REAL-WORLD EXPERIENCE WITH PROLONGED TEDIZOLID TREATMENT: EFFECTIVENESS AND SAFETY

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BACKGROUND AND IMPORTANCE

Tedizolid demonstrated to be effective and safe in clinical trials. However, evidence concerning its effectiveness and tolerability in long-term treatments is still scarce.

AIM AND OBJECTIVES

To assess the use of tedizolid and its effectiveness and safaty in long term treatments.

MATERIAL AND METHODS

Observational and retrospective study (1 November 2018 - 31 May 2021)

Inclusion criteria

- > 18 years-of-age.
- Tedizolid treatment for seven or more days

Variables

Collected from electronic medical record

- Demographic
- Clinical
- Safety

Assessment of effectiveness



Clinical resolution of infection and absence of recurrence

Assessment of safety



- Incidence of adverse drug events
- Changes in haemoglobin and platelet count

RESULTS

Demographics

N = 39 patients

• 51% female

Median age: 71 (IQR 52-76)

Tedizolid treatment

Median duration: 15 days (IQR 10-20)

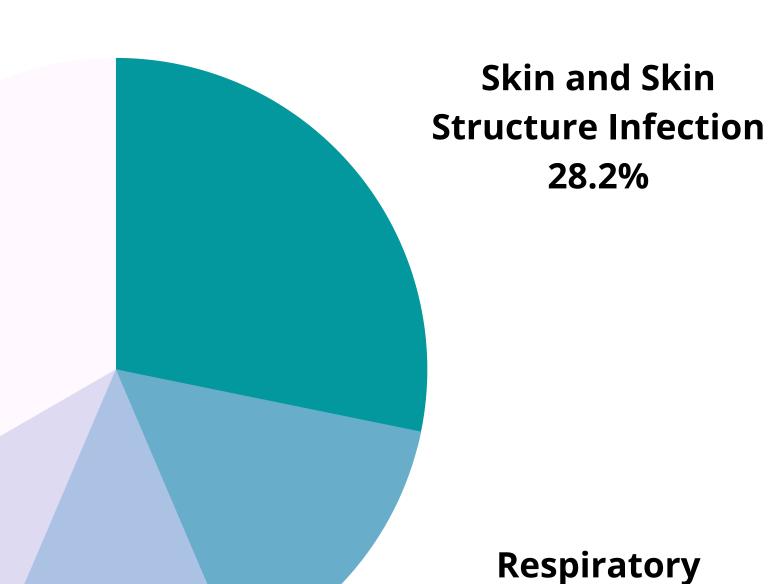
Previous linezolid: 30% (n=12)

Main reason for tedizolid selection:

Drug-drug interactions (53,8%, n=21)

Previous cytopenia (43,5%, n=17)

Treatment indication

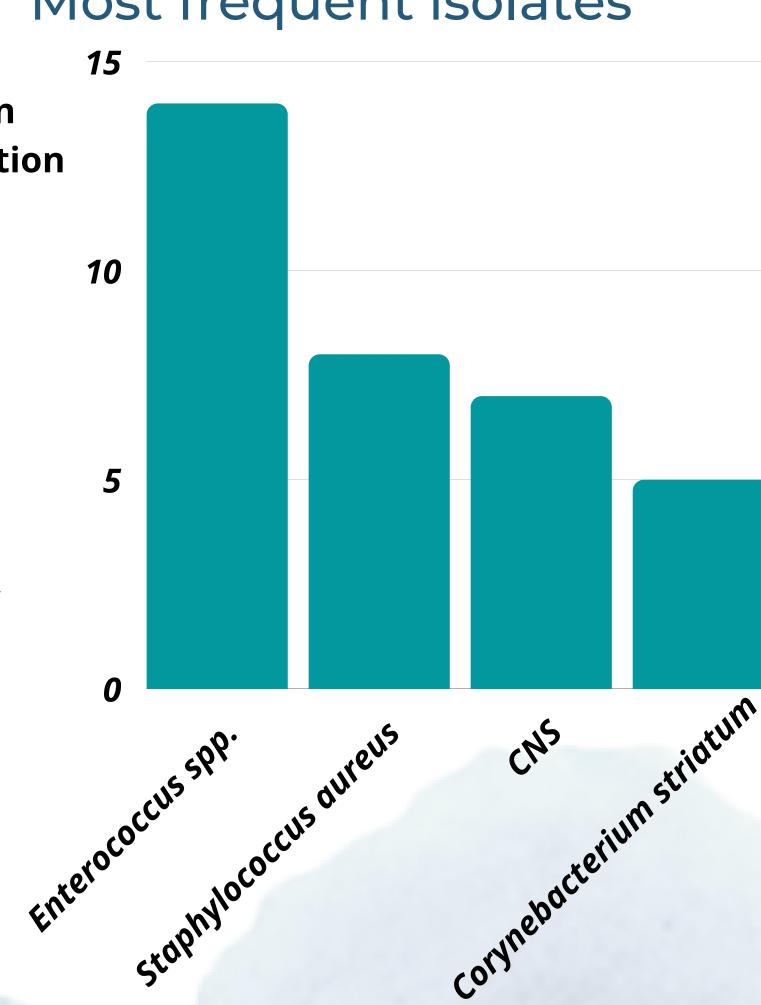


Infection

15.4%

Prostetic Joint Infection 12.8%

Most frequent isolates



Safety outcomes

- No serious adverse drug events reported.
- Mean haemoglobin value unchanged (10.2 g/dL to 10.5 g/dL, p=0,18)
- Mean platelet count unchanged (243,9x10^6/mL to 243,7x10^6/mL, p=0,98)
- Absence of serotoninergic syndrome in patients treated with serotoninergic drugs (n=21)

Other

33.3%

Osteomyeliti

10.3%

Clinical outcomes

79,4% (n=27): favourable evolution, 20,6% (n=7): treatment failure
Three patients were lost during follow-up period, two died of unrelated causes

CONCLUSIONS AND RELEVANCE

In most cases, tedizolid was used for off-label indications. Our results suggest that tedizolid can be an effective and safe alternative in the management of diverse gram positive infections, even in prolonged treatments.





