Research Article

Integration of flux measurements to resolve changes in anabolic and catabolic metabolism in cardiac myocytes

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Although ancillary pathways of glucose metabolism are critical for synthesizing cellular building blocks and modulating stress responses, how they are regulated remains unclear. In the present study, we used radiometric glycolysis assays, [13C6]-glucose isotope tracing, and extracellular flux analysis to understand how phosphofructokinase (PFK)-mediated changes in glycolysis regulate glucose carbon partitioning into catabolic and anabolic pathways. Expression of kinase-deficient or phosphatase-deficient 6-phosphofructo-2-kinase/fructose-2,6-bisphosphatase in rat neonatal cardiomyocytes co-ordinately regulated glycolytic rate and lactate production. Nevertheless, in all groups, >40% of glucose consumed by the cells was unaccounted for via catabolism to pyruvate, which suggests entry of glucose carbons into ancillary pathways branching from metabolites formed in the preparatory phase of glycolysis. Analysis of 13C fractional enrichment patterns suggests that PFK activity regulates glucose carbon incorporation directly into the ribose and the glycerol moieties of purines and phospholipids, respectively. Pyrimidines, UDP-N-acetylhexosamine, and the fatty acyl chains of phosphatidylinositol and triglycerides showed lower 13C incorporation under conditions of high PFK activity; the isotopologue 13C enrichment pattern of each metabolite indicated limitations in mitochondrial aspartate, acetyl CoA and fatty acids. Consistent with this notion, high glycolytic rate diminished mitochondrial activity and the coupling of glycolysis to glucose oxidation. These findings suggest that a major portion of intracellular glucose in cardiac myocytes is apportioned for ancillary biosynthetic reactions and that PFK co-ordinates the activities of the pentose phosphate, hexosamine biosynthetic, and glycerolipid synthesis pathways by directly modulating glycolytic intermediate entry into auxiliary glucose metabolism pathways and by indirectly regulating mitochondrial cataplerosis.

Introduction

Glycolysis is the central pathway of intermediary carbon metabolism. Not only does catabolism of glucose via glycolysis provide usable energy for the cell, but it also contributes glycolytic intermediates that serve as branch-point metabolites for the entry of glucose-derived carbon into ancillary pathways of glucose metabolism. These pathways are required for de novo synthesis of cellular components. For example, the activity of the pentose phosphate pathway (PPP) is required for nucleotide biosynthesis and the maintenance of cellular redox state, whereas the hexosamine biosynthetic pathway (HBP) regulates post-translational protein modifications, protein processing, and stress responses. The synthesis of glycerolipids and serine is also dependent on the production of

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glycolytic intermediates [1]. In the heart, the PPP is activated during hypertrophy [2–4] and heart failure [5], and modulating the activity of this pathway regulates the severity of cardiac pathology [6–11]. Similarly, O-linked β-N-acetylglucosamine (O-GlcNAc)-modified proteins, which require synthesis of the sugar donor UDP-GlcNAc via the HBP [12,13], are elevated in hypertrophied and failing hearts [14–17]. Although less is known about the glycerolipid synthesis pathway (GLP), increased activity of this pathway may underlie pathologic cardiac hypertrophy [18]. In addition, numerous studies have identified aberrant ancillary pathway behavior in the context of diabetes (e.g. as reviewed in refs [19,20]).

The activities of ancillary pathways of glucose metabolism are regulated in part through expression of their rate-limiting or committed-step enzymes, cofactor availability, allosterism, and/or post-translational modifications [21–27]. Flux through these pathways may also be regulated by other means. Specific enzymatic reactions in the glycolytic pathway, especially the phosphofructokinase 1 (PFK1) reaction, can co-ordinate the activity of ancillary pathways in several cell types [28–31]; however, the effects of PFK activity and the glycolytic rate on biosynthetic pathway flux in the heart remain unclear. Understanding how PFK regulates ancillary biosynthetic pathway activity is important because its activity is modulated by increased work ex vivo [32,33], and catecholamines [34] and pressure overload in vivo [35].

In the present study, we examined how PFK activity regulates biosynthetic pathways in cardiomyocytes. For this, we altered PFK1 activity by expressing mutant forms of phosphofructo-2-kinase/fructose-2,6-bisphosphatase (PFK2) in primary rat neonatal cardiomyocytes. This method of regulating glycolysis leverages the strong allosteric regulation of PFK1 by F-2,6-P2 [36–39] and is relevant to mechanisms regulating cardiovascular homeostasis. For example, kinases known to be important for cardiac responses to stress [40,41] are strong regulators of PFK2 activity [42–45]. By integrating radiometric assays, stable isotope tracing, and extracellular flux analyses, we demonstrate that the amount of glucose available for entry into ancillary pathways appears relatively high and that the PFK node of glycolysis exerts varying degrees of control over glucose incorporation in the end products of the PPP, the HBP, and the GLP by not only diverting glucose carbon into these pathways, but also by regulating mitochondrial activity and cataplerosis.

**Experimental procedures**

**Materials**

DMEM growth medium, containing l-glutamine, d-glucose, and pyruvate, was purchased from US Biological (Swampscott, MA, U.S.A.). Insulin, transferrin, and selenium (500×) were purchased from Lonza (Walkersville, MD, U.S.A.). Humulin R was from Eli Lilly (Indianapolis, IN, U.S.A.). All other reagents were from Sigma–Aldrich Corp. (St. Louis, MO, U.S.A.), unless indicated otherwise.

**Rodent models**

All procedures were approved by the University of Louisville Institutional Animal Care and Use Committee and were in accordance with the NIH guidelines. The euthanasia procedures were consistent with the AVMA Guidelines for the Euthanasia of Animals. Neonatal rat cardiomyocytes (NRCMs) were isolated from 1- to 2-day-old Sprague–Dawley rats as previously described [46–48]. Culture media were changed to serum-free DMEM 24 h prior to each experiment.

