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Development of methionine synthase, cystathionine-β-synthase and S-adenosyl-homocysteine hydrolase during gestation in rats


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The developmental onset of three homocysteine metabolizing enzymes in the rat conceptus was investigated. Cystathionine-β-synthase and methionine synthase were assayed from day 10 to day 20 of gestation in decidual and placental tissue, from day 10 to day 12 of gestation in embryonic tissue, from day 14 to day 20 of gestation in fetal liver and from day 14 to day 20 of gestation in fetal tissue without liver. On each day, material was obtained from at least four conceptuses from two dams. S-adenosylhomocysteine hydrolase was assayed in neurulating conceptuses in decidual tissue, parietal yolksac plus ectoplacental cone, visceral yolksac plus amnion and embryo proper. Conceptuses were pooled from seven (day 9.5 of gestation) or three (days 10.5 and 11.5 of gestation) dams. In embryonic and fetal tissue cystathionine-β-synthase first occurred in fetal liver. During the organogenic phase it was present only in decidual tissue. Methionine synthase was present in all tissues from the first gestational day investigated and S-adenosylhomocysteine hydrolase was present in all tissues throughout the neurulating period. Our results indicate that the homocysteine—methionine cycle, which is crucial to transmethylation reactions, is functional during the neurulating period in embryonic tissue. Owing to the absence of cystathionine-β-synthase at this stage of development in embryonic tissue, the homocysteinyl moiety is conserved in the homocysteine—methionine cycle.

Introduction

A disturbance of methionine—homocysteine metabolism may play a role in the aetiology of neural tube defects, and also in other obstetrical problems, such as recurrent spontaneous abortion and abruptio placenta (Steegers-Theunissen et al., 1991, 1992; Schorah et al., 1993; Steegers-Theunissen, 1993; Wouters et al., 1993).

L-homocysteine can be metabolized by several enzymes (Fig. 1). Methionine synthase (N⁵-methyltetrahydrofolate-L-glutamate:L-homocysteine S-methyltransferase, EC 2.1.1.13) remethylates L-homocysteine to L-methionine. In this way the homocysteinyl moiety is conserved in methionine—homocysteine metabolism. At the same time methionine synthase is the only enzyme in mammals that can demethylate N⁵-methyltetrahydrofolate. Since N⁵-methyltetrahydrofolate is the major circulating form of folate, methionine synthase is the key enzyme to their active reduced folate supply, which is essential in one-carbon metabolism, and purine and pyrimidine synthesis. Therefore, methionine synthase is considered to be ubiquitous in mammals (Shane and Stokstad, 1983). During gestational development methionine synthase activity seems to progressively decline in several species (Gaul et al., 1973; Sternowsky et al., 1976). However, a developmental study, including the organogenic phase, has never been published.

Betaine:L-homocysteine S-methyltransferase (EC 2.1.2.3) can also remethylate L-homocysteine to L-methionine. Betaine:L-homocysteine S-methyltransferase activity is found in liver tissue of all mammals investigated and also occurs in kidney tissue (Ericson, 1960; Finkelstein et al., 1971). In humans betaine:L-homocysteine S-methyltransferase activity is also present in both fetal and adult brain tissue, although activity is low (Gaul et al., 1973). Since betaine:L-homocysteine S-methyltransferase is a typical liver enzyme, and the liver is not present at the time of neurulation, the period under investigation, no attempt was made in this study to measure its activity in rat embryonic and fetal tissue.

Cystathionine-β-synthase (EC 4.2.1.22) condenses L-serine with L-homocysteine to L-cystathionine. This is an irreversible step in the transsulfuration pathway, the main pathway for methionine and homocysteine catabolism. Generally cystathionine-β-synthase activity increases during fetal development; however, the developmental onset is unknown, and may differ between species (Volpe and Laster, 1970, 1972; Gaul et al., 1972; Sturman et al., 1976; Rassin et al., 1981).

