

## OUTCOME OF PREGNANCY IN WOMEN WITH PREVIOUS CAESAREAN SECTION

Bellad Girija Channabasappa<sup>1</sup>, Rajendra Ramakrishna Godbole<sup>2</sup>

<sup>1</sup>Assistant Professor, Department of Obstetrics and Gynaecology, Belagavi Institute of Medical Sciences, Belagavi.

<sup>2</sup>Associate Professor, Department of Obstetrics and Gynaecology, Belagavi Institute of Medical Sciences, Belagavi.

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### ABSTRACT

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#### BACKGROUND

Carefully selected cases of Vaginal Birth after Caesarean Section (VBAC) is safe and successful. Even though options of elective caesarean section or a trial of labour are given to women with prior caesarean section, the risk is always present. In successful VBACs, morbidity is less compared to repeat caesarean section. That is why this study is conducted to determine the outcome of pregnancy in women with previous CS.

#### OBJECTIVES

1. To evaluate the clinical course of labour in cases with previous caesarean section.
2. To study the perinatal outcome in cases with previous caesarean section either by vaginal delivery or repeat Caesarean section.
3. To study maternal morbidity in these cases.

#### METHOD

A retrospective analysis of medical records of 250 women with a previous caesarean section, who delivered in BIMS Hospital between May 2015 and July 2015 was carried out. Women with recurrent indications for caesarean section and those having non-recurrent indications with any complicating factors in present pregnancy and women with previous two caesarean sections were not given trial for vaginal delivery. Those women with previous section for the non-recurrent indications were given trial for vaginal delivery.

#### STATISTICAL ANALYSIS

Was done by Chi-square test.

#### RESULT

In 250 cases, 132 cases were given trial for vaginal delivery. In these, vaginal delivery was 61.3% and repeat section was 38%. There is an association between maternal morbidity and type of delivery. Birth weight was associated with the type of delivery. There is no association between neonatal outcome and type of delivery.

#### CONCLUSION

In carefully selected patients, appropriate timing and close supervision, trial of vaginal delivery in previous one caesarean section is safe and successful. Individual approach seems to be the best.

#### KEYWORDS

Previous Caesarean Section, Vagina Birth after Caesarean Section, Trial of Vaginal Delivery, Elective Repeat Section.

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#### INTRODUCTION

It is an attempt to analyse the existing trends in our hospital in management of patients with previous section in subsequent pregnancy. All obstetricians encounter increasing number of post caesarean pregnancy, because the number of primary caesarean section for non-recurrent causes is rapidly rising. The procedure is not simple and needs to be performed only when circumstances distinctly require it.<sup>1</sup> For many years, the phrase 'Once a caesarean, always a caesarean' dictated

obstetric practice. Later, because of increasing Caesarean Sections (CS) it was noted that vaginal birth after CS (VBAC) might help in reducing the rate of CS.<sup>2</sup> In an appropriate clinical and proper selected group of women, VBAC is safe and effective.<sup>3,4</sup>

A trial of vaginal delivery after a previous CS is considered safer than the routine repeat CS, because operative risks are completely eliminated. The hospital stay is much shorter and expenses involved are much less. However, several factors increase the likelihood of failed trial, which in turn might lead to increased maternal and perinatal mortality and morbidity rates.<sup>2</sup> There is a controversy in the trial of vaginal delivery in post-caesarean pregnancy. This needs a critical audit.

So, informing and counseling the concerned women and their relatives regarding the benefits and risks involved in VBAC and repeat CS is essential. That is why this study is conducted to determine the outcome of pregnancy in women with previous CS in relation to vaginal delivery, maternal and perinatal complications, to identify the factors which can influence the outcome of trial of vaginal delivery.

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*Corresponding Author:*

*Dr. Bellad Girija Channabasappa,*

*'Simply OM' Apartments,*

*GF-1, 2<sup>nd</sup> Main, 6<sup>th</sup> Cross,*

*Sadashiv Nagar,*

*Belagavi-590001,*

*Karnataka State.*

*E-mail: rajashekarphypub@gmail.com*

*DOI: 10.14260/jemds/2016/642*

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**METHODS**

A retrospective analysis of medical records of 250 women with previous CS from May 2015 to July 2015. Hospital records of patients with previous CS undergoing delivery in our hospital are collected.

