

# BMJ Open Does pre-pregnancy BMI determine blood pressure during pregnancy? A prospective cohort study

Ary I Savitri,<sup>1</sup> Peter Zuithoff,<sup>1</sup> Joyce L Browne,<sup>1</sup> Dwirani Amelia,<sup>2</sup> Mohammad Baharuddin,<sup>2</sup> Diederick E Grobbee,<sup>1</sup> Cuno S P M Uiterwaal<sup>1</sup>

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<sup>1</sup>Julius Center for Health Sciences and Primary Care, Julius Global Health, University Medical Center Utrecht, Utrecht, The Netherlands

<sup>2</sup>Budi Kemuliaan Hospital, Jakarta, Indonesia

#### Correspondence to

Ary I Savitri;  
aryisavitri@gmail.com

#### ABSTRACT

**Objectives:** To evaluate if pre-pregnancy body mass index (BMI) determines blood pressure throughout pregnancy and to explore the role of gestational weight gain in this association. In addition, the effects of pre-pregnancy BMI and gestational weight gain on the occurrence of gestational hypertension and pre-eclampsia were investigated.

**Design:** Prospective cohort study.

**Setting:** Maternal and child health primary care referral centre, Jakarta, Indonesia.

**Population and measurements:** 2252 pregnant women visiting Budi Kemuliaan Hospital and its branch for regular antenatal care visits from July 2012 to April 2015. Pre-pregnancy BMI (kg/m<sup>2</sup>) was based on self-reported pre-pregnancy weight and measured height at first visit. Gestational weight gain was calculated as weight at the day of delivery minus the pre-pregnancy weight. Systolic blood pressure (SBP) and diastolic blood pressure (DBP) were measured during pregnancy at every visit. Linear mixed models were used to analyse this relation with repeated blood pressure measures as the outcome and pre-pregnancy BMI as the predictor. When looking at gestational hypertension and pre-eclampsia as outcomes, (multiple) logistic regression was used in the analysis.

**Results:** Independent of pre-pregnancy BMI, SBP and DBP increased by 0.99 mm Hg/month and 0.46 mm Hg/month, respectively. Higher pre-pregnancy BMI was associated with higher pregnancy SBP (0.25 mm Hg/kg/m<sup>2</sup>; 95% CI 0.17 to 0.34; p<0.01) and DBP (0.18 mm Hg/kg/m<sup>2</sup>; 0.13 to 0.24; p<0.01) in adjusted analysis. Every 1 kg/m<sup>2</sup> higher pre-pregnancy BMI was associated with 6% and 9% higher odds for gestational hypertension (adjusted OR (aOR) 1.06; 95% CI 1.03 to 1.09; p<0.01) and pre-eclampsia (aOR 1.09; 1.04 to 1.14; p<0.01). Accounting for gestational weight gain did not attenuate these associations.

**Conclusions:** Pre-pregnancy BMI determines the level, but not the change, of blood pressure in pregnancy and is linked to higher odds for gestational hypertension and pre-eclampsia, independent of gestational weight gain.

#### INTRODUCTION

Over the past decades, the prevalence of overweight and obesity among women of reproductive age has increased tremendously

#### Strengths and limitations of this study

- This study was conducted in the population with a low prevalence of maternal smoking and alcohol/drug abuse; therefore, the effect of pre-pregnancy body mass index on blood pressures could be analysed in isolation from such factors.
- Analysis was based on the assumption of linearity between blood pressures increase in pregnancy and gestational age at measurement; thus, it may not reflect actual blood pressure trajectory in pregnancy.
- Pre-pregnancy weight was retrospectively asked during interview at the first antenatal care visit; therefore, some inaccuracies may exist.

in developed and developing countries.<sup>1</sup> High body mass index (BMI), together with high systolic blood pressure (SBP), are known as the leading risk factors for mortality and disability-adjusted life-years in women worldwide,<sup>2</sup> mainly through manifestation of cardiovascular diseases and metabolic disorders.<sup>3</sup>

