On-Site Computed Tomography-Derived Fractional Flow Reserve Using a Machine-Learning Algorithm
— Clinical Effectiveness in a Retrospective Multicenter Cohort —

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Background: This study evaluated the diagnostic capability of on-site coronary computed tomography-derived computational fractional flow reserve (CT-FFR) determinations for detecting coronary artery disease (CAD), as assessed by invasive fractional flow reserve (FFR).

Methods and Results: Seventy-four patients with coronary artery calcium scores <1,500 who underwent coronary CT angiography (CTA) and invasive FFR measurements within 90 days were retrospectively reviewed. CT-FFR was computed using a prototype machine-learning (ML) algorithm in 91 vessels; 47 vessels of 42 patients were determined to have significant CAD (FFR ≤0.8). Correlation between CT-FFR and FFR was good (r=0.786, P<0.001). Per-vessel area under the curve was significantly larger for CT-FFR (0.907, 95% confidence interval: 0.828–0.958) than for CTA stenosis ≥50% (0.595, 0.487–0.697) or ≥70% (0.603, 0.495–0.705) (both P<0.001). Standard coronary CTA classifications recommended further functional tests in 57 patients with moderate or worse stenosis on CTA. CT-FFR analysis (mean analysis time: 16.4±7.5 min) corrected the standard coronary CTA classification in 18 of 74 patients and confirmed it in 45 of 74 patients. Thus, the per-patient diagnostic accuracy of the classifications was improved from 66% (54–77%) to 85% (75–92%).

Conclusions: On-site CT-FFR based on a ML algorithm can provide good diagnostic performance for detecting hemodynamically significant CAD, suggesting the high value of coronary CTA for selected patients in clinical practice.

Key Words: Artificial intelligence; Computed tomography; Coronary artery disease; Fractional flow reserve; Machine learning

With rapid technological advances in multidetector row computed tomography (CT), coronary CT angiography (CTA) is widely used in clinical practice as a reliable diagnostic tool for patients with stable coronary artery disease (CAD). Coronary CTA is highly effective for ruling out CAD, with a high negative predictive value (NPV). However, when stenosis severity is assessed using coronary CTA, it does not always correlate with the severity of myocardial ischemia, and current guidelines recommend further cardiac investigations in addition to coronary CTA for moderate or worse coronary artery stenosis. Fractional flow reserve (FFR) has been established as an invasive standard reference for the assessment of myocardial ischemia. When lesion-specific ischemia is assessed by FFR during invasive coronary angiography (ICA), the findings have a great impact on the practical management of patients with CAD, by assisting decision-making and procedural planning for revascularization therapy.

Coronary CTA-derived computational FFR (CT-FFR) determination is a new diagnostic method that uses information obtained from the resting coronary CTA dataset and computational flow dynamics (CFD). Previous studies have shown that CT-FFR has good correlation with invasively measured FFR and improves the diagnostic differentiation between hemodynamically significant CAD and coronary CTA alone, using either a remote service or on-site...
software. Recently, machine-learning (ML)-based CT-FFR applications have been developed. Single- and multicenter trials have reported the diagnostic performance of ML-based CT-FFR simulations for detecting hemodynamically significant CAD. However, the clinical value of on-site ML-based CT-FFR has not been fully investigated. Thus, this study compared the clinical and decision-making value of on-site ML-based CT-FFR with conventional CTA for managing patients with hemodynamically significant CAD. CT-FFR was also compared with invasively measured FFR as a reference standard for myocardial ischemia.

