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Same-day discharge after percutaneous coronary procedures – Consensus statement of the working group of interventional cardiology (AGIK) of the Austrian Society of Cardiology

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Summary

Introduction Percutaneous coronary intervention is a well-established revascularization strategy for patients with coronary artery disease. Recent technical advances such as radial access, third generation drugeluting stents and highly effective antiplatelet therapy have substantially improved the safety profile of coronary procedures. Despite several practice guidelines and a clear patient preference of early hospital discharge, the percentage of coronary procedures performed in an outpatient setting in Austria remains low, mostly due to safety concerns.

Methods The aim of this consensus statement is to provide a practical framework for the safe and effective implementation of coronary outpatient clinics in Austria. Based on a structured literature review and an in-depth analysis of available practice guidelines a consensus statement was developed and peerreviewed within the working group of interventional cardiology (AGIK) of the Austrian Society of Cardiology.

Results Based on the available literature same-day discharge coronary procedures show a favorable safety profile with no increase in the risk of major adverse events compared to an overnight stay. This document provides a detailed consensus in various clinical settings. The most important prerequisite for sameday discharge is, however, adequate selection of suitable patients and a structured peri-interventional and postinterventional management plan.

Conclusion Based on the data analysis this consensus document provides detailed practice guidelines for the safe operation of daycare cathlab programs in Austria.

Keywords Outpatient coronary procedure · Coronary intervention · Ambulatory treatment · Consensus document · Austria

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Abbreviations

ACS	Acute coronary syndrome
AKI	Acute kidney injury
CAD	Coronary artery disease
CIN	Contrast-induced nephropathy
СТО	Chronic total occlusion
DA	Diagnostic angiography
DAPT	Dual antiplatelet therapy
GFR	Glomerular filtration rate
LM	Left main coronary artery
MACCE	Major adverse cardiovascular or cerebral
	event
MACE	Major adverse cardiovascular event
MV	Multivessel
ND	No data
n. s.	Not statistically significant ($p < 0.05$)
NS	Not specified
NSTEMI	Non-ST-segment elevation myocardial in-
	farction
OBS	Observational study
OCT	Optical coherence tomography
OR	Odds ratio
OS	Overnight stay
PCI	Percutaneous coronary intervention
PCP	Percutaneous coronary procedures
RA	Rotational atherectomy
RCT	Randomized controlled trial
SDD	Same-day discharge
STEMI	ST-segment elevation myocardial infarction
SVG	Saphenous vein graft
TLR	Target lesion revascularization
TTE	Transthoracic echocardiography
TVR	Target vessel revascularization

Introduction

Percutaneous coronary procedures (PCP), including diagnostic angiography (DA) and percutaneous coronary interventions (PCI), are evolving procedures with vast recent improvements regarding access site technique, interventional tools, and concomitant medical treatment. Consistently, the complication profile of PCPs has substantially improved, showing a decline in access site-related, intraprocedural and postprocedural complications [1, 2]. The periprocedural management of PCP patients, however, has remained mostly unchanged, with the majority of centers in Austria scheduling at least one mandatory overnight stay after procedure. On the other hand, consistent with other European countries, the frequency of PCPs performed in Austria is constantly rising, binding greater in-patient resources and beds for hospitals, and creating higher costs for healthcare providers. Especially in the aftermath of coronavirus disease 2019 (COVID-19) many hospitals are struggling with staff shortages, compromising the operation of elective catheterization laboratory programs.

Medical care for patients undergoing PCPs should reflect the current technical standard of these procedures and the current complication profile. Therefore, considering the procedure's safety profile the current standard of treating PCP patients as in-patients with at least one overnight stay irrespective of the medical history or procedural complexity is no longer the only possible approach. There are robust data from patients in randomized trials and large observational studies supporting the safety of performing PCPs as same-day discharge (SDD) procedures for selected patients [3-5]. Furthermore, patient preference towards having PCPs performed as SDD procedures is well documented, especially when performed with radial access [6–8]. Therefore, in an approach of shared decision making, patient preference should be taken into consideration when scheduling PCPs as inpatient or SDD procedures. Also, as highlighted by the American College of Cardiology (ACC) [3], 8% of hospitalizations are associated with highly undesirable events as hospital-acquired infections or falls with subsequent injuries [9, 10]. In this context, SDD PCP programs have been initiated by several cardiology departments in Austria not only to provide better service for cardiology patients but also to create a more efficient patient flow around the catheterization laboratories as critical resources.

In Europe as well as in the USA a remarkable heterogeneity of hospital policies with respect to the treatment standard for PCPs and the frequency of SDD PCPs has been documented [11]. Like any other treatment decision, the discharge policy has important consequences for the patient as well as legal implications for the treating physicians. Thus, a consensus on procedural standards reflecting the available scientific evidence that is supported by the Austrian Society of Cardiology is highly desirable. Furthermore, a common consensus incorporating the experience of clinics running SDD PCP programs for selected patients could serve as a valuable reference standard for quality of care.

Aims

The goals of this consensus document are (1) to provide an overview of the current practice for performing PCPs as an outpatient procedure in Austria and across the world, (2) to review the most recent evidence regarding the safety and feasibility of performing SDD and (3) to provide a consensus framework for the safe and efficient implementation of outpatient clinics within cardiology departments to run a daycare/SDD PCP program. The third goal includes specific recommendations, based on currently available data and experience of the clinics involved, to provide guidance on (4) which patients are best suited for SDD PCP, (5) what intraprocedural techniques or precautions can increase the safety of SDD PCP and (6) what is the appropriate observation period before the patient is discharged as an outpatient.

These recommendations represent the common main aspects endorsed by the working group of interventional cardiology (AGIK) of the Austrian Society of Cardiology and are not intended to be finite or binding. Interventional teams and cardiology departments will make individual modifications to meet their specific expertise and experience. There are different approaches to daycare PCP based on the individual hospital structure and staff training. Furthermore, the authors would like to emphasize that the decision to develop a clinical daycare program around a catheterization laboratory to expand the service offered to patients and referring doctors should be considered an individual choice for every cardiology department. A concept that works well in one hospital could easily be unfeasible in other centers with different local contributing factors. It is not the aim of this consensus document to initiate a competition or create peer pressure as to who is performing the most complex interventions via SDD, but rather to provide guidance on current recommendations and a scientific evidence base for the safe and successful operation of interventional outpatient clinics.

