Effects of protein and vitamin E on haemoglobin formation in rats

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1. Young rats were given, for 9 weeks, vitamin E-deficient diets containing either 20% or 10% casein, with and without a dietary supplement of 350 ppm D-a-tocopheryl acetate. For the next 5 weeks the casein content of the low-protein diets was decreased to 7%.

2. The low-protein diets induced severe growth depression.

3. The dialuric acid-induced haemolysis test showed that the rats given the 20% casein vitamin E-deficient diet were depleted of vitamin E, but that the rate of depletion on the low-casein diet was slower.

4. Haemoglobin levels were slightly decreased by the 10% casein diets after 9 weeks, but this difference was not found after 14 weeks, comparing 20% and 7% casein. Dietary vitamin E had no effect on haemoglobin levels or erythrocyte counts.

Bencze, Gerlóczy, Ugrai & Kneiszl (1966) recently reported that vitamin E raised the very low haemoglobin levels of rats given a vitamin E-deficient low-protein diet. The erythrocyte count of these rats was unaffected by protein depletion or vitamin E deficiency or both. The vitamin E-deficient diet used by these authors contained 9% protein, consisting of casein, meat meal and yeast, all previously extracted with acetone or ether, and also an unspecified fat that had been heated to destroy vitamin E. Although they were able to show that vitamin E completely prevented the anaemia they did not investigate the effect of a higher level of the extracted protein. Their 18% protein diet contained only non-extracted proteins.

In view of the importance of the connexion between vitamin E and blood formation and because of the uncertainties in the experimental design of Bencze et al. (1966), we have studied the effect of vitamin E on haemoglobin levels in rats given various dietary concentrations of purified casein.

Experimental

Rats and diets. Forty-one 6-week-old Norwegian hooded rats that had previously received a vitamin E-deficient diet were allocated at random into four groups, with equal distribution of sexes and litter-mates. The groups were given, respectively, a 20% casein diet, that diet with 350 ppm D-a-tocopheryl acetate, a 10% casein diet and that diet with 350 ppm D-a-tocopheryl acetate. The 20% casein diet had the percentage composition: casein ('low vitamin content'; Genatosan Ltd) 20, salt mixture (Green, Diplock, Bunyan, McHale & Muthy, 1967) 4.93, vitamin mixture (Bunyan, Green, Diplock & Robinson, 1967) 0.4, lard 5, sucrose 49.2 and glucose 20.47. The 10% casein diet had the percentage composition: casein ('low vitamin content'; Genatosan Ltd) 10, salt mixture (Green et al. 1967) 4.94, NaH2PO4·2H2O 0.4, vitamin mixture

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(Bunyan et al. 1967) 0.4, lard 5, sucrose 59.2 and glucose 20.07. As described below, the casein content of the latter diet was decreased during the experiment to 7%, with the further addition of 0.54% NaH$_2$PO$_4$.2H$_2$O and 2.46% glucose.

**Blood tests.** Blood samples were obtained from the tips of the rats’ tails. Erythrocytes were counted in the usual way and haemoglobin was estimated by the cyanohaematin method of King & Gilchrist (1947). Dialuric acid-induced haemolysis was measured as described by Bunyan, Green, Edwin & Diplock (1960).

**RESULTS**

At 15 weeks of age the haemoglobin of rats given the 10% casein diets was significantly less than that of rats given the 20% casein diets to the extent of about 0.8 g/100 ml blood (Table 1). Vitamin E had no effect on either haemoglobin level. The rats given vitamin E were fully protected against dialuric acid-induced haemolysis, as expected, and those given the vitamin E-deficient 20% casein diet showed a high degree of haemolysis. However, the rats given the vitamin E-deficient 10% casein diet did not all show haemolysis and so were apparently not all depleted of vitamin E.

For the next 5 weeks the casein content of the low-protein diet was decreased from 10% to 7% in an attempt to produce a more severe anaemia, against which vitamin E might have an effect. It was then found that neither vitamin E nor protein concentration affected haemoglobin levels and erythrocyte counts (Table 1). The haemolysis test showed that the rats given the low-protein vitamin E-deficient diet were then more uniformly depleted than before, but still to a lesser extent than rats given the 20% casein vitamin E-deficient diet. As expected, the low-protein diets decreased weight gain during the experiment.

**DISCUSSION**

Many studies have implicated vitamin E in the formation of red blood cells and in their subsequent integrity and survival. Leucocytosis, anaemia and impaired erythrocyte survival have been found in vitamin E-deficient monkeys (Dinning, 1963). A vitamin E-responsive macrocytic anaemia has been found in children (Majaj, Dinning, Azzam & Darby, 1963) and two recent reports (Binder, Herting, Hurst, Finch & Spiro, 1965; Leonard & Losowsky, 1967) have indicated diminished survival of erythrocytes in adults with vitamin E deficiency due to malabsorption. Vitamin E has also been found to increase capillary resistance in children with purpura (Gerlóczy, Lancos & Szabo, 1966). Haemolytic anaemia occurs in premature infants with low vitamin E reserves (Oski & Barness, 1967) and in alloxan-treated vitamin E-deficient rats (György & Rose, 1948).

Apart from the work of Bencze et al. (1966), several other reports have suggested that the action of vitamin E on blood formation may also be involved with the amount and nature of the dietary protein. Whitaker, Fort, Vimokesant & Dinning (1967) found that the anaemia of Thailand children suffering from protein-calorie malnutrition was cured by a combination of vitamin E and protein, but not by protein alone. In the
Table 1. Effects of dietary protein and vitamin E on haemoglobin concentrations

(6-week-old rats were given the vitamin E-deficient diets containing 20% or 10% casein (see p. 751), with and without a supplement of 350 ppm d-α-tocopheryl acetate. When the rats were 15 weeks old the casein content of the lower-protein diet was decreased to 7%. Results are given as means with standard deviations, except for haemolysis values at 15 weeks of age, where approximate means and the number of analyses are shown.)

