



Cardiac abnormalities in athletes after SARS-CoV-2 infection: a systematic review

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ABSTRACT

Objectives Quantification of pericardial/myocardial involvement and risks of sudden cardiac arrest/sudden cardiac death (SCA/SCD) after SARS-CoV-2 infection in athletes who return to sports.

Design Systematic review on post-SARS-CoV-2 infection pericardial/myocardial manifestations in athletes.

Data sources Combinations of key terms in Medline, Embase and Scopus (through 2 June 2021).

Eligibility criteria for selecting studies Inclusion: athletes, with cardiovascular magnetic resonance (CMR) or echocardiography after recovery from SARS-CoV-2 infection, including arrhythmia outcomes. Exclusion: study population ≥ 1 individual comorbidity and mean age < 18 or > 64 years. Quality assessment was performed using Joanna Briggs Institute Critical Appraisal tools checklists.

Results In total, 12 manuscripts (1650 papers reviewed) comprising 3131 athletes (2198 college/student athletes, 879 professional athletes and 54 elite athletes) were included. The prevalence of myocarditis on echocardiography and/or CMR was 0%–15%, pericardial effusion 0%–58% and late gadolinium enhancement (LGE) 0%–46%. Weighted means of diagnosed myocarditis were 2.1% in college/student athletes and 0% in elite athletes. The prevalence of LGE was markedly lower in studies with high-quality assessment scores (3%–4%) versus low scores (38%–42%). A single study reported reversibility of myocardial involvement in 40.7%. No important arrhythmias were reported. Ten studies (n=4171) reporting postrecovery troponin T/I found no clear relationship with cardiac abnormalities.

Summary/conclusion Athletes have an overall low risk of SARS-CoV-2 pericardial/myocardial involvement, arrhythmias and SCA/SCD. Rates of pericardial/myocardial abnormalities in athletes are highly variable and dependent on study quality. Troponin screenings seem unreliable to identify athletes at risk for myocardial involvement. Prospective athlete studies, with pre-SARS-CoV-2 imaging (CMR), including structured follow-up and arrhythmia monitoring, are urgently needed.

INTRODUCTION

SARS-CoV-2 has spread globally and was declared a pandemic by the WHO on 11 March 2020.¹ Most studies performed early in

Key messages

What is already known

- Rates of sustained cardiac involvement after SARS-CoV-2 recovery have been reported to be highly variable in both symptomatic and asymptomatic athletes.
- Physical exercise is thought to be associated with a worsening of prognosis in viral myocarditis in general.
- Risk of potentially fatal arrhythmias in athletes with cardiac involvement after recovery from SARS-CoV-2 is largely unknown.
- No clear consensus exists on how to screen for post-SARS-CoV-2 cardiac abnormalities, and existing consensus documents include different diagnostic modalities (ECG, imaging and biomarkers) and cut-offs.

What are the new findings

- Athletes have an overall low risk (0%–5%) of SARS-CoV-2 pericardial/myocardial involvement, with no reported arrhythmias or sudden cardiac arrest/sudden cardiac death.
- Rates of pericardial/myocardial abnormalities in athletes are highly variable per study and dependent on study quality, with higher quality studies reporting rates of 3%–4%.
- Troponin screenings are unreliable to identify athletes at risk for myocardial involvement.
- Prospective athlete studies, with pre-SARS-CoV-2 CMR, including structured follow-up and arrhythmia monitoring, are urgently needed.

the pandemic focused on cardiac complications in hospitalised and severely ill patients, but recently, there has been marked interest in SARS-CoV-2 cardiac complications and sequelae in young and healthy individuals, including athletes.²

The initial studies in asymptomatic athletes or athletes with a mild or moderate course of illness reported considerable sustained cardiac involvement after recovery from SARS-CoV-2 infection. Later studies have not replicated these findings. Considering that

physical exercise has been associated with a worsening of prognosis in viral myocarditis,³⁻⁵ cardiac involvement in athletes who have recovered from SARS-CoV-2 infection is of particular relevance, and a clear scope of the problem is needed.

Multiple return-to-play consensus statements have been published to assist clinicians in screening for SARS-CoV-2 cardiac abnormalities.⁶⁻⁹ These consensus statements are likewise highly variable and include different diagnostic modalities (eg, ECGs, exercise tests, Holter monitoring, troponin levels and echocardiography) and cut-offs. Numerous statements place emphasis on serum troponin levels, as they have been observed to be elevated in severely ill patients as compared with those with milder COVID-19¹⁰ and negatively correlated with prognosis.¹¹⁻¹⁴ Yet, little is known about the efficacy of such screenings.^{15 16} Finally, the risk of potentially fatal arrhythmias in athletes with cardiac involvement after recovery from SARS-CoV-2 remains largely unknown.

We therefore aimed to address the following research questions: in athletes recovered from SARS-CoV-2 infection: (1) what is the prevalence of SARS-CoV-2 cardiac

involvement (as found using imaging investigations), stratified by study quality?; (2) what is the incidence of ventricular tachyarrhythmias (and SCA/SCD) in athletes with documented cardiac involvement?

We performed a systematic search, review and quality assessment to provide a comprehensive overview of myocardial and pericardial involvement after SARS-CoV-2 infection, long-term cardiac sequelae after infection and risks of sudden cardiac arrest/sudden cardiac death (SCA/SCD) in athletes.

METHODS

Literature search

We conducted a systematic search strategy according to Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines¹⁷ (PROSPERO ID:CRD42021233964). The comprehensive electronic search was performed in Medline (Ovid), Embase (Ovid) and Scopus (final update 2 June 2021), with combined terms or synonyms for: COVID-19, SARS-CoV-2, cardiovascular imaging, cardiac MRI, echocardiography, SCD, athletes, (young) adults and arrhythmias (figure 1).

