	L CARDIOVASCULAR DISEASE DATABASE (PCI REGISTRY) NOTIFICATION FORM For NCVD Use only: Centre:
	is form to notify all PCI admissions at your centre to NCVD PCI Registry. Where check boxes □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□
A1. Date of Admissio	A2.* Time of Admission :(hh/mm) (in 24hr clock)
SECTION 1 : DEMO	GRAPHICS
. Patient Name : (as per MyKad/Other Docur	2. Hospital RN: ment ID) (if applicable)
. Identification Card	MyKad :
Number :	Other ID document No:  Specify type (eg.passport, armed force ID, Old IC):
. Gender:	- 11 11 11 0
a. Date of Birth:	Male Female 5. Nationality: Malaysian Non Malaysian  6b. Age on admission:
(write DOB as 01/01/yy if age is known)	(dd/mm/yy) (Auto Calculate)
. Ethnic Group:	Malay Punjabi Melanau Bidayuh Foreigner, specify
•	<ul><li>○ Chinese ○ Orang Asli ○ Murut ○ Iban</li><li>○ Indian ○ Kadazan Dusun ○ Bajau ○ Other M'sian, specify :</li></ul>
. Contact Number	(1): (2):
SECTION 2 : STATU	JS BEFORE EVENT
. Smoking Status:	Never Former (quit >30 days) Current (any tobacco use within last 30 days) Not Available
. Medical history :	
a) Dyslipidaemia	
o) Hypertension	(Presence of 50% stenosis on CTA, angiogram, ischaemia on functional cardiac imaging such as nuclear, MRI, echo or positive treadmill test. High calcium score alone is not sufficient.)
c) Diabetes	
OHA Insulir	h) History of heart failure
d) Family history of prema	ature Yes No Not known i) Cerebrovascular disease Yes No Not known
cardiovascular disease	
(1st degree relative with eith	ner MI or stroke; < 55 y/old if Male & <65 y/old if Female)  k) Chronic renal failure  Yes  No  Not known
e) Myocardial infarction hi	story
SECTION 3 · CLINIC	CAL EXAMINATION and BASELINE INVESTIGATION
	a. Height: (kg) Not Available Not Available (Auto Calculate
. Heart rate	3. Blood pressure a. Systolic: (mm/le)
(at start of PCI in the cath lab):	(beats / min)  (beats / min)  (at start of PCI in the cath lab):  (b. Diastolic: (mmHg)
. Fasting Blood Glucose:	Not Available 5. Hb A1c: % Not Available
a. Total cholesterol:	6b. LDL Levels: mmol/L Not Available
. Baseline creatinine :	μmol/L Not Available 6c. HDL Levels: mmol/L Not Available
. Baseline ECG:	Sinus rhythm Atrial Fibrillation 2nd /3rd AVB LBBB RBBB ST Deviation (for GRACE Score)
. Non Invasive Test:	i.
0. Glomerular	a. MDRD: b. Cockcroft-Gault:
Filtration Rate (GFR):	(Auto Calculate) mL/min/1.73m <sup>2</sup> (Auto Calculate) mL/min
GFR (Modification of GFR (Cockcroft-Gau	Diet in Renal Disease (MDRD): 186 x (serum creatinine[micromol/L] / 88.4) <sup>1.154</sup> x (age) -0.203x (0.742 if female)  It formula): Male: 1.23 x (140 - Age) x Weight (kg) / serum Creatinine (micromol/L)
SECTION 4 : PREV	Female : 1.04 x (140 - Age) x Weight (kg) / serum Creatinine (micromol/L)  IOUS INTERVENTIONS
. Previous Yes	○ No 2. Previous ○ Yes ○ No
PCI ·     C	te of most recent PCI (dd/mm/yy):    Date of most recent CABG (dd/mm/yy):   Date of most recent CABG (dd/mm/yy):   Not Available

