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The isoelectric point of proteins influences their translocation to the extrahaustorial matrix of the barley powdery mildew fungus

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ABSTRACT

Many biotrophic fungal plant pathogens develop feeding structures, haustoria, inside living plant cells, which is essential for their success. Extrahaustorial membranes (EHMs) surround haustoria and delimit the extrahaustorial matrices (EHMxs). Little is known about transport mechanisms across EHMs and what properties proteins and nutrients need in order to cross these membranes. To investigate this further, we expressed fluorescent proteins in the cytosol of infected barley leaf epidermal cells after particle bombardment and investigated properties that influenced their localization in the powdery mildew EHMx. We showed that this translocation is favoured by a neutral isoelectric point (pI) between 6.0 and 8.4. However, for proteins larger than 50 kDa, pI alone does not explain their localization, hinting towards a more complex interplay between pI, size and sequence properties. We discuss the possibility that an EHM translocon is involved in protein uptake into the EHMx.

KEYWORDS

Powdery mildew fungus, extrahaustorial membrane, extrahaustorial matrix, protein translocation

INTRODUCTION

Specialization and adaptation ensure survival in nature. In some cases, the specialization and adaptation of a pathogen to a specific plant niche is so high that living without it is no longer possible. This so-called obligate biotrophic interaction is true for the powdery mildew fungus, *Blumeria graminis* f. sp. *hordei* (*Bgh*), which colonizes barley (*Hordeum vulgare*) and can give a yield loss of 10-20% (<https://www.cabi.org/isc/datasheet/22075>). In order to survive, the fungus establishes an intimate symbiosis with the host plant, involving bidirectional transport across membranes allowing communication, manipulation and delivery of molecules. After successful penetration, a fungal feeding structure, the haustorium, is formed inside the host cell. This differentiated hypha becomes surrounded by a host-generated extrahaustorial membrane (EHM) that possesses endoplasmic reticulum (ER) membrane-like properties (Kwaaitaal et al. 2017). Between the haustorium and the EHM, a compartment called the extrahaustorial matrix (EHMx) is located. In contrast to the membranes and organelles in very close proximity to the EHM on the side of the host (Kwaaitaal et al 2017), transmission electron micrographs have shown the EHMx to be easily discernible and up to 3 µm wide (Hippe-Sanwald et al., 1992). Interestingly, the chloroplast protein, PSI-D, was localized in haustoria of *Erysiphe pisi* growing on pea, thus suggesting protein transport across both the EHM and the fungal plasma membrane (Testut, Callow, & Green, 1999). Otherwise, molecular and cellular information available on the haustorial complex is very sparse, despite the obvious importance of this compartment. Therefore, any novel information about it will be very useful as it forms the basis for further studies that eventually could lead to design of disease resistance.

In order to manipulate plant immunity, effector molecules are likely to be transported from the fungus to the plant cell (Lu et al., 2016). The *Bgh* genome encodes

approximately 500 so-called candidate secreted effector proteins (CSEPs) (Pedersen et al., 2012), for some of which there is evidence that they contribute positively to the success of the fungus (e.g. Aguilar et al. 2016; Ahmed et al. 2015; Pennington et al. 2016).

The present study was initially meant to analyse the subcellular localization of CSEPs in barley leaf epidermal cells. Serendipitously, however, our observations instead confirmed our previous finding (Ahmed et al., 2015) that proteins often are translocated from the host cytosol, across the EHM, into the EHMx. Being an entirely unstudied phenomenon, we looked further into it and found interesting details on protein properties influencing uptake into the EHMx. We found that neutral pI is an essential property for proteins to accumulate in the EHMx. However, protein size and sequence also influence uptake.

RESULTS

Extrahaustorial matrix localization of proteins expressed in the plant cytosol is influenced by their pI

We have previously observed that free monomeric (m)Cherry, free monomeric Yellow Fluorescent Protein (mYFP) and mYFP-CSEP0162 (without signal peptide for secretion), expressed in the cytosol of barley epidermal cells with *Bgh* haustoria, were not only present in the plant cell cytosol, but strikingly also entered the EHMx (Ahmed et al., 2015). This compartment is distinct in bright field images obtained by confocal microscopy (Figure 1, Supplemental Figure 1). mCherry is clearly present in the EHMx between the EHM, labelled by GFP-HvRabD2a previously shown to bind to this membrane (Mark Kwaaitaal et al., 2017), and the haustorium (Figure 1). The signal of free mCherry in a region of interest (1) (ROI (1)) in the EHMx, relative to the one in a ROI (2) in the plant cytosol, is 2.51 (mean of 106 cells; SE, 0.148). When expressing free mYFP in the plant cytosol, ROI1/ROI2 equals 2.86 (mean of 53 cells; SE, 0.218). While this indicated that intact proteins are able to cross the EHM, our previous study also demonstrated that certain proteins did not localize to the EHMx, as found for mYFP-CSEP0105 (Ahmed et al., 2015). Here we expanded the analysis, and examined mYFP fusions of 11 *Bgh* CSEPs without signal peptides, thus forcing their targeting to the cytosol. This allowed localization analysis of proteins with diverse sequence, pI and size in barley epidermal cells with *Bgh* haustoria 48 hours post inoculation (hpi). The CSEPs were fused to either the