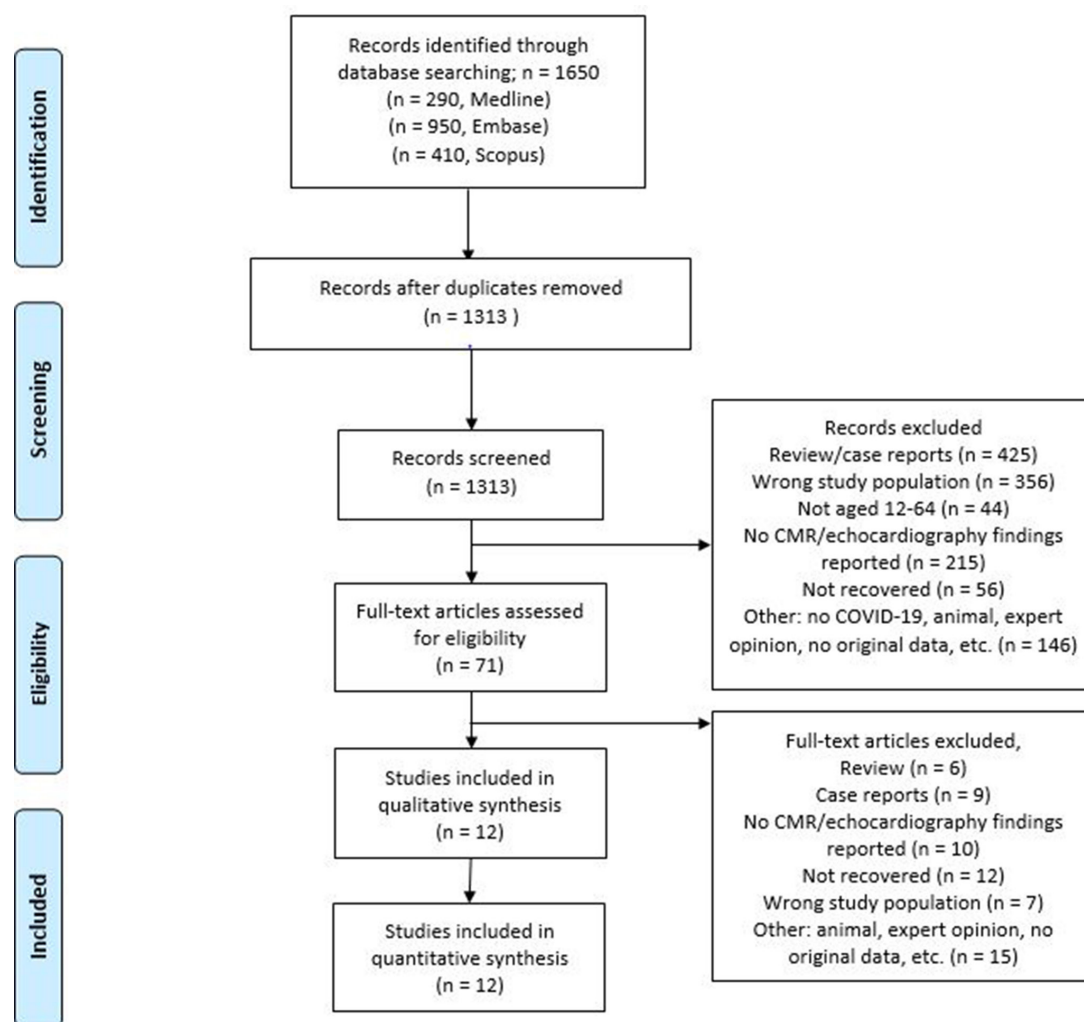


Figure 1 PRISMA 2009 flow chart describing selection of studies included in the systematic review.¹⁷ CMR, cardiovascular magnetic resonance; PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses.

Conference abstracts from Embase (Ovid) were included to identify unpublished studies. Subsequently, we performed manual citation and reference searching of the included studies in Google Scholar. The full search strategy can be found in online supplemental file 1.

Study selection

We included studies when they: 1) contained clinical data on athletes recovered from SARS-CoV-2 infection, (2) reported cardiovascular imaging by either cardiovascular magnetic resonance (CMR) or echocardiography, (3) performed investigations after SARS-CoV-2 infection, with emphasis on arrhythmia outcomes. Athletes were defined as individuals of young or adult age, either amateur or professional, who engage in regular exercise training and participate in official sports competitions. College/student athletes were defined as individuals of young age, who participate in organised competitive sports sponsored by the educational institution in which the student is enrolled, and elite athletes, (national team, Olympians, and professional athletes) were defined as athletes who generally exercise >10 hours/week.¹⁸ Exclusion criteria were: (1) study population with ≥ 1 mean individual comorbidities, (2) mean age of total study population <18 or >64 years, (3) no original data reported and (4) case reports. Two investigators (JCvH and JLS) independently screened and assessed all identified manuscripts. Discrepancies were resolved by discussion between the two reviewers and external team-based expert¹⁹ opinion, until consensus was reached.

Outcomes

Our primary outcome of interest was myocardial and/or pericardial abnormalities/manifestations as documented using CMR or echocardiography, in athletes recovered from SARS-CoV-2 infection. Secondary outcomes included arrhythmias (if reported) and SCA/SCD and potential associations between imaging abnormalities and serum troponin levels.

Data extraction

First, we extracted data on authors, demographic findings, sample sizes, time to imaging, clinical features, elevated troponin levels (according to individual study-defined cut-offs), reported arrhythmias or SCA/SCD and time of investigation/duration of follow-up (FU). Second, we extracted data on CMR findings, which included rates of elevated T1 time and elevated T2 time. In short, both T1 and T2 times are calculated using an MRI technique for myocardial tissue characterisation. An increased T1 time can be indicative for cardiomyopathies and increased T2 times for myocardial inflammation. We also extracted data of pericardial effusion (PE) according to individual study-defined cut-offs, pericardial enhancement and presence of late gadolinium enhancement (LGE). In short, LGE demonstrated regional differences in myocardial extracellular volume, and abnormal uptake and washout patterns (myocardial, non-ischaemic,

pericardial, hinge-point or other) within the extracellular space, such as seen in myocardial inflammation and fibrosis. In addition, we extracted the criteria used to diagnose myocarditis (if reported) and the amount of diagnosed myocarditis. Third, we extracted data on echocardiographic findings, which included left ventricular (LV) end-diastolic diameter (EDD), LV ejection fraction (LVEF), right ventricle (RV) ejection fraction (RVEF), right ventricle function/tricuspid annular plane systolic excursion (TAPSE) (ie, right ventricular longitudinal contraction or shortening, a measure of right ventricular function), LV and RV global longitudinal strain (GLS) (ventricular deformation patterns), early to late diastolic transmitral flow velocity (E/A), ratio of early diastolic mitral inflow velocity/ tissue velocity (E/e'), segmental wall motion abnormalities, PE (according to study-defined cut-offs) and systolic pulmonary artery pressure. Two independent reviewers extracted the relevant data, which was stored in a Microsoft Excel database and exported to R for statistical analysis.

Statistical methods and quality assessment

Data are presented as means and SD, medians with IQR and weighted means (for data synthesis), as appropriate.²⁰ Statistical analyses were performed using R (V.1.3.1993).

Data quality was assessed using the Joanna Briggs Institute Critical Appraisal tools checklists²¹ by two authors independently. Discrepancies were resolved after joint article review and discussion. The relationship between data quality and severe cardiac abnormalities, defined as LGE and PE on CMR, was subsequently plotted.

