Salmonella typhi and Pregnancy: A Case Report

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Abstract

Background: Salmonella typhi may be a cause of significant morbidity and mortality in both the mother and fetus. Febrile illness during pregnancy, especially that associated with hemolysis, is associated with chorioamnionitis, pyelonephritis, or viral syndrome. As such, S. typhi should be considered when a patient presents with a fever and hemolysis. We present a case of S. typhi complicating pregnancy.

Case: A primigravida at 26 weeks gestation presented with persistent spiking temperature and severe hemolysis. Her presenting signs and symptoms included fever, vomiting, cough, earache, jaundice, dark urine, and anemia. After 4 units of blood, her posttransfusion hematocrit was 29%. Hemolysis was evident on a peripheral blood smear. Total bilirubin was 2.5 mg/dl and direct bilirubin was 1.2 mg/dl. Gentamicin and clindamycin were administered empirically. Stool, blood, and urine cultures were obtained. Blood cultures were positive for *S. typhi*. Antibiotic coverage was changed to gentamicin and ceftriaxone. She defervesced on the 5th day and had no further problems. A healthy male infant was delivered vaginally at term.

Conclusion: Typhoid fever is a serious infection, and special concern arises when S. typhi complicates pregnancy. The diagnosis of S. typhi should be considered in a gravida presenting with fever with prompt institution of antibiotic therapy. Appropriate cultures are essential for confirming the diagnosis. © 1994 Wiley-Liss, Inc.

Key words				
Hemolysis,	typhoid f	fever,	microbiologic clues	

In 1901, Dr. Frank W. Lynch¹ reported 1,079 cases of typhoid fever at the Johns Hopkins Hospital. Two hundred eighty-nine of these cases were females but only 5 occurred during pregnancy. Subsequently, Hicks and French² in 1905 reported a 65-85% abortion or premature labor rate when *Salmonella typhi* infection complicated pregnancy. With the advent of antibiotics as well as improved sanitation, food processing, and storage, the incidence of typhoid fever in the United States has declined. Despite this decline, typhoid fever remains a serious complication in pregnant women. We present a case of *S. typhi* complicating pregnancy to emphasize the importance of an evaluation of geographic risks and microbiologic clues as well as hematologic sequelae.

CASE REPORT

A 21-year-old Hispanic primigravida at 26 weeks gestation was transferred to our institution for persistent spiking temperature and hemolysis. She had been admitted to her local hospital with fever to 40°C, vomiting, cough, and an earache. During that admission, the patient was noted to have dark urine, anemia, and jaundice. A presumptive diagnosis of hepatitis was made and she received 4 units of blood. Seven days prior to admission, she was prescribed amoxicillin, even though she had a his-

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tory of penicillin allergy. An obstetric abdominal ultrasound was consistent with a 26-week gestation, as well as a normal liver and gallbladder. The initial vital signs at transfer were a blood pressure of 100/70, pulse of 86, respiratory rate of 18, temperature of 37.8°C. She was jaundiced but had no petechiae or other skin lesions. Her chest was clear and she had no costovertebral angle tenderness. Her abdomen was soft without tenderness and no hepatosplenomegaly. Her fundal height was 25 cm with a soft nontender uterus. A bimanual examination revealed a long, closed cervix and intact membranes. On admission, her hematocrit was 29.3%, platelet count 187,000/mm³, and white blood cell (WBC) count 11,500/mm³ with a differential of 81% polymorphonuclear cells, 17% lymphocytes, 4% monocytes, and 1% basophils. Her peripheral smear revealed evidence of hemolvsis even though it was obtained posttransfusion. A urinalysis was significant for a specific gravity of 1.010, pH of 7.0, bilirubin of 4.0, and urobilinogen of 1.2. The results of serum chemistry revealed a potassium of 3.3 mEq/l, glucose of 184 mg/dl, blood urea nitrogen of 6 mg/dl, and albumin of 2.8 g/dl. An evaluation of liver function revealed an aspartate aminotransferase of 69 U/l, alkaline phosphatase of 143 U/l, total bilirubin of 5.4 mg/dl, direct bilirubin of 1.8 mg/dl, and a γ -glutamyltranspeptidase of 72 U/l. Stool, urine, and blood cultures were obtained. Gentamicin and clindamycin were administered empirically with no laboratory evidence of further hemolysis. Blood cultures were subsequently positive for Salmonella species, despite the fact that the patient had received ampicillin. Antibiotic coverage was changed to gentamicin and ceftriaxone. She defervesced on the 5th hospital day and was discharged on hospital day 10 with a repeat hematocrit of 23.8%. Urine and stool cultures were sterile. The final report on the organism isolated from a blood culture was S. typhi. She delivered vaginally a 2,845 g male infant with an Apgar score of 9 at both 1 and 5 minutes at 38 weeks estimated gestational age. There were no further complications, and both mother and infant left the hospital in good condition.

