PPAR ligands improve impaired metabolic pathways in fetal hearts of diabetic rats

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Abstract

In maternal diabetes, the fetal heart can be structurally and functionally affected. Maternal diets enriched in certain unsaturated fatty acids can activate the nuclear receptors peroxisome proliferator-activated receptors (PPARs) and regulate metabolic and anti-inflammatory pathways during development. Our aim was to investigate whether PPAR\(\alpha\) expression, lipid metabolism, lipoperoxidation, and nitric oxide (NO) production are altered in the fetal hearts of diabetic rats, and to analyze the putative effects of in vivo PPAR activation on these parameters. We found decreased PPAR\(\alpha\) expression in the hearts of male but not female fetuses of diabetic rats when compared with controls. Fetal treatments with the PPAR\(\alpha\) ligand leukotriene B\(4\) upregulated the expression of PPAR\(\alpha\) and target genes involved in fatty acid oxidation in the fetal hearts. Increased concentrations of triglycerides, cholesterol, and phospholipids were found in the hearts of fetuses of diabetic rats. Maternal treatments with diets supplemented with 6% olive oil or 6% safflower oil, enriched in unsaturated fatty acids that can activate PPARs, led to few changes in lipid concentrations, but up-regulated PPAR\(\alpha\) expression in fetal hearts. NO production, which was increased in the hearts of male and female fetuses of diabetic rats when compared with controls. Fetal treatments with the PPAR\(\alpha\) ligand leukotriene B\(4\) upregulated the expression of PPAR\(\alpha\) and target genes involved in fatty acid oxidation in the fetal hearts. Increased concentrations of triglycerides, cholesterol, and phospholipids were found in the hearts of fetuses of diabetic rats. Maternal treatments with diets supplemented with 6% olive oil or 6% safflower oil, enriched in unsaturated fatty acids that can activate PPARs, led to few changes in lipid concentrations, but up-regulated PPAR\(\alpha\) expression in fetal hearts. NO production, which was increased in the hearts of male and female fetuses in the diabetic group, and lipoperoxidation, which was increased in the hearts of male fetuses in the diabetic group, was reduced by the maternal treatments supplemented with safflower oil. In conclusion, impaired PPAR\(\alpha\) expression, altered lipid metabolism, and increased oxidative and nitridergic pathways were evidenced in hearts of fetuses of diabetic rats and were regulated in a gender-dependent manner by treatments enriched with PPAR ligands.

Key Words
- diabetes in pregnancy
- fetus
- heart
- nitric oxide
- PPAR

Introduction

Maternal diabetes can impair the development of the fetus and lead to adverse consequences evident in the offspring both during the perinatal period and in later life (Michael Weindling 2009, Simeoni & Barker 2009, Ali & Dornhorst 2011). The fetal heart is a target organ that can be structurally and functionally affected by maternal diabetes (Molin et al. 2004, Corrigan et al. 2009). In the fetal period, the heart uses glucose and lactate as main oxidative substrate sources and then switches to fatty acids in the neonate, to assure proper energy metabolism according to dietary and physiological conditions (Finck 2007).
Peroxisome proliferator-activated receptor alpha (PPARα), the first of the three PPAR isotypes identified, is an important regulator of myocardial lipid metabolism that also contributes to the control of inflammation and oxidative stress (Wahl & Michalik 2012, Lee et al. 2013, Palomer et al. 2013). PPARα is expressed in the fetal heart and its expression increases after birth (Abbott 2009). In mouse models of diabetes, the fetal heart shows decreased PPARα expression (Lindegaard & Nielsen 2008). In experimental models, both PPARα inactivation and overexpression can lead to metabolic alterations in the heart (Finck 2007, Lindegaard & Nielsen 2008). In adult diabetic patients, chronic inflammation and alterations in energy and lipid homeostasis are involved in cardiac dysfunction (Palomer et al. 2013). In the cardiovascular system, several metabolic and anti-inflammatory pathways are regulated by ligands of the nuclear receptor PPARα both through transactivation and transrepression mechanisms (Lefebvre et al. 2006, Wahl & Michalik 2012). Fibrates are pharmacological ligands of PPARα used to regulate dyslipidemia, which ameliorate cardiovascular diseases and reduce pro-inflammatory markers in the heart of diabetic patients (Lefebvre et al. 2006, Lee et al. 2013). The natural ligands of PPARα include leukotriene (LT) B4, and certain unsaturated fatty acids such as oleic acid and linoleic acid, which can be efficiently transferred through the placenta (Hilh et al. 2002, Herrera et al. 2006, Jawerbaum & Capobianco 2011).

