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ORIGINAL ARTICLE

Dirhamnolipids secreted from *Pseudomonas aeruginosa* modify antifungal susceptibility of *Aspergillus fumigatus* by inhibiting β 1,3 glucan synthase activity

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***Pseudomonas aeruginosa* and *Aspergillus fumigatus* are the two microorganisms responsible for most of the chronic infections in cystic fibrosis patients. *P. aeruginosa* is known to produce quorum-sensing controlled rhamnolipids during chronic infections. Here we show that the dirhamnolipids secreted from *P. aeruginosa* (i) induce *A. fumigatus* to produce an extracellular matrix, rich in galactosaminogalactan, 1,8-dihydroxynaphthalene (DHN)- and pyo-melanin, surrounding their hyphae, which facilitates *P. aeruginosa* binding and (ii) inhibit *A. fumigatus* growth by blocking β 1,3 glucan synthase (GS) activity, thus altering the cell wall architecture. *A. fumigatus* in the presence of diRHls resulted in a growth phenotype similar to that upon its treatment with antifungal echinocandins, showing multibranched hyphae and thicker cell wall rich in chitin. The diRHl structure containing two rhamnose moieties attached to fatty acyl chain is essential for the interaction with β 1,3 GS; however, the site of action of diRHls on GS is different from that of echinocandins, and showed synergistic antifungal effect with azoles.**

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Introduction

Aspergillus fumigatus and *Pseudomonas aeruginosa* are the two microorganisms responsible for most of the chronic infections encountered in cystic fibrosis adult patients. *A. fumigatus* is indeed the major filamentous fungus reported to colonize the airways of these patients, with a prevalence of 12–57% (Paugam *et al.*, 2010), and major risks in developing allergic bronchopulmonary aspergillosis in 15% of patients (Knutsen *et al.*, 2012; Baxter *et al.*, 2013). The most common bacterium isolated from the sputum of patients with cystic fibrosis is *P. aeruginosa*, which chronically colonizes cystic fibrosis airways in up to 75% of adult patients (Hill *et al.*,

2005; Moree *et al.*, 2012; Baxter *et al.*, 2013). The co-colonization by *A. fumigatus* and *P. aeruginosa* is very deleterious for the patients, characterized by a high decline in cystic fibrosis pulmonary functions and the worst prognosis with no access to lung transplantations (Ferreira *et al.*, 2015).

Several recent studies have shown that *P. aeruginosa* and *A. fumigatus* interact during lung colonization. The first evidence for crosstalk's *in vivo* between *A. fumigatus* and *P. aeruginosa* emerged from the fact that antibacterial therapy targeting *P. aeruginosa* colonization is followed by a significant reduction in *A. fumigatus* colonization (Baxter *et al.*, 2013). Two studies have shown a direct interaction between *P. aeruginosa* and *A. fumigatus* (Blyth, 1971; Mowat *et al.*, 2010). *P. aeruginosa* secretes molecules such as homoserine-lactones, phenazines, siderophores, quinolones, which interfere with the behavior of *A. fumigatus* (Mowat *et al.*, 2010; Moree *et al.*, 2012; Briard *et al.*, 2015). Homoserine-lactones reduce the growth of *A. fumigatus*, whereas phenazines have a dual effect on *A. fumigatus*. At low concentrations such as in an iron-starved environment, pyocyanin, phenazine-carboxamide and phenazine-carboxylic acid could

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stimulate the growth of *A. fumigatus* by reducing the Fe(III). However, at high concentrations, all phenazines (including 1-hydroxy-phenazine) penetrated into the cells and induced the production of reactive-oxygen species and reactive-nitrogen species leading to fungal death (Briard *et al.*, 2015). Such interactions were not demonstrated *in vivo*; however, phenazines were detected in the sputum of cystic fibrosis patients at concentrations which stimulated the growth of *A. fumigatus in vitro* (Wilson *et al.*, 1988; Bjarnsholt *et al.*, 2010; Briard *et al.*, 2015). Regarding quinolones, their effect on *A. fumigatus* or on other fungi was never studied (Diggle *et al.*, 2003).

The present study is centered on rhamnolipids (Rhls), which are secreted by *P. aeruginosa* and other *Pseudomonas* species (Abalos *et al.*, 2001; Moree *et al.*, 2012). To date these molecules have been shown to display only tension active properties. Our study demonstrated that the growth inhibition of *A. fumigatus* and an increased thickness of *A. fumigatus* cell wall following the adhesion of *P. aeruginosa* to *A. fumigatus* is due to the secretion of dirhamnolipids (diRhls) which inhibit the fungal β 1,3 glucan synthase (GS).

Materials and methods

Strains and culture conditions

The *A. fumigatus* reference strain used in this study is CEA17 Δ akuB^{KU80} (ku80) deficient in non-homologous end joining (da Silva Ferreira *et al.*, 2006), which originates from the clinical isolate CBS 144-89, and is as pathogenic as CBS 144-89 in experimental murine aspergillosis models.

CEA17 Δ akuB^{KU80} was used to generate the Δ pksP, Δ hppD, Δ sph3 deletion strains and EMFR S678P mutant. Δ pksP is a 1,8-dihydroxynaphthalene (DHN)-melanin-deficient mutant (Jahn *et al.*, 2000); Δ hppD is a pyo-melanin-deficient mutant and is a kind gift of AA Brakhage (Hans Knöll Institute, Jena, Germany) (Schmaler-Ripcke *et al.*, 2009); Δ sph3 is a spherulin-4-like-deleted mutant, deficient in the production of galactosaminogalactan (GAG), constructed as described in the Supplementary Information (Supplementary Figure 1). EMFR S678P is resistant to caspofungin, due to a point mutation for Serine⁶⁷⁸ in proline in the unique FKS gene coding for the β 1,3 GS (Rocha *et al.*, 2007), and is a kind gift from DS Perlin (Public Health Research Institute, Newark, NJ, USA).

All strains were conserved on 2% (w/v) malt agar slants. One-week-old conidia were recovered from the slants by vortexing with 0.05% (v/v) aqueous Tween 20 solution and used for inoculation in 2YT or Minimal Medium (Briard *et al.*, 2015), RPMI (Clavaud *et al.*, 2012) or Sabouraud (Lamarre *et al.*, 2009) media.

The *P. aeruginosa* used in this study is the reference strain PAO1 (Holloway *et al.*, 1994). This strain was used to generate the deletion strain, Δ rhIA, which do not produce Rhls and is a kind gift

of Niels Højby (Department of clinical microbiology, Rigshospitalet, Copenhagen) (Jensen *et al.*, 2007). *P. aeruginosa* strains were conserved in 2YT medium containing 50% glycerol at -80°C .

Co-culture of *A. fumigatus* and *P. aeruginosa*

To determine the best conditions, preliminary experiments were done with different culture agar-media in Petri dishes (90 mm diameter) as described in the Supplementary Information. The selected co-interaction model is 2×10^7 ku80 conidia inoculated on RPMI plates spotted at the same time with $5 \mu\text{l}$ of 2.5×10^5 PAO1 and incubated for 16 h at 37°C .

Culture of *A. fumigatus* in presence of culture filtrate or dirhamnolipids of *P. aeruginosa*

PAO1 ($A_{600} = 0.05$) was inoculated in 50 ml 2YT for 24 h at 37°C . The bacteria were separated by centrifugation at 15 000 r.p.m. for 10 min and the culture filtrate (CF) was filtrated through $0.22 \mu\text{m}$ Minisart filter (Sigma-Aldrich, Saint-Quentin Fallavier, France). *A. fumigatus* conidia (2×10^7) suspended in 3 ml 0.4% agar containing PAO1 CF was overlaid on RPMI-agar and incubated for 16 h at 37°C . The mycelia grown were observed by transmission electron microscopy (TEM) as described below.

