Liver degeneration and lymphoid deficiencies in mice lacking suppressor of cytokine signaling-1

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ABSTRACT SOCS-1, a member of the suppressor of cytokine signaling (SOCS) family, was identified in a genetic screen for inhibitors of interleukin 6 signal transduction. SOCS-1 transcription is induced by cytokines, and the protein binds and inactivates signal transducers and activators of transcription and of the interleukin 6 receptor. Thus, SOCS-1 forms part of a feedback loop that modulates signal transduction from cytokine receptors. To examine the role of SOCS-1 in vivo, we have used gene targeting to generate mice lacking this protein. SOCS-1-deficient mice exhibited stunted growth and died before weaning with fatty degeneration of the liver and monocytic infiltration of several organs. In addition, the thymus of weaning mice was reduced markedly in size, and there was a progressive loss of maturing B lymphocytes in the bone marrow, spleen, and peripheral blood. Thus, SOCS-1 is required for in vivo regulation of multiple cell types and is indispensable for normal postnatal growth and survival.

Cytokines induce a variety of biological responses by binding to specific cell surface receptors and activating cytoplasmic signal transduction pathways (1). Many of the proteins that mediate these signals from the cell surface to the nucleus, such as Janus kinases and signal transducers and activators of transcription (STATs), have been well characterized (2, 3). There is clearly a need for tight control of these signal transduction pathways, but little is known about how these signals are turned off. The adverse consequences of an imbalance between positive and negative signals is evident in the motheaten mouse, which lacks the phosphatase SHP-1. These mice fail to regulate cytokine responses appropriately, resulting in hyperproliferation and accumulation of cells in several hematopoietic lineages and the development of systemic autoimmune disease (4).

A new family of proteins that regulates responses to cytokines has been identified recently. Suppressor of cytokine signaling1 (SOCS-1) was cloned in a functional screen for inhibitors of interleukin 6 signal transduction (SOCS-1) transcription is induced by cytokines, and the protein binds and inactivates signal transducers and activators of transcription and of the interleukin 6 receptor. Thus, SOCS-1 forms part of a feedback loop that modulates signal transduction from cytokine receptors. To examine the role of SOCS-1 in vivo, we have used gene targeting to generate mice lacking this protein. SOCS-1-deficient mice exhibited stunted growth and died before weaning with fatty degeneration of the liver and monocytic infiltration of several organs. In addition, the thymus of weaning mice was reduced markedly in size, and there was a progressive loss of maturing B lymphocytes in the bone marrow, spleen, and peripheral blood. Thus, SOCS-1 is required for in vivo regulation of multiple cell types and is indispensable for normal postnatal growth and survival.

MATERIALS AND METHODS

Generation of Targeted Embryonic Stem (ES) Cells and SOCS-1−/− Mice. A 5′ fragment of the murine SOCS-1 gene extending ~2.5 kilobases (kb) from the protein initiation ATG was generated by PCR. This fragment was ligated directly upstream of the initiation codon of β-galactosidase via a BamHI site in the plasmid pGALαloxneo, which also contains a PGKneo cassette flanked by loxP sites. A 3.2-kb BamHI/EcoRI fragment from the genomic SOCS-1 locus was blunted and ligated into a blunted XhoI site 3′ to the PGK neo cassette. This construct was linearized and electroporated into 129/Sv-derived W9.5 ES cells (12). Clones surviving selection in 175 μg/ml G418 were screened for those in which the targeting vector had recombined with an endogenous SOCS-1 allele by using Southern blots of EcoRI-digested genomic DNA probed with a 1.5-kb EcoRI-HindIII genomic SOCS-1 fragment (Fig. 1, probe A). A targeted ES cell clone was injected into C57BL/6 blastocysts to generate chimeric mice. Male chimeras were mated with C57BL/6 females to yield SOCS-1 heterozygotes, which were interbred to produce wild-type (SOCS-1+/−), heterozygous (SOCS-1+−/−), and homozygous mutant (SOCS-1−−/−) mice. The genotypes of offspring were determined by Southern blot analysis of genomic DNA extracted from tail biopsies as described above. The deletion of SOCS-1 coding sequence and subsequent inability to produce SOCS-1 mRNA in mutant mice was confirmed in nucleic acid blots, which were performed as described (13). Northern blots were probed with probe B (Fig. 1) from the SOCS-1 coding gene expression is induced by cytokines both in vitro and in vivo, and, once produced, they act directly on components of the cytokine signaling pathways to switch them off (5–7, 10). The SOCS proteins suppress cytokine signaling in at least two distinct ways: SOCS-1 inhibits the signal by interacting with Janus kinases and inhibiting their catalytic activity (6, 7) whereas CIS binds directly to the cytokine receptor, competing with STATs for access to docking sites on the activated receptor (9).

