Effects of Taurine Supplementation on Hepatic Markers of Inflammation and Lipid Metabolism in Mothers and Offspring in the Setting of Maternal Obesity

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Abstract

Maternal obesity is associated with obesity and metabolic disorders in offspring. However, intervention strategies to reverse or ameliorate the effects of maternal obesity on offspring health are limited. Following maternal undernutrition, taurine supplementation can improve outcomes in offspring, possibly via effects on glucose homeostasis and insulin secretion. The effects of taurine in mediating inflammatory processes as a protective mechanism has not yet been investigated. Further, the efficacy of taurine supplementation in the setting of maternal obesity is not known. Using a model of maternal obesity, we examined the effects of maternal taurine supplementation on outcomes related to inflammation and lipid metabolism in mothers and neonates. Time-mated Wistar rats were randomised to either: 1) control: control diet during pregnancy and lactation (CON); 2) CON supplemented with 1.5% taurine in drinking water (CT); 3) maternal obesogenic diet (high fat, high fructose) during pregnancy and lactation (MO); or 4) MO supplemented with taurine (MOT). Maternal and neonatal weights, plasma cytokines and hepatic gene expression were analysed. A MO diet resulted in maternal hyperinsulinemia and hyperleptinemia and increased plasma glucose, glutamate and TNF-α concentrations. Taurine normalised maternal plasma TNF-α and glutamate concentrations in MOT animals. Both MO and MOT mothers displayed evidence of fatty liver accompanied by alterations in key markers of hepatic lipid metabolism. MO neonates displayed a pro-inflammatory hepatic profile which was partially rescued in MOT offspring. Conversely, a pro-inflammatory phenotype was observed in MOT mothers suggesting a possible maternal trade-off to protect the neonate. Despite protective effects of taurine in MOT offspring, neonatal mortality was increased in CT neonates, indicating possible adverse effects of taurine in the setting of normal pregnancy. These data suggest that maternal taurine supplementation may ameliorate the adverse effects observed in offspring following a maternal obesogenic diet but these effects are dependent upon prior maternal nutritional background.

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Introduction

Obesity and overweight during pregnancy has become a major emerging issue for maternal and neonatal health over the past decade [1,2]. Periconceptional and gestational obesity are associated with insulin resistance (IR) and low-grade inflammation which increases the incidence of gestational diabetes, preeclampsia, miscarriage, and neonatal mortality and the long-term risk of developing metabolic syndrome [3–5]. A recent clinical study highlighted the relationship between intrahepatic fat and IR in women with previous gestational diabetes (GDM) [6], indicating mild hepatic steatosis in postpartum women may contribute to IR-related metabolic dysfunction.

In addition to metabolic disorders and adverse pregnancy outcomes, maternal obesity has been shown to impact the long term health of the offspring [7]. The developmental origins of health and disease (DOHaD) paradigm proposes that insults such as poor maternal nutrition during critical windows of development, can lead to an increased propensity in offspring to develop obesity and related metabolic and cardiovascular disorders in later life [8]. Both human studies [9,10] and animal models [11,12] clearly show a link between maternal obesity and heightened risk of metabolic disorders in offspring, yet relatively little is known about the mechanisms involved. Therefore, broad lifestyle recommendations remain the most common preventative strategies [7].

A number of studies have reported the effectiveness of taurine (2-aminoethanesulfonic acid) in treating IR [13–15]. Taurine is a sulphonic amino acid derived from methionine and cysteine metabolism and is found ubiquitously in all mammalian tissues. The synthesis and metabolism of taurine has known species-specific differences although taurine can be synthesised in vivo in both the human and rodent [16]. Taurine is involved in bile acid synthesis, osmoregulation, modulation of neurotransmitters, glucose homeostasis and insulin secretion [17,18]. Reports suggest that taurine supplementation can enhance insulin sensitivity.
through modification of insulin signaling enzymes in fructose-fed rats [19]. Furthermore, maternal taurine supplementation to low protein mothers has been documented to normalise pancreatic islet development in offspring with normalisation of glucose and insulin homeostasis in later life [20–22]. These beneficial effects on glucose metabolism have been shown to persist into adult life [23]. Although the effects of maternal taurine supplementation as relates to improved glucose homeostasis and beta-cell function in offspring have been well documented, the direct effects of taurine supplementation on the mother are not well documented. Further, taurine has been proposed to play a role in mediating inflammatory processes but this has yet to be examined as a potential mechanism by which maternal taurine supplementation leads to protective effects in the offspring. Recent work by Lin et al. has shown that taurine can improves obesity-induced inflammatory responses and modulates the unbalanced phenotype of adipose tissue macrophages [24]. Obesity is characterised by a state of low grade inflammation and maternal obesity is well established to lead to obesity and related metabolic disorders in offspring [12,25]. In this context, the efficacy of maternal taurine supplementation as an intervention in the setting of maternal obesity has yet to be investigated. Since most studies in the area of developmental programming focus on offspring outcomes, very little attention is paid to the direct effects on maternal health and wellbeing. The current study therefore investigated the effect of taurine supplementation to pregnant and lactating dams fed either a control or obesogenic diet on both maternal and offspring metabolic and hepatic inflammatory profiles.

