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# **coralME Documentation**

***Release 1.0***

**Juan D. Tibocha Bonilla, Rodrigo Santibanez-Palominos**

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## DESCRIPTION

The **C**omprehensive **R**econstruction **A**lgorithm for **ME**-models (**coralME**) is an automatic pipeline for the reconstruction of ME-models. coralME integrates existing ME-modeling packages [COBRAme](#), [ECOLIme](#), and [solveME](#), generalizes their functions for implementation on any prokaryote, and processes readily available organism-specific inputs for the automatic generation of a working ME-model.

coralME has four main objectives:

1. **Synchronize** input files to remove contradictory entries.
2. **Complement** input files from homology with a template organism to complete the E-matrix.
3. **Build** a working ME-model
4. **Inform** the user about necessary steps to curate the ME-model.

This resource is intended to:

1. Describe basic inputs required for ME-model reconstruction.
2. Describe the architecture of coralME.
3. Demonstrate how to build a ME-model with coralME.
4. Describe how to perform manual curation guided by coralME's curation notes.

## 2.1 Getting started

For details on inputs go to [Description of Inputs](#).

For information about coralME architecture go to [Architecture of coralME](#).

### 2.1.1 Download files from BioCyc

BioCyc files are optional but useful, you should download them after having your gene id consistent M-model and genbank files.

The quickest way to do this is to copy one of the genes from the genbank file into the BioCyc search bar. Your organism should appear in the list if it is available in BioCyc.

To download:

Go to **Tools>Special SmartTables**

The screenshot shows the BioCyc website interface. The top navigation bar includes 'Tools', 'Sites', 'Pathway Tools', and 'Help'. The 'Tools' menu is expanded, showing four main categories: Search, Genome, Metabolism, and Analysis. Under the 'Genome' category, the 'SmartTables' sub-menu is expanded, and 'Special SmartTables' is highlighted with a red box. Below the navigation menu, there is a table titled 'Organism or Sample Properties' with the following data:

Organism or Sample Properties	
Environment:	soil plants
Collection Date:	1981
Relationship to Oxygen:	aerobe
Temperature Range:	mesophile
NCBI Genome Type:	reference

From here you can download the 5 optional BioCyc files for your organism.

**Special SmartTables Directory**

**Welcome to SmartTables**

A SmartTable is a collection of BioCyc objects, such as genes or metabolites, together with associated data, that can be created, edited, manipulated, and shared on the web.

[\[SmartTables Documentation\]](#) [\[Directory of SmartTables Users\]](#)

My SmartTables	Public SmartTables	Shared With Me	Special SmartTables
<b>Special SmartTables</b>			
1			All compounds of P. putida KT2440
2			All genes of P. putida KT2440
3			BioCyc All Databases
4			All pathways of P. putida KT2440
5			All promoters of P. putida KT2440
6			All proteins (polypeptides + protein complexes) of P. putida KT2440
7			All polypeptides of P. putida KT2440
8			All protein complexes of P. putida KT2440
9			All enzymes of P. putida KT2440
10			All ribosomal proteins of P. putida KT2440
11			All transcription factors of P. putida KT2440
12			All transporters of P. putida KT2440
13			All cytosolic proteins of P. putida KT2440
14			All membrane proteins of P. putida KT2440
15			All periplasmic proteins of P. putida KT2440
16			All publications of P. putida KT2440
17			All reactions of P. putida KT2440
18			All riboswitches of P. putida KT2440
19			All RNAs of P. putida KT2440
20			All terminators of P. putida KT2440
21			All transcription factor binding sites of P. putida KT2440
22			All transcription units of P. putida KT2440

Show pagged **Show all**

genes.txt  
sequences.fasta

proteins.txt

RNAs.txt

TUs.txt

## Download genes.txt and sequences.fasta

lida KT2440

Product
rhs family protein
phenylacetyl-CoA dehydrogenase monomer [component of phenylacetyl-CoA dehydrogenase]
carbamate kinase
methyl-accepting chemotaxis protein
leucine-binding DNA-binding transcriptional regulator
inner membrane protein

**Export SmartTable to Spreadsheet F...**

Use the following format for values in spreadsheet:

