

Hyperpolarized ^{13}C -Pyruvate Magnetic Resonance Reveals Rapid Lactate Export in Metastatic Renal Cell Carcinomas

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Abstract

Renal cell carcinomas (RCC) are a heterogeneous group of tumors with a wide range of aggressiveness. Noninvasive methods to confidently predict the tumor biologic behavior and select appropriate treatment are lacking. Here, we investigate the dynamic metabolic flux in living RCC cells using hyperpolarized ^{13}C -pyruvate magnetic resonance spectroscopy (MRS) combined with a bioreactor platform and interrogated the biochemical basis of the MRS data with respect to cancer aggressiveness. RCC cells have significantly higher pyruvate-to-lactate flux than the normal renal tubule cells. Furthermore, a key feature distinguishing the localized from the metastatic RCC cells is the lactate efflux rate, mediated by the monocarboxylate transporter 4 (MCT4). The metastatic RCC cells have significantly higher MCT4 expression and corresponding higher lactate efflux, which is essential for maintaining a high rate of glycolysis. We show that such differential cellular transporter expression and associated metabolic phenotype can be noninvasively assessed via real-time monitoring of hyperpolarized ^{13}C -pyruvate-to-lactate flux. *Cancer Res*; 73(2); 529–38. ©2012 AACR.

Introduction

The incidence of renal tumors, both malignant renal cell carcinomas (RCC) and benign renal tumors, has increased significantly in the last 20 years (1). In the case of renal tumors, biopsies are not routinely done, due to the risk of hemorrhage and high likelihood of indeterminate histology (2, 3). Treatment selection is thus heavily reliant on noninvasive imaging assessment of tumor masses. However, there are significant limitations to the current imaging methods for renal tumor characterization. It is increasingly recognized that RCCs are a heterogeneous group of tumors with a wide range of biologic aggressiveness (4, 5). Emerging active surveillance data have shown that a significant percentage of small RCCs (<4 cm) are indolent with low metastatic risk, and patients may be over-treated if all such RCCs are surgically removed (5, 6). On the other hand, 20% to 40% of patients undergoing nephrectomies for clinically localized RCCs develop metastases with poor outcome (7). Unfortunately, current imaging methods cannot reliably predict the risks of progression from localized RCC to metastatic disease (8). Furthermore, certain benign renal tumors are difficult to distinguish from RCCs by imaging

(9). This diagnostic challenge has resulted in the unnecessary resection of many benign renal tumors, which constitute 20% of all renal tumors less than 4 cm, with the associated surgical risks and potential loss of renal function (10). Therefore, new imaging methods are needed to predict the biologic behavior of renal tumors and select appropriate treatment.

The unique metabolism of cancer cells is central to their malignant behavior. For example, a common property of cancers is altered glucose metabolism with elevated glycolysis and lactate production in the presence of oxygen (11, 12). Increased glycolysis facilitates the uptake and incorporation of nutrients and biomass needed for cell proliferation in cancers (13, 14) and acidifies the extracellular microenvironment-promoting invasion of neighboring tissue and metastasis (15). A number of genomics and proteomics studies have shown increased metabolism to lactate in RCCs (16–19). Specifically, proteomic analysis of RCC tissues and metabolic profiling of serum samples revealed increased levels of glycolytic enzymes in RCC tissues and higher lactate in the serum of patients with RCC (19). Metastatic RCCs have also shown a bioenergetic shift toward aerobic glycolysis and lactate production (18). These studies provide the rationale for metabolic imaging of glycolysis as a noninvasive means to characterize renal tumor aggressiveness.

Hyperpolarized ^{13}C magnetic resonance is a new molecular imaging technique that allows rapid and noninvasive monitoring of dynamic pathway-specific metabolic and physiologic processes. Hyperpolarization, achieved through the dynamic nuclear polarization (DNP) technique (20), can provide dramatic gains in sensitivity (>10,000-fold increase) for imaging ^{13}C -labeled biomolecules. The hyperpolarized ^{13}C -probes can be injected into living systems, and their metabolism can be observed in real-time by chemical shift. The most commonly

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used hyperpolarized ^{13}C probe is ^{13}C -pyruvate, which is at the juncture of several important energy and biosynthetic pathways. For example, pyruvate may be converted to lactate in glycolysis, to acetyl-CoA to support the tricarboxylic acid (TCA) cycle, or to alanine via transamination for protein synthesis. Hyperpolarized ^{13}C -pyruvate magnetic resonance has already been applied to the detection of the presence (21–25) and progression (26, 27) of a number of cancers. The metabolic changes seen in RCCs suggest that hyperpolarized ^{13}C -pyruvate will also be an excellent probe to interrogate these tumors noninvasively.

In this work, we compared the pyruvate metabolism of immortalized cells derived from human renal proximal tubules (the origin of most human RCCs), a localized human RCC, and a metastatic human RCC, with the goal of identifying clinically translatable hyperpolarized biomarkers of renal tumor aggressiveness. After evaluating the steady-state metabolism of these cells, we assessed the dynamic hyperpolarized pyruvate-to-lactate flux using a magnetic resonance compatible bioreactor platform that provides a controlled and physiologic setting for the cells (28). By monitoring the real-time metabolic flux using hyperpolarized magnetic resonance, we showed that RCC cells have significantly higher pyruvate-to-lactate flux than the normal renal proximal tubule cells. Furthermore, we showed that cells derived from the metastatic RCC have more rapid export of lactate to the extracellular space compared with the cells derived from the localized RCC, and that these differences are likely mediated by the differential expression of monocarboxylate transporter 4 (MCT4). These results suggest that using hyperpolarized ^{13}C -pyruvate to assess lactate production and export has the potential to improve the noninvasive characterization of renal tumors.

Materials and Methods

Cell lines

HK-2 is an immortalized proximal tubule epithelial cell line from normal adult human kidney (29) and was obtained from American Type Culture Collection [ATCC; obtained June, 2010; authentication conducted at ATCC was via short tandem repeat (STR) profiling]. UMRC6 cells are representative of localized human clear cell RCC (30), and were a gift from Dr. Bart Grossman (MD Anderson Cancer Center, Houston, TX; obtained January, 2010; authenticated using STR profiling, October 2012). UOK262 cells are derived from a metastasis of the highly aggressive hereditary leiomyomatosis RCC (HLRCC), which is characterized by mutation of the TCA cycle enzyme fumarate hydratase (31). UOK262 cells were a gift from Dr. W. Marston Linehan (National Cancer Institute, Bethesda, MD; obtained May, 2010; authenticated using STR profiling, October 2012). All cells were grown in Dulbecco's Modified Eagle's Medium (DMEM) with 4.5 g/L glucose. The cells were passaged serially and were used for assays and magnetic resonance experiments between passages 2 to 10 and at 60% to 80% confluency.

