The impact of ethnicity on adverse perinatal outcome in women with chronic hypertension: a cohort study

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Running Head: Ethnicity and chronic hypertension in pregnancy

Keywords: pregnancy, chronic hypertension, ethnicity, stillbirth, fetal growth restriction, prematurity

This article has been accepted for publication and undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1002/uog.20132

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ABSTRACT

Objectives: This study aimed to assess the impact of maternal ethnicity on the risk of adverse perinatal outcome in pregnant women with chronic hypertension.

Methods: Demographic and delivery data of women with chronic hypertension and singleton pregnancies from three obstetric units (2000 to 2014) were collated. Multivariable logistic regression models were used to calculate risk ratios (RR) by ethnic group for adverse perinatal outcome in women with chronic hypertension adjusted for other maternal characteristics. The impact of maternal ethnicity on birthweight centile calculation was investigated by comparing customised birthweight centile (GROW) to birthweight centile calculator that does not adjust for maternal ethnicity (Intergrowth 21st).

Results: The cohort included 4045 women (4481 pregnancies) with chronic hypertension. Women of White ethnicity accounted for 47% (n=2122) of the cohort; 36% were Black (n=1601) and 8.5% Asian (n=379). The overall incidence of stillbirth was 1.6%, preterm birth <37 weeks 16%, and fetal growth restriction (birthweight <3rd centile) 11%. Black women, compared to White women, had the highest risk for all adverse perinatal outcomes: stillbirth 3.1% versus 0.6% (adjusted RR 5.56; 95% CI 2.79 to 11.09), preterm birth <37 weeks 21% versus 11% (aRR 1.70; 95% CI 1.43 to 2.01), birthweight <3rd centile 16% versus 7.4% (aRR 2.07; 95% CI 1.71 to 2.51). Asian women, compared to White women, were also at increased risk of adverse perinatal outcomes: stillbirth 1.6% versus 0.6% (aRR 3.03; 95% CI 1.11 to 8.28), preterm birth <37 weeks 20% versus 11% (aRR 1.86; 95% CI 1.44 to 2.40) and birthweight <3rd centile 12% versus 7.4% (aRR 1.69; 95% CI 1.24 to 2.30).

Conclusions: Black ethnicity (compared to White) is associated with the greatest risk of adverse perinatal outcome in women with chronic hypertension even after adjusting for other maternal characteristics. Women of Asian ethnicity are also at increased risk, but to a lesser extent.

Health Regulation Authority Approval: 17/HRA/0021
INTRODUCTION

Ethnic variation in the incidence of adverse maternal and perinatal outcomes has been reported,\textsuperscript{1-4} with women of Black ethnicity being at greater risk of hypertensive disorders of pregnancy compared to White\textsuperscript{1,5} as well as other pregnancy-related morbidity and mortality.\textsuperscript{3,6} Some postulate that these differences relate predominantly to social deprivation and restricted access to healthcare,\textsuperscript{2,7} but given the ethnic variation in pathophysiology observed in conditions such as hypertension outside pregnancy, the mechanisms underpinning these differences in pregnancy warrant further investigation.\textsuperscript{8,9}

Chronic hypertension complicates around 3\% of pregnancies and is rising in incidence due to increasing maternal age and the global obesity epidemic.\textsuperscript{10-14} Superimposed pre-eclampsia, stillbirth, fetal growth restriction, and preterm birth all occur more frequently in pregnancies complicated by chronic hypertension, compared with normotensive pregnancies.\textsuperscript{11,15-18} Previous population studies have found that young and middle-aged women of Black ethnicity demonstrate a steeper age-gradient in prevalence of chronic hypertension than those of White ethnicity and men of Black ethnicity.\textsuperscript{8} Outside pregnancy, ethnic origin is considered when prescribing antihypertensive treatment due to variation in drug response;\textsuperscript{19} the potential benefits of tailoring antihypertensive treatment in pregnancy is yet to be elucidated.