☒ frame IDs

☐ common names  
(frame IDs make re-importing data easier)

Export smarttable

**OPERATIONS**

New ▶

Export ▼

- ▶ to Spreadsheet File... → genes.txt
- ▶ to FASTA File... → sequences.fasta
- ▶ Export to SDF...
- ▶ Export pathways to Pathway Collage

Delete ▶

Column ▶

Rows ▶

Paint Data ▶

- ▶ Rename
- ▶ Extend SmartTable by Uploading File...
- ▶ Edit Description
- ▶ Set Operations ...
- ▶ Filter
- ▶ Browse this SmartTable
- ▶ Sharing...
- ▶ Create Frozen Copy...
- ▶ Get email notification of updates to genes or pathways in this column

## Download proteins.txt, RNAs.txt and TUs.txt

The same process of genes.txt applies to proteins.txt, RNAs.txt and TUs.txt.

Some columns must be added manually using BioCyc's dropdown lists **ADD PROPERTY COLUMN** and **ADD TRANSFORM COLUMN** within the SmartTable editing webpage.

	Proteins
<input type="checkbox"/> 1	electron transfer flavoprotein (a protein CPLX1G01-196)
<input type="checkbox"/> 2	protein dithiol oxidoreductase (disulfide-forming) (a protein G1G01-133-MONOMER)
<input type="checkbox"/> 3	FleQ-cyclic di-3',5'-guanylate (a protein CPLX1G01-186)
<input type="checkbox"/> 4	HutC-urocanate (a protein CPLX1G01-219)
<input type="checkbox"/> 5	apo-FnrA (a protein CPLX1G01-161)

## Download proteins.txt

The index (Proteins Complexes) is in the SmartTable by default, but you need to add the columns Common-Name, Genes of polypeptide, complex, or RNA, and Locations.

- Common-Name is available in the dropdown list **ADD PROPERTY COLUMN**
- Genes of polypeptide, complex, or RNA is available in the dropdown list **ADD TRANSFORM COLUMN**
- Locations is available in the dropdown list **ADD PROPERTY COLUMN**.

## Download RNAs.txt

The index (All-tRNAs Misc-RNAs rRNAs) is in the SmartTable by default, but you need to add the columns Common-Name, and Gene.

- Common-Name is available in the dropdown list **ADD PROPERTY COLUMN**
- Gene is available in the dropdown list **ADD PROPERTY COLUMN**.