To screen for the CF-active fraction during purification (described below), 1.5×10^4 *A. fumigatus* conidia ml^{-1} were inoculated on eight-well glass bottom Ibidi μ -slides in 2YT medium reconstituted using the PAO1 CF instead of water or in presence of purified diRhls (obtained from CF as described below). The culture was incubated at 37°C for 24 h. Mycelium was observed under light microscopy.

Analysis of the binding of *P. aeruginosa* to *A. fumigatus* polysaccharides, melanin and hyphae

P. aeruginosa was incubated with *A. fumigatus* cell wall α 1,3 glucan, β 1,3 glucan, galactomannan, GAG, melanin, chitin (Sigma-Aldrich), and ku80 or Δ sph3 hyphae as described in the Supplementary Information.

Scanning and transmission electron microscopy preparation

The zone of partial fungal-bacterial inhibition (Figure 1a) from *A. fumigatus* and *P. aeruginosa* co-culture was cut in the Durapore filter, fixed in 2.5% glutaraldehyde in 0.1 M sodium cacodylate buffer for 24 h at 4°C and prepared for scanning electron microscopy and TEM (Lamarre *et al.*, 2009; Briard *et al.*, 2015). As a control, the same culture was prepared without *P. aeruginosa*.

The thickness of the cell wall was calculated from 50 hyphae with six measures per hyphae from TEM pictures using the ImageJ/fiji analysis (Schneider *et al.*, 2012).

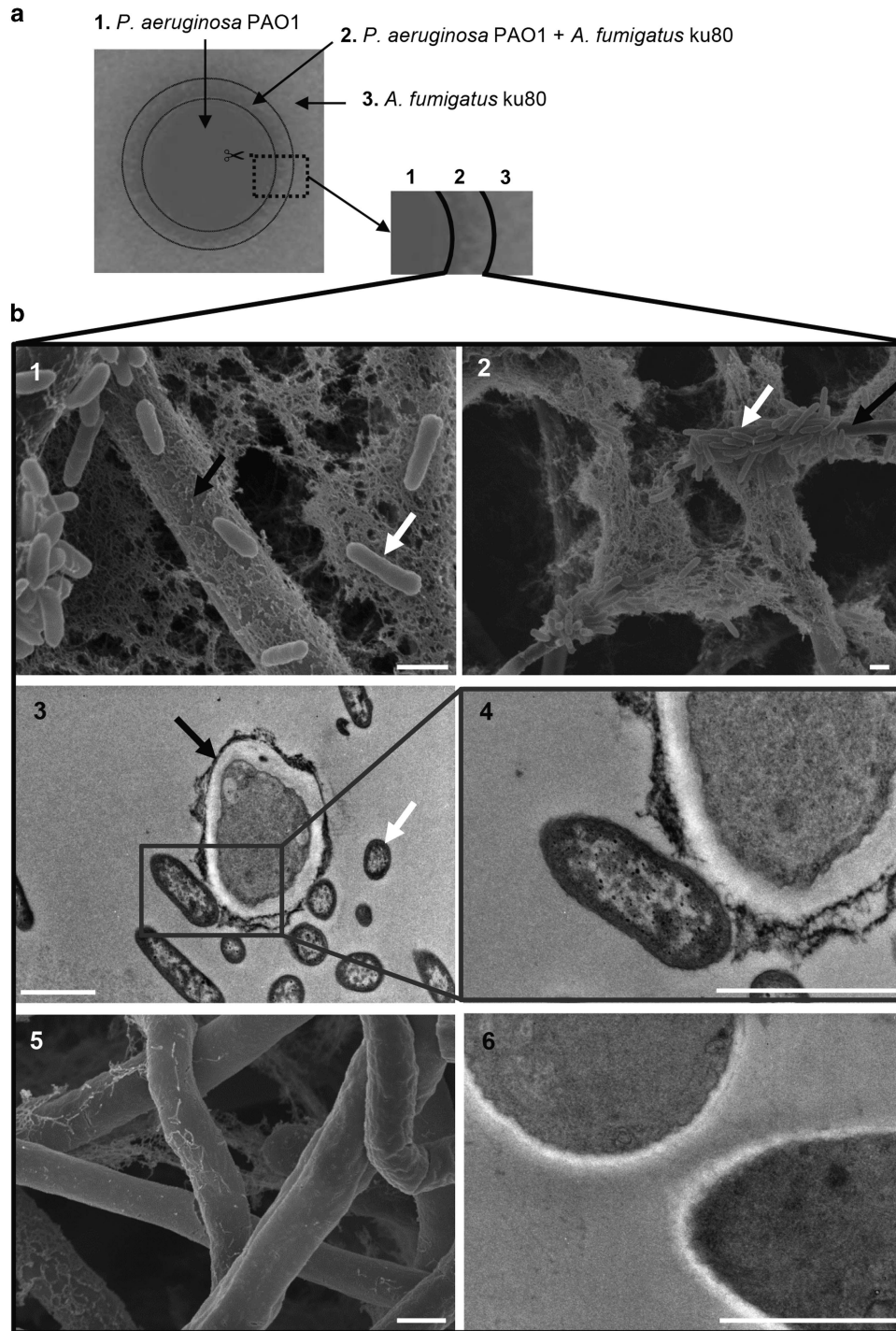


Figure 1 Interaction between *A. fumigatus* ku80 and *P. aeruginosa* PAO1 strains. (a) Zone of partial inhibition (dotted square) between *A. fumigatus* and *P. aeruginosa*. (b) Scanning (b1, 2 and 5) and transmission (b3, 4 and 6) electron microscopy views of the zone of partial inhibition; (b1–4) ku80-PAO1; (b4) enlargement of the interaction zone showed in b3; (b5 and 6) control ku80 in the absence of PAO1. White arrow, bacteria; black arrow, fungus. Note the presence of an increased amount of fungal ECM in presence of *P. aeruginosa* (b1 compared to b5 and b3 compared to b6). Scale bars, 1 μ m.

Extraction and isolation of the active molecules from P. aeruginosa 24 h culture filtrate

The PAO1 24 h CF was extracted twice with ethyl acetate (1:1) and the organic phase was evaporated and dried. A mixed fraction, F-diRhls, containing two to three diRhls depending on the CF batches,

was purified from the organic extract as described in the Supplementary Information. Each fraction was identified by high-resolution mass spectrometry, which correlated with the published data (Sharma *et al.*, 2007; Abdel-Mawgoud *et al.*, 2010; Arutchelvi and Doble, 2010).

