

# The Role of PPAR $\delta$ Agonist GW501516 in Rats with Gestational Diabetes Mellitus

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**Background:** Gestational diabetes mellitus (GDM) is a disorder of glucose metabolism that occurs or is found for the first time during pregnancy. GDM is very harmful and urgently needs drug treatment to improve pregnancy outcome. PPAR $\delta$  is involved in a variety of biological processes related to glycolipid metabolism in the body, suggesting that it may be closely related to insulin resistance and impaired glucose tolerance. The role of PPAR $\delta$  agonist GW501516 in gestational diabetes has not been studied.

**Methods:** Firstly, the rat model of GDM was established. Then, fasting blood-glucose (FGB), fasting insulin (FINS), HOMA-islet resistance index (HOMA-IR) and insulin sensitivity index (ISI) of GDM rats treated with GW501516 were measured on day 3, day 10 and day 17. Glucose tolerance test was performed on the 20th day of gestation to measure glucose tolerance in rats. The expression of PPAR $\delta$  and Angptl8 in islet tissues of rats was detected by Western blot and immunohistochemistry (IHC). Histopathological changes of islet were detected by HE stain; apoptosis rate of islet cells was detected by TUNEL; and expression of apoptosis-related proteins in the cells was detected by Western blot. The biochemical kits were used to detect the expression of lipid metabolism-related factors in blood of GDM rats after the PPAR $\delta$  agonist GW501516 treatment. Finally, the expression of SREBP-1c and GLUT2 in islet tissues was detected by RT-qPCR and IHC.

**Results:** The PPAR $\delta$  agonist GW501516 decreased the expression of FGB, FINS and HOMA-IR in GDM rats, and we found that GW501516 decreased ISI in GDM rats. GW501516 increased glucose tolerance in GDM rats too. In GDM rats, the expression of PPAR $\delta$  in islet decreased and the expression of Angptl8 increased, which was reversed by GW501516. In addition, we also found that GW501516 can improve the damaged islet tissue of GDM rats, reduce the apoptosis rate of islet cells and inhibit the expression of lipid metabolism-related factors in the blood. Finally, we found that GW501516 inhibited the expression of SREBP-1c and promoted the expression of GLUT2 in the islet tissue.

**Conclusion:** The PPAR $\delta$  agonist GW501516 could improve the blood glucose level, damaged islet tissue and increase the insulin content in the rats with GDM, possibly by regulating the SREBP-1c/GLUT2 pathway. Our study provided a new basis for clinical treatment of GDM in pregnant women with PPAR $\delta$  agonist GW501516.

**Keywords:** PPAR $\delta$ , GW501516, gestational diabetes mellitus, GDM, SREBP-1c/GLUT2 pathway

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## Introduction

Gestational diabetes mellitus (GDM) refers to the normal glucose metabolism before pregnancy and the phenomenon of the decline of potential glucose tolerance and diabetes that occurs during pregnancy. More than 80% of pregnant women with diabetes are GDM.<sup>1</sup> GDM is very harmful, not only affect the pregnant woman,

which leads to dystocia, retinopathy and an increased risk of type 2 diabetes, but also leads to fetal malformation, stillbirth and neonatal respiratory disease caused by immature fetal lungs.<sup>2</sup> Therefore, diabetes treatment and intervention should be taken in time to control the blood glucose level and improve their pregnancy outcomes.

Peroxisome proliferators-activated receptor (PPAR), as a class of nuclear transcription factors, can be activated by corresponding ligands and is considered to be a member of the nuclear receptor superfamily. PPAR is divided into three subtypes: PPAR $\alpha$ , PPAR  $\delta$  and PPAR $\gamma$ .<sup>3</sup> PPAR  $\delta$  are widely distributed in adipose tissue, muscle tissue and nerve tissue, especially in tissues related to glucose and lipid metabolism. Activation of PPAR  $\delta$  stimulates the oxidation of fatty acids in adipose tissue and skeletal muscle, which improve dyslipidemia in mice and humans.<sup>4</sup> In BXD mice, the expression of PPAR  $\delta$  in muscle was positively correlated with endurance performance and activation of PPAR  $\delta$  can effectively inhibit glucose catabolic metabolism without affecting muscle fiber type or mitochondrial content.<sup>5</sup> This suggests that PPAR  $\delta$  may be closely related to insulin resistance and impaired glucose tolerance. However, the expression and the role of PPAR  $\delta$  in GDM are unknown.

GW501516 is a selective ligand of PPAR  $\delta$ , which can reduce apoptosis of pancreatic beta cells of mice caused by palmitate (PAM) and LPS,<sup>6,7</sup> and dysfunction of pancreatic beta cells induced by elevated free fatty acid (FFA) cycle now was considered as important pathological reasons for type 2 diabetes.<sup>8</sup> However, the role of PPAR  $\delta$  agonist GW501516 in GDM has not been studied.

Angiopoietin-like protein 8 (Angptl8) is secreted protein factor discovered in recent years, which are closely related to sugar metabolism, lipid metabolism and insulin sensitivity.<sup>9,10</sup> Studies have shown that Angptl8 affects plasma lipoprotein levels by regulating glucose homeostasis and insulin resistance.<sup>11</sup> So, Angptl8 can also be used as a marker for the development of GDM.

