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*Total number of authors:* 12

*Published in:* Molecular Plant Pathology

*Link to article, DOI:* 10.1111/mpp.12960

*Publication date:* 2020

*Document Version*  
Publisher's PDF, also known as Version of record

*Link back to DTU Orbit*

*Citation (APA):* Mentges, M., Glasenapp, A., Boenisch, M., Malz, S., Henrissat, B., Frandsen, R. J. N., Güldener, U., Münsterkötter, M., Bormann, J., Lebrun, M-H., Schäfer, W., & Martinez-Rocha, A. L. (2020). Infection cushions of *Fusarium graminearum* are fungal arsenals for wheat infection. *Molecular Plant Pathology, 21*(8), 1070-1087. https://doi.org/10.1111/mpp.12960
Infection cushions of *Fusarium graminearum* are fungal arsenals for wheat infection

Michael Mentges | Anika Glasenapp | Marike Boenisch | Sascha Malz | Bernard Henriussat | Rasmus J.N. Frandsen | Ulrich Güldener | Martin Münsterkötter | Jörg Bormann | Marc-Henri Lebrun | Wilhelm Schäfer | Ana Lilia Martinez-Rocha

**Abstract**

*Fusarium graminearum* is one of the most destructive plant pathogens worldwide, causing fusarium head blight (FHB) on cereals. *F. graminearum* colonizes wheat plant surfaces with specialized unbranched hyphae called runner hyphae (RH), which develop multicelled complex appressoria called infection cushions (IC). IC generate multiple penetration sites, allowing the fungus to enter the plant cuticle. Complex infection structures are typical for several economically important plant pathogens, yet with unknown molecular basis. In this study, RH and IC formed on the surface of wheat paleae were isolated by laser capture microdissection. RNA-Seq-based transcriptomic analyses were performed on RH and IC and compared to mycelium grown in complete medium (MY). Both RH and IC displayed a high number of infection up-regulated genes (982), encoding, among others, carbohydrate-active enzymes (CAZymes: 140), putative effectors (PE: 88), or secondary metabolism gene clusters (SMC: 12 of 67 clusters). RH specifically up-regulated one SMC corresponding to aurofusarin biosynthesis, a broad activity antibiotic. IC specifically up-regulated 248 genes encoding mostly putative virulence factors such as 7 SMC, including the mycotoxin deoxynivalenol and the newly identified fusaoctaxin A, 33 PE, and 42 CAZymes. Furthermore, we studied selected candidate virulence factors using cellular biology and reverse genetics. Hence, our results demonstrate that IC accumulate an arsenal of proven and putative virulence factors to facilitate the invasion of epidermal cells.

**Keywords**
effectors, *Fusarium graminearum*, infection cushion, runner hyphae, secondary metabolites, transcriptome, wheat infection
1 | INTRODUCTION

Fusarium head blight (FHB), caused by *Fusarium graminearum*, is a devastating disease of cereals including wheat, barley, oats, and rye with large economic impacts (Savary et al., 2012). After infection and colonization of wheat heads, *F. graminearum* reduces wheat yield by interfering with kernel development and by poisoning the remaining kernels with a cocktail of mycotoxins, such as deoxynivalenol (DON) and zearalenone, rendering them unsuitable for food and feed usage (Takemura et al., 2007). To date, there are no wheat cultivars available that are fully resistant to *F. graminearum* infection (Mesterhazy, 1995). Recently, comprehensive transcriptomic analyses of partially resistant and susceptible wheat cultivars inoculated with *F. graminearum* were performed to better understand the host molecular response to FHB (Biselli et al., 2018; Pan et al., 2018; Wang et al., 2018). Additionally, a transcriptional profiling approach separated symptomless and symptomatic aspects of the FHB infection and defined subsets of *F. graminearum* genes expressed in a single cereal host species or across two or more cereal hosts (Brown et al., 2017). Additional transcriptomics-based studies have been conducted, focusing on later stages of infection, using wheat coleoptiles, wheat spikes, and maize stalks (Lysøe et al., 2011; Zhang et al., 2012; Zhang et al., 2016; Kazan and Gardiner, 2018). To date, we are lacking information on the initial stages of fungal infection, from conidial germination to fungal growth on the plant surface and penetration into the plant epidermal cells. FHB starts when conidia of *F. graminearum* adhere to the surface of wheat spikes with the help of hydrophobin proteins (Quarantin et al., 2019). After germination on wheat floral tissues, *F. graminearum* grows epiphytically on the plant surface, using specialized unbranched hyphae, called runner hyphae (RH). These RH differentiate multicellular appressoria, called infection cushions (IC), able to produce several penetration events (Boenisch and Schäfer, 2011; Bormann et al., 2014). We previously showed that the deletion of *F. graminearum* adenylyl cyclase necessary for CAMP production, as well as overexpression of deoxypseudosine hydroxylase, the second activating enzyme of the eukaryotic translation initiation factor 5A, abolish the formation of IC and, therefore, the ability of the fungus to infect wheat (Bormann et al., 2014; Martinez-Rocha et al., 2016).

