

Kardiologia Polska

The Official Peer-reviewed Journal of the Polish Cardiac Society since 1957

Online first

This is a provisional PDF only. Copyedited and fully formatted version will be made available soon

ISSN 0022–9032 e-ISSN 1897–4279

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Article type: Review

Received: July 11, 2024 Accepted: July 30, 2024

Early publication date: July 31, 2024

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When to perform percutaneous coronary interventions in TAVI patients? Recent

advances

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ABSTRACT

Coronary artery disease (CAD) is prevalent in approximately 50% of patients with severe aortic

valve stenosis undergoing transcatheter aortic valve implantation (TAVI). The impact of CAD

on TAVI outcomes and optimal management strategies remains unclear.

This manuscript reviews the latest evidence on assessing and determining the timing for

treating CAD in TAVI patients to optimize clinical outcomes and resource utilization.

We discuss the current methods for CAD diagnosis, including invasive coronary angiography

(ICA), coronary computed tomography angiography, and the role of functional assessment

indices like fractional flow reserve and instantaneous wave-free ratio in guiding

revascularization decisions. While ICA remains the standard for determining CAD severity in

TAVI candidates, coronary computed tomography angiography has shown potential in

reducing unnecessary ICA procedures. When indicated, fractional flow reserve seems more

reliable than instantaneous wave-free ratio in a rtic valve stenosis patients, particularly when

2

evaluated post-TAVI. Recent data suggest that percutaneous coronary intervention post-TAVI may be associated with improved outcomes compared to pre-TAVI interventions.

Concluding, the optimal management of CAD in TAVI patients is still under investigation. Current evidence supports a tailored approach, considering both pre- and post-TAVI percutaneous coronary intervention strategies based on individual patient characteristics and procedural complexities. Further randomized trials are needed to establish definitive guidelines.

Key words: aortic valve stenosis, coronary artery disease, coronary revascularization, fractional flow reserve, instantaneous wave-free ratio, transcatheter aortic valve implantation

INTRODUCTION

The prevalence of coronary artery disease (CAD) in patients with severe aortic valve stenosis (AVS) undergoing transcatheter aortic valve implantation (TAVI) is approximately 50% [1–4], with variations ranging from 15% to 80% in major clinical trials based on patients' age and risk profile [5–9]. Notably, CAD prevalence is lower in low-risk patients compared to those at intermediate and high risk. Among TAVI recipients with CAD, about half have multivessel disease, and there is often involvement of the left main (LM)/left anterior descending artery (LAD) [10, 11].

Early observational studies conducted shortly after the introduction of TAVI produced mixed results regarding the impact of coexisting CAD on patient outcomes. Some studies found no significant difference in survival between patients with and without significant coronary lesions [12, 13], or between those who underwent complete versus incomplete revascularization, while other studies reported the opposite findings [14]. Additionally, higher rates of procedural complications have been observed when percutaneous coronary intervention (PCI) is performed concurrently with TAVI, potentially leading to adverse outcomes in this vulnerable patient population [15]. However, it is important to note that most studies had a limited follow-up period of less than 2 years, and a longer duration may be necessary to accurately assess the impact of CAD on clinical outcomes post-TAVI.

Moreover, there is no consensus on how to evaluate the clinical relevance of identified stenoses in AVS patients with CAD, nor on the timing of eventual revascularization in this setting, with many operators relying solely on angiographic severity to guide PCI decisions.

Currently, these specific issues have not been definitively addressed by the scientific community, and the management of CAD is often left to the experience of individual operators or centres. This manuscript aims to review the latest evidence on the assessment and the potential timing strategies for the treatment of CAD in patients undergoing TAVI, with the goal to optimize patients' outcomes and healthcare resource utilization (Figure 1).

