Clinical Study

Spectrum of Maternofetal Outcomes during Dengue Infection in Pregnancy: An Insight

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Dengue is a vector transmitted viral infection; tropical and subtropical countries see outbreaks of dengue each year. There is a paucity of literature on effects of dengue infection on pregnancy outcome and this prompted us to undertake a study for better understanding of pregnancy implications with dengue infection. Pregnant women admitted during the seasonal outbreak of dengue between September 2015 and October 2015 were studied and maternal and fetal outcomes in sixteen NS1Ag positive women were analysed. Out of sixteen women diagnosed with dengue fever, three had dengue shock syndrome (DSS) and eight had dengue haemorrhagic fever (DHF). The most common obstetric complication seen in 43% of the cases was oligohydramnios. Bleeding manifestations occurred in seven women and there were three maternal deaths. Perinatal complications included three intrauterine deaths, six nursery admissions, and one neonatal death. Thus dengue infection was associated with high maternal and perinatal mortality. In view of poor obstetric outcomes, this viral infection warrants early admission and prompt management.

1. Introduction

Dengue is a mosquito-borne viral disease. It is caused by viruses of the genus Flavivirus, family Flaviviridae, and Group IV ssRNA. Dengue is transmitted to humans by the mosquito Aedes aegypti and is one of the most rapidly spreading viral infections. There has been a 30-fold increase in its incidence in the last 50 years coupled to increasing migration from rural to urban areas. Annually 50 million people acquire dengue infection worldwide [1]. Dengue remains a major health concern for Southeast Asian countries with cyclic epidemics. In India multiple viral serotypes are circulating and some regions have case fatality rates of 3–5% in general population, which is much higher than other Southeast Asian regions (1%) [2].

Complications of dengue in pregnancy have been scarcely studied. Literature suggests an increased risk of maternal hemorrhage, preterm labour, and oligohydramnios [3–5]. Clinical presentation of dengue may be confused with HELLP syndrome, both conditions having elevated liver enzymes, haemolysis, and low platelet counts; however serology helps in distinguishing the two conditions. Low platelet counts pose a risk of postpartum hemorrhage. As reported in previous studies, dengue is associated with preterm births, low birth weight, fetal deaths, and vertical transmission leading to neonatal thrombocytopenia requiring platelet transfusions [6–10].

In a quest to answer vital issues we critically analysed dengue antigen positive cases in our institute and studied maternal and fetal outcomes.

2. Material and Method

This study was conducted in the Department of Obstetrics and Gynaecology of a tertiary level government hospital, Delhi, India. During the months of September and October 2015, 60 pregnant women were seen in the emergency room with complaint of fever and were admitted for evaluation. As per protocol, dengue PCR (NS1Ag) was done in all women. Out of 60, 16 were dengue positive.

Dengue was diagnosed using NS1Ag and/or IgM serology. Clinical grading was done according to WHO classification and case definitions. An acute febrile illness with two or more clinical manifestations like headache, retroorbital
Table I: Clinical profile of NS1Ag positive patients.

