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Harries, Iwan; Weir-McCall, Jonathan R.; Williams, Michelle C.; Shambrook, James; Roditi, Giles; Bull, Russel
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CT imaging prior to transcatheter aortic valve implantation in the UK

Iwan Harries, Jonathan R Weir-McCall, Michelle C Williams, James Shambrook, Giles Roditi, Russel Bull, Gareth J Morgan-Hughes, Edward D Nicol, Alastair J Moss

ABSTRACT

Objective This cross-sectional observational study sought to describe variations in CT in the context of transcatheter aortic valve implantation (CT-TAVI) as currently performed in the UK.

Methods 408 members of the British Society of Cardiovascular Imaging were invited to complete a 27-item online CT-TAVI survey.

Results 47 responses (12% response rate) were received from 40 cardiac centres, 23 (58%) of which performed TAVI on-site (TAVI centres). Only six respondents (13%) performed high-volume activity (>200 scans per year) compared with 13 (28%) performing moderate (100–200 scans per year) and 27 (59%) performing low (0–99 scans per year) volume activity. Acquisition protocols varied (41% retrospective, 12% prospective with wide padding, 47% prospective with narrow padding), as did the phase of reporting (45% systolic, 37% diastolic, 11% both, 6% unreported). Median dose length product was 675 mGy cm (IQR 477–954 mGy cm). Compared with non-TAVI centres, TAVI centres were more likely to report minimum iliofemoral luminal diameter (n=25, 96% vs n=7, 58%, p=0.003) and optimal tube angulation for intervention (n=12, 46% vs n=1, 8%, p=0.02).

Conclusions This national survey formally describes current CT-TAVI practice in the UK. High-volume activity was only present at one in seven cardiac CT centres. There is wide variation in scan acquisition, scan reporting and radiation dose exposure in cardiac CT centres.

INTRODUCTION

Percutaneous transcatheter aortic valve implantation (TAVI) for severe aortic valve disease has become widespread over the past decade. The improved safety profile of TAVI compared with cardiothoracic surgery in lower risk cohorts is likely to result in the further expansion of TAVI in the coming years. Historically, cardiothoracic surgery has offered direct visualisation of the cardiac anatomy to plan and implement appropriate valve replacement, but the advent of TAVI has necessitated a preprocedural road map of the cardiac anatomy to facilitate the safe deployment of appropriately sized aortic valve prostheses. The isotropic resolution of cardiac CT has made it a critical investigation in assessing valve anatomy when deciding which patients may be suitable for TAVI.

While standards of practice exist for cardiac CT in the assessment of coronary artery disease, there is a paucity of data regarding the optimal image acquisition protocols and standardised reporting criteria for CT in TAVI (CT-TAVI) scans. Recently, the Society of Cardiovascular Computed Tomography has published an expert consensus document highlighting the need for standardisation of nomenclature and measurements using CT-TAVI. However, whether these international recommendations reflect the current practice of CT-TAVI acquisition and reporting in the UK is unknown. Additionally, while diagnostic dose reference levels are available for most CT procedures, the acceptable limits of ionising radiation for CT-TAVI are yet to be established.

This study aimed to survey members of a national cardiovascular imaging society who...
perform CT-TAVI to assess the variability in scan acquisition, reporting and dissemination of findings.

METHODS

Survey population
All 408 members of the British Society of Cardiovascular Imaging (BSCI) were invited to complete a 27-item online survey (online supplementary file 1) (www.surveymonkey.com) in April 2018. The BSCI is composed of members of the cardiovascular imaging community, which serves as a Specialist Advisory Group on matters related to cardiovascular imaging to the British Cardiovascular Society and Royal College of Radiologists, UK. While standards of practice for the training and reporting of CT coronary angiography (CTCA) have been produced by the BSCI in line with recommendations from the Francis report, no standards of practice exist for the training and reporting of CT-TAVI scans. Survey questions were constructed to assess the level of activity, and variability in CT-TAVI practice, across centres in the UK currently delivering this service for the National Health Service (NHS) England, NHS Wales, NHS Scotland, and Health and Social Care in Northern Ireland. Patients and the public were not involved in the design of the survey.

