Melatonin Increases Fetal Weight in Wild-Type Mice but Not in Mouse Models of Fetal Growth Restriction

Lewis J. Renshall1,2, Hannah L. Morgan1,2, Hymke Moens1,2, David Cansfield1,2, Sarah L. Finn-Sell1,2, Teresa Tropea1,2, Elizabeth C. Cottrell1,2, Susan Greenwood1,2, Colin P. Sibley1,2, Mark Wareing1,2 and Mark R. Dilworth1,2*

1 Maternal and Fetal Health Research Centre, Division of Developmental Biology and Medicine, Faculty of Biology, Medicine and Health, University of Manchester, Manchester, United Kingdom, 2 Manchester Academic Health Science Centre, Manchester University NHS Foundation Trust, St. Mary’s Hospital, Manchester, United Kingdom

INTRODUCTION

Fetal growth restriction (FGR), the inability of a fetus to achieve its genetic growth potential, affects between 5 and 10% of pregnancies and is a major risk factor for stillbirth (Miller et al., 2008). In addition, FGR infants that survive at higher risk of childhood morbidities such as cerebral palsy (Jarvis et al., 2003). There are also well-established correlations between being born small and an increased risk of a host of adulthood diseases including hypertension, diabetes, and stroke (Barker and Osmond, 1988; Barker et al., 1990; Osmond et al., 2007). Whilst there are many pathologies underlying FGR, the majority of cases are due to placental dysfunction (Baschat et al., 2007).
Despite these significant antenatal and postnatal consequences of FGR, there is no treatment. Current clinical management results in early delivery of the baby which is itself associated with poor fetal and neonatal outcomes (Bernstein et al., 2000; Resnik, 2002). Thus, the need for an effective treatment for FGR remains paramount. This lack of therapies for FGR is compounded by the fact that there is a reluctance to design drugs specifically for obstetric conditions (Fisk and Atun, 2008). Thus, there has been a drive for the re-purposing of therapeutics, licensed for use in other clinical diseases, which may be of translational benefit to obstetric medicine.

In order for potential treatments for FGR to progress toward clinical trials there is a need for well-characterized in vivo models of FGR in which to carry out pre-clinical efficacy testing. Two such models of FGR are the endothelial nitric oxide synthase knockout (eNOS\(^{-/-}\)) mouse and the placental specific insulin-like growth factor 2 knockout mouse (P0\(^{+/+}\)), which have been extensively characterized by ourselves and others (Shesely et al., 1996; Constancia et al., 2002; Dilworth et al., 2010, 2011; Kusinski et al., 2011, 2012; Kulandavelu et al., 2012, 2013; Stanley et al., 2012a). eNOS catalyses the conversion of L-arginine to nitric oxide (NO), a potent vasodilator that acts via smooth muscle cells (Moncada et al., 1991). Deletion of eNOS in mice (eNOS\(^{-/-}\)) results in hypertension, also maintained during pregnancy (Shesely et al., 1996; Kusinski et al., 2012), and growth restricted fetuses that are 10–15% smaller than wild-type (WT) fetuses near term. In common with some cases of human FGR, uterine and umbilical artery blood flow velocity is reduced in eNOS\(^{-/-}\) mice compared with WT (Toal et al., 2008; Ghosh and Gudmundsson, 2009; McCowan et al., 2010) whilst uterine arteries of eNOS\(^{-/-}\) mice exhibit oxidative stress (Stanley et al., 2012a) and reduced system A amino acid transport (Kusinski et al., 2012), akin to human FGR (Glawzier et al., 1996; Biri et al., 2007; Shibata et al., 2008; Mert et al., 2012).

