

TM-25659-Induced Activation of FGF21 Level Decreases Insulin Resistance and Inflammation in Skeletal Muscle via GCN2 Pathways

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The TAZ activator 2-butyl-5-methyl-6-(pyridine-3-yl)-3-[2'-(1H-tetrazole-5-yl)-biphenyl-4-ylmethyl]-3H-imidazo[4,5-b]pyridine (TM-25659) inhibits adipocyte differentiation by interacting with peroxisome proliferator-activated receptor gamma. TM-25659 was previously shown to decrease weight gain in a high fat (HF) diet-induced obesity (DIO) mouse model. However, the fundamental mechanisms underlying the effects of TM-25659 remain unknown. Therefore, we investigated the effects of TM-25659 on skeletal muscle functions in C2 myotubes and C57BL/6J mice. We studied the molecular mechanisms underlying the contribution of TM-25659 to palmitate (PA)-induced insulin resistance in C2 myotubes. TM-25659 improved PA-induced insulin resistance and inflammation in C2 myotubes. In addition, TM-25659 increased FGF21 mRNA expression, protein levels, and FGF21 secretion in C2 myotubes via activation of GCN2 pathways (GCN2-phospho-eIF2 α -ATF4 and FGF21). This beneficial effect of TM-25659 was diminished by FGF21 siRNA. C57BL/6J mice were fed a HF diet for 30 weeks. The HF-diet group was randomly divided into two groups for the next 14 days: the HF-diet and HF-diet + TM-25659 groups. The HF diet + TM-25659-treated mice showed improvements in their fasting blood glucose levels, insulin sensitivity, insulin-stimulated Akt phosphorylation, and inflammation, but neither body weight nor food intake was affected. The HF diet + TM-25659-treated mice also exhibited increased expression of both FGF21 mRNA and protein. These data indicate that

TM-25659 may be beneficial for treating insulin resistance by inducing FGF21 in models of PA-induced insulin resistance and HF diet-induced insulin resistance.

INTRODUCTION

Transcriptional co-activator with PDZ-binding motif (TAZ) functions as a co-activator, and its activity is regulated via interactions with 14-3-3 protein and PDZ domain proteins (Kanai et al., 2000). Increased TAZ activity in the nucleus suppresses adipocyte development and increases osteoblast differentiation by interacting with peroxisome proliferator-activated receptor gamma (PPAR- γ) and runt-related transcription factor 2, respectively (Hong et al., 2005). The TAZ activator 2-butyl-5-methyl-6-(pyridine-3-yl)-3-[2'-(1H-tetrazole-5-yl)-biphenyl-4-ylmethyl]-3H-imidazo[4,5-b]pyridine (TM-25659) inhibits adipocyte development by interacting with PPAR- γ and decreases weight gain in obese mouse models (Jang et al., 2012). However, the fundamental mechanisms underlying the effects of TM-25659 are unknown.

Fibroblast growth factor 21 (FGF21) is a member of the FGF family with multiple metabolic functions (Kharitonov et al., 2005). FGF21 functions as an endocrine hormone with anti-diabetic and anti-obesity effects and is produced in peripheral tissues (e.g., the liver, white and brown adipose tissues, skeletal muscle, and the pancreas) (Fisher et al., 2010). Specifically, the administration of recombinant FGF21 or overexpression of FGF21 ameliorates obesity, insulin sensitivity, inflammation, and glucose clearance in obese or diabetic animals (Kharitonov et al., 2005; Wang et al., 2015; Xu et al., 2009). In contrast, FGF21 deficiency leads to body weight gain, impaired glucose tolerance, elevated blood insulin, and fatty liver development (Badman et al., 2007; Inagaki et al., 2007).

General control nonderepressible 2 (GCN2) is a serine/threonine protein kinase used to detect amino acid deficiency. Decreased consumption of dietary protein leads to a reduction in the amount of amino acids delivered to the liver, activation of GCN2, and an eventual increase in eIF2- α phosphorylation and activation of ATF4 (Anthony et al., 2004; Laeger et al., 2014). ATF4 binds amino acid response elements (AAREs) in the FGF21 promoter, which leads to an increase of FGF21 in the liver and blood circulation. FGF21 increases energy

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expenditure (EE) and decreases body weight gain in mice (Ge et al., 2012; Laeger et al., 2014). However, GCN2-knockout mice fail to exhibit increased eIF2 α phosphorylation during both dietary and pharmacological amino acid deprivation, ultimately resulting in hepatic steatosis and liver injury (Anthony et al., 2004; Guo and Cavener, 2007; Wilson et al., 2013).

We found that TM-25659 improved insulin signaling and inflammation in PA-induced insulin-resistant skeletal muscle cells. TM-25659 also increased FGF21 mRNA, protein, and secretion levels in association with increased GCN2 in C2 myotubes. The TM-25659-increased FGF21 protein levels were reduced by GCN2 siRNA. Additionally, administration of TM-25659 lowered the fasting glucose levels, improved insulin resistance, ameliorated inflammation, and increased FGF21 protein levels in our HF diet-induced obesity (DIO) mouse model.

