

Role of circulating endothelial progenitors in responsiveness to erythropoiesis-
stimulating agents in hemodialysis patients.

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Abstract

Background

Though treatment of anemia in chronic kidney disease (CKD) patients has been improved remarkably by erythropoiesis-stimulating agents (ESA), hypo-responsiveness to ESA poses a persistent problem in a subgroup of CKD patients, especially those on dialysis, with deteriorated mortality or cardiovascular risks. Pathomechanism of this condition including cardiovascular implications has not been sufficiently investigated.

Methods

Clinical parameters of 101 chronic hemodialysis patients which included those hospitalized in our university hospital were examined cross-sectionally. As a marker estimating ESA hypo-responsiveness, erythropoietin resistance index (ERI) was calculated. Numbers of circulating endothelial progenitor cells (EPC) were measured by flow cytometry. Associations among values were analyzed by methods including multiple linear regression.

Results

Majority (93%) of the subjects were hospitalized patients with various comorbidities. ERI (18.4 [7.2-33.0] IU/week/kg/g/dL) and ESA dose (161.6 [75.0-320.9] IU/week/kg) of all the subjects were relatively high. Factors negatively correlated with EPC included age, HD vintage, CRP, pulse rate, ESA dose and ERI, while male sex and systolic blood pressure were positively correlated. By multiple linear regression analysis, age, sex and ERI (standardized coefficient beta -0.202, p=0.039) remained as the independently predicting factors of EPC (log CD133/Flk1+ EPC). Darbepoetin alpha was used in 65 patients, especially predominant when requiring higher ESA doses, but by correlation or multivariable analyses, this did not substantially modify the negative association between ERI and EPC.

Conclusions

ERI was independently associated with EPC paucity in a cohort with various comorbidities. This may suggest a link which connects ESA hypo-responsiveness to compromised cardiovascular prognosis of dialysis patients.

Key Words

Dialysis, Anemia, Erythropoietin, Erythropoiesis-stimulating agent, Endothelial progenitor cell

Introduction

Anemia is a frequent complication of chronic kidney disease (CKD) patients, for which diminished erythropoietin (EPO) production in the kidney is for the most part responsible [1]. While development and clinical use of erythropoiesis-stimulating agents (ESA) have remarkably improved treatment of CKD's anemia, it has become apparent that individual responses to ESA are considerably varied. This hypo-responsiveness in a subgroup of patients continues to pose a persistent problem, associated with deteriorated consequences in mortality and cardiovascular risks [2]. Though iron deficiency, inflammation, malnutrition or secondary hyperparathyroidism are among the reported frequent causes for hypo-responsiveness [1,3], the whole pathological mechanism has not been adequately clarified, including its cardiovascular implications, necessitating

further investigation.

Endothelial damage is observed at relatively early stages of atherosclerotic diseases [4]. It has been extensively shown that diagnosis and treatment of endothelial damage have important implications in cardiovascular prognosis [5]. Endothelial progenitor cells (EPC) were originally identified as circulating cells which co-express CD34 and Flk1 (fetal liver kinase 1) which is also known as VEGFR-2 (vascular endothelial growth factor receptor-2) or KDR (kinase insert domain receptor). EPCs are thought to be produced in the bone marrow, and after being released into systemic circulation, to provide endothelial protection [6]. EPCs have been widely shown to be useful as a marker of endothelial function, and to have therapeutic utility in cardiovascular diseases, including in CKD patients [7].

Not much is known as yet on how closely ESA hypo-responsiveness is related to EPC levels, or more broadly, to endothelial function. In this cross-sectional study, we attempted to shed light on this hypothesized connection by investigating the association of EPC numbers with various clinical parameters including ERI, a marker estimating ESA hypo-responsiveness, in chronic hemodialysis (HD) patients.

Methods

Patient recruitment, blood sample acquisition and data collection

Any patient who underwent chronic HD treatment at Blood Purification Center, Dokkyo

Medical University Hospital, from 2018 to 2020 became a study participant candidate.

Study subjects were recruited from both outpatients and inpatients admitted for various

treatment purposes. Those who had been treated with anti-cancer drugs or

hematological transfusions, or who had undergone major surgeries during three weeks

prior to the EPC measurement were excluded. We got informed consents for participation

from study subjects in accordance with procedures determined by our university hospital

ethics committee, which approved this study. Since we used freshly collected blood

sample at the start of dialysis session for flow cytometry measurement, patient

recruitment was done in so far as was made possible on the basis of measurement

practicability, but without any particular biased intention in selection.

