Mutations in succinate dehydrogenase B (SDHB) enhance neutrophil survival independent of HIF-1α expression

Robert Jones,1,2,* Kate E. McDonald,1,2,* Joseph A. Willson,3 Bart Ghesquière,4,5 David Sammut,1 Eleni Daniel,2 Alison J. Harris,2 Amy Lewis,1 A. A. Roger Thompson,1 Rebecca S. Dickinson,3 Tracie Plant,3 Fiona Murphy,3 Pranvera Sadiku,3 Brian G. Keevil,6 Peter Carmeliet,4,5 Moira K. B. Whyte,3 John Newell-Price,2,† and Sarah R. Walmsley3,†

1Department of Infection, Immunity, and Cardiovascular Disease and 2Department of Oncology and Metabolism, University of Sheffield, Sheffield, United Kingdom; 3Medical Research Council/University of Edinburgh Centre for Inflammation Research, Queen’s Medical Research Institute, University of Edinburgh, Edinburgh, United Kingdom; 4Laboratory of Angiogenesis and Vascular Metabolism, Vascular Research Center, VIB, Leuven, Belgium; 5Department of Oncology, KU Leuven, Leuven, Belgium; and 6School of Medicine, University of Manchester, Manchester, United Kingdom

Neutrophils are unusual in their reliance on glycolysis to maintain their energy requirements despite the presence of mitochondria and tricarboxylic acid (TCA) cycle intermediaries. This metabolic adaptation is thought in part to underpin their survival and anti-microbial function in tissues that are typically hypoxic. Despite their unique metabolism, little is known about the importance of flux between metabolic pathways in determining neutrophil survival responses. Recent work has demonstrated the importance of the hypoxia-inducible factor (HIF)/prolyl hydroxylase domain (PHD) containing enzyme oxygen-sensing pathway in this regard, identifying both HIF-1α and PHD3 as critical regulators of neutrophil survival in hypoxia.[MD1] with the extended survival of neutrophils in hypoxia being dependent on HIF-1α expression. In parallel, an expanding body of work has addressed the role of HIF-1α in coordinating macrophage functional responses to proinflammatory mediators.[MD2] This work led to the observation that, in macrophages, lipopolysaccharide (LPS) causes an intracellular increase in succinate levels, resulting in HIF-1α stabilization and enhanced interleukin-1β signaling.[MD3] Subsequently, the metabolic rewiring of antimicrobial (M1) and tissue repair (M2) macrophages has been elucidated, with important consequences of TCA cycle activity and integrity for regulation of nitric oxide and N-glycosylation signaling, respectively.[MD4] Whether the TCA cycle activity and succinate accumulation regulates HIF-1α and hypoxic survival in neutrophils is unknown.