S-adenosylhomocysteine hydrolase (EC 3.3.1.1) is the only source of L-homocysteine in mammals as it hydrolyses S-adenosylhomocysteine to L-homocysteine and adenosine.
However, when l-homocysteine levels are high, S-adenosylhomocysteine hydrolase will synthesize S-adenosylhomocysteine, as the reaction is reversible and S-adenosylhomocysteine synthesis is energetically more favourable (Ueland, 1982). Since S-adenosylhomocysteine accumulation will inhibit crucial transmethylation reactions, in mammals, S-adenosylhomocysteine hydrolase is assumed to be ubiquitous as well. However, developmental studies have not been published.

The objective of this study was to investigate the gestational development of methionine synthase, cystathionine-β-synthase and S-adenosylhomocysteine hydrolase in the rat conceptus to provide a better understanding of the possible relationship between methionine—homocysteine metabolism and the aetiology of neural tube defects.

Materials and Methods

Chemicals

Bovine serum albumin (BSA), $N^\delta$-methyltetrahydrofolate, complete o-phthalaldehyde reagent solution, Coomassie Brilliant Blue G, dithiothreitol, l-homocysteinethiolactone, l-serine, l-cystathione, S-adenosylhomocysteine, S-adenosylmethionine, lubrol PX and pyridoxal phosphate were purchased from Sigma (St Louis, MO). Ascorbic acid and 2-mercaptoethanol were purchased from Merck (Darmstadt). T61 containing 20% (w/v) embutramide, 5% (w/v) mebezoniumiodide and 0.5% (w/v) tetracaine hydrochloride in water) was obtained from Hoechst Veterinär GmbH (Munich); hydroxycobalamin (vitamin B12b) from Fluka Chemie (Buchs); [U-14C]-l-serine (specific activity 166 mCi mmol$^{-1}$) from DuPont de Nemours (Dreiech); erythro-9[2-hydroxy-3-nonyl]adenine from Burroughs Wellcome (Research Triangle Park, NC); adenosine from USB (Cleveland, OH) and Triton X-100 from Serva Feinbiochemica (Heidelberg). All solutions were made in tap water, demineralized by a Milli-RO 10TS system from Millipore Corporation (Bedford, MA), except for the HPLC buffers, which were prepared with demineralized water additionally filtered by a Nanopure System from Sybron/Barnstaedt (De Bugue, IA). l-Homocysteine was prepared from l-homocysteinethiolactone by alkaline hydrolysis (5 mol NaOH l$^{-1}$, 5 min, 37°C). The solution was neutralized with HCl and diluted with the appropriate assay buffer.

Animals and dissection of tissues

Random bred Cpb-WU (Wistar) rats were housed in pairs or groups of three in cages and allowed to eat (MRH-TM pellets, Hope Farms B.V., Woerden) and drink (tap water) ad libitum. For animals from which material was obtained on days 10, 11, 12, 14, 16, 18 or 20 of gestation, lights were on from 12:00 h to 12:00 h.
until 00:00 h. On day 0, females (11–26 weeks old, 200–300 g) were brought together with males (1:1) from 09:00 h to 12:00 h. Animals from which material was obtained on days 9.5, 10.5 or 11.5 of gestation, were housed under the same conditions, except for the light regimen and the time of mating, which were shifted by 12 h.

