

DOI: <http://dx.doi.org/10.18203/2320-1770.ijrcog20160862>

Research Article

Maternal and fetal outcome in jaundice complicating pregnancy: a prospective study

Swati Sharma, Rupa Aherwar*, Shashikala Jawade

Department of Obstetrics & Gynaecology, Chirayu Medical College, Bhopal, India

Received: 27 January 2016

Accepted: 01 March 2016

***Correspondence:**

Dr. Rupa Aherwar,

E-mail: ahirwar.rupa@gmail.com

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ABSTRACT

Background: The objective of the study was to study maternal and fetal outcome in pregnancy complicated with jaundice.

Methods: 30 antenatal patients with clinical /laboratory evidence of Jaundice were selected for study in period between august 2014 to august 2015.

Results: The peak age of incidence in our study was 21-25 years (66.6%) and majority were primigravida (66.6%). All cases were in third trimester of pregnancy, 93.3% were unbooked, 73.3% were term, 60% were of lower socioeconomic status and 73.3% were urban. All patients presented with jaundice at time of admission. Pruritus was most common presenting symptom present in 60% of patients. Other presenting complaints were nausea, high BP, abdominal pain and petechiae. Viral Hepatitis was most important cause of jaundice in this study found in 46.7% of cases. Preeclampsia and ICP were other causes of jaundice in this study. Hepatitis B was the most common cause of acute hepatitis (26.7%) and incidence of hepatitis E was 13.3% in our study. Maternal mortality was found in 1 case of hepatitis E complicated with hepatic encephalopathy and coagulopathy. 2% of these patients developed FHF. All patients were kept in ICU for intensive monitoring. PPH was most common maternal complication in 60% of patients. There was 1 maternal death in our study. Of 30 patients, 12 had spontaneous onset of labour. All delivered vaginally of which 8(26.7%) were preterm of which 2 died, 4(13.3%) were IUGR, 12 (40%) had fetal distress with meconium stained liquor, 5 (16.6%) had PROM, 2 (6.7%) had fetal growth restriction and 2 (6.7%) delivered uneventfully.

Conclusions: Jaundice in pregnancy results in a very high perinatal as well as maternal morbidity and mortality, and requires an early diagnosis and careful management.

Keywords: Jaundice in pregnancy, Viral hepatitis

INTRODUCTION

Liver dysfunction during pregnancy is multifactorial in origin and diagnosis is often challenging. The key to maternal and fetal wellbeing is an early diagnosis and appropriate management.

Hepatic disorders complicate about 3% of all pregnancies and fall under various categories.¹ First is a heterogeneous group of liver disorders that are unique to pregnancy and occur in patients with a previously healthy

liver. These include intrahepatic cholestasis of pregnancy, acute fatty liver of pregnancy and liver dysfunction associated with hyperemesis gravidarum and preeclampsia. These conditions remit spontaneously in puerperium.

Secondly, pregnancy may occur in background of a pre-existing liver disease e.g. chronic viral hepatitis and cirrhosis liver. Third category is of common viral diseases like acute viral hepatitis which may occur incidentally during pregnancy. The fourth situation is that

of disorders which are probably related to pregnancy e.g. biliary tract disease and Budd-Chiari Syndrome.

Although still enigmatic, there have been recent interesting advances in understanding of these unique pregnancy-related liver diseases. Hyperemesis gravidarum is intractable, dehydrating vomiting in the first trimester of pregnancy; 50% of patients with this condition have liver dysfunction. Intrahepatic cholestasis of pregnancy is pruritus and elevated bile acids in the second half of pregnancy, accompanied by high levels of aminotransferases and mild jaundice. Maternal management is symptomatic with ursodeoxycholic acid; for the fetus, however, this is a high-risk pregnancy requiring close fetal monitoring and early delivery.

Severe preeclampsia itself is the commonest cause of hepatic tenderness and liver dysfunction in pregnancy, and 2%-12% of cases are further complicated by hemolysis (H), elevated liver tests (EL), and low platelet count (LP)-the HELLP syndrome. Immediate delivery is the only definitive therapy, but many maternal complications can occur, including abruptio placentae, renal failure, subcapsular hematomas, and hepatic rupture. Acute fatty liver of pregnancy is a sudden catastrophic illness occurring almost exclusively in the third trimester; microvesicular fatty infiltration of hepatocytes causes acute liver failure with coagulopathy and encephalopathy. Early diagnosis and immediate delivery are essential for maternal and fetal survival.

