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Comparison of adverse perinatal outcomes between Asians and Caucasians: a population-based retrospective cohort study in Ontario

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Abstract

Background: Racial disparities in adverse perinatal outcomes have been studied in other countries, but little has been done for the Canadian population. In this study, we sought to examine the disparities in adverse perinatal outcomes between Asians and Caucasians in Ontario, Canada.

Methods: We conducted a population-based retrospective cohort study that included all Asian and Caucasian women who attended a prenatal screening and resulted in a singleton birth in an Ontario hospital (April 1st, 2015–March 31st, 2017). Generalized estimating equation models were used to estimate the independent adjusted relative risks and adjusted risk difference of adverse perinatal outcomes for Asians compared with Caucasians.

Results: Among 237,293 eligible women, 31% were Asian and 69% were Caucasian. Asians were at an increased risk of gestational diabetes mellitus, placental previa, early preterm birth (< 32 weeks), preterm birth, emergency cesarean section, 3rd and 4th degree perineal tears, low birth weight (< 2500 g, < 1500 g), small-for-gestational-age (<10th percentile, <3rd percentile), neonatal intensive care unit admission, and hyperbilirubinemia requiring treatment, but had lower risks of preeclampsia, macrosomia (birth weight > 4000 g), large-for-gestational-age neonates, 5-min Apgar score < 7, and arterial cord pH ≤ 7.1, as compared with Caucasians. No difference in risk of elective cesarean section was observed between Asians and Caucasians.

Conclusion: There are significant differences in several adverse perinatal outcomes between Asians and Caucasians. These differences should be taken into consideration for clinical practices due to the large Asian population in Canada.

Background

Racial disparities in health outcomes have been widely recognized [1]. Maternal race provides a significant axis for studies investigating perinatal outcomes, including stillbirth, preterm delivery, gestational diabetes mellitus (GDM), preeclampsia, and low or high birth weight [2, 3].

For example, White women are about one and half times more likely to have preterm birth and almost three times more likely to delivery very preterm birth compared to Black women [4].

Infant mortality rate in Blacks is also doubled as compared to Whites [3]. Racial disparities in perinatal outcomes have been believed due to the complexities of social, genetic and environmental factors [1, 5, 6]. Racial disparities in access to health care and prenatal care as well as insurance coverage are also demonstrated to contribute to differential health outcomes [7–10]. A previous study noted that women belonging to a racial/

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ethnic-minority were more likely to be exposed to chronic stressors in their lifetime, enhancing their risk for poor perinatal outcomes [11]. Disparities in perinatal outcomes between Black and Caucasian women have been well documented by a series of studies in the United States (US) [4, 12, 13]. However, health disparities between Asian-Caucasian Americans have been understudied although Asian Americans account for 5.7% of the US population [14], which has promoted more research in the perinatal field to Asian Americans [15]. In Canada, population-based studies examining Asian-Caucasian differences in components of maternal and neonatal outcomes are scarce. Although Canada and the US share some social and economic similarities, results of studies conducted in the US may not be generalizable to the Canadian population due to differing racial composition and population context.

Asian Canadians comprise the largest and fastest growing minority group in Canada. Ontario, as the most populous province of Canada and with Asian origin accounting for 23% of the total population [16], provides a unique opportunity to investigate variations in adverse perinatal outcomes between Asians and Caucasians. We therefore conducted a retrospective cohort study to examine disparities in adverse perinatal outcomes between Asians and Caucasians in Ontario.

Methods

Study design and data source

The study design is a population-based retrospective cohort study. We used data from Better Outcomes Registry & Network (BORN) Ontario birth registry, which contains comprehensive perinatal information covering virtually all hospital deliveries in Ontario, to conduct this study. Data access to the BORN is managed under the Personal Health Information Protection Act, 2004 (PHIPA) [17]. This study received ethical approval from the Children's Hospital of Eastern Ontario Research Ethics Board (16/119X) and the Ottawa Health Science Network Research Ethics Board (20160780-01H).

Study population

All Asian and Caucasian women who attended a prenatal screening and resulted in a singleton birth in any Ontario hospital from April 1st, 2015 to March 31st, 2017, were included in this study. If participants had multiple births during the study period, only the first birth was included. We excluded women with missing information on ethnicity or classified as mixed or other racial groups. Women with a history of hypertension were excluded for analysis of gestational hypertension and preeclampsia. Women who were diagnosed with diabetes prior to the index pregnancy were excluded for analysis of GDM.

Outcome measures

Outcome measures considered in this study consist of a range of adverse maternal and neonatal complications. Maternal outcomes included GDM, gestational hypertension, preeclampsia, placental previa, preterm birth (< 37, < 34, < 32 weeks), spontaneous preterm birth, cesarean section (elective, emergency), assisted vaginal delivery, episiotomy, and 3rd and 4th degree perineal tears. Neonatal outcomes included sentinel congenital anomalies, low birth weight (LBW) (< 2500 g, < 1500 g), macrosomia (> 4000 g), small-for-gestational-age (SGA) neonates (defined as < 10th percentile of birth weight for gestational age) [13], SGA neonates (< 3rd percentile), large-for-gestational-age (LGA) neonates (defined as > 90th percentile of birth weight for gestational age), 5-min Apgar score < 7, cord arterial PH ≤ 7.1 , hyperbilirubinemia require treatment (limiting to live births), and neonatal intensive care unit (NICU) admission. Values of birth weight outside of the range of 250 g–6000 g and values of arterial cord pH outside of the range of 6.6–7.4 were considered as outliers and were set to missing.