Appressoria of *Magnaporthe oryzae* and *Colletotrichum* species are melanized single cells emerging directly from conidial germ tubes (Perfect et al., 1999; Wilson and Talbot, 2009). They are produced due to the perception of a hydrophobic surface and are morphologically very different from *F. graminearum* complex IC that differentiate from specialized RH (Boenisch and Schäfer, 2011). Unicellular appressoria have been widely studied at both the histological and the molecular level, making it most probably the best examined fungal structure (O’Connell et al., 2012; Soanes et al., 2012). Several plant pathogens such as *Botrytis cinerea*, infecting approximately 200 plant species, *Sclerotinia sclerotiorum*, causing white mould mostly on vegetables, and *Rhizoctonia solani*, a wide host range pathogen, penetrate their host plants using complex appressoria similar to *F. graminearum* IC (Armentrout et al., 1987; Backhouse and Willetts, 1987; Garg et al., 2010). Although IC have been historically described several times in recent decades (Dodman et al., 1968; Nikraftar et al., 2013), a molecular description of their development is still pending.

In this study we removed the infecting fungal mycelium from the underlying wheat floral tissue and separated RH from IC using laser capture microdissection. We compared transcriptional changes occurring in these specialized fungal cells (RH and IC) to mycelium (MY) grown in complete medium (CM). This transcriptomic analysis allowed the identification of fungal transcripts specifically detected in the specialized fungal structures for epiphytic growth (RH) and IC. In more detail, we analysed genes encoding carbohydrate-active enzymes, putative secreted effector proteins, and secondary metabolite biosynthetic enzymes. In particular, transcripts detected in IC encode putative virulence factors prior to the invasion of epidermal cells.

2 | RESULTS

2.1 | Identification of *F. graminearum* epiphytic growth on wheat palea tissue

Under favourable conditions, conidia germinate and differentiate into specialized RH that epiphytically colonize the surface of wheat (Figure 1a). RH differentiate into IC that are complex appressoria made of agglomerated hyphae (Figure 1a,b). IC facilitate numerous penetration events and colonization of wheat epidermal cells (Figure 1c). Because the formation of IC is necessary for a successful infection, we performed transcriptomic analysis to identify changes in gene expression at the initial stage of wheat palea infection. RH and IC were detached from the palea surface, separated by laser-assisted microdissection, and used for RNA extraction (Figure 1d). To identify genes differentially expressed during infection, we also extracted RNA from *F. graminearum* hyphae (MY) growing in CM. cDNA libraries of MY, RH, and IC were constructed, Illumina-sequenced, and mapped against the genome of *F. graminearum*. Three replicates of the different libraries, which are highly coherent in a Pearson correlation test (Table S1), were used. The expression patterns of a set of five genes differentially expressed during infection according to RNA-Seq data were confirmed using quantitative reverse transcription PCR (RT-qPCR), proving the reliability of the experimental setup (Figure S1). The validated genes are relevant to our study (trichodiene synthase FgTRI5, polyketide synthase FgPKS12, and the putative effector 1 FgPE1) or are required for wheat virulence (GABA transaminases FgGTA1 and FgGTA2; Bönnighausen et al., 2015).

2.2 | Global gene expression profile of *F. graminearum* during initial infection of wheat floral tissue

Expression patterns of 13,826 predicted genes of *F. graminearum* were compared between MY, RH, and IC (Data S1 and Figure 2a). A total of 12,089 (87%), 11,778 (85%), and 12,504 (90%) transcripts were detected in MY, RH, and IC, respectively (Figure 2b); 870 (6%) transcripts...
were not detected in any cell type (Figure 2b). A comparison of the
differentially expressed genes between the three cell types was per -
fomed (Figure 2c). Genes with a log_2 fold change (log_2 FC) above the
threshold of +2 were classified as “up-regulated”, while genes with a
log_2 FC below −2 were “down-regulated” and genes with a log_2 FC
between −2 and +2 were “nonregulated”. The major differences were
found between IC and MY presenting 839 up-regulated genes and
2,709 down-regulated genes (Figure 2c).

Infection regulated genes were identified by comparing RH
and IC expression of genes to MY (Table 1). In total, we identified
3,916 infection regulated genes, further dissected into infection
up-regulated (982) and infection down-regulated genes (2,934).
Most of the infection up-regulated genes were up-regulated in both
RH and IC (485) or in IC (354, Figure 3a and Table 1). Fewer genes
were specifically up-regulated in RH (143). Similarly, most infec -
tion down-regulated genes were down-regulated in both RH and
IC (1,536) or in IC (1,173) and fewer genes were specifically down-
regulated in RH (225, Figure 3b and Table 1). Heat maps of infection
up-regulated or infection down-regulated gene expression show not
only large differences in expression pattern between RH/IC and MY,
but also between RH and IC (Figure 3c,d). To gain an unbiased view
of transcriptional changes we identified the top 50 up-regulated
Furthermore, genes were classified as encoding secreted or nonsecreted proteins, and sorted into the following gene families: secondary metabolite biosynthesis gene clusters (SMC; Sieber et al., 2014), transcription factors (TF; Son et al., 2011), transmembrane receptors (TMR), histone-modifying proteins (HM), protein kinases/phosphatases (PK), dehydrogenases (DH), carbohydrate-active enzymes (CAZymes; Lombard et al., 2014), genes involved in reactive oxygen species metabolism (ROS), and putative effector (PE) proteins (Data S1 and S2). Heat maps of clustered expression values in MY, RH and IC were generated according to these gene families. While heat maps for SMC, DH, TMR, and PE showed a diverse regulation pattern, genes grouped in nonsecreted proteins, TF, HM, and PK were under-represented in IC. Heat maps for secreted proteins, CAZymes, and ROS showed more over-represented genes in IC compared to MY or RH (Figure S2).

