The Effect of Acute Inflammation on Iron Metabolism in Rats

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UCHIDA, T., IGARASHI, T., SUZUKI, T., KOKUBUN, K., MATSUDA, S. and KARIYONE, S. The Effect of Acute Inflammation on Iron Metabolism in Rats. Tohoku J. exp. Med., 1983, 139 (3), 293-298 — Iron metabolism in rats with acute turpentine-induced inflammation was evaluated. In acute inflammation, reduced plasma iron and total iron-binding capacity values, shortened plasma iron disappearance time and lower plasma iron turnover were observed. The administration of $^{59}$Fe chondroitin ferrous sulfate in order to evaluate the reticuloendothelial (RE) function revealed a significantly increased $^{59}$Fe retention in the liver and lower incorporation into red blood cells. Radioactivity in hepatic RE cells was higher in acute inflammation than in control. These results suggest the possibility of a block in the transfer of iron from RE cells to the plasma iron pool during acute inflammation.

Materials and Methods

Experimental animals. Male rats of the Wistar strain, between 8 and 12 weeks of age, and weighing from 150 to 250 g, were used in all experiments. Sterile inflammation was induced by the injection of 0.25 ml of turpentine oil into the muscle of one thigh during ether anesthesia.

RE and parenchymal iron labeling method. Two kinds of radioactive substances were employed for labeling hepatic RE and parenchymal cells (Cook 1972; Hershko 1973).

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Radioiron labeling of hepatic RE cells was performed by injecting $^{59}$Fe chondroitin ferrous sulfate ($^{59}$Fe-Blutal, RE tag, SA 30.7 mCi per mg Fe, 3.3 $\mu$g Fe per ml, Dainabot Radioisotope Laboratory, Suita, Osaka) i.v. in a dose of 100 $\mu$g of elemental iron.

**Determination of serum iron, TIBC and ferritin level.** Blood removed by exsanguination under ether anesthesia was centrifuged at 1,000 $\times$ g for 20 min. The iron content and TIBC of the supernatant serum was measured by the method of International Committee for Standardization in Hematology (1978a, b). Serum ferritin concentration and ferritin content in hepatic ferritin fraction were measured by the two sites radioimmunometric assay using antibody coupled paper discs as a solid phase (Niitsu et al. 1975).

**Ferrokinetics in rats.** Ferrokinetics in turpentine-induced inflamed rats were investigated. Radioactive iron (1 $\mu$Ci of $^{59}$FeCl$_3$ dissolved in 0.5 ml saline) was injected intravenously. Samples of tail vein blood were obtained 5, 15, 30, 60 and 120 min after the injection with 50 $\mu$l disposable pipettes. Each sample was immediately transferred to a counting tube and radioactivity was determined in a well-type gamma spectrometer. Half-time clearance of $^{59}$Fe was determined by least squared regression analysis. The blood volume was estimated from the factor 5.75 ml per 100 g of body weight (Belcher and Harris 1957). Serum iron levels were determined by sacrificing the animals just after the study. Plasma iron turnover (PIT) was calculated as described by Cook et al. (1970) and was expressed as mg Fe per kg per day.

**Iron release from the RE cells in turpentine-induced inflammation rats.** Iron release from the RE cells was determined after the injection of 1 $\mu$Ci of $^{59}$Fe chondroitin ferrous sulfate in control, turpentine-induced inflammation and transfused rats. The transfused rats were prepared by the intravenous injection of 1 ml whole blood for 4 days. Samples of tail vein blood were withdrawn 1 hr after the injection and then daily for the next 8 days. The radioactivity of each sample was plotted as the percent of counts per min to the radioactivity injected.

**Measurement of tissue radioiron.** In this series of experiments, RE tag was performed in control, turpentine-induced inflammation and RE overloaded rats. Inflamed rats were made by the intramuscular injection of 0.25 ml turpentine oil for 4 days. RE overload was prepared by the injection of 3 mg unlabeled chondroitin ferrous sulfate followed by $^{59}$Fe chondroitin sulfate administration. The animals were sacrificed 1, 3, 7 and 10 days after the injection of $^{59}$Fe-tagged compounds. Radioactivity and iron content of the total liver, ferritin and hemosiderin fraction were measured as follows: Liver uptake was determined from the activity in weighed aliquots of tissue related to total organ weight. Ferritin and hemosiderin fraction was separated by the method of Linder and Munoro (1972), in which liver was homogenized in 9 volumes of saline and this homogenate was heated at 70°C for 10 min, followed by cooling and centrifugation. Sediment was kept as hemosiderin-rich fraction and supernatant as ferritin fraction for the counting of the radioactivity. Iron content of liver homogenate, ferritin and hemosiderin fraction was measured by the method of Barry and Sherlock (1971). Ferritin contents in hepatic ferritin fraction were measured by the two sites radioimmunometric assay.

