

Towards an integrated approach for the annotation of ICEs and IMEs in bacterial genomes

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Julie Lao, Gérard Guédon, Thomas Lacroix, Nathalie N. Leblond-Bourget, Helene H. Chiapello. Towards an integrated approach for the annotation of ICEs and IMEs in bacterial genomes. JOBIM 2018, Jul 2018, Marseille, France. hal-01829606

HAL Id: hal-01829606 https://hal.univ-lorraine.fr/hal-01829606

Submitted on 4 Jul 2018

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Towards an integrated approach for the annotation of ICEs and IMEs in bacterial genomes

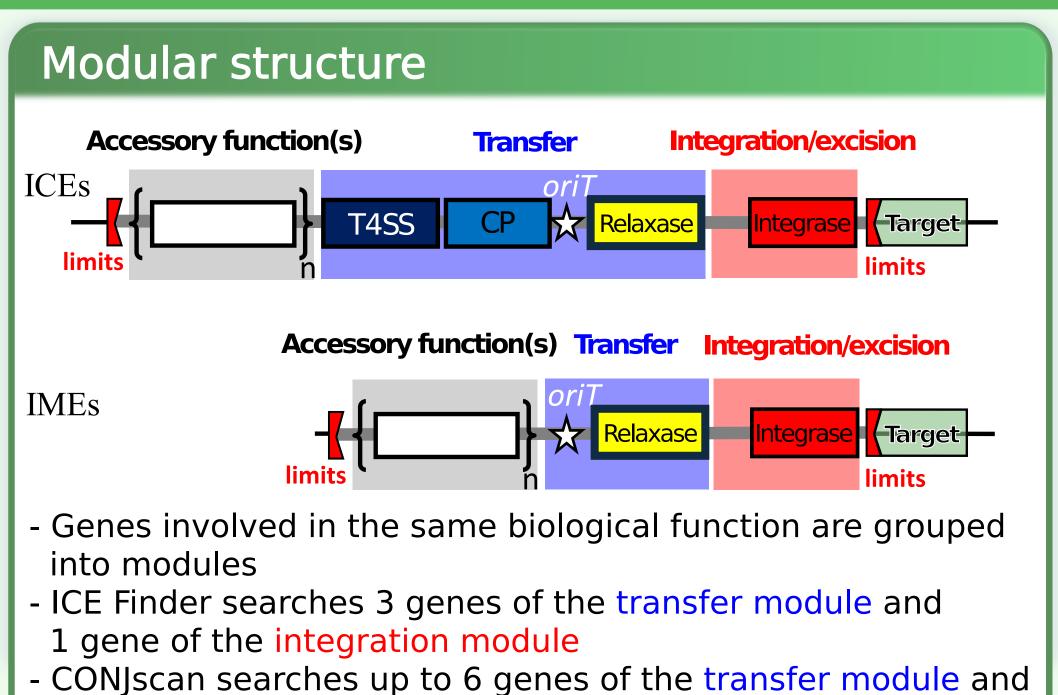
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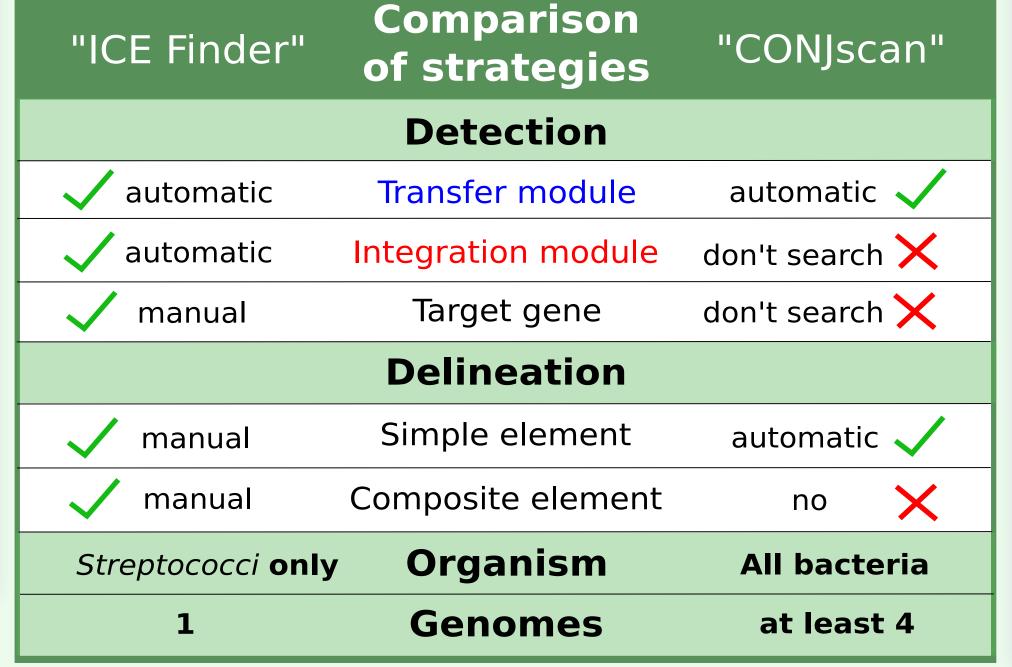
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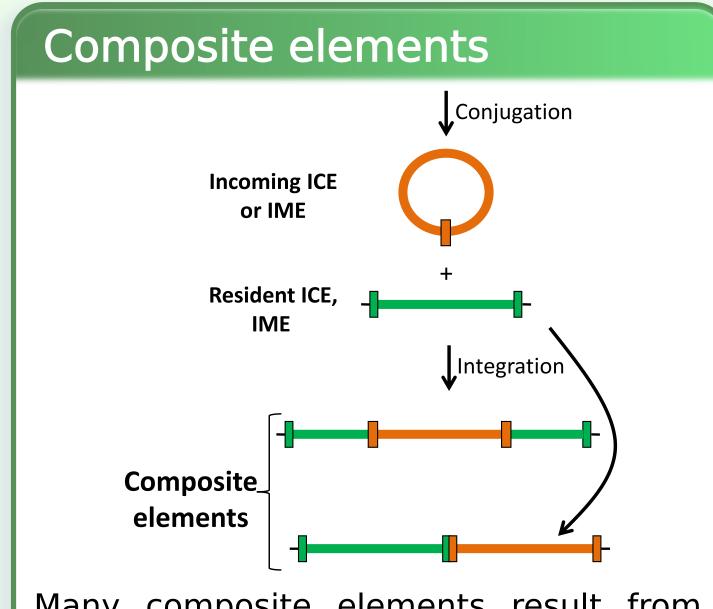
Abstract

