Appendix C: Supplementary Tables

Data Tables for Cardiac Arrest in the Catheterization Lab

Table S1: Incidence and Outcome of Cardiac Arrest in the Cardiac Intervention Laboratory

Author/year	Study design/period	Cardiac arrests in Cath lab n/x (%)	Initial rhythms	Outcomes	Comments
Sharma 2019 ¹	Retrospective cohort study (2012–2016)	63/13,112 (0.5%)	PEA 29 (46%); VF/VT 15 (24%); hypotension 19 (30%)	Survived catheterization lab 42 (67%); 1-year Survival 30 (37%)	
Sprung 2006 ²	Retrospective cohort study (1990–2000)	114/51,985 (0.2%)	VF 72 (63.7%); Asystole 30 (26.6%); PEA 11 (9.7%)	Survived procedure: 88/114 (77.2%) Survival to discharge: 64/114 (56.1%)	Long-Term Survival for survivors after cardiac arrest not significantly worse than those who did not have a cardiac arrest: hazard ratio 1.47 (95% CI 0.88-2.46, p = 0.14)
Elkaryoni 2022 ³	Prospective cohort study (2000–2019)	6865 / ?	Asystole 1337 (19.5%), PEA 2951 (43.0%), Pulseless VT 680 (9.9%), VF 1897 (27.6%)	Overall Survival to discharge: 38.1%	GWTG-R

Table S2: Incidence And Outcome From Cardiac Arrest During PCI In The Cardiac InterventionLaboratory Among Patients With And Without Acute ST-Elevation MI

First Autho r. Year	Study Design / period	Purpose/Pr imary Obiective	Population	Denom inator	Number of cardiac arrests	Initial rhythm	Outcome	comments
Addala 2005 ⁴	Observati onal Retrospe ctive	Incidence of VF	Unselected (Elective and non- elective procedures)	19,497	164 (0.84%)	VF	Survival to discharge 164 (100%)	Single centre in Michigan. Cardiogenic shock excluded. All patients successfully defibrillated within 1 min from initiation of VF. VF developed during right coronary injection in 98 patients, left coronary injection in 64 patients, and during bypass graft injection in 2 patients (p< 0.05)
Webb, 2002 ⁵	Observati onal Retrospe ctive 1996– 1999	Incidence of CA	Unselected but with separate reports for stable angina, CS, MI, Cariogenic Shock	4363	Cardiac arrest during PCI 27 (0.6%) All 57 cardiac arrests incidence:1.3% Elective: 0.02% Unstable angina: 0.5%,	All 57 arrests: VF: 36%, VT 28%, bradycardia : 24% asystole: 8% PEA: 2%	24-h survival for all cardiac arrests 37%	Single centre in Vancouver. 57 patients who had CA either during the procedure or later the same day as the procedure. 47% (N=27) had CA during PCI.

					AMI: 15%, Cardiogenic Shock: 10%			Of those who had CA during PCI CA occurred after the initial injection of radiographic contrast into a coronary artery in 12%, after the initial balloon inflation in 60%, and after stent implantation in 13%
Huang 2002 ⁶	Observati onal Prospecti ve	Incidence of VF	Elective Procedures: angina despite adequate medical therapy, or ischemia demonstrated during stress testing; and (2) diameter stenosis of >75%	Overall 905, LCA 561, RCA 344	Overall: 2%, LCA 0.5%, RCA 4.6%	VF	NR	Single centre in Taiwan. VF more frequent in RCA PCI

Table S3: Mechanical Chest Compression CPR In The Cardiac Intervention Laboratory

First Autho r, Year	Study Design / period	Purpose/Primary Objective	Population	Denomi nator	Numbe r of cardiac arrests	Initi al rhyt hm	Surviva l to dischar ge	comments
Mehta 2009 ⁷	Observat ional Retrospe ctive 2004– 2006	To evaluate risk factors for and outcomes of patients with VT/VF	STEMI adults presenting within 12 h of symptom onset	5745	180 (3%)*	VT/V F	83.8%*	296 hospitals in 17 countries, (APEX AMI trial) Excluded - Patients with isolated inferior STEMI, pregnant, known or suspected complement deficiency or active infection, other serious medical problems likely to hamper their recovery, or fibrinolytic therapy for the treatment of their qualifying events Including all patients with VT/VF until the end of catheterisation
Demid ova, 2015 ⁸	Observati onal Retrospe ctive 2007– 2012	To analyse clinical predictors of reperfusion VF	STEMI	3274	71 (1.9%)	VF	81.7%	Single centre Lund, Sweden All patients who had VF during reperfusion
Giglioli , 2006 ⁹	Observati onal Retrospe ctive 2002– 2003	Evaluate incidence, timing and complications from primary PCI in STEMI patients	STEMI within 12 h of onset of symptoms, or within 24 h if signs of persistent ischemia or shock were present.	689	38 (5.5%)	VF	NR	VF was statistically more frequent in patients with inferior AMI than in those with anterior AMI (47 versus 29 patients, P<0.001).