In this study, we evaluated the reciprocal relationship between TM-25659 and FGF21 in skeletal muscle cells and mice, and examined the hypothesis that FGF21 mediates the metabolic benefits of a TM-25659 by comparing the effects of TM-25659-treated skeletal muscle cells subjected to the small interfering RNA (siRNA)-mediated knockdown of FGF21 (FGF21 siRNA) using PA-induced insulin-resistant skeletal muscle cells as controls.

MATERIALS AND METHODS

Materials

TM-25659 was obtained from the Korea Research Institute of Chemical Technology (Korea). Bovine serum albumin (BSA), insulin, PA, and methyl cellulose were purchased from Sigma-Aldrich (USA). Anti-AKT, anti-phospho-AKT (ser473), anti-eIF2 α , anti-phospho-eIF2 α (ser51), anti-PERK, anti-phospho-PERK (Thr980), and anti-ATF4 antibodies were acquired from Cell Signaling Technology (USA). Anti-GCN2, anti-phospho-GCN2 (Thr898) antibodies were acquired from Biorbyt (UK). Anti-FGF21 antibodies were purchased from Abcam (USA). Anti-actin antibody was obtained from Santa Cruz Biotechnology (USA). Culture media, culture supplements, horse serum, and fetal bovine serum (FBS) were purchased from Gibco-BRL (USA). TM-25659 was dissolved in 0.5% methyl cellulose in water for *in vivo* administration.

Preparation of PA

PA/BSA conjugates were prepared by soaping PA with sodium hydroxide (NaOH) and mixing with BSA. Briefly, 20 mM PA in 10 mM NaOH was incubated at 70°C for 30 min, and the fatty acid soaps were combined with 5% fatty acid-free BSA in phosphate-buffered saline (PBS) at a 1:3 volume ratio. The combined fatty acids consisted of 5 mM PA and 3.75% BSA. The PA/BSA conjugates were diluted in 5% horse serum medium (~0.4% BSA) and administered to cultured cells at a final concentration of 500 μ M PA. The molar ratio of PA to BSA was ~3.5:1, and the concentration of BSA in the 500 μ M PA medium was ~0.7%.

Cell culture

C2 mouse skeletal myoblasts were obtained from Dr. Hey-Sun Kim (Ajou University, Korea) and grown in high-glucose (4.5 g/L glucose) Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% FBS and antibiotics (10 μ g/ml streptomycin and 100 IU/ml penicillin) at 37°C in a humidified atmosphere of 95% air and 5% CO₂. After washing the 70-80% confluent myoblasts with PBS, differentiation to myotubes was initiated by transferring the cells to low-serum differentiation medium [DMEM with low glucose (1 g/L) supplemented with 5% horse

serum]. After a 3-day incubation, differentiation was complete as determined by morphological changes and creatine kinase expression.

Immunoblot analysis

Cells and mouse soleus muscles were suspended in RIPA buffer [150 mM sodium chloride (NaCl), 1% NP-40, 0.5% deoxycholate, 0.1% sodium dodecyl sulfate (SDS), 50 mM Tris-HCl, pH 7.5, and protease inhibitor cocktail (Roche Applied Science, Germany)] and incubated on ice for 20 min. Whole proteins were extracted by differential centrifugation (13,000 \times g, 10 min), and the protein concentrations in the lysates were determined using a protein assay kit (Bio-Rad, USA). An equal volume of 2 \times SDS sample buffer (125 mM Tris-HCl, pH 6.8, 4% SDS, 4% 2-mercaptoethanol, and 20% glycerol) was added to the cell lysates, and equivalent amounts of protein (20 μ g) were loaded onto 8-12% polyacrylamide gels, electrophoresed, and transferred electrophoretically to polyvinylidene fluoride membranes (Millipore, USA). After blocking the membranes with 5% skim milk for 30 min, the target antigens were reacted with primary antibodies, followed by secondary antibodies (horseradish peroxidase-conjugated anti-goat IgG or anti-rabbit IgG). Immuno-reactive bands were visualized by enhanced chemiluminescence (Amersham Pharmacia Biotech, USA).

