Effect of the MTHFR 677C/T Polymorphism on Homocysteinemia in Response to Creatine Supplementation: A Case Study

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Received March 19, 2013
Accepted May 2, 2013
On-line July 17, 2013

Summary
Creatine (Cr) is recommended as a dietary supplement especially for athletes but its therapeutic potential is also discussed. It is assumed that human body uses Cr for the formation of phosphocreatine, which is necessary for muscular work as a source of energy. Production of Cr in a body is closely connected to methionine cycle where guanidinoacetate (GAA) is in a final step methylated from S-adenosylmethionine (SAM). Increased availability of SAM for phosphatidylcholine (PC) and sarcosine synthesis can potentially stimulate endogenous production of betaine a thus methylation of homocysteine (Hcy) to form methionine. Our subject who was methylenetetrahydrofolate reductase (MTHFR) 677TT homozygote lowered plasma Hcy from 33.3 µmol/l to 17.1 µmol/l following one-month Cr supplementation (5 g/day) opposite to 677CC and CT genotypes whose Hcy levels tended to increase (but still in normal ranges). We suppose that Cr supplementation stimulates pathways leading to production of sarcosine which can serve to regenerate tetrahydrofolate (THF) to form 5,10-methylene-THF. This could potentially increase MTHFR enzyme activity which may later result in increased Hcy methylation. Cr supplementation significantly effects metabolism of one carbon unit and potentially lower body’s demands for methyl groups. This could be beneficial as in the case of reduced enzyme activity such as MTHFR 677C/T polymorphism.

Key words
Creatine (Cr) • Homocysteine (Hcy) • Supplementation • MTHFR gene • 677C/T

Introduction
Creatine (Cr) supplementation has been widely used in athletes to support recovery in short term bouts of high-intensity exercise enabling more effective training and performance. Beneficial effect of Cr supplementation in young, healthy males is the enhanced muscle fiber size, strength and increased lean body mass (Greenhaff et al. 1994, Kreider et al. 1998). The greatest improvements in performance have been found in activities, which possibly stress the phosphocreatine (PCr) system such as series of high-power output exercises (Terjung et al. 2000). Studies have shown improved anaerobic exercise performance (Law et al. 2009), and specific performance in many sports such as fin swimming (Juhasz et al. 2009), swimming, all-out cycling, sprinting, repeated jumping, and resistance training (Juhn and Tarnopolsky 1998).

The potential therapeutic value of Cr supplementation has recently been investigated with respect to various neurodegenerative disorders which include; Alzheimer’s, Parkinson’s, amyotrophic lateral sclerosis, and Huntington’s disease patients (Adhihetty and Beal 2008). In patients with diseases that result in atrophy or muscle fatigue secondary to impaired energy production there have shown beneficial effects from Cr supplementation. Cr supplementation has also been investigated in various neuromuscular diseases including mitochondrial cytopathies, neuropathic disorders, dystrophies, congenital myopathies, and inflammatory myopathies (Tarnopolsky and Martin 1999).
The daily turnover of Cr (1-2 g/day) is replaced either through dietary Cr intake (from animal-derived tissues such as meat) or through endogenous synthesis in the liver from amino acid precursors (arginine, glycine and methionine). At the same time and approximately the rate (2 g/day), Cr is broken down to creatinine and excreted to the urine (Stead et al. 2006). The first step of Cr synthesis is the formation of guanidinoacetate (GAA) and ornithine in a reaction catalyzed by glycine amidinotransferase (AGAT), GAA can then be methylated on the original glycine nitrogen using S-adenosylmethionine (SAM) as the methyl donor. This reaction yields Cr and S-adenosylhomocysteine (SAH) and is catalyzed by the enzyme guanidinoacetate N-methyltransferase (GAMT) (Wyss and Kaddurah-Daouk 2000). The rate-limiting step in Cr synthesis is the formation of guanidinoacetate by AGAT and it has been shown that Cr supplementation down-regulates AGAT expression (Guthmiller et al. 1994). An increase in serum levels of Cr resulted in a decrease in AGAT enzyme activity, enzyme level, and mRNA expression in rat kidney (McGuire et al. 1984). On the other hand, the growth hormone up-regulates AGAT expression in rats (Guthmiller et al. 1994). AGAT is highly active in the kidneys, whereas GAMT is highly active in the liver. Therefore it is suggested that GAA is synthesized primarily in the kidney and then transported to the liver where it is methylated to form Cr (Fig. 1) (Wyss and Kaddurah-Daouk 2000). In rats, Van Pilsum et al. (1972) documented high activity of AGAT apart from kidney in pancreas, brain, spleen, and testes. In a study Edison et al. (2007) authors suggested that renal GAA production in a human may account only for about 20% of total GAA synthesis.