Patients with rare germ line mutations in genes encoding the TCA cycle enzyme succinate dehydrogenase (SDH) allow us to directly question the role of the TCA cycle and mitochondrial respiratory chain in neutrophil survival responses. SDH oxidizes succinate to fumarate in the TCA cycle and is a ubiquinone oxidoreductase, also functioning in complex II of the respiratory chain. SDH comprises four subunits (A-D), with inherited mutations of each of the subunits linked to the development of pheochromocytoma (PHEO) and paraganglioma (PGL) after somatic inactivation of the wild-type allele and loss of heterozygosity.[MD5] We questioned whether heterozygous germ line mutations in SDHB (SDHBx) would reduce SDH activity in the peripheral blood neutrophils of these patients, leading to accumulation of intracellular succinate, HIF-1α stabilization, and a pseudohypoxic survival phenotype, given the importance of the B subunit for SDH catalytic function and its high prevalence within PHEO/PGL patient populations.[MD6] To determine whether succinate is implicated in regulating neutrophil survival responses, we isolated peripheral blood neutrophils from patients with heterozygous germ line SDHBx mutations in whom an increase in intracellular succinate would be predicted. In total, 20 individuals with frameshift, splice, missense, or nonsense mutations were studied (supplemental Table 1, available on the Blood Web site). Although all but 1 patient displayed plasma succinate levels within the normal range, a significantly higher plasma succinate level was observed in patients with SDHBx (Figure 1A). To confirm the consequence of SDHBx mutations on intracellular succinate and to measure other TCA cycle and glycolytic intermediaries, peripheral blood neutrophils were isolated from 3 individuals with SDHBx and 3 healthy controls, and relative metabolite abundance was determined by gas chromatography–mass spectrometry (Figure 1B). Succinate was significantly more abundant in neutrophils isolated from patients with SDHBx than from controls. This finding was paralleled by increases in lactic acid and citric acid, but no changes in other TCA cycle intermediaries (α-ketoglutaric acid, fumaric acid, or malic acid) were observed. Thus, neutrophils heterozygous for mutant SDHBx gene expression display the predicted elevation in intracellular succinate, but with no decrease in downstream TCA cycle intermediaries. Citric acid levels were increased, which may reflect an increase in biosynthetic requirements outside the TCA cycle. In keeping with the increased succinate in SDHBx neutrophils, a detectable increase in protein succinylation was also observed (Figure 1C).
The consequence of SDHB heterozygosity for constitutive rates of neutrophil apoptosis and hypoxic survival responses was determined. SDHBx neutrophils displayed both reduced constitutive apoptosis and enhanced survival in hypoxia, as assessed both by cellular morphology (Figure 1D) and Annexin-V positivity (Figure 1E). Given the previous report of succinate-mediated HIF-1α stabilization in bone marrow–derived macrophages,11 and the importance of HIF-1α for hypoxic neutrophil survival,6 we asked whether the reduced apoptosis in SDHBx neutrophils was secondary to increased HIF-1α activity. HIF-1α protein in SDHBx neutrophils was elevated in hypoxia (Figure 2A-B) but undetectable in normoxic cells, in which reduced rates of apoptosis were observed. Thus, the phenotype of enhanced neutrophil survival in the setting of SDHB heterozygosity occurs independently of HIF-1α protein expression. In keeping with unaltered HIF-1α activity in normoxic SDHBx neutrophils, we saw no alterations either in glucose uptake (Figure 2C-D), in which a hypoxic uplift is observed in healthy control cells, or in extracellular acidification rates, an indirect measure of glycolytic activity (Figure 2E). Interestingly, we observed that neutrophils isolated from individuals with SDHB mutations displayed significantly reduced levels of oxidant stress (Figure 2F), a phenotype associated with enhanced neutrophil survival in chronic granulomatous disease patients.19 However, no differences in nicotinamide adenine dinucleotide phosphate (NADP)/reduced NADP (NADPH) redox ratios were detected between patients and controls, suggesting the SDHB phenotype to be independent of altered NADPH oxidase 2 activity (Figure 2G). This finding led us to question whether SDH was regulating apoptosis through its role as a mitochondrial ubiquinone oxidoreductase. SDHBx neutrophils demonstrated an increased ratio of oxidized to reduced nicotinamide adenine dinucleotide (NAD) (Figure 2H), and treatment of healthy human neutrophils with the irreversible SDH inhibitor 3-nitropropionic acid reduced constitutive neutrophil apoptosis (Figure 2I), thus implicating impaired mitochondrial complex II and compensatory changes in the electron transport chain in the enhanced survival of SDHB-mutant neutrophils.
These studies use a valuable patient group with a specific mutation in SDHB as an experimental system in which to delineate the role of the TCA cycle and mitochondrial respiratory chain in neutrophil survival responses. It provides the first description of elevated intracellular succinate levels in neutrophils isolated from patients with heterozygous mutations in SDHB and the first evidence of a dysfunctional TCA cycle in resting-state peripheral blood neutrophils. In marked contrast to the role of succinate in facilitating HIF-1α-dependent inflammatory responses in LPS-stimulated macrophages, we dissociate enhanced neutrophil survival from HIF-1α.
stabilization in the context of germ line mutations in SDH, linking it instead to a phenotype of impaired mitochondrial complex II function and reduced oxidative stress. Taken together, this work identifies key metabolic differences between neutrophils and macrophages and raises further important questions as to the metabolic control of neutrophil function and survival. Future work dissecting the consequences of SDHB mutations for neutrophil host-pathogen responses and inflammation resolution will be key in this regard.

*R.J. and K.E.M. contributed equally to this study.
†J.N.-P. and S.R.W. contributed equally to this study.

The online version of this article contains a data supplement.

Acknowledgments: This work was principally supported by Wellcome Trust Senior Clinical Fellowship Award 098518 (S.R.W.), Medical Research Council (MRC) Clinical Training Fellowship Awards G0802255 (A.A.R.T.) and MR/K023845/1 (R.S.D.), and an Engineering and Physical Sciences Research Council and MRC Centre for Doctoral Training in Optical Medical Imaging PhD studentship (J.A.W.). The MRC/University of Edinburgh Centre for Inflammation Research is supported by an MRC Centre grant. The study was also supported by Methusalem funding from the Flemish Government (P.C.).


Conflict-of-interest disclosure: The authors declare no competing financial interests.

Correspondence: Sarah R. Walmsley, MRC/University of Edinburgh Centre for Inflammation Research, The Queen’s Medical Research Institute, University of Edinburgh, Edinburgh, United Kingdom; e-mail: sarah.walmsley@ed.ac.uk.

References


DOI 10.1182/blood-2016-02-696922

© 2016 by The American Society of Hematology
Mutations in succinate dehydrogenase B (SDHB) enhance neutrophil survival independent of HIF-1α expression

Robert Jones, Kate E. McDonald, Joseph A. Willson, Bart Ghesquière, David Sammut, Eleni Daniel, Alison J. Harris, Amy Lewis, A. A. Roger Thompson, Rebecca S. Dickinson, Tracie Plant, Fiona Murphy, Pranvera Sadiku, Brian G. Keevil, Peter Carmeliet, Moira K. B. Whyte, John Newell-Price and Sarah R. Walmsley