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© 2024 Zaki, Albaroudi, Shaban, Alkahlout, Yigit, Elnabawy, Basharat, Almarri and Azad. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms. Comparison between transthoracic echocardiography and transoesophageal echocardiography in the diagnosis of acute aortic dissection from an emergency perspective. A systematic review and meta-analysis

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**Background:** Acute aortic dissection (AAD) is a life-threatening medical condition with high early fatality. Therefore, a prompt and precise diagnosis, which can be achieved through invasive and non-invasive techniques is vital. Echocardiography, unlike MRI and CT, is accessible in emergency units and bedside-compatible. The recommended echocardiographic techniques for AAD are transthoracic and transoesophageal echocardiography (TTE and TOE). Therefore, our review compares their diagnostic roles in AAD.

**Methods:** Studies relevant to our topic were attained through a database search and manual scrutiny of references lists of articles obtained from the electronic databases. The Quality Assessment of Diagnostic Accuracy Studies tool (QUADAS-2) has been used for quality assessment. All quantitative analyses were performed using either STATA 16 or Comprehensive Meta-Analyst software.

**Results:** The search strategy yielded 1,798 articles, of which only 11 were eligible for inclusion. Our subgroup analysis showed that conventional TTE had a sensitivity and specificity of 85.35% and 84.51% for the diagnosis of Stanford type A AAD and was 45.89% sensitive and 87.05% specific for the diagnosis of type B AAD. However, the subgroup analysis shows that contrast-enhancement of TTE results in a sensitivity and specificity of 93.30% and 97.60% for diagnosis of type A AAD, and 83.60% and 94.50% for diagnosis of type B AAD, respectively. On the other hand, conventional TOE was 93.64% sensitive and 95.50% specific for the diagnosis of type B AAD. Moreover, our analyses show that TTE has pooled false negative and positive rates of 28.6% and 18.6%, while TOE has shown false negative and positive rates of 2.4% and 4.3%, respectively.

**Conclusion:** TOE is the more favorable diagnostic tool for AAD diagnosis than TTE. However, it cannot be used as a stand-alone diagnostic tool since misdiagnosis cases are being reported. Contrast-enhanced TTE can also diagnose AAD since it provides similar results to conventional TOE.

KEYWORDS

aortic dissection, emergency medicine, meta-analysis, point-of-care systems, ultrasound, echocardiography, sensitivity, specificity

# Introduction

Acute aortic dissection (AAD) is the most prevalent lifethreatening condition that impairs the aorta, with a mortality rate that ranges from 1% to 1.4% per hour when untreated (1, 2). The death rate is generally substantial in type A dissection (involving the ascending aorta), with reports suggesting nearly 58% mortality rate in patients without treatment and 26% in those with surgical therapy (3). On the other hand, a lower mortality rate is observed in the type B dissection, with reports showing mortality rates of about 31% in patients who underwent surgery and 11% for those medically treated (3). Given these high mortality rates, accurate and early diagnosis of AAD is essential in reducing morbidity and mortality in these patients.

In the early 1960s, aortography was considered the standard tool for diagnosing AD; however, recent technological advancements have led to the use of noninvasive imaging modalities such as echocardiography, magnetic resonance imaging (MRI), and computed tomography (CT). The European Society of Cardiology and the European Association of echocardiography have suggested that transthoracic echocardiography (TTE), which involves the placement of the ultrasound transducer on the chest, be used as the first-line imaging test for patients with suspected AAD especially when the aortic dissection risk score (ADD-RS) (score ranges of 0 to 3, of which 0 is classified as low risk, 1 moderate risk and 2-3 is high risk) shows a low probability for AAD (4, 5). However, TTE is restricted in patients exhibiting abnormal chest wall configuration, obesity, pulmonary emphysema, and mechanically ventilated patients (4). Therefore, transoesophageal echocardiography (TOE), which involves inserting the transducer in the esophagus, has been used to overcome these limitations (4, 6). Previous research studies have shown that TOE has almost similar sensitivity and specificity as CT and MRI; however, due to its lower availability compared to CT and patient discomfort, it is only used as the first-line imaging test in about 30%-40% of the AAD cases (3, 7, 8).

To the best of our knowledge, there is no systematic review comparing TTE to TOE regarding the diagnosis of AAD. Therefore, the current review will compare the diagnostic performance of TOE and TTE in diagnosing AAD and form a basis for emergency medicine care.

# Methodology

### Protocol & registration

This systematic review and meta-analysis was conducted in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) 2020 guiding principles and protocol registered on PROSPERO article (CRD42023404905).

# Eligibility criteria

Before any study was included in the current review, the following criteria had to be met;

- 1. Studies designed as either observational (prospective, retrospective, or cohort) or randomized trials published in English. This criterion was fundamental to our analysis since we did not want to have direct translations of scientific terms that could undermine our research.
- 2. Studies that evaluated any classification of AAD (either type A or type B in the Stanford classification or type I, II, or III in the DeBakey classification). Type A dissection is defined as dissection proximal to the brachiocephalic artery and involves the ascending aorta, while type B AD originates from the left subclavian artery and involves only the descending aorta. On the other hand, type I AD originates in the ascending aorta to the ascending aorta, and type III originates and is limited to the ascending aorta, and type III starts in the descending aorta and extends distally above or below the diaphragm.
- 3. Studies that evaluated either TOE or TTE in the AAD diagnosis.

On the other hand, the reviewers ensured that studies that met the following criteria were excluded from the current review;

- 1. Studies designed as either abstracts without full articles, systematic reviews, case reports or series, diagnostic algorithms, or letters to the editor.
- 2. Studies that only evaluated other imaging techniques (CT and MRI) and aortography in diagnosing AAD.
- 3. Studies that only included patients with Chronic or Asymptomatic AD.

- 4. Studies with insufficient sample size (less than 30 participants). This criterion was used in this review to help improve the statistical power of our analyses.
- 5. Studies that evaluated patients with the acute aortic syndrome or AD in general but did not distinguish the data for AAD.

#### Literature search

Five electronic databases-Embase, PubMed, ScienceDirect, Scopus, and Google scholar-were scoured for scientific journals published in English. The reviewers created a search strategy that involved combining scientific terms using the Boolean expressions "AND" and "OR". The strategy employed was as follows; ("Transoesophageal echocardiography" OR "transesophageal echocardiography" OR "TOE" OR "TEE") AND ("transthoracic echocardiography" OR "TTE") AND ("Acute aortic dissection" OR "acute Type A aortic dissection" OR "acute Type B aortic dissecting"). Additional studies were obtained by going through the reference lists of relevant studies from the databases mentioned earlier. To avoid undermining our scientific research, grey literature, and close duplicates were not retrieved.

## Quality assessment

The current study was designed as a diagnostic review; therefore, the Quality Assessment of Diagnostic Accuracy Studies tool (QUADAS-2) in the Review Manager software (RevMan 5.4.1) was employed to assess the risk of bias. This tool usually consists of the Assessment of bias category, subdivided into four domains (selection, index test, reference standard, and flow and timing) and applicability concerns category, which consists of three fields, including patient selection, index test, and reference standard. In the quality assessment, low risk of bias was assigned a green color, while high risk and unclear risk of bias were assigned red and yellow, respectively.