^1H NMR experiments

Cells were plated on 150-cm²-coated Petri dishes (Fisher Scientific) and incubated for 24-hours in DMEM media supplemented with [1- ^{13}C] glucose, or for 2-hours in DMEM media

supplemented with [3- ^{13}C] pyruvate (Cambridge Isotope Laboratories). At the end of incubation, an aliquot of medium was collected, and cells were extracted in ice-cold methanol (32). The cell extracts were reconstituted in D₂O with known amounts of trimethyl silyl pentanoate (TSP) for internal reference. The extracts were measured on Bruker Advance III 800 MHz equipped with a cryo-cooled triple-resonance probe. High-resolution water-suppressed proton spectra were obtained with a repetition time of 12 seconds and 64 averages. The metabolite peak areas were quantified against the known TSP peak area.

Hyperpolarized [1- ^{13}C] pyruvate magnetic resonance bioreactor experiment

Cells were electrostatically encapsulated into 2.5% w/v alginate microspheres as previously described (28, 33), and then loaded into a magnetic resonance-compatible bioreactor. Approximately 800 μL of microspheres were perfused in the bioreactor with DMEM H-21 media at a flow rate of 2.5 mL/min. For the flow rate modulation bioreactor experiments, the flow rate was changed to either 1.3 or 3.8 mL/min for the duration of the hyperpolarized scans. The media was kept at 37°C with water-jacketed perfusion lines and was maintained at 95% air/5% CO₂ via gas exchanger. All bioreactor studies were conducted on a 500 MHz Varian Inova (Agilent Technologies) with a 10 mm, triple-tune, direct-detect, broadband probe at 37°C. For the hyperpolarized ^{13}C -pyruvate studies, 2.5 μL of 14.2 mol/L ^{13}C -pyruvate mixed with 15 mmol/L of the trityl radical (GE Healthcare) was polarized on a Hypersense polarizer (Oxford Instruments). This was followed by dissolution in 5 mL of 50 mmol/L phosphate buffer. One milliliter of the resulting 7.5 mmol/L hyperpolarized pyruvate solution was injected into the bioreactor containing the microspheres. Hyperpolarized ^{13}C magnetic resonance data were acquired dynamically with a 10° flip-angle, pulse repetition time of 3 seconds and for a duration of 300 seconds. ^{31}P spectra (repetition time 3 seconds, 1,024 averages, 90° flip-angle) were acquired before and after each hyperpolarized study to assess cell viability.

Cell number determination for bioreactor experiments

Moles of ATP per cell for each cell line were measured using CellTiter-Glo luminescent cell viability assay and Veritas Luminometer (Promega). Moles of ATP corresponding to the β -nucleoside triphosphates (β -NTP) peak area on ^{31}P spectra were determined using a ^{31}P calibration curve. The number of cells in each bioreactor experiment was then calculated by dividing the moles of ATP approximated from β -NTP peak by the moles of ATP per cell as measured by the luminescent assay.

mRNA expression and enzyme activity assay

Total RNA was purified from cells using RNeasy procedure kit (Qiagen), and reverse transcribed using iScript cDNA Synthesis kit (BioRad Laboratories). PCR was conducted in triplicate for the lactate dehydrogenase- α (LDH- α) and the monocarboxylate transporters 1 and 4 (Hs00161826_m1, Hs00358829_m1) on the ABI 7900HT (Applied Biosystems).

Cyclophilin and β -actin (Applied Biosystems) were used as control, and the relative fold difference was calculated for each primer/probe combination.

LDH activity of cell lysates was measured spectrophotometrically by quantifying the linear decrease in NADH absorbance at varying pyruvate concentrations at 339 nm using a microplate reader (Tecan Group Ltd.). The maximum velocity (V_{\max}) and the Michaelis–Menten constant (K_m) were estimated using the Lineweaver–Burke plot.

Data analysis

The hyperpolarized pyruvate-to-lactate flux was calculated using a previously published model (28). The pyruvate-to-lactate flux was normalized by the number of cells in each bioreactor study and the injected amount of hyperpolarized pyruvate. ^{31}P metabolite peaks were integrated and normalized by the number of cells to determine the concentration of phosphocholine (PC), glycerophosphocholine (GPC) and β -NTP. Resonances were corrected for their respective ^{31}P T_1 relaxation times (Supplementary Table S2). One-way ANOVA was used to assess the difference between the 3 groups with Tukey–Kramer method *post hoc* tests using statistical software package JMP (SAS Institute). All values are reported as mean \pm SE.

Results

^1H MRS shows steady-state metabolite concentrations that reflect increased glycolysis and truncated TCA cycle in metastatic UOK262 cells

We first used ^1H magnetic resonance spectroscopy (MRS) to interrogate the steady-state metabolite concentrations in HK2, UMRC6, and UOK262 cells. HK2 cells are derived from human renal proximal tubule cells (doubling time = 72–96 hours; ref. 29). UMRC6 cells (doubling time = 43 hours) originate from a localized human clear cell RCC (30). Finally, UOK262 cells were isolated from a metastasis of HLRCC (doubling time = 23 hours; ref. 31). HLRCC is an aggressive RCC characterized by mutation of the TCA cycle enzyme fumarate hydratase. UOK262 cells therefore have markedly reduced oxidative phosphorylation and are highly glycolytic (31). Figure 1A illustrates the biochemical scheme of glycolysis and TCA cycle. Figure 1B shows the major steady-state intracellular metabolite concentrations, as measured by ^1H MRS, in the 3 cell lines. We found that the steady-state lactate concentration was significantly higher in the UOK262 cells compared with the UMRC6 or HK2 cells (both $P < 0.05$). The increased steady-state lactate in the UOK262 cells is consistent with the fumarate hydratase mutation, which sharply attenuates the mitochondrial TCA cycle and concomitantly drives glycolysis for energy production (31).

