Mode of Delivery in Meconium Stained Amniotic Fluid: A Descriptive Clinical Study

Laxmi N. Itagi*, Harini R.**

Abstract

Introduction: Meconium is likely to cause both mechanical obstruction of the airways and chemical pneumonitis. The free fatty acids in meconium will strip away alveolar surfactant. Atelectasis, consolidation, pneumothorax and pneumomediastinum may occur that would prove to be fatal. Methodology: During study cases were selected with pregnant women at term gestation with cephalic presentation with meconium stained amniotic fluid, keeping in mind the inclusion and exclusion criteria. Results: 75 patients with thin meconium stained amniotic fluid, 34 (45.33%) had normal vaginal delivery, while in thick meconium stained amniotic fluid out of 125 only 36 (28.8%) delivered normally. Conclusion: Incidence of *Assistant Professor, Dept. LSCS was more in thick meconium stained amniotic fluid as compared to with thin Meconium stained amniotic fluid

Keywords:Meconium; Amniotic Fluid; Mode of Delivery.

Introduction

of Obstetrics and Gynaecology, Mahadevappa Rampure Medical College, Kalburgi. **Senior Resident (DNB), Department of OBG, Apollo Hospitals, Chennai

Laxmi N. Itagi Assistant Professor, Dept. of Obstetrics and Gynaecology, Mahadevappa Rampure Medical College, Sedam Road, Mahadevappa Marg, Kalaburagi, Karnataka 585105. rajirajeshwari24@gmail.com

Meconium aspiration remains the most significant cause of morbidity and mortality during neonatal period. The overall incidence of meconium aspiration in live born infants is 1 to 3%. Between 10 and 30% of meconium stained babies develop varying degrees of respiratory difficulties (Brown 1981) [1].

Meconium is likely to cause both mechanical obstruction of the airways and chemical pneumonitis. The free fatty acids in meconium will strip away alveolar surfactant (Clark 1987) [2]. Atelectasis, consolidation, pneumothorax and pneumomediastinum may occur that would prove to be fatal. Meconium has definite toxic properties of a low grade nature, similar to those of bile, but is much more pronounced in their local effect. The pH of meconium ranges between 5.5-7.

Bacsik [3](1977) explained the pathophysiology of MAS as follows -Mechanical airway obstruction by particles of meconium or by squamous epithelial cells probably plays the most important role in the pathophysiology of MAS. A large amount of meconium is capable of completely obstructing the trachea, resulting in rapid death from asphyxia and acute corpulmonale. Smaller amount move quickly to the lung periphery resulting in obstruction of the distal airway resulting to atelectasis followed by hypoxia. Partial airway obstruction would produce a ball valve effect leading to trapping of air.

Infants of either meconium aspiration or staining showed a significant inflammatory response of lung tissue, but several infants with abundant intra-alveolar meconium showed no significant inflammatory response. Hyaline membranes were evident histologically in six of ten live born infants with meconium aspiration (Perlman 1989) [4].

The irritating action of meconium on the pulmonary parenchyma might initiate a chemical pneumonitis helping to compromise pulmonary function and this could explain the inflammatory changes seen histologically in infants dying of MAS, and partially explain the high incidence of effusion seen on chest X-ray.

Methodology

A prospective study of 200 cases of meconium stained amniotic fluid was studied at Teaching & General Hospital. During study cases were selected with pregnant women at term gestation with cephalic presentation with meconium stained amniotic fluid, keeping in mind the inclusion and exclusion criteria.

Methods of Collection of Data

A careful history is taken from all cases particularly about age, parity, gravidity and duration of labour. Previous obstetric history

Previous obstetric complications

A detailed clinical examination and appropriate investigations.

Inclusion Criteria

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All pregnant women in labour with cephalic

presentation with singleton pregnancy with meconium stained liquor irrespective of age, parity and stage of labour.

Artificial rupture of membranes or spontaneous rupture of membranes.

Pregnancy induced hypertension

Previous normal deliveries and previous LSCS.

