National Joint Registry www.njrcentre.org.uk			DS VERS	_	Form: MI	DSv4.0 A2 v1.3
Ankle Single Stage Revision Ankle Stage 1 of 2 Stage Revision Ankle Stage 2 of 2 Stage Revision Ankle Conversion to Arthrodesis Amputation			ent Addressograp			
Important: Please tick relevant boxes. All compositived to the accompanying 'Minimum Labels Sheet'. Please ensure that all she	Dataset Form Cor	nponent				
All fields are Mandatory unless other	wise indicated					
REMEMBER! MAKE A NOTE OF THE NJR I	REFERENCE NUMB	ER WHEN YOU	ENTER THIS DATA	NJR REF:		
PATIENT DETAILS						_
Patient Consent Obtained	Yes □	No □	Not Recorde	ed □		
Patient Hospital ID						
Body Mass Index (enter either H&W OR BMI OR tick Not Available box)	Height (IN CM) Weight (IN KG)		ВМІ	ı	Not Availabl	e □
						_
PATIENT IDENTIFIERS						
Forename						
Surname						
Gender	Male □	Female □	Not Known		Not Specifie	d□
Date of Birth	DD/MM/YYYY					
Patient Postcode			Overseas A	ddress 🗆		
NHS Number (if available)						
OPERATION DETAILS						
Hospital						
Operation Date	DD/MM/YYYY					
Anaesthetic Types	General Regional – Epidu	□ ural □	•	– Nerve Block – Spinal (Intrathe	□ cal) □	
Patient ASA Grade	1 🗆	2 🗆	3 □	4 🗆	5	
Operation Funding	NHS □	Independ	dent 🗆			
SURGEON DETAILS						
Consultant in Charge						
Operating Surgeon						
Operating Surgeon Grade	Consultant	SpR/ST3-8 [☐ F1-ST2 ☐	Specialty Docto	or/SAS □	Other

First Assistant Grade

Consultant □

Other

ANKLE REVI	SION PROCEDURE D	DETAILS						
	Single Stage Revi			Conversion	to Arthrodesi	-		
	•			Amputation	ı			
	Stage 2 of 2 Stage	Revision						
Side	Left □	Righ	nt 🗆					
	Infection			Implant Fr	acture			
	High Suspicion (eg pus or confined micro	o) 🗆	Tibial Co	mponent			
		awaiting micro/histo)		Talar Co	mponent			
l	Aseptic Loosenin	ng			Component			
Indications For /		nt			olyethylene (-	nt 🗆	
Findings at Tim of Revision (sel		nt			nsert Disloca			
all that apply)	Lysis			=	nt Migration/	Dissociati		
	Tibia			Pain (undi	iagnosed)			
	Talus			Stiffness				
	Malalignment				e Impingeme	ent		
				Other				
PRIMARY OF	PERATION DETAILS							
Primary Operat	ion Date OR Year	DD/MM/YYYY	Please ente	er Date if known		Not Avail	able □	
Primary Operat	ion Hospital					Not Avail	able □	
	ΓS REMOVED (Do no		2 of 2 Stage	e Revision)				
Components R	emoved	Tibial Yes □	No □ Tala	r Yes □	No □ M	eniscal	Yes □	No □
Brand					Not Availab	ole 🗆		
Arthrodesis (PPROACH (Used for Patient Procedure ar	nd Surgeon's Notes (Only))					
	Prosthetic Replacement I	-		Fusion (Subtalar	-		itting)	
Tadoric Company								
1 -	Elsewhere (eg Hybrid)		Pantal	ar Fusion		i i C i Naii)		
Approach		Anterior		Anterolatera Other	al 🗆			
		Lateral (transfibular)						
Associated Pro	codures at the time	Subtalar Joint Fusion Talonavicular Fusion			ula Osteotomy		y C	
Associated Procedures at the time of surgery* (select all that apply)			ant Ootootomy		dial Malleolar eral Ligament		=	
	11 37	Calcaneal Displacement Achilles Tendon Length	•		dial Ligament			
*Also select if previously carried out or procedures	Fusion Distal Tibiofibu	•	□ lvied	-	Reconstit			
are planned at the time of index surgery		1 dalon Diatai Hololiba	iiai Joint	□ Otil Nor				
THROMBOPRO	OPHYLAXIS REGIME (ii	ntention to treat)		110.	10			_
	····	Aspirin		Warfarin			None	•
Chemical		LMWH			mbin Inhibito		. 10.10	. –
		Pentasaccharide		Other				
		Foot Pump		Oth	ier 🗆			
Mechanical		Intermittent Calf Comp	oression	Nor	_			
meenamea.		TED Stockings						
BONE GRAFT								
Was bone graft	used?	Yes (structural) □	Yes (no	on-structural) 🛘		No □		
What type of gr	aft was used?	Allograft □	Auto	graft □		Synthetic	c Graft □	
SURGEON'S N								
INTRA-OPER	ATIVE EVENT							
		None		Fr	acture (other)			
	Operative Event	Fracture medial malle			gament Injury			
(select all that apply)	Fracture lateral malled	olus 🗆		ther				

Minimum Dataset Form - COMPONENT LABELS

1.	Please affix any component labels to this sheet and ensure any extra component label sheets are attached to the main
	Minimum Dataset Form.
2.	Ensure all component details are provided, including cement.
3.	The NJR DOES NOT record the following: wire, mesh, cables, plates, screws, surgical tools, endoprostheses or bipolar heads.