CAD SEVERITY ASSESSMENT IN PATIENTS WITH SEVERE AORTIC STENOSIS: INDICATION FOR PCI

Given the fragility of the population being examined, the primary goal is to achieve diagnosis while conserving resources, time, and minimizing the risk of complications. This approach will enhance patient comfort and optimize the use of medical resources. There are several potential methods to investigate the presence and significance of CAD: invasive coronary angiography (ICA), coronary computed tomography angiography (CCTA), or non-invasive ischemia tests. However, TAVI candidates are often not ideal for non-invasive ischemia tests due to their fragility, which prevents them from performing physical stress tests, and the frequent presence of left ventricular (LV) hypertrophy, which can confound results.

Given that TAVI patients necessarily require an ECG-gated computed tomography (CT) prior to TAVI, some authors have suggested using this imaging technique to select patients for ICA based on CT results. A dedicated study showed that this hybrid strategy is feasible and potentially clinically relevant, with CA performed in only a quarter of patients due to the detection of obstructive stenosis on CT, without affecting clinical outcomes when ICA was deferred [16]. As expected from experiences with non-AVS patients, compared to ICA, CCTA has excellent negative predictive value and sensitivity, but low specificity, which decreases further in the presence of previous stented segments or heavy calcifications [17]. Despite these limitations, a recent large meta-analysis also demonstrated that using CT as a gatekeeper for ICA in the TAVI work-up could reduce the number of coronary angiographies by 37%. With an increasing number of low-risk TAVI patients, this reduction is likely to grow, given the lower probability of CAD and calcified lesions in younger patients [18].

However, to date, ICA remains the standard examination for determining the presence and severity of CAD in TAVI candidates. The current European Society of Cardiology guidelines [19] state that myocardial revascularization using PCI should be considered in patients with a primary indication to undergo TAVI and presenting, based on angiography alone, with coronary artery diameter stenosis >70% in proximal segments, although this recommendation has a low level of evidence (C). In contrast, no recommendations are provided

for non-proximal or less severe stenosis. Notably, American Guidelines indicate that investigation of intermediate CAD through invasive physiological assessment is safe even in the presence of AVS [20].

Although an angiographic stenosis >70% seems a reliable threshold for detecting critical stenosis in patients with aortic stenosis [21, 22], coronary physiology studies suggest that 20% of these cases have a normal fractional flow reserve (FFR), a percentage that increases dramatically (up to 65%) with 50%–70% stenosis [23]. While this is well-documented for non-AVS patients, similar conclusions cannot be drawn for AVS patients, where morphological and hemodynamic changes induced by the valve disease may affect functional indexes [24].

Theoretically, LV hypertrophy with interstitial fibrosis, commonly found in AVS, increases LV end-diastolic pressure and induces microvascular dysfunction, both potentially leading to blunted vasodilatory capacity and increased "back pressure" (Pd), resulting in altered FFR values. Although instantaneous wave-free ratio (iFR) does not require pharmacological vasodilation, it may also be affected by LV end-diastolic pressure. High left ventricular filling pressures impact coronary resting flow by increasing myocardial metabolic demand. It has been reported that increasing resting flow reduces the pressure drop along the target vessel, resulting in falsely higher iFR values [25].

Additionally, LV obstruction reduces systemic pressure and consequently the pressure upstream of the stenosis, causing a concomitant decrease in Pa. Based on this hypothesis, when using FFR or iFR, we might expect an increased Pd due to impaired vasodilatory response to adenosine or higher resting flow, which combined with lower Pa, could result in falsely higher values.

However, comparing FFR and iFR with myocardial perfusion imaging (stress single photon emission computed tomography [SPECT]) in AVS patients, it has been shown that these functional indexes perform relatively well. FFR, in particular, had the best agreement with SPECT (85%), with an area under the curve 0.91 and negative predictive value 95% for detecting ischemia. On the other hand, iFR produced a significant proportion of false positives (39% of negative SPECT) using the standard cut-off ≤0.89, while using a pre-specified 0.82 cut-off improved iFR agreement with SPECT to 73% [26].

