



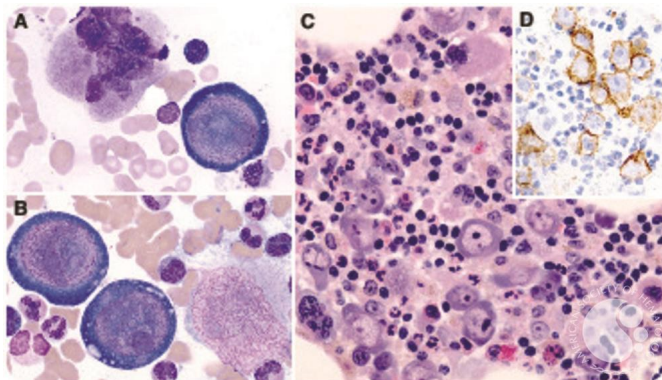
Red cheeks to red cell aplasia: parvovirus B19 in a heart transplant patient

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Category: Infectious Disease > Viruses > Parvovirus B19

Description: A 66-year-old man developed dizziness and fatigue 2 months after orthotopic heart transplant for ischemic cardiomyopathy. His immunosuppressive regimen included tacrolimus, mycophenolate mofetil, and prednisone. He was found to be severely anemic (hemoglobin, 6.6 g/dL). Clinical studies excluded acute blood loss or hemolysis. Reticulocyte count was 0.2%, and erythropoietin level was 750 mIU/mL (range, 3-19 mIU/mL). A bone marrow aspirate smear revealed large atypical cells with prominent intranuclear inclusions and vacuolated cytoplasm (panels A and B; original magnification $\times 100$, Wright-Giemsa stain), which were also evident on a hematoxylin and eosin stain of the aspirate clot (panel C; original magnification $\times 40$). Megakaryocytes and myeloid precursors were easily identified; however, there was essentially no erythroid maturation. E-cadherin stain confirmed that atypical cells were virally transformed erythroid precursors (panel D; original magnification $\times 40$). Serum parvovirus B19 polymerase chain reaction was positive. Anti-parvoviral immunoglobulin M (IgM) and IgG titers were within normal limits. Giant virally transformed proerythroblasts and erythroid maturation arrest are classic findings of parvovirus B19–associated pure red cell aplasia. Although most adults acquire lifelong immunity to parvovirus B19 from exposure in childhood, reinfection can occur in the setting of immunocompromise, leading to prolonged severe anemia. In retrospect, our patient had had contact with a granddaughter with high fevers and red cheeks several weeks before his initial presentation. His hemoglobin recovered after supportive transfusions and 5 days of IV immunoglobulin. He remains well 2 years posttransplant.





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