### Data extraction and definitions

The data extraction process was carried out by two reviewers who later compiled the data in a tabular form. The data retrieved was as follows; Author ID (first author's surname and the publishment year), patient characteristics (patients enrolled and patients with AAD in the final diagnosis), study design, study location, and reference test, and the main outcomes. The primary outcome of the current review was to compare the sensitivity and specificity of TTE to TOE, while the secondary outcome was to compare misdiagnosis between the two techniques. All the discrepancies in the collected data were initially resolved through interactive discussion between the two reviewers and if a consensus could not be reached a third reviewer was consulted.

According to accepted conventions, AAD represents patients with the sudden onset of symptoms within 48 h, while subacute represents patients with symptoms occurring for at least two weeks. On the other hand, chronic AD represents patients whose symptoms have occurred for more than two weeks. However, for this study, we considered AAD to be patients with symptoms occurring between 1 and 14 days.

### Data synthesis

The diagnostic accuracy analysis was done using STATA 16, while the rate of misdiagnosis was carried out using Comprehensive Meta-analyst software. In these analyses, we employed the randoms effect model to counter the expected heterogeneity, calculated using the I<sup>2</sup> statistics. The heterogeneity values between 0%-40%, 41%-60%, and 61%-100% were considered low, moderate, and substantial, respectively. All the results related to diagnostic accuracy (specificity and sensitivity) were presented in the form of forest plots. To further enhance our analysis, we subdivided the outcomes based on the type of AAD. The Stanford classification was used, of which the DeBarkley classifications I and II were considered type A AAD, and classification III was considered type B AAD. On the other hand, the results of misdiagnosis, including false negatives and false positives, were recorded in a tabular form. In all these analyses, a 95% confidence interval was applied.

# Results

## Study selection

We applied the search strategy mentioned earlier in the five electronic databases and attained 1,798 articles related to our topic. The articles were then screened for duplicates, of which 815 close (articles with nearly identical content i.e., multiple versions of the same study with minor differences) or exact (articles with identical content) duplicates were excluded. Afterward, the remaining 983 articles were screened by looking at their titles and abstracts, of which 896 were excluded. Out of the 87 remaining articles, we did not retrieve 44 because they were either abstracts without full evidence, letters to the editor, Guidelines, case reports, systematic reviews, or ongoing trials. Finally, 11 articles met the criteria for inclusion while the other 32 articles were excluded as follows; 10 evaluated other imaging techniques in AAD diagnosis, 8 were published in different languages, 2 only included patients with chronic or asymptomatic AD, and 12 included patients with acute aortic syndrome or AD in general but did not distinguish data for AAD. A summarized literature selection criterion is shown in the PRISMA diagram below (Figure 1).

### Summary of study characteristics

Out of the 11 studies, 8 were carried out in single centers (9–16), while the other 3 were conducted in multiple centers (17–19). In these studies, a total of 3,046 patients suspected of AAD were enrolled, of which a majority were men (65% (1995/3,046). Out of the 2,418 patients suspected of AAD in 10 studies, AAD was confirmed in 775 (32%) patients during the final diagnosis, of which type A AAD was confirmed in 661 patient and type B AAD was



confirmed in 114 patients (Table 1). The Study by Moore and colleagues (19) did not provide the data for patients with AAD in the final diagnosis; therefore, it was not used to calculated the ratio of patients with AAD in the final diagnosis.

# Quality assessment results

The risk of bias summary is presented in Figure 2 below. From the summary, it is evident that all the studies satisfy at least four of the seven assessment criteria, meaning that they had a low risk of bias. Considering the patient selection criteria all the studies had a low risk of bias and low concern except for one study which used a non-consecutive patient selection and included about 50% with an already established suspicion for AAD. The index and reference test assessment criteria also showed low risk and low concern; however, it is worth noting that in most studies it was unclear whether the reference test was conducted without knowing the results of the index test. On the other hand, with regard to flow and timing, all studies except two showed a high risk of bias. The high risk of bias was because these studies evaluated more than one reference test.

Main outcomes			Overall, TTE was 90% sensitive and 96% specific for the diagnosis of CAAAD. Of the 9 false negatives recorded in the TTE studies, 6 were patients with CAAAD.	The presence of at least one sign (indirect or direct) in the TTE studies had a sensitivity and specificity of 88% and 56% for the diagnosis of TAAAD.	TTE detected the initial dissection flap in the ascending aorta in 159 patients, while the intraoperative inspection detected TAAAD in 171 patients.	Of the 24 patients who underwent TTE studies, 50% (12/24) were true positives, and 50% (12/24) were false negatives	Overall specificity and sensitivity for AAD diagnosis using TTE were 71.2% and 73.7%, respectively, while for TOE were 94.2% and 97.3%, respectively. TTE showed a sensitivity and specificity of 82.2% and 89.2% for diagnosing Ascending aorta diasection, 79.1% and 88.2% for diagnosing aortic arch diagnosis of descending aorta dissection, and 65.6.3% and 81.8% for the diagnosis of descending aorta dissection, 90.7% and 95.6% and 96.4% for the diagnosis of dissection, 90.7% and 98.8% for the diagnosis of descending aorta dissection.	TOE had a sensitivity of 100% for AD diagnosis, while TTE was 82.7% sensitive. TOE had a false-positive result in 7 patients, thus resulting in a specificity of 68.2% TTE showed a sensitivity and specificity of 85% and 78.1% for the diagnosis of type A AD, while TOE had a sensitivity and specificity of 100% and 78.8%, respectively. TTE showed a sensitivity and specificity of
Reference	test		Surgical inspection, autopsy, CT, or MRI.	TOE and CTA.	CT and actual surgical findings	Chest x-ray, CT, and Coronary angiography	CT, MRI, and surgical findings	MRI and surgical findings.
Device used			Sonos 5,500 ultrasound system with an S3 probe and harmonic imaging.	MyLab30 Gold and HD7 with 2–5 MHz phased array probe.	Vivid I (GE) or CX 50 (Phillips) ultrasound system with an S3 3.5–5 MHz transducer.	NR	GE Vivid 1 or Vivid 7 with a 1.5–4.3 MHZ transducer using harmonic imaging in TTE studies and multiplanar probe with a 2.9–8 MHz transducer using harmonic imaging in TOE studies	2.25-3.5 MHz transducers attached to V3400 R CV60, HP77065 or HP Sonos 1,000 sector scanner
Index	test		T'TE	TTE	TTE	TTE	TOE and TOE	TOE and TOE
	AAD in osis	Type B ( <i>n</i> )	1	~	1	9		11
ics	ts with <i>i</i> al diagne	Type A ( <i>n</i> )	62	50	171	43	45	20
racterist	Patien fini	Total ( <i>n</i> )	62	57	171	49	26	31
ient cha	olled	Mean age (years)	66 ± 15	NR	58.69 ± 12.61	54.8 ± −9.2	61 (49- 74)	52 ± 15
Pat	its enr	М	191/ 79	155/ 126	131/ 47	41/8	36	35/ 18
	Patier	Sample ( <i>n</i> )	270	281	178	49	128	53
Location			Italy	Italy	Poland	Greece	Spain	Germany
Study design			Retrospective study	Prospective single-center cohort study	Retrospective single-center study	Retrospective single-center study	Retrospective single-center study	Prospective single-center study
Author ID			Cecconi et al. (11)	Nazerian et al. (12)	Sobczyk et al. (13)	Asouhidou et al. (14)	Evangelista et al. (15)	Nienaber et al. (16)

