

Isolation of *Sporisorium scitamineum* wild-type for the production of glycolipid biosurfactants from hydrophilic carbon sources

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Background – Sporisorium scitamineum

- <u>Sporisorium scitamineum</u> is a fungus which systemically infects sugarcane tissue.
- Sugarcane smut is one of the <u>main diseases of sugarcane</u>, causing significant losses in productivity of this perennial crop.
- However, it has been observed that <u>S. scitamineum excretes useful products</u>, including different <u>glycolipid biosurfactants</u>.
- Traditionally, vegetable oils are required for glycolipid production. However, vegetable oils are expensive and complicates the downstream processing.
- S. scitamineum is well adapted to growth on sugars, the main carbon source available in sugarcane tissue.
- Therefore, this project investigated the production of glycolipid biosurfactants by cultivating *S. scitamineum* on <u>various sugars</u>.

Methodology

- Drop spores from infected sugarcane samples on a specifically formulated agar solution.
- Sub-culture fungal growth spots.

Isolate

Grow

- Verify the isolation by sequencing the genes of the isolated fungus.
- Grow S. scitamineum wild-type on various pure hydrophilic carbon sources.
- Grow S. scitamineum on sugarcane molasses.

• Investigate the production of MELs from the various hydrophilic carbon source-

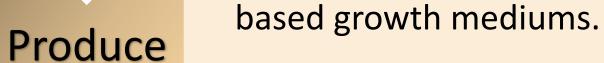


Figure 1: Life cycle of <u>Sporisorium scitamineum</u> (Taniguti et al., 2015)

Background – Glycolipid biosurfactants

Mannosylerythritol lipids:

- MELs possess a hydrophilic moiety, consisting of 4-O- β -D-mannopyranosyl-Derythritol, attached to various fatty acids.
- MELs are ideal for cosmetic applications.



Results

Isolation:

- S. scitamineum wild-type was successfully isolate from sugarcane samples provided by SASRI.
- The genes were amplified with a MyTaq Plant PCR kit. and
- The amplified genes were sequenced and imported into the Basic Local Alignment Search Tool (BLAST) to verify the presence of *S. scitamineum*.



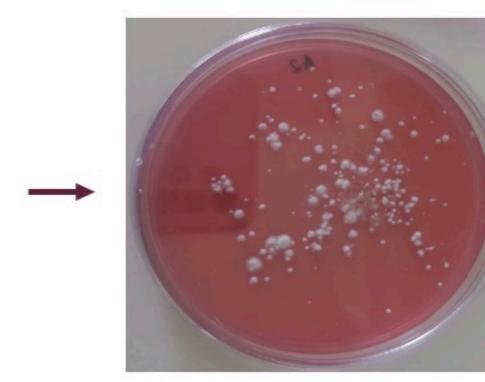




Figure 3: Experimental procedure implemented for the isolation of <u>*S. scitamineum wild-type*</u>

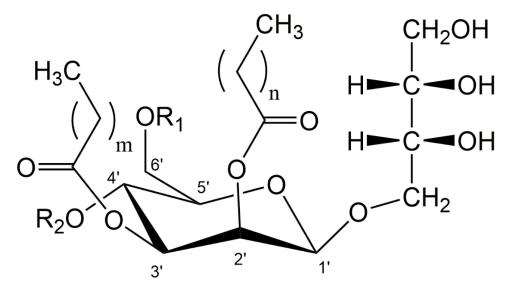


Figure 2: General structure of MELs (Beck et al., 2020)

Cellobiose lipids:

- CBLs possess a cellobiose moiety, consisting of two glycosidically linked glucose molecules, attached to a fatty acid chain.
- CBLs have excellent antifungal properties.

