Usefulness of machine learning in COVID-19 for the detection and prognosis of cardiovascular complications

Allison Zimmerman¹ and Dinesh Kalra¹,*

¹Division of Cardiology, Rush University Medical Center, Chicago, IL 60612, USA
*Correspondence: dinesh_kalra@rush.edu (Dinesh K. Kalra)

DOI: 10.31083/j.rcm.2020.03.120

Since January 2020, coronavirus disease 2019 (COVID-19) has rapidly become a global concern, and its cardiovascular manifestations have highlighted the need for fast, sensitive and specific tools for early identification and risk stratification. Machine learning is a software solution with the ability to analyze large amounts of data and make predictions without prior programming. When faced with new problems with unique challenges as evident in the COVID-19 pandemic, machine learning can offer solutions that are not apparent on the surface by sifting quickly through massive quantities of data and making associations that may have been missed. Artificial intelligence is a broad term that encompasses different tools, including various types of machine learning and deep learning. Here, we review several cardiovascular applications of machine learning and artificial intelligence and their potential applications to cardiovascular diagnosis, prognosis, and therapy in COVID-19 infection.

Keywords
COVID-19; artificial intelligence; machine learning; cardiovascular

1. Introduction

As the coronavirus disease 2019 (COVID-19) pandemic quickly spread across the globe in the winter and spring of 2020, the cardiovascular (CV) manifestations induced by this viral infection became an important facet of this multisystem disease. Several publications have noted myocarditis, cardiomyopathy, systolic and diastolic dysfunction, pump failure, pericardial involvement cardiac arrhythmias including sudden cardiac death and pulseless electrical activity as possible cardiac complications in patients with COVID-19 (Bin et al., 2016; Guo et al., 2020; Ruan et al., 2020; S. Shi et al., 2020) (Table 1). Historically, it is well known that a higher incidence of cardiac events is also associated with seasonal influenza and other respiratory viral outbreaks, suggesting that acute viral infections can elicit similar cardiac effects via proinflammatory pathways and prothrombotic changes although the intensity and extent of both seems to be higher in COVID-19 (Nguyen et al., 2016). In a study of CV complications of severe acute respiratory syndrome (SARS) in 121 patients, 50.4% developed hypertension in the hospital (P. Zhou et al., 2020). Of these patients, 71.9% developed persistent tachycardia, with 40% exhibiting continued tachycardia at outpatient follow up. Although tachycardic CV complications were common in SARS, they were usually self-limiting and not associated with increased mortality. There is a case report of middle eastern respiratory syndrome–related coronavirus (MERSCoV) acute myocarditis and acute onset heart failure, manifesting as acute myocardial edema and injury of the apical and lateral walls of the left ventricle, but this was an isolated case (Alhogbani, 2016).

Most individuals with COVID-19 present with fever, cough, respiratory symptoms, headache, diarrhea and loss of smell/taste (Guan et al., 2020). Nonetheless, accurately distinguishing between infection by COVID-19 versus other infectious and non-infectious etiologies can be difficult at presentation. Herein, diagnostic algorithms based on constellation of symptoms/signs and initial testing incorporating biomarkers, serology, PCR, chest imaging may assist in increasing diagnostic accuracy so that patients may be quickly diagnosed correctly and triaged expeditiously. This will allow unburdening of busy emergency rooms and reduce pressure on healthcare providers. Artificial intelligence (AI) and machine learning (ML) can potentially be of significant utility in this field. Chest computed tomography (CT) is widely used for the management of COVID-19 pneumonia because of its wide availability and short scan time. AI techniques utilizing CT imaging data have been applied to screen for and assess the severity of COVID-19 pneumonia as well as distinguish it from other forms of pneumonia (F. Shi et al., 2020). For example, a deep learning (DL) based method was developed based on analysis of thoracic CT images that lead to automated detection and monitoring of COVID-19 patients over time (Gozes et al., 2020). Another DL method was able to detect COVID-19 pneumonia and distinguish it from community acquired pneumonia using chest CT (J. Chen et al., 2020). When it comes to CV involvement, just as in pulmonary involvement with COVID-19, there is considerable clinical variability in involvement of the CV system also between different individuals with similar baseline clinical risk features, and even individuals within the same family. This may have to do with host factors, genetics, immune responsiveness, viral load and other yet unidentified disease modifiers. Currently, there are no predictive models that allow physicians to quantify CV risk in patients with COVID-19. As the experience with the COVID-19 pandemic grows, the treatment pathway for cardiac presentations will likely evolve. More advanced approaches involving data collection that can be used to train diagnostic algorithms via ML and AI technologies are currently underway as healthcare professionals struggle with practical difficulties of working in high stress and...
The noise in the training data that the actual dataset does not gener-
ate, the challenge is to perform all these steps efficiently while avoid-
ing over-fitting (occurs when a model learns the detail and
ultimate, the external test can be used to assess its generalizability once optimized. Ulti-
mal, the challenge is to perform all these steps efficiently while avoiding over-fitting (occurs when a model learns the detail and

