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# Biofilm formation of Staphylococcus lugdunensis strains is associated with clonal lineage and dependent on iron availability

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## Background

Staphylococcus lugdunensis is a virulent coagulase-negative staphylococcal species<sup>1</sup>. The ability to form biofilm appears to be one of its main virulence factors<sup>2</sup>, and is promoted by iron-limitation condition<sup>3</sup>. Nevertheless, little is known about biofilm-associated genotypes. In this context, the objective of this study was to compare the impact of iron restriction on fitness and biofilm formation of strains belonging to all clonal complexes (CCs) defined

by Multilocus sequence typing<sup>4</sup> to date.





The study included **49** clinical strains lacksquarefrom different clonal complexes and geographical origins (French and Swedish cities) (Table 1)

Clonal complex	<b>CC1</b>	CC2	CC3	CC4	CC5	CC6	<b>CC7</b>	Singleton*	_
Number of strains	11	5	11	4	5	5	4	4	
Geographical origins	Strasbourg, Rouen, Tours, Kronoberg, Nancy	Strasbourg, Montpellier, Nantes, Nancy, Rouen	Rouen, Strasbourg	Strasbourg, Rouen, Bordeaux	Strasbourg, Bordeaux, Rouen	Kronoberg, Nantes, Strasbourg, Rouen	Strasbourg, Rouen	Rouen, Strasbourg	

**Table 1:** S. lugdunensis strains categorized by clonal complexes and geographical origins. \*Singleton Sequence Type (ST)

## **\*** Fitness

- kinetics Growth 24 ulletover hours performed in triplicates for all strains (Tecan spark<sup>®</sup>):
  - ✓ In **rich media** (Tryptic Soy Broth [TSB])
  - ✓ In iron-restricted condition

(RPMI 1640 with 1% casamino acids)

✓ Determination of generation time (G=ln2/ $\mu$ ) and **maximal growth rate**  $(\mu)$  (Figure 1)



## Biofilm formation



*microtiter plates* 

- 24h biofilm formation studied using crystal violet staining method on 96well polystyrene microtiter plates in quadruplicates:
  - ✓ In **rich media** (TSB with 1% glucose) for all strains
  - ✓ In **iron-restricted condition** for 34 strains (Figure 2)
- ✓ Using *S. epidermidis* ATCC 35984 as biofilm positive control



Results

Figure 3: Biofilm formation of 49 S. lugdunensis strains after a 24h incubation at 37°C in rich medium (TSB + 1% glucose) according to the CC (OD mean values). Error bars represent the standard deviation within each CC.

- All strains were biofilm producers (DO<sub>590nm</sub> > 0,1) in rich medium after 24h
- **CC2** strains produced significantly more biofilm than CC4 (*p*= 0.0203), CC5 (p=0.0215) and singleton STs (p=0.0045) strains (Figure 3)

Figure 4: Biofilm formation of 34 S. lugdunensis strains after a 24h incubation at 37°C in rich medium (TSB + 1% glucose [full bars]) or iron-restricted medium (RPMI + 1% casamino acids [hatched bars]) according to the CC (OD mean values). Error bars represent the standard deviation within each CC.

- In iron-restricted conditions, CC1 (p= 0.0079), CC2 (p= 0.0079) and CC3 (p= 0.026) strains produced significantly less biofilm than in rich medium
- On the contrary, CC6 (p= 0.0079) strains were significantly more productive (Figure 4)

### Conclusions

- This study shows for the first time that the **biofilm-forming ability** of *S. lugdunensis* strains is strongly **correlated with CC affiliation**.
- In addition, two different **CC-dependent behaviors** are observed in an **iron-restricted environment**. These results are to be confirmed with a larger number of strains.

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