**Gene transfer and expression analysis**

Replication-deficient adenoviral vectors were constructed and purified by Vector BioLabs (Malvern, PA, U.S.A.) using cDNA for mutated forms of rat liver 6-phosphofructo-2-kinase/Fru-2,6-P2 bisphosphatase [PFKFB1 (6-phosphofructo-2-kinase/fructo-2,6-bisphosphatase isoform 1) isoform of PFK2]. The enzyme was engineered to have single amino acid point mutations to foster a kinase-deficient (S32D and T55V; Glyco Lo) or phosphatase-deficient (S32A and H258A; Glyco Hi) form of PFK2 [37,38,49–51]. A 1.4 kb BamHI/Hne1 fragment of a pLenti6-3×FLAG-pd-PFK2 plasmid was subcloned into a Dual-CCM(+) shuttle vector, which has dual CMV promoters to drive both green fluorescent protein (GFP) and the insert. The backbone of the adenoviral vector is type 5 (dE1/E3). An Ad-GFP control virus, in which GFP is driven by the CMV promoter, was also purchased from Vector BioLabs.

Cardiac myocytes were transduced with adenovirus for 4 h in fresh media, followed by PBS wash and the addition of fresh media. For all viral treatments, a multiplicity of infection (MOI) of 50 was used. Viral transduction and functional expression of the target protein was confirmed by eGFP fluorescence, immunoblotting,
and glycolysis assays. For measuring protein abundance, cells were lysed in a common lysis buffer containing 20 mM HEPES, 110 mM KCl, 1 mM EDTA, 1% NP-40, 0.1% SDS, and phosphatase and protease inhibitors. Cell lysates were then centrifuged at 13,000g for 20 min and the supernatant was collected. Protein was measured using the Lowry DC assay kit (Bio-Rad Laboratories, Hercules, CA, U.S.A.) and separated by SDS-PAGE, electroblotted onto PVDF membranes, and probed for PFKFB1 (Novus Biologicals, Littleton, CO, U.S.A.) according to the manufacturer’s protocol. A horseradish peroxidase-linked secondary antibody (Cell Signaling Technology, Danvers, MA, U.S.A.) was used to detect and visualize the protein band with a Typhoon 9400 variable mode imager (GE Healthcare, Chicago, IL, U.S.A.).

Radiolabeled glycolysis assay
The rate of glucose utilization was determined using [5-3H]-glucose as previously described [52–54]. Briefly, after 48 h of transduction, the media were replaced with 2 ml of fresh, serum-free media containing 5.5 or 25 mM glucose with the addition of 2 μCi/ml [5-3H]-glucose (Moravek Biochemicals, Brea, CA, U.S.A.). Following incubation for 3 h, 100 μl of medium was collected and added to 100 μl of 0.2 M HCl in a microcentrifuge tube. This tube, with the tube cap removed, was placed in a scintillation vial containing 500 μl of dH2O to allow for evaporative diffusion of [3H]2O in the microcentrifuge tubes into the scintillation vials. To account for incomplete equilibration of [3H]2O and background, known amounts (μCi) of [5-3H]-glucose and [13C][D2]-glucose (Moravek Biochemicals) were placed into microcentrifuge tubes and these were placed into separate scintillation vials containing 500 μl of dH2O. After incubation at 37°C for 72 h, the microcentrifuge tube was removed from the vial, 10 ml of scintillation fluid was added, and scintillation counting was performed using a Tri-Carb 2900TR Liquid Scintillation Analyzer (Packard Bioscience Company, Meriden, CT, U.S.A.). Glucose utilization was then calculated as reported by Ashcroft et al. [52], with considerations for the specific activity of [5-3H]-glucose, incomplete equilibration and background, dilution of [5-3H]- unlabeled glucose, and scintillation counter efficiency. To normalize glucose utilization to total protein, the cells were first washed with PBS (to remove adherent radioactivity) and then lysed in common lysis buffer. After centrifugation, protein concentration was measured using the Lowry assay kit (Bio-Rad Laboratories).

Stable isotope tracing
NRCMs were incubated in six-well plates for 18 h in glucose-free DMEM containing 1 mM pyruvate, 4 mM glutamine, and 25 mM [13C6]-glucose. Cell reactions were then quenched in cold acetone and extracted in acetonitrile:water:chloroform (v/v/v, 2:1.5:1), as described previously [39,55–57], to obtain the polar, nonpolar, and insoluble proteinaceous fractions. The nonpolar (lipid) layer was collected, dried under a stream of nitrogen gas, and reconstituted in 0.1 ml of chloroform:methanol:butylated hydroxytoluene (2:1 + 1 mM) mixture. The extract was diluted 10x with 1 mM butylated hydroxytoluene solution in methanol and used for Fourier transform ion cyclotron resonance-mass spectrometry (FTICR-MS) analysis.

For stable isotope nucleotide analysis, the samples were prepared using a previously published protocol [39,57], with slight modifications. Briefly, lyophilized polar extracts were reconstituted in 50 μl of 5 mM aqueous hexylamine, adjusted to a pH of 6.3 with acetic acid (solvent A). Samples were then loaded onto a 100-μl capacity C18 tip (Thermo Fisher Scientific, Waltham, MA, U.S.A.) followed by washing twice with 50 μl of solvent A. The metabolites were eluted with 70% solvent A and 30% 1 mM ammonium acetate in 90% methanol (pH 8.5; solvent B). The resulting eluates were diluted 3x with methanol and analyzed via FTICR-MS.

[13C6]-glucose and [13C3]-lactate measurements in the media
For quantification of [13C6]-glucose and [13C3]-lactate in cell media, 10 μl of media were mixed with 9990 μl of 5 mM NH4OH in methanol containing 5 μM [D2]-glucose and 5 μM [D2]-lactate internal standard and used without any additional pretreatment for FTICR-MS analysis and quantification. Blank incubations without cells were run in parallel. The concentrations of [13C]-labeled glucose and lactate were assessed using seven-point calibration curves (0.3125–20 μM).