Data from contemporaneous cohorts of women with chronic hypertension in pregnancy investigating the relationship between ethnicity and the incidence of adverse perinatal outcome are lacking and could inform research prioritisation, surveillance pathways and stratified treatment options. The objectives of this large multicentre cohort study were to assess the impact of ethnicity on the risk of adverse perinatal outcome in pregnancy complicated by chronic hypertension amongst a population with access to free healthcare.
METHODS

The cohort was collated from three obstetric units in the UK: Guy’s and St Thomas’ National Health Service (NHS) Foundation Trust (London), St George’s University Hospitals NHS Foundation Trust (London), and Central Manchester University Hospitals NHS Foundation Trust (Manchester). All deliveries after 20 weeks’ gestation recorded on maternity databases between 2000 and 2014 with maternal history of chronic hypertension (diagnosed outside of pregnancy) or a documented blood pressure greater than or equal to 140 mmHg systolic and/or 90 mmHg diastolic recorded before 20 weeks’ gestation (as per the International Society for the Study of Hypertension in Pregnancy classification) were extracted for analysis. Multifetal pregnancies were then excluded from the cohort due to a risk of confounding perinatal outcomes. Demographic and delivery data were recorded for all the remaining singleton pregnancies complicated by chronic hypertension. Only pregnancies with a complete baseline demographic dataset (comprising maternal age, body mass index (BMI), parity, smoking, ethnicity, and deprivation index) were included in the analysis. The size of the study was dictated by the earliest year maternal demographic and delivery data were recorded electronically at each centre.

Ethnicity (as recorded at antenatal booking) was assigned using four ethnic groups, White, Black, Asian, and Other, based on the grouping used by the UK Office for National Statistics. For example women of European origin were assigned to the White ethnic group, women of West African or Caribbean origin were assigned to the Black group, and women of Indian, Pakistani or Bangladeshi origin were assigned to the Asian group. Women of mixed ethnic origin were included in the group that they shared heritage with in the following order of priority: Black, Asian, White and Other. Socioeconomic status was classified using data from the 2010 UK census (updated in 2015) regarding deprivation in seven domains: Income, Employment, Health, Education, Housing and Services, Crime, and the Living Environment.
Women were linked to one of 32,844 census areas using their full 8-character postcode; participants were then categorised into the five groups with one being the least deprived and five being the most deprived. Where the participant had no fixed abode they were included in deprivation index group five.

Birthweight centiles were calculated using birthweight centile charts (Gestation Related Optimal Weight (version 6.7.5.1 2014) and Intergrowth-21st (http://intergrowth21.ndog.ox.ac.uk/)). The GROW customised birthweight centiles adjust for maternal height, maternal weight, maternal ethnicity, parity, gestation at delivery, infant sex and infant birthweight in its calculation. The Intergrowth-21st birthweight centiles adjust for gestation at delivery, infant sex and infant birthweight. Infants were then categorised into those with a birthweight less than the 10th centile and less than the 3rd centile (the latter group characterised as fetal growth restriction (FGR) by a recent Delphi consensus). GROW birthweight centiles were used in the primary analysis and Intergrowth-21st for subsequent comparison. Preterm births (spontaneous and iatrogenic) were categorised as those born before 37 weeks’ gestation and 34 weeks’ gestation. Stillbirths were defined as infants born without signs of life after 20 weeks’ gestation. Neonatal unit admission included infants requiring neonatal intensive care unit and/or special care baby unit admission.
Statistical Analysis

Means (SD) or medians (IQR) were calculated for continuous variables and numbers with percentages were calculated for categorical data. Within the cohort of pregnancies complicated by chronic hypertension, unadjusted risk ratios and associated 95% CIs were calculated by generalised linear models using the statistical package Stata (version 14.1) for baseline demographic factors and subsequent perinatal outcomes (including stillbirth, preterm birth, neonatal unit admission, birthweight <10th and <3rd centile). Adjusted risk ratios were then calculated using a multivariable regression model comparing ethnicity with other demographic factors that could be explanatory or confounding in association with adverse perinatal outcome: deprivation index, maternal age, parity, BMI, smoking history, and year of delivery. Allowance was made for women having more than one pregnancy within the cohort duration by adjusting the standard errors for clustering by hospital identification number. The validity of the model was confirmed by repeating the multivariable regression only including the significant risk factors for each perinatal outcome. Further associations of maternal characteristics and adverse outcome were explored via chi-squared test or linear regression models. Investigation of possible confounding included assessing the impact of centre of delivery on each perinatal outcome.