RNA isolation and quantitative real-time polymerase chain reaction (PCR)

Total RNA was isolated from cell or muscle tissues and prepared using RNAiso Plus reagent (TaKaRa Bio Inc., Japan) according to the manufacturer's instructions. Briefly, C2 myotube cDNA was synthesized with avian myeloblastosis virus reverse transcriptase using random 9-mers. The cDNA was amplified by PCR with primer sets for FGF21 [CCC CAA ACC AGG ATG CAA CA (forward, F) and ATG CTG CTT GGT CTT GGG GC (reverse, R)], tumor necrosis factor (TNF)- α [GGC ACT CCC CCA AAA GAT GG (F) and CCC TGC CAC AAG CAG GAA TG (R)], interleukin (IL)-6 [CCA TCC AGT TGC CTT CTT GGG (F) and GCC GTG GTT CTC ACC AGC AT (R)], IL-1 β [TCT CGC AGC AGC ACA TCA ACA (F) and CCT GGA AGG TCC ACG GGA AA (R)], monocyte chemoattractant protein (MCP)-1 [CAG CCA GAT GCA GTT AAC GC (F) and GCC TAC TCA TTG GGA TCA TCT (R)], and RPL32 [AAG CGA AAC TGG CGG AAA CC (F) and CCC ATA ACC GAT GTT GGG CA (R)]. Quantitative real-time PCR was performed using SYBR Green (TaKaRa Bio Inc.) with a TaKaRa TP-815 instrument. All expression values were normalized to RPL32 mRNA levels.

siRNAs

Twenty-one nucleotide siRNA duplexes were designed and created by Bioneer (Korea). The siRNA sequences were as follows: green fluorescent protein, 5'-GUU CAG CGU GUC CGG CGA GTT-3', FGF21, 5'-CUG AUG GAA UGG AUG AGA U-3', and GCN2, 5'-GUG GAA GCU AAA UUC UGU A-3'. C2 myotubes were transfected with siRNA oligonucleotides using a pipette-type electroporator (Microporator-Mini; Digital Biotechnology, Korea) according to the manufacturer's instructions. Briefly, 1 \times 10⁶ C2 myotubes were transfected with 60 nM siRNA (1 μ g in 10 μ l of R buffer) by microporation at a pulse voltage of 1005 V, pulse width of 35 ms, and pulse number of 2. After transfection, the C2 myotubes were differentiated by switching the medium to differentiation medium for 3 days. Following differentiation, the cells were seeded in 12-well plates at a concentration of 1 \times 10⁵ cells/well and treated with or without TM-25659 for 16 h.

Uptake of 2-N-(7-nitrobenz-2-oxa-1,3-diazol-4-yl) amino-2-deoxyglucose (2-NBDG)

C2 myotubes were pretreated with PA (500 μ M) and TM-25659 (25 or 50 μ M) for 16 h. Next, the cells were starved for 4 h and then preincubated in Krebs-Ringer bicarbonate buffer (pH 7.4) containing 2% BSA at 37°C for 30 min. They were then treated with 500 μ M 2-NBDG (cat. no. N13195; Invitrogen, USA) with or without 100 nM insulin at 37°C for 2 h. The cells were washed with ice-cold PBS three times and lysed with cell lysis buffer. The lysates were centrifuged at 13,000 rpm and 4°C for 20 min. The supernatants were measured for fluorescence (excitation 475 nm, emission 550 nm) using a SpectraMax Gemini EM microplate reader (Molecular Devices, USA). Protein concentrations were determined by the Bradford assay.

Animal experiments

Six-week-old male C57BL/6J mice were purchased from Japan SLC Inc. (Japan). All animal experiments were approved by the Animal Ethics Committee of the Laboratory Animal Research Center, Ajou University Medical Center (Korea). The mice were housed in a temperature-controlled room (22 \pm 2°C) with a 12/12-h light/dark cycle and fed *ad libitum*. Eight-week-old male C57BL/6J mice were fed a HF diet (n = 12) for 30 weeks. The mice were provided a HF diet with 60% kcal from fat (Research Diets, USA) *ad libitum*. The HF diet

group was randomly assigned during the additional 14 days to two groups: HF and HF + 50 mg/kg TM-25659 (TM). The mice were orally administered either vehicle or TM-25659 (50 mg/kg) every other day for 14 days, and their body weight was measured every other day.

Insulin tolerance tests

The intraperitoneal insulin tolerance test (ITT) was performed by intraperitoneally injecting regular human insulin (0.5 U/kg) after a 6-h fast. Blood samples were taken at different time points (0, 30, 60, and 120 min after insulin loading) from a tail vein. Plasma glucose was measured using Accu-chek (Korea Roche Diagnostics, Korea).

FGF21 secretion

The concentration of FGF21 in the culture medium of C2 myotubes was determined using a mouse/rat FGF21 enzyme-linked immunosorbent assay (ELISA) (R&D Systems, USA). The concentration of albumin in the culture medium was determined using a mouse albumin ELISA kit (BioTek, USA).

Statistical analysis

Data are presented as the mean \pm standard error of at least three independent experiments. Statistical differences between the groups were determined using Student's *t*-test and Fisher's exact test. *P*-values < 0.05 were considered significant.