RESULTS

The initial search yielded 1650 papers. After extensive review, we included a total of 12 manuscripts comprising 3131 recovered post-SARS-CoV-2 athletes. Except for the study by Gervasi *et al*,²² all studies were cross-sectional or retrospective and did not include pre-SARS-CoV-2 imaging. Study and demographic characteristics of the respective studies are shown in [table 1](#) for studies using CMR as their primary imaging modality, and in [table 2](#) for studies using echocardiography as their primary imaging modality. Overall, median age of all study participants was 22 years (IQR 18–41); 36% were women. In five studies comprising 268 athletes including study controls CMR was performed; in three studies comprising 897 athletes including study controls echocardiography was performed; in four studies comprising 1968 athletes including study controls both CMR and echocardiography were performed. COVID-19 was diagnosed by real-time reverse transcriptase PCR in seven studies, by antibody testing in two studies and in two studies by one of the two; one study (Cavigli *et al*)²³ study did not report diagnostic tests. All included athletes had recovered from SARS-CoV-2 infection, of which 0.1% recovered in-hospital and 99.9% at home. Two studies (Gervasi *et al* and Cavigli *et al*) reported arrhythmia outcomes, and one study (Moulson *et al*)²⁴ reported a resuscitated SCA after

Table 1 CMR patient characteristics

Study	Patient selection	Age (years)	Time to CMR (days)	Severity no. (%) [*]	EDV (mL)		EDV index (mL/m ²)		EF (%)		Troponin elevated no. (%)
					LV	RV	LV	RV	LV	RV	
Hendrickson et al ^{§1}	5 SARS-CoV-2 recovered competitive college athletes (68% men)	20 (18–27)†	16 (12–34)	1: 25 (19)† 2: 75 (55)† 3: 37 (27)†	NR	NR	98 (90–111)	92 (82–100)	56 (53–62)	56 (52–62)	4 (3)†
Vago et al ^{‡2}	12 elite athletes recovered from COVID-19 (17% men)	23 (20–23)	17 (17–19)	1: 2 (17) 2: 10 (83)	NR	NR	100 (95–110)	NR	58 (55–61)	NR	0 (0)
					15 age-matched and sex-matched healthy elite athletes	NR	102 (98–109)	NR	57 (54–60)	NR	0 (0)
					15 healthy controls	NR	85 (80–89)	NR	60 (58–63)	NR	0 (0)
Rajpal et al ^{‡6}	26 competitive college athletes with COVID-19 (58% men)	19.5 (±1.5)	11–53	1: 14 (54) 2: 12 (46)	NR	NR	NR	NR	58	57	0 (0)
Malek et al ^{‡3}	26 elite volleyball and soccer athletes with COVID-19 (19% men)	24 (21–27)	32 (22–62)	1: 6 (23) 2: 14 (54) 3: 5 (19) 4: 1 (4)	NR	NR	95 (84–106)	96 (85–105)	61 (60–62)	59 (57–60)	0 (0)
Brito et al ^{‡7}	48 student athletes recovered from COVID-19 (85% men)	19 (19–21)	27 (22–33)	1: 16 (30) 2: 36 (66) 3: 2 (4)	NR	NR	104.7 (93.1–117.2)	92.4 (82.9–107.3)	60.3 (56.7–63.3)	54.6 (47.6–59.8)	1 (3)
Clark et al ^{‡4}	59 college athletes recovered from COVID-19 (37% men)	20 (19–34)	21.5 (13–37)	1: 13 (22) 2: 46 (78)	NR	NR	114.4 (99.5–118.7)	99.8 (95.5–105.3)	59.09 (54.8–61.6)	51.3 (50.8–57.2)	0 (0)
					59 post-COVID-19 athletes	160 (143–213)	93 (84–100)	100 (91–115)	60 (56–63)	53 (50–56)	NR
					27 healthy controls	164 (138–210)	88 (78–99)	89 (78–104)	60 (57–64)	58 (55–60)	NR
					60 athletic controls	195 (167–235)	95 (86–111)	99 (87–117)	58 (56–60)	53 (51–57)	NR
Starekova et al ^{‡8}	145 competitive student athletes recovered from COVID-19 (74% men)	20 (17–23)	15 (11–194)	1: 24 (17) 2: 71 (49) 3: 40 (28)	NR	NR	104 (26)	110 (22)	58 (5)	54 (6)	2 (1)
Moulson et al ^{‡4}	312 SARS-CoV-2 recovered athletes (52% men)	20 (±2)	33 (18–63)	1: 5† 2: 4† 3: 4† 4: 8†	NR	NR	NR	NR	55 (41–67)‡	NR	24 (1)§
Daniels et al ^{‡5}	1597 college athletes recovered from COVID-19 (60% men)	NR	22 (10–77)	NR	NR	NR	NR	NR	NR	NR	6 (0–4)

^{*}1: asymptomatic; 2: mild; 3: moderate; and 4: severe symptoms.

†Values based on the total CMR study population (n=137).

‡Values only based on SARS-CoV-2 pericardial/myocardial involved athletes (n=21); no data were reported for the total CMR study population (n=312).

§Values based on the entire study (n=2719) population, not specified for the CMR study population.

CMR, cardiovascular magnetic resonance; EDV, end-diastolic volume; EF, ejection fraction; LV, left ventricle; RV, right ventricle.

Table 2 Post-SARS-CoV-2 diagnostic echocardiography features

Study	Patient selection	Age (years)	Time to echo (days)	Severity No. (%)	N	Troponin elevated No. (%)	LVEDD (mm/m ²)	LVEF %	RV function/ TAPSE (mm)	Strain (%)			E/A, E/e' ratio No. (%)	Segmental wall motion abnormalities, no. (%)	Pericardial effusion >5 mm, no. (%)	SPAP (mm Hg)	
										LV	RV						
Hendrickson <i>et al</i> ³¹	5 SARS-CoV-2 recovered college athletes	20 (18–27)†	NR	1: 25 (18)† 2: 75 (55)† 3: 37 (27)†	5 SARS-CoV-2 recovered college athletes	4 (3)†	NR	NR	NR	NR	NR	NR	NR	4 (3)†	NR	NR	
Gervasi <i>et al</i> ²²	18 professional soccer players recovered from COVID-19 (63.7% male)	22 (21–27)	15	1: 6 (33) 2: 12 (67)	18 post-COVID-19 professional soccer players	1 (5.6)	55.5 (52–58)	61 (58–63)	19.5 (19–22)	NR	NR	1.71 (1.5–1.9)	NR	NR	NR	NR	
Moulson <i>et al</i> ²⁴	21 SARS-CoV-2 recovered athletes (52% male)	20 (±2)	15 (11–25)	1: 5† 2: 4† 3: 4† 4: 8†	21 SARS-CoV-2 recovered athletes	24 (0.9)§	NR	51 (45–65)	NR	NR	NR	NR	1 (0.04)†	6 (0.27)†	NR	NR	
Brito <i>et al</i> ²⁷	54 student athletes recovered from COVID-19 (85% male)	19 (19–21)	27 (22–33)	1: 16 (30) 2: 36 (66) 3: 2 (4)	38 COVID-19 positive symptomatic athletes 16 COVID-19 positive asymptomatic athletes 20 COVID-19 negative controls	1 (3)	NR	60 (55–64)	NR	NR	6.2 (5.6–7.5)	NR	NR	2 (5)	NR	NR	
Cavigli <i>et al</i> ²³	90 professional athletes recovered from COVID-19 (71% male)	24 (±10)	NR	1: 21 (23) 2: 69 (77)	90 post-COVID-19 professional athletes	NR	NR	NR	NR	NR	NR	NR	0 (0)	3 (3)	NR	NR	
Martinez <i>et al</i> ³⁵	789 professional athletes recovered from COVID-19 (98.5% male)	25 (19–41)	19 (3–156)	1: 329 (42) 2: 460 (58)	789 post-COVID-19 professional athletes	6 (0.8)	NR	49	NR	NR	NR	NR	1 (0.1)	3 (0.4)	NR	NR	
Daniels <i>et al</i> ²⁵	1597 SARS-CoV-2 recovered college athletes	NR	NR	NR	1597 SARS-CoV-2 recovered college athletes	6 (0.4)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR

*1: asymptomatic-, 2: mild-, 3: moderate-, 4: severe symptoms.