The study was registered with the Health Regulation Authority (17/HRA/0021) and reported in line with STROBE guidance for observational studies.27
RESULTS

Data from 4481 singleton pregnancies in 4045 women with chronic hypertension between 2000 and 2014 were included in the analysis. The distribution of ethnicity was 47% (n=2122) White, 36% Black (n=1601), 8.5% Asian (n=379), and 8.5% Other (n=379). The flow diagram of participants is shown in Figure 1. Only pregnancies with all demographic characteristics required for the multivariate regression model were included in the final analysis (86% of singleton pregnancies identified). The most common missing data point was BMI accounting for 13% of pregnancies excluded from the analysis with the majority of these in the earliest years of the dataset. Sixteen postcodes could not be linked to deprivation index as they were incorrectly recorded in the source data; these participants were also excluded from the analysis. Thirteen women had no fixed abode and were included in deprivation index group five. The cohort included 2016 (45%) pregnancies from Guy’s and St Thomas’ NHS Foundation Trust (London), 2029 (45%) from St George’s University Hospitals NHS Foundation Trust (London), and 436 (10%) from Central Manchester University Hospitals NHS Foundation Trust (Manchester). There were a greater number of pregnancies included in the cohort from 2010 onwards as maternity records from Central Manchester University Hospitals NHS Foundation Trust were not available before this date (Table 1).

The proportion of women in this cohort requiring caesarean section was 39% and maternal high dependency unit care was needed for 11%. The incidence of stillbirth was 1.6%, preterm birth before 37 weeks’ gestation 16%, birthweight below the 3rd centile 11% and neonatal unit admission was required for 9.2% of infants. Maternal and perinatal outcomes for the whole cohort and for each ethnic group are presented in Table 2.
Multivariable regression model analysis comparing ethnicity to other risk factors contributing to adverse perinatal outcome

Using stillbirth as an important perinatal endpoint, the impact of ethnicity and other maternal characteristics was assessed in a multivariable regression model. The only maternal characteristics remaining significant in the adjusted model were Black and Asian ethnicity. Black women had a risk ratio of 5.56 (95% CI 2.79 to 11.09) of having a stillbirth compared to White women, and Asian women had risk ratio of 3.03 (95% CI 1.11 to 8.28) compared to White women (Table S1).

When the risks were calculated for birthweight <3rd centile, an increased risk was seen in women of Black (RR 2.07; 95% CI 1.71 to 2.51), Asian (RR 1.69; 95% CI 1.24 to 2.30) or Other ethnicity (RR 1.70, 95% CI 1.26 to 2.31) (compared to White women), women aged 40 years or older (RR 1.53; 95% CI 1.09 to 2.16) (compared to women aged <40 years), and women who smoked (RR 1.53; 95% CI 1.15 to 2.04) (compared to non-smokers) (Table S2). Baseline characteristics significantly increasing the risk of birthweight <10th centile in the adjusted model included Black ethnicity (RR 1.64; 95% CI 1.45 to 1.87), being aged 40 years or older (RR 1.34; 95% CI 1.06 to 1.69), nulliparity (RR 1.16; 95% CI 1.03 to 1.30), smoking (RR 1.52; 95% CI 1.28 to 1.81), and living in an area of greatest deprivation (RR 1.20; 95% CI 1.04 to 1.38) (Table S3).

Factors increasing the risk of preterm birth before 37 weeks’ gestation included: Black or Asian ethnicity (RR 1.70; 95% CI 1.43 to 2.01 and RR 1.82; 95% CI 1.41 to 2.35 respectively), being aged 40 years or older (RR 1.52; 95% CI 1.13 to 2.05), nulliparity (RR 1.17; 95% CI 1.01 to 1.35), and living in an area of greatest deprivation (RR 1.42; 95% CI 1.18 to 1.70) (Table S4). When the analysis was repeated assessing the risk of preterm birth before 34 weeks’ gestation, only women of Black (RR 2.69; 95% CI 2.03 to 3.55), Asian (RR 2.48; 95% CI 1.69 to 3.66) and Other
ethnic groups (RR 1.69; 95% CI 1.07 to 2.68), and women living in the most deprived areas (RR 1.39; 95% CI 1.05 to 1.86) were at significantly increased risk (Table S5). In this cohort, where 15.6% of women delivered preterm, 3.2% followed spontaneous labour whilst 12.4% were iatrogenic. No ethnic difference was seen amongst the spontaneous preterm births, but the proportion of Black versus White women giving birth before 37 weeks’ gestation through iatrogenic delivery was 17% versus 7.9% (p<0.0001).