Blood sample test results, such as blood cell counts or biochemistry values, which were

measured at the time of EPC sample collection, were mostly used, but in some patients, those on the closest other HD day, and within a week were adopted. Some values, which normally are not measured on a daily basis, such as total cholesterol, triglyceride or intact PTH, were adopted within a month of EPC measurement. Blood sugar levels were measured at the start of dialysis, and included non-fasting values. BNP values were adopted irrespective of the timing of dialysis session within a week from EPC measurement. Blood pressure and pulse rate at the start of dialysis session on the same day of EPC measurement were adopted. As the body weight, dry weight which was actually achieved around the time of EPC measurement was adopted. To delineate the comorbidities profiles of the study subjects which included many requiring admission for treatments, Charlson Comorbidity Index was used [8].

EPC measurement by flow cytometry

Mononuclear cell fraction was separated from 8 mL heparinized whole blood by gradient centrifugation using Lymphoprep (Abbot Diagnostics Technologies AS, Abbott Park, Illinois, USA). Of the 1 mL finally suspended mononuclear cells in PBS, 100 microL was

used for a set of antibodies staining. For staining, 10 microL of each antibody solution was added, mixed well, and incubated for 15 min at room temperature, in darkness. As control, same 100 microL mononuclear cells not stained was used. After the staining, sample was washed, by adding 2 mL PBS, and after gently mixing, cell suspension was centrifuged at 1500 rpm for 5 min. Washing was done twice. After the final centrifugation, sample was mixed with 300 microL PBS, and the suspension was subjected to flow cytometry analysis with the use of BD FACSCalibur (BD, Franklin Lakes, NJ, USA). Antibodies used were anti-Human CD133PE (Miltenybiotec, Bergisch Gladbach, Germany), PE Mouse Anti-Human CD34 (BD, Franklin Lakes, NJ, USA) and Anti-Human VEGFR2/KDR PerCP MAb (R&D Systems, Minneapolis, Minnesota, USA). Percentages of cells which were double positive for CD34/Flk1 or CD133/Flk1 were counted as estimates of circulating EPCs number.

Erythropoietin resistance index (ERI)

ERI was calculated as a marker estimating ESA hypo-responsiveness. Weekly ESA dose (in IU epoetin alpha or beta) per body weight kilogram was divided by hemoglobin (g/dl).

ESA dose which had been administered for the longest during the three weeks up to the EPC measurement was adopted. For dose conversion between ESA types, 1 microgram darbepoetin alpha (DA) was regarded as equivalent to 250 IU epoetin (either alpha or beta) and to 1.2 microgram epoetin beta pegol (continuous erythropoietin receptor activator [CERA]) [9].

Statistics

For comparison between two groups, T test or Mann Whitney's U test was used for parametric or non-parametric variables respectively. Chi-square test was used for comparison of categorical variables. Spearman's analysis was used to examine correlation between two variables, except, Pearson's method was used between log CD133/Flk+ EPC and log CD34/Flk1+ EPC. Multiple linear regression analysis with CD133/Flk+ EPC as the dependent variable was performed by incorporating independent variables by forced entry mode. For comparison of ESA doses among three ESA types, one-way ANOVA followed by Bonferroni correction was performed. For all the statistical analyses, SPSS version 26 (IBM Japan, Tokyo, Japan) was used. *P* values

less than 0.05 were considered statistically significant.

Results

Patient characteristics

Patient characteristics according to high and low CD133/Flk1+ EPC groups are shown in Table 1. Majority (93%) of the study subjects were inpatients, with relatively high median ERI (18.4 [7.2-33.0] IU/week/kg/g/dL) and ESA dose (161.6 [75.0-320.9] IU/week/kg) as a whole. In the high EPC group patients, in comparison to the low EPC group, significantly lower values were seen in age, inpatients ratio and CRP, while higher value was seen in CERA use ratio. Though there were no significant difference in total comorbidities burdens by Charlson Comorbidity Index, the low EPC group tended to include more patients in such categories as coronary artery disease (CAD), peripheral vascular disease (PVD) or congestive heart failure (CHF), but not significantly.

Correlation between log CD133/Flk1+ and log CD34/Flk1+ EPCs.

