

Japanese CTO PCI Expert Registry

Date	
Patient Name	
PCI ID	

Please fill in the columns highlighted in yellow with Ball Point pen.

←Please fill in the ID if lesion registration is done online

Patient Basic Information 1

Please note that all fields followed by an asterisk must be filled in.

Patient registration category	<input type="radio"/> Case at the affiliated hospital <input type="radio"/> non-affiliated hospital (in own country) <input type="radio"/> non-affiliated hospital (overseas)			
Hospital Name*				
Hospital Dr. in Charge*	Name*		Title*	
	Tel. *		E-mail*	

Japanese Operator Name		Operator ID		Registration Date	
Patient identification code			Patient name Initials		
Patient Identification memorandum					

Basic Information	Date of Procedure*		Patient DOB		Age	
	SEX	<input type="radio"/> Male <input type="radio"/> Female	Height	cm	Weight*	kg

Basic Information

Medical History	OMI ¹⁾	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown
	Prior CABG ²⁾	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown
	Prior PCI	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown

1) OMI : Please mark "YES" if diagnosed OMI in past and myocardium is not well.

2) CABG History to be diagnosed at when at First PCI.

Risk Factor	Hypertension	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown
	Diabetes	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown Choose "Yes" for diabetes Current Treatment: <input type="radio"/> None <input type="radio"/> Dietary Only <input type="radio"/> Oral Anti diabetic Drug <input type="radio"/> Insulin Therapy
	Dyslipidemia	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown ※If Statin is prescribed, "Yes"
	Smoking	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown If Yes : <input type="radio"/> Currently Smoking <input type="radio"/> Smoking in Past ※"Smoking history" means over 1 month after stop-smoking
	ASO	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown
	Coexisted Heart Disease	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown If Yes : <input type="radio"/> family history <input type="radio"/> congenital heart disease <input type="radio"/> Valve disease <input type="radio"/> pacemaker implanted <input type="radio"/> Arrhythmia <input type="radio"/> Others :
	Comorbidity	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown If yes: <input type="radio"/> COPD <input type="radio"/> Dialysis <input type="radio"/> Cerebral Vascular Disorder <input type="radio"/> Others :

<Definition of CTO>

1. TIMI 0 (no antegrade flow except bridging/ipsilateral collateral flow)
2. more than 3 months / unknown occlusion duration
3. a total occlusion in a major coronary artery with significant territory or a graft occlusion
4. Multi-occlusion in a same coronary artery (tandem CTO) is deemed to be a single CTO.

Case Basic Information

Clinical application for CTO procedure	<input type="radio"/> Asympamatic Isch <input type="radio"/> Stable Angina <input type="radio"/> Unstable Angina <input type="radio"/> OMI <input type="radio"/> AMI		
Angina (CCS Classification)	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown If Yes <input type="radio"/> I <input type="radio"/> II <input type="radio"/> III <input type="radio"/> IV		
cardiac insufficiencyheart failure (NYHA Classification)	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown If Yes <input type="radio"/> I <input type="radio"/> II <input type="radio"/> III <input type="radio"/> IV		
ECG (CTO Area)	<input type="radio"/> Normal <input type="radio"/> Abnormal Q Wave <input type="radio"/> Abnormal ST-T		
Exercise Tolerance Test	<input type="radio"/> Not Tested <input type="radio"/> Negative <input type="radio"/> Positive <input type="radio"/> Unratable		
MSCT	<input type="radio"/> Not Tested <input type="radio"/> Tested <input type="radio"/> Unknown		
Pre Op Serum-Cre	mg/dl (Date:)	eGFR ¹⁾	
LVEF (%)	%	Assessment Procedure	<input type="radio"/> UCG <input type="radio"/> RI <input type="radio"/> MRI <input type="radio"/> LVG <input type="radio"/> MSCT
Asynergy (CTO area)	<input type="radio"/> Normal <input type="radio"/> Hypokinesis <input type="radio"/> Akinesis <input type="radio"/> Dyskinesis		
Residual Myocardium	<input type="radio"/> Existing <input type="radio"/> None <input type="radio"/> Unknown	Assessment Procedure	<input type="radio"/> UCG <input type="radio"/> RI <input type="radio"/> MRI <input type="radio"/> LVG <input type="radio"/> MSCT
EuroScore II ²⁾			