Methods

Animal Model

**Ethics statement.** All procedures described were approved by the Animal Ethics Committee at the University of Auckland (Approval R838).

Virgin Wistar rats were time mated at 100 days of age using an estrous cycle monitor (EC-40, Fine Science Tools, San Francisco, USA). Day 1 of pregnancy was determined by the presence of spermatozoa after a vaginal smear. Pregnant rats were then housed individually with free access to food and water and maintained at 25°C and a 12 h light: 12 h darkness cycle. Animals were randomly assigned to one of four nutritional groups: control group (CON) fed a standard chow diet (Diet 2018, 24% calories from protein, 18% from fat, 58% from carbohydrate, Harlan Teklad, Blackthorn, Bicester, UK) ad libitum throughout pregnancy and lactation (n = 9); control taurine group (CT) fed Harlan chow diet with additional 1.5% w/v taurine supplementation in drinking water [20,22] (n = 7); maternal obesogenic diet group (MO) fed a high-fat high-fructose diet (20% calories from protein, 45% from fat, 35% from carbohydrate) (including 26% from fructose); Diet D05101602, Research Diets, NJ, USA; n = 8); maternal obesogenic diet and taurine group (MOT) fed the obesogenic diet with additional 1.5% w/v taurine supplementation in drinking water (n = 8).

Two discrete time points were investigated. Firstly, effects of maternal taurine supplementation on neonatal outcomes and secondly the direct effects of taurine on maternal lipid and inflammatory profiles at the end of the lactation period. Maternal body weight, food and fluid intake were recorded daily. After birth, litter size was adjusted to 8 pups per litter (post-weaning offspring were utilised in an independent study). Neonatal plasma and liver samples were collected from randomly chosen excluded pups following decapitation. Litter size, sex ratio and birth weight were recorded at the time of birth. At the end of lactation, dams were fasted overnight and killed by decapitation following anaesthesia with sodium pentobarbitone (60 mg/kg IP). Maternal body composition was measured by dual energy X-ray absorptiometry using dedicated small animal software (DEXA, Lunar Prodigy, Madison, WI, USA). Maternal and neonatal blood glucose and β-hydroxybutyrate (BHB) were measured from tail blood samples using a glucose meter (Optium, Abbott Laboratories) at the time of cull.

Plasma Analysis

ELISA kits were used to measure plasma insulin and leptin (CrystalChem, USA), tumour necrosis factor (TNF)-α, interleukin (IL)-1β and IL-6 (Quantikine ELISA; R&D Systems Europe, Abingdon, UK). Plasma uric acid was measured using a commercially sourced assay kit (Cayman Chemical, Ann Arbor, MI, USA). The homeostasis model assessment of insulin resistance (HOMA-IR) was calculated as: Fasting glucose (mmol/l)/fasting insulin (mU/l)/22.5 [26]. Maternal plasma glutamate and taurine were analysed using a Hitachi 902 autoanalyser (Hitachi High Technologies Corporation, Tokyo, Japan). Homocysteine (HcY) was measured by commercial immunoassay (Abbott AxSYM system). Limited plasma sample precluded measurement of taurine, glutamate or HcY in neonatal samples.

Hepatic mRNA Expression

Total RNA was isolated from liver tissue using RNaseasy™ mini kit (QIAGEN, Hilden, Germany) and cDNA synthesized from 2 μg of RNA by using SuperScript® VILO™ cDNA Synthesis Kit (Invitrogen™, Life Technologies Corporation, California, USA). Real-time PCR analysis for maternal hepatic sterol regulatory element-binding protein-1c (SREBP-1c), peroxisome proliferator-activated receptor alpha (PPARα), lipoprotein lipase (LPL), silent mating type information regulation 2 homolog 1 (SIRT1), fructokinase and phosphoenolpyruvate carboxykinase (PEPCK) expression was performed using LightCycler®480 SYBR green I master (Roche Diagnostics; Auckland, New Zealand). The relative amounts of genes were quantitated using standard curve and normalised to the geometric mean of cyclophilin A and β-actin expression. Real time PCR analysis for maternal hepatic fatty acid synthase (FASN), CD36, TNFα, IL-1β, IL-1R1 and all neonatal samples was carried out by using PreDeveloped TaqMan® Assay Reagent Kits in the ABI 7900HT Fast Real-Time PCR System (Applied Biosystems). To control for between-sample variability, mRNA levels were normalized to the geometric mean of cyclophilin A and hypoxanthine phosphoribosyltransferase (HPRT) for each sample by subtracting the Ct of controls from the Ct for the gene of interest producing a Ct value. The ΔCt for each treatment sample was compared to the mean ΔCt for control samples using the relative quantification 2-ΔΔCt method to determine fold-change [27]. Primer details are provided in Table S1.