Sterol regulatory element-binding protein 1c (SREBP-1c) is an important transcription factor regulating lipid synthesis and glucose metabolism. Overexpression of SREBP-1c can lead to increased lipid synthesis and disorder of glucose and lipid metabolism, which plays an important role in the occurrence and development of GDM.<sup>12</sup> Studies have shown that SREBP-1c in liver cells of diabetic rats can mediate the expression of GLUT2 gene stimulated by glucose.<sup>13,14</sup> GLUT2 is mainly expressed in the basolateral membrane of the proximal tubules of the liver and pancreas and plays an important

role in glucose homeostasis in the organism.<sup>15</sup> In addition, Kramer et al<sup>16</sup> reported that GW501516 and GW0742, specific activators of PPAR  $\delta$ , can directly affect the insulin action of skeletal muscles by targeting the regulation of SREBP 1c/GLUT2 pathway. Therefore, it was speculated that whether GW501516 acted through the SREBP 1c/GLUT2 pathway in GDM rats.

Therefore, the purpose of this study was to detect the expression of PPAR  $\delta$  in the pancreas of GDM rats and to investigate the role of its agonist GW501516 in lipid metabolism and insulin resistance in GDM rats. Our study provided a new basis for clinical treatment of GDM in pregnant women with PPAR $\delta$  agonist GW501516.

## Materials and Methods

### Establishment of Animal Model

Fifty healthy female Wistar rats (180 ~ 220 g) were adaptively fed a basic diet for 1 week before modeling. Then, they were given with a high-fat feed (corn starch 24.15%, casein 20%, dextrin 13.2%, sugar 10%, 0.1% cholesterol, 22% lard, 5% cellulose and 3.5% minerals, vitamins, 1% 0.3% l-cystine, choline chloride 0.25%, bile salts, 0.5%) for 8 weeks. The ratio of males to females in the same cage was 1:2, and they were allowed to spend the night together and the female mice whose sperm was observed by microscopic examination were labeled as pregnant mice. Fasting for 12h on the first morning of pregnancy before treatment, a small dose of streptozotocin (STZ) (35mg/kg) was intraperitoneal injection, which was dissolved with citrate-sodium citrate buffer at pH4.4, while the blank control group was only given buffer injection. After the injection for 72h (the 4th day of gestation), fasting blood glucose was measured to be stable above 13.8mmol/L and gestational diabetes mellitus was considered to have been successfully modeled.

### Experimental Grouping and Administration

The animals were divided into normal pregnant female rats (control, n=10), gestational diabetes mellitus (GDM, n=10), GW501516 (1mg/kg, n=10), GW501516 (3mg/kg, n=10), GW501516 (10mg/kg, n=10), and metformin hydrochloride (DMBG, 200mg/kg · d, n=10). Both GW501516 groups and GDM group were feed with high-fat feed. After the successful establishment of the GDM model (the 4th day of gestation), gavage of GW501516 in different doses was

given for 20 days, once a day. The control group and the model group were given normal saline.

## Hematoxylin–Eosin (HE) Stain

Islet tissue was brought into 4% paraformaldehyde solution for several days and subsequently 5 mm longitudinal sections were cut using a rotary microtome (Fa. Leica Biosystems Nussloch GmbH, Germany). These sections were stained with hematoxylin–eosin (HE) according to the manufacturer's instructions.

## Immunohistochemistry (IHC)

The paraffin sections were blocked with 2% milk for 1 h and probed with the primary antibody (1:800) overnight at 4°C. After washing, the sections were incubated for 2 h at room temperature in the appropriate biotinylated secondary antibodies (diluted 1:500; all purchased from Vector Labs). According to the manufacturer's instructions, the slides were incubated with the DAB kit for 10 min. All tissue sections were examined by light microscopy (BX51, Olympus, Japan).

## TUNEL Assay

Paraffin-embedded sections were stained with terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assay (In Situ Cell Death Detection Kit, Roche). In Situ Cell Death Detection Kit with fluorescein-dUTP (Roche) was used to detect TUNEL<sup>+</sup> cells. Islet cells were analyzed using Nikon Eclipse E400 Microscope equipped with a Nikon DS-Ri1 camera using NIS-Elements Microscope Imaging Software or Spot Advance Software. The percentage of apoptotic cells was calculated from randomly selected fields.

## Western Blot Assay

Islet tissue lysates were prepared using tissue protein extraction reagent (T-PER), which were then subjected to centrifugation at 4 °C with a rotation speed of 10,000 rpm for 10 min. The protein concentration was measured using BCA kit (Beyotime). Samples underwent denaturation at 95 °C for 10 min and store overnight at –80°C. The proteins separated on the gel were transferred onto PVDF membranes and after transfer, membrane containing proteins were blocked with 5% BSA for 1 h at room temperature. Then, the blots were washed effectively with TBST for 3 times and incubated with diluted primary antibodies at 4 °C overnight. The next day, blots were washed and treated with respective secondary antibodies for 1 h at room temperature and expression of the proteins of various experimental groups was obtained using enhanced chemiluminescence

reagent (GE Healthcare) and Image J software (version 146; National Institutes of Health, Bethesda) was used to analyze fold-changes of protein levels. Anti-cleaved-caspase3 (1:1000; 9953S), anti-Bcl2 (1:1000; ab32124), anti-Bax (1:1000; ab32503), anti-SREBP-1c (1:1000; ab133125), anti-GLUT2 (1:1000; ab54460), anti-GAPDH (1:1000; ab181602) antibodies were obtained from Abcam.

## ELISA Assay

The levels of high-density lipoprotein (HDL) and low-density lipoprotein cholesterol (LDL) and insulin were measured by ELISA according to the manufacturer's instructions (ThermoFisher, USA). The absorbance was measured using a VersaMax microplate reader (Molecular Devices Corporation, CA, USA).

## Determination of Triglyceride (TG) and Total Cholesterol (TC)

TG and TC were measured in the blood of GDM rats using cholesterol and triglyceride assay kits (Cayman Chemical, Ann Arbor, MI, USA) following instructions from the manufacturer.