Methods

Consensus development

Following the ACC methodology for creating expert consensus decision pathways [12], this consensus document was developed as a draft outline within the working group and circulated among a task force of the AGIK for peer review. Individual practice guidelines from all individual interventional outpatient clinics were integrated. The final document was once again circulated for organizational review and approval from the Austrian Society of Cardiology.

Review of scientific evidence

This position paper is based on three consensus documents on SDD PCI published by the SCAI [4, 5, 13] and one from the ACC [3]. Furthermore, a systematic literature review and meta-analysis was performed by the Austrian Society of Cardiology focusing on SDD after radial access PCI [14].

Current trends for PCP in an outpatient setting

International trends

Reports on PCIs as daycare procedures in selected patients have been published since the late 1980s [15], remarkably in an era where femoral access was still the procedural standard and balloon angioplasty alone the dominant treatment option for coronary artery stenoses. Since then many centers have adopted radial access PCI as the standard treatment option for elective patients and SDD PCI numbers are steadily increasing in the USA and across Europe:

In their 2021 cross-sectional analysis of the American CathPCI Registry Bradley et al. studied the frequency of patients undergoing elective PCI discharged on the day of procedure [16]. From a total of 819,091 elective PCIs in 1716 hospitals, the survey showed a rapid increase of SDD PCI from 4.5% in 2009 to 28.6% in 2017, a trend that was even more prominent in radial access procedures (increase from 9.9% to 39.7%) compared to femoral access PCI (increase from 4.3% to 19.5%). Dividing the study duration into three time intervals (2009-2011, 2012-2014, 2015-2017), the authors also demonstrated a substantial increase in radial access PCI among SDD (21.8%, 43.4% and 58.3%, respectively) and overnight stay patients (7.8%, 20.1% and 32.6%, respectively) [16]. Among SDD PCI procedures, the percentage of proximal LAD and high risk or type C lesion PCIs also significantly increased over the study period. On the other hand, the authors show a large hospital-level variation of SDD frequencies with 25% of the sites studied discharging < 10% of their patients on the day of PCI despite radial access [16].

Similar trends were published in a Canadian study showing data from 35,972 procedures from 17 clinics in and around Ontario, Canada, from 2008 to 2015 [17]. During the study period the total rate of SDD PCI increased from 17% in 2008 to 45% in 2015; however, as in the previous study, a substantial interhospital variation was detected, including 17 clinics with 0-17% SDD PCI. Indicators of high SDD PCI rates were (1) university clinics and university-affiliated teaching hospitals (40.1% vs. 10.7%), (2) presence of onsite cardiac surgery departments (34.9% vs. 9.4%), and (3) centers with >50% radial access use (42.8% vs. 25.9%) [17].

A 10-year single-center study from France (from 2007 to 2016) showed a comparable increase in SDD PCI procedures from 14% in 2008 to 32% in 2015 [18]. Comparing SDD vs. overnight stay patients, the authors found that in the SDD cohort radial access was used more often, two-vessel CAD was less frequent, and the average number of stents implanted per procedure was lower [18]. The most common reason for overnight admission was the wish to monitor patients in the absence of any complications.

It is worth noting that the increase in SDD PCI procedures runs parallel with a general temporal trend of patients referred to the catheterization laboratory being older, having more comorbidities and a higher calculated risk score for coronary interventions [2]. In a retrospective longitudinal analysis from the Veteran's Affairs Clinical Assessment, Reporting and Tracking Program data from 2009 to 2015 Waldo et al. detected a significant increase in the proportion of DA and PCI patients who had a high Framingham risk score. The hazard ratio (HR) for mortality, however, remained constant over time, with a nonsignificant

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decrease among those undergoing intervention (HR 0.983; 95% confidence interval, CI 0.967–1.000). Similar to the trends in SDD procedures cited above, the authors found a steep increase in radial access from 5% to 32% [2].

Current clinical practice in Austria

A substantial number of Austrian cardiology centers have introduced daycare beds or daycare departments in order to offer a specific scope of procedures on a SDD basis. Generally, DA, PCI, right heart catheterization, cardioversion, loop recorder implant, pacemaker/automatic internal cardioverter defibrillator device replacement as well as administration of intravenous therapy, such as iron infusion, diuretic or antibiotic therapies are being offered as SDD procedures. Depending on the inventories and local environments of the clinics, there are individual policies considering the scope and extent of SDD services. In all centers, local guidelines exist detailing conditions when to switch treatment of a patient back to OS and spare bed capacities are reserved for this purpose.

Currently SDD PCPs are offered by 77% of all cardiology centers in Austria. Of these clinics 40% are running the ambulatory procedures via dedicated outpatient departments and 55% have integrated outpatient beds into regular wards (5% missing data). Fixed protocols or checklists for accepting or rejecting patients for ambulatory procedures have been established in 85% of Austrian cardiology centers. While 77% of clinics offer DA as SDD procedures, only 35% are performing PCI in an outpatient setting. There is some heterogeneity of local protocols with single-stent PCI considered feasible for SDD in all centers performing PCI in an ambulatory setting, while bifurcations, multiple stent and multivessel (MV) PCI are switched to OS in the majority of centers. The volume of SDD cases varies from 2 per week up to 6 per day, whereas the coronavirus disease 2019 (COVID-19) pandemic with subsequent shortages on routine wards has led to an increase in the volume of SDD cases in almost all clinics.

