

Special Article - Anemia

New Experimental Anemic Model by Using a Nitrogen-Containing Bisphosphonate

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Abstract

Anemia is a pathological condition associated with various diseases including cancer, and is indicated by a reduction in erythrocyte numbers or the amount of hemoglobin in the peripheral blood. The analysis of changes in hematopoiesis during anemia is important to develop novel therapies and to understand hematopoietic ontogeny. We previously reported that injecting animals with Nitrogen-Containing Bisphosphonate (NBP), which is an inhibitor of osteoclastic bone resorption, decreased erythropoiesis in Bone Marrow (BM). Moreover, we induced severe anemia in a mouse model by injecting NBP in combination with Phenylhydrazine (PHZ), and embryonic type globin mRNA was detected in both the BM and the liver in this anemia model. In addition, wine-colored capsuled structures were unexpectedly observed in the abdominal cavity of this anemic mouse model, and active erythropoiesis was also observed in these structures. Here, we review recent insights into the pathogenic mechanisms underlying various forms of anemia that have been gained through our findings.

Keywords: Nitrogen-containing bisphosphonate; Extramedullary hematopoiesis; Embryonic hemoglobin; G-CSF; SDF-1

Abbreviations

NBP: Nitrogen-Containing Bisphosphonate; G-CSF: Granulocyte-Colony Stimulating Factor; PHZ: Phenylhydrazine; BM: Bone Marrow; SDF-1: Stromal Cell-Derived Factor 1; HSCs: Hematopoietic Stem Cells; HPCs: Hematopoietic Progenitor Cells; EPO: Erythropoietin

Introduction

Anemia is a pathological condition that involves a reduction in the number of erythrocytes or the amount of hemoglobin in the peripheral blood, and is caused by various diseases such as cancers and bone marrow disorders. The pathology of anemia is very complex, and studies that clarify the mechanism of recovery from anemia and its pathology are essential to establishing an understanding of the therapy, and the ontogeny of hematopoiesis and tissue engineering [1]. Many studies have reported experimental anemia models, including phlebotomy, schistosome parasite infection and drug-induction, which are known to be different from the actual disease state (Table 1) [2]. Phenylhydrazine (PHZ) is used to experimentally induce hemolytic anemia in laboratory animals [3], by the mechanism of RBC lipid peroxidation [4,5].

Hematopoiesis occurs in the Bone Marrow (BM) of adult mammals under normal conditions. BM possesses a specialized microenvironment called the 'niche' that maintains Hematopoietic Stem Cells (HSCs) [6,7]. The stem cell niche, which is composed of cellular compartments, produces several cytokines, such as G-CSF, GM-CSF and SDF-1 [8,9].

Nitrogen-Containing Bisphosphonates (NBPs) have strong anti-bone resorption effects, and are used as therapeutic agents against bone resorption disorders such as osteoporosis and metastatic bone

diseases [10-12]. We previously reported that mice injected with NBP developed decreased erythropoiesis in their BM (Figure 1) [13]. We induced severe anemia in a mouse model by injecting NBP in combination with Phenylhydrazine (PHZ), and we then analyzed the erythropoiesis and the levels of different types of hemoglobin in this model. Here, we review recent insights from the established severely

Table 1: The comparing among the experimental anemic models.

	Hb reduction	Reticulocytosis	Proerythroblast
Bled [2]	+	+	++
Parasite [2]	+	+	+
PHZ injection [2]	+	++	+

+: Significant increase compared with normal. ++: Significant increase compared other group.