| Study               | N  | Study Design   | Findings                                                                 |
|---------------------|----|----------------|--------------------------------------------------------------------------|
| Guo et al., 2020    | 187| Case series    | 52 patients suffered acute myocardial injury with a mean CK-MB fraction of 3.34 ng/ml and mean myoglobin of 128.7 μg/ml. 8 patients (4.3%) developed cardiomyopathy with a mean troponin T level of 15.4 pg/ml and mean NT-proBNP of 817.4 pg/ml |
| Arentz et al., 2020 | 21 | Case series    | 7 patients (33%) developed cardiomyopathy                                |
| F. Zhou et al., 2020| 191| Retrospective cohort | 44 patients (23%) developed heart failure. 33 (17%) had acute cardiac injury with 24 patients having a hs-Troponin I level of > 28 pg/ml |
| Punsmann et al., 2020| 100| Prospective cohort | 78% of patients who had recently recovered from COVID had abnormal CMR findings and ongoing inflammation was seen in 60% of patients |
| B. Li et al., 2020  | 1527| Meta-analysis   | 122 patients (8%) showed signs of cardiac injury                         |
| Huang et al., 2020  | 41 | Case series    | 5 patients (12%) were diagnosed with acute myocardial injury with an increased troponin I level > 28 pg/ml |
| Wang et al., 2020   | 138| Case series    | 16.7% of patients developed arrhythmia and 7.2% experienced acute cardiac injury |
| S. Shi et al., 2020 | 416| Prospective cohort | 82 patients (19.7%) developed cardiac injury                             |

Currently, there are a few ongoing clinical trials investigat-
ing COVID-19 and its effects on the CV system using AI (Table 3). An overview of diagnostic testing in patients COVID-19 and CV involvement is provided in Table 4.

2.1 Echocardiography

There are several features of COVID-19 apparent on echocar-
diography that are mainly related to the severity of the disease and its complications. In the early stages following the systemic inflammatory response, there can be evidence of hyperdynamic cardiac function manifested by an increase in cardiac output and left ventricular ejection fraction with or without a decrease in systemic vascular resistance. Other abnormal findings include acute stress-induced cardiomyopathy (Takotsubo) characterized by apical ballooning and left ventricular segmental contraction abnormalities, right ventricular enlargement with associated pulmonary hypertension, and diffuse myocardial inhibition (usually in the later stages resulting from severe hypoxia) (Peng et al., 2020).

Echocardiography is a useful tool quickly identifying a patient’s hemodynamic status, differentiating between different etiologies of shock, and monitoring. It is especially informative and uniquely feasible in critically ill patients, many of whom are proned and on multiple intravenous drips and difficult to transport to CT imaging suites. With the data that can be obtained from clinical echocardiography, there is ample opportunity for the development of echocardiography-based AI platforms. Innovations using AI have the potential to standardize practice, optimize accuracy, and ultimately improve clinical workflow. ML offers the potential to improve the accuracy and reliability of echocardiography by combining clinician interpretation with information derived from ML algorithms and can provide additional predictive information that may be too subtle to be detected by the human eye. High volume data generated from cardiac imaging can be integrated in a multiparametric approach for pattern recognition and imaging data-based disease phenotype characterization (Alsharqi et al., 2018).

The increase in accuracy combined with the time saving ben-
fits demonstrate the value of incorporating ML into the field of echocardiography during this pandemic as well as in general. Currently, there are two studies underway in the United States attempt-

Table 1. Summary of the literature for cardiac acute myocardial injury in COVID19

| Study               | N  | Study Design | Findings                                                                 |
|---------------------|----|--------------|--------------------------------------------------------------------------|
| Guo et al., 2020    | 187| Case series  | 52 patients suffered acute myocardial injury with a mean CK-MB fraction of 3.34 ng/ml and mean myoglobin of 128.7 μg/ml. 8 patients (4.3%) developed cardiomyopathy with a mean troponin T level of 15.4 pg/ml and mean NT-proBNP of 817.4 pg/ml |
| Arentz et al., 2020 | 21 | Case series  | 7 patients (33%) developed cardiomyopathy                                |
| F. Zhou et al., 2020| 191| Retrospective cohort | 44 patients (23%) developed heart failure. 33 (17%) had acute cardiac injury with 24 patients having a hs-Troponin I level of > 28 pg/ml |
| Punsmann et al., 2020| 100| Prospective cohort | 78% of patients who had recently recovered from COVID had abnormal CMR findings and ongoing inflammation was seen in 60% of patients |
| B. Li et al., 2020  | 1527| Meta-analysis | 122 patients (8%) showed signs of cardiac injury                         |
| Huang et al., 2020  | 41 | Case series  | 5 patients (12%) were diagnosed with acute myocardial injury with an increased troponin I level > 28 pg/ml |
| Wang et al., 2020   | 138| Case series  | 16.7% of patients developed arrhythmia and 7.2% experienced acute cardiac injury |
| S. Shi et al., 2020 | 416| Prospective cohort | 82 patients (19.7%) developed cardiac injury                             |
Table 2. Overview of common algorithms used in machine learning

| Algorithm                              | Description                                                                                                                                                                                                 | Illustration |
|----------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------|
| Support vector machine (SVM)           | SVM classifier is constructed by projecting data into higher dimensional space mapping by using kernel functions and devising a new boundary within the new space which can be used for classification, regression, or other tasks such as outlier detection. The disadvantage to SVM is that the classification is dichotomous, and no probability of class membership is given. | ![SVM Illustration](image) |
| k – Nearest neighbor (KNN)             | In KNN, every object being classified to its k, the number of nearest neighbors to include in the estimate of class membership.                                                                                               | ![KNN Illustration](image) |
| Decision trees                         | This algorithm repeatedly splits the data set according to a criterion that maximizes the separation of the data, a process known as recursive partitioning, resulting in a tree-like structure.                                  | ![Decision Tree Illustration](image) |
| Artificial neural networks (ANNs)      | These networks consist of nodes called neurons arranged in a network layout with different layers (input, hidden, and output) that are connected to each other by weighted edges. A neuron in a hidden layer is activated when an input neuron passes a large enough value to trigger the next neuron. Activated neurons continue to pass value to the next layer of neurons until an output is generated upon reaching the final layer. | ![ANN Illustration](image) |
| Regularized regression                 | Allows for introduction of constraints when the number of variables in a system exceeds the number of observations and improve the generalizability the model. Some common forms of regression include LASSO (least absolute shrinkage and selection operator) regression, ridge regression, and elastic net regression. | ![Regularized Regression Illustration](image) |

