



Maternal Morbidity and Mortality Following a Trial of Labor in Women with Previous Cesarean Section at Tertiary Care Teaching Hospitals in India

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ABSTRACT

Background: Repeat caesarean section and planned vaginal birth after caesarean section are both associated with benefits and harms.

Methods: Prospective data was recorded on management practices, associated complications, morbidity and mortality on 15664 consecutive cases of previous cesarean section reporting at 30 medical colleges/teaching hospitals for delivery.

Results: A total of 4035 (25.8%) women out of the 15664 women with a previous cesarean section underwent a trial of labor (TOL). Of these, 2513 (62.0%) had a successful trial of labor (S-TOL) while the rest required an emergency repeat cesarean section. The overall maternal morbidity was 2.3% and 34.0% in women with S-TOL and failed trial of labor (F-TOL) respectively. Blood loss more than 1000ml was seen in 20.6% of cases with F-TOL where as for S-TOL it was 0.3%, blood transfusion was 7.0% in F-TOL where as it was 0.8% in S-TOL, dehiscence of scar in F-TOL was 5.4% as compared to 0.2% in S-TOL, post-operative complication/delivery were seen in 6.8% cases in F-TOL where as in S-TOL it was 0.4%, uterine rupture was 0.7% in F-TOL as compared to 0.1% in S-TOL and was statistically significant. Maternal death was seen in 4 (0.3%) cases of F-TOL as compared to 6 (0.2%) cases in S-TOL ($p=0.45$) and the difference was not significant. There were 27 (1.8%) child deaths who born after F-TOL as compared to 65 (2.6%) born after S-TOL which was highly significant ($P=0.00$).

Conclusions: Women who experience failed trial of labor have higher risk of morbidity as compared to those with a successful trial of labor. More accurate prediction for safe, successful vaginal birth after cesarean delivery is needed.

Keywords: Cesarean section, trail of labor, maternal morbidity, mortality.

INTRODUCTION

Rate of caesarean sections has been increasing over a period of time both in developed and developing countries. Today, it is one of the most

commonly performed surgical procedures, and despite advances in surgical technique and safer anesthesia, it is still associated with a higher maternal morbidity. Due to a rise in the rates of

primary caesarean section globally, repeat caesarean section has also become very common. As caesarean birth rates continue to rise, increasingly obstetricians are faced with the choice of planning a vaginal or caesarean birth after a previous caesarean birth.¹ In an attempt to reduce the rising trend of caesarean delivery worldwide, obstetrician now offer trial of labor more readily to women who have had a caesarean section.²⁻⁴ Several studies both in developed and developing countries have shown that it is not only feasible but safe.⁴⁻⁷ Among women who attempt a trial of labor after a previous caesarean section, 60% to 80% have a vaginal delivery,⁸⁻¹¹ and morbidity is lower among women who have a vaginal delivery after a previous caesarean section than among women who elect to undergo a second caesarean section.^{9,12,13} Although trial of labor is usually successful and relatively safe, it may occasionally be associated with severe maternal morbidity and even mortality and correct management represents one of the most significant and challenging issues in obstetric practice.^{4, 7} With this background the study was conducted prospectively to study the maternal morbidity and mortality amongst women who underwent a trial of labor after a previous caesarean delivery at tertiary care teaching hospitals in India.

METHODS

A hospital based maternal health database was established at 30 medical colleges/teaching hospitals situated all over the country and prospective data was recorded for a period of 8 months on management practices, associated complications, morbidity and mortality in 15664 consecutive cases of previous caesarean section reporting for delivery. Structured case record forms were completed by trained medical research staff.

The study population was divided into 2 groups based on whether the woman underwent a trial of labor (TOL) or an elective repeat caesarean section

(El-RCS) as the mode of delivery. The first group (TOL) was further subdivided into women who successful trial of labor (S-TOL) and women who had a failed trial of labor (F-TOL) and had to undergo an emergency caesarean section. Both groups (S-TOL and F-TOL) were compared with regard to any type of maternal morbidity, uterine rupture/dehiscence, and emergency interventions like blood transfusion and hysterectomy. The data collected were coded and fed into the computer using Epi-Info and analyzed using SPSS V 19.0. Descriptive statistics such as mean, standard deviation and percentage were used and to find association chi square test was used.

