HL Kao 23

Impella-Assisted CHIP-PCI

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Disclosures

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Case Example

- ▶ 76y/o man, NIDDM, HLP, ex-smoker
- CKD with eGFR 30ml/min/1.73M²
- CAD s/p CABG 9yrs ago with LIMA-LAD, SVG-OM, RCA not grafted
- Recurrent angina and CHF
- LVEF 25% with LVEDD 67, apical aneurysm, lateral and inferior hypokinesia

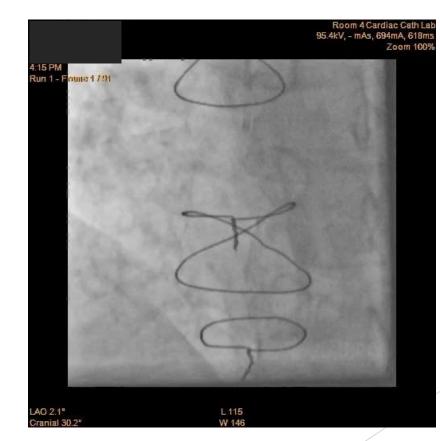
LCA



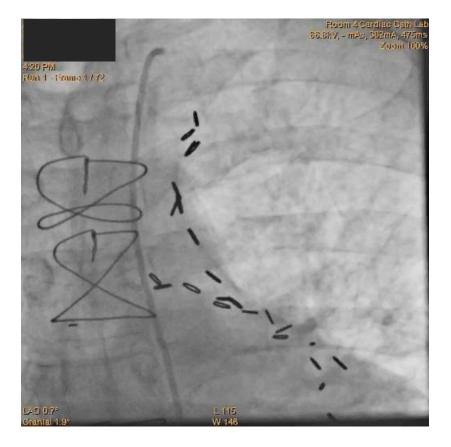


RCA





Baseline LIMA and SVG



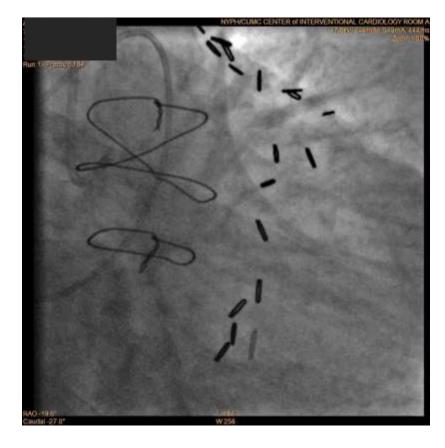


Baseline LVG



- Heart team, surgeon declined
- Medical treatment intensified, partial improvement for 2 months
- But ACS with resting chest pain and troponin elevation occurred

LM and Dx progression

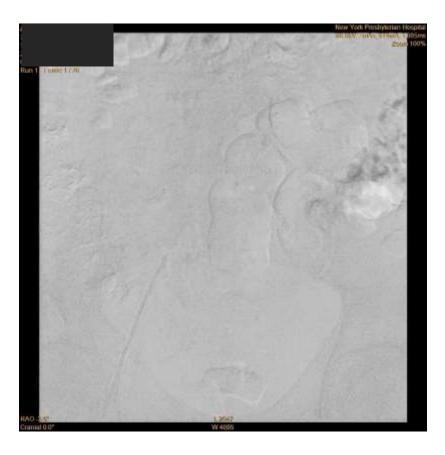


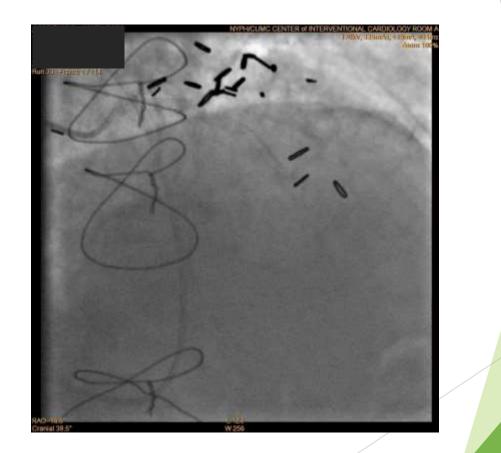


Urgent PCI

- Marked elevation of PAP and wedge pressure with any balloon inflation
- Poor left femoral pulse and no IABP possibility
- Patient had to be sedated for the procedure
- > 2 DES hurried into LM and Dx

