Extramedullary Expansion of Hematopoietic Progenitor Cells in Interleukin (IL)-6-sIL-6R Double Transgenic Mice

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Summary

Soluble cytokine receptors modulate the activity of their cognate ligands. Interleukin (IL)-6 in association with the soluble IL-6 receptor (sIL-6R) can activate cells expressing the gp130 signal transducer lacking the specific IL-6R. To investigate the function of the IL-6–sIL-6R complex in vivo and to discriminate the function of the IL-6–sIL-6R complex from the function of IL-6 alone, we have established a transgenic mouse model. Double-transgenic mice coexpressing IL-6 and sIL-6R were generated and compared with IL-6 and sIL-6R single-transgenic mice. The main phenotype found in IL-6–sIL-6R mice was a dramatic increase of extramedullary hematopoietic progenitor cells in liver and spleen but not in the bone marrow. In IL-6 single-transgenic mice and sIL-6R single-transgenic mice no such effects were observed. The high numbers of hematopoietic progenitor cells were reflected by a strong increase of peripheral blood cell numbers. Therefore, activators of the gp130 signal transducer like the IL-6–IL-6R complex may represent most powerful stimulators for extramedullary hematopoietic progenitor cells. gp130 activators may become important for the expansion of hematopoietic progenitor cells in vivo and in vitro.

IL-6 is a mediator of hematopoietic cell growth and differentiation acting on B cells, T cells, keratinocytes, neuronal cells, osteoclasts, and endothelial cells (for reviews, see references 1 and 2). In the liver, IL-6 modulates the transcription of acute phase response genes during acute and chronic inflammatory states (3).

The IL-6-type cytokine family comprises IL-6, IL-11, ciliary neurotrophic factor (CNTF),1 leukemia inhibitory factor (LIF), oncostatin M (OSM), and cardiotrophin-1 (CT-1). The biological action of these cytokines is mediated by multisubunit cell surface receptors that share a common signaling subunit, the gp130 signal-transducing molecule (4, 5). The subunits of these receptors are members of the cytokine/growth hormone receptor family that have conserved cysteine and tryptophane residues in the extracellular domain and that signal via the JAK–STAT pathway (6–8). IL-6 and IL-11 use a gp130 homodimer and OSM, CNTF, LIF, and CT-1 use a heterodimer of gp130 and the 190 kD LIF receptor for the activation of intracellular signaling cascades (4). The receptor signaling complexes for CNTF, IL-6, IL-11, and probably CT-1 (9) contain an additional ligand specific subunit that is not required for LIF and OSM binding. Recently, a separate OSM-specific receptor has been identified that also forms a heterodimer with gp130 (10).

Many soluble receptors (sR), like TNF-R, IL-1R, and IL-4R, act antagonistically, competing with membrane receptors for the binding of the cytokines (11–13). However, some agonistic cytokine–sR complexes stimulate signal-transducing receptors, rendering cells sensitive to the respective cytokines that lack specific cytokine receptors on their surface. Such cells on their own would not be able to respond to the respective cytokine alone. This pathway, recently termed transsignaling, is used by the cytokines of the IL-6 family (14, 15).

On target cells, IL-6 first binds to a specific IL-6 receptor (IL-6Rα) (16). This IL-6–IL-6R complex induces the homodimerization of two gp130 molecules (17, 18), leading to intracellular signaling events. Soluble forms of the IL-6R complex are generated by limited proteolysis from the cell surface (19, 20). Elevated concentrations can be found in inflammatory diseases (21–24). In vivo, the sIL-6R acts as a serum-binding protein for IL-6 and prolongs the plasma

1Abbreviations used in this paper: CNTF, ciliary neurotrophic factor; CT-1, cardiotrophin-1; LIF, leukemia inhibitory factor; OSM, oncostatin M; PEPCK, phosphoenolpyruvate carboxykinase; sIL-6R, soluble interleukin-6 receptor; TNF-R, tumor necrosis factor receptor.
Materials and Methods

Generation of Transgenic Mice. The generation of human sIL-6R and human IL-6 transgenic mice has been described in detail elsewhere (25, 26). By crossing homozygous sIL-6R and IL-6 mice, hemizygous double transgenic mice (IL-6–sIL-6R mice) were generated that coexpress both transgenes. Expression of the sIL-6R transgene was driven by the neonatal phosphoenolpyruvate carboxykinase (PEPCK) promoter, which is transcriptionally activated at day 3 after birth and does not produce intrauterine developmental effects (25). The IL-6 transgene was driven by the murine metallothionein-1 promoter (26). Serum concentration of IL-6 ranged between 10 and 20 ng/ml in IL-6 and IL-6–sIL-6R mice; concentrations of sIL-6R ranged between 4 and 8 μg/ml in sIL-6R and IL-6–sIL-6R mice (data not shown).

