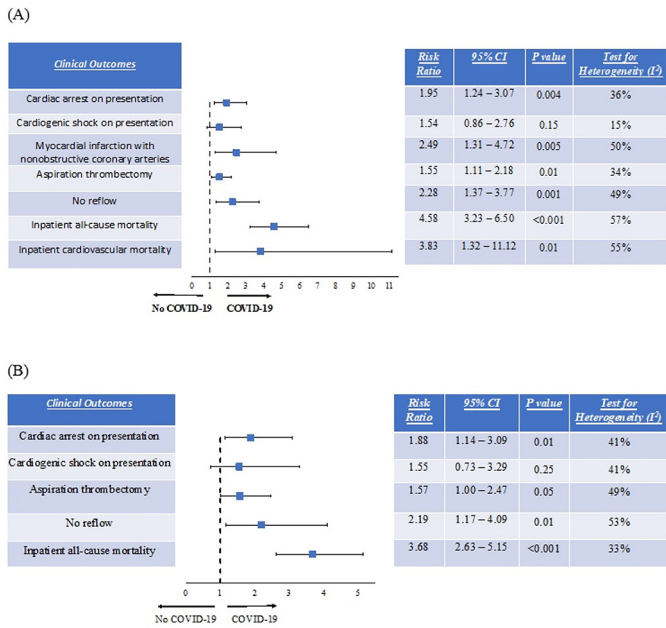




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Background The outcomes of patients with acute coronary syndrome (ACS) and COVID-19 infection are variable. We performed a pooled analysis of studies comparing the outcomes of ACS in patients with COVID-19 versus no COVID-19 infection.

Methods Statistical analysis was performed using Revman V.5.3 and Mantel Haenszel risk ratio. Outcomes studied were 1) in-hospital all-cause and cardiovascular mortality; 2) cardiac arrest on presentation; 3) myocardial infarction with nonobstructive coronary arteries (MINOCA); 4) aspiration thrombectomy use; and 5) no reflow phenomenon.

Results Nine studies (6,664 patients) met the inclusion criteria. Patients with ACS and COVID-19 infection have 4.6 times and 3.8 times higher risk of in-hospital all-cause and cardiovascular mortality (RR 4.58, 95% CI 3.23 - 6.50, p<0.001) (RR 3.83, 95% CI 1.32- 11.12, p=0.01), respectively, compared to patients without COVID-19. They also have a significantly high risk of cardiac arrest on presentation (RR 1.95, 95% CI 1.24 - 3.07, p=0.004). There was an elevated risk of requiring aspiration thrombectomy (RR 1.55, 95% CI 1.11 - 2.18, p=0.01) and no reflow (RR 2.28, 95% CI 1.37 - 3.77, p=0.001), along with higher risk of MINOCA (RR 2.49, 95% CI 1.31 - 4.72, p=0.005) in COVID-19 patients. Subgroup analysis of patients with STEMI also showed a significantly higher risk of in-hospital all-cause mortality, cardiac arrest on presentation, no reflow, and use of aspiration thrombectomy. Serum C-reactive protein (MD 65.33 mg/L, 95% CI 44.42 - 86.23, p<0.001) and D-dimer levels (MD 1.48 mg/L, 95% CI 0.65 - 2.31, p=0.005) were significantly higher in COVID-19 patients.

Conclusions Patients with ACS and COVID-19 have an increased risk of in-hospital all-cause and cardiovascular mortality, as well as higher risk of aspiration thrombectomy use, no reflow and MINOCA compared to no COVID-19 patients.

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200.09 Myocarditis After COVID-19 Vaccination: A Systematic Review of Case Studies

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Background The coronavirus disease of 2019 (COVID-19) is a global pandemic with over 200 million cases and four million deaths

