

## Intravenous administration of iron sucrose for treating anemia in postpartum women

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### Abstract

**Background:** To compare the efficacy of oral and intravenous administration of iron supplements for treating postpartum anemia.

**Methods:** One hundred and four anemic postpartum women were studied prospectively. The criteria for the diagnosis of anemia were Hb <8 gr/dl and ferritin < 10µg/dl. They were randomised into two groups. Group A consisted of 78 women who received i.v. a total amount of 300 mg iron sucrose in three days. Group B consisted of 26 women, who received orally 800 mg iron proteinsuccinylate daily for four weeks.

**Results:** At the end of the study, in group A the increase of Hb mean level was 4.6 gr/dl and of ferritin mean level was 105 mg/L. In group B the increase in hemoglobin mean level was 2.3 gr/dl and ferritin mean level was 68 mg/L. There was significant difference in the increase of hemoglobin level ( $p=0.0001$ ) and also in the increase in ferritin level ( $p=0.0004$ ) between the two groups.

**Conclusion:** Intravenous administration of iron sucrose seems to be safe and it helps postpartum women to recover early from anemia. Hippokratia 2009; 13 (1): 38-40

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Iron deficiency during pregnancy and postpartum could be due to insufficient absorption and to increased needs resulting to chronic iron deficiency and anemia<sup>1</sup>. The human body does not have a mechanism of getting rid of extra iron amount and the mechanism of iron absorption plays a crucial role in iron homeostasis<sup>2</sup>. During pregnancy the needs for iron are increased due to the fetus, the placenta and the increased volume of maternal erythrocytes. Women in the reproductive age frequently have anemia and iron deficiency due to menstrual loss. Frequently these women are already anemic by the time they get pregnant<sup>3,4</sup>.

During the period of pregnancy the extra needs on iron are about 1000mg in total and they should be replaced in order to avoid severe anemia. The usual management is the replacement of iron by oral supplementations. Blood transfusion is the last resort used only in very severe cases of anemia with symptomatic patients<sup>5,6</sup>. The oral therapy though is time consuming and probably not enough in severe cases of anemia. On the other hand blood transfusion although it can promptly and reliably treat anemia, entails a lot of dangers like cross reactions and viral infections. In order to avoid these side effects intravenous administration of iron sucrose could be a convenient and reliable solution.

### Materials and methods

In our prospective study, one hundred and four postpartum women with severe iron deficiency anemia were randomised to receive i.v. or oral iron treatment. Seventy seven of them had a normal vaginal delivery and the rest 27 had a caesarean section. The women were divided randomly in two groups. Demographic data were not matched. Group A consisted of 78 women and group B of 26 women. All the caesarean sections were performed by a consultant obstetrician and were elective. The normal deliveries were all performed by senior midwives. The estimated blood loss was between 500-800 mls for the sections and not significant in all cases of normal deliveries. From the study we excluded cases with a diagnosis of placenta previa, placenta abruption, pre-eclampsia and clotting disorders. All women who participated were lactating postpartum and had amenorrhea during the study period. Also, after laboratory investigations and hematology assistance all other types of anemia hereditary or acquired were excluded from the study. The main criteria in order to administer intravenous iron was hemoglobin levels < 8 gr/L and ferritin levels < 10 µ/L. In group A we used for intravenous infusion 100 mg iron sucrose daily, diluted in NaCl 0.9%, for three days. The administration was performed slowly within two hours, in order to avoid any adverse reactions. Group B was the control group and

we used oral supplementation of iron proteinsuccinylate 800 mg daily for one month. The treatment for group B lasted for 28 days.

One week after the initiation of the treatment we measured hemoglobin, ferritine, SGOT, SGPT blood levels and proteinuria of both groups. The same analysis was performed four weeks later.

For statistical analysis of the results between the two groups we used the unpaired graph-pad t-test.

## Results

Fifty two out of seventy eight women (66%) from the group A attended the outpatient one month later and the rest 26 (34%) failed to attend. In group B only 20 out of 26 (77%) women kept their follow up appointments. One week after the initiation of the treatment in the group A the mean values of hemoglobin and ferritine blood levels were 8.8 gr/dl and 38 µg/l correspondingly. Four weeks later, the mean values of hemoglobin were 12.6 gr/dl and of ferritine 115 µg/l (Table 1). In group B the increase in

group B CI: 9.71-10.88, SD: 0.47, mean Hb increase 2.3 gr/dl) and also in the increase in ferritin levels  $p=0.0004$  between the two groups (group A CI: 101-128, SD: 11.1, mean ferritin increase 105 mg/L and group B CI:66-89, SD: 9, mean ferritin increase 68 mg/L).