Histological Analysis

5 mm thick representative sections from the left lobe of maternal livers were fixed in paraformaldehyde and paraflin embedded. Cross sections were prepared using Leica RM 2135 rotary microtome (Leica Instruments, Nussloch, Germany). Haematoxylin and Eosin (H&E) staining was conducted for general histology. Sections were mounted using d难受te plasticizer xylene (DPX) mounting medium (BioLab ltd, New Zealand) and analysed under light microscope (Nikon 800, Tokyo, Japan) and images taken (Nikon FDX-35, Tokyo, Japan) and processed with NIS Elements-D software (Nikon, Tokyo, Japan). Scoring of steatosis, lobular inflammation, hepatocyte ballooning and overall
NAS score (NAFLD Activity Score) was undertaken by a blinded observer using the methodology of Kleiner et al. [28]. The scoring system comprised four primary features evaluated semi-quantitatively: steatosis (0–3), lobular inflammation (0–3), hepatocellular ballooning (0–2) and NAS score (unweighted sum of steatosis, lobular inflammation, and hepatocellular ballooning scores).

Statistical Analysis

Data analysis was completed using factorial analysis of variance (ANOVA) using SigmaStat software (Systat Software, San Jose, Ca, USA). For the maternal data, maternal diet and taurine were used as factors. For neonatal data, maternal diet, taurine and were used as factors. Where appropriate, post-hoc analyses were performed (Holm-Sidak method) to determine which groups were significantly different from each other. Maternal body weight data was analysed using repeated measures analysis. Maternal liver histology data were analysed via non-parametric methods (Wilcoxon rank sum test followed by Bonferroni correction). Data that failed to meet the criteria required for parametric analysis (normal distribution and equal variance) were transformed where necessary. All data are shown as means ± SEM. A p-value of <0.05 was accepted as statistically significant.

Results

Maternal and Offspring Weights

A maternal obesogenic diet resulted in an overall increase in maternal body weights in both MO and MOT groups (Figure 1) during pregnancy compared to CON and CT groups and was statistically significant from day 8 to day 18 gestation. Of note there was a small decrease in maternal body weights in CT dams compared to CON which was not evident in the MOT group and this was reflected in a significant maternal diet x taurine statistical interaction (p<0.0001). In the immediate post-partum period, MO and MOT dams remained significantly heavier than CON and CT mothers but there were no differences in maternal body weights between groups for the remainder of the lactation period (Figure 1). Birth weights were significantly reduced in female MO and MOT offspring compared to CON and CT groups (Table 1) but were not different between male groups. Interestingly, maternal taurine supplementation significantly increased neonatal mortality in the CT group (CON 1.4±0.1%; CT 7.1±4.0%, p<0.05), without significant difference between CON, MO and MOT groups. Litter size and sex ratios were not affected by maternal diet or taurine supplementation (data not shown). Total maternal fat mass was significantly increased in MO but not in MOT animals when compared with CON at the end of lactation (Table 1). An effect of maternal diet was observed on relative maternal liver weight, increasing significantly in MO and MOT groups compared to CON (Table 1). Weaning weight was significantly increased in MO offspring compared to CON. There was a significant maternal diet x taurine interaction (p = 0.01) in males whereby CT body weights were higher than CON but the reverse holding true in MO groups with MO weights being higher than MOT. Weaning (postnatal day 22) weights were not significantly different between any of the female offspring groups (Table 1).

