Narrative review of cardiac computed tomography perfusion: insights into static rest perfusion

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Abstract: Cardiac or left ventricular perfusion performed with cardiac computed tomography (CCT) is a developing method that may have the potential to complete in a very straightforward way the assessment of ischemic heart disease by means of CT. Myocardial CT perfusion (CTP) can be achieved with a single static scan during the first-pass of the iodinate contrast agent, with the monoenergetic or dual-energy acquisition, or as a dynamic, time-resolved scan during stress by using coronary vasodilator agents. Several methods can be performed, and we focused on static perfusion. CTP may serve as a useful adjunct to coronary CT angiography (CTA) to improve specificity of detecting myocardial ischemia. Technological advances will reduce the radiation dose of myocardial CTP, such as low tube voltage imaging or new reconstruction algorithms, making it a more viable clinical option. The advantages of static first-pass non-stress perfusion are several; the main one is that it can be done to each and every patient who undergoes CCT for the assessment of coronary artery tree. Future advances in CTP will likely improve the diagnostic accuracy of CTP + CTA, and will better estimate the severity of ischemia. Therefore, it is simple and comprehensive. However, it has several limitations. In this review we will discuss the technique with its advantages and limitations.

Keywords: Cardiac computed tomography (CCT); perfusion; static; first-pass; stenosis; ischemia; accuracy

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Introduction

Cardiac computed tomography (CCT) is the non-invasive reference model for the evaluation of coronary arteries (1) and has become the predominant imaging modality to identify coronary artery disease (CAD) with high diagnostic accuracy. CT imaging of the heart has improved rapidly, and it is now possible to perform a comprehensive evaluation using a low amount of radiation dose.

Non-invasive acquisition of patients with suspected CAD has traditionally rested on two pillars: morphologic assessment for coronary artery stenosis and functional/perfusion evaluation for determining the downstream hemodynamic significance of lesions (2). This approach is of fundamental importance as the morphological evaluation of coronary stenoses does not linearly reflect the extension and severity of myocardial ischemia throughout the left ventricle (LV).

Furthermore, by evaluating morphology and function in a combined way, the prognosis and supervision of patients are more reliable and efficient (3,4).

According to the current appropriate use criteria and guidelines, CCT is the method of choice for the exclusion of significant coronary artery stenosis in patients with low
Myocardial perfusion methodologies

Myocardial perfusion can be assessed through different imaging modalities. CCT can be considered a semiquantitative/quantitative method useful in detecting ischemia, and comparable to other perfusion imaging tests.

CCT may be considered in patients with low or intermediate pre-test probability of CAD and non-conclusive stress test. The advantage of CCT over other methods is to detect anatomical details of coronary tree and myocardium. The spatial resolution is very high compared to other imaging modalities but one of the disadvantages is the radiation dose exposure which, however, is gradually improving thanks to technological advancement.

Besides to radiological diagnostics, nuclear cardiology techniques play a main role in the evaluation of heart failure. Myocardial single photon emission computed tomography (SPECT) offer valuable information of ventricular function and myocardial perfusion and commonly used radio-tracers include technetium (Tc)-based agents, such as 99mTc-sestamibi and 99mTc-tetrofosmin. Moreover, positron emission tomography (PET) assess myocardial perfusion, metabolism and viability with high quality images and provide to quantitative analysis. Cardiac PET perfusion radio-tracers are rubidium-82 ($^{82}$Rb), oxygen-15 water ($^{15}$O-H$_2$O) and nitrogen-13 ammonia ($^{13}$N-NH$_3$), most often used with metabolic $^{18}$F-fluorodeoxyglucose ($^{18}$F-FDG) (9).

Although nuclear medicine investigations represent, in the case of PET, the gold standard in the evaluation of myocardial metabolism, CTP has numerous advantages, well described in the next paragraph.

The limitations are in sharp decrease thanks to technological progress and depend largely on the equipment. Dose values delivered will therefore be different based on the performance and technology used.

CTP derives from the CTA acquisition but still remains affected to artifacts, such as beam hardening, which can be overcome thanks to the development of iterative reconstruction algorithms.

Despite the use of multiple protocols and various artifacts, the available data suggests than CTP may have an accuracy which is comparable to SPECT for identifying myocardial ischemia. Ultimately, CTP may serve as a useful adjunct to coronary CTA to improve specificity of detecting myocardial ischemia. Future advances in CTP will likely improve the diagnostic accuracy of CTP + CTA, and will better estimate the severity of ischemia (10).

