SwissEPnet		Extraction Form		
	Hospital _			
Name	ID			
First name	Date of birth			
Address	Sex	Om Of		
	Phone			
Country				
lockerion outtouts				
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				

☐ Active epicardial lead (s)

Overall number of prior device operations

Number of currently implanted intravasc. leads

Date of last device operation _____

☐ Active RA lead

☐ Prior device upgrade

Underlying rhythm >=40/min

☐ Active conduction system lead

Type of currently implanted device O PM O ICD

☐ Active RV lead ☐ Active CS lead

O yes O no

Prior to extraction						
Clinical characteristics						
Heightcm	Weigh	t kg				
☐ Atrial Fibrillation						
Chronic renal insufficiency	(GFR < 60 ml/r	min.) O no	one O yes	O hemodia	alysis	
Comorbidities						
☐ COPD ☐ Hyperte				cardiac surgery		occlusion
CMP CMP other	r details (if 99 =	other)				
Echocardiography						
LVEF %						
☐ Vegetation						
Tricuspid valve insufficiend	cy O nor	ie	O mild	O moderate	O severe	
Medication						
Antiaggregation	O nor	ie	O single	O dual		
Anticoagulation	O nor			errupted >48h	O NOAC co	ontinued
☐ Heparin bridging	O VK	A interrupted	O VKA conti	nued		
Lead extraction general i	nformation					
Date of intervention						
Indication for lead extraction	on LLL	Indication othe	er details (if 99	= other)		
Setting	O EP lab	O Hybrid OR	O OR			
Anesthesia	O conscious s	sedation	O intubation	n		
TEE	O none	O in room	O in situ			
Cardiac surgeon	○ none○ informed & stand-by○ planned as hybrid procedure					
Heart lung machine	O none O informed & stand-by O in the OR					
Bridge occlusion balloon	O none	O prep kit	O balloon i	in situ O b	alloon deploye	ed
Additional intervention during lead extraction						
Additional intervention other details (if 99 = other)						
Re-implantation system O none O temporary O perm. transvenous O perm. non-transvenous O permanent leadless O perm. epicardial						
Re-implantation device	O none O	temporary O	PM O ICE	O S-ICD	O EV-ICD	O leadless
Re-implantation electrodes	O none	temporary O	VVI O DD	D O CRT	O leadless	O CSP
Use of antibiotic envelope	O no O	yes				
Operator 1		O	perator 2			
Total procedure time	_ min Total f	luoro time	_ min	Total fluoro dose	сGу	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			
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			
Туре	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)	noPhilipps TightRailCook evolution	○ no○ Philipps TightRail○ Cook evolution	○ no○ Philipps TightRai○ Cook evolution	○ noI ○ Philipps TightRail○ Cook evolution	○ no○ Philipps TightRail○ Cook evolution
Femoral snaring	NoNeedle's EyeGoose neckEN snare	NoNeedle's EyeGoose neckEN snare	NoNeedle's EyeGoose neckEN 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	SuperiorInferiorboth	SuperiorInferiorboth	Superiorinferiorboth	O superior O inferior O both	SuperiorInferiorboth

Complications until disc	harge					
Total duration of hospital	stav at the	extraction center	days			
·	_	SAUGUION COMO				
Death	Death O no O unrelated 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 s	surgery	☐ HLM in use			
☐ New severe tricuspid ir	nsufficiency		☐ Groin complication (req. intervention)			
☐ Pulmonary embolism			☐ Newly elevated hemidiaphragm			
☐ Anesthesia complication		☐ New diagnosis of thrombosis				
☐ Pocket hematoma (req. intervention)		☐ Transfusion of >=2 packed red blood cells				
☐ Cardiac decompensation		□ SIRS				
☐ other:				·····		
				· · · · · · · · · · · · · · · · · · ·		
30 days outcome						
Total duration of hospital stay all institutions together days						
Complications occuring between discharge from extraction center to day 30						
☐ none ☐ Death] Pulmonary embo	olism	☐ Cardiac decompensation		
☐ other:				· · · · · · · · · · · · · · · · · · ·		
Lead issue O none	C) dislocation	O perforation	O electrical		
Code List						

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 06 = direct re-implant of new device 99 = other