Relationship between vitamin D and gestational diabetes in overweight or obese pregnant women may be mediated by adiponectin

Aya Mousa¹, Sally K Abell¹, Soulmaz Shorakae¹, Cheryce L Harrison¹, Negar Naderpoor¹, Danielle Hiam², Alba Moreno-Asso², Nigel K Stepto^{1,2}, Helena J Teede¹, Barbora de Courten¹

¹ Monash Centre for Health Research and Implementation (MCHRI), School of Public Health and Preventive Medicine, Monash University; Locked Bag 29, Clayton, VIC 3168, Melbourne, Australia

² Institute of Sport, Exercise and Active Living, Victoria University; PO Box 14428, VIC 8001, Melbourne, Australia

Correspondence:

A/Prof. Barbora de Courten, MD PhD MPH FRACP

Monash Centre for Health Research and Implementation (MCHRI), School of Public health and Preventive Medicine, Monash University

Level 1, 43-51 Kanooka Grove, Clayton,
Melbourne, VIC 3168, Australia

Telephone: +61 3 857 22651

Email: barbora.decourten@monash.edu

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Abbreviations: **1,25(OH)₂D₃**, 1,25-dihydroxyvitamin D₃; **25(OH)D**, 25-hydroxyvitamin D; **BMI**, body mass index; **ELISA**, enzyme linked immunosorbent assays; **GDM**, gestational diabetes mellitus; **HDL/LDL**, high-/ low- density lipoprotein; **HMW**, high-molecular weight; **IL-6**, interleukin-6; **MCP-1**, monocyte-chemoattractant protein-1; **OGTT**, oral glucose tolerance test; **PCOS**, polycystic ovary syndrome; **PTB**, preterm birth; **RCT**, randomized controlled trial; **VDR**, vitamin D receptor.

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ABSTRACT

Scope: Maternal vitamin D deficiency has been implicated in adverse pregnancy outcomes. However, the association between vitamin D and inflammation, particularly adipokines, remains unexplored in pregnancy.

Methods and Results: In 102 overweight or obese pregnant women at high-risk of gestational diabetes (GDM), we investigated relationships between maternal 25-hydroxyvitamin D (25(OH)D) concentrations at 12-15 weeks gestation (baseline) and serum lipids, inflammatory markers, novel adipokines (omentin-1, visfatin, high molecular weight (HMW)-adiponectin), and subsequent pregnancy outcomes (GDM, preeclampsia, preterm birth [PTB]). After adjustment for maternal factors (age, BMI, parity, ethnicity, and smoking status), baseline 25(OH)D concentrations were inversely associated with total cholesterol and triglycerides, and positively associated with HMW-adiponectin. Higher baseline 25(OH)D concentrations were associated with decreased fasting and 1-hour post-OGTT glucose and reduced risk of GDM at 26-28 weeks, as well as with longer gestation and reduced risk of PTB upon additional adjustment for caesarean section. Adding HMW-adiponectin to the multivariable models attenuated most associations, and HMW-adiponectin was a significant predictor in the models.

Conclusion: Our findings suggest that lower maternal 25(OH)D concentrations in overweight/obese pregnant women at high-risk of GDM are associated with increased cardiometabolic risks during pregnancy and adverse pregnancy outcomes, and that these associations may be mediated by HMW-adiponectin.

Graphical Abstract –Text

This study investigated associations between vitamin D concentrations in 102 women at 12-15 weeks gestation and cardiometabolic risk factors including novel adipokines (omentin-1, visfatin, high molecular weight (HMW)-adiponectin) as well as subsequent pregnancy outcomes. Lower maternal vitamin D levels were associated with poorer lipid profiles, suboptimal inflammatory and adipokine profiles, and poorer glucose metabolism later in pregnancy, as well as adverse pregnancy outcomes including gestational diabetes and preterm birth. These associations appeared to be mediated by HMW-adiponectin.

1. Introduction

Vitamin D deficiency is prevalent worldwide, and is common during pregnancy [1]. The primary source of vitamin D is through sun exposure to ultraviolet B radiation. Therefore, increased rates of obesity and sedentary indoor lifestyles, as well as sun avoidance behaviors and use of sunscreen to prevent skin cancer have contributed to the rise in vitamin D deficiency [1]. Moreover, few foods are naturally high in vitamin D or are vitamin D-fortified [1]. Although there is currently no consensus regarding optimal vitamin D levels during pregnancy, the Institute of Medicine [2] and National Institutes of Health [3] suggest that circulating 25-hydroxyvitamin D (25(OH)D) concentrations <50 and <25 nmol/l are considered deficient and severely deficient, respectively. It is therefore concerning that 40–98% of pregnant women globally have 25(OH)D concentrations <50 nmol/l and 15–84% have concentrations <25 nmol/l [1].

