

Rates and Determinants of Successful Vaginal Birth after a Previous Caesarean Section: A Prospective Cohort Study

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Abstract

Objectives: To determine the rates and determinants of successful vaginal birth after a previous caesarean section (VBAC) among pregnant women with a previous caesarean section (CS).

Methods: This was a prospective cohort study of pregnant women, who attempted VBACs following one previous lower segment caesarean delivery in Nnamdi Azikiwe University Teaching Hospital (NAUTH), Nnewi, south-east Nigeria. Exclusion criteria were multiple gestations, more than one previous CS, or recurrent indications (clinical evidence of pelvic contraction, previous classical CS, and previous repair of vaginal fistula) for CS. Analysis was done with Epi info 2013 version 7.0 for Windows using descriptive and inferential statistics at 95% level of confidence.

Results: During the study period, 74 women were assessed for eligibility, but nine were excluded with 65 of the 74 (87.8%) women being allowed trial of labour after one caesarean section (TOLAC). Of the 65 women that were allowed TOLAC, 22(33.8%) had successful vaginal delivery, while 43 (66.2%) had failed VBAC; giving a ratio of 1:3. Failed VBAC constituted 10.1% (43/424) of the total CS performed during the study period and the major indications were fetal distress and suspected macrosomia. Maternal age, booking status, socio-economic status, method of onset of labour, gestational age at delivery and birth weight had no significant association with success of VBAC ($p > 0.05$).

Only multiparity, history of successful vaginal delivery before and/or after first caesarean delivery had significant association with successful VBAC ($P < 0.05$). The APGAR scores of the babies that had successful VBAC were not significantly different from those of failed VBAC ($P > 0.05$). There were no maternal death, but there were 3 perinatal deaths, giving a perinatal mortality rate of 2.86/1000. None of the babies with successful VBAC suffered perinatal death.

Conclusion: Approximately one in every three women undergoing TOLAC had successful VBAC and significant determinants were multiparity, history a previous vaginal delivery before and/ or after the first CS. Proper and optimal antenatal evaluation of possible favourable factors for successful VBAC is needed.

Keywords: VBAC; multiparity; TOLAC; vaginal delivery

1. INTRODUCTION

Caesarean delivery has remained an important surgical intervention in contemporary obstetric practice [1-3]. Worldwide, primary and repeat caesarean delivery rates have reached their highest levels [4-6]. Even though, variation exists in the rates across countries; currently the

rate ranges from 10% to 40% [6-8]. Due to increased risk of maternal complications on repeat caesarean section (CS) and safety of vaginal birth after caesarean section (VBAC), trial of labour for selected group of patients with a previous lower segment transverse scar has become a favorite strategy [7].

In 1988, American College of Obstetrician-Gynaecologist (ACOG) recommended that, in the absence of a contraindication, a woman with one previous low-transverse caesarean delivery be counseled to attempt labour in a subsequent pregnancy [7, 9]. ACOG further suggests that a woman with a twin pregnancy or a woman who requires induction of labour may also be considered candidates for vaginal birth after caesarean section (VBAC) with appropriate counseling [9].

In general, 60 to 80% success rate of VBAC has been reported by many authors if the primary caesarean was done for nonrecurring indications [3]. Some of the nonrecurring indications for CS include poor progress of labour, fetal distress, cord prolapse, placenta previa, transverse lie, breech presentation, pregnancy induced hypertension and multiple pregnancy [9,10].

Nevertheless, there is considerable variation in the proportion of women who are offered and attempted VBAC across centres. British figures indicate that among women with a prior caesarean section, 33% will successfully achieve vaginal birth in the subsequent pregnancy and institutional variations ranged from 6% to 64% [11, 12]. The leading indications for the repeat CS were: failure to progress, fetal distress and scar tenderness [13, 14].

The intriguing questions are how to reliably predict successful VBAC, and how to determine and quantify the magnitude of the risk of failure that is acceptable to women. Many studies have addressed methods for identifying women at low and high risk of failure of an attempted VBAC but none of them has resulted in a validated result [1-3]. Even those factors found to be associated with successful VBAC vary from centre to centre. Currently, therefore, there is no single validated tool which holds true for all to predict successful vaginal birth among women with a prior caesarean delivery.