The P0\(^{+/+}\) mouse, in which a placental specific promoter of Igf2 is deleted, is a model of late-onset FGR, with fetuses approximately 20% smaller than WT near term (Constancia et al., 2002; Dilworth et al., 2010, 2011). Placentas of P0\(^{+/+}\) mice also demonstrate a reduction in weight versus WT and this reduction precedes the onset of FGR (Constancia et al., 2002). P0\(^{+/+}\) placentas demonstrate aberrant placental morphology, in common with human FGR, with reduced surface area and increased thickness of the interhemal membrane, both of which contribute to a reduction in permeability to hydrophilic solutes (Mayhew et al., 2003; Sibley et al., 2004). Placental nutrient transfer is also aberrant in P0\(^{+/+}\) mice (Constancia et al., 2002, 2005; Dilworth et al., 2010). We have previously demonstrated that uterine and umbilical artery blood flow velocity in P0 mice\(^{+/+}\) is indistinguishable from WT mice (Dilworth et al., 2013). Together, eNOS\(^{-/-}\) and P0\(^{+/+}\) mice represent abnormal blood flow and defective placental transport phenotypes, respectively, that underlie placental dysfunction in FGR, thus providing suitable models in which to test potential therapeutics.

One such candidate therapeutic is melatonin, a naturally synthesized neurohormone primarily produced by the pineal gland and important in the establishment of circadian rhythms. Melatonin is a powerful antioxidant and in a rat model of FGR, in which dams were undernourished, maternal melatonin treatment normalized birth weights of growth restricted pups (Richter et al., 2009) with associated increases in the level of antioxidant enzymes including catalase and manganese-superoxide dismutase (SOD2) within the placenta. In a separate study, where FGR was induced by lipopolysaccharide (LPS) administration in mice, melatonin normalized fetal weights via reductions in placental oxidative stress and hypoxia (Chen et al., 2006). In pregnant sheep, intravenous infusion of melatonin increased umbilical artery blood flow via NO dependent mechanisms (Thakor et al., 2010).

eNOS\(^{-/-}\) and P0\(^{+/+}\) knockout mice provide two models that map onto differing phenotypes of placental dysfunction. The use of eNOS\(^{-/-}\) mice is akin to those FGR cases that present with abnormal uterine artery blood flow and endothelial dysfunction. The choice of this model also allows a discrete assessment of the importance of eNOS, via its deletion, in the effectiveness of melatonin in increasing fetal growth. The use of the P0\(^{+/+}\) mouse enables an assessment of melatonin in a model of late-onset FGR akin to women that do not present with abnormalities in uteroplacental blood flow but demonstrate placental pathology as evidenced by abnormal morphology and nutrient transfer. We thus tested the hypothesis that antenatal melatonin supplementation would increase fetal growth in the eNOS\(^{-/-}\) and P0\(^{+/+}\) mouse models of FGR, each representing a different placental pathology associated with FGR.

**MATERIALS AND METHODS**

**Animals**

This study was carried out in accordance with the recommendations of the UK Animals (Scientific Procedures) Act of 1986 under Home Office licenses PPL 40/3385 and P9755892D. The protocols were approved by the Local Animal Welfare and Ethical Review Board (AWERB) of the University of Manchester.

Endothelial NO synthase knockout mouse (eNOS\(^{-/-}\)), stock number 002684, were purchased from Jackson Laboratories (Bar Harbor, ME, United States). C57Bl/6J mice (Envigo, United Kingdom), the background strain for eNOS\(^{-/-}\), were used as control mice (wild-type, WT) for the eNOS\(^{-/-}\) studies. Placental specific insulin-like growth factor 2 knockout mouse (P0\(^{+/+}\)) were a kind gift from Wolf Reik and Miguel Constancia (University of Cambridge) (Constancia et al., 2000). eNOS\(^{-/-}\) female mice were mated with eNOS\(^{-/-}\) male mice. WT female mice were mated with WT males. For P0 matings, P0\(^{+/+}\) male mice were mated with C57Bl/6J female mice which resulted in mixed litters of P0\(^{+/+}\) and WT (P0\(^{+/+}\), control) fetuses. All animals were provided with nesting material and housed in
individually ventilated cages maintained under a constant 12 h light/dark cycle at 21–23°C with free access to food (BK001 diet, Special Dietary Services, United Kingdom) and water (Hydropac, Lab products Inc, Seaford, DE, United States).

