Echocardiographic screening for the anomalous aortic origin of coronary arteries

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ABSTRACT

Aims We sought to determine the diagnostic performance, clinical profiles and outcomes of anomalous aortic origin of coronary arteries (AAOCA) using a standardised echocardiographic approach in young adults and athletes.

Methods In 2015–2019, we screened 5998 outpatients (age 16 years (Q1–Q3: 11, 36)), referred for routine echocardiography, using four specific echocardiographic windows: parasternal short/long axis and apical 4/5-chambers view. Coronary CT confirmed AAOCA. For the performance analysis, 300 coronary-CT scans were available; two independent and double-blinded physicians retrospectively reviewed echocardiographic images.

Results A total of 47 AAOCA was diagnosed; the overall prevalence was 0.0078%. Over 5 years, we found a significant increment of AAOCA diagnostic rate (P for trend = 0.002). Syncope (n=17/47) and palpitations (n=6/47) were prevalent symptoms. All patients suspended sports activity at the diagnosis. Twenty-seven patients underwent surgery, while 20 underwent a conservative medical treatment. All patients are alive at a median follow-up of 3±1.6 years; only surgical repairs restarted their activity. Our method showed better sensitivity than traditional short-axis evaluation: 93% vs 83%, p=0.0030 (AUC 0.96 (95% CI 0.92, 0.99) and AUC 0.89 (95% CI 0.83, 0.95), respectively), with a good interobserver agreement (95%, k=0.83, p<0.001).

Conclusions The application of a standardised echocardiographic approach for AAOCA detection led to a significantly increased rate of identified anomalies. This approach demonstrated higher sensitivity than the traditional echocardiographic assessment. Implementing this protocol in clinical practice may help improve the AAOCA diagnosis in young adults and athletes.

Trial registration number NCT04224090.

INTRODUCTION

The anomalous aortic origin of coronary arteries (AAOCA) is a rare congenital condition, accounting for abnormalities of origin, course, destination, size and number of the coronary vessels.1 The prevalence of the AAOCA is unclear; it ranges from 0.021% to 5.79%, depending on the imaging techniques used for the diagnosis: coronary angiography, cardiac magnetic resonance (CMR), CT or echocardiography.1 Though, autopsy-based study and case series of patients that experienced sudden cardiac death (SCD) due to AAOCA generally present higher frequency than imaging ones.1,2

The diagnosis and the screening of AAOCA can be difficult.3 The ECG is often normal or presents unspecific signs.4 Therefore, a high diagnostic suspicion is required to refer children and adolescents to invasive and ionising examinations (CT or angiography).5 In this context, echocardiography has always been considered the method of screening and diagnosis with the most limited accuracy. However, this examination is the first-line test recommended and prescribed by physicians when a congenital cardiac abnormality is suspected.6

To overcome these limitations, we designed, studied and implemented a specific echocardiographic 4-views-based approach.
to diagnose AAOCA. Therefore, our study aimed to assess the diagnostic performance, clinical profiles and outcomes of AAOCA using this standardised echocardiographic approach in young adults and athletes.

METHODS

Study population and design
The present study is a single-centre retrospective analysis. We tested the diagnostic performance of our echocardiographic-based approach for AAOCA detection and examined the clinical profiles and outcomes of the AAOCA diagnosed within these examinations. A comprehensive flow chart is presented in figure 1.

In 2015–2019, 5998 outpatients (16 years old (Q1–Q3: 11, 36)), both young adults and athletes, underwent routine echocardiography in our department due to the mandatory cardiovascular screening programme for the eligibility assessment to non-competitive or competitive sports activity.

The echocardiographic-based approach for AAOCA detection was systematically implemented in the department since January 2015. For our purposes, we considered only patients undergoing their first examination and thus excluded bicuspid aortic valves, known or newly discovered, already known structural or congenital heart disease, cardiomyopathies and all their longitudinal follow-up echocardiographic examinations.

All the suspected AAOCA (N=57) were referred for a coronary-CT confirmation. For the performance analysis, 243 consecutive additional patients were also asked to undergo a coronary-CT scan. A total of 300 coronary-CT scans was achieved. Confirmed AAOCA were then followed-up for a median time of 3±1.6 years.

