

Feto-Maternal Outcome with Meconium Stained Liquor: A Clinical Study from Medical College of Ambala

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Abstract

Introduction: Incidence of meconium stained amniotic fluid (MSAF) is about 10–15% of all pregnancies which is related with maturation of the fetal Gastrointestinal Tract (GIT) but may also be associated with unfavorable neonatal outcome. *Objectives:* Aim of our study was to study the fetomaternal outcome in cases with meconium stained amniotic fluid. *Materials and Methods:* The present study was conducted in Maharishi Markandeshwar Medical College & Research Center, Ambala. Total 100 cases with MSAF fulfilling the inclusion criteria were included in the study. All the parameters were analyzed to identify the various fetomaternal outcome. *Results:* we found that the incidence of MSAF was 10.8%, thin MSAF was seen in 74% cases and thick MSAF in remaining 26%. MSAF was present frequently in primigravidas and with increase gestational age. The incidence of abnormal FHR pattern was common in the thick meconium group (69.2%) as compared to thin meconium group (18.9%). The most common mode of delivery in cases with thin MSAF was normal vaginal delivery (51.5%) while in cases with thick MSAF was by lower segment caesarean section (LSCS) (61.5%). The incidence of NICU admission in thin MSAF group was 18.9% while in thick MSAF 46.1% required NICU admission. *Conclusion:* As the incidence of perinatal morbidity and mortality is more in cases with meconium stained amniotic fluid, henceforth the patients with meconium staining of amniotic fluid needs to be monitored vigorously with timely interventions and proper neonatal resuscitation.

Keywords: MSAF-Meconium Stained Amniotic Fluid; LSCS; Fetal Gastrointestinal Tract (GIT).

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Introduction

Assessment of Fetal well being has traditionally been evaluated on the basis of FHR patterns, fetal movements and colour of the amniotic fluid.¹

The presence of meconium-stained amniotic fluid (MSAF) during labor has been long considered the predictor of adverse fetal outcomes such as meconium aspiration syndrome and perinatal asphyxia, which leads to perinatal and neonatal

morbidity and mortality.² The incidence of Meconium staining of amniotic fluid (MSAF) is present in Almost 10-15% of all pregnancies and is comparatively common in term births particularly in postdated deliveries.³ The etio-pathogenesis of MSAF is poorly understood.⁴ Passage of meconium normally occurs within 24–48 hours after birth. In utero passage of meconium by the fetus represents the GI tract maturation but pathologically it has been associated to fetal hypoxia in which there occurs relaxation of the anal sphincter and increased peristalsis due to vaginal stimulation. Hence passage of meconium in utero may simply symbolize the normally developed fetal gastro intestinal tract (GIT) as in postdated pregnancies. Meconium passage is infrequent before 34 weeks of gestation and incidence raises steadily beyond 37 weeks of gestation. According to Royal College of Obstetricians and Gynecologists (RCOG) intrapartum care guideline, meconium stained amniotic fluid is classified as significant MSL and non-significant MSL. Non-significant MSL is defined as a thin yellow or greenish tinged fluid; containing non-particulate meconium whereas significant MSL is explained as dark green

or black amniotic fluid that is thick and tenacious and consists lumps of meconium.⁵ Factors such as placental insufficiency, maternal hypertension, pre-eclampsia, oligohydramnios or maternal drug abuse (tobacco, cocaine) might lead to in-utero passage of meconium.⁶ So the existence of MSAF in other circumstances is an indicator of fetal distress and may be associated with unfavorable fetal and neonatal outcome.^{7,8}