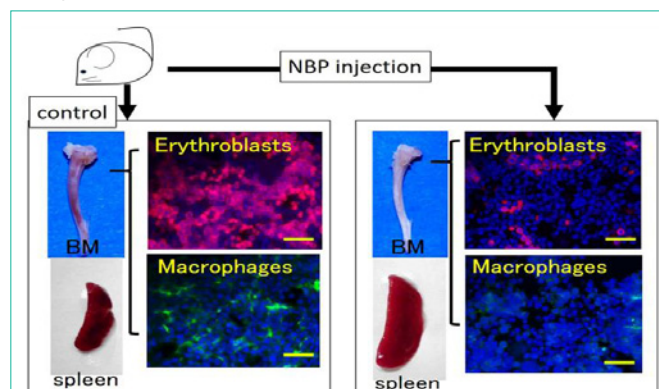
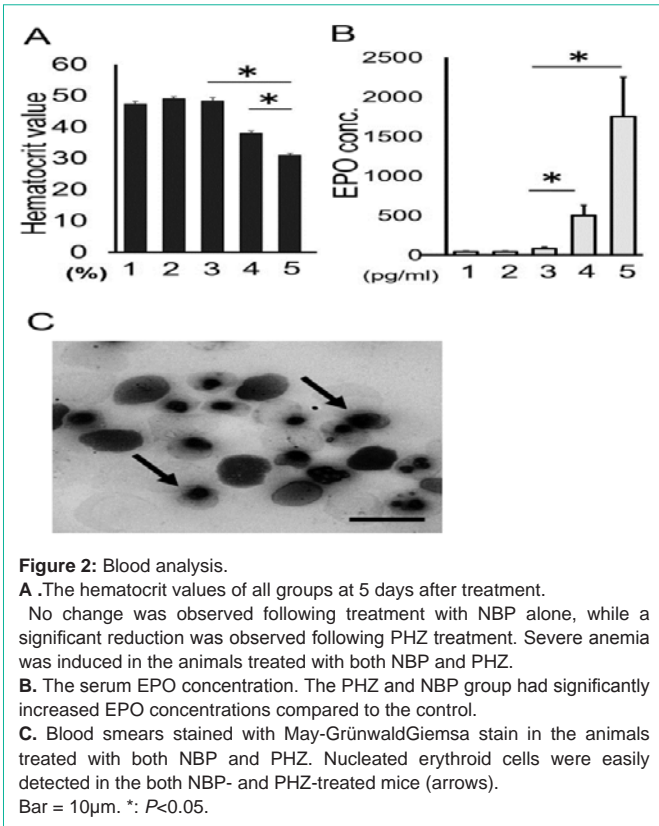


Figure 1: NBP injection impacts hematopoiesis.

The BM from NBP treated mice become white compared to that of control mice, and splenomegaly is induced by NBP injection. NBP injection also caused a reduction in the number of erythroblasts and resident macrophages in the BM. Bars = 50µm.



anemic mouse model, and provide further understanding of the new proposed mechanism for recovery from anemia.

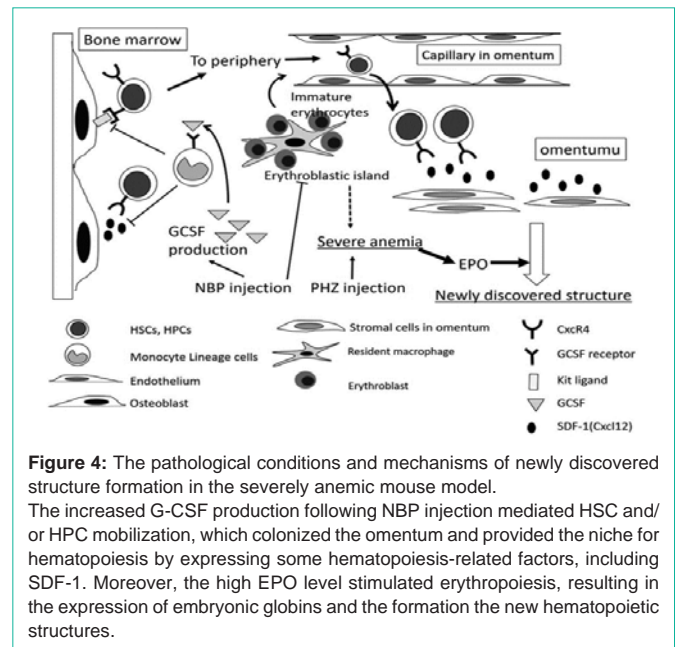
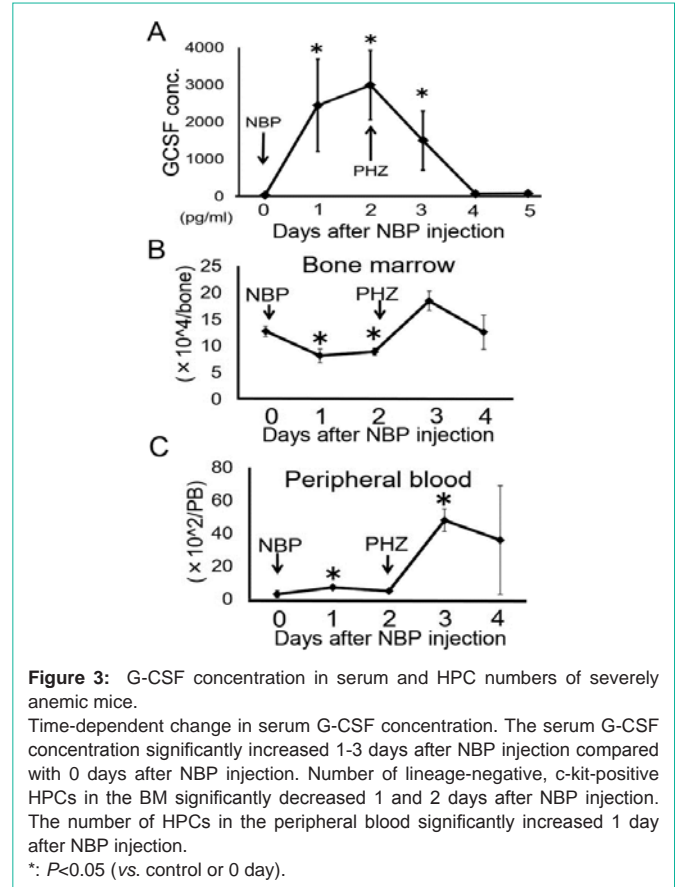
Effects of nitrogen-containing bisphosphonate on normal and splenectomized mice

