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Patients With Acute Myocarditis Following mRNA COVID-19 Vaccination

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IMPORTANCE Vaccine-associated myocarditis is an unusual entity that has been described for the smallpox vaccine, but only anecdotal case reports have been described for other vaccines. Whether COVID-19 vaccination may be linked to the occurrence of myocarditis is unknown.

OBJECTIVE To describe a group of 7 patients with acute myocarditis over 3 months, 4 of whom had recent messenger RNA (mRNA) COVID-19 vaccination.

DESIGN, SETTING, AND PARTICIPANTS All patients referred for cardiovascular magnetic resonance imaging at Duke University Medical Center were asked to participate in a prospective outcomes registry. Two searches of the registry database were performed: first, to identify patients with acute myocarditis for the 3-month period between February 1 and April 30 for 2017 through 2021, and second, to identify all patients with possible vaccine-associated myocarditis for the past 20 years. Once patients with possible vaccine-associated myocarditis were identified, data available in the registry were supplemented by additional data collection from the electronic health record and a telephone interview.

EXPOSURES mRNA COVID-19 vaccine.

MAIN OUTCOMES AND MEASURES Occurrence of acute myocarditis by cardiovascular magnetic resonance imaging.

RESULTS In the 3-month period between February 1 and April 30, 2021, 7 patients with acute myocarditis were identified, of which 4 occurred within 5 days of COVID-19 vaccination. Three were younger male individuals (age, 23-36 years) and 1 was a 70-year-old female individual. All 4 had received the second dose of an mRNA vaccine (2 received mRNA-1273 [Moderna], and 2 received BNT162b2 [Pfizer]). All presented with severe chest pain, had biomarker evidence of myocardial injury, and were hospitalized. Coincident testing for COVID-19 and respiratory viruses provided no alternative explanation. Cardiac magnetic resonance imaging findings were typical for myocarditis, including regional dysfunction, late gadolinium enhancement, and elevated native T1 and T2.

CONCLUSIONS AND RELEVANCE In this study, magnetic resonance imaging findings were found to be consistent with acute myocarditis in 7 patients; 4 of whom had preceding COVID-19 vaccination. Further investigation is needed to determine associations of COVID-19 vaccination and myocarditis.

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JAMA Cardiol. 2021;6(10):1196-1201. doi:10.1001/jamacardio.2021.2828 Published online June 29, 2021. accine-associated myocarditis is an unusual entity that has been described for the smallpox vaccine,¹ but otherwise only anecdotal case reports have been described for other vaccines. Among 416 629 adults receiving live measles, mumps, and rubella; varicella; oral polio; or yellow fever viral vaccinations in the Vaccine Safety Datalink, there were no patients with myocarditis in the 42 days following vaccination.² There are a few case reports of myocarditis following the seasonal influenza vaccine in otherwise healthy adults,^{3,4} but a causal relationship is difficult to establish and the case reports could have been due to chance.

In late December 2020, COVID-19 vaccination began in the US, and on April 7, 2021, vaccination was opened to all adults 16 years or older in North Carolina. Here, we report cardiac magnetic resonance (CMR) imaging findings in 4 patients consistent with acute myocarditis at our institution; all 4 had recent vaccination for COVID-19. All presented with severe chest pain associated with biomarker evidence of myocardial injury and were hospitalized. Included are data on clinical presentation and results from in-hospital testing. To provide context, we also report the prevalence of acute myocarditis by CMR imaging at our institution during the same 3-month period for each of the past 5 years.

Methods

All patients referred for CMR imaging at Duke University Medical Center are asked to participate in a prospective CMR imaging outcomes registry, which contains clinical data, finalized clinical CMR imaging reports, and full Digital Imaging and Communications in Medicine image data sets.⁵ The registry has been approved by the institutional review board at Duke University Medical Center, and all patients sign informed consent prior to participating.