Neonatal unit admission was associated with Black, Asian and Other ethnicity compared to White (RR 1.55; 95% CI 1.25 to 1.93, RR 1.58; 95% CI 1.12 to 2.23, and RR 1.46; 95% CI 1.06 to 2.04 respectively), and nulliparity compared to multiparity (RR 1.32; 95% CI 1.08 to 1.61) (Table S6). Year of delivery also affected the risk of neonatal unit admission with babies born between 2000-2004 and 2005-2009 at greater risk than those born between 2010-2014 (RR 1.35; 95% CI 1.05 to 1.74, and RR 1.30; 95% CI 1.06 to 1.60 respectively).

The regression model for each perinatal outcome was repeated including only the data from women who delivered their babies in the last five years of the study as a sensitivity analysis. This ensured that all centres were included for the same time period. The characteristics associated with the greatest increased risk of adverse outcome remained significant. Non-White ethnicity, living in an area of greater deprivation, smoking and primiparity were all associated with and increased risk of adverse outcome in this model (Table S7).
A summary of all the significant adjusted risk ratios and related confidence intervals for the maternal characteristics associated with adverse perinatal outcome in women with chronic hypertension was collated (Table 3).

Further investigation into potential aetiology underpinning disparity in outcome within the cohort of pregnancies complicated by chronic hypertension

Investigation of a surrogate for severity of maternal disease was conducted by comparing the proportion of mothers admitted to the high dependency unit or intensive care unit after birth.

Black mothers versus White mothers were more likely to require high-level care after birth, 14% versus 8.2% (odds ratio 1.83; 95% CI 1.46-2.29).

Of stillborn babies, 77% had a birthweight <10th centile and 63% had a birthweight <3rd centile.

Most stillbirths occurred before 37 weeks’ gestation (93%). No impact of ethnicity on the proportion of stillbirths with birthweight <10th centile or the gestation at delivery was found.

Within the entire cohort, the proportion of neonates with birthweight <3rd centile was higher amongst those born preterm (54% of births before 34 weeks’ gestation and 19% of births 34 to 37 weeks’ gestation) compared to 6.9% of births from 37 weeks’ gestation (Figure S1). Just over half the infants requiring neonatal admission had birthweights below the 10th centile and 40% of infants requiring neonatal unit admission had birthweight <3rd centile.
Further investigation of the importance of ethnicity within birthweight centile calculation was assessed by comparing the Intergrowth-21st birthweight centiles, which do not include an adjustment for maternal ethnicity, with the GROW birthweight centiles.\textsuperscript{24, 25} The proportion of infants classed as <3rd birthweight centile was 11% using the GROW versus 5.1% using Intergrowth-21st birthweight centiles; those with birthweight <10th centile were 23% versus 13% respectively. No infants classed as birthweight <3rd centile by Intergrowth-21st but not by GROW were admitted to the neonatal care unit; however, 47 infants (12%) requiring neonatal unit admission were classed as birthweight <3rd centile by GROW but birthweight >10th by Intergrowth-21st. The sensitivity and specificity for infants requiring neonatal unit admission were 40% and 93% for those with birthweight <3rd centile calculated using GROW, compared to 16% and 96% respectively for those with birthweight <3rd centile calculated using Intergrowth-21st.
DISCUSSION

Main Findings

This large contemporary cohort study of women with chronic hypertension in pregnancy demonstrates that non-White ethnicity (especially Black) is associated with an increased risk of adverse maternal and perinatal outcome. Other maternal characteristics associated with adverse perinatal outcome have also been identified and are consistent with findings of previous studies; advanced maternal age, smoking status, primiparity and deprivation all impact perinatal complications to differing extents.