Characteristics	N (%)	Laboratory and Testing Variables	N (%)	Treatment	N (%)
Age, mean (range), year	25.0 (14-70)	Troponin		NSAID	25 (34.2)
Sex		Elevated	29 (39.7)	Colchicine	17 (23.3)
Male	69 (94.5)	Not elevated	0	Steroids	7 (9.6)
Female	4 (5.5)	Not reported	44 (60.3)	Beta-blocker	7 (9.6)
Vaccine type		cTnI, mean (SD), ng/mL	8.3 (8.7)	IVIG	6 (8.2)
BNT-162b2	47 (64.4)	Peak cTnI, mean (SD), ng/mL	18.1 (15.3)	Aspirin	4 (5.5)
mRNA-1273	25 (34.2)	hs-cTnI, mean (SD), ng/mL	2.081 (2.459)	ACEI/ARB	3 (4.1)
Ad.26.COV2.S	1 (1.4)	Peak hs-cTnI, mean (SD), pg/mL	6.028 (2.098)	Diuretics	1 (1.4)
Vaccine dose		cTnI, mean (SD), ng/L	373.1 (463.5)	Statins	1 (1.4)
First	9 (12.3)	Peak cTnT, mean (SD), ng/L	688.1 (564.9)	Clopidogrel	1 (1.5)
Second	64 (87.7)	WBC		Not reported	39 (53.4)
Time to onset, mean (SD), day	3.5 (3.82)	Normal	15 (20.6)	*Fictitious atrial rhythm, sinus tachycardia, non-sustained ventricular tachycardia, left axis deviation, and incomplete right bundle branch block	
Length of stay, mean (SD), day	5.2 (1.80)	Abnormal	8 (10.9)		
Not reported	50 (68.5)	WBC count, mean (SD), /µL	8,987 (3,943)		
CRP		Elevated	34 (46.6)		
Yes	20 (27.4)	Not elevated	2 (2.6)		
No	20 (27.4)	Not reported	37 (50.7)		
Not reported	33 (45.2)	CRP, mean (SD), mg/L	46.3 (41.0)		
Chest pain		ESR			
Yes	64 (87.7)	Elevated	14 (19.2)		
No	9 (12.3)	Not elevated	13 (17.8)		
Not reported	9 (12.3)	Not reported	46 (63.0)		
Chills		ESR, mean (SD), mm/h	52 (71.2)		
Yes	12 (16.4)	BNP			
No	52 (71.2)	Elevated	2 (2.7)		
Not reported	9 (12.3)	Not elevated	7 (9.6)		
Myalgia		Not reported	64 (87.7)		
Yes	12 (16.4)	BNP, mean (SD), pg/mL	71.2 (60.4)		
No	52 (71.2)	EKG			
Not reported	9 (12.3)	Normal	3 (4.1)		
Headache		ST elevation	57 (78.1)		
Yes	7 (9.6)	ST depression	2 (2.7)		
No	57 (78.1)	PR depression	8 (11.0)		
Not reported	9 (12.3)	T wave inversion	27 (37.0)		
Dyspnea		Others*	8 (11.0)		
Yes	7 (9.6)	Not reported	3 (4.1)		
No	57 (78.1)	LVEF			
Not reported	9 (12.3)	Temperature, °C			
Vital Signs	Mean (SD)	SBP, mmHg	37.8 (0.78)		
Temperature, °C		DBP, mmHg	119.8 (14.1)		
<50%		PR, beats per minute	71.6 (11.1)		
50%		RR, beats per minute	91.4 (18.5)		
Not reported		SPO ₂ , %	18.0 (1.31)		
53.3 (4.7)		SPO ₂ , %	98.7 (1.32)		

worldwide. Anti-COVID-19 vaccinations have had exceptional success in subduing the incidence, prevalence, and disease severity of COVID-19, but rare cases of myocarditis have been reported after COVID-19 vaccinations.

Methods We performed a systematic literature search on PUBMED, MEDLINE, EMBASE, and Cochrane Reviews database from inception to July 18, 2021. Studies were analyzed based on predetermined eligibility criteria.

Results A total of 19 studies containing 73 cases of COVID-19 vaccine-associated myocarditis were catalogued. Mean age was 25 years, and male to female ratio was 17:1. For 87.7% of patients, myocarditis occurred after the second dose. Average time to onset and length of hospitalization were 3.5 days and 5.2 days, respectively. Prognosis was benign with 100% recovery. Chest pain (100%); elevation of troponin (100%) and CRP (94.4%); and ST elevation on EKG (81.4%) were common. NSAIDs (73.5%) were the most used medication, followed by colchicine (50%).

Conclusions Patients with COVID-19 vaccine-associated myocarditis are usually younger males presenting with chest pain 3.5 days after receiving their second dose. Work-up typically shows elevation of troponin and CRP with ST changes in EKG. Diagnosis is made after excluding all other etiologies. Given significant population benefit from COVID-19 vaccination, physicians should continue to encourage vaccination while remaining vigilant of the very rare occurrence of myocarditis following COVID-19 vaccination.

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ENDOVASCULAR – Critical Limb Ischemia

300.01 Utilization of EKOS (Ultrasound-Accelerated Thrombolysis) in the Treatment of Acute Limb Ischemia: One-Year Outcome Follow-Up
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Background Acute limb ischemia (ALI) is a serious complication of peripheral arterial disease and critical limb ischemia. ALI can be difficult to treat since it often involves many vessels. If detected early enough, ALI can be treated by various endovascular techniques to quickly revascularize the affected vessels to prevent negative outcomes and future complications. Improvements in endovascular technologies are key to fast and efficient re-perfusion. The purpose of this case series is to analyze the efficacy of EKOS ultrasound-accelerated thrombolysis (Boston Scientific, USA) as a treatment