



# Maintaining Myocardial Glucose Utilization in Diabetic Cardiomyopathy Accelerates Mitochondrial Dysfunction

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**Cardiac glucose uptake and oxidation are reduced in diabetes despite hyperglycemia. Mitochondrial dysfunction contributes to heart failure in diabetes. It is unclear whether these changes are adaptive or maladaptive. To directly evaluate the relationship between glucose delivery and mitochondrial dysfunction in diabetic cardiomyopathy, we generated transgenic mice with inducible cardiomyocyte-specific expression of the GLUT4. We examined mice rendered hyperglycemic following low-dose streptozotocin prior to increasing cardiomyocyte glucose uptake by transgene induction. Enhanced myocardial glucose in nondiabetic mice decreased mitochondrial ATP generation and was associated with echocardiographic evidence of diastolic dysfunction. Increasing myocardial glucose delivery after short-term diabetes onset exacerbated mitochondrial oxidative dysfunction. Transcriptomic analysis revealed that the largest changes, driven by glucose and diabetes, were in genes involved in mitochondrial function. This glucose-dependent transcriptional repression was in part mediated by O-GlcNAcylation of the transcription factor Sp1. Increased glucose uptake induced direct O-GlcNAcylation of many electron transport chain subunits and other**

**mitochondrial proteins. These findings identify mitochondria as a major target of glucotoxicity. They also suggest that reduced glucose utilization in diabetic cardiomyopathy might defend against glucotoxicity and caution that restoring glucose delivery to the heart in the context of diabetes could accelerate mitochondrial dysfunction by disrupting protective metabolic adaptations.**

Of the numerous complications associated with diabetes, cardiovascular diseases remain the major cause of death. Both insulin-deficient (type 1) and insulin-resistant (type 2) diabetes are accompanied by a complex milieu of systemic changes including hyperlipidemia and hyperglycemia. It has long been known that despite increased availability of multiple myocardial substrates, in uncontrolled diabetes the heart shows increased fatty acid (FA) utilization and reduced glucose utilization (1). These metabolic changes acting in concert with mitochondrial dysfunction, oxidative stress, and aberrant intracellular signaling are believed to contribute to ventricular dysfunction even in the absence of coronary artery disease and hypertension (2).

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Typically, the healthy heart shows flexibility in its ability to utilize ketone bodies, lactate, FA, and glucose to meet its constant energy demands. These substrates are utilized in a concentration-dependent manner, with a preference for FA. The relative balance between glucose and FA utilization is maintained via allosteric regulation by intermediates of glucose and FA metabolism (3). In diabetes, impaired insulin signaling increases circulating free FA as a result of increased lipolysis (4), which contributes to increased myocardial FA utilization. The mechanisms linking increased FA utilization, lipotoxicity, mitochondrial dysfunction, and diabetic cardiomyopathy are well studied (5). In contrast, many mechanisms including reduced cardiac GLUT4 protein levels contribute to impaired myocardial glucose utilization in diabetes (6), but the contribution of GLUT4 remains incompletely defined. Furthermore, the contribution of glucose toxicity in diabetic cardiomyopathy is less well understood, particularly in light of the relative reduction in myocardial glucose utilization in diabetes. Support for a causative role of increased glucose has been suggested by the emerging relationship between certain glucose-lowering therapies and reduction of heart failure in diabetes (7). Specifically, evidence linking glucose lowering by sodium–glucose cotransporter 2 inhibitors and the reduction of heart failure has reignited interest in the relationship between myocardial energetics and glucose availability in modulating heart failure risk in diabetes (8–10). Thus, increased understanding of mechanisms by which glucose may directly contribute to myocardial dysfunction and mitochondrial impairment, characteristic of diabetic cardiomyopathy, may further inform the complex interaction between myocardial metabolism and heart failure, particularly in the context of diabetes.

*Slc2a4* (a.k.a. GLUT4) is the major mediator of myocardial glucose uptake in the contracting heart (11). Overexpression of GLUT4 in skeletal muscle, adipose tissue, and heart prevents the metabolic abnormalities characteristic of diabetic cardiomyopathy; however, it has not been clear whether these benefits are directly attributable to GLUT4 in cardiomyocytes or improved systemic metabolic homeostasis (12). Increased expression from birth of the related family member *Slc2a1* (i.e., GLUT1) protected against pressure overload–induced heart failure (13). We have more recently reported that inducible GLUT1 overexpression in the context of pressure overload hypertrophy prevented mitochondrial dysfunction without rescuing contractile dysfunction (14). Conversely, loss of GLUT4 expression predisposes the heart to failure in response to either chronic pressure overload or intermittent exercise training (15). Together, these and other studies strongly support the concept that glucose utilization and GLUT expression may play an important role in the myocardial adaptation to hemodynamic stress. In contrast, constitutive increases in glucose uptake might exacerbate ventricular dysfunction in response to diet-induced obesity (16). However, it remains to be determined whether maintenance of normal rates of myocardial glucose uptake and

utilization in the context of diabetes could be cardioprotective. Under conditions of sustained glucose exposure in cultured cardiomyocytes, impaired mitochondrial function has been correlated with the protein posttranslational modification of O-GlcNAcylation that may contribute to decreased mitochondrial complex activity (17,18). In addition to direct modification of mitochondrial proteins, glucose also regulates gene expression by modifying transcription factors and modulating pathways that induce epigenetic changes (19). These data support that under certain circumstances restoring glucose delivery in the context of high extracellular glucose may actually impair cellular function. Thus, increased understanding of mechanisms by which glucose may directly contribute to altered gene expression and differential protein modifications will inform mechanisms linking myocardial mitochondrial impairment to the complex interaction between myocardial metabolism and heart failure in the context of diabetes.

In the current study, we use an inducible cardiac-specific, myc-tagged GLUT4 overexpression mouse model (mG4H) that normalizes myocardial glucose uptake in the context of diabetes. This model also enabled us to define glucose-specific changes in gene expression, mitochondrial protein content, and mitochondrial bioenergetics. We observed that restoration of glucose delivery by maintaining or increasing GLUT4 expression in the context of diabetes enhances glycolytic function but exacerbates the decline in glucose oxidation, prevents the increase in FA oxidation, and accelerates the decline in mitochondrial oxidative phosphorylation (OXPHOS) capacity. These changes were associated with posttranslational modifications (i.e., O-GlcNAcylation) of mitochondrial proteins and transcription factors that contribute to changes in gene expression that would predict mitochondrial impairment. Thus, the decline in cardiac glucose utilization in the context of diabetes may play an adaptive role in reducing glucotoxicity, and circumventing this could worsen outcomes.

## RESEARCH DESIGN AND METHODS

### Generation of Inducible Cardiomyocyte-Specific GLUT4 Overexpression in Mouse

Mice were studied in accordance with protocols approved by the Institutional Animal Care and Use Committee of the University of Utah and University of Alabama at Birmingham. To generate the inducible mycGLUT4 transgene, a *c-myc* epitope-tagged *Rattus norvegicus* GLUT4 cDNA (20) was subcloned into the pTRE2 vector (Clontech Laboratories) that contains a promoter region under regulation of tetracycline response elements and injected into oocytes from FVB/NJ mice. Offspring with germ line transmission were crossed with mice expressing  $\alpha$ MHC-tON (21), for inducible cardiomyocyte expression. For all studies, 8- to 10-week-old double transgenic mice (mG4H) were the experimental group with  $\alpha$ MHC-tON as controls (Con). Both male and female mice were used for a subset of studies, e.g., echocardiography and isolated working heart. As no differences between the sexes were seen, data were

combined. For all other studies, male mice were used. The mice were housed at 22°C on a 12 h–12 h light-dark cycle with ad libitum access to food (8656; Harlan Teklad) and water. The transgene was induced by injection with 100 µg doxycycline (DOX) (500 µg/mL in 0.9% NaCl) (Sigma-Aldrich) and switched to 1 g/kg DOX-containing chow (F5820; Bio-Serv).

### Cell Culture, Promoter Reporter Construction, and Luciferase Assays

#### Cell Culture

C<sub>2</sub>C<sub>12</sub> *Mus musculus* muscle myoblasts were purchased from ATCC (Manassas, VA). Low passage cells (<10) were maintained at 37°C under 5% CO<sub>2</sub> in DMEM containing 4.5 g/L glucose supplemented with 10% FBS. When cells were confluent, cell medium was changed to DMEM supplemented with 2% horse serum for myotube differentiation.

#### Adenoviral Expression Vectors

Adenoviral green fluorescent protein (GFP) and O-GlcNAcase (OGA) have previously been described (22).

#### Mammalian Expression Vectors

pCMV-Sp1 and empty control plasmids were provided by Dr. Jonathan M. Horowitz, and their construction has previously been described (23).

#### Reporter Constructs

The mouse *Ndufa9* gene promoter was generated by PCR amplification of FVB/NJ genomic DNA followed by cloning into the pGL4.1 as previously described (24). All promoter deletions had the same 3′-primer: 5′-GAGAGCTCCAAA TATCCCTTCTCTGC-3′, which has a SacI-cloning site 994 bp downstream from the transcription start site (TSS). The −2 kb, “full-length,” promoter reporter 5′-primer, 5′-ACTGGCCGGTACCACAGGAAG-3′, has a KpnI-cloning site 1,855 bp upstream from the TSS. The deletions had KpnI sites for the −0.5-kb fragment 504 bp, 5′-AGGTACCGT GAACACTAA-3′, and the −0.3-kb fragment 326 bp, 5′-AGG TACCCTGCAGAGCT-3′, upstream from the TSS, respectively.

#### Site-Directed Mutagenesis

Site-directed mutagenesis of the candidate Sp1 response elements (REs) was performed using the QuikChangeII Kit (Agilent, Santa Clara, CA) following the manufacturer’s protocol with the following modifications. Candidate Sp1-REs were mutated as shown in Fig. 5A. The sites were mutated with the following primers: Sp1-RE1, 5′-CAGTGACTA GATGTGTGGGAGCTCGGGTGATACTCAGAGGGGA-3′/5′-TC CCCTCTGAGTATACCCGAGCTCCACACATCTAGTCACTG-3′; Sp1-RE2, 5′-GCGTGCGAGGGGCTTAGAGCTCGGGCGCCTCTGC AT-3′/5′-ATGCAGAGGCGCCGAGCTCTAAGCCCCTCGCACGC-3′; and Sp1-RE3 (M3) 5′-CAGGGGAAAGGGAGCTCGGACGGCA GGAC-3′/5′-GTCCTGCCGTCCGAGCTCCCTTTCCCTG-3′.

