INTRODUCTION
Meconium is the first intestinal secretions from fetus. It starts as early as ten weeks of gestation and incidence of intrauterine passage of meconium increases with the gestational age. Meconium reduces the antibacterial property of amniotic fluid by altering the level of zinc in it which leads to intra amniotic infections. Meconium stained amniotic fluid is a common obstetric situation occurring in 07-22% of women in labour. 5% in preterm pregnancy, up to 20% in term and>20% in post term pregnancy.1,2 Meconium aspiration syndrome is believed to result from aspiration of meconium during intrauterine gasping or at the time of first breath. In case of hypoxia, gasping of fetus results in meconium aspiration which neutralizes the surfactant action and promotes inflammation of lung tissues, whereas persistent hypoxia after birth, aspirated meconium results in pulmonary hypertension.3 The aim of this study was to assess feto-maternal outcome following intrapartum amnioinfusion in patients with meconium stained amniotic fluid.

ABSTRACT
Background: In our country a major cause of perinatal mortality and morbidity is meconium aspiration syndrome (MAS) in new-born. The aim of this study is to assess feto-maternal outcome following intrapartum amnioinfusion in patients with meconium stained amniotic fluid and the rate of cesarean deliveries following intrapartum amnioinfusion in patients with meconium stained amniotic fluid.
Methods: This prospective observational study was conducted on 252 patients with pregnancy at or beyond 37 weeks in active labour with moderate to thick meconium stained amniotic fluid following spontaneous rupture or ARM. In such cases amnioinfusion was performed. Continuous electronic FHR monitoring was performed. Emergency LSCS was done when fetal Bradycardia was recorded or in case of non-progress of labor. Fetomaternal outcome will be noted.
Results: In present study most, women had normal vaginal delivery 157 (62.30%) followed by LSCS 93 (36.91%) followed by forceps/vacuum delivery 02 (0.07%). No maternal complication was seen in 230 women (91.26%). Accidenatal hemorrhage was seen in 01 (0.39%) which was managed by emergency LSCS. Out of 252 neonates, 183 asymptomatic neonates at birth, 69 needed resuscitation and in which 52 neonates recovered and shifted back to mother and 17 neonates referred and admitted in NICU for MAS in which 07 were recovered and 10 neonates (3.96%) died due to MAS.
Conclusions: Intrapartum amnio infusion in meconium stained amniotic fluid by diluting the meconium and by decreasing the cord compression decreases the incidence of foetal distress and there by decreases the rate of caesarean section, these all leads to decrease the incidence of maternal and perinatal morbidity and mortality.
Keywords: Meconium, Amniotic fluid, MAS, Amnioinfusion
amnioinfusion in patients with meconium stained amniotic fluid.

METHODS

Detailed history taking, general physical examination and obstetric examination were done. All relevant investigations were sent. After explaining the procedure & written informed consent amnioinfusion was done with a sterile catheter was introduced transcervically to a depth of 30 mm, and a bolus of 600 ml of sterile saline at room temperature was infused under the force of gravity at a rate of 20 ml/min over a period of 30 min. More fluid was infused at the same rate till the returning fluid is clear or up to a maximum of 1,000 ml. Women were assessed by uterine palpation at 15-min intervals for uterine hypertonic contractions. In such cases, amnioinfusion was discontinued. Continuous electronic FHR monitoring was performed.

Augmentation for inadequate uterine contraction was done with use of oxytocin if there was a delay in the progress of labor and no fetal bradycardia recorded. Emergency LSCS was done when fetal Bradycardia was recorded or in case of non-progress of labor. Apgar scoring at 1 minute and 5 minutes was done.

The composite primary outcome measure was the occurrence of perinatal death, moderate or severe meconium aspiration syndrome, or both. In accordance with clinical criteria, the meconium aspiration syndrome was defined as respiratory distress in the first 4 h after birth and categorized as severe (requiring assisted mechanical ventilation) or moderate (requiring oxygen for at least 48 h at a concentration of 40% or greater but without mechanical ventilation). Secondary outcomes included perinatal death or maternal death or serious morbidity or both. Serious perinatal morbidity included moderate or severe meconium aspiration syndrome; hypotonia; assisted ventilation or intubation of more than 5 min duration; a 5 min Apgar score below 7; an umbilical-artery blood pH value below 7.05; abnormal consciousness; the need for tube feeding; convulsions; and a blood or lumbar culture positive for bacteria. Serious maternal morbidity included the presence of any of the following: uterine rupture, amniotic-fluid embolism, accidental hemorrhage requiring urgent delivery, postpartum hemorrhage requiring transfusion, hysterectomy, admission to the intensive care unit, or disseminated intravascular coagulation. All statistical analysis was done by SPSS 21 software.