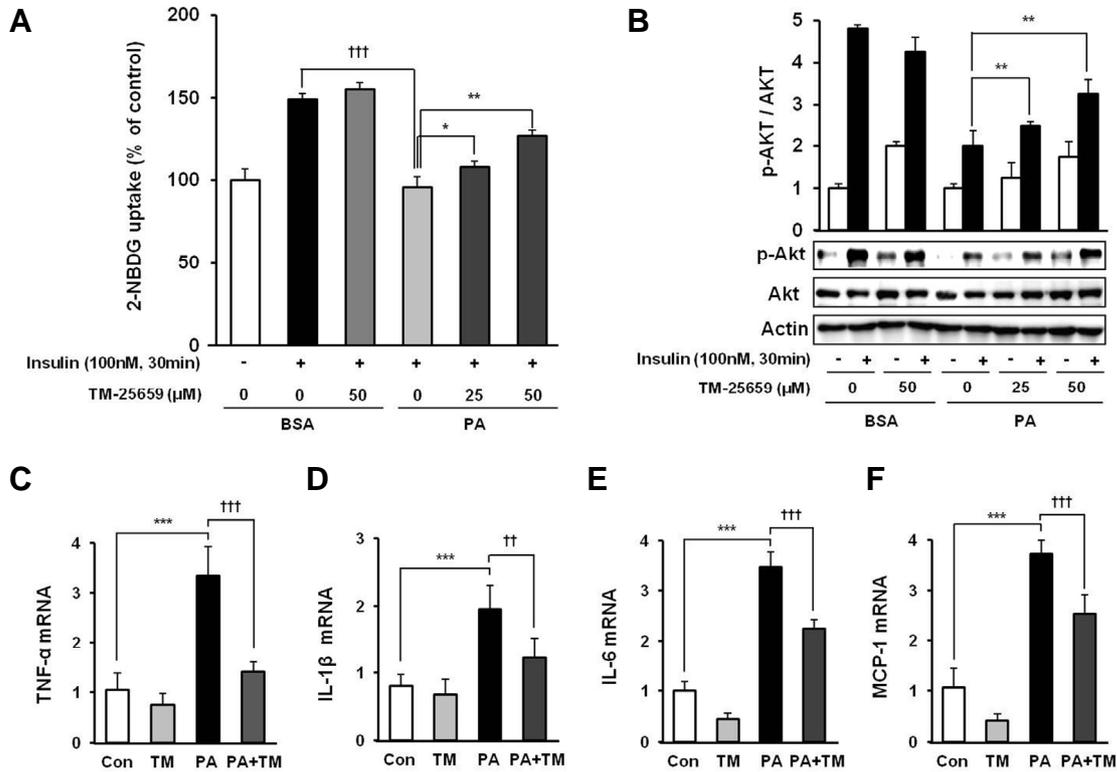


Fig. 1. Beneficial effects of TM-25659 on insulin resistance and inflammation caused by palmitate (PA) in skeletal muscle cells. Skeletal muscle cells were harvested 30 min after insulin (100 nM) treatment of PA-, TM-25659-, or vehicle-treated cells for 16 h. (A) TM-25659 prevented PA-induced impaired glucose uptake by 2-NBDG in C2 myotubes. (B) Representative immunoblots showing phospho (Ser 473) and total AKT in skeletal muscle cells of the control (Con), TAZ modulator (TM), PA, and PA + TM groups, respectively. (C-F) TNF- α , IL-1 β , IL-6, and MCP-1 mRNA expression as quantified by real-time PCR. [Con, BSA; TM, 50 μ M TM-25659; PA, 500 μ M PA for 16 h]. The data are given as the mean \pm standard error. **P* < 0.05, ***P* < 0.01, ****P* < 0.001, ††*P* < 0.01, †††*P* < 0.001.

