



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

Fetal Growth in Diabetic Pregnancies

Citation for published version:

Anblagan, D, Deshpande, R, Jones, N, Pitiot, A, Costigan, C, Allcock, K, Raine-Fenning, N, Leach, L, Bugg, G, Mansell, P & Gowland, PA 2012, 'Fetal Growth in Diabetic Pregnancies', The International society for Magnetic Resonance in Medicine, Melbourne, United Kingdom, 5/05/12 - 11/05/12.

Link:

[Link to publication record in Edinburgh Research Explorer](#)

Document Version:

Publisher's PDF, also known as Version of record

General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.



Fetal Growth in Diabetic Pregnancies

Devasuda Anblagan¹, Ruta Deshpande², Nia W Jones², Alain Pitiot³, Carolyn Costigan¹, Kirsty Allcock⁴, Nick Raine Fenning⁵, Lopa Leach⁶, George Bugg², Peter Mansell², and Penny A Gowland¹

¹Sir Peter Mansfield Magnetic Resonance Centre, University of Nottingham, Nottingham, Nottinghamshire, United Kingdom, ²Nottingham University Hospitals NHS Trust, United Kingdom, ³School of Psychology, University of Nottingham, United Kingdom, ⁴Medical Schools and Nursing Centres, University of Nottingham, United Kingdom, ⁵School of Clinical Sciences, University of Nottingham, United Kingdom, ⁶School of Biomedical Sciences, University of Nottingham, United Kingdom

Introduction: Pregnancy complicated by pre-existing or gestational diabetes often results in macrosomic infants (birth weight > 4000 g) with increased body fat, but not increased head or brain size¹. They are at increased risk of shoulder dystocia², brachial plexus injury, prolonged labor and caesarean section. In the long term, the fetuses are at higher risk of becoming obese and developing Type 2 diabetes³. We previously demonstrated MRI measures of fetal fat increase in a small number of diabetic pregnancies at late gestation⁴. We now extend this to a larger group at different gestational ages, and include fetal length measurements.

Aim: To compare measurements of fetal weight, fetal length and fetal fat composition (made using a novel scanning/analysis protocol) in normal and diabetic pregnancies.

Methods: Scanning: 14 diabetic and 12 control pregnant women were recruited with ethics approval; all gave informed consent. They were scanned twice at 22–26 weeks and 33–37 weeks gestational age (GA), using 1.5 T Philips Achieva MRI scanner with 5-element SENSE cardiac coil or 4-element SENSE torso coil, depending on size. Women lay on their right side in the decubitus position to avoid vena caval compression. All scans were conducted with a specific absorption rate of <2.0 W kg⁻¹. Two sequences were acquired in 3 orthogonal blocks of images to allow motion effects to average out. To measure fetal weight and length a Half Fourier Single Shot Turbo Spin-Echo (HASTE: 123 slices in 147 s, 0.78×0.78×6.00 mm³ voxels, TE = 120 ms) and Balanced Fast Field Echo (bFFE, 130 slices in 167 s, TR = 5.8 ms, TE = 2.3 ms, flip angle = 70°, 0.78×0.78×6.00 mm³ voxels) sequences were acquired. To measure fetal adiposity a water suppressed fast field echo (FFE, 12 transverse slices encompassing the whole fetus in 12.4 s, slice gap = 20 mm, TR = 147 ms, TE = 4.6 ms, 1.88×1.86×6.00 mm³ and FOV = 402 mm) sequence was acquired. **Analysis:** A semi-automated segmentation approach based upon pixel intensity was used to quantify the fat volume, taking account of noise and correcting somewhat for partial volume effects; this has previously proved to give reliable estimates^{4,5}. The freehand mask drawn around the fetus on GIMP (2.2.13) and fat images were loaded into MATLAB (R2010a) to compute the intensity histogram (h_j , $j=1 \rightarrow N$; where N is number of bins) within the masked region. Histograms were often bimodal with one peak corresponding to background noise and a second corresponding to fat voxels. The fetal fat volume (V_f) was calculated using:

$$V_f = XY(Z+G) \sum_j h_j S_j \quad \text{and} \quad S_j = (1 - e^{-\frac{(j-2i_p)}{i_p/2}}) \text{ for } 2i_p < j < N \text{ or } S_j = 0 \text{ for } 2i_p < j < N$$

where X, Y, Z are the voxel dimensions, G is the slice gap and S_j is a scaling factor designed to take account of noise and partial volume effects. Total fetal volume was measured on bFFE or HASTE images by drawing a freehand mask around the fetus using Analyze 9.0. Percentage fetal fat was measured from total and fat volumes. The fetal length was found from the sums of the length of the fetal spine (measured in two curved sections), thigh, lower leg and skull height (crown to base of brain stem) on the bFFE and HASTE images using Analyze 9.0; in order to measure the length of the spine the images were resampled obliquely.