Recent evidence suggests that vitamin D deficiency may contribute to unfavorable cardiometabolic profiles during pregnancy and adverse pregnancy outcomes [4, 5]. The nuclear vitamin D receptor (VDR) and metabolizing enzyme, 1α -hydroxylase, are present in the decidua, placenta, ovary, and endometrium [6]. *In vitro* studies have shown that active vitamin D ($1,25(\text{OH})_2\text{D}_3$) has potent anti-inflammatory properties [7], and directly activates transcription of the insulin receptor gene [8], as well as regulating transcription of genes associated with placental invasion, normal implantation, and angiogenesis [9]. Therefore, vitamin D deficiency may lead to increased inflammation and decreased insulin action, which could contribute to increased risk of adverse pregnancy outcomes including gestational diabetes mellitus (GDM), preeclampsia, and preterm birth (PTB). Indeed, observational studies have reported inverse associations between maternal 25(OH)D concentrations and cardiometabolic risk factors during pregnancy, including insulin resistance [10], dysglycemia

[11], dyslipidemia [12], hypertension [13], and inflammation [14, 15]. Prospectively, maternal vitamin D deficiency has been linked to increased risk of GDM [16], pre-eclampsia [17], and PTB [18]; however results are inconsistent [19-21]. Moreover, most existing studies are limited by small sample sizes and lack of adjustment for potential confounders including obesity, parity, ethnicity, and smoking status.

It is proposed that maternal vitamin D deficiency may contribute to increased cardiometabolic risk and adverse pregnancy outcomes via pro-inflammatory pathways [22] and there is growing interest in the potential role of adipokines [23]. Studies have reported associations between vitamin D deficiency and altered adipokine concentrations in patients with type 2 diabetes [24] and polycystic ovary syndrome (PCOS) [25]; however these relationships have not been explored in pregnancy. To our knowledge, no previous studies have examined associations between vitamin D and novel adipokines including visfatin or omentin-1 in pregnancy, and few have examined high-molecular weight (HMW) adiponectin [26]. These adipokines are believed to be involved in the pathophysiology of adverse pregnancy outcomes including GDM [16, 23], thus further investigation of the relationship between vitamin D deficiency and these adipokines in pregnancy is warranted.

We aimed to address existing knowledge gaps by examining: (i) associations between 25(OH)D concentrations and serum lipids, inflammatory markers, and novel adipokines in early pregnancy (12-15 weeks gestation); (ii) associations between 25(OH)D concentrations in early pregnancy (12-15 weeks gestation) and risk of subsequent adverse pregnancy outcomes (GDM at 26-28 weeks gestation, preeclampsia during pregnancy, and PTB at delivery); and (iii) whether these associations may be mediated by circulating adipokine concentrations.

2. Materials and methods

2.1. Study design and participants

This study involved post-hoc analyses of pre-collected bio-banked serum samples from a large randomized controlled trial (RCT) in high risk pregnancies [27]. The RCT aimed to prevent excess gestational weight gain in women identified as high risk for GDM based on a validated risk prediction tool [28]. Methods for the RCT have been reported previously [27]. Briefly, 228 women were recruited from three large tertiary teaching hospitals in Melbourne, Australia (2008-2010). Women were included in the study if they met the following criteria: age >18 years old; with a singleton pregnancy; <15 weeks gestation; overweight (body mass index (BMI) $>25\text{kg/m}^2$ or $>23\text{ kg/m}^2$ if high risk ethnicity [29]) or obese (BMI $\geq 30\text{ kg/m}^2$) at baseline (12-15 weeks gestation); with an increased risk of GDM (score ≥ 3 based on risk prediction tool [28]). Exclusion criteria included a BMI $\geq 45\text{ kg/m}^2$, non-English speaking women, known pre-existing type 1 or type 2 diabetes, or any other chronic medical condition. The study population was well characterized including demographics, antenatal characteristics, and longitudinal evaluation of pregnancy outcomes. Women were randomized to an antenatal lifestyle intervention or a control group. Samples from the control group (n=102), who were assigned to routine antenatal care with standard dietary and lifestyle advice, were analyzed in the current study.

2.2. Data collection

As previously reported [27], demographic characteristics were collected using a baseline questionnaire. BMI was calculated at the baseline visit (12-15 weeks) by a registered nurse independent to the research staff. Fasting venous blood samples were drawn at 12-15 weeks gestation for all participants. Serum was stored in multiple aliquots under sterile conditions at

-80°C. GDM screening was performed at 26-28 weeks gestation with an oral glucose tolerance test (OGTT). GDM was diagnosed based on the Australasian Diabetes in Pregnancy Society (ADIPS 1998) criteria at the time of the study (fasting glucose ≥ 5.5 mmol/l and/or 2-hour glucose ≥ 8.0 mmol/l) [30]. Maternal and obstetric outcomes were obtained from the Birthing Outcomes System database, which is commonly used for research purposes [28, 31] and includes standardized information for routine reporting of perinatal data in Victoria, Australia. Pregnancy outcomes assessed in this study included preeclampsia diagnosed based on the Royal Australia New Zealand College of Obstetricians and Gynecologists guidelines [32], and PTB defined as <37 weeks gestation [32]. The primary study was approved by the Monash Health Research Advisory and Ethics Committee (Project ID: 07216C) and all participants gave written informed consent [27].