Being overweight or obese also puts women at risk for developing various pregnancy complications, including gestational hypertension and pre-eclampsia,<sup>4–10</sup> and higher risk for cardiovascular events later in life.<sup>11</sup> As compared with normal weight women, overweight and obese women have 3–5 times increased risk for pre-eclampsia.<sup>3</sup> Higher baseline BMI recorded at the first antenatal visit has also consistently been linked to higher risk of hypertensive disorders of pregnancy (HDP).<sup>12–13</sup> Several studies reported that pre-pregnancy BMI influences SBP and diastolic blood pressure (DBP) levels during pregnancy.<sup>5–14–16</sup> However, studies that reported the effect of pre-pregnancy BMI on the rate of change in SBP and DBP are limited. Furthermore, the role of weight gain on the relation between pre-pregnancy BMI and blood pressure during pregnancy is also less consistent. Several studies reported that excessive

gestational weight gain was associated with higher risk of gestational hypertension and pre-eclampsia,<sup>5 16 17</sup> but some other studies did not find such associations.<sup>18 19</sup> It is also unclear whether the relation between obesity and HDP was mediated by excessive weight gain, as overweight or obese women were shown to be more likely to experience excessive weight gain as compared with women with normal BMI.<sup>20</sup>

In the present study, we investigated the effect of pre-pregnancy BMI, as a proxy for maternal adiposity, on several blood pressure measures throughout pregnancy and on the occurrence of gestational hypertension and pre-eclampsia. Furthermore, we explored the influence of gestational weight gain in these associations.

## METHODS

### Study population

This study was based on a prospective cohort of 2252 pregnant women in the private mother-child health Budi Kemuliaan Hospital and its branch (Budi Kemuliaan Petojo) in Jakarta, Indonesia. Women were recruited at first visits for antenatal care (ANC) between July 2012 and October 2014 and followed up until delivery, the last in April 2015. All women who attended ANC visits were invited and asked to sign written informed consent.

Midwives obtained interview information regarding women's demography including age and socioeconomic background (monthly family income, education, employment status). Women's education was categorised as low if the women had finished elementary or junior high school, medium if they finished senior high school or high if they attained education from university. Women's employment status was classified as not working, working in a formal job, or working in a non-formal job (trader, nanny, cleaning service, etc). Women's clinical data included medical history, information on current pregnancy (last menstrual period (LMP), pre-pregnancy weight and weight at every ANC visit), women's obstetrical history (parity, previous pregnancy complications, previous mode(s) of delivery) and family history of diseases (including family history of hypertension and diabetes mellitus type 2).

### Anthropometric measures

Pre-pregnancy weight was self-reported at the first ANC and height was measured using standardised stadiometers. Pre-pregnancy BMI was then calculated as pre-pregnancy weight in kilograms divided by the square of height in metres. Women's weight was measured at each ANC visit using standard scales.

### Outcome measurements

Women's SBP and DBP were routinely measured by midwives at every ANC visit. SBP and DBP were measured once per visit in seated position, after a minimum 30 min rest, using a mercury sphygmomanometer.

Gestational hypertension and pre-eclampsia were classified according to the definitions from the International Society for the Study of Hypertension in Pregnancy (ISSHP).<sup>21</sup> Gestational hypertension was defined as SBP of 140 mm Hg or more and/or DBP of 90 mm Hg or more on two occasions in a woman with no hypertension prior to 20 weeks of gestation. Pre-eclampsia was defined as gestational hypertension with proteinuria.

### Confounding variables and intermediate variables

Women's age, parity, secondhand smoking exposure and monthly family income (proxy for socioeconomic status) were a priori considered as possible confounders. Age was calculated as the difference between the dates of birth and of women's first ANC visit. Parity was classified as nulliparous or multiparous. Data on secondhand smoking exposure were used instead of active smoking because few (<1%) of these pregnant women smoke. Monthly family income was asked to the women as an estimate range. Gestational age at delivery was derived from the date of LMP and the date of delivery.

As a possible intermediate factor, gestational weight gain was calculated as weight on the day of delivery minus pre-pregnancy weight in kilograms.

