

UMass Chan Medical School

eScholarship@UMassChan

UMass Center for Clinical and Translational
Science Research Retreat

2014 UMass Center for Clinical and
Translational Science Research Retreat

May 20th, 12:30 PM

Adipose Tissue Inflammation in Diabetic versus Non-diabetic Pregnancies

Tiffany A. Moore Simas
University of Massachusetts Medical School

Et al.

Let us know how access to this document benefits you.

Follow this and additional works at: https://escholarship.umassmed.edu/cts_retreat



Part of the Cellular and Molecular Physiology Commons, Endocrine System Diseases Commons, Endocrinology Commons, Endocrinology, Diabetes, and Metabolism Commons, Maternal and Child Health Commons, Nutritional and Metabolic Diseases Commons, Obstetrics and Gynecology Commons, Reproductive and Urinary Physiology Commons, Translational Medical Research Commons, and the Women's Health Commons

Repository Citation

Moore Simas TA, Rojas-Rodriguez R, Draper JT, Sert A, Kruse M, Boeras C, Rosenthal B, Gaelikman O, Min S, Leung KG, Corvera S. (2014). Adipose Tissue Inflammation in Diabetic versus Non-diabetic Pregnancies. UMass Center for Clinical and Translational Science Research Retreat. <https://doi.org/10.13028/0ttf-dz76>. Retrieved from https://escholarship.umassmed.edu/cts_retreat/2014/posters/69

Creative Commons License



This work is licensed under a [Creative Commons Attribution-NonCommercial-Share Alike 3.0 License](https://creativecommons.org/licenses/by-nc-sa/3.0/). This material is brought to you by eScholarship@UMassChan. It has been accepted for inclusion in UMass Center for Clinical and Translational Science Research Retreat by an authorized administrator of eScholarship@UMassChan. For more information, please contact Lisa.Palmer@umassmed.edu.

Adipose Tissue Inflammation in Diabetic versus Non-diabetic Pregnancies

Tiffany A. Moore Simas, MD, MPH, MEd
UMass Medical School/UMMHC
508-334-6678
Tiffanya.mooresimas@umassmemorial.org

Tiffany A. Moore Simas, MD, MPH, MEd^{1a}; Raziell Rojas-Rodriguez, BS^{1b}; Jacqueline Tessa Draper, BS¹; Aylin Sert, MEd¹; Maxwell Kruse^{1b}; Crina Boeras, MD^{1a}; Brittany Rosenthal^{1b}; Olga Gealekman, PhD^{1b}; So Yun Min, MSc^{1b}; Kathy Leung, MPH^{1a}; Silvia Corvera, MD^{1b}

¹University of Massachusetts Medical School; ^a Obstetrics & Gynecology, ^b Program in Molecular Medicine

Objective: Beyond weight associated with fetus, placenta, amniotic fluid and increased blood volume, adipose tissue (AT) expansion is an accepted and expected component of pregnancy weight gain. Normal pregnancy is associated with relative insulin resistance (IR). In non-pregnant humans, AT expansion has been associated with IR and AT inflammation. However, it is not known whether AT expansion and IR in pregnancy are also associated with AT inflammation. This pilot study examined relationships between AT expansion and inflammation in control versus diabetic pregnancies.

Methods: Eligible subjects undergoing scheduled Cesarean delivery for obstetric indications were prospectively enrolled. Subjects provided demographic and anthropometric data, and biologic specimens. Immunofluorescence microscopy was performed on subcutaneous (SQ) and omental (OM) AT samples to evaluate macrophage infiltration. Included gravidas had paired AT samples and either negative glucola screening (controls) or gestational or pre-gestational Type 2 diabetes mellitus (T2DM).

Results: 13 subjects with SQ and OM AT samples were evaluated (10-controls, 3-diabetics (2-T2DM and 1-GDM)). Mean BMI and gestational weight gain of controls was 27.8 kg/m² (range 19.5-42) and 27.6 pounds (range 15-36) and of diabetics was 30.6 kg/m² (range 30-33) and 19 pounds (range -3-30), respectively. Macrophage infiltration was seen in OM AT from 2/ 3 diabetics and 0/ 10 controls (see figure).

Conclusions: These results indicate that AT expansion in non-diabetic pregnancies is not accompanied by macrophage infiltration. Thus, the IR of normal pregnancy is unlikely to be related to AT inflammation, and AT expansion per se does not lead to AT inflammation. However, as has been reported for T2DM in non-pregnant humans, the presence (T2DM) or development (GDM) of diabetes in pregnancy is associated with macrophage infiltration of AT. Despite the small sample size, the observed large differences in macrophage infiltration between controls and diabetics suggest that these findings will persist in a larger cohort.