Fig. 1. Biochemical pathways related to Cr metabolism and methionine cycle. Creatine supplementation leads to inhibition of its endogenous synthesis. Simultaneously PEMT and GNMT activity is stimulated leading to produce choline, betaine and sarcosine respectively. Production and degradation of sarcosine utilize THF to form 5,10-methylene-THF stimulating MTHFR activity (bold arrows). GAA, Guanidinoacetate; AGAT, Glycine amidinotransferase; SAM, S-adenosylmethionine; SAH, S-adenosylhomocysteine; GAMT, guanidinoacetate N-methyltransferase; PEMT, Phosphatidylethanolamine N-methyltransferase; GNMT, Glycine N-methyltransferase; PC, Phosphatidylcholine; MTR, Methionine synthase; MTRR, Methionine synthase reductase; MTHFR, Methyleneetetrahydrofolate reductase; BHMT, Betaine homocysteine methyltransferase; CHDH, Choline dehydrogenase; THF, Tetrahydrofolate; SARDH, Sarcosine dehydrogenase; DMG, Dimethylglycine; Met, Methionine; AHCY, S-adenosylhomocysteine hydrolase; P, Phosphatidylcholine; MTR, Methionine synthase; MTRR, Methionine synthase reducetase; MTHFR, Methyleneetetrahydrofolate reductase; BHMT, Betaine homocysteine methyltransferase; CHDH, Choline dehydrogenase; THF, Tetrahydrofolate; SARDH, Sarcosine dehydrogenase; DMG, Dimethylglycine; Met, Methionine; AHCY, S-adenosylhomocysteine hydrolase.
Cr metabolism disorders have so far been described at the level of two synthetic steps, GAMT and AGAT, and at the level of the Cr transporter 1 (CrT1). The most common GAMT and AGAT deficiency symptoms and signs are delayed language development, learning disorders, autistic behavior, epileptic seizures, and movement disorders (Gordon 2010). GAMT and AGAT deficiency are treatable by oral Cr supplementation, but patients with Cr transporter deficiency do not respond to this type of treatment (Evangelou et al. 2009). Neither nine months of L-arginine supplementation did not showed effectiveness in the four patients affected with Cr transporter deficiency (Fons et al. 2008).

In a human body, more than 90 % of SAM is used for methylation reactions by at least 50 different methyltransferases (Stead et al. 2006). There are three major methyltransferases playing an important role in the generation of homocysteine (HCy) and regulation of methyl group metabolism; except for GAMT they include phosphatidylethanolamine N-methyltransferase (PEMT), and glycine N-methyltransferase (GNMT) (Mudd et al. 2007) (Fig. 1). Production of Cr via GAMT and phosphatidylcholine (PC) via PEMT is considered to be the largest consumer of methyl groups derived from SAM. Stead et al. (2006) suggests opposite of original investment where GAMT has been proposed to consume up to 70 % of methyl groups (Mudd et al. 1980), and considers PEMT to be the primary consumer of methyl groups, having the greatest impact on HCY levels. It was found that PEMT –/- mice have lower choline pools in liver despite being fed sufficient or supplemental amounts of dietary choline (Zhu et al. 2003). When PEMT is deleted in mice, plasma HCY concentrations fall 50 % and, when it is over expressed, plasma HCY concentrations increase 40 %, demonstrating that PEMT activity is a very major consumer of SAM (and thereby a producer of HCY) (Jacobs et al. 2005). Furthermore, methylation of macromolecules such as DNA, RNA, histones, and other proteins play critical roles in cellular metabolism. Therefore, the level of SAM must be carefully regulated to maintain cellular homeostasis (Luka et al. 2009).