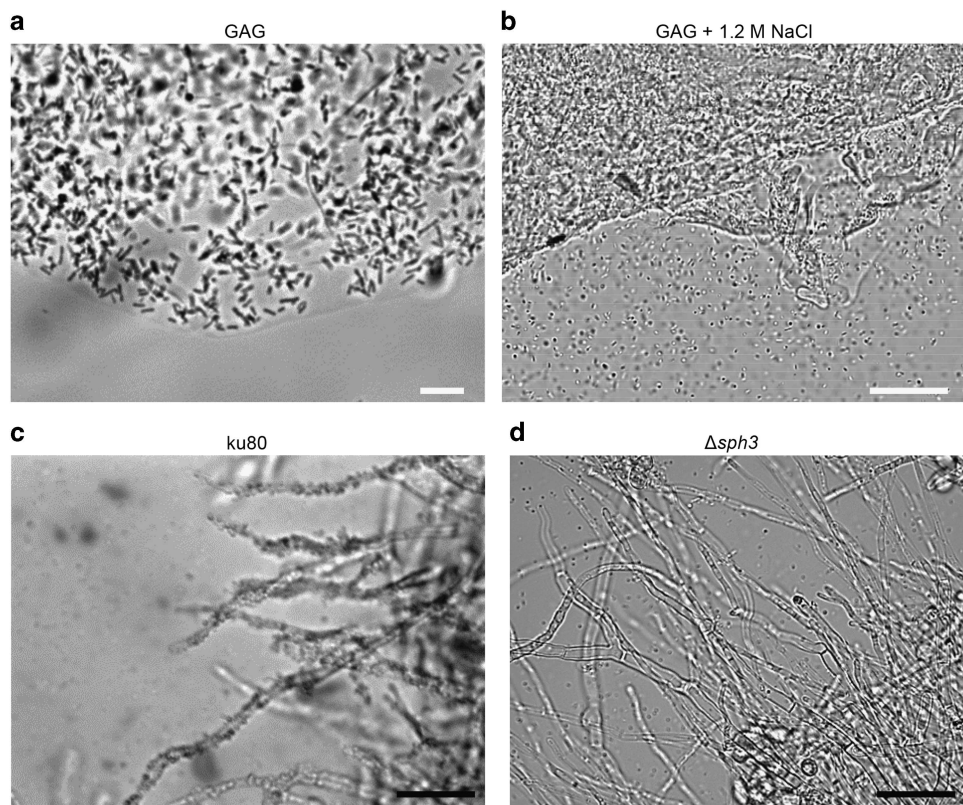


Figure 2 Binding of *P. aeruginosa* to *A. fumigatus* GAG. (a) incubation of PAO1 with purified GAG. (b) Incubation of PAO1 with purified GAG in presence of 1.2 M NaCl. (c) Adhesion of PAO1 to *A. fumigatus* ku80 hyphae. (d) Absence of adhesion of PAO1 to *A. fumigatus* $\Delta sph3$ mutant hyphae, unable to produce GAG. Note that the binding of PAO1 to GAG is inhibited under high salt concentration (b). Scale bars, 25 μm .

Chemical synthesis of dirhamnolipid analogs diRhamnose- C_3 and diRhamnose- C_8

The synthesis of two diRhls analogs, diRhamnose- C_3 (diRha- C_3) and diRhamnose- C_8 (diRha- C_8) bearing a simpler aglycon—a propyl and an octyl chains, respectively, is described in the Supplementary Information.

Method for susceptibility testing and biomass quantification by crystal violet

Minimal effective concentration (MEC) of the molecules was determined according to the Clinical Laboratory Standards Institute M38-A2 protocol (NCCM) by microdilution method in 96-well plates using crystal violet biomass quantification as previously described (Briard *et al.*, 2015).

Determination of the fractional inhibitory concentration (FIC) index (FIC_i)

The FIC_i of drug interactions was determined via checkerboard titration assays (Clavaud *et al.*, 2012). The concentrations of the antifungal agents ranged from 0.008 to 1 $\mu\text{g ml}^{-1}$, 0.004 to 0.5 $\mu\text{g ml}^{-1}$ and 0.031 to 4 $\mu\text{g ml}^{-1}$ for caspofungin, voriconazole and itraconazole, respectively. The concentrations of F-diRhls ranged from 0.03 to 4 mM. Serial twofold dilutions of antifungals were prepared with the assay culture in 2YT in 96-wells flat bottom plates as

described above. The microtiter plates were incubated at 37 °C for 20 h and the growth biomass was assessed using crystal violet and resazurin methods (Clavaud *et al.*, 2012; Briard *et al.*, 2015). The pharmacodynamic interaction analysis was done as described (Mavridou *et al.*, 2015). Briefly, the synergistic, additive or antagonistic effect of paired combinations of drugs was captured by the FIC_i: (a) $\text{FIC} = \text{FIC}_{\text{drugA}} + \text{FIC}_{\text{drugB}}$; (b) $\text{FIC}_{\text{drugA}} = \text{MIC}_{\text{drugsA+B}} / \text{MIC}_{\text{drugA}}$; and (c) $\text{FIC}_{\text{drugB}} = \text{MIC}_{\text{drugsA+B}} / \text{MIC}_{\text{drugB}}$. If the $\text{FIC} \leq \text{FIC}_{\text{min}}$ value is lower than 1, this indicates synergistic interactions between two drugs; if $\text{FIC} \geq \text{FIC}_{\text{max}}$ value is higher than 1.25, then an antagonistic interaction exists and between these two values, the interactions between the two drugs are additive.

Effect of diRhI on cell wall

Glucosamine and galactosamine determination in ku80 cell wall in presence and absence of F-diRhls and preparation of cell-free glucan and chitin synthases extracts are described in the Supplementary Information.

Statistical analysis

Data are reported as means \pm s.e.m. Comparisons, performed with Graph Pad Prism 3.0 software (<http://graphpad-prism.software.informer.com/3.0/>) and analysis of variance statistical test.

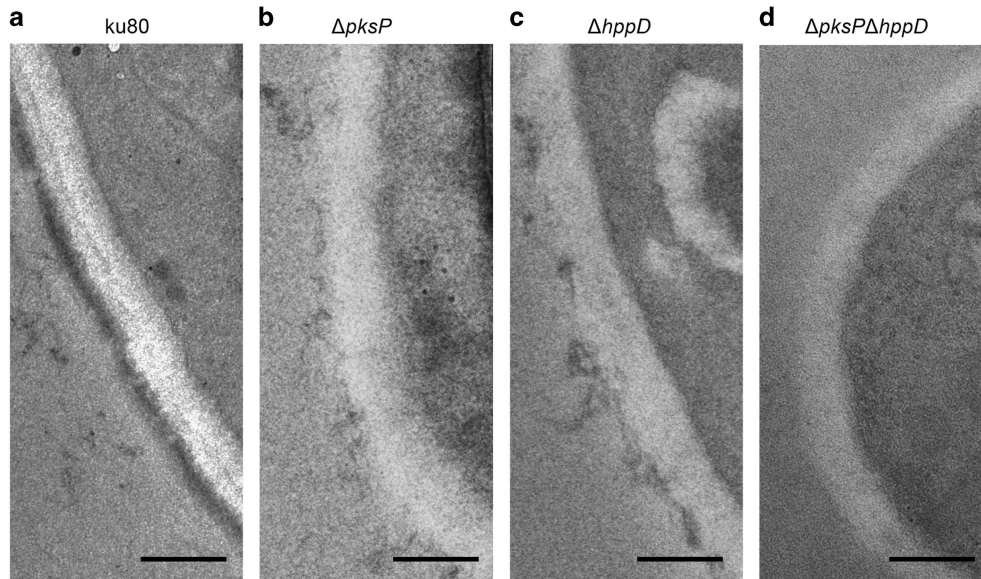


Figure 3 Presence of melanin in the ECM of *A. fumigatus* ku80 during co-cultivation with *P. aeruginosa* PAO1 in RPMI. (a) Parental strain ku80. (b) *AfΔpksP* mutant unable to produce DHN-melanin. (c) *AfΔhppD* mutant unable to produce pyo-melanin. (d) Double *AfΔpksPΔhppD* mutant. Note the absence of melanin in *AfΔpksPΔhppD*, showing that melanin in ECM is a mixture of DHN and pyo-melanin. Scale bars, 250 nm.

