



BRAZILIAN JOURNAL
OF MEDICAL AND BIOLOGICAL RESEARCH

www.bjournal.com.br

ISSN 0100-879X
Volume 43 (01) 1-123 January 2010

BIOMEDICAL SCIENCES
AND
CLINICAL INVESTIGATION

Braz J Med Biol Res, January 2010, Volume 43(1) 1-7

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The Brazilian Journal of Medical and Biological Research is partially financed by



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Genetics of homocysteine metabolism and associated disorders

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Abstract

Homocysteine is a sulfur-containing amino acid derived from the metabolism of methionine, an essential amino acid, and is metabolized by one of two pathways: remethylation or transsulfuration. Abnormalities of these pathways lead to hyperhomocysteinemia. Hyperhomocysteinemia is observed in approximately 5% of the general population and is associated with an increased risk for many disorders, including vascular and neurodegenerative diseases, autoimmune disorders, birth defects, diabetes, renal disease, osteoporosis, neuropsychiatric disorders, and cancer. We review here the correlation between homocysteine metabolism and the disorders described above with genetic variants on genes coding for enzymes of homocysteine metabolism relevant to clinical practice, especially common variants of the *MTHFR* gene, 677C>T and 1298A>C. We also discuss the management of hyperhomocysteinemia with folic acid supplementation and fortification of folic acid and the impact of a decrease in the prevalence of congenital anomalies and a decline in the incidence of stroke mortality.

Key words: Homocysteine; Hyperhomocysteinemia; Folate metabolism; Single nucleotide polymorphism; Susceptibility genes

Introduction

Homocysteine (Hcy) is the demethylated derivative of methionine, which, after conversion to S-adenosylmethionine, is the most important methyl group donor in the body. Hcy is metabolized by one of the two following pathways, i.e., remethylation or transsulfuration (Figure 1). In remethylation, Hcy is transformed to methionine by the addition of a methyl group from 5-methyltetrahydrofolate or betaine. 5-Methyltetrahydrofolate is the result of the conversion of folic acid from the diet to 5,10-methyltetrahydrofolate and finally to 5-methyltetrahydrofolate by the enzyme 5,10-methyltetrahydrofolate reductase (MTHFR). In all tissues, the cofactor vitamin B12 participates in the remethylation reaction with 5-methyltetrahydrofolate, whereas the reaction with betaine is restricted to the liver and is independent of vitamin B12. Methionine is then activated by ATP to form S-adenosylmethionine, a universal methyl donor to a variety of acceptors. In the transsulfuration pathway, Hcy is converted to cystathionine by cystathionine β -synthase and finally to cysteine using vitamin B6 as a cofactor (1).

Therefore, plasma Hcy levels are determined by several factors such as the cofactors vitamin B12, vitamin B6 and folic acid and enzymes involved in methionine metabolism. The levels of these cofactors are inversely correlated, with

plasma Hcy levels increasing with decreasing vitamin B, folic acid and vitamin B6 concentrations (1).

In the circulation, 80-90% of Hcy is protein bound especially to albumin, its main carrier in plasma. The remaining 10-20% of total Hcy (tHcy) is acid-soluble free Hcy presenting as Hcy-cysteine mixed disulfide and homocystine (a dimer of Hcy) and less than the remaining 1% is present in the free reduced form (2).

Homocysteine and MTHFR

MTHFR is an important enzyme in Hcy metabolism, which catalyzes the conversion of 5,10-methylenetetrahydrofolate to 5-methyltetrahydrofolate, the predominant circulating form of folate. The *MTHFR* gene has at least two functional polymorphisms, 677C>T and 1298A>C. The *MTHFR* 677T allele is associated with reduced enzymatic activity, decreased concentrations of folate in serum, plasma, and red blood cells, and mildly increased plasma tHcy concentrations (3,4).

The *MTHFR* polymorphism 1298A>C also affects MTHFR activity but is not associated with higher plasma Hcy or lower folate levels (5). Normal MTHFR activity is

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Received April 29, 2009. Accepted November 26, 2009. Available online December 7, 2009. Published January 11, 2010.