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e Main outcomes			55.5% and 100% for the diagnosis of type B AD while TOE had a sensitivity and specificity of 90.9% and 97.6%, respectively.	The standard TOE diagnosed aortic dissection with a sensitivity and specificity of 91% and 95%, respectively.	<ul> <li>phy, The specificity and sensitivity of TOE in and identifying AD was 90% (79-100) and 98% (95-100) for type A and 100% and 99% (97%-100%) for type B.</li> <li>The TOE findings were false negative in 3 patients (2 with type A and 1 with type B).</li> </ul>	Mhen any signs of TTE were considered for the diagnosis of TAAAD, a 97% sensitivity and 78% specificity was recorded.	IRA, The sensitivity and specificity of TTE associated with any sign was 96.5% and 70.2% for diagnosis of type A AAD.	<ul> <li>TOE had an overall sensitivity of 88% for AD diagnosis (90% for Type A and 80% for type B).</li> <li>The TOE findings were false negative in 23 patients (14 with type A and 7 with type B).</li> <li>A acute aortic dissection; CAAAD; classic type A</li> </ul>
Referenc	lest			Gated-CT angiography	CT, Angiograf MRI, surgery, autopsy.	Surgical findin	CTA, TOE, M surgery and autopsy	CT, MRI, and angiography. ng: TAAAD; type
Device used				IE33 Philips with 2–7 MHz Omni III probe.	Sonos 1,000 and 1,500	NR	The following multiprobe machines with a 2– 5 MHz phased array probe were used: two MyLab 5, two MyLab30 Gold, two MyLab alpha (Esaote, Genova, Italy), one HD7 (Koninklijke Philips, Amsterdam, Netherlands), three Vivid S5, and one Vivid S6 (GE Healthcare, Wauwatosa, WI, USA)	NR omography: MRI, magnetic resonance imagir
Index	lest			TOE	TOE	TTE	TTE	TOE computed t
	AAD in osis	Type B ( <i>n</i> )		13	19	I	27	NR aphy; CT, c
ics	ts with <sub>.</sub> al diagn	Type A ( <i>n</i> )		6	30	146	85	NR
racterist	Patien fin	Total ( <i>n</i> )		22	49	146	112	NR ageal ech
tient cha	olled	Mean age (years)		67 ± 13.3	60.6 ± 16.8	NR	62 ± 16.7	63.6 ± 13.7 transoesoph
Ра	its en	Яч		44/ 22	80/ 32	272/ 170	540/ 299	414/ 214 y; TOE,
	Patier	Sample ( <i>n</i> )		66	112	442	839	628 cardiograph
Location				Italy	United States	China	Brazil, Germany, Italy, and Switzerland	United States, Italy, Japan, Germany, Spain, and Israel transthoracic echc
Study design				Single center retrospective study	A single center retrospective study.	Retrospective study of two centers	A multicenter prospective study	A multicenter retrospective study. tic dissection; TTE,
Author ID				Agricola et al. (17)	Keren et al. (18)	Hui et al. (19)	Nazerian et al. (20)	Moore et al. (7) AAD, acute aor



#### TTE diagnostic accuracy

Eight of the 11 studies evaluated the diagnostic role of TTE in type A AAD. A subgroup analysis of data from these studies showed that conventional TTE had a pooled sensitivity and specificity of 85.35% (95% CI: 78.40–92.30) (Figure 3) and 84.51% (95% CI: 74.22–94.80) (Figure 4), respectively. Conversely, subgroup analysis of data in one of the studies showed that contrast-enhanced TTE had a sensitivity and specificity of 93.30% (95% CI: 85.75–100) (Figure 3) and 97.60% (95% CI: 94.45–100) (Figure 4).

On the other hand, 3 studies evaluated the role of TTE in diagnosing type B AAD. A subgroup analysis of data from these studies indicated that conventional TTE had pooled sensitivity and specificity of 45.89% (95% CI: 20.98–70.80) (Figure 5) and 87.05 (95% CI: 66.47–100) (Figure 6), respectively. In contrast, the subgroup analysis suggested that contrast-enhanced TTE had a sensitivity and specificity 83.60% (95% CI: 74.40–92.80) (Figure 5) and 94.50 (86–100) (Figure 6).

# the diagnosis of type A AAD, of which the subgroup analysis showed that conventional TOE was 93.64% sensitive and 95.50% specific, while contrast-enhanced TOE was 95.60% sensitive and 100% specific (Figures 7, 8).

On the other hand, 4 of the 5 studies evaluated the diagnosis of Type B AAD, of which the overall sensitivity and specificity of conventional TOE was 99.80% and 99.87%, while contrastenhanced TOE was 100% sensitive and specific (Figures 9, 10).

#### Misdiagnosis

The pooled data from studies evaluating the role of TTE in diagnosing AAD shows that false negatives occur at a rate of 28.6%, while false positives occur at a rate of 18.6%. On the other hand, the pooled data for studies evaluating TOE shows that false positives and negatives occur in about 2.4% and 4.3% of the patients undergoing this diagnostic criterion for AAD (Table 2).

#### TOE diagnostic accuracy

The Diagnostic accuracy of TOE in diagnosing AAD was discussed in 5 included studies. Four out of 5 studies evaluated

# Discussion

AAD represents a medical emergency with a high mortality rate within the first few hours of symptom onset. Therefore, it is

Study						Effect S with 959	ize % Cl	Weight (%)
Contrast-enhanced TTE								
Evangelista et al., 2010a				-	<b>—</b> — 93.3	80 [ 85.75	, 100.00]	12.77
Heterogeneity: $T^2 = 0.00$ , $I^2 = \%$ , $H^2 = .$					<b>93</b> .3	80 [ 85.75	, 100.85]	
Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .								
Conventional TTE								
Asouhidou et al., 2009			-	_	52.1	7 [ 30.88,	73.47]	5.47
Hui et al., 2022					97.0	0 [ 94.00,	100.00]	15.23
Cecconi et al., 2012				_	- 90.0	0[81.50	98.50]	12.15
Nazerian et al., 2014			-	-	54.0	0 [ 39.50	68.50]	8.42
Nienaber et al., 1992			-		— 85.0	0 [ 67.66	100.00]	7.02
Sobczyk et al., 2015					- 92.9	8 [ 88.85	97.11]	14.75
Evangelista et al., 2010b					— 82.2	0 [ 69.90	94.50]	9.69
Nazerian et al., 2019						0[91.85	100.00]	14.50
Heterogeneity: T <sup>2</sup> = 71.65, I <sup>2</sup> = 87.18%, H <sup>2</sup> = 7.	80			-	85.3	5 [ 78.40	92.30]	
Test of $\theta_i = \theta_j$ ; Q(7) = 54.61, p = 0.00								
Overall				•	86.6	6 [ 80.50	92.82]	
Heterogeneity: $T^2 = 62.49$ , $I^2 = 85.36\%$ , $H^2 = 6$ .	83							
Test of $\theta_i = \theta_i$ : Q(8) = 54.63, p = 0.00								
Test of group differences: $Q_b(1) = 2.31$ , $p = 0.1$	3							
Random-effects DerSimonian-Laird model	20	40	60	80	100			
<b>;</b> plot of pooled sensitivity of TTE in diagnosing type A A	AD.							

essential to have rapid and accurate imaging tools to diagnose this condition. In the present study, we have evaluated the role of TTE and TOE in the diagnosis of AAD and found that conventional TOE has a better sensitivity and specificity for the diagnosis of both Type A and type B AAD compared to conventional TTE. However, our subgroup analysis has shown that contrastenhanced TTE results in similar diagnostic accuracy as the conventional TOE. Moreover, pooled results show that misdiagnosis is lower when TOE is used for AAD diagnosis than when TTE is used.