Exposure and covariates

Maternal race (Asian/Caucasian) was the main exposure measure, self-reported and recorded by care providers at the prenatal screening. We considered a series of relevant factors which could be potential confounders for the association between maternal race and perinatal outcomes, including maternal age at delivery ($\leq 18, 19-24, 25-29, 30-34, 35-39, \geq 40$ years) [1, 2, 18–20], neighbourhood household income (lowest, 2nd, 3th, 4th, highest), parity (0, ≥ 1) [1, 18, 19, 21], pre-existing physical health problems (hypertension or diabetes or heart disease or pulmonary disease) [1], pre-existing mental health problems (a composite measure of depression and anxiety), previous cesarean section (yes, no) [1], pre-pregnancy body mass index (BMI) (defined as height in kilograms (kg) divided by weight in meters squared (m^2) (< 18.5, 18.5–24.9, 25.0–29.9, 30–34.9, 35–39.9, ≥ 40 kg/ m^2) [1, 7], assisted reproductive technology (ART) (yes, no) [1], substance use/alcohol exposure/smoking during pregnancy (yes, no) [1, 7, 8, 20], maternal residence area (rural, urban) [2, 18], obstetrician in antenatal care team (yes, no) [2, 7, 18, 19], and hospital level of maternal care at delivery (I, II, III) [22]. We derived neighbourhood household income and maternal residence area data from the 2011 Canadian census using Statistics Canada's Postal Code Conversion File (PCCF) through the maternal residence postal code because BORN does not collect data on social economic status [23].

Statistical analysis

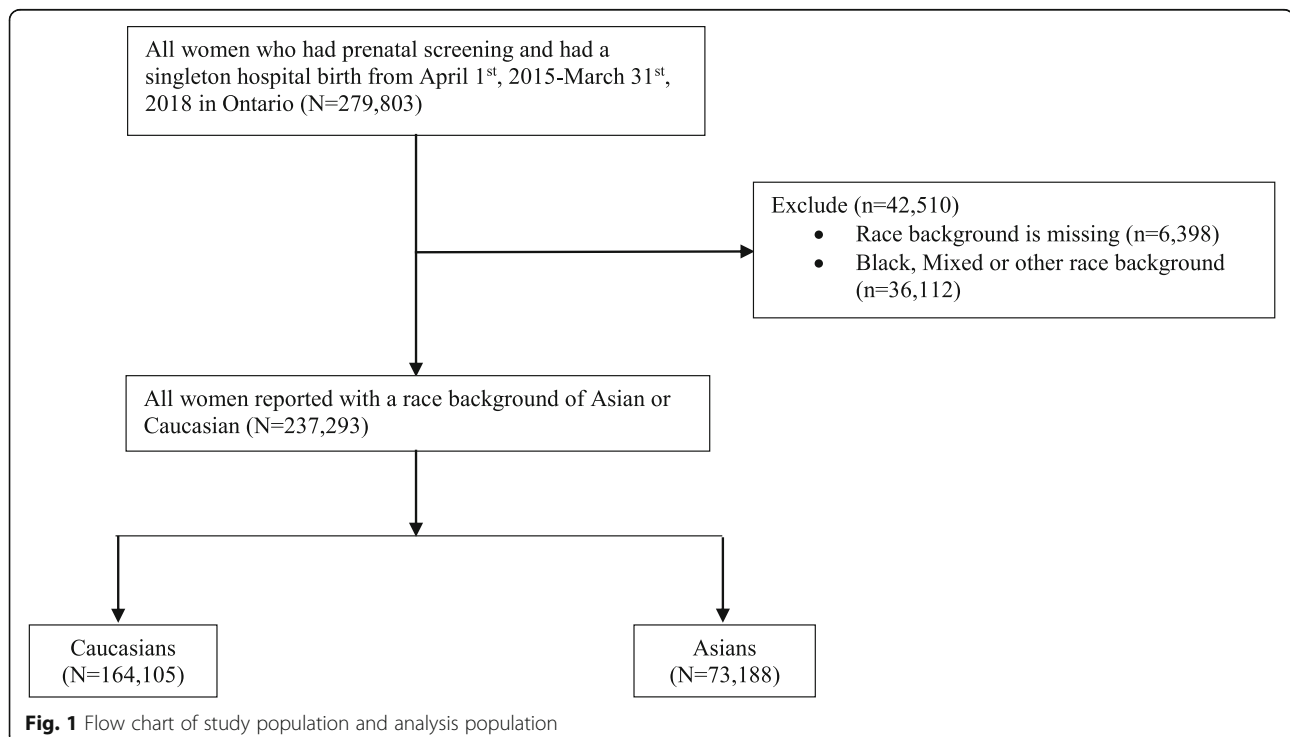
We first compared baseline characteristics between Asians and Caucasians. Continuously distributed variables were

presented by mean \pm standard deviation (SD) and compared by using a t-test. Categorical variables were displayed by counts and percentages and compared by using a chi-square test. We then compared adverse perinatal outcomes between Caucasians and Asians. Generalized estimating equations (GEE) model with a log link function and a Poisson distribution were used to estimate the adjusted relative risks (aRR) and adjusted risk difference (aRD) with their 95% confidence intervals (CI) of perinatal outcomes for Asians, with Caucasians as the reference [24, 25]. Potential confounding variables included in the GEE models were maternal age at delivery, neighborhood household income, pre-existing physical health problems, pre-existing mental health problems, previous caesarean section, pre-pregnancy BMI, parity, ART, substance use/alcohol exposure/smoking during pregnancy. For procedure-related outcomes (assisted vaginal delivery, episiotomy, cesarean section, episiotomy, 3rd and 4th degree vaginal tears, NICU admission) were further adjusted for maternal residence area, obstetrician in antenatal care team, and hospital level of maternal care at delivery, in addition to the aforementioned covariates. Confounders are carefully selected to be adjusted in the multivariable models for each perinatal outcome separately to avoid the occurrence of overadjustment [26, 27]. All confounders in the multivariate regression analysis were selected by ensuring they are independently associated with both race in our source population and perinatal outcomes among Caucasian women (the reference group in this study) only