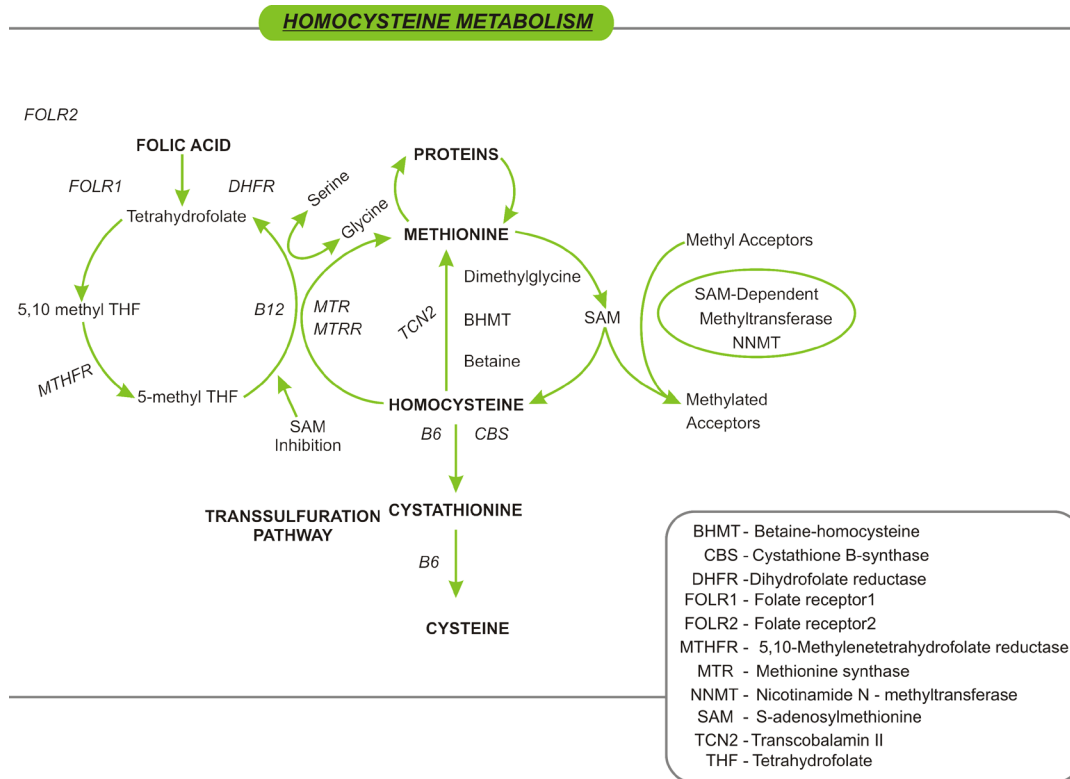


Figure 1. Homocysteine metabolism.

crucial to maintain the pool of circulating folate and methionine and to prevent the accumulation of Hcy (3).

Double heterozygosity for *MTHFR* 677C>T and 1298A>C polymorphisms results in lower MTHFR activity as compared to heterozygosity for either *MTHFR* variant separately (5). Individuals with the 677TT genotype have approximately 30% the MTHFR enzyme activity of those with the 677CC genotype, whereas 677CT heterozygotes have about 65% enzymatic activity (6).

Hyperhomocysteinemia

Blood levels of hyperhomocysteinemia (HHcy) are age

and gender related. Plasma Hcy levels are higher in men than in women (7) and increase from 10.8 $\mu\text{mol/L}$ at age 40-42 up to 12.4 $\mu\text{mol/L}$ between 65-67 years (8). HHcy is classified as moderate (15-30 $\mu\text{mol/L}$), intermediate (31-100 $\mu\text{mol/L}$) or severe (>100 $\mu\text{mol/L}$) (9).

In humans, HHcy can also be distinguished by category, including cause, prevalence and severity. The more severe cases are due to homozygous defects in genes encoding enzymes of Hcy metabolism (Table 1).

HHcy is observed in approximately 5% of the general population and has been associated with many disorders. Below we review several disorders that have been related to abnormal Hcy metabolism.

Table 1. Classification of hyperhomocysteinemia according to Selhub (1).