a. Patient Name :	b. M	lyKad / Other ID No:		c. Date of Procedure:						
SECTION 5 : CARDIAC STATUS AT PCI PROCEDURE										
1. Angina type:   None  Atypical  Typical										
2. Canadian Cardiovascular	9	CCS 0 = Asympto		© CCS 2 © CCS 3	© CCS 4					
3. NYHA:	○ NYHA I ○ NY	, ,	NYHA III	© NYHA						
4. Killip class :	I No clinical signs of HF									
(STEMI & NSTEMI)	<ul> <li>☐ I No clinical signs of HF</li> <li>☐ III Acute Pulmonary Oedema (APO)</li> <li>☐ IV Cardiogenic Shock</li> <li>☐ Not Applicable/Not Available</li> </ul>									
5. Coronary Artery  * Disease (CAD)	○ STEMI ○ NS	STEMI	◯ UA	Chronic Stable Angi	na					
Presentation:	* Anterior Posterior	<ul><li>Lateral</li><li>Right sided</li></ul>	<ul><li>Inferior</li><li>Left Main Stem</li></ul>	Others, specify:						
6. STEMI Event : (Please complete if <24 hours since onset of STEMI symptoms)	a) STEMI onset :	i. Date: d d / r	m m / y y	ii. Time: h h i m m (i	n 24 hr clock)					
	b) Arrival at first hospital (non PCI hospital) :	i. Date: d d / r	m m / y y	ii. Time: h h i m m (i	n 24 hr clock)					
	c) Arrival at PCI hospital:	i. Date: dd/n	m m / y y	ii. Time: h h i m m (i	n 24 hr clock)					
	d) First device activation (balloon inflation / stent / DCB / aspiration) :	i. Date: d d / n  Not Applicable	m m / y y	ii. Time: h h • m m (i	n 24 hr clock)					
	e) In hospital STEMI onset	i. Date: dd/	m_m / _y _y	ii. Time: h h i m m (i	n 24 hr clock)					
7. EF Status (at time of PCI procedure):	(Do not use % (Do not use % '>' or '<' symbol)    Not Available    8. Cardiac Arrest:    Arrest:   Out of hospital (only got STEMI & NSTEMI)    (Auto Calculate)									
SECTION 6 : CATH LA	B VISIT									
1.a) Date of procedure:	/ / (de	d/mm/yy)	1.b). Time of proced	dure:	(in 24 hr clock)					
2. PCI status:		PCI Ad hoc	⊚ STEMI	→ ○ Primary ○ Delaye	d Routine PCI					
*	<ul> <li>NSTEMI/UA → Urgent (within 24hrs)</li></ul>		Rescue Delayed Selective PCI Pharmacoinvasive							
2a) Door to balloon time (auto calculate) mins										
3. Medication:	*a) Thrombolytics	i) Time <pre> &lt;3hrs Duration: </pre> 3-6hrs <pre> 6-12l</pre>	rs >24hrs	ii) Types: Streptokina Tenectepla: Others:	se ( tPA					
	*b) Ilb / Illa Blockad	Prior During	g After	⊚ No						
	*c) Heparin Yes	○ No	* <u>d) LMWH</u>	◯ Yes ◯ No						
	* <u>e) Fondaparin</u>	○ No	*f) Aspirin							
	*g) Ticagrelor	○ No	*h) Clopidogrel		ose:					
	* <u>i) Others</u>	ecify: No		1	<ul><li>300mg</li><li>≥ 1200mg</li></ul>					
4. Planned duration of DAPT:		>12 months Not Available	cutaneous entry:	Brachial Femora Ulnar	l Distal radial					
6. Closure device:	No Suture Seal Other,specify:		onary disease % stenosis:	LAD LCx Graft LMX	RCA					
8. Fluoroscopy time:	Not Available 9. Total Dose:			. mGy	Not Available					
10. Contrast Volume :	ml 🔳	Not Available								

a. Patient Name : b. MyKad / Other ID No: c. Date of Procedure: Instructions: 1. For skip lesion - to document as different lesions.If two 3. For long lesion, please document as one single lesion. 4. Please document intervention involves side branch as a second lesion. lesions treated, please fill up two Section 7 forms. **SECTION 7: PCI PROCEDURE DETAILS** (Complete for ALL interventions. Please use one form per lesion treated.) Total no.of lesion treated: NATIVE 2. Dominance: Left Right Co-dominance Dominance: Right **Dominance: Left** 3. Lesion Code to (if applicable) (1-25): 4.i. Graft If Graft treated (lesion codes 18-25), record the grafted native coronary vessel **Target** Target **Target** Vessel Vessel Vessel Graft Graft Graft 20 SVG 1 18 LIMA 23 RAD 1 21 SVG 2 19 RIMA 24 RAD 2 22 SVG 3 25 RAD 3 4.ii. Location in Ostial Body Native Anastomosis graft: (for graft PCI only) 5. Coronary De novo Restenosis (No prior stent) lesion: Stent thrombosis 15. Slow Yes → a Transient Persistent a.Type: Acute Late Flow / No b. Intra coronary medication for no reflow Sub acute Very late No Reflow: In stent restenosis Yes 

N i. Adenosine v.Nicarpidine i.Duration: ii Verapamil Yes N Yes
N Year(s) Month(s) (\*Duration from the known iii. Nitrates vii. Others Not available Yes 