#### Luciferase Assays

The constructs described above and shown in Fig. 5A were used for promoter reporter luciferase assays as previously described (24). C<sub>2</sub>C<sub>12</sub> cells were plated at a density of

140,000 cells/mL in 12-well plates and transfected using Lipofectamine 2000 (Invitrogen, Waltham, MA) as previously described (24). Firefly luciferase reporter plasmids were cotransfected with CMV promoter-driven Renilla luciferase to control for transfection efficiency and the appropriate mammalian expression vector (e.g., pCMV-Sp1) or its corresponding empty vector. For luciferase reporter assays, cells were harvested and analyzed using Dual-Glo (Promega, Madison, WI) and were read according to the manufacturer’s protocols.

#### Electron Microscopy

Tissue samples were fixed and processed for histology and electron microscopy as previously described (25). Mitochondrial volume density was determined as the mitochondrion-containing fraction of a 32-by-32 grid placed on pictures of ×3,500 magnification (*n* = 3 hearts and 2–3 pictures/heart). Mitochondrial number was determined in identical-size pictures of ×18,000 magnification (*n* = 3 hearts and 2–3 pictures/heart).

#### Histological Analysis

Fresh hearts were fixed by immersion in 10% buffered formalin and analyzed as previously described (15). Tissue was embedded in paraffin, sectioned, and stained by the manufacturer’s protocol with Masson trichrome for visualization of fibrotic tissue on 5–10 images per group. Light microscopy was performed using a Leica DMRB inverted microscope (Leica Microsystems, Wetzlar, Germany) and captured using XnView software (XnSoft, Reims, France).

#### Protein Analysis by Immunoprecipitation and Western Blotting

Whole-cell, mitochondria-enriched, or nuclear-enriched protein extracts were resolved by SDS-PAGE and following transfer to polyvinylidene fluoride membranes were probed with antibodies as defined below. Detection and quantification were performed by measuring fluorescently labeled secondary antibodies using the Odyssey Infrared Imaging System and accompanying software (version 3.0; LI-COR Biosciences, Lincoln, NE). Immunoprecipitations were performed as follows. Nuclear fractions were isolated from left ventricular tissue, and protein was quantified. A total of 5% of the lysate was saved for input. Anti-O-GlcNAc antibody (5 µg) (RL2, ab2735; Abcam, Cambridge, U.K.) was conjugated with magnetic beads (cat. no. 10004D, Dynabeads; Thermo Fisher Scientific, Waltham, MA) in PBS with 0.02% Tween-20 at room temperature for 90 min. Protein lysate (1 mg) was diluted with 1 mL radioimmunoprecipitation assay buffer and added to the antibody-beads conjugate and incubated overnight at 4°C. Beads were washed five times with radioimmunoprecipitation assay buffer, and protein was eluted with Tris-Glycine SDS sample buffer (cat. no. LC2676; Invitrogen, Carlsbad, CA) containing NuPAGE sample reducing agent (NP009; Invitrogen) and blotted (immunoblot) with the anti-Sp1 antibody (07-645; EMD Millipore, Burlington, MA). Other antibodies used were for 4-aminobutyrates

aminotransferase (ABAT) (ab108259; Abcam), complex IV (CIV) subunit COX4I1 (A21348; Invitrogen), GAPDH (CS-2118; Cell Signaling Technology, Danvers, MA), GlcNAc (CTD110.6, MMS-248R; Covance, Princeton, NJ), GlcNAc (RL2, ab2735; Abcam), GLUT4 (ab654; Abcam), Myc (sc-13922; Santa Cruz Biotechnology, Santa Cruz, CA), complex I (CI) subunit NDUFA9 (459100; Invitrogen), Sp1 (07-645; EMD Millipore), complex II (CII) subunit, SDHB (A21345; Invitrogen), complex III (CIII) subunit UQCRC1 (ab110252; Abcam), and VDAC (PA1-954A; Affinity BioReagents, Golden, CO).

### **In Vivo Cardiac Contractile Function**

#### **Echocardiography**

Echocardiography was performed in a subset of the mice prior to respiration studies as previously described (26). Mice were anesthetized with isoflurane, first at an induction dose of 2%, and then transferred to a warmed platform with a nose cone at 1%–1.5% maintenance isoflurane and imaged in the left lateral decubitus position with a linear 13-MHz probe (Vivid V echocardiograph; GE Healthcare). Cardiac dimensions and function were calculated from these digital images as previously described (27). Diastolic function was assessed with the VisualSonics Vevo 2100 Imaging System (Toronto, Canada) with the mouse under isoflurane anesthesia using the same protocol as above. Spectral Doppler was used to determine transmitral early (E) and atrial (A) wave peak velocities with the ratio of E to A (E/A) calculated. Peak early (E') annular velocities were recorded in tissue Doppler mode.

#### **Left Ventricular Catheterization**

Left ventricular catheterization was performed on an overlapping subset of animals, invasive left ventricular hemodynamic measurements were performed with a temperature-calibrated 1.4-Fr micromanometer-tipped catheter (Millar Instruments, Houston, TX) inserted through the right carotid artery in anesthetized mice and analyzed as previously described (27).

#### **Microarray and Bioinformatics Analysis**

For transcriptomic array-based analysis, left ventricular tissue was harvested from transgenic mG4H or Con mice following 4 weeks of treatment as outlined above ( $n = 6$ ). RNA isolation was performed as described below. Extracted RNA was analyzed to ensure RNA quality, with RNA integrity number  $>7$ . Gene expression intensity was quantified using the Mouse GE 4x44K v2 Microarray (Agilent) by the Health Sciences Center (HSC) Core at the University of Utah as previously described (25). Significant changes in gene expression were then analyzed for functional and network gene set enrichment analysis, along with curated literature-supported candidate upstream regulators, using Ingenuity Pathway Analysis (QIAGEN, Hilden, Germany) unless otherwise specified.

#### **Mitochondrial Purification**

Hearts were freshly excised; the atria were separated from the ventricles, and the right ventricle was dissected from

the left ventricle. Mitochondria were isolated from the left ventricle by differential centrifugation as previously described (28).

#### **OXPHOS Complex Activity Assays**

Activities of individual complexes of the respiratory chain were determined spectrophotometrically, corrected to mitochondrial protein levels, and presented as percent of vehicle (Veh)-treated Con. CI activity, NADH dehydrogenase, was measured as reduction of 2,6-dichloroindophenol by electrons from decylubiquinol following oxidation of NADH by CI (29); CII activity, succinate dehydrogenase, CIII activity, coenzyme Q-cytochrome c reductase, and CIV activity, cytochrome c oxidase, were measured as previously described (30).

#### **Preparation of Isolated Cardiomyocytes and Glucose Uptake/GLUT4 Translocation Assays**

Cardiac myocytes were isolated as previously described (31). Perfused hearts were digested with type I collagenase, and once the heart became translucent and soft, it was minced and myocytes were dissociated by sequential washing in buffer. Cells were pelleted by centrifugation and used for glucose transport assays. Studies commenced after waiting for 90 min to allow cardiomyocytes to attach. Cells were washed and then treated with or without insulin, 2-deoxyglucose (2DG), and 1  $\mu\text{Ci}/\mu\text{L}$  [ $^3\text{H}$ ]2DG (NEN Life Science Products, Boston, MA), and glucose uptake was determined as previously described (14). To measure GLUT4 translocation by exofacial exposure of myc epitope in nonpermeabilized cardiomyocytes, we performed immunofluorescence assay as previously described (32).

#### **Respiration and ATP Measurements in Saponin-Permeabilized Cardiac Fibers**

The respiratory rates of saponin-permeabilized fibers were determined using a fiber-optic oxygen sensor (Ocean Optics, Largo, FL) as previously described (33). Studies were performed with three independent substrates: 1) glutamate/malate, 2) pyruvate/malate, or 3) palmitoylcarnitine/malate. Respiratory rates are defined as follows:  $V_0$ , permeabilized fibers in the presence of substrate;  $V_{\text{ADP}}$ , maximally stimulated respiration following addition of ADP; and  $V_{\text{Oligo}}$ , oligomycin uncoupled, following addition of oligomycin, which inhibits ATP synthase. ATP generation was determined with the ENLITEN ATP Assay System (Promega). ATP/O was calculated as the ratio of ATP synthesis rates and  $V_{\text{ADP}}$ .

#### **RNA Isolation and Quantification**

For each group, six individual male mice with at least one sibling pair in another group were harvested between 6:00 A.M. and 7:00 A.M., right ventricle was dissected away, and left ventricle was immediately placed in RNA, later followed by isolation, using the RNeasy Fibrous Tissue Mini Kit (QIAGEN), purification, cDNA synthesis, and labeling. Either microarray quantification (as described above) or

quantitative PCR (qPCR) was performed as previously described (25). Primer pairs were designed based on GenBank accession numbers with default settings. Dissociation curves followed by agarose gel size confirmation were analyzed for all primer pairs to ensure single product amplification. Oligos for qPCR analysis were used as follows: for transforming growth factor  $\beta$ 1 (*Tgfb1*) NM\_011577.2 forward 5'-CGGAATACAGGGCTTTCGAT-3', reverse 5'-TTCATGTCATGGATGGTGCC-3'; collagen type I  $\alpha$ 1 (*Col1a1*) NM\_007743.3 forward 5'-TACAGTGGATTGCAGGGTCT-3', reverse 5'-TCTACCATCTTTGCCAACGG-3'; transgelin (i.e., SM22 $\alpha$ , *Tagln*) NM\_011526.5 forward 5'-GCGACTAGTGAGTGGATTG-3', reverse 5'-GATCCCTCAGGATACAGGCT-3'; actin  $\alpha$ 2 ( $\alpha$ SMA, *Acta2*) NM\_007392.3 forward 5'-CCGCATGTATGTGGCTATT-3', reverse 5'-AGATAGGCACGTTGTGAGTC-3'; elastin (*Eln*) NM\_012751.1 forward 5'-GCAGAGGAGCAAAGCTTATTTTC-3', reverse 5'-CCAGCACAGCCAAGACATTGT-3'; ribosomal protein lateral stalk subunit P0 (36B4, *Rplp0*) NM\_007475.5 forward 5'-ACCTCCTTCTCCGGCTTT-3', reverse 5'-CTCCAGTCTTTATCAGCTGC-3'; 4-aminobutyrate aminotransferase (*Abat*) NM\_001170978.1 forward 5'-CCAGGAAGCCTTACCGAT-3', reverse 5'-TTGTTGAGCAGGTCTTCCC-3'; and peptidylprolyl isomerase (*Cphn*, *Ppia*) NM\_008907.2 forward 5'-AGCACTGGAGAGAAAGGATTTGG-3', reverse 5'-TCTTCTGTGGTCTTGCCATT-3'.