Maternal Plasma Profile

Maternal plasma taurine was increased in CT and MOT groups compared to CON and MO groups (Table 2). Maternal fasting glucose was increased while BHB concentrations decreased in MO and MOT groups compared to CON and CT groups and there was no effect of taurine (Table 2). Maternal plasma uric acid was significantly decreased in MOT compared to MO groups (Table 2). MO and MOT groups displayed significant hyperinsulinemia and increased HOMA-IR indices when compared to CON and CT groups (Table 2 and Figure 2a). Hyperleptinemia was observed in MO but not MOT groups although there was no significant overall effect of taurine supplementation. TNF-α was significantly increased in MO compared to CON, CT and MOT groups. Plasma IL-1β and IL-6 concentrations were not affected by diet or taurine supplementation (Table 2). Maternal plasma homocysteine (HcY) concentrations were significantly increased in response to the maternal obesogenic diet in MO and MOT dams (Figure 2b). Plasma glutamate concentrations were significantly increased in MO dams compared to all other groups (Figure 2b, maternal diet x taurine interaction p<0.05).

Maternal Hepatic Morphology

Given the changes observed in maternal liver weights in response to diet as shown in Table 1, analysis of the histological features associated with NAFLD was undertaken using the method of Kleiner et al. [28]. Scores related to steatosis, lobular inflammation, hepatocyte ballooning and overall NAS score were significantly increased in MO and MOT groups compared to CON and CT groups. There was no significant effect of maternal taurine supplementation on any of the markers analysed (Table 3 and Figure 3).

Maternal Hepatic Gene Expression

To further assess the fatty liver phenotype observed via hepatic morphology, we conducted gene expression analysis for genes related to lipogenesis and inflammatory responsiveness. Hepatic SREBP-1c expression was significantly increased in the MO and MOT groups when compared to the CON and CT groups (Figure 4a). FASN expression was increased in both MO and MOT groups compared to CON and CT (Figure 4b). PPARγ was significantly down-regulated in MO and MOT groups compared...
to CON and CT groups (Figure 4c). LPL was significantly increased in MO and MOT groups compared to CON and CT and increased in CT versus CON groups (Figure 4d). Fructokinase expression was significantly reduced in MOT group compared to CON, CT and MO groups (Figure 4e). CD36 expression was significantly increased in CT and MOT groups when compared to CON and MO groups (Figure 4f). SIRT1 expression was increased in CT versus CON and MOT groups (Figure 4g). PEPCK was significantly reduced in both MO and MOT groups compared to CON and CT (Figure 4h). Hepatic TNF-α, IL-1β and IL-1R1 expression was significantly increased in MOT animals compared to CON, CT and MO groups (Figures 5a–c). No effect was observed on maternal hepatic TNFR1 expression (Figure 5d). Overall main effect statistical data are provided in Table S2.

**Neonatal Plasma and Physiology Profile**

There were no significant effects of maternal diet or taurine supplementation on neonatal plasma insulin or leptin concentrations although there was a trend toward reduced leptin concentrations in MO and MOT offspring versus controls (p = 0.075 and p = 0.07 for males and females respectively, Table 4). Plasma BHB concentrations were significantly reduced in CT, MO and MOT offspring compared to CON and there was a significant diet and taurine interaction in both male and female neonates (CON>CT but MO<MOT) (Table 4). Female offspring relative liver weights were significantly increased in MO and MOT compared to CON.

**Table 1. Maternal, neonatal and weaning weight data.**

| Groups   | Effect | Diet | Taurine | Interaction |
|----------|--------|------|---------|-------------|
| Maternal weight (g) | F = 8.461 | 0.007 | F = 0.00835 | 0.926 |
| Maternal liver weight (% body weight) | F = 53.203 | <0.001 | F = 0.00149 | 0.969 |
| Maternal total fat (%) | F = 6.140 | 0.020 | F = 0.0465 | 1.118 |
| Birthweights (male, g) | F = 0.895 | 0.348 | F = 3.902 | 2.410 |
| Birthweights (female, g) | F = 0.003 | 0.003 | F = 0.481 | 0.015 |
| Weaning weight (male, g) | F = 29.175 | 0.003 | F = 0.491 | 0.318 |
| Weaning weight (female, g) | F = 1.804 | 0.182 | F = 0.451 | 5.612 |

Values are presented as means ± SEM, n = 7–9 per group for maternal data, minimum 20 per group for birth and weaning weights. Bold font indicates effect P value <0.05 via two-way ANOVA. Upper case letter (A, B) superscripts indicate comparison procedures were conducted between all groups fed MO diet and all groups fed CON diet. Lower case letter superscripts (a,b,c) indicate multiple comparison procedures were conducted for diet and taurine interaction. Groups that do not share the same letter are significantly different from each other (p<0.05).
and CT groups (CON 3.9 ± 0.1%; CT 3.8 ± 0.2%; MO 4.0 ± 0.1%; MOT 4.2 ± 0.1%). Interestingly, male offspring relative liver weights did not differ among groups (data not shown).

