



Patient hospital label	Hospital patient identification number	
	Surname	Date of birth dd.mm.yyyy
	First name	Biological sex <input type="radio"/> f <input type="radio"/> m

Baseline characteristics

cvRF <input type="radio"/> None <input type="checkbox"/> Arterial hypertension (on treatment) <input type="checkbox"/> Diabetes requiring medication	<input type="checkbox"/> Dyslipidemia (on statin or LDL-C > 2.6 mmol/L) <input type="checkbox"/> Smoking (active) <input type="checkbox"/> Family history <input type="radio"/> cvRF unknown / not documented
Patient history <input type="radio"/> None <input type="checkbox"/> Prior PCI (with or without stents) <input type="checkbox"/> Prior CABG <input type="radio"/> Unknown / not documented	LVEF (By laevugram or non-invasive imaging test closest to CA) <input type="radio"/> Normal (50-70%) <input type="radio"/> Unknown / not documented <input type="radio"/> Mildly reduced (40-49%) <input type="radio"/> Moderately reduced (30-39%) <input type="radio"/> Severely reduced (<30%)
Out of hospital cardiac arrest <input type="radio"/> Yes <input type="radio"/> No	Cardiogenic shock (SBP < 90 mmHg for 30 min or vasopressor to maintain, and lactate > 2 mmol/L) <input type="radio"/> Yes <input type="radio"/> No
Intubated <input type="radio"/> Yes <input type="radio"/> No	Mechanical support device(s) (in case of either cardiogenic shock or high-risk PCI) <input type="radio"/> None <input type="checkbox"/> Impella <input type="checkbox"/> ECMO

Indication

Indication (only one answer) <input type="radio"/> Clinical presentation as ACS <input type="radio"/> Suspected CAD or suspected progression of known CAD <input type="radio"/> Planned valvular heart disease intervention <input type="radio"/> Follow-up after HTX <input type="radio"/> Heart failure (LVEF < 40%) <input type="radio"/> Routine follow-up (e.g. after left main PCI) <input type="radio"/> Tachy- or bradyarrhythmia <input type="radio"/> Planned vascular surgery <input type="radio"/> Planned major surgery	If ACS, please specify (Cardiac arrest patients without STEMI count as NSTEMI) <input type="radio"/> Unstable angina pectoris (new or progressive angina pectoris, according to Braunwald definition) <input type="radio"/> Non-STEMI <input type="radio"/> STEMI If suspected CAD was indicated, please choose the following Coronary angiography-preceding test(s) suggesting CAD (high-pretest likelihood) <input type="radio"/> None <input type="checkbox"/> Treadmill indicating ischemia (or other exercise stress tests) <input type="checkbox"/> CCTA with significant stenoses <input type="checkbox"/> Stress MRI with ischemia <input type="checkbox"/> Stress TTE with ischemia <input type="checkbox"/> TTE with hypokinesia <input type="checkbox"/> Nuclear test with ischemia
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Symptoms if suspected or stable CAD

Dyspnoe

☐ Yes ☐ No

Angina pectoris

☐ Yes ☐ CCS ☐ I ☐ II ☐ III ☐ IV

☐ No

Anti-anginal drugs (prior or current, betablocker, Calcium channel blocker, long-acting nitrate, Ivabradine, Ranolazine, Trimetazidine)

☐ Yes ☐ No

STEMI-related referral times

Date of symptoms onset

dd.mm.yyyy

Date of hospital entry

dd.mm.yyyy

Date of recanalization

dd.mm.yyyy

Time of symptoms onset

hh:mm

☐ Symptoms onset unknown / not documented

Time of hospital entry

hh:mm

Time of recanalization

hh:mm

Non-STEMI-related referral times

Time between first ECG and coronary angiography

☐ <24 hours ☐ 24-48 hours ☐ >48 hours ☐ unknown / not documented

General procedural characteristics

Date of procedure

dd.mm.yyyy

Access (access through which the PCI was finalized)

☐ Femoral (Radial attempt converted to femoral counts as femoral)
☐ Radial (Brachial counts as radial)

Dose unit

☐ mGycm2
☐ cGycm2 or uGym2

Dose (round to whole number)

PCI-specific procedural characteristics

- ☐ **PCI aborted** Only complete CTO procedure, intracoronary physiology and imaging

Number of treated lesions (Coronary artery lesion undergoing PCI. If two lesions are covered by two or more overlapping stents, this counts as one lesion)

Number of treated vessels

LM, LAD, LCX, RCA, or graft. If a lesion extends over several vessels only the main vessel is counted

Unprotected left main stenosis treated

☐ Yes ☐ No

Number of implanted stents

Number of used drug-eluting balloons

Intervention(s) for stent failure

- ☐ None
☐ Restenosis
☐ Stent thrombosis

CTO procedure

(Chronic occlusion presumably > 3 months, Subacute ACS presentation excluded)

- ☐ None ☐ Antegrade ☐ Retrograde

Calcium modification

- ☐ None ☐ Rotablation
☐ Cutting balloon ☐ Lithotripsy
☐ Scoring balloon ☐ Orbital atherectomy
☐ High-pressure balloon (RBP \geq 30 atm)

Bifurcation

- ☐ None
☐ 1 stent
☐ 2 stents

Intracoronary physiology

- ☐ None ☐ Resting index (iFR/RFR) ☐ FFR

Intracoronary imaging

- ☐ None ☐ IVUS ☐ OCT

Complication

Major complication

- ☐ None ☐ Clinically overt stroke
☐ Emergency open heart surgery ☐ Emergency vascular surgery (non-cardiac)
☐ Procedural death ☐ Pericardial tamponade

Prescribed medication

Prescribed medication at the time of discharge from CathLab, including previously prescribed medication that will be continued

Antithrombotic medication

- ☐ ASA ☐ P2Y12 inhibitor ☐ N/OAC

If P2Y12 inhibitor, specify

- ☐ Clopidogrel ☐ Ticagrelor
☐ Prasugrel ☐ Other

Statin therapy

- ☐ None
☐ Low-intensity (or moderate-intensity)
☐ High-intensity (Rosuvastatin \geq 20mg, Atorvastatin \geq 40mg)

If dual antiplatelet therapy, specify duration (select closest answer)

- ☐ <1 month ☐ 6 months
☐ 1 month ☐ 12 months
☐ 3 months ☐ >12 months

If triple antithrombotic therapy, specify duration (select closest answer)

- ☐ <1 week ☐ 3 months
☐ 1 month ☐ 6 months