### Streptozotocin-Induced Diabetes

Mice were made diabetic using the low-dose streptozotocin (STZ) protocol as outlined by the Animal Models of Diabetic Complications Consortium. After 4–6 h of fasting, mice were injected with a stock solution of STZ (7.5 mg/mL) at 55 mg/kg body wt or an equivalent volume of sodium citrate buffer (0.1 mol/L, pH 4.5) for five consecutive days. On day 7 and weekly thereafter, blood glucose was monitored using a standard Contour glucometer (Ascensia, Basel, Switzerland). Mice were considered diabetic if their blood glucose was >200 mg/dL.

### Substrate Metabolism and Contractile Function in Isolated Working Hearts

Cardiac substrate metabolism was measured in hearts isolated from Con and mG4H mice 4 weeks after STZ administration and 2 weeks of DOX treatment. Hearts were prepared and perfused using protocols previously described (15). For determination of metabolism, rates of glycolysis for exogenous glucose, glucose oxidation, and palmitate oxidation were measured over a 60-min period in working hearts, and aortic pressures and cardiac performance including coronary flow were monitored by collection of coronary sinus effluent, with the caveat that Thebesian vein drainage may be included in the measurement, as we have previously described (34).

### Tissue and Serum Metabolite Levels

Blood was obtained from the submandibular facial vein following Goldenrod Animal Lancet (MEDipoint, Inc., Mineola, NY) puncture. Serum was used for measurement of free FA (FFA) and triglyceride (TG) levels. Circulating TG

and FFA concentrations were determined using kits (Wako Diagnostics, Mountain View, CA). Blood glucose was measured from tail vein puncture using the Contour glucometer. Tissue glycogen levels were measured in extract from liver, gastrocnemius, or heart as previously described (35). Results are presented as glucose released from glycogen corrected to tissue weight.

### Two-Dimensional Electrophoresis Analysis and Identification of Glycosylated Mitochondrial Proteins

Two-dimensional PAGE (2D-PAGE) was performed as previously described with the following modifications (36). In brief, following protein separation, gels were stained with Pro-Q Emerald 488 Glycoprotein Gel and Blot Stain Kit (P21875; Thermo Fisher Scientific) following the manufacturer's protocol. Quantitative differences in the protein patterns between the Con-Veh group and the three treatment groups (i.e., Con-STZ, mG4H-Veh, and mG4H-STZ) were analyzed. Spots that significantly changed in these groups were selected using the statistical module of the MELANIE III software package.

### In-Gel Digestion and Mass Spectrometry

Gel spots were destained and subjected to trypsin (20 ng/ $\mu$ L; Promega) digest. Further extraction of peptides from the gel material was performed twice, and these solutions were collected and combined. The supernatant solutions of extracted peptides were combined and dried in a vacuum centrifuge. The samples were then subjected to liquid chromatography (LC)–tandem mass spectrometry (MS/MS) analysis by the HSC Core at the University of Utah. Specifically, peptides were analyzed using a nano-LC-MS/MS system comprised of a nano-LC pump (Eksigent, AB SCIEX, Framingham, MA) and an LTQ-FT mass spectrometer (Thermo Electron Corporation, Waltham, MA). LTQ-FT MS raw data files were processed to peak lists with BioWorks Browser 3.2 software (Thermo Electron Corporation). Resulting DTA files from each data acquisition file were merged, and the data file was searched for identified proteins against the National Center for Biotechnology Information (NCBI) database, mouse taxonomy subdatabase, using Mascot search engine (version 2.2.1; Matrix Science). Identified peptides were accepted only when the Mascot ion score value exceeded 20.

### Quantification and Statistical Analysis

Data are presented as means  $\pm$  SEM. Sample number ( $n$ ) indicates the number of independent biological samples (individual mice or wells of cells) in each experiment. Unpaired Student  $t$  test was used to analyze comparisons between two groups. Data sets with more than two groups were analyzed by two-way ANOVA, and significance was assessed using Tukey post hoc analysis. Statistical calculations were performed using the JMP Pro 9.0 software package (SAS Institute, Cary, NC) or GraphPad Prism 8 software (GraphPad, San Diego, CA), and significance was set at  $P < 0.05$ . For array-based gene expression

analysis, statistical significance was assessed with a Benjamini-Hochberg *P* value adjustment.

### Data and Resource Availability

All data from this article are available from the corresponding authors upon request. The gene expression data discussed in this publication have been deposited in NCBI's Gene Expression Omnibus (GEO) (37) and are accessible through GEO Series accession number GSE123975 (<https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE123975>).

## RESULTS

### Development of Inducible Cardiomyocyte-Specific GLUT4 Transgenic Mice

To modulate glucose uptake in the adult mouse heart, we generated double transgenic mice with DOX-inducible cardiac-specific expression of a myc-tagged GLUT4 transgene (mG4H). Western blot analysis revealed a fivefold increase in GLUT4 protein following 2 weeks of transgene induction (Fig. 1A). Increases in GLUT4 protein levels were observed as early as 2 days after introduction of DOX, reaching maximum levels at about 1 week (Supplementary Fig. 1A). Transgene induction was not observed in single transgenic animals, in double transgenic animals without DOX, or in other tissues examined (Fig. 1A, Supplementary Fig. 1A and B, and data not shown).

### Enhanced Glucose Uptake and Storage Following GLUT4 Transgene Induction

To assess transgene function, we isolated cardiac myocytes from mG4H mice following 2 weeks of DOX treatment. We performed immunofluorescence histochemistry (IHC) for detection of the exofacial myc-tag epitope of the transgene in nonpermeabilized cells. There was patchy GLUT4 protein on the sarcolemma of cardiomyocytes under basal conditions, and recruitment to the sarcolemma was markedly increased in response to insulin treatment (Fig. 1B). Using similarly prepared cells from both littermate Con and mG4H mice, we measured [<sup>3</sup>H]2DG uptake. 2DG uptake was increased 3.2-fold by transgene induction alone, which was similar to insulin-treated cells isolated from Con mice (Fig. 1C). Insulin further increased 2DG uptake by another 2.2-fold above the increased baseline uptake in cells isolated from mG4H mice (Fig. 1C). Compared with hearts of Con mice, mG4H hearts contained ~2.6-fold more glucose stored as glycogen (Fig. 1D) but not in other tissues examined including liver and skeletal muscle (Supplementary Fig. 1C and D).

### Development of Insulin-Deficient Diabetes Is Not Altered by Cardiac GLUT4 Transgene Induction

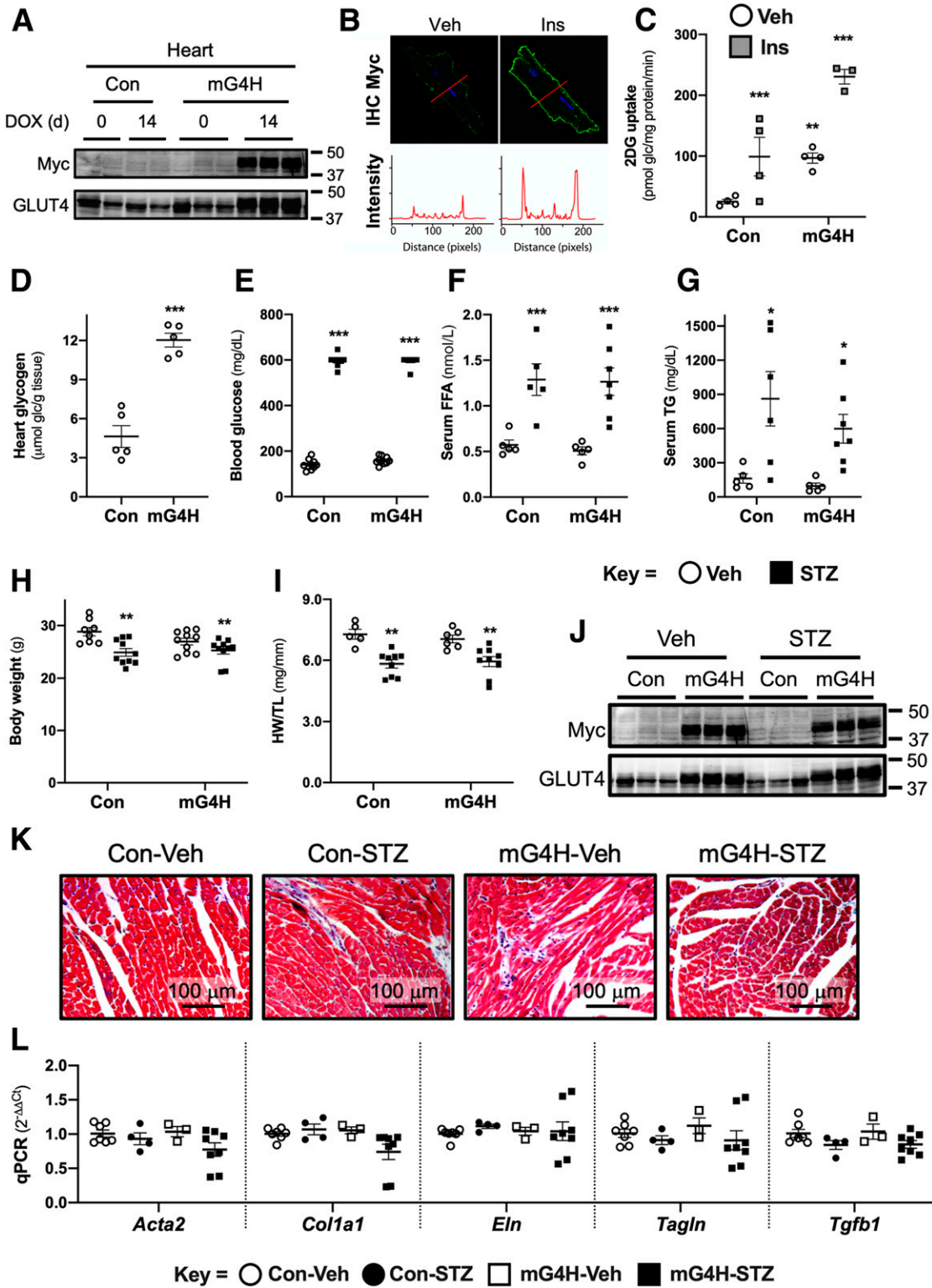
To determine the role of restoring glucose delivery to the heart under diabetic conditions, we used the low-dose STZ model of insulin-deficient diabetes. By 1 week after the initial STZ injection, all treated mice were hyperglycemic (blood glucose >200 mg/dL) compared with Veh-injected