When the conceptuses had reached the gestational age desired, the dams were killed by an intracardial injection of 0.4 ml T61. The uterus was removed and the conceptuses were explanted and placed in Hank's balanced salt solution. Conceptuses at day 10, 11 or 12 of gestation were dissected into (i) embryo proper and (ii) decidua. Extraembryonic membranes (amnion, visceral and parietal yolk sac and ectoplacental cone) were discarded. Conceptuses at day 14, 16, 18 and 20 of gestation were dissected into (i) fetal liver, (ii) rest of fetus and (iii) placenta plus extraembryonic membranes plus residual decidual tissue. Conceptuses at day 9.5 of gestation were dissected into (i) embryo proper plus amnion plus visceral yolk sac, (ii) parietal yolk sac plus ectoplacental cone and (iii) decidua. Conceptuses at day 10.5 or 11.5 of gestation were dissected into (i) embryo proper, (ii) amnion plus visceral yolk sac, (iii) parietal yolk sac plus ectoplacental cone and (iv) decidua. For reference, maternal liver (day 10 of gestation) was used. After dissection, tissues were immediately frozen in liquid nitrogen and stored at −80°C until further treatment.

**Enzyme assays**

Protein concentrations were assayed using Coomassie Brilliant Blue G according to Bradford (1965). BSA was used as a standard. All samples were analysed in triplicate.

The methionine synthase and the cystathionine-β-synthase assays were performed with all tissues obtained from dams on days 10, 11, 12, 14, 16, 18 or 20 of gestation. At each stage, material was obtained from two animals and at least two conceptuses from each animal were assayed in duplicate. The methionine synthase assay and measurement of the α-phthalaldehyde derived methionine by HPLC and fluorometric detection were performed according to Garras et al. (1991).

The cystathionine-β-synthase assay was performed according to Fowler et al. (1978), except that tissues were homogenized (1:10, w:v) in ice-cold 50 mmol potassium phosphate buffer 1−1 (pH 7.4) containing 0.1% (w/v) lubrol PX and 1 mmol Na₂EDTA·2H₂O 1−1, 20 μmol erythro-[2-hydroxy-3-nonyl]adenine 1−1, 1.64 ml Triton X-100 1−1 and 1.1 mmol dithiothreitol 1−1. The assay was performed at 37°C for 20 min in the same buffer (10 μl homogenate in a final volume of 200 μl) additionally containing 0.5 mmol L-homocysteine 1−1 and 0.5 mmol adenosine 1−1. After the incubation, protein was precipitated with 12 μl 8 mol perchloric acid 1−1. After 10 min on ice, the mixture was centrifuged in an Eppendorf centrifuge for 5 min (13 000 g, 4°C) and 25 μl 4 mol dipotassium hydrogenphosphate 1−1 was added to the supernatant. After another 10 min on ice, the precipitate was removed by centrifugation and the supernatant was frozen in liquid nitrogen. Samples were stored at −80°C until further analysis. S-adenosylhomocysteine was measured in duplicate by using HPLC and spectrophotometric detection according to DeAbreu et al. (1982). All enzyme activities were expressed in μU (pmol product formed min−1 mg−1 protein).

**Statistical analyses**

Linear regression analysis was used to test whether methionine synthase activity or cystathionine-β-synthase activity increased or decreased during gestation. A decrease or increase was regarded as significant if P < 0.05 for the null-hypothesis that the regression coefficient equalled zero. Differences in enzyme activity between gestational days and differences in S-adenosylhomocysteine hydrodase activity between tissues for each gestational day were tested using Student's t test with Bonferroni correction. With a nominal α of 0.05, differences were regarded as statistically significant at P < 0.05 k−1, where k is the number of comparisons made. All differences, increases and decreases, in enzyme activities mentioned in the results section met the degree of statistical significance as defined above.

**Results**

Methionine synthase was present in all tissues investigated. The highest specific activity was found in embryonic tissue on day 12 of gestation. On days 10 and 11 of gestation it was lower and from day 12 onward the activity decreased at an average rate of 33 μU day−1. Methionine synthase specific activity of maternal liver was 180 μU, which was comparable to the average methionine synthase specific activity of fetal liver. Methionine synthase activity of the fetal liver decreased during gestational development at a rate of 26 μU day−1. The lowest methionine synthase specific activity was found in decidual and placental tissue, and here it did not change significantly during gestation (Fig. 2).