N- or C-terminus of mYFP. However, independent of the orientation of the fusion, seven of the CSEP fusions were visible in the EHMx and four were not (Figure 2. and Table 1).

We considered if protein size could determine whether they localize to the matrix or not. However, we initially declined this idea as the largest fusion proteins (up to 74 kDa) generally localized to the EHMx, while some of the smallest (down to 41 kDa) generally did not (Table 1). Instead, we turned to examine the protein's calculated pI-values. Here we noted that the CSEP-fusions that entered the EHMx have a predicted $pI \geq 6.0$, and those that did not, have a predicted $pI \leq 5.8$, with the exception of CSEP0162 (pI 5.6), which did localize to the EHMx (Table 1). To test the causality behind this correlation, we generated a series of constructs encoding pI-modified mCherry, with varying numbers of aspartates (Asp), lysines (Lys) and alanines (Ala) added to its C-terminus (Table 2). After co-expressing those pI-modified mCherry versions with mYFP (pI 6.2) in barley leaf epidermal cells with *Bgh* haustoria, we quantified their accumulation in the EHMx. Here the mCherry signal at a ROI (1) in the matrix, normalized to the signal of mYFP in the same ROI, was recorded relative to the mCherry signal at a ROI (2) in the plant cytosol as well normalized to the mYFP signal (Figure 3). The results indeed suggested an effect of pI on protein localization in the matrix. The quantifications demonstrated that proteins with pI values between 6.0 and 8.4 were efficiently translocated from the plant cell cytosol to the matrix, while this feature was gradually lost when proteins possessed more extreme pIs.

To confirm these observations, we reversed the fluorescent proteins and generated constructs encoding pI-modified mYFP (Table 2), which we co-expressed with mCherry (pI 5.9) for normalization. The data obtained with these constructs corroborated the data from the pI-modified mCherry (Figure 3b). As mCherry and mYFP have low amino acid identity (29%), our data suggested that the modified pI due to the added aspartates and lysines at the C-terminus determined localization of these proteins in the EHMx.

As our initial non-quantitative examination suggested that high pI CSEP-mYFP fusion proteins readily localized to the EHMx, unlike the high pI mCherry and mYFP, we re-tested seven mYFP-CSEPs and quantified their distribution between the cytosol and EHMx. The data are included in Figure 3b and align well with the mCherry and mYFP-data (with the exceptions of CSEP0422, pI 5.8, and CSEP0244, pI 8.7), again validating that efficient EHMx targeting depends on the pI. In this case, except for mCherry or mYFP no amino acids were artificially

attached to the C- or N-terminus, suggesting that EHMx recruitment depends on the overall protein pI and not our modifications.

Protein size, sequence and pI affect translocation to the extrahaustorial matrix

We found that pHusion, a pH sensor with a size of 52 kDa was targeted poorly to the EHMx, even after we increased its pI to 6.2 or 6.5, which according to our findings with modified mCherry and mYFP proteins would facilitate take-up into the EHMx. In this case, the quantification was done with without normalization since pHusion is a GFP-RFP fusion protein (Gjetting, Ytting, Schulz, & Fuglsang, 2012), leaving little option for use of a fluorescent reference protein (Table 3). A CFP-YFP fusion protein having a similar size was tested to confirm this observation, and here mCherry could be used as reference protein. Also, in this case, a very low signal was detected in the EHMx even though the pI of 6 was predicted to permit EHMx targeting (Table 3). In summary, the matrix uptake of the double-fluorescent proteins (~52 kDa) was only 10-20% of that of single fluorescent proteins (~27 kDa), suggesting that also protein size influences translocation into the EHMx.

