## Intraosseous Vancomycin Reduces the Rate of Periprosthetic Joint Infection Following Aseptic Revision Total Knee Arthroplasty

Colin A McNamara, Austin Wininger<sup>1</sup>, Thomas Castlen Sullivan, Timothy Scott Brown, Terry A Clyburn, Stephen J Incavo<sup>2</sup>, Kwan Park<sup>2</sup>

<sup>1</sup>Houston Methodist Orthopedics and Sports Medicine, <sup>2</sup>Houston Methodist Hospital

INTRODUCTION: Periprosthetic joint infection (PJI) is a devastating complication following total knee arthroplasty (TKA). Prior literature supports the intraosseous (IO) delivery of vancomycin as a safe and effective technique for primary TKA. The purpose of this study was to evaluate its efficacy for aseptic revision TKA.

METHODS: A single-institution retrospective review was performed on patients who underwent aseptic revision TKA from May 2016 to October 2023. Vancomycin was administered through an intravenous (IV) route in 386 cases and via an IO infusion in 333 cases. The IV cohort received a 15mg/kg dose of vancomycin prior to skin incision. The IO cohort received a 500mg dose of vancomycin infused into the tibia after tourniquet inflation. All patients also received a weight-based dose of IV cefazolin perioperatively. Patient characteristics, surgical details, and infection-related data were extracted during chart review. PJI diagnosis was based on the 2018 Musculoskeletal Infection Society criteria. Fisher's exact tests and chi-square analyses were used to compare categorical outcomes.

RESULTS: The incidence of PJI was significantly lower in the IO cohort compared to the IV cohort at 30-day (0.3% vs 2.1%, P = 0.03), 90-day (0.9% vs 3.1%, P = 0.04), and 1-year follow-up (1.6% vs 4.9%, P = 0.04). There were no reported adverse reactions to vancomycin. There were no differences in the incidence of acute kidney injury (2.7% vs 2.9%, P =0.90), deep venous thrombosis (1.2% vs 1.8%, P = 0.56), or pulmonary embolism (0% vs 0.3%, P = 1.0) between groups.

DISCUSSION AND CONCLUSION: Intraosseous vancomycin infusion is a safe and effective alternative to IV administration for patients undergoing aseptic revision TKA. Furthermore, IO vancomycin optimized the efficiency of vancomycin administration in this high-risk surgical cohort and resulted in a significant reduction in the rate of PJI through 1-year follow-up.

Asep Ca Eli	tic Revision TKA ses Assessed for gibility (N=843)	
		<ul> <li>Divided.</li> <li>- Previously underwent revision surgery for PII of the same knee (N= 57).</li> <li>- Prosperative diagnosis of periprosthetic fracture (N= 30).</li> <li>- Postoperative follow-up &lt;50 days (N= 20).</li> <li>- Review follow-up &lt;50 days (N= 20).</li> <li>- Review follow-up &lt;50 days (N= 20).</li> <li>- Review follow-up &lt;50 days (N= 20).</li> <li>- Trianta to the operative knee caused an arthrotomy requiring resperation (N= 4).</li> <li>- PII offend a state hematogenous infection (N= 2).</li> </ul>
Ir	icluded (N=719)	]
IO Vancomycin (N=333)	IV	Vancomycin (N=386)

Organism	IV	IO	Outcome	Post- Operative	IV. N (%)	IO. N (%)	P-value
Methicillin-sensitive	4	2		Time Frame			
Staphylococcus aureus			Periprosthetic	30 Day	8 (2.1%)	1 (0.3%)	0.0331
Coagulase-negative	2		Joint Infection	90 Day 1 Year	12 (3.1%) 15 (4.9%)	3 (0.9%) 4 (1.6%)	0.0389
Stanhylococcus aureus	2	1	Wound	30 Day	7 (1.8%)	2 (0.6%)	0.1877
Methicillin-resistant	1	1	Complications Not Requiring	90 Day	9 (2.3%)	2 (0.6%)	0.0713
Staphylococcus aureus			Reoperation	1 Year	9 (2.9%)	2 (0.8%)	0.1229
Enterobacter cloacae	2	0	Wound Complications	30 Day 90 Day	6 (1.6%) 9 (2.3%)	3 (0.9%) 5 (1.5%)	0.5158 0.4218
Group B Streptococcus	2	0	Requiring Reoperation	1 Year	9 (2.9%)	5 (2.0%)	0.5093
Pseudomonas aeruginosa	2	0	Cumulative Infections	30 Day	21 (5.4%)	6 (1.8%)	0.0105
Citrobacter koseri	1	0		90 Day 1 Year	30 (7.8%)	10 (3.0%) 11 (4.5%)	0.0054
Proteus mirabilis	1	0			(10.8%)		I