Diabetes induces a pro-oxidant and pro-inflammatory state in both the non-pregnant and the pregnant states (Calkin & Thomas 2008, Lappas et al. 2011). In previous studies, in a mild diabetic rat model during pregnancy we have evaluated and found alterations in PPARα expression and signaling in different fetal organs such as the liver and the lungs (Jawerbaum & White 2010, Martinez et al. 2011a, Kurtz et al. 2012). In the current study, we used the same diabetic rat model to address whether the fetal heart has altered expression of PPARα and lipid-oxidizing enzymes, lipid content, nitric oxide (NO) production, and lipid peroxidation. We also carried out fetal treatments with LTB4 and evaluated maternal diets enriched with 6% olive oil or 6% safflower oil to identify putative changes in PPARα expression, lipid metabolism, and oxidative and nitridergic pathways. Considering the known sexual differences in PPAR signaling and in metabolic and heart diseases, the hearts from both female and male fetuses were separately analyzed (Kautzky-Willer & Handisurya 2009, Yoon 2009, Garcia-Patterson et al. 2011, Mosca et al. 2011).

Materials and methods

Animals

Albino Wistar rats bred in our animal facility were fed ad libitum with commercial rat chow (Asociación Cooperativa Argentina, Buenos Aires, Argentina). To induce diabetes, at 2 days of age, neonates were injected with streptozotocin (90 mg/kg, s.c., Sigma–Aldrich) diluted in citrate buffer (0.05 M, pH 4.5, Sigma–Aldrich), as previously described (Jawerbaum & White 2010, Martinez et al. 2011a). The control animals were injected with citrate buffer alone. The diabetic state was confirmed in 2-month-old rats before mating. The rats were considered diabetic when they presented fasting glycemia values higher than 130 mg/dl. The guidelines for the care and use of animals approved by the local institution were followed (EXP-UBA 0011821), according to the Principles of Laboratory Animal Care (NIH publication number 85-23, revised 1985, http://grants1.nih.gov/grants/olaw/references/phspol.htm).

Control and diabetic female rats were mated with control males. The first day of pregnancy was confirmed by the presence of sperm cells in vaginal smears. On this day, both control and diabetic animals were randomized into two groups: group 1, animals whose fetuses were injected with the PPARα ligand LTB4 or vehicle on days 19, 20, and 21 of pregnancy and group 2, animals fed with diets supplemented with 6% olive oil or 6% safflower oil, enriched in natural PPAR ligands, from days 1 to 21 of pregnancy.

In group 1, to inject the fetuses, the animals (n=9 control and n=9 diabetic rats) were anesthetized in a CO2 chamber on days 19, 20, and 21 of pregnancy and a slight anesthesia maintained with ether vapors. An abdominal incision was performed and the left horn of the uterus was exposed. The fetuses were numbered from the ovary and alternatively injected with LTB4 (0.1 nmol/fetus dissolved in vehicle; Cayman Chemical Co., Ann Arbor, MI, USA) or vehicle (0.3 µl ethanol/fetus, dissolved in saline solution) subcutaneously on their backs through the uterine wall, as described previously (Kurtz et al. 2012). The entire surgery lasted <10 min and the animals were completely recovered after 15 min. On day 21 of pregnancy and after 3 h of the last injection, the rats were killed and the hearts from female and male fetuses were explanted and preserved as described below.

In group 2, control and diabetic mothers were fed from days 1 to 21 of pregnancy either with a standard diet or with diets enriched in unsaturated fatty acids that activate PPARs: 6% olive oil (354% enriched in oleic acid) and 6% safflower oil (226% enriched in linoleic acid).
Expression of PPARα and rate-limiting enzymes in lipid oxidation