Despite the reduced EHMx localization of double-fluorescent proteins, we observed that larger CSEP-mYFP fusion proteins with a size above 70 kDa were taken up into the EHMx. Additionally, as indicated above, CSEP0422-mYFP had a low relative intensity in the matrix even though it has a low molecular weight (41 kDa) and a pI of 5.8 (Figure 3b). This shows that not only pI and size influence the translocation into the EHMx, but also the amino acid sequence and how this may affect protein structure of the translocated protein seems to be important.

Extreme pI hardly drives fungal effectors across the extrahaustorial membrane into the host cytosol

We know little on the directionality of the protein transfer across the EHM, as we have only made snap shots at single time-points and no studies of the transfer dynamics were possible. Yet, protein transfer may be bi-directional, and what we observe are equilibria of proteins moving in and out of the matrix. At the same time, it is unclear how effector proteins, secreted

by the fungal haustoria into the matrix, make their way to the host cytosol. If matrix chemical or physical conditions are important for the pI-influenced EHMx localization, it could be that effectors with extreme pIs are driven to the host cytosol as a result of a pI-dependent exclusion from the EHMx.

To investigate this possibility, we profiled the predicted pI-values of 491 CSEPs (Pedersen et al., 2012). The resulting multimodal pI-profile indeed suggested that most effector candidates (~60%) have pI-values outside the 6.0-8.4 interval (Figure S2a) that permits EHMx accumulation of host cell-expressed proteins. This would suggest that the extreme pI-effectors are driven out of the matrix. However, data suggests that this effector candidate multimodal pI-profile is unbiased and not selected for, as pI-values of theoretical proteins generated to reflect the amino acid occurrence broadly found in eukaryotes, show an identical multimodal profile (Weiller, Caraux, & Sylvester, 2004). If a pI-based transfer would be in play, selection of proteins with permissive pI-values would be expected to skew the profile away from the one of theoretical non-selected proteins, like it is the case for extremophiles (Weiller et al., 2004). Even if we restrict the analysis to the 96 CSEPs, that have a transcript expression level in the haustorium above the mean (Pedersen et al., 2012), we found no skewing of their pI profile either (Figure S2b). Therefore, we have no reason to suggest that effector pI may be important for their expected transfer to the host cytosol.

Nuclear export signal does not alter translocation to the extrahaustorial matrix

The modified mCherry and mYFP, that are efficiently localized to the EHMx, have stretches of lysines reminiscent of nuclear localization signals (NLS), which are rich in lysines and arginines (Freitas & Cunha, 2009). Consequently, a translocation mechanism through the EHM similar to nuclear import arose as a possibility. To test this idea, we fused a nuclear export signal (NES), identified in Haasen et al. (1999) and adapted to monocot plants (Scott, 2013), to the C-terminus of mCherry and examined its signal intensities in the nucleus, the EHMx and the cytosol relative to co-expressed mYFP. The signal in the nucleus, but not in the EHMx, was strongly reduced (Figure 4, Figure S3). This showed that the NES functioned as expected in protein export from the nucleus; however, the high fluorescent signal in the EHMx makes it unlikely that it NES-mediated nuclear export plays a role in protein translocation across the EHM. Yet it cannot be excluded that a mechanism using an NLS-like signal mediates protein transport across the EHM.

Neutral pH of the EHMx

The observation that extreme pI proteins have poor uptake in the EHMx, suggests that highly charged proteins cannot access this compartment. As the pH of the environment, together with the pI, determines the overall charge of the protein, it is very relevant to know the pH inside the EHMx. Therefore, we expressed two pH sensors in barley epidermal cells with haustoria, namely pHusion (see above) and pHluorin (Gao, Knight, Trewavas, Sattelmacher, & Plieth, 2004; Miesenböck, De Angelis, & Rothman, 1998). The 490/510 emission ratio of pHluorin and GFP/RFP emission ratio of pHusion did not differ between the nucleus and the matrix, suggesting that the two compartments have a similar pH (Figure S4). Meanwhile, the 490/510 emission ratio of pHluorin was significantly higher in the cytosol compared to the nucleus and the EHMx (mean of 15 measurements). This suggests that the pH in the matrix might be slightly higher than the cytosol. However, this was not observed when using the pHusion sensor. Even though the detection maximum of both pH sensors is determined to be pH 8 (Gao et al. 2004; Gjetting et al. 2012; Miesenböck et al. 1998), other factors in the EHMx environment might cause the differences between the two sensors. Technical limitations prevented us from obtaining a reliable pH calibration curve. Nevertheless, our data might suggest a peripherally higher pH in the matrix relative to the cytosol.