This finding was further confirmed in another study, which showed that the conventional iFR cut-off had lower diagnostic agreement with FFR classification of coronary lesions in the presence of AVS, compared to non-AVS patients. According to the data, the best iFR cut-off for predicting FFR \leq 0.8 was lower (0.83) than the standard one (0.89) [27, 28].

Further studies have investigated the variability of FFR and iFR after TAVI, once LV obstruction was removed, reflecting a more physiologic condition. Regarding FFR, only minor variations were observed after TAVI compared to baseline. Notably, the direction of these variations (improvement or worsening) depended on pre-TAVI FFR results: positive FFR values tended to worsen post-TAVI, while negative FFR values tended to improve post-TAVI [24]. TAVI, by inducing an immediate decrease in hyperaemic microvascular resistance and an increase in hyperaemic flow velocity, is associated with an immediate improvement in coronary microcirculation's vasodilator capacity. This pathophysiologic assumption may be one factor underlying FFR variations post-TAVI, although dedicated studies are needed [29].

Similar patterns were not observed for iFR, which showed wide individual variations after TAVI, with higher delta (iFR after TAVI — iFR before TAVI) associated with a greater drop in transacrtic gradient after valve intervention [22].

Considering that TAVI patients will live without AVS, it is intuitive that the functional significance of intermediate coronary lesions should be evaluated in the absence of this condition, hence after valve implantation. Available literature supports FFR as a more reliable parameter than iFR, and even suggests measuring it post-TAVI for a more accurate assessment of the need for myocardial revascularization, especially for borderline FFR values that may decrease after AVS removal [24].

Moreover, the recent advent of image-based functional assessment techniques (e.g., quantitative flow ratio) is timely, considering TAVI candidates represent an ideal population for their application. The ability to examine the functional significance of coronary stenosis without wiring the vessel is appealing, particularly when post-TAVI measurement is desirable. Mejía-Rentería et al. [30] analyzed the diagnostic performance of quantitative flow ratio in AVS patients against FFR, reporting per-vessel sensitivity, specificity, area under the ROC curve, and accuracy of 84% (95% CI, 71%–92%), 80% (95% CI, 69%–88%), 0.88 (95% CI, 0.82–0.93), and 81%, respectively [30, 31].

Beyond these peculiarities, clinical outcomes after physio-guided revascularization of intermediate stenosis in TAVI patients support the validity of this strategy. An observational study including 216 patients compared FFR-guidance to angio-guidance for myocardial revascularizations in the TAVI context: patients evaluated with FFR had significantly better outcomes at 2 years compared to those guided by angiography alone (major adverse cardiac and cerebrovascular events free-survival 92.6% vs. 82.0%; P = 0.035), mainly due to a higher rate of periprocedural myocardial infarction in the angio-group. Most lesions assessed by FFR were negative (72%), and these patients, treated medically, performed better than those treated

by angio-guided PCI [32]. In line with this, investigating the residual functional SYNTAX score (rFSS) after TAVI revealed that functional incomplete revascularization (rFSS >0) was associated with worse event-free survival at follow-up [33].

Despite the observational nature of these studies, the data reassure about the safety of PCI deferral based on negative FFR, while cautioning against the risks of angio-driven PCI. The upcoming results of randomized clinical trials (FAITAVI — NCT03360591, NOTION-3 — NCT03058627, TAVI-PET — NCT04882488) focused on this specific topic will further elucidate the role of invasive functional assessment in determining when to perform PCI in TAVI patients (Table 1).

PCI TIMING IN PATIENTS UNDERGOING TAVI

Once the indication for PCI is established, it is common practice to perform it upstream of TAVI, still in the presence of severe AVS, through a staged procedure or during TAVI before valve deployment, with initial studies supporting such a strategy [15, 19, 34–37]. This is also the recommendation provided in the present American Guidelines [20], while European guidelines do not specify timing for coronary revascularization, suggesting it should be based on clinical presentation and coronary anatomy complexity [19]. These studies have generally been interpreted as proof of the feasibility of a PCI pre-TAVI approach, rather than offering a comparison between different timings for PCI (pre- and post-TAVI).

