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Cutting Edge: IL-6–Driven Immune Dysregulation Is Strictly Dependent on IL-6R α-Chain Expression

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IL-6 binds to the IL-6R α-chain (IL-6Rα) and signals via the signal transducer gp130. Recently, IL-6 was found to also bind to the cell surface glycoprotein CD5, which would then engage gp130 in the absence of IL-6Rα. However, the biological relevance of this alternative pathway is under debate. In this study, we developed a mouse model, in which murine IL-6 is overexpressed in a CD11c-Cre–dependent manner. Transgenic mice developed a lethal immune dysregulation syndrome with increased numbers of Ly-6G+ neutrophils and Ly-6C hi monocytes/macrophages. IL-6 overexpression promoted activation of CD4+ T cells while suppressing CD5+ B-1a cell development. However, additional ablation of IL-6Rα protected IL-6–overexpressing mice from IL-6–triggered inflammation and fully phenocopied IL-6Rα–deficient mice without IL-6 overexpression. Mechanistically, IL-6Rα deficiency completely prevented downstream activation of STAT3 in response to IL-6. Altogether, our data clarify that IL-6Rα is the only biologically relevant receptor for IL-6 in mice.

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E levated levels of IL-6 have been observed in numerous pathological conditions, and several drugs are successfully used in a series of human diseases, including rheumatoid arthritis, Castleman disease, and giant cell arteritis to target IL-6 and its receptor, IL-6R α-chain (IL-6Rα) (1). For other diseases like multiple myeloma and neuromyelitis optica, the clinical program for the development of IL-6–neutralizing agents is very advanced, and drugs targeting IL-6 or IL-6Rα might be licensed for these diseases in the near future (2).

IL-6, together with IL-1, IL-27, CNTF, LIF, OSM, CT-1, and CLC, belongs to the group of cytokines that use gp130 for signal transduction (3). Since its discovery in 1990, it was believed that the IL-6–signaling complex consists of a unique IL-6–binding receptor IL-6Rα (also known as CD126) and the signal transducer gp130 (4). The assembly of IL-6, IL-6Rα, and gp130 leads to activation of STAT3-mediated intracellular signaling pathways, which control cell survival, activation, and proliferation (2). IL-6Rα also exists in a soluble form, which is generated via secretion or shedding of membrane-bound IL-6Rα. The complex of IL-6 and soluble IL-6Rα is able to bind to gp130 and then induce STAT3 phosphorylation in cells that do not express IL-6Rα themselves. This signaling modality of IL-6 was termed IL-6 trans-signaling (5). A third modality of IL-6 signaling, called IL-6 cluster signaling, is mediated by cell-bound presentation of the IL-6/IL-6Rα complex in trans by a donor cell to a receiving cell that expresses gp130 (2, 6). In both IL-6 trans-signaling and IL-6 cluster signaling, cells that express gp130, but lack IL-6Rα, still respond to IL-6.

In 2016, an alternative pathway of IL-6 signal transduction was proposed in which IL-6 binds to the membrane-anchored

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Abbreviations used in this article: DC, dendritic cell; IL-6Rα, IL-6R α-chain; OE, overexpression; VD, viability dye-negative.

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glycoprotein CD5 instead of IL-6Rα and, via gp130, initiates STAT3 phosphorylation in B cells (7). This study suggested the CD5-dependent pathway of IL-6 signaling was critical in the promotion of cancer progression (7). However, the mechanism for this novel IL-6 binding was not elucidated. In fact, soluble CD5, despite binding to IL-6, is unable to induce IL-6 trans-signaling (8). Massive overproduction of IL-6 is observed in sepsis (9) and has been reported during the cytokine release syndrome in response to CAR-T cell immune therapies (10). To keep exaggerated IL-6 responses in check, any design of therapeutic intervention needs to consider potential alternative signaling pathways of IL-6. In particular, the idea of CD5 as a molecule to substitute for IL-6Rα is indispensable for IL-6 signaling.

