

# Hematopoiesis

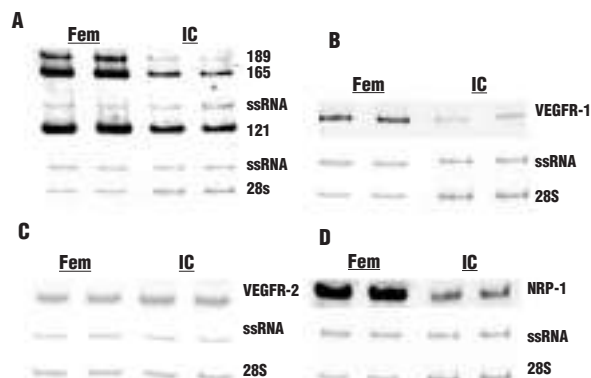
## Differential expression of vascular endothelial growth factor and its receptors in hematopoietic and fatty bone marrow: evidence that neuropilin-1 is produced by fat cells

**Vascular endothelial growth factor (VEGF), its receptors (VEGFR-1, VEGFR-2) and neuropilin-1 (NRP-1) are expressed at variable levels in bone marrow. NRP-1 expression is higher in fatty bone marrow than in hematopoietic marrow. Adipocytes are responsible for NRP-1 expression suggesting that they may play a role in hematopoiesis by producing NRP-1 or that NRP-1 may regulate adipocyte activity.**

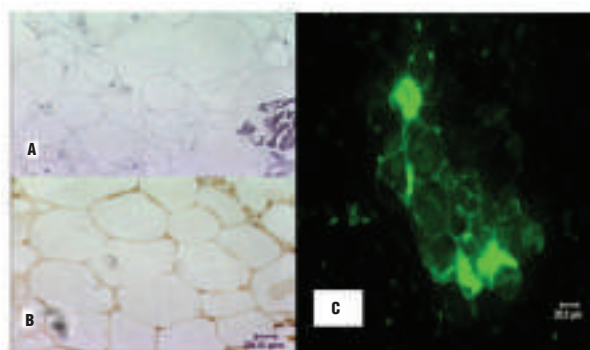
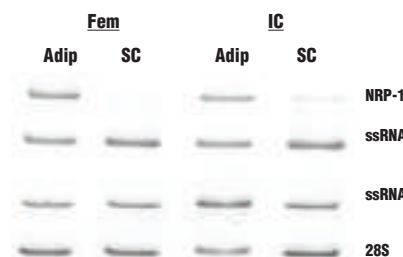
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Recent evidence suggests that vascular endothelial growth factor (VEGF) is involved in hematopoiesis and in the pathogenesis of hematopoietic malignancies.<sup>1</sup> Neuropilin-1 (NRP1) was recently recognized as an isoform-specific receptor for VEGF165<sup>2</sup> acting as coreceptor for the VEGFR-2. Our primary goal was to evaluate the expression of NRP-1 in human hematopoietic marrow and to identify cells responsible for its expression *in vivo*. Key partners of NRP-1 such as VEGF, VEGFR-1 and VEGFR-2 were also studied. Samples from iliac bone (rich in hematopoietic marrow), and from femurs (mostly composed of fatty cells) were from 18 human bone marrow core samples obtained from autopsies performed at the Department of Pathology and from 8 human core biopsies obtained from patients undergoing hip surgery at the Department of Orthopedic Surgery. The total RNA was purified by centrifugation on a cesium chloride cushion<sup>3</sup> and from cell suspensions using the High Pure RNA Isolation kit (Roche Diagnostics, Mannheim, Germany). Reverse transcription-polymerase chain reaction (RT-PCR) was performed under non-competitive conditions in the presence of a synthetic RNA used as internal standard in order to make the procedure quantitative.<sup>4</sup> To evaluate a possible correlation between the hematopoietic activity and the mRNA level of the VEGF isoforms and their receptors, total RNA was isolated from femoral fatty bone marrow and from iliac crest hematopoietic bone marrow collected from the same donor and RT-PCR was performed as above. The hematopoietic activity was evaluated by calculating the relative percentages of the tissue hematopoietic cells versus adipose cells.<sup>5</sup> In all tested specimens, the main isoforms found were VEGF165 and VEGF121 expressed at variable levels. This pattern of expression seems to be specific to bone marrow and is in agreement with the data reporting the RT-PCR analysis of the VEGF121 and VEGF165 in a series of human hematopoietic cell lines.<sup>6</sup> A significant expression of the 189 isoform was observed in femoral bone marrow whereas it was barely detectable in iliac bone marrow (Figure 1A). In most of samples of femoral bone marrow, the total VEGF mRNA was higher than in iliac crest bone marrow:



**Figure 1.** A: Representative examples of the electrophoretic patterns of RT-PCR products of A: VEGF, B: VEGFR-1, C: VEGFR-2 and D: NRP-1 mRNA measured in duplicate using total RNA isolated from the femoral (fem) and the iliac crest (IC) bone marrow of the same donor. The total RNA in the sample solution was evaluated by measuring the 28S rRNA. A synthetic RNA (ssRNA) was added in each tube reaction to monitor the yield of both the reverse transcription and the amplification steps (only for 28S rRNA, VEGF and NRP-1).