Mienta Observati To examine the incidence, STEMI, >=18 years, 3065 133 VI/V 129/13 Patients enrolled in 4 Primary 2004 ¹⁰ Prospecti incidence, <12 h form symptom (4.3%) F 3 Angioplasty in MI trials in the 2004 ¹⁰ Prospecti predictors, and outcomes of onset onset 99.2% North America/South America/Europe/Middle East and Asia in the 1990's the cardiac catheterization catheterization Iboratory among Excluded - Patients with: Contraindications to	N.A. J. I	01	T	CTENAL AD	2005	422	$\lambda = \lambda $	420/42	Built and a small shire 4 Built
7, Onal Incidence, <12 n form symptom	ivienta	Observati	To examine the	STEIVII, >=18 years,	3065	133	V 1/V	129/13	Patients enrolled in 4 Primary
2004 ⁴⁰ Prospecti predictors, and onset 99.2% North America/South ve outcomes of (1990s) VT/VF occurring in the cardiac catheterization	,	onal	incidence,	<12 h form symptom		(4.3%)	F	3	Angioplasty in MI trials in the
ve outcomes of America/Europe/Middle East (1990s) VT/VF occurring in the cardiac and Asia in the 1990's catheterization Excluded - Patients with: laboratory among Contraindications to	200410	Prospecti	predictors, and	onset				99.2%	North America/South
(1990s) VT/VF occurring in the cardiac catheterization and Asia in the 1990's Isolation Excluded - Patients with: Isolation Contraindications to		ve	outcomes of						America/Europe/Middle East
the cardiac catheterization laboratory among		(1990s)	VT/VF occurring in						and Asia in the 1990's
catheterization Excluded - Patients with:			the cardiac						
laboratory among			catheterization						Excluded - Patients with:
			laboratory among						Contraindications to
natients reperfusion thrombolytic			natients						reperfusion, thrombolytic
undergoing the strength of the			undergoing						therapy for index STEML renal
primary PCI for			nrimary PCI for						failure cardiogenic shock life
stream of the st			CTENAL						ovpostancy <1 year shild
STEIVIL expectation states and the state of									expectaticy <1 year, child-
bearing potential,									bearing potential,
contraindications to aspirin,									contraindications to aspirin,
heparin, or ticlopidine in later									heparin, or ticlopidine in later
PAMI trials.									PAMI trials.
Sustained VT as well as VF									Sustained VT as well as VF
included									included
Patients randomized to the									Patients randomized to the
thrombolytic arm in PAMI-1									thrombolytic arm in PAMI-1
79% of these patients required									79% of these patients required
defibrillation. So at least 21%									defibrillation. So at least 21%
									were not in CA
Were not in ex.									were not in eA.
CPP was percessary in only 8									CPP was necessary in only 8
Patients ((V)									criticate (COV)
			-	0751 AL	2622	74 (20)			patients (8%)
Henriq Observati To compare STEMI 2628 74 (3%) VF NR Single centre in the	Henriq	Observati	To compare	STEMI	2628	74 (3%)	VF	NR	Single centre in the
ues, onal characteristics of Netherlands.	ues,	onal	characteristics of						Netherlands.
2005 ¹¹ Prospecti patients with VF	200511	Prospecti	patients with VF						
ve during PCI versus Patients with out-of-hospital		ve	during PCI versus						Patients with out-of-hospital
1995 – patients with VF resuscitation and VF on arrival		1995 –	patients with VF						resuscitation and VF on arrival
2001 before PCI of the ambulance were		2001	before PCI						of the ambulance were
excluded.									excluded.
Patients with VF during PCI had									Patients with VF during PCI had
more often RCA-related Mi and									more often RCA-related MI and
more frequently TIMI 0 hefore									more frequently TIMI 0 before
									PCI

Table S4: ECPR In The Cardiac Intervention Laboratory

Auth or & Year	Study Design/p eriod	Number of cardiac arrests in Cath Lab	Initial rhythm	Outcomes	Comments				
Mechar	Mechanical piston device (LUCAS)								
Wagn er, 2016 ¹ 2	Prospecti ve observati onal study 2009– 2013	n=32 LUCAS MCC application in cath lab n=10 historical control with manual CPR in cath lab (1999- 2003)	LUCAS MCC group PEA 22/32 (69%) Asystole 5/32 (16%) VF/VT 5/32 (16%) Historical control PEA 4/10 (40%) Asystole 2/10 (20%) VF/VT 2/10 (20%)	LUCAS MCC group Hospital Discharge CPC 1-2: 8/32 (25%) 1 year survival CPC 1–2: 7/32 (22%) Historical control Survival (unspecified) 1/10 (10%)	Prospective vs historical groups not readily comparable due to different treatment periods and other characteristics				

Wagn er, 2010 ¹ 3	Retrospe ctive registry analysis 2004– 2008	n=43 LUCAS MCC	PEA 28/43 (65%) Asystole 9/43 (21%) VF/VT 6/43 (14%)	Survived cath lab 17 patients (39%) (16 with ROSC and 1 with ongoing MCC). Survival to Hospital Discharge: 12/43 (28%) Survival to Hospital Discharge CPC 1-2: 11/43 (26%)	All 43 included patients arrested in the cath lab Use of LUCAS MCC during PCI or pericardiocentesis after cardiac arrest in the cath-lab Procedure success rate: 27/42 (76%) of the PCI procedures done during MCC were successful. Complications: All survivors experienced rib fractures, and one patient suffered a ruptured spleen due to incorrect application of the device.
Larse n, 2007 ¹ 4	Retrospe ctive study - LUCAS	n=6 LUCAS MCC applied in cath lab	n=6 (VF 2, PEA 1, hypotension 3) LUCAS applied in the cath lab	n=3 patients survived the intervention no patients were	4 patients were treated with manual chest compressions and all 4 died. LUCAS was applied on the cardiac cath table for 6 (for VF in 2, for PEA in 1 & for bradycardia or hypotension in 3).
	MCC applicatio n during PCI 2005– 2006	n=6 LUCAS MCC applied after OHCA before cath lab arrival	n=6 (VF 5, asystole 1) resuscitated from OHCA before cath lab arrival	discharged alive.	LUCAS was applied after resuscitation from OHCA & before coronary angiography for 6 : for severe hypotension & bradycardia in 5 & VF in 1. LUCAS was applied prehospital for 1 OHCA. Autopsies were performed in 11 of the cases, revealing sternal and costal fractures in 7/11 patients and liver laceration in 1/11 patient. In 2/11 patients small sub-capsular haematomas in the liver were reported. There was 1 case of excessive intra-thoracic bleeding in the catheterisation laboratory (managed by thoracotomy).
Ventu rini, 2017 ¹ 5	Retrospe ctive registry analysis 2011– 2016	n=43 total cardiac arrests n=20 cath lab arrests (15 LUCAS MCC, 5 Manual CPR). n=11 OHCA (9 LUCAS CPR, 2 Manual CPR). n= 8 ED cardiac arrest (4 LUCAS MCC, 4 Manual CPR) n=3 LUCAS MCC not specified by initial site but eventually received in cath lab n=1 manual CPR not specified by initial site but	LUCAS MCC VF/VT 13/31 (42%) Manual CPR VF/VT 5/12 (42%)	ROSC: LUCAS MCC 22/31 (74%) versus manual CPR 5/12 (42%) 30-day Survival: LUCAS MCC 19% versus manual CPR 8% Survival to Hospital Discharge: LUCAS MCC 13% versus manual CPR 8%	 strap securement. 43 patients required chest compressions for cardiac arrest in the cath lab (12 manual CPR, 31 received LUCAS MCC) All patients had the possibility of transitioning to percutaneous mechanical circulatory support (MCS) (MCS - IABP or Impella), or ECLS during resuscitation. 22/31 patients with LUCAS MCC received MCS: 95% of patients who received MCS achieved ROSC compared to 11% without MCS (p = 0.004). 14/31 (45%) with LUCAS were bridged to ECLS. Patients receiving ECLS were more likely to achieve ROSC (100% vs. 53%, p = 0.003).