## Quantitative Real-Time Polymerase Chain Reaction (RT-qPCR)

Total RNA was extracted from cultured cells using RNAzol RT (Sigma-Aldrich, USA), following the manufacturer's protocol. The RNA concentration and quantification were assessed using a Nanodrop spectrophotometer (Thermo Fisher Scientific, Inc.). After DNase I digestion, QuantiTect Reverse Transcription Kit (QIAGEN China, Shanghai, China) was used to carry out reverse transcriptions to synthesize cDNA, according to the manufacturer's protocols. Subsequently, cDNA was analyzed using QuantiTect SYBR Green PCR Kit (QIAGEN China, Shanghai, China), according to the manufacturer's protocols. Amplification conditions were as follows: 95°C for 10 min, followed by 40 cycles of 95°C for 10 sec and 60°C for 60 sec. Primer sequences used in PCR were obtained from GenScript (Piscataway, NJ, USA). The primer sequences are as follows: SREBP-1c forward: 5'- AGTAGCAGCGGTGGAAGT –3'; SREBP-1c reverse: 5'-CTTCCTGTATGTGCACTGGCGGTCT-3'. GLUT2 forward: 5'- GTCAGAAGACAAGATCACCGGA-3'; GLUT2 reverse: 5'- AGGTGCATTGATCACACCGA-3'. GAPDH forward: 5'-AGCCACATCGCTCAGACAC-3'; GAPDH reverse: 5'-GCCCAATACGACCAAATCC-3'.

GAPDH was included as an internal control and  $2^{-\Delta\Delta C_q}$  method was used for statistical analysis.<sup>17</sup>

## Statistical Analysis

Data are expressed as mean  $\pm$  SD. Statistically significant values were compared using *T*-test, and *P*-values of less than 0.05 were considered statistically significant.

## Results

### GW501516 Decreased Fasting Blood Glucose (FBG), Fasting Insulin (FINS), HOMA-Islet Resistance Index (HOMA-IR) Expression and Increased Insulin Sensitivity Index (ISI) Expression in GDM Rats

FBG and FINS in each group were examined when GDM rats were treated with GW501516 on 3rd, 10th, and 17th day, as shown in Figure 1, we found that compared with the control group, the level of FBG, FINS and HOMA-IR in GDM rats increased and the level of ISI decreased significantly; Compared with the GDM group, the levels of FBG, FINS and HOMA-IR in rats treated with GW501516 decreased in dose dependence, while the levels of ISI increased significantly. No difference in FBG, FINS, HOMA-IR and ISI levels

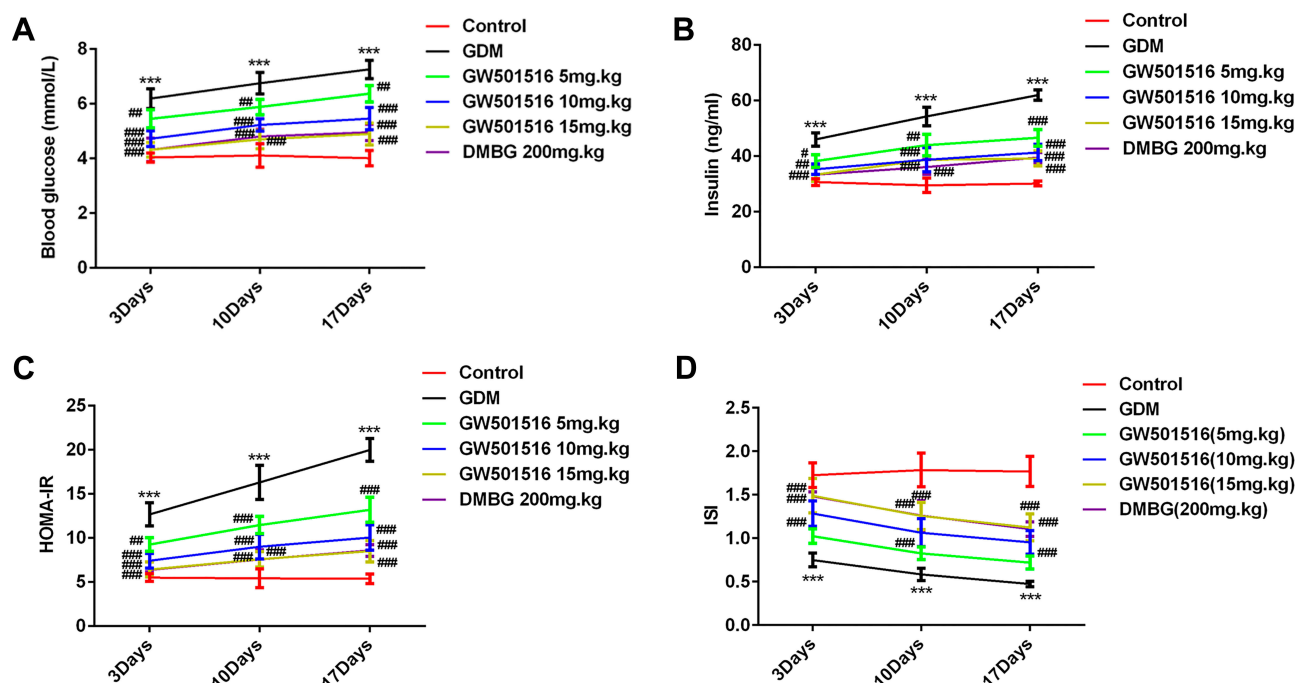
between the GW501516 (15mg/kg) and DMBG was observed.

