Notizen 1023

Correlation between Diagnostic 59Fe2+-Absorption and Serum Ferritin Concentration in Man

H. C. Heinrich, J. Brüggemann, E. E. Gabbe, and M. Gläser

with the technical assistance of

Fatima Icagic and Ellen Pape

Division of Medical Biochemistry, Institute of Physiological Chemistry, University Hospital Eppendorf, University of Hamburg

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Iron Absorption, Serum Ferritin, Iron Stores, Iron Deficiency

A high correlation coefficient r = -0.832 ($P_{r \pm 0} \le 0.0001$) was estimated in man for the inverse relationship between the diagnostic ⁵⁹Fe²⁺-absorption and the serum ferritin concentration which is very close to the correlation r = -0.88as described for the relationship between the diagnostic ⁵⁹Fe²⁺-absorption and the diffuse cytoplasmic storage iron in the bone marrow macrophages. The increase of the diagnostic 59Fe2+-absorption seems to be an earlier and more sensitive indicator of depleted iron stores whereas the serum ferritin decreases somewhat later during the development of iron deficiency.

Depleted iron stores are characterized and can be diagnosed by the increased 59Fe-absorption from a small 10 μ mol (= 0.56 mg) ⁵⁹Fe²⁺-dose ^{1, 2, 4} since a correlation coefficient of r = -0.88 was estimated for the inverse relationship between intestinal ⁵⁹Fe²⁺-absorption and the "Berlin-Blue"-reactive diffuse cytoplasmatic non-haeme storage iron in the bone marrow reticuloendothelial cells 3, 4. The traditional invasive method of Berlin-Blue staining the bone marrow for assessing body iron stores is however refused by many patients and has the disadvantage of being at the best a semiquantitative and subjective method with a limited reproducibility and capacity which requires some experience for a relevant evaluation by the observer. In order to investigate the possibility of substituting this classical method of assessing body iron stores by the recently introduced quantitative and reproducible immunoradiometric assay for serum ferritin 5, 6 we have studied the correlation between the diagnostic $^{59}\mathrm{Fe^{2^+}}\text{-}absorption$ and the serum ferritin concentration in subjects with well defined stages of iron stores or iron deficiency 2-4.

The 59 Fe-absorption from a diagnostic $10~\mu mol$ (= $0.56 \,\mathrm{mg}$) $^{59}\mathrm{Fe^{2+}}$ -dose was calculated from the whole body retention of absorbed ⁵⁹Fe which was

Requests for reprints should be sent to Prof. Dr. H. C. Heinrich, Universitäts-Krankenhaus Eppendorf, Physiologisch Chemisches Institut, Medizinische Biochemie, Martinistr. 52, D-2000 Hamburg 20.

Table I. Diagnostic 59 Fe ²⁺ absorption (in $^{\%}$) and serum ferritin concentration (in 6 ml) in subjects with normal Fe-stores, prelatent, latent and manifest in deficiency. ($n = \text{number of subjects}$, $\overline{X}_g = \text{geometric mean}$, $C_{\text{S.D.}}$ and $C_{\text{S.E.}} = \text{coefficients of standard deviation and standard error.)}$	²⁺ absor _j r of sub	ption $(in \%)$ bjects, $\overline{X}_g =$	and seri geometr	ım ferri ic mean	tin concent	tration 1 Cs.E.	(in ng/ml) = coefficie	in subje nts of st	cts with andard	n normal F deviation a	e-store	and serum ferritin concentration (in ng/ml) in subjects with normal Fe-stores, prelatent, geometric mean, Cs.D. and Cs.E. = coefficients of standard deviation and standard error.)	latent a	ınd maı	nifest in
Subjects with	и	Range	Totals $\overline{X}_{\mathbf{g}}$	Cs.D. Cs.E.	Cs.E.	и	Range	Males $\overline{X}_{m{g}}$	Cs.D. Cs.E.	Cs.E.	и	Range	Females $\overline{X}_{\mathbf{g}}$		Cs.D. Cs.E.
normal Fe-stores ⁵⁹ Fe ²⁺ -absorption serum ferritin	79	10 - 53 27 -221	29	1.45	1.05	29	10 - 47 28 -221	26 106	1.47	1.14	38	16 - 53 27 -185	32 69	1.38	1.05 1.08
prelatent Fe-deficiency ⁵⁹ Fe-absorption serum ferritin	31	$\begin{array}{c} 52 & -100 \\ 7.8 - & 64 \end{array}$	76 27	1.20	1.03	4 4	55 - 100 $16 - 30$	77 24	1.30 1.34	1.14	27 27	52 - 96 $7.8 - 64$	76 27	1.19	$\frac{1.03}{1.12}$
latent Fe-deficiency 59Fe-absorption serum ferritin	15	$63 - 100 \\ 5.3 - 32$	89 14	1.17	1.04 1.13	44	$96 - 97 \\ 5.3 - 14$	96 9.6	1.00	1.00	===	$\frac{63}{9.5} - 100$	85 16	1.19	1.06
manifest Fe-deficiency 59Fe-absorption serum ferritin	45 45	63 - 100 $2.7 - 12$	87 6.0	1.17	1.02 1.05	111	$^{71}_{2.7-12}$	91 5.1	1.13	1.04	34 34	63 - 100 $3.4 - 10$	86 6.3	1.17	1.03