<pretreatment therapeutic strategy>

Target revascularization ¹⁾	<input type="radio"/> Plan for Target revascularization <input type="radio"/> non complete revascularization If choose "No" for target revascularization residual artery: <input type="checkbox"/> RCA <input type="checkbox"/> LAD <input type="checkbox"/> LCX <input type="checkbox"/> LMT <input type="checkbox"/> Graft Reason of incomplete revascularization : <input type="checkbox"/> Residual myocardium <input type="checkbox"/> small vessel / small Myocardial perfusion <input type="checkbox"/> Technically difficult <input type="checkbox"/> clinically significance
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- 1) Plan for Target revascularization : It has a significant lesion (main coronary branch) for complete revascularization
- 2) Treating period (staged PCI) regardless of the times of hospitalization

<Coronary lesion Basic information>

Multivessel disease ¹⁾	<input type="checkbox"/> SVD <input type="checkbox"/> DVD <input type="checkbox"/> TVD <input type="checkbox"/> LMT + MVD				
Significant lesions of Coronary artery	<input type="checkbox"/> RCA	<input type="checkbox"/> 1	<input type="checkbox"/> 2	<input type="checkbox"/> 3	<input type="checkbox"/> others
	<input type="checkbox"/> LAD	<input type="checkbox"/> 6	<input type="checkbox"/> 7	<input type="checkbox"/> 8	<input type="checkbox"/> 9 <input type="checkbox"/> 10
	<input type="checkbox"/> LCX	<input type="checkbox"/> 11	<input type="checkbox"/> 12	<input type="checkbox"/> 13	<input type="checkbox"/> 14 <input type="checkbox"/> 15
	<input type="checkbox"/> LMT	<input type="checkbox"/> ostium <input type="checkbox"/> body <input type="checkbox"/> bifurcation			
	<input type="checkbox"/> Graft	<input type="checkbox"/> LITA	<input type="checkbox"/> RITA	<input type="checkbox"/> RA	<input type="checkbox"/> SVG <input type="checkbox"/> GEA
	Graft Anastomosis destination: <input type="checkbox"/> →RCA <input type="checkbox"/> →LAD <input type="checkbox"/> →LCX				
	Comment				
CTO artery	<input type="checkbox"/> RCA <input type="checkbox"/> LAD <input type="checkbox"/> LCX <input type="checkbox"/> LMT <input type="checkbox"/> Graft				

(if graft is intact, grafted native artery non-CTO)

Jeopardized Collateral	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unknown
	If yes, have been treated with donor artery before CTO_PCI: <input type="radio"/> Yes <input type="radio"/> No

標的 CTO 病変数	<input type="radio"/> 1 lesion <input type="radio"/> 2 lesion
標的病変枝	<input type="checkbox"/> RCA <input type="checkbox"/> LAD <input type="checkbox"/> LCX <input type="checkbox"/> LMT <input type="checkbox"/> Graft
	Graft anastomosis: <input type="checkbox"/> →RCA <input type="checkbox"/> →LAD <input type="checkbox"/> →LCX

Only select by Core laboratory (Operator is NOT required)	
Preparative angio image	<input type="radio"/> Yes <input type="radio"/> No
Syntax Score ²⁾	<input type="radio"/> acalculia
The propriety of Anatomical adaption	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> unevaluable
	「No」:
	<input type="radio"/> Did not admit the same current level as main coronary artery or main coronary artery branch
	<input type="radio"/> TIMI ≥ 1
	<input type="radio"/> Tandem OR same main coronary artery branch
	<input type="radio"/> Other ()

1) staged PCI の場合、一連の PCI 前の罹患枝数

2) Syntax Score:(<http://www.syntaxscore.com/>)※ : Needs to be filled out by both operator and core laboratory※ : Needs to be filled out by core laboratory (operator is not required)

- | |
|--|
| <input type="checkbox"/> Check box: Select from plurality items
<input type="radio"/> Radio button: Choose only one |
|--|

② **Retrograde(+Antegrade)**

Including Case in which Retrograde channel crossed and prepared for Retrograde but wire continuously crossed antegrade Retro +Ante technique (Kissing wire Cross,Reverse CART,CART etc.)