The alanine concentration was lower in the 2 RCC cell lines compared with the HK2 cells, likely due to increased flux of pyruvate to lactate. We also found significantly increased glutamate and decreased aspartate concentration in the UOK262 cells compared with the other 2 cell lines. Glutamate is reversibly formed from α -ketoglutarate, a TCA cycle intermediate proximal to fumarate. Aspartate is formed reversibly from oxaloacetate, a TCA intermediate distal to fumarate. In

the UOK262 cells, the increased steady-state glutamate and reduced aspartate are consistent with truncation of TCA cycle metabolism beyond fumarate, due to the fumarate hydratase mutation. The concentration of GPC, an abundant renal osmolyte (34), was similar among the 3 cell lines. Interestingly, we found that phosphocholine was significantly higher in the UMRC6 cells than the UOK262 cells. While phosphocholine has been used as a biomarker of tumor proliferation and aggressiveness in other types of cancers (35, 36), the levels of phosphocholine did not correlate with aggressiveness in the RCC cell lines in our study. Phosphocholine is converted from choline by the enzyme choline kinase- α (CHKA) in the phosphatidylcholine synthesis (Kennedy) pathway. A recent study reported that CHKA forms a complex with EGF receptor (EGFR) in a c-Src–dependent manner, and functions cooperatively with EGFR and c-Src in regulating pathways critical to cell proliferation (37). Such required functional interaction among the 3 enzymes for cancer cell proliferation may in part explain the lack of direct correlation between the phosphocholine levels and proliferation rates/aggressiveness of the 2 RCC cell lines in our study.

Twenty-four-hour labeling of cells with $[1-^{13}\text{C}]$ glucose or $[3-^{13}\text{C}]$ pyruvate in 2D cell cultures shows increased flux to lactate in the metastatic UOK262 cells

To further characterize glycolysis and lactate production in the RCC cells, we investigated the flux from labeled $[1-^{13}\text{C}]$ glucose to lactate in 2-dimensional (2D) cell cultures following 24-hour incubation. Figure 2A shows the scheme of ^{13}C -labeled carbon atom transitions used to detect glucose metabolism to lactate. Glucose, the primary fuel for energy in cells, is taken up primarily via the glucose transporter 1 (GLUT1) and is converted to pyruvate and then lactate during glycolysis. Lactate is preferentially exported out of the cells via the MCT4 (38). Figure 2B and C show representative MRS of metabolites in the culture medium and intracellular compartment, respectively, of UOK262 cells following 24-hour labeling with $[1-^{13}\text{C}]$ glucose. Figure 2D and E show the concentrations of ^{13}C -labeled lactate in the medium and the intracellular compartment, respectively, of the 3 cell lines following incubation with $[1-^{13}\text{C}]$ glucose. After 24 hour of incubation with $[1-^{13}\text{C}]$ glucose, 99% of lactate was found in the extracellular medium. The concentration of ^{13}C lactate in the medium increased progressively from HK2 to UMRC6 to UOK262 cells, with the lactate concentration being nearly 3-fold higher in the medium of UOK262 cells compared with that of the HK2 cells. The fractional enrichment of lactate was defined as the ^{13}C -labeled lactate/ ^{13}C -labeled lactate + unlabeled lactate. The fractional enrichment of lactate in the medium was $76\% \pm 1\%$, $65\% \pm 2\%$, and $84\% \pm 1\%$ in the HK2, UMRC6, and UOK262 cells, respectively. This implies that the predominant source of lactate in these cells is glucose although there is a contribution from other carbon sources as well. The intracellular concentration of ^{13}C lactate was also significantly higher in the UOK262 cells compared with the UMRC6 or the HK2 cells ($P < 0.05$). The intracellular fractional enrichment of lactate was $62.5\% \pm 3.1\%$, $61.8\% \pm 1.5\%$, and $78.6\% \pm 0.4\%$ in the HK2, UMRC6, and UOK262 cells, respectively. The differential lactate