Antibodies. The following antibodies were used: anti-Gr-1 (R B6-8C5), Mac-1 (M1/70), B220 (RA3-6B2), CD4 (H129.19), CD8a (53-6.7) (all PharMingen, San Diego, CA), mAbs were used as lineage markers. Anti-c-kit (2B8) and Sca-1 (E13-161.7) (both PharMingen) were used as hematopoietic stem cell markers. FITC- or PE-conjugated antibodies were used.

Cell Sorting and Flow Cytometric Analysis. Freshly isolated spleen, liver, and bone marrow cells were obtained according to methods previously described (27). Cells were washed three times with RPMI 1640 medium and suspended at a concentration of 2 × 10^6 cells/ml of PBS containing 2.5% FCS and 0.05% sodium azide before use. Double staining using anti-CD45 (30F11.1) (PharMingen) mAbs and various lineage and hematopoietic stem cell–specific mAbs were used.

Result

IL-6–sIL-6R Mice Are Smaller and Have a Reduced Body Weight. The IL-6 transgene expression was regulated by the mouse metallothionein-1 promoter. The sIL-6R transgene expression was driven by the rat PEPCK promoter, which acts neonatally and permits evaluation of the influence of gene products after birth without intrauterine developmental effects. A distinct phenotype was observed in IL-6–sIL-6R mice compared with single-transgenic mice: IL-6 and sIL-6R single-transgenic and nontransgenic littermates (Fig. 1A, left; data not shown) developed normally with respect to weight gain, food and water intake, appearance, and behavior. However, IL-6–sIL-6R mice (Fig. 1A, right) were remarkably smaller compared with single-transgenic and nontransgenic littermates. When body weights were measured over a period of 20 wk, it became evident that the IL-6–sIL-6R mice developed reduced body weights as soon as 6 wk after birth (Fig. 1B). At the age of 20 wk, mean values obtained in IL-6–sIL-6R mice were around 20–25 g compared with 40 g in single-transgenic and nontransgenic littermates (Fig. 1B). Moreover, at autopsy and at histopathological analysis, IL-6–sIL-6R mice were found to have markedly reduced body fat in all regions of the organism compared with single-transgenic littermates (data not shown).

Massive Increase of Liver and Spleen Weight in IL-6–sIL-6R Mice. Most remarkably, at the age of 4–6 wk, IL-6–sIL-6R mice displayed distended abdominal regions (Fig. 1A). Single-transgenic and nontransgenic littermates were normal in this respect. The abdominal distension was caused by a marked increase of liver and spleen weights relative to the total body weight. Time course experiments (Fig. 2, A and B) revealed steadily increasing relative liver and spleen weights in IL-6–sIL-6R mice, while the respective relative organ weights in single-transgenic and nontransgenic littermates were unchanged. At wk 20, the relative liver weight (Fig. 2A) had duplicated and the relative spleen weight (Fig. 2B) had increased by a factor of 5 when compared with single-transgenic and nontransgenic littermates.

Immunohistochemistry. Immunohistochemistry was performed using the indirect peroxidase staining method. Rat anti–mouse Ly-6, rat anti–mouse CD3, rat anti–mouse B220, (PharMingen), and rat anti–mouse Ia (Boehringer Mannheim, Germany) were used as primary antibodies. As a secondary antibody, peroxidase–conjugated goat anti–rat IgG (H + L) (Dianova, Hamburg, Germany) was used.

In Vitro Colony-forming Assays. The number of hematopoietic progenitor cells was determined in clonogenic assays as described previously (28). Cell samples were obtained from femurs and suspensions of spleen and liver derived from one individual animal at the age of 8–12 wk. The culture mixture for the assays consisted of 0.8% methyl cellulose, 1% BSA, 10% FCS, 5% human plasma and growth factors. Cells were seeded in agar cultures at a concentration of 10^5/well, and CFU-GM and BFU-E were stimulated with G-CSF (50 ng/ml) plus murine IL-3 (300 U/ml) or stem cell factor (150 ng/ml) plus IL-3 plus erythropoietin (2 U/ml), respectively. The total number of colonies per organ was calculated using the cellularity obtained for each organ and for the bone marrow, assuming that each femur represents 6% of the total marrow.

