Assessing the pros and cons of vaginal birth after caesarean relative to elective repeat caesarean section

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Received: 15 February 2020
Accepted: 11 March 2020

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ABSTRACT

Background: Recent years have witnessed a rise in rate of primary caesarean section (CS). No. of women reporting with a previous CS scar is also increasing. Judicious trial of labor in such patients can prevent repeat caesarean section. Aim of this study was to assessing the safety and success rate of vaginal birth after caesarean (VBAC) in selected cases of patients who have undergone previous lower segment CS (LSCS) is the main aim of this study.

Methods: In this prospective observational study carried out in a tertiary care teaching hospital over a period of 1 year. 375 pregnant women with a history of one previous LSCS for non-recurrent indications were enrolled. The statistical technique of t-test was administered for relative comparison with respect to maternal and neonatal complications across the two groups, i.e. repeat LSCS and vaginal delivery.

Results: Out of 375 patients 187 patients (49.9%) underwent elective LSCS for recurrent indication and for non-recurrent indication associated with some complicating factor. Trial of labor in 188 (50.10%) was given out of which 59.3% had spontaneous vaginal delivery,7.20% had instrumental delivery and 33.50% landed into emergency CS. Commonest cause of Em. LSCS being Fetal distress. As regards maternal complications, no statistically significant difference was found between the Repeat LSCS and Vaginal delivery groups (t = 0.779, p > 0.05). On similar lines, there was no statistically significant difference across both groups as regards neonatal complications (t = 0.632, p > 0.05).

Conclusions: Taking into account the increased trend of primary CS, trial of VBAC in selected cases is very important. It can be concluded that VBAC has chances of success in cases with previous one LSCS but it must be carefully investigated and monitored.

Keywords: Elective caesarean section, Lower segment cesarean section, Vaginal birth after cesarean section

INTRODUCTION

To start with, this paper is hinged on the dictum as quoted by the eminent Edwin Bradford Cragin, an obstetrician in 1916 that is, - “Once a caesarean, always a caesarean” and it is true historically.1,2 But now there are other perspectives that call in for discussion and so it has formed the crux of this research study. Of late there has been a rapid increase in the rate of C-section, in India and world across, which may be unwarranted despite the medical complications that might ensue after the C-section. The World Health Organization has stated that caesarean rates higher than 10- 15% indicate unnecessary maternal risk. This paper also presents a critique analyzing the risks and benefits associated with VBAC. It argues the case in point whether vaginal birth after CS (VBAC) might help in reducing the rates of C-section.

All post caesarean pregnancies do not require repeat CS and a majority of them may have uncomplicated vaginal
delivery. In fact, a trial of vaginal birth after a previous CS (VBAC) is considered safer than a routine repeat CS. VBAC has distinct advantages over a repeat CS. However, in cases of failed trial of labor there are increased chances of maternal and perinatal morbidity &mortality. However, trial of labor is also not bereft with complications which should be taken into account. Basic prerequisite being the availability of a well-equipped emergency care in order to deal with any risk /complications which might arise during trial of labor.

The pertinent implication for practitioners is to answer the question as to whether a previous caesarean is a medical indication for a repeat caesarean. And if the answer is “NO” then, what is the prerequisites for successful subsequent deliveries in such cases. It is for the physicians to predict the cases for VBAC based on evaluation of factors that significantly influence the risk of failed attempted VBAC, such as, obstetric history, e.g. uterine scar type (lower segment/classical), uterine closure technique (single/double layer closure), number of previous caesareans. The indication for caesarean, prior vaginal births, interval between 2 deliveries, and maternal age, in addition to other factors, such as current labor management, including induction or augmentation of prostaglandins or oxytocin.2-4

In view of this, trial of vaginal delivery in women with post caesarean pregnancy remains controversial and continuous critical audit of the trends is imperative. Women and their relatives should be informed and counselled regarding the safety and the risk involved in both the modes of delivery†.

Risks and benefits as per delivery method

Benefits of VBAC

Women who choose VBAC compared to Elective repeat CS have reported greater satisfaction with their delivery, which is attributed to feelings of selflessness quicker recovery time from VBAC has been reported.5 Maternal morbidity and mortality associated with VBAC is also low.