DISCUSSION

In this study, we clearly show that proteins produced in the plant cytosol can enter the EHMx, which is easily discernible at haustoria of the barley powdery mildew fungus. The haustorial complex is delimited by an extrahaustorial membrane, which shares features with the ER membrane (Kwaaitaal et al. 2017), and the present data indicate that this membrane has a translocation mechanism. We suggest that accumulation in the EHMx is affected by 1) protein pI. With the addition of amino acids, the predicted isoelectric point of mCherry and mYFP was altered and their localization within the plant cell was investigated. We show that medium pI proteins (pI 6.0-8.4) are efficiently translocated to the EHMx, whereas low and high pI proteins (pI < 6.0 and pI > 8.4) are gradually less abundant in the EHMx (Figure 3b). We also find that

accumulation depends on 2) protein size, as a single mYFP protein gave a strong matrix signal whereas double-fluorescent proteins did not. Finally, 3) protein sequence plays an important role, as CSEP0422-mYFP localized poorly, despite its small size of 40 kDa, while the larger CSEP0247-mYFP, CSEP0244-mYFP and CSEP0025-mYFP of approximately 70 kDa accumulated well within the EHMx.

We cannot completely exclude that the observed differences in fluorescence intensities may be caused by differential degradation of the tested proteins in the matrix, and indeed, addition of lysines may increase proteolysis resistance (Markert, Koditz, Ulbrich-Hofmann, & Arnold, 2003). However, the fact that the highest number of lysines (12) added to the fluorescent proteins reduced the matrix signal, and the fact that the mYFP single fluorescent protein gave a stronger matrix signal than the double-fluorescent proteins, suggest that a translocation mechanism is determining protein targeting to the EHMx.

Co-translational and post-translational translocation mechanisms exist for uptake of soluble proteins into plant cell organelles. These mechanisms are all based on translocon proteins. We will disregard ER co-translational translocation, as all proteins in this study lack signal peptides. Different mechanisms are responsible for post-translational translocation into the ER, mitochondria, chloroplasts and peroxisomes (Kunze & Berger, 2015). However, since we do not observe our marker proteins in any of these compartments, we have no reason to believe that one of these mechanisms is used directly for translocation across the EHM. Besides, we have previously found that the ER translocon protein, Sec61 β , is absent from the EHM (Zhang, Hanisch, Kwaaitaal, Pedersen, & Thordal-Christensen, 2013). This agrees with the fact that the unmodified, cytosolic mCherry and mYFP, which are easily taken up, lack exposed alpha-helix structures (Soleja, Manzoor, Khan, Ahmad, & Mohsin, 2018), which are required for targeting into these compartments (Kunze et al. 2015). We consider that the fact that size, pI and sequence influenced the EHMx uptake and suggest that a gated protein translocon is involved. We also consider that proteins may be taken up as globular proteins, as mYFP and mCherry quickly fold into their globular structure after translation, and we doubt that cells with haustoria have a chaperone system dedicated to unfold proteins for this uptake purpose. Therefore, we speculate that an EHM translocon may be flexible like the peroxisome import machinery is suggested to be (Emmanouilidis, Gopalswamy, Passon, Wilmanns, & Sattler, 2016; Kunze & Berger, 2015). Additionally, a pore complex similar to the nuclear pore complex is not likely, as the nuclear export signal showed no effect on the localization within

the matrix (Figure 4). Yet, we cannot exclude that a translocation mechanism similar to nuclear import, but without the nuclear export function, may exist. We have searched CSEP0247, CSEP0244 and CSEP0025 for consensus sequence tags potentially guiding these large and easily taken-up proteins to a translocon. However, our efforts using <https://wolfpsort.hgc.jp/>, <http://www.cbs.dtu.dk/services/DeepLoc/> and <http://localizer.csiro.au/> have been in vain. Therefore, we hope that future sequence break-down analyses of CSEP0247, CSEP0244 or CSEP0025 will help us to identify a consensus sequence determining the up-take and perhaps also translocon proteins.

It is not obvious why proteins with extreme pI were poorly localized to the EHMx. We attempted to estimate the pH in the EHMx and found that it may be marginally higher than the cytosol pH, which does not provide an explanation for the pI-effect. However, extreme pI proteins have a strong net charge that, irrespective of it being positive or negative, may hamper passage through a translocon pore.

Regardless of the translocation mechanism, our results suggest that proteins with an isoelectric point close to physiological pH accumulate more effectively after translocation to the EHMx than charged ones. The question arises whether there is a biological relevance of the matrix localization of plant expressed proteins. As described above, it may not be relevant for transfer of effectors to the plant cytosol. However, it may play a role in fungal nutrient uptake, where host proteins can reach the haustorium. Here they may be degraded on the outside of the haustorium PM and taken up as amino acids or small peptides, or they may be endocytosed as more or less intact proteins.

