ROLE OF ADMISSION TEST IN PREDICTING ADVERSE FOETAL OUTCOME

M. Mumtaj¹

¹Assistant Professor, Department of OBG, KAPV Government Medical College, Trichy.

ABSTRACT

BACKGROUND

The aim of the evaluation of admission test in high-risk and low-risk groups. Evaluation of role of admission test in predicting adverse outcome of foetus in both low and high-risk groups'.

MATERIALS AND METHODS

This is a cross-sectional study done in Tirunelveli Medical College in a period of six months for 100 patients with equally divided low and high-risk groups.

RESULTS

With normal tracing, 96.9% of babies have no asphyxia. With suspicious trace, 65.2% had no asphyxia. With ominous trace, 58.3% had no asphyxia. In high-risk cases with normal tracings, 85% had no asphyxia. With suspicious tracing, 68.4% had no asphyxia. With ominous tracing, 54.5% had no asphyxia. In short, it is 78.6% sensitive in high-risk cases. In low-risk cases with normal tracing, 97.8% had no asphyxia. With suspicious tracing, 100% had no asphyxia. In short, AT is 91.8% specific in low-risk cases.

CONCLUSION

Admission test is a good intrapartum test both in high-risk and low-risk groups. It is simple, highly acceptable and also it can be repeated. It has 78.6% sensitivity in high-risk cases, 91.8% specificity in low-risk cases and over all negative predictive value is 91%.

KEYWORDS

Admission Test, CTG, Foetal Outcome, Maternal Outcome.

HOW TO CITE THIS ARTICLE: Mumtaj M. Role of admission test in predicting adverse foetal outcome. J. Evid. Based Med. Healthc. 2016; 3(90), 4890-4894. DOI: 10.18410/jebmh/2016/1031

BACKGROUND

The goal of antepartum foetal surveillance is to prevent foetal death. Each and every foetus has a potential risk of intrapartum hypoxia or birth injury and an optimal outcome can be concluded only at the end of labour. However, any definite insult due to the process of labour can only be identified on long-term follow-up.

AIM OF THE STUDY

Evaluation of admission test in high-risk and low-risk groups. Evaluation of role of admission test in predicting adverse outcome of foetus at risk in both low-risk and high-risk groups.

OBJECTIVE

In promoting safe motherhood as defined by the World Health Organization, our objectives must be to optimise the

- 1. Health of the mother.
- 2. Health of the offspring.
- 3. Emotional satisfaction of the mother and her family.

Financial or Other, Competing Interest: None. Submission 29-08-2016, Peer Review 10-09-2016, Acceptance 07-10-2016, Published 08-11-2016. Corresponding Author: Dr. M. Mumtaj, #7/4, Ramarayar Street, Tennur High Road, Tirchy-17. E-mail: mumtajanifa@gmail.com DOI: 10.18410/jebmh/2016/1031



Inclusion Criteria Low-Risk Cases

- Pregnant patients with gestational age of 37 weeks up to 40 weeks.
- With labour pains either spontaneous (or) accelerated.
- With cephalic presentation.

High-Risk Cases

Post-dated pregnancy, pregnancy-induced hypertension, IUGR, oligohydramnios, Rh negative pregnancy, long period of primary infertility, bad obstetric history, heart disease, anaemia, previous LSCS, face presentation.

Exclusion Criteria

Antepartum haemorrhage, multiple pregnancy, major anomalies of a foetus <30 wks.

MATERIALS AND METHODS

In this admissions, test was done for 100 patients in the labour ward at the time of admission.

The patients were followed up according to the AT results.

Patients with normal tracings were followed up by intermittent auscultation and electronic monitoring done once in 4 to 5 hours during monitoring. When we suspected foetal distress, emergency interventions was made according to the stage of labour.⁽¹⁾ In patients with suspicious ominous tracings, immediate ARM done and colour of the liquor was assessed.

Jebmh.com

In patients with thin meconium-stained amniotic fluid, amnioinfusion was given and the labour was allowed to progress. They were followed up carefully by intermittent auscultation and CTG monitoring. When there is a change of colour of liquor or when ominous pattern appears on CTG record according to the stage of labour, the labour was terminated by either forceps or caesarean section. The findings of the admission test are correlated with the outcome of the pregnancy. To evaluate the outcome of pregnancy, foetal distress was considered to be present when ominous FHR changes led to caesarean section or forceps delivery and the newborn had an APGAR score <7 at 5 minutes following spontaneous delivery⁽²⁾ (Arul Kumaran-GIBB).

