Use of *in vivo* magnetic resonance spectroscopy for studying metabolic diseases

Jong-Hee Hwang and Cheol Soo Choi

Owing to the worldwide obesity epidemic and the sedentary lifestyle in industrialized countries, the number of people with metabolic diseases is explosively increasing. Magnetic resonance spectroscopy (MRS), which is fundamentally similar to magnetic resonance imaging, can detect metabolic changes *in vivo* noninvasively. With its noninvasive nature, $^1$H, $^{13}$C and $^{31}$P MRS are being actively utilized in clinical and biomedical metabolic studies to detect lipids and important metabolites without ionizing radiation. $^1$H MRS can quantify lipid content in liver and muscle and can detect other metabolites, such as 2-hydroxyglutarate, *in vivo*. Of interest, many studies have indicated that hepatic and intramyocellular lipid content is inversely correlated with insulin sensitivity in humans. Thus, lipid content can be utilized as an *in vivo* biomarker for detecting early insulin resistance. Employing $^{13}$C MRS, hepatic glycogen synthesis and breakdown can be directly detected, whereas $^{31}$P MRS provides *in vivo* adenosine triphosphate (ATP) synthesis rates by saturation transfer methods in addition to ATP content. These *in vivo* data can be very difficult to assess by other methods and offer a critical piece of metabolic information. To aid the reader in understanding these new methods, fundamentals of MRS are described in this review in addition to promising future applications of MRS and its limitations.

**INTRODUCTION**

Diabetes mellitus is one of the most common metabolic diseases. According to the World Health Organization (WHO), 347 million people worldwide currently have been diagnosed with diabetes. In industrialized countries with a sedentary lifestyle, the number of diabetic patients is expected to increase at a higher rate. Insulin resistance is a good predictor for the clinical onset of type 2 diabetes mellitus (T2DM) and precedes the disease by many years. To detect early metabolic changes such as developing insulin resistance, noninvasive magnetic resonance (MR) techniques can be an ideal tool for noninvasively monitoring alterations in insulin sensitivity using the lipid content of the liver or muscle as a biomarker in humans and animal models.$^{1-3}$ With the advantage of its noninvasiveness, nuclear magnetic resonance (NMR) has evolved as a powerful medical imaging tool since the development of MR imaging (MRI) in the early 1970s by Lauterbur and Mansfield.$^{4,5}$ *In vivo* MR spectroscopy (MRS) shares the same fundamentals as MRI and has developed in parallel. It is used to investigate the details of biochemistry and physiology in metabolic diseases in addition to the morphological information assessed by MRI.

To examine metabolic disorders, many clinical and biomedical studies have been conducted using MRS methods, for example, the quantification of hepatic lipid and glycogen content as well as tricarboxylic acid (TCA) cycle or adenosine triphosphate (ATP) synthesis rates *in vivo* in real time using $^1$H, $^{31}$P and $^{13}$C MRS. These data are difficult to assess by other modalities. Together these noninvasive methods can provide crucial information for metabolic studies and clinical research. For clinical studies, the noninvasiveness is a prime advantage of the methods; however, for preclinical research, the ability to perform longitudinal studies without killing animals adds value due to the high cost of transgenic and knockout mice. Thus, this noninvasive approach also helps to reduce the number of animals being killed and to eliminate interanimal variances by longitudinal follow-up of the same animal.

In this review, the most commonly used and promising MRS methods are described, including $^1$H, $^{31}$P and $^{13}$C MRS, with a particular emphasis on metabolic diseases.

---

1Korea Mouse Metabolic Phenotyping Center (KMMPC), Lee Gil Ya Cancer and Diabetes Institute, Gachon University, Incheon, Korea and 2Division of Endocrinology, Department of Internal Medicine, Gachon University Gil Medical Center, Incheon, Korea

Correspondence: Professor J-H Hwang or Dr CS Choi, Department of Molecular Medicine, Lee Gil Ya Cancer and Diabetes Institute, Gachon University, Yeonsu-Gu, Songdo-Dong 7-45, Incheon 406-840, Korea.