During gestation, cystathionine-β-synthase activity was not present in embryonic tissue. In this organogenic phase of development, it was present only in the surrounding decidual or placental tissue. In the latter tissue, it decreased during gestation at a rate of 15 μU day−1 and was virtually absent at day 16 of gestation. In fetal tissue, activity first occurred in the liver, where it was present from the first day (day 14) that it
Fig. 2. Methionine synthase activity (---) and cystathionine-β-synthase activity (----) in (a) embryonic and fetal tissue without liver, (b) fetal liver and (c) decidual and placental tissue during the second half of gestation in rats. Each point represents the mean of four conceptuses taken from two dams. Error bars represent SEM.

Fig. 3. S-adenosylhomocysteine hydrolase activity in neurulating rat conceptuses. Enzyme activity was measured in the direction of synthesis. Conceptuses were separated into embryo proper (+ amnion) (□), parietal yolksac (+ ectoplacental cone) (■) and decidua (●), except for embryos from day 9.5 of gestation, in which embryos and visceral yolksac were pooled together (△). Bars and error bars represent the mean and SEM of three pools of conceptuses.

Discussion

The data on the developmental onset of cystathionine-β-synthase in rat fetuses extend those of Volpe and Laster (1972) and show that this enzyme first appears in the fetal liver and that, before this organ is formed, cystathionine-β-synthase activity is present only in the surrounding decidual and
placental tissue. The presence of methionine synthase activity in rat embryos and fetal liver (Baden et al., 1984, 1987; Hansen and Billings, 1986) is confirmed; however, owing to the decrease of methionine synthase activity of fetal liver towards the end of gestation, it appears that the observation from Baden et al. (1984, 1987) that fetal liver activity is 50–65% of maternal liver activity is valid only for fetal liver at the end of gestation. The activity of methionine synthase in embryonic and fetal tissues from day 12 to day 18 of gestation was higher relative to maternal liver activity, which is in contrast to the results of Hansen and Billings (1985) that, at day 14 of gestation, activity in embryonic and fetal tissues was the same as the activity of maternal liver. The observations presented here extend those of Hansen and Billings (1985) and, for the first time, demonstrate that methionine synthase activity is present in neurulating rat embryos. In studies on the teratogenicity of nitrous oxide, a methionine synthase inhibitor, the presence of methionine synthase in neurulating embryos was only presumed (Baden et al., 1983; Baden and Fujinaga, 1991; Fujinaga and Baden, 1994).

This is the first report to demonstrate S-adenosylhomocysteine hydrolase activity in neurulating rat embryos. The higher activity of this enzyme in embryonic and visceral yolk sac tissue relative to the activity in the surrounding parietal yolk sac and decidua indicates that S-adenosylhomocysteine metabolism occurs mainly in the embryonic and visceral yolk sac compartments. The presence of S-adenosylhomocysteine hydrolase and methionine synthase, and the absence of cystathionine-β-synthase, in the embryo proper indicates that the homocysteine–methionine cycle is functional at this stage of development, but that there is no diversion into the transsulfuration pathway in the embryo. It is calculated that in adult rat liver tissue about half of the homocysteine is transsulfurated, whereas the other half is remethylated to methionine (Finkelstein and Martin, 1986; Finkelstein, 1990). The proportion of homocysteine that will be remethylated can be increased or decreased depending on the need for S-adenosylmethionine. The conservation of the homocysteinyl moiety in the homocysteine–methionine cycle as a result of the absence of cystathionine-β-synthase in the neurulating embryo emphasizes the importance of this cycle in embryonic development at this stage. The relevance of this notion is also supported by the work of Klein and coworkers (Coelho et al., 1989; Coelho and Klein, 1990; Ferrari et al., 1993), who show that methionine deficiency leads to neural tube defects in cultured rat embryos and that the teratogenicity of human sera in the whole embryo culture system is related to numbers of previous spontaneous abortions and nutritional factors.}

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