On E12.5, animals were randomly assigned, using an online blocked randomization tool, into either control or treated groups. Researchers were not blinded to treatment group. Both treated and control groups were assessed concurrently. Those mice in the treated group were given 5 µg/ml melatonin (Sigma-Aldrich, United Kingdom) via drinking water. This dose was chosen according to another rodent study showing beneficial effects on fetal/birth weight (Richter et al., 2009). This study resulted in plasma concentrations of melatonin equivalent to those following the use of melatonin for jet lag (Herxheimer and Petrie, 2002) and similar to the proposed dose for the phase 1 pilot clinical trial in an FGR cohort (Alers et al., 2013). Due to poor solubility in water, melatonin was initially dissolved in 100% ethanol. The final concentration of ethanol in the drinking water was 0.05%. Mice were dosed until E18.5 with a fresh bottle made up at E15.5. Control animals had access to 0.05% ethanol in standard drinking water (vehicle). All dosing, including bottle changes, and harvesting of tissue took place in the morning between 8 am and 12 pm.

In total, 21 WT and 18 eNOS−/− mice were placed on vehicle, and 21 WT and 20 eNOS−/− mice were placed on melatonin treatment. For P0+/+ studies, 13 mice were placed on vehicle and 14 mice placed on melatonin. P0−/− mice produce mixed litters of P0+/+ and WT pups (referred to as P0+/+ to distinguish from WT controls used as a comparison for eNOS−/− mice). All mice were humanely euthanased at E18.5. Fetal and placental weights were recorded from all litters following weighing on a Mettler AC100 analytical balance (Mettler-Toledo, Leicester, United Kingdom). Fetal biometric measures (crown:rump length, abdominal circumference and head circumference) were taken as previously described and treatment, followed by Bonferroni’s post-test to compare differences between individual groups. For uterine and umbilical artery wire myography, a two-way ANOVA was used to test for differences between groups. For urine and umbilical artery wire myography, a two-way ANOVA was used to test for differences between groups. For uterine and umbilical artery wire myography, a two-way ANOVA was used to test for differences between groups. For uterine and umbilical artery wire myography, a two-way ANOVA was used to test for differences between groups.
RESULTS

Fluid Consumption

Following commencement of melatonin supplementation, amount of water drunk (with either melatonin or vehicle added) was measured until E18.5. There was no difference in amount of fluid drunk (ml/day) according to either genotype or treatment in eNOS−/− and WT mice (WT vehicle 7.6 ± 0.4, WT melatonin 6.6 ± 0.2, eNOS−/− vehicle 7.2 ± 0.5, eNOS−/− melatonin 6.8 ± 0.4, two-way ANOVA). There was no difference in amount of fluid drunk between treatment groups in dams carrying P0+/+ and P0+/− (6.7 ± 0.3 vehicle group versus 7.1 ± 0.5 in the melatonin treated group, Mann–Whitney test).