with a cutoff of 0.05 [28]. Multiple imputation method was used to account for missing data in the regression analysis, in which five datasets were imputed by using fully conditional specification (FCS) logistic regression method [29–31], assuming a joint distribution for all variables. Specifically, linear regression model was used for maternal age and pre-pregnancy BMI (kg/m^2). Generalized logit model was used for household income quintile, parity, previous caesarean section, assisted reproductive technologies, substance use during pregnancy, mental health, and urban/rural residence. All variables used in multivariable analysis were included in imputation models. Statistical Analysis System (SAS) for windows, version 9.4 (SAS Institute, Cary, NC) was used to perform all of the analysis in this study, the criteria for statistical significance was set at $\alpha = 0.05$.

Results

A total of 237,293 eligible women (30.9% Asians and 69.1% Caucasians) were included in the final analysis (Fig. 1). Compared to Caucasian women, Asian women tended to be older and had a significantly higher rate of being in the lowest income quintile level, living in urban areas, being underweight, having a previous caesarean section, and having an obstetrician on the antenatal care team. On the other hand, Asian women were less likely to be nulliparous, be overweight/obese, partake in alcohol consumption/substance use/smoking during pregnancy, have pre-existing disease, have mental health



problems, and also less likely to deliver in lower maternal level of care hospital (Table 1).

Compared with Caucasian women, Asian women had higher risks of GDM, placental previa, preterm birth (< 37, < 34, < 32 weeks), spontaneous preterm birth, emergency cesarean section, episiotomy and 3rd and 4th degree perineal tears, but lower risks of gestational hypertension, preeclampsia after adjusting for relevant confounders (Table 2). No difference was found in risk of elective caesarean section between these two groups.

Compared with Caucasians, Asians had higher risks of low birth weight (< 2500 g, < 1500 g), SGA neonates (< 10th percentile, < 3rd percentile), NICU admission, and hyperbilirubinemia requiring treatment, but had lower risks of sentinel congenital anomalies, macrosomia, LGA neonates, 5-min Apgar score < 7, and arterial cord pH ≤ 7.1 after adjusting potential confounders (Table 3).

Discussion

In this population-based study, we have several principal findings. First, we found that compared with Caucasians, Asians had elevated risks of GDM, placental previa, preterm birth, and emergency cesarean section, whereas had lower risks of gestational hypertension, preeclampsia. Second, Asians have elevated risk of LBW, SGA, NICU admission, and hyperbilirubinemia requiring treatment, compared to Caucasians, but are less likely to have macrosomia, LGA, 5-min Apgar score < 7, sentinel congenital anomalies, and arterial cord pH ≤ 7.1 . We find no difference in risk of elective cesarean section was observed between the two groups.

The most substantial difference in adverse maternal outcomes between Asian and Caucasian women observed in this study was GDM. Asian background has been associated with markedly increased risk of GDM [32], and our study finding provided additional evidence supporting an increased risk of GDM in Asians. A recent systematic review and meta-analysis showed that the pooled rate of GDM in Asians was 11.5% (95% CI 10.9–12.1) [33], which is close to the rate of GDM (13.7%) in Asians in this study. Another important difference in adverse maternal outcomes observed in this study was that Asian women had a higher rate of 3rd and 4th degree perineal tear than Caucasian women, which is consistent with previous studies [34, 35]. In our study, Asian women were less likely to have macrosomic babies and less likely to be nulliparous, which are protective against perineal tears [36]. This phenomenon is likely associated with the smaller stature and shorter perineum of Asian women relative to Caucasian women [37]. Asian women had higher risk of preterm birth, which is similar to previous studies [38, 39]. This is more pronounced in the subgroup of early preterm birth (e.g., < 32 weeks). The findings of Asians having an elevated risk of placenta

previa is also consistent with our previous studies [40] which may be explained by cultural influence (such as stress), nutrition or true genetic differences. Asian women are observed to have a relatively larger placenta even though the reasons are still not elucidated [41]. The higher rate of emergency cesarean section among Asians may be explained by social deprivation or communication difficulties [42]. We speculate that the lower risk of preeclampsia and gestational hypertension for Asian women, may in part be explained by their lower risk of health behaviours such as recreational drugs, alcohol, and cigarette smoking during pregnancy that was observed in this study and many previous studies [43–45], although it is still unclear.

The most striking difference in neonatal outcomes between Asians and Caucasians was the size of the newborns, in which Asians had higher risks of low birth weight and SGA but lower risk of macrosomia and LGA than Caucasians. These findings are in general consistent with previous studies [20, 21, 46, 47], although somewhat different from an earlier study by our team comparing birth weight distribution between Caucasian and East Asian (Chinese) [48]. Specifically, we found that while the mean birth weight in Chinese was substantially lower than that of in Caucasians with lower rate of macrosomia, the rate of low birth weight was also lower in Chinese infants [48]. As we have stated earlier, Asians in this study were from different regions in Asia with some distinctive features and lumping them together limited our ability to identify specific differences among them.