2.2 Genetics

One of the main aims within the field of genomics is to characterize gene function and identify connections between genotypes and phenotypes, which is crucial for the development of prediction models and the development of precision medicine. DL can be used to make accurate and quick large-scale genomic association studies (Ho et al., 2019).

Human angiotensin converting enzyme (ACE2) is an entry receptor for the SARS-CoV-2 Spiked glycoprotein (Fang et al., 2020; P. Zhou et al., 2020). Fang et al. hypothesized that ACE2 stimulating drugs could worsen clinical outcomes in COVID-19 infections by increasing the expression of ACE2. A recent single cell atlas of the human heart showed that pericytes exhibit high levels of ACE2, which implies that the local inflammation generated during COVID-19 infection of the pericytes may lead to microvascular dysfunction leading to myocardial infarction with nonobstructive coronaries (L. Chen et al., 2020). ACE2 genetic polymorphism has been shown to affect the binding ability of the virus, which could suggest a possible genetic predisposition to COVID-19 infection (Cao et al., 2020). Therefore, it is possible that ML approaches may be able to identify reliable features for using AI techniques to generate predictive models that can forecast COVID-19 related CV complications using echocardiography. A multi-site study lead by the Mayo Clinic is attempting to pioneer an AI platform with EchoGO Core AI software that will be able to automatically analyze images obtained via echocardiography in order to map how the virus attacks the heart (DAIC., 2020). This study will include 500 COVID-19 positive men and women between ages 18 and 89. The objective is the assessment of automated cardiac measurements, ejection fraction, and global longitudinal strain to classify cardiac outcomes in COVID-19 patients. Another study at Johns Hopkins University is currently collecting data from 300 confirmed COVID-19 patients including cardiac specific laboratory markers, continuously obtained vital signs, and imaging data from echocardiography and CT scans to train an ML algorithm (Stempniak, 2020). The goal of the study is to develop a predictive risk score that will be able to predict cardiac events up to 24 hours ahead of time and perform baseline risk stratification on new patients. Given the vast amount and complexity of data associated with echocardiography, AI platforms will need to include many studies that encompass a wide array of clinical characteristics, pathologic features, image quality, and ultrasound vendors.
2.3 Risk assessment

Severe COVID-19 is characterized by rapidly progressive systemic inflammation, sepsis, multiorgan failure, and death. There is often a delay between onset of symptoms and CV manifestations, which presents a challenge when it comes to characterizing the risk for myocardial damage and CV complications in patients with COVID-19. This requires a broad evaluation of numerous variables in order to identify patients at risk that would not be identified with standard statistical analysis. ML is a robust tool that can incorporate nontraditional and unknown risk factors that may be used in CV risk stratification. A few studies have demonstrated a potential role for ML in this regard. An important feature of ML is that it can aid cardiologists in making accurate predictions regarding CV risk in different settings. A study by Kwon et al. developed a DL algorithm that was able to detect in-hospital cardiac arrest and death without attempted resuscitation (Kwon et al., 2018). It performed better than standard methods and had a higher sensitivity and lower false alarm rates. Another study by Alaa et al. (2019) created an ML automated tool based on a dataset of more than 400,000 people with over 450 different variables to risk stratify patients without a history of CV disease. This tool improved CV risk prediction compared to the standard Framingham risk score and discovered new CV risk factors and interactions between different features of an individual. Ongoing efforts have been to develop novel diagnostic tools using ML algorithms.

Table 3. Clinical trials on COVID-19 and the CV system using artificial intelligence

| Study (year) | N   | Primary outcome measures                                                                 | Location                                           | Status  |
|-------------|-----|------------------------------------------------------------------------------------------|----------------------------------------------------|---------|
| **QT-Logs**: a Clinical Study to Monitor Cardiac Safety, With Artificial Intelligence for QT Interval Analysis of ECG Data From Smartwatches, in Patients Receiving Hydroxychloroquine Treatment for COVID-19 (2020) (Assistance Publique Hopitaux De Marseille, 2020) | 100 | Measurement of QTc using an artificial intelligence (AI)-based solution and ECG data collected via smartwatches, compare to standard 12 leads EKG reviewed by cardiologist | Assistance Publique Hôpitaux de Marseille, France | Complete |
| **CARDICoVRISK**: Cardiovascular Risk in COVID-19 Patients: Metabolic, Prothrombotic and Proinflammatory Mechanisms Associated with Outcome and with Cardiorespiratory Features During the Acute Viral Disease and at Short Term Follow-up (2020) (Parati, 2020) | 5500 | To Identify the clinical, immunological, inflammatory, viral, cardiorespiratory consequences of COVID-19 that may persist a few months after discharge and may affect mid- and long-term prognosis | Istituto Auxologico Italiano, Milan, Italy | Not yet recruiting |
| **Coviva**: Coronavirus Disease-19 Survival - The COVIVA Study (Twerenbold and Pfister, 2020) | 1500 | i) perform extensive clinical and biomarker phenotyping in COVID-19 suspects presenting to the emergency department (ED) and in COVID-19 patients with subsequent ICU admission, ii) compare clinical and biomarker profiles of COVID-19 patients with a control group, iii) derive and validate personalized risk prediction models for early clinical decision support, and iv) explore pathophysiological mechanisms including inflammatory and CV pathways | University Hospital Basel, Basel, Switzerland | Recruiting |

