Implementation of 3D Printing in Medical Care for Preoperative Planning of Complex Ventricular Septal Defect

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ABSTRACT

Three-dimensional (3D) modeling and printing in medicine have emerged to encompass every aspect of medical applications. This ranges from education, illustration, and treatment, as well as patient care whether for purposes of diagnosis or treatment and surgical planning. In the past few decades, these novel tools have shown promising utility to help radiologists and the medical team to improve quality of patient care and outcomes via 3D printing application and utilization. This workflow will be illustrated through a ventricular septal defect (VSD) case at which 3D analysis was critical in the assessment and treatment planning of the patient’s underlying medical condition.

CASE REPORT

An 82-year-old female with history of diabetes, coronary artery disease, and hypertension was referred to our institution for further assessment of multiple cardiac comorbidities. Her past medical history was significant for postoperative chest pain after a lumbar spine surgery for treatment of spinal stenosis. A 12-lead electrocardiogram (EKG) showed normal sinus rhythm and right bundle branch block, which could be related the patient’s reported history of prior myocardial infarction (MI) (Figure 1). Transesophageal echocardiogram (TEE) at that time showed aortic valve dysfunction, left ventricular ejection fraction of 55-60%, normal left ventricular wall motion, and ventricular septal defect (VSD) involving the inferior aspect of the interventricular muscular septum (Figure 2, 3). The aortic valve dysfunction showed: moderate stenosis with mild to moderate regurgitation; peak velocity: 261 cm/s; and the ratio of left ventricular outflow tract (LVOT) peak velocity to aortic valve (AV) peak velocity, also known as LVOT/AV ratio: 0.26—indicating moderate aortic stenosis. The VSD measured 1.1 cm in diameter with multiple suspected connections to right ventricular cavity, which showed left to right shunt flow with a peak velocity of 515 cm/s and a peak gradient of 106 mmHg (Figure 3). The VSD was located in the inferior aspect of the interventricular muscular septum 3.6 cm from the mitral valve annulus and 4.1 cm from the left ventricular apex (Figure 4). Upon further investigation, the patient reported 15-year chronic history of incidentally found heart murmur during routine outpatient visit that she never followed-up on, possibly related to the patient’s VSD, which was presumed to be post-MI in etiology. The patient’s comorbidities and recent spine surgery prevented
immediate cardiac catheterization at the outside hospital facility.

The patient was stabilized medically and discharged home but continued to report progressively worsening mild dyspnea and low activity tolerance. The patient was therefore referred to our institution cardiology service for further workup. A gated cardiac computed tomography angiogram (CTA) was performed to assess for transcatheter aortic valve repair (TAVR) planning and accessibility of VSD for closure device. The CTA showed aortic valvarus sclerosis, mitral annular calcifications, inferior interventricular muscular septum VSD, dilated pulmonary arteries and an enlarged left atrium (Figure 4). A diagnostic and therapeutic left heart catheterization (LHC) was then performed, which showed diffuse right coronary artery (RCA) atherosclerosis causing moderate stenosis near the origin, which was treated with three Xience drug eluting stents (DES) (figure 5). In addition, the patient received an aortic valve TAVR using a 26 mm Sapien aortic valve bioprosthesis (Figure 6). The patient’s procedure was complicated by left femoral artery access occlusion necessitating vascular surgical repair using ipsilateral great saphenous vein (GSV) patch. The patient also suffered post-TAVR complete atroventricular block (Figure 7), which required placement of cardiac pacing device using ACCOLADETM pacemaker (Boston Scientific, model L301) via right femoral vein access (figure 8). These complications had led to prolonged hospitalization and further deferment of the VSD closure procedure.

Post-intervention EKG demonstrated post-procedural changes of cardiac pacemaker device placement, which showed expected electronic atrial pacemaker signal on the EKG tracer, normal sinus rhythm, along with resolution of the previously seen right bundle branch block (Figure 9).

In addition, postprocedural echocardiogram showed proper positioning of TAVR without valvular dysfunction, and normal biometrics of the aortic valve prosthesis (figure 10). The echocardiogram showed no signs of valvular stenosis or regurgitation; transvalvular peak velocity: 102 cm/s; mean velocity: 67.5 cm/s; and postoperative LVOT/AV ratio: 0.76 (previously measured 0.25 on preoperative echocardiogram), which indicates resolution of the aortic stenosis (Figure 10).