Results

Morphological modifications of A. fumigatus mycelium in presence of P. aeruginosa

Preliminary experiments determined the best culture conditions allowing the fungus and the bacterium to grow together: 2×10^7 ku80 *A. fumigatus* conidia were inoculated on an RPMI agar plates and at the same time $5 \mu\text{l}$ of 2.5×10^5 PAO1 *P. aeruginosa* was spotted in the center of the Petri-dish. Under these experimental conditions, a zone of partial fungal inhibition was observed at the junction between the bacterial and the fungal colonies (Figure 1a). The zone of partial inhibition was observed under scanning electron microscopy and TEM (Figure 1b). The first observation under scanning electron microscopy showed an electron-dense extracellular matrix (ECM) embedding the hyphae of *A. fumigatus* which was more abundant in the presence of the bacteria than in its absence (Figures 1b1, b2 and b5). PAO1 cells adhered to the hyphae, mainly at their apex and to the ECM (Figures 1b1, b2). To identify the fungal cell wall molecule to which *P. aeruginosa* was binding, the *A. fumigatus* cell wall polysaccharides α 1,3 glucan, β 1,3 glucan, galactomannan, chitin, GAG and melanin (Latzg , 2010) were incubated with the PAO1 cells and scored the binding under light microscopy. The bacteria only bound to GAG (Figure 2; Supplementary Figure 2 and data not shown). Addition of 1.2M NaCl prevented the binding of the bacteria to GAG showing that PAO1 binding was due to ionic interactions (Figures 2a and b). To confirm that GAG is responsible for the binding of *P. aeruginosa* to *A. fumigatus* ku80 hyphae, the spherulin-4 like gene, coding for Sph3 involved in the synthesis of GAG in *A. fumigatus*

(Bamford *et al.*, 2015) was deleted in ku80 (Supplementary Figure 1) and the Δsph3 strain was incubated with *P. aeruginosa*. As shown in Figures 2c and d, *P. aeruginosa* did not bind to Δsph3 hyphae.

The interaction between bacterium and fungus induced formation of an electron-dense ECM around the fungal hyphae. This electron-dense material was absent in the control cultures (in absence of bacteria) as well as at the contact point between the fungal cell wall and *P. aeruginosa* PAO1 (Figures 1b3). This electron-dense material is reminiscent of melanin observed around hyphae in the *A. fumigatus* biofilm (Beauvais *et al.*, 2007). Three *A. fumigatus* deletion mutants ΔpksP , ΔhppD and $\Delta\text{pksP}\Delta\text{hppD}$ affected in DHN-, pyo- or both melanin synthesis, respectively (Jahn *et al.*, 2000; Schmalzer-Ripcke *et al.*, 2009), were grown in with the presence of PAO1. As shown in the Figure 3, this electron-dense material was partially lost in ΔpksP and ΔhppD , and was absent in $\Delta\text{pksP}\Delta\text{hppD}$. Therefore, the electron-dense material present in the ECM in presence of PAO1 was composed of both DHN- and pyo-melanin.

TEM also showed that *A. fumigatus* presented a thicker cell wall ($104\text{ nm} \pm 3.2$) when grown in the presence of *P. aeruginosa* (Supplementary Figure 3), compared to that in its absence ($72\text{ nm} \pm 1.19$). This observation suggested that bacterial co-culture induced modification of the fungal cell wall structure.

P. aeruginosa secreted dirhamnolipids are responsible for the *A. fumigatus* cell wall modification
To test whether *P. aeruginosa* induced cell wall modifications in *A. fumigatus* requires its direct

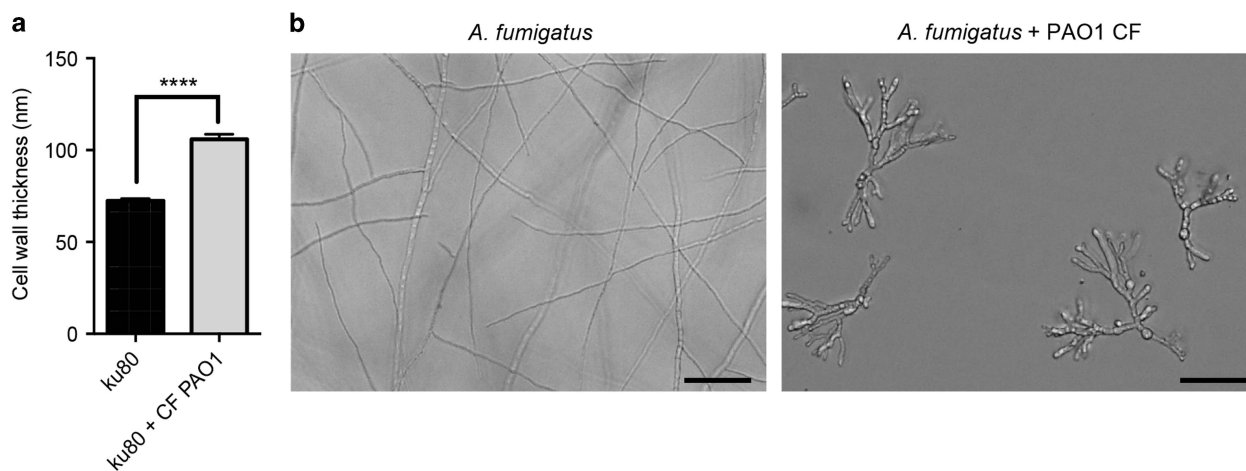


Figure 4 *A. fumigatus* hyphal morphology grown in presence of 24 h PAO1 CF. (a) Thickness of *A. fumigatus* ku80 cell wall in presence of CF in RPMI, calculated from 50 hyphae with six measures per hyphae from TEM pictures using the ImageJ/fiji analysis (Schneider et al, 2012). (b) Light microscopy images showing the increased branching of the fungal hyphae in presence of CF. Scale bar 50 μ m.