We performed 2 searches of the registry. First, to identify patients with acute myocarditis for February, March, and April of 2017 through 2021, a search for the term *myocarditis* was performed within the report summary and limited to the relevant time period. Second, to identify all patients with possible vaccine-associated myocarditis for the past 20 years, we searched for the terms *vaccine* or *vaccination* in the report

Key Points

Question Is COVID-19 vaccination linked to the occurrence of myocarditis?

Findings In this study of 7 patients with acute myocarditis, 4 occurred within 5 days of COVID-19 vaccination between February 1 and April 30, 2021. All 4 patients had received the second dose of a messenger RNA (mRNA) vaccine, presented with severe chest pain, had biomarker evidence of myocardial injury, were hospitalized, and had cardiac magnetic resonance imaging findings typical of myocarditis.

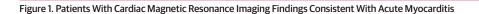
Meaning Although causality cannot be established, the findings raise the possibility of an association between mRNA COVID-19 vaccination and acute myocarditis.

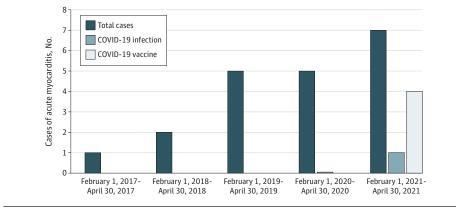
summary and the history fields without limitation in the time period. A second independent reviewer performed the same searches and confirmed the counts. Once patients with possible vaccine-associated myocarditis were identified, data available in the registry were supplemented with information from the electronic health record and telephone interview.

CMR imaging was performed on 1.5-T or 3-T scanners. A standard protocol for myocarditis was used, which included cine, T1 and T2 mapping, and late gadolinium enhancement (LGE).^{6,7} Native T1 and T2 times were measured on pixelwise maps with regions drawn to match areas with and without LGE. Cutoffs for abnormally elevated T1 and T2 were based on 2 SDs above the respective means in a healthy population imaged on the same scanners.

Results

In the 3-month period between February 1 and April 30, 2021, we identified 7 patients with acute myocarditis of whom 4 had recent COVID-19 vaccination (**Figure 1**). A search of the entire registry identified a single additional patient with myocarditis diagnosed by CMR imaging within days following quadrivalent influenza vaccination. The patient was a member of the military, and this was his second episode; he had documented myocarditis associated with smallpox vaccination 4 years earlier.





The total number of patients with acute myocarditis is highest for 2021 with 4 associated with recent COVID-19 vaccination. Only 1 patient had myocarditis associated with acute COVID-19 infection.