All-in-all, we consider the present description of protein translocation into the EHMx to be a valuable contribution to the otherwise sparse insight available on the haustorial complex.

EXPERIMENTAL PROCEDURES

Plant and fungal growth conditions

For bombardments, the barley line Ingrid *ror2*, hypersusceptible due to its mutation in the syntaxin gene, *ROR2* (Collins et al., 2003), was used. Plants were grown for 7 days with a

photoperiod of 16 h (150 μ Es-1m⁻²) and 20/15°C day/night. *Blumeria graminis* f. sp. *hordei* isolate B103 was propagated on barley P-10 plants in a cycle of one week.

Cloning of constructs

Ubiquitin-promoter driven DNA constructs encoding CSEP0105, CSEP0162 and CSEP0254 (without signal peptides) fused to the C-terminus of mYFP were generated previously (Ahmed et al., 2015). Similar constructs were generated here for CSEP0025, CSEP0062, CSEP0244, CSEP0247, CSEP0345, CSEP0420, CSEP0422 and CSEP0443 (Pedersen et al., 2012). However, in this case the fusions were on the N-terminus of mYFP. In short, the CSEP coding sequences were amplified, without their signal peptide coding sequences, via PCR from cDNA of plant material infected with *Bgh*, and cloned into pENTR/D-TOPO® vectors (Invitrogen). After validation, the inserts were transferred in frame to a destination vector via a Gateway LR reaction, thus fusing the encoded protein to the N-terminus of mYFP. For constructs encoding fluorescent proteins with different pls, empty Gateway destination vectors (pUbi-mCherry-Gateway and pUbi-mYFP-Gateway) (Kwaaitaal et al. 2010) were used as template for initial PCR amplification, using Q5 high-fidelity polymerase (New England Biolabs). In order to alter the pl of the fluorescent proteins (mCherry, mYFP and pHusion), codons for selected amino acids were added to the reverse primers (Table S1). The PCR products were cloned into the entry vector, pENTR/D-TOPO. The YC3.6 sequence was amplified from the pcDNA3 vector (Nagai, Yamada, Tominaga, Ichikawa, & Miyawaki, 2004) using Gateway compatible primers (Table S1) by PCR, and using a Gateway BP reaction recombined into pDONR201. Followed by an LR reaction, the coding sequences were transferred into the destination vector, pUbi-Gateway-nos, containing a ubiquitin promoter and a NOS terminator. In the process of cloning, heat shock transformation of TOP10 competent cells was used. All clones obtained were validated by sequencing.

Particle bombardment

For transient expression of the fluorescent proteins, the Biolistic PDS-1000/He Particle Delivery System from Bio-Rad was used. Several modifications to the manufacturer's manual were carried out: for each bombardment, seven detached barley leaves of 7-day-old plants were used. The particle coating was performed using 7 μ g DNA for each construct together

with 2.4 mg gold, 1 µg/µl protamine (Sivamani, Delong, & Qu, 2009) and 0.625 M CaCl₂ (Rasco-Gaunt, Riley, Barcelo, & Lazzeri, 1999). For the bombardments, a hepta adapter and rupture discs bursting at a helium pressure of 1.100 psi were used. After bombardment, the leaves were transferred onto 1% phytoagar petri dishes containing 40 mg/mL benzimidazole and immediately inoculated with powdery mildew spores. Microscopic evaluation was performed 48 h later.

Confocal microscopy

Confocal imaging was performed using a Leica SP5-X laser-scanning microscope. Bombarded barley leaves were mounted with Perfluorodecalin (Alfa Aesar A18288) and imaging was performed with a 63x water immersion lens and a numerical aperture of 1.20. The fluorescent protein mYFP was excited at 514 nm and the emission collected between 527 nm and 586 nm, whereas mCherry was excited at 582 nm and emission collected between 599 nm and 650 nm. In order to restrict bleed-through and to avoid overlapping excitation-emission signals, the sequential scan mode between lines was activated. Settings for pHusion, pHluorin and CFP/YFP were according to Gjetting et al. (2012), Gao et al. (2004) and Kwaaitaal et al. (2017), respectively.

Protein size and isoelectric point

The size and pI of proteins were calculated using the Create Sequence Statistics tool of the CLC Main Workbench 7.0.2 by QIAGEN.