③ **Retrograde(+Antegrade)Unsuccessful → Antegrade**

Cases in which; Failed to cross Retrograde channel / Unable to complete operation due to complication

(EX; perforation, dissection, Thrombus) of Retrograde system/ Operation continued after removing retrograde system

In this chosen category, retrograde+antegrade items are required to be filled out

<CTO Procedure information (Lesion1)>

【STENT information lesion1】	
Stenting	<input type="radio"/> Yes <input type="radio"/> No
Used STENT	<input type="checkbox"/> BMS <input type="checkbox"/> DES <input type="checkbox"/> BVS
Bifurcation stenting	<input type="radio"/> Yes <input type="radio"/> No
	If "Yes", bifurcation stenting method: <input type="checkbox"/> single stenting <input type="checkbox"/> double stenting
Number of stent- CTO branch only	()
	For over 2 stents- stent overlap: <input type="radio"/> Yes <input type="radio"/> No
Maximum diameter of the used stent	<input type="radio"/> <2.5mm <input type="radio"/> 2.5~2.9 <input type="radio"/> 3.0~3.4 <input type="radio"/> >3.5mm
Sum of stent length	mm (The sum of stent length used for CTO branch)
Used DES	<input type="checkbox"/> Promus <input type="checkbox"/> Xience <input type="checkbox"/> Nobori <input type="checkbox"/> Resolute <input type="checkbox"/> Endeavor <input type="checkbox"/> Ultimaster <input type="checkbox"/> Synergy <input type="checkbox"/> Other product name ()
post STENT balloon	Post dilatation (MAX) mm
Usage of DEB	<input type="radio"/> Yes <input type="radio"/> No If yes, DEB diameter: mm

【Antegrade only】 *Only if "Antegrade only" was chosen at strategy.	
Usage of contrast medium	<input type="radio"/> Yes <input type="radio"/> No
Crossed guide wire via Antegrade	<input type="radio"/> Successful <input type="radio"/> Unsuccessful
	Reason of "Unsuccessful": <input type="checkbox"/> GW unable to pass CTO <input type="checkbox"/> device failed to reach CTO <input type="checkbox"/> Suspension of PCI due to complication <input type="checkbox"/> Other()
1 st guide wire via antegrade	Name of product:
1 st guide wire via antegrade for crossing CTO	Name of product:
Last wire /Crossed wire	<input type="radio"/> Last (Finally used GW) <input type="radio"/> Crossed (GW used to cross)
	Name of product:

Method or special device for Last wire/Crossed wire	<input type="checkbox"/> single wire <input type="checkbox"/> double wire method <input type="checkbox"/> IVUS guide crossing <input type="checkbox"/> STAR <input type="checkbox"/> Stingray <input type="checkbox"/> other technic / devices ()
GW Usage of step up / step down method	<input type="radio"/> Yes <input type="radio"/> No If "Yes": <input type="radio"/> step up <input type="radio"/> step down <input type="radio"/> both
Used OTW or Penetration catheter	<input type="radio"/> Yes <input type="radio"/> No If "Yes", name of product: <input type="checkbox"/> Corsair <input type="checkbox"/> Finecross <input type="checkbox"/> Crusade <input type="checkbox"/> Tornus <input type="checkbox"/> Caravel <input type="checkbox"/> GuideLiner <input type="checkbox"/> Guidezila <input type="checkbox"/> OTW <input type="checkbox"/> Other()

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<CTO procedure info(Lesion1)>

【Retrograde + Antegrade】	
*Fill in if chosen Retrograde + Antegrade and retro+ante unsuccessful → ante in strategy.	
Retrograde indication	<input type="radio"/> Accessed Retrograde for Antegrade backup, but crossed with Antegrade *If this item is chosen, fill in only the *1 parts <input type="radio"/> Changed course to Retrograde due to failure of Antegrade <input type="radio"/> Initially scheduled to pursue retrograde (Ante/Retro order regardless)
Antegrade GW 1)	<input type="radio"/> Yes <input type="radio"/> No If "yes" name of product: ()
Antegrade start CTO GW 1)	<input type="radio"/> Yes <input type="radio"/> No If "yes" name of product: ()
Completed /Crossed Antegrade GW 1)	<input type="radio"/> Completed (Final GW used) <input type="radio"/> Crossed (crossed GW) <input type="radio"/> None If "Completed" or "Crossed", Name of product: ()
Procedure method or special device used to complete/cross GW 1)	<input type="checkbox"/> single wire <input type="checkbox"/> double wire method <input type="checkbox"/> IVUS guide crossing <input type="checkbox"/> STAR <input type="checkbox"/> Stingray <input type="checkbox"/> other technic / devices ()
GW step up / step down method 1)	<input type="radio"/> Yes <input type="radio"/> No If "Yes": <input type="radio"/> step up <input type="radio"/> step down <input type="radio"/> both
Antegrade-Usage of Penetration catheter /Special device 1)	<input type="radio"/> Yes <input type="radio"/> No If "Yes", Name of product: <input type="checkbox"/> Corsair <input type="checkbox"/> Finecross <input type="checkbox"/> Crusade <input type="checkbox"/> Tornus <input type="checkbox"/> Caravel <input type="checkbox"/> GuideLiner <input type="checkbox"/> Guidezila <input type="checkbox"/> OTW <input type="checkbox"/> Other catheter() <input type="checkbox"/> Other special device()
fluoro time for Antegrade GW 1)	<input type="radio"/> min. <input type="radio"/> Unknown

