

Stenotrophomonas maltophilia treatment

Treatment of [Stenotrophomonas maltophilia infections](#) can be difficult, as *S. maltophilia* is inherently resistant to many classes of [antibiotics](#) ¹⁾.

A bacterial strain of the Gram-negative opportunistic pathogen *Stenotrophomonas maltophilia* capable of degrading [colistin](#) and exhibiting a high-level colistin resistance was isolated from the soil environment. A colistin-degrading protease (Cdp) was identified in this strain, and its contribution to colistin resistance was demonstrated by growth inhibition experiments using knock-out (Δ cdp) and complemented (Δ cdp::cdp) mutants ²⁾

Stenotrophomonas maltophilia is an urgent global threat due to its increasing incidence and intrinsic antibiotic resistance. Antibiotic development has focused on carbapenem-resistant Enterobacteriaceae, *Pseudomonas*, and *Acinetobacter*, with approved antibiotics in recent years having limited activity for *Stenotrophomonas*. Accordingly, novel treatment strategies for *Stenotrophomonas* are desperately needed ³⁾.

[Fluoroquinolone](#) and trimethoprim-sulfamethoxazole (SXT) monotherapies may be equally effective for the treatment of *S. maltophilia* infections. Resistance was documented in subsequent isolates of *S. maltophilia* in both groups ⁴⁾.

Antibiotics with in vitro activity against *S. maltophilia* include [trimethoprim-sulfamethoxazole](#) (SXT), [fluoroquinolones](#) (FQs), [tetracyclines](#), ticarcillin-clavulanate, and [ceftazidime](#); however, there are limited clinical data on the use of these agents ^{5) 6) 7) 8) 9) 10)}

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