

<div>E2</div> <div> Single stage revision (includes DAIR with modular exchange and modular exchange for indications other than infection) Stage 1 of 2 stage revision Stage 2 of 2 stage revision Conversion to arthrodesis Partial excision arthroplasty (i.e. removal of radial head prosthesis) Excision arthroplasty Amputation </div>	Patient addressograph
Important: Please tick relevant boxes. All component stickers should be affixed to the accompanying 'Minimum Data Set form component labels sheet'. Please ensure that all sheets are stapled together.	

All fields are mandatory unless otherwise indicated

Remember! Make a note of the NJR reference number when you enter the data	NJR ref:
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Patient details			
NJR patient consent obtained	Yes <input type="checkbox"/>	No <input type="checkbox"/>	Not recorded <input type="checkbox"/>
If 'Yes' or 'No' was selected for patient consent above, was consent provided by a consultee on behalf of the patient?	Yes <input type="checkbox"/>	No/Not known <input type="checkbox"/>	This refers to NJR consent being provided by a third party on behalf of the patient, not the 'consent to operate' normally obtained by the consultant. If an 'NJR Patient Consultee Declaration Form' has not been completed, this section should be completed as 'No/Not known'
Body Mass Index (enter either H&W or BMI or tick Not available box)	Height (In M) Weight (In KG)	BMI	Not available <input type="checkbox"/>
Handedness	Left <input type="checkbox"/>	Right <input type="checkbox"/>	Ambidextrous <input type="checkbox"/> Unknown <input type="checkbox"/>

Patient identifiers			
Forename(s)			
Surname			
Sex	Male <input type="checkbox"/>	Female <input type="checkbox"/>	Indeterminate <input type="checkbox"/>
Date of birth (DD/MM/YYYY)			
Patient postcode		Overseas address <input type="checkbox"/>	
NHS number or National Patient Identifier (if available)			
Patient hospital ID			
Patient email address (if provided)			
Patient mobile phone number (if provided)			

Operation details				
Hospital				
Operation date (DD/MM/YYYY)				
Anaesthetic types	General <input type="checkbox"/>	<input type="checkbox"/>	Regional – nerve block <input type="checkbox"/>	<input type="checkbox"/>
Patient ASA grade	1 <input type="checkbox"/>	2 <input type="checkbox"/>	3 <input type="checkbox"/>	4 <input type="checkbox"/> 5 <input type="checkbox"/>
Operation funding	NHS <input type="checkbox"/>	Independent <input type="checkbox"/>		

Surgeon details													
Consultant in charge													
Operating surgeon number one													
Operating surgeon number one Grade		Consultant <input type="checkbox"/>	SPR/ST3-8 <input type="checkbox"/>	Senior Fellow (Post-CCT or equivalent) <input type="checkbox"/>	F1-ST2 <input type="checkbox"/>	Specialty doctor/SAS <input type="checkbox"/>	Other <input type="checkbox"/>						
Dual consultant operation?		Yes <input type="checkbox"/>		No <input type="checkbox"/>									
Operating consultant number two (if dual consultant operation)													
First assistant Grade		Consultant <input type="checkbox"/>		Other <input type="checkbox"/>									
Elbow revision procedure details													
Procedure type	Single stage revision (includes DAIR with modular exchange and modular exchange for indications other than infection)	<input type="checkbox"/>	Conversion to arthrodesis		<input type="checkbox"/>								
			Partial excision arthroplasty (i.e. removal of radial head prosthesis)		<input type="checkbox"/>								
			Excision arthroplasty		<input type="checkbox"/>								
	Stage 1 of 2 stage revision		<input type="checkbox"/>	Amputation		<input type="checkbox"/>							
	Stage 2 of 2 stage revision		<input type="checkbox"/>										
Revision of		Primary arthroplasty		<input type="checkbox"/>	Previous revision arthroplasty		<input type="checkbox"/>						
Side		Left <input type="checkbox"/>		Right <input type="checkbox"/>									
Indications for/findings at time of revision (select all that apply)		Infection		<input type="checkbox"/>									
		Instability		<input type="checkbox"/>									
		Failed Hemiarthroplasty		<input type="checkbox"/>									
					Ulnar	Humeral	Radial head						
		Breakage/dissociation of prosthesis			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>						
		Aseptic loosening			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>						
		Lysis			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>						
		Periprosthetic fracture			<input type="checkbox"/>								
		Unified Classification System (UCS) site											
		Humerus, distal			A	B1	B2	B3	C	D	E	F	N/A
		Ulna/radius, proximal			A	B1	B2	B3	C	D	E	F	N/A
		<i>A Apophyseal or extraarticular/periarticular</i> <i>B1 Bed of the implant or around the implant – prosthesis stable, good bone</i> <i>B2 Bed of the implant or around the implant – prosthesis loose, good bone</i> <i>B3 Bed of the implant or around the implant – prosthesis loose, poor bone or bone defect</i>			<i>C Clear of or distant to the implant</i> <i>D Dividing the bone between two implants</i> <i>E Each of two bones supporting arthroplasty</i> <i>F Facing and articulating with a hemiarthroplasty</i>								
		Other: <input type="checkbox"/> If "Other" selected, please enter text (max 25 characters):											