RESULTS

Of the 15664 women with a previous caesarean section, 4035 (25.8%) women underwent a trial of labor (TOL) and an elective repeat caesarean section (El-RCS) was carried out in 5399 (34.5%) cases. For various indications, an emergency repeat caesarean section was carried out on the remaining 6230 (39.7%) women. Among the women who underwent a TOL, 2513 (62.0%) had a S-TOL while in 1522 (38.0%) women, there was a failed trial of labor (F-TOL) requiring an emergency repeat caesarean section. The total number of women who therefore underwent an emergency repeat caesarean section was 7752. (Fig 1)

There was no significant difference ($P=0.98$) in the mean age of women with F-TOL (26.0 ± 3.8 years) and S-TOL (25.9 ± 3.8 years). Majority of the caesarean section in women with F-TOL (81%) and S-TOL (80.4%) were done in the age group 20-30 years. The mean parity of women who had F-TOL was 1.2 ± 0.6 and the value for those who had S-TOL was 1.5 ± 0.8 . The difference was statistically significant ($P=0.00$). A trial of labor was more likely to fail in 80% if the infant weight was 2500 g or more. (Table 1)

There was a statistically significant difference ($P=0.00$) in any maternal morbidity which was found higher among women who underwent F-TOL (34%) as compared to S-TOL (2.3%). Blood

loss more than 1000 ml was seen in 314 (20.60%) women with F-TOL and was significantly higher than that seen in 7 (0.3%) women with S-TOL (OR: 0.01, CI: 0.0-0.02, p=0.00). Similarly, a significantly higher proportion of women with F-TOL received blood transfusion (7.0%) in contrast to women with S-TOL (0.8%) (OR: 0.24, CI: 0.16-0.34, p=0.00). Dehiscence of scar in F-TOL was present in 82 (5.4%) cases as compared to 6 (0.2%) in S-TOL and was highly significant (OR: 0.04, CI: 0.02-0.1, p=0.00). Post-operative/delivery complications were present in 104 (6.8%) women who had F-TOL whereas it was only seen in 10 (0.4%) women with S-TOL (OR: 18.36, CI: 9.22-37.67, p=0.00). Uterine rupture was 0.7% versus 0.1% in F-TOL and S-TOL respectively

(OR: 0.12, CI: 0.02-0.59, p=0.001). Maternal death was reported in 4 (0.3%) cases of F-TOL as compared to 6 (0.2%) cases in S-TOL (p=0.45) which was not statistically significant. The average duration of hospital stay for S-TOL was significantly less (4.5±3.9 days) as compared to (10.6±5.0) days for F-TOL and was highly significant (p=0.000)(Table 2). There was no significant difference in the rates of admission to a neonatal intensive care unit between the two groups. It was 12.5% in F-TOL whereas it was 11.0% in children born after S-TOL (OR: 1.15, CI: 0.94-1.41, p=0.16). However, child death reported after F-TOL was 27 (1.8%) as compared to 65 (2.6%) in S-TOL cases which was highly significant (P=0.00).