2.3. Biochemical analyses

Fasting serum lipids (triglycerides, total cholesterol, and high- and low-density lipoprotein cholesterol [HDL, LDL]) as well as plasma glucose, and 25(OH)D concentrations were analyzed by accredited commercial pathology providers (Melbourne Pathology and Monash Pathology, Melbourne, Australia). Lipids and glucose were analyzed using a Hitachi modular analyzer (Roche Diagnostics, Mannheim, Germany) and 25(OH)D concentrations were measured using direct competitive chemiluminescent immunoassays (CLIA) (DiaSorin Inc., MN, USA) with inter- and intra-assay coefficients of variation (CVs) of $<10\%$ and $<4\%$, respectively. Serum inflammatory biomarkers and adipokines were analyzed using sandwich-type enzyme linked immunosorbent assays (ELISA). All assays were performed in duplicate according to manufacturers' instructions. Pilot studies were performed to calculate dilutions and ensure concentrations fit within the standard curve for each assay. Samples with values outside the standard curve were excluded from analysis for that specific biomarker. Analyte

concentrations were calculated from best-fit standard curves multiplied by the dilution factor to determine target protein concentrations. HMW-adiponectin was measured using the ALPCO ELISA (47-ADPHU-E01; sensitivity= 0.019 ng/ml), which required pre-treatment with protease II to selectively digest low- and middle-molecular weight adiponectin. HMW-adiponectin was measured instead of total adiponectin in this study as it is considered the active form of adiponectin and a more sensitive marker of glucose intolerance and insulin resistance than total adiponectin [33, 34]. Visfatin was measured using the BioVendor ELISA (RAG004R; sensitivity=30 pg/ml), and Omentin-1 was measured using the Millipore ELISA (EZH0MNTN1-29K; sensitivity= 0.23 ng/ml). Interleukin-6 (IL-6) was measured using the Abcam high sensitivity ELISA (ab46042; sensitivity=0.81 pg/ml). Monocyte chemoattractant protein-1 (MCP-1) was measured using the Picokine ELISA (EK0441; sensitivity <1 pg/ml). Inter- and intra-assay CVs for all biomarkers were <10%.

2.4. Statistical analyses

Statistical analyses were performed using Stata Version 12.0 (StatCorp LP, USA). Participant characteristics and clinical and biochemical parameters are presented as mean $\pm$ standard deviation (SD), frequencies (n [%]), or median (interquartile range [IQR]) for non-normally distributed variables, unless otherwise indicated. Variables which were not normally distributed were logarithmically transformed to the base 10 to approximate normality prior to analysis. Univariable associations between 25(OH)D concentrations and continuous and categorical outcomes were assessed using Pearson correlations and simple logistic regression, with results presented as correlation coefficients for continuous outcomes or odds ratios with 95% confidence intervals for binary outcome variables. Differences in mean 25(OH)D concentrations between groups (eg: women with or without GDM) were assessed using independent Student's *t*-tests. Variables which were statistically significant on univariable

analyses ($p < 0.05$) were included as covariates in multiple linear and logistic regression models. In first model, we adjusted for predetermined maternal factors considered to be clinically relevant to the outcomes, including: age, BMI, parity, ethnicity, and smoking status. In the second model, we performed exploratory analyses to further adjust for adipokines which were significantly associated with the outcomes of interest to examine their potential mediatory role. Findings were considered statistically significant at a two-tailed level of $p < 0.05$.

3. Results

3.1. Sample characteristics

Demographic, biochemical, and clinical outcome data are presented in Table 1. The mean age of participants was 31.9 ± 4.5 years (mean $\pm$ SD) (range= 25 - 43 years), with a median BMI of 29 (26 - 34) kg/m^2 (median [IQR]) at baseline (12-15 weeks). Mean 25(OH)D concentration of participants at baseline was 48.0 ± 16.0 nmol/l, with over half ($n = 53$; 52%) being classified as vitamin D-deficient (25(OH)D ≤ 50 nmol/l). At 26-28 weeks gestation, 25 of the 102 women (24.5%) were diagnosed with GDM. Four (3.9%) of the 102 women were diagnosed with preeclampsia during the course of their pregnancy, and six women (5.8%) experienced a PTB (< 37 weeks gestation).

3.2. Cross-sectional associations with serum 25(OH)D concentrations

Results of cross-sectional univariable analyses at baseline are presented in Table 2. Serum 25(OH)D concentrations were not associated with maternal characteristics including age ($p = 0.2$) or BMI ($p = 0.1$) (Table 2) and there were no differences in 25(OH)D concentrations by parity ($p = 0.8$), ethnicity ($p = 0.4$), or smoking status ($p = 0.3$) (data not shown).

At baseline, serum 25(OH)D concentrations were inversely associated with total cholesterol ($r=-0.22$ mmol/l, $p=0.03$), triglycerides ($r=-0.30$ mmol/l, $p=0.003$), and IL-6 concentrations ($r=-0.20$ pg/ml, $p=0.048$), and positively associated with HMW-adiponectin ($r=0.27$ μ g/ml, $p=0.007$). There were no associations between 25(OH)D concentrations and HDL or LDL cholesterol or the other inflammatory markers and adipokines measured including MCP-1, omentin-1, and visfatin (all $p>0.05$; Table 2).