Results: Figure 1 shows a fat image from a diabetic pregnancy. At 22–26 weeks very little fetal fat was visible with no significant difference between diabetics and controls. At 33–37 weeks the data showed significant increase in fetal fat volume ($p = 0.006$), percentage fetal fat ($p = 0.009$) and fetal length ($p = 0.017$) in fetuses of diabetic mothers compared with controls (Figure 2). The total fetal volume of fetuses of diabetic mothers is greater than controls, but this difference was not statistically significant. The subjective intra-abdominal fat scores were 0.13 for diabetics and 0.3 for controls at 22–26 weeks, increasing to 1.46 for diabetics and 0.9 for controls at 33–37 weeks.

Discussion: MRI can measure the overall increase in fetal size and the increase in fetal fat fraction in diabetic pregnancies. In utero assessment of fetal size and fat distribution would allow clinicians to identify fetuses at risk of complicated delivery and study the effect of maternal diabetes on metabolic changes in the infant.

References: [1] P.M. Catalano, et al. (1998), *Diabetes Care* 21 Suppl 2 B85-90. [2] D.S. Acker, et al. (1985), *Obstetrics & Gynecology* 66 762-768. [3] S.P. Chauhan et al. (2005), *Am J Obstet Gynecol* 193 332-346. [4] D. Anblagan, *Proc ISMRM* 20 (2011) 4183. [5] D. Anblagan, *Proc ISMRM* 19 (2010) 5910.

FUNDED BY DIABETES UK.

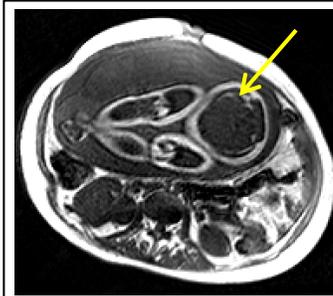


Figure 1: Image of a transverse plane through the fetal thorax showing fat only. Fetal fat surrounding thorax and legs can be seen.

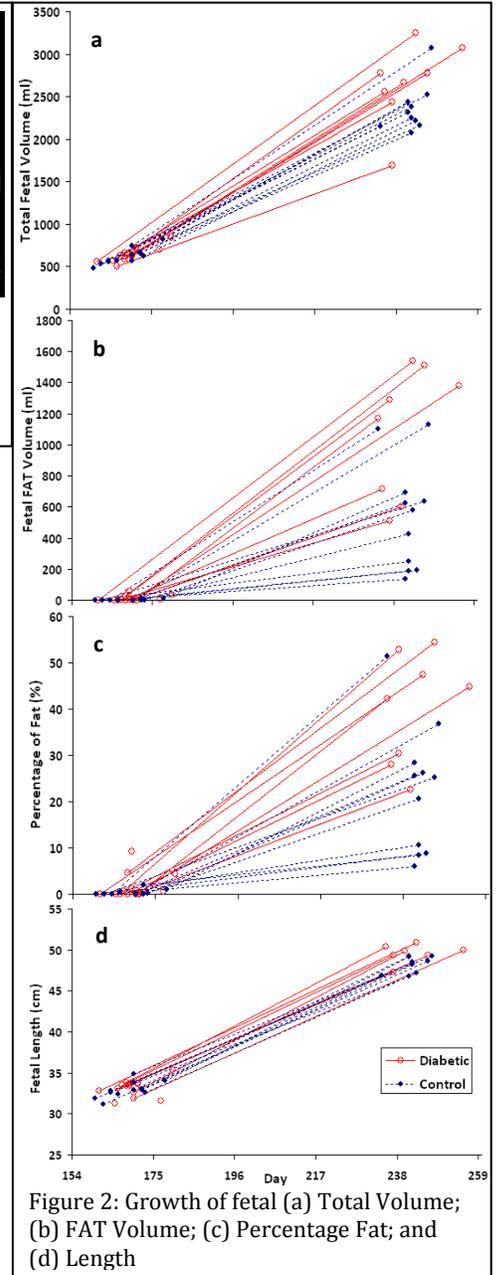


Figure 2: Growth of fetal (a) Total Volume; (b) FAT Volume; (c) Percentage Fat; and (d) Length