Results

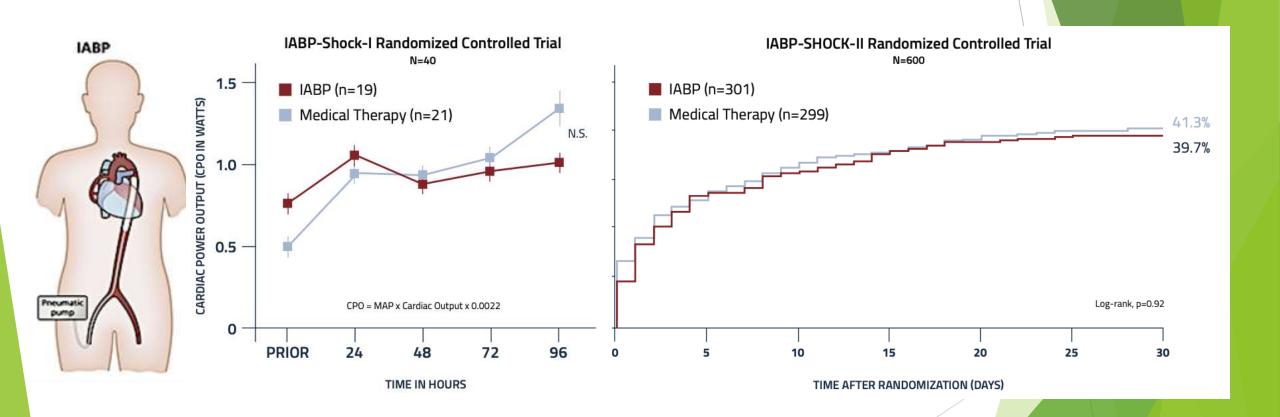




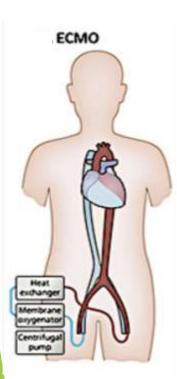
Post-PCI course

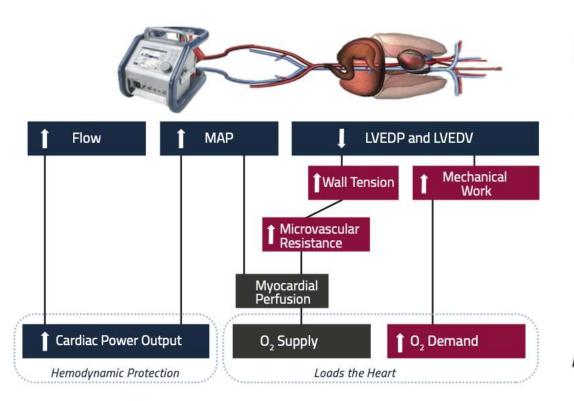
- Still Fc 4 angina and dyspnea, bed-ridden
- Surgeon consulted again, and refused again
- ► Tl201 scan showed inferior viable but ischemic myocardium
- MCS-assisted CHIP-PCI is the only solution, but which?