In agreement with the Declaration of Helsinki, all patients enrolled gave written informed consent at the time of their evaluation, stating that data and images may be subsequently used for research purposes. For participants under 18 years old, a parent and/or the legal tutor gave informed consent.

Since the observational nature of the study, no ethical committee approval was required.

Anomalous aortic origin of coronary arteries definition
Coronary arteries are traditionally defined as normal when arising from the appropriate sinus of valsalva (right or left), presented a proper course (the right coronary artery (RCA) in the right atrioventricular groove; the left anterior descending (LAD) in the anterior interventricular groove, and the left circumflex (LCX) in the left atrioventricular groove) and terminated in the capillary bed.7,8

For the present analysis, AAOCA were defined as anatomic variants differing from the anatomy mentioned above. In particular, the anomalous origin of the right coronary artery (AORCA) when RCA originated from the left coronary sinus of the aorta, or the non-coronary sinus, and the anomalous origin of the left coronary artery (AOLCA) when LAD or LCX originated from the right coronary sinus of the aorta or the non-coronary sinus. AAOCA, both AORCA and AOLCA, were also considered the coronary arteries arising from the aorta by a separate ostium, shared or common ostium, or as branch vessels. Besides, AAOCA were also defined as the coronary arteries that originated directly from the ascending aorta instead of valsalva’s sinus.8,9

The aortic coronary arteries from the pulmonary artery (ACAPA) were also identified and defined as the coronary artery with an origin from the pulmonary artery: ARCAPA for the right coronary artery and ALCAPA for the left one.3,10

Echocardiography
All the echocardiographic studies were performed using a commercially available Philips iE33 ultrasound systems (Philips Medical Systems, Amsterdam, The Netherlands), using a 2.5 or 3.5 MHz transducer, as appropriate. Images were acquired at a frame rate of 70–80 frames/s, applying a standard two-dimensional greyscale. Images of three consecutive heart cycles were collected from the parasternal short axis (PSAX) and long-axis (PLAX), and the 4/5-chambers apical views.10,11

A standardised approach for the assessment of AAOCA
Our method consisted of the visualisation of the coronary arteries from four focused-specific views: PSAX, the traditional approach,3,11 along with the PLAX, and the apical 4-chambers and 5-chambers view (figure 2).3,12 Both two-dimensional echocardiography and color-Doppler imaging were used to visualise the coronary arteries. Since the coronaries usually have low-velocity flow, the Nyquist limit was decreased to 20–40 cm/s, as recommended.11

A hallmark of AAOCA was considered the absence of a proper visualisation of the origin of the coronary arteries from the aortic root (figure 3, panels A and B) at the PSAX.3,12 In addition, their origin and the proximal course can be evaluated from the apical 5-chambers view, focused again on the aortic root (figure 3, panel E).
We also evaluated the presence of the so-called ring sign; if present, it can be visualised from the PLAX (figure 3, panel D) and should be interpreted as atypical.\textsuperscript{11,13} This sign is of interest when an aberrant left or right AAOCA is suspected. A coronary artery originating from the contralateral aortic sinus, along its intramural course, crosses the mid-anterior line of the circular aorta to reach the sinus of valsalva of its intended origin. Consequently, its cross-section should be visible in an orthogonal view traversing the mid-long axis of the aorta, namely, the parasternal long-axis view.\textsuperscript{3,12,13}

In normal conditions, by tilting the transducer posteriorly from the PLAX, and focusing on the RVOT, it is possible to image the left main and the left anterior descending (figure 3, panel C). An AOLCA with the LAD originating from the right sinus of valsalva should be suspected from this view in the presence of LAD and
RCA displayed contemporarily (figure 3, panel C). Besides, its absence can be pathognomonic LAD agenesis or hypoplasia.\textsuperscript{11}

From the apical 4-chambers view, if a coronary artery is visualised running through the left atrioventricular groove (figure 3, panel F), instead of the right atrioventricular one (figure 2, panel D), a coronary retro aortic course could be suspected in both cases: LCX or LAD from the right coronary sinus.\textsuperscript{6,11}