Fetal and Placental Weights and Fetal Biometric Measurements

**eNOS−/− Mice**

Litter sizes (mean ± SEM) were as follows; WT vehicle 7.1 ± 0.4, WT melatonin 7.0 ± 0.4, eNOS−/− vehicle 6.9 ± 0.4, eNOS−/− melatonin 7.0 ± 0.4. Litter size was not significantly affected by either genotype or melatonin treatment. Fetal and placental weights are shown in Figure 1. Fetal weight (Figure 1A) was significantly lower in eNOS−/− versus WT, independent of treatment (P < 0.001). Additionally, melatonin significantly increased fetal weight in WT mice (mean fetal weight 1.15 ± 0.02 g in WT vehicle group versus 1.20 ± 0.01 g in WT melatonin, P < 0.05) but not in eNOS−/− mice (0.99 ± 0.02 g in eNOS−/− versus 1.01 ± 0.01 g in eNOS−/− melatonin). To explore this effect on fetal weight in further detail, fetal weight distribution curves (Figure 1B) were constructed. The curves for eNOS−/− mice treated with vehicle and melatonin were similar, but the curve for WT mice treated with melatonin was shifted to the right of the curve for WT mice receiving vehicle. In this WT vehicle treated group, 0.968 g and 1.007 g were representative of the 5th and 10th fetal weight centiles, respectively. When assessing the percentage of fetuses with weights falling below the 5th centile of WT vehicle treated mice, it was observed that 2% of WT melatonin treated, 30% of eNOS−/− vehicle and 30% of eNOS−/− melatonin treated fetuses fell below this 5th weight centile. When assessing the percentage of fetuses weighing below the 10th centile of WT mice treated with vehicle, typically defined as the small for gestational age threshold in humans, there was a statistically significant reduction in the percentage of melatonin treated fetuses (3%) below this 10th centile value compared with WT vehicle (P < 0.01, chi-squared test). There was no difference in the percentage of fetuses weighing below the 10th centile of the WT vehicle treated population between eNOS−/− mice receiving melatonin versus eNOS−/− treated with vehicle (45% versus 47%, respectively). Placental weights (Figure 1C) were significantly elevated in eNOS−/− versus WT mice, independent of treatment (P < 0.01). Melatonin treatment had no significant effect on placental weight in either eNOS−/− or WT mice. The fetal:placental weight ratio (Figure 1D) was significantly reduced in eNOS−/− mice independent of treatment and was unaffected by melatonin treatment in both WT and eNOS−/− mice. Fetal biometric measurements are shown in Table 1. Fetal crown:rump length, abdominal circumference and head circumference were all significantly reduced in eNOS−/− versus WT mice, independent of treatment (P < 0.001); melatonin treatment did not alter crown:rump length or abdominal circumference in either WT or eNOS−/− fetuses but head circumference was increased following melatonin treatment in WT fetuses only (P < 0.01).

**P0+/− Mice**

Litter size (mean ± SEM) was comparable between vehicle (8.0 ± 0.3) and melatonin treated (8.0 ± 0.4) mice. Fetal and placental weights are presented in Figure 2. Fetal weight was significantly reduced in P0+/− mice compared with P0+/+ controls (P < 0.0001) with this effect being independent of treatment group (Figure 2A). Melatonin supplementation had no significant effect on fetal weight in either genotype. Fetal weight distribution curves appeared comparable between P0+/− treated and untreated groups (Figure 2B) but there was a suggestion that melatonin supplementation had shifted the P0+/− curve to the left. In order to explore this further, we assessed the number of fetuses falling below the 5th and 10th centile of P0+/− fetal weights. Values for the 5th and 10th centiles were 1.02 g and 1.06 g, respectively, with 4% and 7% of the P0+/− melatonin population falling below these values, not significantly different from the P0+/+ controls. Placental weights were significantly reduced in P0+/− versus P0+/+ mice (P < 0.0001, Figure 2C) and this effect was independent of treatment group. Melatonin supplementation did not alter placental weight in either P0+/− or P0+/+ mice. Fetal-placental weight ratio (Figure 2D) was significantly increased in P0+/− mice independent of treatment group (P < 0.0001) but melatonin supplementation did not alter F:P ratio in either P0+/− or P0+/+ mice. Crown:rump lengths, abdominal and head circumferences are shown in Table 2. There was no effect of melatonin treatment on crown-rump length in P0+/+ fetuses. Crown:rump length in P0+/− fetuses supplemented with melatonin in utero also failed to reach statistical significance (P = 0.06). Melatonin supplementation increased abdominal circumference in P0+/− mice (P < 0.05) but there was no effect on P0+/− fetuses. Genotype, independent of treatment, had no significant effect on head circumference; melatonin supplementation did not alter head circumference in either P0+/− or P0+/+ fetuses.