Overview of myocardial CTP

CT evaluation of myocardial perfusion is based on the distribution of iodinated contrast agent during its first pass over the myocardium. Because the arterial blood flow determines the contrast material’s distribution, myocardial perfusion defects can be identified as hypoattenuating areas containing less concentration of contrast material (11). Static CTP imaging occurs at or near the peak contrast opacification of the left ventricle and includes the acquisition of a single dataset. Dynamic CTP imaging requires sequential datasets during the initial pass of iodinated contrast from the venous to the arterial circulation. On both static and dynamic CTP imaging, regions of LV hypoperfusion will appear as low attenuation regions within an otherwise normal contrast distribution (i.e., perfusion), typically worse in the subendocardial layer than the epicardial layer.

A comprehensive myocardial CTP examination should include both rest and stress acquisitions to differentiate reversible from fixed myocardial perfusion defects (12).
Rest acquisition is derived from datasets obtained for coronary anatomy (13). It is less sensitive to the detection of myocardial ischemia since blood flow, and therefore perfusion is compromised when the stenosis is hemodynamically significant and exceeds 85–90% of the luminal diameter. A subendocardial or transmural hyperenhancement in the rest phase may indicate a critical flow-limiting coronary artery stenosis or a myocardial necrosis/scar from infarction.

Moreover, being derived from the CCT acquisition, it does not require further scans, thus decreasing the radiation dose delivered to the patient.

Besides, available post-processing software may allow for the generation of attenuation-based color mapping and a semi-quantitative assessment using a transmural perfusion ratio (TPR). TPR is just the ratio of the mean Hounsfield unit (HU) of a region of interest (ROI) within the subendocardial layer compared with the mean HU attenuation within the same ROI of the epicardial layer (Figure 1). This evaluation highlights the well-described aspects of an ischemic wave, which is worse in the subendocardial myocardial layers and gradually improves moving closer to the epicardial coronaries (i.e., creating a gradient). The use of TPR in static CTP significantly improves...
diagnostic accuracy compared to other techniques (14).

Other CT imaging signs of necrosis/fibrosis may be identified at the rest phase, such as myocardial wall thinning, lipomatous metaplasia (<10 HU), myocardial calcifications, aneurysmal/pseudoaneurysmal dilation, and mural thrombus.

These data highlight the significance of rest imaging in myocardial perfusion evaluation (12).

Acquisition techniques and image analysis

Static CTP imaging allows for qualitative evaluation of myocardial iodine contrast attenuation of a single acquisition. This image is taken during the arterial phase of first-pass contrast enhancement. The flow rate of injected contrast medium should be at least 5 mL/s, to provide for notable differences in contrast enhancement. It is of paramount importance the timing for first-pass single-phase stress CTP to increase diagnostic accuracy (15). Because the same acquisition can be used for the evaluation of coronary artery stenosis and myocardial blood supply, static CTP at rest can be achieved employing any CT scanner able to obtain diagnostic-quality CCT (16), hence being less technology-dependent.

Static CTP evaluation at rest is generally performed using CCT image datasets. Thus, no additional acquisition time or radiation is necessary. It can further be categorized into single- and dual-energy techniques.

Single energy acquisition

Generally, the CCT examination allows generating rest myocardial CTP imaging ECG-gating of coronary CTA or CTP can be retrospective but also prospective, which is another feature of most advanced multislice CT scanners (64 or more slices), providing a significant reduction in terms of radiation dose (less than 5 mSv), without any significant decreasing in image quality (17).

A prospective ECG-triggered high-pitch spiral acquisition performed with the 2nd-generation 128-slice dual-source CT (DSCT) scanner concedes the volumetric acquisition data of the heart in a single cardiac cycle with radiation dose as low as 1 mSv (18,19).

The analysis of myocardial perfusion is qualitative and consists in the identification of areas of hypodensity corresponding to areas of hypoperfusion. For myocardial perfusion analysis, multiplanar reconstructions should be carried out at 5 to 8 mm to improve the contrast-to-noise ratio (17). Myocardial contrast enhancement rises proportionally with iodine concentration, so perfusion defects appear as a hypodense area with subendocardial or transmural distribution concerning the normal myocardium (Figure 1).

Automated software providing modeling of the perfusion data into a 3D volume-rendering of the LV and a 17-segment polar map, are readily available.

Finally, automated software applications allow for evaluation of the TPR (an endocardial-to-mean epicardial ratio of attenuation values) using the 17-segment bulls s-eye scheme. However, this semi-quantitative index parameter seems not reliable in the presence of prior infarction or significant beam-hardening. To better distinguish between artifacts and true perfusion defects, myocardial perfusion in multiphase reconstructions are recommended. Artifacts may affect only a single cardiac phase, whereas true defects should persist throughout the entire cardiac cycle (16).

