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## Pseudomonas-Candida interaction in dual-species biofilms

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Bacteria and fungi co-inhabit in a wide variety of environments and the interactions between them can result in huge medical and economic impacts. Pseudomonas aeruginosa, a Gram-negative bacterium, and Candida albicans, a dimorphic fungus. are two important opportunistic pathogens frequently identified as the major causes of nosocomial infections, mainly due to their ability to form virulent biofilms. A pathogenic interaction between P. aeruginosa and C. albicans was recently identified and, it was also found that C. albicans' morphology and virulence are significantly affected in the presence of P. aeruginosa. In the present work, the interaction between P. aeruginosa and C. albicans in dual-species biofilms was studied. Biofilm formation was carried out in 24-well microplates containing 1 ml of Yeast Peptone Dextrose medium and 10 ul of each cellular suspension with an OD<sub>600nm</sub> of 1. Biofilms were formed during 24 and 48 hours with medium renewal every 12 hours. The results revealed that in mixed biofilms C. albicans proliferation was inhibited by the presence of both P. aeruginosa ATCC 10145 and PAO1 strains. The number of C. albicans viable cells was reduced by 2 and 3 logs in 24 and 48 hours old biofilms compared to single C. albicans biofilms, Conversely, P. aeruginosa was not influenced by the presence of C. albicans and so, the amount of viable cells of P. aeruginosa was similar in single and dualspecies biofilms for both P. aeruginosa strains studied. To better understand the cause of C. albicans inhibition, biofilms of C. albicans with P. aeruginosa LPS mutant strains were also studied. These results showed that the LPS chain of P. aeruginosa has a great impact on C. albicans proliferation - mutants with full LPS chain inhibit the greatest while mutants with truncated A and B chains and also truncated outer core allow the growth of Candida. According to this study, the inhibition of C. albicans biofilm formation is directly correlated with the composition of the P. aeruginosa LPS chain.

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