E-mail: jonghtwang@gachon.ac.kr or cschoi@gachon.ac.kr

Received 14 October 2014; accepted 29 October 2014
GENERAL OVERVIEW OF MRI AND MRS: SIMILARITIES AND DIFFERENCES

MRI and MRS data can be obtained using a spectrometer, as shown in Figure 1a, or MRI scanners in hospitals. Among all nuclei, only nonzero nuclear spins ($I \neq 0$) are detectable by NMR, for example, $^1$H, $^{31}$P and $^{13}$C. As shown in Figure 1b, without a magnetic field (when an animal is outside the range of a magnet), all of the spins are randomly oriented. However, within a magnetic field (inside of a magnet, Figure 1c), all of the spins are aligned with or against the field. In other words, nuclear spin energy levels are quantized at different energy levels. When the magnetic fields of all spins are summed, a net longitudinal magnetization is generated parallel to the magnetic field, as shown in Figure 1c.

A radiofrequency coil is a special device that can emit energy and receive signals. A small amount of energy is applied to this net magnetization, causing the magnetization to flip to another orientation, as demonstrated in Figure 1d. However, as soon as this transition occurs, the magnetization tries to return to its original orientation (the thermal equilibrium state as in Figure 1c) with precession. While in precession, the magnetization also emits energy (d). This energy is detected by a receiver, which generates nuclear magnetic resonance signals.

![Figure 1](image_url)

**Figure 1** (a) Bruker 9.4T magnetic resonance spectrometer in Lee Gil Ya Cancer and Diabetes Institute, Gachon University. (b, c) Without a magnetic field (outside of the magnet), all of the spins are randomly oriented (b), but within a field (inside of the magnet), all of the spins are aligned with or against the field (quantized at different energy levels, c). When all of the spins are combined, a longitudinal magnetization is generated parallel to the magnetic field. (d) When a small amount of energy is applied to the magnetization by a radiofrequency (r.f.) coil, the magnetization is flipped to another orientation. However, the magnetization tries to return to its original position (the thermal equilibrium (c)) with precession. While in precession, the magnetization also emits energy (d). This energy is detected by a receiver, which generates nuclear magnetic resonance signals.

MRI and MRS: their similarities and differences

Proton ($^1$H) MRI and MRS are the most widely utilized methods in metabolic diseases because the signal-to-noise ratio (SNR) of proton MRS/MRI is generally higher than other nuclei. Although the cost for $^1$H MRS/MRI examinations is higher than computed tomography and ultrasound, this modality is widely used due to the accurate assessment of anatomy and metabolites, such as liver lipids expressed as a percentage by $^1$H MRS.

As previously mentioned, MRI and MRS share a common physical background. MRI is an imaging technique employing NMR, while MRS also uses NMR, but is spectroscopy.
As demonstrated in Figure 4b, separate water and lipid signals for insulin sensitivity. The principle of this method is simple. 1H MRS for hepatic lipid quantification. Related research. Techniques provide important information for medicine and morphological information can be assessed. Thus, these are combined, biochemical data of a specific location with morphological information can be assessed. Thus, these techniques provide important information for medicine and its related research.

1H MRS

1H MRS for hepatic lipid quantification

Currently, hepatic lipid content by 1H MRS is the gold standard measurement and is often utilized for clinical trials to develop new hepatic lipid-lowering drugs and employed in preclinical studies. This technique has received attention in the medical community particularly due to several pragmatic applications. Liver and intracellular muscular lipid content reportedly correlate with insulin resistance, which is the best predictor for the clinical onset of T2DM. Thus, lipid content, as assessed by 1H MRS, can serve as a surrogate in vivo marker for insulin sensitivity. The principle of this method is simple. As demonstrated in Figure 4b, separate water and lipid signals are easily detected and quantified by 1H MRS. Lipid signals can be converted to percent (%) lipid relative to water. MRI is also used in addition to this 1H MRS method to quantify adipose tissue volume such as visceral adipose tissue (sagittal view (Figure 4c) and transverse view (Figure 4d)), which cannot be easily assessed by other imaging methods, for example, dual-energy X-ray absorptiometry.