Usually, ACAPA are rare and do not present with a predictable pattern imaged with specific echocardiographic views.\textsuperscript{3,11} For ACAPA, the absence of one coronary artery from the aortic root and its appearance at the pulmonary artery level is very suggestive and used for this analysis.\textsuperscript{3,11}

Coronary CT
All coronary-CT examinations were obtained using a 384 (192\times2)-slices third-generation scanner (SOMATOM Force CT; Siemens Healthineers, Forchheim, Germany). Images were acquired at the isotropic resolution of 0.5–0.75 mm in a 25 cm field of view. Patients received a non-ionic low-osmolality contrast agent (iodixanol; GE Healthcare Life Sciences, Chalfont, UK), with a volume of 30–120 mL (1–1.5 mL/kg) and flow injection rate of 3–5 mL/s, followed by saline injection administered intravenously using a power injector at same volume and velocity.\textsuperscript{5} A bolus triggering was used to scan initiation. The scans were acquired with a 70–120 kV of tube voltage with an auto-bolus triggering was used to scan initiation. The scans were available and were considered as the gold standard echocardiographic images and re-examined them in a double-blinded fashion from the coronary-CT examinations and diagnosis, ignoring the aim of this re-examination. For the performance analysis, 300 coronary-CT scans were available and were considered as the gold standard.

Separately for each patient, the physicians reviewed both the short-axis view, focused on the aortic root (two-dimensional images only and images with colour, the traditional approach for AAOCA detection)\textsuperscript{1}\textsuperscript{6} and our proposed method (based on the visualisation of four standardised echocardiographic views); then, were asked to decide if the coronary anatomy can be considered as usual or not. The physicians can access the full echocardiographic examination, except the specific 4-views, while examining the traditional approach. A QLab software (Philips Medical Systems, Amsterdam, The Netherlands) was used for the offline retrospective analysis.

Performance analysis
According to our internal policy, echocardiographic and CT images are stored and available in our Institutes’ Core Lab for further analysis. Two independent physicians (MP and ACB), both experts in echocardiography and AAOCA, retrospectively reviewed the 5998 echocardiographic images and re-examined them in a double-blinded fashion from the coronary-CT examinations and diagnosis, ignoring the aim of this re-examination. For the performance analysis, 300 coronary-CT scans were available and were considered as the gold standard.

RESULTS
General characteristics
In our cohort, coronary abnormalities were suspected in 62 patients using echocardiography. Of them, five were diagnosed as ARCAPA or ALCAPA; the echocardiograms were considered as exhaustive, without the need for further investigations. The latter underwent surgery for coronary repair. Besides, 57 patients underwent a coronary-CT scan for the diagnosis confirmation of AAOCA. These examinations discovered 10 false positives, while a total of 47 AAOCA were confirmed.

The overall prevalence of AAOCA was 0.0078%, with 0.0015% of incidence. The 22.8% (n=13/47) AAOLCA, while the 59.6% (n=34/47) were AAORCA. The RCA from the left coronary sinus with an inter-arterial course was the most frequent AAOCa 56.1% (N=37). Regarding AAOLCA, the LCX from the right coronary sinus with a retro-aortic course (8.8%, N=5) was the most frequent.

Echocardiographic screening results
After the implementation in 2015 of our echocardiographic-based method of diagnosis, we found a progressive increment of the number of the AAOCa identified (P for trend=0.002), growing progressively through the years 2015–2019 (table 1). A rise not accompanied by an increased number of misdiagnosis (P for trend=0.20). Of note, we observed that our method determined a significant progression of the AAOCA diagnosis: both AOLCA...
We acknowledged 10 misdiagnoses of the total 57 suspected AAOCA that underwent the coronary-CT scan for confirmation. Most of them (N=5/10) was due to the para-commissural coronary position to the aortic valve (p=0.029). The overlap with nearby structure at the echocardiography and the presence of a separate conal ramus were also found to be secondary causes of echocardiographic images misinterpretation.