Items surveyed
All participants were surveyed individually to prevent collaboration bias between institutions. All responses were included in the analysis. Parameters recorded included the type of institution (centres performing TAVI (TAVI centre) vs those without a TAVI (non-TAVI centre) programme), make of CT scanner, total number of CT-TAVI scans performed annually, scan acquisition protocol, use of medication (beta-blockers) during scan acquisition, contrast volume and radiation doses of the last five scans performed, structured reporting metrics of the aortic valve annulus and vascular access, and access to and distribution of findings to the TAVI multidisciplinary team.

Statistical analysis
Continuous variables are reported as mean±SD or median and IQR, for normally and non-normally distributed variables, respectively. The survey questions were built using skip logic to save respondents time and ensure a high response rate. The skip or ‘conditional branching’ pattern varies depending on the respondent’s answers and this results in incomplete data sets for some survey questions. Consequently, categorical variables are presented as counts and percentages of the total number of complete responses for each question. The survey was designed as a descriptive exploration of current practices and thus not designed nor powered to detect specific differences. Where differences in practice were apparent, continuous variables were assessed with Student’s t-test and categorical variables were assessed with Fisher’s exact test in cases where χ² test was not valid. A p value <0.05 was taken to indicate statistical significance. The analysis was carried out in SPSS (IBM SPSS Statistics for Macintosh, V.25.0. IBM).

RESULTS

CT-TAVI annualised activity in TAVI and non-TAVI centres
Forty-seven responses (12% response rate) from 40 cardiac centres in the UK were received (figure 1). Fifty-three per cent of respondents (n=25/47) were radiologists and 47% (n=22/47) were cardiologists with a subspecialty interest in cardiac CT. Most respondents worked in centres which included a TAVI programme on-site (n=30/47, 63.8%) (figure 1). Annualised volume of CT-TAVI activity for acquisition and reporting varied considerably (n=1, no data provided), 59% of respondents (n=27/46) reported a low volume of activity (<100 CT-TAVI scans per annum), compared with 28% (n=13/46) performing a moderate level of activity (100–200 scans per annum) and 13% (n=6/46) performing a high level of scan activity (>200 scans per annum) (figure 1). Respondents working in TAVI centres performed more CT-TAVI activities per annum compared with those in non-TAVI centres (p<0.001).

Scan acquisition and radiation dose in TAVI and non-TAVI centres
Overall, retrospective ECG-gated acquisitions were performed by 41% (n=14/34) of respondents, compared with 47% (n=16/34) performing prospective ECG-triggered acquisitions with narrow padding, and 12% (n=4/34) performing prospective ECG-triggered acquisitions with wide padding (figure 2). There was no difference between TAVI and non-TAVI centres regarding acquisition protocol (p=0.72). Median contrast volumes and dose length product (DLP) of the five most recently performed CT-TAVI scans did not vary between TAVI and non-TAVI centres (table 1). The median CT-TAVI DLP across all participating centres was 675 mGy.cm (IQR 477–954 mGy.cm). Using wide padding (30%–80% R-R interval) more than doubles the CT-TAVI DLP from 423 to 921 mGy.cm for prospectively triggered CT-TAVI, whereas retrospective ECG gating yielded a median DLP of 882 mGy.cm (table 1). Beta-blockade was administered by 15% (n=7/47) of respondents. Furthermore, cardiologists were significantly more likely to administer beta-blockers compared with radiologists (n=6/25, 24% vs n=1/22, 5%, p=0.01).

CT-TAVI reporting in TAVI and non-TAVI centres
Thirty-five centres provided parameters on reporting. Two centres acquired scans but did not analyse preprocedural measurements (6%). Forty-five per cent (n=16/35) of centres reported systolic-only aortic annulus measurements, 37% (n=13/35) reported diastolic-only aortic annulus measurements and 11% (n=4/35) reported both systolic and diastolic aortic annulus measurements. Five centres did not report the phase of reconstruction. With respect to the detailed assessment of aortic root measurements, there were no standard metrics which
were reported by all centres (table 2). Interestingly, coronary artery luminal stenosis severity was included in all CT-TAVI reports at 24% (n=9/38) of centres and sometimes reported in 50% (n=19/38) of centres. Compared with non-TAVI centres, TAVI centres were more likely to report minimum iliofemoral luminal diameter (n=25/26, 96% vs n=7/12, 58%, p=0.003) and optimal tube angulation for intervention (n=12/26, 46% vs n=1/12, 8%, p=0.02) (table 2). There were no significant differences in reporting parameters when compared by specialty.