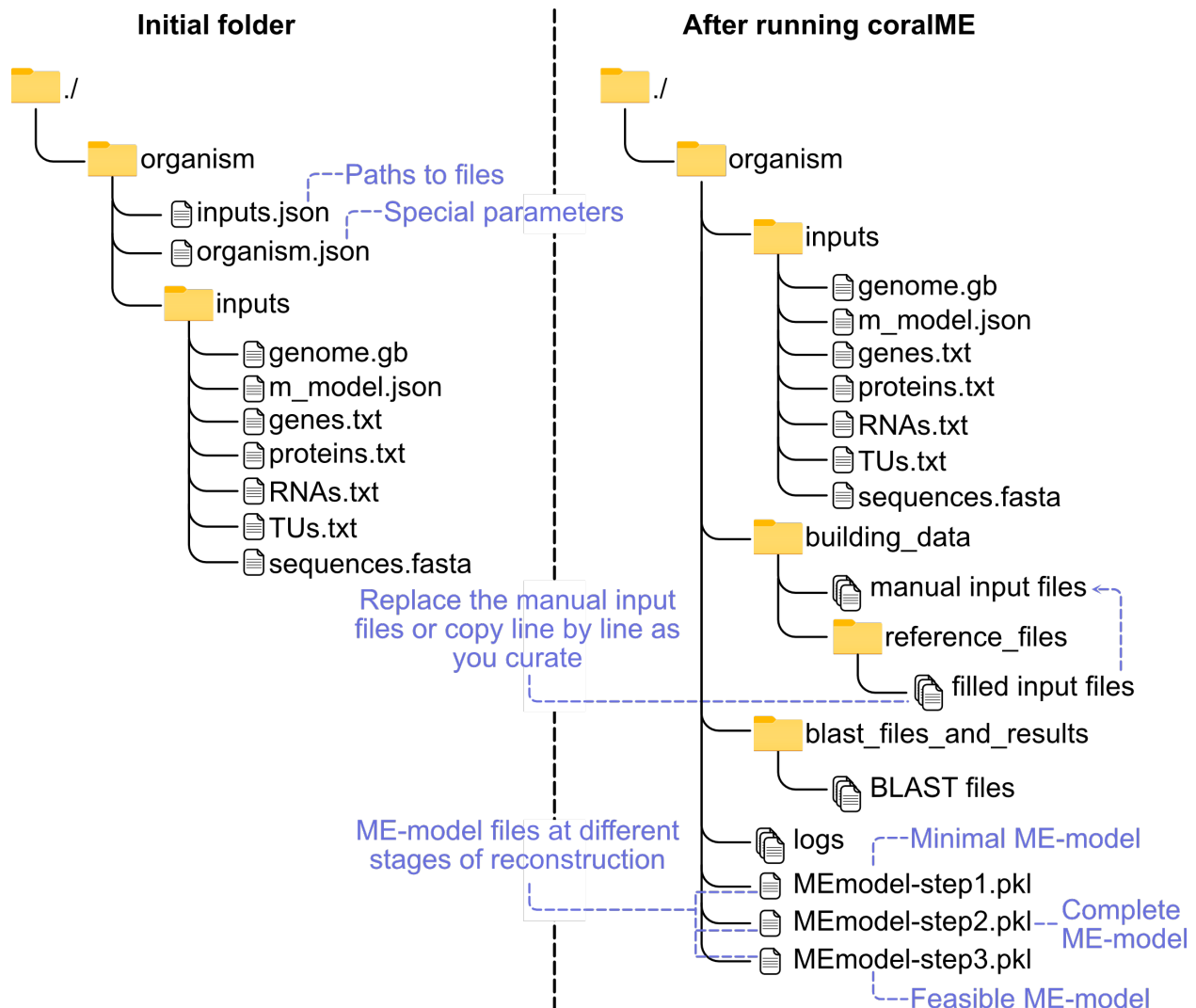
## Download TUs.txt

The index Transcription-Units is in the SmartTable by default, but you need to add the columns Genes of transcription unit, and Direction.

- Genes of transcription unit is available in the dropdown list **ADD TRANSFORM COLUMN**
- Direction is available in the dropdown list **ADD PROPERTY COLUMN**.

### 2.1.2 Initialize the folder for your organism

Copy your files to create your initial folder



### Define inputs in input.json.

See an example of `input.json`

### Define parameters in organism.json.

See an example of `organism.json`

---

**Note:** You do not need to modify these parameters right away. But once you are at the curation stage you will have to ensure these parameters are applicable to your organism.

---

## 2.1.3 Reconstruct with coralME

Here we show an example to reconstruct a dME-model of *B. subtilis*

### Import packages

```
[ ]: from coralme.builder.main import MEBuilder
```

### Define organism and inputs

```
[ ]: org = "./helper_files/tutorial/"
```

Load configuration files

```
[ ]: import os
os.chdir(org)
```

```
[ ]: organism = '{}/organism.json'.format(org)
inputs = '{}input.json'.format(org)
```

### Create builder

```
class coralme.builder.main.MEBuilder(*args, **kwargs)
```

MEBuilder class to coordinate the reconstruction of ME-models.

#### Parameters

- **\*args** – Positional arguments are passed as paths to JSON files that update the configuration of the parent class.
- **\*\*kwargs** – Further keyword arguments are passed on as dictionaries to update the configuration of the parent class.

**generate\_files**(*overwrite=True*)

Performs the Synchronize and Complement steps of the reconstruction.

This function will read the Organism and the Reference. It will synchronize the input files, complement them, and finally build the OSM for the Organism.



**Parameters**

**overwrite** (*bool*) – If True, overwrite the OSM using the defined path in the configuration.

**get\_homology**(*evaluate=1e-10*)

Calculates homology between Organism and Reference.