1H MRS for intramyocellular lipids

Another major advantage of muscular lipid measurements by 1H MRS is the intramyocellular (IMCL) and extramyocellular (EMCL) lipid quantitation without the manual cleaning of EMCL adipose tissue. This separation of the two components is achieved by a bulk magnetic susceptibility effect in soleus and tibialis anterior muscles (Figures 5a and b), resulting in distinct chemical shifts between IMCL and EMCL due to the different characteristics of fat states, that is, IMCL lipids have a spherical shape (like a lipid droplet), and EMCL lipids form in fiber-like bundles (like a semisolid), which show a strong orientation in parallel with the muscle fibers. Thus, this modality provides not only a noninvasive approach, but it also removes potential quantification errors that originated from incomplete manual EMCL cleaning. Numerous studies have demonstrated a correlation of hepatic lipid content and IMCL with insulin resistance. Furthermore, IMCL assessment by 1H MRS can also be utilized as an indicator of muscular lipid metabolism during exercise.

Compositional analysis of fat by 1H MRS

Recently, fat quantity and composition are receiving attention because of their relevance to metabolism and cardiac diseases. Some previous publications noted that saturated and unsaturated fats can have a different impact on metabolism and are potentially related to cardiac disorders. Thus, noninvasive assessments of fat composition by MRS could also become a powerful tool in relevant research. In tissues where lipid concentrations are very high, such as in bone marrow, fat compositional analysis by 1H MRS can be powerful. More discussion on this topic will be described in the 13C MRS section.

31P MRS

Overview of 31P MRS

Historically, 31P MRS has been utilized for the detection of ATP and phosphocreatinine (PCr) as early as MRI, in the 1970s. Next to protons, 31P nuclei demonstrate a high SNR due to the 99% natural abundance of the 31P isotope. Rates of ATP synthase and forward creatine kinase activity can be assessed in humans and animals using magnetization transfer 31P MRS. Recently, ATP and other phosphorus metabolite quantities have been related to metabolic alterations, such as those that occur in insulin-resistant T2DM. Moreover, using saturation transfer 31P MRS, ATP synthesis rates were reportedly reduced in muscles from the elderly and in the offspring of T2DM patients. Thus, both absolute phosphorus metabolite quantities and kinetic information are
Magnetic Resonance Spectroscopy (MRS) vs. Magnetic Resonance Imaging (MRI)

**Figure 3** Comparison between MRS and MRI. (a) For MRS, a schematic explanation on how to obtain an MR spectrum and its example in the human brain. Time domain data (free induction decay: FID) were acquired and they were converted to frequency domain data through fast Fourier transformation (FFT). The resulting spectrum demonstrated a peak/peaks as shown in the example of a tumor. Within the tumor, choline (Cho), creatine (Cre), NAA(N-acetyl aspartate) and lactate doublet peaks are clearly demonstrated. (b) For MRI, a schematic explanation for how to obtain an MRI is provided. 2D/3D time domain data were converted to 2D/3D spatial domain data via FFT. The resulting images can exhibit T1- or T2-weighted contrast depending on the sequence and the parameters employed. MRI, magnetic resonance imaging; MRS, magnetic resonance spectroscopy; 2D, two dimensional; 3D, three dimensional.