Among the 870 transcripts not detected (FPKM = 0), 604 encode proteins with unknown function, 74 are PE, 48 are genes involved in SMC, 36 in ROS, 21 in TP, and 19 in TF, among others (Data S1, sheet TND). Within the infection regulated genes, 634 encode secreted proteins, of which 306 were infection down-regulated, while 328 were infection up-regulated genes (Table S4). The highest transcriptional changes on infection regulated secreted proteins were on CAZymes, ROS, and PE (Table S4). Within the 3,378 infection-regulated nonsecreted proteins, 694 were infection up-regulated, while 2,684 were infection down-regulated. Functional categories with the highest transcriptional changes in infection-regulated nonsecreted proteins were the TF, TP, PK, and ROS (Table S4). In addition, 32 of the 67 known SMC were differentially expressed on infection, with 19 SMC being down-regulated and 13 SMC being up-regulated (Table 2). In conclusion, plant colonization triggers a wide range of changes in the \textit{F. graminearum} transcriptome.
2.3 | Transcriptional changes specific for RH and IC

A total of 573 genes were differentially expressed in IC compared to RH, of which 238 were down- and 335 up-regulated in IC (Figure 2c). Some 248 genes were infection and IC up-regulated compared to RH, while only 44 were infection and RH up-regulated genes (Table S5). The main gene families corresponding to infection up-regulated genes in IC were SMC (38 genes, corresponding to 16 clusters), CAZymes (42), ROS (34), and PE (33) (Table 3). This comparison shows that IC expressed a specific set of putative virulence factors (CAZymes, PE, and SMC), which we further investigated.

2.4 | IC are enriched in plant cell wall-degrading enzymes

CAZymes are proteins involved in cleavage, modification, or synthesis of glycosidic bonds (Lombard et al., 2014). The *F. graminearum* genome harbours 518 CAZyme encoding genes, of which 168 are involved in plant cell wall degradation (PCWDC), 95 in fungal cell wall modification (FCM), and 18 in starch biosynthesis/degradation (SDC). Genes involved in FCM corresponding essentially to chitin-binding proteins, α-mannan and β-glucans biosynthesis or modification, and enzymes for starch/glycogen processing are enriched in the set of infection down-regulated CAZymes (Table S6). Among the infection down-regulated CAZyme genes, only six supposedly target the plant cell wall. On the contrary, 102 PCWDC are predominantly infection up-regulated (Table S7). Among the 140 infection up-regulated degradative CAZymes, 35 PCWDC and four FCM were specifically up-regulated in IC, while only three genes (with yet unknown pathway annotation) were specifically up-regulated in RH (Table S8).

To test whether the production of PCWDC by IC has an impact on plant tissue, we assessed the status of the plant cell walls underneath IC by confocal laser microscopy. The intact plant epidermal cells showed a strong autofluorescence. Plant cells under and in the vicinity of IC completely lacked this autofluorescence (Figure 4a–c and Movie S1). Scanning electron microscopy showed that, in a later stage of infection in which fungal hyphae have already penetrated the plant underneath the IC, the cuticle is ruptured; however, an opening in the cell wall larger than the cushion itself was not observed (Figure 1c), as previously described (Bormann et al., 2014). Therefore, IC produce enzymes that mask or digest wheat cell wall compounds naturally emitting fluorescence after UV excitation like phenolic substances. Interestingly, 22 of the 61 redox CAZymes with proposed functions in degradation of phenolic compounds are infection up-regulated (Data S3). In particular, seven of them are IC up-regulated (Figure 4d).

2.5 | IC are enriched in infection up-regulated putative effectors

Secreted fungal effector proteins modulate host immune response to facilitate infection (Petre and Kamoun, 2014). Here, PE proteins were defined as secreted proteins, without transmembrane domains and a maximum size of 1,000 amino acids. Using this definition, 524 PE were identified (Data S4 and Figure 5a). Furthermore, 199 PE smaller than 200 amino acids and with a cysteine content higher than 2% were identified (Table S9). PE were classified as known effectors (PE: 44) containing previously identified domains and/or were defined as effectors in fungi or bacteria (Table S10), and as unknown effectors (PE: 480) with no predicted domains (Data S4 and Figure 5a). 77 PE were infection down-regulated (10 known and 67 unknown), while 88 were infection up-regulated genes (eight known and 80 unknown). Of the 88 infection up-regulated PE, four knowns and 29 unknowns were specifically up-regulated in IC compared to RH (Figure 5b,c). The 80 unknown infection up-regulated PE were further classified according to their taxonomic specificity into 20 PE conserved across kingdoms and 60 fungal-specific PE, including 14 *Fusarium*-specific and 10 *F. graminearum*-specific (Table S11).
Putative effector FgPE1 is up-regulated in IC, localized at the interface between fungal and plant cell walls, and dispensable for pathogenicity

FgPE1 (FGSG_04213), a putative effector up-regulated in IC compared to RH, was selected for further functional studies. FgPE1 gene expression was monitored by visualization of mCherry fluorescence driven by the FgPE1 promoter (FgPE1_prom:mCherry, Figure S3a). FgPE1 was expressed in both RH and IC on wheat palea, with the highest expression in IC at 4 days post-inoculation (dpi) (Figure S3b,c). A cross-section of an IC demonstrates high level of expression of FgPE1 (Figure S3d,e) during tissue invasion. Interestingly, also mycelia grown in wheat medium displayed strong FgPE1 expression at 1 and 3 dpi (Figure S3f,g). Mycelia grown in CM, in contrast, exhibited only faint mCherry signals at 1 dpi, slowly ramping up until 3 dpi (Figure S3h,i), indicating time- and nutrient-dependent regulation.