<table>
<thead>
<tr>
<th>Dietary casein (%)</th>
<th>Dietary vitamin E (ppm)</th>
<th>Haemoglobin (g/100 ml)*</th>
<th>Erythrocyte count (millions/mm³)*</th>
<th>Dialuric acid-induced haemolysis (%)</th>
<th>Haemoglobin (g/100 ml)†</th>
<th>Erythrocyte count (millions/mm³)†</th>
<th>Dialuric acid-induced haemolysis (%)†</th>
<th>Weight gain (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>None</td>
<td>15.7 ± 0.64 (7)</td>
<td>c. 80 (10)</td>
<td>14.9 ± 1.0 (5)</td>
<td>9.6 ± 1.2 (2)</td>
<td>8.2 ± 0.3 (5)</td>
<td>166 ± 22 (6)</td>
<td>Males: 97 ± 5 (4)</td>
</tr>
<tr>
<td>20 (later, 7)†</td>
<td>350</td>
<td>15.2 ± 0.8 (7)</td>
<td>c. 5 (6)</td>
<td>15.2 ± 0.9 (5)</td>
<td>9.9 ± 0.9 (3)</td>
<td>0 ± 0 (5)</td>
<td>173 ± 28 (5)</td>
<td>Females: 104 ± 16 (5)</td>
</tr>
<tr>
<td>10 (later, 7)†</td>
<td>None</td>
<td>14.4 ± 1.5 (8)</td>
<td>c. 80 (5); c. 5 (4)</td>
<td>14.3 ± 1.6 (5)</td>
<td>10.5 ± 0.4 (3)</td>
<td>608 ± 12.8 (5)</td>
<td>73 ± 34 (5)‖</td>
<td></td>
</tr>
<tr>
<td></td>
<td>350</td>
<td>14.8 ± 1.1 (9)†</td>
<td>c. 5 (10)</td>
<td>14.6 ± 1.5 (9)</td>
<td>10.6 ± 0.6 (3)</td>
<td>0 ± 0 (5)</td>
<td>107 ± 33 (7)†</td>
<td></td>
</tr>
</tbody>
</table>

* No. of rats shown in parentheses.
† No. of tests, each on combined blood samples from one male and one female rat, shown in parentheses.
‡ Decreased from 10% to 7%, when the rats were 15 weeks old.
§ Results for 10% casein diets were significantly lower than the others (P < 0.05).

‖ Significantly lower than the result for the corresponding 20% casein diet (P < 0.01).
* Significantly lower than the result for the corresponding 20% casein diet (P < 0.05).
experiments of Peragello & Fiori (1938–9), vitamin E accelerated the post-haemorrhagic regeneration of haemoglobin in dogs given a low-protein diet, but Jalavisto (1954) did not find such an effect in rabbits. Moore (1949) observed signs of anaemia in a small group of rats each given only 0.2 g casein and 1 g yeast per day; a test on one rat showed leucocytosis and a very low erythrocyte count, but the haemoglobin level was reported to be normal. Anaemia was not observed in the rats receiving vitamin E or adequate casein. More recently, Aterman & Norkin (1963) reported lowered haemoglobin levels in rats given an (unspecified) yeast diet. Both vitamin E and Se prevented the liver necrosis induced by the diet, but only vitamin E prevented the anaemia.

Lazier & Beveridge (1964) found that the haemoglobin of rats given a torula yeast diet was depressed 12–20% below that of rats given the diet supplemented with a combination of methionine, cystine, Se and vitamin E. Rats given supplements of either methionine or Se or vitamin E had 10% less haemoglobin than fully supplemented controls. However, anaemia is not usually found in vitamin E-deficient rats given a diet of normal protein content. Thus, Dinning, Young, Simmons & Day (1954) found no change in the erythrocyte count or haemoglobin level of rats made deficient of vitamin E or pyridoxine or both over a period of 15 weeks, although there was a considerable rise in the neutrophil count. Tests on 9-month-old vitamin E-deficient and vitamin E-supplemented rats from our own colony showed 15.2 ± 1.9 and 14.2 ± 1.1 g haemoglobin/100 ml blood, respectively. Both diets contained 15% protein as casein.

We have been unable to confirm the finding by Bencze et al. (1966) of a 70% loss of haemoglobin in vitamin E-deficient rats given a low-protein diet or any effect of vitamin E on haemoglobin levels. We used diets containing lard and purified casein and adequate supplements of the B-vitamins and vitamins A, D₃ and K. However, haemoglobin levels were not depressed by more than 2–8%, even though the rats were clearly depleted of protein, as evidenced by their poor weight gains, and of vitamin E, as evidenced by the positive haemolysis test. The severe anaemia produced in the experiments of Bencze et al. (1966) may have been caused by several factors other than depletion of protein and vitamin E. Toxic substances may have been produced by the solvent extraction of the protein component of the diet and also by heating the fat component. If a secondary B-vitamin deficiency was involved it is unlikely that it was pyridoxine deficiency, in the light of the work of Dinning et al. (1954) referred to earlier.

It would be of great interest to discover the true cause of the anaemia induced by Bencze et al. (1966) in view of the complete activity of vitamin E in preventing it.

REFERENCES

Protein and vitamin E effects on haemoglobin


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