The higher rate of NICU admission for neonates delivered by Asian women might be associated with the increased risk of early preterm birth among Asian women found in our study. The result of lower rate of 5-min Apgar score < 7 among Asian population compared with Caucasian in our study, which is similar to a CDC US report covering 3,163,441 live births from 48 Reporting States and the District of Columbia [49]. It is also consistent with results from a prior study (monthly vital statistics report_1995) [50]. However, the reasons for lower rate of 5-min Apgar score < 7 are still needed to be explored.

The finding of higher risk of hyperbilirubinemia required treatment in Asian infants was consistent with previous studies [51, 52]. One of the reasons the phenomenon of increased risk of hyperbilirubinemia required treatment among Asians may be caused by a common DNA-sequence variant (Gly71Arg) carried by Asians, resulting in an amino acid change in the uridine diphosphate glucuronosyl-transferase protein [53]. Asian women were found to have a slightly lower risk of sentinel congenital anomalies, whereas a previous study found no difference in overall congenital anomalies

Table 1 Comparison of characteristics between Asians and Caucasians, Ontario, Canada, April 1st, 2015-March 31st, 2017 (N = 237,293)

Characteristics	Asian		Caucasian		P value
	n	%	n	%	
	73,188	30.9	164,105	69.1	
Maternal Age at delivery (years) (Mean ± SD)	32.07 ± 4.5		31.08 ± 4.98		<.0001
≤ 18	49	0.1	1221	0.7	< 0.001
19–24	3207	4.4	15,229	9.3	
25–29	17,792	24.3	41,652	25.4	
30–34	30,428	41.6	65,448	39.9	
35–39	18,019	24.7	34,207	20.9	
≥ 40	3592	4.9	6116	3.7	
Missing	101	0.1	232	0.1	
Neighbourhood median household income quintiles (link to 2011 Canadian Census data)					<.0001
Quintile 1 (lowest)	17,117	23.6	26,877	16.6	
Quintile 2	16,540	22.8	30,678	18.9	
Quintile 3	16,544	22.8	33,781	20.8	
Quintile 4	13,679	18.9	37,563	23.2	
Quintile 5 (highest)	8556	11.8	33,148	20.5	
Missing	752	1.0	2058	1.3	
Maternal pre-existing disease ^a					<.0001
No	70,579	96.4	151,491	92.4	
Yes	2609	3.6	12,544	7.6	
Mental health Condition					<.0001
No	66,363	96.6	123,975	80.8	
Yes	2366	3.4	29,465	19.2	
Missing	4459	6.1	10,665	6.5	
Previous cesarean section					<.0001
Yes	60,129	83.8	138,671	86.5	
No	11,656	16.2	21,624	13.5	
Missing	1403	1.9	3810	2.3	
Pre-pregnancy BMI (kg/m ²) (Mean ± SD)	23.4 ± 4.5		25.74 ± 6.17		<.0001
Underweight (< 18.5)	5958	9.7	6537	4.4	<.0001
Normal (18.5–24.9)	36,655	59.7	75,072	50.8	
Overweight (25.0–29.9)	13,409	21.8	36,235	24.5	
Obese (30–34.9)	3982	6.5	17,044	11.5	
Obese (35–39.9)	1001	1.6	7982	5.4	
Obese (≥40)	378	0.6	4991	3.4	
Missing	11,805	16.1	16,244	9.9	
Parity					<.0001
0	31,289	43.1	75,996	46.7	
≥ 1	41,386	56.9	86,738	53.3	
Missing	513	0.7	1371	0.8	
Conception by assisted reproductive technology					<.0001
No	65,565	96.2	145,616	95.4	
Yes	2622	3.8	6978	4.6	

Table 1 Comparison of characteristics between Asians and Caucasians, Ontario, Canada, April 1st, 2015-March 31st, 2017 (N = 237,293) (Continued)

Characteristics	Asian		Caucasian		P value
	n	%	n	%	
	73,188	30.9	164,105	69.1	
Missing	5001	6.8	11,511	7.0	
Drug use during pregnancy					<.0001
No	69,375	97.9	137,147	86.7	
Yes	1458	2.1	21,012	13.3	
Missing	2355	3.2	5946	3.6	
Alcohol exposure during pregnancy					<.0001
No	69,833	99.1	152,805	97.2	
Yes	612	0.9	4470	2.8	
Missing	2743	3.7	6830	4.2	
Smoking during pregnancy (any time)					<.0001
No	69,485	98.9	140,296	89.4	
Yes	771	1.1	16,575	10.6	
Missing	2932	4.0	7234	4.4	
Maternal residence area					
Urban	72,386	99.2	139,566	85.4	
Rural	593	0.8	23,841	14.6	
Missing	209	0.3	698	0.4	
Obstetrician in antenatal care team					<.0001
No	11,171	15.3	49,579	30.2	
Yes	62,017	84.7	114,456	69.8	
Hospital level of maternal care					< 0.001
level I	1383	1.9	18,597	11.3	
level II	60,006	82.0	101,777	62.0	
level III	11,799	16.1	43,661	26.6	

^aMaternal pre-existing disease includes any of hypertension, diabetes, heart disease, and pulmonary disease

1. Missing data represents missing values for neighborhood household income level and education level, parity, previous caesarean section, drug use, alcohol use, birth weight and antenatal health care provider were excluded from the percentage calculation

2. Bold values mean the risk factor favouring corresponding race group

(including sentinel congenital anomalies, down syndrome etc.) between Asians and Caucasians [39].