**Neonatal Hepatic Inflammatory Profile**

IL-1R1 expression in male and female neonates was significantly increased in MO offspring compared to all other groups (Figure 6a). Hepatic IL-1β expression was increased in female MO neonates compared to all other female groups. In male neonates, IL-1β was decreased in CT versus all other groups (Figure 6b). TNFR1 expression was decreased in MOT female neonates compared to the MO group (Figure 6c). There were no significant differences in TNFR1 between the male neonatal groups. There were no differences in neonatal hepatic TNF-α expression (Figure 6d). There were no significant differences in markers related to IR or lipid metabolism (Figure S1).

**Discussion**

The present study demonstrated significant metabolic and inflammatory changes in dams and neonates in response to a maternal high-fat-high-fructose diet. Dams fed the obesogenic diet displayed an adverse metabolic phenotype which included increased weight gain, hyperglycaemia, hyperhomocysteinemia, IR, and evidence of hepatic steatosis and inflammation. Maternal taurine supplementation significantly attenuated the proinflammatory plasma profile and plasma glutamate induced by obesogenic diet. Despite these beneficial systemic effects, maternal hepatic lipid metabolism and inflammatory profile were further impaired in response to taurine supplementation. Conversely, the neonatal hepatic immunophenotype, which was worsened by maternal obesogenic diet, was normalised by maternal taurine supplementation. However, it must be noted that the neonatal and maternal time-points represent independent observations and causative associations can only be speculated upon. Nonetheless, even as independent observations, both the maternal and neonatal data point to a marked effect of taurine supplementation on modifying maternal and neonatal outcomes in the setting of a maternal obesogenic diet.

This study demonstrated that maternal obesogenic diet induced significant hepatic steatosis in dams. Hepatic lipogenic gene expression was altered by the obesogenic diet. Increased SREBP-1c, FASN, LPL and decreased PPARα expression was observed in MO dams. There are three possible main pathways by which these altered gene expression may contribute to hepatic steatosis. Firstly, SREBP-1c is an important transcriptional regulator of fatty acid synthesis and regulates the expression of key enzymes such as acetyl coenzyme-A carboxylase-1 (ACC-1), Stearoyl-CoA desaturase-1 (SCD1) and FASN [29,30]. SREBP-1c expression is insulin sensitive [31], and overexpression can induce de novo lipogenesis via upregulation of FASN [29,32] potentially contributing to lipotoxicity. Secondly, PPARα is a mediator of hepatic fatty acid β-oxidation [33]. As a member of the nuclear hormone receptor superfamily, PPARα can be attenuated by insulin in hepatocytes [34]. In our model, elevated circulating insulin may contribute to the downregulation of hepatic PPARα and associated reduction in BHB levels indicating reduced fatty acid utilization, a hypothesis supported by others [35]. Thirdly, upregulated LPL expression may contribute to hepatic steatosis by increasing intracellular free fatty acid accumulation through hydrolysis of lipoprotein triglyceride [36,37]. Interestingly, it has been shown that fructose can induce hepatic steatosis via fructokinase-mediated fructose phosphorylation [38,39]. However, in the current study fructokinase was downregulated in high-fat high-fructose diet group, suggesting high-fat, rather than high-fructose component of the maternal diet

Table 2. Maternal plasma profile.

| Groups | Effect | Diet | Taurine | Interaction |
|--------|--------|------|---------|-------------|
| CON    |        |      |         |             |
| CT     |        |      |         |             |
| MO     |        |      |         |             |
| MOT    |        |      |         |             |
| Taurine (umol/l) | 328±23  | 436±40 | 341±20 | 415±32 |
| Glucose (mmol/l) | 6.11±0.49 | 7.0±0.40 | 8.2±0.56 | 7.79±0.30 |
| BHB (mmol/l)    | 1.09±0.12 | 0.91±0.11 | 0.73±0.09 | 0.64±0.04 |
| Uric acid (µM)  | 2.983±0.454 | 3.797±0.593 | 2.636±0.812 | 1.845±0.301 |
| Leptin (ng/ml)  | 1.420±0.129 | 1.611±0.176 | 1.994±0.161 | 1.671±0.150 |
| Insulin (ng/ml) | 1.247±0.139 | 1.728±0.381 | 2.665±0.402 | 2.980±0.564 |
| TNF-α (µg/l)   | 0.111±0.008 | 0.129±0.013 | 0.155±0.017 | 0.102±0.006 |
| IL-1β (µg/l)   | 0.716±0.006 | 0.718±0.006 | 0.718±0.006 | 0.718±0.006 |
| IL-6 (µg/l)    | 0.172±0.013 | 0.165±0.016 | 0.141±0.017 | 0.185±0.019 |

Values are presented as mean ± SEM, n=7–9 per group. Bold font indicates effect P value <0.05 via two-way ANOVA, main effects were not indicated when a significant interaction was determined. Comparisons between groups for each significant effect where applicable are denoted by 3 sets of superscripts. Upper case letter (A,B) superscripts indicate comparison procedures were conducted between all groups fed MO diet and all groups fed CON diet. Upper case letter (A,B) superscripts indicate comparison procedures were conducted between groups with taurine supplementation and groups without taurine supplementation. Lower case letter superscripts (a,b,c) indicate multiple comparison procedures were conducted for diet and taurine interaction. Groups that do not share the same letter are significantly different from each other (p<0.05).