Severe hyperhomocysteinemia

High total homocysteine (tHcy) levels at all times (31 to >100 $\mu\text{mol/L}$) caused, for example, by deficiencies in cystathionine beta synthase (CBS), methylenetetrahydrofolate reductase (MTHFR), or in enzymes of B12 metabolism

Mild hyperhomocysteinemia

Moderately high tHcy levels (15-30 $\mu\text{mol/L}$) under fasting conditions; reflects impaired homocysteine methylation (folate, B12 or moderate enzyme defects, e.g., thermolabile MTHFR)

Post-methionine load

Abnormal increase in tHcy (>15 $\mu\text{mol/L}$) after a methionine load (100 mg/kg); reflects impaired homocysteine transsulfuration (heterozygous CBS defects, B6 deficiency)

Hyperhomocysteinemia and neural tube defects

Neural tube defects (NTD) are among the most common birth defects worldwide. NTD result from failure of neural tube formation or closure during the first 28 days of pregnancy. Studies have supported the use of folic acid in the prevention of the first occurrence of NTD (10). These findings suggest that impaired folic acid metabolism plays a role in NTD.

Several studies on different populations have reported that high maternal Hcy levels are associated with an increased risk for offspring with NTD (11-17). A case-control study conducted in Brazil showed that mothers with NTD children had increased tHcy levels when compared with control mothers. Case mothers also had low vitamin B12 levels compared to control mothers. No differences in *MTHFR* polymorphisms were identified in mothers, children with NTD or controls (18). Another Brazilian study was also unable to identify *MTHFR* polymorphism and NTD, suggesting that this gene does not play a role in NTD in the Brazilian population (19).

Hyperhomocysteinemia and nonsyndromic oral clefts

Oral clefts are common birth defects whose prevalence differs between geographical regions and ethnic groups. They have a complex, multifactorial inheritance involving genetic and environmental factors.

A case-control study showed that mothers of children with clefts had higher plasma tHcy concentrations both under fasting conditions and after a methionine afterload test compared to control mothers, suggesting that impaired folic acid metabolism plays a role in clefting (20). A recent study has indicated that periconceptional use of folate could prevent the occurrence of cleft lip and palate (CL/P) but not of cleft palate alone (21).

A study demonstrated a significantly increased risk of having a child with CL/P if the mother was heterozygous for both common *MTHFR* polymorphisms (677C>T, 1298A>C) (22). The risk of a child with CL/P was only significantly increased if mothers carrying the *MTHFR* 677TT or *MTHFR* 1298CC genotype also had a low periconceptional intake of dietary folate and/or folic acid supplements, suggesting that the two *MTHFR* polymorphisms were independent risk factors for CL/P (23).

Hyperhomocysteinemia and congenital heart defects

Congenital heart defects (CHD) occur in 11/1000 live births in the United States and annually account for approximately 6000 total deaths and one-tenth of infant deaths in the USA. CHD develop during the first 3 weeks

after conception and are common congenital anomalies of multifactorial origin influenced by genetic and environmental factors (24). A meta-analysis demonstrated that maternal HHcy was significantly associated with an increased risk of having a child with a CHD, suggesting that maternal HHcy is an important risk factor for CHD (23).

Hyperhomocysteinemia and cardiovascular disorders

Vascular diseases are commonly associated with traditional risk factors such as systemic arterial hypertension, diabetes mellitus and smoking, but in the last decade other risk markers have been identified, one of them being Hcy. Various studies had identified HHcy as an independent risk factor for coronary artery disease (CAD). The relationship between *MTHFR* polymorphism and the severity of CAD in patients undergoing coronary artery bypass surgery showed that Hcy levels were significantly higher in patients with CAD than in control subjects and the genotype of *MTHFR* 677C>T was associated with the extent of CAD in patients at high risk for this pathology (25).

A study comparing post-mortem serum Hcy levels of men who died suddenly of severe CAD and controls showed that serum Hcy levels were elevated in men who died with coronary thrombus compared to controls. This study also showed that the risk was increased if there was concomitant diabetes mellitus and the presence of fibrous plaque atheromas. Whether elevated serum Hcy might be causally associated with sudden death or merely a marker of a process related to sudden death is still unclear (26).

Hyperhomocysteinemia and atherosclerosis

Atherosclerosis is a chronic inflammatory disease of the arteries, in which deposits of fatty substances, cholesterol, calcium and other substances build up in the endothelial layer of the arteries. Several studies have identified moderate HHcy as an independent risk factor for atherosclerotic disease (9).