Representative flow cytometry scatter plots for EPC percentage measurement are shown in Figure 1. Raw data distributions were skewed, but after log-transformation, both CD133/Flk1+ and CD34/Flk1+ EPC showed normal distributions, with Kolmogorov-Smirnov statistic of 0.053 and 0.051 respectively. Histogram of log CD133/Flk1+ EPC distribution is shown in Supplementary Figure 1. Log CD133/Flk1+ EPC and log CD34/Flk1+ EPC were closely correlated with each other ($r=0.842$, $p<0.001$, Figure 2). We used these log-transformed values of EPCs in the following analyses.

Clinical variables correlated with log CD133/Flk1+ EPC and log CD34/Flk1+ EPC.

We then examined factors associated with log CD133/Flk1+ EPC and log CD34/Flk1+ EPC by correlation analysis (Table 2). Significant factors negatively correlated ($p<0.05$) with either log CD133/Flk1+ EPC or log CD34/Flk1+ EPC were age, HD vintage, CRP, pulse rate, ESA dose, ERI and presence of PVD, while significant positively correlated

factors were male sex, systolic blood pressure and CERA use. Log CD133/Flk1+ EPC, in comparison to log CD34/Flk1+ EPC, tended to be more closely correlated with variables such as age, HD vintage, CRP or ERI. Overall, however, both EPCs showed similar tendencies of correlation.

Multivariable association of factors with log CD133/Flk1+ EPC.

Since log CD133/Flk1+ EPCs tended to be more associated with the factors above examined, we further examined this association by multiple linear regression analysis with log CD133/Flk1+ EPC as the dependent variable. As independent variables, we chose, from among the above significantly correlated factors, those which are likely to be the cause rather than the result, excluding those which are mechanistically expected to have duplicity. Accordingly, we incorporated into the model age, male sex, HD vintage, CRP, systolic blood pressure, pulse rate and ERI, by forced entry mode. As the result, age, sex and ERI remained as the only independent predicting factors (Table 3). Residuals of the regression showed normal distribution with Kolmogorov-Smirnov statistic of 0.054, suggesting validity of the inference (Supplementary Figure 2).

Similar tendencies of epoetin and DA in ERI's association with EPC.

We then examined if types of ESA influenced the above associations. ESA dose overall was closely correlated with ERI ($r=0.989$, $p<0.001$), largely because ESA dose is incorporated in the calculation of ERI. Distribution of administered ESA types across the range of ESA doses were uneven, with DA predominantly used when dose requirements were higher (Figure 3). When ESA doses among the three ESA types were compared, only difference between epoetin and DA was significant ($p<0.001$) (Figure 3). We tried to crudely determine, separately for each ESA type, in which direction ESA doses and ERI were associated with log CD133/Flk1+ EPC by Spearman's correlation. As the result, for each of the three ESA types, ESA doses and ERI tended to be negatively associated with EPC, in the same way as the aggregate of all ESAs, though not significantly (Table 4). Also, to confirm this in multivariable context, we added use of epoetin, as opposed to DA and CERA, as one of the variables predicting log CD133/Flk1+ EPC number into the same linear multiple regression model of Table 3, and as the result, ERI's standardized coefficient beta was changed from -0.202 to -0.214, still negatively

associated and significant ($p=0.046$) (Supplementary Table 1). When choice of DA, as opposed to epoetin and CERA, was instead added as one of the variables predicting log CD133/Flk1+ EPC number, ERI's standardized coefficient beta was changed from -0.202 to -0.148, no longer statistically significant ($p=0.168$), but still association was in the negative direction (Supplementary Table 2). These results suggested that though whole ERIs' negative association with EPC may have been more influenced by DA's larger share of contribution especially at higher doses, both epoetin and DA seemed to have similar tendencies in these associations. Since CERA was used in a very small number of patients ($n=5$), its association may be affected by coincidence, and was regarded not worth examining.

No involvement of iron status in EPC levels or EPCs' association with ERI.

As shown in Table 2, values reflecting iron status, namely serum iron, TSAT and ferritin, were not correlated with EPC levels. We further compared subgroups of subjects with or without iron deficiency. Iron deficiency was defined as TSAT <20% and ferritin <100ng/ml, according to 2015 Japanese Society for Dialysis Therapy: Guidelines for

Renal Anemia in Chronic Kidney Disease.[10] As shown in Figure 4, distributions of subjects according to log CD133/Flk1+ EPC levels in subgroup with iron deficiency (n=24) and subgroup without iron deficiency (n=77) were similar by Mann-Whitney U test ($p=0.473$). As shown in Supplementary Table 3, tendencies of correlations between log CD133/Flk1+ EPC with ERI were similarly in negative direction in these subgroups.