RESULTS

SI.	Casas	Total	Percentage	
No.	Cases	No.	(%)	
1.	Low-risk cases	50	50	
2.	High-risk cases	50	50	
	Total	100	100	
Break	up of high-risk cases			
	a) Post-dated Pregnancy	17	34	
	b) IUGR/Oligohydramnios	2	4	
	c) Long period of infertility	4	8	
	d) Pre-term labour	1	2	
	e) Bad obstetric history		8	
	f) Heart disease	3	6	
	g) PIH	7	14	
	h) Rh-ve	5	10	
	i) Anaemia	5	10	
	j) Pr. LSCS	1	2	
	k) Face	1	2	
	Table 1. Breakup of High-Risk Cases			

This table shows various types of cases on whom admission test were performed. Among these, low-risk cases form 50% and high-risk cases form 50%.

Age	Total Number of	Percentage	
Group	Patients	(%)	
18-24	64	64	
25-29	30	30	
30-34	3	3	
35-39	3	3	
Total	100	100	
Table 2. Age-Wise Distribution			

Majority of them, 64% fall under 18-24 yrs. Remaining 30% of it is formed by 25-29 yrs. age group persons. 3% is formed by 30-34 yrs. age group persons, 3% is formed by persons between 35-39 yrs. of age.

Gravity	Total Number	Percentage (%)	
Primi	52	52	
G2	29	29	
G3	16	16	
G4	3	3	
Table 3. Obstetric Index			

Among 100 patients, 52 were primi, second gravida was 29%, third gravida was 16% and fourth gravida and above forms 3%.

CTG Patterns	Total Number of Women	Percentage (%)	
Normal tracing	65	65	
Suspicious tracing	23	23	
Ominous pattern	12	12	
Total	100	200	
Table 4. CTG Tracing Pattern in all Cases			

Among 100 cases, normal tracing was observed in 65% of cases, suspicious tracing in 23% of cases and ominous tracing in 12% of cases.

High-Risk	Normal	Suspicious	Ominous		
Cases	Normai	Suspicious	Ommous		
Post-dated	7	6	2		
pregnancy	,	0	2		
IUGR/Oligohyd	_	_	2		
ramnios		_	2		
Long period of	_	2	2		
infertility	-	2	2		
Preterm labour	-	2	-		
Bad obstetric	n	2			
history	2	2	-		
Heart disease	4	1	-		
Pregnancy-					
induced	4	3	2		
hypertension					
Rh negative	3	-	2		
Anaemia	2	2	1		
Previous LSCS	-	1	-		
Face	_	_	_		
presentation	-	-	-		
Total	Total 20 19 11				
Table 5. CTG Tracings in High-Risk Cases					

Among the high-risk cases, 20 cases had normal tracing, 19 cases had suspicious tracing, 11 cases had ominous tracing.

Mode of	Number of	Percentage		
Delivery	Women	(%)		
Labour natural	65	65		
Forceps delivery	15	15		
Caesarean section	20	20		
Total	100	100		
Table 6. Mode of Delivery in all Cases				

Jebmh.com

Among 100 patients, 65% delivered by labour natural, 15 delivered by forceps, 20% delivered by caesarean section.

CTG Tracing	Labour Natural	Forceps Delivery	Caesarean Section	
Normal (n=65)	60	5	-	
Suspicious (n=23)	5	9	9	
Ominous (n=12)	-	1	11	
Total	65	15	20	
Table 7. Mode of Delivery According to CTG Findings				

In normal tracings, out of 65 cases, 60 delivered by labour natural, 5 delivered by forceps. All the cases were delivered by forceps for non-foetal distress indication. Foetal distress may develop during the course of labour due to various reasons like hyperstimulation of uterus, short cord, cord around the neck and intrapartum abruption of placenta.²

CTG Tracing	Labour Natural	Forceps Delivery	Caesarean Section	
Normal (n=20)	15 (75%)	5 (25%)	-	
Suspicious (n=19)	3 (15.8%)	7 (36.8%)	9 (47.4%)	
Ominous (n=11)	0%	1 (9.1%)	10 (90.9%)	
<i>Table 8. Mode of Delivery According to CTG</i> <i>Tracing in High-Risk Cases</i>				

In those with normal tracing (20), 15 (75%) delivered by labour natural, 5 (25%) delivered by forceps. In those with suspicious tracing 3 (15.8%) delivered by labour natural, 7 (36.8%) delivered by forceps, 9 (47.4%) delivered by LSCS.

The patients were taken up for emergency LSCS in view of the high-risk factors such as long period of infertility in (2) case, PIH (3), Anaemia (1), Post dated pregnancy (1), Pr. LSCS (1), BOH (1). In those with ominous tracing, 1 (9.1%) delivered by forceps 10 (90.9%) were delivered by LSCS to avoid foetal distress.