To determine the subcellular localization, FgPE1 was translationally fused to mCherry (FgPE1_prom:FgPE1:mCherry, Figure S4a) and transformed into a wild-type- (WT) strain expressing cytoplasmic green fluorescent protein (GFP) constitutively. A high level of FgPE1-mCherry was found in conidia produced on wheat medium (Figure S4b,c). In addition, FgPE1-mCherry localized around old hyphae but not in young growing hyphae, both on palea (6 dpi; Figure S4f–h) and in wheat medium (1 and 3 dpi; Figure S4i–l). On CM,
however, no FgPE1 signal was observed at 1 and 3 days of growth (Figure S4m–p), suggesting that FgPE1 expression is induced in the presence of plant-derived compounds. To determine whether FgPE1 was localized at the fungal cell wall or plasma membrane, protoplasts were generated and incubated on wheat medium for 20 hr. Confocal laser microscopy of undigested mycelia and partially digested hyphae showed that the mCherry signal was located around hyphae (Figure S5a–f) while protoplasts lacked a mCherry signal, demonstrating that FgPE1 localizes to the fungal cell wall and not to the plasma membrane (Figure S5).

When inoculated on paleae, mCherry signals were first observed around IC (Figure 6a,b). Six days after inoculation, vacuolated subcuticular hyphae were produced beneath IC that displayed a strong mCherry signal (Figure 6c,d and Movie S2). In addition, a mCherry signal was present in the fungal–plant cell wall interface in the vicinity of fungal infection hyphae (Figure 6c,d and Movie S2). The WT strain expressing cytosolic GFP was used as control and lacked a fluorescence signal in the detection range of mCherry, proving that the FgPE-mCherry signal is specific and not due to plant-derived autofluorescence (Figure 6e–h).

FgPE1 was deleted by gene replacement (Figure S6a,b). In virulence assays on wheat the deletion mutants as well as an ectopic mutant and WT exhibited full infection, indicating that FgPE1 is dispensable for virulence (Figure S6c,d).

2.7 | Wheat infection triggers secondary metabolite production

In this study, 53 of 67 SMC (Sieber et al., 2014) were expressed in any of the tested conditions (Table 2). The remaining 14 SMC, including those involved in biosynthesis of zearalenone (C15), butenolide (C49), fusarin C (C42), and fusarielin (C60), were neither expressed in CM nor during infection (Table 2). Twenty-three SMC were infection down-regulated, among them the carotenoid (C28) and siderophore ferricrocin clusters (C33, Table 2). Interestingly, there was one unknown SMC (C09) specifically down-regulated in IC, while similarly expressed in RH and MY. Two undescribed SMC (C37 and C40) were specifically down-regulated in RH and similarly expressed in IC and MY (Table 2). Twelve SMC were infection up-regulated. Among the six SMC up-regulated in RH and IC were two known siderophore clusters, triacylfusarin (C21) and malonichrome (C63). Five SMC were specifically up-regulated in IC, including the known virulence factors trichothecene (TRI, C23, Figure 7a) and fusaoctaxin A (C64, Table 2). To determine the impact of DON on the initial plant colonization, we quantified and compared spikelet infection of the DON-deficient trichodiene synthase deletion mutant (∆tri5) and the WT strain, both expressing GFP constitutively (Jansen et al., 2005). Fluorescence microscopy of longitudinal cuts through inoculated

| TABLE 2 | Comparative expression analysis of 67 secondary metabolite gene clusters (SMC) |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| **Infection up-regulated** | **RH and IC up-regulated** | 6 | C03, C21 (triacyt fusarine), C25, C36, C57, C63 (malonichrome) |
| | **Specifically RH up-regulated** | 1 | C13 (aurofusarin) |
| | **Specifically IC up-regulated** | 5 | C16, C22, C23 (trichothecene), C64 (fusaoxacin A), C66 |
| **Similar expression** | 18 | C08, C12, C17, C18 (arcinol), C19, C26, C30, C32, C34, C35, C43, C44, C45, C50, C56, C59 (culmorin), C65, C67 |
| **Infection down-regulated** | **RH and IC down-regulated** | 20 | C04, C05, C06, C10, C11, C20, C27, C28 (carotenoid), C29, C31, C33 (ferricrocin), C38, C39, C41, C47, C54, C55, C58, C61, C62 |
| | **Specifically RH down-regulated** | 2 | C37, C40 |
| | **Specifically IC down-regulated** | 1 | C09 |
| **Not expressed** | 14 | C01, C02, C07, C14, C15 (zearalenone), C24, C42 (fusarin C), C46, C48, C49 (butenolide), C51, C52, C53 (precursor of insoluble perithecial pigment), C60 (fuscarielin) |

*Gene clusters expression analysis was determined as mentioned in the methods section, estimation of secondary metabolite gene cluster (SMC) regulation.