Discussion

In this analysis of a relatively small-size cohort of dialysis patients, age, sex and ERI were the independent predicting factors of circulating EPC number. Though involvement of EPO or ESA in endothelial function has been previously demonstrated, our present study may be one of the few, if any, to have shown the association between a marker of ESA hypo-responsiveness and decreased EPC.

Comparison of ERI and ESA dose in terms of overall associations with variables

ESA hypo-responsiveness would be, as a concept, an index reflecting degree of body's inability to react to secreted EPO, just like insulin resistance in diabetes. But in actual clinical situations, increased EPO dose itself in hypo-responsive patients may also further exacerbate the condition. ERI has been generally accepted as a marker which is static in nature but capable of estimating dynamic ESA hypo-responsiveness. Though prognostic utility of ERI is extensively shown [11], it is by definition correlated with ESA dose itself. It may be in reality difficult to distinguish effects of ESA dose from ESA hypo-responsiveness. In our present study, when correlations with various clinical parameters were compared between ESA dose and ERI, patterns were similar, almost identical, though there were slight differences in association levels with variables like albumin (negative correlation), or CRP (positive correlation) (Supplementary Table 4). With such variables, ERI tended to be associated slightly more strongly, than ESA dose was. This may be because systemic conditions like inflammation or malnutrition, in addition to ESA dose itself, contributed to hypo-responsiveness to ESA.

ERI of the study subjects in relation to definition of ESA hypo-responsiveness

“ESA hypo-responsiveness” is a term generally used for patients who require relatively high doses of ESAs to increase or maintain hemoglobin levels. Currently there is no established quantitative definition of ESA hypo-responsiveness. KDIGO (Kidney Disease: Improving Global Outcomes) guidelines define those patients as hypo-responsive whose hemoglobin levels do not increase within the first month of ESA treatment even with appropriate weight-based dosing [12]. In a clinical practice population-based study, approximately 20% of incident ESA users, irrespective of ESA type including originator or biosimilar, were non-responders defined as with less than zero increase in hemoglobin level in comparison to the baseline during the first three months of treatment [13]. With respect to significance of ERI, which is a static, thus estimating marker of dynamic ESA hypo-responsiveness, a study which observed HD patients for a median of 4.6 years suggested baseline ERI of 11.31 as a prognostic hypo-responsiveness threshold predicting combined cardiovascular events and mortality outcome [14]. This may indicate that the median ERI of 18.4 (7.2-33.0) in our present cohort was relatively high, with the majority in the suggested hypo-responsive category.

Endothelial pathophysiology and ESA hypo-responsiveness

The exact mechanism linking ESA hypo-responsiveness to fewer EPCs cannot be determined from our cross-sectional study. Possibility of direct effects of EPO on EPCs has been shown previously, some studies suggesting that EPO may be beneficial by promoting EPC release. EPO has been shown to mobilize EPC release from the bone marrow and to promote neovascularization in mice [15]. Four-week DA administration to stable coronary artery disease patients increased CD34/CD133+ EPCs and improved FMD (flow-mediated dilatation) findings, a marker of endothelial function [16]. Evidence is limited in CKD patients, but one study has reported administration of epoetin, but not of DA, to stage 5 patients either on HD or not yet, increased circulating EPC numbers [17]. More broadly, EPO's role in endothelial function in general is ambiguous. It has been showed that EPO directly improved FMD findings in diabetic rats [18], while there was no correlation between EPO concentrations and endothelial function in healthy humans [19].

In contrast to the above potentially beneficial effects of EPO on endothelium, there is accumulated evidence that EPO causes hypertension, where proposed mechanism are

wide-ranging, including deranged NO regulation, endothelins, sympathoadrenal system, or renin-angiotensin system [20]. Overall, judging from numerous mutually conflicting reports, EPO may exert variable cardiovascular effects, possibly depending on clinical contexts. Though it is only speculative, the role of EPO may be altered depending on the pro-inflammatory conditions of CKD, as in our study subjects, with its harmful aspects against the endothelium or cardiovascular organs being accentuated.