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represented the predominant contributor to the maternal hepatic steatosis.

Contrary to obesity studies in non-pregnant animals [40–42], maternal diet-induced hepatic steatosis was not reversed, but further aggravated by taurine supplementation. The steatosis observed in the present study represents an instance of non-alcoholic fatty liver disease (NAFLD) which is a chronic condition which is characterized by two distinct phases [43]. Firstly, hyperglycemia and IR contribute to ectopic fat deposition in the liver contributing to intracellular lipid accumulation in hepatocytes increasing oxidative stress and proinflammatory cytokines production which initiate the second phase characterized by hepatocyte apoptosis and progressive fibrosis. In the current study, exacerbated hepatic steatosis in MOT group was evidenced by an elevated hepatic immunophenotype which included upregulated TNF-α and IL-1β expression. It is well established, that TNF-α and IL-1β expression is correlated to the severity of steatohepatitis [44–46]. Increasing free fatty acid accumulation promotes this

Figure 3. Maternal liver histology. Left column, H&E, 10x magnification; right column 40x magnification. CON: control, n = 9; CT: control with taurine, n = 7; MO: maternal obesogenic diet, n = 8; MOT: MO maternal obesogenic diet with taurine, n = 7. Arrows indicate ballooned hepatocytes. doi:10.1371/journal.pone.0076961.g003
proinflammatory phenotype through activation of the TLR4 signaling pathway which culminates in NF-κB activation [17]. In our study, in response to taurine supplementation, maternal hepatic CD36 expression was markedly increased. CD36 is a long chain fatty acid transporter [48] which can directly bind free fatty acid from plasma [37]. Increased fatty acid uptake via CD36, together with potential de novo lipogenesis resulting from increased FASN expression in MOT mothers may contribute to intracellular hepatic fatty acid accumulation. Therefore, this ectopic fatty acid deposition may exacerbate inflammatory processes triggering NAFLD in taurine supplemented high-fat fed mothers.

Although hepatic proinflammatory profile deteriorated in the MOT group, circulating TNF-α was significantly reduced. Maternal taurine supplementation has been previously shown to down-regulate systemic inflammation in acute trauma, sepsis and other immune deficient conditions in non-pregnant subjects [49]. Adipose tissue represents a major contributor to circulating TNF-α concentrations [50]. In the present study, we observed reduced plasma TNF-α concentrations in response to maternal taurine supplementation concomitant with reduced adiposity in the MOT mothers which is in agreement with previous observations whereby weight loss is associated with improvements in inflammatory markers [51].

We noted a reduction in maternal weight gain in taurine supplemented control and MO dams in the last stage of gestation. This was not reflected in an overall change in birth weights or a change in maternal water or food intake. There is evidence for a role of taurine in reducing fat mass in the rodent [52,53] and in the present study we observed a reduction in fat mass in MOT dams but this was at the time of lactation; whether taurine had an effect on modifying maternal body composition in late pregnancy is not known.

Given the link between high fat diets and changes in glutamate metabolism [54], we also examine circulating glutamate levels in the lactating dams. Glutamate at high levels is well known to be toxic to the central nervous system [55], retinal neurons [56] and pancreatic islets [57]. As the fetal blood brain barrier is relatively permeable, even a slight increase in circulating glutamate has been shown to overstimulate neurons in the arcuate nucleus that can lead to metabolic dysregulation in later life [58]. Given the ability of glutamate to cross the placenta [59], it is possible that the elevated maternal glutamate levels observed in the present study may lead to significant adverse effects on later metabolic function in offspring. Taurine supplementation normalized circulating glutamate levels; previous studies have demonstrated that tissue specific levels of glutamate can be decreased by taurine supplementation. This may be due to taurine-mediated alterations in calcium influx via cysteine/glutamate antiport systems [60–62] and may be a mechanism by which beneficial effects of taurine are exerted in the present study but warrants further investigation. Of note in the present study was the sex-specific effects of maternal taurine supplementation on outcomes in the neonate. We have recently reported on sex-specific differences in placental weights in a model of maternal fructose intake [63] and it is possible that the sex-specific effects observed in offspring in the current study may in part be mediated by alterations in, for example, glutamate transfer across the placenta. In a recent paper by Tang et al. in the setting of a maternal low protein diet, it was shown that maternal taurine supplementation led to a reduction in insulin sensitivity in female but not male offspring although the mechanisms underpinning this sex dimorphism are not clear [25].