The gene expression of PPARα and the rate-limiting enzymes in lipid oxidation of acyl CoA oxidase (ACO) and carnitine palmitoyltransferase 1 (CPT1) were evaluated by RT-PCR, a semi-quantitative method, as described previously (Kurtz et al. 2012). Briefly, RNA was extracted from the hearts of one female and one male fetus from each rat (n = 9 rats in each experimental group) with TRI reagent (Genbiotech, Buenos Aires, Argentina) in accordance with the manufacturer’s instructions. cDNA was synthesized by incubating 1 μg of extracted RNA in a first-strand buffer containing MMLV enzyme (Promega), random primer hexamers, and each of all four dNTPs (Invitrogen), in accordance with the MMLV manufacturer’s instructions. cDNA (2 μl, selected to work within the linear range) was amplified by PCR in a buffer containing dNTPs, magnesium chloride solution, Taq polymerase (GoTaq Polymerase, Promega), and each specific primer in accordance with the Taq polymerase manufacturer’s instructions.

Primers for PPARα were as follows: forward, 5′-TCACAAATGCAATCCGT-3′ and reverse: 5′-GGCTCTTGACCTTTGTT-3′, whose amplification product is a 177-bp fragment (Kurtz et al. 2012). The primers for Aco were as follows: forward: 5′-CCAATACGCAA-TAGTCTCG-3′ and reverse, 5′-CGCTTGATCTGTA-TGGCGAT-3′, whose amplification product is a 363-bp fragment (Lillycrop et al. 2005). The primers for Cpt1 were as follows: forward, 5′-TATCGTCACATTA-GACCGT-3′ and reverse, 5′-CATCTATGACCTCCTG-GACCT-3′, whose amplification product is a 715-bp fragment (Cheng et al. 2004). The primers for the ribosomal protein L30, used as an internal control, were as follows: forward, 5′-CCATCTTGCCGTCTGATC-3′ and reverse, 5′-GGCGAGGATAACCAATTTC-3′, whose amplification product is a 201-bp fragment (Primer 3 Software, Cambridge, MA, USA). The initial step in the reaction was 95 °C for 5 min, followed by 33 cycles for PPARα, 33 cycles for ACO, 34 cycles for CPT1, and 26 cycles for L30, as selected to work within the linear range. Each cycle consisted of denaturation at 95 °C for 15 s, primer annealing at 58 °C for 30 s, and extension at 72 °C for 15 s. The resulting products were separated on a 2% agarose gel and stained with SYBR safe (Invitrogen). The images were taken with an ImageQuant spectrophotometer (GE Healthcare, Buckinghamshire, UK) and the density of the bands was quantified with the Image J software (NIH, Bethesda, MD, USA) and normalized to L30.

Analysis of lipid content

Lipid content was determined by thin layer chromatography, as described previously (Martinez et al. 2011a). Briefly, hearts from three female or three male fetuses from each rat were pooled and homogenized (n = 9 rats in each experimental group) in 500 μl PBS and protein content in the homogenates was measured by the Bradford assay. The tissue lipids were extracted from 470 μl of each homogenate by three rounds of organic extraction in methanol/chloroform (2:1). The lipids were developed by thin layer chromatography in 0.2 mm silica gel plates (Merck), using hexane:ether:acetic acid (80:20:2, v:v:v) as the developing solvent mixture. The lipid species were stained with iodine vapors, identified, and quantified by comparison with known the amounts of standards on the same plate, and densitometric analysis was performed using the Image J software.
Evaluation of NO production

NO production was evaluated by measuring the concentration of its stable metabolite nitrates/nitrites, as previously determined (Kurtz et al. 2012), by using a commercial assay kit (Cayman Chemical Co.). For this, hearts of two female or two male fetuses from each rat were pooled and homogenized \((n=9\) rats in each experimental group) in 1 ml Tris–HCl buffer of pH 7.6, and an aliquot was separated for protein analysis. The nitrates in the supernatant were reduced to nitrites by nitrate reductase, and the total nitrites were measured using nitrates as standard. Analysis of lipoperoxidation

Lipoperoxidation was assessed as described previously (Martinez et al. 2011a), by measuring the concentrations of thiobarbituric acid reactive substances (TBARS), a method widely used to assess peroxidation of fatty acids (Ohkawa et al. 1979). Briefly, hearts from two female or two male fetuses from each rat were pooled and homogenized \((n=9\) rats in each experimental group) in 100 mM Tris–HCl buffer (0.1 mM, pH 7.4). The homogenate was added with 40% trichloroacetic acid (Merck). After centrifugation, the supernatant was added with an equal volume of thiobarbituric acid (46 mM; Sigma–Aldrich), the solution was heated at 95°C and, after cooling, quantified spectrophotometrically at 540 nm. Different concentrations of malondialdehyde (Sigma–Aldrich) subjected to the same conditions as the tissue homogenates were used as standards.