However, operators choosing this strategy must contend with several drawbacks: an increased risk of acute kidney injury, dual antiplatelet therapy before TAVI which might increase bleeding and vascular complications during TAVI [38], and systemic hypoperfusion derived from aortic valve obstruction which might complicate some complex PCI procedures. On the other hand, performing PCI before TAVI is desirable in the presence of ongoing ischemia (i.e., acute coronary syndromes), it allows easier coronary access, and it might be preferred in case of ostial/proximal lesions to prevent large myocardial ischemia during LV pacing (Table 2).

An appealing alternative consists in alleviating LV overload by performing a balloon aortic valvuloplasty followed by PCI during the same procedure [39, 40]. This approach, when tolerated, reduces systemic hypoperfusion and potential secondary ischemic PCI complications, serving as a bridge to TAVI in patients with temporary TAVI contraindications.

Conversely, few studies have investigated the potential advantages of performing PCI after TAVI in the absence of AVS, hampered by the risk of ischemic and hemodynamic

complications potentially inducible during the TAVI procedure due to the presence of a large ischemic territory [41–44].

Ochiai et al. [45], using balloon-expandable valves only, reported similar clinical outcomes between pre-TAVI (n = 143), concomitant (n = 77), or post-TAVI (n = 38) PCI groups at 2 years. However, the use of balloon-expandable valves only limits the generalization of such results to the real-world population, where self-expandable valves notably hinder coronary access.

Another study examined the feasibility, periprocedural complications, and 2-year outcomes of PCI pre- vs. post-TAVI in 144 patients, 73% (n = 105) of whom underwent PCI of LM, proximal LAD, or proximal dominant right coronary artery (RCA). Both self- and balloon-expandable valves were included [44]. While procedural success was achieved in all cases and overall periprocedural complications were similar, a higher incidence of in-hospital stroke was reported among the PCI pre-TAVI group. Additionally, at 2 years, the major adverse cardiac and cerebrovascular events free-survival was lower in patients who underwent PCI before TAVI.

These results have recently been corroborated by the REVASC-TAVI registry, a large multicentre international study. Based on data from 1603 patients, the study concluded that performing PCI after TAVI seems to be associated with improved 2-year clinical outcomes compared to other revascularization timing strategies, significantly reducing all-cause deaths, as well as the composite of all-cause death, stroke, MI, or unplanned rehospitalization for heart failure [46] (Table 3).

These findings offer interesting insights: firstly, PCI after TAVI appears safe and feasible, at least as a PCI-first approach. The similar rate of periprocedural complications suggests that even the presence of "high-risk" lesions (i.e., LM or proximal LAD stenosis), with a large ischemic burden, does not compromise the TAVI procedure when coronary lesions are treated after TAVI, regardless of the valve type.

Available data on the feasibility of coronary access after TAVI suggest greater difficulties with high frame, supra-annular prostheses compared to low frame, intra-annular valves [47, 48]. However, even considering only supra-annular high frame prostheses (e.g., Medtronic Evolut device), the inability to cannulate the coronaries ranges between 0 to 15%, with most studies reporting almost 100% success [43, 49–53].

While the type of prosthesis plays a key role in this setting, other risk factors for difficult coronary canulation include patients' anatomical characteristics and procedural techniques.

Procedural factors are particularly relevant today, as there is increasing attention from prosthesis manufacturers and operators on developing enhanced devices to achieve commissural alignment and the ideal implant height during valve implantation. It has been demonstrated that orienting the Medtronic Evolut device with the flush port placed at 3 o'clock and keeping the Evolut hat marker at the outer curve of the thoracic aorta reduced the misalignment between device commissures and coronary ostia from 38 to 24% [54]. Similarly, the COMALIGN study showed that orienting the specific fluoroscopic markers of any prosthesis according to the cusp overlap angiographic view (left and right cusps) can achieve correct commissural alignment in up to 9 out of 10 patients [55].