Different clinics have established various local arrangements to integrate a daycare facility into existing cardiology departments. There are two dominant forms of implementation: (1) SDD beds integrated into regular wards where the same doctors, nurses and service staff treat and monitor outpatients, and (2) dedicated outpatient clinics, sometimes shared between different departments (e.g., cardiology combined with gastroenterology, surgery or dermatology), where specific outpatient doctors and nurses are on duty. One benefit of the first form of outpatient clinics is greater flexibility in discharge times, especially considering the waiting period with wound compression after PCI (sometimes up to 6h). As staff are present on the regular wards virtually 24 h/7 days per week, discharge can take place without any limitation. On

the other hand, SDD capacities would be equally compromised in cases of staff shortages or bed reallocation due to COVID-19. Independent outpatient departments (implementation 2) are somewhat more limited in operating hours, on the other hand the general appearance as a non-ward department is preferred by some patients. The staff are also focused on outpatients only and free from many duties connected with generally more immobile patients on regular wards, which could improve the general treatment atmosphere.

Safety profile and outcome of SDD PCP

Review of clinical trials and meta-analyses

As stated above, the Austrian Society of Cardiology has performed a systematic literature review and metaanalysis to substantiate its practice recommendations [14]. In short, four large meta-analyses have published outcome data for SDD PCI [19–22], summarizing 38,785 patients with SDD and 256,049 patients staying overnight. All four meta-analyses are consistent in demonstrating no added risk of SDD practice assessed by MACE at 24 h and/or 30 days.

It could be argued that in all trials including RCTs patients deemed suitable for SDD represent a selected patient population, generally being younger, with a lower burden of chronic medical conditions, less severe degree of CAD and residence not far from the interventional clinic; however, it is the very same selection process that is unanimously being recommended by current advisories to identify potential patients suitable for SDD PCP [3, 4]. It was never intended to offer SDD PCPs to all patients with an indication for coronary angiography or PCI. Conversely, the application of current SDD selection criteria has been shown to be effective: A comparison between randomized and non-randomized (excluded) patients from the RCT by Bertrand et al. [23] showed that the rate of MACE (death, MI, TVR) in patients excluded from randomization was significantly higher at 30 days (10.2% vs. 1.6%), 6 months (17.5% vs. 5.6%), and 12 months (24.5% vs. 9%) as compared with randomized patients [13, 23].

Finally, it should be pointed out that none of the observational studies, RCTs or meta-analyses discussed above featured a protocol where only DA was considered suitable for SDD or where PCI/stent implantation was by default an exclusion criterion for SDD. The safety and outcome data extracted exclusively represent PCI on a SDD basis.

Data on safety of SDD in specific patients and settings

With growing experience of SDD PCPs and especially PCIs, some centers have extended the SDD concept to more challenging PCI indications, once again highlighting the robust safety profile of this treatment approach [24–29]. We have reviewed the evidence base for SDD PCI in higher-risk settings. This includes complex PCI, elderly patients (age >75 years), rotational atherectomy, chronic total occlusion PCI, and left main PCI. Briefly, currently available data from a multitude of trials shows no additional risk for these challenging settings. It must be acknowledged, however, that a large number of trials are retrospective analyses from registries and that a significant degree of heterogeneity exists considering the frequency of higher-risk PCIs on a SDD basis.

Patient satisfaction

Patient preference should be taken into account when scheduling invasive procedures. There is a well-documented patient preference towards radial access for PCP, which leads to less discomfort, less frequent hematomas and better quality of life postprocedure [30, 31].

Across different countries, different study settings and decades of interventional experience SDD after PCP has been shown to be the preferred treatment mode as opposed to OS [6–8, 32–36], satisfying another clear patient preference for short hospital stay, earlier and easier ambulation.

Cost effectiveness

Numerous international studies on cost efficacy of SDD departments have demonstrated a potential for significant cost savings when performing PCPs in an ambulatory setting, while maintaining the existing catheterization laboratory inventories and staff at the center [35–38].

Data from Austria confirm that for the individual cardiology departments, the setup of an interventional outpatient clinic into an existing infrastructure does not involve large investments. A calculation from the University Clinic Innsbruck has shown that after implementation of a cardiology outpatient clinic, estimated annual treatment costs could be reduced by about 560,000 \in , the largest component being cost for personnel (University Clinic for Internal Medicine III, Medical University of Innsbruck; unpublished data), consistent with reductions in fixed hospital costs reported before [37].

Guidelines and international consensus statements

The concept of SDD PCI is being adopted more frequently in cardiology centers around the world and several cardiology societies have developed consensus documents outlining practice guidelines for a safe and effective performance of PCI via outpatient departments. Table 1 shows a synopsis of (1) the Society for Cardiovascular Angiography and Intervention (SCAI) 2009 expert consensus document focusing on length of stay post-SDD PCI [13], (2) the SCAI update from 2018 [5], (3) the SCAI 2020 update position statement for ambulatory surgical centers [4] and (4) the 2021 American College of Cardiology (ACC) expert consensus decision pathway for SDD PCI [3]. On the far right is (5) the corresponding equivalent for each category from the present Austrian Society of Cardiology consensus.

Comparing exclusion criteria against SDD extracted from the four different sources shows that the 2009 SCAI consensus, being the earliest, is the strictest among all five. Potential patients have to be almost free from any chronic disease irrespective of clinical stability. PCI is limited to a single DES of maximally 28 mm length. Proximal LAD, bifurcations, multivessel PCI (MV) are excluded [13]. The updates from 2018, 2020 and 2021 take into account newer trial results supporting a very favorable safety profile of SDD PCI, therefore widening the range of acceptable patient comorbidities and procedure complexities. Chronic conditions like diabetes mellitus, COPD, chronic kidney disease or heart failure are acceptable if under adequate treatment and clinically stable [3–5]. Instead of defining individual exclusion criteria regarding number of stents, cumulative stent length or number of vessels treated, the newer recommendations emphasize the importance of a complicationfree procedural outcome and stable clinical condition of the patient relative to the baseline level preprocedure. Even challenging PCI settings (MV, LM, RA, CTO) are not categorically excluded as long as no periprocedural complications occurred and no exacerbation of any underlying disease of the patient was detected [3]. On the other hand, even in a complication-free postinterventional setting, the patient's willingness to be discharged on the same day, to understand postinterventional DAPT and to be accompanied by a caregiver at home are crucial criteria before proceeding to discharge.