Image processing and statistical analysis

Planar focused images were analysed with the Leica Application Suite (LAS) Advanced Fluorescence Lite Software (Leica Microsystems GmbH, Germany) with a clear focus on the EHM to make sure that the fluorescence in the EHMx was measured. Quantification of the signal from the protein of interest was performed with the Line Profile Tool that provides grey-scale values along a linear region of interest (ROI) or the stack profile tool measuring an area of interest. The Line Profile Statistics showed the mean value for each ROI separated for the two channels (*e.g.* mYFP channel 1 /mCherry channel 2). For each ROI, a protein of interest-value, normalized to an internal control (either mYFP or mCherry), was used for further analysis. Two ROIs were defined for each epidermal cell with a haustorium; one in the EHMx (ROI1) and one in the plant cytosol (ROI2). By dividing the ROI1 value by the ROI2 value, a

single value for each image was obtained, displaying the relative intensity in the EHMx. For each protein of interest, data from at least 10 EHMxs were included in the statistical analysis (unless stated otherwise). The data are provided in Supplemental Figure S4 and Supplemental Table S3. Indirect normalization of pHusion (GFP-RFP) proteins was obtained by dividing their relative EHMx intensity by the experienced averaged relative EHMx intensity of mCherry, which is 2.86 (see above). The statistics were performed in R (<https://www.r-project.org/>) by using the R stats package (version 3.5.1) followed by a multiple comparison (package *agricolae* version 1.2-8).

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FIGURES

FIGURE 1 Free mCherry is translocated from the plant cytosol to the extrahaustorial matrix of *Bgh*-infected barley cells. The extrahaustorial membrane is labelled with GFP-HvRabD2a. The localization was investigated 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells. hb, haustorial body; EHMx, extrahaustorial matrix; EHM, extrahaustorial membrane; BF, bright field. Images are recorded in a single plane. Scale bar, 10 μ m.

FIGURE 2 The protein sequence influences the uptake into the extrahaustorial matrix. The localization of CSEP-mYFP fusions and co-expressed mCherry was investigated 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells. (a) CSEP0420-mYFP (59 kDa) does not localize within the extrahaustorial matrix (EHMx) as only signal from mCherry is visible. (b) CSEP0247-mYFP (72 kDa) localizes in the EHMx together with mCherry. hb, haustorial body; hf, haustorial fingers; arrowhead, extrahaustorial matrix; BF, bright field. Images are recorded in a single plane. Scale bar, 10 μ m.

FIGURE 3 Localization of mCherry, mYFP (both 27 kDa) and CSEP-mYFP fusion proteins in the EHMx is influenced by their pI. The localizations were investigated 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells. (a) Examples of EHMx localization of mCherry, altered in its pI by addition of amino acids (see Table 2). BF, bright

field. Images are recorded in a single plane. Scale bar, 10 μm . (b) Relative EHMx fluorescence of mCherry (red) and mYFP (yellow) pl-altered (see Table 2) variants was tested and normalized to unaltered mYFP and mCherry, respectively. Each data-point is based on at least 10 cells, except for the protein represented in stripes (see Table S2). Roman numbers refer to (a). CSEP-mYFP and mYFP-CSEP fusions (see Table 1) were tested as well and normalized to unaltered mCherry. The statistics were performed in R using ANOVA followed by a multiple comparison showing the least significant difference (LSD). Different letters indicate statistically significant differences ($p < 0.01$). Error bars, standard error.

FIGURE 4 A nuclear export signal (NES) has no influence on the accumulation of mCherry in the extrahaustorial matrix. The localizations were investigated 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells. (a,b) Accumulation of mCherry-NES is significantly reduced, relative to mYFP, in the barley nucleus compared to the cytosol, whereas the signal in the extrahaustorial matrix (arrow) is not different. hb, haustorial body; hf, haustorial fingers; nc, nucleus; arrowhead, extrahaustorial matrix. Quantifications based on 10 images are shown. Statistics were performed in R using ANOVA followed by a multiple comparison showing the least significant difference (LSD) with $p > 0.05$. Error bars, standard error. Pictures are taken in a single plane. Scale bar, 10 μm .

Table 1. Visual scoring of the EHMx-localization of CSEP-mYFP and mYFP-CSEP fusion proteins suggests pl-dependency. Confocal microscopy-localizations of CSEPs (without signal peptide) fused to mYFP were investigated 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells.

