Short Communication

Association between adiposity and inflammatory markers in maternal and fetal blood in a group of Mexican pregnant women

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In the present pilot study, we evaluated the effect of maternal adiposity on the plasma concentration of adipocytokines in pregnant women and their newborns. Twenty patients with term gestations without labour were initially selected by pregestational BMI and then classified into two study groups (n=10 each), according to their median value of adiposity (total body fat). Concentrations of TNF-α, IL-1β, IL-6, leptin and adiponectin in plasma of maternal peripheral blood and fetal cord blood were measured and correlated to maternal adiposity. Maternal adiposity showed a significant negative correlation with fetal adiponectin (r2=0.587, P=0.01) and IL-6 (r2=0.466, P=0.05), a significant positive correlation with maternal leptin (r2=0.527, P=0.02) and no correlation with TNF-α or IL-1β. Adiponectin was higher in fetal plasma than in maternal plasma (P=0.043), but significantly lower in newborns from women with high adiposity than in newborns from women with low adiposity (P=0.040). Our results suggest that fetuses from obese women may be less able to control inflammation, due to lower circulating anti-inflammatory adipocytokines, which could limit their optimal development or even increase the risk of abortion or preterm labour.

Pregnancy: Adiposity: Cytokines: Inflammation: Obesity

Obesity represents one of the major public health problems worldwide. In Mexico alone, it affects almost 70% of people between 30 and 60 years old13.

Obesity directly contributes to an increase in proinflammatory adipocytokines, such as leptin, TNF-α, IL-6 and IL-1β, and a decrease in adiponectin. Alterations in the concentrations of these cytokines are known to result from greater fat mass (adiposity), causing the chronic inflammation associated with type 2 diabetes and CVD, among other complications12–4.

In pregnant women, obesity may exacerbate the chronic inflammation associated with gestation, particularly at term, increasing the mother’s risk of presenting several complications such as gestational diabetes, pre-eclampsia, infection and preterm labour, among others5–7.

Furthermore, high concentrations of proinflammatory cytokines can also affect fetal development by causing major alterations in bronchopulmonary and neurological development and predisposing to a number of childhood and adult obesity-related conditions4,8,9.

However, the impact that greater maternal adiposity might have on the concentration of pro-inflammatory cytokines in fetal circulation is still uncertain. Therefore, the purpose of the present pilot study was to identify the possible associations between maternal adiposity and the concentration of various pro-inflammatory markers at term gestation both in maternal and fetal circulations.

Experimental methods

Patient selection

The present study was conducted according to the guidelines laid down in the Declaration of Helsinki and all procedures...
involving patients were approved by the Internal Research and Ethics Committees of the National Institute of Perinatology in Mexico City. A written informed consent was obtained from all patients.

Twenty women with term pregnancies (>37 weeks of gestation) who delivered by elective caesarean section at the National Institute of Perinatology in Mexico City were included in the present study, together with their newborns. Patients were carefully selected to discard labour, infection, metabolic or autoimmune pathologies.

Patients were programmed for elective caesarean section because of a personal or familial history of pregnancy-related or other complications, according to institutional policies. Most of the patients had a history of pre-eclampsia, abortions and preterm deliveries; however, these were all conditions of previous pregnancies and patients presented no complications during the current gestation.

For the present pilot study, women were initially selected using their pregestational BMI to include ten patients diagnosed as ‘normal’ and ten diagnosed as ‘overweight/obese’ according to the BMI classification of the World Health Organization (10).

The main characteristics of participant women and their newborns are presented in Table 1. No differences were found in maternal age, parity or gestational age. Maternal adiposity was significantly different between groups (as expected). No differences were found in newborn characteristics between groups.

**Anthropometric measurements**

Maternal measurements were made at the time of their admission, 12 h before surgery. The present weight of the participants was measured using a Tanita scale model 1631 (Tanita, Arlington Heights, IL, USA). Height was measured with a Seca 206 instrument (Seca Corporation, Hanover, MD, USA). Tricipital, subcapular and leg skinfolds were measured using a Harpender skinfold caliper (Baty International, West Sussex, UK). Resistance was measured with a bioimpedance system (RJL Systems, Clinton Township, MI, USA) according to the manufacturer’s instructions. All measurements were made by the same standardised person.