The use of NBP results in strong anti-bone resorption effects. In addition, NBP has inflammatory side effects including fever, jaw osteomyelitis, osteonecrosis and extramedullary erythropoiesis [13-15]. Our previous study reported that a single injection of a relatively large dose of NBP into mice induced Extramedullary erythropoiesis by depleting resident BM macrophages and increasing the number of granulocytes in the peripheral blood [13]. Moreover, the injection of NBP into splenectomized mice induced extramedullary erythropoiesis without anemia and caused changes in EPO concentrations in the liver [16].

A severely anemic mouse model

There are many experimental models of anemia that have been used to study this disease. We analyzed the changes in hematopoiesis after induction of anemia by inhibiting erythropoiesis in the BM. Splenectomized mice were treated with NBP to inhibit erythropoiesis in the BM, and with PHZ to induce hemolytic anemia. Treating animals with both NBP and PHZ induced more serious anemia than administering PHZ alone, and the concentration of EPO in the serum was also significantly increased (Figure 2A & 2B). Moreover, numerous nucleated erythrocytes and reticulocytes were observed in the peripheral blood of mice treated with both NBP and PHZ (Figure 2C).

The number of erythroid lineage cells in the BM was significantly



decreased following NBP injection. However, the proportion of these cells in the mice treated with NBP and PHZ was significantly increased despite the use of NBP [17]. This indicates that splenectomized mice treated with both NBP and PHZ become critically anemic and display a significant increase in EPO levels, which enhances erythropoiesis.

The presence of nucleated erythrocytes in a blood smear suggested that abnormal erythropoiesis was induced, because nucleated erythrocytes are not normally observed in the blood in mammals, except during the early embryonic stages.

Expression of types of embryonic hemoglobin

Mammalian hematopoiesis occurs in two distinct waves, commonly referred to as the primitive and definitive waves, which originate in the yolk sac, and in the fetal liver and BM, respectively [18]. Yolk sac-derived primitive erythroid cells remain nucleated and enucleated terminally in circulation, whereas definitive erythroid cells produced in the fetal liver are released into circulation after complete maturation [5,6,19,20]. The components of the globin tetramer are encoded by the α - and β -globin gene loci. There are 3 functional α -globins (ζ -, $\alpha 1$ - and $\alpha 2$) and 4 β -globins (Ey-, $\beta h 1$ -, $\beta 1$ - and $\beta 2$ -) in mice [6,21,22]. Primitive erythrocytes express the embryonic complement of globin chains, which initially consist of ζ - and $\beta h 1$ -globins, followed by $\alpha 1$ - and $\alpha 2$ -, and Ey-globins at the primitive proerythroblast stage [6,21-23]. Definitive erythroid cells complete their maturation and enucleation at erythropoietic sites, the fetal liver and BM, where the adult complement of globin chains consisting of $\alpha 1$ -, $\alpha 2$ -, $\beta 1$ -, and $\beta 2$ - are expressed.

Since we observed nucleated erythrocytes in the peripheral blood of severely anemic mice in our anemia model, we determined the expression of embryonic globins. The embryonic globins ζ -, $\beta h 1$ - and Ey- and the adult globins α - and β major were expressed in the BM and the liver in the severely anemic mice. We also identified the cell clusters that expressed the embryonic globins ζ -, $\beta h 1$ - and Ey- in the BM and liver of severely anemic mice [17]. These results suggest that abnormal erythropoiesis may occur in this critical anemia model, and that embryonic globins may be activated as a response to hypoxemia.