‘Heart Team’ dissemination of findings
A ‘Heart Team’ discussion involving a cardiac imaging specialist is more frequent in TAVI centres compared with non-TAVI centres (96%, n=24/25 vs 33%, n=4/12, p<0.001) (figure 3). However, to facilitate interpretation of CT-TAVI findings, a cardiac imaging specialist is ‘always present’ at 57% (n=16/28) of centres overall (54%, n=13/24 TAVI centres vs 75%, n=3/4 non-TAVI centres), and ‘most of the time’ at a further 21% (n=6/28) of
### Table 1  Estimated contrast volumes and dose length product (DLP) for the last five CT-TAVI scans performed at each centre and radiation doses according to scan protocol

<table>
<thead>
<tr>
<th></th>
<th>Overall (n=22)</th>
<th>TAVI centres (n=22)</th>
<th>Non-TAVI centres (n=9)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Contrast volume (mL)</strong></td>
<td>100 (80–130)</td>
<td>95 (74–120)</td>
<td>110 (88–160)</td>
<td>0.06</td>
</tr>
<tr>
<td>Min 50, max 170</td>
<td>Min 60, max 150</td>
<td>Min 50, max 170</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>DLP (mGy.cm)</strong></td>
<td>675 (477–954)</td>
<td>550 (423–939)</td>
<td>700 (538–1075)</td>
<td>0.66</td>
</tr>
<tr>
<td>Min 139, max 2000</td>
<td>Min 145, max 2000</td>
<td>Min 139, max 1750</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### CT-TAVI scan protocol

<table>
<thead>
<tr>
<th></th>
<th>Retrospective—whole cardiac cycle (n=12)</th>
<th>Prospective ECG-gated narrow padding (n=12)</th>
<th>Prospective ECG-gated wide padding (n=4)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>DLP (mGy.cm)</strong></td>
<td>882 (524–1688)*</td>
<td>423 (230–631)</td>
<td>921 (850–1138)**</td>
</tr>
</tbody>
</table>

Values are median and IQR.

*P=0.003, **p=0.002, both versus prospective ECG-gated narrow padding.

TAVI, transcatheter aortic valve implantation.