Little is understood about specificity within the SOCS family of proteins. Constitutive over-expression of SOCS-1 inhibits cellular responses in vitro to a variety of cytokines, including IL-6, leukemia inhibitory factor, interferon-γ, and thrombopoietin (5–7, 11), but the primary targets and biological consequences of SOCS-1 action in vivo are unknown. To address these questions, we have generated mice that lack the SOCS-1 protein. These animals exhibited stunted growth and abnormalities in a diverse range of organs and died before weaning, indicating an essential role of this negative regulator in postnatal growth and survival.

Abbreviations: SOCS, suppressor of cytokine signaling; STAT, signal transducers and activators of transcription; IL, interleukin; kb, kilobase; ES, embryonic stem.

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14395
Production of Mice Lacking SOCS-1. A targeting construct was assembled to enable homologous recombination to replace the entire SOCS-1 coding region with β-galactosidase in ES cells (Fig. 1a). An ES cell clone bearing a disrupted SOCS-1 allele was used to generate chimeric mice, from which animals lacking a functional SOCS-1 gene were bred. Southern blot analysis of tail biopsies from offspring of heterozygous parents (Fig. 1b) revealed that SOCS-1<sup>−/−</sup> mice were born in similar numbers to their wild-type littermates (30:88:32 for SOCS-1<sup>−/−</sup>;SOCS-1<sup>−/−</sup>;SOCS-1<sup>−/−</sup>), suggesting that SOCS-1 was not essential for normal embryonic development. Northern blots of RNA extracted from the thymus of SOCS-1<sup>−/−</sup> mice confirmed that SOCS-1 mRNA was not transcribed from the disrupted allele (Fig. 1c).

Postnatal Lethality and Histopathology of SOCS-1<sup>−/−</sup> Mice. At birth, SOCS-1-deficient mice were indistinguishable from their normal littermates (birth weights: SOCS-1<sup>−/−</sup>, 1.90 ± 0.29 g; SOCS-1<sup>−/−</sup>, 1.89 ± 0.29 g; and SOCS-1<sup>−/−</sup>, 1.94 ± 0.10 g, n = 2–8). However, within 10 days the SOCS-1<sup>−/−</sup> mice were significantly smaller (body weight at days 9–12: SOCS-1<sup>−/−</sup> or SOCS-1<sup>−/−</sup>, 6.27 ± 1.58 g; and SOCS-1<sup>−/−</sup>, 4.06 ± 1.12 g; n = 9–11) and became ill and died before they reached 3 weeks of age (Fig. 2a and b). Histological examination of the organs of SOCS-1<sup>−/−</sup> mice at 7–20 days of age revealed major abnormalities in the liver. Parenchymal cells commonly contained an accumulation of lipid-containing vacuoles, resulting in cells with expanded cytoplasm and shrunken nuclei (Fig. 2c and d). The fatty degeneration either involved local areas not related to portal vessels or was generalized throughout the whole organ. In the latter case, which represented half of the SOCS-1<sup>−/−</sup> livers examined, areas of subcapsular necrosis were present. Unlike livers from normal neonatal mice, which showed focal aggregates of erythroid and, less commonly, granulocytic cells, the SOCS-1<sup>−/−</sup> livers exhibited both focal and generalized infiltration, predominantly by immature and mature monocytic and granulocytic cells (Fig. 2c and d). Less frequently, megakaryocytes and eosinophils were present and were more numerous in SOCS-1-deficient mice than in their heterozygous or normal littersmates. Abnormalities were also evident in three other organs of SOCS-1<sup>−/−</sup> mice. In seven of eight mice examined, the pancreas was infiltrated by monocytes invading septal regions between acini, associated with some reduction in acinar size (Fig. 2e and f). In the majority of SOCS-1<sup>−/−</sup> mice, infiltration of the heart by monocytes and less numerous granulocytes also was observed (Fig. 2g and h), as was increased cellularity of alveolar walls in the lung, often associated with macrophage cuffing around major vessels (data not shown). Other than the overall reduction in organ size, paralleling the stunted growth of these animals, no abnormalities were observed in skeletal muscle, gut, bladder, kidney, gonads, or brain of SOCS-1<sup>−/−</sup> mice.

