Opioid agonist treatment and risk of death or rehospitalisation following injection drug use-associated bacterial and fungal infections

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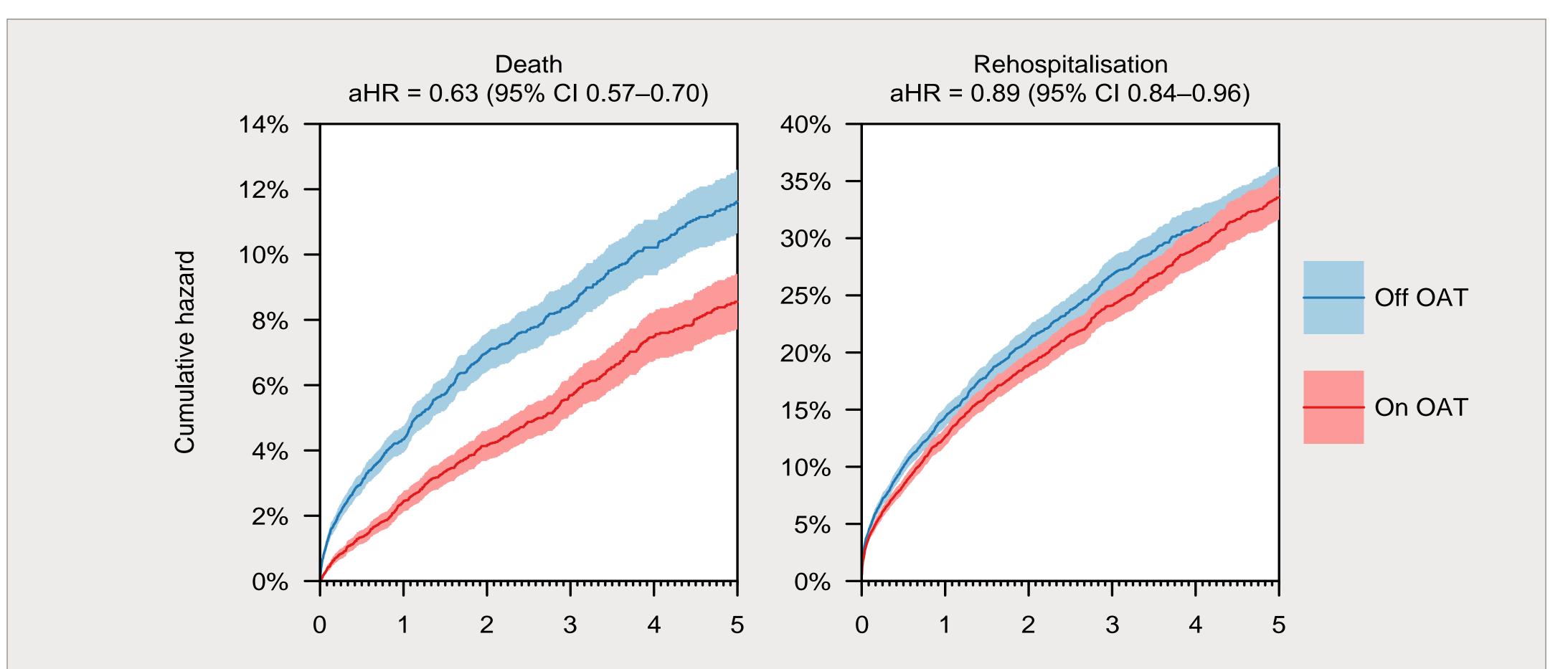
Research question: Is opioid agonist treatment (methadone or buprenorphine) associated with reduced death and reinfection after injecting-related infections?

Sample: 8,943 people with severe injecting-related infections in New South Wales, Australia. This was defined as patients admitted to hospital with a skin or soft tissue infection (79% of patients) or an invasive infection (21% of patients), and at least one prior episode of opioid agonist treatment. Mean age at admission was 39 (sd. 11); 34% were women.

Exposure: Opioid agonist treatment after discharge (time-varying).

Outcomes: (1) All-cause death; (2) Reshospitalisation with an injecting-related infection.

Analysis: Survival analysis, with adjustment for age, sex, Aboriginal or Torres Strait Islander status, comorbidity at time of admission, prior incarceration, and prior records of stimulant use.



Years after hospital discharge

Figure: Extended Kaplan–Meier curves for time to death and time to rehospitalisation among participants in the OATS study who survived an initial hospitalization with injecting-related bacterial or fungal infection

Results: 4,292 (48%) were receiving OAT at the time of discharge. During median 6.5 years followup, 1,481 patients died and opioid agonist treatment was associated with a large reduction in risk. During median 3.4 years follow-up, 3,653 were re-hospitalised and opioid agonist therapy was associated with a small reduction in risk.

Conclusion: Opioid agonist therapy should be considered as part of a multicomponent treatment strategy for injecting-related infections, aiming to reduce death and reinfection.

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