### Data analysis

Baseline characteristics were described by pre-pregnancy BMI tertiles and group differences were tested using  $\chi^2$ , one-way analysis of variance or Kruskal-Wallis tests where appropriate. Skewed data were reported as median and IQR. Associations between pre-pregnancy BMI and blood pressures were analysed using (un)adjusted linear mixed models for repeated measures, with corresponding 95% CI and p values. Furthermore, to investigate whether gestational weight gain acted as an intermediate factor in this association, we developed an explanatory model which included variables on gestational weight gain. Analyses about the effects of pre-pregnancy BMI on women's risk of developing gestational hypertension and pre-eclampsia were performed using (multiple) logistic regression.

The pattern of blood pressure changes during pregnancy was plotted for each pre-pregnancy BMI tertile by fitting (adjusted) linear mixed models separately with SBP and DBP as the outcomes. The estimated marginal means of SBP and DBP at 13, 16, 19, 22, 25, 28, 31, 34, 37 and 39 weeks of gestation were used to describe the blood pressure trajectory during pregnancy. The baseline gestational age was set at 13 weeks because this was the median gestational age of the first measurement at the ANC visit. Thirty-nine weeks was set as the last BP measurement since this was the median gestational age at delivery. The relation between blood pressure and gestational age (time) was assumed to be linear over the course of pregnancy, as our main goal in this analysis was the mutual ranking of women's blood pressure. In addition, the pattern of weight increase during pregnancy was also presented separately for women in each

pre-pregnancy BMI tertile using the same approach. Analyses were performed based on non-missing data. All statistical analyses were performed using IBM SPSS (V.21 for Windows).

## RESULTS

Figure 1 provides the study flow chart. Owing to missing data on pre-pregnancy BMI, the analysis concerned 2031 women. Baseline characteristics of women according to tertiles of pre-pregnancy BMI are presented in table 1. As compared with the highest BMI tertile, women in the first and second tertiles were younger, more likely nulliparous and had higher total gestational weight gain. Fewer women in the lowest tertile had attained high education. More women in the highest tertile did not work or worked in a non-formal sector. Monthly family income slightly differed with pre-pregnancy BMI. No difference was found with respect to women's smoking status, secondhand smoking exposure and gestational age at the first visit and at delivery.

Table 2 shows that higher pre-pregnancy BMI was associated with a significantly higher SBP and DBP, both crude and confounding adjusted. The explanatory model shows that further accounting for total gestational weight gain did not attenuate the associations. Adjustments for other measures of weight gain were also performed, by adding a variable of weight gain measured at each ANC visit or a variable of weight gain for

gestational age and women's pre-pregnancy weight into the adjusted model. However, the use of these measures did not change the findings compared with the presented measurement for gestational weight gain (data not shown).

In online supplementary material 1, we present the effect estimates if twin pregnancies were excluded from the analysis (n=26). There were no significant differences found as compared with the analysis with twin pregnancies included.

There were 264 women (13.0%) in our study who were diagnosed with gestational hypertension and 81 women (4.0%) with pre-eclampsia. In table 3, the associations between pre-pregnancy BMI and occurrence of gestational hypertension and pre-eclampsia were presented. Higher pre-pregnancy BMI was associated with higher odds for gestational hypertension and pre-eclampsia, in the crude and adjusted models. Adjustment for total gestational weight gain in the explanatory model did not attenuate the associations.

Throughout pregnancy, SBP was estimated to increase by 0.99 mm Hg/month while DBP increased by 0.46 mm Hg/month in all women, independent of their pre-pregnancy BMI tertiles and other possible confounders. Figure 2 shows that mean blood pressures were consistently higher with higher pre-pregnancy BMI, although the pace of increase did not seem to be affected. The lines that depict the pattern of change in SBP and DBP during pregnancy appeared to be parallel in women from different pre-pregnancy BMI tertiles. In addition, no interaction was found between pre-pregnancy BMI and gestational age (p=0.91 in the confounder-adjusted model).

Figure 3 shows the pattern of weight gain in pregnancy by pre-pregnancy BMI tertile. The level of gestational weight gain differed according to women's pre-pregnancy BMI tertiles. The mean gestational weight gain of women in the first, second and third tertile pre-pregnancy BMI was 14.3, 12.8 and 10.2 kg consecutively.