risk assessment and classification by using biochemical and clinical data (e.g. ACE2 presence and expression level). ML could also be used to analyze genetic variants from patients with mild to severe COVID-19 infections to risk stratify patients based on their vulnerability or resistance to infection.

Interleukin-6 (IL-6) is another possible molecular target contributing to CV morbidity and mortality in COVID-19 patients. There is experimental evidence supporting its role in atherosclerosis as well as cardiac fibrosis and failure (Brauner et al., 2018; Kusters et al., 2018; van der Heijden et al., 2018; Wang et al., 2019). There is also evidence demonstrating a possible role in plaque formation and with the development and progression of abdominal aortic aneurysm (Huber et al., 1999; Nishihara et al., 2017). Additionally, genetic variants with increased circulating levels of IL-6 receptors (and therefore reduced levels of IL-6) have been shown to protect against coronary artery disease (CAD) (Sarwar et al., 2012). Other cytokine moieties involved in COVID-19 related cytokine release syndrome may also contribute to myocardial tissue damage and are potential targets for large scale data analysis. These include pathways mediated by granulocyte-macrophage colony-stimulating factor (GM-CSF), tumor necrosis factor α (TNF-α), interleukin-17 (IL-17), interleukin 18 (IL-18), and interferon γ (IFN-γ) (Guzik et al., 2020).
Table 4. Diagnostic tests in patients with COVID-19 and cardiovascular involvement

| Test                | Diagnostic considerations in COVID-19 patients |
|---------------------|------------------------------------------------|
| NT-pro BNP/BNP      | Frequently elevated in patients with COVID-19 even in the absence of clinical heart failure or elevated filling pressures and associated with unfavorable course among patients with ARDS. Should not trigger evaluation for heart failure unless there is clinical evidence per ACC guidelines. |
| Troponin            | May be helpful in risk assessment in patients requiring ICU level care and can be an indicator of myocardial injury. Should be considered if the diagnosis of MI is being considered on clinical grounds per ACC guidelines. |
| Procalcitonin       | A marker of bacterial infection. Reports from Wuhan demonstrated an association between elevation in its levels with cardiac injury and requiring ICU care |
| IL-6                | High concentrations associated with worse outcome |
| Complete blood count| Acute cardiac injury and higher mortality are associated with lymphopenia and thrombocytopenia |
| EKG                 | Dynamic ischemic appearing changes when there is myocardial involvement |
| Echocardiography    | May show hyperdynamic changes in the early stages, and global or regional systolic and diastolic dysfunction later on. Stress cardiomyopathy like wall motion abnormalities and pericardial effusion may occur. |
| CMR                 | Patients often show lower LVEF, higher left ventricular volumes, higher left ventricular mass, raised native T1 and T2 values, late gadolinium enhancement (LGE). Likely useful in monitoring the long-term CV consequences of COVID-19 |
| Cardiac CT          | May be helpful in the setting of elevated troponin when CAD needs to be ruled out noninvasively |

that can identify patients infected with COVID-19. For example, ML based screening of COVID-19 assay designs using a CRISPR-based system for detecting the virus showed a higher sensitivity and speed (Metsky et al., 2020). Currently, there are no approaches available to predict CV outcomes in COVID-19 patients. There is a prospective cohort study in process that is examining data to develop a predictive model in northern Italy, which is one of the regions in the world most affected by COVID-19 (Parati, 2020). This study is enrolling 5500 COVID-19 patients and will be using ML techniques to develop multivariate models for risk stratification. They will be collecting bio-humoral and imaging data to explore the prognostic and pathophysiological role of immunologic factors, activation of coagulation, endothelial dysfunction, inflammatory response, genetic, hormonal, and metabolic factors and acute cardiac damage. The study is expected to complete in September 2020.

2.4 Electrophysiology

The contribution of COVID-19 to the development of arrhythmias is well described but specific mechanisms are still being elucidated. However, most patients with COVID-19 or in whom COVID-19 is suspected will have a baseline electrocardiogram (EKG) performed at the time of entry into the healthcare system, particularly those with severe disease or those in whom QT prolonging medications may be used (Gandhi et al., 2020). AI-enhanced, cost-effectively acquired EKG may allow for opportunities to screen for disorders not typically associated with ECG (Lopez-Jimenez et al., 2020). One of the major aims for AI technologies applied to electrophysiologic data is to improve population health via interpretable insight. This is based on the idea that ECGs often contain subtleties that may not always be readily interpretable by humans (Attia et al., 2016). As an example, one study reported that an AI enabled ECG alone was able to identify low EF with a high degree of accuracy (Attia et al., 2019). It is possible that other conditions may be similarly identified by AI enabled ECGs, which would allow for improved risk stratification at a low cost. Additionally, with the use of consumer smartphone or smartwatch enabled ECG devices, we may be able to offer more cost-effective screening at a general population level, especially for individuals infected with SARS-CoV-2 but with milder disease severity not requiring hospitalization and inpatient telemetry monitoring. However, the expertise in reading ECGs is limited to healthcare settings. Preliminary data suggests that AI techniques may improve interpretation of these strips and facilitate appropriate triaging to those needed to be seen by a physician (Hagiwara et al., 2018; Tison et al., 2018). There is a prospective cohort trial currently taking place in France that is evaluating a new method for remote monitoring of corrected QT measurements using AI based solution and ECG data collected via smartwatches (AI-QTc) in patients being treated with hydroxychloroquine (Assistance Publique Hopitaux De Marseille., 2020). AI-QTc data will be compared with the standard manual QTc reviewed by a cardiologist.

