Aortitis and Atrioventricular Block following COVID-19 Infection

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Abstract

Introduction: Several types of cardiovascular (CV) complications have been reported with COVID-19. In this report, we describe the first known case of concomitant aortitis and atrioventricular block (AVB) following COVID-19 infection.

Case presentation: A 40-year-old previously healthy African male presented with pericarditis, six weeks after a minimally symptomatic COVID-19 infection. He also developed transient AVB. He was started on high dose steroids and did not suffer any complications from his aortitis. The patient did not require any pacemaker insertion.

Conclusions: AVB and aortitis are two uncommon and distinct complications of COVID-19 infection. As these complications can occur in the subacute phase of a minimally symptomatic infection, it is of paramount importance that physicians remain vigilant beyond the acute infection, even in initially mild cases.

Keywords: Aortitis; Atrioventricular-block; AVB; COVID; B.1.1.7; UK-variant

Abbreviations: ACE2: Angiotensin Converting Enzyme2; AVB: Atrioventricular Block; CRP: C-Reactive Protein; CT: Computed Tomography; CV: Cardiovascular; FDG: Fluorodeoxyglucose; HR: Heart Rate; PET: Positron Emission Tomography

Background

Significant morbidity and mortality can arise from COVID-19 infection. COVID-19 infection has been associated with several types of cardiovascular (CV) complications including myocardial injury, myocarditis, acute myocardial infarction, heart failure and arrhythmia [1], which often manifest weeks following initial infection [2]. COVID-19 has also been associated rarely with aortitis and atrioventricular block (AVB) separately [3, 4]. To the best of our knowledge, we are the first to report a case of concurrent aortitis and AVB following a COVID-19 infection. In this report, we described the management and outcome of concurrent aortitis and AVB.

Case Presentation

Mr. C is a 40-year-old previously healthy African male who worked as a construction site supervisor. The patient acquired COVID-19 in March 2021 and had only minimal fatigue in the initial weeks of his infection.

Four weeks after his initial infection, Mr. C presented with pleuritic chest pain with diffuse ST-segment elevations on his electrocardiogram. Additionally, he also developed high grade AVB with a heart rate (HR) of 45 per minute.

Investigations

Laboratory investigation revealed an elevated C-reactive protein (CRP) at 98 milligrams/L. Tuberculosis Quantiferon, cultures for histoplasma, Blastomyces, leishmania, HIV and Lyme serologies, four blood cultures, autoimmune screen and vasculitis work-up were all negative. High sensitivity troponin and thyroid function tests were normal. Transthoracic echocardiogram and a cardiac computed tomography (CT) were normal. A cardiac magnetic resonance imaging (MRI) showed acute left ventricular myocardial edema without fibrosis and a small circumferential pericardial effusion. Chest computed tomography was negative for active sarcoidosis. Fluorodeoxyglucose (FDG)-positron emission tomography (PET) scan (done for sarcoidosis work-up) showed a moderate degree of metabolic activity in the ascending aorta, suspicious for a vasculitis (Appendix 2). An initial syphilis enzyme immunoassay was positive, yet a rapid plasma regain test was negative. A confirmatory test was positive for endemic treponema syphilis.

Treatment

Due to the spontaneous improvement of HR to 70/min within a few days, the patient did not require a pacemaker. The patient was started on ibuprofen and colchicine for pericarditis. He was initiated on a tapering course of prednisone 40mg orally daily over four weeks for the aortitis. He also received intramuscular penicillin injections to treat the suspected endemic treponema.
Outcome A and Follow-Up
At seven weeks after his initial COVID-19 infection, the patient still had mild intermittent chest pain without any other active CV, neurological or rheumatologic symptoms.

Discussion
To the best of our knowledge, this is the first reported case of aortitis complicated by AVB following a COVID-19 infection. The AVB was likely due to extension of the inflammation of the aorta to the AV node (adjacent to the aortic root).

Wang et al. reported a prevalence of 23% of cardiac arrhythmia in the initial Wuhan cohort of COVID-19 infected individuals [5]. The most frequent arrhythmia were atrial fibrillation, atrial flutter and ventricular tachycardia [6]. AVB were rare following COVID-19 infection [7-9]. These conduction abnormalities were thought to be due to acute inflammation of the intrinsic conduction system [7] and/or from direct viral injury to the myocardium [10]. AVB was generally transient (less than 1 week) and did not require a permanent pacemaker [6,7,11]. Temporary pacemakers were required in rare instances [12].

Kawasaki’s disease and cutaneous vasculitis following COVID-19 infection have been observed in children [13,14]. In adults, the vasculitis phenomena manifested mostly as thromboembolism, cutaneous vasculitis, and myopericarditis [15]. Four cases of inflammatory aortitis due to COVID-19 have been reported [3, 16-18]. The spectrum of COVID-19 related aortitis symptoms varied widely from asymptomatic to fatigue, weight loss, chest, back and abdominal pain, claudication, neurological deficits, life-threatening aortic aneurysm and aortic dissection [3]. Acute COVID-19 aortitis is suspected to be due to an infiltration of virions into the aortic endothelium mediated by Angiotensin-Converting Enzyme 2 (ACE2) receptors. The cellular viral invasion leads to acute endothelitis and
leukocytoclastic vasculitis. This inflammatory cascade leads to deposition of immune complexes with a type-3 hypersensitivity reaction [17, 19-21]. Prior COVID-19 related aortitis have been successfully treated with two-to-four-week courses of 40-60mg of prednisone [3,16].

FDG PET is recommended for the diagnosis of suspected aortitis, regardless of etiology. There are no current guidelines regarding PET use in the management of COVID-19 infection. Repeating the FDG PET three to six months following the infection may be helpful to evaluate the response to therapy. Additionally, FDG PET can also confirm relapse of large vessel vasculitis [22-24].

Although our patient tested positive for syphilis, it was unlikely that his aortitis was due to active venereal syphilis. He was from west Africa where the prevalence of endemic treponema was 33% [25]. It was most likely that he had endemic treponema infection, known as Bejel infection, which is not associated with any CV complication [25]. Furthermore, concomitant AVB and aortitis is exceedingly rare in venereal syphilis. Veneralsyphilis is associated with gummata (granulomatous) infiltrating the conduction system [26]. The gummata would not have resolved spontaneously, as in our patient. Furthermore, this patient had no other finding suggestive of tertiary or CV syphilis.

The development of subacute CV complications following the initial COVID-19 infection emphasizes the need for health care providers to remain vigilant beyond the first few weeks of the infection. The median onset of CV symptoms following COVID-19 infection is 47 days [2]. Of note, our patient had only minimal symptoms with the initial COVID-19 infection. Therefore, mild COVID-19 infection does not preclude late life-threatening complications.

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