SwissEPnet
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Hospital			
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Name	ID						
First name	Date of birth						
Address	Sex	Om Of					
	Phone						
Country							
Inclusion criteria							
☐ Planned extraction procedure for at least one transvenous lead with dwell duration >=12 months or use of dedicated extraction tools (locking stylet, sheath, snare etc)							
Existing device prior to extraction							
Date of last device operation	Overall r	number of prior device operations					
Type of currently implanted device O PM O ICD Number of currently implanted intravasc. leads							
☐ Active RA lead ☐ Active RV lead ☐ Active CS lead							
☐ Active conduction system lead	☐ Active epicardial lea	d (s)					
☐ Prior device upgrade							
_ ·							

Prior to extraction							
Clinical characteristics							
Heightcm	Weight _	kg					
☐ Atrial Fibrillation							
Chronic renal insufficiency (GFR < 60 ml/mir	n.) O non	ie O	yes	O hemodia	lysis	
Comorbidities							
☐ COPD ☐ Hyperter					iac surgery		cclusion
CMP LLL CMP other	details (if 99 = ot	ther)					· · · · · · · · · · · · · · · · · · ·
Echocardiography							
LVEF %							
☐ Vegetation							
Tricuspid valve insufficiency	O none	(O mild	0 1	moderate	O severe	
Medication							
Antiaggregation	O none	(O single	e O	dual		
Anticoagulation	O none	(O NOA	.C interrup	ted >48h	O NOAC co	ntinued
☐ Heparin bridging	O VKA i	nterrupted (O VKA	continued			
ш перапп bridging							
Lead extraction general in	formation						
Date of intervention							
Indication for lead extraction	ı LLL Inc	dication other	details	(if 99 = oth	ner)		
Setting	O EP lab	O Hybrid OR	₹	O OR			
Anesthesia	O conscious s	edation) intubat	ion		
TEE	O none	O in room		O in situ			
Cardiac surgeon	○ none○ planned as I	informed &		-by	O in the C	DR	
Heart lung machine	O none	O informed &	& stand-	-by	O in the C)R	
Bridge occlusion balloon	O none	O prep kit		O balloor	n in situ	O balloon	deployed
Additional intervention during lead extraction							
Additional intervention other details (if 99 = other)							
Re-implantation system	O none O permanent l	○ temporary eadless		- •	nent transven nent epicardia		
Re-implantation device	O none	O temporary	,	ОРМ	O ICD	O S-ICD	O leadless
Re-implantation electrode	O none	O temporary	′	O VVI	O DDD	O CRT	O leadless
Use of antibiotic envelope	O no	O yes					
Operator 1 Operator 2							
Total procedure time	min Total fluc	oro time	min	Total	fluoro dose _	cGy :	x cm ²

Details on leads targeted for extraction						
Number of leads targeted for extraction						
	Lead 1	Lead 2	Lead 3	Lead 4	Lead 5	
Date of implant						
Serial number						
Manufacturer						
Model						
Fixation	O active O passive	O active O passive	O active O passive	O active O passive	O active O passive	
Location of fixation	O RA O RV O CS O Cond. system O Leadless	O RA O RV O CS O Cond. system O Leadless	O RA O RV O CS O Cond. system O Leadless	O RA O RV O CS O Cond. System O Leadless	O RA O RV O CS O Cond. system O Leadless	
Lead access	O left O right	O left O right	O left O right	O left O right	O left O right	
Туре	O pace/sense O single-coil O dual-coil	O pace/sense O single-coil O dual-coil	O pace/sense O single-coil O dual-coil	O pace/sense O single-coil O dual-coil	O pace/sense O single-coil O dual-coil	
Simple traction						
Locking stylet						
Teflon sheath						
Laser sheath						
Mech. rot. sheath (Select the last sheath used for successful extraction)	O no O Philipps TightRai O Cook evolution	○ no○ Philipps TightRai○ Cook evolution	○ noI ○ Philipps TightRai○ Cook evolution	○ noI ○ Philipps TightRai○ Cook evolution	○ noI ○ Philipps TightRail○ Cook evolution	
Femoral snaring	O No O Needle's Eye O Goose neck O EN snare	O No O Needle's Eye O Goose neck O EN snare	O No O Needle's Eye O Goose neck O EN snare	O No O Needle's Eye O Goose neck O EN snare	O No O Needle's Eye O Goose neck O EN snare	
Result					O compl. success O clinical success O failure	
Approach	O superior O inferior O both	superiorinferiorboth	superiorinferiorboth	O superior O inferior O both	O superior O inferior O both	

Complications until discharge						
Total duration of hospital stay at the extraction center days						
Total duration of nospital	stay at the t	extraction center	days			
Death	O no O unrelat	ed to extraction	O complication of the lead extraction procedure O due to the disease which indicated extraction			
SVC laceration	O no	O yes				
Hemato-/Pneumothorax	O no		O interventional drainage	O surgical drainage		
Cardiac tamponade	O no		O interventional drainage	O surgical drainage		
Lead issue	O none	O dislocation	O perforation	O electrical		
☐ none			☐ Stroke			
☐ Unplanned conversion	to cardiac	surgery	☐ HLM in use			
☐ New severe tricuspid ir	nsufficiency		☐ Groin complication (req. ir	ntervention)		
☐ Pulmonary embolism			☐ Newly elevated hemidiaphragm			
☐ Anesthesia complication	on		☐ New diagnosis of thrombosis			
☐ Pocket hematoma (req	ı. interventio	on)	☐ Transfusion of >=2 packed red blood cells			
☐ Cardiac decompensati	on		□SIRS			
☐ other:						
30 days outcome						
Total duration of hospital stay all institutions together days						
Complications occuring	between c	lischarge from e	xtraction center to day 30			
☐ none ☐ Death] Pulmonary emb		☐ Cardiac decompensation		
☐ other:		<u>-</u>		·		
Lead issue O none	C) dislocation	O perforation	O electrical		
Code List	Code List					
Clinical characteristics prior to extraction Lead extraction general information						

Clinical characteristics prior to extraction	Lead extraction general information	
CMP 01 = none 02 = ICM 03 = DCM 04 = valvular 05 = primary electrical disease 06 = GUCH 99 = other	Indication for lead extraction 01 = CIED-related endocarditis without pocket infection 02 = isolated pocket infection 03 = pocket infection with bacteraemia 04 = occult bacteraemia with probable CIED infection 05 = lead dysfunction 06 = upgrade 07 = vascular occlusion 08 = pain 99 = other	Additional intervention during lead extraction 01 = none 02 = PFO occlusion 03 = ASD occlusion 04 = transvenous vegetation aspiration 05 = PTA 99 = other