Peripheral Blood Counts. In general anesthesia, blood was obtained by cardiac puncture and analyzed in a CellDyn 3500 hemocounter (Abbott, Delkenheim, Germany).
Figure 1. Weight abnormalities in IL-6-sIL-6R mice. (A) An IL-6-sIL-6R double-transgenic mouse on the right side and a sIL-6R single-transgenic mouse on the left side at the age of 8 wk. (B) Total body weight recorded of IL-6-sIL-6R mice, of single-transgenic mice, and of nontransgenic littermates.
The gross morphology of livers and spleens were examined in mice at 8 wk of age (Fig. 2 C). Livers and spleens of 8-wk-old IL-6R mice (Fig. 2 C, left) and IL-6 mice and non-transgenic littermates (data not shown) were completely normal. Livers of 8-wk-old IL-6–sIL-6R mice (Fig. 2 C, right), however, appeared dark in color and had a rugged and humpy surface. This appearance was caused by round, whitish, and with respect to the liver surface elevated foci. The spleens of IL-6–sIL-6R mice were clearly enlarged in size (Fig. 2 C, right). The enlargement of the spleen is even more
Figure 3. Histomorphological analysis of livers and spleens in IL-6–sIL-6R (A–D), and IL-6 (A'–D') mice. (A) Liver tissue with several hematopoietic foci containing predominantly granulopoietic precursor cells and terminally differentiated segmental granulocytes (IL-6–sIL-6R, 8 wk). (A') Normal liver tissue with absence of hematopoietic foci in an 8-wk-old IL-6 transgenic mouse. (B) Spleen tissue with activated extramedullary hematopoiesis showing enlarged granulopoietic and erythropoietic areas as well as increased number of megakaryocytes (IL-6–sIL-6R, 8 wk). (B') Spleen tissue from an IL-6 transgenic mouse at 8 wk showing regular distribution of red and white pulp and little hematopoiesis. (C) Liver tissue with large hematopoietic focus displaying blastic precursor cells and terminally differentiated granulocytes (IL-6–sIL-6R, 16 wk). (C') Liver tissue from an IL-6 transgenic mouse at 16 wk with periportal plasmacytoma infiltrate. (D) Spleen tissue with destroyed architecture and complete involvement by hematopoiesis (IL-6–sIL-6R, 16 wk). (D') Spleen tissue completely infiltrated by plasmacytoma (IL-6, 16 wk). The scale bars indicate 100 μm.
remarkable in light of the fact that IL-6-IL-6R mice had half the body weight of single-transgenic and nontransgenic littersmates (see Fig. 1 B).

Increase of Liver and Spleen Weight in IL-6-sIL-6R Mice Is Caused by Extramedullary Hematopoiesis. The increase of liver and spleen weight in IL-6-sIL-6R mice was caused by a marked extramedullary proliferation of hematopoietic cells in both organs (Fig. 3, A-D), which was not detected in any other parenchymal organ. No extramedullary hematopoiesis was found in single-transgenic and nontransgenic littersmates (Fig. 3, A'-D'; data not shown). In the livers and spleens of IL-6-sIL-6R mice, multiple hematopoietic foci were present, which increased in size with time. At 8 wk, there were occasional foci (Figs. 3, A and B) present in both

Figure 4. Immunohistochemical analysis of the Ly-6 (A-C), and la (A'-C') surface protein expression on hematopoietic liver cells in IL-6-sIL-6R double-transgenic mice. Granulocytic (A) and monocytic/macrophage (A') cells are almost completely absent in the liver at the age of 4 wk. At 6 wk, scattered granulocytic (B) and small clusters of monocytic/macrophage cells (B') appear in the liver, representing early hepatic extramedullary hematopoiesis. At the age of 12 wk, there are large confluent hepatic foci showing predominantly granulopoietic differentiation (C), but also significant monopoietic cells (C'). The scale bars indicate 100 μm.
ocytes were found as soon as the age of 8 wk (Fig. 3). The number of hematopoietic progenitor cells, which are unresponsive to IL-6 alone. The IL-6–sIL-6R complex leads to extramedullary accumulation of hematopoietic progenitor cells in the bone marrow, spleen, and liver of IL-6–sIL-6R mice (Table 1, upper panel), the number of granulocytes, macrophages, and Sca-1+ progenitor cells increased by a factor of 2.3, 1.5, and 2, respectively, compared with single-transgenic or nontransgenic littermates. B cells were increased 2.3-fold in both IL-6 single-transgenic and in IL-6–sIL-6R mice compared with nontransgenic littermates. CD4+ and CD8+ lymphocytes were not different among the groups of mice tested.