- Integrative Conjugative Elements (ICEs) are mobile genetic elements (MGE) that are integrated in bacterial chromosomes which encode their own transfer by conjugation.
- Integrative Mobilizable Elements (IMEs) are another type of bacterial integrated MGE, they lack a full conjugation system and hijack co-resident conjugative elements to promote their own transfer.
- ICEs and IMES are poorly known and annotated but play a key role in bacterial genome evolution and the very few systematic investigations of these MGEs in bacterial genomes suggest that they have a very high prevalence.
- Dedicated bioinformatics approaches are needed for their identification and precise delineation.

In this poster, we present the first results from the comparison of the two currently existing strategies for the detection and delineation of ICEs: "ICE Finder" strategy^a and "CONJscan" strategy^b







Many composite elements result from the insertion of an incoming element into another or into one of its ends

"ICE Finder" strategy

- Automatic detection of "signature" proteins of the transfer and integration modules using **BlastP**. A manually cured database of known elements in *streptococci*^a is used.

don't search the integration module

- **Manually annotate and delineate** *simple* and composite elements at the nucleotide **level** through the research of the *target gene*.
- In many cases, the target gene contains one of the definite boundaries of the element.

"CONJscan" strategy

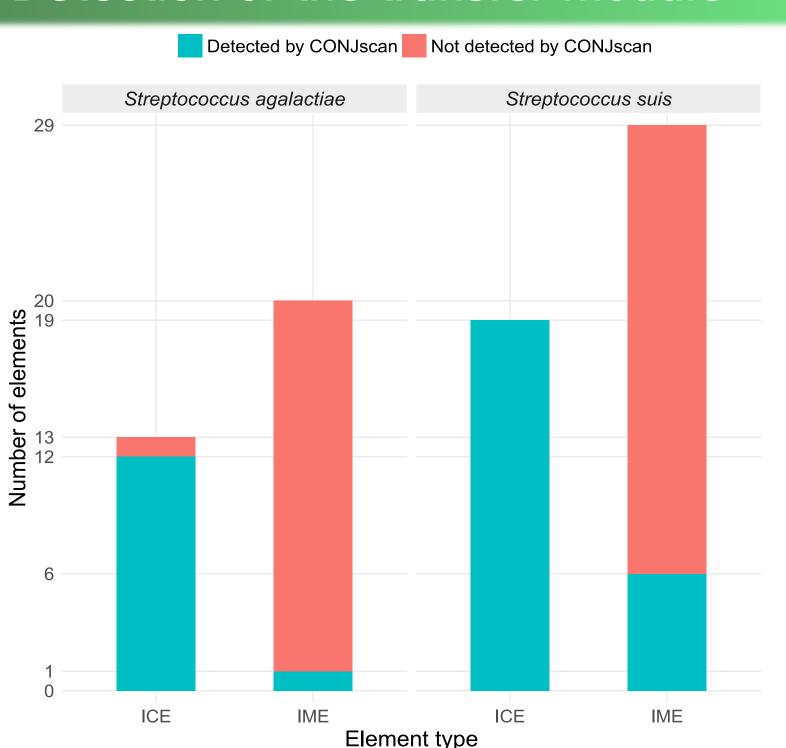
- It is a pipeline that delineates ICEs only in all bacterial genomes by using the core genes that surrounds them^{b,c}.
- This procedure is based on the detection of one module only (transfer module) using the CONJscan module of MacSyFinder software^{d,e}.
- It needs at least 4 different closely-related genomes to enable ICE annotation.