Con. At 2 weeks after the initial STZ or Veh injection, all mice were switched to DOX-containing chow to induce GLUT4 transgene expression in mG4H mice and followed an additional 2 weeks. At this 4-week time point, all STZ-treated mice had blood glucose >500 mg/dL compared with values <150 mg/dL in Veh-injected mice (Fig. 1E). Blood was collected, mice were euthanized, weights were collected, and tissue was snap-frozen for molecular analysis at this 4-week time point. Consistent with the development of diabetes, STZ-treated mice were also hyperlipidemic as shown by significantly elevated serum FFA (Fig. 1F) and TG (Fig. 1G) concentrations. As expected, STZ-induced diabetes resulted in lower body weight (Fig. 1H) and a lower heart weight-to-tibia length ratio (Fig. 1I). Cardiomyocyte-specific overexpression of GLUT4 had no effect on these parameters. Furthermore, Western blot analysis revealed that STZ treatment in turn had no significant effect on DOX-induced GLUT4 overexpression (Fig. 1J). Interestingly, GLUT4 overexpression had relatively little effect on cardiac histology (Fig. 1K) or fibrotic marker gene expression (Fig. 1L).

### Induction of GLUT4 Transgene Following Short-term Hyperglycemia Did Not Alter In Vivo Systolic Cardiac Function or Exacerbate Diastolic Dysfunction

Systolic function was assessed by echocardiography (Fig. 2A). The primary difference observed was reduced cardiac output in STZ-treated animals irrespective of genotype, with a trend toward lower cardiac output in Veh-treated mG4H mice and no differences in ejection fraction (Fig. 2B and Supplementary Table 1). The change in cardiac output may reflect the smaller heart size and body weight. Invasive left ventricular catheterization revealed equivalent degrees of systolic dysfunction in STZ-treated Con and mG4H mice (Supplementary Table 2). Two independent echocardiographic measurements were used to estimate diastolic function: E/A ratio and the E/E' ratio. GLUT4 induction was characterized by decreased A-wave amplitude leading to an increase in the E/A ratio that was independent of STZ treatment (Fig. 2C and D). In Con animals, the mean E/A ratio was higher following STZ treatment, but overlapping values in a subset of Veh-treated Con animals precluded statistical significance. Tissue Doppler imaging showed reduced E'-wave amplitude in both STZ-treated groups leading to comparable elevations in E/E' ratios. Although the mean E/E' ratio was higher in Veh-treated mG4H mice, values were partially overlapping in Veh-treated Con animals, precluding statistical significance (Fig. 2E and F). With results taken together, it appears that short-term STZ-induced diabetes does not impair systolic dysfunction in vivo, but evidence for some degree of diastolic dysfunction is apparent, and depending on the measurement used, transgene induction might also induce diastolic dysfunction. However, there is little evidence of synergistic exacerbation by transgene induction and diabetes.





**Figure 1**—mG4H mice exhibit inducible cardiomyocyte-specific expression of the mycGLUT4 transgene that enhances glucose (glc) uptake and storage in the heart without influencing systemic metabolic homeostasis or cellular structure. **A**: Western blot analysis of whole cell extracts for Myc-tag or GLUT4 protein from biventricular heart tissue in the hearts of single transgenic (Con) and double transgenic (mG4H) mice in the absence or presence of DOX. d, days (additional conditions and tissues described in Supplementary Fig. 1A and B). **B**: IHC for Myc tag of primary adult cardiomyocytes isolated from mG4H mice 2 weeks following DOX induction in the absence or presence of insulin. **C**: Basal and insulin-stimulated 2DG uptake in primary adult cardiomyocytes isolated from Con and mG4H mice following 2 weeks of DOX treatment ( $n = 3-4$ ). Ins, 0.1 nmol/L insulin. **D**: Heart glycogen content of Con and mG4H mice following 2 weeks of DOX treatment ( $n = 5$ ), with additional tissues in Supplementary Fig. 1C and D. **E**: Development of hyperglycemia following STZ treatment compared with Veh-injected mice. Serum FFAs (**F**) ( $n = 5-7$ ) and TG (**G**) ( $n = 5-7$ ). Body weight (**H**) ( $n = 11-14$ ) and heart weight (HW)-to-tibia length (TL) ratio (**I**) ( $n = 5-9$ ) at 4 weeks following Veh or STZ treatment. **J**: Western blot analysis as in A, with STZ treatment ( $n = 3$ ). Histological examination of

### Substrate Metabolism and Ex Vivo Function in Diabetes Are Altered by GLUT4 Transgene Induction

Cardiac glucose utilization by either glycolysis or oxidation was increased in perfused isolated working heart (IWH), following GLUT4 overexpression in mG4H (Fig. 3A and B). As expected, glucose utilization was decreased in hearts of Con-STZ mice. Diabetes-induced repression of glycolysis was reversed with GLUT4 overexpression (Fig. 3A), but impairment in glucose oxidation was exacerbated (Fig. 3B). The expected compensatory increase in FA utilization, as measured by palmitate oxidation in IWH, was observed in hearts of Con-STZ mice (Fig. 3C). Consistent with mitochondrial oxidative impairment as suggested by reduced glucose oxidation, palmitate oxidation failed to increase in hearts of mG4H-STZ mice (Fig. 3C). There were subtle, but statistically significant, increases in basal ex vivo contractile function in hearts of mG4H-Veh mice including aortic flow, cardiac output, and cardiac power, which were reversed in diabetic animals (Fig. 3D–F and Supplementary Table 3).

### Mitochondrial Oxygen Consumption and Respiratory Coupling

Mitochondrial respiratory function was further assessed using saponin-permeabilized cardiac fibers from hearts of Con and mG4H mice in the presence or absence of diabetes. Basal respiration ( $V_o$ ) was significantly reduced only in the mG4H-STZ group with glutamate or pyruvate as substrates (Fig. 4A and B). Maximal glutamate-supported ADP-stimulated mitochondrial oxygen consumption ( $V_{ADP}$ ) was equivalently suppressed by diabetes in Con and mG4H mice and in animals in the mG4H-STZ group. In contrast,  $V_{ADP}$  was suppressed in pyruvate- and palmitoylcarnitine-treated samples only in the mG4H-STZ group (Fig. 4A–C). ATP synthesis was suppressed in the mG4H-STZ group with either pyruvate or palmitoylcarnitine as substrates (Fig. 4E and F) but was unchanged with glutamate (Fig. 4D). Interestingly, the only change in ATP/O ratio was in the mG4H-Veh treatment group incubated with palmitoylcarnitine (Fig. 4G–I). There were no statistically significant changes from GLUT4 overexpression (mG4H) in mitochondrial number, and a modest, but significant, increase in mitochondrial volume was observed in both STZ groups (Fig. 4J–L).

### Mitochondrial Respiratory Complex Activity and O-GlcNAcylation of OXPHOS Proteins and Mitochondrial Enzymes

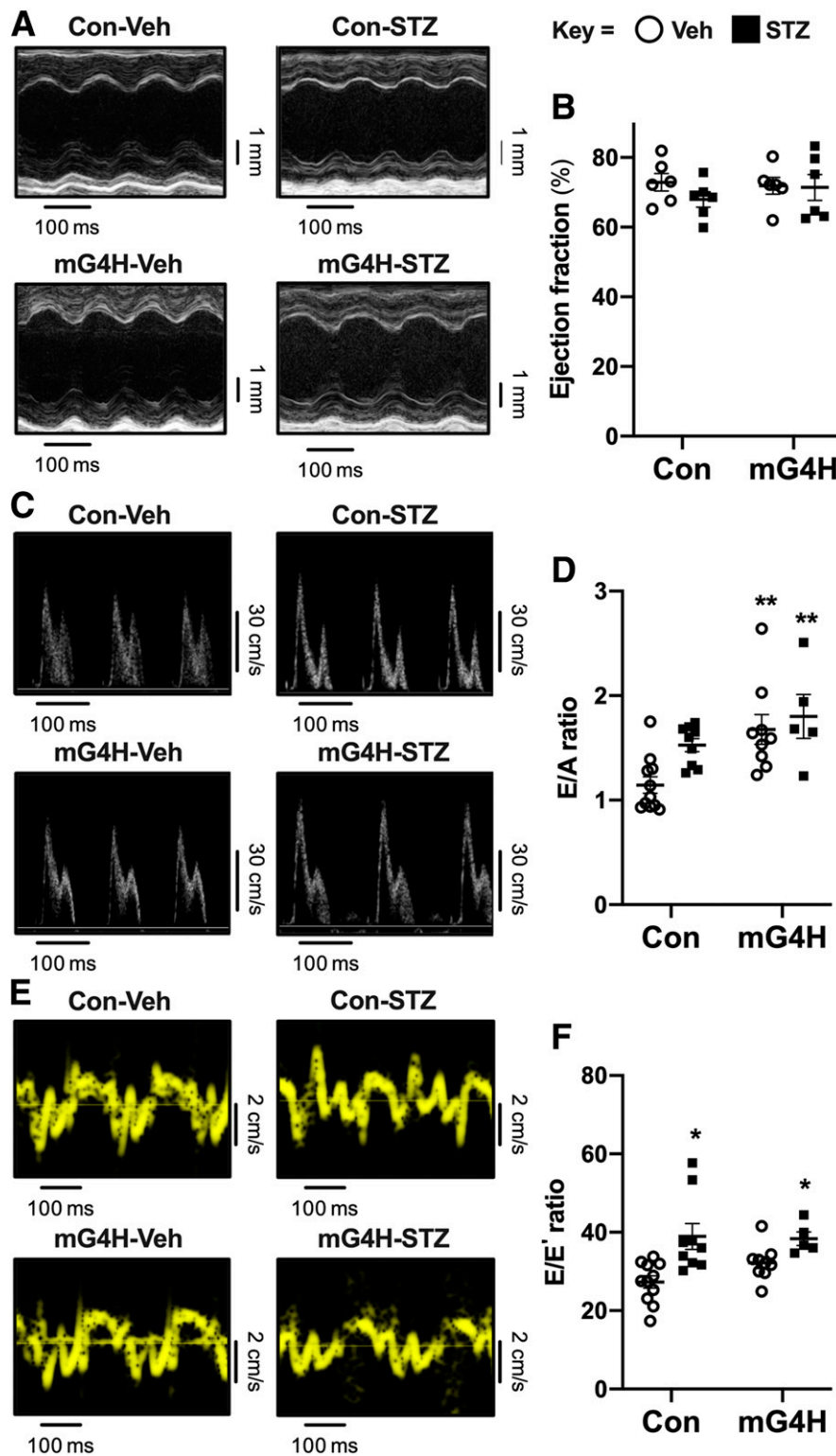
For further elucidation of potential mechanisms for the mitochondrial respiratory defects observed, mitochondria were isolated from the four groups. In mG4H-STZ mice,