The most prominent findings were seen in the liver of IL-6–sIL-6R mice (Table 1, lower panel). Granulocytes, macrophages, Sca-1+ progenitor cells, and B cells were absent or present only scarcely in single-transgenic sIL-6R mice and nontransgenic littermates. Macrophages and B cells were increased 15-fold and 45-fold in IL-6 single-transgenic mice, respectively, compared with sIL-6R single-transgenic or nontransgenic littermates. All cell lines (granulocytes, macrophages, progenitor cells, and B cells) were increased by a factor of 200–300 in IL-6–sIL-6R mice compared with IL-6R single-transgenic or nontransgenic littermates.

These results show that hematopoietic cells in the liver and spleen of IL-6–sIL-6R mice were mainly granulocytes and monocytes, to a lesser extent B cells, and not T cells.

Massive Hematopoietic Progenitor Cell Proliferation in the Liver and Spleen.

In view of the extramedullary hematopoiesis observed in liver and spleen, we investigated whether the continuous stimulation of the gp130 signal transducer affected committed hematopoietic progenitor cells of the granulocytic-monocytic and erythocyte lineages in different hematopoietic organs. The frequency of hematopoietic progenitor cells in the bone marrow, spleen, and liver of transgenic and nontransgenic mice was determined using in vitro clonogenic assays. The IL-6–sIL-6R-mediated gp130 stimulation had distinct effects on different hematopoietic lineages in extramedullary organs: the total number of CFU-GM and BFU-E in spleen and liver derived from IL-6–sIL-6R mice was increased ten times when compared

| Organ | Marker | Nontransgenic | sIL-6R | IL-6 | IL-6–sIL-6R |
|-------|--------|---------------|--------|------|-------------|
| Spleen | GR-1   | 10.0           | 10.5   | 10.7 | 23.6        |
|       | MAC-1  | 12             | 14     | 6.7  | 21.6        |
|       | SCA-1  | 20             | 21     | 23.4 | 42          |
|       | B220   | 14             | 15     | 35   | 36          |
|       | CD8    | 2.15           | 2.11   | 2.34 | 3           |
|       | CD4    | 6.8            | 8.8    | 3.8  | 5.8         |
| Liver | GR-1   | 0.003          |        | 0.004| 1.14        |
|       | MAC-1  |                | 0.006  | 0.08 | 1.04        |
|       | SCA-1  |                |        | 0    | 0.932       |
|       | B220   | 0.001          | 0.002  | 0.09 | 0.665       |

Freshly isolated cells obtained from organs as indicated in the table were subjected to FACS® analysis. Values are expressed as total numbers of cells × 10^6 per organ. One representative experiment out of three is shown.

Table 1. Distribution of Phenotypes of H ematopoietic C eils Isolated from the Spleen and Liver

Characters of H ematopoietic C eils by Immunohistochemistry and FACS® Analysis. To characterize further the nature of the hematopoietic cells in the liver, immunohistochemistry was performed with antibodies to surface antigens of neutrophils (Ly-6), monocytes/macrophages (Ia), B cells (B220), and T cells (CD3). As early as 4 wk after birth, IL-6–sIL-6R mice showed single positive neutrophils (Fig. 4 A) and monocytic/macrophages (Fig. 4 A') in the liver. At the age of 6 wk, there were increased numbers of scattered neutrophilic granulocytes (Fig. 4 B) and small clusters of monocytic cells (Fig. 4 B') present in the liver. Large confluent foci of both cell types were detected at the age of 16 wk (Figs. 4, C and D). Immunohistochemistry using antibodies directed against B and T cell epitopes revealed that B cells were also present, but to a lesser extent, and that T cells were absent (data not shown). In single-transgenic and nontransgenic littermates, immunohistochemistry using the above mentioned antibodies gave negative results (data not shown).