**Figure 4** (a) Axial image of the liver and the voxel where spectroscopic measurements were obtained. (b) ¹H-magnetic resonance (MR) spectrum from the liver clearly shows the methylene peak of triglycerides. (c) Sagittal image of an abdomen; the lumbar spine was set as a landmark to select the location for subcutaneous and visceral fat analysis in transverse images. Lumbar spine L3 and L4 are denoted. (d) Transverse image of an abdomen for fat analysis. Subcutaneous and visceral fat were separated (bottom) and the fat content was quantified after converting those images to black-and-white binary images.¹
useful to examine changes in metabolism, aging and diseases by assessing enzyme activities and mitochondrial function.\textsuperscript{19–24} \textsuperscript{31}P MRS has also been utilized in animal models to evaluate the effect of diets on animal models.\textsuperscript{24,25} Another key player in energy metabolism, nicotinamide adenine dinucleotide, can also be detected and analyzed using \textsuperscript{31}P MRS.\textsuperscript{26}

Quantification of phosphorus metabolites and their role as biomarkers

In \textsuperscript{31}P MRS, many phosphorus metabolites can be detected, including ATP, PCr, phosphocholine, glycerophosphocholine, glucose-6-phosphate (G6P) and others.\textsuperscript{27} A family of phosphomonoesters was detected at ~6.4–6.6 p.p.m. in Figure 6b, which includes G6P. G6P is an important intermediate for glucose oxidation. There is a defect in glycogen synthesis in muscles of T2DM patients.\textsuperscript{28} To find the mechanism(s) behind the defect, Shulman et al measured G6P and found lower G6P concentrations in muscles of T2D patients using \textsuperscript{31}P MRS. From the data, they concluded that there was a defect in glucose transport and/or phosphorylation in T2DM patients.\textsuperscript{29} Later, a problem in glucose transport was detected in T2DM using \textsuperscript{31}P MRS with additional studies.\textsuperscript{30}

Traditionally, phosphorus metabolites have been well utilized to examine myopathy. For example, Barnes et al.\textsuperscript{31}

CONVENTIONAL \textsuperscript{13}C MRS AND HYPERPOLARIZED \textsuperscript{13}C DETECTION

General overview of \textsuperscript{13}C MRS

The stable carbon isotope \textsuperscript{13}C is NMR detectible but naturally occurs at only 1.1% among all carbon isotopes. Thus, the intrinsic SNR for \textsuperscript{13}C is much lower than \textsuperscript{1}H and \textsuperscript{31}P MRS. However, \textsuperscript{13}C MRS has a wide chemical shift range, which makes many different carbons easily distinguishable. This characteristic adds to the value of \textsuperscript{13}C MRS, such as for glycogen detection. In \textsuperscript{1}H MRS, the differentiation of protons attached to C-1 glucose and C-1 glycogen is very challenging due to the proximity of the two signals and overlapping nearby strong water signals. However, in \textsuperscript{13}C MRS, the separate detection of C-1 carbons in a glucosyl unit in glycogen and glucose are detectible due to the wide range of chemical shifts (Figure 7a). Moreover, the time course change in hepatic observed increased Pi, a reduced PCr/ATP ratio and elevated pH in muscles of patients with myotonic dystrophy, and Kemp et al.\textsuperscript{32} reported mitochondrial dysfunction in muscles using \textsuperscript{31}P MRS combined with exercise.
glycogen was clearly depicted in Figure 7b without any exogenous labeling. Due to low endogenous natural abundance of $^{13}$C, an exogenously labeled $^{13}$C signal can be easily detectible, enabling pulse-chase-type experiments to be performed. However, without exogenous labeling, $^{13}$C MRS struggles with low SNR. Recently, to overcome low SNR, hyperpolarizing methods have been introduced and demonstrate that SNR enhancement can be achieved by $10^3$–$10^4$ times in a few selected metabolites. This method potentially will resolve the low SNR issue and contribute to a further broadening of $^{13}$C MRS usage. This will be discussed more in the hyperpolarized $^{13}$C MRS section.