there was a decrease in electron transport chain complex activity for all four electron transport chain complexes measured spectrophotometrically (Fig. 5A–D). Diabetes alone reduced CII activity in Con mice (Fig. 5B). Analysis of mitochondrial OXPHOS protein content revealed a modest but significant decrease in CI protein NDUFA9 in mG4H-STZ mice (Fig. 5E and F). There were no significant changes in CII or CIII proteins, but a significant increase in CIV protein COX4I1 was observed in the mG4H-STZ group (Fig. 5G–I). Various mechanisms have been evaluated to determine the role of glucose in regulating mitochondrial protein function, including changes in the posttranslational modification O-GlcNAcylation (38). Western blot analysis of isolated cardiac mitochondrial proteins (as in Fig. 5E) revealed a modest increase in protein O-GlcNAcylation in mG4H-Veh and a robust increase in STZ groups (Fig. 5J). To begin to identify O-GlcNAc-modified proteins, we first used a candidate approach and examined the previously identified NDUFA9 (17). There was a modest increase in protein O-GlcNAcylation of immunoprecipitated NDUFA9 by transgene induction and diabetes (Fig. 5K). We next performed targeted glycoproteomics to identify additional proteins with GlcNAc modifications. Specifically, we used isolated mitochondrial fractions (as in Fig. 5) and separated proteins by two-dimensional PAGE followed by Pro-Q Emerald 488 glycoprotein staining. Fourteen spots were then selected that showed similar staining in each of the three treatment groups relative to Con-Veh and were subjected to MS identification. All 14 proteins had one predominant peptide identified by Mascot database searches (Table 1). Furthermore, all proteins identified are known to be associated with mitochondria, and the majority have previously been shown to possess O-GlcNAcylation sites on proteins in the diabetic rat heart (39). The latter finding supports the conclusion that increasing glucose uptake alone (mG4H) may replicate certain changes in protein posttranslational regulation associated with the complex metabolic milieu that occurs in diabetes.

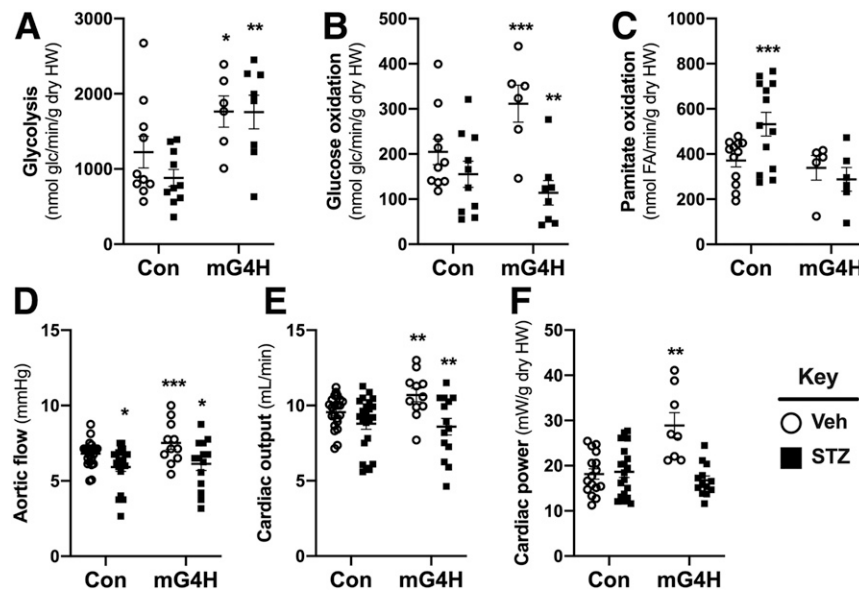
### Expression Levels of the Mitochondrial CI Subunit Ndufa9 Is Directly Regulated by Glucose and O-GlcNAcylation

To further explore the mechanisms by which glucose availability may regulate mitochondrial oxidative capacity, we used a candidate approach to define the molecular pathways by which glucose can alter gene expression. We reasoned that given the widespread changes in mitochondrial protein O-GlcNAcylation, parallel changes could be taking place in the nucleus. We focused initially on the transcription factor Sp1, a known target of regulation by





**Figure 2**—Echocardiographic analysis of in vivo cardiac dimensions and function in diabetic mice with short-term overexpression of GLUT4. Representative M-mode images (A) show preserved systolic function following STZ treatment or GLUT4 transgene induction. B: Quantification of ejection fraction from M-mode; additional parameters are reported in Supplementary Table 1. Transmittal Doppler flow reveals diastolic dysfunction in mG4H mice manifested by increased E/A ratio ( $n = 5-11$ ); representative images (C) and quantification (D). Tissue Doppler imaging also identified an elevated E/E' ratio in STZ-treated diabetic mice independent of genotype ( $n = 5-11$ ); representative images (E) and quantification (F). Quantitative data are means  $\pm$  SEM. \* $P < 0.05$  or \*\* $P < 0.01$  vs. Con-Veh.



**Figure 3**—Cardiac function and substrate metabolism in IWH is altered in diabetic mice with short-term overexpression of GLUT4. Glycolysis (A) ( $n = 6-10$ ), glucose (glc) oxidation (B) ( $n = 6-10$ ), and palmitate oxidation (C) ( $n = 5-13$ ) in IWH from mice 4 weeks after STZ treatment and DOX-induced transgene expression for 2 weeks. DOX was administered 2 weeks after STZ treatment (heart weight [HW]). Ex vivo contractile function of data pooled from both glucose and palmitate metabolism studies for aortic flow (D), cardiac output (E), and cardiac power (F). Additional contractile function data associated with these perfusions are shown in Supplementary Table 3. Quantitative data are means  $\pm$  SEM. \* $P < 0.05$ , \*\* $P < 0.01$ , or \*\*\* $P < 0.001$  vs. Con-Veh.

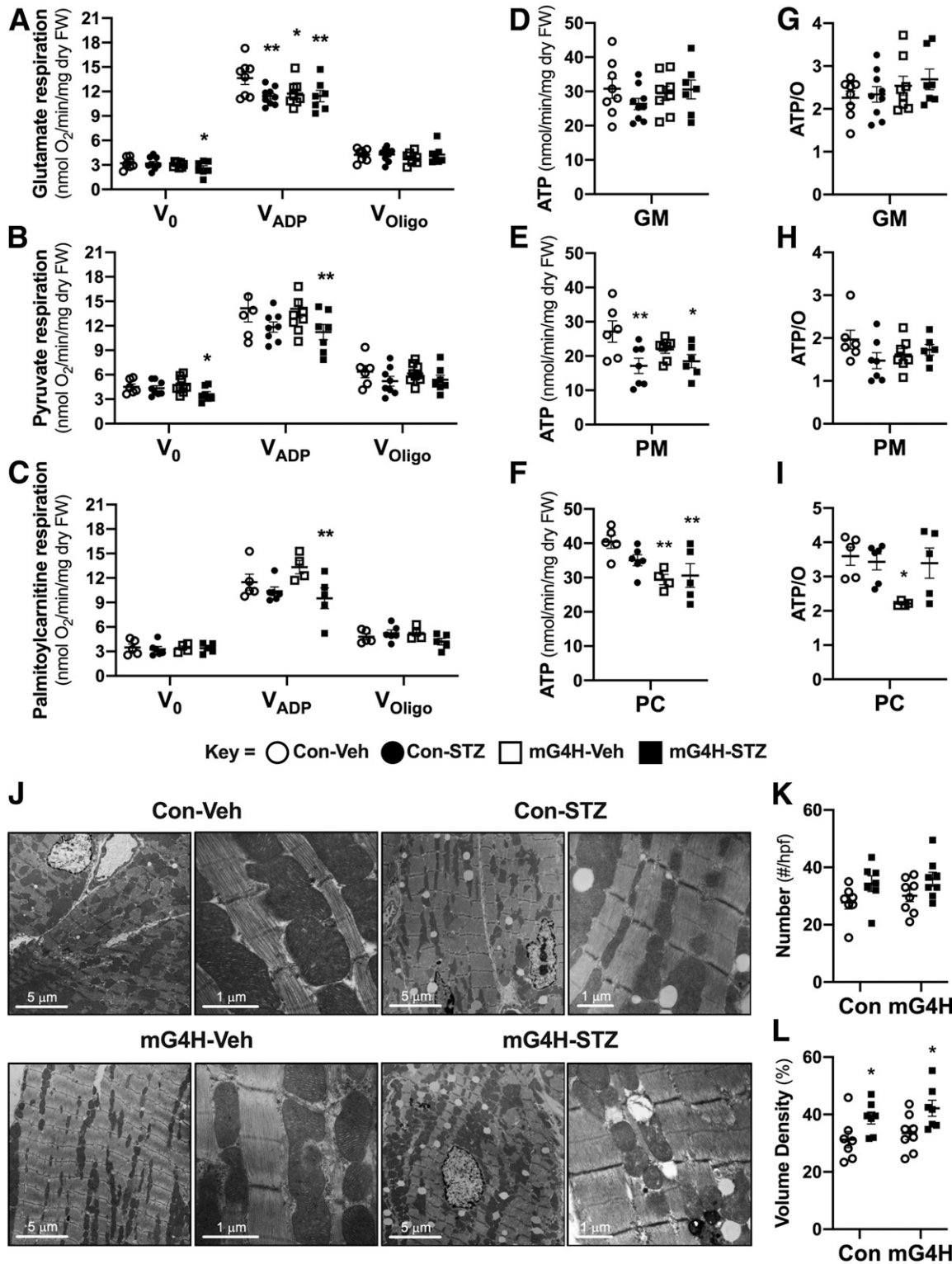
O-GlcNAcylation (40), and determined whether its regulation of mitochondrial OXPHOS gene expression could be modulated by O-GlcNAcylation in the hearts of diabetic and mG4H mice. Transcript levels of the *Ndufa9* subunit of OXPHOS CI were repressed in mG4H-STZ mice (Fig. 5F), suggesting a potential role for glucose-dependent repression of gene expression. Many regions of the *Ndufa9* gene are evolutionarily conserved from human to mouse (Fig. 6A) and harbor candidate Sp1 REs. We cloned a large portion of the mouse *Ndufa9* gene promoter ( $-2$  to  $1$  kb) into a luciferase reporter and generated two deletion mutants that removed some of these candidate REs. Following transfection of the reporter plasmids into  $C_2C_{12}$  myoblasts, cells were differentiated into myotubes. Promoter activity was assessed, revealing it was suppressed by high-glucose treatment ( $25$  mmol/L) in the full-length and first deletion ( $-0.5$  kb). However, this repression was lost upon further truncation (Fig. 6B). We then sequentially mutated each of the three candidate Sp1 REs in the glucose-responsive regions and found that only one site, between the  $-0.5$  and  $-0.3$  kb promoter fragments, was actually required for glucose-mediated transcriptional suppression (Fig. 6C and data not shown). Lastly, we used an Sp1 overexpression plasmid to demonstrate that Sp1 was sufficient to confer transcriptional suppression and further enhanced glucose-mediated repression of promoter activity (Fig. 6D).