**Parameters**

**value** (*float*, *default 1e-10*) – Sets the E-value cutoff for calling protein homologs using BLAST.

**get\_trna\_to\_codon**()

Gets tRNA to codon association from the Genome.

**prepare\_model**()

Performs initial preparation of the M-model.

This function will fix some known issues that M-models can

**Parameters**

**overwrite** (*bool*) – If True, overwrite the OSM using the defined path in the configuration.

**troubleshoot**(*growth\_key\_and\_value=None*, *skip={}*, *guesses={}*, *platform=None*, *solver='gurobi'*, *savefile=None*, *gapfill\_cofactors=False*)

**growth\_key\_and\_value:**

dictionary of Sympy.Symbol and value to replace

**skip:**

set of ME-components to not evaluate during gapfilling

**guesses:**

set of ME-components to try first before any other set of components

**platform:**

'win32' to use gurobi (default) or cplex as solver

**solver:**

'gurobi' (default, if platform is 'win32') or 'cplex'

**savefile:**

file path (absolute or relative) to save the ME-model as a pickle file

```
[ ]: builder = MEBuilder(*[organism, inputs])
```

**Generate files**

```
[ ]: builder.generate_files(overwrite=True)
```

## Build ME-model

```
[ ]: builder.build_me_model(overwrite=False)
```

## Troubleshoot ME-model

```
[ ]: builder.troubleshoot(growth_key_and_value = { builder.me_model.mu : 0.001 })
```

---

**Note:** We set 0.001 as a standard value for feasibility checking, but feel free to modify it! Sometimes too high a value could put a significant strain on the model and give too many gaps to start with. Too low a value might not show you all the gaps needed.

---

### 2.1.4 Curate manually

For details on manual curation go to *How to manually curate a ME-model using coralME*.

## 2.2 Description of Inputs

coralME takes a total of 7 inputs, 2 required and 5 optional:

### 2.2.1 Types of inputs

#### Required

1. **Genome file** (**genome.gb**)
2. **M-model** (**m\_model.json** or **m\_model.xml**)

