

Automated image analysis of the host-pathogen interaction between phagocytes and *Aspergillus fumigatus*.

Mech F*, Thywissen A*, Guthke R, Brakhage AA, Figge MT (2011) Automated image analysis of the host-pathogen interaction between phagocytes and *Aspergillus fumigatus*. *PLOS ONE* 6(5), e19591, *authors contributed equally.

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Abstract

Aspergillus fumigatus is a ubiquitous airborne fungus and opportunistic human pathogen. In immunocompromised hosts, the fungus can cause life-threatening diseases like invasive pulmonary aspergillosis. Since the incidence of fungal systemic infections drastically increased over the last years, it is a major goal to investigate the pathobiology of *A. fumigatus* and in particular the interactions of *A. fumigatus* conidia with immune cells. Many of these studies include the activity of immune effector cells, in particular of macrophages, when they are confronted with conidia of *A. fumigatus* wild-type and mutant strains. Here, we report the development of an

automated analysis of confocal laser scanning microscopy images from macrophages coincubated with different *A. fumigatus* strains. At present, microscopy images are often analysed manually, including cell counting and determination of interrelations between cells, which is very time consuming and error-prone. Automation of this process overcomes these disadvantages and standardises the analysis, which is a prerequisite for further systems biological studies including mathematical modeling of the infection process. For this purpose, the cells in our experimental setup were differentially stained and monitored by confocal laser scanning microscopy. To perform the image analysis in an automatic fashion, we developed a ruleset that is generally applicable to phagocytosis assays and in the present case was processed by the software Definiens Developer XD. As a result of a complete image analysis we obtained features such as size, shape, number of cells and cell-cell contacts. The analysis reported here, reveals that different mutants of *A. fumigatus* have a major influence on the ability of macrophages to adhere and to phagocytose the respective conidia. In particular, we observe that the phagocytosis ratio and the aggregation behaviour of *pksP* mutant compared to wild-type conidia are both significantly increased.

Beteiligte Forschungseinheiten

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Identifizier

doi: 10.1371/journal.pone.0019591

PMID: 21573171