Association between first birth caesarean delivery and adverse maternal-perinatal outcomes in the second pregnancy: a registry-based study in Northern Tanzania

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Abstract

Background: Caesarean delivery (CD) is the commonest obstetric surgery and surgical intervention to save lives of the mother and/or the new-borns. Despite been accepted as safe procedure, caesarean delivery has an increased risk of adverse maternal and fetal outcomes. The rising rate of caesarean delivery has been a major public health concern worldwide and the consequences that come along with it urgently need to be assessed, especially in resource limited settings. We aimed to examine the relationship between first birth caesarean delivery and adverse maternal and perinatal outcomes in the second pregnancy among women who delivered at a tertiary hospital in Northern Tanzania.

Methods: A retrospective cohort study was conducted using maternally-linked data from Kilimanjaro Christian Medical Centre. All women who had singleton second delivery between the years 2011 to 2015 were studied. A total of 5,984 women with singleton second delivery were analysed. Multivariable log-binomial regression was used to determine the association between first caesarean delivery and maternal-perinatal outcomes in the second pregnancy.

Results: Caesarean delivery in the first birth was associated with an increased risk of adverse maternal and perinatal outcomes in the second pregnancy. These included repeated CD (ARR 1.19; 95% CI: 1.05–1.34), pre/eclampsia (ARR 1.38; 95% CI: 1.06–1.78), gestational diabetes mellitus (ARR 2.80; 95% CI: 1.07–7.36), uterine rupture (ARR 1.56; CI: 1.05–2.32), peri-partum hysterectomy (ARR 2.28; CI: 1.04–5.02) and preterm birth (ARR 1.21; CI: 1.05–1.38).

Conclusion: Caesarean delivery in their first pregnancy had an increased risk of repeated caesarean delivery and other adverse maternal-perinatal outcomes in the following pregnancy. Findings from this study highlight the importance of devising regional specific measures to mitigate unnecessary primary caesarean delivery. Additionally, these findings may help both clinicians and women in deciding against or for trial of labor after previous caesarean delivery in an event of absent direct obstetric indication.

Keywords: Caesarean delivery, Maternal -Perinatal outcomes, Tanzania

Introduction

Globally, the rising rate of caesarean delivery (CD) has been a major public health concern to the public health worldwide [1]. There are wide variations in respect to the management of pregnancy in women with previous caesarean delivery and more specifically...
CD in the first birth between high-income and low-income countries [2]. Over the three decades, evidence showed a worldwide spiked rate of CD ranging from 6–40% [3], and primary CD being the most common indicator among women with previous CD [4, 5]. Most recent CD rate in Tanzania was estimated to be 6%, with an estimated rate of 11% for Kilimanjaro region while the CD rate at KCMC hospital, a tertiary hospital for Northern zone of Tanzania is reported ranging between 29.9–35.5% [6–8].

In 2015, two thirds of the global maternal deaths were reported to occur in Sub-Saharan Africa (SSA), and the maternal mortality ratio was estimated to be twice the global average of 546 per 100,000 live births [9]. The CD rate in SSA has been stagnant at 3.5% compared to the increasing CD rate globally, yet the maternal death after CD is fifty times higher compared to high income countries [9].

A major challenge is that previous CD is one of the leading indication for the repeated caesarean delivery in the subsequent delivery [10]. Inadequate counseling offered by physicians and midwives to women who had previous one CD might have been attributed to the increased rate of repeat CD. Indeed, many of these women report in labor with limited or no knowledge regarding the possibility of a trial of labor [10]. In addition to this, inability to adequately monitor the fetus and safely augment the progress of labor when the choice is to proceed with a trial of labor is challenging [10, 11].

Several studies conducted in developing countries in Europe, Asia and America have demonstrated the complex association of first CD with maternal and fetal morbidity and mortality in the second pregnancy. These include increased risks of repeated CD, preeclampsia, placenta previa, placenta abruptio, postpartum haemorrhage, uterine rupture, peri-partum hysterectomy, preterm birth, unexplained antepartum fetal death, and low birth weight [3, 12–18]. However, in most sub-Saharan Africa countries including Tanzania, where limited resources coupled with a relatively higher maternal mortality ratio is overwhelming, the extensiveness of this association has not much been widely studied. Additionally, with the increasing rate of CD in these settings, it is imperative to study the risks of the first CD which can then be communicated to the patient for their future reproductive planning. This study aimed to examine the association between first birth CD and maternal-perinatal outcomes in the second pregnancy among women who delivered at Kilimanjaro Christian Medical Centre a tertiary health hospital in Northern Tanzania.