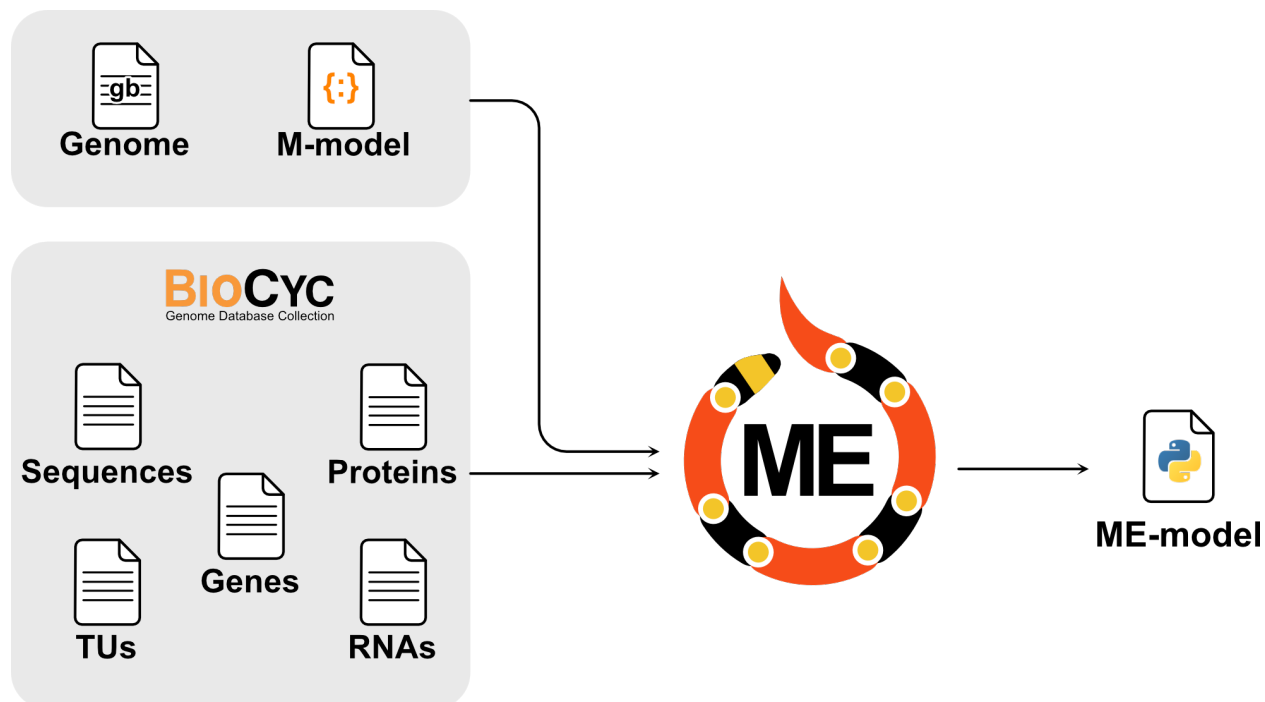
#### Optional

Downloadable from an existing **BioCyc** database under **Special SmartTables**. If no optional files are provided, coralME complements them with **genome.gb**

3. **Genes file**, by default: **genes.txt**
4. **RNAs file**, by default: **RNAs.txt**
5. **Proteins file**, by default: **proteins.txt**
6. **TUs file**, by default: **TUs.txt**.
7. **Sequences file**, by default: **sequences.fasta**

## Configuration

8. **Paths file**, by default: **inputs.json**
9. **Parameters file**, by default: **organism.json**



### 2.2.2 Description

#### Genome (genome.gb)

##### Description

The genome file contains provides coralME with:

- Gene annotations.
- Gene sequences.

##### Requirements

1. Locus tags (locus\_tag or old\_locus\_tag) **MUST** be consistent with **m\_model.json**. Make sure you download the same genome file that was used to reconstruct the M-model.
2. Has name **genome.gb**.
3. Genbank-compliant file. Must be read by BioPython correctly.
4. It must contain the entire genome sequence. Make sure to enable **Customize View>Show Sequence** before downloading the genbank file from NCBI.

See an example of [genome.gb](#) and [sequences.fasta](#)

## M-model (`m_model.json`)

### Description

The M-model provides coralME with the metabolic model components:

- Metabolic network (M-matrix)
- Gene-protein-reaction associations
- Environmental and internal constraints
- Reaction subsystems
- Biomass composition

### Requirements

1. Gene identifiers MUST be consistent with **genome.gb** locus\_tag or old\_locus\_tag. Make sure you download the same genome file that was used to reconstruct the M-model.
2. Has name **m\_model.json**.
3. COBRApy-compliant. Must be read by cobrapy-0.25.0.

See an example of [m\\_model.json](#)

## Gene dictionary (`genes.txt`) [optional]

### Description

**genes.txt** is a gene information table that can be downloaded from the **All genes of organism SmartTable** of the BioCyc database. Click **Export>to Spreadsheet File>frame IDs**. This file is optional and is meant to complement the information from **genome.gb** in case the latter is missing genes.

**genes.txt** provides coralME with:

- Gene locus tags
- Gene names
- Gene annotations
- Gene positions
- Gene products (protein, tRNA, etc.)

### Requirements

1. Contains the index **Gene Name** and columns **Accession-1**, **Left-End-Position**, **Right-End-Position**, and **Product**.
2. **Accession-1** MUST be consistent with the gene IDs in the GPRs of **m\_model.json** and with the locus\_tag (or old\_locus\_tag) in **genome.gb**.
3. **Gene Name** is consistent with:
  - Column **Genes of polypeptide, complex, or RNA of proteins.txt**
  - Column **Gene of RNAs.txt**

- Column **Genes of transcription unit** of **TUs.txt**
- Gene identifiers in **sequences.fasta**

4. **Product** is consistent with:

- Index of **proteins.txt**
- Index of **RNAs.txt**

5. Must be tab-separated

See an example of [genes.txt](#)

---

**Note: Requirement 2** regarding ID consistency should be directly met if the files are downloaded from the correct BioCyc database.

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**Note: Requirements 3, 4 and 5** regarding ID consistency should be directly met if the files are downloaded from the same BioCyc database.