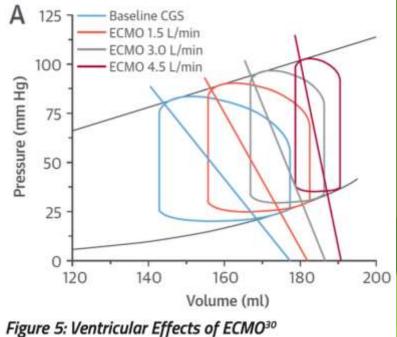
IABP



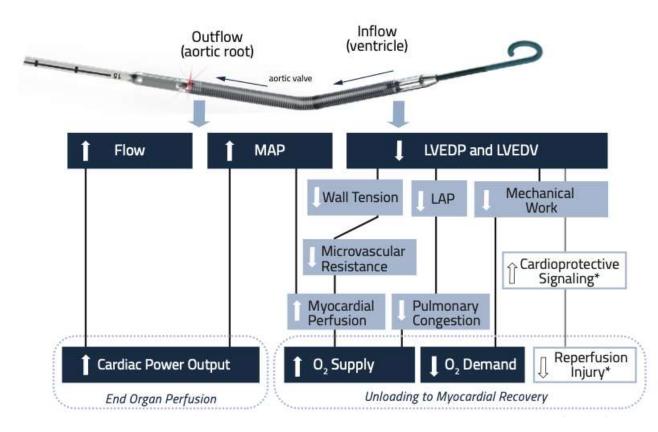
ECMO



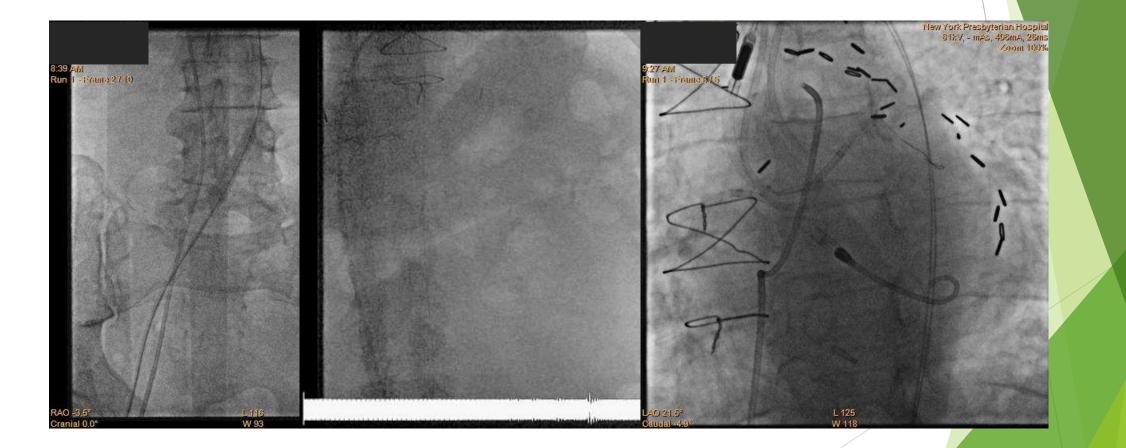




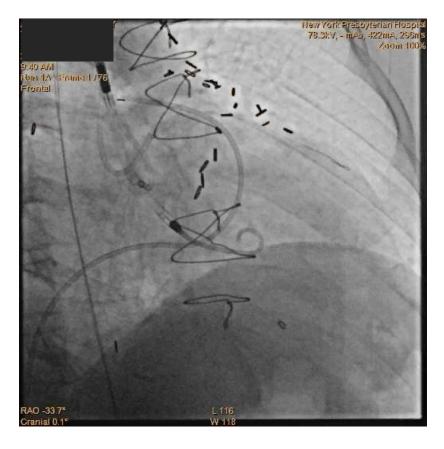
Hemodynamic features of Impella



PTA, Impella and PAC in position

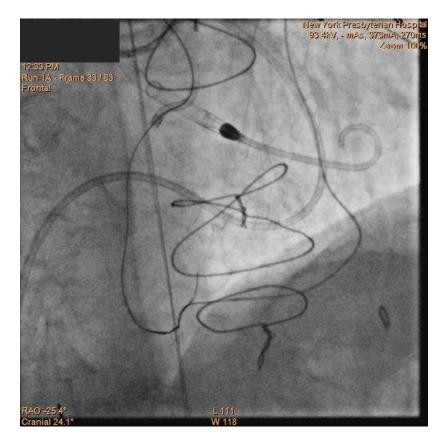


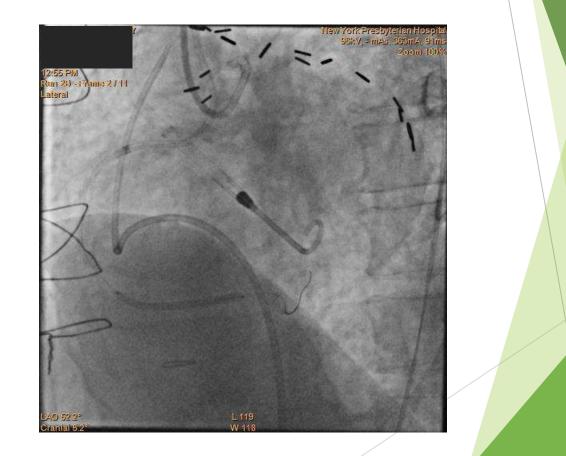
Bi-radial approach



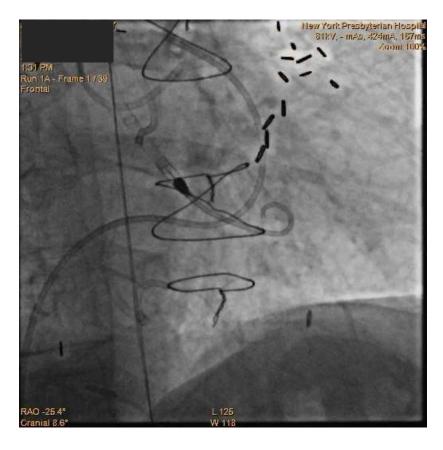


Retrograde successful





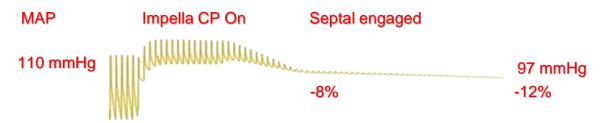
Final angiogram



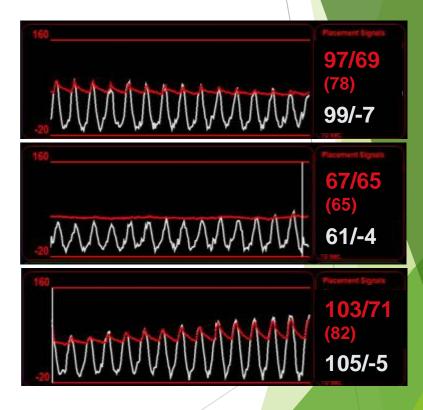