Cells harvested from liver, spleen, and bone marrow from single-transgenic and nontransgenic littermates and from IL-6–sIL-6R mice were subjected to FACS® analysis to determine the percentage of macrophages, granulocytes, progenitor cells, and B and T lymphocytes present in these organs. In the bone marrow, there were no significant differences detected between IL-6–sIL-6R mice and single-transgenic and nontransgenic littermates with respect to the various surface antigens tested (data not shown). In the spleen of IL-6–sIL-6R mice (Table 1, upper panel), the number of granulocytes, macrophages, and Sca-1+ progenitor cells increased by a factor of 2.3, 1.5, and 2, respectively, compared with single-transgenic or nontransgenic littermates. B cells were increased 2.3-fold in both IL-6 single-transgenic and in IL-6–sIL-6R mice compared with nontransgenic littermates. CD4+ and CD8+ lymphocytes were not different among the groups of mice tested.

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Extramedullary hematopoietic progenitor cells are functionally normal. Peripheral blood cell numbers were determined at the age of 8, 16, and 24 wk in IL-6–sIL-6R mice and compared with numbers measured in IL-6 and sIL-6R single-transgenic mice and nontransgenic littermates. Whereas the peripheral blood cell numbers of single-transgenic and nontransgenic littermates were unchanged over the period of time tested, a most dramatic increase of white blood cells consisting mainly of neutrophilic granulocytes was observed in IL-6–sIL-6R mice (Fig. 6, A and B). At the age of 8 wk, there was no difference observed between IL-6–sIL-6R mice and single-transgenic mice and nontransgenic littermates. However, at the age of 16 wk, white blood cells and neutrophils increased by a factor of 10 and 17, respectively, and at the age of 24 wk by a factor of 27 and 48, respectively, in IL-6–sIL-6R mice. Platelets (Fig. 6 C), red blood cells (Fig. 6 D), and hemoglobin values (Fig. 6 E) of IL-6–sIL-6R mice increased by a factor of 2.1, 1.5, and 1.5, respectively, when compared with age-matched nontransgenic littermates and did not change in 16- and 24-wk-old mice. In single-transgenic animals, peripheral blood cell counts and hemoglobin values were not altered compared with nontransgenic mice (Fig. 6, C–E). Thus, the expanded pool of hematopoietic progenitor cells in IL-6–sIL-6R mice appears to retain its capacity to undergo proliferation and differentiation into mature blood cells, resulting in expansion of circulating blood cells.

**Discussion**

Our study demonstrates that the activation of the gp130 signal transducer by a complex of IL-6 and the sIL-6R represents a major stimulation of growth and differentiation of hematopoietic progenitor cells in vivo. The presence of IL-6 alone is not sufficient to stimulate these cells. As demonstrated by histomorphological and immunohistochemical data and in vitro clonogenic assays, IL-6–sIL-6R mice exhibit an expansion of hematopoietic progenitor cells of the granulocytic-monocytic, megakaryocytic, and erythroid lineages, whereas peripheral blood cell counts and hemoglobin values were not altered compared with nontransgenic mice (Fig. 6, C–E).

We conclude that these hematopoietic progenitor cells express gp130 signal-transducing molecules on their surface and lack the specific IL-6R. This notion is underscored by a recently published study using gp130-deficient mice, which were shown to exhibit a greatly diminished number of hematopoietic progenitor cells (29). Thus, gp130 activation is required to stimulate hematopoietic progenitor cells. Because IL-6 and IL-11 act via a gp130 homodimer, loss of IL-6 can be compensated by IL-11, possibly explaining why IL-6-deficient mice are viable (30, 31). In mice overexpressing cytokines like LIF (32) or OSM (33), requiring the gp130/LIF-R heterodimer for signal transduction, hematopoietic effects are scarce, indicating that hematopoietic progenitor cells express little or no LIF-R on their surface. This hypothesis is supported by findings from mice in which the LIF-R has been deleted by targeted disruption (34). These mice show no significant abnormalities of the hematopoietic system.
The fact that increased B cell numbers are detected in spleen and liver of IL-6 and IL-6-sIL-6R transgenic mice supports the notion that B cells express the specific IL-6R on their surface and thus may be activated by IL-6 alone (35). Accordingly, IL-6 induced plasmacytosis in IL-6 and IL-6-sIL-6R mice by stimulating proliferation and maturation of B cells (36, 37).