**Materials and methods**

**Study design and setting**

A medical birth registry-based retrospective cohort study was designed using maternally-linked data from Kilimanjaro Christian Medical Centre (KCMC). A five years’ data from 1st January, 2011 to 31st December 2015 was reviewed. KCMC hospital tertiary referral hospitals and a university teaching hospital located in the Northern zone of Tanzania. The hospital receives women from the local community and referrals from the nearby regions including Arusha, Manyara, Tanga, and Singida. The average number of deliveries per year is between 4000–4800 deliveries. Over the period of 6 years (2005–2010), the rate of CD at KCMC is 29.9–35% [7].

**Data source and data collection**

The KCMC medical birth registry was established in 2000 to serve clinical, administrative and research purposes and to date, over 70,000 births have been recorded. All women who delivered at KCMC undergo a prospective interview with a standardized questionnaire within 24-h of delivery or later in case of any delivery complications. The interview is conducted by the trained midwives at the department of Obstetrics and Gynaecology. The interview is conducted by the trained midwives at the department of Obstetrics and Gynaecology. The details of the interview procedures has been described elsewhere [19]. Information regarding birth outcomes, delivery mode, obstetric history and socio-demographic is recorded in the birth registry, including information of neonates admitted to neonatal care unit. For women who delivered their first birth at KCMC, we linked the mother’s record with the child’s by unique number, which is assigned to every woman who delivers at KCMC.

**Study population**

Data for 19,670 women who delivered within the years 2011 to 2015 were obtained from KCMC birth registry. Of these, 13,211 women were multiparous. The study sample included only data for women with complete information of the second singleton delivery within the study period, and whose records were available at KCMC medical birth registry. These women were further classified into two groups; those with first birth CD and those with spontaneous vaginal delivery (SVD). Women with missing information on the mode of delivery in their first and second pregnancies were excluded. The total of 13,722 women were excluded according to our exclusion criteria. The remaining, 5,948 women had singleton deliveries in the second birth, of which 4,367(73.4%) and 1,581(26.6%) women had SVD and CD respectively in their first birth, therefore they constituted the final sample size and were analysed (Fig. 1).
Study variables and definitions

**Dependent variable**
The primary outcome variables in this study were adverse maternal and perinatal outcomes in the second pregnancy. Adverse maternal outcomes included caesarean delivery, Abruption placenta, Placenta Previa, Postpartum haemorrhage, Pre/eclampsia, Gestational Diabetes Mellitus, Uterine Rupture and Peri-partum Hysterectomy. Adverse perinatal outcomes included Apgar score less than 7 at 5th minutes, admission to a neonatal care unit, low birth weight (LBW), fetal macrosomia, preterm birth, still birth and neonatal death before 24 h.

**Independent variables**
The main exposure of interest was first birth caesarean delivery. It was defined as caesarean delivery preformed during the first pregnancy for the first time. Other covariates included maternal age which originally were recorded as continuous and then was categorized into 15–19 years, 20–29 years, 30–39 years and ≥ 40 years, maternal education, Body Mass Index (BMI) was calculated as ante-natal care booking body weight in kg/height in metres squared, then was categorized according to the World Health Organization (WHO) standard; Normal weight 18.5–24.9 kg/m², underweight < 18.5 kg/m², overweight 25.0–29.9 kg/m² and obese ≥ 30 kg/m². Gestational age at delivery was recorded in weeks as continuous, and then was categorized as 28–32 weeks, 33–36 weeks and ≥ 37 weeks. Inter-pregnancy interval was calculated from year of delivery in the first pregnancy to the year of second pregnancy which was recorded as continuous, and then was converted into months: < 24 months, 24–36 months, 37–60 months and > 60 months. Referral status was divided into three categories: home, district hospital and regional hospital. Obstetric outcomes in first birth such as Pre/eclampsia, Epilepsy, GDM, anaemia, heart disease, still birth, macrosomia, low birth weight and preterm birth were assessed in both groups.

