

Genome-Guided strategy to access Burkholderia diversity from environment and its chemistry.

João Luiz Baldim Zanin (PG)^{1,2*}, Emerson Glassey (PG)², Jessica Ochoa (PG)², Weng Ruh Wong (PG)², Roger G. Linington (PQ)², Marisi G. Soares (PG)¹.

jotaelebaldim@gmail.com

¹Chemistry institute of Federal University of Alfenas, Alfenas, Minas Gerais, Brazil²

²Department of Chemistry and Biochemistry, University of California, Santa Cruz

Palavras Chave: *Burkholderia*, Genome, Isolation Method, Chemical Prediction.

Introduction

According to recent discoveries, the genus *Burkholderia*, being part of the phylum Proteobacteria, has been recognized to contain a substantial number of clusters encoding putatively novel natural products, but to date has not been well studied.¹ Since there are few efficient methods to isolate *Burkholderia* species from environmental sources, this project aims to develop a new isolation method using genomic, metagenomic and metabolic information to take advantage of unique properties of *Burkholderia* metabolism. This approach relies on examination of important metabolic properties, including antibiotic resistance, metal resistance, as well as the carbon and nitrogen metabolism profiles from *Burkholderia* and untargeted species in bacterial communities.

Thus, this strategy can generate the possibility to construct differential culture media to access this genus in diverse types of environmental samples. Ultimately, the objective of this research program is to examine environmental samples of the genus *Burkholderia* in order to discover next-generation antibiotics, guided by chemical predictions, with unique molecular architectures and biological modes of action for the treatment of drug resistant infections.

Results

All experiments have demonstrated that this method is reliable for specific isolation for the genus *Burkholderia*. The results presented 100% of efficacy when the method was complete (Table 1). Once there is the possibility of using bioinformatics tools guiding the secondary metabolites production, all the genomic predictions were created for future isolation and identification of compounds.² The structure's core and its amino acids; nrps sequences were found (Figure 1) and these results show all the sources of compounds that possibly can be produced by the biosynthetic gene clusters in *Burkholderia* species.

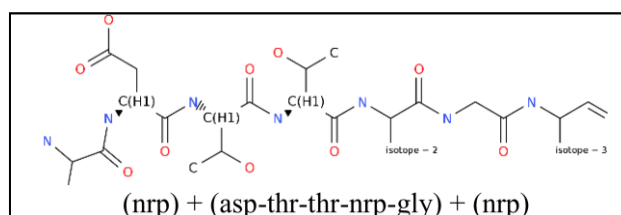


Figure 1. One example of core structure predicted for *Burkholderia* sp. CCGE1003 from Nrps biosynthetic cluster and its subunits sequence.

Table 1. Species isolated by using the genome-guide method from one environmental spot.

SAMPLE NAME	SPECIES
RL14-064-BSI-A	<i>Burkholderia gladioli</i> strain CACua-73 16S
RL14-064-BSJ-A	<i>Burkholderia</i> sp. FSGSA12 16S ribosomal
RL14-065-BSA-A	<i>Burkholderia multivorans</i> ATCC 17616
RL14-065-BSB-A	<i>Burkholderia gladioli</i> strain 2002721590
RL14-065-BSC-A	<i>Burkholderia gladioli</i> strain 2002721590
RL14-065-BSD-A	<i>Burkholderia</i> sp. FSGSA12 16S

Conclusions

All the results demonstrated that this genome-guided method is efficient for isolating microorganisms of chemical interest.

Agradecimentos

We thank other members of Linington Lab; S. Lokey and W. Bray for biological screening facilities; The fellowship from CAPES-Brasil (Process number 8074135). Unifal-MG. This work was funded by NIH-TW006634.

1.Cimermancic, P. *et al.* Insights into Secondary Metabolism from a Global Analysis of Prokaryotic Biosynthetic Gene Clusters. *Cell* **158**, 412–421 (2014).

2.Medema, M. H. *et al.* antiSMASH: rapid identification, annotation and analysis of secondary metabolite biosynthesis gene clusters in bacterial and fungal genome sequences. *Nucleic Acids Res.* **39**, W339–46 (2011).