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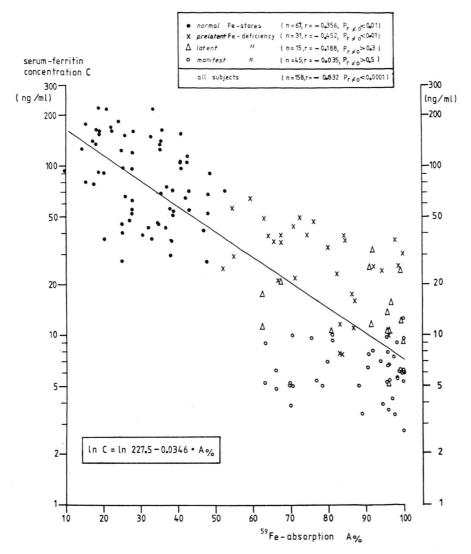


Fig. 1.
Correlation between diagnostic ⁵⁹Fe²⁺-absorption and serum ferritin concentration.

measured in starved subjects after 14 days within the 4 π -counting geometry of a whole body radioactivity detector with liquid organic scintillator 7. Serum ferritin was estimated with an immunoradiometric assay using a modification 8 of the two-site or sandwich solid phase system 6. According to their individual diagnostic $^{59}\text{Fe}^{2+}$ -absorption, serum iron, total iron binding capacity and haemoglobin concentration the subjects were subdivided into 4 groups with normal iron stores (diagnostic $^{59}\text{Fe}^{2+}$ -absorption normal: 10-53%; all other parameters normal), prelatent iron deficiency (diagnostic $^{59}\text{Fe}^{2+}$ -absorption increased to 52-100%, $\overline{X}_g=76\%$; all other parameters within the normal range), latent iron deficiency (in addition to increased

diagnostic $^{59}{\rm Fe^{2^+}}\text{-}absorption}$ serum iron reduced below $60~\mu{\rm g}/100~{\rm ml}$ and the total iron binding capacity increased above $400~\mu{\rm g}/100~{\rm ml};$ transferrin-Fe-saturation $<\!20\%$) and manifest iron deficiency (haemoglobin concentration $<\!12~{\rm g}/100~{\rm ml},$ Hb $_{\rm E}$ $<\!27~{\rm pg/erythrocyte},$ transferrin-Fe-saturation $<\!15\%$).

The inverse relationship between the diagnostic $^{59}\text{Fe}^{2+}$ -absorption A_{N_0} and the serum ferritin concentration C is described by the regression line $\ln C = \ln 227.5 - 0.0346 \cdot A_{\text{N}_0}$ (Fig. 1). A high correlation coefficient r = -0.832 ($P_{\text{r}\pm 0} < 0.0001$) was estimated for this inverse relationship if the data of all the 158 patients were considered. This value is very close to the correlation of r = -0.88

Notizen 1025

as described earlier for the inverse relationship between the diagnostic $^{59}\text{Fe}^{2^+}$ -absorption and the diffuse cytoplasmatic storage iron in the bone marrow RE-cells 3,4 . The negative correlation between the diagnostic ^{59}Fe -absorption and the serum ferritin concentration was also still significant for the group of 67 subjects with normal iron stores $(r=-0.356; 0.01>P_{r})>0.001$ and the group of 31 subjects with prelatent iron deficiency $(r=-0.452, 0.01>P_{r})>0.001$. No significant correlation, however, was observed within the group of 15 subjects with latent iron deficiency $(r=-0.188, P_{r})>0.3$ and within the group of 45 subjects with manifest iron deficiency $(r=-0.035, P_{r})>0.5$.

Lower correlation coefficients of -0.58^9 and -0.398 10 have been described for the inverse relationship between serum ferritin and ⁵⁹Fe-absorption from larger amounts of 2-3 mg iron which were added to a test meal and administered to normal subjects with non-defined iron stores. Because of the existing dose relationship of intestinal iron absorption in relation to iron stores smaller doses of ⁵⁹Fe (e.g. 0.56 mg Fe²⁺) are more suitable for a non-overlapping separation between subjects with normal and depleted iron stores than larger doses of several mg Fe 11. Many food components do furthermore interfere with the diagnostic reliability of intestinal ⁵⁹Fe²⁺-absorption so that a higher and more reliable correlation has to be expected between serum ferritin and the absorption from a small diagnostic dose of only 0.56 mg Fe²⁺ in starved subjects.

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It seems that an increase of the diagnostic ⁵⁹Fe²⁺absorption is an earlier and more sensitive indicator of iron stores depletion and that the serum ferritin levels decrease somewhat later. This explains that more than half of the serum ferritin levels in subjects with prelatent iron deficiency (total range 7.8 -64; $\overline{X}_{\rm g}=27$ ng/ml, $C_{\rm S.D.}=1.77$, Table I) are still within the lower normal range of 27-64 ng/ml whereas the other half is below the lower normal border line of 27 ng/ml and within the range of 7.8 and 27 ng/ml (Fig. 1 and Table I). The depleted iron stores in prelatent iron deficiency which are characterized by the high negative correlation (r =-0.88) between the increase of diagnostic ⁵⁹Feabsorption and the diminution of the diffuse cytoplasmatic storage iron in bone marrow RE-cells 2-4 therefore cannot be diagnosed only on the basis of reduced serum ferritin concentrations. Serum ferritin levels between 27 and 64 ng/ml are only suspect of being caused by depleted iron stores whereas serum ferritin levels below 27 ng/ml do definitely indicate the consumption of storage iron in subjects with prelatent, latent or manifest iron deficiency. If a serum ferritin rise due to increased release in infection or inflammation can be excluded by the simultaneous estimation of normal ervthrocyte sedimentation rate, leucocyte count and body temperature serum ferritin levels above 64 ng/ml do exclude exhausted iron stores or prelatent iron deficiency.

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