Results of multivariable linear regression analyses are presented in Table 3A. After adjusting for maternal factors age, BMI, parity, ethnicity, and smoking status, 25(OH)D concentrations remained inversely associated with total cholesterol ($\beta=-0.23$, $p=0.04$), and triglycerides ($\beta=-0.28$, $p=0.02$), and positively associated with HMW-adiponectin ($\beta=0.25$, $p=0.02$); however the association with IL-6 was no longer significant ($p=0.6$; Table 3A).

3.3. Longitudinal associations with serum 25(OH)D concentrations

Results of longitudinal univariable analyses are presented in Table 2. Baseline 25(OH)D concentrations (at 12-15 weeks) were associated with glycemic outcomes at 26-28 weeks, including fasting glucose ($r=-0.35$ mmol/l, $p=0.0008$) and 1-hour post-OGTT glucose levels ($r=-0.22$ mmol/l, $p=0.04$), but were not associated with 2-hour post-OGTT glucose levels ($p=0.1$; Table 2). Women who developed GDM at 26-28 weeks ($n=25$) had a significantly lower 25(OH)D concentration at baseline compared with women who did not develop GDM ($n=77$; 42.1 ± 14.7 versus 49.8 ± 16.1 nmol/l, respectively, $p=0.03$). In simple logistic regression, higher 25(OH)D concentrations at baseline were associated with a reduced risk of GDM at 26-28 weeks gestation (OR [95%CI]= 0.97 [0.94-0.99], $p=0.04$; Table 2).

Baseline 25(OH)D concentrations were also positively associated with length of gestation ($r=0.32$ weeks, $p=0.001$; Table 2). A trend was observed for lower baseline 25(OH)D concentrations in women who subsequently had a PTB ($n=6$) compared to women who

reached term (n=96); however this did not reach statistical significance (36.6 versus 48.3 nmol/l, respectively, $p=0.08$). In simple logistic regression, a trend was observed for an association between baseline 25(OH)D concentrations and risk of PTB (OR [95%CI]= 0.95 [0.90-1.00], $p=0.09$). There were no associations between 25(OH)D concentrations and preeclampsia ($p=0.6$; Table 2) or caesarean section ($p=0.7$; Table 2).

After adjustment for maternal factors age, BMI, parity, ethnicity, and smoking status, baseline 25(OH)D concentrations remained inversely associated with fasting and 1-hour post-OGTT glucose levels at 26-28 weeks ($\beta=-0.38$, $p=0.002$, and $\beta=-0.26$, $p=0.02$, respectively; Table 3A). Similarly, baseline 25(OH)D concentrations remained associated with a reduced risk of GDM at 26-28 weeks after adjustment for maternal factors (OR [95%CI]= 0.96 [0.93-0.99], $p=0.02$; Table 3B). The positive association between 25(OH)D concentration and length of gestation also persisted in the multivariable model ($\beta=0.33$, $p=0.005$; Table 3A). A non-significant trend was observed for an association between higher baseline 25(OH)D concentrations and reduced risk of PTB after adjustment for maternal factors ($p=0.06$; data not shown), which became significant after additional adjustment for caesarean section (OR [95%CI]= 0.89 [0.79-0.99], $p=0.03$; Table 3B).

3.4. Exploratory analysis of potential role of adipokines

Results of exploratory analyses are presented in Table 3. HMW-adiponectin was the only adipokine associated with the outcome measures on univariable analyses (all $p<0.05$; data not shown), hence it was added as a covariate in the multivariable models. In cross-sectional analyses, after adjusting for HMW-adiponectin and maternal factors age, BMI, parity, ethnicity and smoking status, the associations between 25(OH)D concentrations and total cholesterol and triglycerides at baseline were no longer significant ($p=0.08$ and $p=0.1$,

respectively; Table 3A). HMW-adiponectin was the only significant predictor in the model for triglycerides at baseline ($p=0.005$).

In longitudinal analyses, baseline 25(OH)D concentrations remained associated with fasting glucose at 26-28 weeks after adjustment for HMW-adiponectin in addition to maternal factors ($\beta=-0.27$, $p=0.03$; Table 3A). However, associations with 1-hour post-OGTT glucose levels as well as GDM risk were attenuated ($p=0.2$ and $p=0.1$, respectively; Table 3). In these models, HMW-adiponectin was the only significant predictor associated with risk of GDM ($p=0.02$), and showed a trend toward a significant association with 1-hour post-OGTT glucose levels ($p=0.08$).

Baseline 25(OH)D concentrations remained associated with length of gestation after adjustment for HMW-adiponectin in addition to maternal factors ($\beta=0.32$, $p=0.01$; Table 3A); however the association with risk of PTB was attenuated ($p=0.09$; Table 3B). HMW-adiponectin was not a significant predictor in the model for PTB ($p=0.3$).

4. Discussion

To the best of our knowledge, this is the first study to investigate the relationships between maternal vitamin D levels and cardiometabolic risk factors and pregnancy outcomes in relation to novel adipokines. We found that in overweight or obese pregnant women at high risk of GDM, 25(OH)D concentrations in early pregnancy were inversely associated with cardiometabolic risk factors and had small but significant inverse associations with risk of adverse pregnancy outcomes including GDM. All relationships remained significant after adjusting for relevant maternal factors. However, most associations were attenuated after adjustment for HMW-adiponectin, suggesting a potential mediatory role for this adipokine.

