An Unusual Cause of Chest Pain: Mycobacterium Avium Complex and the Immune Reconstitution Inflammatory Syndrome

Richard J. Lin, MD, PhD1
Jie Song, MD2

1 Department of Medicine, Weill Cornell Medical Center, New York, New York.
2 Department of Pathology, University of Chicago Medical Center, Chicago, Illinois.

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The HIV-associated immune reconstitution inflammatory syndrome usually manifests as new infections or worsening of pre-existing infections during the first few months of initiating anti-retroviral therapy. It is commonly associated with local or systemic inflammation, presumably due to rapid reconstitution of host immune system. Here we describe a unique case of the immune reconstitution inflammatory syndrome presenting as acute pericarditis and pericardial effusion caused by mycobacterium avium complex. We also demonstrate that judicious use of steroids, along with pathogen specific antimicrobial therapy, can prevent local complications of the inflammatory response. Journal of Hospital Medicine 2011;6:309–311. © 2011 Society of Hospital Medicine.

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Although antiretroviral therapy for human immunodeficiency virus (HIV)-infected patients reduces viral load dramatically and improves immune function, some patients experience a clinical deterioration within the first few months of therapy because of an exuberant and dysregulated immune response— the immune reconstitution inflammatory syndrome (IRIS). The exaggerated immune response associated with this syndrome can be stimulated by either antigens from infectious agents (typically a mycobacterium or cryptococcus) or from autoantigens, giving rise to a heterogeneous range of clinical manifestations.1 IRIS may present as an inflammatory reaction that unmasks a previously untreated infection or as a paradoxical worsening of an infection that is being treated appropriately. Although most cases of IRIS are mild and self-limited, some patients require aggressive treatment.3

Case Report

A 53-year-old man was evaluated for a 5-day history of intermittent chest pain. He had been diagnosed with HIV/acquired immune deficiency syndrome (AIDS) 11 years ago but he had not been compliant with therapy. Seven years earlier he had been treated for 9 months with isoniazid for a positive tuberculosis skin test. Three months before admission, he developed methicillin-resistant Staphylococcus aureus skin abscesses and was found to have a CD4 count of 1/μL and a HIV viral load of over 400,000 copies/mL. He finished a course of vancomycin, and was started on lopinavir, ritonavir, abacavir, lamivudine, and zidovudine. Five days before admission, he was evaluated in the emergency department for intermittent chest pain and described using cocaine. There was only J-point elevation on the electrocardiogram (Figure 1A), serial cardiac enzymes were negative, and he was discharged home. However, despite discontinuation of cocaine use, his chest pain worsened, became pleuritic, and was associated with dyspnea, which prompted this admission. Physical examination was remarkable only for tachycardia, although the electrocardiogram now revealed diffuse ST segment (ST) elevation and PR segment (PR) depression, consistent with acute pericarditis (Figure 1B).

Serial cardiac enzymes, viral studies, and bacterial, fungal, and mycobacterial blood cultures were negative. His CD4 count was 16/μL, and the HIV viral load was 870 copies/mL.

The patient was treated with high-dose ibuprofen and colchicine, but mild chest pain and electrocardiogram changes persisted, and he developed a friction rub. A chest computed tomography (CT) scan was negative for pulmonary embolism and revealed no significant infra-thoracic pathology, except for a moderate pericardial effusion that was confirmed by transthoracic echocardiogram (Figure 2A). There was no echocardiographic evidence of tamponade. He underwent thoroscopic pericardial and mediastinal lymph node biopsy, along with drainage of the pericardial effusion. Pericardial biopsy showed acute on chronic inflammation consistent with pericarditis (Figure 2B) and culture was positive for Mycobacterium Avium Complex (MAC). He was treated with clarithromycin, ethambutol, and prednisone, and his antiretroviral medications were continued. At 2, 6, and 12 months follow-up, he was asymptomatic, the electrocardiogram had normalized (Figure 1C), and the echocardiogram showed no effusion or evidence of pericardial constriction.

Discussion

This case demonstrates a unique manifestation of the IRIS associated with MAC infection, which more typically presents as peripheral, pulmonary, or intra-abdominal lymphadenopathy.2,3 It usually responds to MAC therapy, although intra-abdominal disease portends a poor prognosis.3,4 This patient has two significant risk factors for the development of IRIS: low CD4 count at the time of antiretroviral therapy and rapid viral clearance.5,6 While his CD4 count...
FIGURE 1. A: 12-lead EKG at initial ER visit showing J-point elevations and early depolarization changes. B: 12-lead EKG at diagnosis showing changes consistent with acute pericarditis. C: 12-lead EKG at 2-month follow-up showing resolution of changes of acute pericarditis. EKG, electrocardiogram. [Color figure can be viewed in the online issue, which is available at wileyonlinelibrary.com.]

FIGURE 2. A: Echocardiographic evidence of a moderate pericardial effusion. B: Hematoxylin-eosin (H&E) stain of the pericardial biopsy showing acute on chronic inflammation (×200). [Color figure can be viewed in the online issue, which is available at wileyonlinelibrary.com.]
response is lower than expected for IRIS, previous studies have shown that functional immune recovery usually precedes quantitative CD4 count recovery, and that IRIS could happen at low CD4 count.\(^1,7\) Finally, we believe that the use of corticosteroids accounted for his rapid clinical improvement and favorable long-term outcome, consistent with previous experience of corticosteroid use in MAC-associated IRIS.\(^3,4\) To our knowledge, this is the first reported case of MAC-associated IRIS presenting as isolated acute pericarditis and pericardial effusion. In conclusion, our case illustrates that IRIS can present as an abnormal immune response to an opportunistic infection in an unusual location. Clinicians must be aware that after starting antiretroviral therapy, new symptoms, including chest pain, might represent 1 of the IRISs, and that corticosteroids might be beneficial when inflammation is severe.

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Address for correspondence and reprint requests:
Richard J. Lin, MD, PhD, Division of Hospital Medicine, Department of Medicine, Weill Cornell Medical College and New York-Presbyterian Hospital, 525 East 68th Street, Box 130, New York, NY, 10065; Telephone: 212-746-9832; Fax: 212-746-4734; E-mail: ril9016@med.cornell.edu Received 5 October 2008; revision received 26 January 2010; accepted 26 January 2010.

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