A successful VBAC also increases the probability of future vaginal deliveries.6 Less maternal depression has been reported with vaginal deliveries.5 Additionally, successful lactation is more likely with VBAC.

Benefits of elective CS

Elective CS can be scheduled at a convenient time; hence the outcome is more certain and there is less painful delivery.

The parents can schedule days off from work.3 Since 30-50% of VBAC land up in repeat caesareaen if trial of labor fails, so many expectant mothers and obstetricians opt for elective caesarean rather than facing the risk an emergent caesarean, which is more dangerous than planned caesareans.

For women elective caesarean is theorized to be protective for the pelvic floor. However, this is controversial, because pregnancy itself predisposes to pelvic floor weakness and not the method of delivery. But the most important advantage is avoidance of risk of scar rupture. The true risk of scar rupture with VBAC is uncertain as many factors affect it. Several studies have been done to examine risk factors and predictors of uterine scar rupture in pregnancies following prior caesarean deliveries, however clear correlations have proven elusive.7

Risk to fetus in VBAC

VBAC has a times higher risk of perinatal deaths as compared to elective CS but this risk is shown to be equivalent to that of a fetus of a primigravida mother.8,11 The absolute risk being 4.5 per 10,000 births and with wide confidence limits. Risk of neonatal birth trauma and postpartum hemorrhage if operative vaginal delivery, like vacuum or forceps assisted delivery is required is increased. Rates of wet lung syndrome and neonatal convulsions are lesser in such deliveries.9

METHODS

In order to examine the potential cases for successful VBAC after the first caesarean, an examination was conducted for 375 women with previous one lower segment caesarean section (LSCS) in time period of 18th May 2018 to 31th May 2019. The statistical technique of paired t-test was run to relatively assess the neonatal complications and maternal complications across the two groups i.e. repeat LSCS and vaginal delivery.

Patient selection was guided by both medical and non-medical grounds and it included factors, such as incision type, and number of previous caesarean sections, physical factors such as cervical ripening, effacement, and progress of labour, and obstetric history like gravidity, parity, and prior vaginal delivery.8,10

Certain non-medical factors which were taken into account include, preference of patients, provider comfort regarding the mode of delivery. If during trial of labour there was any suspicion of scar dehiscence or fetal distress or if progress of labour was unsatisfactory, then trial was abandoned. All women were always prepared for emergency CS, if need arose.

Complete history including indication of previous CS, intra and postoperative complications during previous surgery, the details of the present pregnancy, scar tenderness and any other disorder were recorded. However, those who had presented with intrauterine fetal death, two previous CS and scar of other uterine surgery were excluded from the study. Women with recurrent
indications for CS or those having non recurrent indications with any complicating factors in present pregnancy were taken for elective LSCS (n = 188). Those women with previous one LSCS for the non-recurrent indications were given a trial for vaginal delivery (n = 197).

*Case selection for the trial for vaginal delivery was done as per ACOG guidelines*

- Singleton pregnancy
- Gestational age > 34 weeks
- History of previous one LSCS
- Non recurrent indication for the previous LSCS.

*The labor was monitored with*

- Maternal vital parameters were monitored hourly particularly pulse and BP
- Fetal heart rate monitoring by intermittent auscultation - every 15 minutes in the first stage and every 5 minutes in second stage of labor
- Progress of labor as per WHO partograph
- A close watch for the early recognition of scar dehiscence by identifying signs such as maternal tachycardia, vaginal bleeding, scars tenderness and fetal distress.

Attempt at vaginal delivery was abandoned if there was any suspicion of scar dehiscence or fetal distress or unsatisfactory progress of labor. All women were always prepared for emergency CS if need arose.

**RESULTS**

During the period of study, a total of 375 women with previous one CS were included in the study protocol. Looking into the previous details and present findings, in 188 (50.10%) women (Table 1) vaginal delivery was contemplated and in 187 (49.9%) women elective CS was planned.

| Category of female respondents | Frequency | Percentage |
|--------------------------------|-----------|------------|
| Vaginal delivery               | 188       | 50.10%     |
| Elective CS                    | 187       | 49.90%     |
| **Total**                      | **375**   | **100.00%**|

Table 2 shows the demographic profile in terms of age group. It was found that most of the women (63.7%) belong to 26-30 years of age. Further, from Table 3, it can be inferred from the above table that majority of the women delivered at the gestational age of 37-40 weeks (77.6%).