Procedure summary

- Stable hemodynamics throughout the procedure
 - ▶ PAP <30mmHg, LVEDP <10mmHg
 - No inotropic/vasopressor needed
 - Patient conscious and comfortable with only nasal O2



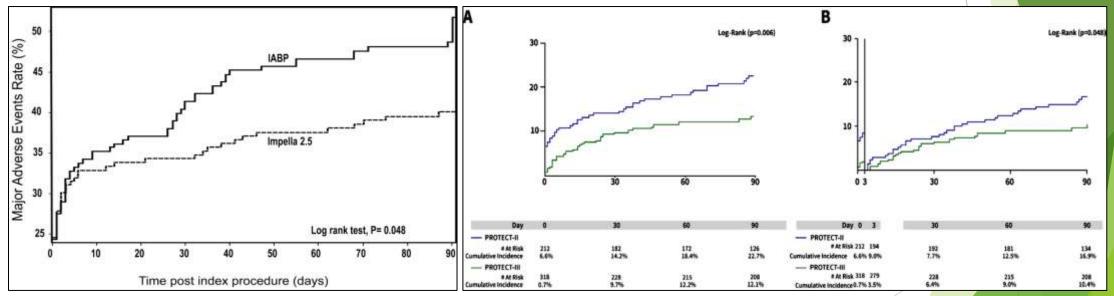
- No access complication and ambulatory the next day
- Discharged on D2 without CIN, remained CCS/NYHA Fc 1



PROTECT II and III results

Lower 90-day MACE with Impella2.5 vs. IABP

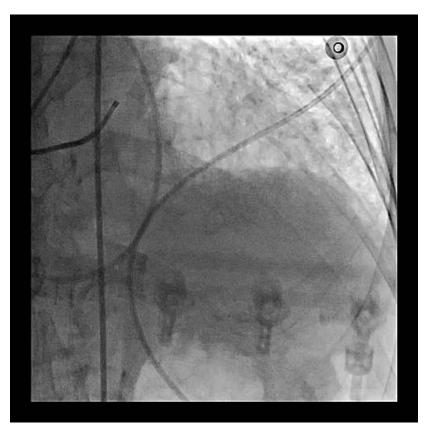
Significant MACE improvement with ImpellaCP vs. 2.5