## DISCUSSION

This study shows that pre-pregnancy BMI determines the level of SBP and DBP in pregnancy, but not the pace of blood pressure increase. Pre-pregnancy BMI was also positively associated with the risk of gestational hypertension and pre-eclampsia. The effect was independent of gestational weight gain.

The strength of this study is its prospective design. Furthermore, our study differed from other studies largely conducted in the Western countries where poor health behaviours, such as smoking, alcohol consumption and drug abuse, were more common among obese/overweight women.<sup>22 23</sup> The proportion of women who have these habits is very low in Indonesia,<sup>24 25</sup> as shown by the data from the women involved in this study. This enabled us to evaluate the effect of pre-pregnancy BMI, as a measure of adiposity, in isolation from such factors.

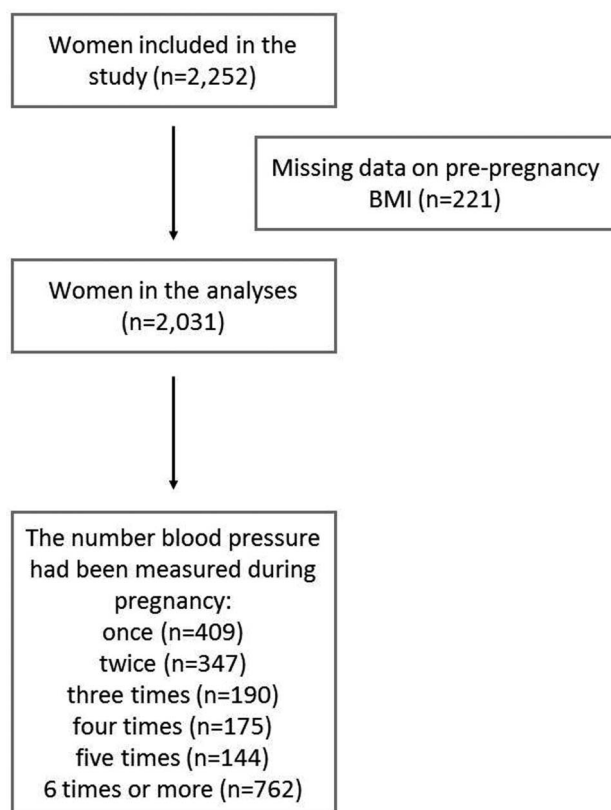


Figure 1 Flow chart of study participants. BMI, body mass index.

**Table 1** Baseline characteristics of women by tertiles of pre-pregnancy BMI

Characteristics	Tertiles of pre-pregnancy BMI			p Value
	Low	Middle	High	
Number (%)	675 (33.2)	684 (33.7)	672 (33.1)	
BMI (kg/m <sup>2</sup> ), median (minimum, maximum)	18.6 (12.5, 20.3)	22.1 (20.3, 23.9)	26.7 (23.9, 47.3)	<0.01*
Age (years), mean (SD)	26.6 (5.6)	28.6 (5.6)	30.5 (5.7)	<0.01†
Women's education				0.03‡
Low education (%)	170 (25.2)	144 (21.1)	132 (19.7)	
Middle education (%)	413 (61.3)	415 (60.8)	415 (61.9)	
High education (%)	91 (13.5)	124 (18.2)	123 (18.4)	
Women's employment status				0.01‡
Not working (%)	385 (57.2)	398 (58.6)	428 (63.9)	
Non-formal job (%)	44 (6.5)	52 (7.7)	57 (8.5)	
Formal job (%)	244 (36.3)	229 (33.7)	185 (27.6)	
Monthly family income				0.01‡
<IDR1 million (%)	68 (10.1)	63 (9.2)	68 (10.1)	
IDR1–2.5 million (%)	320 (47.4)	308 (45.0)	291 (43.3)	
IDR2.5–5 million (%)	204 (30.2)	216 (31.6)	203 (30.2)	
More than IDR5 million (%)	29 (4.3)	59 (8.6)	45 (6.7)	
Refused to answer (%)	54 (8.0)	38 (5.6)	65 (9.7)	
Nulliparity (%)	251 (55.8)	182 (36.6)	132 (25.7)	<0.01‡
Women's smoking status				0.46‡
Yes, sometimes (%)	3 (1.1)	2 (0.7)	3 (1.0)	
Yes, everyday (%)	4 (1.5)	1 (0.3)	1 (0.3)	
Partner's smoking status				0.62‡
Yes, sometimes (%)	42 (15.3)	40 (14.0)	35 (11.8)	
Yes, everyday (%)	125 (45.6)	136 (47.7)	153 (51.5)	
Total gestational weight gain (kg), mean (SD)	14.3 (7.0)	12.8 (6.5)	10.2 (7.1)	<0.01†
Gestational age at first visit (weeks), median (IQR)	13.7 (17.0)	12.1 (17.3)	12.7 (16.8)	0.27*
Gestational age at delivery (weeks), median (IQR)	39.1 (2.3)	39.3 (2.1)	39.3 (2.3)	0.10*
Preterm birth (<37 weeks gestation) (%)	50 (7.4)	39 (5.7)	34 (5.1)	0.18‡
Post-term birth (>42 weeks gestation) (%)	23 (3.4)	30 (4.4)	32 (4.8)	0.44‡

