AI can now identify atrial fibrillation through sinus rhythm
Hendriks, Jeroen; Fabritz, Larissa

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Comment

AI can now identify atrial fibrillation through sinus rhythm

Atrial fibrillation is a substantial health-care challenge and is considered to be a global pandemic, as prevalence rates have increased greatly1 and atrial fibrillation-related hospitalisations outnumber those of major cardiac conditions such as heart failure and myocardial infarction.2 Atrial fibrillation confers an increased risk of stroke and mortality; it therefore needs to be detected not only to manage the arrhythmia but also to prevent comorbidities and death.1 A 10-second, 12-lead electrocardiograph (ECG) in current clinical practice is unlikely to reveal possible atrial fibrillation if not present in this short monitoring time. Silent or undetected atrial fibrillation is common and the few screening methods available are demanding in terms of time and resources. Continuous monitoring by means of loop recorders is often indicated, particularly in case of embolic stroke of undetermined source (ESUS).4 Novel and user-friendly wearables to identify arrhythmias have emerged with recent digital advances: wearable ECG technology using automated photoplethysmography algorithms have shown feasible and accurate cardiac rhythm detection and can aid in monitoring the dynamic burden of time spent in atrial fibrillation,5 while mobile atrial fibrillation applications are available for patients and health-care professionals for education and guidance in management.6

In The Lancet, Zachi Attia and colleagues7 report a study in which they aimed to develop and validate an artificial intelligence (AI)-enabled ECG using a trained neural network to detect the electrocardiographic signature of atrial fibrillation during sinus rhythm. Structural changes in the atria predispose to atrial arrhythmias.8 Deducting atrial fibrillation in a sinus rhythm ECG has been attempted previously by using P wave and PR interval traces to describe phenomena such as interatrial block.9 Here, Attia and colleagues hypothesised that the signature of atrial fibrillation due to the structural changes in the atria could be identified by a trained network, using a standard 10-second, 12-lead ECG recorded during sinus rhythm. Rather than trying to observe atrial fibrillation by prolonged monitoring of sinus rhythm, the authors suggest that AI can avoid this needle-in-a-haystack scenario and instead identify from as few as one normal sinus rhythm the absence of atrial fibrillation. Rather than trying to observe atrial fibrillation by prolonged monitoring of sinus rhythm, the authors suggest that AI can avoid this needle-in-a-haystack scenario and instead identify from as few as one normal sinus rhythm the absence of atrial fibrillation.

AI can now identify atrial fibrillation through sinus rhythm. Structural changes in the atria predispose to atrial arrhythmias.8 Deducting atrial fibrillation in a sinus rhythm ECG has been attempted previously by using P wave and PR interval traces to describe phenomena such as interatrial block.9 Here, Attia and colleagues hypothesised that the signature of atrial fibrillation due to the structural changes in the atria could be identified by a trained network, using a standard 10-second, 12-lead ECG recorded during sinus rhythm. Rather than trying to observe atrial fibrillation by prolonged monitoring of sinus rhythm, the authors suggest that AI can avoid this needle-in-a-haystack scenario and instead identify from as few as one normal sinus rhythm ECG if there is indeed a needle hidden within. P wave characteristics are likely to be picked up by the network, but no criteria are predefined or revealed in retrospect. In total, almost 650 000 ECGs from a cohort of 180 922 patients aged 18 years or older with at least one normal sinus rhythm, standard 10-second, 12-lead ECG from the Mayo Clinic ECG laboratory, were used to develop, test, and validate the network. Patients and their digitally available ECGs were randomly assigned to three datasets: a training dataset (70% of the patient cohort) used to train the network, an internal validation dataset (10% of the patient cohort) to optimise the network, and a testing dataset (20% of the patient cohort) to identify the ability of the AI-enabled ECG to detect atrial fibrillation. When using a single AI-enabled ECG, mathematical performance of the network showed an impressive area under the curve of the operating receiver curve of 0·87 (95% CI
In summary, Atta and colleagues are to be congratulated for their innovative approach and the thorough development and local validation of the AI-enabled ECG. Given that AI algorithms have recently reached cardiologist level in diagnostic performance, this AI-ECG interpretation is ground-breaking in creating an algorithm to reveal the likelihood of atrial fibrillation in ECGs showing sinus rhythm.

Comment

Jeroen M L Hendriks, Larissa Fabritz

Centre for Heart Rhythm Disorders, South Australian Health and Medical Research Institute, University of Adelaide and Royal Adelaide Hospital, Adelaide, SA 5000, Australia (JMLH); Institute of Cardiovascular Science, University of Birmingham, Birmingham, UK (LF); and Department of Cardiology, University Hospital Birmingham, Birmingham, UK (LF)

jeroen.hendriks@adelaide.edu.au

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