The serum concentration of HCY is positively associated with the risk of ischemic heart disease, deep vein thrombosis and pulmonary embolism, and stroke (Boushey et al. 1995). A meta-analysis by Boushey et al. (1995) of 27 studies showed that HCY was an independent, graded risk factor for atherosclerotic disease. Total HCY (tHCy) measured in blood is usually the sum of free (reduced) HCY and protein-bound HCY (Brosnan et al. 2004). There are two methylation pathways to form methionine from HCY, both of which result in lowering HCY concentrations (Olthof et al. 2003). In the first, vitamins B12 and folic acid are involved in a reaction catalyzed by methionine synthase (MTR) (Weisberg et al. 2001). Deficiencies of these vitamins can result in elevated plasma HCY concentrations (Bailey et al. 2002, He et al. 2010). Also single nucleotide polymorphisms in genes coding enzymes involved in this pathway can result in elevated HCY in blood. Examples are substitutions as MTR 2756A/G (Barbosa et al. 2008), methionine synthase reductase (MTRR 66A/G) (Naushad et al. 2008), methylenetetrahydrofolate reductase (MTHFR 677C/T and 1298A/C) (Weisberg et al. 2001), or methylenetetrahydrofolate dehydrogenase (MTHFD1 1958G/A) (Brody et al. 2002). The most profound effect on HCY levels in blood was found in MTHFR 677C/T and/or in MTR 2756A/G (Barbosa et al. 2008). The 677C/T variant which leads to the substitution of Ala-222 by valine produces MTHFR enzyme with reduced activity, resulting in an elevation of serum HCY concentrations of about 20 % (Brattstrom et al. 1998). The 677C/T polymorphism is surprisingly common, with about 10 % of people in the population being homozygous affected (TT), 47 % homozygous unaffected (CC), and 43 % heterozygotes (CT) (Brattstrom et al. 1998). In the Czech population similar frequencies were elicited (CC – 41 %, CT – 49 %, TT – 10 %) (Vesela et al. 2005).

An association with hyperhomocysteinemia was also found in MTHFR 1298A/C polymorphism (Barbosa et al. 2008), although its contribution seems to be lesser compared to 677C/T (Bailey et al. 2002, Barbosa et al. 2008). In additional studies, the activity of enzyme produced by the heterozygotes for both mutations was lower compared to individuals who carried only 677T/T (Chango et al. 2000). The genetic influence of the MTHFR polymorphism on HCY levels is attenuated in females in premenopausal age and is not significant in subjects who exhibit serum levels of folate and/or vitamin B12 above the 50th percentile of distribution in the general population (Cortese and Motti 2001). A decrease in serum HCY of 3 mmol/l (achievable by daily intake of about 0.8 mg folic acid) should reduce the risk of ischemic heart disease by 16 %, deep vein thrombosis by 25 % and stroke by 24 % (Wald et al. 2002).

In the second pathway, the methylation of HCY to form methionine is catalyzed by betaine homocysteine
methyltransferase (BHMT) which is in addition to choline dehydrogenase (CHDH), and PEMT the most important enzyme in choline metabolism (da Costa et al. 2006). Betaine, once formed from choline via CHDH, donates its methyl group to Hcy via BHMT to form methionine. Administration of betaine can lower plasma Hcy concentrations (Steenge et al. 2003). Activity of BHMT is increased during methionine excess and plasma betaine was shown to be a strong determinant after methionine increase in tHCy in subjects not supplemented with B-vitamins (Holm et al. 2004). On the other hand, neither a common variant 742G/A, nor other variants in BHMT gene seem to play a significant role in plasma Hcy (Heil et al. 2000, Morin et al. 2003).

The aim of the present post-hoc analysis is to evaluate homocysteinemia following Cr supplementation in relation to MTHFR 677C/T genotype.