### GW501516 Increased Glucose Tolerance in GDM Rats

After 21 days of treatment with GW501516, the FBG levels at 0, 15, 30, 60, 90 and 120 min in the GDM group were higher than those in the control group. The FBG levels at all of the above time points were lower in the GW501516 group than in the GDM group (Figure 2A). For the insulin test, rats in the GDM group had higher FINS levels compared with the control group; It was obvious that this was reversed with GW501516 treatment (Figure 2B).

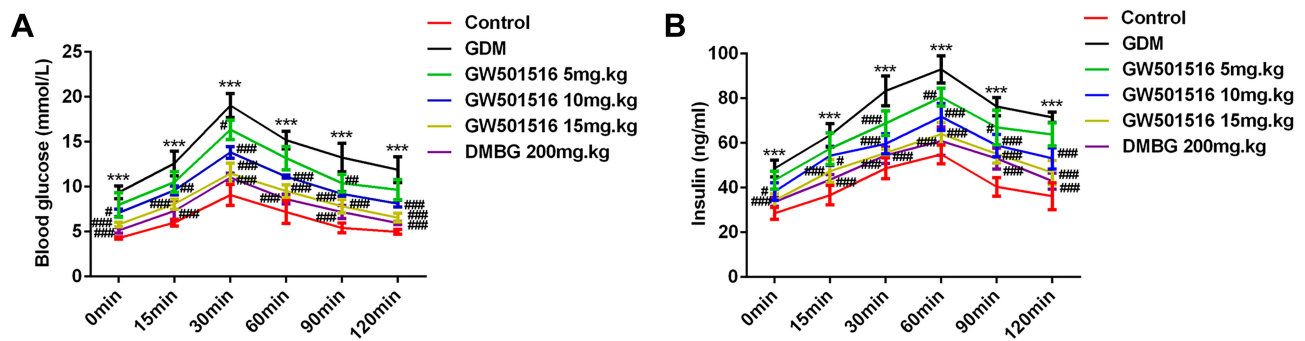
### GW501516 Increased the Expression of PPAR $\delta$ in GDM Rats

We examined the expression of PPAR $\delta$  in the islet tissues of rats and found that compared with the control group, the expression of PPAR $\delta$  in the GDM group was significantly decreased, while GW501516 showed a dose-dependent improvement in the decline of PPAR  $\delta$ , and DMBG also improved the decline (Figure 3A and B). In addition, the expression of angiopoietin-like protein 8 (Angptl 8), which is closely related to carbohydrate metabolism, lipid metabolism

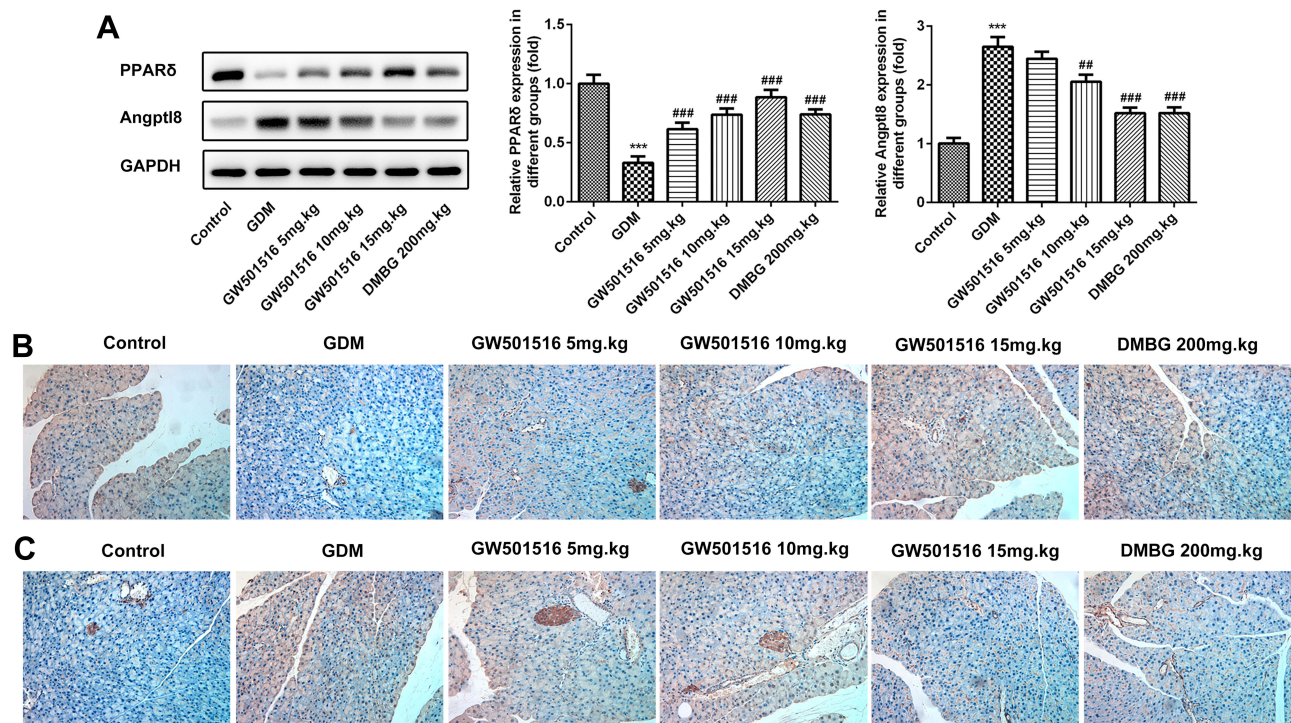


**Figure 1** GW501516 decreased fasting blood glucose (FBG), fasting insulin (FINS), HOMA-islet resistance index (HOMA-IR) expression and increased insulin sensitivity index (ISI) expression in GDM rats. (A) Expression level of FBG in rats. (B) Expression level of FINS in rats. (C) Expression level of HOMA-IR. Calculation method: HOMA-IR = FBG  $\times$  FINS/22.5. (D) Expression level of ISI. Calculation method: ISI = 1/(FBG  $\times$  FINS), where insulin unit was converted as (1 ng/mL = 21.2 mIU/L). n=10, \*\*\**p*<0.001 vs control; #*p*<0.05, ##*p*<0.01, ###*p*<0.001 vs GDM.