†Percentage based on total echocardiography population (n=2231).

‡Values only based on SARS-CoV-2 perimyocardial/myocardial involved athletes (n=21); no data were reported for the total echocardiography study population (n=2231).

§Values based on the entire study (n=2719) population, not specified for the echocardiography study population.

LV, left ventricle; LVEDD, LV end-diastolic diameter; LVEF, LV ejection fraction; NR, not reported; RV, right ventricle; SPAP, systolic pulmonary artery pressure; TAPSE, tricuspid annular plane systolic excursion.

recovery. Only one study (Daniels *et al*²⁵) reported structured FU.

Risk of bias

Relevant checklists according to the Joanna Briggs Institute Critical Appraisal tools were applied (eight cohort studies, two cross-sectional studies, one case series and one case-control study). Detailed quality assessment for individual studies can be found in online supplemental file 1. The risk of bias was the highest in the cohort studies of Rajpal *et al*²⁶ and the cross-sectional study of Brito *et al*.²⁷ The risk of bias was the lowest in the case-control study of Gervasi *et al*²² and the case-series study of Starekova *et al*.²⁸ Other prevalent methodological issues in the cohort studies were not adequately reporting if the participants were free of outcomes at the start of the study (8/8), not reporting completion of FU or reasons for incomplete FU (7/8) and not reporting strategies used to address incomplete FU (7/8).

CMR findings

Studies including CMR are outlined in table 1; main myocardial and pericardial findings are summarised in table 3. Figure 2 illustrates the prevalence of total LGE stratified by risk of bias, and figure 3 illustrates the prevalence of PE stratified by risk of bias. Median time interval from symptom onset to CMR examination was 32 (IQR 11–194) days. In nine studies in 2477 athletes and study controls, the weighted mean for the prevalence of elevated (according to study-defined cut-offs) T1 was 1.5%, elevated T2 was 2.5%, LGE including hinge-point fibrosis was 4.9%, LGE excluding hinge-point fibrosis was 2.6% and PE 11.3%. The weighted mean for the prevalence of active myocarditis according to the Lake Louise criteria (LLC)²⁹ or modified LLC³⁰ was 1.9%.

Athletes

In total nine studies reported findings in 2360 athletes (2326 student/college athlete, and 34 elite athletes), whereof 22% were asymptomatic, 59% had mild, 19% had moderate and 0.2% severe symptoms. The weighted overall mean for the prevalence of elevated T1 was 1.5%, T2 2.6%, LGE 5.4% (six studies reported only total LGE with no specification whether this was myocardial or pericardial), LGE excluding hinge-point fibrosis 2.6%, PE 11.3%, myocardial LGE (if reported) 1.9% and pericardial LGE (if reported) 18.7%. Overall, 2.0% met the LLC for myocarditis, of which 2.1% in college/student athletes, and 0% in elite athletes. Daniels *et al* and Starekova *et al* reported both one athlete with pericarditis, and Moulson *et al* reported one athlete with peri-myocarditis. We observed considerable differences between individual studies, ranging from no myocardial involvement (Vago *et al*) to 15% (Rajpal *et al*²⁶). Hendrickson *et al*³¹ reported CMR findings in only five athletes, with no abnormal findings detected by CMR.

Most studies reported the presence of PE using a cut-off of 10 mm; Brito *et al*²⁷ applied 5 mm as their cut-off.

Consequently, Brito *et al* reported (in college athletes) the highest prevalence of PE (58%). Of note, they also reported significantly higher rates of pericardial LGE in asymptomatic athletes as compared with symptomatic athletes (82% vs 27%, $p=0.003$). In elite athletes specifically, Vago *et al*³² reported no abnormal T1, T2 or LGE, or myocarditis (LLC). Also in elite athletes, Malek *et al*³³ reported no abnormal T1 but elevated T2 in 4%, LGE in 4% and PE in 8%. Moulson *et al* only reported CMR values of 21 athletes with pericardial/myocardial involvement and not in the total CMR population ($n=312$). The largest (survey) study (Daniels *et al*²⁵) reported that 37 (2.3%) athletes met diagnostic criteria for myocarditis. Interestingly, these 37 CMR-identified subclinical potential cases of myocarditis included 20 athletes without cardiovascular symptoms and with normal ECGs, echocardiography and troponin test results. When compared with alternative diagnostic strategies, the CMR strategy resulted in a 7.4-fold diagnostic increase compared with the symptom driven strategy, and a 2.8-fold increase compared with the ECG, echocardiogram and troponin strategy. In addition, the survey study by Daniels *et al*²⁵ was the only study to report FU CMRs in 73% of individuals with clinical or subclinical myocarditis, with the second CMR taking place between 4 and 14 weeks after confirmation of SARS-CoV-2 infection. Complete resolution of both T2 mapping and LGE was found in 40.7% of these individuals; resolution of T2 mapping abnormalities but persistent LGE was found in 59.3%. No other FU data, such as results from arrhythmia monitoring, was reported in this study.

Study controls (healthy and/or athletes)

In total, two studies reported findings in 117 controls (15 age-matched and sex-matched elite athletes, 42 healthy controls and 60 athletic controls). The weighted overall mean for the prevalence of elevated T1 was 0%, T2 was 0%, LGE including hinge-point fibrosis was 8.5%, LGE excluding hinge-point fibrosis was 0%, PE not reported and 0% met the LLC for myocarditis. Clark *et al*,³⁴ reported no values for T1 and T2 but 24% ($n=10$) athletic controls with hinge-point fibrosis.

Echocardiography findings

Study and demographic characteristics of studies using echocardiography as their main imaging modality with level of evidence are summarised in table 2. The median time interval from symptom onset to echocardiography was 27 (IQR 3–156) days. The most frequently reported echocardiography parameters were LVEF, TAPSE and PE. In seven studies comprising 2606 athletes and study controls, the weighted mean for LVEF was 50.8%, TAPSE 19.9 mm and the prevalence of PE 0.6%.