$^{13}$C MRS for glycogen detection
As C-1 liver and muscle glycogen is detectible, this method is utilized to quantify the muscular and hepatic glycogen content during nutrient ingestion or exercise in metabolic studies. Because hepatic glycogen concentration is reasonably high (300–400 mmol of glucosyl unit per liter of tissue in humans), this method can be performed by either natural abundance or exogenous $^{13}$C labeling. Using this method as described in Figures 7a and b, glycogen breakdown and synthesis rates were determined, and altered glycogen metabolism in T1D was identified.\(^{36,37}\) Glycogen storage disease can also be noninvasively identified using the same method.\(^{38}\) Furthermore, combining these studies with $^{13}$C glucose labeling, further kinetic studies were conducted to estimate liver glycogen rates.\(^{39}\)

Compositional lipid analysis
By taking advantage of the wide spectral range, many carbon signals in lipids also can be separately detected using $^{13}$C MRS.\(^{40}\) As demonstrated in previous publications, the effects on body fat composition by different types of diets were studied using natural abundance $^{13}$C MRS.\(^{40}\) Differences of each diet were clearly detected: fish oil (polyunsaturated fatty acid; Figure 8a), normal diet (Figure 8b) and Lorenzo oil (Figure 8c). Thus, $^{13}$C MRS can be utilized for noninvasively monitoring dietary effects.

$^{13}$C MRS for TCA cycle rates
Using exogenous labeled substrates such as C1 glucose and C1 acetate, labeled intermediates in the TCA cycle or relevant intermediates such as glutamate/glutamine can be measured by $^{13}$C MRS, and TCA cycle rates and other important fluxes can be assessed.\(^{41}\) Moreover, by combining saturated transfer $^{31}$P MRS and TCA cycling, muscle oxidative capacity was reduced in the offspring of T2D compared with controls.\(^{20,21}\)

Hyperpolarized $^{13}$C MRS
Despite many advantages of $^{13}$C MRS, this method suffers from low SNR due to its low natural abundance (1.1%) and low gyromagnetic ratio. Thus, increasing SNR is the prime goal for improving the utility of $^{13}$C MRS.

**Figure 7** $^{13}$C NMR spectra from the liver. (a) C-1 glycogen of a doublet peak at 100.1 p.p.m. clearly exhibited together with the lipid peaks from the hepatic lipid carbons. The inlayed trace is zoomed clearly demonstrating the C-1 glycogen doublet. (b) The time course spectra of glycogen peaks at 100.1 p.p.m. demonstrate a clear reduction of C-1 glycogen during a hypoglycemic clamp on a control subject. NMR, nuclear magnetic resonance.\(^{37}\)

**Figure 8** Dietary effect of PUFA and MUFA diets. The differences in PUFA levels are clearly delineated in peak 2 (double-bond peaks in PUFA) and 3 (methylene peaks in between two double bonds in PUFA) in response to the diets. (a) Spectrum from a subject on PUFA (fish oil) diet for >7 years. (b) Spectrum from a control subject who was not on any special diet. (c) Spectrum from a subject on a MUFA (Lorenzo’s oil) diet for 8 years.\(^{40}\) All of the PUFA peaks (2 and 3 peaks) were increased in a subject that had been on a fish oil diet compared with normal and MUFA diets with reference to peak 1 (all double-bond peaks, including MUFA). MUFA, monounsaturated fatty acid; PUFA, polyunsaturated fatty acid.\(^{40}\)
13C MRS users. By hyperpolarizing 13C spin distribution, the SNR of 13C MRS can be enhanced on the order of 10^3–10^4. With recent technical advances in nuclear dynamic polarizers, applications of hyperpolarized 13C MRS have been further demonstrated in animal models, particularly in cancer research due to rapid turnover to lactate with an injection of hyperpolarized C-1 pyruvate. There are two different approaches to hyperpolarize 13C nuclei: one approach dynamically polarizes 13C using a dynamic nuclear polarizer and the other approach utilizes chemically induced parahydrogens. Currently, most published in vivo studies have utilized a dynamic nuclear polarizer approach. In brief, this process artificially increases excitable 13C spins by 10^3–10^4 times; however, the polarized spins are unstable and will relax back to the thermal equilibrium state. Thus, all MR examinations with hyperpolarized substrates have to compete within an order of the T1 relaxation times to fully exploit the advantage of this method. Because tumor cells predominantly choose anaerobic processes, the most recent publications using hyperpolarized techniques described detecting anaerobic cycle products such as lactate. Because metabolic diseases do not normally demonstrate high concentrations of anaerobic products, aerobic products involving the TCA cycle are of more importance. A recent publication demonstrated that 2-oxyglutarate in the TCA cycle was detected using hyperpolarized pyruvate, which is the first evidence of utilization of this method in aerobic processes. In the future, the utility of this method might increase outside of oncology applications.