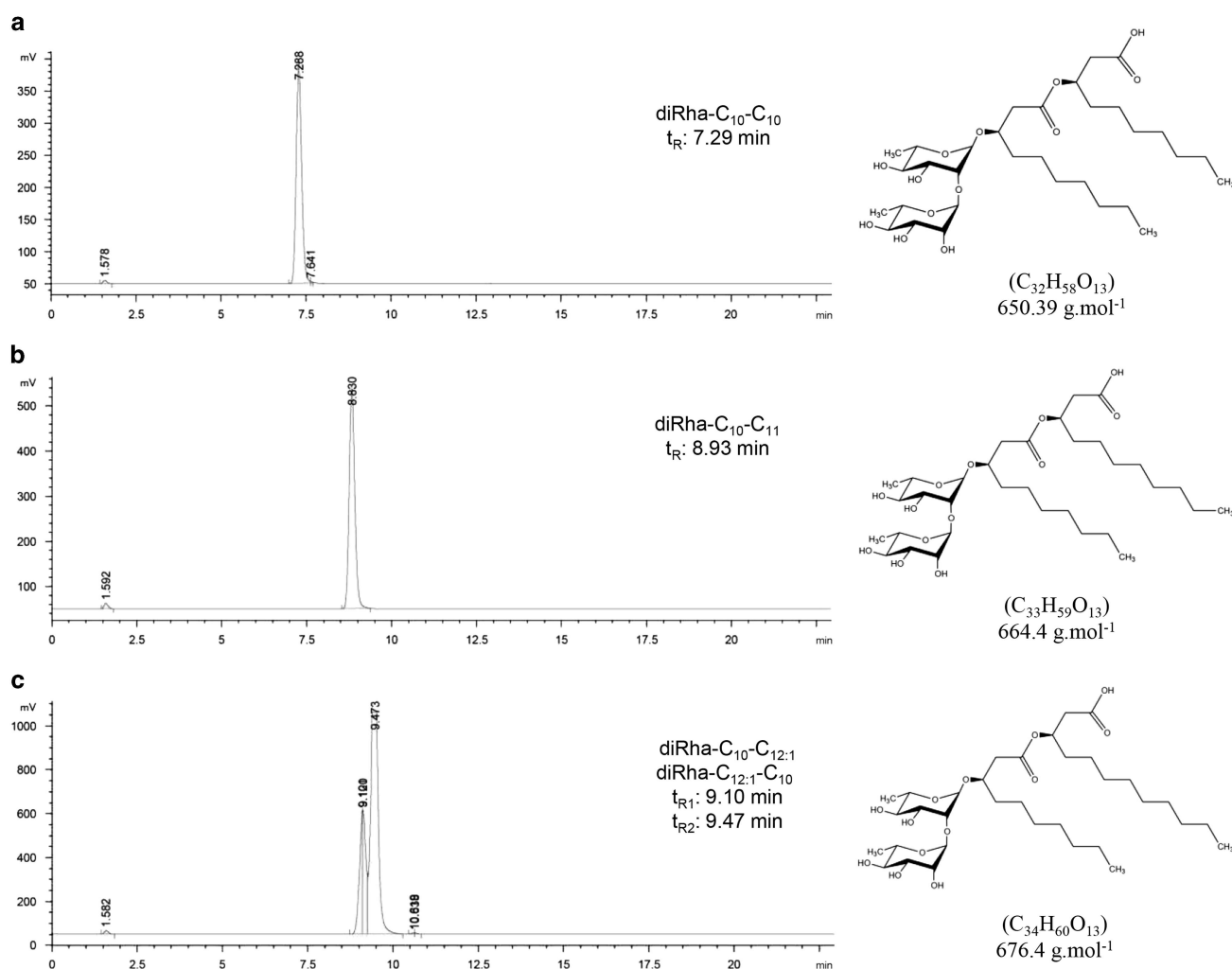


Figure 5 Analytical RP-HPLC profiles of the dirhamnolipids isolated following RP-HPLC purification of the 24 h culture filtrate of *P. aeruginosa*. Elution from a C18 Kromasil column (4.6 \times 150 mm, 5 μ m, 100 \AA) at 1.0 ml min⁻¹ with CH₃CN in 0.08% aqueous trifluoroacetic acid (60 \rightarrow 100% over 15 min.), detection with an evaporative light scattering detector and identification according to the literature (Sharma et al., 2007; Abdel-Mawgoud et al., 2010; Arutchelvi and Doble, 2010). (a) diRha-C₁₀-C₁₀. (b) diRha-C₁₀-C₁₀-CH₃ (diRha-C₁₀-C₁₁) corresponding to the methyl ester of diRha-C₁₀-C₁₀. (c) diRha-C₁₀-C_{12:1} and diRha-C_{12:1}-C₁₀ bearing an unsaturated fatty acid.

contact or due to molecules secreted from bacteria, *A. fumigatus* conidia were incubated in the presence of bacterial CF. The CF from either 2YT or RPMI media promoted the formation of an ECM around fungal hyphae and induced thickening of the fungal cell wall (Figure 4a). Increase in the fungal cell wall thickness (106 nm) was comparable to that observed when bacteria are directly in contact with the fungus. In addition, *A. fumigatus* growth was reduced with multibranched hyphae, the phenotype similar when this fungus is grown in the presence of antifungal echinocandins (Figure 4b; Perlin, 2011).

These results suggested that diffusible molecules in *P. aeruginosa* PAO1 CF were responsible for the fungal cell wall modifications.

The molecules responsible for the cell wall phenotype were then purified from a 2YT CF of *P. aeruginosa* after 24 h of growth. Only one high-performance liquid chromatography fraction gave the cell wall phenotype mentioned above when incubated with *A. fumigatus*. MS analyses (data not shown) showed that this fraction contained a mixture of diRhls (F-diRhls). Following purification, three rhamnobioides differing by the length of their

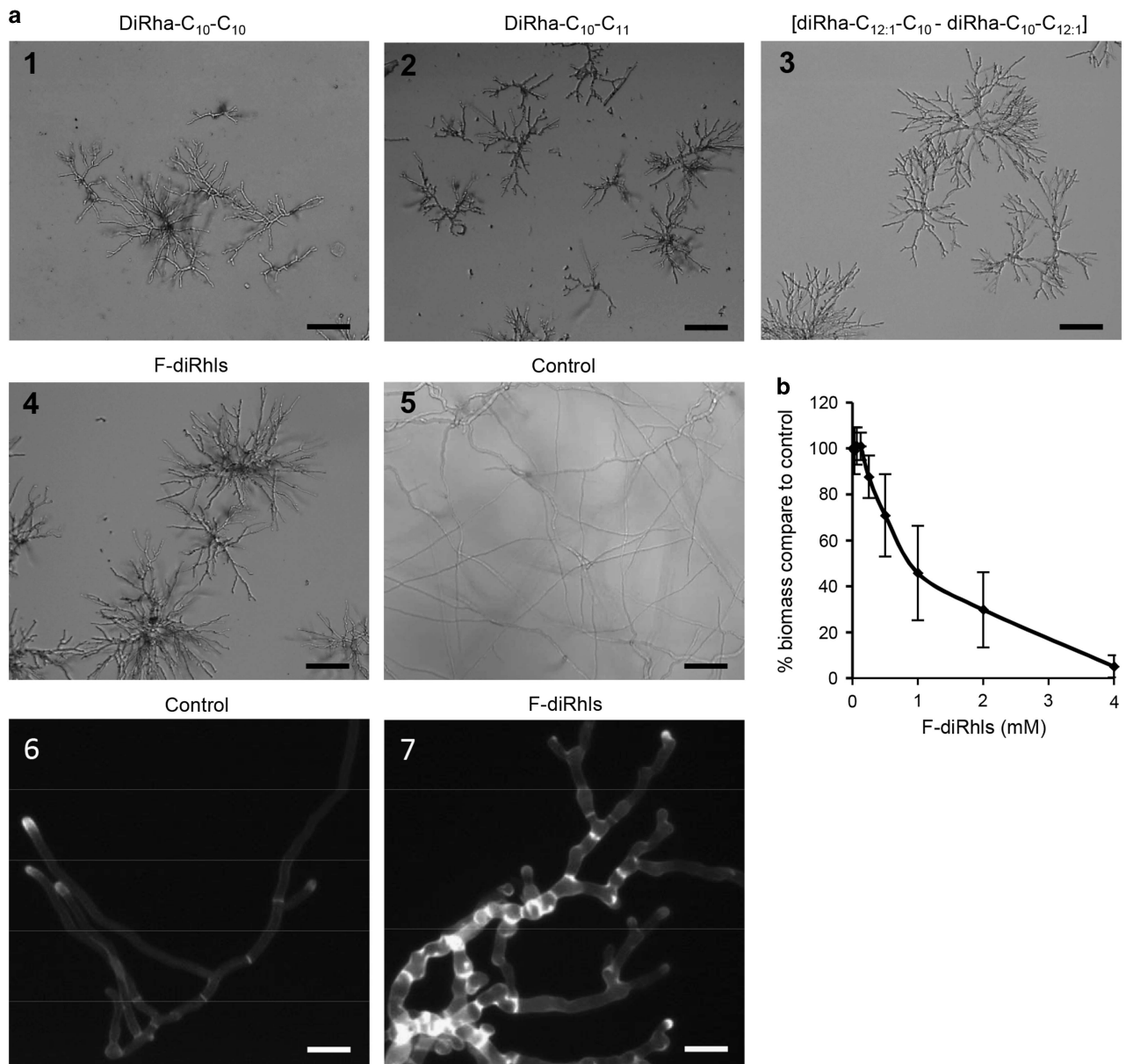


Figure 6 Impact of the diRhls on the *A. fumigatus* mycelial growth. (a1–4) Light microscopy of *A. fumigatus* grown for 24 h in the presence of diRha-C₁₀-C₁₀, diRha-C₁₀-C₁₁, [diRha-C_{12;1}-C₁₀ - diRha-C₁₀-C_{12;1}] or F-diRhls at 4 mM. (a5) *A. fumigatus* grown in the absence of diRhls. Note the increased branching of mycelium after growth with diRhls. (a6 and 7) UV microscopy of calcofluor white labeling of *A. fumigatus* in the absence or presence of diRhls. Note the increased amount of fluorescence in the hyphae in presence of F-diRhls, which is representative of an increase in the chitin. (b) Reduction of mycelial growth by different concentrations of F-diRhls. Scale bars, 50 μ m (a1–a5); 10 μ m (a6 and a7).