Data can be reconstructed with an iterative beam hardening correction (IBHC) algorithm to improve reliability. IBHC is a raw data-based beam hardening correction algorithm that raises image quality using 3D forward projection and exploits an additional two-compartment iodine/water model, to reduce beam hardening artifacts while having proper representation of anatomical structures (20).

The decisive move for the study of a CTP is the match of perfusion defects with the anatomic localization of coronary stenosis, which is crucial for understanding the hemodynamic significance of CAD (8,13).

Dual-energy acquisition

Dual-energy computed tomography (DECT) was first introduced in the late 1970s and provided for the differentiation of materials based on their X-ray attenuation at different tube voltages. The DECT technique permits the simultaneous acquisition of CT data using two different photon spectra (at 80/90/100 and 140/150 kV) of distinctly different mean energies within a single helical acquisition (21).

Based on the specific attenuation characteristics of the different tissues, DECT enables identifying the features of the tissue and evaluating the myocardial blood supply by mapping iodine distribution within the myocardium (22).

Different vendors propose specific CT technologies to perform dual-energy acquisitions. Dual X-ray source system (Siemens Healthineers) is the most commonly used technology: there are two independent tubes paired with two detectors that concurrently emit high (140–150 kV) and
A second modality is based on single-source CT with rapid (about 0.25 ms) switching of tube voltage between 80 and 140 kV either in a single gantry rotation (GSI Cardiac, GE Healthcare) or in sequential rotations (Acquilion One, Toshiba) (13,17).

Another method is represented by the dual-layer (“sandwich”) detector (Philips Healthcare). Two different materials can discriminate between low and high energy photons, with the source working at constant tube voltage; however, this system is not yet available in clinical practice (21).

DSCT-based dual-energy CT can generate an iodine distribution map. Because myocardial iodine concentration is a surrogate for myocardial perfusion, its volume reflects the myocardial blood pool (23). A color-coded map represents different myocardial territories and reflects iodine distribution in all segments, according to American Heart Association (AHA) classification (17 segments). Advanced post-processing workstations automatically compute perfusion maps, and then the relative normalized myocardial attenuation density and TPR are obtained.

DECT myocardial perfusion studies performed during rest or stress protocols showed good accuracy compared with various reference modalities (24-29).

Ruzsics et al. (25) studied the feasibility of rest myocardial perfusion imaging using a first-generation DECT system matched with SPECT in 36 patients and reported a sensitivity and specificity of 92% and 93%, respectively.

A study of 31 patients showed improved sensitivity and specificity in detecting CAD by supplying coronary CTA with DECT myocardial perfusion (29).

De Cecco and colleagues (30) reported a significant increase in specificity from 33% to 67% when DECT myocardial perfusion was combined with coronary CTA when compared with invasive coronary angiography (31-33) (Table 1).

Like the single-energy static technique, DECT generates a snapshot of the myocardial iodine distribution at a single time point. As with single-energy perfusion imaging, different myocardial regions may be scanned in different phases of the iodine contrast uptake, thus narrowing the comparability of myocardial iodine content among regions. An additional limitation is that the non-standardized assessment of the iodine distribution maps makes the evaluation highly user-dependent. Beam hardening artifacts can compromise the correct evaluation of iodine distribution maps that could affect areas of myocardial hypoperfusion and so false-positive findings. The development of new reconstruction algorithms aims to prevent these artifacts that affecting the evaluation of myocardial perfusion, such as the IBHC mentioned above.

### Clinical implementation and role

At present, a variety of protocols are available for myocardial perfusion imaging. Progress will reduce the radiation dose of myocardial CTP, such as low tube voltage imaging or new reconstruction algorithms, making it a more viable clinical option (7).

The rest static CTP technique has the potential to become the most widely used in the clinical setting, given the simplicity of execution, speed, and tolerability with a minimum dose delivery. In clinical routine, identifying normal coronary artery morphology and related normal myocardial perfusion may not require stress imaging. It

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**Table 1** Summary of main static perfusion studies with CCT