Statistical analysis

Data are presented as means ± s.e.m. Groups were compared by two-way ANOVA in conjunction with Bonferroni’s test. A \(P\) value <0.05 was considered statistically significant.

Results

Decreased expression of PPARα and enzymes involved in lipid oxidation in the hearts of fetuses from diabetic rats: effects of fetal treatments with a PPARα activator

We first analyzed the metabolic parameters of experimental group 1 and found that glycemia and triglyceridemia were increased in both mothers and fetuses from diabetic rats compared with controls on day 21 of pregnancy \((P<0.01)\), with no gender differences and no effects of injecting with the PPARα activator LTB4 on fetuses in the groups evaluated (Table 1).

When we evaluated the expression of PPARα, a regulator of lipid metabolism and anti-inflammatory processes in the heart of adults and in different fetal tissues (Martinez et al. 2011a, Kurtz et al. 2012, Lee et al. 2013), we found a decrease in PPARα expression in the hearts of male but not of female fetuses in the diabetic group compared with controls \((P<0.01, \text{Fig. 1A})\). We also evidenced the ability of the PPARα ligand LTB4 (0.1 nmol, injected into the fetuses through the uterine wall on days 19, 20, and 21 of pregnancy) to upregulate PPARα expression in the hearts of female and male fetuses in the diabetic group and of male fetuses in the control group \((P<0.01, \text{Fig. 1A})\). We next evaluated the expression of Aco and Cpt1, relevant PPARα-target genes that code for rate-limiting enzymes in lipid oxidation. We found a decrease in Aco expression in the hearts of male fetuses but not of female fetuses in the diabetic group compared with controls \((P<0.05, \text{Fig. 1B})\), and a decrease in Cpt1 expression in the hearts of male and female fetuses in the diabetic group compared with controls \((P<0.01, \text{Fig. 1C})\). The fetal injections with the PPARα ligand LTB4 (0.1 nmol) led to an increase in Aco expression in the hearts of female and male fetuses in the diabetic group as well as in the hearts of female fetuses in the control group.

Table 1 Maternal and fetal metabolic parameters in control and diabetic rats from experimental group 1 whose fetuses were injected with \(\text{LTB}_4\) or vehicle on days 19, 20, and 21 of pregnancy. Values represent mean ± s.e.m., obtained from mothers and female or male fetuses from \(n=9\) rats in experimental group 1. Two-way ANOVA in conjunction with Bonferroni’s test was performed

<table>
<thead>
<tr>
<th>Maternal data</th>
<th>Glycemia (mg/dl)</th>
<th>Triglyceridemia (g/l)</th>
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<tbody>
<tr>
<td>Control</td>
<td>92 ± 8</td>
<td>1.9 ± 0.3</td>
</tr>
<tr>
<td>Diabetic</td>
<td>214 ± 24</td>
<td>4.0 ± 0.3</td>
</tr>
<tr>
<td>Fetal data (combined males/females)</td>
<td></td>
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<tr>
<td>Control vehicle</td>
<td>47 ± 5</td>
<td>0.54 ± 0.06</td>
</tr>
<tr>
<td>Control (\text{LTB}_4)</td>
<td>49 ± 10</td>
<td>0.60 ± 0.05</td>
</tr>
<tr>
<td>Diabetic vehicle</td>
<td>146 ± 11</td>
<td>0.89 ± 0.07</td>
</tr>
<tr>
<td>Diabetic (\text{LTB}_4)</td>
<td>135 ± 9</td>
<td>0.88 ± 0.04</td>
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\(\text{aP}<0.001 \text{ diabetic vs control}; \text{bP}<0.01, \text{cP}<0.001 \text{ diabetic with vehicle vs control with vehicle.}\)
Effects of maternal diet in the fetal heart

M KURTZ and others

Male fetuses from control and diabetic rats. We found lipid concentrations in the hearts of female and oxidation in the fetal hearts of diabetic rats, we evaluated expression of PPAR considering the sex-dependent changes observed in the safflower oil on lipid concentrations

