

Relation between some folate-dependent metabolic pathways and dietary folate content in chicks

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Responses of several folate-metabolizing pathways to dietary folic acid were studied in 2-week-old chicks. Oxidation of a histidine load to carbon dioxide was impaired in folate-deficient chicks. There was a curvilinear relation between oxidation and dietary folate, and maximum oxidation occurred with 2 mg supplemental folic acid/kg. Hepatic activities of glutamic acid formiminotransferase (*EC* 2.1.2.5) and glycine *N*-methyltransferase (*EC* 2.1.1.20) were not affected significantly ($P > 0.05$) by dietary folic acid. The activity of dihydrofolate reductase (*EC* 1.5.1.3) in erythrocytes was elevated in folate-deficient chicks. These studies show that the activities of two folate-dependent pathways can be used as biochemical criteria of folate status in chicks.

Dietary folate: Folate-dependent metabolic pathways: Chicken.

Specific metabolic criteria of status can be very helpful for assessing requirements for individual vitamins in poultry. For instance, pyruvate carboxylase (*EC* 6.4.1.1) and glutathione reductase (*EC* 1.6.4.2) have provided useful information on the status and requirements of young chickens or turkeys for biotin (Whitehead & Bannister, 1978) and riboflavin (Lee, 1982) respectively. However, comparable criteria are lacking in the case of folic acid.

Previous studies (Maxwell *et al.* 1988) have identified haematological abnormalities in folate-deficient chicks and found that the dietary folate levels needed to prevent some of these abnormalities were higher than current estimates of chick folate requirements (National Research Council, 1984). However, these haematological abnormalities may not have been completely specific for folate status. The studies also established relations between blood and dietary folate levels.

Metabolic pathways directly dependent on folate might provide more specific information on folate status. Burns & Jackson (1976) have reported altered activities of some folate-metabolizing enzymes in livers from folate-deficient chicks. The present report describes the results of further investigations on the effect of folate deficiency on several folate-dependent pathways in chicks and the response of these pathways to graded dietary supplements of folate.

MATERIALS AND METHODS

Birds and husbandry

Male broiler chicks (1-d-old; D. B. Marshall Ltd, Newbridge, Midlothian) were allocated at random to five dietary treatments. The basal folate-deficient diet was similar to the low vitamin casein – gelatin – wheat-starch diet A previously used by Maxwell *et al.* (1988) with the addition (/kg diet) of L-arginine (2.7 g), L-tryptophan (130 mg) and L-cystine (520 mg) to improve its amino acid balance. Four other diets were derived from the basal diet by supplementation with 1, 1.5, 2 or 4 mg folic acid/kg using a commercial preparation containing 800 g folic acid/kg on a dextrin support (Rovimix Folic 80 SD; Hoffmann-La Roche, Basle). The level of 4 mg/kg was considered to be an adequate control in relation

to the estimated requirement of 0.55 mg/kg (National Research Council, 1984). The chicks were housed in compartments of wire-floored electrically heated tier brooders and had unrestricted access to food and water. They were reared in several batches until approximately 14 d of age when they were used in the following studies.

Reagents

Radiochemicals were obtained from Amersham International plc, Amersham, Bucks. The specific activities of L-[1-¹⁴C]histidine, S-adenosyl-L-[methyl-¹⁴C]methionine and [3',5',7',9'-³H]pteroylmonoglutamic acid were 56 mCi/mmol, 0.5 mCi/mmol and 51 Ci/mmol respectively. All other reagents were of analytical or scintillation grade.

Histidine oxidation

These experiments were performed using a metabolism chamber and carbon dioxide-trapping system of ethanolamine: 2-methoxyethanol described by Saunderson & Whitehead (1987). In a preliminary experiment to compare rates of oxidation, one bird from each of the diets containing 0 and 4 mg supplemental folic acid/kg was injected intraperitoneally with a dose of 4 μ Ci L-[1-¹⁴C]histidine/kg body-weight, contained in 0.258 M-L-histidine (10 ml/kg body-weight). Each bird was placed in the chamber and the trapping solution was changed every 30 min over a 3 h collection period. The ¹⁴CO₂ collected in the solution was measured as described by Saunderson (1985).

In the main experiment, between ten and fifteen birds, representing each of the five treatments, were injected intraperitoneally with 1 μ Ci labelled histidine/kg in the same dose of unlabelled histidine as before. Evolved ¹⁴CO₂ was collected for 1.5 h, with the trapping solution being changed at 15 min intervals.