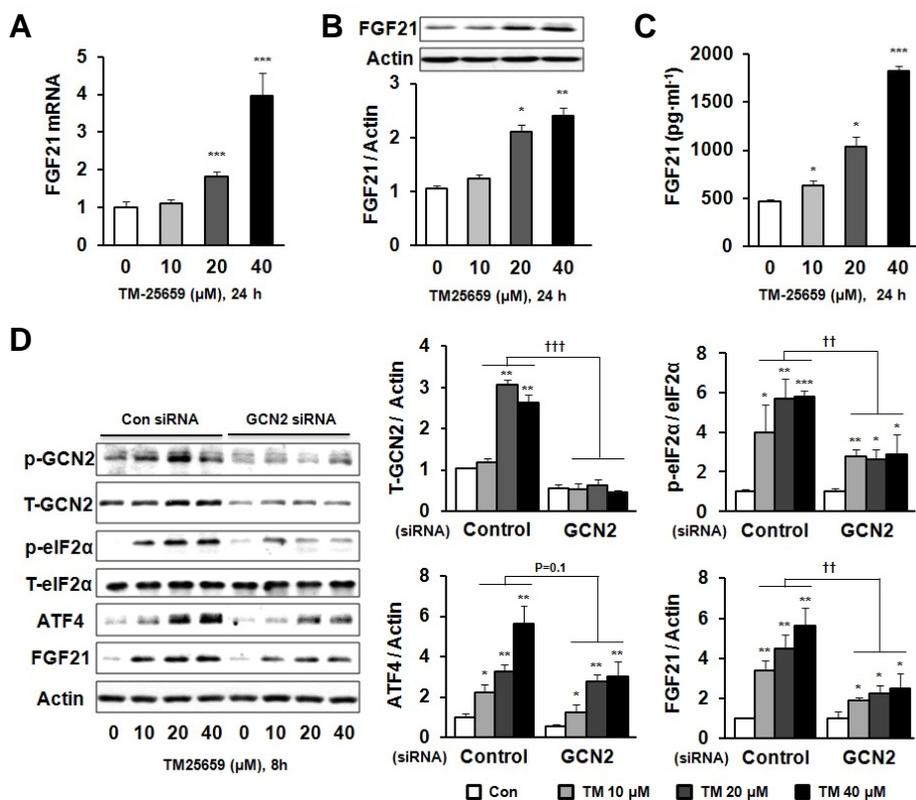


Fig. 2. Effects of TM-25659 on FGF21 expression in skeletal muscle cells. C2 myoblasts were differentiated into C2 myotubes and treated with 10, 20, or 40 μM of TM-25659 or vehicle. (A) FGF21 mRNA expression levels as quantified by real-time PCR. (B) FGF21 protein levels as determined by immunoblotting. (C) Secretion of FGF21 as quantified by an enzyme-linked immunosorbent assay (ELISA). (D) GCN2 pathway in TM-25659-treated skeletal muscle cells with or without GCN2 siRNA. The data are given as the mean ± standard error. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, †† $P < 0.01$, ††† $P < 0.001$.

RESULTS

TM-25659 restored PA-induced insulin resistance and inflammation in skeletal muscle cells

In previous experiments, we found 500 μM PA decreased insulin-stimulated glucose uptake in C2 myotubes (Jung et al., 2011). In this study, we investigated the effects of TM-25659 on PA-induced insulin resistance in C2 myotubes. TM-25659 prevented the decrease in insulin-stimulated glucose uptake in C2 myotubes (Fig. 1A). Additionally, we investigated the effects of TM-25659 on the reduction of insulin-stimulated Akt phosphorylation by PA in C2 myotubes. TM-25659 significantly inhibited the PA-induced reduction in insulin-stimulated Akt phosphorylation (Fig. 1B). These data suggest that TM-25659 restored the PA-induced decrease in insulin sensitivity.

PA increased pro-inflammatory cytokine gene expression in C2 myotubes, while TM-25659 markedly decreased PA-induced expression of TNF-α, IL-1β, IL-6, and MCP-1 in C2 myotubes (Figs. 1C-1F). Additionally, the PA-induced IL-1β maturation levels decreased in C2 myotubes in the PA + TM-25659 (TM) group (data not shown). These data demonstrate that TM-25659 prevented the PA-induced expression of pro-inflammatory cytokines in C2 myotubes.

TM-25659 increased FGF21 levels by GCN2 pathways in skeletal muscle cells

We found that TM-25659 increased FGF21 mRNA, protein, and secretion levels in C2 myotubes. Importantly, TM-25659 significantly increased FGF21 mRNA expression in C2 myotubes in a dose-dependent manner (Fig. 2A). We also observed increased FGF21 protein levels in C2 myotubes treated with TM-25659 (Fig. 2B). Moreover, TM-25659 increased FGF21 secretion in C2 myotubes (Fig. 2C).

TM-25659-induced increased FGF21 protein levels through GCN2 pathways (GCN2-phospho-eIF2α-ATF4 and FGF21) (Fig. 2D). Therefore, we next investigated whether TM-25659 increased GCN2 pathways in C2 myotubes that were and were not treated with GCN2 siRNA. We found that TM-25659 induced increased GCN2 pathways, but this induction was diminished by GCN2 siRNA (Fig. 2D). We also investigated the involvement of other pathways i.e., PERK and PKR, in TM-25659-induced FGF21 expression. We found that TM-25659 induced the induction of PERK and PKR, while FGF21 induction was not reduced by PERK or PKR siRNA treatment. (Supplementary Fig. 1). These data indicate TM-25659-induced increases in FGF21 levels through GCN2 pathways in C2 myotubes.

Effects of TM-25659-reduced insulin resistance and inflammation in skeletal muscle cells treated with or without FGF21 siRNA

We previously showed that TM-25659 markedly increased FGF21 mRNA and protein expression in TM-25659-treated skeletal muscle cells (Figs. 2A and 2B). We investigated the effects of TM-25659-reduced insulin resistance and inflammation in C2 myotubes with or without FGF21 siRNA treatment. FGF21 mRNA and protein expression decreased following FGF21 siRNA treatment (Figs. 3A and 3B). TM-25659 prevented the PA-induced decrease in insulin-stimulated Akt phosphorylation, and the beneficial effects of TM-25659 decreased following FGF21 siRNA treatment (Fig. 3C). We next investigated whether TM-25659 prevented the PA-increased levels of pro-inflammatory cytokines in C2 myotubes treated with or without FGF21 siRNA. We found that the beneficial effects of TM-25659 prevented PA-increased pro-inflammatory cytokine expression (MCP-1, TNF-α, IL-1β, and IL-6), but the decrease in induction was lowered by FGF21 siRNA (Fig. 3D). Consistent