**Figure 2** GW501516 increased glucose tolerance in GDM rats. OGTT assay measured the expression levels of FBG (A) and FINS (B) in rats.  $n=10$ , \*\*\* $p<0.001$  vs control; # $p<0.05$ , ## $p<0.01$ , ### $p<0.001$  vs GDM.



**Figure 3** GW501516 increased the expression of PPAR $\delta$  in GDM rats. The expression levels of PPAR  $\delta$  and Angptl8 in the pancreas were detected by Western blot (A) and IHC of PPAR  $\delta$  (B) and Angptl8 (C).  $n=3$ , \*\*\* $p<0.001$  vs control; ## $p<0.01$ , ### $p<0.001$  vs GDM.

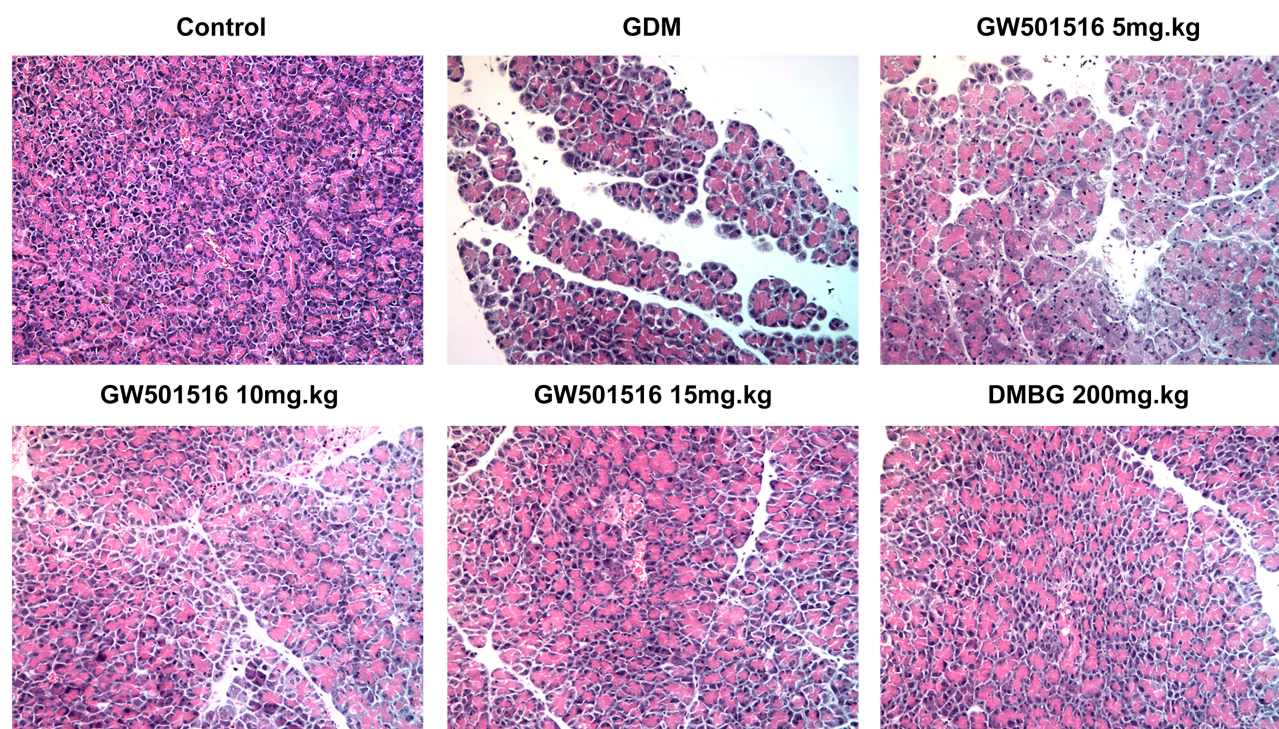
and insulin sensitivity, was also detected in pancreatic tissues. We found that compared with the control group, Angptl8 expression increased significantly in the GDM group, while the expression of Angptl8 decreased in a dose-dependent manner after GW501516 treatment, which was consistent with that of the DMBG group (Figure 3A and C).