Newborn characteristics, including weight, length, cephalic, thoracic and abdominal perimeters, were obtained from the clinical files. Pregestational BMI was calculated using pregestational weight referred by the patients.

**Estimation of maternal adiposity**

Although pregestational BMI was used to initially select the patients, we believe that maternal adiposity (measured at the time of cytokine quantification) would be a much better parameter to relate with the inflammatory environment of term gestation.

Maternal adiposity, defined as total body fat, was estimated for each patient using the equation developed by Villar et al. (11), which considers present weight, body surface, subscapular skinfold, leg skinfold and resistance.

**Biological samples**

Maternal blood (5–10 ml) was drained from the patient’s forearm 12 h before surgery and fetal blood was taken from the umbilical cord vein immediately after the placenta was removed. Samples were collected in 10 ml plastic tubes with heparin, and plasma was separated by centrifuging at room temperature for 15 min at 3000 rpm and stored at −70°C until used. Hb concentrations were determined to ensure equal blood volumes (Table 1).

### Table 1. Characteristics of participant women and their newborns

<table>
<thead>
<tr>
<th></th>
<th>Low adiposity (n 10)</th>
<th>High adiposity (n 10)</th>
<th>P</th>
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</thead>
<tbody>
<tr>
<td><strong>Maternal characteristics</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maternal age (years)</td>
<td>28·5  21–41</td>
<td>30  22–36</td>
<td>0·631</td>
</tr>
<tr>
<td>Height (m)</td>
<td>1·54  1·41–1·65</td>
<td>1·56  1·40–1·65</td>
<td>0·631</td>
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<td>Parity</td>
<td>2·0  1–5</td>
<td>2·5  1–5</td>
<td>0·835</td>
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<tr>
<td>Gestational age (weeks)</td>
<td>38·4  37·3–41·4</td>
<td>38·3  36·6–40·5</td>
<td>0·912</td>
</tr>
<tr>
<td>preBMI (kg/m²)</td>
<td>23·8  21·7–28·5</td>
<td>28·4  22·8–35·2</td>
<td>&lt;0·001*</td>
</tr>
<tr>
<td>Total body fat (kg)†</td>
<td>24·7  17·8–28·6</td>
<td>32·6  29·3–43·5</td>
<td>&lt;0·001*</td>
</tr>
<tr>
<td>Hb (g/l)</td>
<td>130·0  108·0–139·0</td>
<td>124·0  107·0–133·0</td>
<td>0·105</td>
</tr>
<tr>
<td><strong>Newborn characteristics</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sex (n)</td>
<td>Male 6</td>
<td>Female 4</td>
<td>0·656</td>
</tr>
<tr>
<td>Birth weight (g)</td>
<td>3170  2860–3456</td>
<td>2908  2560–3755</td>
<td>0·086</td>
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<td>Birth length (cm)</td>
<td>50·5  48·0–52·0</td>
<td>50·0  47·0–51·0</td>
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<tr>
<td>Cephalic perimeter (cm)</td>
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<td>34·5  32·0–36·5</td>
<td>0·633</td>
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<tr>
<td>Thoracic perimeter (cm)</td>
<td>33·2  31·5–34·0</td>
<td>33·0  31·0–34·5</td>
<td>0·897</td>
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<tr>
<td>Abdominal perimeter (cm)</td>
<td>31·0  29·0–33·0</td>
<td>31·5  30·0–34·5</td>
<td>0·274</td>
</tr>
</tbody>
</table>

* Values were statistically different using Mann–Whitney’s test.
† Total body fat calculated with the equation from Villar et al. (11).
Cytokine quantification

Leptin, adiponectin, TNF-α, IL-6 and IL-1β were chosen because of their known implication in the onset of inflammatory processes, which may affect fetal development and gestational outcome. These cytokines were quantified simultaneously using Fluorokine MAP MultiAnalyte Profiling kit (R&D Systems, Minneapolis, MN, USA) with molecule-specific antibodies in a Bio-Rad system (Bio-Rad, Hercules, CA, USA) following the manufacturer’s protocol. Samples were diluted fourfold as recommended by the manufacturer, and the results were multiplied by the dilution factor.