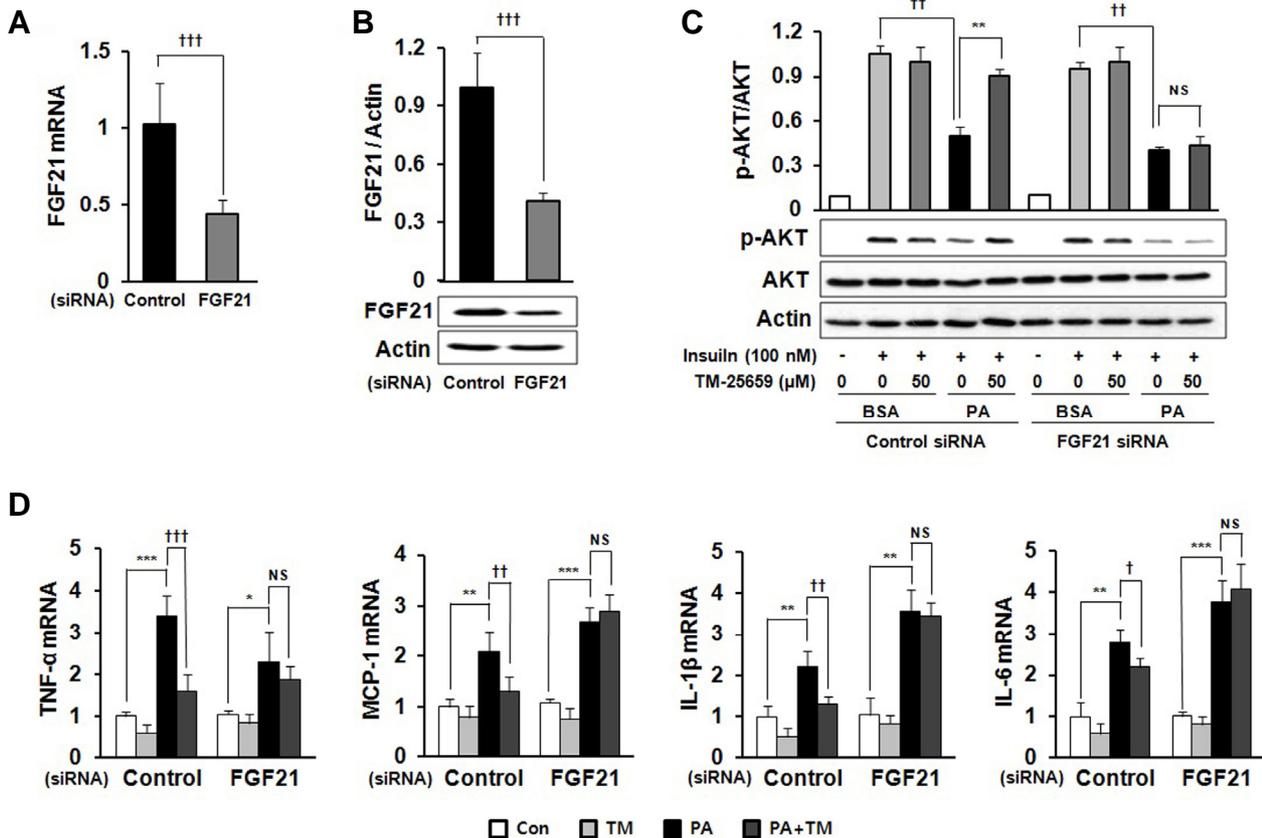


Fig. 3. TM-25659 inhibited insulin resistance and inflammation in skeletal muscle cells in an FGF21-dependent manner. (A) FGF21 mRNA expression with or without FGF21 siRNA as quantified by real-time PCR. (B) Representative immunoblots showing FGF21 in skeletal muscle cells with or without FGF21 siRNA. (C) Representative immunoblots showing phospho (Ser 473) and total AKT in skeletal muscle cells of the control (BSA) and palmitate (PA) groups with or without FGF21. (D) TNF- α , IL-1 β , IL-6, and MCP-1 mRNA expression with or without FGF21 siRNA as quantified by real-time PCR (control (Con), TM-25659 (TM), palmitate (PA), and PA + TM). The data are given as the mean \pm standard error. * P < 0.05, ** P < 0.01, *** P < 0.001, † P < 0.05, †† P < 0.01, ††† P < 0.001.

with our observations in C2 myotubes, TM-25659 significantly decreased the PA-induced expression of pro-inflammatory cytokines in C2 myotubes, whereas these effects were abolished by FGF21 siRNA.