**Ex vivo Uterine Artery Function**

**eNOS−/− Mice**

Uterine artery diameters (μM, mean ± SEM) were as follows; WT vehicle 211 ± 17, WT melatonin 217 ± 25, eNOS−/− vehicle 147 ± 13, eNOS−/− melatonin 182 ± 25. Diameters were reduced in eNOS−/− versus WT, independent of treatment group (P < 0.05, two-way ANOVA). There was no difference in uterine artery diameter following melatonin treatment in either WT or eNOS−/− mice. Maximal constriction to 10^{-5} M PE was greater in uterine arteries of eNOS−/− versus WT mice, independent of treatment group (P < 0.05). Melatonin treatment had no effect on constriction in either WT or eNOS−/− mice (Figure 3A). ACh caused significant
relaxation of uterine arteries from WT mice following pre-constriction with PE; this relaxation was significantly blunted in arteries of eNOS−/− mice independent of treatment \( (P < 0.0001) \). Melatonin had no significant effect on uterine artery relaxation to Ach in either WT or eNOS−/− mice (Figure 3B).

**PO0+/− Mice**

Uterine artery diameters (μM, mean ± SEM) were not significantly different between vehicle (253 ± 19) and melatonin treated dams (238 ± 11). Uterine arteries from melatonin treated mice demonstrated greater maximal constriction to PE compared with the vehicle group \( (P < 0.05, \text{Figure 3C}) \). There was no

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**TABLE 1** | Fetal biometric measures at E18.5 in WT and eNOS−/− mice.

|                         | WT vehicle | WT melatonin | eNOS−/− vehicle | eNOS−/− melatonin |
|-------------------------|------------|--------------|-----------------|------------------|
| Crown:rump length (mm)  | 28.2 ± 0.5 | 28.4 ± 0.5   | 26.8 ± 0.7a,b   | 26.6 ± 0.6a,b    |
| Abdominal circumference (mm) | 25.8 ± 0.3 | 26.0 ± 0.3   | 24.0 ± 0.5a,b   | 23.9 ± 0.3a,b    |
| Head circumference (mm)  | 25.0 ± 0.2 | 24.7 ± 0.3   | 24.4 ± 0.4a,b   | 24.2 ± 0.3a,b    |

Number of litters as follows; WT vehicle \( N = 18 \), WT melatonin \( N = 20 \), eNOS−/− vehicle \( N = 13 \), eNOS−/− melatonin \( N = 17 \). \( aP < 0.001 \) compared with WT vehicle, \( bP < 0.001 \) compared with WT melatonin, generalized linear mixed models with sequential Sidak post-test.
significant difference in relaxation of uterine arteries to Ach (as % of constriction to PE) between vehicle and melatonin treated mice (Figure 3D).

**Ex vivo Umbilical Artery Function**

**eNOS−/− Mice**

Umbilical artery diameters (µM, mean ± SEM) were as follows; WT vehicle 474 ± 26, WT melatonin 455 ± 38, eNOS−/− vehicle 468 ± 25, eNOS−/− melatonin 416 ± 17. There was no significant effect of genotype or treatment on umbilical artery diameter. There was no effect of genotype or treatment on constriction of umbilical arteries to U46619 (Figure 4A). Relaxation of umbilical arteries to SNP (as % EC80 U46619 preconstriction) was genotype-dependent with eNOS−/− mice, independent of treatment group, showing increased relaxation to SNP versus WT (P < 0.05). Melatonin treatment had no effect on umbilical artery relaxation to SNP (Figure 4B).