\*Kruskal-Wallis test.

†ANOVA.

‡ $\chi^2$  test.

ANOVA, analysis of variance; BMI, body mass index.

Pre-pregnancy BMI was calculated from the pre-pregnancy weight that was based on interview. We are aware that it could have limited accuracy. The main analysis was based on the assumption of linearity of the association between increase in blood pressures during pregnancy with gestational age. Therefore, although the graph could accurately describe the relative effect and the rankings of pre-pregnancy BMI tertiles on SBP and DBP, it may not fully reflect the actual absolute trajectory of blood pressure changes during pregnancy. Previous studies reported that blood pressure decreases from early pregnancy to mid-pregnancy and then increases until late pregnancy.<sup>14 19 26</sup> However, our BP measurement began at around 13 weeks of gestation, when SBP and DBP are about to approach their lowest level. Owing to missing data, analyses were limited to women with complete data on the determinant, pre-pregnancy BMI, and outcomes, SBP and DBP, but both were fairly complete. About 20% of the women in this study had only one BP measurement. Half of these women were referred to our hospital at a later stage of pregnancy and therefore assessments from the referring healthcare facilities were taken into

account in diagnosing gestational hypertension and pre-eclampsia. The other women only came once to our hospital for consultation and continued their ANC visits at other healthcare facilities; consequently, gestational hypertension and pre-eclampsia could not be ruled out. This may, to some extent, have led to the underestimated prevalence of both gestational hypertension and pre-eclampsia reported in this study. Sensitivity analyses, by including women with at least two, three and four BP measurements, have been done (shown in online supplementary materials 2–7) and gave results that are comparable to the overall analyses (tables 2 and 3). The same results also appeared when we restricted the analysis to women with two or more BP measurements who were not diagnosed with gestational hypertension or pre-eclampsia (see online supplementary material 8). This suggests that the effect estimates presented in this study are relatively unbiased. Finally, owing to very few women who had first trimester ultrasound crown-rump length measurement, gestational age was based on the LMP.

Classification of pre-pregnancy BMI was based on the tertile, rather than using the WHO cut-off criteria.

**Table 2** Effect of pre-pregnancy BMI on systolic and diastolic blood pressure during pregnancy

	Coefficient	95% CI	p Value
<b>Systolic blood pressure</b>			
Crude model	0.32	0.24 to 0.40	<0.01
Adjusted model*	0.25	0.17 to 0.34	<0.01
Explanatory model†	0.41	0.15 to 0.66	<0.01
<b>Diastolic blood pressure</b>			
Crude model	0.22	0.16 to 0.27	<0.01
Adjusted model*	0.18	0.13 to 0.24	<0.01
Explanatory model†	0.27	0.08 to 0.45	<0.01

Results are expressed as mixed model coefficients of systolic and diastolic blood pressure in mm Hg for every 1 kg/m<sup>2</sup> increase in pre-pregnancy BMI.

\*Adjusted for maternal age, parity, second-hand smoking exposure and socioeconomic status (monthly family income).

†As \* and for total gestational weight gain (difference between maternal weight measured at the day of delivery from the pre-pregnancy weight).  
BMI, body mass index.