Comparison design

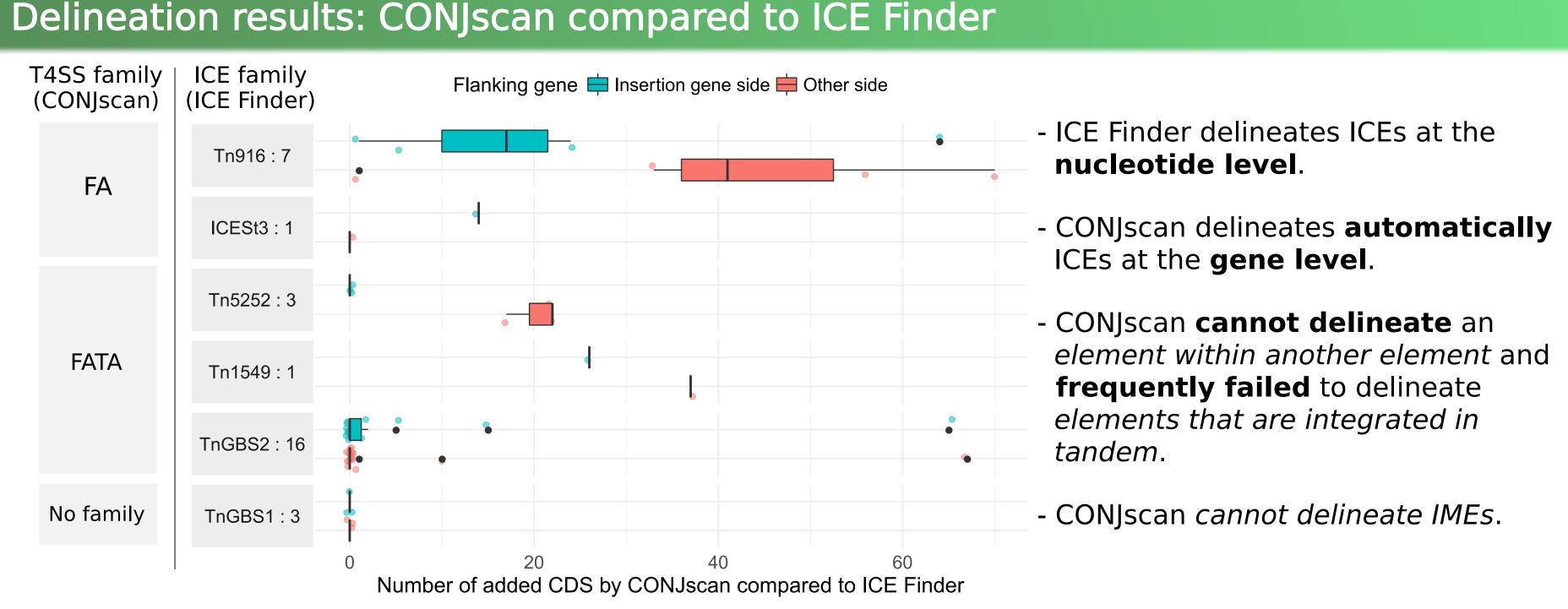
Results obtained with "ICE Finder" strategy are the reference

- **Dataset**: 15 genomes of *Streptococcus suis* and 7 genomes of Streptococcus agalactiae
- **Detection of transfer module by "CONJscan" strategy** is assessed: Is T4SS families of CONJscan consistent with ICE families of ICE Finder?
- Is delineation of ICEs with "CONJscan" strategy and "ICE Finder" strategy similar?

Detection of the transfer module



- ICE Finder detects the transfer module of all known ICEs and IMEs of Streptococci.
- CONJscan detects the transfer module of most ICEs and of some IMEs. It does not have dedicated method for IME detection.



Integration module and target gene

- -"CONJscan" strategy cannot find the target gene because dedicated research of the insertion module is lacking.
- Unlike CONJscan, "ICE Finder" strategy search specifically the integration module (by BlastP of the integrase protein).







Conclusion

- CONJscan has a more specific definition of the T4SS.
- CONJscan does not specifically detect the integration module and does not detect most of transfer module from IMEs in *Streptococci*.
- Boundaries obtained with "CONJscan" strategy are not as precise as the ones obtained with "ICE Finder" strategy. But CONJscan always delineate elements with transfer module.

New strategy proposed:

- 1) Detection of transfer module AND integration module with MacSyFinder^d:
- Development of a mixed model "CONJscan + Integrase" including more HMM profiles of transfer protein (of IMEs) that are not presently searched by CONJscan.
- 2) Get first approximate delineation with "CONJscan" strategy.
- 3) **Refine delineation** with the "ICE Finder" strategy.

References

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