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Characteristic	Patient 1	Patient 2	Patient 3	Patient 4
Age, y	36	23	70	24
Sex	Male	Male	Female	Male
Comorbidities				
Hypertension	No	No	Yes	No
Diabetes	No	No	No	No
Hypercholesterolemia	No	No	Yes	No
Cigarette smoking	No	No	Yes	No
History of CAD or MI	No	No	No	No
Prior myocarditis	No	No	No	No
Prior COVID-19 infection	No	No	No	No
Vaccine received	mRNA-1273	BNT162b2	mRNA-1273	BNT162b2
Doses received	2	2	2	2
	Z	2	2	2
Symptoms prior to vaccination (last dose)		N	N	
Viral prodrome ^a	No	No	No	No
Symptoms within 24 h of vaccination				
Injection site discomfort	Yes	Yes	No	Yes
Fatigue/muscle ache	Yes	Yes	No	Yes
Fever/chills	Yes	Yes	No	Yes
Headache	No	No	No	Yes
Nausea	No	No	No	No
Chest pain	No	No	Yes	No
Hospitalization				
Interval after vaccination, d	3	5	1	2
Symptoms leading to hospitalization				
Chest pain	Yes	Yes	Yes	Yes
Chest pain severity	Severe	Severe	Severe	Severe
Symptom onset	<12 h of Hospitalization	<12 h of Hospitalization	<12 h of Hospitalization	<24 h of Hospitalization
Shortness of breath	Yes	Yes	Yes	No
Diaphoresis	No	No	Yes	No
Syncope/presyncope	No	Yes	No	No
Palpitations	No	No	No	Yes
In-hospital testing	110	110	110	105
in nospital testing	Diffuse ST elevation	Lateral	Anterolateral	Diffuse ST elevation
Electrocardiogram	PR depression	ST elevation	ST elevation	PR depression
Deale transmin no //	•			•
Peak troponin, ng/L	hs-Tn T: 230 (abnormal)	hs-Tn I: 7452 (abnormal)	Tn I: 2.34 (abnormal)	hs-Tn T: 698 (abnormal
CRP, mg/dL	6.32 (Abnormal)	2.2 (Abnormal)	NP	6.08 (Abnormal)
ESR, mm/h	6	30 (Abnormal)	NP	12
ProBNP, pg/mL	NP	780 (Abnormal)	5194 (Abnormal)	65
WBC, /µL	10 200	10 800	16 700 (Abnormal)	13 500 (Abnormal)
COVID-19 testing ^b	Negative ^c	Negative ^d	NP	Negative ^c
Respiratory virus PCR ^e	Negative	Negative	NP	NP
Chest radiography	Normal	Normal	Normal	Normal
Chest CT	Negative for PE	Negative for PE	NP	NP
Coronary angiography	NP	NP	Normal coronaries	NP
Cardiac MRI				
Interval after vaccination, d	3	5	3	3
LVEF, %	53	58	40	59
Regional wall motion abnormality	Yes	Yes	Yes	Yes
Pericardial effusion	Trace	Small	Small	Trace
Pericardial thickness	Normal	Normal	Normal	Normal
LGE present	Yes	Yes	Yes	Yes
Location	Apical lateral	Multiple	Multiple	Lateral
Myocardial pattern	Epicardial	Epicardial	Patchy, diffuse	Epicardial, patchy
wiyocai ulat pattern	Epicalulat	Epicaruidi	ratery, diffuse	Epicalulat, pattily

(continued)

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Table. Characteristics of Patients Who Received COVID-19 Vaccination (continued)

Characteristic	Patient 1	Patient 2	Patient 3	Patient 4
Native T1				
In region of LGE	Abnormal	Abnormal	Abnormal	Abnormal
In region without LGE	Normal	Normal	Normal	Normal
Native T2				
In region of LGE	Not imaged	Abnormal	Abnormal	Abnormal
In region without LGE	Normal	Normal	Normal	Normal
herapy				
Corticosteroids	No	Yes	No	No
Colchicine	Yes	Yes	No	Yes
NSAIDs	Yes	No	No	Yes

Abbreviations: CAD, coronary artery disease; CRP, C-reactive protein; CT, computed tomography; ESR, erythrocyte sedimentation rate; hs-Tn, high-sensitivity troponin; LGE, late gadolinium enhancement; LVEF, left ventricular ejection fraction; MI, myocardial infarction; MRI, magnetic resonance imaging; NP, not performed; NSAIDs, nonsteroidal anti-inflammatory drugs; PCR, polymerase chain reaction; PE, pulmonary emboli; proBNP, pro-brain-type natriuretic peptide; WBC, white blood cell count.

SI conversion factors: To convert CRP to milligrams per liter, multiply by 10; ESR to millimeters per hour, multiply by 1; troponin to micrograms per liter, multiply by 1; WBC to $\times 10^9$ per liter, multiply by 0.001.

The clinical characteristics of the 4 patients with myocarditis following COVID-19 vaccination are shown in the **Table**. Data on race and ethnicity were not available. Three were younger male individuals (age, 23-36 years) and 1 was a 70year-old female individual. All 4 had received the second dose of a messenger RNA (mRNA) vaccine (2 received mRNA-1273 [Moderna], and 2 received BNT162b2 [Pfizer-BioNTech]) between 1 and 5 days before hospitalization. None had a viral prodrome or prior COVID-19 infection. COVID-19 and respiratory virus polymerase chain reaction test results were negative in 3 and 2 patients, respectively, who underwent testing. None had acute pulmonary disease on chest radiograph.

