Enhancing the evolutionary robustness of probiotics via experimental evolution FWO DMP (Flemish Standard DMP)

1. Research Data Summary

List and describe all datasets or research materials that you plan to generate/collect or reuse during your research project. For each dataset or data type (observational, experimental etc.), provide a short name & description (sufficient for yourself to know what data it is about), indicate whether the data are newly generated/collected or reused, digital or physical, also indicate the type of the data (the kind of content), its technical format (file extension), and an estimate of the upper limit of the volume of the data.

| Dataset Name Description New or reused Digital or Physical Digital Data Type Please Choose from the following options: Please choose from the following options: Please choose from the following options: Observational Experimental Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the following options: Other Simulation data Simulation data Simulation data Software Choose from the foll | | | | | | Only for | Only for | Only for |
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| Please choose from the following options: Please choose from the following options: Observational Experimental Read-out for all the bacterial competition glow cytometery for all the bacterial competition gounts via Generate new data of bacterial optical density population measurements of bacterial sequencing Please choose from the following options: Observational Experimental Observational Competition data Software Softwar | Dataset Name | Description | New or reused | | Digital Data Type | | volume | Physical volume |
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| | Evolved bacteria | bacterial populations and isolated clones from the evolved | Generate new data | Physical | Experimental | | | samples, stored in 96-well plate |
| | | | | | | | | |
| | | | | | | | | |

If you reuse existing data, please specify the source, preferably by using a persistent identifier (e.g. DOI, Handle, URL etc.) per dataset or data type:

We will make use of a published reference genomes of *L. rhamnosus* GG (NCBI RefSeq assembly GCF_000011045.1), *L. reuteri* RC-14 (NCBI RefSeq assembly GCF_002762415.1), and *L. fermentum* (NCBI RefSeq assembly GCF_029961225.1) to map our whole genome and RNA sequencing data.

Are there any ethical issues concerning the creation and/or use of the data (e.g. experiments on humans or animals, dual use)? Describe these issues in the comment section. Please refer to specific datasets or data types when appropriate.

No

Will you process personal data? If so, briefly describe the kind of personal data you will use in the comment section. Please refer to specific datasets or data types when appropriate.

No

Does your work have potential for commercial valorization (e.g. tech transfer, for example spin-offs, commercial exploitation, ...)? If so, please comment per dataset or data type where appropriate.

Yes

Both the improved probiotic strains and the experimental evolution platform will be valorized. IP will be protected by copyright and patent applications.

Do existing 3rd party agreements restrict exploitation or dissemination of the data you (re)use (e.g. Material/Data transfer agreements/ research collaboration agreements)? If so, please explain in the comment section to what data they relate and what restrictions are in place.

No

Are there any other legal issues, such as intellectual property rights and ownership, to be managed related to the data you (re)use? If so, please explain in the comment section to what data they relate and which restrictions will be asserted.

• No

We have FTO since patents covering the use of Lacticaseibacillus in the envisioned application are either expired or aborted.

2. Documentation and Metadata

Clearly describe what approach will be followed to capture the accompanying information necessary to keep data understandable and usable, for yourself and others, now and in the future (e.g., in terms of documentation levels and types required, procedures used, Electronic Lab Notebooks, README.txt files, Codebook.tsv etc. where this information is recorded).

1. The data will be accompanied by a detailed metadata txt file denoting important characteristics (e.g. strain, material, time point,..) necessary for interpretation of the results. The key characteristics will also be denoted in the filename of the data files.

- 2. A detailed experimental protocol will be added to the directory of the corresponding experimental results. This step-wise description will facilitate potential future reproduction of the experiments.
- 3. For every deliverable in the project, a general outline txt file will be created. This file provides an overview of all the available data, the design of the experiment and the structure of the data saving.

Will a metadata standard be used to make it easier to find and reuse the data? If so, please specify (where appropriate per dataset or data type) which metadata standard will be used. If not, please specify (where appropriate per dataset or data type) which metadata will be created to make the data easier to find and reuse.