While this consensus endorses many criteria introduced by the 2021 ACC publication, some decisions require further comment:

Age:

It does not seem practicable to introduce a formal age limit as the individual fitness and coping with provisions necessary to organize a SDD treatment may vary substantially despite the numerical age.

Social factors:

The term "reliable person for transit home" does also include a commercial transport service as long as a qualified caregiver is ready and waiting at the patient's home residence.

Access site:

With respect to the magnitude of clinical trials confirming the safety of SDD after femoral access PCI [6, 20], this approach is considered safe if vascular closure devices are used and can be implanted without complications. If, however, the access site has to be

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Table 1 Syn	opsis—Patient or proc	edure characteristics u	Infavorable for SDD			
	SCAI Expert Consensus Document	SCAI Expert Consensus Update	SCAI Position Statement 2020 for ASC	ACC Expert Consensus Decision Pathway	Austrian Society of Cardiol- ogy Consensus	
Publication	Chambers et al. (2009) [13]	Seto et al. (2018) [5]	Box et al. (2020) [4]	Rao et al. (2021) [3]	(This publication)	
-	Exclusion criteria defined by	/ each consensus/position pa	per grouped by category			
Patient						
Age	Age > 70 years	Age: no limit	Age: no limit	Age: no limit	Age: no limit	
Clinical setting	NSTEMI or STEMI	NSTEMI or STEMI	ACS	NSTEMI or STEMI	Staged PCI for NSTEMI or STEMI no exclusion	
	Elevation of cardiac biomarkers	Continuing angina		Staged PCI for NSTEMI or STEMI no exclusion		
Clinical condition	Unstable patient	Not clinically stable	Any signs of clinical insta- bility	Any disease exacerbation of COPD/HF/hypertension	Any signs of clinical insta- bility	
Mental state	<ns></ns>	<ns></ns>	<ns></ns>	Changes in mental state	Patient is mentally incapac- itated	
Diabetes	Insulin-dependent diabetes mellitus	Diabetes mellitus not clinically stable	<ns></ns>	Diabetes mellitus: no concern if clinically stable	Diabetes mellitus: no con- cern if clinically stable especially no risk of hypo- glycemia	
COPD	COPD if significant or requiring medication	COPD if not clinically stable	Advanced COPD requiring oxygen	COPD: no concern if clini- cally stable	COPD: no concern if clini- cally stable	
			Severe pulmonary hyper- tension			
Heart failure	LVEF < 55%	Decompensated HF, fluid overload (No LVEF cut-off)	Decompensated HF (NYHA classes 3 and 4)	HF: no concern if clinically stable (No LVEF cut-off)	HF: no concern if clinically stable (No LVEF cut-off)	
			LVEF < 30%			
Valvular heart disease	Any valvular regurgitation	<ns></ns>	Severe aortic stenosis	<ns></ns>	<ns></ns>	
Renal function	eGFR < 60 ml/min	CKD not clinically stable	CKD with eGFR	CKD: no concern if clini- cally stable	CKD no concern if clinically	
	On dialysis	CKD requiring prolonged hydration	<45 ml/min/1.73 m ² BSA		Stable	
Peripheral vas-	Symptomatic PVD	PVD not clinically stable	Significant PVD limiting femoral or radial access	<ns></ns>	PVD not clinically stable	
(PVD)					PVD limiting radial/ulnar access	
Cerebro-vascular disease	<ns></ns>	<ns></ns>	Recent TIA or stroke	<ns></ns>	<ns></ns>	
Contrast allergy	Any contrast allergy	Contrast reaction with ongoing symptoms	Severe contrast allergy	<ns></ns>	Contrast reaction with ongoing symptoms	
Anemia	<ns></ns>	<ns></ns>	Anemia with Hb $< 9 \text{ g/dl}$	<ns></ns>	<ns></ns>	
Coagulation	<ns></ns>	<ns></ns>	Coagulopathy with INR > 1.5 or platelet count < 100,000	<ns></ns>	<ns></ns>	
Social factors	No adequate home support	No caregiver for 24 h post- procedure	No caregiver for 24 h post- procedure	No reliable person for transit home	No adequate home suppo	
			No adult present for dis- charge and at home	No caregiver for 24 h post- procedure		
Distance to PCI center	Lives or stays > 20 miles from PCI facility	No transportation home	<ns></ns>	Inadequate home support	<ns></ns>	
	No adequate home support	Inadequate home support				
Access to emer- gency medical care/service	Inadequate local emer- gency medical care	Inadequate access to emergency medical care	\geq 30 min drive time to hospital capable of providing emergency medical care	No access to emergency services	\geq 30 min drive time to hospital capable of providing emergency medical care	

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switched from radial to femoral due to complications, resulting in two compromised body areas, the patient should be rescheduled to OS. This is further supported by data indicating that femoral access is associated with a higher complication rate when used as a secondary access site [39].

PCI location:

While no exclusion criteria are defined based on stent location, LM and bifurcation PCI should match the interventional team's expertise. Consistent with current guidelines, especially in LM PCI intravascular imaging techniques is mandatory to document proper stent apposition and absence of stent edge dissections.