We found inverse cross-sectional associations between 25(OH)D concentrations and total cholesterol and triglyceride levels at 12-15 weeks gestation, which persisted after adjusting for maternal factors. This is consistent with a previous study where 25(OH)D <75 nmol/l in the first trimester was associated with higher total and LDL cholesterol [35]. In contrast, Al-Ajlan et al. [36] reported that first trimester 25(OH)D concentrations were positively associated with serum lipids, which was thought to be due to increased lipid synthesis in response to high metabolic demands in early pregnancy [36]. Importantly, this study did not adjust for covariates which may have influenced their results. From a mechanistic perspective, vitamin D may act via the VDR to prevent foam cell formation [37], reduce acetylated LDL cholesterol uptake [38] and regulate serum apolipoprotein A-1 levels [39], thereby enhancing cholesterol transport and improving lipid profiles [38-40].

We report that 25(OH)D concentrations were positively associated with HMW-adiponectin, but not with IL-6, MCP-1, omentin-1, or visfatin after adjustment for maternal factors. Inflammation and adipose tissue failure (characterized by altered adipokine concentrations), have been implicated in the pathophysiology of GDM [41], PTB [42], and preeclampsia [43]. Studies show that omentin-1 has insulin-sensitizing effects [44], while visfatin activates the insulin receptor [45] and MCP-1 deficiency ameliorates insulin resistance [46]. Associations have been reported between 25(OH)D and these markers in studies of healthy women [47] and women with PCOS [25], but not in pregnant women. Thus, further investigation of these relationships in pregnancy is warranted. Only two previous studies investigated relationships between 25(OH)D and IL-6 in pregnancy, and both reported no association after adjustment for covariates, consistent with our finding [14, 15]. Similarly, one previous study examined relationships between 25(OH)D and HMW-adiponectin in pregnancy and reported no association [26]. Disagreement with our finding of a positive association may be due to the

smaller sample size [26] (n=36-37 compared to n=102 in our study); the sampling time-point of 31 weeks gestation (since maternal adiponectin levels decline progressively with advancing gestation [48]); and/or the lack of adjustment for ethnicity (since lower total [49] and HMW-adiponectin [50] have been reported in certain ethnic groups, such as South-Asian women [49, 50]). The mechanism by which vitamin D may increase adiponectin is thought to be via suppression of the TNF- α gene and the adipose tissue renin-angiotensin system, both of which regulate adiponectin production [51, 52]. In turn, increased adiponectin may improve insulin sensitivity, lipid profiles, and inflammation, as has been reported in patients with type 2 diabetes, metabolic syndrome, and cardiovascular disease [53, 54].

In longitudinal analyses, early pregnancy 25(OH)D concentrations (12-15 weeks) were inversely associated with fasting and 1-hour post-OGTT glucose levels and had a small but significant inverse association with risk of GDM at 26-28 weeks. It is important to note that women in this study were at high-risk for GDM, hence our results should be interpreted in light of this. Nevertheless, our findings are similar to previous studies where third trimester 25(OH)D concentrations were inversely associated with fasting [10, 11, 22, 55-57] and post-OGTT glucose levels [58] [11, 56]. Four separate meta-analyses [5, 16, 59, 60] of 7-20 observational studies (n=2100-9200), all reported that women with GDM had lower 25(OH)D levels compared with normoglycemic women, and that GDM risk increased by 40–60% in women with vitamin D deficiency (25(OH)D<50 nmol/l).

However, recent large-scale studies not included in these meta-analyses reported no relationship between 25(OH)D and post-OGTT glucose levels [57] or GDM risk [19-21]. Most of these studies [19, 20, 57] involved predominantly non-vitamin D-deficient women (25(OH)D >70-132 nmol/l). This may explain their null findings since relationships between vitamin D and glucose metabolism are thought to be more pronounced in vitamin D-deficient

women [16, 58, 59]. Moreover, most studies examining vitamin D and glycemic outcomes were conducted at 24-38 weeks gestation [5] and <25% had adjusted for ethnicity, parity, or smoking status [5]. Here, we show that early pregnancy 25(OH)D concentrations (12-15 weeks) are associated with fasting and 1-hour glucose and risk of GDM at 26-28 weeks, which persisted after adjusting for these covariates. Vitamin D may reduce GDM risk by elevating intracellular calcium which is vital for β -cell glycolysis and glucose signaling [60], or by acting on the VDR to upregulate the insulin receptor and facilitate basal- and insulin-mediated glucose oxidation and transport [61-65]. Vitamin D may also decrease insulin resistance and GDM risk by downregulating pro-inflammatory cytokines and upregulating anti-inflammatory cytokines [66-68]. While our findings are consistent with most previous literature, we note that our study cohort comprised high-risk women, and that associations between 25(OH)D and GDM risk were small. Large-scale trials are needed to confirm our findings and to establish whether use of vitamin D supplementation may mitigate GDM risk. Early pregnancy 25(OH)D concentrations in our study were associated with increased length of gestation, as well as with reduced risk of PTB in adjusted analyses. Observational studies examining vitamin D and PTB have produced conflicting results [4, 21, 69-72]. Nevertheless, our findings are consistent with a meta-analysis of 10 observational studies (n=10,098) reporting a 1.3 times greater risk of PTB at 25(OH)D<50 nmol/l compared to higher levels [18]. Additionally, we show that 25(OH)D was associated with length of gestation and PTB risk after adjusting for caesarean section and for the combined effects of age, BMI, ethnicity, parity and smoking status, which has not been shown previously [18]. We were unable to explore sub-categories of PTB (very/moderate/late PTB) due to lack of statistical power; however we report on total length of gestation to show that 25(OH)D is related to length of gestation even when a defined PTB cut-off (<37 weeks) is not applied. Mechanistically,