**The following women were not given trial of vaginal delivery, but were taken for caesarean section (n=118)**

1. Women with recurrent indications for CS.
2. Those having non-recurrent indications with any complicating factors in present pregnancy.
3. Patient with previous two CS.
4. Mal-presentations.
5. Details of previous CS not available.

The following women (n=132) were given trial vaginal delivery: i) Previous one CS for the non-recurrent indications; ii) Singleton pregnancy; iii) Gestation age more than 37 weeks; iv) History of one CS. According to the case record, labour was monitored by every half hourly maternal pulse, foetal heart rate and uterine contractions and 4<sup>th</sup> hourly BP, cervical dilation, descent of head and moulding. A close watch on the early recognition for scar dehiscence by identifying the maternal tachycardia, scar tenderness, foetal distress and vaginal bleeding. Attempt of vaginal delivery was abandoned if there was any suspicion of scar dehiscence or foetal distress or unsatisfactory progress of labour. Such women were taken for CS.

Institutional ethical clearance for the study was obtained on 05-10-2015.

**Statistical Analysis**

Qualitative data were analysed using Chi-square test (X<sup>2</sup>); p value less than 0.05 means statistically significant; p value less than 0.001 means highly significant; p value more than 0.05 is insignificant.

**RESULTS**

Out of the total of 250 patients who were included in the study, 132 were on trial vaginal delivery (52.8%) and on 118 (47.2%) patients' elective Caesarean section was done (Table 1).

Most of the women belonged to 20–25 years of age (183 cases, i.e. 73.2%). Overall age group number of patients is not uniformly distributed among all the parameters. Most of the patients (213 cases, i.e. 85.2%) were in the 37–40 weeks of gestation. Here also number of patients are not uniformly distributed among both the groups, i.e. age and period of gestation groups (Table 2).

Women delivered vaginally in the trial group were 81, i.e. 61.36%. In this trial group 50% had spontaneous vaginal delivery, 11.36% required outlet forceps either for prolonged 2<sup>nd</sup> stage or foetal distress in 2<sup>nd</sup> stage and 38.63% needed emergency LSCS (Table 3).

Foetal distress cases were 24 in number (47%). The number of patients were not distributed uniformly among all the parameters. In 10 women scar tenderness was the indication, but during surgery 2 cases had scar dehiscence and in 1 case there was bladder rupture which was repaired (Table 4).

There was no maternal mortality. Morbidity like pyrexia, wound gaping, UTI, wound infection, bladder rupture and requirement of blood transfusion was more in repeat CS. Para-urethral tear, episiotomy wound gaping and cervical tear were

common in VBAC group. There is association between maternal complication and type of delivery (Table 5).

There is an association between birth weight (2.6 to 4.0 kg) and type of delivery (vaginal delivery, abdominal delivery). Among these, abdominal delivery is significantly higher than the vaginal delivery. So birth weight is associated with the type of delivery. There is no association between type of delivery (vaginal delivery and abdominal delivery) with living children, still births and neonatal deaths (Table 6).

The success rate of VBAC in our study is 61.36% (Table 7). Out of 250 cases, 81 cases (32.4%) of cases delivered vaginally either spontaneously or by forceps and 169 cases (67.6%) required caesarean section. Emergency caesarean section (51 cases) were less than elective caesarean section (118 cases who were not in labour). One case who underwent subtotal hysterectomy was diagnosed as ruptured uterus, since patient came late in labour (Table 8).