FTICR-MS analysis
Lipid and nucleotide spectra and [13C]-glucose/lactate spectra were acquired using a hybrid linear ion trap-FTICR mass spectrometer (Finnigan LTQ FT; Thermo Electron, Bremen, Germany), equipped with a TriVersa NanoMate ion source (Advion Bio-Sciences, Ithaca, NY, U.S.A.) with an A electrospray chip (nozzle...
inner diameter, 5.5 μm). The TriVersa NanoMate was operated by applying 1.5 kV with 0.5 psi head pressure in the positive ion mode and 1.6 kV and 0.7 psi in the negative ion mode. High mass accuracy data were collected using the FTICR analyzer over a mass range from 150 to 1600 Da (lipids; + and − mode) for 15 min at a target mass resolution of 400 000 at 400 m/z. For quantification of 13C6-glucose and 13C3-lactate in cell media, data were collected in the negative ion mode for 2 min over a mass range from 50 to 250 Da at a target mass resolution of 100 000 at 400 m/z. Prior to acquisition, the LTQ-FT was tuned and calibrated according to the manufacturer’s default recommendations to achieve a mass accuracy of 2 ppm or less.

**Stable isotope data analyses**

The FTICR MS data were exported as exact mass lists into a spreadsheet file using Qual-Browser 2.0 (Thermo Electron). Isotopologue peak deconvolution, assignment, natural isotope abundance stripping, and quantification were performed using MetSign software, as described previously [39,58].

**Extracellular flux analysis of cellular energetics**

To measure cellular energetics in intact cardiomyocytes, the Seahorse Bioscience XF24 extracellular flux analyzer was used as described previously [59]. For this, 75 000 cells were plated into each well. An hour before each experiment, the media were replaced with 675 μl of assay media, i.e. unbuffered DMEM containing glucose (5.5 mM), glutamine (4 mM), and pyruvate (1 mM, pH 7.4). Following 1 h in a 37°C, CO2-free incubator, the cells were placed in the instrument for analysis. Basal oxygen consumption rates (OCRs) and extracellular acidification rates (ECARs) were measured using a programmed protocol: 3 cycles of 2 min mix, 2 min wait, and 3 min measure. To interrogate the effects of each treatment on cardiomyocyte bioenergetics, the following compounds were injected into each well with 2 cycles of 2 min mix, 2 min wait, 3 min measure following each injection: Port A, oligomycin (Oligo; 1 μM); Port B, carbonyl cyanide-p-(trifluoromethoxy)phenylhydrazone (FCCP; 1 μM); and Port C, antimycin A, and rotenone (AA/Rot; 10 μM/1 μM). The OCR and ECAR values were normalized to the total amount of protein in each well.

**Statistical analysis**

All values are mean ± SEM. All comparisons were made to the relevant control group and performed using one-way analysis of variance (ANOVA) with Dunnett’s correction for multiple comparisons. The null hypothesis was rejected when \( P < 0.05 \).

**Results**

**PFK regulates glycolysis and the distribution of glucose**

To understand the role of PFK activity in cardiac myocyte metabolism, we expressed kinase-deficient (GlycoLo) or phosphatase-deficient (GlycoHi) PFK2 mutant constructs in NRCMs, which are conducive to well-controlled and detailed metabolic flux analyses. The mutant genes were driven by a CMV promoter (Figure 1A), and the bicistronic vector also contained GFP, which was used to determine relative transduction efficiency. As shown in Figure 1B, the vast majority of NRCMs were transduced with virus. These PFK2 mutants were intentionally derived from the liver-specific isoform of PFK2 (i.e. PFKFB1), so that expression of the proteins could be assessed by immunoblotting, without obfuscation by other isoforms (e.g. PFKFB2/PPFKFB3). Myocytes transduced with the GlycoLo or the GlycoHi constructs showed strong bands at the predicted molecular mass for the mutant forms of PFK2, whereas cells transduced with Ad-GFP showed no expression of PFKFB1 (Figure 1C).

Although GlycoHi-expressing cells routinely displayed a higher level of mutant protein expression relative to GlycoLo-expressing cells, the strength of the relative effect of each PFK2 mutant on the cellular glycolytic rate was equal, with a 2.1-fold increase and 2.2-fold decrease in glycolytic rate for the GlycoHi and GlycoLo constructs, respectively (Figure 1D). This effect on glycolytic rate was independent of glucose concentration, as rates were similar for groups incubated with either 5.5 or 25 mM glucose.

To examine the effect of each mutant PFK2 enzyme on aerobic glycolysis, we replaced the normal growth media with media containing 13C6-glucose and, after 18 h, measured the relative levels of 13C6-glucose and 13C3-lactate in the media. As shown in Figure 1E, the levels of 13C6-glucose consumed by the cell were 1.8-fold higher in GlycoHi-expressing cells compared with those in Ad-GFP, with no difference in GlycoLo-expressing cells. Levels of 13C3-lactate extruded into the media were 2.5-fold higher in GlycoHi cells compared with GFP control cells and >10-fold higher compared with GlycoLo virus-transduced cells.
As shown in Figure 1G, 29% of [13C₆]-glucose was metabolized to [13C₃]-lactate in GFP control cells, whereas in GlycoLo-expressing cells, 6% of labeled glucose was metabolized to lactate; in GlycoHi-expressing cells, 44% of labeled glucose was metabolized to lactate. The assessment of [13C₆]-glucose uptake, [13C₃]-lactate production, and [5-3H]-glucose utilization provides estimates of not only how much glucose entered the cell and was later extruded as lactate, but also how much made it through the enolase step of glycolysis, where the tritiated label is eliminated as 3H₂O. The mean values of glucose uptake and lactate production (provided by stable isotope tracing) and of radiometric measurements were used to calculate the amount of glucose unaccounted for by catabolism to pyruvate. In GFP control cardiomyocytes, only 44% of glucose was catabolized past the enolase step in glycolysis; of this fraction, the majority (65%) was metabolized to lactate (Figure 2A,B). These data show that, under control conditions, a substantial amount of glucose taken up by NRCMs is not catabolized to pyruvate, and that the majority of glucose that is catabolized to pyruvate is fated for lactate production.