In a previous retrospective study in NAUTH, Nnewi, in Nigeria, Ikechebelu et al [15] reported a successful vaginal delivery of 46.7% among women that underwent trial of labour after one caesarean section (TOLAC) between 2001 and 2005 while Obiechina et al [16] reported successful vaginal delivery of 59.5% among women that underwent TOLAC between 2002 and 2006. As at the time of their study, the average CS rate in NAUTH was 18.4% but since then the CS rate has been rising such that Eleje et al [17] in 2010 reported a CS rate of

26.9% with one previous CS scar being the commonest indication. Therefore, while it does appear that the rate of successful VBAC is increasing over the years in Nnewi, Nigeria, there is unexpected corresponding increase in CS rate with failed VBAC being the most common culprit. Additionally, CS rate may be increasing in Nigeria because of the flourishing private hospitals in major towns and possibly increasing cases of litigation. Even though teaching hospitals offer trial of labour for mothers with one scar, there is no prospective study done which shows the rate of VBAC acceptance and success in Nigerian hospitals.

Therefore, the objective of the present study was to prospectively identify maternal demographic, past and present obstetric determinants of successful VBAC in teaching hospital in Nnewi, Nigeria. This will be of great help for obstetricians in the joint obstetrician-patient decision while offering TOLAC in Nigeria.

2. SUBJECTS AND METHOD

2.1. Study Area

This study was conducted in Nnamdi Azikiwe University Teaching Hospital (NAUTH), Nnewi, south-east Nigeria. This teaching hospital provides a 24 hour specialty care. Most of the deliveries and evaluations are made by nurse midwives, resident doctors with a consultant supervision. NAUTH offers trial of labour (TOL) with informed consent from the women if the following conditions are fulfilled: the mother has one previous lower uterine segment scar, nonrecurring previous indications, singleton pregnancy, cephalic presentation, estimated fetal weight less than or equal to 4.0 kg, and no current indication for CS.

2.2. Study Period

This study included mothers accepting a TOL and delivering in the time period of data collection from July 1st 2013 to June 30th 2014.

2.3. Study Design

This study was a prospective cohort study.

2.4. Study Population

This included all consenting pregnant women seen in the labour ward during the period of the study and who met the inclusion criteria.

2.5. Inclusion Criteria

The inclusion criteria included consenting pregnant women with one previous lower

uterine segment CS scar who came for induction of labour or with spontaneous labour or leakage of liquor, with no contraindication for VBAC and allowed to undergo trial of labour by the managing obstetrician and delivered through the vaginal route or emergency CS due to failed VBAC.

2.6. Exclusion Criteria

Those women who had multiple gestations and women with more than one previous CS were excluded. Women who had CS for a reason that would typically lead to another caesarean delivery in subsequent pregnancies (recurrent indications) were also excluded. Such recurrent indications included women with clinical evidence of pelvic contraction, previous classical CS, and previous repair of vaginal fistula.

2.7. Data Collection Method

Factors associated with successful VBAC among mothers with one previous caesarean section were identified. Women were allowed for VBAC according to the hospitals protocol such as women having spontaneous/ induced onset of labour and delivered by vaginal route. Ethical clearance was obtained from NAUTH Ethics committee. Women for the study were interviewed and data collected immediately after the patients arrived at labour ward or prenatal ward for delivery. All women included in the study were followed through delivery and for at least 48 hours after delivery. Information was sought directly from the women and recorded in a structured proforma which included maternal socio demographic, past and present obstetric experience, mode of delivery and birth outcomes variables.

2.8. Variables

Socio demographic variables included: maternal age, marital status, parity, gestational age and booking status. The past obstetric variables included: gestational age at labour or delivery, history of prior successful VBAC and spontaneous vaginal delivery (SVD).