Pregnancy causes very few alterations in the results of standard liver tests. The aminotransferases (AST and ALT), -glutamyltranspeptidase (GGTP), total bilirubin, and serum bile acid level remain within the normal range. Alkaline phosphatase rises modestly in the third trimester. The albumin level is lower than in nonpregnant women, and the cholesterol level higher Guntupalli et al, Maryam et al.^{1,2} Thus, elevations in aminotransferases or GGTP signify pathology, and should prompt a search for disease.

Making the correct diagnosis is of paramount importance, as failure to do so can result in morbidity or mortality for not only the mother, but also for her fetus. This study has been carried out to evaluate causes and maternal fetal outcome in pregnancies complicated with jaundice.

METHODS

This is a prospective study conducted in department of obstetrics and gynecology in a tertiary care referral hospital; Chirayu medical college & Hospital, Bhopal during one year period from august 2014 to august 2015. During this period 1524 pregnant patients were admitted of which 30 patients with clinical /laboratory evidence of icterus were selected for study.

A detailed history was taken and general, systemic and obstetric examinations were carried out. Liver function

tests including serum bilirubin, SGOT, SGPT, alkaline phosphatase, Australia antigen, prothrombin time (PT), partial thromboplastin time (PTT), bleeding time (BT), clotting time (CT) and platelet count were done. The maternal outcome was noted in terms of the mode of termination of pregnancy, maternal complications and maternal end result. Fetal outcome was assessed by perinatal morbidity and mortality, neonatal intensive care need.

The results were tabulated and data analysed as frequencies, percentages and descriptive statistics.

RESULTS

Table 1: Demographic profile of patients (n=30).

Age in years	Number of cases (n = 30)	percentage
15 -20	2	6.7
21 - 25	20	66.66
26 - 30	8	26.7
Socioeconomic status		
Low income group	18	60
Medium income group	8	26.7
High income group	4	13.3
Residence		
Urban	22	73.3
Rural	8	26.7
ANC status		
booked	2	6.7
unbooked	28	93.3
Gravida		
1	20	66.66
2	8	26.7
3	2	6.7
Gestational age		
term	22	73.3
preterm	8	26.7

Table 1 shows the peak age of incidence between 21-25 years (66.6%) and majority were primigravida (66.6%). All cases were in third trimester of pregnancy, 93.3% were unbooked, 73.3% were term, 60% were of lower socioeconomic status and 73.3% were urban.

Table 2 shows that all patients presented with jaundice at the time of admission. Pruritus was the most common presenting symptom present in 60% of patients. Other presenting complaints were nausea, high BP, abdominal pain and petechiae.

Table 3 shows viral Hepatitis was the most important cause of jaundice in this study found in 46.7% of cases. Preeclampsia and ICP were other causes of jaundice in this study. Hepatitis B was the most common cause of acute hepatitis (26.7%) and incidence of hepatitis E was 13.3% in our study. Maternal mortality was found in 1

case of hepatitis E complicated with hepatic encephalopathy and coagulopathy.

Table 2: Clinical presentation at the time of admission (n=30).

Signs and symptoms	Number of cases	percentage
Nausea /vomiting	14	46.67
Pruritus	18	60
Yellow discoloration of skin, eye & urine	20	60.67
Abdominal pain	7	23.33
Pallor	25	83.33
Icterus	30	100
Edema	9	30
Preeclampsia	10	33.33
Petechiae	6	20

Table 3: Aetiology of jaundice in pregnancy (n=30).

Aetiology of jaundice	Number of cases	percentage
HELLP	4	13.3
PIH	10	33.3
ICP(intrahepatic cholestasis of pregnancy)	2	6.7
Hepatitis A	2	6.7
Hepatitis B	8	26.7
Hepatitis E	4	13.3
Cirrhosis liver	0	0
Acute fatty liver of pregnancy	0	0

Table 4: Maternal complications.

Maternal complication	Number of cases	percentage
Preeclampsia-Eclampsia	10	33.3
Preterm labour	8	26.7
ARF	4	13.3
DIC	6	20
PPH	18	60
Encephalopathy	1	3.33
Fever	10	33.3
Multiorgan failure	4	13.3
ICU admission	30	100
Blood/Blood products transfusion	18	60
Shock	2	6.7
Maternal death	1	3.33

Table 4 shows that all patients were kept in ICU for intensive monitoring. PPH was most common maternal complication in 60% of patients. 60% of patient received blood and component therapy and 13.3% developed

multiorgan failure. There was 1 maternal death in our study.

Table 5: Fetal complications.

Fetal complication	Number of cases	percentage
Preterm	8	26.7
IUFD	4	13.3
Meconium stained liquor	12	40
Uneventful	2	6.7
Premature rupture of membranes	5	16.66
Fetal growth restriction	2	6.7

Table 5 shows that out of 30 patients, 12 had spontaneous onset of labour. All delivered vaginally of which 8 (26.7%) were preterm of which 2 died, 4 (13.3%) were IUFD, 12 (40%) had fetal distress with meconium stained liquor, 5 (16.6%) had PROM, 2 (6.7%) had fetal growth restriction and 2 (6.7%) delivered uneventfully.