We have shown that TTE is inferior to TOE in diagnosing AAD. This finding aligns with a previous French multicenter study which recorded sensitivity and specificity of 55% and 89% for TTE and 96% and 94% for TOE. Furthermore, the study concluded that TOE is far superior to TTE when the thoracic descending aorta and the aortic arch are involved (20). These results are further reinforced by a previous meta-analysis which did not distinguish the data for AAD patients. That study showed that TOE had a sensitivity and specificity of 98% and 95%, respectively, and was comparable to other imaging techniques such as MRI and Helical CT (7). Even though our results have shown very high sensitivity and specificity for the TOE in diagnosing AAD, specificity as low as 78% and

sensitivity of 80% have been recorded. However, this does not mean that the results of those studies are not accurate since they fall within the range outlined in the 2014 guidelines on the diagnosis and treatment of aortic diseases by the European society of cardiology (4).

It is worth noting that several variables can influence the sensitivity and specificity of TTE to match the accuracy of TOE. The first variable is the echocardiography level of training. In the study by Cecconi and colleagues, where the TTE was carried out by well-trained operators, the sensitivity and specificity for diagnosing classical type A AAD (CAAAD) was very high (90% and 96%, respectively) (9). Similarly, Sobczyk and Krzysztof reported that the echocardiographic examinations were performed by an experienced echocardiographer, and the reported sensitivity for AAD was very high (99.42%) (11). These results indicate that the TTE should be carried out by well-trained, experienced echocardiographers who can achieve optimal images and interpret them correctly to improve the diagnostic accuracy of this imaging technique in AAD. Evidence also shows that patient selection impacts the outcomes of TTE in diagnosing AAD. Sobczyk and Krzysztof reported that the sensitivity of TTE reached 99.4% when diagnosing type A AAD because the data was retrieved from patients transferred to their center with an already established

Study						with	n 95%	b Cl	(%)
Contrast-enhanced TTE									
Evangelista et al., 2010a				-	- 97.	.60 [ 9	4.45,	100.00]	15.12
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$					<b>9</b> 7	.60 [ 9	4.45,	100.75]	
Test of $\theta_{i} = \theta_{j}$ : Q(0) = 0.00, p = .									
Conventional TTE									
Hui et al., 2022		-	-		78.	00 [ 73	3.50,	82.50]	14.84
Cecconi et al., 2012				-	— 96.	00 [ 93	3.00,	99.00]	15.14
Nazerian et al., 2014				_	— 94.	00 [ 90	0.50,	97.50]	15.05
Nienaber et al., 1992	_				78.	10[62	2.76,	93.44 ]	10.73
Evangelista et al., 2010b					- 89.	20[8	1.90,	96.50]	14.02
Nazerian et al., 2019		-			70.	20[66	6.90,	73.50]	15.09
Heterogeneity: $T^2 = 152.69$ , $I^2 = 96.96\%$ , $H^2 = 32.94$					84.	51[74	4.22,	94.80]	
Test of $\theta_i = \theta_j$ ; Q(5) = 164.68, p = 0.00									
Overall					86.	50 [ 77.	.36, 9	5.64]	
Heterogeneity: $T^2 = 141.19$ , $I^2 = 97.09\%$ , $H^2 = 34.40$									
Test of $\theta_i = \theta_j$ : Q(6) = 206.39, p = 0.00									
Test of group differences: $Q_b(1) = 5.69$ , $p = 0.02$									
Random-effects DerSimonian-Laird model	60	70	80	90	100				

Study				Ef wi	fect Size th 95% (	e Cl	Weight (%)
Contrast-enhanced TTE							
Evangelista et al., 2010a				- 83.60	74.40,	92.80]	32.18
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$				83.60	74.40,	92.80]	
Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .							
Conventional TTE							
Asouhidou et al., 2009				— 0.00 [	0.00,	97.50]	14.33
Nienaber et al., 1992				55.50 [	25.56,	85.44]	22.33
Evangelista et al., 2010b			<u> </u>	56.20 [	44.15,	68.25]	31.16
Heterogeneity: $T^2 = 281.25$ , $I^2 = 58.64\%$ , $H^2 = 2.42$				45.89 [	20.98,	70.80]	
Test of $\theta_i = \theta_j$ : Q(2) = 4.84, p = 0.09							
Overall				56.81 [	32.48,	81.14]	
Heterogeneity: $T^2 = 456.88$ , $I^2 = 86.47\%$ , $H^2 = 7.39$							
Test of $\theta_i = \theta_i$ : Q(3) = 22.18, p = 0.00							
Test of group differences: $Q_b(1) = 7.75$ , p = 0.01							
	0	50		100			
Bandom-effects DerSimonian-Laird model		50					

Study				Effect Size with 95% Cl	W	eight (%)
Contrast-enhanced TTE						
Evangelista et al., 2010a				94.50 [ 86.00, 10	00.00] 7	72.11
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$				<b>94.50</b> [ 81.70, 10	07.30]	
Test of $\theta_i = \theta_j$ : Q(0) = -0.00, p = .						
Conventional TTE						
Nienaber et al., 1992				99.99 [ 91.59, 1	00.00]	8.05
Evangelista et al., 2010b				81.80 [ 70.70, 9	92.90] 1	19.84
Heterogeneity: $T^2 = 0.00$ , $I^2 = 0.00\%$ , $H^2 = 1.00$				87.05 [ 66.47, 1	07.63]	
Test of $\theta_i = \theta_j$ : Q(1) = 0.62, p = 0.43						
Overall				92.42 [ 81.55, 1	03.29]	
Heterogeneity: $T^2 = 0.00$ , $I^2 = 0.00\%$ , $H^2 = 1.00$						
Test of $\theta_i = \theta_i$ : Q(2) = 0.98, p = 0.61						
Test of group differences: $Q_b(1) = 0.36$ , p = 0.55						
Random-effects DerSimonian-Laird model	70	80	90	100		