CSEP	0420	0062	0162	0105	0422	0254	0443	0345	0247	0244	0025
EHMx localization	-	-	+	-	-	+	+	+	+	+	+
Fusion protein size (kDa)	59	44	45	41	41	39	38	61	72	74	73
pI of fusion proteins	4.7 (CSEP-mYFP)	4.9 (mYFP-CSEP)	5.7 (mYFP-CSEP)	5.7 (mYFP-CSEP)	5.8 (CSEP-mYFP)	6.0 (mYFP-CSEP)	6.0 (CSEP-mYFP)	6.5 (CSEP-mYFP)	8.0 (CSEP-mYFP)	8.7 (CSEP-mYFP)	8.8 (CSEP-mYFP)

Table 2. Addition of amino acids at the C-terminus alters the pI of mCherry and mYFP proteins.

	12 Asp	8 Asp 4 Lys	12 Ala	4 Asp 8 Lys	5 Lys 7 Ala	6 Ala 6 Lys	7 Lys 5 Ala	12 Lys	Unal- tered
mCherry pI	5.03	5.47	5.85	6.62	6.96	7.70	8.41	9.25	5.85
mYFP pI	5.16	5.73	6.16	6.84	-	-	-	9.17	6.16

Table 3. Several double-fluorescent proteins localize poorly to the extrahaustorial matrix. Confocal microscopy was used to investigate accumulation 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells. Yellow Cameleon and CFP-YFP were normalized using co-expressed mCherry. For pHusion, no internal control was possible due to technical limitations. Therefore, indirect normalization using an overall relative EHMx intensity of mCherry (2.86) was employed.

Construct	pI	Size (kDa)	Rel. intensity in EHMx (no. of cells)	Normalized directly (d) or indirectly (i)	Reference
Yellow Cameleon (YC3.6)	5.5	73	0.28 (7)	0.13 (d)	Nagai et al., 2004
pHusion	5.9	52	0.49 (7)	0.17 (i)	Gjetting et al., 2012
pHusion pI 6.2 4D8K	6.2	52	0.37 (6)	0.13 (i)	this work
pHusion pI 6.5 7K5A	6.5	52	0.33 (2)	0.11 (i)	this work
CFP-YFP	6	52	0.40 (7)	0.13 (d)	Bethke et al., 2009

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SUPPLEMENTALS

SUPPLEMENTAL FIGURE S1 The extrahaustorial matrix (EHMx) is a compartment surrounding the haustorial body (hb) delimited by the extrahaustorial membrane (EHM). Bright field (BF) image of a *Bgh*-haustorium 48 hours after inoculation of barley plants. hf, haustorial fingers; BF, bright field. Picture is taken in a single plane. Scale bar, 10 μ m.

SUPPLEMENTAL FIGURE S2 Number of candidate secreted effector proteins (CSEPs) with indicated isoelectric points (pI's). (a) All 491 *Bgh* CSEPs from Pedersen et al. (2012). (b) 96 *Bgh* CSEPs highly expressed in the haustorium (Pedersen et al., 2012).

SUPPLEMENTAL FIGURE S3 A nuclear export signal (NES) efficiently excludes mCherry from the nucleus. The localizations were investigated 48 h after transient transformation of barley leaf epidermal cells. Scale bar, 10 μ m.

SUPPLEMENTAL FIGURE S4 Little differentiation in pH between cytosol and extrahaustorial matrix. Fluorescence emission ratios of pH sensors in the nucleus, matrix and cytosol investigated 48 h after transient transformation and *Bgh* inoculation of barley leaf epidermal cells. (a) F490/F510 emission ratio of pHluorin (mean of data from 15 cells). (b) GFP/RFP emission ratio of pHusion (mean of data from 10 cells, except for the nucleus, where only four measurements were obtained). An ANOVA was performed in R followed by a multiple comparison showing the least significant difference (LSD) with $p > 0.05$. In both cases the stack profile tool was used. Error bars, standard error.

SUPPLEMENTAL TABLE S1. Primer sequences used for PCR reactions.