The above-reported rates of successful coronary engagement after TAVI also referred to earlier TAVI periods when operators paid less attention to commissural alignment. Considering the current improved implantation techniques and devices, today we may expect a lower failure rate of coronary access after TAVI. These data can help operators either guide the decision to perform PCI before or after TAVI based on the prosthesis selected for a given patient, or choose a specific valve type (e.g., low frame, intra-annular) to facilitate post-TAVI PCI.

Additional advantages of a TAVI-first approach are worth mentioning, as they might be relevant in selected settings, such as complex PCI procedures, chronic kidney disease, high bleeding risk, and borderline stenosis severity (Table 2). The opportunity to remove LV obstruction and improve systemic perfusion before PCI is particularly relevant in the case of potential ischemic or mechanical complications related to complex PCI that could further impair cardiac output, even if only transiently, affecting organs with low ischemic thresholds such as kidneys and the brain that are already hypoperfused in patients with severe AVS. The immediate favourable hemodynamic effect of TAVI might explain the better kidney tolerance and the lower stroke rate observed [56].

At this regard, it has been demonstrated that TAVI has a protective effect on contrast induced-acute kidney injury incidence (odds ratio, 0.334; 95% CI, 0.193–0.579; P < 0.001) [57], when compared to any other coronary procedures (either diagnostic angiograms or PCI).

This observation suggests the impact of contrast administration on kidney function in patients who had undergone TAVI may be better tolerated because of the hemodynamic changes following aortic valve replacement. This supports an eventual PCI post-TAVI strategy in patients at higher risk of contrast induced-acute kidney injury.

On the other side, explaining the lower incidence of strokes is less straightforward, considering such events might derive from either prolonged brain hypoperfusion in the context

of a diffuse cerebral vascular disease, or thromboembolisms. The latter deserves particular attention, as there exist numerous cerebral protection devices [58], potentially helpful preventing that event. However considering the still limited — and not conclusive — evidence about their use during TAVI [19], the decision to use such devices should be taken on a case-by-case basis. In the setting of PCI and TAVI, the higher risk of strokes observed when PCI is performed firstly (hence while not intervening on the calcified aortic valve) suggests the former physiopathology the most likely. Hence, the eventual use of cerebral protection devices should be limited to cases considered at higher risk of strokes, on the sole basis of the aortic valve disease anatomy and the expected TAVI complexity, regardless to the subsequent PCI.

Considering CAD in AVS patients is often associated with a high burden of coronary calcification, complex procedures are sometimes needed, including debulking techniques (e.g., intravascular lithotripsy and rotational atherectomy [RA]), which could be better tolerated thanks to improve myocardial contractile reserve and global hemodynamics. When implemented before TAVI, RA required the use of balloon pumps and/or inotropic support in a considerable number of patients, testifying to the higher ischemic burden in the presence of AVS [59, 60]. Conversely, other observational data reported the feasibility and safety of RA after TAVI, either through a self-expandable or a balloon-expandable valve [61–63], without periprocedural major cardiovascular or cerebral adverse events.

Importantly, with regard to bleeding, performing TAVI after PCI, in most cases implies the use of dual anti-platelet therapy, or even triple antithrombotic therapy in case of patients requiring anticoagulation. Although TAVI can be deferred until tiple therapy is shifted to anticoagulant plus clopidogrel, this association has been demonstrated to increment the risk of bleeding and vascular complications (1.6-fold risk compared to anticoagulation alone) according to the recent POPULAR TAVI study [38]. This evidence strongly supports operators performing TAVI first in patients under anticoagulant medications (or at high bleeding risk), in order to secure vascular accesses healing before staring any additional antithrombotic agent.

Furthermore, the absence of AVS permits a more accurate diagnosis of the ischemic potential of a given angiographic intermediate stenosis by physiological assessment, as previously reported [22, 24, 64].