vitamin D deficiency may promote proximal muscle weakness, thus having a possible role in the initiation of early labor [73]. Vitamin D also induces antimicrobial peptides including cathelicidin in placental cells, thereby reducing bacterial infections and infection-induced PTB [18]. The anti-inflammatory properties of vitamin D discussed earlier may also reduce risk of PTB, since chronic inflammation can promote premature membrane rupture and spontaneous labor [74].

Finally, addition of HMW-adiponectin to the multivariable models in our study attenuated most associations, and HMW-adiponectin was the only significant predictor variable in the models for GDM and triglycerides, with a trend toward significance in the model for 1-hour glucose. We speculate that HMW-adiponectin may be a potential mediator of these associations. Human cell studies and animal models show that adiponectin modulates several metabolic processes, and has anti-atherosclerotic, anti-inflammatory, and anti-diabetic properties [48, 75, 76]. Reduced levels of the HMW multimeric form of adiponectin is believed to be pathophysiologically involved in the declining insulin sensitivity that takes place with advancing gestation [48], and has been shown to promote dyslipidemia [53, 54, 77], inflammation [53, 54], beta-cell dysfunction [48], and contribute to the development of both GDM [23, 78] and PTB [79]. Therefore, it is possible that low HMW-adiponectin levels secondary to vitamin D deficiency may contribute to increased cardiometabolic risks and adverse pregnancy outcomes. However, mechanistic and intervention studies are needed to confirm our finding, and to establish whether vitamin D supplementation increases HMW-adiponectin levels and whether this would translate into improved pregnancy outcomes. Our study has several strengths. This was the first study examining associations between vitamin D and novel adipokines in pregnancy and the potential mediatory role of these adipokines in relation to cardiometabolic risk and pregnancy outcomes. The sample

comprised a well-characterized cohort of overweight or obese women at high risk of GDM, which allowed examination of a high-risk group where there was no confounding by comorbidities or medication use. We were able to explore other potential confounding by clinically relevant factors including age, BMI, parity, smoking status, and ethnicity, which have seldom been incorporated in previous studies. Serum 25(OH)D was measured earlier in pregnancy than most previous studies, which enabled assessment of longitudinal relationships between 25(OH)D and pregnancy outcomes. We were able to show that associations between 25(OH)D and cardiometabolic risk and pregnancy outcomes may be mediated by HMW-adiponectin, which has not previously been reported.

Limitations include the observational nature of the study which precludes assessment of causality. Because this was a secondary post-hoc analysis of data collected for a larger study [27], there was no formal power calculation. Thus, our sample size may have been underpowered to detect associations between 25(OH)D and other maternal outcomes including preeclampsia, and may explain the non-significant trends observed initially for PTB. Participants were overweight or obese women at high risk of GDM, thus results may not be generalizable. The single measurement of 25(OH)D at 12-15 weeks gestation meant we could not determine temporal changes in 25(OH)D or account for potential supplement use which may have altered vitamin D status over the course of pregnancy. We were not able to measure 25(OH)D using gold-standard liquid chromatography mass-spectrometry, and instead used Diasorin assays which measure both 25(OH)D₂ and 25(OH)D₃ concentrations. Moreover, because vitamin D is fat-soluble, use of a more sensitive measure of adiposity such as percentage body fat instead of BMI may have yielded different results. Finally, although we adjusted for multiple confounders, we did not obtain data on vitamin D intake

from diet or supplements or on sun exposure, seasonality, or sunscreen use, which may have influenced our results.

In summary, early pregnancy 25(OH)D concentrations in overweight or obese pregnant women at high risk of GDM were inversely associated with cardiometabolic risk factors during pregnancy and had small but significant associations with GDM and PTB, after adjustment for covariates. We showed that these associations may be mediated by HMW-adiponectin; however the exact mechanism by which 25(OH)D may interact with HMW-adiponectin to influence pregnancy outcomes remains unknown. Further clinical trials and mechanistic studies are needed, with inclusion of: (i) adequate power for pregnancy outcomes as primary endpoints; (ii) adjustment for covariates which affect both vitamin D status and cardiometabolic risk; and (iii) frequent sampling to assess whether temporal changes in 25(OH)D over the course of pregnancy are related to different clinical outcomes. Such studies may identify “critical windows” of opportunity for measuring and treating vitamin D deficiency and potentially improving pregnancy outcomes.