Accelerated atherosclerosis had been demonstrated in apolipoprotein E-deficient (apoE^{-/-}) mice on an Hcy diet and in cystathionine-synthase (CBS)/apoE^{-/-} double-knockout mice (27,28). These findings suggest that HHcy accelerates atherogenesis in a hyperlipidemic mouse model that develops spontaneous atherosclerosis. However, a causal relationship between atherogenesis and HHcy had not been established in other species, including rats, rabbits, pigs, and primates (29,30).

Hyperhomocysteinemia and Down syndrome

Down syndrome (DS) occurs due to three copies of

chromosome 21, in most cases due to the failure of chromosomal segregation during maternal meiosis. The risk of DS is directly correlated to increased maternal age. An important factor that could modulate the maternal risk for DS is plasma HHcy concentration. An elevated risk for DS has been observed in the presence of the *MTR* 2756G allele in combination with elevated HHcy concentration (31). However, the association of this polymorphism with the risk for DS was not confirmed by an additional study (32).

A study conducted in Brazil with mothers of children with DS and control mothers analyzed tHcy and several variants in the folate pathway (*MTHFR* 677C>T, *MTHFR* 1298A>C, *MTRR* 66A>G, *MTR* 2756A>G, and *CBS* 844ins68), showing that Hcy levels were higher among DS mothers than among controls. Only the 677T allele was associated with altered tHcy levels in the case group. All genotype distributions were similar in the two groups, but the frequency of the 677T allele was significantly higher in the case group. None of the other polymorphisms showed an association with risk for DS when evaluated separately. However, when the presence of the alleles was evaluated as a whole, the mothers of children with DS tended to have a higher number of variant alleles than control mothers (33).

When the effect of plasma HHcy concentrations on maternal risk for DS was investigated, Hcy concentrations were found to be significantly higher in DS mothers compared to control. The study also showed that Hcy concentrations were significantly higher in DS mothers with the *MTHFR* 1298CC genotype compared to the same genotype of control mothers (34).

Hyperhomocysteinemia and smoking or pregnancy

Cigarette smoking has been associated with oxidative stress and an increased risk of many chronic diseases. Smoking induces depletion of cellular antioxidants and is also known to be associated with increased Hcy levels. Several studies have observed an increase in plasma tHcy among smokers compared to non-smokers (35,36).

During pregnancy there is an increase in Hcy levels for each trimester of pregnancy and a decrease of folate levels. Additionally, folate levels continue to decrease after pregnancy (37). Maternal smoking also alters total Hcy concentration in infants. Serum Hcy concentrations were significantly higher in smoking than non-smoking pregnant women and also in the umbilical cord blood of their newborns (38).

Hyperhomocysteinemia and Alzheimer's disease and vascular dementia

Low folate and increased Hcy concentrations in blood are associated with poor cognitive performance in the general population. Elevated levels of Hcy result in neurotoxic

and vasotoxic effects in dementia and Alzheimer's disease, suggesting that Hcy is a direct marker of early cognitive decline (39). However, the pathogenesis of HHcy is still not clear in vascular dementia. The levels of serum folate, vitamin B12 and plasma Hcy were studied in vascular dementia and Alzheimer's disease, and it was found that Hcy was increased, while folate and vitamin B12 were decreased in both of these diseases. This suggests that supplementation of folic acid and vitamin B12 could be of benefit in Alzheimer's disease and dementia (40).

A randomized, double-blind, placebo-controlled study was conducted in the Netherlands on 818 subjects, men and post-menopausal women aged 50-70 years who used placebo or 800 µg/day folic acid for 3 years. Serum folate concentrations increased and plasma Hcy concentrations decreased in participants taking folic acid compared with those taking placebo. This study also showed that supplementation with folic acid significantly improved the decline of cognitive function related to age (41).

Hyperhomocysteinemia and breast cancer

A high intake of folate, which is plentiful in vegetables and fruits, has been associated with reduced risk of several cancers. Folate deficiency has been suggested to increase the risk of cancer through impaired DNA repair synthesis and disruption of DNA methylation that may lead to proto-oncogene activation (42). It is biologically plausible that polymorphisms or gene-environment interactions rather than folate intake alone would have an impact on breast cancer risk since functional polymorphisms in folate-related genes contribute to the alteration of folate metabolism. Therefore, *MTHFR* polymorphisms have been intensively studied in breast cancer but the results are inconsistent (43,44).