Methods

Study Population
This study was approved as a retrospective multicenter observational study by the review board of each institution, and the need for informed consent was waived. The inclusion criteria included (1) age of ≥20 years, (2) maximum interval of 90 days between coronary CTA and invasive FFR measurement, and (3) adequate image quality of coronary CTA. The exclusion criteria included (1) patient refusal to participate, (2) arrhythmia inappropriate for ECG-triggering data acquisition for coronary CTA, (3) poor image quality on coronary CTA (artifact, calcification, and misregistration) for CT-FFR analysis, (4) coronary artery calcium (CAC) Agatston score ≥1,500, (5) prior revascularization, (6) myocardial infarction (MI) (based on clinical information, ECG, echocardiography, and tissue characterization of myocardium using the coronary CTA dataset), (7) total coronary artery occlusion, (8) coronary anomalies (aneurysm or dissection), (9) presence of collateral flow, (10) small-vessel disease and severe, diffuse stenosis (<1.5 mm), (11) low cardiac function (left ventricular [LV] ejection fraction <30%), (12) heart valve disease (moderate or severe aortic stenosis and regurgitation), (13) cardiomyopathy, (14) uncompensated congestive heart failure, (15) implanted intracardiac metallic device (pacemaker, implantable cardioverter defibrillator, prosthetic valve, and sternal clip) and history of cardiovascular surgery, and (16) major cardiovascular event (MI, worsening angina, and hospitalization for heart disease or stoke) during the study period.

Coronary CTA
Two dual-source CT scanners (SOMATOM Force or SOMATOM Definition Flash, Siemens Healthineers, Forchheim, Germany) were used according to the institutional scan protocol. First, a CAC score scan was performed using a prospective ECG-triggering axial scan mode (tube voltage, 120 kV; tube current time product, 80 mA/rotation for a 70-kg body weight as a reference, 3-mm slice thickness). Sublingual nitroglycerin (0.6 mg, Myocor; Astellas Pharma, Tokyo, Japan) was used before the coronary CTA examination, but no additional β-blockers or other heart rate-modifying medication were administered. Coronary CTA was performed using a retrospective ECG-triggering spiral scan mode or prospective ECG-triggering axial scan mode according to the scan heart rate. The scan parameters were as follows: gantry rotation time of 0.25 s/rotation (force), 0.28 s/rotation (definition flash); 90-, 100- or 120-kV tube voltage; 2×192×0.6 mm (force) or 2×128×0.6 mm (definition flash) detector collimation with z-flying focal spot; tube current 228 (145–292) mA per rotation with X-ray tube modulation that depended on patient size, variable pitch, and heart rate. Contrast material (iopamidol, 370 mg iodine/mL; Bayer Yakuhin, Osaka, Japan) was intravenously administered for 15 s (40–60 mL), followed by a 20-mL saline chaser. The scan timing was optimized using bolus tracking or timing bolus scan according to the institution’s scan protocol.

ICA and FFR
ICA was performed using 6- or 7-Fr coronary catheters. Isosorbide dinitrate was given via catheter before angiography and FFR measurements. Invasive FFR measurements were performed by 3 experienced interventional cardiologists for clinical indications to assess whether the CAD lesions were causing hemodynamically relevant myocardial ischemia. An FFR pressure wire (Verrata, Phillips, MA, USA, or AERIS, Abbott, CA, USA) was placed distal to the stenosis of interest. The invasive FFR was measured during hyperemia induced by intravenous infusion of adenosine triphosphate at a rate of 0.14–0.16 μg·kg⁻¹·min⁻¹ via the forearm or femoral vein as previously reported. An FFR ≤0.80 was considered as hemodynamically significant CAD.

For the anatomic locations of the CT-FFR measurements, 3 experienced cardiologists, who were blinded to other functional tests and coronary CTA stenosis severity, transferred the corresponding landmarks for invasive FFR sample locations to the volume-rendered images of coronary CTA with reference to the FFR wire on the fluoroscopic images.

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CT-FFR Analysis
CT-FFR calculations were performed on the coronary CTA datasets using prototype research software (cFFR version 3.0.0, Siemens Healthcare) that uses artificial intelligence (AI) ML. During CT-FFR preparation, a 3D anatomic model of the coronary arteries (luminal model) was semi-automatically created from the coronary CTA dataset by tracing the coronary artery centerline and lumen. The anatomic positions of the coronary stenoses were defined for the CT-FFR computation and then the LV mass was defined using a threshold value from the attenuation of the LV myocardium. The CT-FFR computation provided a coronary tree of color-coded CT-FFR values, with red representing CT-FFR values ≤0.80.