### GW501516 Improved the Damaged Islet Tissue and Reduced the Apoptosis Rate of Islet Cells in GDM Rats

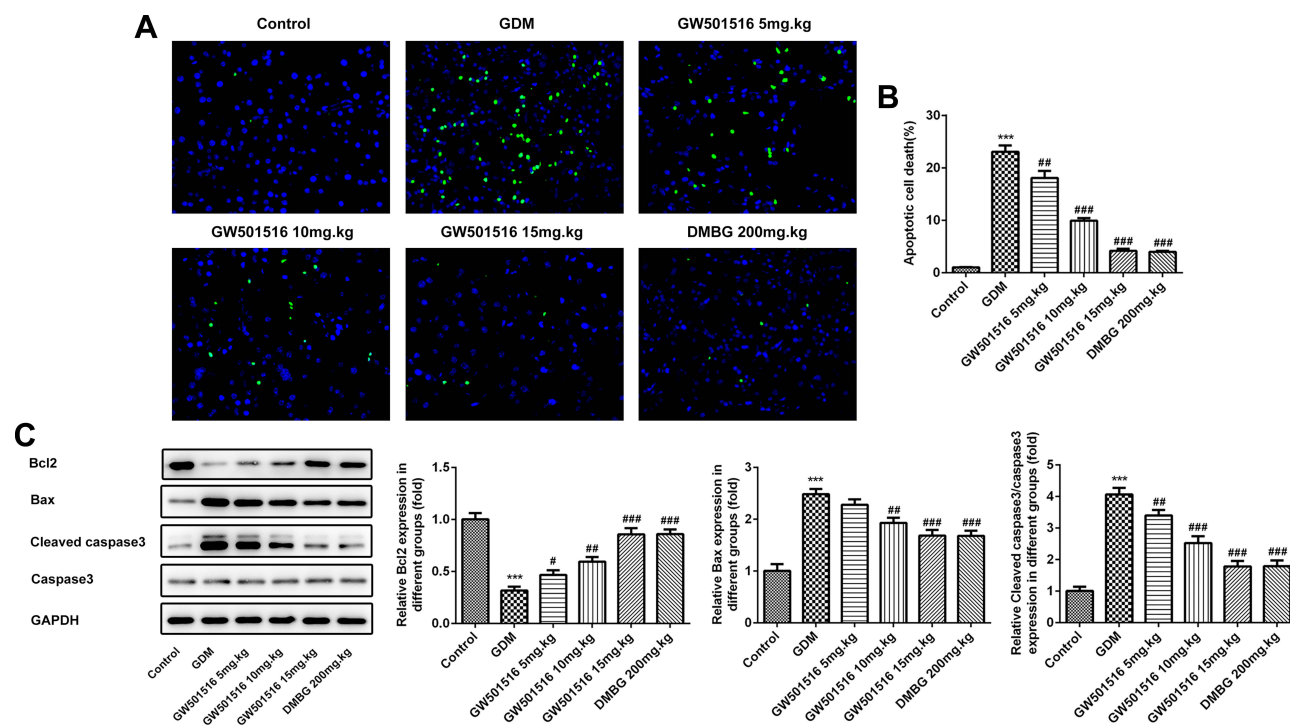
We detected the histopathological changes of islet in rats by HE stain, and we found that the islet tissues of the

GDM group were severely damaged, while the islet tissues of GDM rats were improved after treatment with GW501516 and DMBG (Figure 4). Subsequently, we detected the apoptosis of islet cells. We found that compared with the control group, the apoptosis of islet cells in the GDM group was severe (Figure 5A and B), accompanied by increased expression of Bax and Cleaved caspase3, and decreased expression of Bcl-2 (Figure 5C). Compared with the GDM group, GW501516 and DMBG showed improved apoptosis, decreased expression of Bax and Cleaved caspase3, and increased expression of Bcl-2.





**Figure 4** GW501516 improved the damaged islet tissue in GDM rats. Histopathological changes of rat islet tissue were detected by HE staining.  $n=3$ .



**Figure 5** GW501516 reduced the apoptosis rate of islet cells in GDM rats. **(A)** TUNEL staining detected changes in apoptosis level of islet cells. **(B)** Statistical analysis of apoptosis rate. **(C)** Western blot was used to detect the expression of apoptosis-related proteins.  $n=3$ , \*\*\* $p<0.001$  vs control; # $p<0.05$ , ### $p<0.01$ , #### $p<0.001$  vs GDM.

## GW501516 Inhibited the Expression of Lipid Metabolism-Related Factors in Blood

We tested the expression of lipid metabolism-related factors: low-density lipoprotein cholesterol (LDL), triglyceride (TG), total cholesterol (TC) and high-density lipoprotein (HDL). The results showed that the expressions of LDL, TG and TC in GDM rats were significantly increased, while the expressions of HDL were decreased. Compared with GDM group, after GW501516 and DMBG treatment, the expressions of LDL, TG and TC decreased, while the expressions of HDL increased obviously (Figure 6).