2.5 Advanced imaging

There are studies examining the role of AI to detect COVID-19 on nuclear imaging with chest CT, but there has been only one study to date looking at advanced cardiac imaging in patients with COVID-19 infection identified from a local testing center who underwent voluntary evaluation for cardiac involvement with cardiac MRI (CMR). They demonstrated cardiac involvement in 78% of patients and ongoing myocardial inflammation in 60% with recent COVID-19 disease, independent of pre-existing comorbid conditions, severity of acute illness, and time since original diagnosis. There was no significant trend toward reduction of imaging
or serologic findings during the recovery period. These findings suggest an indication of considerable inflammatory disease burden and urgently require confirmation in a larger cohort.

DL methods have shown promising performance for automated calculation of left ventricular function. One study used a convolutional neural network on a dataset of 596 CMR examinations acquired in different institutions and on scanners from different vendors to create a tool that outperformed manual segmentation, and its accuracy increased with the heterogeneity of the patients that were included (Tao et al., 2019). ML can also be used for automated segmentation of the heart. This is important because segmentation of the left ventricle from the epicardium and endocardium is important for the assessment of CV function. A study by Ngo et al. (2017) used ML for automated segmentation of the heart with a dataset of 45 cardiac cine magnetic resonance with ischemic and nonischemic cardiomyopathy, left ventricular hypertrophy, and normal LV function. The accuracy was comparable to standard methods.

Evaluation of myocardial perfusion can be performed using single-photon emission computed tomography (SPECT). ML has been used in the past to improve diagnostic performance and combine imaging modalities to maximize discriminatory capability. In one instance, an SVM approach was utilized to localize the mitral valve during SPECT acquisition, which is important for the assessment of myocardial perfusion. 392 SPECT scans were used for training and validation, and the ML model exhibited an AUC (area under the curve) of 82 for the detection of obstructive CAD in comparison to two experts (AUC of 0.79 and 0.81) and unadjusted mitral valve plane (AUC 0.63) (Betancur et al., 2017). Based on their study results, the authors suggested that AI could lead to integration of clinical and imaging data and create personalized MACE risk calculations in patients undergoing SPECT MPI. Another large multi-center study that included 1,638 patients applied DL directly to nuclear cardiology polar map images (Betancur et al., 2018). In this study, they demonstrated that DL could outperform the clinical standard (total perfusion deficit) for the identification of obstructive CAD. An AI-driven algorithm has been incorporated into an FDA approved nuclear imaging software that uses a CDS tool and natural language for automated report generation. The system includes over 230 rules of perfusion, reversibility, function, and additional information such as prone vs. supine positioning (Garcia et al., 2018). There was found to be no significant difference between the AI driven structured report and the experts’ impressions of CAD or ischemia. ML has also been used to assess coronary artery dysfunction using obstructive emission tomography (PET) in combination with coronary computed tomography angiography (CCTA) with an algorithm that combined quantitative stenosis, plaque burden, and myocardial mass (Dey et al., 2015). This algorithm generated a risk score to predict impaired myocardial flow reserve quantified by PET and showed that the ability to discriminate coronary dysfunction was superior with the ML derived risk score with an AUC of 0.83 compared to an AUC of 0.66 with quantitative stenosis.

2.6 Biohumoral exams

CV comorbidity seems to be correlated with impaired outcomes in COVID-19 patients. A direct causal relationship or pathophysiological mechanism and the clinical value of established and emerging biomarkers remains unknown. S. Shi et al. (2020) demonstrated that approximately 20% of patients diagnosed with COVID-19 showed signs of cardiac injury as measured by high-sensitivity troponin I (hs-TNI) levels. Additionally, these patients presented with higher levels of C-reactive protein (CRP), NT-proBNP, procollatonin, leukocytes, and creatinine. Whether this occurs as a result of underlying CV disease that confers risk in the setting of hypoxia and viral sepsis vs. direct viral myocarditis is unknown. Furthermore, ethnic and racial variation will bring new challenges to the US in terms of diagnosis and management of patients with COVID-19. The Chinese experience with COVID-19 patients has shown high comorbidity with hypertension, diabetes, and cerebrovascular disease (Huang et al., 2020). Shi et al. noted common comorbidities associated with cardiac injury including hypertension, diabetes, CAD, heart failure, and cerebrovascular disease — all more common in African American patients. Subclinical CV disease tends to occur earlier in African American patients, and they have higher circulating biomarkers of systemic inflammation (Hackler et al., 2019; Lefferts et al., 2017). As such, African Americans in the US may be a particularly vulnerable group to the effects of COVID-19. Pathophysiological mechanisms, differences in mild and severe variants of COVID-19, and clinical factors including ethnicity and underlying comorbidities can be studied extensively using ML techniques. An early and reliable personalized risk prediction represents a major unmet clinical need as it could allow evidence based clinical decision aid for effective resource allocation during the COVID-19 pandemic. Currently, there is a prospective case-control study underway in Switzerland aiming to deliver a platform using ML techniques to develop personalized risk prediction models (Twerenbold and Pfister, 2020). The study will enroll 1500 patients and will be looking at patient clinical phenotyping (e.g. comorbidities, medications, symptoms, vitals, ECG, and imaging data) and extended laboratory analysis and blood sampling. The study will be complete in June 2021.