**Materials and Methods**

**Mice**

Mice with conditional Il6ra allele (Il6ra flox) (11) and CD11c-Cre mice (12) have been previously described. IL-6Rα full-knockout (Il6ra / −) mice were selected from the breeding between Il6ra flox mice and CD11c-Cre mice, which displayed occasional spontaneous germline Cre activity. Principles used to generate mice with the knock-in of a transgene into Rosa26 locus will be reported elsewhere. For breeding strategies involving Rosa26 cDNA in the CAG–(loxP)STOP(loxP)–IL-6–IRES–eGFP transgene inserted to generate mice with the knock-in of a transgene into which displayed occasional spontaneous germline Cre activity. Principles used to generate mice with the knock-in of a transgene into Rosa26 locus have been described (13). Details of the generation of mice carrying murine cDNA in the CAG–(loxP)STOP(loxP)–IL-6–IRES–eGFP transgene inserted in the Rosa26 locus will be reported elsewhere. For breeding strategies involving IL-6 overexpression and mouse group abbreviations, refer to Supplemental Fig. 1A. All mice were on C57BL/6 background and were bred in-house at specific pathogen-free conditions. For experiments, 5–9-wk-old, gender-matched mice were used in accordance with the guidelines of the central animal facility institution (Translational Animal Research Center, University of Mainz).

**Mouse sample collection**

Single-cell suspensions from spleen and lymph nodes were prepared by mechanical dissociation in PBS supplemented with 2% FCS. Peritoneal lavage was performed on lethally anesthetized mice with 5 ml of PBS supplemented with 3% FCS. Peripheral blood was collected from the tail vein. RBCs were removed using ACK lysis buffer.

**IL-6 ELISA**

A murine IL-6 ELISA Kit (BD Biosciences) was used to analyze IL-6 levels in the blood serum. Plates were measured with the Infinite M200 PRO NanoQuant reader (Tecan).

**Flow cytometry**

Single-cell suspensions were stained with Abs together with viability dyes (Supplemental Table I). Stained cells were acquired on FACS Canto II (BD Biosciences), and data were analyzed with FlowJo software. Gating strategy always considered cell size, excluded duplets, and defined living cells as viability dye-negative (VIT+) population.

**p-STAT3 detection by flow cytometry**

Peritoneal lavage preparations pooled from four to five mice per genotype were FACS sorted for CD4+ F4/80+/CD19−CD5+ cells and for CD4+ F4/80−/CD19−CD5− cells. Sorted cells from Il6ra / − mice (CD4+CD5+ cells). Sorted cells from Il6ra / − mice (CD4+CD5− cells). Sorted cells from Il6ra / − mice (CD4+CD5+ cells). Sorted cells from Il6ra / − mice (CD4+CD5− cells). Sorted cells from Il6ra / − mice (CD4+CD5+ cells). Sorted cells from Il6ra / − mice (CD4+CD5− cells). Sorted cells from Il6ra / − mice (CD4+CD5+ cells). Sorted cells from Il6ra / − mice (CD4+CD5− cells).

**Results and Discussion**

The cytokine IL-6 uses various signaling modalities (i.e., classic signaling, trans-signaling, and cluster signaling [also called trans-presentation]) (2) and might even signal through an alternative receptor complex composed of CD5 and gp130 (7). To assess the signaling modality of IL-6 under conditions of exaggerated IL-6 production, we created a murine system of sterile IL-6 overexpression in vivo. To this end, we generated mice carrying a CAG–(loxP)STOP(loxP)–IL-6–IRES–eGFP construct in the Rosa26 locus. Next, we used dendritic cell (DC)–directed CD11c-Cre transgenic mice (12) to activate IL-6 overexpression (OE) from a single copy of our Il6 transgene in mice on an IL-6Rα–sufficient background (IL-6DC-OE Il6ra / + and IL-6DC-OE Il6ra / − [i.e., IL-6DC-OE −/−] and on an IL-6Rα–deficient background (IL-6DC-OE Il6ra / −). As further controls, we used IL-6Rα–sufficient (control) and IL-6Rα–deficient (Il6ra / −) mice without IL-6 overexpression (Supplemental Fig. 1A).

To directly assess IL-6 overexpression, we measured IL-6 levels in the serum of 5–6-wk-old mutant mice. As expected, both groups of mice with IL-6 overexpression showed elevated levels of IL-6 in comparison with wild-type controls and Il6ra / − mice (Fig. 1A). IL-6 levels were significantly higher in IL-6DC-OE −/− mice than in IL-6DC-OE +/− mice, suggesting a positive feed-forward loop for IL-6 expression in mice sufficient for IL-6Rα. Mechanistically, transgenic IL-6 can promote IL-6 expression in IL-6DC-OE −/− mice, from the endogenous Il6 locus, which remained responsive to its natural regulation. In addition, IL-6 forms a complex with the soluble IL-6Rα and gp130 in the blood, which can prolong its half-life only in IL-6Rα–sufficient, but not in IL-6Rα–deficient, mice (14, 15).