DISCUSSION

Liver disease in pregnancy can manifest as a benign disease with abnormal elevation of liver enzyme levels and a good outcome, or it can manifest as a serious entity affecting hepatobiliary function and resulting in liver failure and death to the mother and her fetus. There are no clinical markers that predict the course of a pregnancy and the pathophysiologic mechanisms are not always understood. The overall mortality attributed to liver disorders in pregnancy has dramatically decreased in the past few years because of clinicians' understanding of the physiologic changes that occur during pregnancy, their vigilance in recognizing clinical and laboratory abnormalities, identifying the aetiology and its effective management in a timely manner. A coordinated team approach that involves the primary care physician, obstetrician, hepatologist, is often required to promote good maternal and fetal outcomes.

This study was done in department of Obstetrics and Gynecology in Chirayu medical college from August 2014 to August 15. During this period 1524 pregnant patients were admitted of which 30 patients with clinical/laboratory evidence of icterus was included for study.

The incidence of liver disorders in pregnancy varies in different parts of the world. Liver disease in pregnancy can present with subtle changes in liver biochemical profile or with fulminant hepatic failure (FHF). The overall incidence of liver disorder in pregnancy in our institution (1.96%) was comparable with incidence of 0.4% in study of Acharya N et al and 0.3% in Oladokun et al study.^{3,4} Liver disorders affect at younger age group of patients, the peak age being 21-25 years

(66.6%) in our study. Majority (66.6%) of affection was found in primigravida in third trimester of pregnancy. Study done by Aparajita et al showed incidence of 52.9% in younger age group and 51% were primigravidas.⁵ In our study, 93.3% patients were unbooked, 73.3% were term, 60% were of lower socioeconomic status and 73.3% were urban.

Jaundice as a result of Viral Hepatitis was most important cause in our study; was found in 46.7% of cases. Preeclampsia 33.33% and ICP 6.7% were other causes found. Cholestatic jaundice was found to be the most common cause (54.9%) of liver dysfunction associated with pregnancy in Aparajita et al study.⁵

All patients presented with jaundice at time of admission. Pruritus was most common presenting symptom present in 60% of patients. Other presenting complaints were nausea, high BP, abdominal pain and petechiae. Incidence of Pruritus is consistent with studies of Aparajita et al they found it in 76.5% of cases.⁵

Hepatitis B was the most common cause of acute hepatitis (26.7%) and incidence of hepatitis E was 13.3% in our study; maternal mortality was found in 1 case of hepatitis E complicated with hepatic encephalopathy and coagulopathy. Hepatitis E was the most common cause of acute hepatitis in Aparajita et al study.⁵ It was commonly associated with FHF and high maternal and perinatal morbidity and mortality with 2% patients developing FHF in Kumar A, Beniwal et al study and 16.66% maternal mortality in study of Reddy et al.^{6,7}

All patients were kept in ICU for intensive monitoring. PPH was most common maternal complication in 60% of patients for which uterine balloon tamponade was done and blood products (FFP) were given. There was 1 maternal death in our study. Nearly 2% of the patients required ICU admission in Aparajita et al study study.⁵ Intensive care is a necessity in these cases and various studies had ICU admissions ranging from 4.3% to 62.6% in Pollock et al study.⁸

Of 30 patients, 12 had spontaneous onset of labour. All delivered vaginally of which 8(26.7%) were preterm of which 2 died, 4 (13.3%) were IUFD, 12 (40%) had fetal distress with meconium stained liquor, 5 (16.6%) had PROM, 2 (6.7%) had fetal growth restriction and 2

(6.7%) delivered uneventfully. In Aparajita et al study (5), 94.2% were live births and 5.7% fresh still birth and incidence of prematurity was 13.7 %.

Our present study reemphasizes on the fact that there is increased maternal and fetal morbidity and mortality in pregnancy complicated with jaundice and hence, requires early interventions as timely inductions, PPH prophylaxis assuring availability of adequate blood products for overcoming associated coagulopathy and intensive monitoring of both mother and fetus which requires team work of obstetrician, neonatologist, intensivist, hepatologist and haematologist.

Funding: Not required

Conflict of interest: None declared

Ethical approval: The study was approved by the Institutional Ethics Committee

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Cite this article as: Sharma S, Aherwar R, Jawade S. Maternal and fetal outcome in jaundice complicating pregnancy: a prospective study. *Int J Reprod Contracept Obstet Gynecol* 2016;5:1084-7.