**Statistical analysis**
Data analysis was performed using STATA version 13.0 (StataCorp.2013.College Station, TX: StataCorp LP). Descriptive statistics were summarized using mean and standard deviation for continuous variables while frequency and proportion was used to summarize the categorical variables. The chi-square test was used to determine the association between the first birth CD with baseline maternal and obstetric characteristics in bivariate analysis. Both crude relative risk (CRR) and adjusted relative risk (ARR) with 95% confidence intervals for the association between the first birth CD and maternal-perinatal outcomes in the second delivery was estimated using multivariable log-binomial regression.
model. A $p$-value of $<0.05$ was considered statistically significant.

**Ethical considerations**

The study was approved the Kilimanjaro Christian Medical University College Research and Ethics Committee (ethical clearance certificate number: 2346). Permission to use the medical birth registry data was obtained from the KCMC hospital administration.

**Results**

**Characteristics of the study participants**

The demographic characteristics of the study participants are shown in Table 1. The mean age among both groups was 28.3 (SD = 5.4) years. The women with first birth CD, 653 (41.3%) were overweight and in respect to gestation age at time of delivery, both grouped women delivered at term 3408 (78.0%) and 1187 (75.1%) for SVD and CD respectively. The inter-pregnancy interval ranged between 37–60 months among women with CD (62.4%) in the first birth which was higher compared to SVD (55.5%) group and had significant difference.

**First birth obstetric characteristic of study participants**

Table 2 displays the obstetric characteristics of the first birth among women with CD and SVD. Both grouped women delivered at term (73.1% vs. 68.8%, for those with CD and SVD in their first pregnancy) respectively. Women with CD were more likely to have Pre/eclampsia and GDM [(5.5% vs. 4.0% and (0.8% vs. 0.3%) respectively, in the first birth than those with SVD, although it was of no significance. In respect to the fetal characteristics, women with first birth CD were more likely to have macrosomia (11.0% vs.7.5%, $P<0.0001$) than the contrast group. However, the women with first birth CD were less likely to have preterm birth and stillbirths [(5.1% vs. 3.4%) and (29.0% versus 25.4%)], respectively than those with SVD.

**Association between first birth caesarean delivery and adverse perinatal outcomes in the second pregnancy**

The relationship between first birth caesarean delivery and adverse maternal outcomes has been displayed in Table 3. Women with first birth CD had an increased risk of having CD [ARR1.19 (95% CI: 1.05–1.34)]; pre/eclampsia [ARR1.38 (95% CI: 1.06–1.78)], GDM [ARR 2.80 (95% CI: 1.07–7.36)], uterine rupture [ARR1.56 (95% CI: 1.05–2.32)] and peri-partum hysterectomy [ARR2.28 (95% CI: 1.04–5.02)] in their second pregnancy as compared to their counterparts who had SVD. The association between CD in the first pregnancy and placenta abruption, placenta previa, and PPH was not statistically significant (Table 3).

**Discussion**

In the present study, CD in the first pregnancy was associated with higher risk of adverse maternal and perinatal outcomes in the second pregnancy. We found that women with initial CD had nearly two fold increased risk of repeated CD in their second pregnancy.

The finding in our study is consistent with previous studies done in China and Germany [3, 12]. The reason for high repeated CD for example in China was due to ‘two child policy’, leading to increase in maternal request for CD, to get the next precious baby. In the present study, the high repeated CD could be explained by the nature of the studied population and being conducted at the tertiary hospital that receives pregnant women in different state and conditions of labor, at which to perform CD maybe best possible form of mode of delivery for the attending physician. In addition to this the dilemma and pressure on the doctors is fact that the physician may have not attended to the patient prior and also inadequate information on the first CD before the labor.