A pair-feeding experiment was carried out on birds fed on the basal and control diets. The chicks were fed *ad lib.* as before for 1 week and then transferred to individual small wire cages in a heated room. The birds were allocated to six groups of three birds on the basis of similar body-weight. In each group, two birds continued to be fed on the basal and control diets *ad lib.*, while the third bird remained on the control diet but was pair-fed with the bird given the basal diet. Histidine oxidation was measured as before when the birds were approximately 14 d old.

Enzyme activities

Dihydrofolate reductase (*EC* 1.5.1.3; DHFR) activity was measured in erythrocytes. Blood samples (0.5 ml) were taken from each chick by heart puncture using heparin as an anticoagulant. After centrifugation the plasma and leucocyte layer were drawn off and the erythrocytes were resuspended in ice-cold saline (9 g sodium chloride/l; 1.5 ml). This process was repeated twice more. The erythrocytes were finally suspended in 0.05 M-sodium citrate (0.3 ml) at pH 7.2 and 0°. A small portion of this solution was taken for cell counting using a Coulter counter. The cells in the remainder of the solution were lysed by freeze-thawing twice and insoluble debris was removed by centrifugation at 20000 g for 30 min at 4°.

DHFR was measured by the method of Rothenberg *et al.* (1980) with slight modifications. ³H-labelled folic acid (50 pmol) was added to 250 μ l of a solution of sodium citrate (0.05 M), mercaptoethanol (0.01 M) and sodium dithionite (6 mM) at pH 7.2 in 1.5 ml capped Eppendorf tubes. This mixture was incubated at 37° for 15 min. The erythrocyte preparations (40 μ l) and NADPH (50 μ l, 1.3 mM) were then added to each tube. A blank did not contain any erythrocyte preparation. The volume in each tube was made up to 500 μ l with a solution of sodium citrate (0.05 M) and mercaptoethanol (0.01 M) at pH 7.2. The reaction was stopped by the sequential addition of 200 μ l freshly prepared 0.027 M-folic acid in 0.1 M-sodium hydroxide, 100 μ l 0.2 M-hydrochloric acid and 200 μ l 0.3 M-zinc

sulphate to precipitate selectively both unconverted [^3H]folic acid and unreacted [^3H]dihydrofolic acid. After removal of folate and protein precipitates by centrifugation, a 100 μl portion of supernatant fraction was added to 5 ml Optiphase X for scintillation counting. The coefficient of variation of repeat measurements using this method was 5.7%.

Glycine *N*-methyltransferase (*EC* 2.1.1.20; GNMT) and glutamic acid formimino-transferase (*EC* 2.1.2.5; GFT) activities were measured in liver samples. The chicks were killed by cervical dislocation and livers were removed and frozen quickly to -20° . Homogenates for both assays were prepared by homogenizing 0.5 g frozen liver with 2.5 ml ice-cold 10 mM-potassium phosphate and 1 mM-EDTA at pH 7.2 using a Polytron homogenizer. This homogenate was used directly in the assay for GFT (Tabor, 1962). For GNMT, the homogenate was centrifuged at 10000 *g* for 30 min at 4° and the supernatant fraction was assayed by the method of Cook & Wagner (1984) modified by the use of *S*-adenosylmethionine labelled with ^{14}C instead of ^3H . Protein contents of the preparations were measured by the method of Lowry *et al.* (1951). The coefficients of variation of repeat measurements were 5.7 and 2.1% for GFT and GNMT respectively.

Statistical analysis

Values were assessed using analysis of variance. Means for different treatments were compared using Student's *t* test with pooled variance obtained from the analysis. The histidine-oxidation response curve was fitted to the individual values by a modified Newton method of maximizing the likelihood using standard non-linear models available in a Genstat package (Genstat, 1987).

RESULTS

Body-weights at 2 weeks of age of birds fed on the different diets are shown in Table 1. Birds fed on the basal diet were significantly ($P < 0.05$) lighter than those on all other diets, but differences between bird weights with the other diets were not significant ($P > 0.05$). As was also observed in an earlier study (Maxwell *et al.* 1988), growth depression on the basal diet did not occur until after 7 d of age.

Histidine oxidation

Results from the preliminary comparison of rates of oxidation of histidine showed marked differences between the folate-deficient and control chicks (Fig. 1). Over the 3 h collection period, 0.6% of administered radioactivity was released as $^{14}\text{CO}_2$ by the deficient birds compared with 1.4% by the control birds. Peak rate of release of $^{14}\text{CO}_2$ was shown by the control after 1.5 h and this was taken as the collection time in the main experiment.

The percentages of label released as $^{14}\text{CO}_2$ by birds fed on different dietary levels of folate in the main experiment are given in Table 1. There was a significant ($P < 0.05$) suppression in release of $^{14}\text{CO}_2$ by birds fed on the basal diet. The dose-response relation was investigated and an asymptotic curve with the equation given in Fig. 2 was found to give the best fit (R^2 0.5). This curve suggested that about 2 mg supplemental folate/kg were needed for the maximum rate of catabolism of histidine to CO_2 .