Other metabolic biomarkers using 1H and 13C MRS
As mentioned before, traditional 1H MRS has been used for various applications, for example, central nervous system diseases, cancer and metabolic diseases. Recently, as a new tumor biomarker, the detection of 2-hydroxylutarate has been reported by the elegant optimization of 1H MRS. Mutations in isocitrate dehydrogenases 1 and 2 relate to certain types of gliomas. Indeed, elevated 2-hydroxyglutarate was detected by 1H MRS in gliomas with isocitrate dehydrogenase mutations. Thus, one can further utilize 1H(13P/13C) MRS in the future for detecting specific biomarkers based on unusual enzymatic activities in diseases.

LIMITATIONS AND OUTLOOK
In addition to its noninvasive nature, 1H, 31P and 13C MRS methods have provided unique biochemical information that would be difficult to obtain by other methods. It is anticipated that these modalities will serve to develop more complete phenotypic data in humans and animal models, which will become crucial for personalized medicine in the future. Although MRS is costly compared with other imaging modalities, it has found its specific role in metabolic research due to its biochemical specificity. Furthermore, this method can also detect biochemical intermediates, which can connect biochemistry with systemic physiology, for example, G6P, 2-hydroxyglutarate.

The limitation of this method is that it is often not easy to provide standardized protocols in a clinical setting or animal model studies compared with MRI. These modalities also often require two-channel scanners for proton decoupling, whereas routine clinical MRI scanners are equipped with only one channel.

Moreover, due to low sensitivity, the acquisition time tends to be longer compared with MRI. However, with further technical development in hardware and software, the required time for these methods can be shortened in the future. New specific biomarkers for a disease/metabolic change can also be developed to extend the applications of MRS methods. Furthermore, effective combinations of these MRS methods with other modalities may bring more comprehensive and powerful examinations of metabolic changes.

CONFLICT OF INTEREST
The authors declare no conflict of interest.

ACKNOWLEDGEMENTS
This work was supported by the Bio & Medical Technology Development Program of the National Research Foundation (NRF) funded by the Ministry of Science, ICT & Future Planning (2013M3A9D5072563), the Korea Healthcare Technology R&D Project, Ministry for Health, Welfare & Family Affairs, Korea (A102060) and the Gachon University Gil Medical Center (Grant #2013-33).
et al.

12. Insull W, Lang PD, Hsi B, Yoshimura SJ. Studies of arteriosclerosis in Japanese and American men: I. Comparison of fatty acid composition of adipose tissue. J Clin Invest 1969; 48: 1313–1327.

13. Ren J, Dimitrov I, Sherry AD, Malloy CR. Composition of adipose tissue and marrow fat in humans by 1H NMR at 7 tesla. J Lipid Res 2008; 49: 2055–2062.

14. Strobel K, van den Hoff J, Pietzsch J. Localized proton magnetic resonance spectroscopy of lipids in adipose tissue at high spatial resolution in mice in vivo. J Lipid Res 2008; 49: 473–480.

15. Lee Y, Jee HJ, Noh H, Kang GH, Park J, Cho J et al. In vivo (1) H-MRS hepatic lipid profiling in nonalcoholic fatty liver disease: an animal study at 9.4 T. Magn Reson Med 2013; 70: 620–629.