**Figure 2.** A. Representative examples of electrophoretic patterns of RT-PCR products of NRP-1 mRNA using total RNA isolated from adipocytes (adip) and sedimented cells (SC) from femoral (Fem) and iliac crest (IC) bone marrow. B. NRP-1 immunoreactivity on bone marrow sections (a and b) and adipocytes smears (c). NRP-1 immunoperoxidase staining in femoral bone marrow adipocytes (b). (a) are negative controls for the specific signal using normal goat immunoglobulins. NRP-1 immunoreactivity is revealed by immunofluorescence in isolated adipocytes smears (c).

this could be related to the predominance of adipocytes in the femoral samples. Indeed, it has been shown that VEGF mRNA is upregulated during the conversion of 3T3 preadipocytes to adipocytes.<sup>7</sup> The VEGFR-1, VEGFR-2 and NRP-1 were measured in the same series of samples. VEGFR-1 mRNA levels were quite variable from case to case. VEGFR-1 mRNA was either absent in iliac crest and femoral bone marrow or expressed at the same level in both tissues or expressed only in femoral bone marrow or was expressed at higher level in femoral bone marrow than in iliac crest bone marrow (Figure 1B). Values for VEGFR-2 were available in five samples only: there was no significant difference between femoral and iliac crest marrow (Figure 1C).

VEGFR-2 is essential for the development of hematopoietic stem cells during early embryonic development, it may be redundant in adult bone marrows. Since activation of VEGFR-1 is fully sufficient to rescue hematopoietic stem cell survival *in vitro* and hematopoietic repopulation *in vivo*,<sup>8</sup> the presence of VEGFR-2 may be related to the maintenance of bone marrow vasculature. NRP-1 was expressed at higher level in femoral bone marrow than in iliac crest in each donor (Figure 1D) and it seems to be inversely correlated with the hematopoietic activity. The cellular origin of NRP-1 was assessed on isolated cell populations by a floatation/sedimentation procedure. By contrast to sedimented cells (hematopoietic and stromal cells), high levels of NRP-1 mRNA were detected in the adipocytic population (Figure 2A). This was confirmed by *in situ* hybridization (not shown) and at the protein level by immunohistochemistry (Figure 2B). This is the first report demonstrating neuropilin-1 expression in bone marrow *in vivo*. The capacity of adipocytes to produce NRP-1, previously suspected to play an interactive role with hematopoietic cells<sup>9</sup> suggests that adipocytes may contribute to the regulation of hematopoiesis and/or that NRP-1 may be a novel regulator of adipocyte activity in the bone marrow, possibly as a receptor for VEGF. Although this study does not provide a definitive link between NRP-1, adipocyte function and hematopoiesis, such a relationship may exist and deserves further studies.

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## Red Cell Disorders

### Decreased plasma endothelin-1 levels in children with sickle cell disease treated with hydroxyurea

**Plasma endothelin-1 (ET-1) is elevated in patients with sickle cell disease (SCD). Hydroxyurea (HU) is the only drug with demonstrated clinical efficacy in SCD. Here we show that treatment with HU results in a decreased concentration of circulating ET-1 which is not correlated with the HU-induced increase in HbF level. Blunting of the ET-1 vasoconstrictive stimulus could contribute to the beneficial effects of HU.**

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Sickle cell disease (SCD) is characterized by unpredictable painful crises resulting from vaso-occlusion by rigid, sickled red blood cells (RBC). Still, factor(s) initiating vasoocclusive crises (VOC) remain largely unknown. In SCD, the vascular endothelium is chronically activated and expresses various adhesion molecules on its surface. Exacerbation of this activation, in particular within an inflammatory context, is believed to be (one of) the major triggering factor(s) of VOC by promoting the abnormal adhesion of RBC and other circulating cells to the endothelium.<sup>1</sup> The concentration of endothelin-1 is elevated in the plasma of SCD patients, especially during bouts of acute chest syndrome (ACS) and other complications of VOC.<sup>2</sup> Given that it is a powerful vasoconstrictor and pro-inflammatory agonist, ET-1 might be also play an important role in VOC.

Hydroxyurea (HU) significantly reduces the incidence of VOC and ACS, as well as global morbidity and mortality.<sup>3,4</sup> Its initial intended use was to induce fetal hemoglobin (HbF). However, the increment in HbF levels is not constant and it appears that effects of HU are the results of multi-targeted actions.<sup>5</sup> We recently demonstrated that HU down-regulates *ET-1* gene expression by endothelial cells in culture both in basal conditions and after stimulation with pro-inflammatory cytokines.<sup>6</sup> The aim of the