In consistent with other studies, this study revealed that first birth CD is associated with significant increased risk of adverse outcomes: pre/eclampsia, GDM, uterine rupture and peri-partum hysterectomy. Supporting the finding to this study, specifically on pre/eclampsia, study in USA and Peru has shown three folds high and almost one half high risk respectively in the subsequent pregnancy [14, 20]. Hu et al. had contrary to this evidence, did not identify any risk in this relationship [3]. Although the mechanism underlying this association is unclear, the most likely explanation is that cesarean section scar leads to change in the endometrium; hence the pathophysiology for pre/eclampsia is supported by poor trophoblast invasion, less vascularization and incomplete remodeling of spiral arteries [21, 22]. However, in our study we could not do subgroup analysis in respect to indications of cesarean delivery especially for the first delivery and difference in race and ethnicity as incidence of pre/eclampsia is higher in African-American women [23]. Another finding is the increased risk of uterine rupture...
## Table 1  Baseline characteristics of the study participants with singleton birth in the second pregnancy (N = 5948)

| Characteristics                  | Women with first birth SVD (n = 4367) | Women with first birth CD (n = 1581) | P-value |
|----------------------------------|--------------------------------------|-------------------------------------|---------|
| **Age [Mean, SD]**               | [28.3, 5.4]                          | [28.3, 5.4]                         |         |
| Maternal Age                     |                                      |                                     |         |
| 15–19                            | 163.0 3.7%                           | 63.0 4.0%                           | 0.822   |
| 20–29                            | 2483.56.9%                          | 911.57.6%                          |         |
| 30–39                            | 1617.37.0%                          | 574.36.3%                          |         |
| ≥40                              | 104 2.4%                            | 33 2.1%                            |         |
| **Education**                    |                                      |                                     |         |
| None formal                      | 64 1.5%                             | 27 1.7%                            | 0.685   |
| Primary education                | 1844.42.2%                          | 689 43.6%                          |         |
| Secondary education              | 1126 25.8%                          | 398 25.2%                          |         |
| Higher education                 | 1333 30.5%                          | 467 29.5%                          |         |
| **Religion**                     |                                      |                                     |         |
| Christian                        | 3476 79.6%                          | 1243 78.6%                         | 0.604   |
| Muslim                           | 865 19.8%                           | 326 20.6%                          |         |
| Others                           | 26 0.6%                             | 12 0.8%                            |         |
| **Occupation**                   |                                      |                                     |         |
| Housewife                        | 517 11.8%                           | 186 11.8%                          | 0.444   |
| Farmer                           | 650 14.9%                           | 209 13.2%                          |         |
| Business                         | 1454 33.3%                          | 554 35.0%                          |         |
| Employed                         | 1376 31.5%                          | 490 31.0%                          |         |
| Others e.g. students, retired    | 370 8.5%                            | 142 9.0%                           |         |
| **Residency**                    |                                      |                                     |         |
| Urban                            | 2584 59.2%                          | 937 59.3%                          | 0.947   |
| Rural                            | 1783 40.8%                          | 644 40.7%                          |         |
| **Tribe**                        |                                      |                                     |         |
| Chagga                           | 2242 51.3%                          | 793 50.2%                          | 0.857   |
| Pare                             | 527 12.1%                           | 200 12.7%                          |         |
| Maasai                           | 93 2.1%                             | 35 2.2%                            |         |
| Others                           | 1505 34.5%                          | 553 35.0%                          |         |
| **Booking BMI (kg/m²)**          |                                      |                                     |         |
| Underweight                      | 131 3.0%                            | 70 4.4%                            | <0.0001 |
| Normal                           | 2058 47.1%                          | 649 41.0%                          |         |
| Overweight                       | 1762 40.3%                          | 653 41.3%                          |         |
| Obesity                          | 416 9.5%                            | 209 13.2%                          |         |
| [Mean, SD]                       | [24.7, 3.5]                         | [25.3, 4.4]                        |         |
| **ANC visit**                    |                                      |                                     |         |
| ≥4                               | 2882 66.0%                          | 1057 66.9%                         | 0.535   |
| 4                                | 1485 34.0%                          | 524 33.1%                          |         |
| **Gestation age at delivery**    |                                      |                                     |         |
| 28–32                            | 130 3.0%                            | 49 3.1%                            | 0.047   |
| 33–36                            | 829 19.0%                           | 345 21.8%                          |         |
| ≥37                              | 3408 78.0%                          | 1187 75.1%                         |         |
| [Mean, SD]                       | [37.9, 2.3]                         | [37.8, 2.3]                        |         |
| **Inter-pregnancy Interval**     |                                      |                                     |         |
| < 24 months                      | 15 0.3%                             | 7 0.4%                             | <0.0001 |
| 24–36 months                     | 1200 27.5%                          | 338 21.4%                          |         |
| 37–60 months                     | 2423 55.5%                          | 987 62.4%                          |         |
| > 60 months                      | 729 16.7%                           | 249 15.7%                          |         |
Worth mentioning, is higher number of uterine ruptures with limited peri-partum hysterectomy. This disparity is possibly due to repairs of some uterine rupture which were not captured in the registry [13, 14]. Despite uterine scar being a major risk factor for uterine rupture, unscarred uterus may similarly present with uterine rupture as observed in the current study. Previous studies have documented similar findings [24, 25].

