# SH61: What is the Optimal Timing of Second Stage Revision Surgery in the Treatment of PJI? Is this Dependent on Organism?

**Liaison:** Benjamin Zmistowski **Lead delegate:** Joideep Phadnis

Supportive delegates: Jessica Seidelman, Jason Ho

**Recommendation:** Based on the literature an optimal time to reimplantation (TTR) during two stage exchange to treat shoulder PJI cannot be recommended regardless of pathogen. However, there is evidence to suggest that increasing TTR is associated with adverse outcomes. Clinicians should ideally undertake re-implantation in a timely manner once it is safe to do so following antibiotic completion and when the patient has been adequately optimised for further surgery with the knowledge that delay may affect outcome.

#### Strength of recommendation: Limited

**Rationale:** While there is contention as to whether single or two stage revision is the optimal treatment for shoulder prosthetic joint infection (PJI), two-stage replacement remains the gold standard for many patients, particularly in the context of chronic and subacute infection[1].

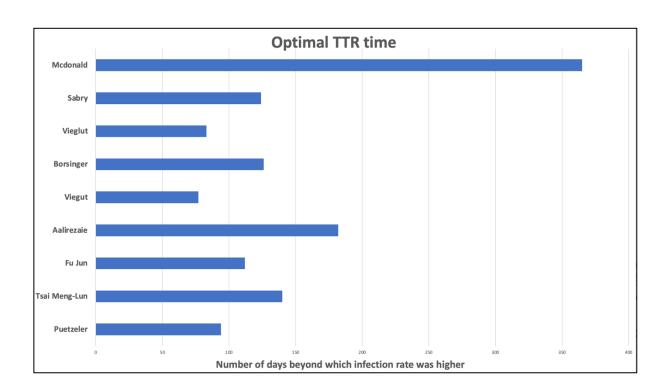
Multiple factors need to be considered when planning second-stage re-implantation. These include biochemical markers, soft tissue envelope, and virulence of the organism to name a few. Variation in time to reimplantation (TTR) has been suggested as a factor that may influence outcome, and because TTR is controllable in most cases, it is appealing to understand what the optimal TTR is for two-stage revision.

A clinical librarian performed a literature review using Embase, Ovid Medline, Pubmed and UpToDate databases to determine the answer to the question. The full search strategy is outlined in the appendix. There were no articles identified in the shoulder literature investigating the research question, hence relevant articles from the hip and knee literature were included in the literature synthesis to provide some guidance on TTR. Even these articles were all retrospective in nature with some case series and cohort studies, representing evidence subject to considerable bias.

Table 1 summarises articles of interest in the available literature and the below chart visually displays their recommended upper limit for TTR.

Article	Number of Arthroplasties	Optimal TTR beyond which re- infection rate was higher	Magnitude of effect	Effect by organism
Puetzler[2]	163	>94 days	2.8 x higher re-infection $p=0.004$	No
Tsai Meng- Lun[3]	361	112-140 days	1.27 odds ratio Not significant	No
Fu Jun[4]	81	84-112 days	7.5% vs 17% re-infection <i>p</i> =0.001	No
Aalirezaie[5]	282	>182 days	21% vs 44% re-infection <i>p</i> =0.057	No

Viegut[6]	76	28-77 days	90% success vs	No
			0% (<28 days)	
			and 52% (>77	
			days) $p = 0.01$	
Winkler[7]	38	<28 days and >28	5% vs 0%	No
		days equivalent	re-infection	
		outcome	No significant	
			difference	
Borsinger[8]	101	>126 days	Odds ratio 7.0	No
			p=0.002	
Vielgut [9]	77	>83 days	Odds ratio 6.1	No
			p=0.007	
Sabry	314	>96 days	Increased median	Gram negative
[10]			TTR in re-	higher failure
			infection group	but not
			(124 vs 96 days)	assessed in
			p=0.01	relation to time
				to
				reimplantation
McDonald	82	>365 days	27% vs 7%	Gram negative
[11]			re-infection rate	higher failure
			p=0.05	but not
				assessed in
				relation to time
				to
				reimplantation



A common theme running through all studies was that increasing TTR was associated with an adverse effect on re-infection rate. However, the included studies are subject to bias because of their retrospective nature and due to the multiple confounding variables that influence failure after two-stage revision for PJI. An example, is that there is little information offered in any of the studies as to why some patients had longer TTR than others within the same series. It is likely that these patients may have had other reasons for delay in reimplantation that would have adversely affected their re-infection rate. Similarly, there is marked heterogeneity within and between studies regarding antibiotic treatment and reporting of bacterial virulence. This makes independently attributing variation in TTR with increased infection rates unreliable and determining the effect of bacterial virulence very difficult. For clinical practice recommendation and accounting for the limitations of the evidence, the scattergram above suggests that surgeons should probably try to avoid extending TTR beyond 3-4 months following the first stage revision. Only two studies looked at the lower limit for

scattergram above suggests that surgeons should probably try to avoid extending TTR beyond 3-4 months following the first stage revision. Only two studies looked at the lower limit for safe TTR. Vielgut concluded that reimplantation at <4 weeks was associated with a higher reinfection rate than between 4 and 11 weeks[6], whereas Borsinger et al. found that the lowest risk of re-infection was between 0-12 weeks and highest >18 weeks in their series[8]. While these findings are far from conclusive, they do suggest that there is no advantage to very early re-implantation.

### **References:**

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#### Appendix 1

#### **Search Strategy and Terms Used**

Data bases searched: Embase, Ovid medline, Pubmed, UpToDate Date of Search: September 30<sup>th</sup> 2024 performed by Lucy Wells (medical librarian). 52 Articles Identified in initial search

## **Full Search methodology:**

Ovid MEDLINE(R) ALL <1946 to September 26, 2024>								
1	Prosthesis-Relate	ed	Ir	nfections/	15299			
2 (pro	osthe*	adj2		infect*).ti,a	b. 6338			
3 ((periprosthe*	or peri-prosthe*	or peri	prosthe*)	adj2 joint	infect*).ti,ab. 4218			
4 1	or	2	O	r	3 20441			
5		Reoperation	on/		97842			
6 ((two-stage or t	wo stage or seco	ond-stage	or second	stage) adj2	revision).ti,ab. 1127			
7	5	or		6	98385			
8 ((optimal or pr	oper or evaluat	* or inter	rval) adj5	(time or ti	ming)).ti,ab. 129476			
9 4 and 7 and 8 78								
Embase <1974 to 20	024 Week 39>							
1 ex	p p	rosthesis		infection/	14536			
		11.0						
2 (pro	osthe*	adj2		infect*).ti,a	b. 7969			
•	osthe* or peri-prosthe*	3	prosthe*)	, ,				
•		3	prosthe*)	adj2 joint				
3 ((periprosthe*	or peri-prosthe* or	or peri	0	adj2 joint	infect*).ti,ab. 4330			
3 ((periprosthe* 4 1 5	or peri-prosthe* or	or peri 2 reoperatio	o n/	adj2 joint r	infect*).ti,ab. 4330 3 19266			
3 ((periprosthe* 4 1 5	or peri-prosthe* or	or peri 2 reoperatio	o n/	adj2 joint r	infect*).ti,ab. 4330 3 19266 111934			
3 ((periprosthe* 4 1 5 6 ((two-stage or t 7	or peri-prosthe* or wo stage or second	or peri 2 reoperatio ond-stage or	on/ or second	adj2 joint r stage) adj2	infect*).ti,ab. 4330 3 19266 111934 revision).ti,ab. 1191			