



Don't Shrug At Lug: Spotting *Staphylococcus lugdunensis*

Staphylococcus lugdunensis is a coagulase-negative Staphylococcus (CoNS) that has historically been misidentified as *Staphylococcus aureus* or classified as a nonspecific CoNS prior to the adoption of mass-spectrometry microbiological identification.¹ Although it is a CoNS, *S. lugdunensis* differs in its virulence. How should a clinician approach a clinical culture with *S. lugdunensis*?

Where does *Staphylococcus lugdunensis* typically reside?

S. lugdunensis is a colonizing skin flora found in 30-50% of individuals.² It most frequently colonizes the lower parts of the body, such as inguinal folds, perineal sites, or lower extremities.³

Why is *Staphylococcus lugdunensis* different from other coagulase-negative *Staphylococci*?

Unlike other CoNS, *S. lugdunensis* produces virulence factors, including toxin-like peptides and proteins that bind host coagulation factors. Due to increased adhesion capabilities, this organism is associated with endocarditis, implanted cardiac electrical device infections, skin/soft tissue infections, bone/joint infections, and vascular catheter-related infections.⁴ In a study comparing 30-day mortality among patients with both native and prosthetic valve endocarditis, patients with *S. lugdunensis* endocarditis had a higher rate of mortality than those with other CoNS endocarditis (20% vs. 7%; p=0.016).⁵

Which antimicrobials should be used on this organism?

Treatment options for infections due to *S. lugdunensis* are similar to those of *S. aureus*. However, rates of methicillin resistance in *S. lugdunensis* are typically lower than *S. aureus* and other CoNS. Methicillin resistance in *S. lugdunensis* due to *mecA* gene variants (SCCmec types I, IV, and V) has been found to be between 3-8%.^{6,7} Anti-staphylococcal agents such as cefazolin, vancomycin, and daptomycin can be used for *S. lugdunensis* infections.⁸

Key Takeaway: *Staphylococcus lugdunensis* is more virulent than other coagulase-negative *Staphylococcus* species, and should be clinically managed like *S. aureus*.

References:

1. Heilbronner S, Foster TJ. *Staphylococcus lugdunensis*: a Skin Commensal with Invasive Pathogenic Potential. *Clin Microbiol Rev*. 2020;34(2):e00205-20. doi:10.1128/CMR.00205-20
2. Herry Y, et al. *Staphylococcus lugdunensis* prosthetic joint infection: A multicentric cohort study. *J Infect*. 2022;85(6):652-659. doi:10.1016/j.jinf.2022.10.025
3. Bieber L, Kahlmeter G. *Staphylococcus lugdunensis* in several niches of the normal skin flora. *Clin Microbiol Infect*. 2010;16(4):385-388. doi:10.1111/j.1469-0691.2009.02813.x
4. Zaaroura H, Geffen Y, Bergman R, Avitan-Hersh E. Clinical and microbiological properties of *Staphylococcus lugdunensis* skin infections. *J Dermatol*. 2018;45(8):994-999. doi:10.1111/1346-8138.14496
5. Aldman MH, Rasmussen M, Olaison L, Pählman LI. Endocarditis due to *Staphylococcus lugdunensis*-a retrospective national registry-based study. *Eur J Clin Microbiol Infect Dis*. 2021;40(5):1103-1106. doi:10.1007/s10096-020-04134-w
6. Argemi X, Hansmann Y, Riegel P, Prévost G. Is *Staphylococcus lugdunensis* Significant in Clinical Samples?. *J Clin Microbiol*. 2017;55(11):3167-3174. doi:10.1128/JCM.00846-1
7. Ho PL, et al. Carriage niches and molecular epidemiology of *Staphylococcus lugdunensis* and methicillin-resistant *S. lugdunensis* among patients undergoing long-term renal replacement therapy. *Diagn Microbiol Infect Dis*. 2015;81(2):141-144. doi:10.1016/j.diagmicrobio.2014.10.004
8. Taha L, Stegger M, Söderquist B. *Staphylococcus lugdunensis*: antimicrobial susceptibility and optimal treatment options. *Eur J Clin Microbiol Infect Dis*. 2019;38(8):1449-1455. doi:10.1007/s10096-019-03571-6