Offspring hepatic inflammatory profile was significantly rescued as a result of maternal taurine supplementation. Data from the present study indicates that maternal obeseogenic diet had adverse effects on neonatal hepatic inflammatory profiles. Increased expression of IL-1β and TNF-α receptors in MO offspring suggest that maternal developmental programming induces a predisposition to hepatic inflammatory responses which may contribute to long term risk of hepatic IR, steatosis and fibrosis. Notably, a recent study by Chiappini et al. demonstrated the importance of IL-1R1 overexpression in response to early development of obesity-induced NAFLD [64]. Furthermore, in other studies, IL-1-R1 and IL-1β knockout mice display attenuated hepatic steatosis and inflammation when exposed to alcohol and high cholesterol diet [65,66]. This suggests that reversal of IL-1-R1 and TNFR1 in neonates from the MOT group may contribute to an overall improvement of long term metabolic health. We also examined parameters relating to IR and lipid metabolism (Figure S1); however these were unaffected in neonates which is unsurprising given the age and lack of metabolic challenge in these animals. Nevertheless, inflammation is a key pathological factor which we have shown can exacerbate hepatic steatosis. Therefore, the finding that maternal taurine supplementation reversed maternal diet-induced inflammatory receptor overexpression in neonates would suggest that taurine is having a protective effect on offspring liver inflammatory response and may confer protection against fatty liver disease in later life.

Additionally, we observed that in the control pregnancies, taurine supplementation increased neonatal mortality, despite no such an effect in the maternal obeseogenic groups. Animal studies have demonstrated that taurine can prevent mortality in STD-induced diabetic adult rats [67]. However, there are limited data on possible adverse effects of taurine in normal pregnancies. Earlier work by Boujendar et al. reported adverse effects of taurine supplementation in offspring of control pregnancies as reflected in fetal hypoglycaemia and decreased pancreatic and postnatal body weights [21]. We did not observe any significant changes in neonatal glucose concentrations in the present study but there was a significant reduction in birthweight in female but not male offspring of taurine supplemented dams. Taurine supplementation in vivo has been reported to protect islets in offspring from low protein fed dams from cytokine toxicity, but increase islet sensitivity to cytokines and impair pancreatic development in control animals [21,22]. Therefore, possible taurine toxicity in the setting of normal pregnancy outcomes should be further investigated.

In conclusion, a maternal obeseogenic diet-induced postpartum impaired insulin sensitivity, hepatic steatosis and contributed to a programmed neonatal hepatic inflammatory profile. Maternal taurine supplementation exacerbated maternal hepatic steatosis

### Table 3. Maternal liver scoring.

| Groups          | CON | CT  | MO  | MOT |
|-----------------|-----|-----|-----|-----|
| Steatosis       | 0   | 0   | 1.28±0.18a | 1.29±0.18a |
| Lobular inflammmation | 0.3±0.15b | 0.2±0.10 | 1.88±0.30a | 1.43±0.20a |
| Hepatocyte ballooning | 0±0 | 0±0 | 1.38±0.18a | 1.14±0.14a |
| NAS score       | 0.3±0.15b | 0±0 | 4.63±0.53a | 3.86±0.34a |

Values are presented as means ± SEM, n=7–9 per group. Pairwise comparisons between groups were conducted via Wilcoxon rank sum test with Bonferroni correction, groups that did not share the same letter are significantly different from each other (p<0.05).

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Figure 4. Maternal hepatic lipid and glucose metabolism related gene expression. Values are presented as mean ± SEM, n = 7–9 per group. Bold font indicates effect P value <0.05 via two-way ANOVA, main effects were not indicated when significant interaction was determined. Comparisons between groups for each effect where significant are denoted by 3 sets of superscripts. Upper case letter (A, B) superscripts indicate comparison procedures were conducted between all groups fed MO diet and all groups fed CON diet. Upper case letter (A1, B1) superscripts indicate comparison procedures were conducted between groups with taurine supplementation and groups without taurine supplementation. Lower case letter superscripts (a, b, c) indicate multiple comparison procedures were conducted for diet and taurine interaction. Groups that do not share the same letter are significantly different from each other (p < 0.05).