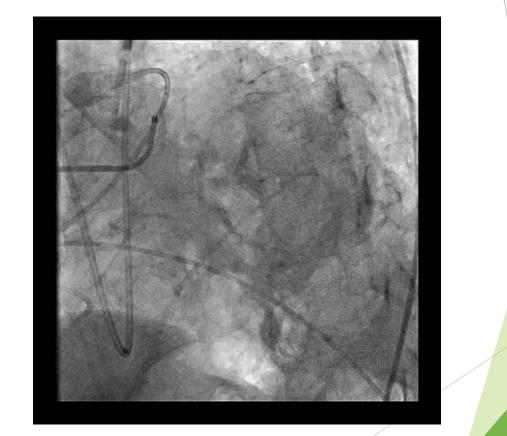


O'Neill WW, et al.Circulation 2012

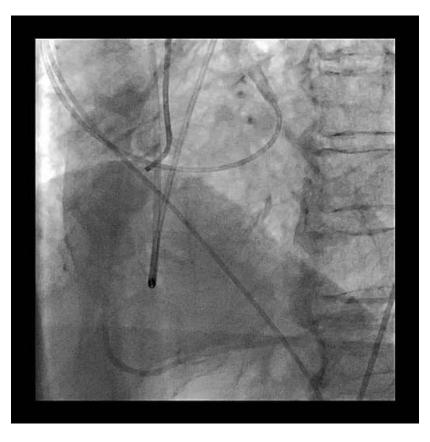
O'Neill WW, et al.AHJ 2022

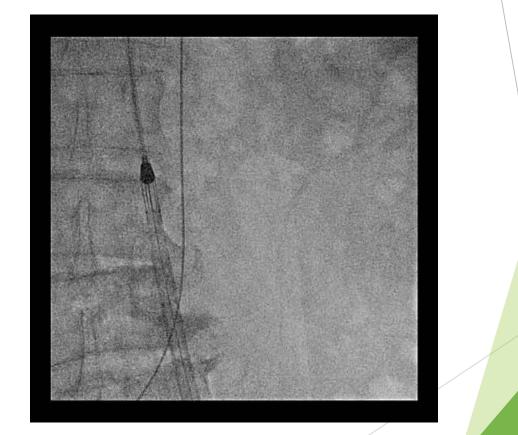
Another case of surgical refusal



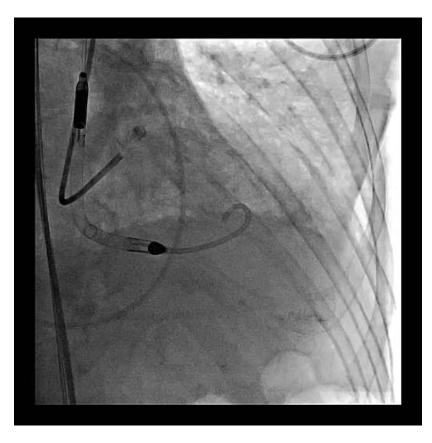


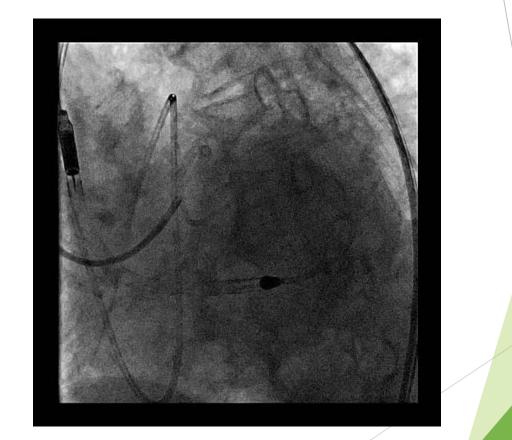
Impella-assisted CHIP-PCI Starting PCWP 35mmHg CO 3.6L/min CPO 0.6W