*Name of known secondary metabolites is given in parentheses. IC, infection cushions; RH, runner hyphae; MY, mycelium grown in complete medium. Secondary metabolite cluster nomenclature according to Sieber et al., 2014.
wheat spikes revealed less mycelial growth inside spikelets inoculated with the Δtri5 mutant compared to the ones inoculated with the WT (Figure 7b). For molecular quantification, fungal DNA from inoculated spikelets was extracted and the relative amount measured by quantitative PCR (qPCR) as previously described (Voigt et al., 2007). Results revealed that Δtri5 grew 60% and 75% less than the WT at 3 and 5 dpi, respectively (Figure 7c). Hence, DON facilitates rapid colonization of plant tissues at the initial stage of infection.

2.8 | Aurofusarin is a RH-specific antibiotic active against a wide range of microorganisms

The single SMC specifically up-regulated in RH is involved in aurofusarin biosynthesis (C13) and is strongly down-regulated in IC (Table 2 and Figure 8a). To test whether aurofusarin could act as an antibiotic against microbial competitors growing on the wheat floral tissue’s surface, the toxicity of aurofusarin on bacteria and fungi was assessed using mycelium extracts from either WT or the aurofusarin-deficient Δpks12 mutant (Figure 8b and Table S12). The presence or absence of aurofusarin in extracts was determined by LC-MS (Figure S7). WT extract was highly toxic to the gram-positive bacteria Bacillus subtilis and Micrococcus luteus (98%–100% growth inhibition [GI]), while other bacteria such as Escherichia coli, Pseudomonas aeruginosa, Pseudomonas fluorescens, Janthinobacteria HH102, and Rhizobium sp. NG234 were insensitive (Figure 8b). WT extract was also highly toxic to Pyrenophora teres (100% GI), Candida albicans (100% GI), and Pichia pastoris (85% GI), while moderately toxic to Candida parapsilosis (50% GI) and Saccharomyces cerevisiae (50% GI). F. graminearum and, closely related, Nectria haematococca were totally insensitive to WT extracts (Figure 8b). Extracts from Δpks12 were completely nontoxic to all tested organisms, demonstrating that aurofusarin is indeed responsible for the observed toxicity of WT extract to bacteria and fungi.

3 | DISCUSSION

Although FHB is a devastating cereals disease that occurs worldwide, the molecular basis of the initial steps of infection are basically unknown. On the contrary, the large-scale reprogramming of appressorial gene expression of the plant pathogens M. oryzae and Colletotrichum species is well known (O’Connell et al., 2012; Soanes et al., 2012). Here, the transcriptomes of hyphae grown in culture (MY) and epiphytically grown RH revealed fundamental transcriptional differences. Interestingly, the majority of these genes (73% out of 2,390 genes) are down-regulated in RH compared to MY. These transcriptional changes highlight the huge differences between fungal hyphae growing in a nutritious broth and under very restrictive conditions on palea. Similar results were reported from the plant pathogen Colletotrichum higginsianum where transcriptomes of appressoria built in culture and on leaves were significantly different, though morphologically indistinguishable (O’Connell et al., 2012), stressing the necessity to elucidate the different aspects of fungal development in its natural environment.
The majority of up-regulated genes in RH or IC compared to MY encode hypothetical proteins with unknown function, indicating a requirement for characterization of such proteins (Tables S2 and S3). The next more up-regulated genes encode for CAZyme degradative enzymes, suggesting preparation to break the plant cell wall (Tables S2 and S3). Thirty-five plant cell wall-degrading enzymes (PCWDC) are specifically induced in IC encoding enzymes with predicted cellulolytic, xylanolytic, pectinolytic, and oxidoreductase activity. The activity of these PCWDC may cause the observed loss of blue fluorescence around the developed IC, preceding penetration. This fluorescence of the plant epidermal cell wall is attributed especially to phenolic substances such as chlorogenic acid, caffeic acid,
coumarins, stilbenes, and ferulic acid of the stress-induced phenylpropanoid pathway, which are also known to be involved in plant defence (Dixon and Paiva, 1995; Lang et al., 1991). Cell wall-bound ferulic acid is the major substance causing blue light emission in grasses like wheat (Lichtenthaler and Schweiger, 1998). Interestingly, ferulic acid inhibits in-culture DON biosynthesis at the transcriptional level (Boutigny et al., 2009). DON, in turn, does not contribute to the loss of fluorescence, as a DON-deficient mutant causes the same phenotypes as the wild type (Boenisch and Schäfer, 2011).