RESULTS

In our study out of 252 pregnant women, 159 (63.09%) were in more than 40 week gestational age group and 93 (36.90%) were in 37-40 week gestational age group. It showed that postdatism causes passage of meconium during labour due to fetal maturity (Table 1).
It was observed that Out of 252 pregnant women, among 149 (59.12%) labour was induced and among 103 (40.86%) labor was spontaneous (Figure 3). In our study most of women are post dated in which induction of labour was done, so it shows that induction of labour itself increases the risk of passage of meconium. Meconium in Amniotic fluid was detected at cervical dilatation of 5-6 cms in 142(56.34%) women and at cervical dilatation of 3-4 cms in 110(43.63%) women (Table 2).

There was statistically significant difference found in amnioinfusion to delivery interval (Hrs) and obstetrics index (p=0.028) (Table 3).

There was statistically no significant difference found in distribution of study subjects according to mode of delivery and obstetrics index (p=0.243) (Table 4).

Out of 252 neonates, 183 were asymptomatic and required only routine care at birth, 69 neonates needed resuscitation out of which 52 recovered and could be shifted back to mother however 17 neonates had to be admitted in NICU for meconium aspiration syndrome. Out of these 17 neonates, 7 recovered and shifted back to mother . Neonatal death was seen in 10 neonates (3.96%) due to MAS (Table 5).

There was statistically highly significant difference in APGAR score at 1 and 5 minutes (p=0.001) (Table 6).

### Table 2: Cervical dilatation at the time of meconium detection.

| Cervical dilatation (cm) | Number of cases (n=252) | Percentage (%) |
|--------------------------|--------------------------|-----------------|
| 3-4                      | 110                      | 43.63           |
| 5-6                      | 142                      | 56.34           |
| Total                    | 252                      | 100.0           |

| Amnioinfusion to delivery interval (hrs) | Primi gravida | Multi gravida | Grand multi | Total |
|----------------------------------------|---------------|---------------|-------------|-------|
| 1-2                                    | 79            | 45            | 07          | 131   |
| 2-3                                    | 54            | 30            | 00          | 84    |
| 3-4                                    | 30            | 07            | 00          | 37    |
| Total                                  | 163           | 82            | 07          | 252   |

| Chi square value | 10.9 |
|------------------|------|
| Significant P value | 0.028 (S) |

### Table 6: APGAR score at 1 and 5 minutes.

| APGAR Score | At 1 minute | At 5 minutes |
|-------------|-------------|--------------|
|             | Number      | %            |
|             | Number      | %            |
| <7          | 106         | 42.06        |
| >7          | 146         | 57.93        |
| Total       | 252         | 100.0        |

| Chi square value | 19.9 |
|------------------|------|
| Significance 'p' value | 0.00 (HS) |

### Table 5: Fetal outcome.

| Mode of delivery | Primigravida | Multi gravida | Grand multi |
|------------------|--------------|---------------|-------------|
| NVD              | 94           | 57            | 06          |
| Forceps/vacuum   | 01           | 01            | 00          |
| LSCS             | 68           | 24            | 01          |
| Total            | 163          | 82            | 07          |

| Chi square value | 5.46 |
|------------------|------|
| Significant P value | 0.243 (NS) |

### DISCUSSION

In contrast to our study, a Hospital based cross sectional study was conducted at Hiwat comprehensive specialized referral hospital in 2018 showed that mothers whose age greater than 30 years were 5.6 times more likely to develop MSAF during labour than those less than 30 years (AOR=5.6,95% CI=3.35-9.44). Oyelese et al conducted a retrospective cohort study and showed that rates of meconium stained amniotic fluid increased from 1.2% at 32 weeks to 100% at 42 weeks and concluded that rising incidence of meconium stained amniotic fluid with gestational age is consistent with the hypothesis that fetal maturation is a major etiologic factor in meconium passage. It was seen that passage of meconium was more in primigravida, 163 (64.68%) as compared to multigravida 62 (24.60%). Osava et al found that being multiparous was associated as a protective factor against the presence of meconium stained amniotic fluid (33% lower risk), but only for women with a history of normal births.

