Magnetic Resonance Imaging Detects Placental Hypoxia and Acidosis in Mouse Models of Perturbed Pregnancies

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Introduction

The placenta is central in the aetiology of preeclampsia, a complication of pregnancy characterised by gestational hypertension and proteinuria and a leading cause of morbidity and mortality in both mothers and infants. It has been postulated that reduced placental perfusion as a result of abnormal placental implantation is the initiating event that leads to the maternal syndrome [1]. More recently, endothelial dysfunction has emerged as the proximal cause for the maternal clinical symptoms, with increased levels of soluble fms-like tyrosine kinase-1 (sFlt-1) secreted by the placenta interfering with the bioavailability of vascular endothelial growth factor (VEGF) and VEGF signalling across the maternal endothelium leading to the maternal hypertensive response [2]. The question however remains concerning the cause of the dysregulation of sFlt-1. Although placental ischaemia and oxidative stress have been shown to have a significant role it remains unclear whether this is an outcome or a cause of structural changes in the placenta and how this links with release of the anti-angiogenic molecules.

Placental vascular visualisation has been difficult and limited to mainly ultrasound Doppler in live animals or recently in vivo multiphoton microscopy [3]. Magnetic Resonance Imaging (MRI) studies of placental anatomy and perfusion have been conducted in humans [4–6], mice [7–9] and rats [10]. While dynamic contrast-enhanced MRI, using injected contrast agents, has yielded estimates of mean blood flow in the placenta of animals [11] non-invasive techniques such as arterial spin labelling, diffusion imaging and measurements of T1 (spin-lattice or longitudinal) and T2 (spin-spin or transverse) relaxation times have been investigated to provide alternative safe techniques for assessing human placental structure and function [12].

A major source of image contrast in MRI studies performed without the use of contrast agents comes from the inherent variation in relaxation times between tissues, as well as contributions from proton density, diffusion and flow. Relaxation results from local fluctuations in magnetic fields, which in turn result from predominantly reorientational and to a lesser extent translational motion of the species containing the nuclear spins. T1 and T2 relaxation times are tissue specific with T1 relaxation times tending to be longest when the protons are “bound” to macromolecules, shorter when they are “free” in solution and shortest when they are in intermediate “structured” states. T2 relaxation times tend to be longest where protons are “free” in solution and shortest when “bound” to macromolecules [13].

In previous studies of T1 and T2 relaxation times, human placenta has been reported as appearing homogeneous, with no internal morphology apparent [18–20]. These studies involved field strengths of 0.5 or 1.5 Tesla and have shown a correlation of...
$T_1$ and $T_2$ relaxation times with gestational age and a trend for shorter $T_1$ and $T_2$ times in pregnancies complicated by preeclampsia and fetal growth restriction [19].

In this study we have used much higher field strength (11.74 Tesla) to investigate whether structural heterogeneities in the placenta could be discerned by $T_2$ mapping and to determine if perfusion is a determinant of any observed differences in $T_2$ relaxation times. Further we have investigated whether $T_2$ mapping is capable of detecting changes in morphology or perfusion in two mouse models of perturbed pregnancy. The reduced uterine perfusion pressure (RUPP) model, where perfusion is deliberately reduced in order to mimic the oxidative and inflammatory stress in the first “stage” of preeclampsia [21–22], and the inflammatory cytokine imbalance (TNF-$\alpha$ infusion) model [23–24] were utilized in order to enable the examination of links between structural abnormalities in the placenta, altered perfusion, and the downstream effects that lead to the maternal hypertensive response.

Methods

Ethics Statement

All procedures were approved by the University of Western Sydney Animal Care and Ethics Committee (Animal Research Authority #A6668) and follow the “Guidelines to Promote the Wellbeing of Animals used for Scientific Purposes” as laid out by the National Health and Medical Research Council of Australia. Animals were monitored twice daily post surgical interventions and additional analgesia was provided if required. Animals (dams and fetuses) were euthanized by cervical dislocation prior to tissue collection.