		eventually received in cath lab			
Chyrc hel, 2022 ¹ 6	Retrospe ctive cohort study 2013– 2020	n= 48 total cardiac arrests received LUCAS MCC 23/48 (48%) had cardiac arrest in the cath lab	PEA 22/48 (46%) Asystole 8/48 (17%) VF/VT 13/48 (27%) Unknown 5/48 (10%)	ROSC was achieved in 31% of patients (15/48). Survival to hospital discharge = 17% (8/48).	Small single hospital group of patients who arrested either before or during angiography. 30 patients who could have been analysed were excluded due to lack of data and clinical history in the notes. In patients with hyperkalemia, survival rate was 50%, while survival rate for those with potassium < 5.0 mmol/L was only 4% (p = 0.0007).
Mads en Hardi g, 2019 ¹ 7	Retrospe ctive observati onal study 2004– 2013	n=35 total cardiac arrests received LUCAS MCC n=27/35 (77%) in cath lab n=8/35 (23%) taken to cath lab with ongoing CPR.	PEA 17/35 (49%) Asystole 4/35 (11%) Bradycardia 7/35 (20%) VF/VT 7/35 (20%)	ROSC 18/35 (51%) ROSC 14/27 (53%) for arrest in cath lab Survival CPC 1–2 9/35 (26%) Survival CPC 1-2 9/27 (33%) among cath lab arrests ROSC and survival did not differ across presenting rhythms of VF/VT, PEA, asystole or bradycardia. No survivors among those with ongoing CPR on arrival in cath lab	No patient survived who arrived at the cath-lab still requiring CPR (potential candidate for ECPR) The median time of MCC in the cath-lab for those who did survive was 10min versus 45min for those patients not surviving. Initial arrest rhythm did not predict outcome. If a diastolic arterial BP of 30 mmHg can't be achieved, consider escalation to ECPR. This decision should be made within the first 10–20 min of resuscitation efforts in the cath-lab, as longer periods are associated with a decrease in survival. LUCAS CPR time was shorter for those who gained ROSC and survived. Those that arrived at the cath-lab with ongoing CPR had a lower chance of obtaining ROSC than if the arrest occurred in the cath lab (22% vs 53%,p = 0.086). None of the patients survived if resuscitation was ongoing when they were admitted to the cath-lab.
Load-dis	l stributing bar	d device (Autopuls	e)		
Spiro 2015 ¹ ⁸	Retrospe ctive observati onal study in- hospital cardiac arrest 2011– 2013	n=25 received Autopulse-CPR during in- hospital cardiac arrest 15/25 cardiac arrest during invasive procedures (14/15 in cath lab)	14 patients had Autopulse -CPR started in the cath lab due to: VF/T 7/14 (50%) PEA 7/14 (50%)	ROSC 12/25 (48%) with Autopulse-CPR 7/25 (28%) survived to hospital discharge 3/9 (33%) patients who received Autopulse CPR with simultaneous PCI survived to hospital discharge with normal cerebral function (CPC 1- 2)	 15/25 (60%) of patients received Autopulse-CPR at some stage during an invasive procedure (14/15 in cath lab). In 9/15 (60%) Autopulse -CPR received simultaneous with invasive procedure (4/9 with PCI, 4/9 with angiography+TEE+temp pacer, and 1/9 pericardial drainage) & in 6/15 there was a pause in the invasive procedure whilst the A-Pulse was attached. Complications: Battery depletion (1), difficult backboard placement (1) compression band twist (1), clip detachment (1)

Table S5: Mechanical Circulatory Support In The Cardiac Intervention Laboratory

Author	Study	Number of	Initial rhythm	Outcomes	Comments
/ year	Design	arrests			
Shawl, 1990 ¹⁹	Case series (time frame not reported)	7	VF or asystole	4 out of 7 patients survived (57%). All survivors NYHA class 1 at 6 months.	Percutaneous cardiopulmonary bypass was instituted for cardiac arrest in cath lab refractory to ACLS (mean cannulation time from cardiac arrest: 21 min). Subsequent interventions included coronary bypass surgery (n=3) and coronary angioplasty (n=2)
Moone y, 1991 ²⁰	Case series (1988 - 1989)	11 (5 arrests in cath lab)	N/A	5/5 (100%) survival (cath lab) 2/6 (33%) survival (outside cath lab)	Percutaneous cardiopulmonary bypass was instituted in 5 patients with cardiac arrest during percutaneous coronary procedures, and 6 patients with cardiac arrest outside the cath lab. Subsequent interventions included coronary bypass surgery (5/5 in the cath lab group, 2/6 in the non-cath lab group.
Gramb ow, 1994 ²¹	Retrospe ctive observati onal study (1988 - 1992)	7 cardiac arrests 23 cardiogenic shock	N/A	0/7 (0%) survival in cardiac arrest 6/23 (26%) survival in cardiogenic shock	Percutaneous cardiopulmonary bypass was initiated after cardiac arrest (mean time: 21 min) or shock during diagnostic or therapeutic cardiac procedures in the cath lab (mean cannulation time 17 min) using portable Bard PCB system - via FA and FV access Subsequent interventions included emergent cardiac surgery (n=14), coronary angioplasty (n=13) and medical therapy (n=3).
Nagao, 1999 ²²	Prospecti ve observati onal study (1994– 1997)	32 with refractory VF and STEMI 19 OHCA 13 IHCA (ER)	Refractory VF	ROSC: 28/32 (87.5%) Weaned from ECMO 6 (18.8%) Good Neurological outcome 3/32 (9.4%)	Percutaneous coronary bypass was instituted in 32/32 patients with refractory VF complicating STEMI who were transferred to the cath lab of the Emergency Room. Intravenous rTPA was administered to those patients with MI diagnosed before VF. Subsequent interventions included coronary angiography/ angioplasty if adequate reperfusion had not been achieved.
Goslar, 2016 ²³	Retrospe ctive observati onal study (2010- 2015)	12 cardiac arrests in cath lab 11 cardiogenic shock	N/A	Weaned off ECMO 6/12 (50%) Survived to hospital discharge 2 (17%)	VA-ECMO was instituted in patients with refractory cardiac arrest (n=12) or cardiogenic shock (n=11) in the cath lab, and in 33 patients outside the cath lab (ICU, n=8; operating room, n=25). Subsequent interventions included coronary angiography/angioplasty, pulmonary angiography or CT followed by thrombolysis if pulmonary embolism. N=9/23 patients (39%) had concomitant IABP
Parr, 2020 ²⁴	Retrospe ctive observati onal study (2010– 2018)	39 cardiac arrests in the cath lab 23 Cardiogenic shock	VF/VT 19/39 (48.7%) PEA 20/39 (51.3%)	30-day Survival: 17/39 (44%) cardiac arrest; 12/23 (52%) cardiogenic shock 1-year Survival: 16/39 (44%) cardiac arrest; 11/23 (48%) cardiogenic shock	VA-ECMO was instituted in patients with cardiac arrest or cardiogenic shock during percutaneous procedures in the cath lab (median cannulation time from collapse: 38 min). Complications included stroke (32.3%), hemorrhage at the cannula site (33.9%), extremity malperfusion (19.4%), and new acute kidney insufficiency requiring renal replacement therapy (38.7%). Subsequent interventions included IABP (n=13 cardiac arrest, n=3 shock), ventricular assist device (n=2 cardiac arrest, n=4 shock) and coronary bypass surgery (n=7 cardiac arrest, n=5 shock).