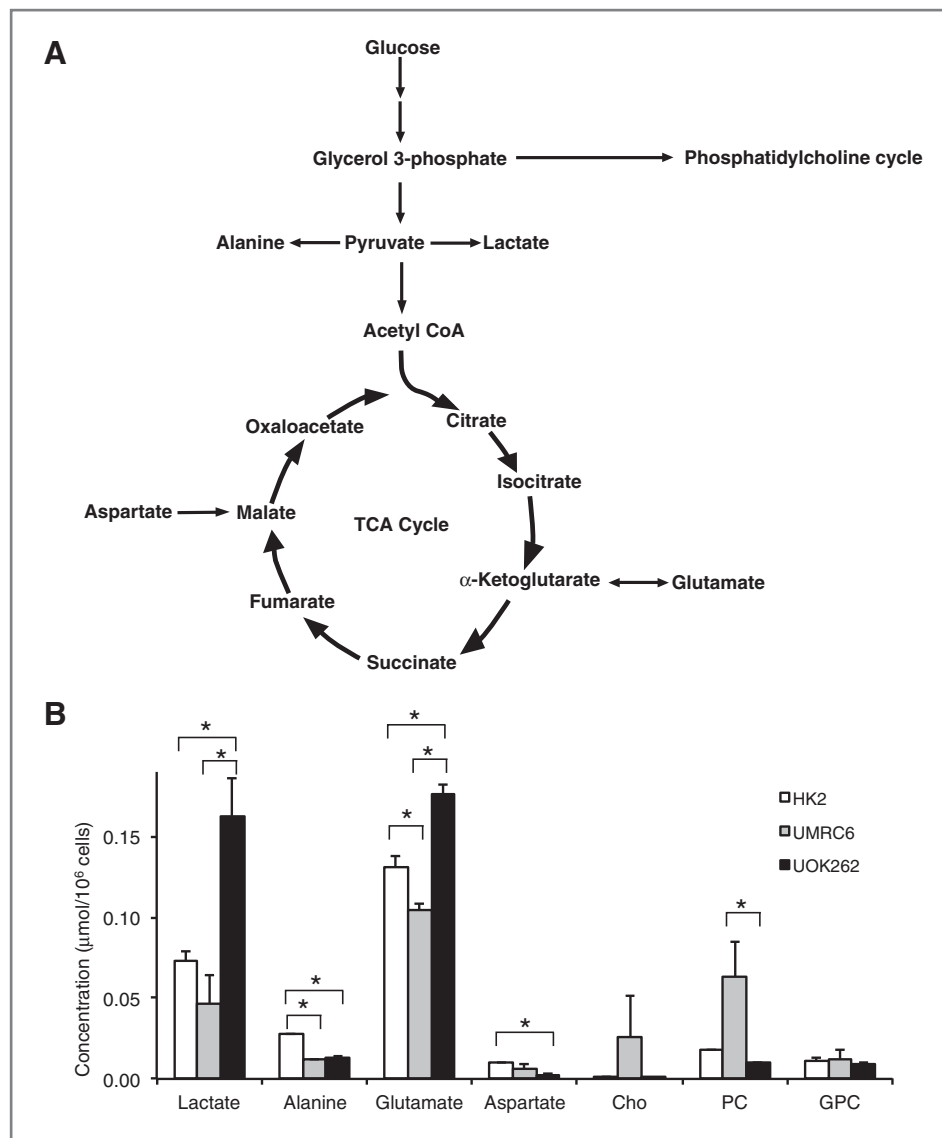


Figure 1. Steady-state major metabolites in the 3 cell lines. A, biochemical scheme of glycolysis and TCA cycle. B, steady-state concentrations of major metabolites, as measured by ¹H MRS, in the 3 cell lines (*N* = 5 each). All values are reported as mean ± SE. *, significant difference (*P* < 0.05).

fractional enrichment in the intracellular compartment and the medium might be related to lactate compartmentalization in the cells. Such compartmentalization has been reported to exist in the brain and myocardium (39, 40). It is possible that one compartment of intracellular lactate originates predominantly from ¹³C-labeled glucose, and the subsequently labeled lactate is preferentially exported into the medium. Another compartment of lactate may derive from other sources such as from glutamine via glutaminolysis (41), and this compartment of lactate may not be as readily exported into the medium as that from glucose. The presence of lactate compartmentalization may also in part explain the higher fractional enrichment of intracellular lactate in the UOK262 cells compared with the other cells. Because UOK262 have fumarate hydratase mutation with reduced TCA cycle metabolism, they are more likely to produce lactate from glycolysis than from other pathways such as glutaminolysis that contains parts of the TCA cycle. This may explain the higher fractional enrichment

of the intracellular lactate from labeled glucose in the UOK262 cells. Taken together, the earlier findings confirmed that UOK262 cells are highly glycolytic with increased production of lactate. Interestingly, the ¹³C lactate concentration was lower in the UMRC6 RCC cells relative to both HK2 and UOK262 cells. This was in agreement with the steady-state intracellular lactate concentration data, which also showed a decreased lactate pool in UMRC6 cells compared with the other 2 cell lines (Fig. 1B).

Hyperpolarized ¹³C MRS shows higher real-time pyruvate-to-lactate flux in RCC cells compared with normal HK2 cells but lower pyruvate-to-lactate flux in the metastatic UOK262 cells than the localized UMRC6 cells

Given the dynamic nature of cellular metabolism, we then investigated the real-time pyruvate metabolism in the 3 cell lines using hyperpolarized ¹³C magnetic resonance. We carried

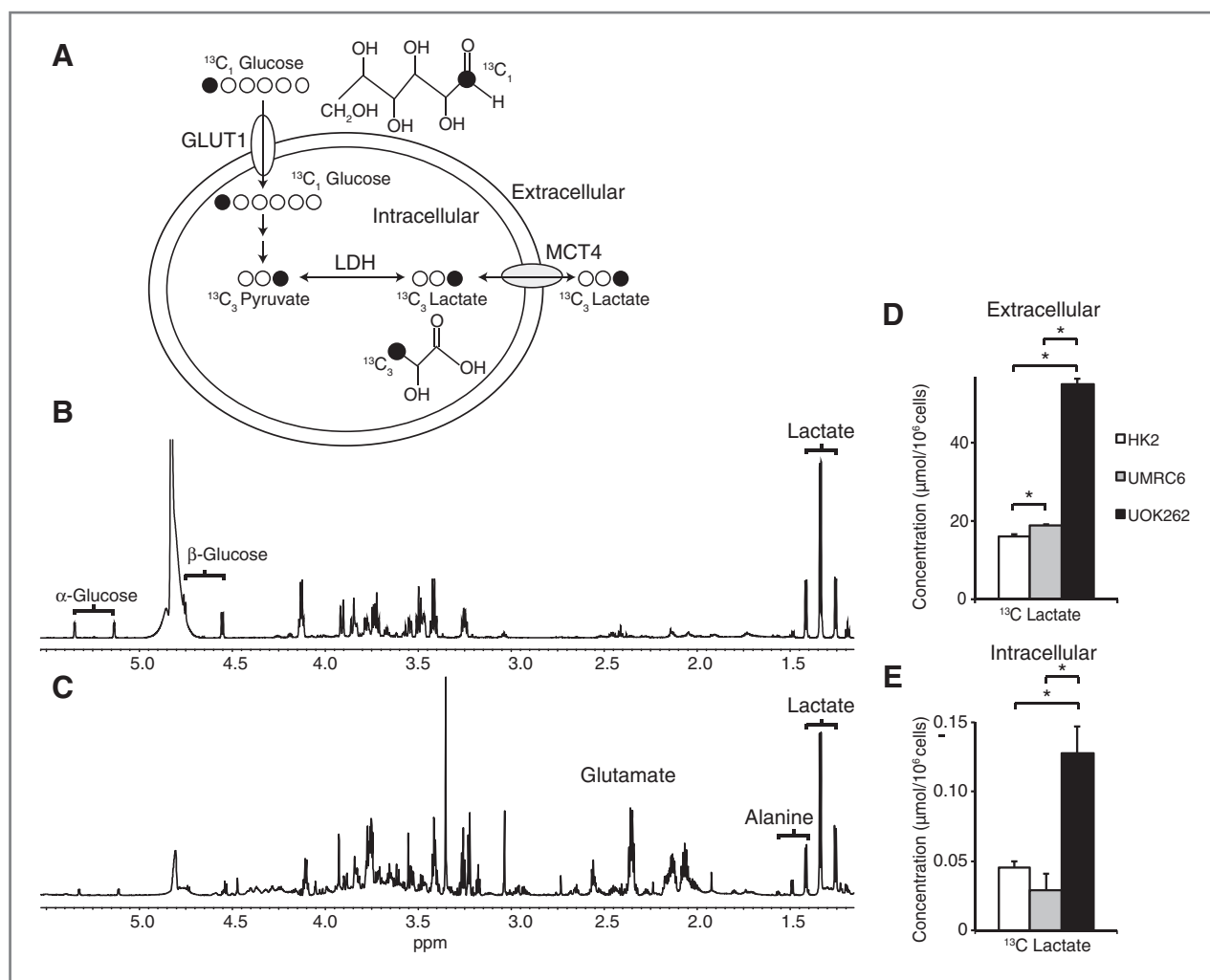


Figure 2. ^{13}C -labeled lactate in the media and intracellular compartment of the 3 cell lines following 24-hour incubation with $[1-^{13}\text{C}]$ glucose. A, biochemical scheme illustrating ^{13}C -labeled carbon atom transitions used to detect glucose metabolism to lactate. Representative ^1H MRS of metabolites in the medium (B) and intracellular compartments (C) of UOK262 following 24-hour labeling with $[1-^{13}\text{C}]$ glucose. The brackets indicate the ^{13}C satellites of each metabolite. Concentrations of ^{13}C -labeled lactate in the media (D) and intracellular compartment (E) of the 3 cell lines following incubation with $[1-^{13}\text{C}]$ glucose ($N = 5$ each). All values are reported as mean \pm SE. *, significant difference ($P < 0.05$).