As shown in Figure 1E, glucose uptake was not different between GFP and GlycoLo cells, yet GlycoLo myocytes demonstrated 2-fold lower glucose utilization through the enolase step of glycolysis and 5-fold lower lactate production. In these cells, 78% of glucose-derived carbon was partitioned upstream of the enolase reaction, leaving 22% available for pathways downstream from enolase (Figure 2B). Although lactate production was markedly suppressed in GlycoLo cells, the absolute amount of glucose-derived carbon available for either mitochondrial oxidation, transamination to alanine, or carboxylation to oxaloacetate was identical with that of...
control cells. Interestingly, while GlycoHi cells showed 80% higher levels of glucose uptake compared with GFP control cells (see Figure 1E), 44% of glucose (by mass) was not catabolized via glycolysis. Nevertheless, a portion of glucose equivalent to that apportioned upstream of enolase (0.67 nmol/h/µg protein) taken into the cell could be accounted for as lactate (0.66 nmol/h/µg protein), and 21% of glucose carbon that was catabolized past the enolase reaction was available for fates other than lactate production (Figure 2B).

**Figure 2. Phosphofructokinase-mediated glucose distribution in isolated cardiomyocytes.**
Schematics illustrating glucose utilization in myocytes transduced with GFP, GlycoLo, or GlycoHi adenoviruses. Measurements from radiometric and isotopic glucose utilization assays were used to delineate glucose distribution in neonatal rat cardiomyocytes transduced with each virus. (A) Schematic illustrating glucose distribution in control (GFP) myocytes. Distributions of glucose are drawn to relative scale. (B) Pie chart illustrating the distribution of glucose: numbers (in nmol/h/µg protein) and percentages (%) indicate the absolute and relative distribution of glucose ascribed to the portions of glucose metabolism: upstream of enolase (light red), through enolase (gray shading), converted into lactate (yellow), and available for transamination to alanine, anaplerotic formation of oxaloacetate (OAA), or glucose oxidation (light blue). Pie charts are drawn to scale. Both schematics consider glucose uptake and lactate production provided by [13C6]-glucose tracing and by glucose utilization measured by radiometric assays, respectively.

**PFK activity regulates glucose carbon partitioning into the PPP in NRCMs**
To determine how the activity of PFK regulates critical anabolic reactions derived from glycolysis, we modulated the glycolytic rate by expressing the GlycoLo or GlycoHi PFK2 mutants in cardiomyocytes and then assessed carbon incorporation into end products of ancillary biosynthetic pathways using stable isotope-resolved metabolomics [60]. Expression of the GlycoHi mutant modestly diminished incorporation of glucose-derived carbon into ATP. Lower fractional enrichment of 13C in GlycoHi-expressing cells is attributable to decreased carbon allocation into the phosphoribosyl ring of the nucleotide (m+5 isotopologue; Figure 3A,B), suggesting diminished flux of 13C-labeled glucose-6-phosphate (G6P) through the oxidative PPP (oxPPP). The
higher level of the \(m+0\) isotopologue of ATP (i.e. the unlabeled ATP pool) in Glyco\(^{11}\) cells further indicates that the flux of \(^{13}\)C\(_6\)-G6P through the oxPPP was lower in Glyco\(^{11}\) cells. Similar changes were observed in other purines such as GTP (Figure 3C). Myocytes expressing the Glyco\(^{0}\) mutant showed \(^{13}\)C fractional enrichment values into ATP that were similar to GFP-transduced control cells (Figure 3A,B), suggesting that low PFK activity does not strongly affect purine synthesis in these cells.

For the pyrimidines CTP and UTP, we found a higher \(m+0\) isotopologue value in Glyco\(^{11}\) cells and a lower \(m+0\) isotopologue value in Glyco\(^{0}\) cells compared with GFP controls. These data suggest that high rates of

![Figure 3. Phosphofructokinase co-ordinates PPP activity.](image)

Isolated cardiomyocytes were transduced with GFP, Glyco\(^{10}\), or Glyco\(^{11}\) adenoviruses and incubated with \([^{13}\text{C}_6]\)-glucose for 18 h. Stable isotope tracing of nucleotides in NRCMs incubated with U-[\(^{13}\text{C}_6\)]-glucose: (A) atom-resolved map illustrating the biological and biochemical history of \(^{13}\)C incorporation into the purine ATP; fractional enrichment values of \(^{13}\)C in (B) ATP and (C) GTP after incubation with [\(^{13}\text{C}_6\)]-glucose; (D) atom-resolved map illustrating the biological and biochemical history of \(^{13}\)C incorporation into the pyrimidine UTP; fractional enrichment values of \(^{13}\)C in (E) UTP and (F) CTP after incubation with [\(^{13}\text{C}_6\)]-glucose. Colored -axis labels correlate to the labeled molecule biosynthetic components indicated in the atom-resolved maps. Graph represents three replicates per group from one isolation, \(^*P<0.05\), \(^{**}P<0.01\), \(^{***}P<0.001\), \(^{****}P<0.0001\).