3. Conclusions

The applications of AI, including ML, have the capability to exploit data-rich platforms and transform approaches to diagnose, risk stratify, prevent, and treat CV disease. There have been advances in AI over the years within different areas of cardiology, and the extrapolation of this technology towards the evaluation of COVID-19 patients would also prove useful. Advances with AI will require close collaboration amongst computer scientists, clinicians and investigators in order to identify the most appropriate and relevant problems to be solved and the best approaches to do so.

Abbreviations

COVID-19: Coronavirus disease 2019; AI: Artificial intelligence; ML: Machine learning; DL: Deep learning; ACE2: Angiotensin converting enzyme; IL-6: Interleukin 6; GM-CSF: Granulocyte-macrophage colony stimulating factor; TNF α: Tumor necrosis factor α; IL-17: Interleukin 17; IL-18: Interleukin 18; IFN-γ: Interferon

Authors' contributions

Both authors wrote and contributed to the article.
Acknowledgments

We would like to express our gratitude to all those who helped
me during the writing of this manuscript.

Conflict of interest

The authors declare no conflicts of interest statement.

Submitted: June 28, 2020
Revised: September 06, 2020
Accepted: September 08, 2020
Published: September 30, 2020

References

Ali, S. J., Anouchke, K., Singh, G., Slomka, P. J., Kolli, K. K., Ku-
mar, A., Pandey, M., Malialak, G., van Rosendael, A. R., Beeey, A.
N., Berman, D. S., Leipsic, J., Nieman, K., Andreini, D., Pontone, G.,
Schoepf, U. J., Shaw, L. J., Chang, H., Narula, J., Bax, J. J., Guran,
Y. and Min, J. K. (2019) Clinical applications of machine learning in
cardiovascular disease and its relevance to cardiac imaging. European
Heart Journal 40, 1975-1986.

Ala, A. M., Bolton, T., Di Angelantonio, E., Rudd, J. H. F. and van der
Schaar, M. (2019) Cardiovascular disease risk prediction using auto-
460
mated machine learning: A prospective study of 423,604 UK Biobank
participants. Plos one 14, e0213653.

Alharghani, T. (2016) Acute myocarditis associated with novel Middle East
respiratory syndrome coronavirus. Annals of Saudi Medicine 36, 78-80.

Alsharqi, M., Woodward, W. J., Mumith, J. A., Markham, D. C., Upton,
and Lee, P. (2018) Artificial intelligence and echocardiography. Echo
Research and Practice 161, R115-R125.

Arentz, M., Yim, E., Klaff, L., Lohmandwala, S., Riedo, F. X., Chong, M.
and Lee, P. (2020) Characteristics and outcomes of 21 Critically Ill
patients with COVID-19 in Washington State. JAMA 323, 1612.

Assistance Publique Hopitaux De Marseille. (2020) QT-Logs: Artifi-
cial Intelligence in patient receiving treatment for COVID-19 (QT-Logs).
Available at: https://clinicaltrials.gov/ct2/show/NCT04371744 (Accessed:
June 4, 2020).

Attia, Z. I., Electrical and Computer Engineering, Ben-Gurion Univer-
sity of the Negev, Beer Sheva, Israel, DeSimone, C. V., Dillon, J. J., Sapir,
Y., Somers, V. K., Digan, J. L., Bruce, C. J., Ackerman, M. J., Asir-
vatham, S. J., Strieter, B. L., Bakrytj, J., Scott, C. G., Bennett, K.
E., Ladewig, D. J., Giles, E. J., Sadot, D., Geva, A. B. and Friedman,
P. A. (2016) Novel bloodless potassium determination using a signal-
processed single-lead ECG. Journal of the American Heart Association
5, e002746.

Attia, Z. I., Kapa, S., Lopez-Jimenez, F., McKie, P. M., Ladewig, D. J.,
Satam, G., Pellikka, P. A., Enriquez-Sarano, M., Noseworthy, P. A.,
Munger, T. M., Asirvatham, S. J., Scott, C. G., Carter, R. E. and Fried-
man, P. A. (2019) Screening for cardiac contractile dysfunction using an
artificial intelligence–enabled electrocardiogram. Nature Medicine
25, 70-74.

Betancur, J., Commandeur, F., Motlaff, M., Sharir, T., Einstein, A. J.,
Bokhari, S., Fish, M. B., Ruddly, T. D., Kaufmann, P., Sinunas, A. J.,
Miller, E. J., Bateman, T. M., Durbala, S., Di Carli, M., Germano, G.,
Otaki, Y., Tamarapooro, B. K., Dey, D., Berman, D. S. and Slomka,
P. Y. (2018) Deep learning for prediction of obstructive disease from
fast myocardial perfusion SPECT. JACC: Cardiovascular Imaging 11,
1654-1663.