fatty acid chains were identified: diRha-C₁₀-C₁₀ eluting at t_R: 7.29 min had *m/z* 650.39 (HRMS-ESI⁺ for C₃₂H₅₈O₁₃ ([M+Na]⁺, 673.3775), diRha-C₁₀-C₁₁ eluting at t_R: 8.93 min had *m/z* 664.4 (HRMS-ESI⁺ for C₃₃H₆₀O₁₃ ([M+Na]⁺, 687.3932), diRha-C₁₀-C_{12:1} and diRha-C_{12:1}-C₁₀ eluting at t_R: 9.10 min and 9.47 min, respectively, had *m/z* 676.4 (HRMS-ESI⁺ for C₃₄H₆₀O₁₃ ([M+Na]⁺, 699.3932) (Figure 5). F-diRhls was majorly composed of diRha-C₁₀-C₁₀ (60–100% depending of the CF batches). Each diRhls was either tested individually or in combination (F-diRhls) on *A. fumigatus*.

TEM observation of the hyphae after treatment with 4 mM diRha-C₁₀-C₁₀ showed an increased thickness of the cell wall (215 nm ± 5,6) and ECM as observed previously with *P. aeruginosa* PAO1 and its CF (Supplementary Figures 4a and c). This increased thickness of the cell wall could be correlated to an increased concentration of the GAG adhesin at the cell surface. This hypothesis was verified biochemically by determination of the amount of galactosamine monomers, representative of GAG, in the extracted cell wall in absence and presence of 4 mM F-diRhls; 1.74 µg (± 0.44)

galactosamine per mg mycelium dry weight was found in *A. fumigatus* cell wall in the absence of F-diRhls which was significantly increased to 11.26 µg (± 0.14) galactosamine per mg mycelium dry weight in the presence of F-diRhls.

To confirm that F-diRhls were responsible for the morphological changes in *A. fumigatus* cell wall, co-culturing of the fungus and the *P. aeruginosa* $\Delta rhIA$ mutant, which is unable to synthesize Rhls, was undertaken. The $\Delta rhIA$ mutant bound to ku80 hyphae similar to *P. aeruginosa* PAO1 but no melanin was present as in the control without bacteria (Figure 1b6; Supplementary Figure 4b). Moreover, the cell wall thickness was not modified in presence of $\Delta rhIA$ (Supplementary Figure 4c). These results confirmed that the cell wall phenotype induced by the bacterium was due to the presence of diRhls secreted by PAO1.

Fungal growth phenotype is due to the inhibition of β 1,3 glucan synthase by diRhls

A. fumigatus had a reduced growth and altered morphology in presence of diRhls similar to that

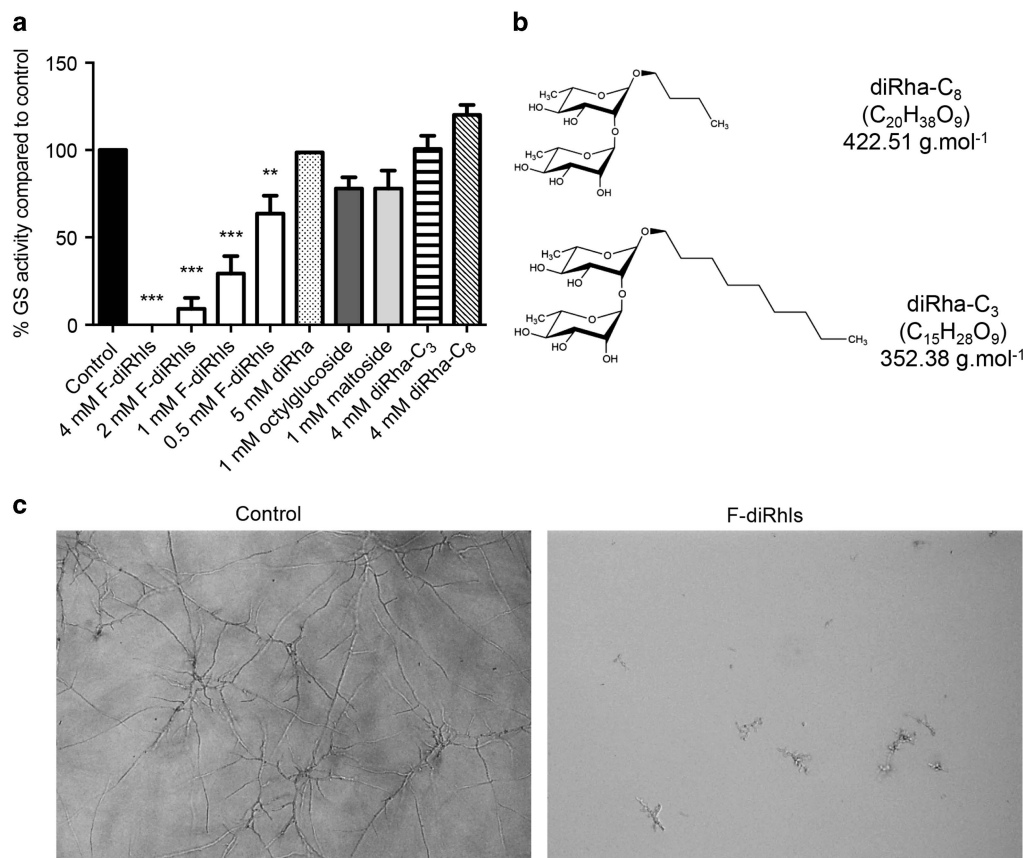


Figure 7 Specific inhibition of *A. fumigatus* ku80 β 1,3 glucan synthase activity (***P* = 0.0036; ****P* < 0.0001); 0.5–4 mM F-diRhls, 5 mM rhamnobiose (diRha), 1 mM octylglucoside and maltoside, 4 mM propyl- and octyl-rhamnobioside analogs (diRha-C₃ and diRha-C₈, respectively) were added in the glucan synthase reaction mixture (*P* > 0.05). (b) Structure of the chemically synthesized diRha-C₃ and diRha-C₈. (c) phenotype of *A. fumigatus* EMFR S678P in absence and presence of 0.5 mM F-diRhls.