During this current admission and prior to the patient’s discharge, the patient’s case was discussed at interventional cardiology conference and the radiology department was consulted for further characterization of the patient’s VSD given the presumed complexity on CTA images. Our department’s 3D Clinical Applications Division and Advanced Imaging Lab was able to provide a 3D print of the patient’s VSD by using the CTA dataset. The 3D print revealed that the patient’s VSD had complex anatomy with three fenestrations involving the right aspect of the VSD and was not simply a single defect (Figures 11, 12). Treatment of such VSD would require one large or multiple amplatzer devices potentially leading to multiple complications; such as, prolonged procedure time, amplatzer device detachment, formation of blood clots, which can lead to embolic strokes, or bacterial colonization within the small fenestrations, which increases the risk for infectious endocarditis and subsequently postoperative septic emboli. Our multidisciplinary team concluded that having an anatomically complex VSD with multiple fenestrations would pose more long-term postoperative complications to the patient than if left untreated, and that the postoperative risks in these situations would outweigh the benefits of attempting VSD closure. Due to the complex anatomy of the VSD and the patient’s multiple co-morbidities, the patient was therefore deemed a poor candidate for VSD closure and further treatment was deferred.

The patient was discharged home and a six-month outpatient clinic visit follow-up showed overall improvement in the patient’s symptoms, increase exercise tolerability, and reduction in the frequency of reported dyspnea or chest pain. This case highlights the critical role 3D printing can play in management of complex patients and in providing value added care especially in perioperative planning.

MATERIAL AND METHODS:
Imaging acquisition was performed via Toshiba Aquilion Prime, 0.5 mm slice thickness, with 219.65 mm field of view (FOV) using a gated cardiac CTA. We used 3D software assembly: Mimics Innovation Suite Medical 20.0 and 3-matic 12.0, by Materialise®–FDA cleared medical application. Implementation was done using Form2–SLA Printer, offered by Formlabs, which employs liquid resin, which allows printing flexible or rigid material. Post-processing of flexible print was performed using isopropyl alcohol wash and then was cured using ultraviolet light bath.

DISCUSSION

Etiology & Demographics:
VSD is the most common congenital heart disease in children, and is considered the second most common form of adult congenital heart disease following bicuspid aortic valve (1, 2). The congenital form of VSD can happen in isolation in about 0.3% of all newborns, or as a part of complex congenital heart disease (CHD), which accounts for 37% of all CHD (1, 3). The etiology of congenital VSD is failure of formation and closure of the interventricular (IV) septum during embryologic development. The acquired form of VSD in adulthood occurs due to various etiologies and is commonly encountered secondary to myocardial infarction, infectious endocarditis, or iatrogenic injury (4-6).

Clinical & Imaging Findings:
Color Doppler ultrasound transthoracic echocardiography (TTE) is considered the gold standard of VSD diagnosis and evaluation with 95% sensitivity. TTE is also important in evaluating the size, shape, number, and location of VSD and providing hemodynamic information of the direction of the jet size and shunt direction (7, 8).

In addition to ultrasound (US), computed tomography (CT) and magnetic resonance imaging (MRI) cardiac imaging are considered superior imaging modalities that can provide valuable information regarding the anatomy and morphology
of the VSD. In particular, cardiac CT higher resolution capabilities can further delineate any complex anatomy of the VSD shape and morphology, and cardiac MRI can help in evaluating the jet size and direction, along with underlying cardiac pathology such as MI, infectious endocarditis, or various types of congenital or restrictive heart diseases (9, 10).

Three-dimensional (3D) modeling and printing have been utilized in the recent years as advanced imaging tools in medical care and diagnostic workup of unusual cases with complex anatomy and pathology. This emerging technology is revolutionizing patient-centered care by providing virtual and physical representation of the patient’s own complex anatomy and pathology in 3D (11-13). Through this technology, radiologists can provide additional diagnostic information that can improve patient quality and clinical outcomes (14-16).

New technologies like 3D modeling and printing also provide radiologists with novel tools to use in their practice to improve quality of patient care and outcomes (11-13, 17). Due to unfamiliarity with the technology by the majority of practicing radiologists, it may appear that it is not a tangible option for their practice. However, radiology has the opportunity to drive advancement of this technology to meet their unique needs, make it more feasible for every day clinical use, and add value to clinical practices (14, 15, 18, 19). In addition, 3D modeling and printing are promising tools that have the potential to revolutionize surgical planning and intervention in order to improve patient care and outcomes (14, 16, 17, 20, 21). This emerging technology can also potentially impact the future of medical training for surgical residents by providing a 3D printed model for preoperative planning (21-24).