Study					E W	ffect Size ith 95%	e Cl	Weight (%)
Contrast-enhanced TOE								
Evangelista et al., 2010a					95.60	89.80,	100.00]	23.47
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$					95.60	89.80,	101.40]	
Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .								
Conventional TOE								
Keren et al., 1996					90.00 [	79.50,	100.00]	9.53
Evangelista et al., 2010b					95.60 [	89.80,	100.00]	23.47
Nienaber et al., 1992			_		100.00 [	91.58,	100.00]	13.72
Moore et al., 2002					90.00 [	85.25,	94.75]	29.80
Heterogeneity: T <sup>2</sup> = 9.07, I <sup>2</sup> = 43.04%, H <sup>2</sup> = 1.76			$\langle$		93.64 [	89.10,	98.19]	
Test of $\theta_i = \theta_j$ : Q(3) = 5.27, p = 0.15								
Overall					94.00 [	90.49,	97.51]	
Heterogeneity: T <sup>2</sup> = 4.88, I <sup>2</sup> = 30.88%, H <sup>2</sup> = 1.45								
Test of $\theta_1 = \theta_1$ : Q(4) = 5.79, p = 0.22								
Test of group differences: $Q_{b}(1) = 0.27$ , p = 0.60								
Random-effects DerSimonian-Laird model	80	85	90	95	100			
	Study         Contrast-enhanced TOE         Evangelista et al., 2010a         Heterogeneity: $T^2 = 0.00$ , $l^2 = .%$ , $H^2 = .$ Test of $\theta_i = \theta_j$ : $Q(0) = 0.00$ , $p = .$ Conventional TOE         Keren et al., 1996         Evangelista et al., 2010b         Nienaber et al., 1992         Moore et al., 2002         Heterogeneity: $T^2 = 9.07$ , $l^2 = 43.04\%$ , $H^2 = 1.76$ Test of $\theta_i = \theta_j$ : $Q(3) = 5.27$ , $p = 0.15$ Overall         Heterogeneity: $T^2 = 4.88$ , $l^2 = 30.88\%$ , $H^2 = 1.45$ Test of $\theta_i = \theta_j$ : $Q(4) = 5.79$ , $p = 0.22$ Test of $group$ differences: $Q_b(1) = 0.27$ , $p = 0.60$ Random-effects DerSimonian-Laird model	Study         Contrast-enhanced TOE         Evangelista et al., 2010a         Heterogeneity: $T^2 = 0.00$ , $I^2 = .%$ , $H^2 = .$ Test of $\theta_i = \theta_j$ : $Q(0) = 0.00$ , $p = .$ Conventional TOE         Keren et al., 1996         Evangelista et al., 2010b         Nienaber et al., 1992         Moore et al., 2002         Heterogeneity: $T^2 = 9.07$ , $I^2 = 43.04\%$ , $H^2 = 1.76$ Test of $\theta_i = \theta_j$ : Q(3) = 5.27, p = 0.15         Overall         Heterogeneity: $T^2 = 4.88$ , $I^2 = 30.88\%$ , $H^2 = 1.45$ Test of $\theta_i = \theta_j$ : Q(4) = 5.79, p = 0.22         Test of group differences: $Q_b(1) = 0.27$ , p = 0.60         80         Random-effects DerSimonian-Laird model	Study         Contrast-enhanced TOE         Evangelista et al., 2010a         Heterogeneity: $T^2 = 0.00$ , $I^2 = .%$ , $H^2 = .$ Test of $\theta_i = \theta_j$ : $Q(0) = 0.00$ , $p = .$ Conventional TOE         Keren et al., 1996         Evangelista et al., 2010b         Nienaber et al., 1992         Moore et al., 2002         Heterogeneity: $T^2 = 9.07$ , $I^2 = 43.04\%$ , $H^2 = 1.76$ Test of $\theta_i = \theta_j$ : $Q(3) = 5.27$ , $p = 0.15$ Overall         Heterogeneity: $T^2 = 4.88$ , $I^2 = 30.88\%$ , $H^2 = 1.45$ Test of $\theta_i = \theta_j$ : $Q(4) = 5.79$ , $p = 0.22$ Test of $group$ differences: $Q_b(1) = 0.27$ , $p = 0.60$ $80$ $80$	Study         Contrast-enhanced TOE         Evangelista et al., 2010a         Heterogeneity: $T^2 = 0.00$ , $l^2 = .%$ , $H^2 = .$ Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .         Conventional TOE         Keren et al., 1996         Evangelista et al., 2010b         Nienaber et al., 1992         Moore et al., 2002         Heterogeneity: $T^2 = 9.07$ , $l^2 = 43.04\%$ , $H^2 = 1.76$ Test of $\theta_i = \theta_j$ : Q(3) = 5.27, p = 0.15         Overall         Heterogeneity: $T^2 = 4.88$ , $l^2 = 30.88\%$ , $H^2 = 1.45$ Test of $\theta_i = \theta_j$ : Q(4) = 5.79, p = 0.22         Test of group differences: $Q_b(1) = 0.27$ , p = 0.60         80       85         90         Random-effects DerSimonian-Laird model	Study         Contrast-enhanced TOE         Evangelista et al., 2010a         Heterogeneity: $T^2 = 0.00$ , $I^2 = .%$ , $H^2 = .$ Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .         Conventional TOE         Keren et al., 1996         Evangelista et al., 2010b         Nienaber et al., 1992         Moore et al., 2002         Heterogeneity: $T^2 = 9.07$ , $I^2 = 43.04\%$ , $H^2 = 1.76$ Test of $\theta_i = \theta_j$ : Q(3) = 5.27, p = 0.15         Overall         Heterogeneity: $T^2 = 4.88$ , $I^2 = 30.88\%$ , $H^2 = 1.45$ Test of $\theta_i = \theta_j$ : Q(4) = 5.79, p = 0.22         Test of group differences: $Q_b(1) = 0.27$ , p = 0.60         80       85       90       95         Random-effects DerSimonian-Laird model	Study       with         Contrast-enhanced TOE       Evangelista et al., 2010a       95.60 [         Heterogeneity: $T^2 = 0.00$ , $I^2 = .%$ , $H^2 = .$ 95.60 [         Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .       95.60 [         Conventional TOE       90.00 [         Keren et al., 1996       90.00 [         Evangelista et al., 2010b       95.60 [         Nienaber et al., 1992       90.00 [         Moore et al., 2002       90.00 [         Heterogeneity: $T^2 = 9.07$ , $I^2 = 43.04\%$ , $H^2 = 1.76$ 93.64 [         Test of $\theta_i = \theta_j$ : Q(3) = 5.27, p = 0.15       94.00 [         Overall       94.00 [         Heterogeneity: $T^2 = 4.88$ , $I^2 = 30.88\%$ , $H^2 = 1.45$ 94.00 [         Test of $\theta_i = \theta_j$ : Q(4) = 5.79, p = 0.22       90.00 [         Test of group differences: $Q_b(1) = 0.27$ , p = 0.60       90.95 100         80       85       90       95	Study       with 95%         Contrast-enhanced TOE       Evangelista et al., 2010a       95.60 [ 89.80,         Heterogeneity: $T^2 = 0.00, l^2 = .%, H^2 = .$ 95.60 [ 89.80,         Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .       95.60 [ 89.80,         Conventional TOE       90.00 [ 79.50,         Keren et al., 1996       90.00 [ 79.50,         Evangelista et al., 2010b       95.60 [ 89.80,         Nienaber et al., 2002       90.00 [ 79.50,         Heterogeneity: $T^2 = 9.07, l^2 = 43.04\%, H^2 = 1.76$ 90.00 [ 85.25,         Test of $\theta_i = \theta_j$ : Q(3) = 5.27, p = 0.15       94.00 [ 90.49,         Overall       94.00 [ 90.49,         Heterogeneity: $T^2 = 4.88, l^2 = 30.88\%, H^2 = 1.45$ 94.00 [ 90.49,         Test of $\theta_i = \theta_j$ : Q(4) = 5.79, p = 0.22       94.00 [ 90.49,         Reterogeneity: $T^2 = 4.88, l^2 = 30.88\%, H^2 = 1.45$ 94.00 [ 90.49,         Generalize the standard model       95.00 [ 90.49,         More et al., 2002       90.00 [ 90.49,         Heterogeneity: $T^2 = 4.88, l^2 = 30.88\%, H^2 = 1.45$ 94.00 [ 90.49,         Generalize the standard model       95.00 [ 90.49,	Study       with 95% CI         Contrast-enhanced TOE       Evangelista et al., 2010a       95.60 [ 89.80, 100.00]         Heterogeneity: $T^2 = 0.00$ , $I^2 = .%$ , $H^2 = .$ 95.60 [ 89.80, 101.40]         Test of $\theta_1 = \theta_j$ : Q(0) = 0.00, p = .       90.00 [ 79.50, 100.00]         Conventional TOE       90.00 [ 79.50, 100.00]         Keren et al., 1996       90.00 [ 79.50, 100.00]         Evangelista et al., 2010b       95.60 [ 89.80, 100.00]         Nienaber et al., 1992       90.00 [ 79.50, 100.00]         Moore et al., 2002       90.00 [ 85.25, 94.75]         Heterogeneity: $T^2 = 9.07$ , $I^2 = 43.04\%$ , $H^2 = 1.76$ 93.64 [ 89.10, 98.19]         Test of $\theta_1 = \theta_j$ : Q(3) = 5.27, p = 0.15       94.00 [ 90.49, 97.51]         Overall       94.00 [ 90.49, 97.51]         Heterogeneity: $T^2 = 4.88$ , $I^2 = 30.88\%$ , $H^2 = 1.45$ 94.00 [ 90.49, 97.51]         Test of $\theta_1 = \theta_j$ : Q(4) = 5.79, p = 0.22       90.95 100         Random-effects DerSimonian-Laird model       90.95 100