Exclusion Criteria

Malpresentation

Multiple pregnancies

Preterm and post-term pregnancy

Maternal medical diseases

Fetal malformation

Intrauterine fetal demise

Obstetric complications: Eclampsia, antepartum hemorrhage

Results

The data collected in this study is presented in the following tables as seen in 200 patients with meconium stained amniotic fluid.

Table 1: Incidence of MSAF					
Total	MSAF	Percent			
5075	704	13.87			

Table 2: Relationship of maternal factors with Meconium Stained Amniotic Fluid

Out of 5075 deliveries conducted, 704 i.e., 13.87% were meconium stained out of which 200 cases were selected for the present study which had inclusion

criteria.

Of the total number of 200 cases, 125 cases had thick meconium and 75 cases thin meconium.

Maternal factors		No. of Deliv	veries	Percentage	
PIH		33		16.50	
Anem	ia	23		11.50	
PROM	Λ	20		10.00	
Non-progress	of labour	6		3.00	
Previous	LSCS	21		10.50	
Total		103		51.5%	
able 3: Incidence o	f gravidity in mec	onium stained amniotic	fluid		
Gravidity	Т	hin		Thick	
,	No.	Percent	No.	Perce	ent
Primi	45	60.00	83	66.4	0
Multigravida	30	40.00	42	33.6	0
Total	75	100.00	125	100.0)0

Out of 200 cases with Meconium stained amniotic fluid 103 cases i.e. 51.5% were associated with maternal complication (both antepartum and intrapartum). Of these 33 were PIH (16.5%) followed by anemia 23 cases (11.5%), PROM – 10%, previous LSCS 10.5% and non-progress of labour in 3% of cases

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because of non-progress of labour.

Incidence of thin meconium is more in primigravida 60.0% as compared to multigravida i.e., 40%. Incidence of thick meconium stained inprimigravida i.e., 66.4% as compared with

multigravida which is 33.6%. There was no difference in the type of meconium with respect to gravidity but overall incidence of meconium stained amniotic fluid was more in primi because period of gestation and duration of labour is more.

Stage of labour when meconium detected	Type of	No. of cases	Mode of delivery				
	meconium		NVD	Vaccum	Forceps	LSCS	
Latent phase	Thin	13	5			8	
-	Thick	33	5	1		27	
Active phase	Thin	20	10	4		6	
	Thick	38	13	5	1	19	
2 nd Stage	Thin	12	5	3	1	3	
	Thick	12	5	4	1	2	
Total		128	43	17	3	65	

Table 4: Meconium first observed and mode of delivery in case of primi

Table 5: Meconium first observed and mode of delivery in case of Multipara

Stage of labour when	Type of	No. of	Mode of delivery				
meconium detected	meconium	cases	NVD	Vaccum	Forceps	LSCS	
Latent phase	Thin	11	1			10	
1	Thick	18	1	3		14	
Active phase	Thin	9	8			1	
1	Thick	23	11	1	1	10	
2 nd Stage	Thin	10	5	4	1		
0	Thick	1	1				
Total		72	27	7	2	35	
Table 6: Mode of delive	ry						
Mode of delivery	Thin	Percent	Thick	Percent	Total	Percent	
Vaginal delivery	34	45.33	36	28.8	70	35.00	
Forceps assisted	2	2.67	3	2.4	5	2.50	
vaginal delivery							
Vaccum assisted	11	14.67	14	11.20	25	12.50	
vaginal delivery							
LSCS	28	37.33	72	57.60	100	50.00	
Total	75	100.00	125	100.00	200	100.00	
c ² =26.089	p<0.001 Highly significant						

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The number of emergency LSCS was more in primi, in whom, meconium was detected early in labour and was thick compared to thin and other stages of labour.

The number of emergency LSCS was more in multi, in whom, meconium was detected early in labour and it was thick compared to thin and other stages of labour. Trial of labour was shortened with cases of previous LSCS with MSAF as indication for LSCS involved in both the conditions.

75 patients with thin meconium stained amniotic fluid, 34 (45.33%) had normal vaginal delivery, while in thick meconium stained amniotic fluid out of 125 only 36 (28.8%) delivered normally. Incidence of LSCS was more in thick meconium stained amniotic fluid i.e., 57.6% as compared to 37.33% with thin Meconium stained amniotic fluid.