name	forward	reverse
mCherry pl 5.03/mYFP pl 5.16	CACCATGGTGAGCAAGGGCGAG	TCAATCGTCATCGTCATCGTCATCATCGTCGTCATCATCTTGTACAGCTCGTCCAT
mCherry pl 5.47/mYFP pl 5.73	CACCATGGTGAGCAAGGGCGAG	TCACCTTCTCTTTTATCATCGTCGTCATCATCGTCGTCCTTGTACAGCTCGTCCAT
mCherry pl 5.85/mYFP pl 6.16	CACCATGGTGAGCAAGGGCGAG	TCACGCCGCGGCGGCCGCCGCCGAGCCGCCGAGCAGCCTTGTACAGCTCGTCCAT
mCherry pl 6.62/mYFP 6.84	CACCATGGTGAGCAAGGGCGAG	TCACCTTCTCTTCTTCTCTTTTCTTGTGTCGTCATCGTCCTGTACAGCTCGTCCAT
mCherry pl 6.96	CACCATGGTGAGCAAGGGCGAG	TCACGCCGCGGCGGCCGCTTCTCTTCTTCTTCTTCTTTTCTGTACAGCTCGTCCAT
mCherry pl 7.7	CACCATGGTGAGCAAGGGCGAG	TCACGCCGCGGCGGCCGCCGCTTCTTCTTCTTCTTGTACAGCTCGTCCAT
mCherry pl 8.41	CACCATGGTGAGCAAGGGCGAG	TCACGCCGCGGCGGCCGCCGCCGCTTCTTCTTCTTCTTGTACAGCTCGTCCAT
mCherry pl 9.25/mYFP 9.17	CACCATGGTGAGCAAGGGCGAG	TCACCTTCTTTTTTCTTCTTCTTCTTCTTCTTCTTCTTGTACAGCTCGTCCAT
mCherry_NES	CACCATGGTGAGCAAGGGCGAG	TCACTCCACGGTCAGCTTCTCCAGCAGGCCGGCGGTCTTCTTGTACAGCTCGTCCAT
CSEP0420	CACCATGGCACCTCTGTCACTTTAAAGTC	CTCGGCTTCTTGTCACTCG
CSEP0062	CACCAAAAGCAACTACGTGTGCC	ATGTGTTTGGGTTTTCCTCG
CSEP0422	CACCATGTTTGTAAAGAATTACGTACCATACG	AAATTGGACGGTTCTCTTC
CSEP0443	CACCATGGCTAAATATTATACATGTGGCG	TTTACAATCCACAGATACGTT
CSEP0345	CACCATGTGGAAGCTGTTCCATC	GCCCAACCAACCTTTTGC
CSEP0247	CACCATGACCAATGTTCCATACTC	GGATTCTGTGACTTGACC
CSEP0244	CACCATGGCAAATATCCCATACTCG	AGAATCGGTGGCTGGTCG
CSEP0025	CACCATGGTAATATACCATACTCGG	GAGCTGTATGACTGAGCT
pHusion pl 6.2	CACCATGGCCTCCTCCGAGGAC	TCACCTTCTTCTTCTTCTTTTCTTGTGTCGTCATCGTCCTTGTACAGCTCGTCCAT
pHusion pl 6.5	CACCATGGCCTCCTCCGAGGAC	TCACGCCGCGGCGGCCGCCGCCGCTTCTTCTTCTTCTTGTACAGCTCGTCCAT
GWYC36	GGGGACAAGTTTGTACAAAAAGCAGGCTTAGCGGCCG CCACCATGGTG	GGGGACCACCTTTGTACAAGAAAGCTGGGTCAATCTGCAGAAATTTTACTC

SUPPLEMENTAL TABLE S2. Raw data of localization study in Figure 3.

	pI¹	No. of images	No. of exp.	Relative intensity	SD²	SE³
mCherry	5.03	11	4	0.311	0.126	0.038
mYFP	5.16	13	3	0.442	0.136	0.039
mCherry	5.47	18	3	0.483	0.209	0.049
mYFP	5.73	13	2	0.701	0.172	0.050
mCherry	5.85	12	2	0.957	0.304	0.088
mYFP	6.16	5	5	1.447	0.862	0.386
mCherry	6.62	13	4	1.455	0.435	0.121
mYFP	6.84	10	5	1.082	0.309	0.098
mCherry	6.96	20	3	1.229	0.303	0.068
mCherry	7.7	15	2	1.415	0.428	0.111
mCherry	8.41	12	2	1.017	0.341	0.098
mYFP	9.17	12	3	0.370	0.149	0.043
mCherry	9.25	5	2	0.382	0.104	0.046

¹Calculated isoelectric point.²Standard deviation.³Standard error.

SUPPLEMENTAL TABLE S3 Raw data of localization study in Supplemental Figure S4.

Name	No. of images	No. of exp.	Relative intensity	SD ¹	SE ²
mCherry-NES					
• Nucleus	11	2	0.175	0.153	0.048
• EHMx			0.581	0.547	0.173
• Cytosol			0.660	0.542	0.171
pHluorin					
• Nucleus	15	3	0.895	0.178	0.0460
• EHMx			0.802	0.162	0.0418
• Cytosol			1.151	0.144	0.0374
pHusion					
• Nucleus	4	2	2.929	0.5273	0.264
• EHMx	10	3	2.975	1.31	0.414
• Cytosol	10	3	2.980	1.123	0.355

¹Standard deviation.²Standard error.