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glycogenolysis diminish incorporation, and that low rates of glycogenolysis increase incorporation of glucose-derived carbon into pyrimidines (Figure 3D,E). Interestingly, both the GlycoLo and the GlycoHi NRCMs showed increased levels of CTP and UTP having only the ribose ring labeled (m+5 isotopologue). These data could indicate that either decreasing or increasing glycogenolysis augments oxPPP flux. This interpretation would be consistent with the observation that both GlycoLo and GlycoHi cells show more glucose available for entry into ancillary biosynthetic pathways (see Figure 2); however, it is also possible that higher levels of m+5 labeling could be due to an accumulation of these pyrimidine isotopologues, where the contribution of [13C6]-glucose to de novo synthesis of the uridine ring is diminished. Indeed, examination of the m+7 and m+8 isotopologues of UTP, which probably correspond to the addition of carbons from 13C-labeled aspartate to the molecule (see Figure 3D), shows that both the GlycoLo and GlycoHi cells have less incorporation of 13C, which is known to derive from Krebs cycle-(m+7) or pyruvate carboxylase-engendered (m+8) oxaloacetate. The m+6 isotopologue probably corresponds to complete 13C3-labeling of the ribose ring and 13C1 labeling of aspartate, occurring as a result of scrambling during the second turn of the Krebs cycle, as suggested previously [61]. Overall, these data suggest that high PFK activity modestly diminishes oxPPP flux and 13C incorporation into purines. Pyrimidine synthesis appears to be more robustly affected by PFK activity, with high rates of glycogenolysis diminishing incorporation of glucose-derived carbon into UTP and CTP molecules.

**High PFK activity diminishes glucose-derived carbon incorporation into UDP-N-acetylhexosamine**

The molecular synthesis of UDP-N-acetyl hexosamines requires end products or intermediates of several metabolic pathways, including those from the Krebs cycle, the pyrimidine biosynthetic pathway, glutamine, and glycogenolysis; hence, these pathways collectively contribute to UDP-N-acetylhexosamine (UDP-HexNAc) synthesis (Figure 4A). To determine how the glycogenolytic activity affects glucose-derived carbon allocation into the HBP, we measured fractional enrichment of 13C from [13C6]-glucose into UDP-HexNAc in NRCMs transduced with GlycoLo or GlycoHi virus. After 18 h, >90% of the UDP-HexNAc pool integrated 13C-glucose-derived carbon (Figure 4B), which suggests that UDP-HexNAc has a faster rate of glucose carbon incorporation relative to purines and pyrimidines. That the m+0 isotopologue was higher in GlycoHi cells indicates that high rates of glycogenolysis diminish glucose carbon incorporation into UDP-HexNAc. The lower 13C enrichment of many isotopologues (e.g. m+7, m+14, m+15, and m+16) appeared to be due to diminished input of metabolites that require mitochondrial transformation of 13C-glucose-derived carbon; however, the m +5, m+6, and m+11 isotopologues (which are the forms of UDP-HexNAc with the ribose, hexose, and ribose + hexose moieties constituted with 13C, respectively) demonstrate significantly higher fractional enrichment in GlycoHi cells. Given that both the GlycoLo and the GlycoHi cells had similar levels of glucose available for reactions upstream of enolase (see Figure 2), the simplest explanation for this result is that the mitochondrial cataplerotic metabolites (derived initially from [13C6]-glucose-derived) are limiting for the synthesis of UDP-HexNAc; as a result, the m+5 and m+6 isotopologues of the metabolite accumulate as 13C incorporation from cataplerotic metabolites slows down. This would be consistent with findings in pyrimidines, which suggest that cataplerotic generation of aspartate is limiting for the incorporation of glucose-derived carbon into CTP and UTP.

Although low rates of glycogenolysis did not affect the overall level of incorporation of glucose-derived carbon into UDP-HexNAc, that the m+11 isotopologue in GlycoLo cells was higher than that in control cells suggests that low PFK activity promotes channeling of F6P (fructose-6-phosphate) not only into the HBP (contributing six 13C), but, because the ribose ring is included (contributing five 13C), also into the PPP (Figure 4A,B). Changes in the m+8 isotopologue are difficult to interpret because this isotopologue could consist of 13C-labeled N-acetylhexosamine and stably labeled UDP. Altogether, these results suggest that high PFK activity decreases glucose carbon allocation into UDP-HexNAc primarily via limiting the formation of cataplerotic intermediates (e.g. acetyl CoA, aspartate) derived initially from [13C6]-glucose.

**PKF regulates glycerolipid synthesis in cardiomyocytes**

In cardiac myocytes, glycerolipid synthesis requires free fatty acids (FFAs) and glycerol-3-phosphate (G3P). While FFAs can be derived from transporter-mediated uptake across the plasma membrane [62], myocytes have a relatively low capacity for free glycerol uptake; therefore, there is little contribution of glycerol kinase to G3P synthesis [63,64]. However, G3P can be generated from dihydroxyacetone phosphate (DHAP) by...
glycerol-3-phosphate dehydrogenase (Figure 5A). Nevertheless, it is unclear whether or how the rate of glycolysis affects glycerolipid synthesis. We found that in comparison with cells transduced with GFP virus, the GlycoLo cells showed diminished incorporation of \(^{13}\text{C}\) from \(^{13}\text{C}_6\)-glucose into the glycerol (m+3) moiety of phosphatidylcholine (PC; Figure 5B), phosphatidylinositol (PI; Figure 5C), phosphatidylethanolamine (PE; Figure 5D), and phosphatidylserine (PS; Figure 5E). Incorporation of \(^{13}\text{C}_6\)-glucose-derived carbon into fatty acids in PC, PI, PE, and PS was not different in GlycoLo cells from that in control cells, suggesting that only the synthesis of the glycerol backbone portion of these phospholipids is affected by low rates of glycolysis. High glycolytic rates affected \(^{13}\text{C}\) incorporation into phospholipids to a lesser extent. Although PE synthesis showed a slight increase in \(^{13}\text{C}\) fractional enrichment in glycerol (and a concomitant decrease in fractional enrichment of the unlabeled pool; Figure 5D), PI showed diminished labeling characterized by lower incorporation of \(^{13}\text{C}\) into fatty acids. Glycerol \(^{13}\text{C}\) labeling in PI from GlycoHi cells was identical with that of control cells. This dichotomy in labeling patterns in different phospholipids could suggest the presence of unique enzyme complexes that participate in metabolic channeling for the synthesis of distinct glycerolipids.