To test the hypothesis that O-GlcNAcylation mediates the glucose-dependent transcriptional repression of *Ndufa9* via Sp1, we repeated experiments following adenoviral overexpression of a control GFP or OGA (the enzyme that

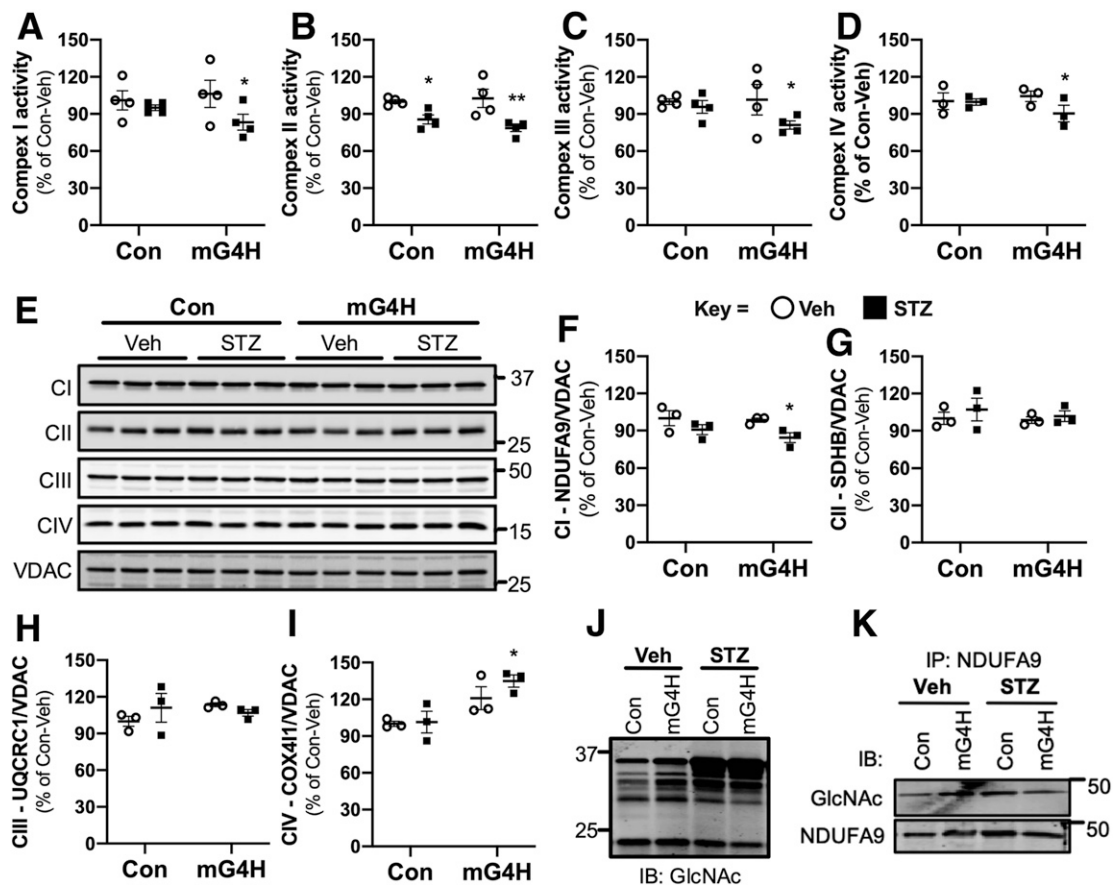
removes protein O-GlcNAcylation). OGA-mediated O-GlcNAc removal prevented glucose-mediated transcriptional repression, which was present when GFP alone was overexpressed (Fig. 6E). To determine whether a similar mechanism existed in the hearts of our mouse models, we performed protein immunoprecipitations from left ventricular nuclear lysates of mice treated as in Fig. 5, using an O-GlcNAc-specific antibody (RL2), and Sp1 was detected by using anti-Sp1 antibody (Fig. 6F). Diabetes induced a modest enrichment of cardiac Sp1 O-GlcNAcylation compared with nondiabetic mice that was enhanced in mG4H-STZ hearts (Fig. 6G). mG4H mice also showed increased glycosylation of cardiac Sp1, further supporting a glucose-specific role for O-GlcNAcylation of Sp1. Collectively, these results suggest that increased glucose availability may independently regulate *Ndufa9* expression via protein O-GlcNAcylation.

#### Identification of Glucose-Regulated Gene Expression Profile in the Heart

To globally identify gene expression pathways impacted by increased myocardial cardiac glucose delivery versus STZ-induced diabetes, we used microarray-based gene expression analysis. Heatmap and hierarchical clustering of differentially expressed genes (DEGs) by STZ treatment ( $P < 0.01$ ) revealed a distinct clustering by genotype within the Veh-treated mice, whereas samples failed to cluster by genotype within the STZ-treated group (Fig. 7A). To examine the genes that were regulated in the mG4H-STZ group, we used a volcano plot of mG4H-STZ DEGs (Fig. 7B). This approach revealed robust changes in numerous



**Figure 4**—Mitochondrial respiratory parameters and ATP synthesis rates are decreased by induction of GLUT4 following short-term hyperglycemia with minor changes in mitochondrial number and structure. Mitochondrial respiration in saponin-permeabilized cardiac fibers examining V<sub>0</sub>, V<sub>ADP</sub>, and V<sub>Oligo</sub> oxygen utilization with glutamate/malate (GM) (A) (n = 7–9), pyruvate/malate (PM) (B) (n = 6–9), or palmitoylcarnitine/malate (PC) (C) (n = 4–6) as substrates (fiber weight [FW]), ATP synthesis rates (D–F) and ATP/O ratios (G–I) in comparably treated fibers from the same mice as in A–C. J: Representative electron micrographs of cardiac tissue; left, low magnification, and right, high magnification of different heart sections. K: Mitochondrial number per high-power field (hpf) (n = 3 hearts). L: Mitochondrial volume density as a percentage (n = 3 hearts). Quantitative data are means ± SEM. \*P < 0.05 or \*\*P < 0.01 vs. Con-Veh.



**Figure 5**—Induction of cardiac GLUT4 in diabetes decreases respiratory chain complex activities in parallel with O-GlcNAc modifications of mitochondrial proteins. Specific respiratory complex activities in mitochondria isolated from Con and mG4H mice treated as in Fig. 1E and beyond. NADH dehydrogenase CI activity (A) ( $n = 4$ ), succinate dehydrogenase CII activity (B) ( $n = 4$ ), coenzyme Q-cytochrome c reductase CIII activity (C) ( $n = 4$ ), and cytochrome c oxidase CIV activity (D) ( $n = 3$ ). E: Representative subunit protein levels from CI to CIV in isolated mitochondria corrected to VDAC. (CI, NDUFA9; CII, SDHB; CIII, UQCRC1; CIV, COX5I1.) F–I: Quantification of data presented in E. J: Mitochondrial protein O-GlcNAcylation in isolated mitochondria. K: O-GlcNAcylation of NDUFA9 as assessed by immunoprecipitation (IP) followed by O-GlcNAc immunoblot (IB). Quantitative data are means  $\pm$  SEM. \* $P < 0.05$  or \*\* $P < 0.01$  vs. Con-Veh.