In the pair-feeding experiment, the mean percentages of label released as $^{14}\text{CO}_2$ in 1.5 h were 6.38 (SE 0.71), 9.67 (SE 0.65) and 11.49 (SE 1.33) for the groups given diets containing no supplemental folate, 4 mg folate/kg (pair-fed with the first group) and 4 mg folate/kg *ad lib*. The values for the latter groups given supplemental folate did not differ significantly ($P > 0.05$) from each other, but both were higher ($P < 0.05$) than the value for birds fed on the basal diet.

Table 1. *Body-weight, oxidation of histidine to carbon dioxide and activities of dihydrofolate reductase (EC 1.5.1.3; DHFR), glycine N-methyltransferase (EC 2.1.1.20; GNMT) and glutamic acid formiminotransferase (EC 2.1.2.5; GFT) at 2 weeks of age in chicks fed on a purified diet supplemented with graded levels of folic acid**

(No. of observations shown in parentheses)

Supplemental folic acid (mg/kg)...	0	1	1.5	2	4	Standard deviation (from ANOVA table)
Body-wt (g)	146 ^a	200 ^b	208 ^b	181 ^b	207 ^b	45
Histidine oxidation (% of label released as ¹⁴ CO ₂ in 1.5 h)	4.67 ^a (13)	8.41 ^b (10)	7.13 ^b (14)	8.75 ^b (15)	8.66 ^b (13)	2.62
DHFR (pmol tetrahydrofolate/10 ⁹ erythrocytes in 20 min)	0.51 ^a (9)	0.28 ^b (10)	0.32 ^b (5)	0.34 ^b (6)	0.35 ^b (2)	0.14
GNMT (nmol sarcosine/mg protein in 15 min)	2.00 ^a (5)	2.03 ^a (8)	1.16 ^a (7)	1.46 ^a (8)	1.47 ^a (8)	0.73
GFT (μmol 5,10-methylenetetrahydrofolate/mg protein/min)	4.55 ^a (7)	5.70 ^a (9)	5.35 ^a (7)	5.35 ^a (10)	4.95 ^a (7)	0.85

^{a, b} Within a row, mean values with different superscript letters were significantly different ($P < 0.05$).

* For details of diets, see p. 203.

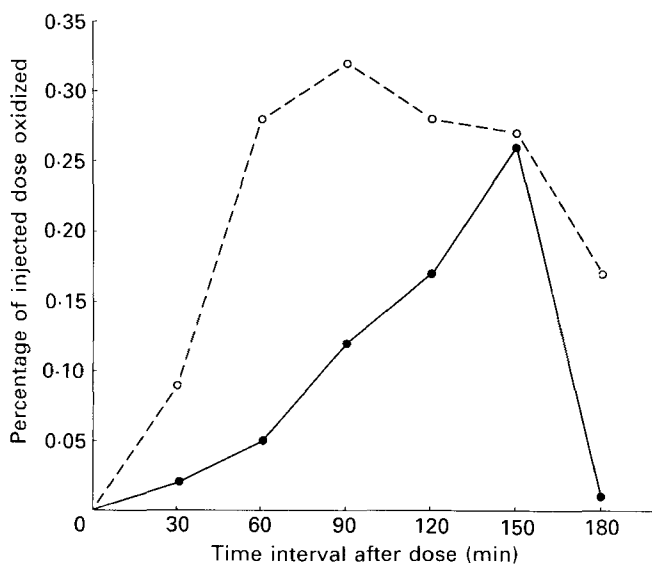


Fig. 1. Percentage of [1-¹⁴C]histidine released as ¹⁴CO₂ by chicks fed on diets containing 0 (●—●) and 4 (○--○) mg supplemental folic acid/kg (one chick per treatment). For details of diets, see p. 203.

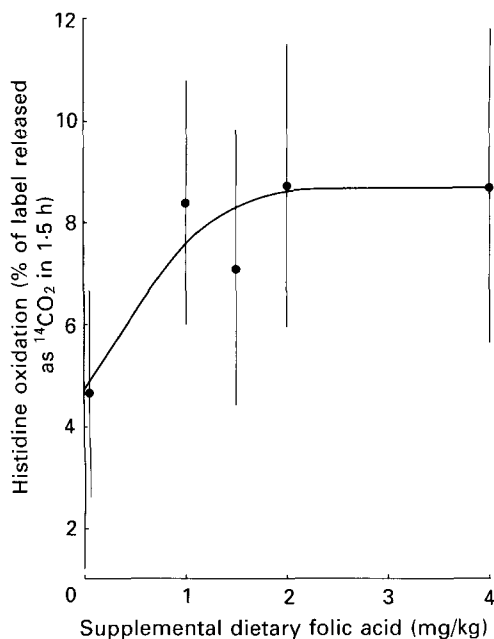


Fig. 2. Relation between histidine oxidation and dietary folic acid in chicks. Points are mean values for groups of chicks (standard deviations represented by vertical bars). The equation of the fitted curve is $Y = 8.58 - 3.87(0.24)^x$. For no. of chicks per treatment, see Table 1. For details of diets, see p. 203.