5. Acknowledgements

5.1. Author contributions

AM performed data analysis and interpretation and wrote the first draft of the manuscript. HJT and CLH were the lead investigators on the original RCT. SKA, SS, DH and AMA performed laboratory biomarker analyses and contributed to data interpretation and writing and editing the manuscript. NN and NKS contributed to data interpretation and writing and editing the manuscript. BdC, CLH and HJT planned and designed the study and contributed to data interpretation and writing and editing the manuscript. BdC is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the

integrity of the data and the accuracy of the data analysis. All authors meet the ICMJE criteria for authorship, and have approved the final version of the manuscript.

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5.3. Conflict of interest statement: There are no conflicts of interest to declare.

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Table 1. Sample characteristics:

Variable	<i>n</i>	Mean ± SD or median (interquartile range)
Maternal Characteristics (at baseline: 12 – 15 weeks)		
Age (years)	101	31.9 ± 4.5
BMI (kg/m ²)	97	29 (26 – 34)
Parity, <i>n</i> (%) Primiparous Parous	90	35 (38.9) 55 (61.1)
Ethnicity ^a , <i>n</i> (%) Australian South East and North East Asian Southern and Central Asian Other (African, European, Polynesian)	101	41 (40.6) 14 (13.9) 35 (34.6) 11 (10.9)
Smoker, <i>n</i> (%)	102	7 (6.9)
Family history of diabetes ^b , <i>n</i> (%)	78	35 (44.9)
Serum 25(OH)D (nmol/l)	102	48.0 ± 16.0
Serum lipids and inflammatory markers/ adipokines (at baseline: 12 – 15 weeks)		
Total cholesterol (mmol/l)	102	5.4 ± 0.8
HDL cholesterol (mmol/l)	102	1.7 (1.1 – 1.9)
LDL cholesterol (mmol/l)	102	2.9 ± 0.6
Triglycerides (mmol/l)	102	1.5 (1.2 – 2.1)
IL-6 (pg/ml)	100	2.0 (1.7 – 2.6)
MCP-1 (pg/ml)	102	86.9 (38.4 – 127.9)
HMW-adiponectin (µg/ml)	102	2.4 (1.5 – 3.2)
Visfatin (ng/ml)	97	0.13 (0.07 – 0.20)

Omentin-1 (ng/ml)	99	50.0 (38.0 – 68.0)
<i>Glycemic Outcomes (at OGTT: 26 – 28 weeks)</i>		
Fasting glucose- OGTT^c (mmol/l)	102	4.5 (4.2 – 4.9)
1-hour glucose post- OGTT^c (mmol/l)	102	8.5 ± 2.3
2-hour glucose post^c (mmol/l)	102	6.7 (5.8 – 7.8)
Gestational Diabetes Mellitus^c, n (%)	102	25 (24.5)
<i>Pregnancy Outcomes (during pregnancy/ at delivery)</i>		
Pre-eclampsia^c, n (%)	102	4 (3.9)
Preterm birth (gestation <37 weeks)^c, n (%)	102	6 (5.9)
Caesarean section^c, n (%)	102	25 (24.5)
Length of gestation^c (weeks)	97	39.4 (38.4 – 40.2)

Data are presented as mean ± standard deviation and n (%), or median (interquartile range) for non-normally distributed variables, unless otherwise specified;

^a ethnicity was derived from self-reported country of birth;

^b family history refers to first degree relatives only.

^c all variables measured at 12-15 weeks gestation except OGTT (26-28 weeks), length of gestation, and pregnancy outcomes (during pregnancy/ at delivery);

Abbreviations: 25(OH)D, 25-hydroxyvitamin D; BMI, body mass index; IL-6, interleukin-6; MCP-1, monocyte-chemoattractant protein-1; HMW, high-molecular weight; HDL/LDL, high-/ low- density lipoprotein; OGTT, oral glucose tolerance test; C-section, caesarean section.

Table 2. Univariable associations between 25-hydroxyvitamin D concentrations and cardiometabolic risk factors and pregnancy outcomes:

<i>Variable</i>	<i>n</i>	<i>Correlation coefficient (r) or odds ratio (95% CI)</i>	<i>p</i>
<i>Maternal Characteristics (at baseline: 12 – 15 weeks)</i>			
Age (years)	101	0.13	0.2
BMI (kg/m ²)	97	-0.15	0.1
<i>Serum lipids and inflammatory markers / adipokines (at baseline: 12 – 15 weeks)</i>			
Total cholesterol (mmol/l)	98	-0.22	0.03
HDL cholesterol (mmol/l)	98	0.004	0.9
LDL cholesterol (mmol/l)	98	-0.07	0.5
Triglycerides (mmol/l)	98	-0.30	0.003
IL-6 (pg/ml)	100	-0.20	0.048
MCP-1 (pg/ml)	102	-0.09	0.4
HMW-adiponectin (µg/ml)	102	0.27	0.007
Visfatin (ng/ml)	97	0.03	0.2
Omentin-1 (ng/ml)	99	0.13	0.2
<i>Glycemic outcomes (at OGTT: 26 – 28 weeks)</i>			
Fasting glucose ^a (mmol/l)	89	-0.35	0.0008
1-hour glucose post-OGTT ^a (mmol/l)	88	-0.22	0.04
2-hour glucose post-OGTT ^a (mmol/l)	89	-0.16	0.1
Gestational Diabetes Mellitus ^a	102	0.97 (0.94 – 0.99)	0.04
<i>Pregnancy outcomes (during pregnancy / at delivery)</i>			
Pre-eclampsia ^a	97	1.02 (1.00 – 1.09)	0.6
Length of gestation ^a (weeks)	97	0.32	0.001
Preterm birth ^a (gestation <37 weeks)	97	0.95 (0.90 – 1.00)	0.09
Caesarean section ^a	97	1.00 (0.97 – 1.03)	0.7

