Two cases of Kawasaki disease presented with acute febrile jaundice

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Kawasaki disease is an acute, systemic vasculitis of unknown etiology. Although gastrointestinal involvement does not belong to the classic diagnostic criteria; diarrhea, abdominal pain, hepatic dysfunction, hydrops of gallbladder, and acute febrile cholestatic jaundice are reported in patients with Kawasaki disease. We describe here two cases presented with fever, and acute jaundice as initial features of Kawasaki disease.

Key words: children, cholestasis, jaundice, Kawasaki disease.

Kawasaki disease (KD) is an acute, systemic vasculitis of unknown etiology that generally affects children younger than five years of age. The diagnosis of KD is based on the clinical criteria that include the most frequent and characteristic features of the disease. The presentation with atypical or incomplete KD findings, is a leading cause of delayed diagnosis. Gastrointestinal tract involvement is defined among the noncardiac findings of disease.¹ In this report we described two patients with an unusual onset of KD, characterized by fever, and acute jaundice.

Case Reports

Case 1

A six-year-old boy was admitted to hospital with complaints of fever for seven days, abdominal pain and jaundice. Physical examination revealed a fever of 38.5 °C, irritability, jaundice and abdominal tenderness. The medical history of the patient was unremarkable. Laboratory findings were as follows: hemoglobin concentration 9.1 g/dl; white blood cell count (WBC) 8700/mm³; platelet count 182.000/mm³; erythrocyte sedimentation rate (ESR), 55 mm/h (0-10 mm/h) and C-reactive protein (CRP), 160 mg/dl (0-8 mg/dl), serum alanin aminotransferase (ALT) 42 U/L (5-40 U/L), aspartate aminotransferase (AST) 29 U/L (8-33 U/L), total bilirubin 4.8 mg/dl (0-2 mg/dl), direct bilirubin 4 mg/dl (0-0.5 mg/dl), gamma glutamyl transferase (GGT) 168 U/L (<23 U/L), albumin 2.8 g/dl (3.1-4.8 g/dl). Urinalysis revealed the presence of pyuria. Serological tests for hepatitis A, B, C, Epstein-Barr virus (EBV) and cytomegalovirus were negative for acute infections. Abdominal ultrasonography showed that subhepatic and pelvic fluid, normal gallbladder, bile ducts and pancreas. Minimal bilateral pleural effusion was detected on chest ultrasonography. Ceftriaxone treatment was commenced for possible enteric fever or urinary tract infection. Cultures of blood, urine and throat had resulted as negative thereafter but fever of the patient persisted despite broad-spectrum antibiotic treatment. Echocardiography was performed in suspicion of atypical KD on the 9th day of the illness because of persisting clinical conditions including fever and jaundice. Bilateral diffuse dilatation of the coronary arteries (CAs) (3.3 mm left and 3.6 mm right) and minimal mitral valve insufficiency were detected on echocardiography. Patient was treated with single dose of intravenous immunoglobulin (IVIG, 2 g/kg/day) and acetylsalicylic acid (ASA, 80 mg/kg/day for the first 10 days). On the 11th day of illness desquamation of fingers and bilateral conjunctival erythema...
had occurred. His fever resolved after IVIG treatment within 24 hours and bilirubin and GGT levels returned to normal levels. Repeated abdominal and chest ultrasonography revealed no pelvic or pleural fluid. After 8th day of high dose ASA treatment control echocardiography demonstrated an increase of CAs diameters (left 5.6 mm and right 3.5 mm). Low-molecular-weight heparin (LMWH) was added to the treatment. He was discharged after 32 days of hospitalization with the antiaggregant dose of ASA (5 mg/kg/day) and LMWH. The dilated CAs gradually improved on repeated echocardiographic investigations. A one year outpatient follow-up revealed no remarkable additional morbidity or recurrence of KD.

Case 2
A previously healthy 2.5-year-old boy was admitted to hospital with seven days of fever, swelling and redness of upper surface of foot. He had 39 °C fever, tenderness and indurative edema of feet and icter on physical examination. The medical history of the patient was unremarkable. Laboratory findings were as follows; hemoglobin 10.1 g/dl, WBC 22,400/mm$^3$, platelet count 413,000/mm$^3$, ESR 145 mm/h, CRP 360 mg/dl, ALT 51 U/L, AST 44 U/L, total bilirubin 5.4 mg/dl, direct bilirubin 4.1 mg/dl, GGT 444 U/L. Urinalysis revealed the presence of pyuria. Cultures of blood, urine and throat were negative. Abdominal ultrasonography was normal. Serological tests for hepatitis A, B, C, EBV and cytomegalovirus were negative for acute infections. In light of the findings of fever for seven days and cholestasis without apparent causes, tenderness and indurative edema of feet, atypical KD was considered. The echocardiographic investigation that was performed on the second day of hospitalization demonstrated irregularity and dilatation of left CA ostia and mitral valve insufficiency. A single dose of IVIG (2 g/kg/day) and ASA (80 mg/kg/day, for the first 10 days) were administered. Thereafter, bilateral conjunctival erythema developed. After 24 hours of treatment, clinical and laboratory improvement was rapidly obtained with regression of fever and bilirubin levels. ASA was reduced to an antiaggregant dose after 10 days. The patient was discharged with sequela of mild left CA dilatation and minimal mitral valve insufficiency. A two month follow-up revealed no recurrence of fever or additional increase of CA diameters with antiaggregant dose of ASA.