The strong association of stillborn infants, abnormal FHR tracings, neonatal encephalopathy, respiratory distress (Meconium aspiration syndrome, MAS) and abnormal neurological outcome with presence of MSAF has been testified in the literature.⁹ Various indicators of fetal distress are associated with the existence of MSAF as meconium stained new-borns have lower scalp pH and lower umbilical cord artery pH in comparison with infants born with clear amniotic fluid.^{10,11} There have been conflicting consequences reported in labours, complicated by meconium staining of the amniotic fluid, fluctuating with degree of meconium staining.^{12,13} However, in vast majority of cases, no major problem occurs in infants born through MSAF.¹⁴ Presence of meconium below the vocal cords is termed as meconium aspiration syndrome (MAS).¹⁵ Meconium aspiration syndrome occurs all over the world about 5–10.5% of neonates with

MSAF; which accounts around 12% of neonatal mortality (as much as 40% case fatality rate for the neonate and around 2% of perinatal mortality). Furthermore, the rates of severe mental retardation and cerebral palsy are significantly greater among infants born with MSAF.^{16,17}

Materials & Methods

The present study was conducted among patients delivering in department of Obstetrics & Gynaecology of Maharishi Markandeshwar Medical College & Research Center, Ambala over a period of 10 months. Total 100 cases were included in the study who were fulfilling the inclusion criteria and gave consent to participate in the study. Approval from institutional ethical committee was taken

Study duration: 1 Jan 2017 to 31 Oct 2017

Study design: Cross-sectional observational study

Inclusion criteria

1. Term pregnancies (>37 completed weeks)
2. Cephalic presentation
3. Singleton pregnancies

Exclusion criteria

1. Preterm labour
2. Congenital anomalies
3. Multiple gestations
4. Intrauterine fetal death
5. Breech presentation

After taking informed consent from patient fulfilling inclusion criteria detailed history was taken, followed by thorough obstetrical examination, patients were managed as per standard Institutional protocol. Obstetrical & neonatal outcomes were entered as per predesigned Proforma. All collected data was tabulated and appropriate statistical tests were applied. *p*-value <0.05 was taken as statistically significant.

Results

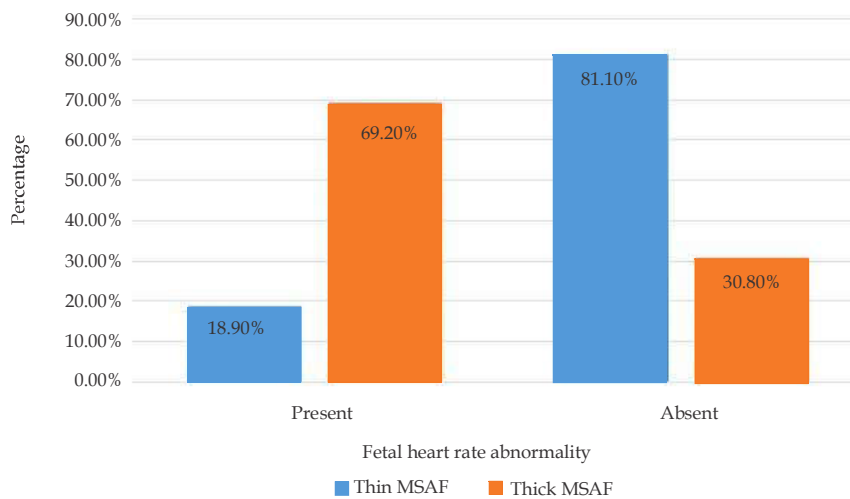
During the study period 2512 deliveries took place in our institute out of which 272 i.e., 10.8% had MSAF out of which 100 were included. MSAF was seen more commonly in primigravidas (56%), The incidence of MSAF was seen to increase with

increase in gestational age i.e., 14% in 37-38 weeks age group, 38% in 39-40 weeks and it increases upto 48% in 41-42 weeks pregnancy (Table 1). Amongst 100 cases with MSAF 74% cases had thin meconium and 26% had thick meconium. As depicted in Figure 1 fetal heart rate abnormalities were observed more commonly in the group with thick meconium group i.e., 69.2% while in the thin meconium group abnormalities present in 18.9% cases. This difference was highly significant