Attempted channel <input type="checkbox"/>	<input type="checkbox"/> septal <input type="checkbox"/> epicardial <input type="checkbox"/> atrial <input type="checkbox"/> graft If "septal", Septal surfing: <input type="radio"/> unattempt <input type="radio"/> attempted but unsuccessful <input type="radio"/> Successful
Used channel to cross GW for Retrograde operation <input type="checkbox"/>	<input type="radio"/> Unsuccessful <input type="radio"/> septal <input type="radio"/> epicardial <input type="radio"/> atrial <input type="radio"/> graft <input type="radio"/> Other()
Approach for Retrograde channel <input type="checkbox"/>	<input type="radio"/> Successful <input type="radio"/> Unsuccessful If "Successful": <input type="radio"/> Both GW and catheter passed <input type="radio"/> Only GW passed
Access site (Retrograde)	<input type="checkbox"/> femoral <input type="checkbox"/> radial <input type="checkbox"/> brachial
Guiding catheter (for Retrograde)	<input type="checkbox"/> 5Fr <input type="checkbox"/> 6Fr <input type="checkbox"/> 7Fr <input type="checkbox"/> 8Fr <input type="checkbox"/> Retro access to ipsicollateral
GW crossed for Retro channel <input type="checkbox"/>	Name of product()
Starting GW for Retrograde to CTO	Name of product()
GW used up to the balloon at reverse CART	Name of product()
Completed /Crossed Retrograde GW	<input type="radio"/> Completed(last GW used) <input type="radio"/> Crossed (Crossed GW) <input type="radio"/> None
	If "Completed" or "Crossed", name of product: ()
Penetration catheter/Special device used for retrograde	<input type="radio"/> Yes <input type="radio"/> No
	If "Yes", Name of product: <input type="checkbox"/> Corsair <input type="checkbox"/> Finecross <input type="checkbox"/> Crusade <input type="checkbox"/> Tornus <input type="checkbox"/> Caravel <input type="checkbox"/> GuideLiner <input type="checkbox"/> Guidezilla <input type="checkbox"/> OTW <input type="checkbox"/> Other catheter() <input type="checkbox"/> Other special device()
CTO cross method / Special device (Final /Crossed)	<input type="radio"/> Kissing wire Cross <input type="radio"/> Retrograde Wire Cross <input type="radio"/> CART <input type="radio"/> Reverse CART <input type="radio"/> Other technique () <input type="radio"/> Special device to cross GW()
	GW cross direction: <input type="checkbox"/> Antegrade <input type="checkbox"/> Retrograde
After Retrograde GW crossing, alteration for Antegrade GW	<input type="radio"/> Successful <input type="radio"/> Unsuccessful <input type="radio"/> Unattempt
	Alteration method: <input type="radio"/> externalization <input type="radio"/> rendezvous <input type="radio"/> antegrade parallel wire <input type="radio"/> induced by retrograde balloon expansion <input type="radio"/> Other ()
Device pass	<input type="radio"/> Device passed <input type="radio"/> Unsuccessful

【Retrograde indication】 if choice input item

① Accessed Retrograde for Antegrade backup, but crossed with Antegrade *If this item is chosen, fill in only parts

② Changed course to retrograde due to failure of Antegrade ③Initially scheduled to pursue Retrograde (Ante/Retro order regardless)

If ②,③ is chosen fill in all 【Retrograde + Antegrade】 items.