Methods

Subjects

This is a secondary analysis of data from 11 athletes participating in our previous study (Navratil et al. 2010) who provided DNA from buccal cells. Of those participants, 10 submitted DNA testing. All subjects were young, aged 24-28 years old, healthy, physically active persons, dealing with sportive activities (ice hockey, football, horsemanship, and athletics) on a professional level. Subjects' height, weight, and body composition via Bioelectrical Impedance Analysis (Multi-frequency analyzer In Body 3.0, Korea) was measured. All 10 men were Caucasian. Subject characteristics are shown in Table 1. Written informed consent was obtained from all subjects under protocols approved by the Institutional Ethics Committee of the Charles University of Faculty of Physical Education and Sport.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>24.6±2.1</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>181.7±4.8</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>82.8±11.0</td>
</tr>
<tr>
<td>Body fat (%)</td>
<td>12.0±2.9</td>
</tr>
<tr>
<td>BMI (kg.m⁻²)</td>
<td>25.0±2.4</td>
</tr>
<tr>
<td>Fat Free Mass (kg)</td>
<td>75.6±8.0</td>
</tr>
</tbody>
</table>

Values are means ± SD.

Intervention

As previously published, all subjects ingested 5 g of CR-monohydrate (Plutino, Czech Republic) a day, diluted in tepid water. These doses were administered every morning (at about 8 a.m.) for 30 days. Participants were not allowed to consume any other supplement, especially those containing vitamin B and folic acid, and they were advised to maintain their usual dietary habits and physical activity during the study.

Biochemical assays

Fasting blood and urine samples were collected at baseline and the next morning after completion 30-day Cr supplementation and further analyzed for several metabolites including Hcy as described previously (Navratil et al. 2010, Petr et al. 2011).

Genotyping

Genomic DNA was isolated from buccal cells collected with cheek brushes (Whatman, USA). Samples were lysed and DNA was stabilized with DNA Extract All Reagents Kit (Applied Biosystems, USA) according to the manufacturer's protocol. Samples (5 μl) were genotyped according to the manufacturer's protocol on an Illumina BeadStation 500G Golden Gate genotyping platform using a custom panel (GS0011351-OPA) of 384 candidate single-nucleotide polymorphisms. From the whole set the genotypes for the MTHFR C677T polymorphism (rs1801133) were extracted.

Results

Of 10 subjects, 9 individuals were carrying 677CC+CT, and 1 individual the 677TT genotype. Pre-test levels of plasma Hcy were normal among those carrying 677CC+CT genotype (6.3±1.3 μmol/l), but strongly elevated in 677TT carrier (33.2 μmol/l). After 30-day Cr supplementation individuals with 677CC+CT genotype mildly elevated Hcy levels, but completely different response was registered in 677TT carrier who lowered Hcy almost to normal levels (Table 2).
In spite of that, we assume our outcomes to be interesting, as they should contribute to better understanding how supplemented Cr can affect methyl donor balance and consequent plasma HCy levels.

Table 2. Pre-test and post-test HCy levels in different MTHFR 677C/T genotypes.

<table>
<thead>
<tr>
<th></th>
<th>677CC (n=4)</th>
<th>677CT (n=5)</th>
<th>677TT (n=1)</th>
<th>677CC+CT</th>
</tr>
</thead>
<tbody>
<tr>
<td>HCY (µmol/l)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>pre-test</td>
<td>5.9±1.3</td>
<td>6.6±1.3</td>
<td>33.2</td>
<td>6.3±1.3</td>
</tr>
<tr>
<td>post-test</td>
<td>9.9±2.9</td>
<td>11.6±3.3</td>
<td>17.1</td>
<td>10.9±3.2</td>
</tr>
</tbody>
</table>

Values are means ± SD.

A meta-analysis of published cohort studies states that hyperhomocysteinemia moderately increases the risk of a first cardiovascular event, regardless of age and follow-up duration (Bautista et al. 2002). In homocystinuria, which is a rare inherited disorder (most often due to cystathionine β-synthase deficiency, which occurs in 1 from 200,000 people), plasma HCY levels are markedly elevated (>50 µmol/l; normal range, 5 to 15 µmol/l), and patients have severe, widespread vascular disease (Bellamy et al. 1998). In the general population, mild to moderate elevations in plasma HCY (15 to 35 µmol/l) are common and may occur due to inherited enzyme variants and/or a relative deficiency of folate, vitamin B₁₂, or vitamin B₆, which are required for the normal metabolism of HCY (Ubbink et al. 1993).