| Parity | POG weeks + days | IgM | Bleeding manifestations | Obstetrics complication | ICU admission | Maternal morbidity | Neonatal outcome |
|--------|-----------------|-----|-------------------------|-------------------------|---------------|-------------------|-----------------|
| 1      | G₁P₂L₂          | 39  | +                       | N                       | Oligo, IUD    | Y                 | NICU (neonatal death-meconium aspiration syndrome with respiratory failure) |
| 2      | G₁P₁L₁          | 38 + 4 | –                      | Y (PPH)                | Oligo, LSCS  | Y                 | PPH, death      |
| 3      | Primi           | 34 + 4 | +                       | N                       | PTVD          | Y                 | Preterm delivery, death NICU (prematurity) |
| 4      | Primi           | 39   | +                       | N                       | Oligo, NVD   | N                 | None NICU (respiratory distress) |
| 5      | G₁P₁L₁          | 36 + 4 | –                      | N                       | Oligo, NVD   | N                 | None Normal     |
| 6      | G₁P₁L₁          | 24 + 4 | –                      | N                       | Oligo, NVD   | N                 | None Normal     |
| 7      | G₁P₁L₁          | 36   | –                       | N                       | Oligo, FGR, NVD | N             | None NICU (LBW) |
| 8      | G₁P₁L₁          | 38   | +                       | N                       | Oligo, IUD   | N                 | None IUD        |
| 9      | G₁P₁L₁          | 17 + 4 | –                      | Y (BPV)                | Spontaneous abortion | N     | Abortion          |
| 10     | P₂L₂            | Post LSCS | –                      | Y (hematoma)           | Rectus sheath hematoma | N     | Hematoma drainage Normal |
| 11     | G₁P₁L₁          | 40   | –                       | Y (2° PPH)             | NVD           | N                 | 2° PPH NICU (meconium aspiration) |
| 12     | Primi           | 34 + 2 | –                      | Y (epistaxis)          | NVD           | N                 | Epistaxis Normal |
| 13     | Primi           | 39   | –                       | Y                       | Forceps delivery | N     | PPH Normal       |
| 14     | G₁P₁L₁          | 32 + 2 | +                      | Y                       | PTVD          | N                 | Preterm delivery NICU (prematurity) |
| 15     | G₂P₂L₂          | 37 + 5 | –                       | N (bleeding P/R, petechiae, and subconjunctival hemorrhage) | NVD           | N                 | None Normal     |
| 16     | Primi           | 38 + 5 | –                       | Y (hematuria)          | IUD           | N                 | None IUD        |

G: gravidae, P: parity, L: living issue, VD: vaginal delivery, NVD: normal vaginal delivery, IUD: intrauterine death, PTVD: preterm vaginal delivery, PPH: postpartum hemorrhage, 2° PPH: secondary PPH, LBW: low birth weight, oligo: oligohydramnios, Y: yes, N: no, and NICU: neonatal intensive care unit.

pain, myalgia, arthralgia, rash, hemorrhagic manifestation, or leukopenia and positive serology or occurrence at a time of dengue outbreak was taken as the definition of dengue fever (DF). Dengue hemorrhagic fever (DHF) was classified as fever, hemorrhagic tendencies, thrombocytopenia, evidence of plasma leakage, association of hepatomegaly, and circulatory disturbances. Increase in serial hematocrit was taken as evidence of plasma leakage and ultrasound showing pleural effusion, ascites, and gall bladder edema were taken as supportive evidence. Dengue shock syndrome (DSS) was classified when DHF symptoms included rapid and weak pulse, narrow pulse pressure of less than 20 mmHg, and hypotension [2].

Patients were managed with antipyretics, adequate hydration, and blood product transfusion as necessary. Strict maternal and fetal surveillance were done to identify complications early. Platelet counts were done two to three times per day depending on clinical profile. Obstetric data, clinical,
3. Results and Discussion

A majority of women were unbooked and presented with classical symptoms of high grade fever, myalgia, headache, and rashes to the emergency room. All women were young, age ranging from 22 to 32 years; mean age was 25 years. Out of 16 cases 13 presented in third trimester and two in second trimester and one was referred after LSCS was done in another hospital. She presented with large rectus sheath hematoma following caesarean section needing surgical exploration and drainage. None of the women had any underlying diseases or comorbidity except one (#case 5) who had moderate anemia on admission.

In the current study, eight women (50%) had dengue haemorrhagic fever (DHF). Ten women required transfusion to maintain platelet count in desired range; on average each woman required six platelet transfusions. Most women had low platelet counts at the time of admission (mean 81,000/mm$^3$); 62% had platelets less than 50,000/mm$^3$. Platelet count less than 30,000/mm$^3$ was seen in dengue shock syndrome with increased risk of bleeding manifestations. A Sri Lankan case series reported 10 women with DHF but only eight required platelet transfusion [5]. Three women succumbed to dengue shock syndrome in current study with a mortality rate of 18% compared to 3–5% case fatality rate in rest of the India in nonpregnant population [2]. In another study done in India by Agrawal et al. maternal mortality rate was 12% [3].

Symptomatic dengue infection may increase the risk of preterm labour and, hence, low birth weight (LBW) as suggested by previous studies [3–5, II]. In the study under discussion two women had preterm delivery (12%). Most women presented near term ($n = 10$) and of the four who presented in early gestation two had preterm labour. Preterm births reported were as low as 11% in the French Guiana study versus 41% in the study by Basurko et al. [4, II]. One woman had spontaneous abortion (6%) and seven (43%) pregnancies were complicated with oligohydramnios in present study compared to 52% rate of oligohydramnios from a study in India by Agrawal et al. [3].