Hrynie wicz, 2021 ²⁵	Retrospe ctive observati onal study (2012 – 2017)	8 cardiac arrests in the cath lab 11 in other in-hospital locations 7 OHCA	Cath lab: VF/VT 6 (75%); PEA/asystole 2 (25%) IHCA: VF/VT 7 (64%); PEA/asystole 4 (36%) OHCA: VF/VT 4 (57%); PEA/asystole 3 (43%)	Cath lab: 7/8 (88%) survival to discharge and at 6 months; 88% with CPC 1-2 Other IHCA: 6/11 (55%) survival to discharge and at 6 months; 45% with CPC 1-2 OHCA: 5/7 (71%) survival to discharge and at 6 months 71% with CPC 1-2	VA-ECMO was instituted in 8 patients with cardiac arrest in the cath lab, 11 patients in other in-hospital locations and 7 patients with OHCA who were transferred to the cath lab (median cannulation time: 39 min cath lab, 45 min IHCA, 72 min OHCA). Subsequent interventions for patients with cardiac arrest in the cath lab included revascularization (n=7/8 patients), hypothermia (n=2/8 patients). Outcomes were better in patients with initial rhythm VF/VT versus PEA/asystole.
Radsel, 2021 ²⁶	Prospecti ve observati onal study (2010– 2020)	52 cardiac arrests (n=36, 69.2% were cannulated in cath lab) 78 cardiogenic shocks	N/A	Cardiac arrest survival to discharge (CPC 1 or 2) 15/52 (29%).	 VA-ECMO was instituted in n=23 patients before percutaneous or surgical interventions and n=17 inmediately after these. Settings of cardiac arrests included OHCA or IHCA (Emergency Room, cardiac ICU, the ward or the cath lab). N=36 patients (69.2%) were cannulated in the cath lab. Interventions before VA-ECMO E-CPR included PCI (15/52, 28.8%), TAVI (1, 1.9%) and electrophysiology procedure (1, 1.9%). Subsequent interventions on VA-ECMO included coronary angiography (38, 73.1%), PCI (n=24, 46.2%)), CABG (4, 7.7%), aortic surgery (3, 5.8%), pericardiotomy, left ventricular assist device, pulmonary embolectomy, abdominal surgery (1 each, 1.9%).
Mazzeff i, 2024 ²⁷	Retrospe ctive cohort study (2020– 2023	Total 2515 cardiac arrests 602 cardiac arrests in cath lab	Cath lab arrests: VT 68 (11.3% VF 167 (27.7%) PEA 222 (36.9%) Asystole 57 (9.5%) Unknown 88 (14.6%)	Survival to discharge 235 (39%)	Extracorporeal Life Support Organization (ELSO) registry data. Objective: to explore whether ECPR mortality differs by IHCA location and whether moving patients for cannulation impacts outcome. Pre-ECPR interventions for cardiac arrest in cath lab group: IABP 51 (8.5%) RV assist device 2 (0.3%) Impella 65 (10.8%) Conventional CPR time 25 (14–39) minutes. Adjusted odds ratio (aOR) for mortality higher in patients with cardiac arrest in the ICU (aOR, 1.85; 95% Cl, 1.45– 2.38; p <0.001) and in patients with cardiac arrest in acute care bed (aOR, 1.68; 95% Cl, 1.09–2.58; p = 0.02) compared with in the cath lab. Survival to discharge for cardiac arrests in other locations (and ECPR): ICU = 243/939 (25.9%) Acute care bed = 67/242 (27.7%)