(p -value <0.0001). In cases with thin meconium stained liquor the commonest mode of delivery was normal vaginal delivery (51.5%) followed by LSCS (27.1%) followed by assisted vaginal delivery in 21.6% cases. While in cases with thick MSAF the commonest mode of delivery was by LSCS (61.5%) followed by assisted vaginal delivery. in 23.1% followed by normal vaginal delivery (15.4%). (Figure 2) but the observed difference cannot be considered significant ($p > 0.05$)

Table 1: Distribution as per gestational age

Gest age (in wks)	No. of cases	Percentage
37-38 wks	7	14%
39-40 wks	19	38%
41-42 wks	24	48%



p -value <0.001

Fig. 1: Abnormal fetal heart

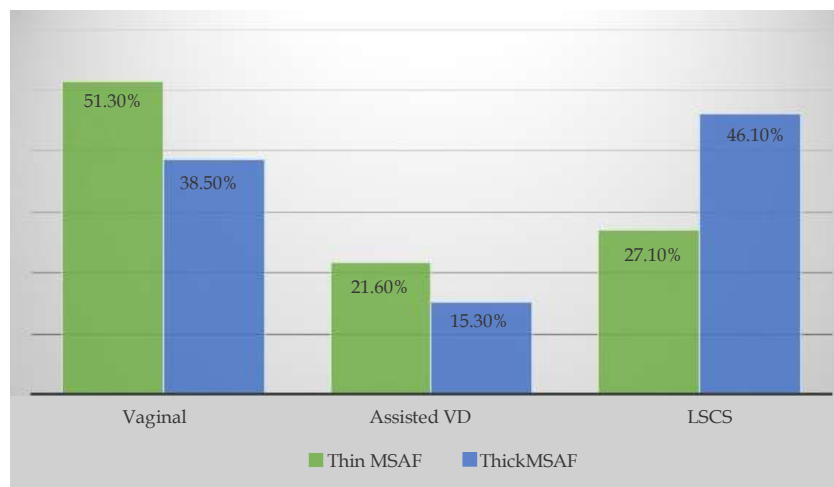


Fig. 2: Mode of delivery in cases of MSAF

Table 2 illustrates that in the group with thin MSAF the Apgar score of less than 5 at 1 minute was seen in 35.1% cases while in 64.9% cases the Apgar score was more than 5. While the Apgar score of less than 5 at 5 minutes was seen in only in 18.9% cases and more than 5 in 81.1% case. On the contrary in the group with thick MSAF Apgar score at one minute was less than 5 in 46.2% cases and more than 5 in 53.8% cases. On assessing the Apgar score at 5 minute 38.4% cases had Apgar score of less than 5 while 61.6% cases had Apgar score more than 5. Cases with thin MSAF 18.9% required NICU admission which amongst cases with thick MSAF 46.1% required NICU admission. On comparing

the perinatal outcome in both the groups in the cases with thin MSAF the morbidity was 18.9% while in the thick MSAF group morbidity was as high as 61.5%. The morbidities were in the form of respiratory distress 62%, seizures 25%, sepsis 1%. Only one mortality was observed in the thick MSAF group the cause of which was meconium pneumonitis developing pulmonary hypertension. Out of cases undergoing lower segment caesarean Section 3 cases had wound infection out of which in one case had to undergo resuturing. All the cases were from the thick meconium group. This can be explained due to increased operative interference in the thick meconium group.

Table 2: Apgar scores in 1 and 5 minutes

	Thin MSAF		Thick MSAF	
	No.	Percentage	No.	Percentage
At 1 minute				
<5	13	35.1%	6	46.2%
>5	24	64.9%	7	53.8%
At 5 minute				
<5	7	18.9%	5	38.4%
>5	30	81.1%	8	61.6%