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· ·	SCAI Expert Consensus Document	SCAI Expert Consensus Update	SCAI Position Statement 2020 for ASC	ACC Expert Consensus Decision Pathway	Austrian Society of Cardiol- ogy Consensus	
Procedure						
Access site	Femoral access no exclu- sion with closure device	Brachial access	<ns></ns>	Femoral access no exclu- sion with closure device	Femoral access no exclu- sion with closure device	
	Brachial no exclusion					
Sheath	<ns></ns>	Sheath size \ge 9 French	<ns></ns>	Sheath size > 7 French	<ns></ns>	
				Sheathless guide 6.5 French no exclusion		
PCI location	1-vessel disease with LM PCI	Last remaining artery	Unprotected LM	No exclusions on location (including LM)	No exclusions on location	
	Prox. LAD	No exclusion if LM or bifurcation	Bifurcation with significant side branch involvement	Bifurcation no exclusion		
	Any bifurcation		Bypass graft			
	SVG		Last remaining conduit			
	IMA		Extreme prox. angulation/ tortuosity			
Multivessel	>1 vessel PCI	No limit on number of vessels	3-vessel CAD	No limit on number of vessels	No limit on number of vessels	
Stent length/	>1 Stent	No limit on stent number	No limit on stent number	No limit on stent number	No limit on stent number	
number	>28 mm	No limit on cumulative stent length	No limit on cumulative stent length	No limit on cumulative stent length	No limit on cumulative stent length	
Periprocedural adverse events	<ns></ns>	Periprocedural MI	Any cardiac or non-cardiac instability during PCI	Any periprocedural compli- cations	Any cardiac or non-cardiac instability during PCI	
PCI success	Balloon angioplasty without stent	Balloon angioplasty without stent	<ns></ns>	Unsuccessful stent deploy- ment	<ns></ns>	
<u> </u>	10	Inability to deliver stent	D 11 1 1 1 000/	D		
Residual stenosis	<ns></ns>	<ns></ns>	Residual stenosis > 30%	Residual stenosis > 50%	<ns></ns>	
TIMI flow	<ns></ns>	<ns></ns>	TIMI flow < Grade 3	TIMI flow < Grade 3	TIMI flow < Grade 3	
			I ransient vessel closure		I ransient vessel closure	
Side branches	Any SB loss	Any SB closure	Significant SB involvement	SB: no diameter limit	Any SB loss if clinically	
(50)	Compromised SB flow		SB loss > 1 mm		persistent AP, arrhythmia)	
Dissection	Any dissection	Any dissection	Types B–F dissection in target vessel at the end of procedure	<ns></ns>	Any dissection not covered by DES or treated with DEB	
Thrombotic events	Distal embolization	<ns></ns>	Any intracoronary throm- bus	<ns></ns>	Any intracoronary throm- bus	
CTO-PCI	CTO attempt	CTO attempt	Any CTO	CTO no exclusion if clini- cally stable	CTO no exclusion if clini- cally stable	
Rotational atherectomy	Any rotational atherectomy	Any rotational atherectomy	Any rotational atherectomy	Rotational atherectomy no exclusion	Calcium modifying therapy no exclusion if clinically stable	
Left-ventricular	<ns></ns>	Any LVAD	<ns></ns>	<ns></ns>	Any LVAD	
assist device (LVAD)					Any circulatory support (inotropes etc.)	
Anticoagulation	Use of GPIIb/IIIa inhibitors	Use of GPIIb/Illa inhibitors	<ns></ns>	<ns></ns>	<ns></ns>	
Contrast medium	Large volume of contrast medium (> 500 ml)	Large contrast volume	<ns></ns>	<ns></ns>	Large volume of contrast medium (> 500 ml)	

Table 1 (Continued)

Cumulative stent length:

Consistent with the ACC 2021 consensus [3], the introduction of a limit on number of stents or vessels treated in SDD PCIs does not seem to be warranted. A multitude of studies in this review have shown no added risk for complex interventions including MV and multistent PCIs. Signalling extensive atherosclerotic disease, cumulative stent length does have an adverse effect on short-term and long-term MACE in BMS [40] and 1st generation DES [41]. In 2nd and 3rd

generation DES this correlation has been shown to be either absent [42, 43], or only affecting late outcomes in women [44].

Dissections:

Dissections not treated with DES or DEB should be considered an exclusion criterion for SDD post-PCI. If OCT is being used, this criterion has to be balanced with the fact that small edge dissections are found in up to 40% of OCTs post-DES implant [45]. Of these 80% are not detectable via angiography [46] and have

	Table I (Cor	ilinuea)				
		SCAI Expert Consensus Document	SCAI Expert Consensus Update	SCAI Position Statement 2020 for ASC	ACC Expert Consensus Decision Pathway	Austrian Society of Cardiology Consensus
	Post-procedure					
İ	Access site	Hematoma	Bleeding complications	Access site hematoma	Bleeding	Clinically relevant bleeding or hematoma
					Vascular complications	
	ECG	Any rhythm disorders	(Postprocedure ECG "if ordered")	ECG abnormalities or rhythm disorder prior to discharge	Persistent ischemic ECG changes	Dynamic ECG changes
					Dysrhythmia	Dysrhythmia
	Pain	Continuing chest pain	Continuing angina	Chest pain	Unresolved/severe chest pain	Persistent chest pain
	Interventionalist, nursing staff	<ns></ns>	Discomfort of caregiver/ physician about SDD	Operator judgement favors OS	<ns></ns>	Discomfort of caregiver/ physician about SDD
	Patient's deci- sion	Patient and family not willing to consider early discharge	Discomfort of patient about SDD	<ns></ns>	Patient not willing to be discharged	Discomfort of patient about SDD
ļ	Discharge recomi	mendations				
Cli	Clinical follow-up	Involve quality improve- ment committee, assess complications, patient satisfaction	Follow-up appointment	Follow-up appointment within 1–2 weeks	Schedule contact call 1 day after discharge	According to local expertise
		Interaction with patient and family			Referral to cardiac rehabili- tation	
l	ACS acute coronary syndrome, CKD chronic kidney disease, COPD chronic obstructive pulmonary disease, CTO chronic total occlusion, DEB drug-eluting balloon,					

ACS acute coronary syndrome, CKD chronic Kidney disease, CUPD chronic obstructive pulmonary disease, CTO chronic total occlusion, DEB drug-eluting balloon, eGFR estimated glomerular filtration rate, HF heart failure, LM left main coronary artery, LVAD left ventricular assist device, LVEF left ventricular ejection fraction, MI myocardial infarction, <NS> not specified, NSTEMI non ST-elevated myocardial infarction, PCI percutaneous coronary intervention, PVD peripheral vascular disease, SB side branch, STEMI ST-elevated myocardial infarction, SVG saphenous vein graft, TIA transitory ischemic attack, TIMI thrombolysis in myocardial infarction classification

no impact on clinical outcome [47]. Therefore, consistent with current OCT guidelines, clinically relevant edge dissections post-DES implant are defined as either (1)>200 µm in depth into the vessel wall (media), and/or (2) reference luminal area <4.5 mm² at either proximal or distal stent edge, and/or (3) ≥ 3 mm in length from the stent edge, and/or (4) spanning >60° in arc from the center of the vessel [47, 48].