Table S6: Intracoronary Epinephrine In The Cardiac Intervention Laboratory

Author/Y ear	Study design/ period	Cardiac arrests in Cath lab n/x (%)	Initial rhythms	Outcome	Comments
Bagai Y, 2011 ²⁸	Case series 2006–2009	8/8 (100%)	N/R	7/8 (87.5%) successful support in cardiac arrest or cardiogenic shock; 3/7 (43%) survival to hospital discharge for in-lab cardiac arrest patients treated with Multifunctional Percutaneous Heart (MPH)	
Loehn, 2020 ²⁹	Retrospecti ve observation al study (Impella) 2014–2016	43/73 (59%)	Asystole 9/43 (21%), PEA 11/43, VF/VT 23/43 (54%)	Impella implantation during ongoing CPR versus implantation after ROSC had no significant impact on survival to discharge (28.5% vs. 27.2%, p=0.92). Among whole group (those with cardiac arrest and those with cardiogenic shock without cardiac arrest) the overall survival rate at discharge was low in Impella recipients (35.6%) but better when Impella placed pre-PCI (50%) than post PCI (23.1%) p=0.027. In whole group (regardless of when impella was placed), impella was the sole independent predictor of survival at discharge and at 30, 90 & 180 days.	
Vase, 2017 ³⁰	Observatio nal study (Impella) 2014–2016	8	VF 5 (62%), PEA 3 (37%).	For those with rCA, survival to discharge was 4/8 (50%) compared with 7/12 (58%) for those with cardiogenic shock. All patients survived 6 months and were CPC 1–2,	8 patients with refractory CA & 12 with cardiogenic shock. At the time of receiving the Impella device all cardiac arrests were in PEA.
Gerfer, 2023 ³¹	Retrospecti ve cohort study 2014–2016	59/729 (8%)	37% of CPR patients underwent defibrillation but no further details given about the arrest rhythms	49/59 (83%) survived to hospital discharge	 16/59 (27%) cardiac arrest patients required 'heart-lung circulatory support' but their outcomes are not reported separately. Patients who required CPR had a lower ejection fraction and lower aortic gradients than those who did not. Patients with intra-procedural complications such as tamponade and valve displacement had higher incidence of CPR. All complications were higher in the group needing CPR.
Almajed, 2023 ³²	Retrospecti ve cohort study (Impella) 2013–2022	6 (5 during TAVR; 1 during BAV)	Not stated	30-day survival 5/6 (83%) 4/5 for TAVR and 1/1 for BAV	2680 procedures: 1965 TAVR and 715 BAV. 120 used Impella support, 26 TAVR and 94 BAV, but only 5 and 1 for cardiac arrest. Cardiogenic shock TAVR Impella cases mortality 35.7 % and cardiogenic shock BAV 44.2%

Orvin,	Observatio	41/87	Not stated	Overall survival to hospital	For all 87 cases (75.9% VA ECMO, 19.5%
2021 ³³	nal study	(47%)		discharge after TAVI with pMCS	Impella CP, 4.6% TandemHeart).
	(Impella,			insertion was 72.5%.	
	VA ECMO,				No separate data for the 41 cardiac arrest
	TandemHea			1 yr survival was close to 50%.	cases.
	rt)				
	2011-2020				

Resuscitation of Patients with Durable Mechanical Circulatory Support with Acutely Altered Perfusion or Cardiac Arrest: supplementary data tables

Study	Publication	Study Type	Continent	Total	Population	Mechanical	Chest
•	Year			number of		Support	Compression
				patients		Device(s)	s Described
				with			
				acutely			
				altered			
				perfusion			
Senman et al ³⁴	2024	Case Series	North	58	Both In-	LVAD	Yes
			America		and-Out of		
					nospital		
	Case series o	of 58 LVAD suppo	rted patients a	t a single institu	tion who suffer	ed cardiac arres	st. Of these, 24
	received ches	t compressions a	and 34 received	d no chest comp	pressions. Per i	eview of the no	tes, the most
	common reas	on for withholding	g of chest com	pressions was a	a perceived con	itraindication to	chest
	compressions	s in LVAD support	ted patients. If	nere were no do	cumented case	es of device disi	odgement.
	Survival was s	similar between ti	nose who rocoiv	and did not rece	ive chest comp	ressions, but ne	eurologic
	outcomes we			ed chest compre			
Sande et al ³⁵ l	2023	Case Report	North	1	In-hospital	LVAD	No
			America				
	Case report of	f a patient experi	encing device	alarms after und	dergoing an abl	ation procedure	shortly after a
	percutaneous	LVAD placement	t. A bedside ec	ho showed a la	rge circumferer	ntial pericardial	effusion with
	right ventricula	ar collapse and ta	amponade. The	e patient underv	vent bedside pe	ericardiocentesi	s with improved
	physiology.						
Victor et al ³⁶	2022	Case Report	North	1	In-hospital	LVAD	No
			America				
	Case report of	f a patient experi	encing increas	ing dyspnea an	d hemodynami	c instability 6 da	vs after I VAD
	placement. LV	AD flow rate adi	ustments and v	asopressor utili	zation were un	successful. and	ultrasound
	identified a pe	ricardial effusion	. Successful o	perative manage	ement was per	formed.	
				Ũ			
Akin et al ³⁷	2022	Case Report	Europe	1	In-hospital		No
Annota	2022		Europe		in-nospital	LVAD	110
	A case report	of a patient 10 da	ays after LVAD	placement invo	lved a researcl	n study for a sul	olingual
	microcirculato	ory imaging tool fo	or microvascula	ar circulation an	d perfusion. Th	e device reveal	ed severe failure
	of the microcil	rculation, and the	e patient later d	eveloped hemo	dynamic compi	romise and sign	S OT
	nypoperiusion	1. Cardiac tampoi	nade was iden	lined that was s	ubsequently su	rgically correcte	ed.
Ratman et al ³⁸	2022	Case Report	Europe	1	Out-of-	LVAD	No
					Hospital		
	Case report o	f a patient admitt	ed with low LV	AD flows and m	ultiple organ fa	ilure. Pump flow	s and evidence
	of organ injury	y improved with fl	uids.				
Doita et al ³⁹	2022	Case Report	Asia	1	In-hospital	LVAD	Yes