Using this data-driven approach, a relatively comparable number of women were classified in each group, ensuring sufficient power for statistical testing. Furthermore, there has been uncertainty about an appropriate BMI cut-off for the Asian population. Several studies recommend a lower cut-off for overweight and obesity in the Asian population, but some others do not.<sup>27</sup>

Our study showed that pre-pregnancy BMI influences the level of SBP and DBP during pregnancy. This effect was present consistently over the course of pregnancy. It was in accordance with the reports from previous studies.<sup>5 14 15 28</sup> Almost similar to our findings, a study which explored the relation between blood pressure during pregnancy with gestational age and pre-pregnancy BMI as continuous scale measurements also

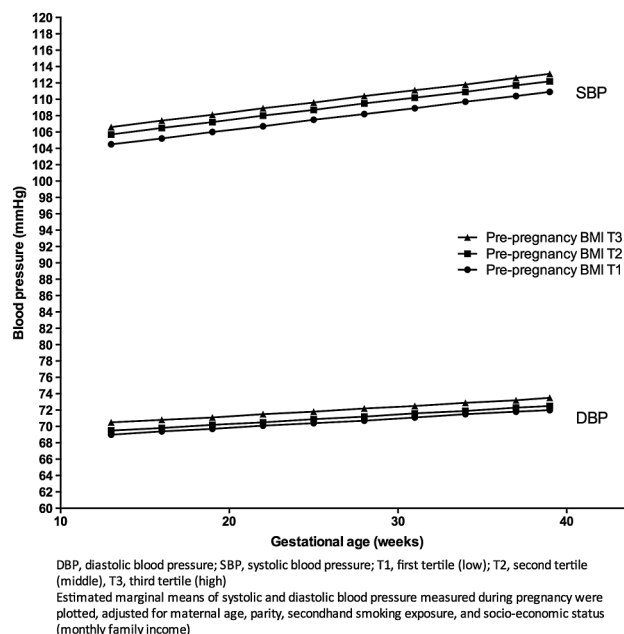
**Table 3** Effect of pre-pregnancy BMI on the risk for developing gestational hypertension and pre-eclampsia

	OR	95% CI	p Value
<b>Gestational hypertension</b>			
Crude model	1.07	1.04 to 1.10	<0.01
Adjusted model*	1.06	1.03 to 1.09	<0.01
Explanatory model†	1.10	1.05 to 1.14	<0.01
<b>Pre-eclampsia</b>			
Crude model	1.09	1.05 to 1.14	<0.01
Adjusted model*	1.09	1.04 to 1.14	<0.01
Explanatory model†	1.19	1.11 to 1.27	<0.01

Results are expressed as OR (95% CI) for every 1 kg/m<sup>2</sup> increase in pre-pregnancy BMI from (multiple) logistic regression.

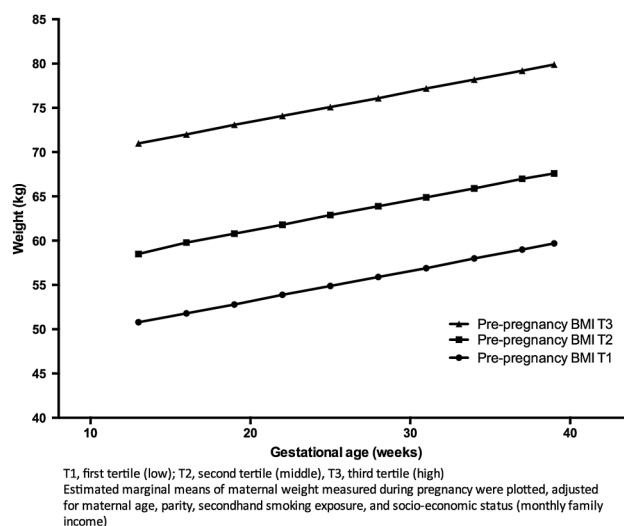
\*Adjusted for maternal age, parity, secondhand smoking exposure and socioeconomic status (monthly family income).

†As \* and for total gestational weight gain (difference between maternal weight measured at the day of delivery from the pre-pregnancy weight).  
BMI, body mass index.