Discussion

In the present study the incidence of MSAF was 10.8% which is slightly higher than the study done by Nirmala *et al.*¹⁸ and is similar to other studies (Table 3). The incidence of MSAF was more commonly seen in primigravidas than multigravidas which is similar to study done by Becker *et al.* which shows that meconium is common in nulliparous females.¹⁹ In our study the incidence

of MSAF increased with increase in gestational age which is similar to study done by Fisher *et al.* in 2012.²⁰ On dividing the cases into thick and thin meconium group, thin meconium was seen in 74% cases and thick meconium in remaining 26% cases out of total 100 cases with MSAF. The incidence of abnormal FHR pattern was observed more commonly in the thick meconium group (69.2%) as compared to thin meconium group (7%). These findings are similar to that of Gupta *et al.* and Odongo *et al.*^{21,22}

Table 3: Incidence of MSAF in earlier studies

Nirmala <i>et al.</i> ¹²	7.89%
Haribhaskar ¹³	11.2%
Sauda <i>et al.</i> ¹⁴	11.9%
Present study	10.8%

p-value 0.445

In cases with thin meconium stained liquor the commonest mode of delivery was normal vaginal delivery (51.5%) followed by LSCS (27.1%) followed by assisted vaginal delivery in 21.6% cases. While in cases with thick MSAF the commonest mode of delivery was by LSCS (61.5%) followed by assisted vaginal delivery in 23.1% followed by normal vaginal delivery (15.4%). In a study done by Becker *et al.*¹⁹ and Naqui *et al.*²³ also shows

increased incidence of caesarean section in cases with meconium stained amniotic fluid.

In the group with thin MSAF the Apgar score of less than 5 at 1 minute was seen in 35.1% cases while in 64.9% cases the Apgar score was more than 5. On assessing the Apgar score at 5 minute 38.4% cases had Apgar score of less than 5 while 61.6% cases had Apgar score more than 5. Wong *et al.*²⁴ and

Naqui *et al.*²³ also found similar results while Becker *et al.*¹⁹ found no statistically significant difference between two groups. On comparing the perinatal outcome in both the groups in the cases with thin MSAF the morbidity was 18.9% while in the thick MSAF group morbidity was as high as 61.5%. The morbidities were in the form of respiratory distress 62%, seizures 25%, sepsis 1%. Only one mortality was observed in the thick MSAF group the cause of which was meconium pneumonitis developing pulmonary hypertension. Increased perinatal morbidity and mortality was also observed in studies done by Naqui *et al.*²³ and Patil *et al.*²⁵ The incidence of postpartum complications are also increased in the group with thick MSAF as a result of increased operative interventions.

Conclusion

We conclude that the incidence of FHR abnormalities, operative interventions, NICU admissions, perinatal morbidity and mortality and maternal morbidities are higher in the group with thick MSAF as compared with thin MSAF. So to achieve the best outcome; the patients with meconium staining of amniotic fluid require to be monitored vigorously with timely interventions and proper neonatal resuscitation.