Treatment & Prognosis:
About 90% of isolated VSD cases are asymptomatic incidentally diagnosed during routine physical examination prior to spontaneous closure during childhood (1). Persistent VSD in adulthood can be treated with prophylactic endocarditis medications or surgical closure (25-27). Untreated cases are associated with minimal increased risk for congestive heart failure, arrhythmia, or endocarditis, compared to the general population (28). Various literature resources recommended surgical or endovascular intervention if a patient with VSD presents with symptoms, along with other factors such as, the location, size, and direction of the VSD shunt flow. A commonly used convention for classifying VSD based on size is comparing the VSD diameter to the aortic annulus diameter (AAD). For example, a small VSD measures less or equal to 25% of the AAD and is considered large when it is greater than 75% of the AAD. For patients with untreated chronic large left-to-right VSD shunt, the patient may develop pulmonary arterial hypertension secondary to irreversible vascular endothelial hypertrophy (25-28). Eisenmenger syndrome, which accounts for 10% to 15% of patients with untreated VSD, develops when the pulmonary pressure exceeds the systemic pressure leading right-to-left VSD shunt, which leads to admixture of unoxygenated blood into the systemic circulation. If the shunt volume is large enough, the patient may develop cyanotic heart disease (28).

Differential Diagnoses:
Differential diagnosis of cardiac wall defects includes: pseudoaneurysm, aneurysm, or diverticulum (29). Cardiac pseudoaneurysms are almost exclusively encountered in the left ventricular (LV) free wall secondary to prior infection, trauma, or ischemia. Pseudoaneurysms represent LV rupture contained by overlying adherent pericardium, which prevents frank tamponade (30). Approximately 10% of LV pseudoaneurysms are asymptomatic with symptomatic patients reporting occasional chest pain, palpitations, dyspnea, or syncope. This entity can be incidentally detected on echocardiogram, CT, or MRI as abrupt discontinuity of cardiac wall with orifice-to false aneurysm diameter ratio of more than 0.5 with internal turbulent blood flow, lack of aneurysmal wall calcification, and presence of a thrombus on MRI and US (31, 32).

Conversely, true cardiac aneurysms are thin myocardial wall outpouchings composed of fibrous tissue and necrotic cardiomyocytes, commonly encountered in the LV anterior wall, occasionally with septal involvement. Various etiologies have been linked to true aneurysms, including ischemia, infection, or trauma, with associated symptoms of chest pain, fatigue and palpitations (33). True aneurysms can be complicated by intramural thrombus formation leading to stroke or limb ischemia in the setting of thromboembolic disease. While left ventriculography conventional angiography is the gold standard, MRI, CT, and US can offer further details regarding the presence of wall calcification, transmural thrombus, wall dyskinesia, or ejection fraction abnormality (32). On US and CT, cardiac aneurysms would demonstrate thinning of normal myocardium replaced by thin fibrous tissue, which occasionally calcifies; while on MRI cardiac aneurysm usually presents as T2 hypointense thinning of the left ventricular wall with diffuse post-gadolinium delayed enhancement (23, 33).

Myocardial diverticulum is a congenital protrusion of the cardiac wall characterized by the presence of normal cardiomyocytes that can affect the left ventricular free wall, right ventricular apex, or rarely the IV septum (34). The majority are asymptomatic, but surgical resection is recommended to avoid long term complications such as heart failure, infection, or thromboembolism (35). Imaging findings on US usually demonstrate subtle anechoic wall defect without focal bulging or surrounding myocardial wall abnormal echogenicity (34). On CT, a diverticulum presents as an incomplete, noncalcified defect in the ventricular wall that follows blood pool attenuation. Similarly, on MRI it usually shows normal T1 and T2 signal of the myocardial signal, preserved normal myocardial thickness and enhancement, with focal defect that follows blood pool signal on all MRI sequences (36).

TEACHING POINT
3D Printing is a useful medical tool for the visualization of complex cardiac anatomy such as ventricular septal defects, which can impact the patient’s clinical diagnosis and perioperative planning.
REFERENCES


Figure 1 (bottom): An 82-year-old female with history of multiple cardiac comorbidities presented with chest pain after lumbar spine surgery at an outside hospital. FINDINGS: 12-lead electrocardiogram showing sinus rhythm and right bundle branch block, which could be related the patient's reported history of prior myocardial infarction. TECHNIQUE: 12 lead Electrocardiogram.
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**Figure 2 (left):** An 82-year-old female with history of multiple cardiac comorbidities.

FINDINGS: 2D echocardiogram ultrasound at the level of the aortic valve demonstrates thickening of the aortic valve leaflets indicating aortic stenosis. A blue jet of color flow in the proximal aorta suggesting a superimposed component of aortic regurgitation.

TECHNIQUE: 2D Doppler ultrasound echocardiogram at the level of the aortic valve, using 12Hz frequency probe.