CTG Tracing	Neonatal Outcome (Apgar)			
	No Asphyxia (7-10)	Moderate Asphyxia (6-4)	Severe Asphyxia (<4)	
Normal (n=65)	63 (96.9%)	2 (3.1%)	0	
Suspicious (n=23)	15 (65.2%)	7 (30.4%)	1 (4.3%)	
Ominous (n=12)	5 (58.3)%	4 (33.3%)	1 (8.3%)	
Table 9. Apgar Score According to CTG (All)				

p = 0.0004

There is statistically significant relationship (p<0.05) between CTG findings and Apgar score. Cases with normal CTG findings have high Apgar score. Suspicious and ominous cases give birth to more children with moderate and severe asphyxia.

Out of 100 cases, in those with normal tracings, 63 (96.9%) developed asphyxia 2 (3.1%) developed moderate asphyxia. There were no cases of severe asphyxia. In those with suspicious tracings, 15 (65.2%) developed no asphyxia 7 (30.4%) developed moderate asphyxia 1 (4.3%) developed severe asphyxia. In these ominous tracings, 5 (58.3%) developed no asphyxia. 4 (33.3%) developed moderate asphyxia and 1 (8.3%) developed severe asphyxia.

CTG Tracing	Neonatal Outcome (APGAR)			
	NoModerateSevereAsphyxiaAsphyxiaAsphyxia(7-10)(6-4)(<4)			
Normal (n=20)	17 (85%)	3 (15%)	0%	
Suspicious (n=19)	13 (68.4%)	5 (26.3%)	1 (5.3%)	
Ominous (n=11)	6 (54.5)%	4 (36.4%)	1 (9.1%)	
Table 10. Apgar Score According to CTG				
(in High-Risk Groups)				

The relationship between CTG pattern and Apgar scores is statistically significant (p < 0.05) among high-risk cases.

In high-risk cases in those with normal tracing, 17 (85%) has no asphyxia 3 (15%) has asphyxia. This is attributable to be inherent risk factor in the high-risk groups.

In those with suspicious tracing, 13 (68.4%) has no asphyxia 6 (31.6%) had asphyxia.

In those with ominous tracing, 6 (54.5%) had no asphyxia 5 (45.5%) in asphyxia.

CTG Tracing	Neonatal Outcome (Apgar)			
	No Asphyxia (7-10)	Moderate Asphyxia (6-4)	Severe Asphyxia (<4)	
Normal (n=46)	45 (97.8%)	1 (2.2%)	-	
Suspicious (n=3)	3 (100%)	-	-	
Ominous (n=1)	1 (100%)	-	-	
Table 11. Apgar Score According to CTG (in Low-Risk Groups)				

Jebmh.com

CTG Pattern	Admission Test (n)	Foetal Distress	
Normal tracing	65	4 (6.15%)	
Suspicious tracing	23	6 (26.1%)	
Ominous tracing	12	5 (41.7%)	
Table 12. Results of AT in Relation to the Incidence of Foetal Distress			

p - 0.0001

There exists statistically significant relationship (p<0.05) between the results of AT and incidence of foetal distress among the total study cases.

In all cases, in those with normal tracing, 4 (6.15%) developed distress. In those with suspicious tracing 6 (26.1%) developed foetal distress. In those with ominous tracing, 5 (41.7%) developed foetal distress.

CTG Pattern	Admission Test (n)	Foetal Distress
Normal tracing	20	3 (15%)
Suspicious tracing	19	6 (31.6%)
Ominous tracing	11	5 (45.5%)
Total	50	14
<i>Table 13. Results of AT in Relation to the Incidence of Foetal Distress</i>		

p = 0.0116

There exists statistically significant relationship (p<0.05) between the results of AT and incidence of foetal distress among the high-risk cases.

In high-risk group, in those with normal tracing, 3 (15%) developed distress. In those with suspicious tracing, 6 (31.6%) developed foetal distress. In those with ominous tracing, 5 (45.5%) developed foetal distress.³

CTG Pattern	Admission Test (n)	Foetal Distress	
Normal tracing	46	1 (2.2%)	
Suspicious tracing	3	-	
Ominous tracing	1	-	
Total	50	1	
Table 14. Results of AT in Relations to the Incidence of Total Distress in Low-Risk Group			

p = 0.0002

There exists statistically significant relationship (p<0.05) between the results of AT and incidence of foetal distress among the low-risk cases. With normal tracings, 1 (2.2%) developed foetal distress with suspicious tracing (0%) developed foetal distress with ominous tracing (0%) developed foetal distress.

CTG Pattern	Total No. of Cases	Admitted in ICU for Asphyxia	Percentage (%)	
Normal	65	_	0	
tracing	05	-	0	
Suspicious	22	3	13	
tracing	25	5	15	
Caesarean	12	1	22.2	
tracing	12	т	22.2	
Table 15. Neonatal ICU Admission				

'p' = 0.0006

The relationship between ICU admission and CTG patterns is significant among the total study cases. Normal tracings, admission is nil. With suspicious tracing, admission is 13%. With ominous tracing, admission is 33.3%.