out our hyperpolarized ^{13}C magnetic resonance experiments using a bioreactor, a continuously perfused 3-dimensional (3D) cell culture system that provides a controlled and physiologic setting for the cells. This system has been shown to produce highly reproducible hyperpolarized magnetic resonance data (28), and facilitates the characterization of hyperpolarized substrate to metabolite conversion. ^{31}P MRS was used to monitor changes in cell bioenergetics during the bioreactor studies. Representative ^{31}P spectra of the cells are shown in Supplementary Fig. S1. Nuclear magnetic resonance (NMR) signals for the NTPs (γ -NTP, α -NTP, and β -NTP), phosphocholine, inorganic phosphate (P_i), and GPC were readily visible. The total NTP content was unchanged following the injection of hyperpolarized ^{13}C -pyruvate, indicating maintenance of cell viability during the course of the hyperpolarized experiments. Figure 3 shows the PC/GPC ratios and PC+GPC concentration in the 3 cell lines. We found significantly higher

PC+GPC concentration in the UMRC6 cells compared with the UOK262 cells ($P < 0.05$), which was in agreement with the steady-state ^1H data from 2D cell culture. In addition to monitoring cell energetics, ^{31}P spectroscopy also enables quantitative hyperpolarized data analysis by normalizing the hyperpolarized magnetic resonance data with respect to the number of viable cells, through concomitant measurements of β -NTP concentration via ^{31}P MRS.

Figure 4A illustrates the scheme of ^{13}C -labeled carbon atom transitions used to detect ^{13}C -pyruvate metabolism during the hyperpolarized magnetic resonance experiment. After the injection of hyperpolarized ^{13}C -pyruvate into the bioreactor, the real-time pyruvate-to-lactate flux was assessed for all 3 cell lines. The data were fit to a 2-state model of interconversion of pyruvate to lactate and the metabolic fluxes were calculated (28). Figure 4B shows fitted pyruvate-to-lactate flux and representative spectra of ^{13}C -pyruvate and lactate. The average

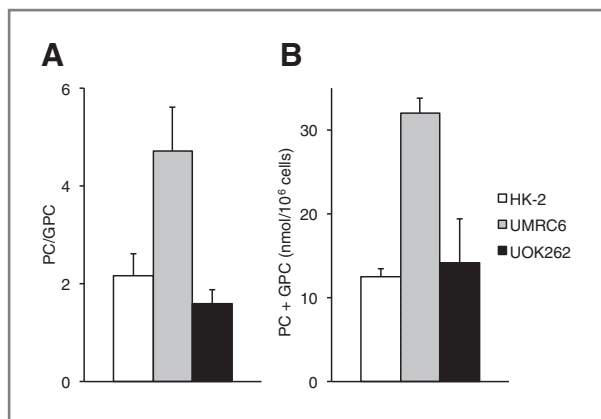


Figure 3. Phosphocholine metabolites in the 3 cell lines. PC/GPC ratios (A) and PC+GPC concentration (B) of cells encapsulated and perfused in a bioreactor ($N = 5$ each). All values are reported as mean \pm SE.

fluxes, at a flow rate of 2.5 mL/min in the bioreactor, for each of these cell lines are shown in Fig. 4C. The observed flux rate was significantly higher in the 2 RCC cell lines (UMRC6 and UOK262) as compared with the renal tubule cell line HK2 (UMRC6 vs. HK2, $P < 0.0001$; UOK262 vs. HK2, $P = 0.003$). Unexpectedly, the observed real-time hyperpolarized pyruvate-to-lactate flux for UOK262 cells (representative of metastatic RCC) was lower than that of the UMRC6 cells (representative of localized RCC). Similar to the flux data, the area under the curve for the ^{13}C lactate was higher in the RCC cells than the normal renal tubule cells but was lower in the UOK262 RCC cells than the UMRC6 RCC cells (Supplementary

Table S1). Additional analysis of hyperpolarized ^{13}C dynamics of the cells perfused in the bioreactors was summarized in Supplementary Table S1.

mRNA expression analysis shows increased MCT4 in the metastatic UOK262 cells

To better understand the cellular processes underlying the hyperpolarized pyruvate flux results, we then assayed the mRNA expression and enzyme activity level of LDHA, and the mRNA expression of MCT1 and MCT4 in the 3 cell lines. LDHA encodes the predominantly M isoform of LDH, which catalyzes the conversion between pyruvate and lactate. MCT1 mediates the pyruvate transport into the cells, and MCT4 mediates the efflux of the lactate out of the cells (42). We found that the mRNA expression of LDHA was significantly higher in the UOK262 cells than the other 2 cell lines (Fig. 5). For the LDH activity, K_m of the 2 RCC cell lines was significantly higher than that of the HK2 cells ($P < 0.03$) but not significantly different between the UMRC6 and UOK262 RCC cells. The V_{max} of UOK262 cells was significantly higher than that of HK2 cells ($P < 0.05$). The mRNA expression of MCT1 was significantly higher in the UMRC6 cells (UMRC6 vs. HK2, $P = 0.0004$; UMRC6 vs. UOK262, $P = 0.0002$), whereas the MCT4 expression was significantly elevated in the UOK262 cells (UOK262 vs. HK2, $P = 0.001$; UOK262 vs. UMRC6, $P = 0.02$).