Lymphocyte Deficiencies in SOCS-1<sup>−/−</sup> Mice. In comparison with wild-type littermates, the peripheral blood of SOCS-1<sup>−/−</sup> mice revealed a consistent and severe reduction in lymphocyte levels, a moderate reduction in eosinophil and platelet numbers, a consistently lower hematocrit, and a variable elevation of blood granulocytes (Table 1 and Fig. 2i). The smaller body weight of SOCS-1-deficient mice was asso-
associated with a femoral marrow cell count that was not that of littermate mice, and cytological analysis revealed a consistent deficit of lymphocytes in this population (Fig. 2k). Although spleen weight appeared to be normal in SOCS-1−/− mice, this organ also exhibited a relative deficit in lymphocytes, which became more evident with increasing age (Fig. 2j). Histological analysis of the spleen showed that, in SOCS-1−/− mice, lymphoid follicles either were completely absent or were composed of immature cells, often mixed with monocytes and granulocytes. Most SOCS-1-deficient spleens showed expanded areas of red pulp with nucleated erythroid cells as the dominant population. The most striking hematopoietic abnormality in SOCS-1−/− mice was the small size of the thymus (Fig. 2j). Unlike normal neonatal mice, the number of cortical lymphocytes failed to increase with age, and, instead, the cortex became progressively depleted of lymphoid cells. The lymph nodes showed a similar deficit in lymphoid follicle formation, and, although Peyer’s patches were present, these often contained few lymphocytes.

The reduction in thymic cellularity in SOCS-1−/− mice was not associated with selective loss of a specific subset of thymocytes. Fluorescence-activated cell sorting revealed normal ratios of T cells expressing the CD4 and/or CD8 cell surface markers (Fig. 3a). In contrast, selective depletion of maturing B-lymphoid cells was evident in the bone marrow and

Table 1. Hematological profile of peripheral blood in SOCS-1−/− mice

<table>
<thead>
<tr>
<th>Genotype</th>
<th>SOCS-1+/− or +/+</th>
<th>SOCS-1−/−</th>
</tr>
</thead>
<tbody>
<tr>
<td>Platelets, ×10^9/ml</td>
<td>737 ± 22</td>
<td>435 ± 251</td>
</tr>
<tr>
<td>Hematocrit, %</td>
<td>37 ± 3</td>
<td>28 ± 3</td>
</tr>
<tr>
<td>White cell count, ×10^9/ml</td>
<td>5.55 ± 2.46</td>
<td>4.08 ± 2.64</td>
</tr>
<tr>
<td>Neutrophils</td>
<td>0.95 ± 0.64</td>
<td>2.42 ± 1.93</td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>3.63 ± 1.85</td>
<td>0.73 ± 0.36</td>
</tr>
<tr>
<td>Monocytes</td>
<td>0.73 ± 0.36</td>
<td>0.94 ± 1.22</td>
</tr>
<tr>
<td>Eosinophils</td>
<td>0.16 ± 0.24</td>
<td>0.01 ± 0.02</td>
</tr>
</tbody>
</table>