Animals

C57BL/6JArc mice were obtained from the Animal Resource Centre (Canning Vale, WA, Aus) and were housed in a temperature controlled room in individually ventilated cages (up to 5 per cage), maintained in a 12:12-h light-dark cycle with ad libitum access to water and standard rodent chow. Animals were time-mated and on day 13.5 of gestation were randomly assigned to either RUPP ($n=3$), TNF-$\alpha$ infusion ($n=3$) or normal pregnant ($n=3$), sham surgery ($n=2$) or saline infusion ($n=2$) control groups and housed individually until termination of the experiment. Animals were subject to MRI on day 17.5 of gestation unless otherwise indicated.

Reduced utero-placental perfusion pressure (RUPP) procedure

The RUPP procedure has been described in both rats [21] and baboons [22] to induce placental ischaemia and hypertension. For the study undertaken here the procedure was altered to perform a unilateral ligation of the right uterine artery only as ligation of both branches of uterine artery and the lower abdominal aorta resulted in a high rate of abortion in the mice. Briefly, a silk suture was tied around the right uterine artery proximal to the ovarian artery on day 13.5 of gestation. Normal pregnant mice underwent a sham surgery on the same day of pregnancy. Surgical anaesthesia was induced with a subcutaneous implantation of a mini-osmotic pump (Model 1007/D, Alzet, Cupertino, CA) designed to deliver a constant release of TNF-$\alpha$ (500 ng/kg/day). Briefly, an incision was made into the skin below the right scapulae on day 13.5 of gestation and the mini-osmotic pump primed with TNF-$\alpha$ was inserted into a subcutaneous pocket. Control animals received a saline implant. Anaesthesia and analgesia was as described for RUPP surgery.

Magnetic Resonance Imaging (MRI)

$^1$H MRI images were taken of anaesthetised mice placed in a vertical animal probe using a Bruker Avance 11.74 Tesla wide-bore spectrometer with micro-imaging probe capable of generating gradients of 0.45 T/m. Anaesthesia was induced with 4% isoflurane in a chamber before the animals were transferred supine to the animal imaging probe. Isoflurane (lowered to 2%) was continuously delivered via a nose cone at an air flow of 150 ml/min. A small collar was used to maintain their head in a vertical position during scanning. A pressure sensitive pillow was taped to their abdomen to monitor respiration, the mice were wrapped for insulation and the animal chamber of the imaging probe was maintained at around 28 °C. The probe was inserted vertically into the scanner and the isoflurane concentration reduced to 1.5–1.7% and titrated to a respiration rate of approximately 50–60 breaths per minute. Sequence acquisition was gated on respiration (Model 1025 Small Animal Monitoring and Gating System, SA Instruments Inc, Stony Brook, NY, USA) in order to reduce motion artifacts. A Gradient Echo (GEI) sequence protocol was used to obtain a series of localising images across the abdominal region. Thirty contiguous 1 mm slices, with an in plane resolution of 0.25 mm in either axial or coronal plane and high resolution images (in plane resolution of 0.12 mm) were taken of selected slices. $T_2$ measurements using the same geometry were also acquired using a Multi Slice Multi Echo (MSME) sequence protocol (Bruker MSME-$T_2$-map) with a 10 ms echo time and an in-plane resolution of 0.1–0.2 mm. MATLAB (The Mathworks, Natick, MA, USA) was used to generate $R_2$ ($1/T_2$) maps from the acquired data using non-linear least squares regression using the Levenberg-Marquardt-Fletcher algorithm. For quantification, $T_2$ values were calculated from three points in each selected region of interest within 2–5 individual placentas. $T_1$ maps were also produced for some placenta using MATLAB from data acquired using the TrueFISP sequence in FID mode with a flip angle of 5 degrees, 16 frames and a repetition time of 5 ms [25]. Additional $T_2$ measurements were acquired on the same slices of one normal pregnant mouse immediately after blood flow was reduced to zero by terminal anaesthesia.

Histology

Following MRI at gestational day 17.5, animals were euthanased, placentas collected, rinsed in PBS and fixed for 24 h in 10% formalin (Sigma) at 4 °C and processed into paraffin blocks prior to sectioning into 5 μm slices. Antigen retrieval was performed using proteinase K for 7 min (Dako Aust Campbellfield, Vic, Aus) and sections were immunostained for mouse cytokeratin (pAb Z0622; 1/1000, Dako Aust, Campbellfield, Vic, Aus) using proteinase K for 7 min (Dako Aust Campbellfield, Vic, Aus), visualised with DAB (Envision kit, Dako Aust) and counterstained with haematoxylin. Microscopy was performed with a Zeiss LSM 510 confocal microscope.