**Figure 2** Effect of pre-pregnancy BMI tertiles on systolic and diastolic blood pressure during pregnancy. BMI, body mass index.

found that SBP and DBP levels generally increased with higher pre-pregnancy BMI at any gestational age.<sup>16</sup>

In this study, SBP and DBP were increased by 0.25 and 0.18 mm Hg per BMI unit, respectively. These findings were similar to those in a previous study<sup>28</sup> which reported an increased mean arterial pressure (MAP) of 0.21 mm Hg per BMI unit, but were smaller compared with another study<sup>5</sup> which reported the mean differences of blood pressure in the first, second and third trimesters of 1.03, 0.98 and 0.89 mm Hg for SBP per BMI unit and 0.83, 0.81 and 0.74 mm Hg for DBP per BMI

**Figure 3** Increase in maternal weight during pregnancy according to pre-pregnancy BMI tertiles. BMI, body mass index.

unit. These discrepancies could be attributed to differences in variables included in the adjustment model, where in the latter study gestational weight gain was taken into account in the adjusted model but not in our analysis. Furthermore, women's BMI in that study may be higher since there were more overweight and obese women (27.6%) than in our study (24.8%). Finally, differences in women's ethnicity could also contribute to these differences. To the best of our knowledge, we are among the first to study this in Asian women.

HDP are a spectrum of diseases that encompass chronic hypertension, gestational hypertension, (superimposed) pre-eclampsia and HELLP (haemolysis, elevated liver enzymes and low platelets) syndrome.<sup>21–29</sup> Our findings were also in accordance with several studies which demonstrated the relation between higher pre-pregnancy BMI and increased risk of gestational hypertension and pre-eclampsia.<sup>10–17, 30–34</sup> These were also in line with studies that took obesity or BMI recorded at the first ANC as the determinant.<sup>3–4, 6–8, 12–13, 35–42</sup>

In this study, we found that pre-pregnancy BMI did not materially affect the pace of increase in blood pressure during pregnancy. An interaction term between BMI tertiles and gestational age that was introduced into the model showed non-significant *p* values, suggesting that the relation between BMI and blood pressures did not depend on gestational age. The increases in SBP and DBP were running in parallel independent of women's pre-pregnancy BMI. A previous study<sup>15</sup> also supports our finding as it showed relatively unchanged mean differences of SBP and DBP between various pre-pregnancy BMI groups in all trimesters. In contrast, another study showed that the pace of increase in BP did vary across BMI group and gestational age, with a ceiling effect which attenuated the increase in BP in higher BMI groups, particularly in later stages of pregnancy.<sup>16</sup>

As compared with women with normal range pre-pregnancy BMI, women who were overweight or obese before pregnancy had higher risk for excessive weight gain during pregnancy.<sup>20</sup> Further, this excessive weight gain could synergistically amplify the risk of pre-eclampsia in obese women.<sup>32</sup> Our finding in unselected pregnant women shows that gestational weight gain did not mediate the effect of pre-pregnancy BMI on blood pressure level during pregnancy, as shown by the unattenuated effect estimates. Our overall finding is in accordance with a study in Sweden showing correlation between DBP during pregnancy with women's baseline BMI, but not with their gestational weight gain.<sup>19</sup> Another study also showed that excessive weight gain did not increase the risk of pre-eclampsia,<sup>18</sup> although it raised the risk of transient hypertension. Many other studies, however, reported a positive relation between excessive weight gain and higher risk of HDP that was independent of pre-pregnancy BMI.<sup>5–17, 30–34, 43</sup> Nevertheless, it is important to note that these studies used excessive gestational weight gain as a determinant,

while in our study gestational weight gain was evaluated as an explanatory variable in a continuous scale measurement.