Study					Effect Size with 95% Cl	Weight (%)
Contrast-enhanced TOE						
Evangelista et al., 2010a					100.00 [ 99.40, 10	0.00] 40.37
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$					<b>♦</b> 100.00 [ 99.40, 100	0.60]
Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .						
Conventional TOE						
Keren et al., 1996						0.00] 32.20
Evangelista et al., 2010b					96.40 [ 92.30, 100	0.00] 23.63
Nienaber et al., 1992	-		-		78.79 [ 63.83, 93	3.75] 3.80
Heterogeneity: $T^2 = 12.09$ , $I^2 = 68.52\%$ , $H^2 = 3.18$					95.50 [ 90.40, 100	).59]
Test of $\theta_i = \theta_j$ ; Q(2) = 6.35, p = 0.04						
Overall				•	• 97.70 [ 94.64, 10	0.76]
Heterogeneity: $T^2 = 5.95$ , $I^2 = 76.25\%$ , $H^2 = 4.21$						
Test of $\theta_i = \theta_i$ : Q(3) = 12.63, p = 0.01						
Test of group differences: $Q_{b}(1) = 2.96$ , p = 0.09						
Pandom offects DerSimonian Laird model	60	70	80	90	100	

Study				Effect Size with 95% Cl	Weight (%)
Contrast-enhanced TOE					
Evangelista et al., 2010a				100.00 [ 99.65, 100.00	)] 41.68
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$				<b>♦</b> 100.00 [ 99.65, 100.35	5]
Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .					
Conventional TOE					
Moore et al., 2002				80.00 [ 65.75, 94.25	5] 0.12
Keren et al., 1996				■100.00 [ 99.00, 100.00	)] 16.46
Evangelista et al., 2010b				<b>1</b> 00.00 [ 99.65, 100.00	)] 41.68
Nienaber et al., 1992				90.90 [ 70.38, 100.00	0.06 [0
Heterogeneity: T <sup>2</sup> = 0.77, I <sup>2</sup> = 63.93%, H <sup>2</sup> = 2.77				99.80 [ 98.49, 101.1	1]
Test of $\theta_i = \theta_j$ : Q(3) = 8.32, p = 0.04					
Overall				♦ 99.97 [ 99.48, 100.46	5]
Heterogeneity: T <sup>2</sup> = 0.12, I <sup>2</sup> = 51.92%, H <sup>2</sup> = 2.08					
Test of $\theta = \theta$ : O(4) = 8.32, p = 0.08					
Test of aroup differences: O. (1) = 0.09. $p = 0.77$					
	70	80	90	100	
Random-effects DerSimonian-Laird model					

Study					E	ffect Size ith 95% Cl		Weight (%)
Contrast-enhanced TOE								
Evangelista et al., 2010a					- 100.00 [	99.55, 10	0.00]	47.73
Heterogeneity: $T^2 = 0.00$ , $I^2 = .\%$ , $H^2 = .$					<b>•</b> 100.00 [	99.55, 10	00.45]	
Test of $\theta_i = \theta_j$ : Q(0) = 0.00, p = .								
Conventional TOE								
Keren et al., 1996					99.00 [	97.50, 10	0.00]	4.30
Evangelista et al., 2010b					100.00 [	99.55, 10	0.00]	47.73
Nienaber et al., 1992					97.60 [	91.34, 10	0.00]	0.25
Heterogeneity: $T^2 = 0.03$ , $I^2 = 4.34\%$ , $H^2 = 1.05$					<b>9</b> 9.87 [	99.35, 10	0.39]	
Test of $\theta_{i} = \theta_{j}$ ; Q(2) = 2.09, p = 0.35								
Overall					🔶 99.95 [	99.64, 10	0.26]	
Heterogeneity: $T^2 = 0.00$ , $I^2 = 0.00\%$ , $H^2 = 1.00$								
Test of $\theta_i = \theta_j$ : Q(3) = 2.18, p = 0.54								
Test of group differences: $Q_b(1) = 0.14$ , p = 0.71								
Random-effects DerSimonian-Laird model	92	94	96	98	100			
. <b>0</b> plot of pooled specificity of TOE in diagnosing type B .	AAD.							

suspicion of AD in about 50% of the patients (11). Moreover, with the recent technological advancements, harmonic imaging in TTE has been employed to improve diagnostic accuracy. In harmonic imaging TTE investigations, the frequency is generally double the fundamental frequency to decrease the artifactual echoes emanating from hazy tissue-tissue and tissue-blood interfaces. Thus, this approach offers a better echo graphic signal with an enhanced noise ratio and improved lateral resolution. Owing to this, the overall clarity of pictures is increased, and better identification of thin linear features like aortic intimal flaps or distinct wall thickening may be noticed (9, 21, 22).

In recent years contrast echocardiography has also been used to improve the accuracy of diagnosing AAD (23). Our subgroup analysis has shown that transthoracic contrast echocardiography significantly improved the accuracy of diagnosing AAD and produced similar outcomes as the conventional TOE. This finding is supported by a previous research article which reported that the specificity of conventional TTE increased from

TABLE 2 Pooled echocardiography false negatives and false positives.

Index test	Number of studies	Event rate	95% Cl
TTE			
False negatives	4	0.286	0.087-0.629
False positives	4	0.186	0.051-0.492
TOE			
False negatives	3	0.024	0.11-0.049
False positives	4	0.043	0.016-0.111

73.68% to 100% after contrast enhancement (24). Moreover, that study found that contrast-enhanced TTE correctly diagnosed AAD in all patients, while the conventional TTE had 5 false positive results. In addition, our subgroup analysis has shown a slight improvement in the diagnosis of AAD when using contrast TOE as opposed to conventional TOE. Similarly, Agricola and colleagues found that the sensitivity and specificity of conventional TOE were slightly improved using contrast enhancement (91% and 95% vs. 100%, without and with contrast enhancement, respectively) (15).

Even with the superiority of TOE in diagnosing AAD, our results have shown that false negatives and positives are observed during the diagnosis. This shows that the AAD diagnosis cannot be excluded confidently based on a single TOE test finding. It is strongly recommended that additional imaging tests are performed if the initial diagnostic tests do not identify AD despite the diagnosis being clinically suspected. It is also essential to have other confirmatory imaging tests for patients diagnosed with AAD using the TOE test. Evidence from the study by Agricola and colleagues seems to suggest that contrast-enhanced TOE can eliminate the rate of false positives and negatives (15). According to this study, the conventional TOE had 2 false negatives and 2 false positives in diagnosing AAD, but no false results were recorded when contrast TOE was used. However, the results of this study cannot be used to guide clinical care since it included a very small sample with AAD. Additionally, the study excluded some patients with type A and B AAD for various reasons, including shock. Therefore, the study population was limited, meaning their results might have been influenced.