In IL-6-sIL-6R double transgenic mice a marked reduction of body weight and body size associated with a reduction of body fat was observed compared with single-transgenic and nontransgenic littermates. Currently, we can not explain the reduced body size observed in these animals. However, the reduced body fat that we observed in IL-6-sIL-6R mice could be explained by the fact that the recently identified leptin receptor (OB-R) (38) is a signal-transducing protein similar to gp130, involving the same intracellular signaling molecules (39, 40). The OB-R is located in many tissues, including the hypothalamus, and is
involved in the regulation of the size of adipose tissue mass through effects on satiety and energy metabolism. The natural ligand of the O B-R, leptin, a 16-kD protein, is expressed primarily in the white adipose tissue (41). It might be possible that the IL-6–sIL-6R complex activates gp130 molecules on O B-R–expressing cells, thereby activating leptin-specific signaling pathways.

We have provided evidence that the IL-6–sIL-6R complex in vivo induces extramedullary hematopoiesis in liver and spleen but not in bone marrow. It is known that there are resident hematopoietic stem cells in the adult liver (42). These cells might be expanded after stimulation by IL-6–sIL-6R in a concentration-dependent manner. Rather high concentrations of IL-6 (2 nM) and sIL-6R (>20 nM) are required for the activation of gp130-expressing cells in vitro (43). This is somewhat surprising considering that the K\text{D} of the IL-6–sIL-6R interaction is in the range of 1 nM (16, 44). A possible explanation could be that the average half-life of the IL-6–IL-6R complex is shorter than the time required to assemble the functionally active IL-6–IL-6R/gp130 complex. We have recently demonstrated that the sIL-6R acts predominantly at the site of its generation (M. Peters et al., submitted for publication). Because liver-specific promoters drive the expression of the transgenes used in the present study, one could reason that sufficient concentrations of IL-6 and sIL-6R required for an effective stimulation of hematopoietic progenitor cells are found predominantly in the liver. Recently, transgenic mice coexpressing IL-6 and the membrane-bound IL-6R have been generated (45) in which soluble IL-6R was generated by shedding at levels 100-fold lower compared with IL-6–sIL-6R mice of the present study. No hematopoietic abnormalities were found in these mice, indicating that high levels of the IL-6–sIL-6R complex are required to stimulate hematopoietic progenitor cells.

Extramedullary hematopoiesis in the spleen has been recognized after administration of recombinant G-CSF to mice (46). Splenic stimulation of progenitors in IL-6–sIL-6R mice indicates that cytokines of the IL-6 family can contribute to hematopoiesis in the spleen. Recently, it has been demonstrated that LIF overexpression results in moderate splenomegaly and slightly activated extramedullary hematopoiesis (32, 47). LIF-deficient mice show a reduction of hematopoietic progenitors in the spleen and bone marrow. However, hematopoietic spleen and bone marrow progenitor cells from those mice transplanted into lethally irradiated normal mice retained their normal function, indicating that LIF is required in the microenvironment to maintain stem cell numbers, rather than affecting the potential of hematopoietic progenitors (48). In light of these results, one might speculate, that similar to LIF, the IL-6–sIL-6R complex contributes to the microenvironment required by hematopoietic cells in spleen and liver. However, it is of interest that stimulation by IL-6–sIL-6R leads to far stronger hematopoiesis than stimulation by LIF.

It is possible that liver and spleen cells express other important growth factors like stem cell factor (49, 50), chemokines as pre-B cell–stimulating factor (51), or flt-3 (52), either in membrane-bound or soluble form. These cytokines might be involved as cofactors in the growth of hematopoietic progenitor cells in these organs.

Hematopoietic progenitor cells stimulated by the IL-6–sIL-6R complex are functionally normal. This is reflected by the highly elevated numbers of granulocytes, megakaryocytes, and erythrocytes in the peripheral blood. Thus, the expanded pool of hematopoietic progenitor cells appears to retain its capacity to undergo proliferation and differentiation into mature blood cells, resulting in expansion of circulating blood cells.

In summary, there are three major findings of this study. First, we provided evidence that continuous activation of the gp130 signal transducer by molecules like the IL-6–sIL-6R leads to a most effective stimulation of hematopoietic progenitor cells. Progenitor cells of the granulocytic–monocytic, megakaryocytic, and erythroid lines are involved. Second, IL-6–sIL-6R causes a stimulation of predominantly extramedullary and not bone marrow hematopoietic progenitor cells. Third, progenitor cells are functionally normal, retaining their capacity to undergo maturation into mature blood cells. Apart from expanding CD34\text{+} cells obtained from cord blood in vitro (43), continuous activation of the gp130 signal transducer by agents like the hIL-6–hsIL-6R complex could be the most effective mechanism to reconstitute hematopoiesis in vivo or ex vivo in patients with cytopenic conditions.
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