Athletes

In total, six studies reported findings in 2574 athletes (1677 student athletes, 879 professional athletes and 18 elite athletes), in which 36% were asymptomatic, 60%

Table 3 Post-SARS-CoV-2 diagnostic CMR features

Study	N	T1, no. increased (%)	T2, no. increased (%)	PE, no. (%)	LGE present, no. (%)	LGE pattern, no. (%)			Criteria myocarditis	Confirmed myocarditis, no. (%)
						Myocardial	Non-Ischaemic	Pericardial		
Hendrickson <i>et al</i> ³¹	5 SARS-CoV-2 recovered college athletes	NR	0 (0)	2 (40)	0 (0)	NR	NR	NR	NR	0 (0)
Vago <i>et al</i> ³²	10 elite female athletes	0 (0)	0 (0)	NR	0 (0)	NR	NR	NR	NR	0 (0)
	15 age-matched and sex-matched healthy elite athletes	0 (0)	0 (0)	NR	0 (0)	NR	NR	NR	NR	
	15 healthy controls	0 (0)	0 (0)	NR	0 (0)	NR	NR	NR	NR	
Rajpal <i>et al</i> ²⁶	26 post-COVID-19 college athletes	9 (45)	13 (50)	2 (8)	12 (46)	NR	NR	NR	Lake Louise Criteria	4 (15)
									1 (4) epicardial 4 (15) linear 5 (19) patchy 2 (8) RV insertion	
Malek <i>et al</i> ³³	26 post-COVID-19 elite volleyball and soccer athletes	0 (0)	1 (4)	2 (8)	1 (4)	NR	NR	NR	Lake Louise Criteria	0 (0)
Brito <i>et al</i> ²⁷	37 symptomatic athletes	8 (22)	0 (0)	20* (54)	1 (3)	1 (3)	0 (0)	10 (27)	Lake Louise Criteria	NR
	11 asymptomatic athletes	1 (9)	0 (0)	8* (73)	0 (0)	0 (0)	0 (0)	9 (82)	Lake Louise Criteria	NR
Clark <i>et al</i> ³⁴	59 post-COVID-19 athletes	NR	NR	NR	16 (27)	1 (2)	0 (0)	1 (2)	Lake Louise Criteria (modified)	2 (3)
	27 healthy controls	NR	NR	NR	0 (0)	0 (0)	0 (0)	0 (0)	Lake Louise Criteria (modified)	NR
	60 athletic controls	NR	NR	NR	10 (24)	10 (24)	0 (0)	0 (0)	Lake Louise Criteria (modified)	NR
Starekova <i>et al</i> ²⁸	145 post-COVID-19 student athletes	NR	NR	NR	42 (29)	NR	NR	NR	Lake Louise Criteria	2 (1)
Moulson <i>et al</i> ²⁴	312 SARS-CoV-2 recovered athletes	7 (2)	7 (2)	13 (4)	12 (4)	NR	NR	NR	Lake Louise Criteria (modified)	5 (1.6)
									11 (4) inferior segments 1 (0.3) anterolateral segments	
Daniels <i>et al</i> ²⁵	1597 SARS-CoV-2 recovered college athletes	5 (0.3)	31 (1.9)	NR	37 (2.3)	NR	NR	NR	Lake Louise Criteria (modified)	31 (1.9)

 * ≥ 5 mm PE.

CMR, cardiovascular magnetic resonance; LGE, late gadolinium enhancement; PE, pericardial effusion.

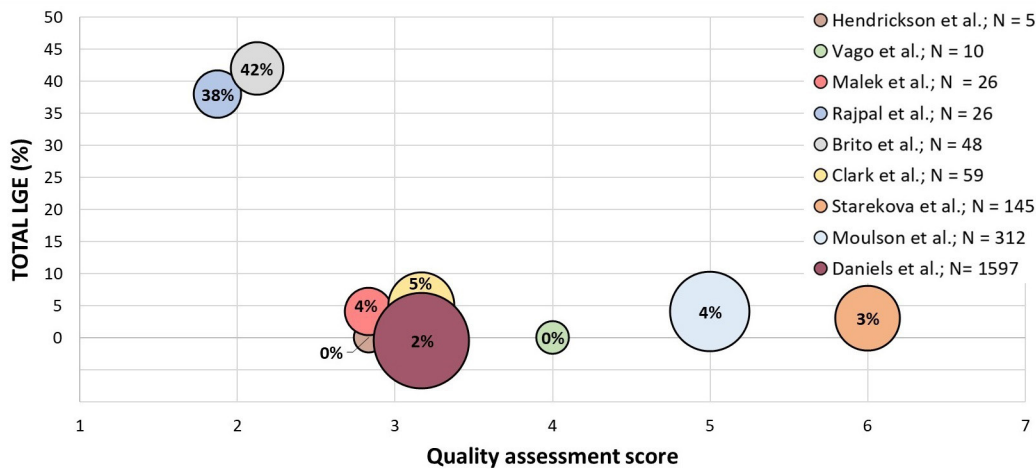


Figure 2 Total % of late gadolinium enhancement (LGE) detected on CMR stratified by risk of bias based on the quality assessment score. Each circle represents a single study. The area of the sphere represents the size of the study population. CMR, cardiovascular magnetic resonance.

had mild symptoms, 4% had moderate symptoms and 0 had severe symptoms. The weighted mean for LVEF was 50.7%, for TAPSE 19.5 mm and for the prevalence of PE 0.6%. Brito *et al.*²⁷ reported no significant differences between asymptomatic and symptomatic student athletes for LVEF ($p=0.5$) and PE ($p=0.5$). They identified three so-called phenotypic student athlete groups: mild pericardial disease, pericardial disease and mild myocardial disease on CMR. In a hierarchical clustering analysis, these groups were associated with GLS $<-16\%$ on echocardiography. Furthermore, Hendrickson *et al.*³¹ reported only a trace of PE in 4 (2.9%) athletes, with further normal evaluations. Gervasi *et al.*²⁴ reported no significant differences between COVID-19 positive soccer players and COVID-19 negative soccer players for LVEF ($p=0.55$) and TAPSE ($p=0.69$). In addition, Moulson *et al.*²⁴ only reported echocardiography data in 21 athletes with pericardial/myocardial involvement. Only those athletes

could therefore be included in the data synthesis, instead of the total echocardiography population ($n=2231$). The study of Daniels *et al.*²⁵ reported one athlete with symptoms of dyspnoea, with decreased LVEF (35%–40%) on echocardiography and CMR. Additionally, Martinez *et al.*³⁵ reported six cases of pericarditis/myocarditis (confirmed on CMR) of which four had abnormal echocardiography findings ('small' (not further defined) PE, LVEF 50%, new region wall abnormalities/preserved LVEF and dilated RV). Finally, Cavigli *et al.*²³ reported PE in three athletes with symptomatic SARS-CoV-2 infections, of which one confirmed myopericarditis and two cases of pericarditis.