The higher hyperpolarized pyruvate-to-lactate flux in UMRC6 cells, as compared with UOK262, was likely due, in part, to the higher MCT1 expression rather than the lactate pool size in the UMRC6 cells, as both the steady-state and 24-hour labeling data showed lower lactate pool size in the UMRC6 cells (Figs. 1B and 2E). Importantly, the differential

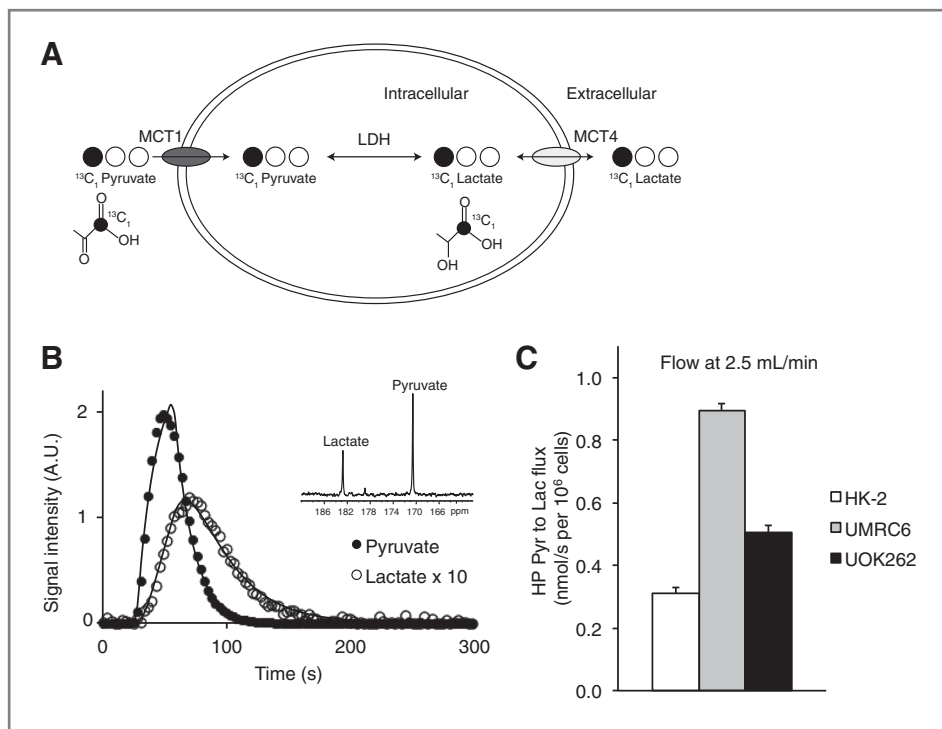
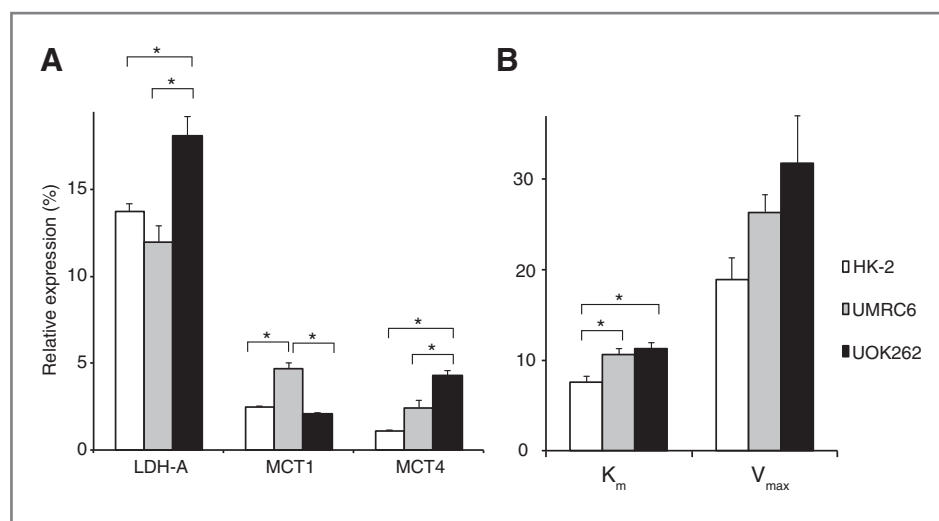


Figure 4. Dynamic hyperpolarized pyruvate-to-lactate flux in the 3 cell lines. A, scheme of ^{13}C -labeled carbon atom transitions used to detect C_1 -labeled pyruvate metabolism during the hyperpolarized experiment. B, fitted pyruvate-to-lactate flux and representative spectra (inset) of ^{13}C -pyruvate and lactate in the UMRC6 cells. C, comparison of observed real-time hyperpolarized pyruvate-to-lactate flux in the 3 cell lines at a flow rate of 2.5 mL/min in the bioreactor ($N = 5$ each). All values are reported as mean \pm SE.

Figure 5. Analysis of relevant enzyme expression/activity and transporter expression in the 3 cell lines. A, mRNA expression of LDHA and monocarboxylate transporters 1 and 4 (MCT1 and MCT4), relative to internal β -actin expression, in the 3 cell lines ($N = 6$ each). B, LDH activity as measured by K_m ($\mu\text{mol pyruvate}/10^6$ cells) and V_{max} ($\mu\text{mol NADH/s}/10^6$ cells) in the 3 cell lines ($N = 6$ each). All values are reported as mean \pm SE. *, significant difference ($P < 0.05$).



expression of MCT4 may explain the apparent discrepancy between the real-time hyperpolarized pyruvate-to-lactate flux and the 24-hours labeling of lactate in the UOK262 cells compared with the UMRC6 cells. The UOK262 cells have an almost 2-fold higher MCT4 expression compared with the UMRC6 cells, suggesting that they likely have more rapid MCT4-mediated export of lactate out of the cells. Rapid lactate efflux is essential for maintaining a neutral intracellular pH, and a high rate of glycolysis and lactate production over time. In contrast, UMRC6 cells have lower MCT4 expression, and likely slower rate of lactate export. Although UMRC6 cells have higher MCT1, these cells would be less able to maintain a high rate of lactate production over time due to buildup of intracellular lactate. Therefore, while the real-time flux of pyruvate to lactate during the timeframe of the hyperpolarized experiment was lower in the UOK262 cells than the UMRC6 cells, the higher MCT4 expression in the UOK262 cells likely resulted in more rapid lactate efflux and accounted for the significantly higher ^{13}C -labeled lactate accumulated in the medium in the 24-hour labeling experiment. Over time, the large amount of labeled lactate accumulated in the medium of UOK262 cells likely diffused back into the cells down a gradient, and may explain the higher intracellular-labeled lactate in the UOK262 cells compared with the UMRC6 cells. We postulate that, while such diffusion of lactate back into the cells may reduce further generation of labeled lactate, this process occurs after a large amount of lactate has already accumulated in the medium of the UOK262 cells. This accumulation of medium lactate and diffusion back into the UOK262 cells were likely accentuated in the 2D cell cultures in which the extracellular lactate was not removed, in contrast to the bioreactor in which the medium was continuously exchanged.