### Table 2  Measurements and structures typically included in the CT-TAVI report at each centre

<table>
<thead>
<tr>
<th></th>
<th>Overall (n=38)</th>
<th>TAVI centres (n=26)</th>
<th>Non-TAVI centres (n=12)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Aortic annulus circumference</strong></td>
<td>21 (55%)</td>
<td>16 (62%)</td>
<td>5 (42%)</td>
<td>0.25</td>
</tr>
<tr>
<td><strong>Aortic annulus area</strong></td>
<td>24 (63%)</td>
<td>17 (65%)</td>
<td>7 (58%)</td>
<td>0.675</td>
</tr>
<tr>
<td><strong>Aortic annulus diameter (perimeter derived)</strong></td>
<td>23 (61%)</td>
<td>17 (65%)</td>
<td>6 (50%)</td>
<td>0.367</td>
</tr>
<tr>
<td><strong>Aortic annulus diameter (area derived)</strong></td>
<td>19 (50%)</td>
<td>13 (50%)</td>
<td>6 (50%)</td>
<td>1.0</td>
</tr>
<tr>
<td><strong>Minimum short-axis measurement</strong></td>
<td>20 (53%)</td>
<td>14 (54%)</td>
<td>6 (50%)</td>
<td>0.82</td>
</tr>
<tr>
<td>Sinus of Valsalva diameter (cusp-cusp)</td>
<td>20 (53%)</td>
<td>13 (50%)</td>
<td>7 (58%)</td>
<td>0.63</td>
</tr>
<tr>
<td><strong>Extent and distribution of aortic root calcification</strong></td>
<td>31 (82%)</td>
<td>22 (85%)</td>
<td>9 (75%)</td>
<td>0.48</td>
</tr>
<tr>
<td><strong>Number of valve cusps (eg, bicuspid, tricuspid)</strong></td>
<td>32 (84%)</td>
<td>22 (85%)</td>
<td>10 (83%)</td>
<td>0.92</td>
</tr>
<tr>
<td><strong>Distance (height) from aortic annulus to LMS ostium</strong></td>
<td>30 (79%)</td>
<td>21 (81%)</td>
<td>9 (75%)</td>
<td>0.69</td>
</tr>
<tr>
<td><strong>Distance (height) from aortic annulus to RCA ostium</strong></td>
<td>30 (79%)</td>
<td>21 (81%)</td>
<td>9 (75%)</td>
<td>0.69</td>
</tr>
<tr>
<td><strong>Distance (height) from aortic annulus to sinotubular junction (left cusp)</strong></td>
<td>11 (29%)</td>
<td>6 (23%)</td>
<td>5 (42%)</td>
<td>0.24</td>
</tr>
<tr>
<td><strong>Distance (height) from aortic annulus to sinotubular junction (right cusp)</strong></td>
<td>9 (24%)</td>
<td>5 (19%)</td>
<td>4 (33%)</td>
<td>0.34</td>
</tr>
<tr>
<td><strong>Distance (height) from aortic annulus to sinotubular junction (non-coronary cusp)</strong></td>
<td>7 (18%)</td>
<td>4 (15%)</td>
<td>3 (25%)</td>
<td>0.48</td>
</tr>
<tr>
<td><strong>Presence of LVH</strong></td>
<td>23 (61%)</td>
<td>18 (69%)</td>
<td>5 (42%)</td>
<td>0.17</td>
</tr>
<tr>
<td><strong>Ascending aorta diameter</strong></td>
<td>31 (82%)</td>
<td>21 (81%)</td>
<td>10 (83%)</td>
<td>0.85</td>
</tr>
<tr>
<td><strong>Descending and abdominal aorta (tortuosity, intraluminal obstruction, calcification)</strong></td>
<td>31 (82%)</td>
<td>23 (88%)</td>
<td>8 (67%)</td>
<td>0.11</td>
</tr>
<tr>
<td><strong>Subclavian and brachiocephalic artery diameter</strong></td>
<td>18 (47%)</td>
<td>16 (62%)</td>
<td>2 (17%)</td>
<td>0.01</td>
</tr>
<tr>
<td><strong>Minimum iliofemoral artery luminal diameter</strong></td>
<td>32 (84%)</td>
<td>25 (96%)</td>
<td>7 (58%)</td>
<td>0.003</td>
</tr>
<tr>
<td><strong>Iliofemoral artery patency</strong></td>
<td>32 (84%)</td>
<td>24 (92%)</td>
<td>8 (67%)</td>
<td>0.04</td>
</tr>
<tr>
<td><strong>Iliofemoral artery tortuosity</strong></td>
<td>31 (82%)</td>
<td>24 (92%)</td>
<td>7 (58%)</td>
<td>0.01</td>
</tr>
<tr>
<td><strong>Optimal tube angulation data to inform fluoroscopic projection for device deployment</strong></td>
<td>13 (34%)</td>
<td>12 (46%)</td>
<td>1 (8%)</td>
<td>0.02</td>
</tr>
</tbody>
</table>

Values are count (n) and percentage (%).

Bold values indicate statistical significance at a p value <0.05.

LMS, left main stem; LVH, left ventricular hypertrophy; RCA, right coronary artery; TAVI, transcatheter aortic valve implantation.
Valvular heart disease

Figure 3  Dissemination of findings to Heart Team. Heart Team multidisciplinary discussions were present at 76% of centres overall (A). When Heart Team discussions were in place, a cardiac imaging specialist was ‘always present’ at 57% of meetings (B). There was a significant difference in the provision of Heart Team meetings at TAVI and non-TAVI centres (96%, n=24/25 vs 33%, n=4/12, p<0.001) (C). TAVI, transcatheter aortic valve implantation.

centres overall (21%, n=5/24 TAVI centres vs 25%, n=1/4 non-TAVI centres). A cardiac imaging specialist is ‘never’ present at 11% (n=3/28) of centres (13%, n=3/24 TAVI centres vs 0%, n=0 non-TAVI centres).