Furthermore, this study also found association of first birth CD and increased risk of developing GDM and peri-partum hysterectomy in the second pregnancy. Although this association is statically significant, the explanation that could account for this, especially for GDM is the small number of events, thus wide confidence interval. Conversely, Hu et al. in China had similar association with larger events [3]. Similar to pre/eclampsia, GDM could be associated with increasing placenta mass which directly influenced anti-insulin hormones production [23].

In relation to maternal outcomes, our findings show that there were some association between first birth CD and placenta abruption, placenta previa and PPH, however not significant, which is consistent with other analysis in respect to placenta previa and PPH [13], however contrary to Hu et al. and Kennare et al. [3, 16]. Possible explanation that may have influenced our finding might be missing information on complication in the first birth which may have influenced the current pregnancy and furthermore our data did not classify the types of placenta previa.

On the aspect of perinatal outcomes, we found that, women with first birth CD had 1.21(CI 95%: 1.05–1.38) higher risk of having preterm birth in the second pregnancy similar to a systemic review and meta-analysis, which involved ten retrospective cohort studies with more than ten million women, showing CD in the first pregnancy increased risk of preterm birth (ARR 1.12, 95%CI 1.01–1.24) in the subsequent pregnancies [26]. This association is explained by the changes in the intra-uterine structure and its microenvironment, although the pathogenesis of this event is unclear [26]. However, inability to control for other possible confounders such as

| Table 1 (continued) |
|---------------------|
| Characteristics     | Women with first birth SVD | Women with first birth CD | P-value |
|                     | (n = 4367) | n | % | (n = 1581) | n | % |       |
| Referral            |           |   |   |           |   |   |       |
| Home                | 3508       | 80.3 | | 1227       | 77.6 | | 0.566 |
| District hospital   | 707        | 16.2 | | 289        | 18.3 | |       |
| Regional hospital   | 152        | 3.5  | | 65         | 4.1  | |       |

| Table 2 First birth obstetric characteristic of study participants (N=5948) |
|-------------------------------------------------------------|
| Characteristics | Women with first birth SVD | Women with first birth CD | P-value |
|                 | (n = 4367) | (n = 1581) |       |
| Maternal Characteristics |       |           |       |
| Gestation age |           |           |       |
| 28–32         | 95        | 2.2       | 25     | 1.6 | 0.005 |
| 33–36         | 1269      | 29.1      | 401    | 25.4 |       |
| ≥37           | 3003      | 68.8      | 1155   | 73.1 |       |
| Medical Conditions |       |           |       |
| Pre/Eclampsia |           |           |       |
| Yes           | 173       | 4.0       | 87     | 5.5  | 0.010 |
| No            | 4194      | 96.0      | 1494   | 94.5 |       |
| Epilepsy      |           |           |       |
| Yes           | 37        | 0.8       | 11     | 0.7  | 0.564 |
| No            | 4330      | 99.2      | 1570   | 99.3 |       |
| GDM           |           |           |       |
| Yes           | 11        | 0.3       | 13     | 0.8  | 0.002 |
| No            | 4356      | 99.7      | 1568   | 99.2 |       |
| Anaemia       |           |           |       |
| Yes           | 48        | 1.1       | 20     | 1.3  | 0.595 |
| No            | 4319      | 98.9      | 1561   | 98.7 |       |
| Heart Disease |           |           |       |
| Yes           | 47        | 1.1       | 20     | 1.3  | 0.542 |
| No            | 4320      | 98.9      | 1561   | 98.7 |       |
| Fetal characteristics |       |           |       |
| Still birth   |           |           |       |
| Yes           | 222       | 5.1       | 54     | 3.4  | 0.007 |
| No            | 4145      | 94.9      | 1527   | 96.6 |       |
| Macrosomiaa   |           |           |       |
| Yes           | 326       | 7.5       | 174    | 11.0 | <0.0001 |
| No            | 4041      | 92.5      | 1407   | 89.0 |       |
| Low birth weightb |       |           |       |
| Yes           | 371       | 8.5       | 138    | 8.7  | 0.776 |
| No            | 3996      | 91.5      | 1443   | 91.3 |       |
| Preterm birth |           |           |       |
| Yes           | 1268      | 29.0      | 401    | 25.4 | 0.005 |
| No            | 3099      | 71.0      | 1180   | 74.6 |       |