There are several strengths of this study. First, this study is based on a large population with comprehensive demographic and health care information, allowing an investigation of a number of adverse perinatal outcomes with appropriate adjustment for potential confounding factors. Second, our study has a large sample size of Asian women, enabling a robust comparison between Caucasians and Asians with greater than at least 90% power to detect the difference for each perinatal outcome with a two-tailed alpha (type 1 error) of 5%, where previous studies had smaller samples of Asians [1, 7, 18, 54, 55]. Third, universal access to quality maternity care helped to isolate maternal factors from health care factors.

Limitations of this study should be acknowledged. First, Asians in this study included women from a variety of regions in Asia. Although these women share some common demographic and cultural background, there are major differences in genetic and environmental factors among them. However, although grouping different Asians together may have limited our ability to reveal some specific differences from Caucasians and to properly interpret specific results, it gives us an overall sense of discrepancies in perinatal outcomes between Asians and Caucasians, which will direct us to focus on some specific outcomes in future work. Second, as race status is considered to be a subjective assessment, which might generate misclassification of race status leading to unavoidable bias. Third, as our study population included women who had undergone prenatal screening

Table 2 Comparison of risks of adverse maternal outcomes between Asians and Caucasians, Ontario, Canada, April 1st, 2015-March 31st, 2017 (N = 69,734)

Maternal Outcomes	Asian		Caucasian (reference)		Adjusted RR (95% CI)	Adjusted RD (95%CI)
	n	%	n	%		
Gestational diabetes	9793	13.38	9077	5.53	2.71 (2.68, 2.74)	1.00 (0.97, 1.03)
Gestational hypertension	2012	2.75	6461	3.94	0.93 (0.88, 0.98)	-0.07 (-0.12, -0.02)
Preeclampsia	1945	2.66	7045	4.29	0.84 (0.78, 0.89)	-0.18 (-0.23, -0.13)
Placental previa	700	0.96	1097	0.67	1.30 (1.21, 1.40)	0.26 (0.17, 0.36)
Preterm birth (< 37 weeks)	5070	6.93	10,419	6.35	1.23 (1.20, 1.27)	0.21 (0.17, 0.24)
Preterm birth (< 34 weeks)	1376	1.88	2603	1.59	1.37 (1.29, 1.44)	0.31 (0.24, 0.38)
Preterm birth (< 32 weeks)	901	1.23	1582	0.96	1.49 (1.39, 1.58)	0.40 (0.30, 0.49)
Spontaneous preterm birth	1994	2.72	4103	2.50	1.25 (1.20, 1.31)	0.23 (0.17, 0.28)
Cesarean section	21,694	29.64	46,416	28.30	1.03 (1.01, 1.05)	0.03 (0.01, 0.05)
Elective cesarean section	11,542	15.77	24,968	15.22	0.91 (0.89, 0.94)	-0.09 (-0.11, -0.07)
Emergency cesarean section	10,146	13.86	21,443	13.07	1.22 (1.20, 1.25)	0.20 (0.18, 0.22)
Assisted vaginal delivery	8521	11.64	14,601	8.90	1.29 (1.26, 1.31)	0.25 (0.23, 0.28)
Episiotomy	10,443	14.27	14,552	8.87	1.40 (1.38, 1.43)	0.34 (0.31, 0.36)
3 rd and 4 th degree perineal tears	2756	4.19	3836	2.65	1.57 (1.52, 1.62)	0.45 (0.40, 0.50)

RR relative risk, CI confidence interval

1. Generalized estimating equations with a log link function and a poisson distribution were used to estimate the relative risks of the outcomes.
2. Covariates included in the adjusted models for each outcome were selected covariates that showed univariate association of $P < 0.05$ with both the exposure and the outcome were included in the adjusted model. The covariate for each outcome was fit separately
3. A fully conditional specification method was used to impute missing values, assuming a joint distribution for all variables. Five imputed datasets were created

Table 3 Comparison of risks of adverse neonatal outcomes between Asians and Caucasians, Ontario, Canada, April 1st, 2015-March 31st, 2017 (N = 69,734)

Neonatal outcome	Asian		Caucasian (reference)		Adjusted RR (95% CI)	Adjusted RD (95%CI)
	n	%	n	%		
Sentinel Congenital Anomalies	1046	1.43	2938	1.79	0.90 (0.83, 0.98)	-0.10 (-0.18, -0.03)
Low birth weight (< 2500 g)	5089	7.00	7160	4.40	1.81 (1.77, 1.85)	0.59 (0.55, 0.63)
Low birth weight (< 1500 g)	756	1.04	1229	0.76	1.59 (1.49, 1.69)	0.46 (0.36, 0.56)
Macrosomia (> 4000 g)	3459	4.76	19,214	11.80	0.43 (0.39, 0.46)	-0.85 (-0.89, -0.82)
Small-for-gestational-age neonates (<10 th percentile)	10,396	14.35	12,229	7.56	1.93 (1.91, 1.96)	0.66 (0.63, 0.69)
Small-for-gestational-age neonates (<3 rd percentile)	2703	3.73	2879	1.78	2.19 (2.14, 2.25)	0.78 (0.73, 0.84)
Large-for-gestational-age neonates (>90 th percentile)	3610	4.98	17,660	10.92	0.50 (0.46, 0.53)	-0.70 (-0.73, -0.66)
5- min Apgar score < 7	1219	1.69	3742	2.32	0.89 (0.82, 0.96)	-0.11 (-0.18, -0.05)
Arterial cord pH \leq 7.1	2595	4.04	8704	6.11	0.71 (0.66, 0.76)	-0.34 (-0.39, -0.30)
NICU admission	8529	11.65	18,796	11.46	1.18 (1.16, 1.21)	0.17 (0.14, 0.19)
Hyperbilirubinemia requiring treatment	4300	6.95	8132	5.52	1.41 (1.37, 1.45)	0.35 (0.31, 0.38)