Detection ranges of the assays were 53.8–43,668.9 pg/ml for leptin, 347.4–255,722.9 pg/ml for adiponectin, 3.8–27,935.5 pg/ml for TNF-α, 4.7–34,875.5 pg/ml for IL-6 and 2.2–15,961.9 pg/ml for IL-1β. Intra-assay coefficients of variance were ≤5% for all cytokines.

Statistical analyses

Maternal adiposity was arbitrarily categorised into two groups according to the median value of total body fat (group 1 ‘low adiposity’ = total body fat < 50 percentile; group 2 ‘high adiposity’ = total body fat > 50 percentile).

Correlations between maternal adiposity (uncategorised) and cytokine concentrations were evaluated with Spearman’s test. Only strong, significant correlations are reported.

Differences in cytokine concentrations were analysed with Mann–Whitney’s test. Differences with \( P \leq 0.05 \) were considered significant.

Differences in sex frequencies of newborns between adiposity groups were analysed with a \( \chi^2 \) test, since fetal sex has been correlated with higher amounts of cytokines.

All statistical analyses were made with Statistical Package for Social Sciences version 12.0 software (Chicago, IL, USA).

Results

No differences were found in maternal age, gestational age, newborn birth weight and length, and newborn cephalic, thoracic or abdominal perimeters between low-adiposity and high-adiposity groups.

Fig. 1. Plasma concentration of cytokines in maternal and fetal blood. (a) TNF-α, (b) IL-6, (c) leptin and (d) adiponectin. Samples were categorised in to ‘low-adiposity’ group (<50 percentile [ ] of total body fat) and ‘high-adiposity’ group (>50 percentile [ ] of total body fat), according to the median value of maternal adiposity. The values represent median concentrations with interquartile ranges. Outlier values are represented by . Differences tested with Mann–Whitney’s test. NS (\( P > 0.05 \)).
Maternal adiposity showed a significant negative correlation with fetal plasma adiponectin ($r = -0.587, P=0.01$) not observed in maternal plasma. Adiponectin concentration was significantly higher in fetal blood than in maternal blood in both groups. Interestingly, this cytokine was significantly lower in fetal blood of newborns from women in the high-adiposity group compared with newborns in the low-adiposity group, a difference not observed in maternal blood (Fig. 1).

IL-6 also showed a significant negative correlation with maternal adiposity ($r = -0.466, P=0.05$). However, its concentration showed no difference between blood (maternal/fetal) or adiposity (low/high) groups (Fig. 1).

As expected, there was a significant positive correlation between maternal adiposity and leptin plasma concentration in maternal peripheral blood ($r 0.527, P=0.02$). This was not observed in fetal blood. Leptin concentrations were not different between the low-adiposity and high-adiposity groups, although they clearly tended to increase in the high-adiposity group. Its concentrations were significantly higher in maternal blood compared with fetal blood (Fig. 1).

Plasma concentration of TNF-α did not correlate with maternal adiposity and was not different between low-adiposity and high-adiposity groups. However, it was significantly higher in fetal blood than in maternal blood (Fig. 1).

IL-1β could not be quantified in any sample, since all of the measurements were below the assay sensitivity limit (0.05 pg/ml).

Even with the small sample size of the present pilot study, some interesting associations between adipocytokines and maternal adiposity could be observed. A larger study, with sample size and power calculated from the results of the present pilot study, would allow us to divide the patients in more groups and further strengthen the associations.

In summary, the present results suggest that, due to lower circulating anti-inflammatory cytokines, fetuses from obese women may be less able to control the inflammatory process (e.g. during intra-uterine infections or even during normal labour), which could limit optimal fetal development or even increase the risk of abortion or preterm labour.

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**References**