CTG Pattern	Total No. of Cases	Admitted in ICU for Asphyxia	Percentage (%)	
Normal	20	_	0%	
tracing	20	-	0%	
Suspicious	10	3	15 80%	
tracing	19	5	13.070	
Ominous	11	4	36.4%	
tracing	11	т Т	50.770	
Table 16. Neonatal ICU Admission in High-Risk Group				

p = 0.0251

The relationship between ICU admission and CTG patterns is statistically significant among the high-risk cases.

With normal tracing, admission is nil. With suspicious tracing, admission is 15.8%. With ominous tracing, admission is 36.4%.

Screening Test Results	Foetal Distress Present	Foetal Distress Absent	Total
Positive	11	24	35
pattern)	11	27	55
Negative (normal CTG)	4	61	65
Table 17. Prediction of Foetal Distress			

p = 0.00001

Foetal distress prediction results are significantly related to screening test findings (p<0.05) among total cases.¹

Among total study cases, admission test in prediction of foetal distress of sensitivity of 73.3%, specificity 71.8%, positive predictive value of 31.5% and negative predictive value of 93.8%.

Screening Test Results	Foetal Distress Present	Foetal Distress Absent	Total
Positive (abnormal CTG pattern)	1	19	30
Negative (normal CTG)	3	17	20
Table 17. A: Prediction of Foetal Distress in High-Risk Group			

p = 0.0232

Foetal distress prediction results are significantly related to screening test findings (p<0.05) among total cases.⁴ Among total study cases, admission test in prediction of foetal distress of sensitivity of 78.6%, specificity 52.8%, positive predictive value of 36.7% and negative predictive value of 85%.

Screening Test Results	Foetal Distress Present	Foetal Distress Absent	Total
Positive (abnormal CTG pattern)	0	4	4
Negative (normal CTG)	1	45	46
Table 17. B: Prediction of Foetal Distress in Low-Risk Group			

p = 0.0002

Foetal distress prediction results are significantly related to screening test findings (p<0.05) among total cases.

DISCUSSION

AT is used to detect foetal wellbeing and foetal distress if present on admission. This helps us in identifying the group of women who will require continuous electronic monitoring or intermittent auscultation during the course of labour.⁵

Antepartum risk factors are not accurate as predictors of foetal outcome. As foetal heart changes and acidosis occur in same frequency in high-risk as well as low-risk group during the course of labour.⁶

Bearing the acute events during the course of labour, AT will be a good predictor of foetal wellbeing at the time of

admission and during the next few hours of labour in term foetus. It will not predict the development of foetal distress that develops several hours later. (Ingemarsson 1993) Therefore, it can be safely assumed that if the AT is normal, it is enough to perform intermittent auscultation and CTG monitoring once in 4-5 hours. But, abnormal tracings should have continuous monitoring throughout labour to diagnose foetal distress earlier. LLAVENO et al (1990) criticises that the policy of continuous foetal monitoring led to increase in caesarean section with no evidence of foetal benefits. To improve the sensitivity and positive predictive value, false positives and false negatives are to be reduced. This can be done by doing additional tests like Foetal Scalp Blood Sampling (FSBS), Foetal Acoustic Stimulation Test (FAST) to

CONCLUSION

diagnose exactly the foetal distress.

Admission test is a good intrapartum test both in high-risk and low-risk groups. It is simple, highly acceptable and also it can be repeated. It has 78.6% sensitivity in high-risk cases, 91.8% specificity in low-risk cases and overall negative predictive value is 91%. A short recording immediately after admission can detect foetal distress if present and predict wellbeing for next few hours.

REFERENCES

- 1. FIGO Guidelines for the use of fetal monitoring. J Gynecology and Obstet 1987;25:159-167.
- Crawford D, Chapman M, Allan L. The assessment of persistent bradycardia, in prenatal life. Br J Obstet Gynaecol 1985;92(9):941-944.
- Harris JL, Krueger TR, Parer JT. Mechanism of late decelerations of the fetal heart rate during hypoxia. Am J Obstet Gynecol 1982;144(5):491-496.
- Fleischer A, Schulman H, Jagani N, et al. The development of fetal acidosis in the presence of an abnormal fetal heart rate tracing. I. The average for gestational age fetus. Am J Obstet Gynecol 1982;144(1):55-60.
- Ingemarsson I, Arulkumaran S, Ingemarsson E, et al. Admission test: a screening test for a fetal distress in labour. Obstet Gynecol 1986;68(6):800-806.
- Leveno KJ, Cunningham FG, Nelson SRN, et al. A prospective comparison of selective and universal electronic fetal monitoring in 34,995 pregnancies. N Engl J Med 1986;315:615-619.