It was observed that meconium stained amniotic fluid was more common in women in which labour was induced 149 (59.12%) as compared to spontaneous onset of labour 103 (40.86%). Samiyappa et al found that...
MSAF was more common in 118 women (52.2%), in which Induction of labour was done as compare to 108 women (47.8%) in which onset of labour was spontaneous. According to Hospital based cross sectional study was at Hiiwat comprehensive specialized referral Hospital Obstetrics and Gynecology department, from March 02- May 27, 2018 showed that women who had induced labour were 2.6 times more likely to develop MSAF as compared to the spontaneous onset of labour (AOR=2.6, 95% CI=1.39-4.87). It was observed that in present study MSAF was more common in women with cervical dilatation 5-6 cm, 142 (56.34%) followed by 3-4 cm, 110 (43.65%). According to a study conducted by Parth et al average cervical dilatation at the time of detection of MSAF was 5.4 cm. In present study most women had normal vaginal delivery 157 (62.30%) followed by LSCS 93 (36.91%) followed by forceps/vacuum delivery 02 (0.07%). A lower caesarean section rate in the current study is possibly due to the fact that Amnioinfusion increases the volume of fluid around the fetus therefore decreasing the probability of foetal distress related to cord compression. This coincides with the studies by Pierce et al showed lower incidence of caesarean section in the amnioinfusion group 19.7% and instrumental delivery was 18%. Similar observations of lower incidence of operative deliveries have been made by several authors, Das et al showed 18% caesarean section rate in amnio infused group. Choudhary et al reported lower caesarean section rate of 31% in amnio infused group. Sahu et al also showed reduced caesarean section rate in amnio infused group. Das et al reported similar perinatal mortality rate of 1% in women with Amnioinfusion. Partha et al shows 1% perinatal mortality in Amnio infused group. In our study out of 252 women, no maternal complication was seen in 230 women (91.26%). Accidental hemorrhage was seen in 01 (0.39%) which was managed by emergency LSCS. Postpartum hemorrhage was seen in 04 women (1.58%) which was medically managed in 03 women and 01 were required bakri balloon insertion. Blood transfusion was done in 05 cases (1.98%). Mahomed reported no complication related to Amnioinfusion procedure. Pupeeral pyrexia seen in only 11 patients (4.36%) out of 252 patients. The decrease in incidence of pupeeral pyrexia probably due to dilutional effect on bacteria that enter the uterus. Rathore et al and Hofmeyr also found similar results. There was no cases of Amniotic fluid embolism and uterine hyperstimulation/ rupture seen in our study. In our study out of 252 neonates 183 neonates (72.61%) are asymptomatic and required routine care at birth. 69 neonates (27.38%) needed resuscitation. Asnani et al reported that 67.5% neonates required routine care and resuscitative measures required in only in 32.5% neonates in amnioinfused group. 49 out of 69 neonates 52 (20.6%) neonates recovered and shifted back to mother and 17 (6.7%) neonates needed admission in NICU for MAS. The presence of thick meconium amniotic fluid is associated with increased perinatal morbidity and mortality. Though sample size in our study is less but as per our study perinatal mortality can be reduced by amnioinfusion.

CONCLUSION

In developing countries with limited intrapartum facilities Amnioinfusion for meconium stained amniotic fluid improves perinatal outcome and decreases the caesarean section rates. In our study intrapartum amnioinfusion in meconium stained amniotic fluid resulted in reduction in caesarean section rate, significant reduction in meconium aspiration syndrome, low Apgar score and neodeath. So, from our study we can conclude that intrapartum amnioinfusion in meconium stained amniotic fluid by diluting the meconium and by decreasing the cord compression decreases the incidence of foetal distress and there by decreases the rate of caesarean section, these all leads to decrease the incidence of maternal and perinatal morbidity and mortality.

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