Acute kidney injury (AKI)/contrast-induced nephropathy (CIN):

In CIN serum creatinine levels tend to rise at the end of a 24 h window postcontrast exposure, reaching a maximum at 48–72 h [49]. Therefore, it is very unlikely that CIN could be detected within the 3–6 h post-procedural observation period even with serial creatinine laboratory tests. For this reason, the cut-off for contrast use mandating a switch from SDD to OS strategy should be set at 500 ml as in [13]. A more conservative, dynamic threshold of $3 \times$ calculated eGFR (Cockroft-Gault formula), which accounts for the elevated risk for CIN in patients with renal failure [50] can be considered in individual patients at high risk for CIN.

Practical recommendations

Hospital environment

All SDD PCPs should be performed by clinics and medical centers with a high annual workload of coronary procedures. Consistent with the 2020 SCAI recommendations for ambulatory surgical centers, we recommend that only operators with an expertise in interventional cardiology, a personal experience beyond a total of 500 procedures and an annual workload of >50 procedures should be performing PCPs in the outpatient setting [4]. A specialized hospital program or SOP for SDD PCPs should be in place including secretaries, nurses and catheterization laboratory personnel, defining the relevant steps from assessment of referrals, scheduling of appointments for outpatient evaluation, daycare admission, cathlab procedure, postprocedure patient care, discharge criteria checklist, and clinical follow-up protocol.

At any time from referral to discharge, including the patient's own condition as well as observations by all levels of personnel, the goal of SDD should be critically re-evaluated.

An experienced colleague should be available at all times to assist in the preprocedural and postprocedural care and to manage discharge formalities for interventional daycare patients. If these tasks are simply added to the list of responsibilities for the interventionalists in the catheterization laboratory, a lack of patient supervision and delays in the discharge process, especially in high volume interventional centers, will be the consequence.

In the outpatient clinic either beds or lounge chairs enabling the patients to relax and enjoy some privacy should be present. While live telemetry of ECG and oxygen saturation as in intermediate care units is not considered necessary, nurses should be present at all times during opening hours and there should be a standard protocol detailing the frequency for preinterventional and postinterventional surveillance. Emergency medical equipment for intubation/ resuscitation, pericardiocentesis, and at least one echocardiography device and one ECG printer should be present in the outpatient clinic to facilitate swift diagnostics and intervention if a patient's condition should deteriorate. If the outpatient beds are included into the inventory of a regular ward, the facilities and technical outfit of the normal ward are sufficient.

Preprocedural considerations

The plan to perform a coronary procedure in an outpatient setting should be discussed with the patient as early as possible and personal concerns and questions should be addressed in an effort of shared decision making.

Relevant social and medical patient characteristics preventing the planned procedure from being performed in a SDD setting are summarized as a checklist in Table 2. One of the critical criteria at this time is the presence of a partner or relative who could assist in transporting the patient to the clinic and back home for the preclinical outpatient assessment and on the day of the procedure. Furthermore, advanced age or serious medical conditions should be taken into consideration. These include decompensated heart failure with LVEF <30%, severe kidney disease and uncontrolled diabetes mellitus. If clinically stable and compensated, however, these chronic conditions do not exclude SDD PCP.

 Table 2
 Checklist for SDD evaluation—Unfavorable factors before admission (adapted from [3–5])

 Social factors
 Social factors

- 1. No adequate home support after the planned procedure
- 2. Patient or caregiver not able to reach emergency medical support if necessary
- 3. Patient is mentally incapacitated

4. Language barrier compromising comprehension of instructions *Medical history*

- 1. LVEF < 30% or decompensated heart failure (NYHA class 3–4)
- 2. Decompensated kidney disease (eGFR $< 30 \text{ ml/min}/1.73 \text{ m}^2 \text{ BSA})$
- 3. Uncontrolled diabetes mellitus, risk of hypoglycemia
- Uncontrolled hypertension (systolic BP > 160 mm Hg despite 3 medications)
- 5. Uncontrolled/exacerbated COPD with home oxygen therapy
- Severe peripheral artery disease compromising radial/femoral access
 Severe contrast dye allergy
- Acute myocardial ischemia (ACS, NSTEMI, STEMI), same day transfer to other hospital is possible

Checklist to assess feasibility of SDD strategy depending on patient's social background and past medical history

Coronary angiography and PCI

Specific early slots in the catheterization laboratory program are required for SDD patients, so that the mandatory period of 3–6h radial compression and clinical observation can be completed without compromising discharge in the afternoon/evening.

All SDD coronary procedures should only be performed by experienced operators with an expertise in non-femoral puncture techniques including radial or ulnar access. Radial access facilitates early mobilization and is associated with lower mortality, fewer adverse cardiovascular events, and bleeding complications [51, 52]. Furthermore, there is a clear patient preference of radial versus femoral access PCPs [8, 30] (see above). The interventional team should be experienced in treating challenging radial anatomies and potential access site complications. If the access site needs to be switched to brachial or femoral due to vascular calcifications or complications during radial puncture, the patient should be scheduled for OS.

As highlighted by the 2021 ACC [3] and the 2018 SCAI [5] expert consensus documents, the clinical condition of the patient during and postprocedure has a greater impact on the final decision on SDD or the length of stay than individual features of the intervention. As summarized in our literature review [14], there is a wide range of RCTs and observational studies supporting the safety of SDD post-PCI even in challenging settings. Consistent with the ACC and SCAI recommendations, we consider SDD postelective PCI feasible and safe, including multivessel DES implantation, bifurcations, proximal/ostial lesions, and successful CTO. If complications or periprocedural adverse events summarized as a checklist in Table 3 occur, SDD is no longer safely possible and the clinical management should be altered to OS, potentially with constant surveillance and treatment depending on the nature of the adverse event. It should be emphasized that the scope of interventions offered within a daycare clinic should reflect the individual interventional center's expertise and scope of interventions.