Consistent with this idea, triacylglycerol (TAG) labeling was remarkably different from that of other glycerolipids. As shown in Figure 5F, TAG species collectively showed little \(^{13}\text{C}\) in the glycerol moiety. Furthermore, low rates of glycolysis had little effect on glucose carbon incorporation into TAGs; however, GlycoHi cells showed lower \(^{13}\text{C}\) incorporation into fatty acids. Collectively, these results suggest that, in isolated NRCMs, low rates of glycolysis diminish phospholipid synthesis by the DHAP \(\rightarrow\) G3P pathway and that high rates of glycolysis decrease incorporation of glucose-derived carbon into the fatty acyl chains of PI and TAGs.

**PFK activity regulates mitochondrial activity in NRCMs**

Stable isotope labeling patterns of several end products of the PPP, HBP, and GLP suggest that high PFK activity diminishes the relative abundance of those \(^{13}\text{C}\) isotopologues that require metabolites derived from mitochondria. This suggests that cataplerosis [65] is diminished under conditions of high PFK activity. Because cataplerotic reactions depend on mitochondrial activity, we next measured respiration in cells expressing either the control construct (GFP) or the GlycoHi or GlycoLo construct. GlycoHi-expressing cells showed 50% lower basal oxygen consumption, whereas GlycoLo-expressing cells showed significantly higher basal OCR values (Figure 6A,C). These PFK-mediated changes in mitochondrial OCR are probably due to the Crabtree effect, which is characterized by increases in mitochondrial activity under conditions of increased glucose availability or glycolytic activity [54,66–68]. Although ATP-linked OCR was not significantly different between the groups, proton leak was significantly diminished in GlycoHi cells. Upon the addition of a mitochondrial uncoupler, GlycoLo and GlycoHi cells showed relative changes in maximal respiratory rate and in reserve capacity that...
mirrored the pattern found for basal OCR and proton leak (Figure 6A,C). These data are consistent with the notion that high PFK activity diminishes mitochondrial activity, which limits the cataplerotic generation of metabolites required for the synthesis of end products of the PPP, HBP, and GLP.

Although cells transduced with the GlycoHi virus had >2-fold elevation in basal ECAR compared with Ad-GFP controls, we observed no differences in ECAR in NRCMs expressing the Glyco Lo PFK2 mutant (Figure 6B). The addition of oligomycin increased ECAR to similar values in all groups, suggesting that inhibition of mitochondrial ATP production overrides PFK2-mediated changes in glycolysis. Although extracellular acidification is a surrogate measure of lactate extrusion and aerobic glycolysis, protons are also generated during respiration via pyruvate dehydrogenase-mediated production of CO2 (i.e. CO2 $\rightarrow$ HCO$_3^-$ + H$^+$) [69,70]; thus, the inconsistency of ECAR with absolute measurements of the glycolytic rate and $^{13}$C$_3$-lactate in GlycoLo cells (see Figure 1) is probably due to an increase in mitochondrial activity. Taken together, measurements

Figure 5. Phosphofructokinase regulates glycerolipid biosynthesis.
Stable isotope tracing of phospholipids and triacylglycerols in cardiomyocytes incubated with media containing $^{13}$C$_6$-glucose for 18 h: (A) atom-resolved map illustrating the biological and biochemical history of $^{13}$C incorporation into glycerolipids; fractional enrichment values of $^{13}$C into: (B) PC; (C) PI; (D) PE; (E) PS; and (F) TAG. Graph represents three replicates per group from one isolation. *$^P$ < 0.05, **$^P$ < 0.01, ***$^P$ < 0.0001.
from radiometric assays, stable isotope tracing, and extracellular flux analyses indicate that PFK regulates ancillary pathways by not only direct partitioning of glucose (e.g. in the PPP and the GLP) but also regulating mitochondrial cataplerotic reactions.

**Discussion**

Most studies of cardiac metabolism have focused primarily on the catabolic reactions required for energy provision [71]. Considerably less effort has been devoted to understanding the regulation of ancillary pathways of glucose metabolism in the heart and how these pathways integrate with catabolic reactions. In the present study, we examined how PFK activity, which immediately precedes the 'lysis' step in glycolysis, regulates several 6-carbon- and 3-carbon-initiated anabolic pathways that require glycolytic intermediates. Our results demonstrate that, in cardiac myocytes, PFK regulates glucose carbon incorporation into ancillary biosynthetic pathways by: (1) directly regulating carbon flow into purines and phospholipids and (2) indirectly influencing mitochondrial cataplerotic activity required for the generation and subsequent incorporation of glucose-derived carbon into products of the PPP, HBP, and GLP. These studies also illustrate how radiometric measurements, stable isotope tracing, and extracellular flux analyses can be integrated to examine substrate fate and to develop detailed understanding of how interventions simultaneously affect anabolic and catabolic metabolism.

We chose to study the integration of anabolism and catabolism in the context of glycolytic regulation at the PFK node. This enzyme controls 65% of the flux through glycolysis in the heart [72], making it the major regulator of glycolytic activity. It is unlikely that, under the experimental conditions used, glucose entry was a major limiting step for its metabolism because NRCMs express both GLUT1 and GLUT4 [73]. Our stable and

![Figure 6. Phosphofructokinase activity regulates mitochondrial activity in isolated cardiomyocytes.](image)