TM-25659 regulates insulin sensitivity and FGF21 mRNA and protein expression in DIO mice

The mice were fed a HF diet for 30 weeks to examine the metabolic effects of TM-25659 treatment before any effects occurred on body weight. Mice in the HF-diet group were randomly divided into two groups for the next 14 days: the HF-diet and HF-diet + TM-25659 groups. After 14 days, no significant changes were observed in body weight or food intake (Figs. 4A and 4B).

The glucose level in the HF + TM group decreased from 9.8 ± 0.5 to 7.6 ± 0.4 mg/dl with fasting (Fig. 4C). TM-25659 treatment improved insulin sensitivity after 14 days, as evaluated by a fasting blood glucose and ITT; a significant decrease in the glucose level occurred postinjection (Fig. 4D).

TM-25659 prevented the HF diet-induced decrease in insulin-stimulated Akt phosphorylation in mouse skeletal muscles (Fig. 4E). We next investigated whether TM-25659 prevented the HF diet-induced increases in the pro-inflammatory cytokine levels in mouse skeletal muscles. We found beneficial effects of

TM-25659 in that it decreased the HF diet-increased pro-inflammatory cytokine expression levels (MCP-1, TNF- α , and IL-1) (Fig. 4F). Because FGF21 was markedly expressed in skeletal muscle cells (C2 myotubes), we investigated the effect of TM-25659 on FGF21 expression in mouse skeletal muscle. After TM-25659 treatment, the FGF21 mRNA expression significantly increased by >2.5-fold, and the FGF21 protein level significantly increased by >1.8-fold (Figs. 4G and 4H). Therefore, TM-25659 is able to induce FGF21 expression and insulin sensitivity without changes in body weight.

DISCUSSION

In this study, treating skeletal insulin-resistant muscle cells with the TAZ activator TM-25659 protected against insulin resistance and inflammation. TM-25659 increased the FGF21 mRNA and protein levels in HF diet-induced mouse skeletal muscle and C2 myotubes. Additionally, TM-25659 increased FGF21 mRNA expression and stimulated FGF21 secretion in C2 myotubes. When skeletal muscle cells were treated with FGF21 siRNA, TM-25659 did not protect against PA-induced insulin resistance and inflammation. Additionally, we demonstrated that TM-25659 administration to HF-induced DIO mice improved the fasting glucose level and insulin sensitivity. Thus,