It is striking that Black ethnicity was consistently associated with the highest increased risk for all adverse outcomes compared with other baseline characteristics in women with chronic hypertension in pregnancy, even after controlling for deprivation, maternal age, BMI, parity, year of delivery, and smoking. The explanation for this is likely to be multifactorial with biological and non-biological factors contributing. Women with chronic hypertension who were of Black (versus White) ethnicity had a more than a five-fold increase in stillbirth risk. A disparity in incidence of stillbirth in women born in African/Caribbean countries (0.7%) compared to women born in the UK (0.5%) is reported by the Office for National Statistics, but these data suggest that the disparity between ethnicities among women with chronic hypertension is much greater. In this cohort, there was greater sensitivity (40% versus 16%) for infants needing neonatal unit admission identified as <3rd birthweight centile using centiles customised for ethnicity (GROW) compared to Intergrowth-21st birthweight centile (without such customisation), and only a small difference in specificity (93% versus 96%). The surrogate outcome of high dependency unit admission was used to assess severity of disease and women of Black ethnicity were significantly more likely to require this level of care after delivery (compared to White women), suggesting a correlation with increased disease severity amongst this group, and supported by a significantly higher proportion of preterm births that
were iatrogenic among mothers of Black ethnicity compared to White. This corresponds with the findings of another study that found an association between Black ethnicity and iatrogenic preterm delivery.\textsuperscript{31}

**Strengths and Limitations**

To our knowledge, this is the largest cohort study of women with chronic hypertension in pregnancy examining the impact of ethnicity on adverse perinatal outcome in the UK. The data were collated from three centres, which reduces the risk of confounding found in single-centre studies. Robust statistical methodology using regression modelling has allowed for identification and quantification of the impact of maternal factors on subsequent perinatal outcome.

The study has some limitations; it was not possible to assess the severity or duration of chronic hypertension within the cohort. Coding for superimposed pre-eclampsia was not sufficiently reliable given the complexities of this diagnosis in women with pre-existing hypertension\textsuperscript{20} and so the additional impact of this diagnosis on adverse perinatal outcome could not be assessed. Additionally, adequately detailed data on blood pressure control, antihypertensive treatment, aspirin prescription, mode of conception and other maternal co-morbidity were not available. Comparison of temporal changes in care was not possible due to restrictions in the availability of this data; the only outcome that appeared to improve over time within the cohort was neonatal unit admission. Postcode data was utilised to assign socioeconomic status; this methodology may have led to incorrect attribution of level of deprivation, but as English postcode data relates to relatively small size areas compared to similar metrics in other countries, it is unlikely that use of this variable will have led to substantial inaccuracies. It is also likely that other characteristics included in the regression model will have allowed some adjustment for incorrect attribution of deprivation.
Although the cohort had free access to healthcare, it was not possible to assess adherence or uptake of antenatal care amongst differing ethnic groups. There may be cultural or language barriers that influence healthcare access, but the FASTER trial examined variation in perinatal outcomes in women who accessed antenatal care in the first trimester and found Black women had an increased risk of perinatal morbidity compared to White (OR 3.5; 95% CI 2.5 to 4.9), so this factor cannot entirely explain the difference in outcome.  

**Interpretation**

Women of Black ethnicity with chronic hypertension are at significantly greater risk of all adverse perinatal outcomes than White women with chronic hypertension in pregnancy. Women of Asian and Other ethnicities are also at increased risk, but to a lesser extent. There are well described differences in pathophysiology causing hypertension that relate to ethnicity. Variation in socioeconomic circumstances have previously been linked to these differences, but these cannot account entirely for the variances given that the model presented here included deprivation score as a baseline characteristic and the participants had free point-of-care access to a healthcare system for their antenatal care. Severity of maternal disease may also have contributed to these findings and highlights the need to optimise treatment strategies for these women. Outside pregnancy, national recommendations in the UK and US are that first-line antihypertensive agents are stratified based on ethnicity with Black women receiving calcium-channel blockers rather than angiotensin-converting enzyme inhibitors. Consideration of the potential role of first-line antihypertensive agent choice in women of African/Caribbean family origin, as is used outside pregnancy, is recommended, in order to evaluate whether this strategy might improve maternal and perinatal outcomes. Investigation of the impact of genetics in addition to environmental and cultural variations...
Comparison of customised and non-customised birthweight centiles has been discussed in recent studies in the general population.\textsuperscript{35, 36} The importance of accounting for maternal ethnicity in the calculation of birthweight centiles is unclear. Ethnicity has a strong association with adverse outcome within this population of women with chronic hypertension and birthweight centiles customised for ethnicity have greater sensitivity and comparable specificity in association with an important outcome indicative of adverse perinatal outcome. Classification of fetal and hence neonatal growth restriction has recently been examined by a Delphi consensus outlining criteria for diagnosis of fetal growth restriction.\textsuperscript{26} Further investigation into classification of neonatal growth restriction and the role of including ethnicity within birthweight centile calculation is warranted.\textsuperscript{26, 37}