DISCUSSION

In this study, we found that only one in seven respondents perform a high annualised volume of CT-TAVI. As such, there is wide variation in practice among UK cardiac centres in the acquisition and reporting of CT-TAVI. Median radiation dose exposure was 675 mGy.cm in UK cardiac centres, however significant reductions in the DLP were obtained by using prospective ECG gating with narrow-padded acquisitions. To ensure optimal reporting standards in all UK cardiac centres, further collaboration between TAVI and non-TAVI centres should be encouraged.

Following the expansion of TAVI over the past decade, the provision of non-invasive cardiac imaging to support structural heart disease programmes can no longer be solely delivered in tertiary cardiac centres. This comes at a time when there has been considerable service expansion in cardiac CT following the implementation of the revised National Institute for Health and Care Excellence Clinical Guideline 95 (update 2016) necessitating that the UK, as a whole, provides over 350,000 cardiac CT angiograms per annum. This compares to the estimated 42,340 scans reported by 198 accredited individuals on 304 CTCA-capable scanners recently being performed. This has resulted in even the highest volume cardiac CT
centres needing to double their service provision for coronary artery disease assessment, and thus hindering further expansion of cardiac CT in the assessment of structural heart disease in tertiary cardiac centres. However, an increasing number of new centres which offer cardiac CT has meant that experienced imaging cardiologists and radiologists can also provide lower numbers of CT-TAVI in house remote from TAVI centres, thereby performing CT-TAVI at a location which may be more convenient for the patient. Whether a ‘hub and spoke’ model that acquires CT imaging at the local centre and offers shared reporting with structural imaging specialists at TAVI centres will improve CT interpretation in non-TAVI centres is unknown. However, this model has been successfully employed in other collaborative networks using CT imaging.10

With respect to radiation dose, the ‘as low as reasonably achievable’ principle is a central tenant of legislation stating that ionising radiation should be optimised to use the minimum amount of radiation to attain adequate diagnostic information.11 For coronary CT angiography, the UK has the lowest international diagnostic reference levels set by Ionising Radiation (Medical Exposure) Regulations at 380 mGy.cm for retrospective acquisitions with ECG gating and 170 mGy.cm for prospective acquisitions with no padding.12 Focusing attention on the optimal scan acquisition in the field of coronary CT angiography has resulted in many centres switching from retrospective to prospective acquisitions and reducing tube current potential, thereby reducing the median radiation exposure by 78%.13 In this study, the wide variation in scanning techniques for acquiring CT-TAVI data sets would suggest that there is scope to lower radiation dose exposure through improved standardisation of practice across UK cardiac CT centres. Indeed, this would fall under the remit of the ‘Getting it right first time’ model that aims to reduce unwarranted variation in UK clinical practice. Of note, radiation doses reported by the members surveyed varied 14-fold between the lowest and highest reported DLPs (range 139–2000 mGy.cm). Some of this variation may be attributed to the perceived need to analyse the aortic annulus in both ventricular systole and diastole. While there is no consensus on the optimal acquisition technique, international guidelines advocate a preference for systolic scan coverage, as this is when the aortic annulus is most circular and has the largest orifice area for accurate valve sizing. However, reverse dynamism of the interventricular septum which is frequently seen in patients with sigmoid hypertrophy may favour dimensions measured in ventricular diastole, thus necessitating retrospective ECG-gated reconstructions throughout the entire cardiac cycle. Further investigation of the optimal CT-TAVI acquisition for routine clinical practice is warranted.