Betancur, J., Rubeau, M., Fuchs, T. A., Otaki, Y., Aronso, Y., Slipczuk,
L., Benz, D. C., Germano, G., Dey, D., Lin, C., Berman, D. S.,
Kaufmann, P. A. and Slomka, P. J. (2017) Automatic valve plane local-
ization in myocardial perfusion SPECT/CT by machine learning: anatomic and clinical validation. Journal of Nuclear Medicine 58, 961-
967.

Bin, S. Y., Heo, J. Y., Song, M., Lee, J., Kim, E., Park, S., Kwon, H., Kim,
S. M., Kim, Y., Si, Y., Lee, I., Baek, Y. H., Choi, W., Min, J., Jeong,
H. W. and Choi, Y. K. (2016) Environmental Contamination and Viral

Shedding in MERS Patients During MERS-CoV Outbreak in South
Korea. Clinical Infectious Diseases 62, 755-760.

Brauner, S., Jiang, X., Thiruvenkata, G. E., Lundberg, A. M., Ostberg, T.
Yan, Z., Kuchroo, V. K., Hansson, G. K. and Wahren-Herlenius, M.
(2018) Augmented Th17 differentiation in Trim21 deficiency promotes
a stable phenotype of atherosclerotic plaques with high collagen con-
tent. Cardiovascular Research 114, 158-167.

Cao, Y., Li, L., Feng, Z., Wan, S., Huang, P., Sun, X., Wen, F., Huang,
X., Ning, G. and Wang, W. (2020) Comparative genetic analysis of
the novel coronavirus (2019-nCoV/SARS-CoV-2) receptor ACE2 in
different populations. Cell Discovery 6, 11.

Chen, L., Wu, L., Zhang, J., Zhang, L., Dong, D., Zhao, Y., Su, S., Wang,
Y., Hu, X., Zheng, B., Zhang, K., Wu, H., Dong, Z., Xu, Y., Zhub,
Y., Chen, X., Yu, L. and Yu, H. (2020) Deep learning-based model for
detecting 2019 novel coronavirus pneumonia on high-resolution com-
puted tomography: a prospective study. medRxiv (in press).

Chen, L., Li, X., Chen, M., Feng, Y. and Xiong, C. (2020) The ACE2
expression in human heart indicates new potential mechanism of heart
injury among patients infected with SARS-CoV-2. Cardiovascular Re-
search 116, 1097-1100.

DAIC. (2020) Mayo clinic uses artificial intelligence to help assess
cardiac danger from COVID-19. Diagnostic and Interventional
Cardiology. Available at: https://www.dicardiology.com/content/
mayo-clinic-uses-artificial-intelligence-help-assess-cardiac-
danger-covid-19 (Accessed: May 19, 2020).

Dey, D., Diaz Zamudio, M., Schubbaeck, A., Juarez Orozco, L. E., Otaki,
Y., Gransnar, H., D., Germano, G., Achenbach, S., Berman, D. S.,
Manchester, A., Alexanderson, E. and Slomka, P. J. (2015) Relationship
between quantitative adverse plaque features from coronary computed
angiography and downstream impaired myocardial flow reserve by 13N-ammonia positron emission tomography: a pilot study. Circulation.
Cardiovascular Imaging 8, e003255.

Fang, L., Karakulakis, G. and Roth, M. (2020) Are patients with hyperten-
sion and diabetes mellitus at increased risk for COVID-19 infection?
The Lancet Respiratory Medicine 8, 221.

Gandhi, R. T., Lynch, J. B. and del Rio, C. (2020) Mild or moderate Covid-
19. New England Journal of Medicine (in press).

Garcia, V. E., Klein, J. L., Moncayo, V., Cooke, C. D., Del’Aune, C., Folks,
R., Moreiras, L. V. and Esteves, F. (2018) Diagnostic performance of
an artificial intelligence-driven cardiac-structured reporting system for
myocardial perfusion SPECT imaging. Journal of Nuclear Cardiology
(in press).

Gozes, O., Frid-Adar, M., Greenspan, H., Patrick, D., Zhang, H., Ji, W.,
Berrheim, A. and Siegel, E. (2020) Rapid AI development cycle for the Coronavirus (COVID-19) pandemic: initial results for automated
detection & patient monitoring using deep learning CT image analysis.
arXiv (in press).

Guan, W., Ni, Z., Hu, Y., Liang, W., Ou, C., He, J., Liu, L., Shan, H., Lei,
C., Hui, D. S. C., et al. (2020) Clinical characteristics of Coronavirus
disease 2019 in China. New England Journal of Medicine 382, 1708-
1720.

Guo, T., Fan, Y., Chen, M., Wu, X., Zhang, L., He, T., Wang, H., Wan, J.,
Wang, X. and Lu, Z. (2020) Cardiovascular implications of fatal out-
comes of patients with Coronavirus disease 2019 (COVID-19). JAMA Cardiology 5, 811.

Guzik, T. J., Mohiddin, S. A., Dimarco, A., Patel, V., Savvatis, K.,
Marelli-Berg, F. M., Madhur, M. S., Tomaszewski, M., Maffia, P.,
D’Acquisto, F., Nicklin, S. A., Marian, A. J., Nosalski, R., Murray,
E. C., Guzik, B., Berry, C., Touyz, R. M., Kreutz, R., Wang, D. W.,
Blenn, D., Sagliocco, O., Creo, F., Thomson, E. C. and McElmes, I.
(2020) COVID-19 and the cardiovascular system: implications for risk
assessment, diagnosis, and treatment options. Cardiovascular Re-
search 116, 1666-1687.