However, in a study performed with 456 breast cancer cases and 912 controls, breast cancer risk was inversely associated with consumption of dietary folate and none of the polymorphisms (*MTHFR*, *MTRR*, *MTR*) showed any significant impact on breast cancer risk. In postmenopausal women, a significantly increased risk of breast cancer was found among individuals with the *MTHFR* 677TT genotype compared with those with the *MTHFR* 677CC genotype. The study showed a significant interaction between *MTRR* A66G and folate intake. The *MTRR* 66GG genotype increased the risk among postmenopausal women with a low folate intake (45).

Hyperhomocysteinemia and depression

Folate and vitamin B12 deficiency, HHcy and the 677T allele of the *MTHFR* gene, which cause impaired methylation reactions in the central nervous system, have been associated with depressive disorders (46). However other studies did not find this association (47,48).

A prospective study on 732 Korean subjects investigated associations between folate, vitamin B12, Hcy, and late-life depression. The incident depression was predicted by lower folate and vitamin B12 levels and higher Hcy levels determined 2 years previously, and was associated with a decline in vitamin B12 levels and an increase in Hcy levels over the intervening period. However, incident depression was not associated with *MTHFR* genotype (49).

Hyperhomocysteinemia and diabetes

Mild HHcy has been observed in type I diabetic patients with microalbuminuria and nephropathy and may explain the increased risk of vascular disease in this high-risk population (50).

A study conducted in Spain on 46 patients with diabetes mellitus between 4 and 19 years of age showed that children with strict control of diabetes type I had normal tHcy, with no significant difference compared to healthy individuals of the same age and social environment (51). It was hypothesized that oral administration of folic acid would reduce plasma Hcy levels, improving endothelial function and oxidative stress in patients with type I diabetes and microalbuminuria. A case study showed a 25% reduction in plasma Hcy after folic acid supplementation in diabetic subjects (52).

Hyperhomocysteinemia and drugs

There is evidence supporting the unfavorable effects of some anti-epileptic drugs on plasma Hcy concentrations. Approximately 10-40% of epileptic patients develop HHcy (53). The levels of HHcy were increased in monotherapy consisting of phenytoin, carbamazepine or valproic acid (54).

A prospective observational study was conducted in Italy in 2007 to evaluate the influence of antiepileptic drugs, *MTHFR* polymorphism and Hcy levels. A higher prevalence of the *MTHFR* 1298C allele was observed in epileptic patients compared to controls. After folate therapy, plasma tHcy and folate were normal in all patients (55).

Management of hyperhomocysteinemia

Elevations in plasma Hcy are common in the general population, particularly in the elderly. Vitamin status is a

primary determinant of mild-to-moderate HHcy and accounts for approximately two thirds of all such cases. Vitamin supplementation results in near normalization of plasma Hcy in most cases. Hcy metabolism requires the participation of folate as well as vitamin B12 (cobalamin) and vitamin B6 (pyridoxal phosphate) coenzymes and, therefore, reduction of Hcy levels in plasma requires supplementation with all three vitamins (1). The minimum daily requirement of folic acid is 50 µg, although the current recommended intake is 400 µg/day for the average adult and 600 µg/day during pregnancy (56).

A large study on 750 males observed that dietary intervention with increased focus and availability of vegetables, fruits and bread, significantly reduced the concentration of tHcy. These findings suggest that the changes in the concentration of cysteine, folate and flavin mononucleotide seem to be predictors of changes in tHcy concentration (57).

Fortification of grain food with folic acid to decrease the incidence of NTD was started in the USA and Canada in 1998. This public health strategy reduced tHcy concentrations, leading not only to a decrease of the prevalence of congenital anomalies but also to decline in the incidence of stroke mortality (58).

In the last decade several studies have been conducted to uncover the direct or indirect influence of increased Hcy levels in several conditions.

It was demonstrated that several polymorphisms in genes of the Hcy-methionine pathway result in HHcy, suggesting that these variants may play a role in many multifactorial disorders of high prevalence in the general population.

Vitamin supplementation is a cost-effective way to decrease HHcy and would prevent at least some of these disorders.

Despite these robust indications about the role of HHcy as a factor associated with several conditions, long-term studies on large samples are still lacking to prove that decreasing the Hcy levels in subjects with HHcy results in better results in terms of health outcomes.

Acknowledgments

Research supported by FIRCA/NIH TW007644-02 (T.M. Félix) and CNPq. R. Giugliani and S. Brustolin are recipients of fellowships from CNPq.

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