Two experienced radiologists (with 4 and 16 years’ experience, respectively, in cardiac imaging) who were blinded to other results, performed the CT-FFR analysis. First, they were involved in a pilot study to assess the interobserver variance for CT-FFR calculations for 20 independently selected patients who were not included in the study population. Next, they carefully prepared for the process of CT-FFR calculation by setting the coronary artery centerline and lumen according to the anatomic positions of the stenosis in a semi-automated manner. The window settings (Hounsfield units, HU) were adjusted per-patient and fixed up to the completion of the 3D coronary artery model (window level: 187 HU, 173–193 HU; window width 868 HU, 778–967 HU). To obtain the lesion-specific CT-FFR results, a third observer (with 15 years’ experience in cardiac imaging) deployed the sample points to the color-coded CT-FFR trees and obtained the CT-FFR values, according to the landmark images previously described. Processing time for editing the centerline and the lumen of the coronary arteries and the calculation time for the virtual FFR simulation were measured and added to determine the total processing time.

Statistical Analysis
Categorical variables are expressed as proportions and continuous variables are expressed as mean±standard deviation or median (interquartile range), as appropriate. The interobserver variabilities were calculated using a weighted kappa statistic and interclass correlation coefficient. Correlation and differences between CTA-FFR and invasive FFR were evaluated with the Spearman’s test and the Bland-Altman test.

Coronary CTA was assessed using both a threshold of 50% and 70% luminal stenosis. A cutoff value ≤0.80 was regarded as hemodynamically significant for both CT-FFR and invasive FFR. On a per-vessel basis, the diagnostic performance of anatomic coronary CTA assessments (≥50% and ≥70% luminal stenosis) and CT-FFR (≤0.80) to detect vessels with invasive FFR ≤0.80 is reported as sensitivity, specificity, positive and negative predictive values (PPVs and NPVs), and accuracy, with their 95% confidence intervals (CIs). Sensitivity and specificity were compared using the McNemar test with Bonferroni correction. Diagnostic accuracy was compared using the area under the curve (AUC) of the receiver-operating characteristic curve analysis. The AUCs were compared using the method described by DeLong et al. On a per-patient basis, patients with at least 1 diseased vessel confirmed by invasive FFR were classified as disease positive and those without any diseased vessels were classified as disease negative. Similar analyses were conducted regarding diagnostic performance and the AUC analysis.

To investigate the clinical value of CT-FFR compared with conventional anatomic CTA-based diagnosis and recommendations (CAD-RAD™ classification) for the indication of patients requiring further evaluation for CAD
Study Population

Figure 1 is a flowchart of patient selection. Initially, 206 patients were collected from collaborating facilities using the inclusion criteria. Of them, data for 128 patients were excluded for various reasons related to patient factors, including the ruling out of invasive FFR assessment, CT image quality factors, and CT-FFR analysis-related factors. In this study, regardless of the target or non-target lesions for CT-FFR analysis, 50 patients with poor image quality coronary CTA (artifact, n=5; calcification, n=24; compound reasons, n=21) and 41 patients with a history of MI or revascularization were excluded. Finally, 74 patients with known CAD or suspected of CAD (56 males; mean age, 70.2±10.3 years) were eligible for analysis. Patients’ characteristics are shown in Table 1. No major cardiovascular events were seen during the study period, which was a median of 31 days (17–47 days) between coronary CTA and invasive FFR measurement. A total of 91 vessels were investigated using a pressure wire. The overall median invasive FFR was 0.80 (0.72–0.86), and 47 vessels (52%) were diagnosed with hemodynamically significant stenosis (FFR ≤ 0.8). Thus, 42 patients (57%) were deemed to have hemodynamically significant stenosis.

Coronary CTA

The median scan heart rate during coronary CTA was 59 (56–67) beats/min. The median CAC Agatston score was 180 (58–510). The median total radiation dose was 7.4 mSv (5.7–11.2 mSv), by summing the CAC score scan and coronary CTA.

