

# Treatment of prosthetic joint infection: DAIR with short duration of rifampicin

### Introduction

Evidence for prolonged rifampicin therapy for prosthetic joint infections (PJI) is limited and treatment limiting adverse events are significant. We hypothesized that the role of rifampicin is most relevant in the early phase of treatment, immediately after surgical debridement. The outcome of PJI treated with Debridement, Antibiotics and Implant Retention including short duration of rifampicin (5 days) was evaluated.



#### Baseline characteristics of 67 patients with PJI Demographics All (n=67) Age at diagnosis (mean, range) 58 (15-92) Sex (male, %) 36 (54%) Implant site (n, %) 33 (49%) Hip Knee 25 (37%) Shoulder, elbow, ankle 1,6,2 (14%) Revision<sup>#</sup> (n, %) 25 (37%) **Comorbidities** (n, %) Diabetes mellitus 7 (10%) Rheumatoid arthritis 16 (24%) Orthopaedic oncology<sup>\$</sup> 17 (25%) Use of immunosuppressant's<sup>^</sup> 18 (27%) **Clinical characteristics** Duration of symptoms <8 davs 46 (69%) 11 (16%) 8-20 days 2 (3%) 21-27 days >27 days 8 (12%) Fever 30 (45%) Fistula 6 (9%) ESR (mean, range) 75 (13-140) Number cultures taken (mean) 4.8 (1-10) Number of positive cultures per 3.6 (0-10) patient (mean, range) Polymicrobial infection 16 (24%)



### Conclusions

- In a selected patient population, the outcome of acute PJI treated with DAIR including only 5 days of rifampicin was 87% (comparable to published cohort studies and a randomized trial using at least 3 months of rifampicin combination therapy).
- No selection of rifampicin-resistant staphylococci was found in relapses
- The overall cure rate reflects our frail patient population.
- To the best of our knowledge, no studies have been published in which a shorter treatment duration with rifampicin was investigated
- A RCT comparing long term rifampicin with short term rifampicin therapy is needed.

# Methods and treatment protocol

All patients with PJI who were treated with surgical debridement and antimicrobial combination therapy including rifampicin - which in our center included only 5 days of rifampicin - followed by monotherapy based on culture and antibiotic susceptibility (e.g. flucloxacillin for *S aureus*) were enrolled in a cohort study (2003-2014). Treatment consisted of prompt, extensive surgical debridement, rinsing with povidone iodine and pulsed lavage. Rifampicine was started immediately after debridement. Outcomes and risk factors for treatment failure were assessed.



# **Definition of cure and failure:**

**Cure**: absence of infection and a stable implant for at least six months after stopping antibiotic therapy.

**Failure**: either chronic suppressive antibiotic therapy with retention of the prosthesis or progression or relapse of infection leading to removal of the prosthesis

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ubgroup outcomes of PJI: DAIR with antibiotics ncluding only 5 days rifampicin		
	n	Cure
ll patients	67	58%
uration of symptoms		
Acute (< 21 days)	57	63%
Chronic (≥ 21 days)	10	30%
atients with steroids/anti-TNF/MTX	18	39%
atients with tumour prosthesis	17	47%
cute* staphylococcal hip or knee PJI	26	69%
cute* staphylococcal hip PJI	14	86%
Ionomicrobial staphylococcal hip PJI	15	87%
Acute = symptoms or last operation < 3 week	(5	

# **Risk factors for failure**

During 2003-2012 liners were generally not exchanged. Chronic PJI (RR 1.90, 95%CI 1.12-3.23) and immunosuppressive therapy (RR 1.76, 95%CI 1.03-3.00) were associated

with increased risk for failure.