RR relative risk, CI confidence interval, RD risk difference, NICU neonatal intensive care unit

1. Generalized estimating equations with a log link function and a poisson distribution were used to estimate the relative risks of the outcomes.
2. Covariates included in the adjusted models for each outcome were selected covariates that showed univariate association of $P < 0.05$ with both the exposure and the outcome were included in the adjusted model. The covariate for each outcome was fit separately
3. A fully conditional specification method was used to impute missing values, assuming a joint distribution for all variables. Five imputed datasets were created

so that it only captures approximately 70% of pregnant women in Ontario [56]. Women who attend prenatal screening tend to live in an urban area and high income neighbourhood, to receive prenatal care from an obstetrician, and are more likely to be an immigrant or a refugee [57]. Fourth, there were significant differences in baseline characteristics between Asians and Caucasians, which might still have impact on our results due to possible residual confounding. Finally, we did not cover some perinatal outcomes, including placenta accreta, postpartum haemorrhage, neonatal asphyxia and infection in our study due to incomplete information in BORN database, and did not report some underpowered outcomes, such as maternal ICU admission, placental abruption, stillbirth, 5-min Apgar score < 4 and neonatal death due to their low incidence rates (less than 1% in Ontario). Despite these limitations, the results of this study are valuable in informing future work on perinatal outcomes in persons from subgroups within the Asian diaspora.

Conclusion

In summary, our population-based study found significant differences in several adverse perinatal outcomes between Asians and Caucasians. Given the heterogeneity in the demographic and social characteristics among different Asian groups, future studies will be valuable to explore these differences among specific Asian groups.

Abbreviations

ART: Assisted Reproductive Technology; BMI: Body Mass Index; BORN: Better Outcomes Registry & Network; CI: Confidence Interval; FCS: Fully Conditional Specification; GDM: Gestational Diabetes Mellitus; GEE: Generalized Estimating Equations; LGA: Large-for-Gestational-Age; NICU: Neonatal Intensive Care Unit; PCCF: Postal Code Conversion File; PHIPA: Personal Health Information Protection Act; RD: Risk Difference; RR: Relative Risk; SAS: Statistical Analysis System; SD: Standard Deviation; SGA: Small-for-Gestational-Age; US: United States

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Authors' contributions

YG and SW contributed to the study concept and design. All of the authors (NZ, EE, WW, DC, SW, YG) were involved in the analysis and interpretation of data and the critical revision of the manuscript for important intellectual content. EE conducted the statistical analysis. NZ wrote the first draft. All of the authors approved the final version to be published. YG is the guarantors of the work.

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Availability of data and materials

The data analyzed during this study are held securely at the prescribed registry BORN Ontario. Data sharing regulations prevent these data from

being made available publicly due to the personal health information in the datasets. Enquiries regarding BORN data must be directed to BORN Ontario (Science@BORNOntario.ca).

Ethics approval and consent to participate

This study received ethical approval from the Children's Hospital of Eastern Ontario Research Ethics Board (16/119X) and the Ottawa Health Science Network Research Ethics Board (20160780-01H). Our research team acquired permissions from BORN local administrator to access the data used in this research.

Consent for publication

Not Applicable.

Competing interests

We have no conflict of interest.

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References

- Soffer MD, Naqvi M, Melka S, Gottlieb A, Romero J, Fox NS. The association between maternal race and adverse outcomes in twin pregnancies with similar healthcare access. *J Matern Fetal Neonatal Med.* 2018;31(18):2424–8.
- Singh GK, Stella MY. Adverse pregnancy outcomes: differences between US- and foreign-born women in major US racial and ethnic groups. *Am J Public Health.* 1996;86(6):837–43.
- Matthews TJ, MacDorman MF. Infant mortality statistics from the 2010 period linked birth/infant death data set. *Natl Vital Stat Rep.* 2013;62(8):1–26.
- McKinnon B, Yang S, Kramer MS, Bushnik T, Sheppard AJ, Kaufman JS. Comparison of black-white disparities in preterm birth between Canada and the United States. *CMAJ.* 2016;188(1):E19–26.
- Hoyert D. Perinatal Mortality in the United States: 1985–91. National Center for Health Statistics. *Vital Health Stat.* 1995;20(26) Accessed August 16, 2020. https://www.cdc.gov/nchs/data/series/sr_20/sr20_026.pdf.
- Alhusen JL, Bower KM, Epstein E, Sharps P. Racial discrimination and adverse birth outcomes: an integrative review. *J Midwifery Womens Heal.* 2016;61(6):707–20.
- Bryant AS, Worjohol A, Caughey AB, Washington AE, Bryant A. Racial/ethnic disparities in obstetrical outcomes and care: prevalence and determinants racial and ethnic disparities in obstetrical outcomes and obstetrical care. *Am J Obstet Gynecol.* 2010;202(4):335–43.
- Culhane JF, Goldenberg RL. Racial disparities in preterm birth. *Semin Perinatol.* 2011;35(4):234–9.
- Creanga AA, Bateman BT, Kuklina EV, Callaghan WM. Racial and ethnic disparities in severe maternal morbidity: A multistate analysis, 2008–2010. *Am J Obstet Gynecol.* 2014;210(5):435.e1–8.
- Bengjamin MI, Capitman JA, Ruwe MB. Disparities in initiation and adherence to prenatal care: impact of insurance, race-ethnicity and nativity. *Matern Child Health J.* 2010;14(4):618–24.
- Latendresse G. The interaction between chronic stress and pregnancy: preterm birth from a biobehavioral perspective. *J Midwifery Womens Heal.* 2009;54(1):8–17.
- Keiser AM, Salinas YD, DeWan AT, Hawley NL, Donohue PK, Strobino DM. Risks of preterm birth among non-Hispanic black and non-Hispanic white women: effect modification by maternal age. *Paediatr Perinat Epidemiol.* 2019;33(5):346–56.
- Kramer MS, Platt RW, Wen SW, Joseph KS, Allen A, Abrahamowicz M, et al. A new and improved population-based Canadian reference for birth weight for gestational age. *Pediatrics.* 2001;108(2):E35.