observed with CF (Figures 4b and 6a). The effect of F-diRhls on *A. fumigatus* was never fungicidal whatever concentration used. The short hyperbranched morphology was reminiscent of the echinocandin effect. Accordingly, only an MEC, which is the lowest drug concentration resulting in aberrant hyphal growth, can be measured (Arikan *et al.*, 2002). The MEC was 4 mM, and resulted in 90% inhibition of *A. fumigatus* growth (Figure 6b). Calcofluor white treatment showed an increase in hyphal fluorescence compared to untreated control after treatment with purified diRhls and F-diRhls at MEC, suggesting an increase in the concentration of cell wall chitin (Figures 6a6 and 7). This hypothesis was verified biochemically by determination of the amount of glucosamine monomers, representative of chitin, in the extracted cell wall treated and untreated with F-diRhls at MEC; in the absence of F-diRhls, 9.15 µg (± 1.55) glucosamine per mg mycelium dry weight was found, which was significantly increased (31.4 µg (± 0.38) glucosamine per mg mycelium dry weight) in the presence of F-diRhls. An increase in the cell wall chitin amount was also observed after incubation of the fungus with echinocandins (Walker *et al.*, 2010). These data suggested that diRhls may have a direct effect on the β1,3 GS activity. Indeed, the F-diRhls inhibited the GS activity in a dose-response manner (Figure 7a). To determine if the inhibition of the GS activity was due to the detergent characteristic of the F-diRhls, other neutral nonionic detergent glycolipids such as octylglucoside and maltoside were tested. These detergents disrupt protein-lipid and lipid-lipid interactions in the membrane, which could disturb their activity. Figure 7a showed a slight statistically insignificant inhibition of GS activity by octylglucoside and maltoside, showing that the dose-dependent inhibition of GS by F-diRhls was mainly due to their GS-specific effect. Moreover, we verified the specificity of F-diRhls on the GS activity by testing the chitin synthase activity in presence of 1 mM F-diRhls, octylglucoside and maltoside. All compounds induced a similar statistically insignificant decrease of the chitin synthase activity (Supplementary Figure 5), showing that in contrast to GS inhibition, their minimal effect on the chitin synthase was not specific, and was due to biosurfactant activity on the membranes.

To investigate the active moiety of F-diRhls on GS activity, we tested the inhibitory effect of rhamnobiase and the propyl- and octyl-rhamnobiase analogs, equipped with a linear three carbons (diRha-C₃) and eight carbons (diRha-C₈) aglycon (Figures 7a and b). None of these molecules inhibited GS activity, suggesting that the branched lipidic tail with β-hydroxy fatty acids of F-diRhls is essential for its activity against GS.

The *A. fumigatus* EMFR S678P strain, which is a mutant resistant to caspofungin consecutively to a point mutation in the binding site of this echinocandin to GS, has a MEC ≥ 16 µg caspofungin ml⁻¹, while the MEC of caspofungin for parental ku80 strain is 0.25 µg ml⁻¹ (Rocha *et al.*, 2007).

Table 1 Pharmacodynamic interaction concentration index (FIC_i) between F-diRhls and CAS, ITR or VTR for *A. fumigatus* ku80

F-diRhls (mM)	MEC ^a or MIC ^b drugs alone			MEC or MIC drugs in combination			ΣFIC _{min} , ΣFIC _{max} index ^c		
	CAS (µg ml ⁻¹)	ITR (µg ml ⁻¹)	VRC (µg ml ⁻¹)	F-diRhls, CAS	F-diRhls, ITR	F-diRhls, VRC	F-diRhls, CAS	F-diRhls, ITR	F-diRhls, VRC
4	0.25	16	0.3	1, 1	2, 1	1, 0.063	2.2, 4.25	0.56, 0.5	0.5, 0.56

Abbreviations: CAS, caspofungin; diRhls, dirhamnolipids; ITR, itraconazole; MEC, minimal effective concentration; VTR, voriconazole.

^aMEC for F-diRhls and CAS.

^bMIC for itraconazole and voriconazole.

^cFIC_i ΣFIC_{min} < 1 = synergy; FIC_i ΣFIC_{max} > 1.25 = antagonism; 1 < FIC_i < 1.25 = additive.

Surprisingly, the MEC of F-diRhls on the EMFR S678P strain was 0.5 mM compared to the 4 mM value with the *A. fumigatus* ku80 strain (Figure 7c). This result indicated that the diRhls and the echinocandin do not bind to the same GS site and that the S678P mutation favored the inhibitory effect of F-diRhls towards GS activity. When F-diRhls and caspofungin were tested in combination on the ku80 strain, the combined effect of two molecules was antagonist with a FIC_i ΣFIC_{max} of 4.25 (Table 1). In contrast to the previously reported additive effect of caspofungin with voriconazole (Elefanti *et al.*, 2013), the results presented in Table 1 showed that the combined effect of F-diRhls and voriconazole or itraconazole was synergistic with a FIC_i ΣFIC_{min} of 0.5 and 0.56, respectively. These data supported the fact that diRhls and caspofungin target the same component of *A. fumigatus*, the GS.

This study shows that the F-diRhls induce several morphological effects: (i) they modify the nature of fungal ECM surrounding the *A. fumigatus* hyphae; (ii) they inhibit the growth of *A. fumigatus* and induce the formation of short multibranched hyphae, which results from the specific inhibition of the fungal GS activity; (iii) they induce a thickening of the cell wall due to compensatory reactions resulting from an increased chitin concentration to palliate the antifungal effect of dirhamnolipids; and (iv) the site of action of diRhls on GS is different from that of echinocandins, which facilitates its synergistic activity with azoles.

Discussion

This study has shown that *P. aeruginosa* binds strongly to the *A. fumigatus* hyphae. We previously reported that the polysaccharides composing *A. fumigatus* ECM were α1,3 glucan, GAG and galactomannan (Beauvais *et al.*, 2007). In the binding experiments, in presence of *P. aeruginosa*, only GAG was found to be the molecule responsible for *P. aeruginosa* to *A. fumigatus* binding. GAG has been reported previously to be a major adhesin of *A. fumigatus* with many functions associated to the invasion of the host tissue (Gresnigt *et al.*, 2014; Robinet *et al.*, 2014). The current study also demonstrated that the production of GAG is increased in response to bacterial assault against the fungus.

Interestingly, the bacteria bind to the fungus principally at the apex that is at the site of the metabolic fungal activity. Increased fixation at the tip of *A. fumigatus* hyphae, where intense production of exudates takes place (Toljander *et al.*, 2007), may be a reason for *P. aeruginosa* to obtain nutrients from the fungus. Similarly, *P. aeruginosa* binding to the hyphal (but not to the yeast) form of *Candida albicans* suggests an interaction via hyphal specific glycoproteins, lectins or adhesins (Hogan and Kolter, 2002). The ligand has not been identified yet.

The present and previous studies from our laboratory and others showed that *P. aeruginosa* produces two types of molecules which can influence *A. fumigatus* growth. The first ones are the toxins which directly kill *A. fumigatus*. Examples of those are phenazines and homoserine lactones (Mowat *et al.*, 2010; Moree *et al.*, 2012; Briard *et al.*, 2015), which display different toxic effects. Phenazines kill the fungus by inducing intracellular reactive-oxygen species and reactive-nitrogen species production (Briard *et al.*, 2015). Homoserine lactones repressed *C. albicans* filamentation (Hogan *et al.*, 2004) and reduced *A. fumigatus* growth, but their mode of action is not yet studied (Mowat *et al.*, 2010).

The second type of compounds is stress molecules which induces a typical anti-stress response from *A. fumigatus*. An example is the diRhls. *A. fumigatus* responds to diRhls in a way which is very similar to any response to environmental molecules including antifungals targeting cell wall such as echinocandins, calcofluor white or congo red (Ram and Klis, 2006; Perlin, 2011; Lee *et al.*, 2012). Interestingly, use of diRhls led to the discovery of a new cell wall compensatory mechanism not previously described, which is the melanin synthesis.