*Extracellular flux analyses of NRCMs: (A) oxygen consumption traces (OCR), (B) extracellular acidification rate (ECAR), and (C) calculated indices of respiration of cardiomyocytes transduced with GFP, GlycoLo, or GlycoHi adenoviruses. n = 3 independent isolations per group. *P < 0.05, **P < 0.01.*
radiometric isotope tracing measurements show that, under control conditions, <50% of glucose taken into myocytes is catabolized past the enolase reaction. This suggests that a majority of the glucose entering the myocyte is not used for energy production, but is probably used by biosynthetic pathways. Although PFK activity was found to be a potent determinant of glycolysis-engendered lactate production, it did not have diametrically co-ordinate effects on the relative amounts of glucose found upstream of the enolase reaction. Surprisingly, both a decrease and an increase in glycolytic activity led to the accumulation of glucose upstream of the enolase reaction; however, the mechanisms by which low and high PFK activity augment the levels of glucose-derived carbon upstream of the enolase reaction appear to differ: when PFK activity was high, glucose uptake was increased, but glucose utilization was elevated; when PFK activity was low, glucose uptake was the same as that found in control cells, but glucose utilization was remarkably diminished. These results suggest that conditions of low PFK activity promote accumulation of metabolites (probably upstream of the PFK step), and that conditions of high glycolytic activity support levels of glucose-derived carbon upstream of the enolase reaction by augmenting glucose uptake. Additionally, our data suggest that the level of glucose available for glucose oxidation, alanine transamination, or pyruvate carboxylation was relatively stable and independent of PFK activity.

Although studies in some cell types such as adipocytes suggest that the HBP is a relatively minor pathway, utilizing only 2–3% of incoming glucose [74], our stable isotope tracing data suggest that the rate of entry of glucose-derived carbon into this pathway may be higher in cardiac myocytes. The unlabeled pool of UDP-HexNAc was below 10% in all groups, suggesting that sugar donors such as UDP-GlcNAc and UDP-GalNAc are synthesized and utilized at high rates. In contrast, >20% of the pool of nucleotides was unlabeled and 13C enrichment of the glycerol moiety of glycerolipids was below 20%. These findings appear to indicate that the HBP utilizes more glucose-derived carbon than either the PPP or the GLP. Given that up to 20% of glucose taken up by the heart could enter the PPP [75], our comparisons of 13C enrichment in end products of these pathways suggest that HBP flux in cardiomyocytes may be much higher than currently appreciated.

Our results are consistent with previous findings in epithelial cells [30], fibroblasts [76,77], and developing embryos [78], which show that glycolysis regulates glucose entry into the PPP. Our findings are also consistent with computational predictions of the role of PFK in regulating the PPP in the intact heart [72]. Similar to findings in neurons [79], in cardiac myocytes, high glycolytic rates driven by PFK decreased the incorporation of glucose carbon into the ribose ring of purines. This suggests that high rates of glycolysis decrease the flux of glucose through the oxPPP. Nevertheless, we identified remarkable differences in 13C labeling in ATP and GTP versus CTP and UTP. Although both classes of nucleotides use the metabolic precursor ribose-5-phosphate to form the ribose ring, purine ring carbon atoms originate from N10-formyltetrahydrofolate, CO2, and glycine, while pyrimidine ring carbons come from carbamoyl phosphate and glutamine, as well as aspartate generated by cataplerosis [80]. In the time frame examined, purines did not demonstrate incorporation of carbons that derive from metabolic products of the serine biosynthetic pathway and one-carbon metabolism. Hence, it is possible that glycine and N10-formyltetrahydrofolate derived from the serine biosynthetic pathway might be inadequate to participate in purine ring synthesis during the 18 h incubation time used in our experiments. Alternatively, unlabeled serine and folate present in the media could be used by these cells for de novo purine biosynthesis.

The decrease in mitochondrial activity detected by extracellular flux analysis in GlycoHi cells, paired with stable isotope tracing results in pyrimidines, suggests low cataplerotic generation of aspartate. Oxaloacetate generated in the Krebs cycle or via pyruvate carboxylation is transaminated to aspartate, which is subsequently used for pyrimidine biosynthesis. That the m+7 and m+8 isotopologues were lower in GlycoHi cells shows that these pathways were diminished under conditions of high PFK activity. From these data, we inferred that the decrease in de novo pyrimidine biosynthesis in the GlycoHi cells may be due to lower rates of cataplerosis. Because tricarboxylic acid anions are delicately balanced by the input and output of intermediates involved in mitochondrial substrate oxidation, anaplerosis, and cataplerosis [65], we reasoned that mitochondrial activity measurements could provide convergent evidence of PFK-mediated regulation of auxiliary glucose biosynthetic pathways. Indeed, the decrease in mitochondrial activity in GlycoHi cells, paired with the observation that 13C-aspartate entry into pyrimidines was low, suggests that inhibition of pyrimidine synthesis under conditions of high glycolytic activity is due to low rates of cataplerosis.

Similar reasoning as well as previous modeling, validation, and biochemical interpretation of 13C isotopologue data for UDP-GlcNAc [81] were used to determine how PFK affects the HBP, which is important given the known role of UDP-hexosamines in modulating N- and O-linked glycosylation [12,13]. The fact that the
unlabeled pool of UDP-HexNAc was higher in Glyco\textsuperscript{H}i cells suggests that the net effect of high PFK activity is a decrease in HBP flux, and, like that found for pyrimidines, this appears to be in part due to diminished mitochondrial activity. Nevertheless, there are limitations for interpreting current data on HBP flux. Because building UDP-HexNAc requires metabolites from many sources, it is possible that the contribution of carbon from non-labeled pools of metabolites may hinder interpretation. A potential solution to this problem would be to use a pulse-chase approach, where isotopic labeling in the system is first saturated (e.g. with both \textsuperscript{13}C and \textsuperscript{15}N) and then the stably labeled substrate(s) are replaced with unlabeled substrate(s). The disappearance of the stable label over time in isotopologues of UDP-HexNAc should, with higher confidence, reveal which intersecting metabolic pathways limit HBP activity and UDP-Glc(Gal)NAC synthesis under a given set of conditions.