The pathophysiological mechanism to explain the association between pre-pregnancy BMI and blood pressure levels during pregnancy remains unclear. It has been speculated that adiposity causes a state of increased inflammation, hyperleptinaemia, hyperinsulinaemia and insulin resistance which further leads to disturbances in autonomic function such as sympathetic nervous activation. These mechanisms appeared to be similar in pregnancy and outside of pregnancy, as has been extensively studied among pre-eclamptic pregnant women<sup>44–46</sup> as well as in obese and hypertensive women.<sup>16, 47–49</sup> Obesity may also contribute to the dysregulation of leptin function, which consequently causes hypertension, and alternatively hyperleptinaemia itself could have a direct effect on the increased level of inflammation and raise in blood pressure.<sup>45</sup> In contrast, several studies in non-pregnant populations suggested that the development of hypertension may precede the weight gain through pregnancy.<sup>50–51</sup> Sympathetic hyperactivity in patients with hypertension might downregulate  $\beta$ -adrenergic receptors, which caused a decrease in thermogenic response and therefore increased their propensity to gain weight.

Notably, an adaptive mechanism, which involves maternal systemic inflammatory response through insulin resistance, occurs in all pregnancies, with pre-eclamptic pregnancies being in the extreme end of the continuum.<sup>52</sup> This is supported by a study which showed that women's pre-pregnancy BMI strongly influenced women's MAP throughout pregnancy in normotensive and chronic hypertensive women alike.<sup>28</sup>

A history of hypertensive pregnancy is known to be associated with higher incidence of high blood pressure, ischaemic heart disease and stroke later in life and occurrence at a younger age.<sup>10, 53–55</sup> It was argued that the positive association between gestational hypertension and cardiovascular diseases in later life was caused by their common pathology.<sup>56</sup> Obesity and chronic medical conditions such as diabetes or hypertension, which are among the most recognised risk factors of pre-eclampsia,<sup>9</sup> also predispose women to later cardiovascular diseases. This could imply that the associations are due largely to pre-pregnancy risk factors, rather than to factors that operate exclusively in pregnancy.<sup>57</sup> Pregnancy could be seen as a metabolic and vascular 'stress test' that could unmask women's pre-existing sub-clinical risks, which may manifest in later life as the effects of ageing and long-term exposure to other classical risk factors emerge.<sup>53–56</sup> In addition, excessive gestational weight gain is also associated with long-term maternal abdominal adiposity, which in turn may increase women's risks of cardiovascular and metabolic disease in later life.<sup>58</sup>

Our study showed that women's SBP and DBP increased with higher pre-pregnancy BMI, suggesting a stable linear relation of adiposity and blood pressures during

pregnancy. This resembles the situations occurring outside of pregnancy, as shown by studies in adults<sup>59,60</sup> and in children and adolescents<sup>61</sup> in which adiposity parameters such as BMI were found to be linearly related to the increase in SBP and DBP. We speculate that the difference in blood pressure levels were actually present between women with different BMI before the pregnancy started, persisted during pregnancy and continuously so in the postpartum period. However, we are unable to confirm this hypothesis fully since we do not have data on women's blood pressure prior to pregnancy and postpartum. In contrast, a small study in Brazil showed that although women with excessive weight had higher SBP and DBP than the normal weight women throughout pregnancy, their blood pressures in the postpartum period were comparable.<sup>14</sup> We do acknowledge that pre-pregnancy BMI is a proxy for body adiposity that may not accurately reflect the distinction between lean body mass and body fat.<sup>62</sup> What the specific contributions of pre-pregnancy lean body mass and body fat to blood pressure in pregnancy are, remain of interest for future research.

In conclusion, pre-pregnancy BMI, as a proxy measure of adiposity, determined women's blood pressure level during pregnancy but not the pace of its increase. Higher pre-pregnancy BMI was also associated with higher risk for developing gestational hypertension and pre-eclampsia. Gestational weight gain does not appear to mediate these effects. Obese and overweight women in reproductive age should be encouraged to practise a healthy lifestyle which includes weight reduction prior to pregnancy. This intervention could protect them against various pregnancy complications and improve their long-term cardiovascular risks. Women who enter pregnancy with high BMI should be monitored more closely for development of hypertensive disorders during pregnancy.

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**Contributors** AIS and CSPMU contributed to the study concept and design; AIS acquired the data; AIS, PZ and CSPMU analysed, interpreted the data and drafted the manuscript; JLB, DA, MB and DEG critically reviewed the manuscript for important intellectual content; AIS and PZ performed the statistical analysis. All authors read and approved the final manuscript.

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