### Clinical profiles and outcomes

In our cohort of AAOCA, the male sex was prevalent (85.1%, n=40/47), while 65.9% (n=31) of participants and AORCA (P for trend=0.007 and P for trend 0.042, respectively).

Data are categorical and described as counts (proportions).

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Total (N=5998)</th>
<th>2015 (N=1065)</th>
<th>2016 (N=1165)</th>
<th>2017 (N=1171)</th>
<th>2018 (N=1562)</th>
<th>2019 (N=1035)</th>
<th>P for trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>AAOCA, n (%)</td>
<td>47 (0.8 %)</td>
<td>3 (0.3 %)</td>
<td>6 (0.5 %)</td>
<td>10 (0.9 %)</td>
<td>12 (0.8 %)</td>
<td>16 (1.5%)</td>
<td>0.002</td>
</tr>
<tr>
<td>Misdiagnosis, n (%)</td>
<td>10 (0.2 %)</td>
<td>3 (0.3 %)</td>
<td>2 (0.2 %)</td>
<td>2 (0.2 %)</td>
<td>3 (0.3 %)</td>
<td>0 (0.0 %)</td>
<td>0.20</td>
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</table>

<table>
<thead>
<tr>
<th>Types of AAOCA</th>
<th></th>
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</tr>
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<tbody>
<tr>
<td>AOLCA, n (%)</td>
<td>13 (0.2 %)</td>
<td>0 (0.0 %)</td>
<td>1 (0.1 %)</td>
<td>3 (0.3 %)</td>
<td>3 (0.2 %)</td>
<td>6 (0.6 %)</td>
<td>0.007</td>
</tr>
<tr>
<td>AORCA, n (%)</td>
<td>34 (0.6 %)</td>
<td>3 (0.3 %)</td>
<td>5 (0.4 %)</td>
<td>7 (0.6 %)</td>
<td>9 (0.6 %)</td>
<td>10 (1.0 %)</td>
<td>0.042</td>
</tr>
</tbody>
</table>

| Table 2 Clinical profiles and demographic characteristics of AAOCA diagnosed in 2015–2019 |
|----------------------------------------|----------------------------------------|----------------------------------------|----------------------------------------|----------------------------------------|----------------------------------------|----------------------------------------|
| AAOCA AORCA AOLCA                      | N=47 N=34 N=13                        | N=47 N=34 N=13                        | N=47 N=34 N=13                        | N=47 N=34 N=13                        | N=47 N=34 N=13                        | N=47 N=34 N=13                        |
| Demographic characteristics            | Age, years 18 (14, 36) 22 (15, 36) 14 (7, 37) | Male sex 40 (85.1%) 28 (82.4%) 12 (92.3%) | Competitive athletes 31 (65.9%) 22 (64.7%) 9 (69.2%) | Sports practiced                        | Athletics 7 (14.9%) 5 (14.7%) 2 (15.4%) | Basket 3 (6.4%) 2 (5.9%) 1 (7.7%) | Cycling 5 (10.6%) 2 (5.9%) 3 (23.1%) | Dance 1 (2.1%) 1 (2.9%) 0 (0.0%) | Gym 6 (12.8%) 4 (11.8%) 2 (15.4%) | Handball 2 (4.3%) 2 (5.9%) 0 (0.0%) | Running 5 (10.6%) 5 (14.7%) 0 (0.0%) | Soccer 9 (19.1%) 5 (14.7%) 4 (30.8%) | Swimming 5 (10.6%) 4 (11.8%) 1 (7.7%) | Tennis 3 (6.4%) 3 (8.8%) 0 (0.0%) | Volleyball 1 (2.1%) 1 (2.9%) 0 (0.0%) | Symptoms                        |
| Asymptomatic                           | 11 (23.4%) 11 (32.4%) 0 (0.0%)          | Arrhythmias                           | 3 (6.4%) 2 (5.9%) 1 (7.7%)            | CP                                      | 5 (10.6%) 3 (8.8%) 2 (15.4%)         | CP and arrhythmias                    | 5 (10.6%) 4 (11.8%) 1 (7.7%)       | Palpitations                        | 6 (12.8%) 4 (11.8%) 2 (15.4%)       | Syncope                           | 17 (36.2%) 10 (29.4%) 7 (53.8%)       | AAOCA, anomalous aortic origin of coronary arteries; AOLCA, anomalous origin of the left coronary artery; AORCA, anomalous origin of the right coronary artery; CP, chest pain.
Table 3  Diagnostic performance of PSAX-view and four-views echocardiographic diagnostic approach for AAOCA