<table>
<thead>
<tr>
<th>Rest-static CTP studies</th>
<th>No. of subjects</th>
<th>CT technology</th>
<th>CT perfusion protocol</th>
<th>CT dose (mSv)</th>
<th>Reference technique</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PPV (%)</th>
<th>NPV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Koonce et al. (23)</td>
<td>36</td>
<td>1st-gen. DSCT</td>
<td>Rest</td>
<td>14</td>
<td>SPECT</td>
<td>92</td>
<td>93</td>
<td>83</td>
<td>97</td>
</tr>
<tr>
<td>Ko et al. (27)</td>
<td>45</td>
<td>1st-gen. DSCT</td>
<td>Rest</td>
<td>4.2</td>
<td>ICA</td>
<td>42</td>
<td>83</td>
<td>59</td>
<td>70</td>
</tr>
<tr>
<td>Wang et al. (29)</td>
<td>31</td>
<td>1st-gen. DSCT</td>
<td>Rest</td>
<td>10.5</td>
<td>SPECT, ICA</td>
<td>89</td>
<td>98</td>
<td>87</td>
<td>98</td>
</tr>
<tr>
<td>Kachenoura et al. (31)</td>
<td>64</td>
<td>64-MDCT</td>
<td>Rest</td>
<td>7–15</td>
<td>ICA</td>
<td>96</td>
<td>68</td>
<td>88</td>
<td>87</td>
</tr>
<tr>
<td>Osawa et al. (33)</td>
<td>145</td>
<td>2nd-gen. DSCT</td>
<td>Rest</td>
<td>14.8</td>
<td>ICA</td>
<td>85</td>
<td>94</td>
<td>79</td>
<td>96</td>
</tr>
<tr>
<td>George et al. (32)</td>
<td>24</td>
<td>64-MDCT</td>
<td>Rest</td>
<td>16.8</td>
<td>SPECT, ICA</td>
<td>86</td>
<td>92</td>
<td>92</td>
<td>85</td>
</tr>
</tbody>
</table>

1st and 2nd gen., generation of equipment; DSCT, dual source computed tomography; MDCT, multi-detector computed tomography; ICA, invasive coronary angiography; SPECT, single photon emission computed tomography; CCT, cardiac computed tomography; CTP, computed tomography perfusion; CT, computed tomography; PPV, positive predictive value; NPV, negative predictive value.
is also possible for those patients diagnosed with critical stenosis of one of the main coronary vessels with resting perfusion defects, where invasive coronary angiography is indicated rather than stress imaging. Several studies have suggested that resting imaging is adequate in the evaluation of myocardial perfusion (32,34). Myocardial infarction can be recognized through low myocardial attenuation areas on resting CT images; however, rest-only protocols have lower diagnostic accuracy than rest-stress protocols (35).

A variety of other methods have also been improved to evaluate the functional signal of coronary artery stenosis by CT imaging. These include the CT-derived fractional flow reserve (36) and the transluminal attenuation gradient of the coronary artery (37).

These methods have the benefit that they are derived from rest images without the necessity of additional imaging, contrast medium, or radiation exposure. Further work is needed to determine whether these methods are better for myocardial perfusion imaging (7).

The main advantage of CT imaging is that accurate information on the coronary arteries can also be obtained. It is also helpful for patients where magnetic resonance imaging (MRI) is contraindicated. CTP imaging can be performed as an add on test after CTA if there is diagnostic doubt or it can be planned for patients with known or suspected heavy calcification or coronary artery stents. Also, CTP imaging can be performed in order to aid the planning of interventions.

### Radiation dose

When considering radiation exposure, static CTP imaging is chosen over dynamic techniques.

The radiation dose for complete static CTP studies varies greatly depending on the specific acquisition protocol. Useful dose values between 2.5 and 21 mSv have been reported with frequent radiation exposure of 12 mSv. Combined dynamic rest and stress CTP studies can require radiation exposure of approximately 18 mSv (2). Also, from standard CCT acquisition, it is possible to obtain rest static CT myocardial perfusion images; this is not the case with dynamic imaging. In this case, no further dose is delivered to the patient.

Moreover, many factors may affect the radiation dose, such as the patient’s biometrical parameters [body mass index (BMI), cardiac output, and heart rate], the type of CT equipment accessible, and the CT protocols used, which have to be tailored to the patient (17).

Current applications are directed to reduce radiation exposure while maintaining high diagnostic performance. In this interest, the use of recent technical findings, including the low voltages (70 to 80 kV) acquisition, automated tube current modulation, and iterative reconstruction, seems to be able to achieve this ambitious goal (13).

### Conclusions

Current experiences suggest that adding perfusion imaging to the standard morphological imaging performed with CTA improves diagnostic accuracy and positive predictive value. It not only provides data on coronary artery atherosclerotic plaques but also functional and hemodynamic information on the myocardium. In various CTP protocols, the evaluation of perfusion is qualitative, quantitative, or semi-quantitative. Implementing this aspect could represent the turning point in the clinical routine to better stratify the patient based on risks and results obtained.

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