Enzyme activities

The specific activities for DHFR in erythrocytes and GFT and GNMT in liver are given in Table 1. DHFR activity was elevated significantly ($P < 0.05$) in the basal group. GFT and GNMT did not show any significant ($P > 0.05$) differences between treatments.

DISCUSSION

The conversion of *N*-formiminoglutarate (Figlu) to glutamate, catalysed by GFT, is the folate-dependent step in the oxidation of histidine. When folate is limiting, Figlu cannot be further metabolized by this route and the increased amount of it excreted in urine can be used as a measure of folate deficiency in mammals such as man and rats (Tabor *et al.* 1953; Luhby *et al.* 1959). This method is impractical in birds, but an alternative approach is to measure CO₂ released from the 1-position of glutamate by oxidation. Folate deficiency should decrease the amount of ¹⁴CO₂ released following the administration of [1-¹⁴C]histidine as part of a histidine load. This response was indeed observed in the present experiment and is consistent with reports of altered patterns of histidine oxidation in other species, e.g. human beings (Fish *et al.* 1963).

Comparisons with pair-fed birds confirmed that the decreased release of label was caused by folate deficiency *per se* rather than by depressed feed intake or growth. The decrease in histidine oxidation was significant ($P < 0.05$) only for the basal diet, containing no added folate, but oxidation was also less with the diets containing 1 and 1.5 mg folate/kg than with the diet containing 4 mg supplemental folate/kg. The response curve fitted to these values (Fig. 2) suggested that about 2 mg supplemental folate/kg was needed for the maximum rate of oxidation. The test, therefore, gives information on the folate status of

chicks. However, since it involves the administration of about half the normal daily histidine intake in a single dose, it is not yet clear to what extent the test also indicates the normal folate requirement of the bird.

Although the oxidation of histidine via the pathway involving GFT is depressed by folate deficiency, the inhibition seems to be caused by a lack of tetrahydrofolate cofactor rather than to a decreased amount of GFT. Thus, measurement of hepatic GFT *in vitro* in a medium containing added folate did not show any effect of dietary folate level on the enzyme specific activity. This result is consistent with previous observations of Burns & Jackson (1976).

GNMT is present in large amounts in mammalian liver: it makes up about 0.5% of the soluble protein in rat liver cytosol (Cook & Wagner, 1981). It catalyses the *S*-adenosylmethionine-dependent conversion of glycine to sarcosine, but may also act as a carrier protein since it binds tightly to, and is inhibited by, 5-methyltetrahydrofolate (Cook & Wagner, 1984). A deficiency of folate might, therefore, be expected to result in an increase in activity of this enzyme. Mean hepatic specific activities of GNMT were indeed found to be over 35% higher at the two lowest dietary levels of folate than at the highest, but these differences were not significant ($P > 0.05$). The high standard deviation (Table 1) suggests there is considerable biological variation in the activity of this enzyme.

DHFR is involved in the conversion of folic acid into one of the main metabolically active folates, tetrahydrofolate. The specific activity of this enzyme was significantly higher in the erythrocytes of chicks given no supplemental folic acid, but showed no differences between the supplemental diets. The elevated activity in the severely folate-deficient chicks is perhaps caused by a feed-back mechanism attempting to normalize reduced folate concentrations. However, this increase in activity of DHFR in erythrocytes contrasts with reported decreases in hepatic activity of DHFR in folate-deficient chicks (Burns & Jackson, 1976) and rats (Pasquali *et al.* 1968). Burns & Jackson (1976) speculated that folate deficiency decreased the concentration of DHFR in chick liver. To reconcile these opposite results, it is therefore necessary to propose that different mechanisms regulate this enzyme in erythrocytes and liver. Further studies are required before the value of DHFR as a criterion of folate status can be properly evaluated.

The results of these investigations confirm that biochemical criteria can be used to give information on folate status in chicks. Folate-metabolizing pathways involving the oxidation of histidine and DHFR seem to be particularly promising. However, before they can be used as a practical means of assessing status, further information is required on the range of dietary folate over which they respond, and influences of other dietary variables.

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