Outcome

Success ¹⁾	<input type="radio"/> success <input type="radio"/> failure	
Complications in a hospital	Death	<input type="radio"/> None <input type="radio"/> Yes (<input type="radio"/> related to PCI <input type="radio"/> NOT related to PCI)
	MI	<input type="radio"/> None <input type="radio"/> Probably None <input type="radio"/> Yes <input type="radio"/> Unknown →If Yes: <input type="radio"/> QMI <input type="radio"/> Non-QMI <input type="radio"/> Unknown *the definition of Yes is more than 3 times CK score compare to normal
	Stent thrombosis	<input type="radio"/> None <input type="radio"/> Yes (<input type="radio"/> coronary occlusion <input type="radio"/> Non coronary occlusion)
	coronary embolization (TIMI 0-1)	<input type="radio"/> None <input type="radio"/> Yes
	cerebral stroke※ (cerebrovascular event)	<input type="radio"/> None <input type="radio"/> Yes (<input type="radio"/> Non bleeding event <input type="radio"/> bleeding event <input type="radio"/> Unknown) ※excluding TIA
	coronary artery perforation	<input type="radio"/> None <input type="radio"/> Minor event <input type="radio"/> pericardial bleeding (No need pericardial drainage) <input type="radio"/> pericardial bleeding (need pericardial drainage)
	Complications at the site of Puncture ²⁾	<input type="radio"/> None <input type="radio"/> Yes (surgical management : <input type="radio"/> None <input type="radio"/> Yes the bleeding calling for blood transfusion or hematoma: <input type="radio"/> None <input type="radio"/> Yes)
	Emergency CABG	<input type="radio"/> None <input type="radio"/> Yes
	Emergency PCI	<input type="radio"/> None <input type="radio"/> Yes
	Serum Cre after procedure	mg/dl (date of measurement :)
	Renal dysfunction (including CIN)	<input type="checkbox"/> None <input type="checkbox"/> more than 25% Increase in s-Cre <input type="checkbox"/> more than 0.5mg/dl increase in s-Cre

- 1) The definition of Success: a success of procedure without death/MI/Emergency CABG/cerebral stroke
2) The definition of Complications at the site of Puncture: blood transfusion(more than 2 units) or complications calling for surgical management

<Outcomes for acute phase>

★The information is not required if it is difficult to obtain for the overseas lesion★

Procedure ³⁾		min.	Contrast medium dose ⁴⁾		ml
Fluoroscopy time [Time]	Frontal(tube) (↓)	Time	Lesion 1	Time	
			Lesion 2	Time	
	Lateral(tube) (→)	Time	Lesion 1	Time	
			Lesion 2	Time	
Adsorbed dose (mGy)	Frontal(tube) (↓)	mGy	Lesion 1	mGy	
			Lesion 2	mGy	
	Lateral(tube) (→)	mGy	Lesion 1	mGy	
			Lesion 2	mGy	

- 3) Procedure time is defined as the time from the start of wire operation to the final angiography in the CTO procedure
- 4) Contrast medium dose is defined as the dose administered only in the CTO procedure
- Fluoroscopy time is defined as the time spent only for the CTO procedure
 - Adsorbed dose is defined as the dose deposited only in the CTO procedure

The condition within a month after PCI procedure

The visit within a month(or Two) after discharge	<input type="radio"/> Yes <input type="radio"/> None <input type="radio"/> Unknown
	In the case of non-visit, the contact hospital and Doctor
	Address
	Hospital
	TEL
Doctor	
CCS	<input type="radio"/> 0 <input type="radio"/> I <input type="radio"/> II <input type="radio"/> III <input type="radio"/> IV <input type="radio"/> Unknown
Radiating Dermatitis	<input type="radio"/> None <input type="radio"/> Yes <input type="radio"/> Unknown If yes: <input type="radio"/> Only blushing <input type="radio"/> Ulceration <input type="radio"/> Requirement for surgery <input type="radio"/> Unknown
Serum Cre after procedure (24hr<<1month)	mg/dl (date of measurement:)
Renal dysfunction (including CIN)	<input type="checkbox"/> None <input type="checkbox"/> more than 25% Increase in s-Cre <input type="checkbox"/> more than 0.5mg/dl increase in s-Cre
Non-fatal MI	<input type="radio"/> None <input type="radio"/> Yes <input type="radio"/> Unknown
Revascularization	<input type="radio"/> None <input type="radio"/> re-PCI <input type="radio"/> CABG <input type="radio"/> Unknown
Cerebrovascular Disorder	<input type="radio"/> None <input type="radio"/> Yes <input type="radio"/> Unknown
Stent Thrombosis	<input type="radio"/> None <input type="radio"/> possible <input type="radio"/> definite ST <input type="radio"/> Unknown (days)
Other complications	<input type="radio"/> None <input type="radio"/> Related to CTO-PCI <input type="radio"/> Non-related to CTO-PCI Comments:) <input type="radio"/> Unknown
Death	<input type="radio"/> None <input type="radio"/> Non-cardiac death <input type="radio"/> Cardiac death <input type="radio"/> Unknown Comments:) <input type="radio"/> Unknown
Follow-up imaging	<input type="radio"/> None <input type="radio"/> Yes <input type="radio"/> Unknown If yes: <input type="radio"/> CAG <input type="radio"/> MSCT, days after procedure restenosis: <input type="radio"/> restenosis(QCA DS>50%) <input type="radio"/> reocclusion <input type="radio"/> none