Study controls (healthy and/or athletes)

In total two studies reported findings in 32 athletic controls. The weighted overall mean LVEF was 60.2%, for TAPSE -20.5 and for the prevalence of PE 0%. Gervasi *et al.* did not report strain and PE values.

Arrhythmias and SCA/SCD

Within a setting of a non-systematic FU period of 113 days, the large cohort of Moulson *et al.* ($n=2820$) demonstrated no instances of SCD, and only one (0.04%) resuscitated SCA. The case of SCA had a prior CMR, performed 17 days after onset of symptoms of COVID-19, without findings suggestive of acute cardiac involvement. The authors conclude that the aetiology of this event was considered uncertain and likely unrelated to COVID-19. No arrhythmia monitoring was performed during FU. Gervasi *et al.* reported arrhythmia outcomes: one athlete (5.6%) had ventricular premature beats (VPBs) during exercise and one athlete (5.6%) had supraventricular premature beats (SVPBs) during exercise, which were not present at exercise testing prior to SARS-CoV-2 infection. Holter monitoring was only performed post-recovery and demonstrated rare or occasional, isolated SVPB and/or VBP in all athletes. In addition, Cavigli *et al.* reported arrhythmia outcomes: 24-hour ambulatory ECG

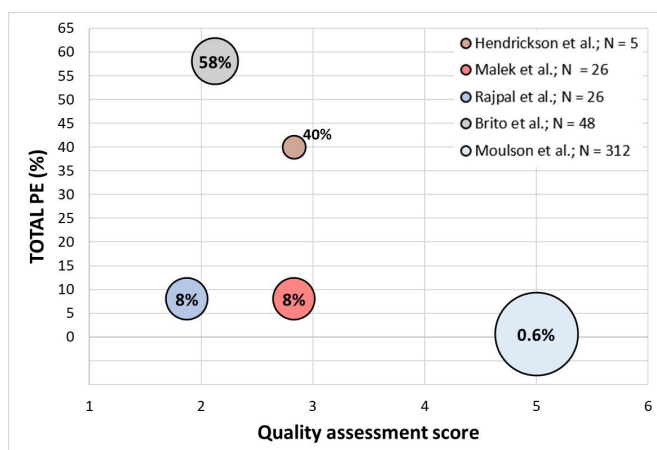


Figure 3 Total % of pericardial effusion (PE) detected on CMR stratified by risk of bias based on the quality assessment score. Each circle represents a single study. The area of the sphere represents the size of the study population. PE, pericardial effusion.

monitoring showed isolated VPBs and SVPBs in 53.3% and 52.5%, respectively, with no malignant arrhythmias. During CPET, neither VPBs nor ST segment or T wave abnormalities were found, except for an athlete showing isolated VPBs and couplets (RBBB configuration with wide QRS-complex) at peak exercise and immediately after exercise. In this athlete, echocardiography demonstrated the presence of mild pericardial effusion and CMR was found to be suggestive for myopericarditis.

Troponin and cardiac abnormalities

In total, 10 studies comprising 4171 athletes and study controls reported serum troponin T or I levels. In studies reporting CMR-documented cardiac abnormalities, the weighted mean of elevated troponin levels (according to individual study cut-offs) was 0.65% in athletes and 0% in controls. The weighted mean for elevated troponin levels in echocardiography studies (at time of echocardiography) was 0.25% in athletes and 0% in controls. Moulson *et al* reported an elevated troponin level in 0.9% of the entire cohort (n=2719) but did not report any association with CMR or echocardiography abnormalities. Gervasi *et al* and Clark *et al* did not report troponin levels in study controls. No study reported a clear relationship between elevated troponin levels and cardiac abnormalities on imaging investigations.

DISCUSSION

Our systematic review addressing cardiac abnormalities in athletes recovered from SARS-CoV-2 infection found marked heterogeneity in investigations, study populations and methodology. The majority of abnormal pericardial/myocardial findings were reported by studies performing CMR as part of their investigations. Overall, myocardial abnormalities on CMR such as myocardial LGE and confirmed myocarditis (LLC) was reported in 0%–15% of cases. Pericardial abnormalities such as PE and pericardial LGE were reported in 0%–57% (CMR) of cases. In the total population, myocarditis was reported on echocardiography and/or CMR in 0%–15%, PE in 8%–58% and LGE in 0%–46%. Importantly, our findings also demonstrate that lower quality studies tended to report considerably higher rates of abnormalities (38%–42%). Studies with higher quality assessment scores showed a somewhat greater agreement in proportions of athletes with LGE (0%–5%) after SARS-CoV-2 infection. As such, our review highlights that emergent, small, low-quality studies should be interpreted with caution and a need for prospective, large athlete cohort studies with a low risk of bias.

Clinical implications

This review demonstrates that the incidence of post-SARS-CoV-2 myocardial abnormalities detected on CMR is low and that CMR is the most sensitive diagnostic modality to detect myocardial abnormalities in athletes. As such, using CMR as a screening modality to detect potential SARS-CoV-2 myocardial abnormalities warrants further

investigation. Additionally, while no studies systematically performed extensive arrhythmia monitoring in individuals with clear myocardial abnormalities, these studies also report no clear cases of SCA/SCD. Therefore, until prospective studies have been published, systematic (arrhythmia) monitoring is recommended in athletes with myocardial abnormalities after SARS-CoV-2. Finally, and perhaps most relevant for sports physicians directly involved in RTS in athletes, troponin screenings outside of specific clinical contexts (ie, cardiac complaints or other diagnostic abnormalities) should not occupy a central place in return-to-play strategies.

FU CMR was reported in only one study (Daniels *et al*) and demonstrated complete reversibility in approximately 40% of individuals diagnosed with myocarditis. Two studies (Gervasi *et al* and Cavigli *et al*) investigated arrhythmia outcomes, with only a single athlete demonstrating couplets at peak effort and in early recovery; this athlete was consequently diagnosed with myopericarditis. One study reported a single instance of resuscitated SCA likely unrelated to COVID-19, with no abnormalities on a prior CMR performed after SARS-CoV-2 infection. No study reported a clear relationship between elevated troponin levels and cardiac abnormalities on imaging investigations. Based on the currently available data, the short-term risk for post-COVID-19 SCA/SCD due to arrhythmias caused by myocardial inflammation appears to be poorly investigated, yet reported numbers are low, and the propensity for SARS-CoV-2 pericardial and myocardial involvement variable.

Although physical activity is thought to negatively influence prognosis in athletes with pericarditis and myocarditis,⁵ the rate of myocarditis in athletes after SARS-CoV-2 infection appears low. Overall, athletes were asymptomatic (36%) or had mild symptoms (59%), which may be reflected in the low rates of myocardial injury. FU data in athletes who continue to perform sports after SARS-CoV-2 infection is needed to further elucidate a potential interaction between physical activity and myocardial involvement.