Yes

Sequencing data: MIxS metadata standard

Flow cytometry data: MIFlowCyt metadata standard

A metadata template will be constructed in the frame of the project for data types where no general metadata standard is available.

3. Data storage & back-up during the research project

Where will the data be stored?

All digital data will be registered using an electronic lab notebook (ELN) based on the Sharepoint platform and backed up on the internal server of KU Leuven, which is maintained by the IT service of KU Leuven. Due to the size of the raw sequencing data, this data will be exempt from this rule and only be stored on an archive network, managed by KU Leuven.

Bacterial strains and populations will be stored in a secured -80°C freezer at the facility.

How will the data be backed up?

The Sharepoint is backed up three times a day. The internal servers are managed by the KU Leuven IT department and backed-up according to their procedures.

Is there currently sufficient storage & backup capacity during the project? If yes, specify concisely. If no or insufficient storage or backup capacities are available, then explain how this will be taken care of.

Yes

How will you ensure that the data are securely stored and not accessed or modified by unauthorized persons?

- 1. Access to the digital data will be limited to members of the research group who contribute to the research.
- 2. Critical documents can be (temporarily) locked by the author(s).
- 3. Sharepoint provides a changelog for detecting and reverting possible unauthorized changes.
- 4. The internal storage provides a back-up for the sharepoint and vice versa.
- 5. Physical data is stored in a secured -80°C freezer at the facility with limited-access.

What are the expected costs for data storage and backup during the research project? How will these costs be covered?

The total cost of storage are estimated on ~€1000-€1500 per year, mainly for the raw sequencing data (~1TB). The costs will be covered by the budget for consumables.

4. Data preservation after the end of the research project

Which data will be retained for at least five years (or longer, in agreement with other retention policies that are applicable) after the end of the project? In case some data cannot be preserved, clearly state the reasons for this (e.g. legal or contractual restrictions, storage/budget issues, institutional policies...).

All of the generated data will be stored for minimum 10 years after the end of the project.

Where will these data be archived (stored and curated for the long-term)?

All data will be archived on the archive network (K:) drives provided by the KU Leuven.

What are the expected costs for data preservation during the expected retention period? How will these costs be covered?

In light of the size of the sequencing data (\sim 1TB), the current storage volume of the archive drive needs to be increased. The costs for an expansion of 1 TB for 10 years are estimated on \leq 2,000, and will be covered by the allocated project budget.

5. Data sharing and reuse

Will the data (or part of the data) be made available for reuse after/during the project? In the comment section please explain per dataset or data type which data will be made available.

· Yes, in an Open Access repository

All digital data will be made available upon publication.

If access is restricted, please specify who will be able to access the data and under what conditions.

n/a

Are there any factors that restrict or prevent the sharing of (some of) the data (e.g. as defined in an agreement with a 3rd party, legal restrictions)? Please explain in the comment section per dataset or data type where appropriate.

· Yes, Intellectual Property Rights

The evolved bacterial strains will not be made available in order to protect the valorisation potential of these strains.

Where will the data be made available? If already known, please provide a repository per dataset or data type.

The sequencing data will be deposited International Nucleotide Sequence Database Collaboration (INSDC).

When will the data be made available?

The data will be made available upon publication of the corresponding research results.

Which data usage licenses are you going to provide? If none, please explain why.

We will make use of open licenses such as CC-BY or their equivalent as our data is not considered sensitive and open licenses are often required for high impact publications.

Do you intend to add a PID/DOI/accession number to your dataset(s)? If already available, you have the option to provide it in the comment section.

Yes

What are the expected costs for data sharing? How will these costs be covered?

No costs are expected, the cost will be paid by the researchers requesting the material.

6. Responsibilities

Who will manage data documentation and metadata during the research project?

The researchers who generate the data are responsible for the correct documentation of the data and metadata.

Who will manage data storage and backup during the research project?

The researchers who generate the data are responsible for the storage and back-up, supervised by Hans Steenackers

Who will manage data preservation and sharing?

Hans Steenackers has the end responsibility and manages long term preservation and sharing.

Who will update and implement this DMP?

Hans Steenackers