Effect of maternal diets enriched with olive oil or safflower oil on lipid concentrations

Considering the sex-dependent changes observed in the expression of PPAR and its target genes involved in lipid oxidation in the fetal hearts of diabetic rats, we evaluated lipid concentrations in the hearts of female and male fetuses from control and diabetic rats. We found increased concentrations of triglycerides (P<0.001), cholesterol (P<0.05), and phospholipids (P<0.05) and no changes in cholesteryl esters in the heart of female and male fetuses in the diabetic group compared with controls (Fig. 2). As we have previously shown that dietary supplementation with 6% olive oil and 6% safflower oil can regulate lipid content in different fetal tissues and through developmental stages (Capobianco et al. 2008a, b, Kurtz et al. 2014), in this study we analyzed the effect of these dietary supplementations, administered from days 1 to 21 of pregnancy. The metabolic parameters evaluated in these animals (experimental group 2) are shown in Table 2. On day 21 of pregnancy, increased glycemia and triglyceridemia were observed in both diabetic mother and fetus rats fed the standard diet compared with controls fed the standard diet (P<0.001), with no gender differences, and no effects of the diets supplemented with 6% olive oil or 6% safflower oil in the groups evaluated (Table 2).

When we analyzed the effect of the maternal diets supplemented with 6% olive and 6% safflower oils on lipid content in the fetal heart, we observed only a few changes, all of them evidenced in the hearts of female fetuses: an increase in triglycerides in the diabetic group fed the olive oil-supplemented diet, a decrease in cholesterol in the control group fed the olive oil-supplemented diet, and an increase in cholesteryl esters in the control group fed the safflower oil-supplemented diet (P<0.05, Fig. 2).

The absence of marked effects on lipid content was evidenced even when the olive oil- and the safflower oil-supplemented diets induced relevant changes in PPAR expression in the hearts of both female and male fetuses (Fig. 3A). Indeed, PPAR expression was negatively regulated by both the olive oil-supplemented diet in the hearts of female fetuses in the diabetic group and positively regulated by the olive oil- and the safflower oil-supplemented diets in the hearts of male fetuses in control and diabetic groups (P<0.05, Fig. 3A). No effects were observed when expression of Aco and Cpt1 was evaluated in the control and the diabetic groups fed olive oil- and safflower oil-supplemented diets compared with the respective groups fed the standard diet (Fig. 3B and C).

Effect of maternal diets enriched with olive oil or safflower oil on NO production

Our previous studies suggest that PPAR is a relevant regulator of NO production in different fetal organs (Martinez et al. 2011b, Kurtz et al. 2012). Excessive
production of NO can affect embryo and fetal development and is a marker of a pro-inflammatory state (Lappas et al. 2011). In this work, we found increased nitrates/nitrites concentrations, an index of NO production, in the hearts of female and male fetuses of diabetic rats when compared with controls ($P < 0.01$, Fig. 4). We also evidenced the ability of the maternal diets supplemented with olive oil to decrease nitrates/nitrites in the hearts of male fetuses in the diabetic group ($P < 0.01$, Fig. 4), and the ability of the maternal diets supplemented with 6% safflower oil to decrease nitrates/nitrites in the hearts of both female and male fetuses in the diabetic group ($P < 0.01$, Fig. 4).

Table 2  Maternal and fetal metabolic parameters in control and diabetic rats from experimental group 2 fed with standard diet supplemented with or without 6% olive oil or 6% safflower oil from days 1 to 21 of pregnancy. Values represent mean ± S.E.M., obtained from mothers and female or male fetuses from $n=9$ rats in experimental group 2. Two-way ANOVA in conjunction with Bonferroni’s test was performed.