Table S7: Details of included studies

	Case report of an LVAD thrombosis leading to left outflow obstruction. The clot was large enough to occupy the LVAD inflow and resulted in nearly no forward flow from the device. The patient suffered cardiac arrest. Chest compressions were administered but the patient could not be resuscitated.								
Barssoum et a ⁴⁰	2022	Retrospective observational cohort	North America	578	In-hospital	LVAD	No		
	Retrospective arrest compar- patients, 226 compressions by potential m were limited.	e analysis of the N ring outcomes of (39.1%) survived s vs. 61% for thos hisclassification as	lational Inpatie those who unc to hospital dis se who did not s only adminis	ent Sample inclu lerwent chest c charge. Mortali achieve chest c trative data was	uding LVAD pati ompressions wi ty was 74% for compressions (p s used and varia	ents who sustai th those who di those receiving o<0.01). This stu ables available f	ined cardiac d not. Of 578 chest Jdy was limited for abstraction		
Pokrajac et al ⁴¹	2022	Retrospective observational cohort	North America	1	In-hospital; Pediatrics Included	LVAD	No		
	Single center, no deaths or shock and org	retrospective rev cardiac arrests in gan failure.	view of 54 eme the ED. 4 pati	rgency departn ents in the coho	nent visits in peo ort died, with on	diatric VAD pati e experiencing	ents. There were cardiogenic		
Esangbedo et al ⁴²	2022	Case Series	North America	4	In-hospital; Pediatrics Included	LVAD; BiVAD	Yes		
	Case series of cardiac arrest brief chest co exploration. P outcomes.	of 5 pediatric patie due to tamponac mpressions. Patie atient 4 had brief	ents who under de and suffered ent 3 had card chest compre	went chest con d severe neurol iac tamponade ssions with tam	npressions with logic injury. Pation and brief chest aponade. Of the	VAD in place. F ent 2 had RVAD compressions p 4 patients, 3 su	Patient 1 had had disconnect and prior to chest urvived with good		
Oates et al ⁴³	2022	Case Report	North America	1	In-hospital	LVAD	No		
	Case report o	f patient who det	eriorated after	attempt at VT a	ablation with hyp	ooxemia from in	tratrial shunt.		
Ziegler et al ⁴⁴	2021	Case Report	North America	1	Out-of- Hospital	LVAD	No		
	Case report o	f an emergency r	epair using sp	licing of a trans	ected driveline i	n a left ventricu	lar assist device.		
lwashita et al ⁴⁵	2020	Case Report	Asia	1	Both In- and-Out of hospital	LVAD; ECMO	Yes		
	Case report of chest compre discovered ar to defibrillatio discontinued.	f cardiac arrest 2 ssions were perfo nd changed after n and VA-ECMO	years post LV. ormed for 40 m 50minutes of t was initiated.	AD placement. ninutes. On arri otal chest comp A complicated o	The device was ival to the hospi pressions. Subs course led to pa	a unknown by re tal a depleted b equent VT was tient death afte	esponders and attery was not responsive r ECMO was		
Eyituoyo et al ⁴⁶	2020	Case Report	North America	1	Both In- and-Out of hospital	LVAD	No		
	A case report hypotension. the emergenc multiorgan fai	of a patient with Upon EMS arriva by department, VF lure and later exp	an LVAD place I, an irregular i ⁻ was noted an bired.	ed 7 years earlie hythm was note id corrected wit	er who develope ed and presume h defibrillation. ⁻	ed altered menta ed to be artifact The patient dev	ation and from LVAD. In eloped		

Saito et al ⁴⁷	2019	Case Report	Asia	1	Out-of- Hospital	LVAD	Yes
	Case report o compressions exchanging e	of a patient who su were performed xternal cables. Th	uffered from glu by paramedic ne patient reco	obal cerebral iso s and LVAD fun vered without a	chemia due to l ction was resto ny neurological	VAD pump stop red after hospita deficit.	opage. Chest al arrival by
Harper et al ⁴⁸	2019	Case Report	North America	1	In-hospital	LVAD	No
	Case report or refractory VT medications a	of a patient with and experiencing and external shoc	n LVAD placed j chest pain, di ks and LVAD fl	3 years earlier zziness and mu low rate was de	presenting to the transformation of the senting to the senting of	ne emergency d s of his ICD. R w better ventricu	epartment in eceived ular filling.
Thiele et al ⁴⁹	2018	Case Report	Europe	1	Out-of- Hospital	LVAD	Unclear
	Case report o	of LVAD driveline of	disconnect. Pa	tient recovered	with re-connec	ting driveline.	
Ornato et al ⁵⁰	2018	Case Report	North America	1	Out-of- Hospital	LVAD	Yes
	Case report o mmHg with co	of a patient with an onfirmation of tub	n LVAD who su e placement. (uffered cardiac a Compressions s	arrest. Patient w tarted and ETC	vas intubated ar O2 rose to 28 n	nd ETCO2 was 0 nmHg.
Godishala et al ⁵¹	2017	Case Series	North America	4	In-hospital	LVAD	No
	Patient 1 rece patient was a complete thro complications diaphoresis s remained syn aortic valve; t	symptomatic. Pat imbotic occlusion and died followir hortness of breat nptom-free. Patien his was removed	in ICD due to V ient 2 also rec of left circumfl ng intracranial h and presynce nt 4 experience but one month	I, attributed to e eived shocks fro ex artery; this w hemorrhage. Pa ope due to coro ed chest pain an h later the throm	Sectrolyte derar om ICD due to ' vas removed bu atient 3 experier nary artery occl nd shortness of bus returned a	ngement; once of VT which was a it the patient suf nced chest pres lusion; once ste breath due to la nd the patient d	corrected the ttributed to ffered sure, nted he arge thrombus in ied.
Yuzefpolskaya et al ⁵²	2016	Case Report	North America	1	In-hospital	LVAD; ECMO	Yes
	This paper pr patients. A ca operative day initially perfor occurred. Che on VA-ECMO	esents an algorith ase study is prese 7 8 after LVAD imp med as the patier est compressions 9. After transfer to	nm for assessin ented by way o plantation deve nt was recently were ultimatel the ICU, the p	nent and manag f rationale for the loped acute alter post-operative ly initiate 15 mir atient was prone	gement of hosp ne algorithm in v ered perfusion. and it was uncl nutes into the ev ounced brain do	italized unrespo which a patient Chest compres lear whether ca vent and the pate ead.	nsive LVAD who was post- sions were not rdiac arrest had tient was placed
Bouchez et al ⁵³	2016	Case Report	Europe	2	Both in and out of hospital	LVAD	No
	Two case rep authors descr electrical stor	orts of patients w ibe a "treatment p m, and defibrillation	ith LVADs who protocol" that in on.	went into VF a ncludes augmei	nd developed d nting MAP, addi	eteriorating RV ressing wall ten	function. The sion, treating
Plymen et al ⁵⁴	2015	Case Report	Europe	1	In-hospital	LVAD; RVAD	Yes
	Case report o treated with a	of a patient with L temporary RVAD	/AD who deve and ultimately	loped RV failure y underwent hea	e and arrhythmi art transplant.	a after embolisr	n. Patient was
Mulukutla et al ⁵⁵	2015	Case Report	North America	1	Out-of- Hospital	BiVAD	No
	Case report o	of patient with BiV	AD who develo	oped sustained,	unstable VT w	ho underwent V	T ablation.