<table>
<thead>
<tr>
<th>Echocardiographic methods</th>
<th>AUC</th>
<th>95% CI</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>LR+</th>
<th>LR-</th>
<th>PPV</th>
<th>NPV</th>
</tr>
</thead>
<tbody>
<tr>
<td>PSAX-view approach</td>
<td>0.89</td>
<td>(0.83, 0.95)</td>
<td>83%</td>
<td>96%</td>
<td>5</td>
<td>0.17</td>
<td>79.5</td>
<td>96.6</td>
</tr>
<tr>
<td>Four-views approach</td>
<td>0.96</td>
<td>(0.92, 0.99)</td>
<td>93%</td>
<td>99%</td>
<td>20</td>
<td>0.06</td>
<td>95.6</td>
<td>98.7</td>
</tr>
</tbody>
</table>

Results of receiver operating characteristic (ROC) curves analysis. AAOCA, anomalous aortic origin of coronary arteries; AUC, area under the curve; LR, likelihood ratio; NPV, negative predictive value; PPV, positive predictive value.

practised competitive sports (table 2). AORCA were older than AOLCA (22 years old vs 14 years old). The majority of AORCA was asymptomatic (n=11, 32.4%). Syncope (n=17/47) and palpitations (n=6/47) were prevalent symptoms in both AORCA and AOLCA.

All patients suspended sports activity at the diagnosis. Twenty-seven patients underwent surgical repair when surgery was applicable due to high-risk anatomy,14 while 20 underwent conservative medical treatment, following currently available guidelines.15 Regarding the surgery techniques, the majority underwent coronary unroofing (n=22/27); three patients experienced coronary reimplantation, one patient a lateral coronary translocation (n=22/27); three patients experienced coronary reimplantation, one patient a lateral coronary translocation and one the Vouhé’s osteoplasty.

To the best of our knowledge, no SCD were reported among the 5998 outpatients that we screened. Regarding the AAOCA that we detected, all patients are alive at a median follow-up of 3±1.6 years. Sport restriction was more frequent in AOLCA than AORCA: 8 (23%) vs 5 (38%), p<0.001 (χ² test). Only patients that underwent surgical repair restarted the sports activity. Besides, a participant developed ischaemic heart disease and underwent percutaneous coronary stenting of the LCX (AAOCA: LCX retro-aortic curse). The latter was not operated because not currently indicated from guidelines.14

Diagnostic performance analysis

The detailed results are shown in table 3. Compared with the traditional echocardiographic approach for the AAOCA detection, centred on the PSAX coronary evaluation and color-Doppler evaluation, the 4-views approach, that we adopted, determined a significant increment of the echocardiographic sensitivity in our examinations: 93% vs 83%, p=0.0030 (AUC 0.96, 95% CI 0.92, 0.99 and AUC 0.89, 95% CI 0.83, 0.95, respectively). When adding the adjunctive echocardiographic views, the positive predictive value of the echocardiographic evaluation augmented from 79.5 up to 96.6, resulting in more conclusive than traditional (positive likelihood ratio >20 and negative <0.1).

Finally, we found a good interobserver agreement between physicians for the adjudicated double-blinded retrospective diagnosis of AAOCA (95%, k=0.83, p<0.001).

DISCUSSION

In the present study, whereas implementing our echocardiographic-based approach for coronary abnormalities detection, we observed a progressive and statistically significant increment in AAOCA diagnosed through 2015–2019. This increment determined an augmentation of the AAOCA identified with the echocardiography. In this cohort, AOLCA were younger and more symptomatic than AORCA; syncope was prevalent, while, after the AAOCA diagnosis, only patients that underwent surgical repair restarted sports activity. No significant events were recorded during a median follow-up of 3±1.6 years. Our method showed better sensitivity than the traditional PSAX evaluation of the origin and proximal course of the coronary arteries, with an excellent interobserver agreement between physicians at the retrospective analysis.