transcripts encoding enzymes that regulate intermediary metabolism, including pyruvate dehydrogenase kinase 4 (*Pdk4*) (2.9-fold increase,  $P = 1.6 \times 10^{-5}$ ), acyl-CoA thioesterase 1 (*Acot1*) (4.4-fold increase,  $P = 4.2 \times 10^{-7}$ ), and  $\beta$ -hydroxybutyrate dehydrogenase (*Bdh1*) (3.5-fold decrease,  $P = 1.3 \times 10^{-5}$ ). The most differentially suppressed gene, *Abat* (4.8-fold decrease,  $P = 4.3 \times 10^{-9}$ ), was further validated via qPCR (Fig. 7C) and Western blotting (Fig. 7D), which confirmed its repression in mG4H to the same extent as observed with Con-STZ. To further identify those genes whose regulation is mediated by glucose alone, we compared differential expression based on mG4H and/or STZ treatment via Venn diagram. This analysis identified 557 DEGs between pairwise differential expression analyses of mG4H-Veh, Con-STZ, and mG4H-STZ relative to Con-Veh (Fig. 7E and Supplementary Table 4). Three-dimensional scatterplot analysis revealed that the ketone metabolic enzyme *Bdh1* was among the most suppressed (6.0-fold decrease in mG4H-STZ,  $P = 1.2 \times 10^{-5}$ ; 3.7-fold decrease in mG4H,  $P = 7.9 \times 10^{-7}$ ; and 6.1-

fold decrease in STZ,  $P = 6.3 \times 10^{-7}$ ), whereas *Ucp2*, *Acot5*, *Acot1*, and *Pdk4* were among the most highly induced (Fig. 7F). Within the minority of DEGs inversely changed by mG4H alone, the gene encoding  $\beta$ MHC (*Myh7*) was the most differentially suppressed in mG4H (3.0-fold decrease,  $P = 0.004$ ) but was induced in both Con-STZ (3.1-fold increase,  $P = 1.7 \times 10^{-6}$ ) and mG4H-STZ (1.9-fold increase,  $P = 0.004$ ) relative to Con-Veh. Pathway analysis of the overlapping and upregulated DEGs (Fig. 7E [yellow 359]) enriched for FA metabolic pathways (Fig. 7G and Supplementary Table 5), whereas downregulated overlapping DEGs (Fig. 7E [blue 168]) enriched for ketone body and amino acid metabolic pathways (Fig. 7G and Supplementary Table 5). Together, these findings support additional metabolic signaling connections between glucose delivery and regulation of cardiac metabolism.

## DISCUSSION

The objective of the current study was to test the hypothesis that restoring glucose delivery to the heart in the

**Table 1—MS-identified mitochondrial proteins with differential glycosylation in Veh- and STZ-treated Con and mG4H mice**

Protein symbol	UniProt (NCBI) identifier	Description	Known protein O-GlcNAc target ref.†
ACADL	P51174	Acyl-CoA dehydrogenase, long-chain	A
ATP5A1	Q03265	ATP synthase F1 complex, $\alpha$ subunit 1	A, B, C
ATP5H	Q9DCX2	ATP synthase F0 complex, subunit D	Novel
ATP5O	Q9DB20	ATP synthase F1 complex, O subunit	A
GATD3A	Q9D172	Glutamine amidotransferase-like class 1 domain-containing protein 3A	Novel
DLAT	Q8BMF4	Dihydrolipoamide S-acetyltransferase (pyruvate dehydrogenase complex)	A, B
ETFB	Q9DCW4	Electron transferring flavoprotein, $\beta$ polypeptide	A
HADHB	Q99JY0	Hydroxyacyl-CoA dehydrogenase trifunctional complex, $\beta$ subunit	A
MDH2	P08249	Malate dehydrogenase 2, NAD	A
OGDH	Q60597	Oxoglutarate ( $\alpha$ -ketoglutarate) dehydrogenase	A, D
PRDX3	P20108	Peroxiredoxin 3	A
SUCLA2	Q9Z219	Succinate-CoA ligase, ADP-forming, $\beta$ subunit	A
UQCRC1	Q9CZ13	Ubiquinol-cytochrome c reductase core protein 1	B
UQCRC1	Q9CR68	Ubiquinol-cytochrome c reductase, Rieske iron-sulfur polypeptide 1	A

†Numbers in reference (ref.) list correspond as follows: A (38), B (54), C (55), D (56).

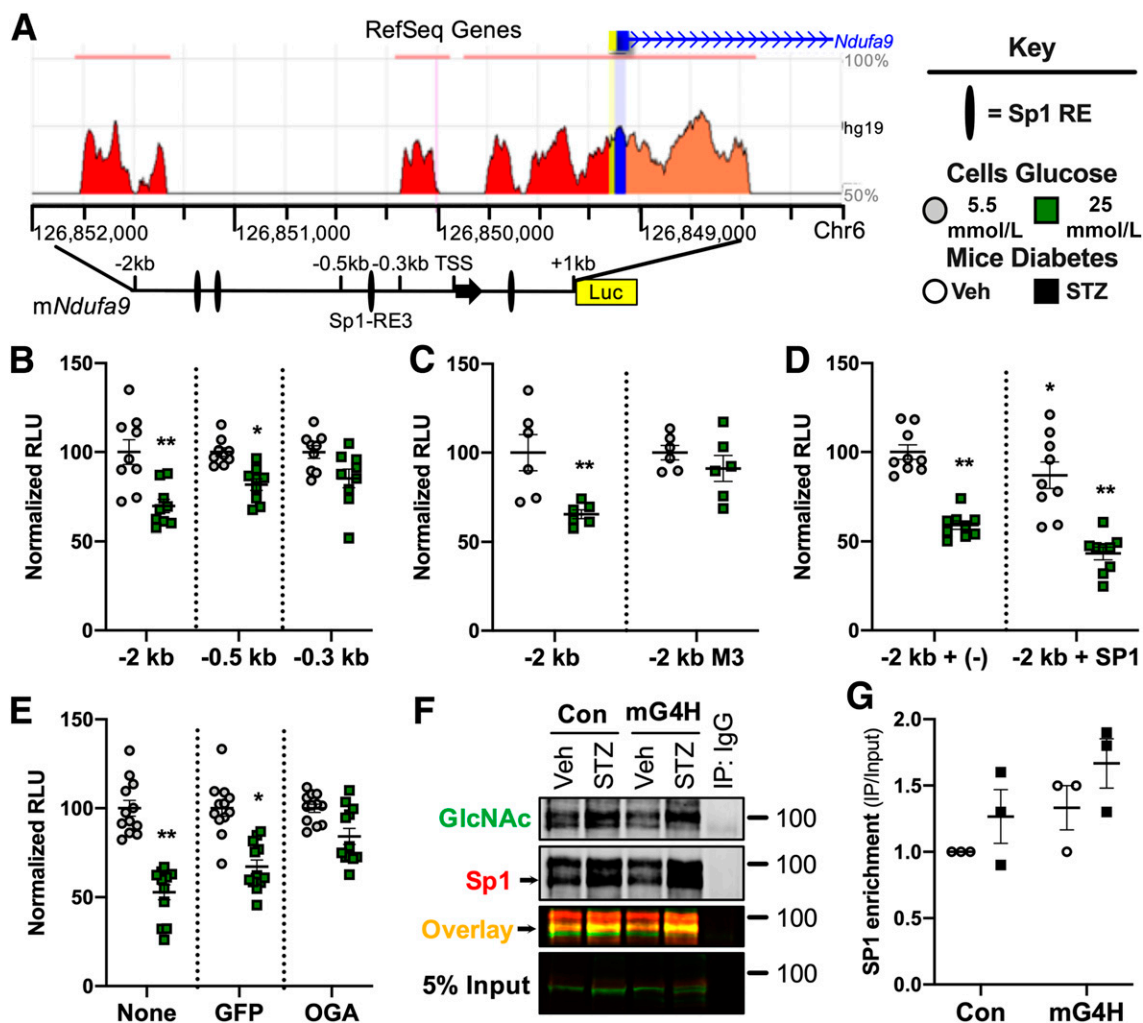
context of diabetes may ameliorate the pathophysiology of diabetic cardiomyopathy. Prior studies revealed that overexpression of GLUT4 in muscle, heart, and adipose tissue from birth prevented the characteristic cardiac metabolic changes and contractile dysfunction in mouse models of diabetes (12,41,42). We more recently reported that loss of cardiac GLUT4 increases susceptibility to heart failure in response to either physiologic or pathologic stress (15). Together, these studies support the hypothesis that maintaining GLUT4-mediated myocardial glucose utilization may be required to prevent diabetes-related ventricular dysfunction and to maintain function following a hemodynamic stress. However, our results indicate that GLUT4 restoration in the context of diabetes not only is unable to rescue either metabolic or contractile dysfunction but also may accelerate the development of diabetic cardiomyopathy. These observations support the perspective that myocardial insulin resistance, as defined by decreased glucose uptake, may provide cardioprotection by defending against fuel overload (43). The current study provides additional molecular insights into mechanisms linking glucotoxicity with mitochondrial dysfunction. Specifically, we observed that increasing myocardial glucose uptake promoted widespread O-GlcNAcylation of core mitochondrial OXPHOS subunits and metabolic enzymes and a representative transcription factor Sp1, which repressed expression of the CI subunit *Ndufa9* in a glucose- and O-GlcNAc-dependent manner. Moreover, increased myocardial glucose alone induced a transcriptional program consistent with induction of FA metabolism and repression of ketone body and amino acid metabolism.

Cardiac-specific overexpression of GLUT1 prevents contractile dysfunction and dilation following pressure overload (13), rescues cardiac dysfunction that is secondary to

PPAR $\alpha$  deficiency (44), and ameliorates ischemic injury associated with aging (45). In contrast, increased myocardial glucose uptake enhances cardiac dysfunction in diet-induced obesity (16), which supports the hypothesis that increasing myocardial glucose delivery in the context of nutrient excess becomes maladaptive.

The short-term duration of hyperglycemia in this study was not expected to induce severe ventricular dysfunction. As such, the current study focused on early mechanisms that could contribute to diabetic cardiomyopathy. We therefore focused on potential mechanisms linking increased myocardial glucose utilization and altered myocardial mitochondrial metabolism. Although not exhaustive, two important modes of regulation were explored. The first was evidence of direct regulation of enzymatic function by posttranslational O-GlcNAcylation of mitochondrial proteins. We identified enzymes and mitochondrial proteins in which a link between O-GlcNAcylation and impaired catalytic activity was previously described in diabetes (17,39). The second was transcriptional regulation, in part, through O-GlcNAcylation of transcriptional regulators (46,47). It should be noted that additional mechanisms likely exist, such as an emerging role of metabolic regulation of epigenetics and its impact on gene expression (19).