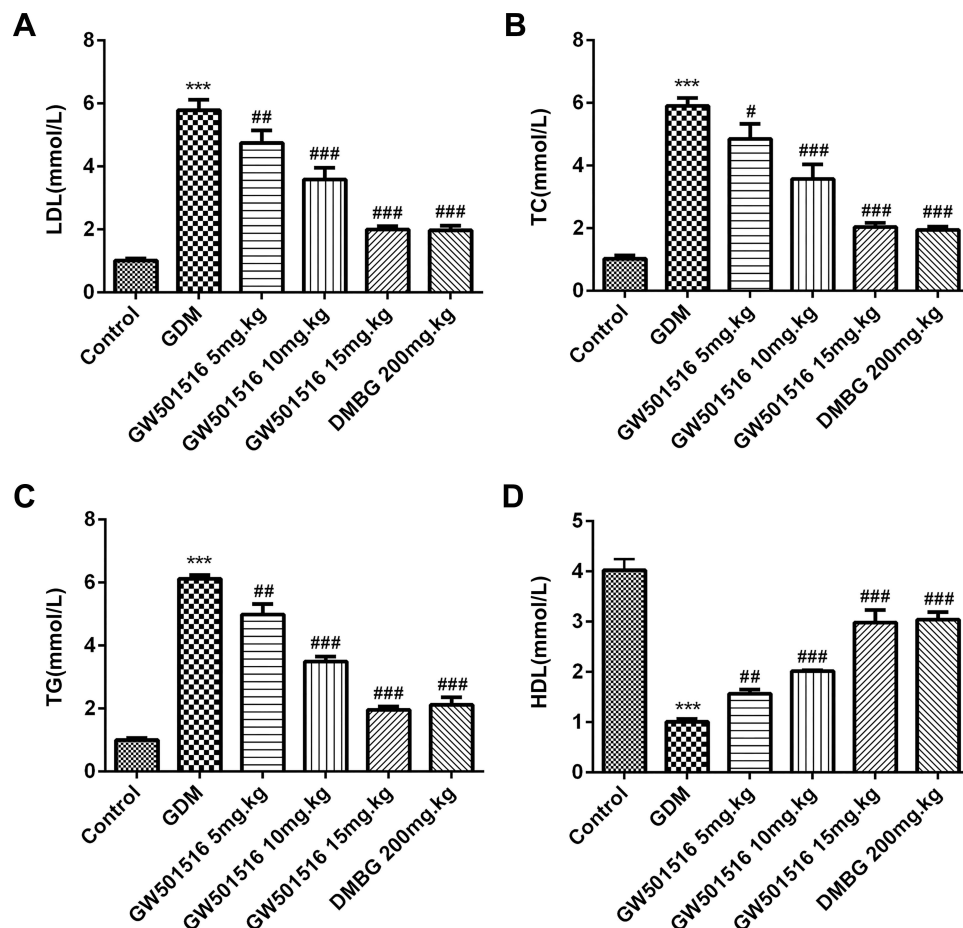
## GW501516 May Have a Therapeutic Effect on GDM Through Regulating the SREBP-1c/GLUT2 Pathway

To further explore the possible targets of GW501516 in the treatment of GDM, we detected the expression of sterol

regulatory element-binding protein 1c (SREBP-1c) and GLUT2. We found that compared with the control group, the expression of SREBP-1c in the GDM group increased, while the expression of SREBP-1c decreased in a dose-dependent manner after GW501516 treatment. The trend of GLUT2 was contrary to that of SREBP-1c (Figure 7A–C). From this, we preliminarily concluded that GW501516 may have a therapeutic effect on GDM by regulating the SREBP-1c/GLUT2 pathway.

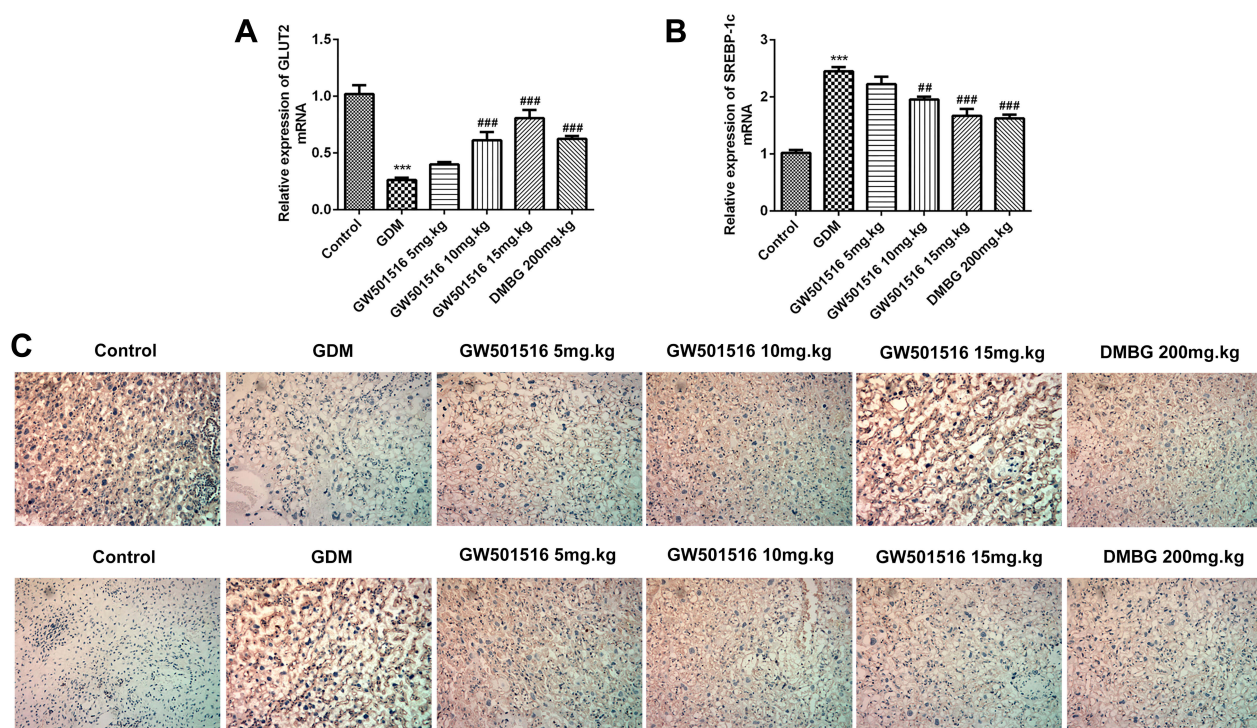
## Discussion

PPAR receptor is a key regulator of adipocyte differentiation and lipid metabolism, which can maintain the stability of blood glucose and lipid, and its activation is necessary for adipogenesis and insulin sensitivity.<sup>18</sup> PPAR is divided into three subtypes: PPAR $\alpha$ , PPAR  $\delta$  and PPAR  $\gamma$ . Studies have shown that the expression of PPAR  $\gamma$  in placenta tissues and adipose tissues is decreased.<sup>19,20</sup> And PPAR  $\gamma$  is involved in the regulation of genes related to lipid synthesis and storage



**Figure 6** GW501516 inhibited the expression of lipid metabolism-related factors in blood. The expressions of LDL (A), TC (B), TG (C) and HDL (D) were determined by the corresponding kit. n=10, \*\*\*p<0.001 vs control; #p<0.05, ###p<0.01, ####p<0.001 vs GDM.