Decisions regarding TAVI are made in the context of a ‘Heart Team’ multidisciplinary discussion which draws on expertise from interventional cardiologists, cardiac surgeons and cardiac imaging specialists.17 While local institutions may differ in their approach to the composition of the ‘Heart Team’, it is unsurprising that these are more likely to occur in TAVI centres than non-TAVI centres. However, it is notable that despite the critical importance of CT-TAVI in the preprocedural planning of transcatheter heart valve procedures, there are some TAVI centres where a cardiac imaging specialist is never present to interpret the imaging findings in the clinical context. This may explain why the aortic annular dimension evaluation, a critical step for transcatheter aortic valve sizing, differs in the level of detailed reporting between sites. In order to maximise the information obtained from CT-TAVI scans both in terms of written reporting and dissemination to interventional cardiologists, cardiac imagers should be integrated into the ‘Heart Team’ discussion. This would help address specific questions in CT-TAVI reporting, such as which parameters constitute a mandatory minimum for a CT-TAVI report. In this regard, the observation that coronary artery stenosis assessment is performed in up to 50% of cardiac CT centres raises the question as to whether this should be routinely reported on CT-TAVI scans.19 The use of CT-TAVI to evaluate the patency of coronary artery bypass grafts and native coronary arteries is feasible, however as periprocedural medication (beta-blocker, glyceryl trinitrate) that is routinely administered for coronary CT angiography is contraindicated in the setting of severe aortic stenosis, optimal images may not always be obtained.20 Interestingly, while selective beta-blockade has been used in the context of severe asymptomatic and symptomatic aortic stenoses, there is a paucity of evidence supporting the safety of intravenous metoprolol administration during CT-TAVI acquisition. If beta1-selective blockade is to be considered on an individual basis, the patient should be counselled on the risks of exacerbating symptoms related to aortic stenosis. Medication use at the time of CT-TAVI will be explored in more detail in future studies.

There are some limitations to this study. Despite attempts to obtain responses from UK cardiac CT centres, some centres that perform CT-TAVI were not captured. Moreover, the survey only elicited responses (response rate only 12%) from members of the BSCI, meaning that an unknown but likely significant number of those reporting CT-TAVI, for example, interventional cardiologists, have not been captured. This introduces bias and influences the external validity of the data, while reliance on self-reporting of data may also introduce error. Furthermore, some responses from different individuals originated from the same centre. Because these data were frequently discordant, reflecting intracentre variation, we elected to include all responses from all respondents, which is also a source of bias. The image quality of CT-TAVI scans obtained at UK cardiac CT centres was not analysed in this study, but we presume that each centre obtains diagnostic CT-TAVI images on the majority of occasions. Finally and perhaps most importantly, no data on complications following CT-TAVI (eg, contrast-induced nephropathy)
were collected. Future studies exploring the relationship between CT-TAVI, periprocedural complications and patient outcomes would be instructive and could inform optimal CT-TAVI scan acquisition protocols.

CONCLUSION
This national survey formally describes current CT-TAVI practice in the UK. High-volume activity was only present at one in seven cardiac CT centres. There is wide variation in scan acquisition, scan reporting and radiation dose exposure in cardiac CT centres.

Contributors IH helped design the research, collected, analysed and interpreted the data, drafted and revised the manuscript critically for important intellectual content. JWM and MCW analysed the data and revised the manuscript critically for important intellectual content. JS, GR, RB and GJMH revised the manuscript critically for important intellectual content. AJM conceived and designed the research, analysed and interpreted the data, interpreted the data and revised the manuscript critically for important intellectual content. EN helped design the research, interpreted the data and revised the manuscript critically for important intellectual content. AJM conceived and designed the research, analysed and interpreted the data, redrafted the manuscript and revised it critically for important intellectual content. All authors approved the final version of the manuscript and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work have been appropriately investigated and resolved.

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Competing interests
None declared.

Patient consent for publication
Not required.

Ethics approval
Owing to the study design, ethical approval was not required prior to distribution of the survey.

Provenance and peer review
Not commissioned; externally peer reviewed.

Data availability statement
Data are deidentified questionnaire responses available from the corresponding author upon reasonable request.

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ORCIDs
Iwan Harries http://orcid.org/0000-0003-0688-3345
Jonathan R Weir-McCall http://orcid.org/0000-0001-5842-842X
Alastair J Moss http://orcid.org/0000-0003-4123-2070

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