Possible role of bone marrow

Otherwise, concerning bone marrow's involvement, secondary hyperparathyroidism, frequently a complication of CKD and an incitement for ESA resistance, is reported to deleteriously affect bone marrow hematopoiesis [21]. Though importance of PTH was not evident in our study, various likewise adverse mediators under the systemic condition of ESA resistance may affect EPC production in the bone marrow. Recently, it was reported that administration of anti-BMP6 (bone morphogenic protein 6) antibody ameliorated mouse models of CKD's anemia, whereby it suppressed hepcidin and increased ferroportin expression in erythroid precursor cells, accelerating erythroid output [22].

Since bone marrow is also involved in the modulation of EPC abundance, our findings may further highlight a potentially pivotal role of bone marrow in ESA hypo-responsiveness, explaining its cardiovascular implications.

ESA types in associations between hypo-responsiveness and EPC

With respect to differences in mortality risks depending on ESA type, a recent study examined effects of “short-acting” epoetin and “long-acting” DA on mortality in close to 200,000 HD patients, in which DA administration, in comparison to epoetin, deteriorated two-year mortality, including cardiovascular deaths [23]. In our present study, we analyzed possible differences between ESA types in the modulation of ERI’s associations with EPC. Since DA, within its range of approved doses, can be used at more potent doses (up to 180 microgram per week) than equivalent epoetin doses, it was more likely to be chosen in patients requiring high doses. This made the direct comparison between ESA types difficult. However, after dose conversion by generally accepted ratios between ESAs, no substantial difference in effects was seen between epoetin and DA, both of which were associated with CD133/Flk1+ EPCs in the same negative direction as the

aggregate of all three ESAs.

EPC subtypes and associations with ESA hypo-responsiveness

Among several definitions of EPCs, we measured CD133/Flk1+ and CD34/Flk1+ cells. They were closely correlated with each other, and demonstrated similar patterns of association with various clinical parameters. However, CD133/Flk1+ EPCs tended to be more strongly associated with a number of factors, especially those related to conditions of inflammation or malnutrition, such as age, HD vintage, albumin, CRP and ERI. ERI was significantly correlated with CD133/Flk1+ EPC, but not with CD34/Flk1+ EPC. CD133Flk1+ EPC is regarded as an immature, younger type, which may have distinct effects depending on disease conditions [24]. In rheumatoid arthritis patients, coronary calcification was more associated with CD133/Flk1+ EPC than with CD34/Flk1+ EPC, where it was speculated to be due to susceptibility of immature CD133/Flk+ EPC to cytokines alteration caused by this disease [25]. Similar mechanism may play a role in HD patients, who are in general in states of chronic inflammation, and further investigation is warranted for elucidation.

Implications for treatment

This study may have implications for the treatment of CKD's anemia. Since current clinical guidelines in general recommend target Hb levels in HD patients not less than 10 to 11.5 g/dL [12], ESA doses are likely to be increased toward that level in cases of hypo-responsiveness, leading to high ERI values. In such conditions, additional increases ESA doses may no longer be much effective in improving anemia, and may exert detrimental effects, including cardiovascular effects, possibly as a result of ESA hypo-responsiveness. In such cases, as is suggested in clinical guidelines [12], balanced, restrained doses of ESA may be appropriate, even if only hemoglobin levels lower than the recommended targets will be achieved.

Alternatively, treatments to improve systemic conditions which promote hypo-responsiveness to ESA, such as chronic inflammation, malnutrition or frailty may be beneficial. In recent years, oral hypoxia-inducible factor prolyl hydroxylase inhibitors (HIF-PHIs), a new class of drugs which increase endogenous erythropoietin production, have become available for clinical use. HIF-PHIs have been shown to improve anemia

not inferiorly in comparison to ESAs, and are presumed to increase EPO to relatively mild levels which may be closer to physiological condition [26]. HIF-PHIs have also been shown to improve iron utilization by mechanisms including down-regulation of hepcidin [27]. These effects of HIF-PHIs may be beneficial for the correction of conditions of ESA hypo-responsiveness, which may be accompanied by disordered endothelial physiology, as suggested in our present study, warranting further investigation.

EPC in HD patients according to sex

In our multivariable analysis, positive independent association of male sex with log CD133/Flk1+ EPC was observed. However, when log CD133/Flk1+ EPC levels were directly compared between the male and female subjects, males overall showed slightly higher values, but there was no significant statistical difference (data not shown). A recent study has shown that in women without cardiovascular diseases, reduced circulating progenitor cells, including those expressing CD34 or CD133, were observed in comparison to men [28]. This study suggested that EPC scarcity, rather than estrogen deficiency, may be responsible for the fact that lower cardiovascular incidence observed

in reproductive-age women, disappears in postmenopausal or diabetic individuals, for whom hormone replacement therapy is not sufficiently effective. Though our finding in respect to EPC in female HD patients may be in agreement with such observations, it may be difficult to draw definitive conclusion since sex is a categorical variable, and especially with the relatively small sample size, other confounding factors may have influenced.