Bleeding manifestations were seen in seven women and three (19%) had postpartum hemorrhage (PPH) (#2, 11, and 13). One of them was diagnosed as DSS and the other two were diagnosed as DF. Both women diagnosed as DF had atonic PPH. One of them had PPH 48 hours following the delivery. Both the patients did not have any other bleeding manifestations and signs of plasma leakage hence were classified as only DF. In retrospective study by Basurko et al. on 53 pregnant women, PPH was reported in 10% of the cases compared to 19% in our study [4].

There were six neonatal intensive care unit (NICU) admissions (37%), three intrauterine deaths (IUD) (18%), and one neonatal death (6%). One neonatal death was caused by meconium aspiration syndrome. None of the neonates in our study showed symptoms of dengue such as thrombocytopenia, anemia, fever, or bleeding manifestations. Diagnostic tests for dengue were not routinely done for asymptomatic neonates. There are a number of studies on poor fetal outcome following dengue infection in parturient females. Vertical transmission of dengue has also been reported [4, 5]. Basurko et al. reported 5.3% vertical transmission rate in their study [4]. They also reported IUD rate of 3.8% and neonatal death rate of 1.9% while Carles et al. also reported high fetal death rate [4, 12].

Pregnant women with dengue fever should be considered for admission early because of its unpredictable course.

| Platelet count ($10^3$/μL) | Hematocrit (%) | Blood product transfusion | Diagnosis |
|-----------------------------|----------------|--------------------------|-----------|
| Admission Least | Admission Maximum | Least | | |
| 1 24 22 | 39 45 | 36  | Y | DSS |
| 2 18 18 | 33 37.6 | 33  | Y | DSS |
| 3 79 19 | 35 37 | 31.1  | Y | DSS |
| 4 123 123 | 29 29 | 28  | N | DF |
| 5 40 24 | 23.4 30 | 23.4  | Y | DHF |
| 6 45 45 | 30 31 | 28  | N | DF |
| 7 170 14 | 32.4 34.9 | 29  | Y | DHF |
| 8 24 20 | 27.8 30 | 27  | Y | DHF |
| 9 18 18 | 42 42 | 31  | Y | DHF |
| 10 32 32 | 38 44 | 33  | Y | DHF |
| 11 90 62 | 33 36 | 33  | N | DF |
| 12 190 43 | 32 39 | 32  | N | DHF |
| 13 76 76 | 29 33 | 29  | N | DF |
| 14 42 39 | 40 41 | 35  | Y | DHF |
| 15 310 160 | 31 32 | 32  | N | DF |
| 16 16 16 | 39 39 | 30  | Y | DHF |
Maintaining normothermia and adequate hydration should be the goal. Low platelet counts should be taken as a marker of severity of the disease. Oligohydramnios is an ominous sign when seen concomitantly with dengue infection because the high fetal mortality was associated with it. The exact cause of oligohydramnios is not known but dehydration associated with dengue fever may be contributory. Hydration should be maintained by encouraging oral intake of oral rehydration solution (ORS), fruit juice, and other fluids containing electrolytes and glucose for replacing losses from fever and associated vomiting [2]. Dengue fever does not warrant any active obstetrical intervention.

Another dilemma faced was in differentiating dengue infection from HELLP syndrome as both have somewhat similar laboratory parameters; however serology will help clinch the diagnosis. Careful examination and high index of suspicion by the treating obstetrician are required to diagnose and manage such cases especially during epidemics.

4. Conclusions

Our knowledge on effect of dengue on pregnancy is somewhat limited. In this study dengue was associated with high maternal and fetal morbidity and mortality. Dengue needs early diagnosis and prompt treatment to avoid its adverse outcomes. Pregnant women should avoid travel to dengue afflicted regions. Vector control methods should be employed during seasonal outbreaks. Awareness programs and medical education programs on the management of the dengue in pregnancy should be initiated especially during outbreaks to provide quality care.

Competing Interests

The authors declare that there are no competing interests regarding the publication of this paper.

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