Among the five secondary metabolite clusters specifically induced in IC are deoxynivalenol (C23, DON) and fusaoctaxin A (C64), both having important roles during infection (Table 2). Fusaoctaxin A facilitates the cell-to-cell movement of the fungus and is important for virulence during coleoptile as well as wheat spike infection (Jia et al., 2019; C64 is FG3_54 in Zhang et al., 2012). DON inhibits the eukaryotic translational machinery and is essential for colonization of the spike, with massive induction during colonization of the developing caryopses and the rachis node (Ilgen et al., 2009). Importantly, both SMC are up-regulated in IC clearly in preparation for the following colonization steps. DON-deficient mutants fail to cross the rachis node (Proctor et al., 1995; Maier et al., 2006; Ilgen et al., 2009), which is accompanied by plant cell wall thickening and jasmonate-related...
defence reactions preventing further fungal colonization (Jansen et al., 2005; Bönnighausen et al., 2019). Our results verify previous fluorescence-microscopy assisted analyses of a specific IC induction of tri5 (FGSG_03537). Yet, this specific induction and subsequent DON biosynthesis are neither necessary for IC development nor plant cell wall penetration (Boenisch and Schäfer, 2011). Quantification of fungal DNA in infected spikelets now revealed a substantially slower spikelet infection of the DON deficient Δtri5 mutant compared to WT-GFP strain. DON, therefore, acts as a virulence factor immediately after penetration. DON was shown to induce programmed cell death (PCD) after infiltration into plant tissues (Desmond et al., 2008; Diamond et al., 2013; Blümke et al., 2015). Therefore, DON-induced PCD may facilitate the release of nutrients during initial infection. Loss of DON, in turn, may result in less nutrients available, causing the observed growth reduction at 3 and 5 dpi. DON deficiency may, therefore, enable successful plant defence reactions that are, in the case of the WT, not initiated or suppressed.

The cluster for aurofusarin biosynthesis is the only one specifically induced in RH. Aurofusarin is a red pigment produced by different Fusarium species, belonging to polyphenol, more accurately bis-naphthopyrone pigments (Frandsen et al., 2006; Xu et al., 2019). In a previous study, deletion of the F. graminearum pks12 gene (FGSG_02324)
led to a loss of red pigment, a higher growth rate, and 10-fold more conidia production than WT but had no impact on pathogenicity on wheat and barley (Malz et al., 2005). Recently, aurofusarin has been described to inhibit *Lactobacillus* and *Bifidobacterium*, but not *E. coli* (Sondergaard et al., 2016). Excitingly, it has been described as an antifeedant that accumulates in high amounts to protect *Fusarium* fungi from a wide range of insects (Xu et al., 2019). The microbiology of the phyllosphere is, in general, not very well understood, but it seems safe to assume that RH of *F. graminearum* ward off other microbes during colonization of the palea’s surface. Among the bacteria found in the microbiome of wheat spikes are *Pseudomonas*, *Bacillus*, *Janthinobacterium*, and *Actinomycetes* (Chen et al., 2018). In this study, we showed that aurofusarin is an inhibitor of different bacterial and fungal species, among them yeast, including most notably the widespread human pathogen *C. albicans*. Another polyphenol pigment found in *F. graminearum* is bostrycothin purpurfusarin, which is also known to have antibiotic properties against *C. albicans* (Frandsen et al., 2016). Further research will show if these secondary metabolites could improve the fight against this widespread human pathogen.

Up-regulated in RH and IC are two infection up-regulated iron-chelating siderophores, triacetyl fusarin and malonichrome, which are necessary for virulence (Oide et al., 2015). A third infection up-regulated iron-chelating siderophore, ferricrocin, is important for sexual development but not for virulence (Oide et al., 2015), explaining why we found this metabolite down-regulated during infection. Fourteen SMCs were not at all expressed, including zearalenone (C15), fusarin C (C42), butenolide (C49), and fusarielin (C60) clusters, all dispensable for wheat infection (Gaffoor et al., 2005; Harris et al., 2007; Sørensen et al.,
2012). *F. graminearum* is a pathogen with a variety of hosts such as wheat, barley, oats, rye, maize, and soybean (Savary *et al.*, 2012; Sella *et al.*, 2014), and many transcripts not detected might be necessary for specific colonization of such hosts (Harris *et al.*, 2016). For instance, the SMC DON necessary for wheat infection, and highly transcribed under our study conditions, does not seem to act as a virulence factor on barley (Maier *et al.*, 2006). Therefore, the lack of transcript detection could be due to specificity or redundancy on their protein function.
Besides CAZymes and SMC, we identified a large number of putative effector proteins (PE) up-regulated in IC. Previously, Lu and Edwards (2016) identified 190 small (≤200 amino acids) and cysteine-rich (≥2%) secreted proteins as candidate effectors in *F. graminearum*. Here, we identified 199 proteins with such characteristics (Table S9). However, several studies reported effectors as secreted proteins with very diverse sizes or cysteine content (Kulkarni et al., 2003; Rooney et al., 2005; Djamei et al., 2011; Frias et al., 2011; Sperschneider et al., 2013; Blümke et al., 2014; Tanaka et al., 2014; Jashni et al., 2015; Quarantin et al., 2016). From the 524 identified PEs, 88 were infection up-regulated and of these, 33 were specifically up-regulated in IC. Four PEs specifically up-regulated in IC contained LysM or CFEM domains, being indicative of key proteins necessary during penetration of the host cell by suppressing fungal recognition and manipulating host functions (Mentlak et al., 2012; Zhang et al., 2012; Takahara et al., 2016). The hemibiotroph *C. higginsianum* transcribes effectors in consecutive waves associated with the transitions in the pathogen’s lifestyle (Kleemann et al., 2012). Hence, it is likely that *F. graminearum* expresses a different set of effectors during later stages of plant colonization. The study of the 80 unknown plant-induced fungal effector-like proteins (Table S11) could lead to the discovery of new targets for fungal control and even specifically *F. graminearum* control. In a different approach using a comparative genome analysis, a set of 2,830 *F. graminearum* genes, presumably associated with pathogenicity, were identified (Sperschneider et al., 2013). We found that roughly 3.9% of these genes (111) are transcriptionally up-regulated during palea colonization, with 35 of them specifically up-regulated in IC and 30 encoding PEs. This set of PEs with high specificity for cereal infection could be of outstanding importance for the initial host–pathogen interaction.