**Figure 7** GW501516 may have a therapeutic effect on GDM through regulating the SREBP-1c/GLUT2 pathway. Expression levels of SREBP-1c (**A**) and GLUT2 (**B**) in the pancreas were detected by RT-qPCR and immunofluorescence staining of SREBP-1c and GLUT2 (**C**). The sample size of RT-qPCR was  $n=5$ , and the sample size of IHC staining was  $n=3$ , \*\*\* $p<0.001$  vs control; ### $p<0.01$ , #### $p<0.001$  vs GDM.

and the production of adipokines and insulin signaling.<sup>21,22</sup> Thiazolidinediones are PPAR  $\gamma$  agonists that can improve the sensitivity of diabetic patients and insulin resistance models.<sup>23</sup> At present, there are few studies on PPAR  $\delta$ , glucose metabolism and lipid metabolism. Doktorova found that activation of PPAR  $\delta$  stimulates the oxidation of fatty acids in adipose tissue and skeletal muscle, improving dyslipidemia in mice and humans.<sup>4</sup> In BXD mice, the expression of PPAR  $\delta$  in muscle was positively correlated with endurance performance and activation of PPAR  $\delta$  can effectively inhibit glucose catabolic metabolism without affecting muscle fiber type or mitochondrial content.<sup>5</sup> However, studies on the expression and role of PPAR  $\delta$  in GDM have not been reported.

After successfully establishing the GDM rat model, we found that the expression of PPAR  $\delta$  was significantly decreased in the GDM rat model, which was consistent with the literature we reviewed, indicating that PPAR  $\delta$  receptor plays an important role in regulating the balance between lipid metabolism and glucose metabolism in GDM. In addition, angiotensin 8 (Angptl8) expression was significantly increased in GDM rats and decreased in a dose-dependent manner after GW501516 treatment. This is because Angptl8 plays an

important role in glucose metabolism during insulin resistance.<sup>24</sup> In 2013, Melton also found that Angptl8 could specifically promote the proliferation of pancreatic  $\beta$  cells and promote the production of insulin, thereby lowering blood glucose.<sup>25</sup>

In this paper, we treated GDM rats with different concentrations of PPAR  $\delta$  receptor agonist GW501516 and found that the expression levels of FBG, FINS and HOMA-IR in GDM rats increased, while ISI levels in GDM rats decreased. Meanwhile, GW501516 can increase the glucose tolerance, improve the damaged islet tissue, and reduce the apoptosis rate of islet cells of GDM rats. Our results are also consistent with those of Li et al.<sup>26</sup> In addition, in our experiment, we only studied the effect of GW501516 on islet tissues and islet cells of GDM rats. Is the effect of GW501516 on other tissues and cells of GDM rats the same as that of islet tissues? Magliano et al.<sup>27</sup> found that short-term administration of GW501516 improves inflammatory state in white adipose tissue and liver damage in high-fructose-fed mice through modulation of the renin-angiotensin system. Early controlled release of peroxisome proliferator-activated receptor  $\beta/\delta$  agonist GW501516 improves diabetic wound healing through redox modulation



of wound microenvironment and GW501516 can facilitate keratinocyte migration.<sup>28</sup> We will continue to study the effects of GW501516 on other organs of GDM rats in later experiments. We also found that GW501516 inhibited the expression of lipid metabolism-related factors in the blood. The results showed that GW501516 had certain therapeutic effect on gestational diabetes.

We investigated the mechanism of GW501516 in the treatment of GDM. Through literature review, we found that GW501516 could restore the expression level of SREBP 1c and GLUT2 in non-alcoholic fatty liver disease to normal. SREBP 1c and GLUT2 play important roles in fat synthesis and glucose metabolism homeostasis. Meanwhile, Kramer et al<sup>16</sup> reported that GW501516 and GW0742, specific activators of PPAR  $\delta$ , can directly affect the insulin action of skeletal muscles by targeting the regulation of SREBP 1c/GLUT2 pathway. Therefore, we speculated whether the therapeutic effect of GW501516 on GDM rats was related to the SREBP 1c/GLUT2 pathway? In our paper, we found that after GDM rats were treated with GW501516, the expression of SREBP 1c decreased in a dose-dependent manner, while the expression of GLUT2 increased in a dose-dependent manner. SREBP 1c is an important transcription factor regulating lipid synthesis and glucose metabolism. Overexpression of SREBP 1c can lead to increased fat synthesis and disorder of glucose and lipid metabolism. GLUT2 plays an important role in glucose homeostasis in organisms. Therefore, we preliminarily concluded that GW501516 may have a therapeutic effect on GDM rats by regulating the SREBP 1c/GLUT2 pathway. The mechanism will be further discussed in our laboratory later.

## Conclusion

The PPAR  $\delta$  agonist GW501516 could improve the blood glucose level, damaged islet tissue and increase the insulin content in the rats with GDM, possibly by regulating the SREBP-1c/GLUT2 pathway.

## Data Sharing Statement

The analyzed data sets generated during the present study are available from the corresponding author on reasonable request.

## Ethics Approval and Consent to Participate

The study protocol was approved by ethics committee of the Shenzhen People's Hospital. All of the procedures

were in compliance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

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## Disclosure

The authors declare that they have no competing interests in this work.

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