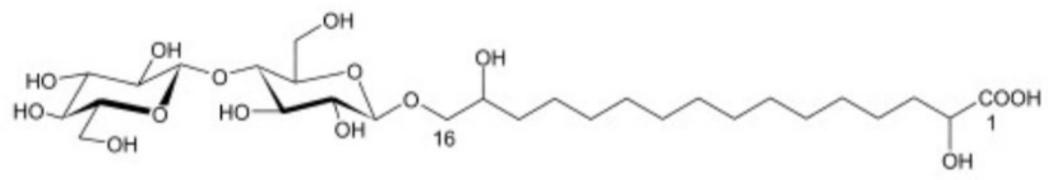
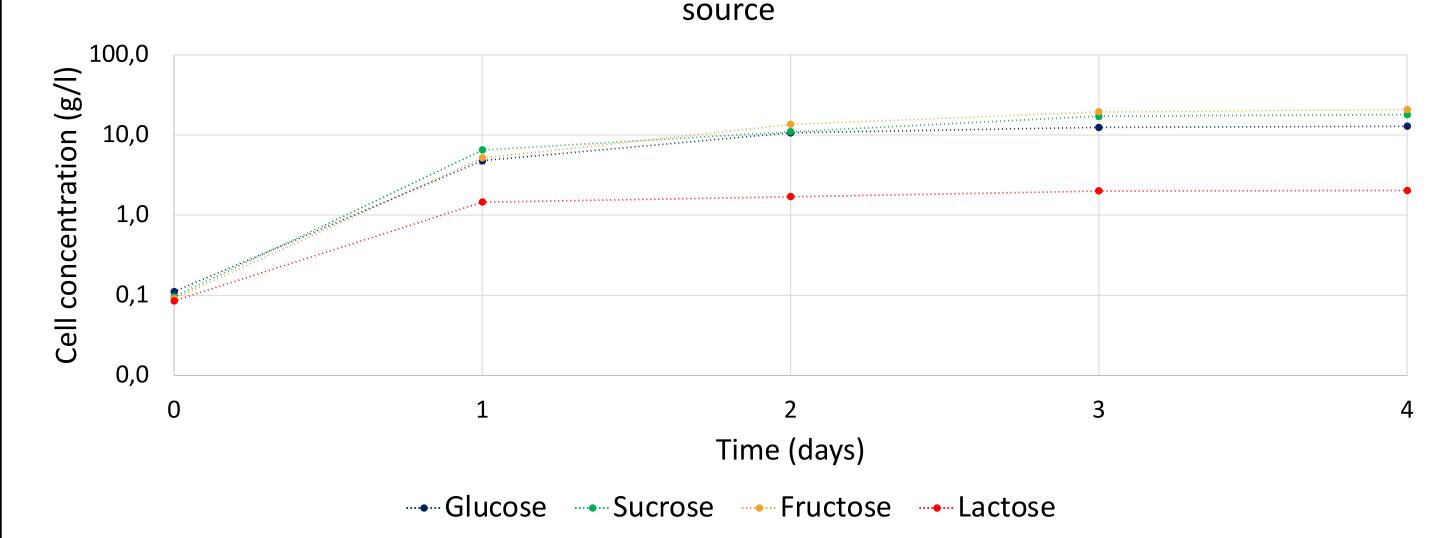


Figure 3: General structure of CBLs (Oraby et al., 2020)