It is also important to note that while MCT1 may affect the hyperpolarized lactate signal (both the intracellular and extracellular hyperpolarized lactate) if it were the rate-limiting step in the pyruvate-to-lactate flux, the relative proportion of the intracellular versus extracellular hyperpolarized lactate would be determined by MCT4, which modulates the lactate efflux. In addition, lactate efflux in general is not expected to be signif-

icantly affected by MCT1, as most of the lactate produced in the cells is derived from glucose (transported via GLUT1) rather than pyruvate (transported via MCT1) uptake into the cells.

Hyperpolarized ^{13}C -pyruvate magnetic resonance combined with flow rate modulation in the bioreactor show rapid efflux of lactate in the metastatic UOK262 cells

We then conducted a second set of hyperpolarized magnetic resonance experiments using different flow rates in the bioreactor to investigate the real-time lactate efflux rate in the 2 RCC cell lines. At high flow rates, the extracellular lactate will more likely flow out of the NMR coil's sensitive volume and will not contribute to the magnetic resonance signal, thereby decreasing the observed pyruvate-to-lactate flux (Fig. 6A). It follows that the relative amount of extracellular lactate (lactate in the medium) of the 2 RCC cell lines, which reflects the lactate efflux rate, can be inferred from the observed hyperpolarized pyruvate-to-lactate flux at different flow rates. Figure 6B shows the pyruvate-to-lactate flux at different flow rates for the 2 RCC cells. For the UMRC6 cells, the mean observed hyperpolarized pyruvate-to-lactate flux was 0.92 nmol/s per 10^6 cells at 1.3 mL/min, 0.90 nmol/s per 10^6 cells at 2.5 mL/min, and 0.94 nmol/s per 10^6 cells at 3.8 mL/min, all of which were not statistically different from one another. For the UOK262 cells, the mean observed hyperpolarized pyruvate-to-lactate flux was 0.56 nmol/s per 10^6 cells at 1.3 mL/min, 0.51 nmol/s per 10^6 cells at 2.5 mL/min, and 0.41 nmol/s per 10^6 cells at 3.8 mL/min. These observed hyperpolarized pyruvate-to-lactate flux for the UOK262 cells progressively decreased at higher flow rate, with a significant 20% decrease in the flux between the 2.5 and 3.8 mL/min flow rate ($P = 0.01$). At the high flow rate of 3.8 mL/min, the decreased pyruvate-to-lactate flux in the UOK262 cells indicated that these cells had more rapid lactate efflux and higher amount of extracellular lactate, which was readily removed from the NMR-sensitive region at high flow rate. The high flow rate should not have significantly limited the MCT1-mediated pyruvate uptake into the

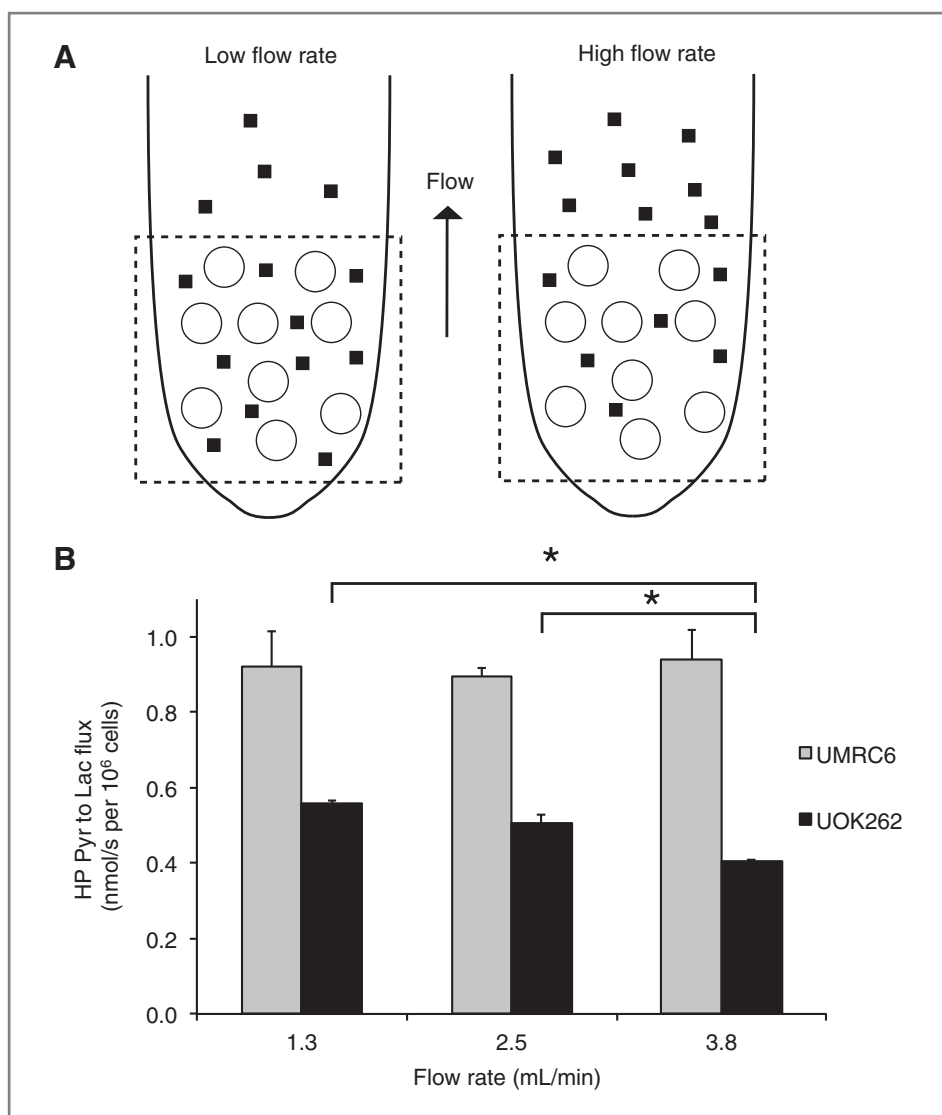


Figure 6. Dynamic hyperpolarized pyruvate-to-lactate flux in the RCC cells following flow rate modulation in the bioreactor. A, schematic illustrating the relationship between flow rates and observed hyperpolarized pyruvate-to-lactate flux in the bioreactor. At high flow rates, the extracellular lactate will more likely flow out of the NMR coil's sensitive volume and will not contribute to the magnetic resonance signal, thereby decreasing the observed pyruvate-to-lactate flux. The dotted square represents NMR sensitive region. ○, encapsulated microspheres containing cells. ■, extracellular lactate. B, hyperpolarized pyruvate-to-lactate flux of UOK262 and UMRC6 cells at 3 different flow rate ($N = 5$ each). There is a decreasing trend in observed pyruvate-to-lactate flux with increasing flow rate for UOK262 cells. All values are reported as mean \pm SE. *, significance ($P < 0.05$).