Statistics

Statistical analysis was carried out using SPSS software (version 20) (SPSS, Inc., Chicago, IL, USA). Generalized linear modelling using linear regression was used to evaluate the differences in $T_2$.
values between regions in 5 placentas before and after blood flow. Generalized Estimating Equation Modelling clustering placenta within animals and animals within treatment groups was used to evaluate the differences in $T_2$ values and the $T_{2\text{lab}}/T_{2\text{junc}}$ ratios. Data are expressed as means ± SE with the level of significance being $p<0.05$. A logarithmic transformation was carried out on ratios prior to statistical analysis.

**Results**

**Distinct regions of $T_2$ contrast in murine placenta**

Three distinct morphological regions of contrast based on $T_2$ relaxation times were discerned in the MRI images of normal pregnant mouse placenta at gestational day 17.5 correlating to the labyrinth, junctional and decidual zones (Figure 1A and B). No variation of contrast within areas of the placenta was observed in the $T_1$ images (Figure 1C). $T_2$ mapping did not differentiate placental regions at gestational day 10.5 prior to the establishment of maternal circulation in the mouse placenta [26] (Figure 1D).

**$T_2$ contrast abolished at loss of blood flow**

To determine the contribution of perfusion to the contrast between regions in the $R_2$ ($1/T_2$) map, additional $T_2$ measurements were acquired on the same slices of one normal pregnant mouse immediately after the blood flow was reduced to zero by terminal anaesthesia. Upon cessation of blood flow the difference in $T_2$ contrast between the three regions was substantially reduced (Figure 2). There was a significant decrease in $T_2$ contrast in the labyrinth ($p<0.001$) and a significant increase in the junctional region ($p=0.003$) upon loss of blood flow, whereas the $T_2$ values of the decidual region remained unchanged ($p=0.21$) (Figure 3). The ratio of $T_{2\text{lab}}/T_{2\text{junc}}$ was calculated to further clarify the observed changes, decreasing from $2.56±0.14$ during blood flow to $1.04±0.14$ after blood flow ceased ($p<0.001$).

**Pattern of $T_2$ contrast altered in perturbed pregnancies**

We examined whether morphological differences could be detected by $T_2$ mapping in the placenta of mice subjected to two experimental models of preeclampsia; namely the RUPP model and the inflammatory cytokine imbalance model (TNF-α). Differences in the pattern of the regions of $T_2$ contrast in the placenta were observed between control, RUPP, and TNF-α treated mice (Figure 4). The ratio of $T_{2\text{lab}}/T_{2\text{junc}}$ was significantly altered in RUPP $1.70±0.16$ (n=3, $p=0.001$) and TNF-α treated, $1.74±0.20$ (n=3, $p=0.001$) animals compared to control animals $2.35±0.06$ (n=8) (Figure 5), and there was a trend for larger $T_2$ values in the junctional zone and deciduas of RUPP and TNF-α treated animals (Figure 6). Sham-operated and saline-infused controls were not significantly different to normal pregnant animals and for the purposes of analysis the three control groups were grouped together.

**Discussion**

This study has shown that higher resolution $T_2$ maps of mouse placenta can clearly differentiate between different regions of the
placenta at time points after the maternal circulation is fully established. This differs from previous studies of $T_2$ relaxation times in humans using much lower field strengths, where the placenta was observed to be homogeneous [18,27]. Our findings show that the labyrinth has a $T_2$ relaxation time twice that of the junctional zone, with the observed contrast between these regions of the placenta being abolished on loss of blood flow. The decreased $T_2_{lab}/T_2_{junc}$ ratio on loss of blood flow was due to both a decrease in the $T_2$ value in the labyrinth and an increase in the $T_2$ value in the junctional zone.