Discussion
Kawasaki disease is characterized by fever for at least five days and four or more of the following five major clinical features; bilateral nonexudative conjunctival injection, erythema of the lips and oral mucosa, changes in the extremities, polymorphous skin rash, and unilateral painless cervical lymphadenopathy.$^{1,2}$ The patients who do not fulfill the major diagnostic criteria yet having echocardiographic findings have been defined as incomplete KD and who present with atypical conditions have been defined as atypical KD.$^1$ Although our first case had none of these classical criteria, febrile cholestasis without an apparent cause was given rise to the diagnosis of KD. The second case had one of the classical criteria additionally.

Although gastrointestinal involvement does not belong to the classic diagnostic criteria; diarrhea, various degrees of abdominal pain, hepatic dysfunction, hydrops of gallbladder up to 15% of patients and acute febrile cholestatic jaundice are reported in patients with KD.$^{1,3}$ The presentation with acute febrile cholestasis are among the atypical KD. Hydrops of the gallbladder as an acute acalculous, noninflammatory condition can occur in infants and children. On the other hand, acute acalculous cholecystitis is uncommon in children and is usually caused by infection especially streptococci, $Staphylococcus aureus$, $Salmonella$ spp, $Leptospira interrogans$, EBV, hepatitis A virus, $Ascaris$ or $Giardia lamblia$. $^{4,5}$ Both of our patients had biochemical findings of acute cholestasis without the evidence of an enlarged, thick-walled gallbladder, without calculi on ultrasonographic investigation. Hepatotropic viral infections and drugs were also excluded. It was postulated that a vasculitis of KD by the inflammatory changes of liver and gallbladder, and obstructive mechanism with enlarged lymph nodes gave cause to cholestasis.$^6$

In a study that reviewed acute febrile cholestatic jaundice, KD was found as the second most frequent (20%) cause among the 24 children. All of these patients had
presented with hepatomegaly, elevated acute phase reactants, aminotransferase levels and cholestatic markers. In an another study from Spain three of the 31 cases with KD had jaundice, higher levels of ALT, AST and GGT without abnormalities in liver, gallbladder and intrahepatic or extrahepatic bile ducts as shown by ultrasound. In a recent study from Turkey, 35 patients with KD were described. None of these patients had acute cholestasis as the initial feature of KD. It is well known that KD is a leading cause of acquired heart disease in children CA aneurysms occurring in 20% to 25% of children who do not receive IVIG treatment. In children who present with incomplete KD, the risk of CA sequelae is higher than classical KD. Records of patients with KD and CA aneurysms were reviewed at four United States centers from 1981 to 2006. Of 195 patients, 137 (70%) had had classical KD, and 53 (27%) had had incomplete KD. Authors investigated the performance of the 2004 American Heart Association (AHA) statement recommendations in this retrospective cohort including 53 eligible patients with incomplete KD for IVIG treatment. They concluded that application of the 2004 AHA recommendations, compared with the classic criteria alone, improves the rate of IVIG treatment for patients with KD who develop CA aneurysms. Standard treatment in the acute stage of KD includes combination of single high dose (2g/kg) IVIG and high-dose aspirin. The patients who respond to initial treatment of IVIG, fever resolves within 24 to 48 hours. However, approximately 10% of the patients have persistent or recurrent fever beyond 36 hours of therapy and require further treatment. In our two patients with incomplete KD and CA aneurysm had responded to initial IVIG treatment and further treatment was not required. Acute management of patients with CA aneurysm should be based on the extent and severity of the lesion. Although low dose ASA is sufficient for mild disease, because of the increased risk of thrombosis through CA, additional therapy such as LMWH, heparin or antiplatelet agent may be required for severe disease management. In conclusion: atypical presentations of KD like acute febrile cholestatic disease should be kept in mind for timely diagnosis and treatment of children with incomplete KD. Early IVIG administration for a febrile syndrome of unknown etiology in a child when jaundice is present, should be considered. The antiaggregant and anticoagulant therapies are useful in the long-term management of KD patients with CA dilatation.

REFERENCES