**Table 3: Gestational age.**

| Period of gestation (weeks) | Frequency | Percentage |
|-----------------------------|-----------|------------|
| 34-37                       | 72        | 19.2%      |
| 37-40                       | 291       | 77.6%      |
| More than 40                | 12        | 3.2%       |
| **Total**                   | **375**   | **100.00%**|

**Table 4: Indications for previous caesarean delivery.**

| Indications for previous caesarean delivery | Frequency | Percentage |
|--------------------------------------------|-----------|------------|
| Cephalopelvic disproportion                | 155       | 41.3%      |
| Fetal distress                             | 79        | 21.1%      |
| Non progression of labor                   | 39        | 10.4%      |
| Malpresentation                            | 34        | 9.1%       |
| Ante partum hemorrhage                     | 18        | 4.8%       |
| Failed trial                               | 19        | 5.1%       |
| Failed induction                           | 9         | 2.4%       |
| Obstructed labor                           | 9         | 2.4%       |
| BOH                                        | 13        | 3.5%       |
| **Total**                                  | **375**   | **100.00%**|

**Table 5: Mode of delivery in trial of labor group (n = 188).**

| Mode of delivery | Frequency | Percentage |
|------------------|-----------|------------|
| Spontaneous vaginal | 111       | 59.30%     |
| Instrumental     | 14        | 7.20%      |
| Emergency repeat LSCS | 63        | 33.50%     |
| **Total**        | **188**   | **100.00%**|

**Table 6: Indications of repeat emergency LSCS (63 cases).**

| Indication of repeat emergency LSCS | Frequency | Percentage |
|------------------------------------|-----------|------------|
| Fetal distress                     | 27        | 42.86%     |
| Non progress of labour             | 22        | 34.92%     |
| Scar tenderness                    | 10        | 15.87%     |
| Abruptio placenta                  | 4         | 6.35%      |
| **Total**                          | **63**    | **100.00%**|

It can be inferred from Table 4 that major indications for previous caesarean delivery included Cephalopelvic disproportion (41.3%), fetal distress (21.1%), non-progression of labor (10.4%), malpresentation (9.1%), failed trial (5.1%) and ante partum hemorrhage (4.8%). Table 5 show that 66.5% women delivered vaginally in the trial group; 59.30% had spontaneous vaginal delivery.
and 7.2% women had to undergo instrumental delivery by outlet forceps or vacuum extraction. The study indicated that the common causes for abandoning the trial of labor and opting for emergency LSCS (Table 6) was fetal distress (42.86%), non-progress of labor (34.92%), scar tenderness (15.87%) and abruptio placenta (6.35%).

**Table 7. Maternal complications.**

| Type of complication | Repeat LSCS group | Vaginal delivery group |
|----------------------|-------------------|------------------------|
| Pyrexia              | 2                 | 1                      |
| Postpartum hemorrhage| 3                 | 2                      |
| Wound gap            | 1                 | 0                      |
| Cervical/vaginal tear| 0                 | 4                      |
| Haematuria           | 1                 | 0                      |
| **Total**            | **7**             | **7**                  |

A relative comparison on maternal complications across the two groups i.e. repeat LSCS and vaginal delivery (Table 7) indicate that pyrexia, PPH, wound gape, haematuria were more in repeat LSCS group while cervical/vaginal tears, traumatic PPH and scar complications were more common in the VBAC group. Further analysis in this study using t-test was conducted to examine the difference of both the group cases, i.e. Repeat LSCS group and the Vaginal delivery group w.r.t Maternal complications (Table 8). The findings show that there was no statistically significant difference between the repeat LSCS group and the vaginal delivery group w.r.t maternal complications (t = 0.779, p > 0.05).

A relative comparison w.r.t Neonatal complication indicates that asphyxia was more common in the vaginal delivery group (Table 9) as compared to the Repeat LSCS group. NICU admissions were more common in the repeat LSCS group (cause being fever, birth asphyxia and jaundice but all babies were discharged in good condition) than in the vaginal delivery group.