* Birth weight ≥ 4000 g; bBirth weight ≤ 2500 g
premature rupture of membranes, infections or cigarette smoking which could lead to preterm delivery was a limitation that could have given different view to our finding. Also the recorded preterm birth was not categorized as either induced or spontaneous and on the other hand, availability of neonatal care facility at KCMC could influence the results in our study, as most of women with preterm labour are referred to KCMC for preterm delivery and neonatal care.

**Strengths and limitations**

Our study included data for five years; hence large scale sample size. Our analysis was strictly restricted to women with first and second singleton pregnancies, thereby eliminating potential confounding effect of parity and multiple gestation pregnancy. Being a secondary data from birth registry, we could not capture many important factors that could have helped us in better analysis, such as indications of previous and current cesarean delivery, history of myomectomy, history of previous placental abruption, where was the first cesarean delivery performed, inability to capture the women with GDM who were tested for OGTT, possibility of leading to inappropriate documentation of certain clinical condition, such as birth-weight, gestational age. These findings may not reflect the situation in the other settings and population in whole, as KCMC being a tertiary hospital, may be leading to referral bias and information retrieved.

Table 3  Association between first birth caesarean delivery and adverse maternal outcomes in the second pregnancy ($N=5948$)

| Maternal outcomes          | Women with first birth SVD ($n = 4367$) | Women with first birth CD ($n = 1581$) | CRR (95% CI)         | ARR (95% CI) |
|----------------------------|----------------------------------------|--------------------------------------|----------------------|--------------|
| Cesarean delivery          | 1568(35.9)                             | 648(41.0)                            | 1.24(1.10–1.39)      | 1.19(1.05–1.34)$^a$ |
| Placental abruption        | 73(1.7)                                | 32(2.0)                              | 1.22(0.80–1.85)      |              |
| Placental previa           | 80(1.8)                                | 37(2.3)                              | 1.28(0.87–1.90)      |              |
| PPH                        | 291(6.7)                               | 122(7.7)                             | 1.17(0.94–1.45)      |              |
| Pre/eclampsia              | 207(4.7)                               | 109(6.9)                             | 1.49(1.17–1.89)      | 1.38(1.06–1.78)$^b$ |
| GDM                        | 15(0.3)                                | 14(0.9)                              | 2.59(1.24–5.38)      | 2.80(1.07–7.36)$^b$ |
| Uterine rupture            | 82(1.9)                                | 43(2.7)                              | 1.46(1.01–2.12)      | 1.56(1.05–2.32)$^b$ |
| Peripartum hysterectomy    | 18(0.4)                                | 15(0.9)                              | 2.31(1.16–4.60)      | 2.28(1.04–5.02)$^a$ |

CRR Crude Relative Risk, ARR Adjusted Relative Risk, CI Confidence Interval

$^a$ Adjusted by Maternal age, Booking BMI, Gestation age, Inter-pregnancy interval, also current and previous pre/eclampsia and GDM

$^b$ adjusted by Maternal age, Gestational age, Booking BMI, Inter-pregnancy interval, previous GDM, and pre/eclampsia