The highly expressed in IC fungal PE, called FgPE1, was characterized. FgPE1 encodes a secreted 151 amino acid protein with an Alt-A1 allergen analog domain present in the Alt-A1 effector from *Alternaria alternata* (Chruszcz et al., 2012). Recently, an Aa1-like protein, PevD1 from *Verticillium dahliae*, has been found to interact and inhibit the antifungal activity of GhPR5 cotton plant protein as a strategy to fight the plant defence and promote fungal infection (Zhang et al., 2018). Here, expression results indicated a transcriptional regulation of FgPE1 depending on plant factors. A previous report showed that FgPE1 is highly expressed at different time points (4, 12, 24, 48, and 72 hpi, and 8 and 14 dpi) during infection of wheat spikes and at 8 days in old minimal medium culture (Lu and Edwards, 2016), supporting the hypothesis that FgPE1 expression is regulated by plant factors and nutrient availability and that it is present not only during early infection but all along the infection process. Replacement of FgPE1 in *F. graminearum* did not affect virulence according to our infection assays (Figure S6). We assume a high degree of functional redundancy within the PE gene family. This is in accordance with studies, for example in *U. maydis*, that, for the most part, failed to identify novel, virulence-specific genes, for example within a pool of potential effector genes (Kämper et al., 2006). FgPE1 is localized at the fungal cell wall. During IC formation, FgPE1 is secreted and at first localized at plant cell walls in the close vicinity around the IC. Once the fungus grows beneath the plant surface, FgPE1 is localized at the fungal–host interface. Effectors with such localization such as ChEC34 and ChEC89 of *C. higginsianum* (Kleemann et al., 2012) or the CFEM1 protein (FGSG_02077) from *F. graminearum* (Zhang et al., 2012) usually are necessary for eliciting or suppressing the plant recognition depending on their lifestyle.

Taken together, we found major transcriptional changes between epiplhical and in-culture hyphae. A complex set of virulence-associated factors, comprising plant cell wall-degrading enzymes, secondary metabolites, and effector proteins among others, are synthesized in IC, preparing the fungal hyphae for successful penetration and subsequent colonization of plant tissue. Therefore, IC are arsenals of fungal combat and the genes expressed in them could provide potentially novel targets for *Fusarium* control.

### 4 | EXPERIMENTAL PROCEDURES

Detailed experimental procedures are described in Methods S1.

#### 4.1 | Fungal growth and conidia production

*F. graminearum* wild type (WT; Fg-8/1; Miedaner et al., 2000) and mutants used and produced in this study were grown, cultured, and transformed as described before (Jansen et al., 2005).

#### 4.2 | Preparation of wheat-infected tissue for laser capture microdissection

Detached wheat palea infection assay was prepared according to Boenisch and Schäfer (2011). Mycelium samples were prepared by inoculating 750 conidia of the WT-GFP strain in 50 ml CM and incubated for 3 days. A mycelium piece of about 1 mm in diameter was used for RNA extraction, amplification, and construction of cDNA libraries.

#### 4.3 | Laser capture microdissection

Paleae containing RH and IC were prepared by cutting off their upper and lower ends and immediately transferred to absolute ethanol on ice according to previous studies (Goldsworthy et al., 1999; Clément-Ziza et al., 2008). RH and IC were prepared and dissected as mentioned in Methods S1.

#### 4.4 | RNA extraction, amplification, and cDNA library construction

RNA extraction, amplification, and cDNA library construction of IC, RH, or MY were performed according to Lê et al. (2005). See Methods S1.
4.5 | Purification of cDNA libraries

For the removal of primers, enzymes, and other substances of the process from the cDNA libraries the NucleoSpin Gel and PCR Clean-up Kit (Macherey-Nagel) was used according to the manufacturer’s instructions.

4.6 | Finalization of cDNA libraries (end-it reaction)

To provide 5′-phosphorylated, blunt-ended cDNAs, the End-It DNA End-Repair Kit (Biozym Biotech Trading Gmbh) was used according to a modified protocol. One microgram of the final cDNA libraries of three independent replicates of mycelia, RH, and compound appressoria, respectively, were sent for RNA-Seq analysis. See Methods S1.

4.7 | RNA-Seq mapping and quantification

RNA-Seq reads were mapped on the reference genome using tophat2 v. 2.0.8. The interval for allowed intron lengths was set to minimum 20 nt and maximum 1 kb (Trapnell et al., 2009). Three highly correlating replicates were used according to the Pearson correlation test (Table S1). We used cufflinks to determine the abundance of transcripts in FPKM (fragments per kilobase of exon per million fragments mapped) and calculated differentially expressed genes using cuffdiff (Trapnell et al., 2009; Trapnell et al., 2012). The gene models were included as raw junctions. The uncorrected p value and the FDR-adjusted q value of the test statistic (q value) were calculated, p and q values per each gene are given in the general part of Data S1. Any given gene of interest can be evaluated by its fold change of transcription and by the resulting p and q values. Genes with a minimum of four-fold increase or decrease in expression (|log2 of the FPKM values + 1| ≥ 2) between two experimental conditions were considered as regulated.