Mean ± SD of data from 10 (SOCS-1−/−) and 14 (SOCS-1+/− or +/+) mice.
Fig. 3. Lymphocyte profiles in SOCS-1−/− mice. (a) Fluorescence-activated cell sorter analysis showing unaltered distribution of thymocytes expressing the CD4 and CD8 cell surface markers and the deficiency in pre-B and B (B220−) as well as mature B cells (B220+IgM+) in the bone marrow of a representative SOCS-1−/− mouse. Profiles of the equivalent populations from age-matched littermate mice (LM) are shown for comparison. (b) Analysis of cultures of bone marrow cells from SOCS-1−/− and littermate (LM) mice showing similar in vitro capacity for B cell (B220+IgM+) maturation. (c) Double-staining with β-galactosidase and Thy-1 (thymus) or B220 (spleen) from heterozygous SOCS-1+/− and wild-type mice demonstrating transcriptional activity of the SOCS-1 locus in thymocytes and B-lymphoid cells.

β-galactosidase was evident in all CD4 and CD8 thymic subsets (data not shown). β-galactosidase activity was also evident in a proportion of pre-B (B220+IgM−) and mature B (B220+IgM+) cells from the spleen (Fig. 3c and data not shown) or the bone marrow (data not shown) of SOCS-1−/− mice. As expected, β-galactosidase activity was negligible in control wild-type lymphoid populations (Fig. 3c). These data confirm that SOCS-1 normally is expressed in the T and B lymphoid lineages.

**DISCUSSION**

Through the generation of mice bearing a targeted disruption of the gene encoding SOCS-1, we have shown that this protein is indispensable for normal postnatal development. Although apparently normal at birth, SOCS-1−/− mice exhibited stunted growth and died within the first 3 weeks of life. Major abnormalities were evident in the liver, with fatty degeneration of parenchymal cells, which may be sufficient to account for the mortality of SOCS-1−/− mice. Deficiencies also were observed in the lymphoid organs, with dramatically reduced cellularity of the thymus and a severe progressive depletion of pre-B and B cells. Extensive focal and generalized hematopoietic infiltration was also evident in the livers of SOCS-1−/− mice, and the pancreas, lungs, and heart of these animals often exhibited a less severe monocytic infiltration (Fig. 2).

The complexity of the phenotypic abnormalities in SOCS-1−/− mice implies that SOCS-1 is indispensable for the normal in vivo regulation of multiple cell types. Although it is feasible that some of the abnormalities observed in SOCS-1−/− mice may be the indirect consequences of a primary defect in one organ, such as the liver, it is striking that the cells that are lost or damaged in SOCS-1−/− mice are among those that normally express this gene (T and B lymphocytes; Fig. 3) or in which expression is induced by cytokine stimulation (liver; ref. 5).

Thus, the lymphocyte deficits, as well as the degeneration of liver parenchymal cells, seem likely to be a direct consequence of the lack of SOCS-1 in these cells. Although the diverse abnormalities that characterize mice lacking SOCS-1 may reflect a common action in divergent cell types, our data do not exclude the possibility that SOCS-1 may mediate differing and independent effects in different organs.

Previous studies in vitro have established that SOCS-1 can act to inhibit cytokine signaling (5), suggesting that abnormalities in mice lacking this protein may result from inappropriate responses to cytokine stimulation. Indeed, STAT1 activation, which normally occurs in response to agents such as IL-6 or interferon γ, appears to occur constitutively in the livers of SOCS-1-deficient mice (data not shown). Overexpression of interferon γ is known to cause liver damage and B-lymphocyte depletion (16, 17), the disease that arises in SOCS-1−/− mice may be the result of dysregulation of signals from such a cytokine. Alternatively, activation of STATs also has been linked to the induction of apoptosis (18, 19), raising the possibility that SOCS-1−/− mice may be unable to regulate apoptotic signals appropriately. To explore these possibilities, we currently are crossing the SOCS-1−/− mice to mice lacking interferon γ or regulators of apoptosis. An additional possibility is raised by the monocytic infiltration observed in multiple organs of SOCS-1−/− mice (Fig. 2). Preliminary data indicate that SOCS-1-deficient granulocyte-macrophage progenitor cells are hyper-responsive to proliferative stimulation by granulocyte–macrophage colony-stimulating factor. If this is associated with the functional activation of monocytes and granulocytes, infiltrating hematopoietic cells also might contribute to tissue damage in SOCS-1−/− mice.

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