To analyze cellular sources of transgenic IL-6, we assessed eGFP reporter expression driven by the CAG promoter after CD11c-Cre mediated excision of the loxP flanked STOP cassette. As expected, we found eGFP expression in DCs in IL-6DC-OE −/− and IL-6DC-OE Il6ra / − mice (Fig. 1B, Supplemental...
Fig. 1C). Because of leakiness of the CD11c-Cre recombinase expression (16), we also noted limited reporter expression in other myeloid cells and in T and B cells of these mice (Supplemental Fig. 1C).

High systemic levels of IL-6 ultimately led to the death of IL-6\(^{DC-OE}\) mice in an IL-6\(^{-}\)–dependent manner (Fig. 1C). IL-6–overexpressing mice carrying both copies of Il6ra died within 6 wk of age, whereas deletion of one Il6ra allele (with one Il6ra allele intact in heterozygous Il6ra\(^{-/-}\) mice) extended the survival of IL-6\(^{DC-OE}\) mice until 11 wk of age. Strikingly, null deficiency in Il6ra restored the survival of IL-6\(^{DC-OE}\) Il6ra\(^{-/-}\) mice with IL-6 overexpression and, thus, fully rescued the phenotype of IL-6\(^{DC-OE}\) Il6ra\(^{-/-}\) mice. Of note, 3 out of 41 Il6ra\(^{-/-}\) mice died during the observation period presumably because of their immunodeficiency (17), irrespective of IL-6 overexpression.

To study the impact of IL-6\(^{Ra}\) expression on IL-6 signaling, we isolated splenocytes from wild-type (Il6ra\(^{+/+}\)), Il6ra\(^{+/−}\), and Il6ra\(^{−−}\) mice. We observed a robust IL-6\(^{Ra}\) expression on CD4\(^{+}\) T cells and, to a lesser extent, on CD11b\(^{+}\) myeloid cells, but very limited expression on CD19\(^{+}\) cells in wild-type mice (Fig. 1D). Deletion of a single Il6ra allele led to reduction in IL-6\(^{Ra}\) expression, suggesting a haploinsufficient gene dose effect (Fig. 1D), which, together with reduced frequencies of IL-6\(^{Ra}\)–positive cells in Il6ra\(^{−−}\) mice (Supplemental Fig. 2A), may contribute to the reduced mortality of IL-6\(^{DC-OE}\) Il6ra\(^{-−}\) mice in comparison with IL-6\(^{DC-OE}\) Il6ra\(^{−−}\) mice (Fig. 1C). To determine the consequence of the lack of IL-6\(^{Ra}\) in downstream signaling events, we stimulated total splenocytes isolated from Il6ra\(^{−−}\), Il6ra\(^{+/−}\), and Il6ra\(^{−−}\) mice with IL-6 and assessed their STAT3 phosphorylation. We observed STAT3 phosphorylation only in wild-type and Il6ra heterozygous cells, whereas Il6ra\(^{−−}\) splenocytes were irresponsive to stimulation with soluble IL-6 (Fig. 1E, Supplemental Fig. 2B). FACS-sorted CD5\(^{-}\) and CD5\(^{+}\) B cells, as well as CD5\(^{+}\) T cells, isolated from Il6ra\(^{−−}\) mice did not respond to soluble IL-6 either, whereas their wild-type counterparts showed a clear STAT3 activation in response to soluble IL-6 (Supplemental Fig. 2C). In contrast, IL-6\(^{Ra}\)–deficient B cells (regardless of CD5 expression), and IL-6\(^{Ra}\)–deficient T cells responded to IL-6 in the presence of exogenous IL-6\(^{Ra}\) in the form of hyper–IL-6, which represents IL-6/IL-6R\(^{Ra}\) fusion protein (Supplemental Fig. 2C). These data indicate that Il6ra\(^{-−}\) cells retain the ability to respond to IL-6 trans-signaling, whereas IL-6\(^{Ra}\) expression on the cell surface is nonredundant for classic IL-6 signaling. Together, this refutes the concept that alternative receptor molecules can compensate for the lack of IL-6\(^{Ra}\) expression.

As a consequence of IL-6 overexpression, CD11b\(^{+}\) myeloid cells massively infiltrated secondary lymphoid organs, resulting in splenomegaly of IL-6\(^{Ra}\)–sufficient IL-6\(^{DC-OE}\) mice (Supplemental Fig. 3). The most dramatic increase in response to transgenic IL-6 was observed in Ly-6G\(^{+}\) neutrophils and Ly-6C\(^{hi}\) monocyes/macrophages (Fig. 2A). Notably, the systemic increase in these myeloid cell subsets was completely abrogated in IL-6\(^{DC-OE}\) Il6ra\(^{-−}\) mice lacking IL-6\(^{Ra}\). It is likely that the systemic inflammatory response in IL-6\(^{DC-OE}\) Il6ra\(^{-−}\) mice is dependent on the massive expansion of myeloid cells.