**Figure 3:** An 82-year-old female with history of multiple cardiac comorbidities.

FINDINGS: 2D echocardiogram ultrasound at the level of the interventricular septum demonstrates ventricular septal defect and left to right ventricular shunt, indicated by a blue color flow jet.

TECHNIQUE: 2D Doppler ultrasound echocardiogram at the level of the interventricular septum using 14Hz frequency probe.
Figure 4: An 82-year-old female with history of multiple cardiac comorbidities.

FINDINGS: Arterial phase coronal contrast enhanced computed tomography image of the heart demonstrates a small interventricular defect involving the inferior muscular aspect of the interventricular septum with complex morphology measuring approximately 1.1 cm in diameter. This finding was presumed to be related to the patient’s reported history of prior myocardial infarction.

TECHNIQUE: 2D reformatted coronal CT, 430 mAs, 100 kV, 1 mm slice thickness, 60 ml of Iovue-370 contrast intravenous contrast.

Figure 5: An 82-year-old female with history of multiple cardiac comorbidities status post left heart catheterization fluoroscopic procedure.

FINDINGS: Left femoral artery catheterization fluoroscopic exam demonstrates multifocal areas of high grade right coronary artery stenosis (yellow arrows) near the origin (Figure 5a). Post-stenting image shows improvement in the sites of focal right coronary artery stenosis (Figure 5b).

TECHNIQUE: Fluoroscopic images of the chest at the level of the right coronary artery after the administration of Iovue-370 intravenous iodinated contrast.
Figure 6: An 82-year-old female with history of multiple cardiac comorbidities status post left heart catheterization fluoroscopic procedure.

FINDINGS: Left femoral artery catheterization fluoroscopic exam demonstrates successful placement of percutaneous transcatheter aortic valve repair using a 26 mm Sapien aortic valve bioprosthesis (Figures 6a-b).

TECHNIQUE: Fluoroscopic images of the chest at the level of the aortic valve after the administration of Isovue-370 intravenous iodinated contrast.

Figure 7: An 82-year-old female with history of multiple cardiac comorbidities status post right coronary artery stenting and percutaneous aortic valve repair procedure.

FINDINGS: 12-lead electrocardiogram demonstrates new complete atrioventricular block likely post-procedural in etiology.

TECHNIQUE: 12 lead Electrocardiogram.
Figure 8 (left): An 82-year-old female with history of multiple cardiac comorbidities status post right coronary artery stenting and percutaneous aortic valve repair procedure. FINDINGS: right femoral venous catheterization fluoroscopic exam demonstrates successful placement of cardiac pacing triple-lead device using ACCOLADE™ pacemaker (Boston Scientific, model L301).
TECHNIQUE: Fluoroscopic images of the chest at the level of the cardiac silhouette.

Figure 9: An 82-year-old female with history of multiple cardiac comorbidities status post cardiac pacing device placement.

FINDINGS: 12-lead electrocardiogram demonstrates electronic atrial pacemaker tracer, intraventricular conduction delay, and resolution of the previously seen right bundle branch block.

TECHNIQUE: 12 lead Electrocardiogram.
Figure 10 (left): An 82-year-old female with history of multiple cardiac comorbidities status post percutaneous aortic valve repair.

FINDINGS: 2D echocardiogram at the level of the aortic valve demonstrates successful placement of transcatheter aortic valve repair using a 26 mm Sapien aortic valve bioprosthesis. Transvalvular peak velocity: 102 cm/s and mean velocity: 67.5 cm/s; left ventricular outflow tract to aortic valve ratio: 0.76, which indicates resolution of the aortic stenosis.

TECHNIQUE: 2D echocardiogram at the level of the aortic valve using high frequency ultrasound probe to assess for percutaneous aortic valve prosthesis placement.

Figure 11: An 82-year-old female with history of multiple cardiac comorbidities.

A 3D heart model designed using software assembly: Mimics Innovation Suite Medical 20.0 and 3-matic 12.0, by Materialise©-FDA cleared medical application, which shows complex anatomy with three ventricular septal defect fenestrations (red labels).
Etiology

- The congenital form of ventricular septal defect occurs secondary to interruption of the interventricular septum formation during embryologic development.
- The acquired form occurs due to various etiologies, commonly including ischemia (myocardial infarction) or infection, such as endocarditis.

Incidence

- Ventricular septal defect is the second most common congenital cardiac anomaly in adults and is also considered the most common congenital abnormality in children accounting for 37% of all congenital heart diseases, with isolated incidence of about 0.3% in newborns.

Symptoms

- Depending on the size of the ventricular septal defect, and the association with other congenital heart diseases, an isolated, small ventricular septal defect is usually asymptomatic and is found incidentally on physical exam.