The current obstetric and fetal factors included: status of the baby at admission of the women in the labour ward, birth weight and outcome of the baby. A failed TOLAC was defined as a repeat caesarean delivery after spontaneous onset of labour or induction or stimulation of labor, while a successful TOLAC was defined as a vaginal birth after CS. Induction of labour

is defined as “the initiation of uterine contractions before the spontaneous onset of labour by medical and/or surgical means for the purpose of delivery,” which includes births induced by oxytocin. For the cases of induction of labour, cervical ripening was done by intra cervical Foley catheter.

2.9. Data and Statistical Analysis

Data were entered after checking completeness, cleaning and coding into EXCEL Spread sheet and analysed using computer EPI-Info 2013 version 7 (v 7; Epi Info, Centers for Disease Control and Prevention, Atlanta, GA). Proportions were compared by Students’ t-test, Chi-square or Fisher exact test where appropriate and the statistical significance of *P*- value was *P*<0.05.

3. RESULTS

During the study period, a total delivery of 1050 occurred. Of the 1050 deliveries, 424(40.4%) women had CS from various indications while 626 (59.6%) women had vaginal delivery, giving a CS rate of 40.4%. Also during the study period, 74 women were recruited. Nine of the recruited women who had elective CS were excluded from the analysis because the women did not undergo TOLAC due to recurring indications, thereby yielding a final achieved sample of 65. This means that 65 of the 74 (87.8%) women were allowed TOLAC.

Table1. *Socio-Demographic Characteristics of the Women*

Variables	Frequency (N=65)	Percentage
Age		
20-24	9	13.8
25-29	25	38.5
30-34	23	35.4
35-39	7	10.8
40-44	1	1.5
Booking Status		
Booked	47	72.3
Un-booked	18	27.7
Parity		
1	31	47.7
2-4	32	49.2
≥5	2	3.1
Educational level		
Primary	1	1.5
Secondary	43	66.2
Tertiary	21	32.3
Social Class		
I	19	29.2
II	40	61.6
III	6	9.2

The mean age of the respondents was 30.5±5.1 years (range=20-44 years). All the women were married and majority, 43 (66.2%) had secondary level of education. The socio-demographic characteristic of the respondents' is shown in table 1.

Of the 65 women, 22(33.8%) had successful vaginal delivery after TOLAC while 43 (66.2%) women had failed VBAC and subsequently had repeat CS; given a ratio of 1:3. Therefore, failed VBAC constituted 10.1% (43/424) of the total CS performed during the study period.

Of the 65 women, 59 (90.8%) women had spontaneous onset of labour while six (9.2%) women had induced labour. Of the six women that had induction of labour, 5 (83.3%) had Foley catheter inserted intracervically for cervical ripening while one (16.7%) unbooked and referred case had misoprostol used for cervical ripening. Only one woman (unbooked) received misoprostol for cervical ripening had uterine rupture.

The mean age of the women with successful VBAC was 31.1±4.1 years (range= 22-44 years) while the mean age of those who had failed VBAC was 30.0±4.9 years (range=20-44 years). The observed difference was not statistically significant (t = 0.70; p = 0.109). Similarly, other factors such as booking status, socio-economic status, method of onset of labour, gestational age at delivery and birth weight had no significant association with success of VBAC (p > 0.05). The association between the successful VBAC and women's socio-demographic, maternal and fetal characteristics is shown in table 2.

Only multiparity, history of successful vaginal delivery before first CS and history of successful vaginal birth after the first CS had significant association with successful VBAC (P <0.05).

There were three cases (1.5%) of uterine rupture (one booked and two unbooked). Two were admitted as cases of uterine rupture with intrauterine fetal death following a trial of labour at home. One patient ruptured her uterus while on the labour ward and her baby was, however, alive. One of the cases (unbooked) had cervical ripening with misoprostol admission at the study hospital.