Non-White ethnicity (especially Black), deprivation, advanced maternal age, primiparity, and smoking status all increase the risk of adverse perinatal outcome in women with chronic hypertension. Ethnicity has the largest impact, with Black women with chronic hypertension at greatest risk. Further research is needed to explore the aetiology underpinning these disparities. An awareness of these differences should be considered to inform stratification of antenatal care and treatment pathways.
ACKNOWLEDGEMENTS

This is independent research supported by the National Institute for Health Research Professorship of Lucy Chappell RP-2014-05-019. The views expressed in this publication are those of the author(s) and not necessarily those of the NHS, the National Institute for Health Research or the Department of Health.

FUNDING

This is independent research supported by the National Institute for Health Research Professorship of Lucy Chappell RP-2014-05-019. Paul Seed is funded by Tommy’s Charity and the CLAHRC South London (NIHR). Dr Jenny Myers is supported by a NIHR Clinician Scientist Fellowship (NIHR-CS-011-020).

CONFLICT OF INTEREST

Professor Nelson-Piercy reports personal fees from Alliance Pharmaceuticals, UCB Pharmaceuticals, LEO Pharmaceuticals, Sanofi Aventis and Warner Chilcott outside the submitted work. The other investigators have no disclosures to report.
REFERENCES


FIGURE LEGENDS

**Figure 1:** Flow diagram of identification of the study cohort
### Tables

**Table 1:** Demographic characteristics of the whole cohort with chronic hypertension and by ethnic group

<table>
<thead>
<tr>
<th>Maternal Characteristic</th>
<th>All pregnancies</th>
<th>White ethnicity n=2122</th>
<th>Black ethnicity n=1601</th>
<th>Asian ethnicity n=379</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age at delivery, years</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mean (SD)</td>
<td>33 (5.8)</td>
<td>33 (5.5)</td>
<td>34 (6.2)</td>
<td>32 (5.3)</td>
</tr>
<tr>
<td><strong>Body Mass Index, kg/m²</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mean (SD)</td>
<td>28 (6.4)</td>
<td>27 (6.4)</td>
<td>30 (6.3)</td>
<td>28 (5.2)</td>
</tr>
<tr>
<td><strong>Nulliparous</strong></td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td></td>
<td>2112 (47%)</td>
<td>1210 (57%)</td>
<td>553 (35%)</td>
<td>135 (36%)</td>
</tr>
<tr>
<td><strong>Smoker</strong></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>312 (7%)</td>
<td>209 (9.9%)</td>
<td>71 (4.4%)</td>
<td>11 (2.9%)</td>
</tr>
<tr>
<td><strong>Deprivation Index</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 (least deprived)</td>
<td>350 (8%)</td>
<td>288 (14%)</td>
<td>21 (1.3%)</td>
<td>19 (5.0%)</td>
</tr>
<tr>
<td>2</td>
<td>462 (10%)</td>
<td>339 (16%)</td>
<td>54 (3.4%)</td>
<td>36 (9.5%)</td>
</tr>
<tr>
<td>3</td>
<td>953 (21%)</td>
<td>522 (25%)</td>
<td>242 (15%)</td>
<td>98 (26%)</td>
</tr>
<tr>
<td>4</td>
<td>1403 (32%)</td>
<td>575 (27%)</td>
<td>574 (36%)</td>
<td>123 (32%)</td>
</tr>
<tr>
<td>5 (most deprived)</td>
<td>1313 (29%)</td>
<td>398 (19%)</td>
<td>710 (44%)</td>
<td>103 (27%)</td>
</tr>
<tr>
<td><strong>Year of pregnancy</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2000-2004</td>
<td>836 (19%)</td>
<td>404 (19%)</td>
<td>318 (20%)</td>
<td>60 (16%)</td>
</tr>
<tr>
<td>2005-2009</td>
<td>1598 (36%)</td>
<td>770 (36%)</td>
<td>575 (36%)</td>
<td>128 (34%)</td>
</tr>
<tr>
<td>2010-2014</td>
<td>2047 (45%)</td>
<td>948 (45%)</td>
<td>708 (44%)</td>
<td>191 (50%)</td>
</tr>
</tbody>
</table>

