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Is ATP Elevated in Patients With GAMT Deficiency?

We read with interest the paper by Bianchi et al,¹ in which the authors report on the treatment monitoring of 2 patients with guanidinoacetate methyltransferase (GAMT) deficiency and 3 patients with an arginine:glycine amidinotransferase defect (AGAT-d). Repetitive MR measurements of these patients are not trivial, and some interesting observations are presented in this paper that may be very useful for their treatment.

Bianchi et al¹ studied the levels of brain creatine (Cr) and phosphocreatine in 1 patient with GAMT deficiency using ³¹P- and ¹H-MR spectroscopy (phosphorous and proton MR spectroscopy) before and after therapy. The authors conclude from the ³¹P-MR spectra that levels of adenosine triphosphate (ATP) are elevated in the brain of this patient before therapy, and that these levels normalize with treatment. Furthermore, they speculate that this may be the result of redistribution in the creatine kinase (CK) reaction ($\text{ATP} + \text{Cr} \rightarrow \text{ADP} + \text{PCr}$) from the reduced levels of Cr in the brains of other patients.¹ In our opinion, this conclusion on increased ATP and the speculation on its origin are debatable.

Tissue levels of ATP are tightly balanced by consumption and production of ATP. Because the brain is a highly oxidative organ, most production of ATP occurs by mitochondrial oxidative phosphorylation, though other processes such as glycolysis also contribute. The primary function of the CK reaction is more to dampen large fluctuations in the ratios of ATP and adenosine diphosphate (ADP) during periods of high energy demand,² all in close relationship to the energy-producing and energy-consuming sites. Therefore, it is unlikely that a mere shift in the concentration of Cr will result in the drastic, global elevation of ATP as reported.

According to Bianchi et al,¹ the supposed elevation of ATP is based on the ³¹P-MR spectrum of GAMT; in patient 1, in particular, the signal intensity for β -ATP (at 16.5 ppm) is increased. However, ATP has 3 phosphate groups, all giving a distinct signal intensity at different positions on the ³¹P-MR spectrum. If ATP were elevated, why are not all 3 signals equally elevated, since they belong to the same molecule? Although the other ATP signals (α -ATP and γ -ATP) also represent other phosphorus compounds such as ADP, these are present at much lower concentrations ($\sim \mu\text{M}$ range) than ATP ($\sim \text{mM}$). Therefore, changes in the levels of these compounds cannot account for a selective elevation in the signal intensity of β -ATP. We believe that the increase in the signal intensity of β -ATP may be because of experimental issues in the ³¹P-MR spectroscopy method. Factors such as incorrect phasing of the MR spectrum, the absence of a first-order phase correction, and insufficient bandwidth of the excitation pulse in combination with a misplaced carrier resonance could all play a role here. Unfortunately, Bianchi et al¹ did not provide sufficient information on data acquisition and quantification of the ³¹P-MR signals to resolve this issue.

Finally, the authors state that my colleagues and I,³ in GAMT-deficient mice, and Schulze et al,⁴ in a patient, display the same abnormality in the signal intensity of ATP and that both my colleagues and I and Schulze et al failed to provide a comment on this. However, in our opinion, our ³¹P-MR spectra of the GAMT-deficient mouse brain did not reveal a similar abnormality in the signal intensity of β -ATP,³ as Bianchi et al¹ observed. As indicated in the legend to Fig 6 of our paper, the ³¹P-MR spectra are scaled arbitrarily. Also, in the paper of Schulze et al,⁴ it seems that signal intensity is scaled arbitrarily. At least when the reference signal intensity (ie, phosphodiester signal intensity) applied by Bianchi et al¹ is used, no significant change in the signal intensity of ATP is observed. Therefore, in our opinion,

the ATP increase in GAMT deficiency before treatment with Cr is not supported by other studies.

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Reply:

On behalf of all of the authors of our study,¹ we thank Dr Renema for his interest in our work and we appreciate his comments. We are fully aware that other studies demonstrate that the cellular pools of adenosine triphosphate (ATP) are very stable during tissue activation and that they are regulated not only by the creatine kinase (CK) system but also by mitochondrial oxidative phosphorylation, glycolysis, and the adenosine monophosphate-cyclase system. Most of these studies have been performed on muscles, and only few data are available for the brain in which the control, coupling, and kinetics of the phosphocreatine/CK/ATP system are much more complex.^{2,3} Acquisition and postprocessing of our ³¹P-MR spectroscopy studies were carried out according to standard procedures for human brain studies. The increased signal intensity of the ATP peaks in the brain of the child with the guanidinoacetate methyltransferase defect (GAMT-d; Fig 2 of our original article) was unique. Due to the different arbitrary units used by Renema et al² and Schulze et al,⁴ we have probably erroneously referenced these articles as supporting our results. Therefore, we believe that the question raised by Renema is still waiting to be confirmed by further ³¹P-MR spectroscopy studies of patients with GAMT-d aimed at measuring absolute concentration of brain ATP.

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