Lastly, one may suppose some conduction disturbances after TAVI might present a ischemic component (e.g., right coronary severe disease), prompting the need for revascularization before TAVI. Of note, the incidence of new-onset conduction disorders requiring permanent pacemaker implantation remains relatively high and represents a cause for concern [65, 66]. Currently, definitive evidence is lacking on the effect of functional

assessment and/or PCI before or after TAVI and its association with post-TAVI conduction disturbances.

However, there are no available randomized clinical trials clearly defining the best timing approach for myocardial revascularization in AVS patients. An ongoing trial (Optimal Timing of Transcatheter Aortic Valve Implantation and Percutaneous Coronary Intervention — The TAVI PCI Trial, NCT04310046) comparing PCI pre- vs. post-TAVI (according to iFR ≤0.89 or angiographic stenosis >90%) will offer more solid evidence on this topic.

For the time being, the available literature suggests both approaches are feasible [67]. Regarding safety, they are mostly comparable, and as to clinical benefits, differences may be subtle and could emerge only after the analysis of a larger number of cases not yet available. Until definitive evidence becomes available, choosing a TAVI-first approach addresses the primary clinical problem, while concomitant stable CAD remains an occasional finding and, as such, the need for treatment is uncertain and can always be considered after a thorough clinical or instrumental evaluation.

CONCLUSIONS AND TAKE HOME MESSAGES (FIGURE)

The best management of CAD in TAVI candidates is still under investigation, and most of the available evidence is based on observational studies from high-volume centres (Tables 1 and 2). Therefore, definitive indications cannot yet be formulated. The following messages should be interpreted as reasonable advice arising from the above-discussed studies:

- **Streamline CAD Detection**: Operators should streamline CAD detection by utilizing CCTA to select patients with abnormal findings to undergo ICA.
- Functional Evaluation for Intermediate Stenosis: Once CAD is detected, functional evaluation is desirable in cases of intermediate stenosis (at least >50%) to guide revascularization and avoid worthless and potentially harmful interventions.
- Timing of Functional Evaluation: Functional evaluation should be carried out after valve deployment to avoid the hemodynamic influence generated by LV obstruction. Among functional indexes, FFR has shown the best correlation with myocardial nuclear imaging in detecting ischemia, providing more reliability when measured before and after TAVI.
- **Timing of PCI**: Ischemia-driven interventions are feasible and safe either pre- or post-TAVI. Potential advantages may arise from postponing the PCI until after valve replacement; however, as there is no one-size-fits-all approach, the best strategy should be tailored on a case-by-case basis.

Article information

Conflict of interest: ML received speaker fees from Medtronic. FBia received a research grant from Abbott Vascular. CA Aurigemma received speaker fees from Abbott Vascular, Abiomed, Medtronic, Terumo, and Daiichi Sankyo. ER received speaker fees from Abiomed, Abbott Vascular and Terumo. LP received speaker fees from Abiomed and Terumo. CT and FBur received speaker fees from Abbott Vascular, Abiomed, Medtronic, and Terumo. The other authors report no conflicts of interest to declare.

Funding: None.

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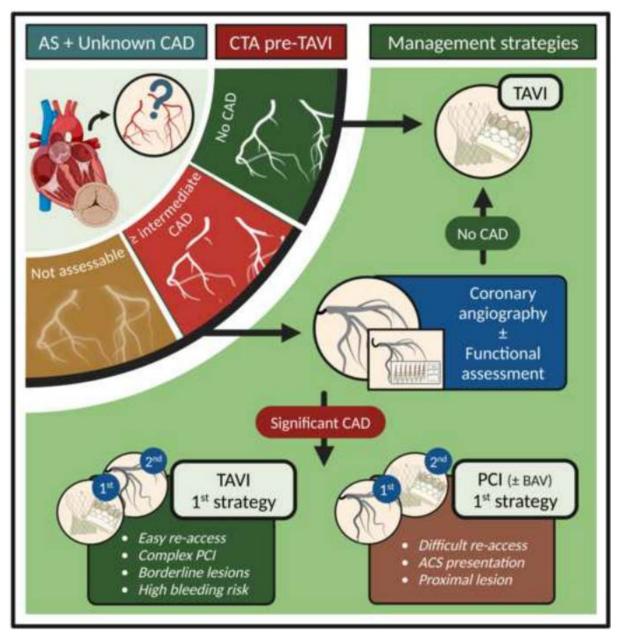