Gender and Age

- It is commonly a congenital heart disease with no gender predilection.
- The acquired form of ventricular septal defect is seen in older population in the setting of myocardial infarction.

Treatment

- Approximately, 90% of small, isolated, congenital ventricular septal defect will close spontaneously during first year of life.
- Patients with persistent adult ventricular septal defect can benefit from surgical closure.

Prognosis

- Persistent adult ventricular septal defect has several comorbidities, such as congestive heart failure, arrhythmia, or endocarditis.

Findings on Imaging

- Cardiac CT or MRI of ventricular septal defect is a muscular defect in the interventricular septum with shunt of blood across the defect. The shunt can be small or large and can be unidirectional or bidirectional. The size and direction of the ventricular septal defect shunt usually correspond to the presence of symptoms and is a good indicator for the assessment of surgical intervention.

Table 1: Summary table of ventricular septal defect.
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#### Table 2: Differential diagnosis table for ventricular septal defect.

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Symptoms</th>
<th>CT Findings</th>
<th>MRI Findings</th>
<th>US Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ventricular septal defect</td>
<td>Isolated small ventricular septal defect are usually asymptomatic.</td>
<td>Focal defect within the normal cardiac muscle, filled with iiodinated contrast. CT is Excellent in delineating the size, location, shape, and complexity of ventricular septal defect.</td>
<td>Focal defect within the normal cardiac muscle which follows the signal of blood pool. MRI imaging can provide valuable information regarding size, location, and direction of jet flow on phase contrast sequences.</td>
<td>TTE is the most sensitive tool in detecting up to 95% of ventricular septal defect by providing information regarding the morphology, location, and size, number of defects, and direction and size of the ventricular septal defect jet flow.</td>
</tr>
<tr>
<td>Cardiac wall aneurysm</td>
<td>Small aneurysms are asymptomatic. Larger aneurysms can cause chest pain, fatigue, and palpitation and are commonly associated with history of prior myocardial infarction.</td>
<td>Fibrous outpouching of the LV anteroapical wall, and occasionally involving the IV septum with calcification.</td>
<td>Thinning of the myocardium which is T2 hypointense with delayed enhancement. Turbulent flow and thrombus are sometimes encountered.</td>
<td>Thinning of the myocardium with turbulent flow. It is commonly seen in the LV anteroapical wall, and occasionally involving the septum. Low ejection fraction and LV wall dyskinesia is commonly seen when it is associated with prior myocardial infarction.</td>
</tr>
<tr>
<td>Cardiac wall pseudoaneurysm</td>
<td>10% are asymptomatic but can cause chest pain, palpitations, dyspnea, and syncope.</td>
<td>Abrupt loss of cardiac wall thickness with adherent pericardium with no calcification, containing non-thrombosed blood.</td>
<td>Abrupt loss of endocardium and myocardial wall with outpouching commonly seen in the LV free wall and apex.</td>
<td>Sudden loss of cardiac wall thickness with turbulent flow commonly seen in the LV free wall.</td>
</tr>
<tr>
<td>Cardiac wall diverticulum</td>
<td>Commonly asymptomatic.</td>
<td>Focal incomplete, noncalcified defect in the interventricular septum that follows blood pool attenuation.</td>
<td>Normal T1 and T2 signal of the myocardium. Preserved myocardial thickness and enhancement with focal defect that follows blood pool signal.</td>
<td>Subtle anechoic wall defect without focal bulging or surrounding myocardial wall abnormal echogenicity.</td>
</tr>
</tbody>
</table>

#### Abbreviations

2D = Two-Dimensional  
3D = Three-Dimensional  
AAD = aortic annulus diameter  
AV = aortic valve  
CHD = congenital heart disease  
CT = computed tomography  
CTA = computed tomography angiogram  
DES = drug eluting stents  
EKG = electrocardiogram  
FOV = field of view  
GSV = great saphenous vein  
IV = interventricular  
LHC = left heart catheterization  
LV = left ventricular  
LVOT = left ventricular outflow tract  
LVOT/AV = left ventricular outflow tract to aortic valve ratio  
MI = myocardial infarction  
MRI = magnetic resonance imaging  
RCA = right coronary artery  
TAVR = transcatheter aortic valve repair  
TEE = Transesophageal echocardiogram  
TTE = transthoracic echocardiography (TTE)  
US = ultrasound  
VSD = ventricular septal defect
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KEYWORDS
Ventricular Septal Defect; 3D Modeling; 3D Printing; Presurgical planning; 3D Printing in Medicine; 3D Printing Applications in Cardiac Imaging

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