Regarding the interobserver variability, the weighted interobserver kappa score was 0.740 (95% CI, 0.601–0.880) for CT image quality and 0.692 (95% CI, 0.414–0.97) for coronary CTA stenosis. The reproducibility was satisfactory. Median CT image quality on a per-patient basis was 3 (2–4).

Results

caused by moderate stenosis or worse stenosis, patients were classified into 3 groups: (1) patients in whom CT-FFR corrected the CTA-based recommendations, (2) patients in whom CT-FFR supported the CTA-based recommendations, and (3) patients in whom CT-FFR evaluation was not effective for the CTA-based recommendations. Subgroups stratified by anatomic CTA stenosis (<50%, 50–69%, and ≥70%) were compared using Fisher’s exact test.

All analyses were performed using statistical software (SPSS version 20.0, IBM, Armonk, NY, USA; MedCalc version 17.9.7, Ostend, Belgium). A 2-tailed P-value <0.05 was deemed significant.

Table 1. Patients Characteristics

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Value</th>
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<tbody>
<tr>
<td>Age (years)</td>
<td>70.2±10.3</td>
</tr>
<tr>
<td>Male/female</td>
<td>56 (76)/18 (24)</td>
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<tr>
<td>Chest symptoms</td>
<td>57 (77)</td>
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<tr>
<td>Coronary risk factors</td>
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<tr>
<td>Hypertension</td>
<td>57 (77)</td>
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<tr>
<td>Hyperlipidemia</td>
<td>40 (54)</td>
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<tr>
<td>Diabetes mellitus</td>
<td>28 (38)</td>
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<tr>
<td>Smoking</td>
<td>23 (31)</td>
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<tr>
<td>Family history of CAD</td>
<td>11 (15)</td>
</tr>
<tr>
<td>CT and FFR inspection period (days)</td>
<td>31 (17–47)</td>
</tr>
<tr>
<td>CAC Agatston score</td>
<td>180 (58–510)</td>
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</table>

Data are presented as n (%), mean±standard deviation or median (interquantile range), unless otherwise stated. CAC, coronary artery calcium; CAD, coronary artery disease; CT, computed tomography; FFR, fractional flow reserve.

Figure 2. Correlation and difference between CTA-FFR and invasive FFR. (A) Correlation of CTA-derived fractional flow reserve (CT-FFR) and invasive FFR on per-vessel basis. There is a moderate Spearman’s correlation coefficient (r=0.786). (B) Bland-Altman plot of CT-FFR and invasive FFR on per-vessel basis. The mean difference between CTA-FFR and invasive FFR was –0.02±0.10. A line is placed at the mean difference value (–0.02) and the corresponding double standard deviation intervals (–0.22 and 0.18). CTA, coronary CT angiography; FFR, fractional flow reserve.
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Of the 91 vessels, coronary CTA depicted 2 minimal stenosis, 18 mild stenoses, 53 moderate stenoses, and 18 severe stenoses. Thus, there were 17 patients with mild stenosis or less, 40 patients with moderate stenosis, and 17 patients with severe stenosis.

**CT-FFR**

For CT-FFR analysis, the mean analysis time per-patient was 16.4\pm 7.5 min (sum of preparation for the 3D coronary artery anatomic model [16.2\pm 7.4 min] and the CT-FFR calculation [16.3\pm 9.9 s]). The overall median CT-FFR was 0.80 (0.68–0.86), and hemodynamically significant stenosis on CT-FFR (≤0.8) was present in 53 vessels (58%) in 45 patients (61%).

Correlation between ML-based CT-FFR and invasive FFR was moderate (P<0.001, r=0.786; **Figure 2A**), with a lower mean outcome for CT-FFR compared with invasive FFR of -0.03\pm 0.10 (Figure 2B).