A large observational study on SDD PCI from 2012 showed excellent safety data in a broad range of patients outside the SCAI/ACC recommendations [13] at that time (2009). These results highlight that any guidelines or position statements should acknowledge further trends in clinical practice standards with developing clinical evidence on safety.

Postprocedural monitoring

Consistent with elective DAs and PCIs in OS patients, postprocedural ECGs can be considered, cardiac enzyme tests are not necessary in the SDD setting, except in the presence of periprocedural complications, persistent symptoms indicative of myocardial ischemia or cardiac decompensation. In this context

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ACS acute coronary syndrome, BP blood pressure, BSA body surface area, COPD chronic obstructive lung disease, eGFR estimated glomerular filtration rate, LVEF left ventricular ejection fraction, NSTEMI non ST-elevated myocardial infarction, NYHA New York Heart Association, STEMI ST-elevated myocardial infarction

 Table 3
 Checklist—Procedural factors against SDD (adapted from [3–5])

 PCI—Local factors

1.	Complication at access site (bleeding, major dissection)			
2.	Contrast dye usage exceeding 500 ml [50]			
PCI—C	Coronary anatomy and procedural factors			
1.	Complications with coronary artery dissection or perforation			
2.	Persistent slow flow, no reflow in target vessel (TIMI < Grade 3)			
3.	Thrombus formation in any coronary vessel			
4.	Any SB loss if clinically significant (ST-segment elevation, persistent AP, arrhythmia)			
5.	Failure to deliver stent post predilatation			
6.	Peri-interventional ischemia with ST-segment elevation			
7.	Hemodynamic instability with inotropic support or left ventricular assist device			
8.	Rhythmologic instability with higher degree AV block or ventricular tachycardia			
9.	Highly complex or prolonged procedure that may put the patient at in- creased risk of adverse outcomes according to the clinical judgement of the operator			
Genera	I clinical condition			
1.	Pulmonary edema requiring oxygen support or diuretic therapy			
2.	Persistent ST segment alteration or chest pain			
3.	Severe contrast allergy requiring medical therapy			
NO cor	traindication against SDD policy [3, 5]			
1.	Single or multivessel PCI including proximal LAD or bifurcation			
2.	Multiple DES implants into one or more target vessels irrespective of stent number or cumulative stent length			
3.	Overstenting side branch > 1 mm with TIMI 3 flow			
4.	Uncomplicated successful CTO attempt			
5.	Ulnar access or distal radial access			
6.	Staged procedures post initial NSTEMI or STEMI			
Intraprocedural and postprocedural adverse events based on local factors, interventional course of events and general medical conditions which are contra-indications against a SDD strategy. The last 6 items are listed to point out specific items compatible with same-day discharge, provided the clinical				

out specific tiems compatible with same-day discharge, provided the clinical surveillance from procedure to discharge is without major adverse events. *CTO* chronic total occlusion, *LAD* left anterior descending coronary artery, *NSTEMI* non ST-elevated myocardial infarction, *PCI* percutaneous coronary intervention, *STEMI* ST-elevated myocardial infarction, *TIMI* thrombolysis in myocardial infarction

it should be noted that two large retrospective studies with cohorts >5000 patients have established that while a preinterventional elevation in serum Trop-T correlates with an adverse outcome, in the absence of periprocedural myocardial infarction a postinterventional rise in Trop-T does not offer any prognostic significance beyond the preinterventional value [53, 54]. In the SDD setting, an age-matched study of 149 SDD versus 154 OS patients showed that post-PCI Trop-T levels >5×upper reference level (URL) did not indicate any incremental short-term or long-term risk [55]. There is no need for bed rest or confinement to a single room in the daycare clinic. Nevertheless, a standardized monitoring protocol should be in place during the 3-6h of compression bandage. The puncture site and blood pressure should be checked and recorded at regular intervals (e.g. $1 \times /h$).

 Table 4
 Checklist—Postprocedural/clinical observation factors against SDD (adapted from [3–5])

Do NOT schedule SDD if:

Proced	ocedure outcome			
1.	Procedural outcome not compatible with SDD (see Table 3)			
2.	Any major concern from the interventional team			
Postpr	ocedural monitoring			
1.	Any clinically relevant decompensation of previous medical condi- tions (COPD, diabetes mellitus, hypertension, heart failure, renal failure, chronic pain)			
2.	Any change in mental state indicative of ischemia or dementia			
3.	Persistent chest pain			
4.	Persistent ECG abnormalities from baseline ECG			
5.	Persistent symptoms of contrast dye allergy			
6.	Major hematoma postpuncture site dressing removal			
7.	Patient not feeling well, unwilling to be discharged before 24 h observation			
Social	factors			
1.	No contact person for transport home or inadequate home support			
2.	Inadequate access to emergency medical care (> 30 min driving time)			
3.	Patient not able to take DAPT responsibly			
<i>COPD</i> chronic obstructive pulmonary disease, <i>ECG</i> electrocardiogram, <i>DAPT</i> dual antiplatelet therapy, <i>SDD</i> same-day discharge Procedural factors, postprocedural monitoring and social factors incompatible with SDD strategy				

In the presence of adverse clinical events, summarized in Table 4, adequate diagnostic measures including BP testing, trans-thoracic echocardiography (TTE), ECG and laboratory tests must be taken. In cases of adverse outcome, the clinical strategy must be changed from SDD to OS and this explained to the patient. This also involves unforeseen social patient factors, e.g., if the personal contact providing transfer from the clinic and home support is no longer available or if the patient may no longer feel comfortable with the concept of SDD.

Discharge decision

The final decision to discharge the patient on the same day should finally be made in agreement with the patient and by nurses, catheterization laboratory personnel, interventionalists and an experienced physician at the ward.

If the outcome of the PCP is successful and postinterventional monitoring was uneventful without changes in pre-existing medical conditions, discharge can be scheduled after removal of the radial pressure bandage. The compression time should be 3 h after DA and 4–6 h after PCI. An experienced doctor, ideally the interventionalist, should inspect the patient's condition and the puncture site after removal of the pressure bandage and give the final decision of discharge.