**P0+/− Mice**

Umbilical artery diameters (µM, mean ± SEM) were as follows; P0+/− vehicle 499 ± 31, P0+/− melatonin 492 ± 36, P0+/− vehicle 499 ± 21, P0+/− melatonin 470 ± 32. There was no significant effect of genotype or treatment on umbilical

| TABLE 2 | Fetal biometric measures at E18.5 in P0+/+ and P0+/− mice. |
|----------|-------------------|-------------------|-------------------|-------------------|
|          | P0+/+ vehicle     | P0+/+ melatonin   | P0+/− vehicle     | P0+/− melatonin   |
| Crown:rump length (mm) | 28.8 ± 0.3        | 28.6 ± 0.2        | 27.4 ± 0.3ab      | 26.7 ± 0.3ab      |
| Abdominal circumference (mm) | 26.4 ± 0.5        | 26.9 ± 0.4        | 23.6 ± 0.4ab      | 24.8 ± 0.3bc      |
| Head circumference (mm) | 25.1 ± 0.4        | 25.1 ± 0.1        | 24.8 ± 0.3        | 24.9 ± 0.2        |

Number of litters as follows; P0+/+ vehicle N = 10, P0+/+ melatonin N = 11, P0+/− vehicle N = 10, P0+/− melatonin N = 11. *P < 0.001 compared with P0+/+ vehicle, **P < 0.001 compared with P0+/− melatonin, #P < 0.05 compared with P0+/− vehicle, generalized linear mixed models with sequential Sidak post-test.
Constriction of umbilical arteries to U46619 was not significantly different between any groups (Figure 5A). Relaxation of umbilical arteries to SNP (expressed as a % EC_{50} U46619 preconstriction, Figure 5B) was not significantly affected by genotype alone but there was a treatment effect. Post hoc tests revealed that umbilical arteries from P0^{+/-} melatonin mice demonstrated significantly increased relaxation to SNP than P0^{+/-} treated by vehicle alone. Additionally, P0^{+/-} mice showed greater relaxation to SNP compared with P0^{+/-} mice in the vehicle and melatonin groups, though this effect was limited to an SNP dose of 10^{-7} M only.

**DISCUSSION**

Contrary to our hypothesis, melatonin failed to increase fetal weight in eNOS^{−/−} or P0^{+/-} mice, two mouse models of FGR which demonstrate different underlying placental pathology, but did increase fetal weight in WT mice. Melatonin supplementation in utero did, however, result in increased fetal abdominal circumference in P0^{+/-} mice. Despite the lack of a vascular phenotype in P0^{+/-} mice, melatonin supplementation increased relaxation of P0^{+/-} umbilical arteries to the NO-donor sodium nitroprusside. Although the mechanisms underpinning the increased fetal weight in WT mice and increased abdominal circumference in P0^{+/-} mice remain to be fully understood; the lack of any effect in eNOS^{−/−} mice suggest that the effects of melatonin are mediated, at least in part, via the presence of eNOS, suggestive of a potential role for NO. In the case of WT mice, melatonin has growth promoting effects even in the absence of a uterine or umbilical blood flow abnormality. Similarly, the data in P0^{+/-} mice suggests that melatonin may alter fetal growth, at least in terms of increased abdominal circumference, in cases of FGR in which there is no detectable uterine or umbilical blood flow abnormality. Consideration of these mechanisms is especially important as a phase 1 pilot clinical
trial to assess the effectiveness of melatonin in pregnancies complicated by severe early onset FGR has commenced (Alers et al., 2013).

Fetal weights in untreated WT and eNOS−/− mice were comparable with previous reports and confirmed the FGR phenotype of eNOS−/− mice at E18.5 (Stanley et al., 2012a; Poudel et al., 2013) whilst the reduced eNOS−/− fetal/placental weight ratio confirmed previous observations suggesting a reduced transport efficiency of the eNOS−/− placenta (Kusinski et al., 2012; Stanley et al., 2012a; Poudel et al., 2013). Fetal and placental weights in P0+/− mice were comparable to previous reports (Constancia et al., 2002; Dilworth et al., 2010, 2011). Previous studies have suggested that melatonin increased fetal weight in a mouse model of LPS-induced FGR (Chen et al., 2006) and FGR following single umbilical artery ligation in sheep (Tare et al., 2014). Additionally, in an undernourished rat model of FGR (Richter et al., 2009), melatonin increased birth weight but this effect was not apparent prior to delivery. In these studies, the effect of increased fetal/birth weight was observed in the FGR groups only, as opposed to the present study where the effect was limited to WT mice. As noted above, this may be explained by a requirement for the presence of eNOS in the mechanism underpinning this increased fetal growth. This is supported by studies showing that the production of NO via NOS enzymes is important in terms of melatonin’s actions of increasing umbilical blood flow in a sheep model of FGR (Thakor et al., 2010). It has also been demonstrated that, following single umbilical artery ligation