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Diabetic</th>
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<tbody>
<tr>
<td></td>
<td>Standard diet</td>
<td>Standard diet supplemented with 6% olive oil</td>
</tr>
<tr>
<td>Maternal data</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glycemia (mg/dl)</td>
<td>93 ± 7</td>
<td>105 ± 9</td>
</tr>
<tr>
<td>Triglyceridemia (g/l)</td>
<td>2.1 ± 0.15</td>
<td>2.03 ± 0.25</td>
</tr>
<tr>
<td>Fetal data</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glycemia (mg/dl)</td>
<td>46 ± 7</td>
<td>45 ± 9</td>
</tr>
<tr>
<td>Triglyceridemia (g/l)</td>
<td>0.52 ± 0.05</td>
<td>0.59 ± 0.06</td>
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$^aP < 0.01$ diabetic with standard diet vs control with standard diet.
6% safflower oil from days 1 to 21 of pregnancy. Lipoperoxidation was observed to be increased in the hearts of male fetuses (\( P < 0.01 \)), although not in the hearts of female fetuses from diabetic rats when compared with controls (Fig. 5). We also evidenced the ability of the maternal diets supplemented with 6% safflower oil, although not of those supplemented with 6% olive oil, to decrease lipoperoxidation in the hearts of male fetuses in the diabetic group (\( P < 0.01 \), Fig. 5).

**Discussion**

In this study, we found sex-dependent increases in lipid content, NO production, and lipoperoxidation, together with decreased expression of PPAR\( \alpha \) and rate-limiting enzymes involved in lipid oxidation in the hearts of term fetuses of diabetic rats. Fetal or maternal treatments with PPAR activators regulated most of the parameters evaluated in a sex-dependent way. PPAR\( \alpha \) expression was upregulated both by fetal treatments with PPAR\( \alpha \) ligands and by maternal treatments with diets enriched with PPAR\( \alpha \) ligands during gestation, thus indicating the ability of these treatments to amplify PPAR\( \alpha \) signaling pathways in the fetal heart.

PPAR\( \alpha \) is a master regulator of lipid oxidation in metabolic tissues such as liver, kidney, and heart (Lefebvre et al. 2006). In this study, we observed a decrease in PPAR\( \alpha \) and its target genes Aco and Cpt1 in the hearts of male fetuses from diabetic rats, although only a decrease in Cpt1 in the hearts of female fetuses in the diabetic group when compared with controls. These decreases, together with

**Effect of maternal diets enriched with olive oil or safflower oil on lipoperoxidation**

As PPAR\( \alpha \) activation can regulate anti-oxidant pathways in different tissues during development (Martinez et al. 2011a,b) and lipoperoxidation is a marker of oxidative stress previously found to be increased in the fetuses and placentas of diabetic rats (Jawerbaum & White 2010), we next evaluated lipoperoxidation in the heart of fetuses from control and diabetic rats that were treated with or without diets supplemented with 6% olive oil or
the increased maternal and fetal triglyceridemia, may be involved in the overaccumulation of lipids in the hearts of fetuses in the diabetes experimental model evaluated. Previous studies performed in Akita mice, a more severe genetic model of diabetes, showed no changes in triglyceride content but a decrease in lipid transporters and in the expression of PPARα in the fetal hearts (Lindegaard & Nielsen 2008). It is known that PPARα regulates lipid oxidation in the hearts of neonates and adults (Finck 2007), but its function as a lipid metabolic sensor is not yet clear in the fetuses. In this work, we found that the PPARα ligand LTB₄ is able to increase the expression of the lipid-oxidizing enzymes ACO and CPT1 in the fetal heart. Nevertheless, maternal diets enriched with PPAR ligands did not increase the expression of ACO and CPT1, and did not lead to a decreased content of lipid species in the fetal heart. It is known that PPAR ligands decrease lipid content in the fetal liver (Martinez et al. 2011a) and that the liver is the main target in treatment with PPAR ligands in adult animals (Finck 2007). However, in this study, we found that both the maternal diets enriched with PPAR ligands and the fetal treatments with the PPARα agonist LTB₄ led to an increased expression of PPARα in the fetal heart, indicating that PPARα signaling is stimulated in this fetal organ by the given treatments. Several studies have shown that different PPAR ligands can induce different conformational changes in PPARs, as well as different biological responses (Hostetler et al. 2005, Gregoire et al. 2009). Indeed, ligand activation stabilizes the interaction between PPAR and RXR, enhances the formation of functional PPAR–RXR heterodimers, and can also regulate the heterodimerization with different partners (Harmon et al. 2011, Balanarasimha et al. 2014). The resulting interaction plays a key role leading to the formation of complexes, which result in gene activation or repression (Harmon et al. 2011). Thus, it is possible that transactivation pathways are those mainly activated by LTB₄ in the fetal heart, leading to the expression of regulation of lipid oxidizing enzymes, whereas transrepression pathways are those mainly activated by the maternal dietary treatments with unsaturated fatty acids. However, further studies are needed to address this point, as this work was limited due to the evaluation of PPAR ligands in different in vivo approaches. The formation of PPAR complexes and the resulting activity are tissue-dependent, and PPARs’ capacity to activate or repress metabolic and pro-inflammatory pathways was similarly evidenced using different in vivo and in vitro approaches in the fetal liver and lung (Harmon et al. 2011, Martinez et al. 2011a, Kurtz et al. 2012).