All 4 patients had abnormal electrocardiogram results and elevated troponin levels. The older female individual underwent coronary angiography, which revealed no atherosclerosis. CMR imaging was performed between 3 and 5 days after vaccination. All had regional wall motion abnormalities on cine imaging, and ejection fraction ranged between 40% and 59%. LGE was present in a nonischemic pattern consistent with myocarditis in all 4 patients (**Figure 2**). Both native T1 and T2 were elevated in the regions with LGE, consistent with acute injury, except in 1 patient who did not have a T2 map acquired at a slice location that included an area with LGE. Pericardial thickness was normal in all, and no patients had pericardial LGE.

The hospital courses for all 4 were uneventful without evidence of arrhythmias or heart failure, and treatment was conservative with nonsteroidal anti-inflammatory drug and colchicine, with 1 receiving corticosteroids. All were discharged within 2 to 4 days of hospitalization.

Discussion

We identified at our institution 4 patients with acute myocarditis occurring within days of mRNA COVID-19

^a Upper respiratory tract symptoms, fever, muscle aches.

^b Performed within 2 days after the onset of chest pain in those who underwent testing.

 $^{\rm c}$ Isothermal COVID-19 test was performed (Abbott ID Now; lower detection limit of 125 genome equivalent/mL).

^d Real-time PCR COVID-19 test was performed (Cepheid Xpert Xpress SARS-CoV-2; lower detection limit of 131 copies/mL).

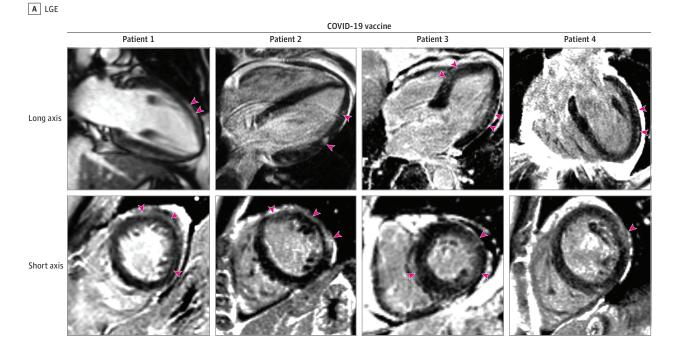
^e Influenza A, influenza B, respiratory syncytial virus.

vaccination. Although a causal relationship cannot be established, we note that none had a viral prodrome or had coincident testing (including COVID-19 polymerase chain reaction and respiratory virus polymerase chain reaction) that revealed an alternative explanation. None had COVID-19 infection in the prior year, suggesting that myocarditis from subsequent multisystem inflammatory syndrome is also unlikely.⁸ Additionally, the diagnosis of acute myocarditis was straightforward. The presenting symptom was acute onset of severe chest pain, and myocardial injury was detected by elevated troponin levels in all patients. CMR imaging abnormalities were observed across multiple techniques (regions with wall motion abnormalities on cine imaging-matched regions with LGE and regions with abnormal native T1 and T2) and cannot be attributed to image artifacts. Moreover, the concordant findings were typical for acute myocarditis. Hence, it is possible that these 4 cases of acute myocarditis represent a rare, potential adverse event linked to mRNA COVID-19 vaccination.

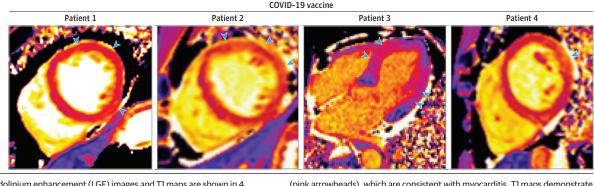
On January 7, 2021, the first patients at our institution began receiving COVID-19 vaccines, and vaccination was opened to all people 16 years and older on April 7, 2021. The first patient at our institution with myocarditis following COVID-19 vaccination was observed in the middle of February 2021, in line with the timing of the second vaccination dose. As of April 30, 2021, there were 561197 individuals in North Carolina living in the 6 counties surrounding our institution who had been fully vaccinated against COVID-19 infection, representing 33% of the population (statewide, 99.1% received an mRNA vaccine).9 Since we have identified 4 patients with myocarditis following vaccination, this indicates that if COVID-19 vaccination is associated with myocarditis, it is quite rare. Nonetheless, these 4 patients represent the majority of patients with acute myocarditis identified in the past 3 months at our institution, and this

