# Hypoxia-inducible factor 1a modulates metabolic activity and cytokine release in anti-Aspergillus fumigatus immune responses initiated by human dendritic cells.

Fliesser M, Morton CO, Bonin M, Ebel F, Hünniger K, Kurzai O, Einsele H, Löffler J (2015) Hypoxia-inducible factor 1a modulates metabolic activity and cytokine release in anti-Aspergillus fumigatus immune responses initiated by human dendritic cells. *Int J Med Microbiol*,

**Details** 

PublMed

#### **Abstract**

The mold Aspergillus fumigatus causes life-threatening infections in immunocompromised patients. Over the past decade, new findings in research have improved our understanding of A. fumigatus-host interactions, including the recent identification of myeloid-expressed hypoxia-inducible factor  $1\alpha$  (HIF- $1\alpha$ ) as a relevant immune-modulating transcription factor and potential therapeutic target in anti-fungal defense. However, the function of HIF- $1\alpha$  signaling for human anti-A. fumigatus immunity is still poorly understood, including its role in dendritic cells (DCs), which are important regulators of anti-fungal immunity. This study investigated the functional relevance of HIF- $1\alpha$  in the anti-A. fumigatus immune response initiated by human DCs. Hypoxic cell culture conditions were included because hypoxic microenvironments occur during A. fumigatus infections and may influence the host immune response. HIF- $1\alpha$  was stabilized in DCs following stimulation with A. fumigatus under normoxic and hypoxic conditions. This stabilization was partially dependent on dectin-1, the major receptor for A. fumigatus on human DCs. Using siRNA-based

HIF-1 $\alpha$  silencing combined with genome-wide transcriptional analysis, a modulatory effect of HIF-1 $\alpha$  on the anti-fungal immune response of human DCs was identified. Specifically, the difference in the transcriptomes of HIF-1 $\alpha$  silenced and non-silenced DCs indicated that HIF-1 $\alpha$  contributes to DC metabolism and cytokine release in response to A. fumigatus under normoxic as well as hypoxic conditions. This was confirmed by further down-stream analyses that included metabolite analysis and cytokine profiling of a time-course infection experiment. Thereby, this study revealed a so far undescribed functional relevance of HIF-1 $\alpha$  in human DC responses against A. fumigatus.

### **Beteiligte Forschungseinheiten**

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## Identifier

doi: S1438-4221(15)30005-9

**PMID:** 26387061