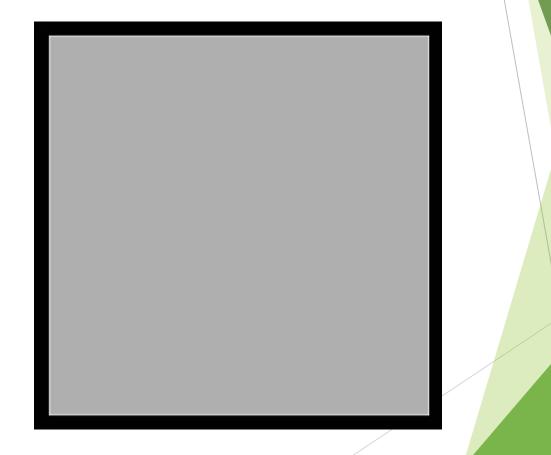
PCWP 12mmHg CO 5.5L/min CPO 1W during MCS LCA final





Patient without any complaint Final RCA and groin angiograms





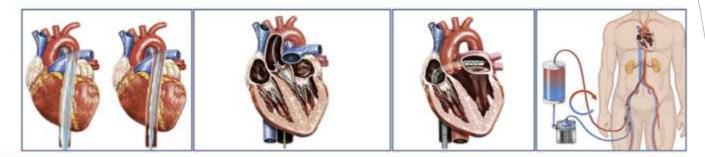
Echocardiography baseline and 1 month

TIS0.8 MI 1.4

75 bpr



Comparison of MCS



	IABP	IMPELLA	TANDEMHEART	VA-ECMO
Cardiac Flow	0.3-0.5 L/ min	1-5L/ min (Impella 2.5, Impella CP, Impella 5)	2.5-5 L/ min	3-7 L-min
Mechanism	Aorta	$LV \rightarrow AO$	$LA \rightarrow AO$	$RA \rightarrow AO$
Maximum implant days	Weeks	7 days	14 days	Weeks
Sheath size	7-8 Fr	13-14 Fr Impella 5.0 - 21 Fr	15-17 Fr Arterial 21 Fr Venous	14-16 Fr Arterial 18-21 Fr Venous
Femoral Artery Size	>4 mm	Impella 2.5 & CP - 5-5.5 mm Impella 5 - 8 mm	8 mm	8 mm
ardiac synchrony or stable rhythm	Yes	No	No	No
Afterload	Ť	(\downarrow)	î	111
MAP	Ŷ	11	↑ ↑	† †
Cardiac Flow	Ť	† †	11	îî.
Cardiac Power	Ŷ	† †	↑ ↑	11
LVEDP	Ť	(1)	11	\leftrightarrow
PCWP	Ļ	<u>++</u>	44	\leftrightarrow
LV Preload	(775)	11	44	Ļ
Coronary Perfusion	Ť	(\uparrow)	777	
Myocardial oxygen demand	Ļ	(++)	⇔↓	\leftrightarrow

Atkinson TM, et al. J Am Coll Cardiol Intv. 2016:9(9):871-83

EAPCI/ACVC expert consensus on percutaneous LVAD 2021

Table 2	Indication for pV	AD-support in HR-PCI ^a	
Device	Indication		Evidence
IABP	Should III used	\triangleright	BCIS-1 ¹⁰
AFP	May be considered	in highly selected patients undergoing HR-PCI in case of	PROTECT II ¹¹ and cohort studies ^{12–15}
	acceptable femo	ral access (>6 mm diameter common femoral artery, no severe tortuosity)	
VA-ECMO	Should III e used		No data available

AFP, microaxial flow pump; HR-PCI, high-risk percutaneous coronary intervention; IABP, intra-aortic balloon pump; VA-ECMO, veno-arterial extracorporeal membrane oxygenation.

^aThere is no common definition of HR-PCI. PCIs might be considered as high risk in patients satisfying the followings clinical and/or anatomical high-risk criteria: clinical characteristics [stable/decompensated LVEF <35%, haemodynamic instability, diabetes mellitus, acute coronary syndromes (ACS), previous cardiac surgery, chronic kidney disease] angiographic characteristics (diffuse CAD, multivessel disease, unprotected left main coronary disease involving bifurcation, severe coronary total occlusion, severely calcified lesions needing rotational atherectomy, last patent conduit).²



REVIEW

Joint EAPCI/ACVC expert consensus document on percutaneous ventricular assist devices

Conclusions

- CHIP-PCI is not just about techniques and devices, but overall planning
- Expect the worst and refresh knowledge on circulatory physiology
- In patients with minimal reserve, preemptive use of MCS is the key to a successful and uncomplicated procedure

Select Your Support Wisely