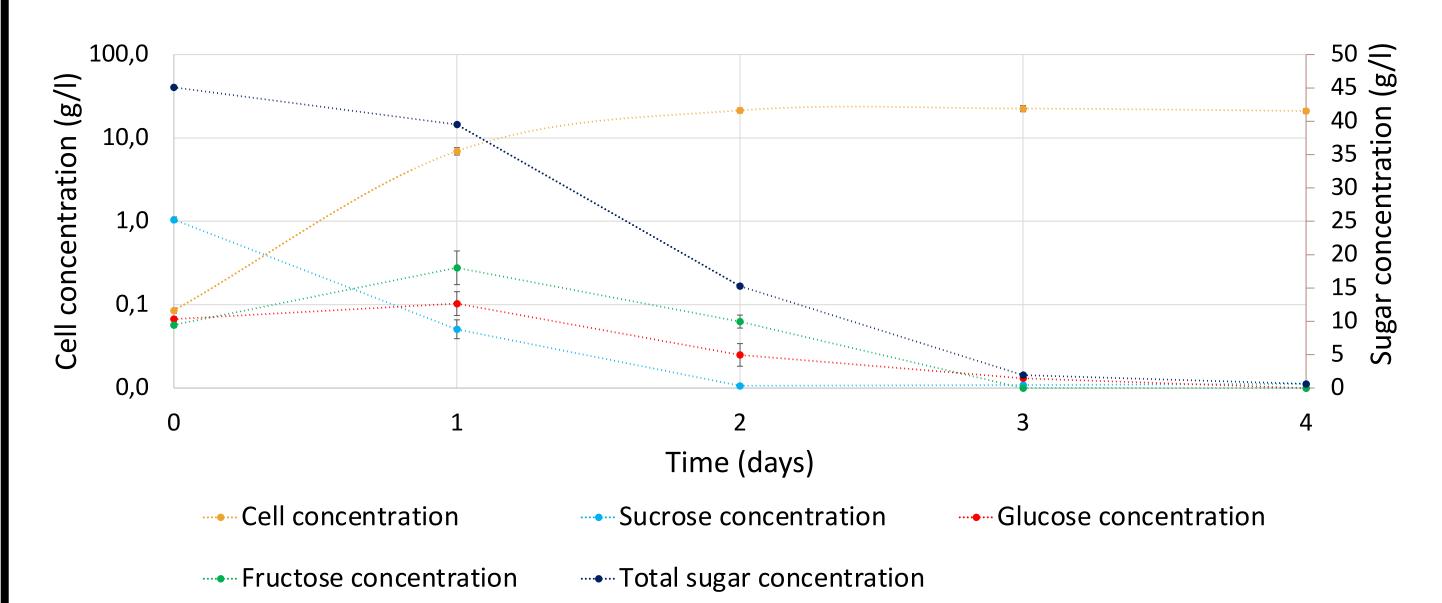
Aim & Objectives Aim: • Investigate the potential of producing glycolipid biosurfactants by cultivating a wild strain of *S. scitamineum* on hydrophilic carbon sources. Objectives:

Growth:

- S. scitamineum grew well on glucose, fructose and sucrose. Lactose didn't perform well.
- Subsequently, S. scitamineum grew well on sugarcane molasses
 S. scitamineum growth with the utilization of different sugars as the carbon



S. Scitamineum growth with sugarcane molasses as the sole carbon source



- Isolate a wild strain of *S. scitamineum* from infected sugarcane samples provided by the South African Sugarcane Research Institute (SASRI).
- Investigate the growth of S. scitamineum on pure hydrophilic carbon sources, as well as sugarcane molasses.
- Investigate the production of glycolipids from the hydrophilic carbon sources.

References

- Beck, A., & Zibek, S. (2020). Growth Behavior of Selected Ustilaginaceae Fungi Used for Mannosylerythritol Lipid (MEL) Biosurfactant Production Evaluation of a Defined Culture Medium. *Frontiers in Bioengineering and Biotechnology*, *8*, 555280. <u>https://doi.org/10.3389/FBIOE.2020.555280/BIBTEX</u>
- Oraby, A., Werner, N., Sungur, Z., & Zibek, S. (2020). Factors Affecting the Synthesis of Cellobiose Lipids by
 Sporisorium scitamineum. *Frontiers in Bioengineering and Biotechnology*, 8.

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• Taniguti, L.M. et al. (2015) 'Complete genome sequence of sporisorium scitamineum and biotrophic interaction transcriptome with sugarcane', PLOS ONE, 10(6). <u>https://doi.org/10.1371/journal.pone.0129318</u>

Production:

- So far, significant glycolipid production (< 0.5 g/l) could not be achieved on production mediums reported in literature.
- We are currently exploring different medium compositions and operating conditions.