SD = standard deviation
Table 2: Maternal and perinatal outcomes of the whole cohort with chronic hypertension and by ethnic group

<table>
<thead>
<tr>
<th></th>
<th>All pregnancies n=4481</th>
<th>White ethnicity n=2122</th>
<th>Black ethnicity n=1601</th>
<th>Asian ethnicity n=379</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Maternal Outcomes</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gestation at delivery, weeks median (IQR)</td>
<td>39.3 (38.0 to 40.4)</td>
<td>39.7 (38.3 to 40.7)</td>
<td>39.0 (37.4 to 40.1)</td>
<td>39.0 (37.0 to 40.0)</td>
</tr>
<tr>
<td><strong>Onset of labour</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Spontaneous</td>
<td>1832 (41%)</td>
<td>942 (45%)</td>
<td>558 (35%)</td>
<td>154 (41%)</td>
</tr>
<tr>
<td>Induction</td>
<td>1471 (33%)</td>
<td>709 (34%)</td>
<td>542 (34%)</td>
<td>91 (24%)</td>
</tr>
<tr>
<td>Pre-labour caesarean section</td>
<td>1007 (23%)</td>
<td>329 (16%)</td>
<td>399 (25%)</td>
<td>101 (27%)</td>
</tr>
<tr>
<td><strong>Mode of delivery</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unassisted vaginal delivery</td>
<td>2152 (48%)</td>
<td>1039 (49%)</td>
<td>758 (47%)</td>
<td>180 (47%)</td>
</tr>
<tr>
<td>Assisted vaginal delivery</td>
<td>580 (13%)</td>
<td>376 (18%)</td>
<td>113 (7.1%)</td>
<td>39 (10%)</td>
</tr>
<tr>
<td>Caesarean section delivery</td>
<td>1749 (39%)</td>
<td>706 (33%)</td>
<td>727 (45%)</td>
<td>160 (42%)</td>
</tr>
<tr>
<td>High dependency unit admission</td>
<td>417 (11%)</td>
<td>151 (7.1%)</td>
<td>201 (13%)</td>
<td>26 (6.9%)</td>
</tr>
<tr>
<td><strong>Fetal outcomes</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stillbirth</td>
<td>72 (1.6%)</td>
<td>12 (0.6%)</td>
<td>49 (3.1%)</td>
<td>6 (1.6%)</td>
</tr>
<tr>
<td>Preterm birth</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;37 weeks</td>
<td>701 (16%)</td>
<td>236 (11%)</td>
<td>335 (21%)</td>
<td>77 (20%)</td>
</tr>
<tr>
<td>&lt;34 weeks</td>
<td>305 (6.8%)</td>
<td>76 (3.6%)</td>
<td>170 (11%)</td>
<td>35 (9.2%)</td>
</tr>
<tr>
<td>Birthweight, g, median (IQR)</td>
<td>3230 (2780 to 3630)</td>
<td>3350 (2960 to 3740)</td>
<td>3100 (2560 to 3520)</td>
<td>3000 (2600 to 3380)</td>
</tr>
<tr>
<td>Birthweight centile (GROW)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;10&lt;sup&gt;th&lt;/sup&gt; centile</td>
<td>1047 (23%)</td>
<td>389 (18%)</td>
<td>492 (31%)</td>
<td>82 (22%)</td>
</tr>
<tr>
<td>&lt;3&lt;sup&gt;rd&lt;/sup&gt; centile</td>
<td>499 (11%)</td>
<td>157 (7.4%)</td>
<td>248 (15%)</td>
<td>46 (12%)</td>
</tr>
<tr>
<td>Neonatal unit admission</td>
<td>413 (9.2%)</td>
<td>156 (7.4%)</td>
<td>176 (11%)</td>
<td>40 (11%)</td>
</tr>
</tbody>
</table>