**Diagnostic Performance of CT-FFR and Coronary CTA**

The diagnostic characteristics of coronary CTA and CT-FFR are shown in Table 2. On a per-vessel basis, for detecting hemodynamically significant CAD (FFR ≤0.8), sensitivities of CTA stenosis ≥50% (87%; 95% CI, 74–95%) and CT-FFR ≤0.8 (89%; 95% CI, 77–96%) were higher than that of CTA stenosis ≥70% (30%; 95% CI, 17–45%) (all P<0.05). Specificity of CTA stenosis ≥70% (91%; 95% CI, 78–98%) and CT-FFR ≤0.80 (75%; 95% CI, 60–87%) were higher than that of CTA stenosis ≥50% (34%; 95% CI, 21–50%) (P<0.05, for each). The AUC to differentiate hemodynamically significant CAD with CT-FFR ≤0.80 (0.907; 95% CI, 0.828–0.958) was greater than those of CTA stenosis ≥50% (0.595; 95% CI, 0.487–0.738) or ≥70% (0.603; 95% CI, 0.495–0.705) (all P<0.001).

On a per-patient basis, sensitivity improved from 33% (95% CI, 20–50%) by CTA stenosis ≥70% to 91% (95% CI, 77–97%) by CT-FFR (P<0.05), and specificity also improved from 38% (95% CI, 24–59%) by CTA stenosis ≥50% to 78% (95% CI, 60–91%) by CT-FFR (P<0.05). The AUC of CT-FFR (0.913; 95% CI, 0.825–0.966) was significantly greater than that of CTA stenosis ≥50% (0.628; 95% CI, 0.508–0.738) or ≥70% (0.620; 95% CI, 0.499–0.730) (both P<0.001).

Regarding the clinical value of on-site ML-based CT-FFR, we evaluated the effectiveness of CT-FFR for standard CTA-based diagnosis and recommendations. According to the CAD-RADSTM classification, further functional tests or ICA was indicated in 57 patients with moderate stenosis or worse, and risk factor modification was indicated in 17 patients with mild stenosis or better. However, additional

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**Table 2. Diagnostic Performance of Coronary CTA and CT-FFR on the Per-Vessel and Per-Patient Levels**

<table>
<thead>
<tr>
<th></th>
<th>Sensitivity, %</th>
<th>Specificity, %</th>
<th>PPV, %</th>
<th>NPV, %</th>
<th>Accuracy, %</th>
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<tbody>
<tr>
<td><strong>All vessels (n=91)</strong></td>
<td></td>
<td></td>
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<tr>
<td><strong>All patients (n=74)</strong></td>
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Data are expressed as n/N, %, and 95% confidence interval, unless otherwise stated. Diagnostic characteristics were assessed for detecting coronary lesions with invasive FFR ≤0.80 as reference. CTA, computed tomography angiography; CT-FFR, computed tomography-derived fractional flow reserve; PPV, positive predictive value; NPV, negative predictive value.
Figure 4. Dataset of a man in his 60s, who presented with exertional angina. Coronary computed tomography angiography (CTA) and a snapshot of CT-derived fractional flow reserve (CT-FFR) analysis for 3D coronary tree image show moderate stenosis (yellow arrow) with partially calcified plaque in the left anterior descending artery (LAD) (A, B), and severe stenosis (red arrow) with non-calcified plaque in the right coronary artery (RCA) (C, D). CT-FFR color-coded map (E) shows a computational FFR of 0.59 at the invasive FFR measurement pressure wire location (white circle) in the LAD, and that of 0.50 in the RCA. During invasive coronary angiography (ICA), FFR measured at the pressure wire location (white circle) was 0.61 distal to the mild stenosis (yellow arrow) on ICA in the LAD (F, G), and 0.62 distal to the moderate stenosis (red arrow) on ICA in the RCA (H, I). He underwent staged percutaneous coronary interventions performed separately for double-vessel disease.