Melanin synthesis is associated with reduced fungal susceptibility to a variety of aggressions. Infection increases the expression of melanin synthesis genes by various fungi and the more virulent strains produce the more melanin (Eisenman and Casadevall, 2012). In the host, melanin interferes with the normal function of phagocytic cells and scavenges reactive-oxygen species (Eisenman and Casadevall, 2012; Chamilos *et al.*, 2016). In *A. fumigatus*, melanin production has been described in biofilm (Beauvais *et al.*, 2007) or *in vivo* the production of DHN-melanin was suggested during experimental aspergillosis under conditions which can be considered to be starved stress conditions (Langfelder *et al.*, 2001). The origin of melanin in *A. fumigatus* biofilm is still unknown. It was hypothesized that melanin resulted from air-oxidation of melanin precursors in the biofilm channels. The use of two *A. fumigatus* mutants unable to produce DHN- and pyo-melanin when co-cultured with *P. aeruginosa* indicated that ECM synthesized by parental *A. fumigatus* in presence of *P. aeruginosa* is composed of DHN- and pyo-melanin (Figure 3). Interestingly the melanin produced by the fungus in response to PAO1 disappeared at the contact region with the bacterium. *Klebsiella aerogenes* synthesizes a dopamine precursor from L-tyrosine that was used by *Cryptococcus neoformans* for melanin production, which required a fungal laccase (Frasés *et al.*, 2006). *A. fumigatus* expressed laccases during vegetative growth and a 4-hydroxyphenylpyruvate dioxygenase (HppD) responsible for the synthesis of pyo-melanin from L-tyrosine present in different media (Sugareva *et al.*, 2006; Schmalzer-Ripcke *et al.*, 2009). Moreover,

A. fumigatus interacting with *P. aeruginosa* Δ rhlA, which does not produce any Rhl, did not synthesize melanin demonstrating that the diRhls were inducing the synthesis of the melanin protecting layer by the fungus. However, the antibacterial function of the fungal melanin and especially the role of the two different melanins have not been investigated.

In *P. aeruginosa*, because of their metabolic synthesis, Rhls are usually produced as a mixture of homologs di- and mono-Rhls diverging in terms of lipid chain length. To our knowledge, diRha-C₁₀-C₁₀ is always the major component (Abalos *et al.*, 2001; Sha *et al.*, 2012; Christova *et al.*, 2013; Singh *et al.*, 2013; Gogoi *et al.*, 2016). DiRha-C₁₀-C₁₀ showed higher antifungal activities on plant pathogens such as *Penicillium juniculosum* and *Bothyitis cinerea* than Rhls (Sha *et al.*, 2012). DiRha-C₁₀-C₁₀ was also active against *C. albicans* with a MIC > 0.15 mM (Singh *et al.*, 2013). At 7 mM, diRha-C₁₀-C₁₀ disrupted the biofilms and their biosurfactant activity inhibited cell adhesion. The presence of diRhls after 24 h in PAO1 CF is in accordance with the kinetics of Rhls synthesis by *P. aeruginosa* after the growth ceased (24–96 h) (Haba *et al.*, 2003; Laabei *et al.*, 2014; Schmidberger *et al.*, 2014). Rhamnolipids are over-produced *in vitro* by *P. aeruginosa* during nutrient limitation such as in iron starvation (Schmidberger *et al.*, 2014). *In vivo* in the host, iron concentration is limited, and 2.4–15.3 $\mu\text{g ml}^{-1}$ Rhls were found in sputa of all cystic fibrosis patients chronically infected by *P. aeruginosa* (Kownatzki *et al.*, 1987). Mucoid isolates from chronically infected patients produced more Rhls than non-mucoid isolates (Bjarnsholt *et al.*, 2010). Indeed, they are components of *P. aeruginosa* biofilm and protect the bacteria against phagocytosis by polymorphonuclear neutrophils by inducing necrosis of the latter (Van Gennip *et al.*, 2009).

Until now the effect of all Rhls, including diRha-C₁₀-C₁₀ and its homologs, was attributed only to their biosurfactant activity, which destabilizes plasma membrane resulting to quantitative changes in phospholipid headgroup (Sha *et al.*, 2012; Sotirova *et al.*, 2012; Singh *et al.*, 2013). Similarly to the role of dihydromaltophilin on plasma membrane sphingolipid biosynthesis (Li *et al.*, 2009), the disorganization of the plasma membrane by diRhls may stimulate cell wall synthesis by activating cell wall integrity signaling pathways, which could explain the thickening of the cell wall. The combination of the rhamnobiase moiety and of the branched aglycon in the diRhls is essential for the interaction with the GS since neither rhamnobiase nor diRha-C₃ and diRha-C₈ analogs showed an inhibitory activity. Similar inhibition of the GS and growth phenotype in *A. fumigatus* has been observed using echinocandins, such as caspofungin, micofungin or anidulafungin, which are lipopeptides differentiated by their aliphatic tails (Perlin, 2011; Clavaud *et al.*, 2012). Enzymatic removal of the aliphatic tails of the

drugs renders them inactive (Perlin, 2011). The antagonistic effect of caspofungin and F-diRhls on *A. fumigatus* confirmed that both lipid conjugates had GS as target. It has been speculated that the tail of the drugs may intercalate into the bilayer resulting in inhibition. Such drug–target interactions would not require the drugs to enter the cell and they may act on the enzyme from the extracellular face of the cell membrane. At the same time, the fact that echinocandin-resistant strain (EMFR S678P) showed high susceptibility to F-diRhls suggested that the inhibitory binding site of F-diRhls differs from that of echinocandin and that the S678P mutation enhances the affinity of F-diRhls for GS. In contrast, the combination of azoles and F-diRhls on *A. fumigatus* was synergic as it was found previously for azoles and echinocandins, which have different inhibitory binding sites (Mavridou *et al.*, 2015).

Other studies showed that *in vitro* *P. aeruginosa* inhibited the growth of *A. fumigatus*, the cystic fibrosis isolates being more inhibitory than non-cystic fibrosis isolates (Ferreira *et al.*, 2015). Moreover, until now it is not known how *A. fumigatus* colonized better in the respiratory tract of cystic fibrosis patients when they got an infection with *P. aeruginosa* and whether the fungus is in close contact with the bacteria. We showed previously that at low ‘*in vivo*’ concentrations, phenazines stimulated the growth of *A. fumigatus* by providing iron uptake (Briard *et al.*, 2015). It is particularly conceivable that *in vivo* *A. fumigatus* establishes in the same niches as *P. aeruginosa* taking advantage of favorable growth conditions due to lung damages. *A. fumigatus* may also benefit from the *P. aeruginosa* biofilm for host immune protection due to the role of Rhls and alginate in the reduction of the host immune responses (Baxter *et al.*, 2013; Briard *et al.*, 2015). Moreover, recent studies from our group also showed that these interactions can be observed at distance. Fungi and bacteria do not have to be in contact and volatiles from *P. aeruginosa* can be used as nutrients by the fungus (Briard *et al.*, 2016). These recent data point out the need to investigate in depth the pulmonary microbiota and the cross-talks between bacteria and fungi. In cystic fibrosis patients, the microbiota is completely disorganized and becomes particularly rich in the pathogenic bacteria *P. aeruginosa* and *A. fumigatus* (Delhaes *et al.*, 2012; Baxter *et al.*, 2013; Armstead *et al.*, 2014). Equilibrium in the different partners of the microbiota may be important to avoid allergic bronchopulmonary aspergillosis, *A. fumigatus* bronchitis and other allergic manifestations as well as controlling cystic fibrosis infections and chronic obstructive pulmonary disorders (Whiteson *et al.*, 2014).

Conflict of Interest

The authors declare no conflict of interest.

Acknowledgements

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