N previous procedure) iv.Adrenalin Yes
N ii. Prior stent type: DES BMS BVS Mg 16. Intracoronary devices used: Others, specify: **IVUS** OCT FFR / IFR / Others Aspiration Penumbra Embolic iii. Clasification: Angiojet catheter Protection Class I (Focal ISR) Rotational Orbital Intravascular Lithotripsy Class II ('Diffuse intrastent' ISR) artherectomy atherect (IVL) Class III ('Diffuse proliferative' ISR) Micro Extension Double lumen micro Class IV (ISR with 'total occlusion') catheter catheter catheter Coil POBA Stent / DCB 6. Lesion type: A B1 B2 C Other, specify: 7. Lesion Ostial CTO > 3mo Calcified lesion \* 17. Stent / DCB details per lesion: description: Thrombus LMS Not Applicable (To fill up ONLY the stent / DCB details) Bifurcation a): SB Treated a. Stent Code Others: b. Diameter(mm) c. Length(mm) (only if SB SB Not Treated = 2.0 mm) b) Medina Classification: a. Stent Code b. Diameter(mm) c. Length(mm) Others: 0 0 1 i) MB prox.: 0 1 ii) MB dist: #2 iii) SB1: 0 1 iv) SB2 1 a. Stent Code Others: b. Diameter(mm) c. Length(mm) 8. Pre PCI % of i.TIMI Flow TIMI-0 TIMI-1 stenosis: a. Stent Code Others: b. Diameter(mm) c. Length(mm) 9. Post PCI % of i.TIMI Flow TIMI-0 TIMI-1 stenosis: b. Diameter(mm) a. Stent Code Others: c. Length(mm) (post): #5 10. Estimated a. Stent Code Others: b. Diameter(mm) c. Length(mm) lesion length: 11. Perforation: ○ No Yes i) Classification 18.Maximum a) Predilatation: b) Postdilatation: Type I (extraluminal crater without extravasation) \* balloon: i. Size: Type II (pericardial or myocardial blushing) Type III (perforation ≥1mm diameter with contrast streaming) ii.Types: ii. Pressure: (atm) Cavity spilling Regular Cutting ■ NC Scoring 12. French Size: Guiding catheter Guiding sheath 19. Other adjunctive procedure: Ventilator Temporary Cardiac Pacing Wire Other, specify: Yes 20. Circulatory support: No iii) Types of guiding catheter PCPS ■ IABP 13. Lesion result: Impella ECMO Unsuccessful Successful Others, specify: 14. Dissection: i. Flow limiting Non flow limiting \*(Post procedure) \* 21. Direct stenting: No

a. Patient Name :	b. MyKad / Other ID No: c. Date of Procedure:						
SECTION 8 : PROCED	URAL COMPLICATION						
1. Outcome:							
*a. Significant Periprocedura	al MI     Yes     No        Not Available	*c. Bail-out CABG	Yes No	)			
Rise in CK/CKMB > x3	3 URL Rise in Troponin > x5 URL	*d. Cardiogenic shock	Yes No	)			
ECG changes		*e. Arrhythmia (VT/VF/Brady)		)			
b. Emergency Reintervention / PCI: Yes No Not Available		* <u>f. TIA / Stroke</u>	Yes	)			
i) Stent thrombosis:	◯ Yes ◯ No	*g. Tamponade		 )			
ii) Dissection:	○ Yes ○ No	*h. Contrast reaction	Yes	)			
iii) Cardiac perforation: iv) Coronary perforation:		* i. New onset / worsened heart failure	○ Yes ○ No				
v) New ischaemia:	O Yes O No	* i. Worsening renal impairment	○ Yes ○ No				
vi) Cardiac tamponade:	Yes      No	(rise of post proecedural creatinine >25% from	_	,			
2. Vascular complications:							
a. Bleeding	Yes No Minimal (Non-CNS bleeding, non-overt b						
	Minor (Non-CNS bleeding, non-overt bleeding with 3-5g/dl						
	Major (Any intracranial bleed or other b	oleeding ≥ 5g/dL Hb drop)					
	Bleeding site Retroperitoneal Percutaneous entry site Others, specify:						
b. RBC / Whole Blood							
Transfusion							
c. Access site occlusion							
d. Loss of distal pulse							
e. Dissection							
f. Pseudoaneurysm	eurysm						
	Ultrasound compression Sur	gery Others, spec	cify:				
g. Perforation							
SECTION 9 : OUTCOM	AF AT DISCHARGE						
1. Outcome:	IL AT DIOCHARGE						
*							
Alive →	a) Date of Discharge (dd/mm/yy):	/					
b	o) Medication: Yes No		Yes N	10			
	Aspirin 🔘 🔘	Statin	0 0	)			
	Clopidogrel	Beta Blocker	0 0	i			
	Ticlodipine	ACE inhibitor	0 0	!			
	Warfarin	ARB	0 0				
	Prasugrel	Other antiplatelet, specify:					
	Ticagrelor	Other, specify:		)			
	NOAC						
○ Death							
	b) Primary cause of  Cardiac  Renal  Others, specify:						
	death:   Infection Neurological						
	○ Vascular ○ Pulmonary						
	c) Location of death:   In Lab  Out of La	ab 					
○ Transferred → *							
to other	a) Date of transfer (dd/mm/yy):  b) Name of hospital:						