Wilson et al ⁵⁶	2014	Case Report	Canada	1	In-hospital	LVAD	No				
	Case report of a single patient with recurrent, brief cardiac arrest and loss of consciousness iso LVAD and fused aortic valve. Underwent aortic valve replacement with improvement.										
Shinar et al ⁵⁷	2014	Case Series	North America	8	Both In- and-Out of hospital	LVAD	Yes				
	Case series of dislodgement chest compre stable. Three patient who u circulation and	of 8 patients who l Eight patient rec ssions. In all case patients underw nderwent 2.5 hou d 4 patients (50%	had LVADs and cords were rev es with return of ent autopsy, w urs of chest con b) survived with	d underwent cho iewed revealing of effective circu ith no device dis mpressions. 6 o n good neurolog	est compressio no apparent d llation, post-arr slodgement fou f 8 (75%) patien ic outcomes.	ns with a focus islodgement afte est pump flows nd—including a nts had return o	on cannula er receiving were reported as in autopsy for a f effective				
Cubillo et al ⁵⁸	2014	Case Report	North America	1	Out-of- Hospital	LVAD	Yes				
	Case report o arrest. Chest taken to the e substantial ne	f emergency repa compressions we mergency depart eurologic injury.	air of an LVAD are initiated by ament where LV	driveline that wa a bystander and /AD flows were	as accidentally d then continue restored, howe	transected resund the by paramedic over patient had	Iting in cardiac s. Patient was sustained				
Garg et al ⁵⁹	2014	Case Series	North America	16	In-hospital	LVAD	No				
	Case series of (56.3%) receit of 9 patients (compression compressions	of 16 patients with ved chest compre 44.4%) who rece initiation. As com a was substantiall	a continuous-fle essions and 2 ived chest con pared to a non y longer.	ow LVADs who s (22.2%) of those npressions had -LVAD cardiac a	suffered in-hos e who received delays of at lea arrest cohort, tii	bital cardiac arre chest compress ist 2 minutes be me to initiation o	est. 9 patients sions survived. 4 fore chest of chest				
Haglund et al ⁶⁰	2014	Case Report	North America	1	In-hospital	LVAD	No				
	Case report o power source cyanotic. LVA compressions	f a patient post-o disconnection fro D power was res s were not provide	perative day 7 om his LVAD le tored with impl ed.	from LVAD imp eading to cardiae roved perfusion,	lantation with a c arrest. He wa , though low flo	cute hyperactiv s found unrespo w alarm continu	e delirium with onsive and led. Chest				
Duff et al ⁶¹	2013	Case Report	North America	2	In-hospital; Pediatrics Included	LVAD; BiVAD	No				
	Case report o LVAD failure a hypercarbia. I	f cardiac arrest ir and circulatory an Patient 2 involvec	n two pediatric rest resulting fi l episodic hypo	patients with ve rom acute pulm operfusion.	entricular assist onary hyperten	devices. Patien sion triggered b	t 1 involved y post-anesthetic				
Schweiger et al ⁶²	2012	Case Report	Europe	1	Out-of- Hospital	LVAD	Yes				
	Case report o response. Par patient's wife	f 2 patients with I ramedics unsure declined.	_VADs—one o of whether to o	f which suffered do CPR and wife	l acutely altered e called VAD sp	l perfusion resu becialist. CPR a	lting in EMS dvised but				
Brenyo et al ⁶³	2011	Case Report	North America	1	Out-of- Hospital	LVAD	No				
	Case report o defibrillated. H	f patient with LVA le was comatose	D who suffere and treated w	d cardiac arrest ith therapeutic l	t with ventricula hypothermia. At	r fibrillation and fter rewarming,	was had neurological				

	recovery other than amnesia around the arrest event.									
Rottenberg et al ⁶⁴	2011	Case Report	North America	1	In-hospital	LVAD; ECMO	Yes			
	Case report o chest compre	Case report of patient sustaining cardiac arrest during redo sternotomy for LVAD exchange. Abdominal chest compressions were performed to avoid damage to inflow cannula.								
Andersen et al ⁶⁵	2009	Case Series	Europe	3	Out-of- Hospital	LVAD	Yes			
	Case series of 23 patients with HeartMate II LVADs describing the incidence of VT/VF during 266 total months of follow up. They noted an incidence of 52%, with external defibrillator or ICD shock in 8 patients and significant hemodynamic instability in 3 patients.									

Table S8: Studies Including Patients who Received Chest Compressions

	Number of		Duration of				
	Patients		Implantation				Documentati
	Receiving		prior to			Duration of	on of MCS
	Chest		Chest			Chest	Dislodgemen
	Compression		Compressio	Cause of		Compressio	t or other
Study	s	Device	ns	arrest	Outcome	ns	Complication
		HVAD,			Hospital		
		Heartmat			survival,		
		e 2,			survival with		
Senman et		Heartmat	See	See	good neurologic	See	
al ³⁴	24	e 3	reference	reference	outcome	reference	None
Theeuwes et		Heartmat					
al ⁶⁶	1	e 3	1.5 years	Unknown	ROSC obtained	2+ hours	None
			-		Expired in		
					hospital after		
					identification of		
					hypoxemic		
		Heartwar		Thrombosi	ischemic		
Doita et al ³⁹	1	e HVAD	1 year	s	encephalopathy	Not reported	None
Barssoum et			Non-index	See	Hospital	See	
al ⁴⁰	578	Unknown	admission	reference	mortality	reference	None
		Patient 1		Patient 1	Patient 1		
		Heartmat		cardiac	hypoxemic		
		e-3;		tamponad	ischemic		
		Patient		e; Patient	encephalopathy		
		2 Jarvik		2	and death;		
		2015		accidental	Patient 2 good		
		LVAD &		disconnect	neurologic		
		PediMag		ion;	outcome and	Patient 1 15	
		RVAD;		Patient 3	transplantation;	minutes;	
		Patient 3	Patient 1 10	hemorrha	Patient 3 good	Patient 2 4	
		HeartMat	days; Patient	ge;	neurologic	minutes;	
		e 3;	2 6 days;	Patient 4	outcome;	Patient 3 2	
		Patient 4	Patient 3 9	cardiac	Patient 4 good	minutes;	
Esangbedo et		Heartwar	days; Patient	tamponad	neurologic	Patient 4 2	
al ⁴²	4	e HVAD	4 14-days	е	outcome	minutes	None
					Hypoxemic		
					ischemic		
Iwashita et		Heartmat		Battery	encephalopathy		
al ⁴⁵	1	e 2	2 years	depletion	and death	120 minutes	None
		Jarvik			Good		
Saito et al ⁴⁷	1	2000	401 days	Unknown	neurologic	Not reported	None