Data are presented as correlation coefficients for continuous outcome variables or odds ratios with 95% CI for categorical outcome variables. Non-normally distributed variables were log-transformed to the base 10 to approximate normality prior to analyses;

p= significance of associations with 25-hydroxyvitamin D using Pearson correlations for continuous variables and simple logistic regression for categorical variables;

^a all variables measured at 12-15 weeks gestation except OGTT (26-28 weeks), length of gestation, and pregnancy outcomes (during pregnancy/ at delivery);

Abbreviations: BMI, body mass index; IL-6, interleukin-6; MCP-1, monocyte-chemoattractant protein-1; HMW, high-molecular weight; OGTT, oral glucose tolerance test; HDL/LDL, high-/ low- density lipoprotein; GDM, gestational diabetes mellitus.

Table 3. Multiple linear and logistic regression models for associations between 25-hydroxyvitamin D concentrations and cardiometabolic risk factors and pregnancy outcomes:

3A. Multiple linear regression analysis for continuous outcomes								
Dependent Variable	Model 1				Model 2			
	Standardized β	R ²	<i>t</i>	<i>p</i>	Standardized β	R ²	<i>t</i>	<i>p</i>
Total cholesterol (mmol/l)	-0.23	0.32	-2.08	0.04	-0.20	0.33	-1.77	0.08
HDL cholesterol (mmol/l)	-0.12	0.27	-1.13	0.3				
LDL cholesterol (mmol/l)	-0.03	0.19	-0.26	0.8				
Triglycerides (mmol/l)	-0.28	0.24	-2.49	0.02	-0.18	0.31	-1.55	0.1 ^c
IL-6 (pg/ml)	-0.06	0.38	-0.57	0.6				
MCP-1 (pg/ml)	-0.04	0.10	-0.38	0.7				
HMW-adiponectin (μg/ml)	0.25	0.30	2.32	0.02	-	-	-	-
Visfatin (ng/ml)	0.08	0.10	0.68	0.5				
Omentin-1 (ng/ml)	0.08	0.13	0.63	0.5				
Fasting glucose ^a (mmol/l)	-0.38	0.27	-3.25	0.002	-0.32	0.30	-2.58	0.01

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1-hour glucose post-OGTT^a (mmol/l)	-0.26	0.30	-2.32	0.02	-0.17	0.32	-1.37	0.2
2-hour glucose post-OGTT^a (mmol/l)	-0.21	0.21	-1.77	0.08				
Length of gestation^a (weeks)	0.33	0.16	2.90	0.005	0.32	0.15	2.65	0.01

3B. Multiple logistic regression analysis for categorical outcomes

Dependent Variable	Model 1			Model 2		
	OR (95% CI)	<i>z</i>	<i>p</i>	OR (95% CI)	<i>z</i>	<i>P</i>
Gestational Diabetes Mellitus^a	0.96 (0.93, 0.99)	-2.35	0.02	0.97 (0.93-1.00)	-1.55	0.1 ^c
Pre-eclampsia^a	1.05 (0.93, 1.20)	0.85	0.4			
Preterm birth^a (gestation <37 weeks)	0.89 (0.79, 0.99)	-2.21	0.03 ^b	0.88 (0.77-1.02)	-1.74	0.08
Caesarean section^a	1.00 (0.97, 1.04)	0.29	0.8			

Data are presented as correlation coefficients for continuous outcome variables or odds ratios with 95% CI for categorical outcome variables. Non-normally distributed variables were log-transformed to the base 10 to approximate normality prior to analyses;

p= significance of associations with 25-hydroxyvitamin D using multiple linear regression for continuous variables (3A) and multiple logistic regression or ANCOVA for categorical variables (3B);

^a all variables measured at 12-15 weeks gestation except OGTT (26-28 weeks), length of gestation, and pregnancy outcomes (during pregnancy/ at delivery);

^b preterm birth was additionally adjusted for caesarean section in both model 1 and model 2;

^c denotes that HMW-adiponectin was a significant predictor in the multivariable model (ie: significantly associated with the outcome).

Model 1: linear and logistic regression models adjusted for maternal factors: age, BMI, parity, ethnicity, and smoking status;

Model 2: linear and logistic regression models adjusted for HMW-adiponectin in addition to maternal factors: age, BMI, parity, ethnicity, and smoking status

Abbreviations: HDL/LDL, high-/ low- density lipoprotein; OGTT, oral glucose tolerance test; IL-6, interleukin-6; MCP-1, monocyte-chemoattractant protein-1; HMW, high-molecular weight; C-section, caesarean section.

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Author/s:

Mousa, A;Abell, SK;Shorakae, S;Harrison, CL;Naderpoor, N;Hiam, D;Moreno-Asso, A;Stepto, NK;Teede, HJ;de Courten, B

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