Table2. Association between Successful VBAC and Women and Fetal' Characteristics

Variables/subgroup	Successful VBAC Group (N=22)	Failed VBAC group (N=43)	P-value
Mean Age	31.1±4.12 years	30.0±4.85 years	0.534
Booking Status			
Booked	19	28	0.061
Unbooked	3	15	
Parity			
Primiparous	5	21	*0.037
Multiparous	17	22	
Socio-economic Class			
High	20	39	0.670
Low	2	4	
Method of Onset of labour			
Spontaneous	20	39	0.670
Induced	2	4	
Previous Successful Vaginal Delivery before the First CS			
Yes	13	4	*<0.001
No	9	3	
Previous Successful VBAC			
Yes	13	3	*<0.001
No	9	39	
Birth Weight of the baby (Kg)			
≤3.5	18	28	0.132
>3.5	4	15	
APGAR Score in 1 min			
≤6	1	3	0.583
>6	21	40	
Uterine Rupture			
Yes	0	3	0.283
No	22	40	
Gestational age at Delivery			
≥40 weeks	10	21	0.502
<40 weeks	12	22	

Table3. Indications for caesarean section in women that had failed VBAC

Indications	Frequency	Percentage
Fetal distress	11	22.9
Suspected macrosomia	11	22.9
Malpresentation	7	14.5
Cephalopelvic disproportion	5	10.4
Borderline pelvis	4	8.3
Imminent uterine rupture	3	6.3
Obstructed labour	3	6.3
Severe oligohydramnios	2	4.2
Cervical stasis	2	4.2
Total	48	100.0

*Five women had more than one indication.

No maternal death was seen in this study. However, there were three perinatal deaths, two of which were intrauterine fetal deaths before admission. All the three perinatal deaths occurred among patients that had failed VBAC. The clinical cause of death was uterine rupture, giving perinatal mortality rate of 2.86/1000. None of the patients that had successful VBAC suffered perinatal death. The APGAR scores of the babies delivered by successful VBAC were not significantly different when compared to babies delivered by emergency CS due to failed VBAC.

4. DISCUSSION

The route of delivery after one previous lower segment CS represents one of the most significant and challenging debates in contemporary obstetric practice [18]. This study has shown that the CS rate in the institution during the study period was 40.4%. CS from failure of successful vaginal birth after one previous CS constituted 10.1% of all cases of CS and 4.1% of all deliveries. A high caesarean section rate of up to 40.4% has also been documented in other studies in Nigeria [4, 5, 18]. This higher rate of caesarean delivery could be explained by the fact that our health care institution is a referral hospital, where many high-risk pregnancies are seen and managed. This is because 87.8% of the women with previous caesarean delivery were allowed a trial of vaginal delivery. Similar high incidences have also been reported by other authors [4, 18-20].

The successful vaginal delivery rate of 33.8% was lower than the range of 60 – 80% reported in the literature [4, 6, 20, 21]. It was also lower than previous study by Ikechebelu et al [15] (46.7%) and Obiechina et al [16] (59.5%) in Nnewi, Nigeria, the location of the hospital of study. However, a similar reduced successful vaginal delivery rate has also been reported by other published reports [21-24]. A dramatic decrease in the rate of VBAC has also been observed in the United States during recent years from 28.3% in 1996 to 9.2% in 2004 [25]. The reduced successful vaginal delivery rate in this study could be a reflection of the increasing trend in the incidence of CS including increasing rate of litigation, the high number of cases of patients who had emergency CS for fetal distress, suspected macrosomia and malpresentations. CS rate may have increased due to use of continuous electronic fetal

monitoring, which was not frequently used a decade earlier. Thus, this reason may explain why the commonest indication for caesarean section was fetal distress. The lower rate was due partly to the fact that trial of vaginal delivery was offered to patients who met the selection criteria, and partly because patients with some indications including fetal macrosomia, malpresentations and placenta praevia who should have elective surgery presented as an emergency in the labour ward, when they could not deliver at home. Patients should therefore be selected on individual merit for trial of vaginal birth.

As seen in this study, successful vaginal delivery has been found to be significantly influenced by prior history of previous vaginal delivery before and after caesarean delivery as reported in various published study [18,19]. It is surprising that there is no significant association between the outcome of trial of vaginal birth after CS and the booking status, socio-economic status, method of onset of labour, gestational age at delivery and birth weight of the babies ($p > 0.05$). These findings contrasts with previous reports in Nigeria by Ilesanmi et al [26], Ola et al [19] and Iyoke et al [4] where spontaneous onset of labour, birth weight of less than 4 kg, and gestational age of less than 40 weeks predicts successful trial of VBAC. This lack of association could be because approximately 30% of women were unbooked and were admitted as emergency after having tried VBAC in a peripheral hospital where the women selection would have been suboptimal. Also their labour would have been either unsupervised or supervised by unskilled and untrained personnel who only ask them out when the situation is bad.