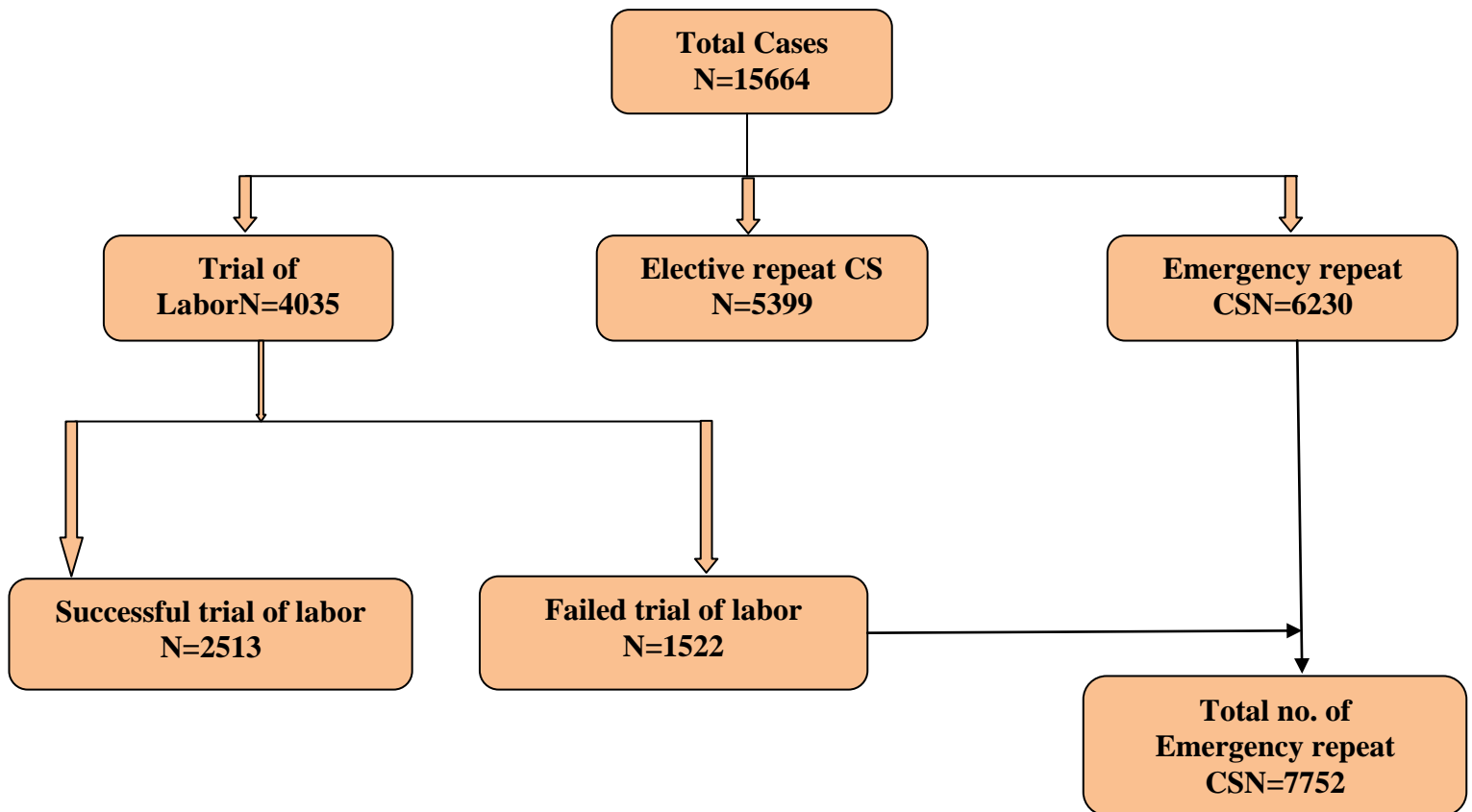


Figure 1: Diagrammatic representation of the selection of the study population.

Table 1. Characteristics of pregnant women who failed to deliver vaginally after a trial of labor and those who delivered successfully vaginally after a trial of labor

Characteristics	Failed trial of labor (F-TOL) (N=1522)	Successful trial of labor (S-TOL) (N=2513)	Odds Ratio/ 95% Confidence interval	P-value
Maternal age (Yrs)				
<=19	12 (0.8)	22 (0.9)	0.90 (0.42-1.92)	0.77
20-24	565 (37.1)	923 (36.7)	1.02 (0.89-1.16)	0.80
25-29	667 (43.8)	1098 (43.7)	1.01 (0.88-1.15)	0.94
30-34	223 (14.7)	382 (15.2)	0.96 (0.80-1.15)	0.64
>=35	55 (3.6)	88 (3.5)	1.03 (0.72-1.40)	0.85
Mean±Sd	26.0±3.8	25.9±3.8		0.98
Parity				
2	1267 (83.2)	1601 (63.7)	2.83 (2.41-3.33)	0.00
3	205 (13.5)	693 (27.6)	0.41 (0.34-0.49)	0.00
4	35 (2.3)	166 (6.6)	0.33 (0.22-0.49)	0.00
5	8 (0.5)	42 (1.7)	0.31 (0.13-0.7)	0.001
>5	7 (0.5)	11(0.4)	1.05 (0.36-2.95)	0.92
Mean±Sd	1.2±0.6	1.5±0.8		0.00
Booking Status				
Booked	1050 (69.0)	1570 (62.5)	1.34 (1.16-1.54)	0.00
Unbooked	472 (31.0)	943 (37.5)		
Infant's birth weight				
<2500	266 (17.5)	722 (28.7)	0.53 (0.45-0.62)	0.00
2500-2999	572 (37.6)	1058 (42.1)	0.83 (0.72-0.95)	0.004
3000-3499	517 (34.0)	588 (23.4)	1.68 (1.46-1.95)	0.00
3500-3999	142 (9.3)	104 (4.1)	2.38 (1.82-3.13)	0.00
>=4000	18 (1.2)	13 (0.5)	2.30 (1.06-5.02)	0.002
Not known	7 (0.5)	28 (1.1)		0.00