In the heart, relevant extra-metabolic functions related with the control of the pro-inflammatory and pro-oxidant environment can be exerted by PPARz activation (Palomer et al. 2013). We found that NO production, which is increased in pro-inflammatory states (Lappas et al. 2011), was increased in the hearts of male and female fetuses from diabetic rats. Previous research has demonstrated that diabetic embryopathy is related with NO overproduction, and that different fetal organs such as liver, lung, and placenta have increased NO production when being developed in a diabetic mother (Martinez et al. 2011a,b, Kurtz et al. 2012). Moreover, in the presence of oxidative stress, NO overproduction leads to the formation of peroxynitrites, which induce severe damage to different organs, including the heart (Mungrue et al. 2002, Lappas et al. 2011). Thus, it was interesting to find that maternal diets enriched with 6% olive oil (only in male fetuses) and 6% safflower oil (in both male and female fetuses) prevented the overproduction of NO in the hearts of fetuses from diabetic rats. Future research is needed to identify whether putative epigenetic changes are involved in the fetal changes observed, and to analyze the effects of these treatments in the heart of neonates.

It is known that cardiac dysfunction is sex-dependent, possibly in part due to the differential ability of estrogens to regulate lipid metabolism (Kauzy-Willer & Handisurya 2009, Oosthuyse & Bosch 2012). PPARz signaling is also sex-dependent due to the effects of estrogens and androgens on PPARz expression and due to the use of common coactivators by different nuclear receptors. Therefore, estrogens and androgens may also play a role in the heart of neonates (Handisurya et al. 2009).
receptors (Collett et al. 2000, Yoon 2009, Benz et al. 2012). Indeed, we have previously found sex-dependent effects of PPARz agonists in different tissues even at fetal stage (Kurtz et al. 2014). Interestingly, in this study, we found that only the heart of male fetuses showed increased lipoperoxidation in the diabetic group. Increased oxidative stress, related with impairments in energy metabolism, stress in endoplasmic reticulum, and apoptosis, has been described in the heart of adult diabetic animals (Li et al. 2007, Palomer et al. 2013).

Previous research has identified the ability of maternal diets enriched with PPAR ligands to reduce oxidative stress during embryo development and to reduce pro-inflammatory markers in the placenta and fetal lungs (Higa et al. 2012, Kurtz et al. 2012, Martinez et al. 2012). A limitation of this study was that, due to the lack of sufficient biological material, we were unable to measure NO production and lipid peroxidation in the LTB4-treated group. Although the 6% olive oil-supplemented diet is enriched with both oleic acid and polyphenols that can exert antioxidant properties (Martin-Pelaez et al. 2013), only the maternal treatments supplemented with 6% safflower oil were able to decrease lipoperoxidation in the hearts of male fetuses from diabetic rats. This may be due to the increased ability of the safflower oil-supplemented diet to amplify PPARz signaling pathways. Indeed, we showed that this diet leads to a greater increase in PPARz expression in the hearts of fetuses from diabetic rats. In addition, safflower oil is enriched with linoleic acid, which can activate PPARs by itself and by the production of eicosanoids that can further activate PPARs (Jawerbaum & Capobianco 2011).

In conclusion, the fetal hearts evaluated in the mild diabetic rat model were profoundly affected, as evidenced by overaccumulation of lipid, increased NO, and lipoperoxidation, which may lead to heart dysfunction later in life. These alterations were more marked in the hearts of male fetuses, which showed reduced expression of PPARz and target genes involved in fatty acid oxidation. Moreover, PPARz signaling in the fetal heart was mainly activated by the maternal dietary treatment enriched with 6% safflower oil, leading to the regulation of lipoperoxidation and NO production, by ameliorating the pro-oxidant and pro-inflammatory environment and thus may prevent the future development of cardiac dysfunction.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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