4.8 | Annotations and databases used

The transcriptome data discussed in this publication have been deposited in NCBI’s Sequence Read Archive (SRA; https://www.ncbi.nlm.nih.gov/sra; SUB3191581; Edgar et al., 2002). The reference genome of F. graminearum PH1 database FGDB (ftp://ftp.mips.gsf.de/fungi/Fusarium/F_graminearum_PH1_v32/) was used to map the cDNA libraries constructed (Wong et al., 2010). Genes were manually grouped in gene families using the tool at https://grn.nim.nih.gov/prime/ with gene family/gene families. See Methods S1.

4.9 | Validation of RNA-Seq data by qPCR

Validated genes were FgTRI5 (FGSG_03537), FgPKS12 (polyketide synthase 12; FGSG_02324), FgPE1 (FGSG_04213; putative effector1), and two GABA-aminotransferases (FgGTA1, FGSG_05554; FgGTA2, FGSG_06751). For relative expression analysis the tool REST (Relative Expression Software Tool) was used (Pfaffl et al., 2002). For evaluation of the housekeeping genes cofolin (FGSG_06245) and ubiquitin (FGSG_10805) we used the comprehensive tool “Ref Finder” (Tables S13 and S14; Vandesompele et al., 2002; Andersen et al., 2004; Pfaffl et al., 2004; Silver et al., 2006). For details see Methods S1.

4.10 | Generation of knock-out, expression, and localization constructs for FgPE1 mutants

All plasmids were constructed using the yeast recombination method (Colot et al., 2006) and the pRS426 background plasmid (Christianson et al., 1992). Amplification of the ORF, and 5′ and 3′ flanks of the genes of interest was performed using primers shown in Table S15 and genomic DNA extracted from the WT strain. The final constructs were excised with the respective restriction enzymes (Table S16) and used to transform F. graminearum WT or WFGP strains. At least two independent mutants were generated and examined. For details see Methods S1.

4.11 | Virulence assay: wheat spikes point inoculation and wheat palea infection

Virulence assays where prepared according to Boenisch and Schäfer (2011) and Frandsen et al. (2006). For details see Methods S1.

4.12 | Quantification of fungal material within inoculated wheat spikes using qPCR

Genomic DNA of inoculated wheat spikes was isolated using the CTAB method and according to Voigt et al. (2007). For details see Methods S1.

4.13 | Fluorescence microscopy

Histological studies of WT-GFP, ∆tri5-GFP, and mutants generated in this study were performed as previously described in Boenisch and Schäfer (2011). For details see Methods S1.

4.14 | Scanning electron microscopy

Scanning electron microscopy (SEM) was done with SEM LEO 1525 at 6 kV using detached paleae of wheat cultivar Nandu inoculated with 5 µl of 2 × 10⁶ conidial suspension of WT-GFP strain and prepared as described (Boenisch and Schäfer, 2011). To identify penetration pores, infection structures were removed from the plant surface of critical point dried paleae using adhesive tape and processed for SEM as previously described (Bormann et al., 2014).
4.15 | Extraction of aurofusarin from *F. graminearum* WT and aurofusarin-deficient mutant Δpks12

Fungal material of the WT strain and the aurofusarin-deficient mutant was harvested after 4 days from 50 ml CM liquid cultures. The respective mycelium was harvested using Miracloth, washed with 100 ml double-distilled water (ddH2O) and semi-dried using a filter paper. Around 1 g of mycelium was transferred into a 2 ml tube and supplemented with 1 ml potassium phosphate-buffer (50 mM, pH 7). After addition of two metal pearls (3 mm diameter), the solution was ground for 15 min using a Retsch mill. After centrifugation at 13,000 rpm for 15 min, the extracted supernatant was filter sterilized using a 0.22 µm Millex GP filter.

4.16 | Analysis of fungal extracts via LC-MS

The extracts of WT strain and Δpks12 mutant were analysed as described in Methods S1.

4.17 | Bioactivity assay

Liquid cultures of the organism listed in Table S12 were used for bioactivity assays. Assays were performed as described in Methods S1.

ACKNOWLEDGMENTS

The project was partially funded by the University of Hamburg, the Deutsche Forschungsgemeinschaft BO4098/2-1 (J.B.), the Austrian Science Fund FWF, and the Special Research Project “Fusarium” F3705/DFGEM1682/6-1 (U.G.). We gratefully acknowledge Birgit Hadeler, Cathrin Kröger, Macarena Iniesta Pallarés, and Dr Joanna Szewinska for excellent technical assistance, Brigitte Doormann for critical reading of the manuscript, and E. Woelken, K. Dehn, R. Walter, and Dr F. Friedrich for technical support with electron microscopy. The authors have declared that no conflict of interests exists.

DATA AVAILABILITY STATEMENT

The data produced in this publication have been deposited in NCBI’s Sequence Read Archive (SRA), https://www.ncbi.nlm.nih.gov/sra; accession SUB3191581.

ORCID

Ana Lilia Martinez-Rocha https://orcid.org/0000-0002-6560-3473

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SUPPORTING INFORMATION

Additional Supporting Information may be found online in the Supporting Information section.

How to cite this article: Mentges M, Glansenapp A, Boenisch M, et al. Infection cushions of Fusarium graminearum are fungal arsenals for wheat infection. Molecular Plant Pathology. 2020;21:1070–1087. https://doi.org/10.1111/mpp.12960