The factors relating to embryonic globin expression

The expression of globin genes is jointly regulated by elements in the promoter regions and an upstream enhancer region called the Locus Control Region (LCR) [6]. A variety of nuclear factors that are involved in transcriptional regulation are thought to be related to globin gene expression and switching between them [24,25]. Members of the KLF (Kruppel-like factor) family are essential transcription factors that bind GC-rich sequences to regulate the biological dynamic changes in globin expression during development [26]. KLF1 (EKLF: Erythroid Kruppel-like Factor) plays an essential role in erythropoiesis and is involved in the expression of β -like embryonic globin by binding to its promoter region [27,28]. KLF2 regulates biological activity in various tissues and enhances the expression of β -like embryonic globin [29,30].

The expression of several embryonic globin transcription factors was analyzed in severely anemic model mice. The expression of *Klf1*, *Klf2* and *Gata1* was up-regulated. Moreover, the $\beta h 1$ - and Ey-globin promoters were bound to KLF1 and KLF2 in the BM and livers of mice treated with NBP and PHZ. These results indicate that KLF1 and/or KLF2 regulate the transcription of embryonic β -like globins in the BM and liver.

Induction of newly discovered hematopoietic structures

Erythropoiesis occurs in the bone marrow and the spleen even in healthy adult mice. Extramedullary erythropoiesis occurs in various

organs, including the lungs, heart, thymus, and hemal nodes, under abnormal conditions such as acute anemia [31-38].

Unexpectedly, wine-colored structures appeared in the omentum or pancreas of our severely anemic model mice [39]. Histological examination identified lymphoid follicle-like cell clusters and well-developed sinusoids, similar to the spleen, and revealed that some hematopoietic cells, including mega karyocytes and erythroblasts, populated these structures [39]. The lymphoid follicle-like clusters were filled with erythroblasts, and most of these cells were proliferating. This suggests that these newly discovered structures in a severely anemic mouse model could be the site of extramedullary hematopoiesis.

HSC mobilization & G-CSF

G-CSF is a cytokine related to granulopoiesis and G-CSF production is enhanced in inflammation and stimulates granulopoiesis. G-CSF has other functions, namely the suppression of osteoblast lineage cells and inhibition of the expression of SDF-1, SCF and VCAM-1, which results in the mobilization of HSCs and HPCs from the BM to the peripheral tissues [40,41]. We examined the cell population in the peripheral blood and BM, and the G-CSF levels in our anemic mouse model.

We first determined the number of lineage-negative and c-kit-positive HPCs in the BM and peripheral blood in order to detect the mobilization of HPCs from the BM to the peripheral tissues. The population of HPCs was significantly decreased in the BM after NBP injection, whereas it was markedly increased in the peripheral blood during the same period following treatment. One day after NBP injection, the G-CSF levels increased precipitously and remained at a high concentration for up to 3 days after treatment, which correlated with the decrease in the number of HPCs in the BM. NBP is known to enhance granulopoiesis and inflammation [13], which might be mediated by G-CSF stimulation. Our results suggest that G-CSF production is stimulated by NBP treatment, which then provoked the mobilization of HPCs from the BM to the peripheral tissue (Figure 3).

Factors relating to extramedullary hematopoiesis

The recruitment of progenitors to the vascular niche involves signaling through SDF-1 [42]. The vascular niche seems to be associated with Extramedullary hematopoiesis in the liver and spleen through the expression of SDF-1, which plays an essential role in the homing of Hematopoietic Stem Cells (HSCs) and HPCs to the BM [43,44].

We detected some cytokines that are related to hematopoiesis, including SCF and SDF-1, in the omentum, even in normal conditions [39]. In particular, SDF-1 plays an essential role in the homing of HSCs and HPCs, and is associated with Extramedullary hematopoiesis [43,44].

The omentum seems to possess a suitable microenvironment for hematopoiesis because it expresses cytokines, including SDF1, even under normal conditions. Increased levels of these cytokines might support the observed mobilization of HPC homing, colonization and differentiation in the peripheral tissues in this study.

Conclusion

We established a severely anemic mouse model by sequential

treatment with NBP and PHZ. This model demonstrated the emergence of nucleated erythrocytes in the peripheral blood, the expression of embryonic types of hemoglobin at hematopoietic sites, and the induction of newly discovered hematopoietic structures (Figure 4). Our results indicate the flexibility of hematopoiesis and suggest that that yolk sac-derived primitive erythroid cells may persist into adulthood in mice. Our insights from this severely anemic mouse model will contribute towards understanding the mechanism of anemia, and our ability to treat affected patients.

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