Our subjects with CC and CT genotypes had pre-test HCY concentration in normal ranges 6.1±1.3 µmol/l with milder individual differences. The only carrier of TT genotype had HCY (33.3 µmol/l). After 30-day Cr supplementation all CC and CT carriers increased plasma HCY to 10.9±3.2 µmol/l opposite to TT carrier who significantly lowered HCY levels to 17.1 µmol/l. An interpretation of these changes is not simple to explain. Reliable description of addressed biochemical pathways should be supported by analysis of associated intermediates e.g. sarcosine in urine.

We assume that supplemented Cr inhibited the synthesis of endogenous Cr and thus increased availability of SAM for PC and sarcosine synthesis via PEMT and GNMT respectively. Concomitantly, the methylation of HCY relied more on BHMT where methyl donor is betaine, stimulated by increased production of PC and choline via increased PEMT activity. Although PEMT is considered to be a major consumer of SAM and a producer of HCY by some authors (Jacobs et al. 2005), provided that PC and choline levels are sufficient from diet, HCY methylation from betaine could be stimulated. Moreover, resulting dimethylglycine demands tetrahydrofolate (THF) for sarcosine formation via dimethylglycine-dehydrogenase. THF is also needed for glycine formation from sarcosine via sarcosine-dehydrogenase (SARDH). Sarcosine production can also be accelerated due to better availability of SAM for GNMT. Based on these facts TFH is dynamically converted to 5,10-methylene-THF (methylene tetrahydrofolate) stimulating MTHFR enzyme to 5-methyl-THF production which is later used for HCY conversion to methionine via MTR. Higher availability of 5,10-methylene-THF could possibly effectively stimulate MTHFR even in 677TT carriers. This can have quite a strong effect on HCY levels as we registered in our one 677TT carrier.

In fact, studies evaluated effects of Cr on HCY levels are not consistent. Cr administration has been shown to decrease plasma tHCy by 25 % in rats (Stead et al. 2001) and by 10 % in humans, according to one report (Korzun 2004); in another report, this reduction in humans was not significant (Steenge et al. 2001). Opposite to these results, Cr supplementation (alone or in combination with L-arginine) was associated with an 11-20 % increase in HCY concentration, which was not attributable to worsened renal function, providing evidence against an effect of Cr on decreasing methylation demand (Jahangir et al. 2009). In rats, plasma HCY was increased up to 2 h after intense anaerobic exercise, but Cr supplementation decreased plasma HCY independent on exercise intensity (Deminice et al. 2011). Recent study shows interesting results about the prevalence of hyperhomocysteinemia (>15 µmol/l) in elite athletes which was 47 % compared to 17 % in...
controls without any correlation between HCy and any of
the other investigated variables, including plasma folate, vitamin B12, blood pressure, lactate dehydrogenase (LDH), creatine kinase (CrK), total and high-density lipoprotein (HDL) cholesterol and interleukin-6 (IL-6) (Borrione et al. 2008).

Our findings demonstrate an increase of HCy following Cr supplementation in MTHFR 677C allele carriers but an average and individual augmentation were all in normal ranges (<15 µmol/l). We suppose, when levels of dietary choline (PC) are sufficient to cover physical needs, a great part of choline from endogenous production of PC should be used to form betaine for further HCy methylation. Provided that choline from diet does not meet body demands, endogenous choline production (via PEMT) is very important. Under these conditions most probably less betaine can be produced to methylate HCy. Pathways following methylation with betaine seem to have a secondary methylation potential in regeneration of THF to form 5,10-methylene-THF. Cr supplementation significantly effects metabolism of one carbon unit and potentially lower body demands for methyl groups. This could be beneficial as in the case of reduced enzyme activity like MTHFR 677C/T polymorphism.

Conflict of Interest
There is no conflict of interest.

Acknowledgements
This study was supported by research grant No. MSM 002160864, SVV 2013-267603 and PRVOUK n. 38 founded by Faculty of Physical Education and Sport, Charles University in Prague.

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