All CMR studies were cross-sectional or retrospective, and only a single study with a small sample size (n=18) performed comparative echocardiography analyses with pre-SARS-CoV-2 infection imaging. While it seems plausible that athletes who fulfil the LLC for myocarditis shortly after infection with SARS-CoV-2 have SARS-CoV-2 myocardial involvement, a recent study in healthcare workers has questioned whether such abnormalities are more common in individuals recovered from SARS-CoV-2 infection when compared with an adequate control group.³⁶ Therefore, caution is advised before attributing (isolated) cardiac abnormalities to SARS-CoV-2. Ideally, to rule out pre-existent abnormalities due to other causes, imaging studies in athletes with pre-existing CMRs should be performed, preferably in well-defined groups as opposed to all comers in imaging centres, with core-lab assessment of all imaging studies. Furthermore,



considering the potential reversibility of pericardial/myocardial abnormalities reported by Daniels *et al*, the extent, and time course and long-term reversibility has yet determined.

Echocardiography findings in our study population did not demonstrate a high prevalence of major cardiac abnormalities, or significant differences compared with in-study controls. Of note, only Brito *et al* reported a high prevalence of pericardial LGE in athletes. Surprisingly, after multimodality adjudication, they demonstrated an association between GLS $<-16\%$ on echocardiography and pericardial/myocardial abnormalities on CMR. First, as GLS is mainly a myocardial parameter, this association hints at possible pericardial/myocardial involvement, not detectable on CMR. Second, while GLS could potentially be a promising tool to identify COVID-19 associated pericardial/myocardial involvement, this finding has not yet been replicated in other studies, and hence such a strategy should not be implemented in routine care for post-COVID-19 athletes. Second, Martinez *et al*³⁵ reported four (of six) cases with abnormal echocardiography (with varying findings) and confirmed pericarditis/myocarditis on CMR. However, they only performed CMRs in athletes with a suspicion of pericardial/myocardial involvement after return-to-play screening, and it is unclear whether athletes without conventional screening abnormalities also had CMR abnormalities. Therefore, according to our findings, screening echocardiography should be used with caution to perform rule-outs for subclinical myocarditis.

Several groups have speculated on causes for SCA/SCD related to COVID-19, such as possible proarrhythmic genetic predisposition or comorbidities that potentially increase the risk of drug-induced and hypoxia-induced ventricular arrhythmias/SCD.^{37 38} Yet, while the studies identified by our systematic review extensively investigated and reported myocardial abnormalities as found using CMR, none of these studies performed a structured investigation of potential arrhythmias. One study (Gervasi *et al*) found no clinically important arrhythmias in 18 football players. However, in addition to the small sample size, this study did not include CMR to document myocardial abnormalities. One study (Moulson *et al*) reported a resuscitated SCA (likely unrelated to SARS-CoV-2) with no abnormal findings on prior CMR after SARS-CoV-2 infection. Hence, these findings cannot readily be extrapolated to athletes with potential or suspected myocardial involvement after COVID-19.

The majority of the studies included in our systematic review determined troponin levels after recovery from COVID-19. Of import, no single paper reported a statistically significant association between elevated serum troponin levels and CMR or echocardiography abnormalities. Therefore, the use of serum troponin after recovery to predict SARS-CoV-2 myocardial involvement remains questionable, with the potential for both false negative and false positive outcomes. According to our findings, troponin testing should not occupy a central

place in return-to-play algorithms, and only be applied in appropriate (expert) clinical settings.

Strengths and limitations

There are several strengths to our study. First, we performed a systematic search through multiple databases, including articles ahead-of-print. Second, data extraction was rigorously performed to report each diagnostic finding. Third, we were able to stratify and weight main findings according to athlete level and recovery location (in-hospital or/and at home). Finally, all studies underwent extensive quality assessment, enabling us to stratify central findings according to study quality.

A number of aspects of our systematic review warrant consideration. First, the majority of the studies included in our review were found to have risk of bias. However, the scarcity of data on this topic in an emerging pandemic makes this a currently unavoidable bias. Second, we did not include case reports in our systematic review, as these included non-athletes with high rates of comorbidities, and because such reports have an intrinsic selection bias. Third, the data across the identified studies was heterogeneous, which limits generalisability. Fourth, the time interval between symptoms onset to cardiac imaging was too variable to allow for a cumulative analysis. Finally, we focused on late complications including late arrhythmias (ie, post-COVID-19) and not on the arrhythmia burden during an active SARS-CoV-2 infection.

CONCLUSION

Our systematic review on athletes recovered from COVID-19 confirms that infection with SARS-CoV-2 is associated with an overall low risk of pericardial/myocardial involvement, arrhythmias and SCA/SCD. Individual studies show a highly variable degree of pericardial and myocardial abnormalities on CMR, which is associated with study quality. CMR appears to be more sensitive to detect (potentially relevant and reversible) post-COVID-19 pericardial/myocardial abnormalities. The extent, time course and potential reversibility of SARS-CoV-2 cardiac abnormalities remain inconclusive. Routine serum troponin screenings after recovery appear unreliable in identifying individuals at risk for such involvement. Prospective studies, preferentially with pre-SARS-CoV-2 imaging (CMR), in larger, well-defined populations, including longer term FU and arrhythmia monitoring, are urgently needed.

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Full search strategy

Database(s): Ovid MEDLINE(R) ALL 1946 to June 02, 2021	
#	Searches
1	*Coronavirus Infections/
2	("covid-19" or "covid 19" or "sars-cov-2" or "sars cov-2" or "sars cov2" or "SARS coronavirus 2").ti,ab,kf.
3	(recover* adj3 ("covid-19" or "covid 19" or "sars-cov-2" or "sars cov-2" or "sars cov2" or "SARS coronavirus 2")).ti,ab.
4	or/1-3
5	exp Athletes/ or exp Adult/ or exp Young Adult/ or exp Middle Aged/ or exp Adolescent/
6	(athlet* or competitive athlet* or elite athlet* or professional* athlet* or adult* or middle age* or adolescent* or ((athlet* or active or healthy or fit) adj3 (population* or community or cohort or competitive or elite or professional* or young))).ti,ab,kf.
7	(sport* adj3 (player* or m?n or wom?n or person*)).ti,ab,kf.
8	(adult* adj3 (young or middel age*)).ti,ab,kf.
9	or/5-8
10	exp Cardiomyopathies/
11	exp Myocarditis/ or exp Pericarditis/ or (myocarditis or pericarditis).ti,ab,kf.
12	exp Death, Sudden, Cardiac/ or ((cardi* or heart) adj3 (arrest or death)).ti,ab,kf.
13	exp Arrhythmias, Cardiac/ or ((heart* or cardi*) adj2 (arrhythmia* or dysrhythmia*)).ti,ab,kf.
14	echocardiography/ or exp Magnetic Resonance Imaging/ or ("cardiac MRI" or cardiac magentic resonance or echocardiogra* or echo-cardiogra*).ti,ab,kf.
15	or/10-14
16	4 and 9 and 15
17	exp Aged/ or exp Child/ or ("aged" or elderly or child*).ti.
18	exp Case Reports/ or exp case study/ or (Case report* or case serie*).pt. or (Case report* or case serie*).ti.
19	or/17-18
20	16 not 19