DISCUSSION

S. typhi is a facultative anaerobic gram-negative enteric rod that causes a wide variety of human infections when ingested in contaminated food or

TABLE I. Common presenting signs and symptoms of typhoid fever^{3,4,6,9}

Persistent fever
Abdominal pain
Diarrhea
Hypothermic response to antipyretics
Headache
Cough
Constipation
Splenomegaly
Rose spots (small petechial hemorrhages)
Generalized malaise
Anorexia
Chills
Relative bradycardia
Leukopenia

water. In addition to typhoid fever, focal systemic infections, septicemia, and gastroenteritis can occur.³ The common presenting findings are summarized in Table 1. Infection with S. typhi is relatively rare and the incidence of infection in the United States has remained unchanged in recent years at 0.2/100,000 population.³ Salmonella is most commonly acquired during travel to countries in which it is endemic, e.g., Mexico, Indian subcontinent, Southeast Asia, and Indonesia.³ Alexandria, Egypt, Jakarta, Indonesia, and Santiago, Chile, are the leading geographic locations. The incubation period ranges from a few days to several months with associated headache, malaise, and chills. A recurrent febrile response is usually seen after the bacteria invade the mucosa of the small intestine which, in turn, causes a transient bacteremia. This is followed by continued multiplication of organisms within phagocytic cells and sustained episodes of secondary bacteremia.

Special concern arises when pregnancy is complicated by *S. typhi* (Table 2). Prior to the antibiotic era, maternal mortality was increased and the fetus was at risk for infection with subsequent abortion.² More recently, Riggall and associates⁴ reported an improved prognosis for both mother and fetus in 7 cases of typhoid fever complicating pregnancy. The diagnosis was confirmed in all cases by blood culture. Therapy consisted of high doses of ampicillin for 2 weeks. All but 1 pregnancy resulted in good outcomes. One pregnancy ended in a spontaneous abortion.

Although typhoid fever is a rare disease in the United States, sporadic infections do occur. Ac-

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TABLE 2.	Various adverse outcomes and morbidity	
associated	with typhoid fever in pregnancy ^{1,2,4}	

Maternal mortality
Blood transfusions
Premature labor
Spontaneous abortion
Fetal infection

TABLE 3. Geographic areas of 203 reported cases of S. *typhi* in the United States from January 1, 1994, to July 23, 1994⁵

Reporting area	No. of cases		
Pacific	50		
Mid-Atlantic	49		
East/North Central	38		
South Atlantic	33		
New England	16		
West/South Central	8		
Mountain	7		
East/South Central	2		
West/North Central	0		

cording to the Centers for Disease Control,⁵ there have been 203 cases of typhoid fever reported in the United States from January 1, 1994, to July 23, 1994. The geographic areas from which cases of typhoid fever have been reported are summarized in Table 3. The states reporting the majority of the cases of typhoid fever include Massachusetts (N = 12), New York (N = 29), New Jersey (N = 14), Illinois (N = 18), Florida (N = 20), and California (N = 50). Moreover, typhoid fever is a serious infection when it does occur. The diagnosis of S. typhi is often not considered when a patient presents with fever. It is of paramount importance that the diagnosis in pregnancy be made with prompt institution of antibiotic therapy which is necessary for improved maternal and fetal prognosis.⁶ Appropriate cultures obtained prior to the initiation of antibiotic therapy are essential for confirming the diagnosis. Blood cultures are positive early in the course of infection, while stool cultures are positive later in the course of the infection. Cultures of multiple sites (urine, amniotic fluid, skin scrapings, bone marrow) improve the yield of a positive culture.

The recommended treatment regimens for S. *typhi* in pregnancy are summarized in Table 4. Chloramphenicol continues to be the treatment of choice; however, ampicillin or trimethoprim/

TABLE 4. Antimicro	obial agents	commonly	used	in
the treatment of S.	typhi in pre	gnancy ^{3,4,8}		

^aMay be given IV if patient cannot tolerate oral medication.

sulfamethoxazole may also be given. Chloramphenicol readily crosses the placenta but has not been shown to be teratogenic when utilized in early pregnancy.⁷ It is unlikely that "gray baby syndrome" (cyanosis, vascular collapse, and death), which has been reported when this antibiotic was given to the premature infant, would be caused by transplacental passage of this drug.⁸ The fluoroquinolones, i.e., ciprofloxacin and ofloxacin, have been shown to be efficacious in the treatment for S. typhi; however, they are currently not recommended during pregnancy. Although they have not been shown to be teratogenic in humans, there are reports of irreversible arthropathy in immature animals. Thus, this class of drugs should be reserved for serious life-threatening infection for which other antibiotics are ineffective or inappropriate.

The present case had the clinical features of unrelenting spiking high fevers, severe hemolysis (for which blood transfusions were given), and obvious illness. She did not have diarrhea which may occur late in the course of the infection. We concur with Dildy and colleagues⁹ that the clinician must also consider geographic risks as well as microbiologic and hematologic clues in order to establish the proper diagnosis.

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