Figure 5. Results of standard coronary CTA assessment, CT-FFR, and invasive FFR. Additional analysis using on-site CT-FFR effectively corrected clinical decision-making made by the standard classification with coronary CTA in 18 patients (red), and supported the decision-making in 45 patients (blue). It did not provide effective information in 11 patients (gray) for the detection of hemodynamically significant coronary artery disease (FFR ≤0.8). CTA, coronary computed tomography angiography; FFR, fractional flow reserve.
assessing with CT-FFR corrected the CAD-RAD\textsuperscript{TM} classification in 18 patients and supported the decision-making in 45 patients (Figures 3.4) or risk factor modification. It did not provide effective information in 11 patients (Figure 5). No significant difference in the effectiveness of CT-FFR was seen among subgroups stratified by coronary CTA stenosis severity (P=0.279992) (Table 3).

### Discussion

The main findings of this study were: (1) the prototype on-site ML-based CT-FFR computation showed good correlation with invasively measured FFR, (2) CT-FFR had high diagnostic performance for the detection of hemodynamically significant CAD on a per-vessel and per-patient basis, and (3) on-site ML-based CT-FFR analysis increased the value of coronary CTA with a short processing time. Based on numerous studies and technological developments, clinical use of coronary CTA has developed from appropriate use criteria to standard guidelines for coronary CTA reporting and recommendations. Meanwhile, because of the relatively low specificity and PPV of semiquantitative coronary CTA assessment, functional assessment is often required to further evaluate the patient.\textsuperscript{22} In current clinical practice, non-invasive myocardial perfusion imaging and invasive FFR measurement have been used for assessing the indication of revascularization therapy and for prediction of various outcomes.\textsuperscript{23–27} However, the additional cost, associated radiation exposure, and diagnostic throughput hamper their use in current practice.

CT-FFR has rapidly established as a new method, with multicenter trials showing that it has good diagnostic capability for detecting hemodynamically significant CAD and providing incremental value with respect to diagnosis, controlling clinical costs, and predicting outcomes.\textsuperscript{10–12,28–30} The Heartflow CFD-based FFR software works using AI with algorithm-based, fully-automated 3D coronary luminal modeling and CFD-based FFR calculations. Nevertheless, because these studies utilized a remote analysis service for CT-FFR computations, clinical application of CT-FFR might not have been fully maximized. The currently available work-in-progress on-site CT-FFR software works on a regular workstation using semi-automated 3D coronary artery modeling and 1 of the following algorithms: (1) the reduced-order CFD-based FFR calculations,\textsuperscript{13,14} (2) CFD-based FFR calculations with an AI algorithm,\textsuperscript{15} or (3) ML-based CT-FFR calculations.\textsuperscript{17–19} The most recently developed on-site ML-based CT-FFR algorithm was developed using a deep learning model to integrate the complex relationship between various anatomic features and the CFD-based FFR results as the ground truth from a synthetically generated database of 12,000 coronary artery models. The ML-based CT-FFR software extracts 28 anatomic input features and calculates the CT-FFR results.\textsuperscript{14} It allows for the quantification of ischemic severity through color-coding of the coronary artery tree for all the analyzed coronary arteries, with greatly shortened CT-FFR calculation time. Single- and multicenter trials have shown comparable diagnostic performance and incremental values for the method in comparison with CFD-based CT-FFR.\textsuperscript{11,15,17–19}

The present study also showed that the on-site ML-based CT-FFR had a high diagnostic capability for detecting hemodynamically significant CAD using a local workstation, and it could immediately provide seamless and effective diagnostic information after standard coronary CTA assessment. Although the CT-FFR analysis was carefully performed by 2 relatively experienced operators independent of the time-cost study, per-patient total processing time (16.4±7.5 min) was clinically reasonable compared with previous studies.\textsuperscript{13–15,19} Additional assessment of CT-FFR identified 4 patients (25%) with hemodynamically significant CAD out of 17 patients with mild stenosis or less on CTA, and ruled out 14 patients (25%) without hemodynamically significant CAD out of 57 patients with moderate or severe stenosis on CTA. Although we thought that clinical effectiveness of CT-FFR analysis compared with standard coronary CTA assessment might vary among subgroups stratified by stenosis severity on coronary CTA, the present results did not show a statistically significant difference.

