0.4
-0.3
-0.2
-0.1
-0.0
0.0

0.1
1.0
10.0

0.1
1.0
10.0

-0.5
-0.4
-0.3
-0.2
-0.1
-0.0
0.0

-0.5
-0.4
-0.3
-0.2
-0.1
-0.0
0.0

-0.5
-0.4
-0.3
-0.2
-0.1
-0.0
0.0

TeA (μM)
(a) 

[Graph showing PMH+ ATPase activity vs. [mM] ATP for Control, 1 μM TeA, and 2.5 μM TeA.]

(b) 

[Graph showing PMH+ ATPase activity vs. [mM] ATP for Control and 5 μM TeA.]

(c) 

[Western blot images showing anti-AHA2 central domain and anti-P-Thr.]
Figure (a) shows the norm. ratio (EGFP/mRFP1) over time (t) under different pH conditions (pH=7, pH=5). The black dots represent the treatment with 10 µM TeA, while the red dots represent the control. The inset graph highlights the pH effects on the norm. ratio eGFP/mRFP1.

Figure (b) illustrates the fluorescence images at different time points (t=0, t=4:30, t=10:00).

The graph indicates a significant difference in the norm. ratio (***).
(a) AHA2

PM H^+-ATPase activity (Percentage of control) vs. log[TeA] (μM)

IC₅₀ 14.3

(b) S. oleracea H^+-ATPase

PM H^+-ATPase activity (Percentage of control) vs. log[TeA] (μM)

IC₅₀ 0.7

(c) aha2Δ92

PM H^+-ATPase activity (Percentage of control) vs. log[TeA] (μM)

(d) aha2Δ77

PM H^+-ATPase activity (Percentage of control) vs. log[TeA] (μM)

IC₅₀ 3.6

(e) aha2Δ40

PM H^+-ATPase activity (Percentage of control) vs. log[TeA] (μM)

(f) PMA1

PM H^+-ATPase activity (Percentage of control) vs. log[TeA] (μM)

IC₅₀ 7.8
### C termini of plant H⁺-ATPases

| Species       | Sequence                        |
|---------------|---------------------------------|
| A. thaliana   | MLTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |
| S. lycopersicum | LMTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |
| B. rapa       | MLTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |
| G. max        | LMTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |

### C termini of fungal H⁺-ATPases

| Species       | Sequence                        |
|---------------|---------------------------------|
| S. cerevisiae | MLTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |
| S. lycopersici| MLTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |
| A. alternata  | MLTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |
| F. oxysporum  | MLTHAIKVLSSVSVVHVGMGLDILSTFSTVT  |