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Figure 5. Maternal hepatic inflammatory gene expression. (a) TNFα; (b) IL-1β; (c) IL-1R1; (d) TNFR1. Lower case letter superscripts (a,b,c) indicate multiple comparison procedures were conducted for diet and taurine interaction. Groups that do not share the same letter are significantly different from each other (p<0.05). Data are means ± SEM, n = 7–9 per group.
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Table 4. Neonatal plasma profile.

| Groups | Leptin (ng/ml) | Insulin (ng/ml) | Glucose (mmol/l) | BHB (mmol/l) |
|--------|----------------|-----------------|------------------|------------|
|        | CON | CT | MO | MOT | CON | CT | MO | MOT | CON | CT | MO | MOT | CON | CT | MO |
| Females |     |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| Leptin (ng/ml) | 1.505±0.304 | 1.147±0.440 | 0.763±0.277 | 0.816±0.221 | F = 3.477 | F = 1.720 | F = 1.709 |
| P = 0.075 | F = 0.203 | F = 0.204 |
| Insulin (ng/ml) | 1.81±0.47 | 0.93±0.25 | 1.08±0.37 | 0.86±0.23 | F = 1.082 | F = 1.980 | F = 0.744 |
| P = 0.309 | F = 0.172 | F = 0.397 |
| Glucose (mmol/l) | 4.413±0.176 | 3.854±0.396 | 4.493±0.383 | 4.433±0.290 | F = 1.176 | F = 1.032 | F = 0.670 |
| P = 0.282 | F = 0.313 | F = 0.416 |
| BHB (mmol/l) | 2.15±0.128a | 1.455±0.225b | 1.693±0.174b | 1.733±0.117b | F = 0.322 | F = 4.370 | F = 5.502 |
| P = 0.572 | F = 0.040 | F = 0.022 |

| Males | Leptin (ng/ml) | Insulin (ng/ml) | Glucose (mmol/l) | BHB (mmol/l) |
|--------|----------------|-----------------|------------------|------------|
|        | CON | CT | MO | MOT | CON | CT | MO | MOT | CON | CT | MO | MOT | CON | CT | MO |
| Leptin (ng/ml) | 1.182±0.242 | 0.527±0.219 | 0.449±0.112 | 0.469±0.129 | F = 3.618 | F = 2.337 | F = 2.631 |
| P = 0.070 | F = 0.140 | F = 0.118 |
| Insulin (ng/ml) | 0.70±0.22 | 0.54±0.25 | 0.70±0.26 | 0.45±0.16 | F = 0.291 | F = 0.685 | F = 0.0372 |
| P = 0.595 | F = 0.416 | F = 0.849 |
| Glucose (mmol/l) | 4.684±0.214 | 5.043±0.244 | 4.994±0.128 | 4.581±0.230 | F = 0.094 | F = 0.012 | F = 2.449 |
| P = 0.760 | F = 0.912 | F = 0.122 |
| BHB (mmol/l) | 2.052±0.082a | 1.414±0.063b | 1.322±0.080b | 1.757±0.106b | F = 3.237 | F = 0.889 | F = 24.877 |
| P = 0.076 | F = 0.349 | F < 0.001 |

Values are presented as mean ± SEM, n = 7–9 per group. Bold font indicates effect P value <0.05 via two-way ANOVA, main effects were not indicated when a significant interaction was determined. Lower case letter superscripts (a,b,c) indicate multiple comparison procedures were conducted for diet and taurine interaction. Groups that do not share the same letter are significantly different from each other (p<0.05).
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yet benefited the circulating proinflammatory profile in dams and reversed the detrimental neonatal hepatic inflammatory cytokine receptor expression. Our findings suggest that maternal taurine supplementation may protect the offspring of obese mothers against developmentally programmed NAFLD despite worsening maternal postpartum fatty liver disease. While a few other studies investigated reversing the programming effects in offspring of undernutrition models [20–22,68–70], our study first reports the reversing effect in a model of maternal obesity. Together, these studies indicate that the early period of life may be a critical window for reversing programming effects. However, while taurine supplementation during pregnancy may modify developmental programming of metabolic dysfunction in offspring, adverse maternal effects in normal pregnancies warrant caution and must be further investigated.

Supporting Information

Figure S1

Table S1 Details of primers used in gene expression analysis. (DOCX)

Table S2 Maternal gene expression main effects. (DOCX)

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Author Contributions

Conceived and designed the experiments: MHV ML DMS. Performed the experiments: ML MHV DMS CR CG. Analyzed the data: MHV ML CR. Contributed reagents/materials/analysis tools: MHV ML CR DMS CG. Wrote the paper: ML MHV CR CG.

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