Figure 1. Proposed algorithm for CAD severity assessment and treatment in patients undergoing TAVI. CAD should be ruled out by CCTA, when available. If not, or if CCTA reveals abnormal findings, coronary angiography is required, before valve implantation. In case of detection of intermediate stenosis, a TAVI-first strategy is recommended when feasible, according to clinical presentation and patients' characteristics (expected easy coronary reaccess, high bleeding risk, etc.). Once TAVI is performed, functional evaluation of intermediate stenosis should be performed with pressure wire or image-based functional assessment to guide eventual revascularization. If a PCI-first approach is desirable (large ischemic myocardium, acute settings, etc.), operators might consider to perform aortic ballon valvuloplasty before PCI as bridge to TAVI

Abbreviations: ACS, acute coronary syndrome; AS, aortic stenosis; BAV, balloon aortic valvuloplasty, CAD, coronary artery disease; CTA, coronary tomography angiography; PCI, percutaneous coronary intervention; TAVI, transcatheter aortic valve replacement

Table 1. Main studies on CAD severity assessment in patients with severe aortic stenosis

Study	Objective	Design	Sample	Publication	Key findings
			Size	Year	
Chieffo A	CCTA to rule	Observational	491	2015	Only 25% of
	out CAD	retrospective			patients underwent
					CA
van den	Diagnostic	Meta-analysis	1275	2018	High NPV and
Boogert	accuracy of				sensitivity, but low
TPW	CCTA for				specificity. CA
	CAD				spared in 37%
	diagnosis				
Scarsini R	Correlation	Observational	28	2019	FFR <0.80 —
	between	prospective			SPECT agreement
	SPECT and				85%, iFR <0.82 —
	FFR/iFR				SPECT 73%
Scarsini R	Comparison	Observational	179	2017	Best iFR cut-off to
	FFR vs. iFR	retrospective			predict FFR <0.80:
	in AS				0.83
Pesarini G	FFR	Observational	133	2016	No significant
	variations	prospective			variations after
	before vs.				TAVR. 6% only
	after TAVR				changed indication
					to treat
Scarsini R	iFR	Observational	145	2018	Erratic individual
	variations	prospective			variations. 15%
	before vs.				changed indication
	after TAVR				to treat

Mejía-	QFR	Observational	138	2020	Accuracy in
Rentería H	diagnostic	retrospective			predicting FFR
	performance				<0.80: 81%
	in AVS				
Lunardi M	Angio- vs.	Observational	216	2021	Higher 2y
	functional-	retrospective			MACCE-free
	guidance for				survival in
	PCI				functional arm
FAITAVI	Angio- vs.	Randomized	320	_	_
trial	functional-	clinical trial			
	guidance for				
	PCI				
NOTION-	FFR-guided	Randomized	452	_	_
3 trial	PCI or	clinical trial			
	medical				
	treatment				
TAVI-	Correlation	Observational	20	_	_
PET trial	of FFR and	prospective			
	iFR with				
	cardiac PET				
	perfusion				

Abbreviations: AS, aortic stenosis; AVS, aortic valve stenosis; CA, coronary angiography; CCTA, coronary computed tomography angiography; FFR, fractional flow reserve; iFR, instantaneous wavefree ratio; MACCE, major adverse cardiac and cerebrovascular events; NPV, negative predictive value; PET, positron emission tomography; QFR, quantitative flow ratio; SPECT, single photon emission computed tomography; TAVR, transcatheter aortic valve replacement; other — see Figure 1

