HHS Public Access

Author manuscript

Am J Transplant. Author manuscript; available in PMC 2020 February 10.

Published in final edited form as:

Am J Transplant. 2019 October; 19(10): 2955–2957. doi:10.1111/ajt.15538.

New skin lesions in a liver transplant recipient

Michela Blain¹, Kara Walter², Lena Sibulesky³, Rachel Bender Ignacio^{1,4}

¹Division of Allergy and Infectious Diseases, Department of Medicine, University of Washington Medical Center, Seattle, Washington

²Division of Gastroenterology, Department of Medicine, University of Washington Medical Center, Seattle, Washington

³Division of Transplant Surgery, Department of Surgery, University of Washington Medical Center, Seattle, Washington

⁴Vaccine and Infectious Diseases Division, Fred Hutchinson Cancer Research Center, Seattle, Washington

Keywords

biopsy; cirrhosis; clinical research/practice; complication: infectious; immunosuppressant - mechanistic target of rapamycin (mTOR); infection and infectious agents - viral: human herpesvirus 8 (HHV-8); infectious disease; liver transplantation/hepatology

A 53-year-old woman from Ukraine underwent orthotopic liver transplant (OLT) for cryptogenic cirrhosis. She received induction immunosuppression with antithymocyte globulin and methylprednisolone, transitioning to tacrolimus monotherapy (troughs 3–12 ng/mL). Pathology of her explanted liver, including stains for infectious and autoimmune processes, revealed no etiology for her cirrhosis. A culture swab from the organ transport media grew *Candida glabrata*, for which she received 4 weeks of prophylactic fluconazole. During the first 45 postoperative days, her course was complicated by biliary stricture, asymptomatic cytomegalovirus viremia for which she received valganciclovir, and *Clostridioides difficile* colitis.

Two months posttransplant, she reported progressing skin lesions on her lower extremities. She first noted lesions 2 days posttransplant, which she had interpreted as bruises. No lesions were noted pretransplant. During the intervening weeks, the lesions progressed proximally, grew in size, and became palpable and tender. A review of systems was otherwise unremarkable. On examination, she had scattered hyperpigmented, tender papules with associated dependent edema and faint erythema of bilateral lower extremities; her examination was otherwise unremarkable. The patient's exposure history was only notable for wading up to her knees into a freshwater lake 1 month post-OLT.

At time of presentation, her anti-infective regimen included valganciclovir, trimethoprim–sulfamethoxazole, and fluconazole. Based on the appearance of the skin lesions, she

Blain et al. Page 2

underwent an urgent skin biopsy, was admitted to the hospital, and was started on intravenous (IV) cefepime and liposomal amphotericin B. Initial fungal, acid-fast, and gram stains from the skin biopsy were negative. An immunohistochemical stain confirmed the diagnosis.

QUESTIONS

- 1. Which of the following diagnoses will the lower extremity skin biopsy likely confirm?
 - a. Disseminated fungal infection
 - b. Kaposi sarcoma
 - c. Bacillary angiomatosis
 - d. Ecthyma gangrenosum
 - e. Cryoglobulinemic vasculitis
- 2. Which infectious agent is associated with the diagnosis?
 - a. Bartonella henselae
 - **b.** Candida glabrata
 - c. Pseudomonas aeruginosa
 - **d.** Human herpesvirus 8
 - e. Hepatitis C virus
- **3.** What is the most likely origin of the pathogen?
 - **a.** Reactivation of latent infection in the recipient
 - **b.** Zoonosis from a domestic animal
 - c. Primary infection acquired exogenously after transplant
 - **d.** Transmission of a donor-derived pathogen
 - e. Contamination of the donor organ transport medium
- **4.** What treatment would you recommend at this time?
 - **a.** Change amphotericin to posaconazole
 - **b.** Decrease immunosuppression
 - c. Initiate IV liposomal doxorubicin
 - d. Begin IV ganciclovir
 - **e.** Initiate oral doxycycline plus rifampin

Blain et al. Page 3



FIGURE 1.The patient demonstrated edema and moderately tender macules and papules on her bilateral lower extremities. The lesions of her left lower extremity are shown