$^c$ Reference group

Table 4  Association between first birth caesarean delivery and adverse perinatal outcomes in the second pregnancy ($N=5948$)

| Adverse outcomes | $^b$Women with first birth SVD ($n = 4367$) | Women with first birth CD ($n = 1581$) | CRR(95%CI)         | ARR(95%CI) |
|------------------|--------------------------------------------|---------------------------------------|--------------------|------------|
| Preterm birth    | 959(22.0)                                  | 394(24.9)                             | 1.18(1.03–1.34)    | 1.21(1.05–1.38)$^a$ |
| Neonatal unit admission | 650(14.9)       | 219(13.9)                             | 0.92(0.77–1.08)    | 0.90(0.74–1.09)$^b$ |
| Macrosomia       | 257(5.9)                                   | 72(4.6)                               | 0.76(0.58–0.99)    | 0.72(0.55–0.95)$^c$ |
| Low birth weight | 417(9.5)                                   | 165(10.4)                             | 1.10(0.91–1.33)    | 0.96(0.78–1.18)$^d$ |
| Low Apgar score in 5 min ($n=5771$) | 89(2.1)                      | 36(2.3)                               | 1.12(0.75–1.65)    | 1.06(0.71–1.58)$^e$ |
| Still birth      | 128(2.9)                                   | 49(3.1)                               | 1.06(0.75–1.47)    | 0.95(0.67–1.36)$^f$ |
| Neonatal death in 24 h | 35(0.8)                    | 20(1.3)                               | 1.58(0.91–2.75)    | 1.72(0.73–4.03)$^g$ |

CRR Crude Relative Risk, ARR Adjusted Relative Risk, CI Confidence Interval

$^a$ adjusted by the current preeclampsia GDM Macrosomia, LBW, and previous preterm

$^b$ adjust by Preeclampsia, GDM, Macrosomia, LBW, Apgar score in 5 and Preterm

$^c$ adjusted by macrosomia in the first birth, preeclampsia, GDM, stillbirth, and preterm in the second birth

$^d$ adjusted by LBW in the first birth, preeclampsia, GDM, and Preterm in the second birth

$^e$ adjusted by preeclampsia, GDM, LBW, Preterm, abortion placenta in the second birth

$^f$ adjusted by stillbirth in the first birth, preeclampsia, GDM, Macrosomia, Preterm and abortion placenta in the second birth

$^g$ adjusted by preeclampsia, GDM, macrosomia, LBW, Apgar score in 5 Preterm, and Abruptio placenta in the second birth

$^h$ Reference group
from birth registry as a secondary data may be limited in its completeness. A multi-centred prospective data collection may enrich the quality of information and true reflection of the findings.

Conclusion and recommendation
In view of our findings, caesarean delivery in the first birth appears to increase the risk of a repeated caesarean delivery and other adverse outcomes in the second pregnancy. We emphasise clinicians to try to balance the risks and benefits of caesarean delivery in the first and future births. However, we should ensure adequate counselling of the women during ante-natal visits on risks versus benefits of and caesarean delivery and vaginal birth after caesarean delivery.

Abbreviations
ANC: Ante-natal care; BMI: Body Mass Index; CD: Caesarean delivery; CI: Confidence Interval; GDM: Gestational Diabetes Mellitus; KCMC: Kilimanjaro Christian Medical Centre; LBW: Low birth weight; PPH: Post-partum Haemorrhage; SSA: Sub-Saharan Africa; SVD: Spontaneous vaginal delivery.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s12884-022-04719-7.

Additional file 1.

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Authors’ contributions
RG and EM conceived and designed the study. MJM and BM had full access to the data. Data validation, data error and inconsistency check and coding was done by RG and MJM. Statistical analysis was performed by RG and MJM. RG wrote the first draft of the manuscript. EM, BM, MJM, BA and LLN reviewed the manuscript. All authors read and approve the final manuscript.

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Availability of data and materials
The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

Declarations
Ethics approval and consent to participate
The Ethical Clearance was approved by Health Research and Ethics committee of Kilimanjaro Christian Medical University College, with ethical clearance certificate No. 2346. As part of the KCMC birth registry protocol, the interviews are usually conducted after the delivery. Oral information is given by the midwives before conducting the interview regarding the birth registry and the data collected and the use of data for research purposes. A verbal informed consent is sought from the adult participants and the parent(s) or guardian of all under 16 years and they are free to opt not to reply any specific questions during the interview. All other methods were carried out in accordance with relevant institutional ethical approval guidelines and regulations.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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