REFERENCES


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(however, other iron chelators were not studied). Increased HIF-2α messenger RNA (mRNA) and protein were found in neutrophils from patients with chronic obstructive pulmonary disease (COPD) and inflammatory arthritis (IA). This chronic HIF-2α expression may elicit unique biological effects not seen in normoxic human neutrophils from patients with a HIF-2α gain-of-function mutation.

Inflammatory cytokines and Toll-like receptor ligands present in COPD and IA may be inducing HIF-2α expression independently of hypoxia. Consistently, patients with gain-of-function HIF-2α mutations express more HIF-2α target genes at steady state, suggesting that hypoxia is not always required for the induction or function of HIF-2α in neutrophils. HIF-2α can antagonize the expression of HIF-1α in nonhematopoietic cells, but it is not known if similar regulatory mechanisms exist in neutrophils.

Previous studies in macrophages demonstrate that lipopolysaccharide can induce HIF-1α mRNA expression but that hypoxia is a prerequisite for HIF-1α protein accumulation via the inhibition of prolyl hydroxylases. Like HIF-1α, the accumulation of HIF-2α protein at sites of inflammation may therefore be dependent on factors controlling both gene transcription and HIF-2α protein half-life.

The key role of HIF-2α in the regulation of macrophage inflammatory cytokine production introduces a possible caveat to this study because the mice used in these studies lacked HIF-2α in neutrophils and macrophages. This was addressed using a fractionated irradiation regimen (3 fractions of 1 Gy/day for 4 days) to ablate bone marrow–derived myeloid cells but not wild-type resident lung macrophages. Following acute lung injury, HIF-2α–deficient neutrophils were recruited to lungs containing wild-type resident macrophages. This recruitment occurred independently of HIF-2α, but the subsequent resolution of neutrophilic inflammation was accelerated in the absence of HIF-2α. Conversely, expression of gain-of-function HIF-2α mutants in zebra fish delayed the clearance of neutrophils from a tail fin wound (see figure). Moreover, HIF-2α deficiency affects neutrophil life span but not phagocytosis or the respiratory burst, suggesting that the diverse functional roles of HIF-2α in macrophages do not extend to neutrophils.

Inflammatory cytokine production by HIF-2α–deficient macrophages and neutrophils was also not altered in an acute lung injury model, supporting the conclusion that the alterations in neutrophil life span are cell intrinsic.

This study reports that human HIF-2α gain-of-function mutations increase basal expression of prolyl hydroxylase-3 (PHD3), an enzyme that helps neutrophils adapt to hypoxia. PHD3 is upregulated in peripheral blood neutrophils from patients with rheumatoid arthritis. This study demonstrates that HIF-2α is also strongly upregulated in peripheral blood neutrophils and lung biopsies from patients with rheumatoid arthritis and COPD, respectively, suggesting that HIF-2α regulates PHD3 expression in these human cells. PHD3 regulates Bcl-xL, a prosurvival protein whose expression is responsive to hypoxia.

However, in this study, no perturbations were noted in Bcl-xL expression in HIF-2α–deficient neutrophils. More studies are needed to determine if Bcl-xL, or other prosurvival proteins such as Mcl-1, are HIF-2α targets during the adaptation to hypoxia. Neutrophil-specific conditional gene-targeting approaches will be essential to identify host factors that shift the balance between neutrophil life and death.

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Comment on Bouvier et al, page 404, and on Bouvier et al, page 414

Preventing pregnancy loss

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In this issue of Blood, 2 articles by The Nimes Obstetricians and Hematologists–Antiphospholipid Syndrome (NOH-APS) Study Group give us new information about the effects of low-molecular-weight heparin (LMWH) on pregnancy complications in women with prior pregnancy loss and either purely obstetric antiphospholipid syndrome (APS) or inherited thrombophilia. The results better define women at risk, suggest a role for LMWH, and confirm the need for further investigation.

APS has been linked to pregnancy complications including fetal loss at all stages of gestation, preeclampsia, eclampsia, placental insufficiency, and premature birth. The inherited thrombophilias, factor V Leiden (FVL) mutation and prothrombin gene (PTG) mutation G20210A, have variable associations with nonthrombotic pregnancy complications; they are believed to play a minimal role in spontaneous abortion or placenta-mediated complications and only a modest role in second- or third-trimester fetal loss.

The NOH-APS study is a large prospective observational study of 618 women with
Neutrophil survival in the death zone

Ben A. Croker