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Research Topic Editorial

Immunosuppressive Amino Acid Catabolizing Enzymes in Health and Disease

Flavia Castellano^{1,2}, Jorge Correale³ and Valérie Molinier-Frenkel^{1,2}

¹Virus-Immunity-Cancer Department, Institut Mondor de Recherche Biomédicale (IMRB), INSERM U955, Université Paris-Est Créteil, F-94010 Créteil, France

² Virus-Immunity-Cancer Department, Institut Mondor de Recherche Biomédicale (IMRB), INSERM U955, Université Paris-Est Créteil, F-94010 Créteil, France

³Department of Neurology, Fleni, Buenos Aires, Argentina

Defective calibration of the immune response is involved in the pathophysiology of a great number of affections beyond chronic infections and autoimmunity. Membrane-expressed immune checkpoint molecules and cytokines are well-known mediators of immune regulation. Control of the level of some essential amino acids and production of bioactive amino acid-derived metabolites represents a less characterized pathway of immune regulation with potential for pharmacological manipulation (1).

Cells sense amino acid levels through at least two different pathways, involving the general control non-derepressible 2 (GCN2) kinase, and the mammalian target of rapamycin (mTOR) kinase, respectively (2). These two related serine/threonine kinases jointly assess nutritional deficiency (GCN2) and adequacy (mTOR). Increased amino acid catabolism leads to amino acid depletion, GCN2 activation and mTOR inactivation, thereby limiting activation of some immune cell populations, in particular T lymphocytes. Hence, basal amino acid catabolism can contribute to immune homeostasis, whereas elevated amino acid catalytic activity reinforces immune suppression. Furthermore, several downstream amino acid metabolites are also important biological mediators of immune response regulation.

Six major amino acid catabolizing enzymes endowed with immunosuppressive properties are known, which catabolize tryptophan (indole 2,3 dioxygenase [IDO] 1, IDO2, and tryptophan dehydrogenase [TDO] 2), arginine (arginase 1 and inducible nitric oxid synthase [iNOS]) and phenylalanine (Interleukin four-induced gene 1 [IL4I1]) (3). These enzymes are genetically unrelated and their mechanisms of action are various and context-dependent. In the immune system, they are mainly produced by subpopulations of myeloid cells stimulated by specific signals, with certain species-related differences that complicate their study. This Research Topic gathers different contributions

highlighting their activities in several pathological conditions, spanning from infection to autoimmunity, cancer and atherosclerosis.

Three contributions address the role of tryptophan-degrading enzymes. The first article by Rafaella Oliveira dos Santos et al links tryptophan metabolism to cytokine production and parasitemia, in patients with malaria caused by *Plasmodium vivax*. In particular, the first malaria episode, which is characterized by a high parasite load, was associated with a stronger IFN γ response and higher levels of kynurenine (a product of the degradation of tryptophan by IDO or TDO) than subsequent episodes occurring after the development of specific IgG. This suggests induction of a regulatory circuit by IFN γ -induced IDO1 and/or TDO, which controls Th1 inflammation to avoid host immunopathology during the period preceding acquisition of protective humoral immunity.

In the second contribution, Lauren Merlo et al, address the complex issue of redundancy and complementarity of the two highly related enzymes, IDO1 and IDO2, by comparing T and B cell-mediated immune responses in double knock-out (KO) and single KOs mouse models in several contexts. Their data importantly show that single deletion of one of these genes leads to significant modifications of the expression of the other. This may affect interpretation of previous results of the literature, as IDO2, which has much weaker tryptophan degrading activity than IDO1, has been shown to play a proinflammatory role. Moreover, these models indicate that B cell responses directly depend on IDO2-induced inflammation, while IDO1 mediates T cell suppression in IDO2 KO mice, an effect that may be related to its enhanced expression in this model.

The last contribution on Trp catabolizing enzymes from Soumya Mohapatra et al concentrates on the activity of TDO2 in the liver, which controls Trp plasmatic levels. They show that hepatocytes reroute Trp metabolism under hypoxic conditions by inhibiting TDO2 expression, which decreases kynurenin production, while tryptamine production by dopa decarboxylase is enhanced. However, AHR activation is maintained under hypoxia, as tryptamine is still capable to activate this important immunomodulatory pathway.

Two research papers reveal some unsuspected particularities of Arg1 and IL4I1. The work of Verena Vonwirth et al focuses on the activation and proliferation of T cells stimulated in supernatants of polymorphonuclear (PMN) cell cultures. Due to arginine depletion by arginase 1, T cell proliferation is abrogated in PMN supernatants. Surprisingly, specific inhibition of arginase 1 in PMN cultures not only releases T cell inhibition but leads to hyperactivation of the T lymphocytes, characterized by increased proliferation, cytotoxicity, and IL-9 and IL-17 production. The authors demonstrate that this hyperactivated state is induced by one or several low molecular weight (<3 kDA), high

temperature-resistant PMN-derived small molecule(s), that are rapidly released after arginase 1 inhibition, although the exact nature of this (these) factor(s) is not identified.

The article from Marie-Line Puiffe et al uses an IL4I1 KO mouse model to dissect the induction of the CD8 T cell response during an acute viral infection. Unexpectedly, the absence of the IL4I1 enzyme is associated with diminished expansion of functional short lived-effector CD8 T cells, but enhanced memory T cell differentiation. These observations are not related to intrinsic differences of CD8 T cells between WT and IL4I1 KO mice, but result from modulation of immune synapse formation and early activation events by IL4I1-expressing DCs. Indeed, IL4I1 enhances the T cell activation threshold, thereby favoring the priming of high-affinity clones, restriction of the response to the most immunodominant peptides and rapid acquisition of effector differentiation.

An extensive review from Bozidarka Zaric et al on the role of amino acids and amino-acid catabolizing enzymes points to their importance in the development of atherosclerosis. The review recapitulates available data indicating that IDO1 and arginase regulate inflammation in this context and should represent promising therapeutic targets in cardiovascular diseases.

Finally, although not included in this Research Topic, another nice review from J Correale, appeared in *Frontier Immunology* in January 2021, describes our knowledge on the role of these enzymes in autoimmunity through the example of multiple sclerosis, a disease that is considered to be mediated by autoreactive Th1, Th17 and B cells.

In conclusion, the six papers of this topic underline the complexity this family of enzymes and increase our knowledge in this complex field. We are grateful to all the authors and reviewers for their precious contributions to this research topic.

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