The pros and cons of clinical application of on-site CT-FFR are currently being discussed. The positive viewpoint suggests that on-site CT-FFR can provide timeliness in clinical practice and promote understanding of the importance of high-quality coronary CTA, and the underlying algorithm and requirements for CT-FFR computations, including the accurate 3D coronary model. The negative viewpoint suggests that CT-FFR may be used incorrectly. Because of the semi-automated post-processing involved in the CT-FFR analysis, such as centerline and coronary lumen tracing, a training period would be required.

### Study Limitations

First, the study population was retrospectively assessed and was limited to patients with good-quality coronary CTA imaging and this excluded a large number of candidates (24%, 50 of 206 cases for poor image quality) from this study, for accurate 3D coronary artery modeling. Thus, for more complex cases, the results may not always be clear as in this study. Second, conventional coronary CTAs were evaluated using the most severe lesions. More dedicated or cumulative assessments corresponding to the landmarks of invasive FFR sample locations might be necessary. Third, the indications for invasive FFR measurement were clinically determined by attending physicians, and therefore,

### Table 3. Effectiveness of CT-FFR to Coronary CTA-Based Decision-Making

<table>
<thead>
<tr>
<th></th>
<th>CT-FFR corrected CTA-based decisions (a)</th>
<th>CT-FFR supported CTA-based decisions (b)</th>
<th>CT-FFR ineffective to CTA-based decisions (c)</th>
<th>Effectiveness, % (a+b) / (a+b+c)</th>
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<tbody>
<tr>
<td>&lt;50% CTA stenosis (n=17)</td>
<td>4</td>
<td>11</td>
<td>2</td>
<td>15/17 (88%)</td>
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<tr>
<td>50–69% CTA stenosis (n=40)</td>
<td>12</td>
<td>20</td>
<td>8</td>
<td>32/40 (80%)</td>
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<tr>
<td>≥70% CTA stenosis (n=17)</td>
<td>2</td>
<td>14</td>
<td>1</td>
<td>16/17 (94%)</td>
</tr>
<tr>
<td>Total</td>
<td>18</td>
<td>45</td>
<td>11</td>
<td>63/74 (85%)</td>
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</table>

Abbreviations as in Table 2.
selection bias may have affected the results. Application of the present results to extremely minimal or severe CTA stenosis, as well as more complex cases with massive calcified lesions, should be performed with caution. Fourth, the present study population was not prescribed additional \( \beta \)-blockers to reduce scanned heart rates. Fifth, cases of MI were clinically excluded. Because the CT-FFR calculation assumes the LV volume obtained from CT as the mass of normal LV myocardium, unrecognized or subendocardial MI might result in a mismatch between the CT-FFR computation and invasive FFR measurement. Thus, both image quality and the various potential factors affecting CT-FFR will require further evaluation.\(^3\) Sixth, on-site CT-FFR analysis requires objectivity and independence. It is desirable that a well-trained single operator can perform standard coronary CTA evaluation and CT-FFR preparation comprising the generation of 3D coronary artery model and color-encoding with CT-FFR values by the CT-FFR calculation. Additionally, the deployment of the sample size was standardized according to invasive FFR or other evidence-based basis (2–3 cm distal to the stenosis).\(^2,3\)\(^1\) blinded to all other patient information except for point-of-care diagnosis. Seventh, ML-based CT-FFR results were not able to be calculated in a small number of patients (5% of the analysis). The possible reasons were investigated by the vendor scientific research group and will be used to improve the present work-in-progress software. Lastly, this study focused on diagnosis, but larger multicenter trials of on-site CT-FFR that investigate interventions to therapeutic strategies, and prognosis are required.\(^3,4,15\)

**Conclusions**

In conclusion, CT-FFR can be assessed using an on-site workstation. ML-based CT-FFR analysis can provide good diagnostic performance for detecting hemodynamically significant CAD during clinically relevant analysis time. Diagnostic factors and various outcomes should be assessed in further clinical investigations.

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**Disclosures**

The authors declare no conflicts of interests. No funding was received for this research.

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