Apart from diagnosing AAD, echocardiography is also essential in diagnosing AD complications and other differential features which increase the risk of mortality in these patients. One of the most frequent complications which can be diagnosed and quantified using echocardiography is aortic regurgitation (AR) which occurs in 40%-76% of patients presenting with type AAD (25). TOE usually provides better information on the mechanisms of AR, which may influence the surgical decision for valve replacement or repair. Nienaber and colleagues reported that TOE and TTE have similar accuracy in diagnosing AR (100% sensitivity and 87.5 specificity for both echocardiography techniques). Further analysis in this study shows that TOE and TTE are even more sensitive than MRI in diagnosing AR; however, more trials should be carried out to establish this finding. Echocardiography is also critical in diagnosing arterial vessel involvement since visceral or peripheral mal perfusion syndrome is highly associated with increased morbidity and mortality. Evidence in the study by Evangelista et al. (13) showed that TTE was more helpful in diagnosing supra-aortic vessel involvement than TOE with and without contrast enhancement. However, it was reported that TOE made the correct diagnosis in all 19 patients with subclavian artery involvement. Further analysis showed that TTE was limited in diagnosing coronary artery and coeliac trunk involvements, while TOE with and without contrast enhancement was able to diagnose 4 patients with coronary dissections confirmed surgically and 19 dissections of coeliac trunk confirmed by CT. The study also showed that none of the echocardiographic techniques could diagnose lower vessel involvement.

Pericardial effusion is another chronic complication that arises in around 40% of individuals with type A acute aortic syndromes (AAS) (25). The existence of pericardial effusion could signal a rupture of the false lumen in the pericardium; however, it might be due to the adventitia irritation induced by aortic hematoma or a minor effusion from the wall. Evidence suggests that echocardiography is a good diagnostic tool for pericardial effusion and cardiac tamponade. For instance, Nienaber and colleagues reported that of the 53 patients with clinically suspected AAD, TTE correctly diagnosed pericardial effusion in 6 patients, while TOE diagnosed pericardial effusion in 7 patients (14). However, TOE was more sensitive than TTE when diagnosing the presence of pericardial effusion. Further analysis showed that the pericardial effusion was associated with type A dissection. This finding is reinforced by Hui and colleagues, who reported that TTE diagnosed pericardial effusion/cardiac tamponade in 48.76% of the patients presenting with type A AAD (17).

In addition to diagnosing AD complications, echocardiography is essential in identifying true and false lumen. The ability to distinguish between the true and false lumen is critical since placing an endoluminal stent graft in the false lumen can have hazardous consequences. However, the distinguishment between the two is not very straightforward, but the following features can be used; the first feature is about the size. Often the false lumen is characterized by a large-sized lumen due to the higher false luminal pressures (26, 27). The second feature is pulsation, of which the true lumen is characterized by systolic expansion while the false lumen is characterized by systolic compression (28). Thirdly, the flow direction can be used to distinguish between the two, of which the true lumen is characterized by systolic antegrade flow while the false lumen is characterized by reduced or absence of systolic antegrade flow or by a retrograde flow (29). Finally, the contrast echo flow in a true lumen is early and fast, while it is delayed and slow in a false lumen. Evidence shows that TTE and TOE are good imaging techniques for identifying false and true lumen. Evangelista and colleagues reported that TOE better visualized the false lumen entry tear than TTE (p < 0.001). However, the conventional TOE could not correctly identify the true lumen in 6 patients (8.8%) (13). In all these cases, the intima was immobile in the M-mode, and the color doppler showed that both lumina had a similar flow pattern. When the contrast-enhanced TOE was applied, all six cases were correctly identified as the true lumen. Further analysis showed that the conventional TTE and TOE could not identify the false lumen flow direction; however, the contrast-enhanced TOE correctly diagnosed 9 cases of false lumen while TTE correctly diagnosed false lumen in 7 patients.

#### Limitations

Like any other scientific research study, our review was limited in some aspects. First, we only included studies published in English, meaning that some of the relevant studies that would have been used in the analysis were excluded, therefore, introducing a selection bias in our study. Secondly, we have a substantial heterogeneity in some of the analyses; however, the heterogeneity did not influence the results of our meta-analyses since most of the studies were of good methodological quality. Thirdly, the studies included in this review did not have one specific reference standard, which might have resulted in the high heterogeneity and influenced the diagnostic accuracy of the index test since not all reference standards provide an accurate diagnosis. Moreover, the studies included in this review are few due to the strict eligibility criteria, which did not include studies that discussed AD but did not differentiate the results for patients with the acute phase. Finally, one study in this review only included patients referred to the medical center, with 50% of the patients already having been diagnosed with AAD; this might have influenced the diagnostic accuracy reported in that study and subsequently influenced our results (11).

# Conclusion

TOE is a favorable diagnostic test with excellent sensitivity and specificity for detecting AAD. However, false positives and negatives are still a problem, meaning that additional imaging tests are required to make a correct diagnosis. Our subgroup analysis also shows that contrast-enhanced TTE can produce similar results to conventional TOE in diagnosing AAD. Therefore, in emergency settings where TOE is not readily available, this imaging modality can be used for the diagnosis. However, it should be noted that both TTE and TOE should be performed by experienced and well-trained operators and interpreters for the correct diagnosis of AAD.

# Author contributions

HA: Conceptualization, Data curation, Formal Analysis, Methodology, Project administration, Resources, Software, Validation, Visualization, Writing - original draft, Writing -BA: Conceptualization, review & editing. Validation. Visualization, Writing - original draft, Writing - review & editing. AA: Conceptualization, Supervision, Writing - review & editing. BA: Conceptualization, Supervision, Writing - review & editing. YY: Data curation, Formal Analysis, Methodology, Project administration, Resources, Software, Writing - review & editing. WE: Data curation, Formal Analysis, Methodology, Project administration, Writing - review & editing. KB: Data curation, Project administration, Resources, Writing - review & editing. NA: Data curation, Writing - review & editing. EE: Data curation, Writing - review & editing.

# References

1. Erbel R, Alfonso F, Boileau C, Dirsch O, Eber B, Haverich A, et al. Diagnosis and management of aortic dissection. *Eur Heart J.* (2001) 22:1642–81. doi: 10.1053/euhj. 2001.2782

2. Braverman AC. Acute aortic dissection. Circulation. (2010) 122:184–8. doi: 10. 1161/CIRCULATIONAHA.110.958975

3. Hagan PG, Nienaber CA, Isselbacher EM, Bruckman D, Karavite DJ, Russman PL, et al. The international registry of acute aortic dissection (IRAD): new insights into an old disease. *JAMA*. (2000) 283:897–903. doi: 10.1001/jama.283.7.897

4. Erbel R, Aboyans V, Boileau C, Bossone E, Bartolomeo RD, Eggebrecht H, et al. 2014 ESC guidelines on the diagnosis and treatment of aortic diseases: document covering acute and chronic aortic diseases of the thoracic and abdominal aorta of the adult. The task force for the diagnosis and treatment of aortic diseases of the European society of cardiology (ESC). *Eur Heart J*. (2014) 35:2873–926. doi: 10. 1093/eurheartj/chu281