Database(s): Embase Classic+Embase 1947 to 2021 June 02	
#	Searches
1	exp coronavirinae/
2	exp Coronavirus infection/
3	("covid-19" or "covid 19" or "sars-cov-2" or "sars cov-2" or "sars cov2" or "SARS coronavirus 2").ti,ab,kw.
4	(recover* adj3 ("covid-19" or "covid 19" or "sars-cov-2" or "sars cov-2" or "sars cov2" or "SARS coronavirus 2")).ti,ab.
5	or/1-4
6	exp athlete/ or exp young adult/ or exp adult/ or exp adolescent/ or exp middle aged/
7	(sport* adj3 (player* or m?n or wom?n or person*)).ti,ab,kw.
8	(athlet* or competitive athlet* or elite athlet* or professional* athlet* or adult* or middle age* or adolescent* or ((athlet* or active or healthy or fit) adj3 (population* or community or cohort* or competitive or elite or professional* or young))).ti,ab,kw.
9	(adult* adj3 (young or middel age*)).ti,ab,kw.
10	or/6-9
11	5 and 10
12	exp cardiomyopathy/ or exp virus myocarditis/ or exp myocarditis/ or exp sudden cardiac death/ or exp heart arrhythmia/ or exp Pericarditis/
13	(pericarditis or myocarditis or ((cardi* or heart) adj3 (death or arrest))).ti,ab,kw.
14	((heart* or cardi*) adj2 (arrhythmia* or dysrhythmia*)).ti,ab,kw.
15	exp echocardiography/ or exp cardiovascular magnetic resonance/ or ("cardi* MRI" or cardi* magentic resonance or echocardiogra* or echo-cardiogra*).ti,ab,kw.
16	or/12-15
17	11 and 16
18	exp aged/ or exp child/ or ("aged" or elderly or child*).ti.
19	exp Case Report/ or exp case study/ or (Case report* or case serie*).pt. or (Case report* or case serie*).ti.
20	or/18-19
21	17 not 20

Scopus (2021-06-02)

TITLE-ABS-KEY-AUTH (("covid-19" OR "covid 19" OR "sars-cov-2" OR "sars cov-2" OR "sars cov2" OR "SARS coronavirus 2") OR (recover* W/3 ("covid-19" OR "covid 19" OR "sars-cov-2" OR "sars cov-2" OR "sars cov2" OR "SARS coronavirus 2"))) AND TITLE-ABS-KEY-AUTH (athlete* OR adult* OR "young adult*" OR adolescent* OR "competitive athlet*" OR "elite athlet*" OR "professional* athlet*" OR "middle age*") OR TITLE-ABS ((athlet* OR active OR healthy OR fit) W/3 (competitive OR professional* OR elite OR population* OR community OR cohort)) OR TITLE-ABS (sport* W/3 (player* OR men OR man OR woman OR women OR person)) AND TITLE-ABS-KEY-AUTH (cardiomyopath* OR myocarditis OR pericarditis OR "sudden cardiac death" OR "heart arrest" OR "cardiac arrest" OR "cardiac arrhythmia*" OR "cardiac MRI" OR "cardiac magentic resonance" OR echocardiogra* OR "echo-cardiogra*" OR ((heart* OR cardiac) W/2 (arrhythmia* OR dysrhythmia*))) AND NOT TITLE-ABS-KEY-AUTH ("aged" OR elderly OR child* OR "case report*" OR "case serie*")

Full quality assessment

Table 1. Critical appraisal included cohort studies ('+' = yes, '-' = no, and '=' = not applicable) according to the Joanna Briggs Institute Critical Appraisal tools checklists.

	Clark et al. (2021)	Cavigli et al. (2021)	Daniels et al. (2021)	Hendrickson et al. (2021)	Malek et al. (2021)	Moulson et al. (2021)	Rajpal et al. (2020)	Vago et al. (2020)
1 Similar groups	+	-	-	-	-	-	-	+
2 Exposures similar in different groups	-	-	-	-	-	-	-	-
3 Exposures validly measured	+	-	+	+	+	+	+	+
4 Identification confounding factors	-	-	-	-	-	-	-	-
5 Strategies confounding factors	-	-	-	-	-	-	-	-
6 Participants free of outcome at the start	-	-	-	-	-	-	-	-
7 Outcomes validly measured	-	+	+	+	+	+	+	+
8 Follow-up long enough	-	-	+	-	-	+	-	-
9 Follow-up complete or reasons described	-	-	-	-	-	+	-	-
10 Strategies incomplete follow-up utilized	-	-	-	-	-	-	-	-
11 Appropriate statistical analysis	+	+	+	+	+	+	-	+
<u>Total yes answers:</u>	3	2	4	3	3	5	2	4

Table 2. Critical appraisal included Case control studies ('+' = yes, '-' = no, and '=' = not applicable) according to the Joanna Briggs Institute Critical Appraisal tools checklists.

	Gervasi et al. (2020)
1 Comparable groups	+
2 Cases and controls matched	+
3 Same criteria used for identification	+
4 Exposure measured in a standard, valid way	+
5 Exposure measured in the same way for both groups	+
6 Confounding factors	-
7 Strategies deal with confounding factors	-
8 Outcomes validly measured	+
9 Exposure period long enough	+
10 Appropriate statistical analysis	+
<u>Total yes answers:</u>	8

Table 3. Critical appraisal included Cross sectional studies ('+' = yes, '-' = no, and '=' = not applicable) according to the Joanna Briggs Institute Critical Appraisal tools checklists.

	Brito et al. (2020)	Martinez et al. (2021)
1 Inclusion criteria clearly defined	-	-
2 Study subjects and setting described in detail	-	-
3 Exposure measured in a standard, valid way	+	+
4 Objective, standard criteria used for measurement of the condition?	-	+
5 Confounding factors	-	-
7 Strategies deal with confounding factors	-	-
7 Outcomes validly measured	-	+
8 Appropriate statistical analysis	+	-
Total yes answers:	2	3

Table 4. Critical appraisal included case series ('+' = yes, '-' = no, and '=' = not applicable) according to the Joanna Briggs Institute Critical Appraisal tools checklists.

	Starekova et al. (2021)
1 Inclusion criteria clearly defined	-
2 Condition measured in a standard, valid way	+
3 Valid methods used for identification of the condition	-
4 Consecutive inclusion	+
5 Complete inclusion	+
6 Clear reporting of demographics	+
7 Clear reporting of clinical information	-
8 Outcomes clearly reported	+
9 Clear reporting of presenting clinic demographic information	-
10 Appropriate statistical analysis	+
Total yes answers:	6