Initially, IL-6 was named “B cell hybridoma growth factor,” to acknowledge its stimulatory function on B cells (18). Although we did not detect robust expression of IL-6\(^{Ra}\) on the surface of B cells (Fig. 1D), IL-6 can pair with soluble IL-6\(^{Ra}\) and act on B cells expressing gp130. Importantly, a recent study suggested that IL-6 can bind to CD5 on B cells and activate STAT3 independently of IL-6\(^{Ra}\) (19). Of note, mice with IL-6 overexpression on an Il6ra\(^{-−}\)–deficient background exhibited a dramatic reduction in B-1a cells in the spleen (Fig. 2B), whereas IL-6\(^{Ra}\) deficiency completely prevented the IL-6–triggered loss of B-1a cells in IL-6\(^{DC-OE}\) Il6ra\(^{-−}\) mice (Fig. 2B).
Because CD4^+ T cells expressed high levels of IL-6Rα (Fig. 1D), we analyzed the T cell compartment in mice with activated IL-6 pathway. We noted a significant shift toward CD44^+CD62L^2 effector CD4^+ T cells in the spleen of IL-6DC-OE mice in comparison with the other groups tested (Fig. 2C). However, the frequencies of effector CD4^+ T cells in response to exaggerated IL-6 expression became normal again after IL-6Rα deletion. In accordance with a previous report (20), we found abundant CD5 expression on CD4^+ T cells, however, irrespective of their IL-6Rα status (Fig. 2D).

To test STAT3 activation in specific lymphocyte subsets in mice with IL-6 overexpression, we performed immunoblotting with lysates from CD19^+ B cells and CD4^+ T cells. Both cell types isolated from IL-6 DC-OE mice displayed STAT3 phosphorylation already in steady state without additional in vitro IL-6 stimulation (Fig. 2E). Importantly, the steady-state STAT3 phosphorylation (and also STAT3 phosphorylation in response to exogenous IL-6 stimulation in vitro) was absolutely dependent on the presence of at least one copy of Il6ra in CD19^+ B cells and in CD4^+ T cells (Fig. 2E).

In the current study, we observed a lethal immunopathology in mice that overexpressed IL-6 as long as they were kept on an IL-6Rα–sufficient background. Among several genetic models with IL-6 overexpression (reviewed in Ref. 21), we activated murine IL-6 overexpression driven by CAG promoter in a CD11c-Cre–dependent manner. By genetic ablation of Il6ra, the phenotype of profound immune dysregulation in mice with transgenic IL-6 was completely rescued. This argues against the biological significance of any IL-6 signaling systems that would be independent of IL-6Rα. In our model, we cannot draw a definite conclusion as to which IL-6 signaling modality (i.e., classic IL-6 signaling, IL-6 trans-signaling, or IL-6 cluster signaling [trans-presentation]), is the most relevant IL-6–signaling modality for the fatal immune dysregulation in IL-6DC-OE mice. All three signaling modalities depend on IL-6Rα, either on the side of the receiving cell (classic IL-6 signaling) or on the side of the donating cell in a soluble manner (IL-6 trans-signaling) or in a cell-bound manner (IL-6 trans-presentation) (2, 6). In steady state, CD4^+ T cells and CD11b^+ myeloid cells express relatively high levels of the membrane-bound IL-6Rα. Thus, we speculate that, at least initially, IL-6 overexpression is operational in lymphocytes and myeloid cells through classic IL-6 signaling. The massive expansion of granulocytes in IL-6DC-OE mice was a robust
observation. However, it remains to be determined whether the expansion of Ly-6G+ cells in IL-6-DC-OE mice is a direct effect of IL-6. Recently, concerns have been raised as to whether circulating granulocytes can directly respond to IL-6 (22).

Taken together, although we did not aim to dissect direct and indirect effects of IL-6 overexpression in developing devastating inflammation, our data compellingly support the absolute requirement for IL-6Rα in mediating IL-6 effects in mice. The fact that our study did not confirm the IL-6–CD5-signaling module proposed for B cells (7) challenges the biological significance of IL-6 signals transduced by an IL-6/CD5/gp130 complex. Our conclusion is in accordance with the recent finding that soluble CD5, despite binding of IL-6, is unable to induce STAT3-mediated signal transduction in gp130-expressing cells (8).

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Disclosures
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