cells. This is because the injected hyperpolarized pyruvate substrate available to the cells was expected to be in excess compared with MCT1, even at the high flow rate of 3.8 mL/min. Indeed, the UMRC6 cells, with 2-fold higher expression of MCT1 compared with the UOK262 cells, showed similar hyperpolarized pyruvate-to-lactate flux at all 3 flow rates, indicating that the flow rates did not limit pyruvate uptake. The flow rate should also not have affected the enzymatic conversion of pyruvate to lactate in the cells. Taken together, the hyperpolarized flux data at different flow rates strongly support the notion that the UOK262 cells have increased MCT4-mediated lactate efflux out of the cells.

In addition, we incubated the UMRC6 and UOK262 cells for 2 hours in medium containing $[3-^{13}\text{C}]$ pyruvate, and observed 0.47 ± 0.05 versus $2.32 \pm 0.27 \mu\text{mol}/10^6$ cells of ^{13}C -labeled lactate in the medium of UMRC6 versus UOK262 cells. This more than 5-fold increase in the extracellular lactate of the UOK262 cells further verifies that lactate derived from labeled pyruvate is produced and transported out of the cells at a

higher rate in the UOK262 cells compared with the UMRC6 cells.

Discussion

There is increasing evidence that RCCs are among those tumors strongly linked to abnormal metabolism, a feature that may be exploited therapeutically. In this work, we investigated the pyruvate metabolism in perfused human RCC cells using a clinically translatable hyperpolarized ^{13}C magnetic resonance probe, and interrogated both the biochemical basis of the observed hyperpolarized magnetic resonance data and its relationship to cancer aggressiveness. We found higher pyruvate-to-lactate flux, consistent with increased glycolysis, in RCC cells compared with normal renal proximal tubule cells. We further noted that a key feature distinguishing the localized UMRC6 from the metastatic UOK262 RCC cells is the lactate efflux rate, and that, importantly, this feature can be noninvasively depicted via real-time monitoring of hyperpolarized ^{13}C -pyruvate-to-lactate flux.

Lactate efflux is predominantly mediated by MCT4, which is a proton-coupled lactate transporter (42), exporting lactate and H⁺ in the same direction out of the cells. Rapid lactate efflux serves to maintain high levels of glycolysis in cancer cells and concurrently acidifies the extracellular environment (11). Low extracellular pH supports invasion and metastasis, perhaps due to pH-dependent activation of cathepsins and metalloproteinases that degrade extracellular matrix and basement membranes (43). In this study, we found that the metastatic UOK262 cells have significantly higher MCT4 expression compared with the localized UMRC6 cells, and also have more rapid export of lactate out of the cells. UOK262 cells have mutations in the TCA enzyme fumarate hydratase, which leads to an uncommon and highly aggressive hereditary RCC. However, recent studies have shown that fumarate hydratase mRNA and protein expression are reduced in clear cell RCC, the most common histologic variant of kidney cancer, promoting tumor migration and invasion (44). The reduced fumarate hydratase leads to accumulation of hypoxia inducible factor-2 alpha (HIF-2 α ; ref. 45), a transcription factor known to promote renal carcinogenesis in part by upregulating glycolysis (17). Thus, the metabolic changes observed in the UOK262 cells are likely not unique to this particular RCC type, and the MCT4-mediated lactate efflux may be an important determinant of RCC aggressiveness in general. Supporting this hypothesis, a recent study showed that MCT4 protein expression in primary clear cell RCCs was associated with poorer relapse-free survival, and correlated with Fuhrman nuclear grade (46). In addition, MCT4 knockdown RCC cell lines had reduced intracellular pH, impaired proliferation, and increased apoptosis (46). These studies indicate that MCT4 targeting may also be an important strategy for the treatment of RCCs.

We showed that the MCT4-mediated lactate efflux in living cells can be explored noninvasively using hyperpolarized ¹³C magnetic resonance. This was accomplished by monitoring the real-time cellular pyruvate-to-lactate fluxes under different flow rates in the bioreactor. While our study used an *ex vivo* system, interrogation of lactate export using hyperpolarized ¹³C magnetic resonance can be achieved *in vivo*. For example, it is possible to measure the tumoral extracellular or interstitial pH, which in part reflects the amount of exported lactate, using hyperpolarized ¹³C bicarbonate magnetic resonance (47, 48). Moreover, it is possible to discriminate the local environment of hyperpolarized metabolites using diffusion weighting *in vivo* (49, 50). Future

studies will develop diffusion-weighted hyperpolarized magnetic resonance that can directly quantify the relative amount of intracellular versus extracellular lactate.

While total lactate levels can also be monitored using ¹H MRS, this approach has limited use in the metabolic evaluation of renal tumors, particularly in the *in vivo* setting. Lactate and lipid peaks usually overlap such that the assessment of lactate is challenging even when methods for lipid suppression are applied. More importantly, the real-time metabolic fluxes, influenced by enzymatic and transporter expression, cannot be captured using ¹H MRS.

In conclusion, we have shown that hyperpolarized ¹³C-pyruvate MRS enables real-time observation of differential lactate efflux, mediated by MCT4, in living RCC cells of varying aggressiveness. Importantly, as MCT4 and lactate efflux are implicated in the pathogenesis of many types of cancers, hyperpolarized ¹³C MRS has the potential to noninvasively interrogate tumor aggressiveness and treatment efficacy in a broad range of cancers.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Authors' Contributions

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Hyperpolarized ^{13}C -Pyruvate Magnetic Resonance Reveals Rapid Lactate Export in Metastatic Renal Cell Carcinomas

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