5. Evangelista A, Flachskampf FA, Erbel R, Antonini-Canterin F, Vlachopoulos C, Rocchi G, et al. Echocardiography in aortic diseases: EAE recommendations for clinical practice. *Eur J Echocardiogr.* (2010) 11:645–58. doi: 10.1093/ejechocard/jeq056

6. Suzuki T, Distante A, Zizza A, Trimarchi S, Villani M, Salerno Uriarte JA, et al. Diagnosis of acute aortic dissection by D-dimer: the international registry of acute aortic dissection substudy on biomarkers (IRAD-bio) experience. *Circulation.* (2009) 119:2702–7. doi: 10.1161/CIRCULATIONAHA.108.833004

7. Shiga T, Wajima Z, Apfel CC, Inoue T, Ohe Y. Diagnostic accuracy of transesophageal echocardiography, helical computed tomography, and magnetic resonance imaging for suspected thoracic aortic dissection: systematic review and meta-analysis. Arch Intern Med. (2006) 166:1350–6. doi: 10.1001/archinte.166.13.1350

8. Sommer T, Fehske W, Holzknecht N, Smekal AV, Keller E, Lutterbey G, et al. Aortic dissection: a comparative study of diagnosis with spiral CT, multiplanar transesophageal echocardiography, and MR imaging. *Radiology*. (1996) 199:347–52. doi: 10.1148/radiology.199.2.8668776

9. Cecconi M, Chirillo F, Costantini C, Iacobone G, Lopez E, Zanoli R, et al. The role of transthoracic echocardiography in the diagnosis and management of acute type A aortic syndrome. *Am Heart J.* (2012) 163:112–8. doi: 10.1016/j.ahj.2011.09.022

10. Nazerian P, Vanni S, Castelli M, Morello F, Tozzetti C, Zagli G, et al. Diagnostic performance of emergency transthoracic focus cardiac ultrasound in suspected acute type A aortic dissection. *Intern Emerg Med.* (2014) 9:665–70. doi: 10.1007/s11739-014-1080-9

11. Sobczyk D, Nycz K. Feasibility and accuracy of bedside transthoracic echocardiography in diagnosis of acute proximal aortic dissection. *Cardiovasc Ultrasound.* (2015) 13:15. doi: 10.1186/s12947-015-0008-5

12. Asouhidou I, Asteri T. Acute aortic dissection: be aware of misdiagnosis. *BMC Res Notes.* (2009) 2:25. doi: 10.1186/1756-0500-2-25

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13. Evangelista A, Avegliano G, Aguilar R, Cuellar H, Igual A, González-Alujas T, et al. Impact of contrast-enhanced echocardiography on the diagnostic algorithm of acute aortic dissection. *Eur Heart J.* (2010) 31:472–9. doi: 10.1093/eurheartj/ehp505

14. Nienaber CA, Spielmann RP, von Kodolitsch Y, Siglow V, Piepho A, Jaup T, et al. Diagnosis of thoracic aortic dissection. Magnetic resonance imaging versus transesophageal echocardiography. *Circulation*. (1992) 85:434–47. doi: 10.1161/01. cir.85.2.434

15. Agricola E, Slavich M, Bertoglio L, Fisicaro A, Oppizzi M, Marone E, et al. The role of contrast enhanced transesophageal echocardiography in the diagnosis and in the morphological and functional characterization of acute aortic syndromes. *Int J Cardiovasc Imaging*. (2014) 30:31–8. doi: 10.1007/s10554-013-0290-y

16. Keren A, Kim CB, Hu BS, Eyngorina I, Billingham ME, Mitchell RS, et al. Accuracy of biplane and multiplane transesophageal echocardiography in diagnosis of typical acute aortic dissection and intramural hematoma. *J Am Coll Cardiol.* (1996) 28:627–36. doi: 10.1016/0735-1097(96)00186-6

17. Hui L, Wang D, Liu T, Liu B, Wang Y, Liu B. Diagnostic performance of transhoracic echocardiography in screening acute type A aortic dissection from ST-segment elevated myocardial infarction. *Cardiovasc Diagn Ther.* (2022) 12:603–13. doi: 10.21037/cdt-22-59

18. Nazerian P, Mueller C, Vanni S, Soeiro AM, Leidel BA, Cerini G, et al. Integration of transthoracic focused cardiac ultrasound in the diagnostic algorithm for suspected acute aortic syndromes. *Eur Heart J.* (2019) 40:1952–60. doi: 10.1093/eurheartj/ehz207

19. Moore AG, Eagle KA, Bruckman D, Moon BS, Malouf JF, Fattori R, et al. Choice of computed tomography, transesophageal echocardiography, magnetic resonance imaging, and aortography in acute aortic dissection: international registry of acute aortic dissection (IRAD). *Am J Cardiol.* (2002) 89:1235–8. doi: 10.1016/s0002-9149(02)02316-0

20. Guéret P, Sénéchal C, Roudaut R, Cormier B, Wolf J, Tribouilloy C, et al. Comparison of transesophageal and transthoracic echocardiography in acute aortic dissection. A multi-center prospective study. *J Am Coll Cardiol.* (1991) 17:A260. doi: 10.1016/0735-1097(91)92008-A

21. Thomas JD, Rubin DN. Tissue harmonic imaging: why does it work? J Am Soc Echocardiogr. (1998) 11:803-8. doi: 10.1016/s0894-7317(98)70055-0

22. Hoffmann R, Marwick TH, Poldermans D, Lethen H, Ciani R, van der Meer P, et al. Refinements in stress echocardiographic techniques improve inter-institutional agreement in interpretation of dobutamine stress echocardiograms. *Eur Heart J.* (2002) 23:821–9. doi: 10.1053/euhj.2001.2968

23. Cursaro M, Simmonds I, Bradley J, Shirazi M, Disney P, Cunnington M. Aortic plaque defined by contrast transthoracic echocardiography. *Australas J Ultrasound Med.* (2012) 15:158–9. doi: 10.1002/j.2205-0140.2012.tb00199.x

24. Lv Q, Wu Y, Wang M. The diagnostic value of contrast enhanced ultrasound in acute aortic dissection. *Ultrasound Med Biol.* (2017) 43:S50. doi: 10.1016/j. ultrasmedbio.2017.08.1107

25. Evangelista A, Maldonado G, Gruosso D, Gutiérrez L, Granato C, Villalva N, et al. The current role of echocardiography in acute aortic syndrome. *Echo Res Pract.* (2019) 6:R53–63. doi: 10.1530/ERP-18-0058

26. Meredith EL, Masani ND. Echocardiography in the emergency assessment of acute aortic syndromes. *Eur J Echocardiogr.* (2009) 10:i31–9. doi: 10.1093/ ejechocard/jen251

27. Hashimoto S, Kumada T, Osakada G, Kubo S, Tokunaga S, Tamaki S, et al. Assessment of transesophageal Doppler echography in dissecting aortic aneurysm. J Am Coll Cardiol. (1989) 14:1253–62. doi: 10.1016/0735-1097(89)90424-5

28. Baliga RR, Nienaber CA, Bossone E, Oh JK, Isselbacher EM, Sechtem U, et al. The role of imaging in aortic dissection and related syndromes. *JACC Cardiovasc Imaging*. (2014) 7:406–24. doi: 10.1016/j.jcmg.2013.10.015

29. Patil TA, Nierich A. Transesophageal echocardiography evaluation of the thoracic aorta. *Ann Card Anaesth.* (2016) 19:S44–55. doi: 10.4103/0971-9784.192623