Table 2. Morbidity in pregnant women who failed to deliver vaginally after a trial of labor and those who delivered successfully vaginally after a trial of labor

MORBIDITY	Failed trial of labor (F-TOL) (N=1522)	Successful trial of labor(S-TOL) (N=2513)	Odds Ratio/ 95% Confidence interval	P-value
Any morbidity	517 (34.0)	57 (2.3)	0.05 (0.03-0.06)	0.0
Anaesthetic complication	16 (1.1)	0 (0.0)	-	-
Dehiscence of the scar	82 (5.4)	6 (0.2)	0.04 (0.02-0.1)	0.0
Uterine rupture	10 (0.7)	2 (0.1)	0.12 (0.02-0.59)	0.001
Blood loss>1000 ml	314 (20.6)	7 (0.3)	0.01 (0.0-0.02)	0.00
Broad ligament hematoma	3 (0.2)	0 (0.0)	-	-
Blood transfusion	107 (7.0)	44 (.8)	0.24 (0.16-0.34)	0.0
Hysterectomy	1 (0.1)	2 (0.1)	0.83 (0.03-11.71)	0.88
Post-operative/delivery complication	104 (6.8)	10 (0.4)	18.36 (9.22-37.67)	0.0

DISCUSSION

Cesarean section can be a lifesaving procedure when medically indicated and is an important indicator of the ability to provide comprehensive obstetric and neonatal care. However, over utilization of this procedure is a growing concern. Over the past decade there has been a gradual increase in the rate of cesarean section even in the developing countries causing considerable concern. In USA rate of abdominal delivery was

29.1%, in England 21.5%, Latin American States it was 40%¹⁴ and 28.1% in tertiary care teaching hospital in India. These concerns are due to the undesirable effects of 'unnecessary' cesarean sections on the health of the mother and child and the economic and health systems impact.

The overall maternal morbidity in our study was 34.0%, 2.3% in F-TOL and S-TOL respectively. For women with a previous cesarean delivery, S-TOL was generally associated with lower

morbidity than scheduled repeat procedures. However, F-TOL were associated with increased rates of the morbidities compared with scheduled repeat procedures, which is consistent with many other studies.¹⁶⁻¹⁸ In this study any morbidity after S-TOL was 2.3% while in F-TOL it was 34.0% which was highly significant ($p=0.00$). Maternal death was reported in 0.3% cases of F-TOL as compared to 0.2% cases in S-TOL ($p=0.45$) which was not statistically significant. The average duration of hospital stay for S-TOL was 4.5 ± 3.9 days as compared to F-TOL 10.6 ± 5.0 . This shows that women who had a S-TOL had a significantly lesser duration of hospital stay as compared to those had a cesarean section ($p=0.000$ highly significant). The rates of admission to a neonatal intensive care unit was 12.5% versus 11.0% (OR: 1.15, CI: 0.94-1.41, $p=0.16$ not statistical significant) in F-TOL and S-TOL respectively. Child death reported in F-TOL was 1.8% as compared to 2.6% in S-TOL which was highly significant ($P=0.00$). Majority of neonates were having NICU admission due to premature rupture of membranes, meconium stained liquor, low birth weight and respiratory distress syndrome. Our study was well comparable with other studies who found that Infants born after successful trial of labor had the lowest rates of NICU admission than those born by failed trial of labor.¹⁹⁻²¹ However, there is as yet no confirmed method of predicting the likelihood that a trial of labor will lead to vaginal delivery for a patient with a previous cesarean section.

CONCLUSION

Women who experience failed trial of labor have higher risk of morbidity as compared to those with a successful trial of labor. More accurate prediction for safe, successful trial of labor is needed.

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