Hacker, E., Lew, J., Gore, M. O., Ayers, C. R., Department of Clinical Sci-
ces UT Southwestern Medical Center Dallas TX, Atzler, D., Khera,
A., Rohatgi, A., Lewis, A., Neeland, I., Omland, T., University of Oslo
Norway and de Lemos, J. A. (2019) Racial differences in cardiovas-
cular biomarkers in the general population. Journal of the American
Heart Association 8, e012729.

Hagiwara, Y., Fujita, H., Oh, S. L., Tan, J. H., Tan, R. S., Ciaccio, E. J. and
Acharya, U. R. (2018) Computer-aided diagnosis of atrial fibrillation based on ECG Signals: A review. Information Sciences 467, 99-114.

Ho, D. S. W., Schnidering, W., Wake, M., Saffery, R. and O’Sullivan, J. (2019) Machine learning SNP based prediction for precision medicine. Frontiers in Genetics 10, 267.

Huang, C., Wang, Y., Li, X., Ren, L., Zhao, J., Hu, Y., Zhang, L., Fan, G., Xu, J., Gu, X., Cheng, Z., Yu, T., Xie, J., Wei, Y., Wu, W., Xie, X., Yin, W., Li, H., Liu, M., Xiao, Y., Gao, H., Guo, L., Xie, J., Wang, G., Jiang, R., Gao, Z., Jin, Q., Wang, J. and Cao, B. (2020) Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. The Lancet 395, 497-506.

Huber, S. A., Sakkinen, P., Conze, D., Hardin, D. and Tracy, R. (1999) Interleukin-6 Exacerbates Early Atherosclerosis in Mice. Arteriosclerosis, Thrombosis, and Vascular Biology 19, 2364-2367.

Johnson, K. W., Torres Soto, J., Glicksberg, B. S., Shaper, K., Miotti, R., Ali, M., Ashley, E. and Dudley, J. T. (2018) Artificial Intelligence in Cardiology. Journal of the American College of Cardiology 71, 2688-2679.

Kusters, P. J. H., Lutgens, E. and Seijkens, T. T. P. (2018) Exploring immune checkpoints as potential therapeutic targets in atherosclerosis. Cardiovascular Research 114, 368-377.

Kwon, J., Lee, Y., Lee, Y. and Park, J. (2018) An algorithm based on deep learning for predicting in-hospital cardiac arrest. Journal of the American Heart Association 7, e008678.

Lefferts, W. K., Augustine, J. A., Spartan, N. L., Atallah-Yunes, N. H., Heffernan, S. and Gump, B. B. (2017) Racial differences in aortic stiffness in children. The Journal of Pediatrics 180, 66-67.

Li, B., Yang, J., Zhao, F., Zhi, L., Wang, X., Liu, L., Bi, Z. and Zhao, Y. (2020) Prevalence and impact of cardiovascular metabolic diseases on COVID-19 in China. Clinical Research in Cardiology 109, 531-538.

Li, L., Qin, X., Z., Yin, Y., Wang, X., Kong, B., Bai, J., Li, Y., Yang, Z., Song, Q., Cao, K., Liu, D., Wang, G., Xu, Q., Fang, X., Zhang, S., Xia, J. and Xia, J. (2020) Using artificial intelligence to detect COVID-19 and community-acquired pneumonia based on pulmonary CT: evaluation of the diagnostic accuracy. Radiology 296, E65-E71.

Lopez-Imenez, F., Attia, Z., Arruda-Olson, A. M., Carrier, R., Chareonthaitawee, P., Jouni, H., Kapa, S., Lerman, A., Luong, C., Medina-Inojosa, J. R., Noseworthy, P. A., Pellika, P. A., Redfield, M. M., Roger, V. L., Sandhu, G. S., Senecal, C. and Friedman, P. A. (2020) Artificial intelligence in cardiology: present and future. Mayo Clinic Proceedings 95, 1015-1039.

Metsky, H. C., Freije, C. A., Kosoko-Thordssen, T.-S. F., Sabeti, P. C. and Myhrvold, C. (2020) CRISPR-based surveillance for COVID-19. bioRxiv.

Neri, E., Miele, V., Coppola, F. and Grassi, R. (2020) Use of CT and magnetic resonance imaging in patients recently recovered from Coronavirus disease 2019 (COVID-19). JAMA Cardiology (in press).

Pan, S., Yang, R., Wang, W., Jiang, L. and Song, J. (2020) Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients from Wuhan, China. Intensive Care Medicine 46, 846-848.

Peng, Q., Chinese Critical Care Ultrasound Study Group (CCCUSG), Wang, X. and Zhang, L. (2020) Using echocardiography to guide the treatment of novel coronavirus pneumonia. Critical Care 24, 143.

Puntmann, V. O., Carej, M. L., Wieters, I., Fahlin, M., Arendt, C., Hoffmann, J., shchendryagina, A., Escher, F., Vasa-Nicotera, M., Zeisher, A. M., Vehreschild, M. and Nagel, E. (2020) Outcomes of cardiovascular magnetic resonance imaging in patients recently recovered from Coronavirus disease 2019 (COVID-19). JAMA Cardiology (in press).

Ruan, Q., Yang, K., Wang, W., Jiang, L. and Song, J. (2020) Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients from Wuhan, China. Intensive Care Medicine 46, 846-848.