Traditionally, echocardiography is considered the imaging technique with the most limited accuracy for the coronary arteries evaluation.1 Despite this belief, echocardiography is the first-line imaging examination that physicians recommend in young patients and athletes when cardiac abnormalities are suspected.16 The echocardiographic assessment of AAOCA is generally done from the PSAX, focused on the aortic root, trying to visualise the origin and the proximal course of the coronary arteries.1 Other acoustic windows are possible, along with a color-Doppler evaluation and probe tilts to off-axis views,11 but they are less implemented in everyday practice, excluding specialised centres in paediatric cardiology and congenital heart diseases.6 The present study adds to previous literature by demonstrating that a standardised and routinely performed approach can ameliorate the sensitivity of the sonographic evaluation of the AAOCA.

The approach that we tested, except the PLAX, focused on the RVOT, is generally part of the routine echocardiographic evaluations.3,10 While the 4-views approach we proposed can be regularly acquired, both by physicians and sonographers. Since this method does not require advanced skills but only little practice (in our experience, no more than 2 or 3 weeks), it can be systematically implemented; these images can be stored, re-examined or sent to experts for a second look, at any time, without altering or interfering with their sensitivity and specificity. Besides, even if in young patients it is infrequent to find poor acoustic windows, it can easily overcome this limitation due to its multiview approach.

Since AAOCA can be associated with SCD, particularly during efforts,6 the diagnosis should be pursued in particular populations and contexts.2,8,9 According
to the previous literature, we found that AOLCA were generally younger and more symptomatic than AORCA; a history of syncpe is frequent, while chest pain and palpitations are less common. On the contrary, AORCA are often asymptomatic, and, accordingly, the age of the diagnosis is usually higher.

Some authors deny the association mentioned above, while others consider this condition as a possible predisposition to accelerated atherosclerosis, but autopsy-based series generally report a relatively more significant number of SCD occurred in young patients and athletes with AAOCA, both AORCA and AOLCA. Consequently, implementing this simple echocardiographic 4-views approach in everyday practice, as we demonstrated, could increase the number of diagnoses within a second level examination, even in subjects asymptomatic, and therefore reduce the amount of some AAOCA-related SCD during sports activities.

Limitations and strengths
One of the principal limitations of this study is that it comprises only a single-centre experience in the AAOCA echocardiographic evaluation. Our analysis’s second limitation is that we do not know the exact amount of the overall false negative, but only 303 cases. By the way, we assume that this number has been little. According to our policy, we usually prescribe a second examination within 8–12 months from the first evaluation, even if the first echocardiogram is considered negative for cardiac abnormalities. So, these double-check between operators and evaluations can reduce misdiagnoses. We believe that some missed AAOCA would have been reported during these years in which we implemented our protocol, but, to date, no missed AAOCA have been reported at these second examinations.

Moreover, other centres across the world usually adopt multiviews and color-Doppler approaches for the AAOCA detection; so, our study suffers from a lack of novelty in this view, but these approaches are not encoded, studied and standardised yet.

Clinical perspectives
AAOCA echocardiographic diagnosis is challenging, and imaging protocols have not been standardised yet. Unlike other approaches, we opted for a standardised method and thus reduced the need for peculiar individual skills and expertise.

This protocol can be easily implemented in sonographic laboratories; the images can be stored and available for further evaluations, especially in the view of telemedicine support, if the examinations are carried out in peripheral centres or sports medicine laboratories. Besides, the method we proposed is not time-consuming since it involves standard echocardiographic windows.

CONCLUSIONS
The application of a new echocardiographic-based method for AAOCA detection led to a significantly increased rate of identified anomalies and AAOLCA. This approach demonstrated better sensitivity than the traditional echocardiographic assessment, with excellent reliability among physicians. Implementing this protocol in clinical practice may reduce AAOCA-related sudden cardiac death in young adults and athletes.

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