16. Burt CT, Gionek T, Bárány M. Analysis of phosphate metabolites, the intracellular pH, and the state of adenosine triphosphate in intact muscle by phosphorus nuclear magnetic resonance. J Biol Chem 1976; 251: 2584–2591.

17. Gadian DG, Houtl DI, Radda GK, Seeley PJ, Chance B, Barlow C. Phosphorus nuclear magnetic resonance studies on normoxic and ischemic cardiac tissue. Proc Natl Acad Sci U S A 1976; 73: 4446–4448.

18. Radda GK, Bore PJ, Gadian DG, Ross BD, Styles P, Taylor DJ et al. 31P NMR examination of two patients with NADH-CoQ reductase deficiency. Nature 1982; 295: 608–609.

19. Szendroedi J, Chmelik M, Schmid AI, Nowotny P, Brehm A, Krssak M et al. Abnormal hepatic energy homeostasis in type 2 diabetes. Hepatology 2009; 50: 1079–1086.

20. Petersen KF, Dufour S, Befroy D, Garcia R, Shulman GI. Impaired mitochondrial activity in the insulin-resistant offspring of patients with type 2 diabetes. N Engl J Med 2004; 350: 664–671.

21. Petersen KF, Befroy D, Dufour S, Dziura J, Arian C, Rothman DL et al. Mitochondrial dysfunction in the elderly: possible role in insulin resistance. Science 2003; 300: 1140–1142.

22. Schmid AI, Szendroedi J, Chmelik M, Kressak M, Moser E, Roden M. Liver ATP synthesis is lower and relates to insulin sensitivity in patients with type 2 diabetes. Diabetes Care 2011; 34: 448–453.

23. Phielix E, Szendroedi J, Roden M. Mitochondrial function and insulin resistance during aging: a mini-review. Gerontology 2011; 57: 387–396.

24. Choi CS, Befroy DE, Codella R, Kim S, Reznick RM, Hwang YJ et al. Paradoxical effects of increased expression of PGC-1alpha on muscle mitochondrial function and insulin-stimulated muscle glycolysis in vivo. Proc Natl Acad Sci USA 2008; 105: 19926–19931.

25. Laurent D, Yerby B, Deacon R, Gajewska S, Hwang JH et al. Role of hepatic glycogen breakdown in defective counterregulation of hypoglycemia in intensively treated type 1 diabetes. Diabetes 2006; 55: 659–666.

26. Roser W, Beckmann N, Wiesmann U, Seelig J. Absolute quantification of the hepatic glycogen content in a patient with glycogen storage disease by 13C-magnetic resonance spectroscopy. Magn Reson Imaging 1996; 14: 1217–1220.

27. Roden M, Perseghin G, Petersen KF, Hwang JH, Cline GW, Gerov K et al. The roles of insulin and glucagon in the regulation of hepatic glycogen synthesis and turnover in humans. J Clin Invest 1996; 97: 642–648.

28. Hwang J-H, Blum J, Leaf A, Ross BD. In vivo characterization of fatty acids in human adipose tissue using natural abundance 1H decoupled 13C MRS at 1.5 T: clinical applications to dietary therapy. NMR Biomed 2003; 16: 160–167.

29. Rothman DL, De Feyter HM, de Graaf RA, Mason GF, Behar KL. 13C MRS studies of neuroenergetics and neurotransmitter cycling in humans. NMR Biomed 2011; 24: 943–957.

30. Mishkosky M, Comment A, Gruetter R. In vivo detection of brain Krebs cycle intermediate by hyperpolarized magnetic resonance. J Cereb Blood Flow Metab 2012; 32: 2108–2113.

31. Choi C, Ganji SK, DeBerardinis RJ, Hatanpaa KJ, Rakheja D, Kovacs Z et al. 2-hydroxylutarate detection by magnetic resonance spectroscopy in IDH-mutated patients with gliomas. Nat Med 2012; 18: 624–629.