**Separation of hepatic RE cells.** Suspensions of hepatic RE cells were prepared by a modification of the method of Cook et al. (1972), in which hepatic parenchymal cells are selectively destroyed by the proteolytic enzyme, pronase (Kaken-Kagaku Ltd., Tokyo). The liver, cut into 2 to 5 mm particles, was suspended in 35 ml pronase solution, freshly prepared by dissolving 250 mg pronase in 100 ml balanced salt solution (BSS). This suspension was shaken in a water bath at 37°C for 30 min and the supernatant collected. After the sedimentation at 200 $\times$ g for 10 min, the sediment was resuspended in 20 ml BSS, followed by the centrifugation at 100 $\times$ g for 5 min. The most of erythrocytes and Kupffer cells were sedimented and the residual erythrocytes were lysed by exposure to distilled water for 25 sec. Final centrifugation was performed at 200 $\times$ g for 5 min and hepatic RE cells were dried in air. The radioactivity was counted and expressed cpm per mg weight of RE cells.
Statistical analyses of data. Values are given in terms of mean±s.e. Differences between mean values were analyzed by Student's t test and taken to be significant when p values were less than 0.05.

RESULTS

Serum iron, TIBC, ferritin concentration and ferrokinetic indices in control and turpentine-induced inflammation rats are shown in Table 1. A low serum iron and TIBC, shortened PID T'_{1/2} and low PIT were evident (p<0.01). Serum ferritin concentrations were higher in inflamed rats than in control, but there was no significant difference.

Iron release from the RE cells in turpentine-induced inflammation rats was evaluated by $^{59}$Fe red blood cell incorporation after the injection of $^{59}$Fe chondroitin ferrous sulfate. There were no significant differences of $^{59}$Fe red blood cell incorporation in control, inflammation (intramuscular injection of 0.25 ml turpentine oil) and blood transfusion groups (Fig. 1).

In other experiments, inflamed rats (four times injections of turpentine-oil 0.25 ml each) and RE iron overloaded rats (injection of unlabeled chondroitin ferrous sulfate 3 mg) were used. $^{59}$Fe incorporation in red blood cells influenced by

| Table 1. Effect of inflammation on the iron metabolism |
|----------------------------------|---------------|
|                                   | Control       | Inflammation |
| Serum iron (µg/100 ml)           | 225±12        | 52±8*         |
| TIBC (µg/100 ml)                 | 300±20        | 148±14*       |
| Ferritin (mg/ml)                 | 83±19         | 114±41†       |
| PID T'_{1/2} (min)               | 195±40        | 120±14*       |
| PIT (mg/kg/day)                  | 0.57±0.08     | 0.16±0.01*    |

n=4. Mean±s.e. * p<0.01, † Not significant.

Fig. 1. $^{59}$Fe red blood cell incorporation after the administration of radioactive chondroitin ferrous sulfate in turpentine-induced inflammation and transfused rats. There were no significant differences of $^{59}$Fe incorporation between them. ○—○, transfusion; △—△, inflammation; ●—●, control.
the release of iron from RE cells was depressed in turpentine-induced inflammation and RE iron overloaded rats (Fig 2). Total liver radioactivity decreased slowly in the inflammation and RE iron overload corresponding with the lower red blood cell utilization. The ratio of ferritin to hemosiderin fraction in $^{59}$Fe activity and in iron content is shown in Fig. 3. The ratio of hemosiderin fraction to total liver was constant in normal and inflammation, but tended to be higher in RE overload. The same phenomena were observed in total liver iron and hemosiderin iron content.

Fig. 2. The ratioactivity of total liver (column) and of red blood cells (solid line) after the injection of $^{59}$Fe chondroitin ferrous sulfate (RE tag) in acute inflammation and RE iron overload. Total liver radioactivity decreased slowly both in inflammation and in RE overload corresponding with the lower red blood cell utilization.

Fig. 3. The radioactivity of hemosiderin (shaded area) and ferritin fraction (unshaded area) after the injection of $^{59}$Fe chondrotin ferrous sulfate. The ratio of hemosiderin fraction to total liver was constant in normal and inflammation, but tended to be higher in RE overload. The same phenomena were observed in total liver iron (○) and hemosiderin iron content (×).
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The radioactivity of hepatic RE cells after the injection of $^{59}$Fe chondroitin ferrous sulfate is shown in Fig. 4. The radioactivity per mg of hepatic RE cells was higher in RE iron overload and turpentine-induced inflammation rats than in control, which suggests that restricted iron supply in inflammation is due to a block in iron release from RE cells.

**DISCUSSION**

The pattern of alteration of iron metabolism in inflammation consists of hypoferremia, a decrease in transferrin, reduced transferrin saturation and normal or increased marrow RE iron (Cartwright 1966; O'Shea et al. 1973). Reutilization of iron from destroyed red cells is impaired due to retention of iron in RE cells (Freireich et al. 1957; Haurani et al. 1965). The average half time of plasma $^{59}$Fe disappearance in the inflammation was 120 min as compared with 195 min in the control. The average PIT was 0.57 mg/kg/day as compared with 0.16 mg/kg/day of control. It is apparent that there is marked inhibition of erythropoiesis in turpentine-induced inflammation rats (Bush et al. 1956; Zucker et al. 1974).

When $^{59}$Fe labeled chondroitin ferrous sulfate was used, a significantly increased retention of radioactivity was observed in the liver during inflammation, which was found to be pronounced in hepatic RE cells by the selective separation of RE cells. $^{59}$Fe incorporation to red blood cells was lower in inflammation than in control group. Fillet et al. (1974) reported the same results using a $^{59}$Fe labeled heat denaturated red blood cells. These findings support the possibility of a block in the transfer of iron from hepatic RE cells to the plasma iron pool during acute inflammation.
References