FIGURE 4 Constriction and relaxation responses of umbilical arteries from WT and eNOS−/− mice at embryonic day 18.5. (A) Dose-dependent constriction to 10−10 to 2 × 10−6 M U46619 shown as a percentage of the maximum constriction to KPSS. (B) Dose dependent relaxation to sodium nitroprusside (SNP) is shown as a percentage of EC80 U46619 constriction. Data are presented as mean ± SEM. N’s as follows (N = litters, n = pups); WT vehicle N = 5, n = 20, WT melatonin N = 5, n = 17 eNOS−/− vehicle N = 5, n = 12, eNOS−/− melatonin N = 5, n = 17. Statistical analysis was carried out by two-way ANOVA to assess the effect of genotype and treatment with Bonferroni post-test to compare individual groups. ∗P < 0.05 WT vs. eNOS−/−.

FIGURE 5 Constriction and relaxation responses of umbilical arteries from P0+/− and P0+/− mice at embryonic day 18.5. (A) Dose-dependent constriction to 10−10 to 2 × 10−6 M U46619 shown as a percentage of the maximum constriction to KPSS. (B) Dose dependent relaxation to sodium nitroprusside (SNP) is shown as a percentage of EC80 U46619 constriction. Data are presented as mean ± SEM. N’s as follows (N = litters, n = pups); P0+/− vehicle N = 7, n = 12, P0+/− melatonin N = 7, n = 12, P0+/− melatonin N = 7, n = 9. Statistical analysis was carried out by two-way ANOVA to assess the effect of genotype and treatment with Bonferroni post-test to compare individual groups. ∗P < 0.05. #P < 0.05 P0+/− melatonin vs. P0+/− vehicle and P0+/− melatonin at 10−7 M.
in sheep, melatonin treatment in utero rescued endothelial-dependent coronary artery function in neonatal FGR lambs by increasing NO bioavailability (Tare et al., 2014). Additionally, melatonin alters fetal cardiometabolic responses in fetal lambs exposed to acute hypoxia in utero, including reductions in fetal arterial blood pressure. These effects were prevented via a NO blockade, providing further evidence of melatonin's effects in terms of increasing NO bioavailability (Thakor et al., 2015).

The increased fetal growth seen here in appropriately grown WT fetuses following maternal melatonin supplementation has been observed before. In sheep, melatonin administered in utero increased fetal ponderal index and abdominal girth, with a trend toward increased fetal weight (Lemley et al., 2012), in lambs that were appropriately grown, with no effect on nutrient-restricted FGR lambs. This increased abdominal girth, or circumference, was noted in the present study in P0+/− pups treated with melatonin but crown:rump length was unaffected ($P = 0.06$). Whilst umbilical blood flow velocity was not measured in the current study, the greater relaxation of umbilical arteries in melatonin treated P0+/− mice is consistent with the increased abdominal circumference (Scorza et al., 1991; Stanley et al., 2012b). Studies investigating other potential therapies for FGR, including sildenafil citrate, have previously reported an increased fetal abdominal circumference growth velocity following treatment (von Dadelszen et al., 2011) and outlined the importance of serial abdominal circumference measures, via ultrasound, as a means of assessing growth velocity before and after treatment. Interestingly, given the increased fetal weight in WT mice following administration of melatonin, one may expect a similar increase in P0+/+ mice, the WT equivalent in a mixed P0 knockout mouse litter. However, melatonin did not alter P0+/+ fetal weight. Whilst the reasons for this can only be speculated upon at this time, it is apparent that P0+/− mice treated with vehicle had a higher mean fetal weight (1.20 ± 0.02 g) than WT vehicle treated mice (1.15 ± 0.02 g) and a weight comparable to WT mice treated with melatonin (1.20 ± 0.02 g). This is presumably because on average, P0 knockout litters consist of 50% growth restricted P0+/− fetuses and there will likely be a relative surplus of maternal nutrients available to enable maximal growth of P0+/+ pups. This phenomenon is similar to the inverse relationship described between litter size and fetal weight in late gestation in the mouse (Ishikawa et al., 2006) which demonstrates that maternal nutrient availability is a limiting factor for fetal growth. It has been reported that melatonin alters utero-placental amino acid flux and increases fetal uptake of branched chain amino acids in a sheep model of FGR following maternal undernutrition (Lemley et al., 2013). Thus, future studies examining materno-fetal amino acid transfer in melatonin treated WT mice may hint toward a potential mechanism for the increased fetal growth observed.