Table 2. Examples of preferred PCI timing based on clinical scenarios

PCI	Clinical scenarios
timing	
Pre-TAVI	Acute coronary syndromes (i.e., when ischemia is the acute issue)

	Ostial/proximal right coronary artery or left main stenosis (possibly)
	leading to large myocardial ischemia during left ventricular pacing
	implanting the valve)
	 Specific anatomies leading to unfavourable interactions between the
	selected valve and the coronary ostia, making coronary canulation
	after TAVI more challenging (e.g., low coronary ostia, ectopic ostia,
	narrow sinuses, valve-in-valve procedures, etc.)
	Following balloon aortic valvuloplasty (bridge to TAVI) when tight
	severe aortic stenosis is present (particularly high aortic-ventricular
	gradients)
Post-	Expected complex and long PCI procedures (e.g., requiring debulking)
TAVI	techniques, etc), when the presence of aortic valve obstruction, thus
	the systemic hypoperfusion, might pose the PCI at higher risk of
	procedural complications
	Presence of severe chronic kidney disease, and high risk of contrast-
	induced acute kidney injury (the normalization of the systemic
	perfusion after TAVI promotes better kidneys perfusion and reduces
	the risk of acute injury during additional contrast medium
	administrations)
	Patients presenting with high bleeding risk or on anticoagulation, to
	avoid performing TAVI on double antithrombotic therapy.
	Borderline coronary stenosis severity at coronary
	angiogram/physiology before TAVI, requiring functional re-evaluation
	after the aortic valve stenosis treatment

Abbreviations: see Figure 1

Table 3. Main studies on PCI timing in patients undergoing TAVR

Study	Objective	Design	Sample	Publication	Key findings
			Size	Year	

Venturi G	Concomitant	Observational	339	2021	Staged CA
	vs. staged	retrospective			increased risk of
	CA				CI-AKI
Ochiai T	Comparison	Observational	258	2020	No 2 years
	of different	retrospective			differences in
	timing for				outcomes
	PCI				between pre-,
					concomitant,
					post-TAVR PCI
Lunardi M	Comparison	Observational	144	2022	Higher in
	of pre-	retrospective			hospital stroke
	TAVR vs.				rate, and 2 years
	post-TAVR				MACCE in pre-
	PCI				TAVR arm
TAVI PCI	Comparison	Randomized	986	_	_
trial	of pre-	clinical trial			
	TAVR vs.				
	post-TAVR				
	PCI (iFR				
	guided)				
Lunardi M	Feasibility	Observational	19	2020	15% inotropic
	and 2 years	retro/prospective			support, 100%
	outcomes of				success, 100%
	RA after				30d survival,
	TAVR				84% 2 years
					survival
Naganuma	Feasibility	Observational	25	2017	40% inotropic
Т.	and 30 days	retrospective			support, 100%
	outcomes of				success, 100%
	RA before				30 days survival
	TAVR				
Lippmann	Feasibility	Observational	29	2017	30% inotropic
M.	and	retrospective			support, 100%

	immediate				success, 100%
	outcomes of				immediate
	RA before				survival
	TAVR				
REVASC-	Comparison	Observational	1603	2023	PCI performed
TAVI	of pre-	retrospective			before, after or
registry	TAVR vs.				concomitantly
	concomitant				with TAVR in
	vs. post-				65.6% (n =
	TAVR PCI				1052), 9.8% (n =
					157) or 24.6% (n
					= 394). 2-year
					all-cause death
					in post-TAVR,
					pre-TAVR,
					concomitant:
					6.8% vs. 20.1%
					vs. 20.6%; P
					<0.001

Abbreviations: CI-AKI, contrast induced-acute kidney injury; RA, rotational atherectomy; TAVR, transcatheter aortic valve replacement; other — see Figure 1 and Table 1