

# **MScCBBi**

MASTER IN
COMPUTATIONAL BIOLOGY
& BIOINFORMATICS

**SPECIALIZATION** Multi-Omics for Life and Health Sciences

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## Benchmarking Causal Reasoning Algorithms for enhanced Drug Discovery

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# BENCHMARKING CAUSAL REASONING ALGORITHMS FOR ENHANCED DRUG DISCOVERY

SOME THOUGHTS ON THE LIFE, THE UNIVERSE, AND EVERYTHING ELSE

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#### ABSTRACT

Regardless of the language in which the dissertation is written, usually there are at least two abstracts: one abstract in the same language as the main text, and another abstract in some other language.

However, this order can be customized by adding one of the following to the file 5\_packages.tex.

```
\ntsetup{abstractorder={<LANG_1>,...,<LANG_N>}}
\ntsetup{abstractorder={<MAIN_LANG>={<LANG_1>,...,<LANG_N>}}}
```

For example, for a main document written in German with abstracts written in German, English and Italian (by this order) use:

```
\ntsetup{abstractorder={de={de,en,it}}}}
```

Concerning its contents, the abstracts should not exceed one page and may answer the following questions (it is essential to adapt to the usual practices of your scientific area):

- 1. What is the problem?
- 2. Why is this problem interesting/challenging?
- 3. What is the proposed approach/solution/contribution?
- 4. What results (implications/consequences) from the solution?

**Keywords:** One keyword, Another keyword, Yet another keyword, One keyword more, The last keyword

#### Resumo

Independentemente da língua em que a dissertação está escrita, geralmente esta contém pelo menos dois resumos: um resumo na mesma língua do texto principal e outro resumo numa outra língua. No entanto, esse pedido pode ser personalizado adicionando um dos seguintes ao arquivo 5\_packages.tex.

```
\abstractorder(<MAIN_LANG>):={<LANG_1>,...,<LANG_N>}
```

Por exemplo, para um documento escrito em Alemão com resumos em Alemão, Inglês e Italiano (por esta ordem), pode usar-se:

```
\ntsetup{abstractorder={de={de,en,it}}}}
```

Relativamente ao seu conteúdo, os resumos não devem ultrapassar uma página e frequentemente tentam responder às seguintes questões (é imprescindível a adaptação às práticas habituais da sua área científica):

- 1. Qual é o problema?
- 2. Porque é que é um problema interessante/desafiante?
- 3. Qual é a proposta de abordagem/solução?
- 4. Quais são as consequências/resultados da solução proposta?

**Palavras-chave:** Primeira palavra-chave, Outra palavra-chave, Mais uma palavra-chave, A última palavra-chave

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### GLOSSARY

CBDD

Computational Biology for Drug Discovery - citation [1] (p. 6)

computer

An electronic device which is capable of receiving information (data) in a particular form and of performing a sequence of operations in accordance with a predetermined but variable set of procedural instructions (program) to produce a result in the form of information or signals. This is a test that adds a citation [1] to the glossary! (*p.* 6)

### ACRONYMS

aaa acronym aaa (p. 6)aab acronym aab (p. 6)aba acronym aba (p. 6)

**abbrev** abbreviation of a longer text (p. 6)

**bbb** acronym bbb (*p. 6*)

xpto xpto xpto xpto xpto xpto (p. 6)

### Symbols

- μ Mu (p. 6)
- $\pi$  the numerical value of pi (p. 6)
- *r* the radius of a circle (*p. 6*)

#### Introduction

This section expounds the underlying motivation, rationale and goals for the study, emphasizing its significance in the field. It provides context by giving some background on the supporting company and the initiative. Furthermore, it outlies a read's guide of this thesis.

#### 1.1 Motivation and Goals

#### 1.2 Scope

This project was conducted within the framework of the Algorithm Benchmarking Consortium (ABC), a subscription-based initiative led by Clarivate for pharmaceutical companies. ABC is dedicated to evaluating a wide range of computational tools for a variety of applications in the life sciences and healthcare field. The topic for this thesis is the development of the ABC's tenth use case – Causal Regulation – which focuses on benchmark and identify the most optimal tools tailored to specific needs within the drug discovery process by identifying key regulators from transcriptomics data and prior knowledge graphs.

#### 1.3 Parallel Contributions

This study expands the state of art in causal reasoning using gene expression data and causal graphs by presenting a robust framework and methods for benchmarking various algorithms designed for this purpose. Beyond this primary focus, several parallel projects with real-world challenges and novel data were developed, and enriched the consultant experience allowing for an expansion in the expertise of bioinformatics. The parallel projects, spanning various domains of computational biology, include:

**Skin Microbiome Atlas** The skin microbiome atlas project involved extensive scientific literature review and dataset curation, followed by a systematic re-analysis of available datasets to ensure consistency and reliability. This was done by pre-processing the raw sequencing data (. . . understand how deep I can go In terms of details – confidentiality issues)

More projects ... More projects ...

#### 1.4 Structure

This study is organized in ? chapters. Chapter 2 introduces

### LITERATURE REVIEW

This section details the

- 2.1 Drug discovery: the importance of the compound's mechanism of action
- 2.2 Causal Reasoning algorithms as an approach for target inference

### Materials and Methods

3.1 Gene expression data

# Adding Support to a New School (work in progress)

The directory uminho contains the customization for all Schools of Universidade do Minho. This university is an example of the case where the regulations are defined at University level and all the schools apply the same thesis layout and organization. So, the all the customization is done in the file uminho/uminho-defaults.ldf, except the definition of the name and logo of each individual school.

As another example, the directory nova contains the customization for all Schools from NOVA University Lisbon. This university grants a lot of freedom in the definition of the thesis layouts. In some cases, they are defined at the School level (e.g., NOVA FCT), while is some other cases they are defined separately for each degree (e.g., NOVA IMS).

- 1. Try all the already supported schools and check which one is closer to your needs;
  - a) Edit Config/1\_novathesis.tex and near line 28 uncomment the line with key \ntsetup{school=<SOMETHING>};
  - b) For each school supported (see the comment), replace <SOMETHING> with the school name, e.g., make it \ntsetup{school=ulisboa/fmv}
  - c) Recompile and check the document. Particularly, check the cover layout, the front-page (second cover) layout, the front-matter contents, the bibliography style;
  - d) Repeat for the next school, until you find one close enough.

2. ...

**CBDD** 

### REFERENCES

- [1] CBDD. Web Page. URL: https://clarivate.com/life-sciences-healthcare/consulting-services/research-and-development-consulting/cbdd/(cit. on p. vi).
- [2] R. J. Dias et al. "Verification of Snapshot Isolation in Transactional Memory Java Programs". In: *Proceedings of the 26th European conference on Object-oriented programming* (*ECOOP'12*). Springer-Verlag, 2012-06 (cit. on p. 10).

A

### NOVATHESIS COVERS SHOWCASE

This Appendix shows examples of covers for some of the supported Schools. When the Schools have very similar covers (e.g., all the schools from Universidade do Minho), just one cover is shown. If the covers for MSc dissertations and PhD thesis are considerable different (e.g., for FCT-NOVA and UMinho), then both are shown.

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### Appendix 2 Lorem Ipsum

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#### Annex 1 Lorem Ipsum

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