IQR= interquartile range

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Table 3: Adjusted risk ratios (95% confidence intervals) of significant maternal characteristics associated with adverse perinatal outcomes in women with chronic hypertension

<table>
<thead>
<tr>
<th>Perinatal outcome</th>
<th>Black versus White women</th>
<th>Asian versus White women</th>
<th>Maternal age &gt;40 years versus &lt;25 years</th>
<th>Smokers versus non-smokers</th>
<th>Primip versus multip</th>
<th>Deprivation Index group 5* versus 1 to 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stillbirth</td>
<td>5.56 (2.79 to 11.09)</td>
<td>3.03 (1.11 to 8.28)</td>
<td>1.07 (0.54 to 2.13)</td>
<td>0.53 (0.12 to 2.20)</td>
<td>1.56 (0.94 to 2.60)</td>
<td>1.31 (0.60 to 2.12)</td>
</tr>
<tr>
<td>Birthweight &lt;10(^{th}) centile</td>
<td>1.64 (1.45 to 1.87)</td>
<td>1.21 (0.97 to 1.51)</td>
<td>1.34 (1.06 to 1.69)</td>
<td>1.52 (1.28 to 1.81)</td>
<td>1.16 (1.03 to 1.30)</td>
<td>1.20 (1.04 to 1.38)</td>
</tr>
<tr>
<td>Birthweight &lt;3(^{rd}) centile</td>
<td>2.07 (1.71 to 2.51)</td>
<td>1.69 (1.24 to 2.30)</td>
<td>1.53 (1.09 to 2.16)</td>
<td>1.53 (1.15 to 2.04)</td>
<td>1.19 (0.99 to 1.42)</td>
<td>1.14 (0.92 to 1.42)</td>
</tr>
<tr>
<td>Preterm &lt;37 weeks</td>
<td>1.70 (1.43 to 2.01)</td>
<td>1.82 (1.41 to 2.35)</td>
<td>1.52 (1.13 to 2.05)</td>
<td>1.26 (0.97 to 1.63)</td>
<td>1.17 (1.01 to 1.35)</td>
<td>1.42 (1.18 to 1.70)</td>
</tr>
<tr>
<td>Preterm &lt;34 weeks</td>
<td>2.69 (2.03 to 3.55)</td>
<td>2.48 (1.69 to 3.66)</td>
<td>1.39 (0.99 to 1.94)</td>
<td>1.34 (0.87 to 2.07)</td>
<td>1.18 (0.94 to 1.49)</td>
<td>1.39 (1.05 to 1.86)</td>
</tr>
<tr>
<td>Neonatal unit admission</td>
<td>1.55 (1.25 to 1.93)</td>
<td>1.58 (1.12 to 2.23)</td>
<td>1.38 (0.93 to 2.04)</td>
<td>1.29 (0.92 to 1.82)</td>
<td>1.32 (1.08 to 1.61)</td>
<td>1.12 (0.88 to 1.43)</td>
</tr>
</tbody>
</table>

*Deprivation index group 5 is associated with the greatest deprivation. Significant risk ratios are in bold type.*
Pregnancies complicated by chronic hypertension 2000 to 2014
n= 5365

Singleton pregnancies complicated by chronic hypertension
2000 to 2014
n= 5231

Singleton pregnancies complicated by chronic hypertension
2000 to 2014 with complete baseline data required for
regression model
n= 4481

Multifetal pregnancies complicated by chronic hypertension 2000 to 2014
n= 134

Pregnancies excluded from analysis due to missing data:
Body Mass Index n=653
Smoking history n= 81
Index of Multiple Deprivation n= 16