Group	No.
Trial vaginal delivery	132 (52.8%)
Elective CS	118 (47.2)

**Table 1: A Total of 250 Patients were Included in the Study**

Parameters	No.	Percentage	
<b>a. Age (Years)</b>			
20-25	183	73.2	X <sup>2</sup> = 32.82 P < 0.001
26-30	53	21.2	
31-35	11	4.4	
36 and above	3	1.2	
<b>b. Period of Gestation (Weeks)</b>			
37 – 40	213	85.2	X <sup>2</sup> = 123.900 P < 0.001
> 40 weeks	37	14.8	
<b>c. Indication for Previous Section</b>			
CPD	105	42	
Foetal distress	61	24.4	
Non-progress of labour	24	9.6	
Mal-presentation	21	8.4	
Severe PIH/ Eclampsia	10	4	
APH	7	2.8	
Failed induction	10	4	
Obstructed labour	8	3.2	
BOH	4	1.6	

**Table 2: Demographic Profile (n=250)**

Mode of Delivery	No.	Percentage
Spontaneous vaginal delivery	66	50
Instrumental	15	11.36
Unsuccessful vaginal delivery	51	38.63

X<sup>2</sup> = 31.23; p < 0.001

**Table 3: Outcome Trial of Labour in Present Pregnancy (n= 132)**

Indication for Repeat Emergency CS (n=51)	No.	Percentage
Foetal distress	24	47.05
Non-progress of labour	13	25.49
Obstructed labour	3	5.88
Scar tenderness	10	19.6
Abruptio placenta	1	1.96

X<sup>2</sup> = 32.82; p < 0.001

**Table 4: Indication for Repeat Emergency CS Maternal Complications**

Type of Complications	Vaginal Delivery (n=81)		Repeat CS (n=169)	
	No	%	No	%
Pyrexia	3	3.7	15	8.87
PPH	1	1.23	2	1.18
Wound gaping	--	--	5	2.96
Cervical tear	1	1.23	--	--
Paraurethral tear	2	2.46	--	--
Episiotomy wound gaping	4	4.93	--	--
Blood transfusion	1	1.23	3	1.77
UTI	2	2.46	10	5.92
Wound infection	--	--	5	2.96
Bladder rupture	--	--	3	1.77
X <sup>2</sup> = 26.87; p < 0.05				

**Table 5: Type of Complication**

Birth Weight (Kg)	Vaginal Delivery	Abdominal Delivery	Total	Analysis
2.0 - 2.5	23 (9.2%)	24 (9.6%)	47 (18.8%)	X <sup>2</sup> = 12.99; p < 0.02
2.6 - 3.0	41 (16.4%)	80 (32.0%)	121 (48.4%)	
3.1 - 3.5	16 (6.4%)	50 (20.0%)	66 (26.4%)	
3.6 - 4.0	1 (0.4%)	15 (6%)	16 (6.4%)	
Living children	79 (97.53%)	166 (98.22%)	245 (98%)	X <sup>2</sup> = 1.05; p > 0.05
Still birth	0	1 (0.56%)	1 (0.4%)	
Neonatal death	2 (2.46%)	2 (1.18%)	4 (1.6%)	

**Table 6: Foetal Outcome in Present Pregnancy (n = 250)**

Sl. No.	Author	Year	Percentage
1	Yadav K. <sup>5</sup>	2000	61.16%
2	Pandey N. <sup>6</sup>	2002	42%
3	Dinsmoor MJ. <sup>7</sup>	2007	76%
4	Turner MJA. <sup>8</sup>	2006	77.8%
5	Chaudhari DR. <sup>9</sup>	2012	67.0%
6	Present Study		61.36%

**Table 7: Comparative Percentage Vaginal Delivery after Trial of Scar of Various Authors**

This table shows VBAC range is 42% - 77.8%. In our study, success rate is 61.36%.

Sl. No.	Outcome in Present Pregnancy	No. of Cases	Percentage
1	Vaginal Delivery		
	i. Normal	66	26.4%
	ii. Forceps	15	6%
	<b>Total</b>	<b>81</b>	<b>32.4%</b>
2	Abdominal Delivery		
	i. Elective caesarean section	118	47.2%
	ii. Emergency caesarean section	51	20.4%
	<b>Total</b>	<b>169</b>	<b>67.6%</b>

**Table 8: Outcome of Labour in Post-Caesarean Section (n=250)**

**DISCUSSION**

Women with prior CS require special care during both antenatal and labour. Before 1970, the term in Obstetric practice was ‘once a Caesarean, always a Caesarean.’ But now it is ‘once a Caesarean, always a hospital delivery.’