## Discussion

Menstrual blood loss, pregnancy and delivery are the main causes of anemia in young women. About 30% of anemic women have iron deficiency anemia with hemoglobin levels below 10 gr/dl, and about 10% of anemic women had hemoglobin levels below 8gr/dl. In about 5% of all deliveries the postpartum hemorrhage is significantly high, more than 1000 mls<sup>8</sup>. In the past, blood transfusion was the only available method in order to treat severe anemic cases. Blood transfusion is a reliable method with excellent results in the treatment of anemia but also with high risk for transmittion of viral infections (HIV, HCV, HBs, CMV) and serious transfusion cross reactions<sup>9</sup>.

**Table 1:** Group A; mean values of blood results before and after intravenous iron treatment.

	Before treatment n=78	One week after treatment n=52	Four weeks after treatment n=52
Hemoglobin mean values (gr/dl)	<8	8.8	12.6
Ferritine (mg/l) mean values	<10	38	115
SGOT,SGPT (mean values)	9	11	14
Urine protein (gr/dl)	0	0.3	0.2

**Table 2:** Group B; mean values of blood results before and after oral iron treatment.

	Before treatment n=26	One week after treatment n=20	4 weeks after treatment n=20
Hemoglobin mean values (gr/dl)	<8	8.1	10.3
Ferritine (mg/l) mean values	<10	19	78
SGOT,SGPT (mean values)	9	12	13
Urine protein (gr/dl)	0.2	0.1	0.2

hemoglobin and ferritine levels were significantly lower. Hemoglobin mean levels were 8.1 gr/dl and 10.3 gr/dl in the first and fourth week respectively. As for ferritin levels the mean values after the first week of treatment were 19 µg/l and after the fourth week 78 µg/l (Table 2). No abnormal blood results have been detected concerning liver function enzymes and there was no proteinuria. The adverse effects from the iron treatment were mild but more prominent in group B mostly constipation and bloating (Table3). There was significant difference in the increase of hemoglobin levels  $p= 0.0001$  (group A CI: 12.04-13.15, SD: 0.44, mean Hb increase 4.6gr/dl and

**Table 3:** Adverse effects of iron treatment.

Adverse effects	Group A	Group B
Headache	1	0
Nausea	1	0
Heartburn	0	2
Hiccup	0	4
Constipation	0	5

Administration of oral iron supplementations is not sufficiently enough in order to reverse anemia promptly, due to the limited absorption, the gastrointestinal symptoms and the poor compliance for long treatment of the patients. Treatments with recombinant human erythropoietin could be useful for cases of chronic inflammation when erythropoiesis is reduced. In cases of iron deficiency anemia the combination of iron supplementations and erythropoietin is not preventing iron loss and is not increasing the endogenous erythropoiesis. On the contrary high iron levels in plasma circulation after simultaneous intravenous administration of iron and erythropoietin, is essential for stimulation of erythropoiesis<sup>10</sup>.

Intravenous iron treatment is indicated for patients with poor compliance in oral supplementations, in cases with poor iron absorption (bowel operations, or diseases), in patients with severe renal impairment, and in postpartum hemorrhage<sup>11,12</sup>. Recent evidence suggest that iron sucrose can be detected in high levels in the liver circulation and marrow within 5 minutes after intravenous administration. The time interval is 5 to 6 hours and the renal metabolism is minimal, less than 5% of the total dose. These data lead to the conclusion that iron sucrose is metabolically available in only a few hours after administration. This way iron is engaged exclusively from the reticulate liver cells, transferrin and apoferritin in the marrow and spleen. Then it is quickly metabolized and it is available for erythropoiesis and inversion of anemia<sup>13</sup>.

In our study after four weeks and with a total dose of i.v. 300 mg of iron sucrose there was complete reversal of anemic status in all women of group A. In group B there was improvement of anemia which was not as significant as in group A, though.

It is already known that intravenous administration of excessive dose of iron might cause liver necrosis, renal, suprarenal and pulmonary damage. The presence of iron sucrose in the plasma circulation is associated with absence of any undesirable effect to the patients. This absence of side effects is partly due to the lower allergic effect of the sucrose complex because of the very slow release of elementary iron from the complex<sup>13,14</sup>. Also the accumulation of iron-sucrose in organic parenchyma is much lower compared to iron-dextran and iron-gluconate<sup>13,14</sup>. In addition, incorporation into the bone marrow for erythropoiesis is faster than other complexes<sup>13,15</sup>. Iron sucrose is quite safe for the liver in daily doses of 100 mg, in comparison with other iron supplementations. Rare anaphylactic reactions because of the use of iron sucrose have been reported in about 0.002% of cases<sup>14</sup>.

## Conclusion

From our study we conclude that i.v. iron sucrose compared to oral iron administration is more efficient in treating postpartum anemia without any toxic effects and with minimal anaphylactic reactions.

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