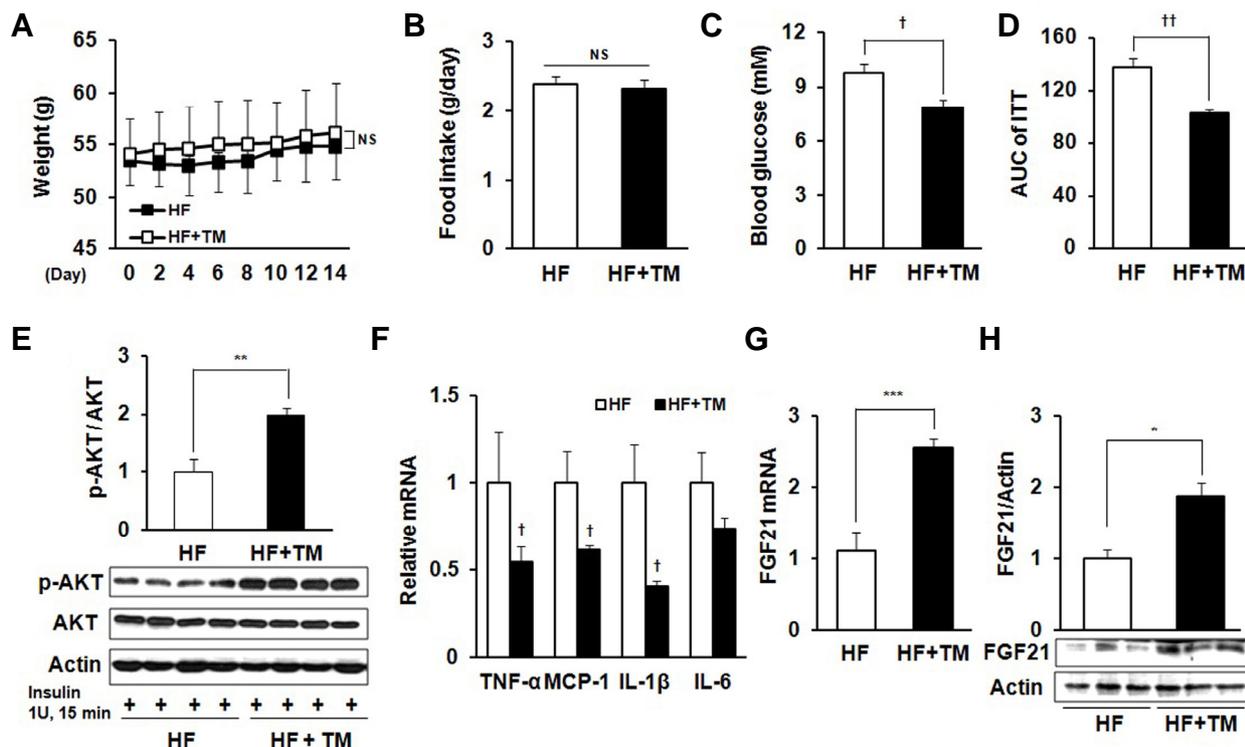


Fig. 4. TM-25659 attenuated fasting glucose and insulin sensitivity in high fat (HF) diet-induced obese mice. C57BL/6J mice were fed a HF diet for 30 weeks. They were then separated into two groups and given either vehicle or TM-25659 (50 mg/kg) orally for 14 days. (A) Body weight was measured every other day. (B) Food intake was similar between the HF and HF + TM groups. (C) Fasting blood glucose levels. (D) Area under the curve of the intraperitoneal insulin tolerance test (ITT) performed 14 days after TM-25659 treatment. (E) Representative immunoblots showing phospho AKT(Ser 473) and total AKT in the skeletal muscle of the HF and HF + TM group mice. (F) TNF- α , IL-1 β , IL-6, and MCP-1 mRNA expression as quantified by real-time PCR. (G) The mRNA expression of FGF21 as quantified by real-time PCR. (H) Protein levels of FGF21 by immunoblotting (HF, HF + TM-25659). The data are given as the mean \pm standard error. * P < 0.05, ** P < 0.01, *** P < 0.001, † P < 0.05, †† P < 0.01, vs. vehicle in the HF group mice; n = 10/group.

the beneficial effects of TM-25659 must be associated with increased FGF21 in skeletal muscle.

Improved body weight following treatment with TM-25659 has been reported (Jang et al., 2012), but neither muscle-specific insulin sensitivity nor the relationship between FGF21 and TM-25659 has been examined. FGF21 is a metabolic hormone produced mainly by the liver, but it is also expressed in muscle and adipose tissue where it regulates glucose and lipid homeostasis via pleiotropic actions (Woo et al., 2013). Here, we first demonstrated that TM-25659 increased FGF21 RNA and protein levels directly as dose- and time-dependent variables, and that FGF21 was secreted at significantly increased levels into mouse skeletal muscle cell medium. To test these effects in mice, TM-25659 was injected into C57BL/6J mice every other day for 14 days. The FGF21 levels were dramatically augmented in biopsied muscles. Treatment of TM-25659 in obese mice led to improved insulin resistance and inflammation by FGF21 induction. For long-term treatment of TM-25659 (more than 8 weeks), it stimulated weight loss (data not shown). Kim et al. (2013) demonstrated that FGF21 was induced through the translocation of activating transcription factor 4 (ATF4) from the cytoplasm to the nucleus in autophagy-deficient skeletal muscle. In addition, FGF21 was found to be induced in the liver under leucine or methionine deprivation through the GCN2-phospho-eIF2 α and ATF4 pathways (De

Sousa-Coelho et al., 2013; Lees et al., 2014). TM-25659-induced FGF21 reacts in the same way in the GCN2-phospho-eIF2 α and ATF4 pathways.

TM-25659 improved insulin resistance in HF diet-fed mouse muscle. Mounting evidence suggests that macrophage infiltration and accumulation of chronic inflammatory cytokines such as TNF- α , IL-1 β , IL-6, or MCP-1 in adipose tissue is positively correlated with insulin resistance (Huh et al., 2014; Ota, 2013; Park et al., 2015; Xu et al., 2003). Because TM-25659 reduced inflammatory cytokines such as TNF- α , IL-1 β , IL-6, and MCP-1, TM-25659 is protective against insulin resistance in mice fed an HFD. Another cause of reduced insulin resistance was FGF21 expression following TM-25659 treatment of muscle. Xu et al. (2009) reported that an FGF21 infusion given to DIO mice protected against insulin resistance by enhancing EE and lipid oxidation. Furthermore, transgenic mice overexpressing hepatic FGF21 were protected from HF diet-induced insulin resistance (Inagaki et al., 2007; Kharitonov et al., 2005). However, FGF21 knockdown mice had reduced adiponectin levels and aggravated chronic inflammation compared to HF diet-fed mice (Lin et al., 2013). The present study demonstrates a similar phenomenon in muscle. TM-25659 treatment prevented PA-induced insulin resistance and the induction of pro-inflammatory processes in skeletal muscle, but FGF21 siRNA eliminated the beneficial effects of TM-25659. Therefore, TM-25659 caused

the accumulation of FGF21 in skeletal muscle and protected against insulin resistance and chronic inflammation in muscles.

In conclusion, our results suggest that TM-25659 administered to HF diet-induced obese mice prevented not only insulin resistance, but also PA-induced impaired insulin signaling. These beneficial effects were mediated via the induction of FGF21 expression.

Note: Supplementary information is available on the Molecules and Cells website (www.molcells.org).

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