The decision for a trial of labour or elective repeat CS is an individual one and that should be based on careful selection and thorough counselling.<sup>4</sup> Maternal characteristics and Obstetric history can provide a rough estimate. Several studies suggest that for appropriately selected women with previous one CS, prior vaginal delivery, a trial for vaginal delivery is safe.

Studies by Yadav K.<sup>5</sup> shows 67.16% vaginal delivery after trial of labour, Pandey N.<sup>6</sup> shows 42.0%, Dinsmoor MJ.<sup>7</sup> shows 76%, Turner MJA.<sup>8</sup> shows 77.8% and Chaudhari DR.<sup>9</sup> shows 67%. Our success rate is 61.36% and is comparable to studies by Yadav K.<sup>5</sup> and Chaudhari DR.<sup>9</sup>

**Mode of Delivery**

In modern obstetrics during second stage of labour, controversy exists regarding use of prophylactic forceps to reduce time. Some studies.<sup>5,9</sup> used prophylactic forceps during second stage labour to cut short the duration. In our study, prophylactic forceps was used in 9 cases (6.81%).

Augmentation of labour using Oxytocin during trial of vaginal delivery in previous Caesarean section is controversial. Singhal P.<sup>10</sup> achieved a success rate of 88%; Iyer S et al<sup>11</sup> found 69% using Oxytocin during trial of labour. In our study, the success rate using Oxytocin in trial of labour is 40.9%.

**Foetal Distress**

Diagnosis of foetal distress was done by noting bradycardia (FSH <100 beats per minute), tachycardia (FSH >160 per minute) and meconium stained liquor. In studies by Yadav K.<sup>5</sup>, Chaudhari DR.<sup>9</sup> and Shakti V.<sup>12</sup>, the foetal distress were 22.72%, 14.15% and 50% of cases respectively. In our study, the foetal distress was 47.05% of cases, which was comparable to study by Shakti V.<sup>12</sup>

**Scar Dehiscence**

It is defined as incomplete or complete separation of uterine scar with membrane intact and foetus in utero. Studies by Choudhari DR.<sup>9</sup>, Iyer S.<sup>11</sup> and Shakti V.<sup>12</sup> and the incidence of scar dehiscence was 5.66%, 0.5% and 1.44% respectively. In our study, the scar dehiscence was 1.5% and is comparable with Iyer S.<sup>11</sup> Factors which have negative impact.<sup>2,13</sup> are cases with labor induction, maternal obesity, maternal age (>35 yrs.), gestational age (>40 weeks), birth weight (>4 kgs) and gap between 2 deliveries (<18 months).

A history of previous success Vaginal Birth After Caesarean Section (VBAC) increases the likelihood of success with future attempts.<sup>14,15</sup> The risk of uterine rupture is higher with an induced labour than with spontaneous labour with trial. Induction and augmentation with Oxytocin is safe in selected cases with standard Obstetric indications. But use of prostaglandins for induction need much caution.

Repeat CS and trial labour are both at risk. Maternal morbidity like pyrexia, atonic post-partum haemorrhage (PPH), urinary tract infection, wound gaping and need for blood transfusion are more common in repeat CS. Cervical tear, traumatic PPH and scar dehiscence are more common in

trial of vaginal delivery. Our study shows an association between maternal complication and type of delivery.

### CONCLUSION

Mode of delivery should be decided depending upon the previous indication, type of scar and associated maternal complications. An attempt for VBAC is well justified for post caesarean pregnancy with non-recurrent indication. Vaginal deliveries are safer than Caesarean section, as there are fewer complications with less maternal morbidity. The ability to predict women who are at high risks for failing the trial of vaginal delivery and those with high probability of successful delivery would help guide the clinician making good clinical decision. This would minimize adverse events.

The key factors to achieve greater degree of success in VBAC are proper selection, appropriate timing and close supervision by competent staff. By this we can eliminate the need for large proportion of repeat Caesarean section. Individualized approach for VBAC seems to be the best.

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