Melatonin treatment in utero increased relaxation of umbilical arteries of P0+/− mice in response to SNP, a NO donor, demonstrating an increased sensitivity that is endothelial-independent. This gives further credence to the notion that NO is important in the actions of melatonin. A recent study in sheep fed a nutrient restricted diet, demonstrated that cotyledonary arteries (described as secondary branches of the umbilical artery) from those sheep on the restricted diet demonstrated increased sensitivity to SNP which was subsequently reversed by melatonin supplementation (Shukla et al., 2014). In the same study, ewes on a normal diet demonstrated that the opposite was true, i.e.; melatonin increased sensitivity to SNP in cotyledonary arteries. These latter data appear to fit with the P0 umbilical data presented here.

In a number of studies, there is direct evidence of melatonin's antioxidant properties. In a study by Chen et al. (2006), melatonin supplementation reduced fetal mortality and increased fetal weight in a model of intrauterine death following LPS administration. This was suggested to be due to a significant reduction in LPS-induced lipid peroxidation and normalization of the LPS-induced decrease in placental glutathione (Chen et al., 2006; Wang et al., 2011). Richter et al. (2009) demonstrated that, in a rat model of induced FGR following maternal undernutrition, melatonin increased fetal weight with an accompanying increase in levels of placental catalase and manganese-superoxide dismutase (MnSOD, also known as SOD2). In the current study, only WT mice demonstrated an increase in fetal weight following melatonin supplementation. WT mice, and thus WT placentas, will not be subject to excess levels of oxidative stress and thus it is unlikely that the effect of melatonin on WT fetal growth is via antioxidant mechanisms. Thus, the mechanism underpinning this increased WT fetal weight remains elusive and future studies examining this present a logical next step.

Whilst the studies discussed above relate to a number of different species, with melatonin administered by a variety of routes (via drinking water, subcutaneous, i.v. infusion), there is a body of evidence that supports the fact that melatonin treatment in utero is able to increase fetal growth (Chen et al., 2006; Richter et al., 2009; Lemley et al., 2012; Tare et al., 2014). However, in the current study, melatonin failed to increase fetal weight in either of two mouse models of FGR each of which demonstrates different placental pathologies observed in cases of human FGR (McCowan et al., 1988; Mayhew et al., 2003; Toal et al., 2008; Gbosh and Gudmundsson, 2009). It is important to note that whilst melatonin did increase fetal weight in WT mice, this is not representative of the severe FGR cases that would be targeted clinically for intervention. The current study does, however, add weight to the importance of NO bioavailability in the actions of melatonin, in particular by targeting a mouse model with a specific deletion of eNOS. Delineating the pathways by which melatonin does act to increase fetal growth will be paramount, particularly as the ability to stratify cases of FGR improves (Audette and Kingdom, 2017; Gaccioli et al., 2017).

**AUTHOR CONTRIBUTIONS**

LR, HLM, HM, DC, SF-S, TT, and EC performed the research. SG and CS contributed to the conception and design of the
work. MW and MD contributed to the conception and design of the work and performed the research. All authors were involved in drafting the paper.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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