Parvovirus B19 was discovered in 1975 during serum screening of normal blood bank donors for hepatitis B antigen in lot B, sample 19, hence the name (1). In 1981 it was shown to have a causative role in transient aplastic crises of patients with chronic hemolytic anemia (2,3), and in 1985 was identified as the agent causing the childhood illness erythema infectiousum, so-called ‘fifth disease’ or ‘slapped cheek syndrome’ (4). Outbreaks of fifth disease primarily affect children in schools or day care centres. In reported outbreaks, 10% of the cases are in those under age five years, 70% in those from ages five to 15 years and 20% of cases are in those older than 15 years (5). Infections tend to be seasonal and are more common in late winter and spring (6). Approximately half of the adult population has been exposed to this agent and acquired immunity between the ages of five and 19 years, with seroprevalence increasing with age (7,8). Adult infections may vary considerably in severity, and the spectrum of illness can include ‘flu-like symptoms, generalized pruritus, arthropathy, joint effusions, anemia, myositis, encephalitis and idiopathic thrombocytopenic purpura (6).
A case of parvovirus B19 infection in an adult is presented. The patient had recent exposure to parvovirus B19 which resulted in acute renal failure, transient elevation of liver enzymes and congestive heart failure.

**CASE PRESENTATION**

A previously healthy 39-year-old female nurse developed a 'flu-like illness with fatigue, malaise, headache, arthralgias and a low grade fever. Her son had developed a similar illness along with a cutaneous rash approximately two weeks earlier and was diagnosed with erythema infectiosum ('fifth disease') by his pediatrician, based on clinical presentation and positive serology. With the exception of persistent fatigue, all other symptoms subsided within 10 days; however, over the next three days she developed generalized edema with 7 kg weight gain. She was evaluated by her family doctor, and laboratory studies revealed acute renal failure (serum creatinine of 251 µmol/L [normal range 60 to 110 µmol/L]), normocytic, normochromic anemia (hemoglobin 93 g/L [115 to 155 g/L]) with reticulocyte count 2.0% (0.9% to 2.6%) and elevated liver enzymes (aspartate aminotransferase 58 U/L [8 to 29 U/L], alanine aminotransferase 80 U/L [1 to 20 U/L], lactate dehydrogenase 311 U/L [108 to 211 U/L]), but normal direct (2 µmol/L [0 to 4 µmol/L]) and total (14 µmol/L [0 to 16 µmol/L]) bilirubin. Urinalysis showed no hematuria and trace proteinuria on dipstick, and the urine sediment was inactive with no red or white blood cells, crystals or casts on microscopy. Renal function had improved two days later when the patient was seen in the nephrologist's office (serum creatinine down to 191 µmol/L); however, by follow-up five days later she had developed exertional dyspnea, orthopnea and paroxysmal nocturnal dyspnea. Examination (20 days after the onset of symptoms) revealed that jugular venous pressure was elevated at 5 cm above the sternal angle, and she had a prominent apical impulse, a third heart sound and bibasilar crepitations findings suggestive of congestive heart failure. Chest x-ray demonstrated an enlarged heart, bilateral blunting at the costophrenic angles, increased interstitial markings, Kerley B lines and perihilar edema, confirming the clinical diagnosis of congestive heart failure (Figure 1). Electrocardiogram demonstrated sinus tachycardia at 110 beats/min with no ST-T changes. Creatine kinase (CK) was slightly elevated at 157 U/L (15 to 20 U/L) with a CK-MB fraction of 1 U/L (relative index of less than 1%). Antinuclear antibody, antiDNA, extractable nuclear antigen and anticardiolipin antibody were all negative. Complement studies showed a normal C4 and marginally reduced C3 at 844 mg/L (870 to 1720 mg/L). Serological studies for syphilis, hepatitis B and C, human immunodeficiency virus, Epstein-Barr virus and *Borrelia burgdorferi* were all negative. An echocardiogram revealed a mild decrease in left ventricular fractional shortening (53%), and wall motion study confirmed a similar abnormality with slight global reduction of her ejection fraction (54%). Ventilation-perfusion scan and Doppler ultrasound examination of her leg veins were negative. Parvovirus B19 serology at this time (day 22) in hospital showed positive titres for both immunoglobulin M.

Figure 1) Chest x-ray demonstrating cardiomegaly, pulmonary vascular redistribution, hilar and pulmonary edema and blunting of costophrenic angles, suggestive of congestive heart failure

Figure 2) Follow-up study nine months later with resolution of patient's symptoms
and G. A diagnosis of parvovirus B19-induced acute renal failure, elevated liver enzymes and congestive heart failure was made.

The patient was treated symptomatically with oxygen and intravenous furosemide. Over the following week her hemoglobin level gradually rose, and her creatinine and liver enzymes level fell towards the normal liver function range. A repeat echocardiogram demonstrated an increase in the left ventricular fractional shortening to 40%. She was discharged home on short term oral furosemide therapy, and her recovery was followed as an out-patient. Six months later she is well without any symptoms. Her creatinine is 82 µmol/L, hemoglobin is 127 g/L, and liver enzymes are completely normal. Her follow up chest x-ray was also completely normal (Figure 2).

**DISCUSSION**

Infection with parvovirus B19 has protean manifestations. In children, the symptoms are similar to many other viral illnesses with fever, nausea, diarrhea, malaise and cough. The bright red facial rash, the so called 'slapped cheek' phenomenon, is a characteristic but not pathognomonic finding.

Adults may develop more severe sequelae from parvovirus B19 infection. Polynarthropathy involving the hands, wrists, knees, ankles and feet is the most common finding and may be painful enough to impair dexterity and mobility (9). Anemia, caused by the lysis of red blood cell precursors, is usually mild because the duration of the illness is typically much shorter than the average erythrocyte lifespan (6). In patients with chronic hemolytic anemias, the anemia may become much more severe, and they may develop life threatening marrow aplasia.

We describe three new, previously unreported clinical manifestations associated with acute parvovirus B19 infection in adults. The acute renal failure in our patient was suspected to be secondary to a postinfectious tubulointerstitial process. This was suggested by a bland urinary sediment without any symptoms. Her creatinine is 82 µmol/L, hemoglobin is 127 g/L, and liver enzymes are completely normal. Her follow up chest x-ray was also completely normal (Figure 2).

**REFERENCES**