Components removed (do not complete for stage 2 of 2 stage revision)				
Radial component removed?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	Not applicable <input type="checkbox"/>	
Humeral component removed?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	Not applicable <input type="checkbox"/>	
Ulnar component removed?	Yes <input type="checkbox"/>	No <input type="checkbox"/>	Not applicable <input type="checkbox"/>	
Surgical approach (used for single stage revision and stage 2 of 2 stage revision)				
Patient procedure (i.e. revision to)	Revision total prosthetic replacement			<input type="checkbox"/>
	Revision radial head replacement			<input type="checkbox"/>
	Revision to lateral resurfacing			<input type="checkbox"/>
	Revision distal humeral hemiarthroplasty			<input type="checkbox"/>
	Debridement And Implant Retention (DAIR) with modular exchange			<input type="checkbox"/>
	Modular exchange for indications other than infection			<input type="checkbox"/>
Fixation type (not applicable for DAIR)	Uncemented <input type="checkbox"/>	Cemented <input type="checkbox"/>	Hybrid <input type="checkbox"/>	
Approach	Lateral	<input type="checkbox"/>	Posterior triceps off	<input type="checkbox"/>
	Posterior triceps on	<input type="checkbox"/>	Medial	<input type="checkbox"/>
Thromboprophylaxis regime (intention to treat)				
Chemical			In hospital	At home
	Aspirin		<input type="checkbox"/>	<input type="checkbox"/>
	LMWH		<input type="checkbox"/>	<input type="checkbox"/>
	Pentasaccharide (e.g. Fondaparinux)		<input type="checkbox"/>	<input type="checkbox"/>
	Warfarin		<input type="checkbox"/>	<input type="checkbox"/>
	Direct thrombin inhibitor (e.g. Dabigatran)		<input type="checkbox"/>	<input type="checkbox"/>
	Factor Xa inhibitor (e.g. Rivaroxaban/Apixaban)		<input type="checkbox"/>	<input type="checkbox"/>
	Other		<input type="checkbox"/>	<input type="checkbox"/>
	None		<input type="checkbox"/>	<input type="checkbox"/>
Mechanical	Foot pump	<input type="checkbox"/>	Other	<input type="checkbox"/>
	Intermittent calf compression	<input type="checkbox"/>	None	<input type="checkbox"/>
	TED stockings	<input type="checkbox"/>		

Bone graft used (not applicable for DAIR)								
Was humeral bone graft used?	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>				
Humeral – form	Structural	<input type="checkbox"/>	Morsellised/chips	<input type="checkbox"/>				
Humeral – type	Autograft	<input type="checkbox"/>	Allograft	<input type="checkbox"/>	Synthetic	<input type="checkbox"/>	Other	<input type="checkbox"/>
Was ulnar bone graft used?	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>				
Ulnar – form	Structural	<input type="checkbox"/>	Morsellised/chips	<input type="checkbox"/>				
Ulnar – type	Autograft	<input type="checkbox"/>	Allograft	<input type="checkbox"/>	Synthetic	<input type="checkbox"/>	Other	<input type="checkbox"/>
Structural implant or other augment(s) used e.g. augment, block, wedge, restrictor, sleeve								
Were humeral structural implant or other augment(s) used?	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>				
Were radial structural implant or other augment(s) used?	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>				
Were ulnar structural implant or other augment(s) used?	Yes	<input type="checkbox"/>	No	<input type="checkbox"/>				
If yes, please affix implant labels under “Accessories”								
Surgeon’s notes								
Intra-operative event								
Untoward intra-operative event (select all that apply)	None	<input type="checkbox"/>	Fracture ulna	<input type="checkbox"/>				
	Shaft penetration humerus	<input type="checkbox"/>	Nerve injury	<input type="checkbox"/>				
	Shaft penetration ulna	<input type="checkbox"/>	Vascular injury	<input type="checkbox"/>				
	Fracture humerus	<input type="checkbox"/>	Other	<input type="checkbox"/>				

Minimum Data Set 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 Data Set form.
2. Ensure all component details are provided, including cement.
3. The NJR **does not** record the following: wire, mesh, cables or surgical tools.

Ulnar component (if used)	Humeral component
Radial component (if used) Required for hemiarthroplasty	Cement (if used)
Accessories	