14. U.S. Department of Commerce. U.S. Census Bureau QuickFacts: United States. Accessed August 16, 2020. <https://www.census.gov/quickfacts/fact/table/US#>.
15. Zhou M. 43 rd Annual Sorokin Lecture Asians in America: The Paradox of "The Model Minority" and "The Perpetual Foreigner." University of Saskatchewan. <https://artsandscience.usask.ca/sociology/documents/43rd%20Annual%20Sorokin%20Lecture.pdf>.
16. Statistics Canada. Census Profile, 2016 Census - Ontario [Province] and Canada [Country]. Accessed August 16, 2020. <https://www12.statcan.gc.ca/census-recensement/2016/dp-pd/prof/details/Page.cfm?Lang=E&Geo1=PR&Code1=35&Geo2=PR&Code2=01&SearchText=Ontario&SearchType=Begins&SearchPR=01&B1=All&GeoLevel=PR&GeoCode=35&type=0>.
17. Dunn S, Sprague AE, Grimshaw JM, Graham ID, Taljaard M, Fell D, et al. A mixed methods evaluation of the maternal-newborn dashboard in Ontario: Dashboard attributes, contextual factors, and facilitators and barriers to use: A study protocol. *Implement Sci*. 2016;11(1):59.
18. Li Q, Keith LG, Kirby RS. Perinatal outcomes among foreign-born and US-born Chinese Americans, 1995-2000. *J Immigr Minor Health*. 2010;12(3):282-9.
19. Alexander GR, Mor JM, Kogan MD, Leland NL, Kieffer E. Pregnancy outcomes of US-born and foreign-born Japanese Americans. *Am J Public Health*. 1996;86(6):820-4.
20. Borrell LN, Rodriguez-Alvarez E, Savitz DA, Baquero MC. Parental race/ethnicity and adverse birth outcomes in New York City: 2000-2010. *Am J Public Health*. 2016;106(8):1491-7.
21. Bowers K, Laughon SK, Kiely M, Brite J, Chen Z, Zhang C. Gestational diabetes, pre-pregnancy obesity and pregnancy weight gain in relation to excess fetal growth: variations by race/ethnicity. *Diabetologia*. 2013;56(6):1263-71.
22. Provincial Council for Maternal and Child Health. Provincial Council for Maternal and Child Health Level of Care Designation. Published October 2019. Accessed August 17, 2020. <https://www.pcmch.on.ca/wp-content/uploads/2019/11/LOC-Website-List-2019.pdf>.
23. Guo Y, Miao Q, Huang T, Fell DB, Harvey ALJ, Wen SW, et al. Racial/ethnic variations in gestational weight gain: a population-based study in Ontario. *Can J Public Health*. 2019;110(5):657-67.
24. Robbins AS, Chao SY, Fonseca VP. What's the relative risk? A method to directly estimate risk ratios in cohort studies of common outcomes. *Ann Epidemiol*. 2002;12(7):452-4.
25. Wacholder S. Binomial regression in glim: estimating risk ratios and risk differences. *Am J Epidemiol*. 1986;123(1):174-84.
26. Hernán MA, Hernández-Díaz S, Werler MM, Mitchell AA. Causal knowledge as a prerequisite for confounding evaluation: an application to birth defects epidemiology. *Am J Epidemiol*. 2002;155(2):176-84.
27. Schisterman EF, Cole SR, Platt RW. Overadjustment bias and unnecessary adjustment in epidemiologic studies. *Epidemiology*. 2009;20(4):488-95.
28. Rothman K, Greenland S, Lash T. *Modern epidemiology*, 3rd edition. Lippincott Williams & Wilkins. 2008. Accessed 16 Aug 2020. <https://www.rti.org/publication/modern-epidemiology-3rd-edition>.
29. Van Buuren S. Multiple imputation of discrete and continuous data by fully conditional specification. *Stat Methods Med Res*. 2007;16(3):219-42.
30. Little RJA, Rubin DB. *Statistical Analysis with Missing Data* (2nd ed.). Wiley; Accessed 10 Aug 2020. <https://www.ebooks.com/en-us/1775204/statistical-analysis-with-missing-data/roderick-j-a-little-donald-b-rubin/>.
31. Rubin DB. *Multiple imputation for nonresponse in surveys*: Wiley; 1987.
32. Savitz DA, Janevic TM, Engel SM, Kaufman JS, Herring AH. Ethnicity and gestational diabetes in New York City, 1995-2003. *BJOG*. 2008;115(8):969-78.
33. Lee KW, Ching SM, Ramachandran V, Yee A, Hoo FK, Chia YC, et al. Prevalence and risk factors of gestational diabetes mellitus in Asia: a systematic review and meta-analysis. *BMC Pregnancy Childbirth*. 2018;18(1):1-20.
34. Edwards H, Grotegut C, Harmanli OH, Rapkin D, Dandolu V. Is severe perineal damage increased in women with prior anal sphincter injury? *J Matern Neonatal Med*. 2006;19(11):723-7.
35. Goldberg J, Hyslop T, Tolosa JE, Sultana C. Racial differences in severe perineal lacerations after vaginal delivery. *Am J Obstet Gynecol*. 2003;188(4):1063-7.
36. Goh R, Goh D, Ellepola H. Perineal tears - a review. *Aust J Gen Pract*. 2018;47(1-2):35-8.