Thus, some of these women would have undergone trial of VBAC even when they ought not to have attempted it in the first place. In our society, there is strong aversion for CS and usually there is general erroneous belief that there is a likelihood of repeat CS if they labour in a hospital with operative facility. Much to our chagrin, some women in this study who had their antenatal care in our facility and received proper counseling absconded to places without maternity services at the mention of possible CS only to present again in the study hospital when things were not progressing as they expected.

Previous published reports in the literature [18, 25] have suggested a significant risk of uterine

rupture with poor outcome for both the woman and her infant following vaginal delivery. Unfortunately, up to date, there are no reliable ways to predict the trial of labour after one CS that will result in emergency CS or successful vaginal birth [5, 9].

In this study there were three (1.5%) cases of uterine rupture (one booked and two unbooked). Two were admitted as cases of uterine rupture with intrauterine fetal death following a trial of labour at home. One patient ruptured her uterus while in the labour ward and her baby was, however, alive. One of the cases (unbooked) had cervical ripening with misoprostol admission at the study hospital. There was salvage for baby whose mother ruptured her uterus in the labour ward compared with mothers who ruptured before admission, because they were unbooked and chose to labour at home. Although, three patients that had induction of labour received intracervical Foley catheter for cervical ripening, ACOG and RCOG agree that induction of labour for maternal or fetal indications is an option for women undergoing TOLAC. There is further agreement that when informing a woman about induction and/or augmentation, clear information should be provided on all potential risks and benefits, especially the potential increased risk of uterine rupture and the potential decreased possibility of achieving VBAC.

Major complications have also been reported following the use of misoprostol [5, 27, 28]. Even though this study reported one uterine rupture following cervical ripening using misoprostol, uterine ruptures were also noted in women who were not augmented, stimulated or induced.

Although previous studies in Nigeria and its sub region had reported maternal mortality in women undergoing TOLAC, the case fatality rate shown from our cohort was 0%. This finding was similar to report by Okpere et al [29] in Benin City, Nigeria. However, the perinatal mortality rate was 2.86/1000. As shown in this study, results of studies on the impact of mode of delivery on low APGAR scores are discordant and do not justify any conclusion. Still, most studies have not found that admission to the intensive care unit differs according to mode of delivery [30, 31]. The APGAR score of babies delivered by emergency caesarean section was not statistically different from that had successful vaginal delivery.

Similarly, Okpere et al. [29] in his earlier report revealed higher perinatal deaths. A number of factors may be responsible to this improved trend. Continuous electronic fetal monitoring and intrapartum care for the duration of planned VBAC to enable prompt identification and management of uterine scar rupture as strongly recommended by ACOG is being practiced in our hospital.

This study has a number of limitations. It is important to acknowledge that this study does not answer the question regarding optimal approach to delivery for women with prior caesarean who must be delivered for maternal or fetal indications. This study has relatively limited sample size and did not compare women undergoing TOLAC with those undergoing planned repeat caesarean section. Such a comparison is needed for further study. We also recognise that categorising the indication for the previous caesarean section into 'recurrent and non-recurrent indications' may be limited by the fact that the obstetrician may not have managed the previous delivery and the previous labour record may not be readily volunteered by the women studied. This is because; recurrent indication for the previous caesarean section is known to be associated with a comparatively lower rate of successful vaginal delivery.

5. CONCLUSION

This study has shown that approximately one third of the women with one previous caesarean section had successful vaginal birth after TOLAC. Significant determinants of successful VBAC in the cohort of women studied were multiparity, history a previous vaginal delivery before and after the first caesarean section VBAC was not risk free for the fetus. It is recommended that consultation to these women should include evaluation for the likelihood of successful TOLAC and the risk of uterine rupture based on the many other previously reported risk factors in order to stratify the risk associated with TOLAC and to allow optimal patient selection.

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