Examining both the structure of the placenta and the influences on $T_2$ relaxation times provides an explanation of the differences between regions and of the changes on loss of blood flow. In the labyrinth, essentially an intermeshed network of independent fetal and maternal blood vessels, there is an abundance of both freely moving protons, in the form of free water, and of blood cells containing mostly highly oxygenated haemoglobin. The junctional zone is dense with spongiotrophoblast and giant trophoblast cells. The longer $T_2$ value measured in the labyrinth, compared to the more cellulary dense junctional zone, can be accounted for by the abundance of freely moving protons in this region which give rise to longer $T_2$ relaxation times [13]. Upon cessation of blood flow, the $T_2$ value in the labyrinth decreased and the $T_2$ value in the junctional zone increased. After cessation of blood flow the tissue continues to metabolise for some time, consuming $O_2$, producing $CO_2$ and generating deoxyhaemoglobin. The decrease in the $T_2$ value in the labyrinth can be accounted for by an increase in the paramagnetic ion deoxyhaemoglobin which gives rise to shorter $T_2$ relaxation times [14–15], however in the junctional zone there are few haemoglobin containing blood cells and hence minimal paramagnetic effect of deoxyhaemoglobin on the $T_2$ value in this region. Conversely, due to the build up of $CO_2$ in the tissue and the consequent acidosis, the increase in free protons would account for the observed increase in the $T_2$ value in the junctional zone. The effects of pH on $T_2$ values has been well documented in both muscle tissue of live patients [16] and in isolated muscle [17], with a clear correlation between decrease of intracellular pH and an increase in the $T_2$ relaxation time. While acidosis would also occur in the labyrinth the predominant effect on the $T_2$ value in this region appears to be that of the paramagnetic deoxyhaemoglobin. Thus, the abolition of contrast between regions of the placenta upon complete loss of blood flow is consistent with the effects on $T_2$ relaxation times by both increases in deoxyhaemoglobin (hypoxia) and decreases in intracellular pH (acidosis).

The RUPP and the TNF-$\alpha$ treated mice also showed a decrease in contrast between the labyrinth and junctional zones, though this is primarily due to a trend for an increase in the $T_2$ value in the
implicating intracellular acidosis in the regulation of signalling pathways leading to the downstream effects on the maternal system. Further studies involving the measurement of intracellular pH using $^{31}\text{P}$ MRI would be warranted to confirm the tissue acidosis in these models.

Structural abnormalities in the placenta have been proposed to result in reduced placental perfusion [20] or ischaemia-reperfusion injury [29], and to lead to placental oxidative stress, cellular damage and inflammation. The subsequent release of anti-angiogenic and other toxic compounds into the maternal circulation has been shown to lead to endothelial dysfunction and the maternal hypertensive response in the preeclamptic pregnancy [30]. The signalling pathways leading to the release of the anti-angiogenic molecules are as yet unclear, but may involve both oxygen dependent elements such as hypoxia inducible factor-1 (HIF-1) [31] or Jumonji domain-containing protein 6 (JmjD6) [32], and non oxygen dependent mechanisms such as toll-like receptor-3 (TLR-3) and NFκB pathways [33]. Our data suggests that pH dependent mechanisms may also play a significant role, and may be as equally important as hypoxia in the perturbed placenta. Recently it has been reported that coupling factor 6 (CF6) activation of ectopic ATP synthase leads to increased sFLT-1 through intracellular acidosis induced c-Src signalling [34]. CF6 is found in the circulation, has higher levels in spontaneously hypertensive rats and its release from the surface of endothelial cells is stimulated by TNF-α [34]. Many other studies have shown that acidosis and hypoxia have a linked role in some signalling pathways [35–38] and our data would be in keeping with these findings.

In conclusion, this study has shown that high resolution $T_2$ maps of mouse placenta can distinguish different regions of the placenta and that, upon loss of blood flow, contrast between regions is abolished consistent with effects on $T_2$ relaxation times by both increases in deoxyhaemoglobin (hypoxia) and decreases in intracellular pH (acidosis).

**Author Contributions**

Conceived and designed the experiments: GB AM AH. Performed the experiments: GB LS TS-G. Analyzed the data: GB TS-G WSP JML. Contributed reagents/materials/analysis tools: WSP AH. Wrote the paper: GB TS-G AM JLM WSP AH.

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**Figure 6. Comparison of the $T_2$ values from different regions of the placenta.** Control animals ($n=8$) are shown in red; RUPP animals ($n=3$) in light blue and TNF-α treated animals ($n=3$) in dark blue. $T_2$ values were calculated from 3 points in each of the labyrinth, junctional zone and decidua regions, from 2–5 placentas from each animal at day 17.5 of gestation. doi:10.1371/journal.pone.0059971.g006
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