37. Green JR, Soohoo SL. Factors associated with rectal injury in spontaneous deliveries. *Obstet Gynecol*. 1989 May;73(5 Pt 1):732-8.
38. Patel RR, Steer P, Doyle P, Little MP, Elliott P. Does gestation vary by ethnic group? A London-based study of over 122 000 pregnancies with spontaneous onset of labour. *Int J Epidemiol*. 2004;33(1):107-13.
39. Egbe A, Lee S, Ho D, Uppu S, Srivastava S. Racial/ethnic differences in the birth prevalence of congenital anomalies in the United States. *J Perinat Med*. 2015;43(1):111-7.
40. Yang Q, Wu Wen S, Caughey S, Krewski D, Sun L, Walker MC. Placenta previa: its relationship with race and the country of origin among Asian women. *Acta Obstet Gynecol Scand*. 2008;87(6):612-6.
41. Taylor VM, Peacock S, Kramer MD, Vaughan TL. Increased risk of placenta previa among women of asian origin. *Obstet Gynecol*. 1995;86(5):805-8.
42. Reddy M, Wallace EM, Mockler JC, Stewart L, Knight M, Hodges R, et al. Maternal Asian ethnicity and obstetric intrapartum intervention: A retrospective cohort study. *BMC Pregnancy Childbirth*. 2017;17(1):7.
43. Hyattsville. Health, United States, 2011: With Special Feature on Socioeconomic Status and Health - PubMed; 2012. Accessed August 16, 2020. <https://pubmed.ncbi.nlm.nih.gov/22812021/>.
44. Ghosh G, Grewal J, Männistö T, Mendola P, Chen Z, Xie Y, et al. Racial/ethnic differences in pregnancy-related hypertensive disease in nulliparous women. *Ethn Dis*. 2014 Summer;24(3):283-9.
45. Manuck TA. Racial and ethnic differences in preterm birth: a complex, multifactorial problem. *Semin Perinatol*. 2017;41(8):511-8.
46. De Wilde JA, Van Buuren S, Middelkoop BJC. Trends in birth weight and the prevalence of low birth weight and small-for-gestational-age in Surinamese south Asian babies since 1974: cross-sectional study of three birth cohorts. *BMC Public Health*. 2013;13(1):931.
47. Tutlam NT, Liu Y, Nelson EJ, Flick LH, Chang JJ. The effects of race and ethnicity on the risk of large-for-gestational-age newborns in women without gestational diabetes by Prepregnancy body mass index categories. *Matern Child Health J*. 2017;21(8):1643-54.
48. Wen SW, Kramer MS, Usher RH. Comparison of birth weight distributions between Chinese and Caucasian infants. *Am J Epidemiol*. 1995;141(12):1177-87.
49. Center for Health Statistics N. Live Births by 5-Minute Apgar Score and Age of Mother, According to Race and Hispanic Origin of Mother: Total of 48 Reporting States and the District of Columbia, 2000. Accessed August 16, 2020. <https://www.cdc.gov/nchs/data/statab/t001x28.pdf>.
50. The Centers for Disease Control and Prevention Es, Divorces, and Deaths for 1994. Accessed August 16, 2020. https://www.cdc.gov/nchs/data/mvsv/mv43_12.pdf.
51. Vangen S, Stoltenberg C, Skjaerven R, Magnus P, Harris JR, Stray-Pedersen B. The heavier the better? Birthweight and perinatal mortality in different ethnic groups. *Int J Epidemiol*. 2002;31(3):654-60.
52. Bhutani VK, Zipursky A, Blencowe H, Khanna R, Sgro M, Ebbesen F, et al. Neonatal hyperbilirubinemia and rhesus disease of the newborn: incidence and impairment estimates for 2010 at regional and global levels. *Pediatr Res*. 2013;74(SUPPL. 1):86-100.
53. Akaba K, Kimura T, Sasaki A, Tanabe S, Ikegami T, Hashimoto M, et al. Neonatal hyperbilirubinemia and mutation of the bilirubin uridine diphosphate-glucuronosyltransferase gene: a common missense mutation among Japanese, Koreans and Chinese. *Biochem Mol Biol Int*. 1998;46(1):21-6.
54. Savitz DA, Stein CR, Siega-Riz AM, Herring AH. Gestational weight gain and birth outcome in relation to Prepregnancy body mass index and ethnicity. *Ann Epidemiol*. 2011;21(2):78-85.
55. Wartko PD, Wong EY, Enquobahrie DA. Maternal birthplace is associated with LowBirth weight within racial/ethnic groups. *Matern Child Health J*. 2017;21(6):1358-66.
56. Sprague A. *Born & Growing Annual Report I 2012-2014 Two Years of Progress*; 2012. Accessed August 16, 2020. <https://www.bornontario.ca/en/publications/resources/Documents/121187-Final---english.pdf>.
57. Hayeems RZ, Campitelli M, Ma X, Huang T, Walker M, Guttman A. Rates of prenatal screening across health care regions in Ontario, Canada: a retrospective cohort study. *CMAJ Open*. 2015;3(2):E236-43.

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