A structured discharge process with a checklist containing all necessary steps is recommended. An example of a discharge checklist is provided in Table 5. The

Table 5 Discharge checklist (adapted from [5])	
Checklist for discharge	
1. Check access site/radial pulse and cover with sterile dressin	g
2. Provide additional wound dressings for ambulatory period	
 Explain procedure result, changes in medication Document instructions on type and duration of DAPT, if PCI v performed Provide prescription of ASS and P2Y12 inhibitor for 30 days 	vas
4. Explain rules of behaviour to protect the puncture site for 7 d	ays
5. Provide discharge letter	
 Instruct the patient on emergency medical service, hand out gency telephone contact to clinic in case of bleeding, chest p other adverse events 	emer- Dain or
7. Notify the patient of a telephone follow-up call on the following day	ng wo
ASS acetyl-salicylic acid, DAPT dual antiplatelet therapy, PCI percuta coronary intervention Concise checklist of clinical assessments and duties to be performed	ineous 1 hv

attending nurses and doctors before discharge from the outpatient clinic

patient needs to be instructed on changes in the medical treatment, especially on the importance of DAPT in case of PCI with DES implantation. After PCI, the patient should leave the daycare clinic with a prescription of DAPT to provide a gapless concomitant therapy poststent implant. Ideally, a telephone follow-up call is arranged for the next working day in order to verify the patient's well-being after discharge.

Clinical follow-up of SDD PCP patients

Consistent with the ACC 2021 expert consensus [3], a clinical follow-up (FU) of SDD PCP patients should be considered in order to collect data on perceived quality of care, potential complications post discharge and to reinforce DAPT medication adherence. This could be done via telephone call on the day post PCP, without the patient having to visit the outpatient department again. It has been documented in prospective studies that DAPT adherence post-SDD PCI is high (87-95%) [8, 56]. In the latter study, nurse-led post-SDD PCI telephone interviews had no influence on P2Y12 inhibitor adherence (telephone 95% vs. control 93%, p=0.627) [56]; however, the authors did find that routine follow-up calls significantly reduced the frequency of unscheduled patient readmittance (8% vs. 16%, p=0.048), as well as self-initiated contacts to general practitioners (29% vs. 42%, p=0.02) [56]. A study from the Netherlands formally assessed the preferred mode of FU post-SDD treatment in 1797 patients treated from 2008 to 2012 [57]. Remarkably, the majority of patients (69.9%) preferred a FU by mail questionnaire, while only a smaller portion of patients preferred a telephone call (13.4%) or email (12.7%; p < 0.001) [57]. It is open to speculation, if this trend has changed in the meantime with a more dominant role of mobile telephones and email access even for older patients. While the AGIK does not mandate strict clinical follow-ups as they bind substantial resources, it must be pointed out that due to the novelty of the concept of SDD and the substantial impact it has on the post-PCI FU process, individual FU data from Austria would be valuable. Alternatively, patient follow-up and aftercare could be performed by resident physicians if close cooperation with the team of the catheter lab is guaranteed. The more critical an individual interventional center views the concept of SDD, the more rigorously it should be collecting clinical FU data.

Conclusion

Due to the rapid evolution of PCPs with predominantly radial access, 3rd generation DES and modern concomitant medical treatment including P2Y12 inhibitors, the safety profile has substantially improved. As discussed above, large longitudinal studies have documented an increase in radial access PCI and a stable low risk for MACE during/post-PCPs despite a trend towards more elderly, more compromized patients and more complex procedures [2]. Despite more recent trials enrolling larger numbers of complex patients [28] and procedures like MV, multistent, bifurcation, CTO [58] and LM PCIs [59, 60], 24-h and 30-day MACE frequencies have remained at low rates comparable to OS patients. Gilchrist et al. performed 100 SDD versus 665 OS PCIs in higher risk patients including complex and MV PCI [61]. Although only 15% of ambulatory patients would have qualified for SDD according to the 2009 SCAI criteria [13], the authors showed excellent safety data with absence of any MACE from 6-24h or at 30 days. These findings highlight the rapid evolution of clinical practice surrounding SDD catheterization laboratory programs.

Austrian interventional cardiology clinics have widely adopted this concept and offer PCPs on a SDD basis. Interventional outpatient departments require experienced personnel at all levels familiar with proper patient stratification preprocedure, PCI via radial access even in complex cases and structured surveillance postprocedure. The Austrian Society of Cardiology has considered recent publications from the ACC [3] and SCAI [4, 5, 13] and a wide scope of publications on radial access SDD PCI to develop an evidence base for its own individual position. Based on data from 4 expert consensus statements [3–5, 13], 4 large meta-analyses [19-22], and a separate literature review performed by the AGIK [14], spanning 21 years of clinical evidence from 2001 to 2022, the practice of SDD DA and PCI can be considered safe and feasible for a subset of CAD patients carefully selected by the criteria detailed above (Table 2). The SDD outpatient clinics provide additional value in fulfilling a well-documented patient preference towards shorter hospital stay following PCPs. Furthermore, SDD has been shown to provide substantial costsaving potential.

Three principles seem to be of paramount importance for a successful outcome: (1) adequate patient



Fig. 1 Concise checklist summarizing inclusion/exclusion criteria for SDD strategy prior to admission, during/after coronary procedure and during clinical follow-up/before discharge

selection with clinical assessment by an experienced cardiologist and (2) adjusting the complexity and scope of SDD procedures to the general experience and scope of the interventional team. (3) Adequate aftercare supplied by experienced medical personnel in or out of hospital should be guaranteed. A concise checklist summarizing pre admission, procedural, post-procedural, and follow-up criteria is illustrated in Fig. 1.

These recommendations are subject to changes with further publications in this area and not intended as finite or binding. The interventional centers' individual policies and interventionalists' expert opinions are the most important domains in transforming these recommendations into daily clinical practice.

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Conflict of interest M.C. Brandt, H. Alber, R. Berger, R.K. Binder, J. Mascherbauer, A. Niessner, M. Schmid and M. Frick declare that they have no competing interests.

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