		LVAD		1	outcome and		
					transplantation		
					Return of		
		Not		Not	spontaneous		
Ornato et al ⁵⁰	1	reported	Not reported	reported	circulation	Not reported	None
					Hypoxemic		
					ischemic		
Yuzefpolskay		Heartmat			encephalopathy		
a et al ⁵²	1	e 2	8 days	Unknown	and death	30 minutes	None
Shinar et al57	8		•	See refere	ence		None
					Hypoxemic		
				Driveline	ischemic		
		HeartWar		transectio	encephalopathy		
Cubillo et al58	1	e LVAD	1.5 years	n	and death	Not reported	None
Garg et al ⁵⁹	9			See refere	ence		None
					Good		
Retherford et		Heartmat		Fractured	neurologic		
al ⁶⁷	1	e 2	3 years	driveline	outcome	30 minutes	None

Mechanical Support for Cardiogenic Shock After Cardiac Arrest: supplementary data tables

Table S9: Evidence Summary for Use of Mechanical Circulatory Support in Post-Cardiac Arrest Patients

Outcomes	Participant	Certainty	Effect	Anticipated absolute	
(importance)	s (studies)	of the	Estimate	e	effect
		evidence	(95% CI)	Risk with	95% CI
		(GRADE)		MCS	
Survival at 30 days /	13 RCTs	low	OR 1.16	37 more	8 fewer to 82
hospital discharge	(n=1842)		(0.97-	per 1000	more
(critical)			1.40)	•	
Cardiac Arrest	6 RCTs ⁶⁸⁻⁷³	low	OR 0.97	8 fewer	78 fewer to
Subgroup	(n=766)		(0.73,1.30)	per 1000	64 more
Survival at 6 or 12	10 RCTs 68-	low	OR 1.18	41 more	13 fewer to
months (critical)	71,73-		(0.95,1.46)	per 1000	94 more
	⁷⁸ (n=1733)				
	10 RCTs ⁶⁹⁻	low	OR 1.21	48 more	34 fewer to
	71,73,75-80		(0.87,1.68)	per 1000	129 more
	(n=757)				
Subgroup microaxial	1 RCT ⁷³	low	OR 1.67		
flow pump			(1.10,2.54)		
Cardiac Arrest	9 RCTs ⁸⁰	low	OR 1.16		
Subgroup (IPMA)			(0.83,1.63)		
Cardiac Arrest			OR 1.56		
Subgroup with STEMI			(1.13,2.16)		
or Resuscitation <10					
minutes					

Survival at longest	14 RCTs ⁶⁸⁻	low	OR 1.17	39 more	7 fewer to 87
available follow-up time	79,81,82		(0.97,1.42)	per 1000	more
	(n=1875)				
Cardiac Arrest	11 RCTs ⁶⁹⁻	low	OR 1.21	41 more	13 fewer to
Subgroup	73,75-79		(0.91,1.60)	per 1000	94 more
	(n=816)				
In-hospital cardiac	1 RC1	low	OR 0.87		
arrest	$^{72}(n=66)$	1	(0.31,2.44)		
	TRU1	IOW	UR 1.67		
	0 PCTe80		(1.10,2.34)		
	9 NO 15-	1000	(0.83.1.63)		
Subgroup with STEM			(0.03, 1.03) OR 1.56		
or Resuscitation <10			$(1\ 13\ 2\ 16)$		
minutes			(1110,2110)		
Favorable Neurological	3 RCTs	low	OR 0.85	37 fewer	109 fewer to
Outcome at Hospital	^{68,76,78} (n=560		(0.60,1.21)	ner 1000	15 more
Discharge / 30 Days)			per 1000	45 11016
(Critical)					
Favorable Neurological	2 RCTs ^{68,78}	low	OR 1.09	21 more	60 fewer to
Outcome at 6 months /	(n=534)		(0.77,1.54)	per 1000	106 more
1 year (Critical)				•	
Favorable Neurological	3 RCTs	low	OR 1.11	25 more	54 fewer to
Outcome at Longest	^{68,76,78} (n=560		(0.79,1.57)	per 1000	111 more
Available Follow-up)				
(Critical)	10 DOT- 68-	1		161 mara	60 more te
Moderate or severe	71.73-	IOW	(1 47 4 02)	164 more	62 more to
(important)	79,81(n=1738)		(1.47,4.02)	per 1000	284 more
Stroke at 30 days	8 RCTs ⁶⁸⁻	low	OR 1 27	6 more	7 fewer to 30
(important)	^{70,73,74,77,78} (n	1011	(0.66.2.45)	0 more	
(=1626)		(0.00,=)	per 1000	more
Hemolysis at 30 days	3 RCTs	low	OR 5.40	39 more	3 fewer to 93
(important)	(n=403)		(0.63,46.0)	ner 1000	more
			00.057		40 1 70
Peripheral Ischemic	11 RC Is ⁰⁰⁻	low	OR 2.57	41 more	16 more to 79
vascular Complications	79.81(p-1710)		(1.60,4.11)	per 1000	more
Sopsis at 20 days	Q	low		17 more	10 fewer to
(important)	0 BCTe68,69,73,7	1000	(0 71 1 70)		40 lewel 10
	7-79,81		(0.71,1.79)	per 1000	93 more
	(n=1565)				
Renal replacement	8 RCTs	low	OR 1.24	34 more	31 fewer to
therapy at 30 days	68,69,73,7		(0.80,1.92)	ner 1000	118 more
(important)	(n=1592)				

Length of stay in ICU	4 RCTs 68-	low	Mean	1.5 days	6.6 shorter to	
(important)	^{70,73} (n=811)		Difference	longer	9.8 longer	
			1.5 days (-		ere renger	
			0.3,3.2)			
Length of stay in	4 RCTs	low	Mean	2.4 days	0.28 shorter	
hospital (important)	^{68,70,72,73} (n=8		difference	longer	to 4.98 longer	
	11)		2.4 days (-		le nee lenge	
			0.3,4.9)			
In-hospital cardiac	1 RCT	low	14 days (IQ	R 2,45) MCS	group vs. 14	
arrest due to acute	⁷² (n=60)		days (IQR 5,29) in standard care			
coronary syndrome			P=0.73			
Quality of life at 1-year	3 RCTs ^{68,83}	low	No			
(important)	⁷⁷ (n=1052)		difference			

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