Further analysis to examine the difference of both the group cases, i.e. repeat LSCS group and the vaginal delivery group w.r.t neonatal complications (Table 10) using t-test revealed that there was no statistically significant difference in neonatal complications across both the groups t = 0.632, p > 0.05.

**Table 8: t-test.**

| Levene's test for equality of variances | t-test for equality of means |
|----------------------------------------|-----------------------------|
| F       | Sig. | T    | df  | Sig. (2-tailed) | Mean difference | Std. error difference | 95% confidence interval of the difference |
|---------|------|------|-----|-----------------|-----------------|-----------------------|----------------------------------------|
| Equal variances assumed                | 0.035 | 0.854 | 0.779 | 12   | 0.451 | 0.57143 | 0.73309 | -1.02583 | 2.16869 |
| Equal variances not assumed            | 0.779 | 11.772 | 0.451 | 0.57143 | 0.73309 | -1.02927 | 2.17213 |

**Table 9: Neonatal complications.**

| Neonatal complications | Repeat LSCS | Vaginal delivery group |
|------------------------|-------------|------------------------|
| Fever                  | 2           | 2                      |
| Asphyxia               | 1           | 4                      |
| Septicaemia            | 1           | 1                      |
| Jaundice               | 3           | 2                      |
| **Total**              | **7**       | **9**                  |

**DISCUSSION**

This study represents our observations over a period of one year. Women with prior one LSCS require special management, both during the antenatal period and during labour. The decision for a trial of labor or the elective repeat LSCS in a patient needs to be based on proper selection of cases and after thorough counseling. Rough idea can be made on the basis of maternal characteristics and obstetric history.

Several studies suggest that for appropriately selected women with previous one LSCS, a trial for vaginal delivery is safe. Published literature shows that there has been a 60-80% success in VBAC. The study success rate (66.5%) is comparable to these studies. Factors that negatively influence the likelihood of successful VBAC are believed to be cases with labor induction and augmentation, maternal obesity, gestational age > 40 weeks, birth weight > 4000 gm and inter delivery interval of less than 19 months. A history of a previous successful VBAC increases the likelihood for success with future attempts.

The risk of uterine rupture is higher with an induced labor than with a spontaneous labor with trial. Induction and augmentation with oxytocin is safe in selected cases
with standard obstetric indications; but use of prostaglandins for induction needs much caution. Bujold E et al have reported that the single layer closure of the previous lower segment incision was associated with a fourfold increase in the risk of uterine rupture compared with a double layer closure.\textsuperscript{14}

| Table 10: t-test on neonatal complications between the test groups. |
|-------------------------------------------------------------|
| **Levene's test** | **t-test for equality of means** |
| for equality of variances | | |
| | F | Sig. | t | Df | Sig. (2-tailed) | Mean difference | Std. error difference | 95% confidence interval of the difference |
| Equal variances assumed | 0.086 | 0.78 | -0.632 | 6 | 0.55 | -0.5 | 0.79057 | -2.43445 | 1.43445 |
| Equal variances not assumed | -0.632 | 5.602 | 0.552 | -0.5 | 0.79057 | -2.46829 | 1.46829 |

Neither repeat cesarean delivery nor trial of labor is risk free. Maternal morbidity in terms of pyrexia, atomic PPH, hematuria, and wound gape are more common in repeat CS group, while cervical and vaginal tear, traumatic PPH and uterine scar problems are more common in trial of vaginal delivery. However, the difference in maternal morbidity rate is not statistically significant, this is evident in our study also.\textsuperscript{15-17} A trial of vaginal delivery may result in small but insignificant increase in the perinatal morbidity and mortality rates, which can be reduced by proper selection of cases.\textsuperscript{15-17}

**CONCLUSION**

Prediction as to whether the trial of labor will succeed or not and to identify patients who are at risk of failing the trial of labor is very important. Many repeat CS can be avoided if proper selection of cases, proper timing and close supervision of labor by competent staff. There is a need for individualized approach to all the cases.

_Funding: No funding sources_  
_Conflict of interest: None declared_  
_Ethical approval: The study was approved by the Institutional Ethics Committee_

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Cite this article as: Prasad SG, Malhotra P. Assessing the pros and cons of vaginal birth after caesarean relative to elective repeat caesarean section. Int J Reprod Contracept Obstet Gynecol 2020;9:1350-5.