To date, little evidence exists for whether or how glycolysis regulates GLP activity. Interestingly, G3P formed via glyceral-3-phosphate dehydrogenase 1 has been shown to inhibit PFK2 and activate fructose biphosphatase 2 \([82]\), suggesting feedback regulation of the GLP on both glycolytic and gluconeogenic activity. An indication that the glycolytic rate regulates GLP flux and glycerolipid synthesis is provided by studies, showing that genetic deletion or overexpression of PFKFB3, a highly active form of PFK2, in adipose tissue co-ordinately regulates fat deposition and obesity \([83,84]\), which is consistent with the notion that glycolysis in adipocytes regulates TAG synthesis and storage. We found that low PFK activity directly limits glucose-derived carbon incorporation into the glycerol backbone of several phospholipids including PC, PI, PE, and PS; however, this was not the case with triacylglycerols, where \textsuperscript{13}C incorporation into glycerol was not different, but fatty acyl chain labeling was diminished in Glyco\textsuperscript{H}i cells. These results suggest that the fatty acid pool derived from catabolic reactions is differentially channeled to route newly synthesized fatty acids for esterification into triacylglycerols rather than phospholipids.

There are several limitations to our study. Although NRCMs are beating cardiac myocytes, they rely on glycolysis for energy to a much greater extent than the adult heart \([85]\). Thus, findings regarding pathways that involve metabolites generated by cataplerosis may not entirely reflect metabolism in the adult heart. Glucose carbon fate in NRCMs appears to differ from the adult heart primarily at the level of coupling between glycolysis and glucose oxidation. In the adult human heart, previous studies show that 26% of extracted glucose undergoes oxidation and 12% is released as lactate \([86]\). In contrast, in NRCMs, we show that <13% of glucose carbon undergoes oxidation and ~24% is released as lactate. Nevertheless, our finding that a large portion of exogenous glucose extracted by cardiac myocytes is apportioned for metabolic pathways other than glycolysis and glucose oxidation is supported by human \textit{in vivo} isotope tracing studies in the myocardium, which demonstrate that 60% of exogenous glucose extracted by contracting myocytes enters a slow turnover pool amenable to entry into ancillary biosynthetic pathways \([86]\). Another consideration is that the NRCM culture medium is devoid of fatty acids, which could influence the coupling between glycolysis and glucose oxidation as well as biosynthetic pathway activity. Finally, it is important to consider that our stable isotope analyses did not include glycogen, glycine, serine, or sorbitol, which limits our understanding of how glycolysis regulates glycogen biosynthesis, serine biosynthesis, and the polyol pathway. Nevertheless, computational models created by Cortassa et al. \([72]\) suggest that the polyol pathway is under strong control by activation or inhibition of PFK. Similarly, previous studies show that PFK activity regulates glycogen abundance in the heart \([37,38]\). Such a behavior would be consistent with co-ordinate regulation of ancillary biosynthetic pathway activity by PFK.

In summary, by using several methods to measure metabolic pathway activity, we found that >40% of glucose consumed by NRCMs is not catabolized to pyruvate, and that PFK has strong effects on the coupling between glucose oxidation and glycolysis. Pairing results from stable isotope-resolved metabolomics with extracellular flux analysis demonstrated that PFK activity regulates glucose carbon incorporation directly into the ribose and the glycerol moieties of purines and phospholipids, respectively, and that high PFK, by indirectly limiting mitochondrial activity, diminishes \textsuperscript{13}C incorporation into pyrimidines, UDP-N-acetylhexosamine(s), and the fatty acyl chains of phosphatidylinositol and triglycerides. These insights could help elucidate how pathological or physiological changes in glucose metabolism regulate the anabolic behavior and remodeling of the myocardium.

**Abbreviations**

CMV, cytomegalovirus; DHAP, dihydroxyacetone phosphate; ECAR, extracellular acidification rate; F6P, fructose-6-phosphate; FFAs, fatty acids; FTICR-MS, Fourier transform ion cyclotron resonance-mass spectrometry; G3P, glycerol-3-phosphate; G6P, glucose-6-phosphate; GFP, green fluorescent protein; GLP, glycerolipid synthesis pathway; HBP, hexosamine biosynthetic pathway; LTQ-FT, hybrid linear ion trap-FTICR.
mass spectrometer; MOI, multiplicity of infection; NRCM, neonatal rat cardiomyocyte; OCR, oxygen consumption rate; oxPPP, oxidative PPP; PC, phosphatidylcholine; PI, phosphatidylinositol; PE, phosphatidylethanolamine; PG, phosphoglycerol; PS, phosphatidylserine; pd-PFK2, phosphatase-deficient PFK2; PFK1, phosphofructokinase 1; PFK2, phosphofructokinase 2 (also denoted as 6-phosphofructo-2-kinase/fructo-2,6-bisphosphatase); PFKFB1, 6-phosphofructo-2-kinase/fructo-2,6-bisphosphatase isoform 1; PFKFB2, 6-phosphofructo-2-kinase/fructo-2,6-bisphosphatase isoform 2; PFKFB3, 6-phosphofructo-2-kinase/fructo-2,6-bisphosphatase isoform 3; PPP, pentose phosphate pathway; TAG, triacylglycerol; UDP-HexNAc, UDP-N-acetylhexasamine.

**Author Contribution**

A.A.G. contributed to financial support, experimental studies, assembly of data, data analysis and interpretation, and manuscript writing, and gave final approval for publication of the manuscript. P.L. contributed to experimental studies, data analysis and interpretation, and manuscript writing, and gave final approval for publication of the manuscript. Y.Z. contributed to experimental studies and gave final approval for publication of the manuscript. X.Z. contributed to experimental studies and gave final approval for publication of the manuscript. A.B. contributed to manuscript writing and gave final approval for publication of the manuscript. B.G.H. contributed to financial support, concept and design, assembly of data, data analysis and interpretation, and manuscript writing, and gave final approval for publication of the manuscript.

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**Competing Interests**

The Authors declare that there are no competing interests associated with the manuscript.

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