There has been resurgent interest in the relationship between cardiac ketone body utilization in heart failure (48,49). In fact, we found that one of the most heavily repressed genes by either GLUT4 expression or diabetes was *Bdh1*, a rate-limiting enzyme in ketone body oxidation. Of note, these changes linked to diabetes or glucotoxicity in rodent hearts are opposite to changes described in models of nondiabetic heart failure. Thus, the extent to which altered ketone metabolism might contribute to the



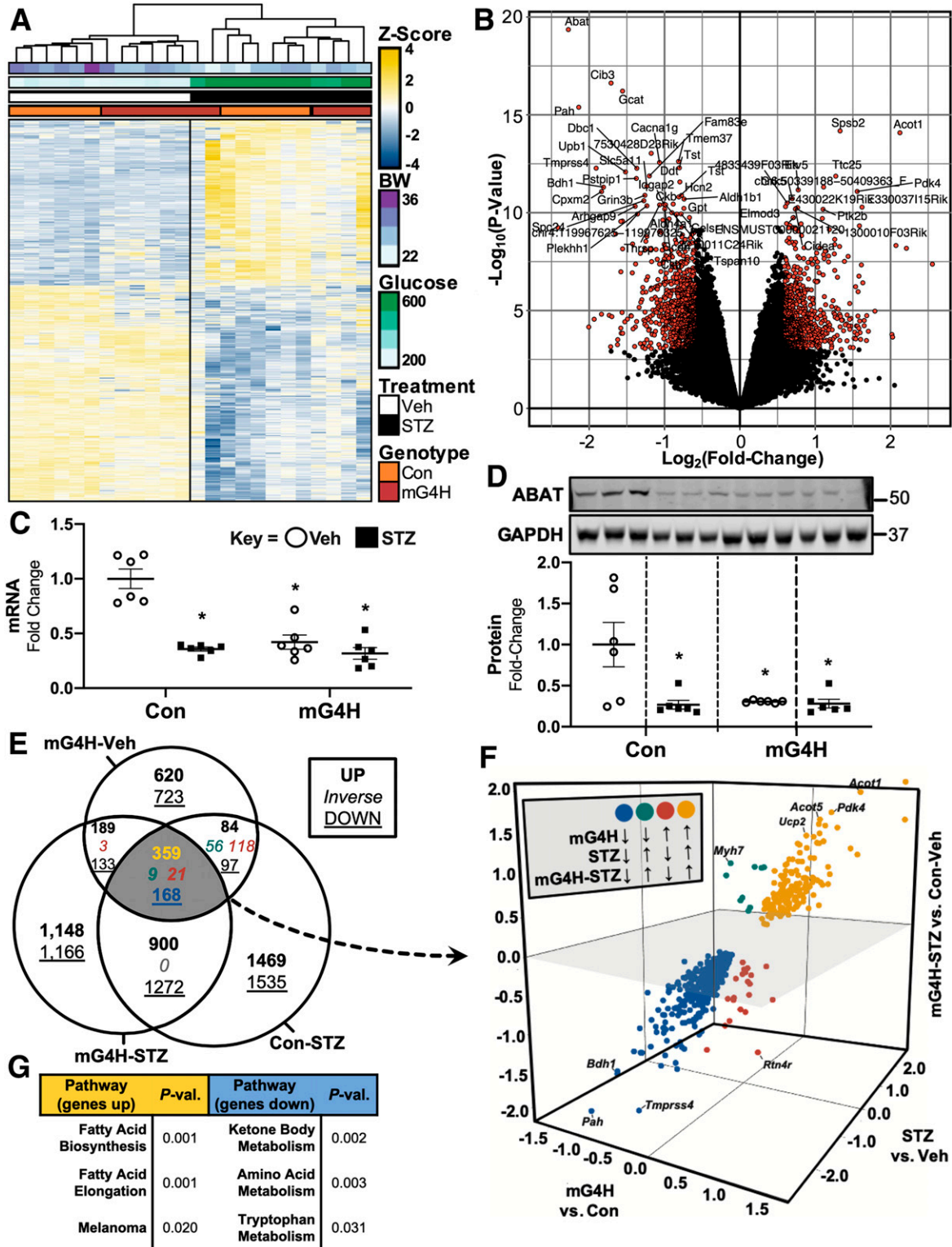
mitochondrial and metabolic phenotypes of mG4H mice remains to be determined.

The robust changes in the regulation of *Abat* are intriguing, and the relationship between this change and glucose-induced metabolic and mitochondrial impairment in response to glucotoxicity in the heart remains to be clarified. *Abat* encodes a metabolic enzyme that catabolizes  $\gamma$ -aminobutyric acid (GABA) for recycling into the tricarboxylic acid cycle as succinate. Prior studies have revealed significantly elevated levels of ABAT protein in failing human dilated cardiomyopathy (1.84-fold) and following trans-aortic constriction (1.95-fold) in the mouse (49). The

impact of altered ABAT on mitochondrial substrate utilization and respiratory function remains to be determined, but its pattern of expression identifies another metabolic pathway that is differentially regulated in diabetic cardiomyopathy relative to heart failure arising from other causes.

Despite its exciting and novel findings, it should be noted that the current study has some limitations that should be carefully considered. First, it has recently been shown that tetracyclines can alter mitochondrial function (50). We observed subtle effects of DOX in our pilot studies and chose to control for them by placing all mice on DOX.





**Figure 7**—Analysis of glucose-dependent vs. diabetes-dependent gene expression patterns in the mouse heart. **A**: Heatmap and hierarchical clustering of DEGs from array-based analysis of left ventricles of mice following 4 weeks of low-dose STZ or Veh treatment. **B**: Volcano plot of mG4H-STZ relative to Con-Veh with the top 50 DEGs labeled. Differential expression of *Abat* in mG4H or Con mice treated with STZ or Veh by qPCR for RNA (**C**) and Western blot for protein levels ( $n = 6$ ) (**D**). **E**: Venn diagram comparing gene expression in mG4H-Veh, Con-STZ, and mG4H-STZ relative to Con-Veh. Genes are listed in Supplementary Table 4. **F**: Coexpressed genes visualized on a three-dimensional scatterplot of fold change for mG4H-Veh vs. Con-Veh, Con-STZ vs. Con-Veh, and mG4H-STZ vs. Con-Veh. **G**: Gene set enrichment analysis of upregulated (yellow) and downregulated (blue) coexpressed DEGs. Genes listed in Supplementary Table 5. Quantitative data are means  $\pm$  SEM. \* $P < 0.05$  vs. Con-Veh. BW, body weight; P-val.,  $P$  value.

As such, our design should primarily reveal differences that can be attributed to increased glucose delivery in a background of DOX exposure. Second, this study focused on candidate proteins such as NDUFA9 and ABAT. We demonstrated that *O*-GlcNAcylation not only might directly regulate the function of NDUFA9 but also may regulate its expression via a similar modification of the Sp1 transcription factor. Although the regulation of gene expression by *O*-GlcNAcylation of transcription factors likely targets multiple pathways as suggested by our microarray pathway analysis, we have not definitively demonstrated that this specific mechanism mediates the global changes in gene expression that were observed. It is also unlikely that the modest change observed in NDUFA9 exclusively accounts for the mitochondrial oxidative impairment observed. Although germ line defects in NDUFA9 alone may lead to neonatal lethality (51), mice with cardiomyocyte-restricted loss of C1 (NDUFS4) are viable, exhibit preserved ventricular function in nonstressed hearts (52), and might even be protected from ischemia/reperfusion injury (53). Thus, it is likely that multiple pathways converge to limit mitochondrial oxidative capacity when glucose availability is increased. Furthermore, we chose to inducibly overexpress GLUT4, while prior studies have focused on constitutive GLUT1 and GLUT4 expression. Despite the known roles of insulin in regulating GLUT4 vesicle trafficking, and the notion that GLUT1 mediates basal levels of glucose uptake, it is now accepted that GLUT4 is the major contributor of basal glucose uptake in the beating heart, given its greater abundance and the role of myocardial contraction in regulating GLUT4 translocation to the sarcolemma. Moreover, given that GLUT4 is repressed in diabetic cardiomyopathy, we believe that increasing or maintaining GLUT4 content in the heart represents a reasonable approach for preserving myocardial glucose utilization in the context of diabetes. We used STZ-induced hyperglycemia for the current study. Although it is possible that other forms of diabetes may result in different findings, studies in GLUT1 transgenic mice following high-fat feeding–induced insulin resistance and diabetes suggest that maintaining glucose uptake in the context of increased FA delivery to the heart that occurs in type 1 and type 2 diabetes will be deleterious.

In conclusion, the current study shows that in the presence of hyperglycemia, increasing myocardial glucose utilization accelerates the decline in glucose and FA oxidative capacity in IWH, reduces mitochondrial oxygen consumption in saponin-permeabilized cardiac fibers, and impacts mitochondrial OXPHOS activity. Our data suggest that the reduction in glucose uptake in the hyperglycemic state likely plays a protective role that limits glucotoxicity. Although increasing glucose delivery to the heart increased glucose uptake, storage, and utilization and was tolerated in the absence of metabolic stress, it clearly sensitized the heart to develop impaired mitochondrial function in the context of nutrient excess. Thus, in the presence of uncontrolled hyperglycemia, normalizing or increasing myocardial

glucose utilization is likely to sensitize the heart to glucotoxicity, which will exacerbate mitochondrial dysfunction and worsen diabetic cardiomyopathy.

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**Duality of Interest.** No potential conflicts of interest relevant to this article were reported.

**Author Contributions.** A.R.W. and E.D.A. secured funding, conceived the study, and wrote the manuscript. All authors reviewed and edited the manuscript. A.R.W. performed mouse studies, left ventricular catheterization and analysis, and promoter cloning and analysis, as well as multiple molecular studies (e.g., RNA quantification, protein quantification, metabolite analysis). J.C.S., M.K.B., L.W., and C.A.A. performed mouse studies, collected samples, and performed Western blots. J.C.S. performed and analyzed the fiber studies. C.M.-H. performed qPCR and echocardiography data analysis and reviewed all statistical analysis of the data as well as graphing of the data. M.E.P. performed microarray analysis and qPCR. O.K. performed and analyzed the respiratory complex activities. H.S. performed and analyzed the two-dimensional PAGE. R.O.P. performed 2DG uptake and the quantification of the mitochondria as appears in the electron micrographs. M.K.B. performed immunoprecipitation studies. J.T. performed IWH perfusion and analysis. A.C.-F. performed IHC. C.D.O., W.E.B., L.J.D., and S.E.L. performed and assisted in analysis of the echocardiography data. W.H.D. provided critical reagents for the *O*-GlcNAc studies. A.R.W. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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