References

1. Samiyappa DP, Ghose S, John LB, *et al.* Maternal and perinatal outcome in meconium stained amniotic fluid at term: a case control study. *Int J Reprod Contracept Obstet Gynecol.* 2016;5:3404-10.
2. Addisu D, Asres A, Gedefaw G, Prevalence of meconium stained amniotic fluid and its associated factors among women who gave birth at term in Felege Hiwot comprehensive specialized referral hospital, North West Ethiopia: a facility based cross-sectional study. *BMC Pregnancy and Child birth.* 2018;18:429
3. Wiswell TE, Tuggle JM, Turner BS. Meconium aspiration syndrome: Have we made a difference? *Pediatrics.* 1990;85:715-21
4. Woods JR, Glantz JC. Significance of amniotic fluid meconium. In Creasy RK Reskin R, Editors. *Maternal fetal medicine: Principles and Practice.* Philadelphia WB: Saunders: 1994;413-422.
5. Sarah M. In: SP SM, *et al.*, editors. *managements of meconium stained liquor.* Cyprus: RCOG; 2016. p .1-10.
6. Mundhra R, Agarwal M. Fetal Outcome in Meconium Stained Deliveries. *Journal of Clinical and Diagnostic Research: JCDR.* 2013;7(12):2874-2876
7. Krebs HB, Petres HE Dunn CJ, Jordann HVF Segreti A. Intrapartum fetal heart monitoring. *Am J Obstet, Gynecol.* 1980;137:936-942
8. Mazor M, Furman B, Wiznitzer A, *et al.* Maternal and perinatal outcome of patients with preterm labour and meconium stained amniotic fluid. *Obstetrics & Gynecology.* 1998;86:830-3.
9. Nathan L, Leveno KJ, Carmody TJ, *et al.* Meconium: A 1990 perspective on an old obstetric hazard. *Obstet Gynecol.* 1994;83:329-332.
10. Ramin K, Leveno K, Kelly M. Observations concerning the pathophysiology of Meconium Aspiration Syndrome. *Am J Obstet Gynecol.* 1994;170:312(#124).
11. Starks C Gregory. Correlation of meconium stained amniotic fluid, early intrapartum fetal pH and Apgar scores as predictors of perinatal outcome. *Obstet and Gynecol.* 1980;56(5):604-9.
12. Low JA, Pancham SR, Worthington O, *et al.* The incidence of fetal asphyxia in 600 high risk monitored pregnancies. *Am J Obstet Gynecol.* 1975;121:456-59.
13. Meis PJ, Hall M, Marshall JR, *et al.* Meconium passage: a new classification for risk assessment during labour. *Am J Obstet Gynecol.* 1978;131:509-13.
14. Cunningham FG, Gant FN, Leveno KJ, Giltrap LL, Haulth CJ, Wentron DK, Williams *Obstetrics 21st ed, Mc GrawHill;* 2001.
15. Ahanya SN, Laksnanan J, Morgan BL, *et al.* Meconium passage in utero: mechanism, consequences and management. *Obstet Gynecol Surv.* 2004;60:45-56.
16. Rajput U, Jain A. Impact of meconium stained amniotic fluid on early neonatal outcome. *J Evol Med Dent Sci.* 2013;2(45):8788-94.
17. Sharma U, Garg S, Tiwari K, *et al.* Perinatal outcome in meconium stained amniotic fluid. *J Evol Med Dent Sci.* 2015;48:8319-27.
18. Nirmala Dhuhan *et al.* Meconium staining of amniotic fluid, a poor predictor of fetal response. *J K Science.* 2010 Oct-Dec;12(4):184-186.
19. Becker S, Solomayer E, Dogan C, *et al.* Meconium stained amniotic fluid- perinatal outcome and obstetrical management in rural and suburban population. *Eur J Obstet Gynecol Reprod Biol.* 2007 May;132(1):46-50.
20. Fischer C, Rybakowski C, Ferdynus C, *et al.* A population based study of MAS in neonates born between 37-43 weeks of

- gestation. *Int J Paediatrics* 2012, & pages doi:10.1155/2012/32154.
21. Kumari S, Gupta SN, Mahato IP, *et al.* Health renaissance September to December, 2012;10(3):198-202.
 22. Odogo BE, Ndavi PM, Gachuno OW, *et al.* Cardiotocography and perinatal outcome in pregnant women of 37-42 weeks gestation. *Pak J Surg.* 2010;27(4):292-8.
 23. Naqui SB, Manzoor S. Association of meconium stained amniotic fluid with perinatal outcome with perinatal outcome in pregnant women of 37-42 weeks gestation. *Pak J Surg.* 2011;27(4):292-298.
 24. Wong WS, Wong KS, Chang A. Epidemiology of meconium staining of amniotic fluid in Hong Kong. *Aust N Z J Obstet Gynaecol.* 1985 May;25(2):90-3
 25. Patil KP, Swamy MK, Samantha K.A one year cross-sectional study of management practices of meconium stained amniotic fluid and perinatal outcome. *J Obstet Gynecol India.* 2006;56(2):128-30.
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