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Figure 2. Cardiac Magnetic Resonance Imaging in Patients With Acute Myocarditis Following COVID-19 Vaccination



B T1 map



Late gadolinium enhancement (LGE) images and T1 maps are shown in 4 patients who recently received COVID-19 vaccine. Patients 1, 2, and 4 demonstrate epicardial LGE, and patient 3 demonstrates patchy, diffuse LGE

(pink arrowheads), which are consistent with myocarditis. TI maps demonstrate abnormal (elevated) native TI in the regions with LGE (blue arrowheads).

led to the highest total number of patients with acute myocarditis compared with the same 3-month period for the past 5 years (Figure 1). Additionally, we identified only those patients with severe unremitting chest pain who sought medical attention. Those with mild or moderate chest pain might not seek medical attention, and it is possible that subclinical myocarditis may occur and could be detected by active surveillance, as has been described with smallpox vaccination.¹⁰

Fortunately, the hospital courses of the 4 patients with myocarditis following COVID-19 vaccination were uneventful, and they were discharged within 2 to 4 days. In brief follow-up, none required rehospitalization (1 was seen in the emergency department for chest pain with negative troponin and pro-brain-type natriuretic peptide levels and was discharged), suggesting that the early prognosis could be benign. This is consistent with the study by Ammirati et al¹¹ that showed that the cardiac mortality plus transplant rate was 0% at 5 years in patients with an uncomplicated presentation of acute myocarditis. Conversely, patients with a complicated presentation (left ventricular ejection fraction, <50% with sustained ventricular arrhythmias or a low cardiac output syndrome requiring inotropes or mechanical circulatory support) had a cardiac mortality plus transplant rate of 14.7% at 5 years. Further study is needed to determine the range of clinical presentations and outcomes for patients with myocarditis following mRNA vaccines.

Limitations

Several caveats should be considered. First, there is no control group, and it is not possible to compare rates of acute myocarditis between those randomly assigned to receive vaccination vs no vaccination. Second, given the media attention, there could be recall or referral bias after COVID-19 vaccination compared with other vaccine exposures, and any temporal link between vaccination and myocarditis could just be due to chance. Third, although tests for COVID-19 infection and respiratory viruses were undertaken in some of the patients, these tests are neither foolproof nor comprehensive. Finally, there is no serological data, which could provide evidence of an excessive response to vaccination or prior subclinical COVID-19 infection (as determined by antinucleocapsid antibodies).

Conclusions

The findings from the present report raise the possibility of an association between mRNA COVID-19 vaccination and acute myocarditis. Since most individuals in North Carolina have received mRNA vaccines, it remains unknown if acute myocarditis will be observed following non-mRNA vaccines. Additionally, whether patients who experience acute myocarditis following COVID-19 vaccination should receive subsequent booster vaccinations for COVID-19 variants (or annual vaccinations for prophylaxis, if needed) is an open question, and the risk vs benefit profile will have to be considered carefully.

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Concept and design: H. W. Kim, Azevedo, Campbell, R. J. Kim.

Acquisition, analysis, or interpretation of data: All authors.

Drafting of the manuscript: H. W. Kim, Jenista, Wendell, Campbell, Darty, Parker, R. J. Kim. Critical revision of the manuscript for important intellectual content: H. W. Kim, Jenista, Azevedo, Campbell.

Administrative, technical, or material support: H. W. Kim, Jenista, Wendell, Azevedo, Campbell, Darty.

Supervision: H. W. Kim, R. J. Kim.

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