Discussion

Table	7:	Incidence	of	MSAF
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Author	Incidence of MSAF (%)			
Kamala G et al⁵	9.37			
Linder et al ⁶	10.40			
Goud and Krishna ⁷	9.80			
Arun ⁸	14.00			
Hari Bhaskar ⁹	11.20			
Present study ¹⁰	13.87			

In the present study the incidence was 13.87%. Out among them 200 cases were randomly selected and studied.

Table 8: Comparative incidence of MSAF with Parity of Mothers

Parity	Kamala et al ¹⁰	Present study
Primi	54.00%	64.00%
2 nd Delivery	24.70%	25.50%
3 rd Delivery	18.00%	5.50%
4th Delivery or more	3.30%	5.00%

Table 9: Comparative incidence of mode of delivery

Mode of delivery	Bhide SS et al ¹¹ (%)		Goud ⁷ (%)		Present study (%)	
	Thin	Thick	Thin	Thick	Thin	Thick
Normal delivery	59.22	49.29	86.48	66.66	45.33	28.28
Forceps	24.27	15.49	4.05	7.84	2.67	2.40
LSCS	16.50	35.21	1.35	21.56	37.33	57.6
Vaccum			5.49	3.92	14.67	11.20

Highest incidence of MSAF was seen in the primigravida i.e., 64%. According to above, present study was correlating with the study done by Kamala Ghokroo et al. More number of cases seen in primigravida i.e., 128 out of 200 cases.

There was increased incidence of operative delivery in our study.

Overall percentage of LSCS was high due to fetal distress as indicated by MSAF and non-reactive NST and is because of cases being referred from peripheral hospital. In the present study, LSCS indication was higher i.e., 57.6%.

Conclusion

Admission to delivery interval is important as well as stage of labour in deciding the mode of delivery and prevention of perinatal morbidity

References

- Brown BL et al, "Intrauterine meconium aspiration" Obstetrics & Gynecology. 1981; 57; 26.
- Clark DA, Nieman GF, Thompson JE, Paskanik AM, Rokhar JE, Bredenberg CE. Surfactant displacement by meconium free fatty acids: An alternative explanation for atelectasis in meconium aspiration syndrome. J. Pediatr. 1987; 110: 765-769.

- Bacsik RD. Meconium aspiration syndrome. Paediatric Clinic of North America. 1977; 24(3): 463-477.
- 4. Perlman EJ, Moore GW, Hutchins GM. The pulmonary vasculature in meconium aspiration. Human pathology. 1989; 20: 701-706.
- Kamala Gokhroo, Usha Sharma et al, "Various maternal factors responsible for meconium stained amniotic fluid", J. Obstetrics & Gynecology of India. 2001; 51: 6.
- Klinger, Mary Celeste MD, Kruse, "Meconium aspiration syndrome: Pathophysiology and prevention", JAM Board FamPract. 1999; 12(6): 450-46.
- Goud P and KrishnaU. Significance of meconium staining of amniotic fluid in labour. Journal of Obstetrics and Gynaecology of India. 1989; 39: 523-526.
- Nayak et al, Asha R Dalal, "meconium staining of amniotic fluid – Significance and fetal outcome", J. of Obstetrics & Gynecology of India. 1991; 41: 480.
- Alchalabi H, Abu-Heija AT, El-Sumnae, Zayed F, Badria LF, Obeidata A. Meconium stained amniotic fluid in term pregnancies – a clinical view. Journal of Obst and Gyn. 1999 May; 19(3): 262-264.
- 10. Kamala Gokhroo, Usha Sharma et al, "Various maternal factors responsible for meconium stained amniotic fluid", J. Obstetrics & Gynecology of India. 2001; 51: 6.
- Bhide SS, Shendurnikar S Aiyer, SR Baxi, "Neonatal outcome after meconium stained amniotic fluid", J. of Obstetrics & Gynecology of India. 1993; 43: 933.