Limitations

There are several limitations to this study. First, due to the relatively small sample size, the possibility may not be ruled out that the results were due to coincidence, though the results of multivariate analysis, reached after taking relevant clinical parameters into account, may well be reasonable. Second, since the study subjects were predominantly inpatients, some with relatively heavy comorbidities burdens and requiring high ESA doses, the findings may not necessarily be applicable to the conditions of more general, stable HD patients. Nonetheless, with an increasing proportion of patients undergoing long-term HD treatments in recent years, the

conditions of this study's subjects may be one aspect of the general tendency, which could only be elucidated for the very reason of this particular character of our cohort. Third, because of the cross-sectional design of this study, causality cannot be conclusively determined. Decreased EPC number may not be the result of ESA hypo-responsiveness, and some other underlying condition may be more dominant, though with the former as the presumed result, our multiple regression analysis results suggested the latter is likely to be a significant independent predictor among various relevant factors.

In conclusion, this study has revealed independent association of ERI, a marker estimating ESA hypo-responsiveness, with EPC paucity in hemodialysis patients, suggesting the possibility of a close link between ESA hypo-responsiveness and endothelial dysfunction, which seems to have not been demonstrated so far. This finding may be useful for devising appropriate treatment of CKD's anemia, especially taking into account cardiovascular implications.

Supplementary Information

The online version contains supplementary material available at

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Authors' contributions

AI wrote the paper with intellectual input from all authors. All authors have approved the manuscript.

Conception and research design: AI, HS, TI; data acquisition: AI, HS, MF, YM; data analysis and interpretation: AI, HS; statistical analysis: AI, HS; supervision and mentorship: AT, TI.

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Availability of data and materials

The dataset generated and/or analyzed during the current study are available from corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate

This study protocol complied with the Declaration of Helsinki and was approved by the

ethics committee of Dokkyo Medical University (acceptance no. R-20-1J). All the study subjects received an explanation of the procedures and possible risks of this study and gave their informed consent to participate.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interest.

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Figure legends.

Figure 1

title: Representative images of flow cytometry scatter plots showing high EPC and low EPC numbers.

legend: Flow cytometry scatter plots showing high (0.028%) and low (0.352%) numbers of EPC. EPC numbers were counted as percentages of cells which were double positive for CD133 and Flk1, in circulating mononuclear cells.

Figure 2

title: Correlation between log CD133/Flk1+ EPC and log CD34/Flk1+ EPC.

legend: Pearson's correlation analysis between log CD133/Flk1+ and log CD34/Flk1+

EPCs.

Figure 3

title: Distribution of ESA doses according to each ESA type.

legend:

Median and interquartile ranges of ESA doses according to ESA type are shown. One microgram darbepoetin alpha (DA) was regarded as equivalent to 250 IU epoetin and to 1.2 microgram epoetin beta pegol (continuous erythropoietin receptor activator [CERA]).

*p<0.001.

Figure 4

title: Distribution of subjects according to log CD133/Flk1+ EPC in iron status subgroups.

legend: Distributions of subjects according to log CD133/Flk1+ EPC levels in subgroup with iron deficiency (n=24) and subgroup without iron deficiency (n=77) were similar by

Mann-Whitney U test ($p=0.473$). Iron deficiency status was defined as TSAT <20% and ferritin <100ng/ml.

Supplementary Figure 1

title: Normal distribution of log-transformed CD133/Flk1+ EPC numbers in percentages.

legend:

CD133/Flk1+ EPC numbers by percentages were multiplied by 1,000 and log-transformed with base of 10. Though distribution of EPC number in percentages was skewed, after log-transformation, data were normally distributed.

Supplementary Figure 2

title: Normal distribution of standardized residuals in multiple linear regression analysis for variables predicting CD133/Flk-1 EPC.

legend: Histogram distribution of standardized residuals in the multiple linear

regression analysis of Table 3, which analyzed independent variables predicting log

CD133/Flk1+ EPC. Kolmogorov-Smirnov statistic: 0.054.