---

## Proteins (proteins.txt) [optional]

### Description

**proteins.txt** is a protein complex information table that can be downloaded from the **All proteins of organism Smart-Table** of the [BioCyc](#) database. Click **Export>to Spreadsheet File>frame IDs**. This file is optional and is meant to complement the information from **genome.gb**.

**proteins.txt** provides coralME with: \* Protein complex compositions

### Requirements

1. Contains the index (**Proteins Complexes**) and columns **Common-Name**, **Genes of polypeptide, complex, or RNA**, and **Locations**.
2. (**Proteins Complexes**) is consistent with:
  - Column **Product** of **genes.txt**
3. **Genes of polypeptide, complex, or RNA** is consistent with:
  - Index **Gene Name** of **genes.txt**
4. Must be tab-separated

See an example of [proteins.txt](#)

---

**Note: Requirements 2, 3 and 4** regarding ID consistency should be directly met if the files are downloaded from the same BioCyc database.

---

## RNAs (RNAs.txt) [optional]

### Description

**RNAs.txt** is an RNA annotation table that can be downloaded from the **All RNAs of organism SmartTable** of the [BioCyc](#) database. Click **Export>to Spreadsheet File>frame IDs**. This file is optional and is meant to complement the information from **genome.gb**.

**RNAs.txt** provides coralME with:

- Genes annotated as RNA products (e.g. tRNA, rRNA, etc.)
- RNA gene annotations (e.g. amino acids - tRNA associations)

### Requirements

1. Contains the index (**All-tRNAs Misc-RNAs rRNAs**) and columns **Common-Name**, and **Gene**
2. (**All-tRNAs Misc-RNAs rRNAs**) is consistent with:
  - Column **Product** of **genes.txt**
3. **Gene** is consistent with:
  - Index **Gene Name** of **genes.txt**
4. Must be tab-separated

See an example of [RNAs.txt](#)

---

**Note: Requirements 2, 3 and 4** regarding ID consistency should be directly met if the files are downloaded from the same BioCyc database.

---

## TUs (TUs.txt) [optional]

### Description

**TUs.txt** is a transcription unit annotation table that can be downloaded from the **All TUs of organism SmartTable** of the [BioCyc](#) database. Click **Export>to Spreadsheet File>frame IDs**. This file is optional and is meant to complement the information from **genome.gb**.

**TUs.txt** provides coralME with:

- Co-transcribed genes (operons).
- Direction of transcription.
- TU IDs.

## Requirements

1. Contains the index **Transcription-Units** and columns **Genes of transcription unit**, and **Direction**
2. **Genes of transcription unit** is consistent with:
  - Index **Gene Name** of **genes.txt**
3. Must be tab-separated

See an example of [TUs.txt](#)

---

**Note: Requirements 2 and 3** regarding ID consistency should be directly met if the files are downloaded from the same BioCyc database.

---

## Gene sequences (sequences.fasta) [optional]

### Description

**sequences.fasta** is a nucleotide FASTA file that can be downloaded from the **All genes of organism SmartTable** of the [BioCyc](#) database. Click **Export>FASTA>Find sequences**. This file is optional and is meant to complement the information from **genome.gb** in case the latter is missing genes.

**sequences.fasta** provides coralME with:

- Gene sequences

## Requirements

1. Gene identifiers are consistent with:
  - Index **Gene Name** of **genes.txt**
2. Must be tab-separated

See an example of [sequences.fasta](#)

---

**Note: Requirements 1, 2 and 3** regarding ID consistency should be directly met if the files are downloaded from the same BioCyc database.

---

## Configuration of paths to files (inputs.json)

### Description

**inputs.json** is a JSON file containing paths to input files for coralME.

**inputs.json** provides coralME with:

- Paths to input files

## Requirements

1. Must be JSON-compliant
2. Must contain paths to required files (M-model and Genome).
3. All defined files must exist.

See an example of [input.json](#)

## Configuration of parameters (organism.json)

### Description

**organism.json** is a JSON file containing paths to input files for coralME.

**organism.json** provides coralME with:

- ME-modeling parameters

### Requirements

1. Must be JSON-compliant
2. Must contain the standard fields.

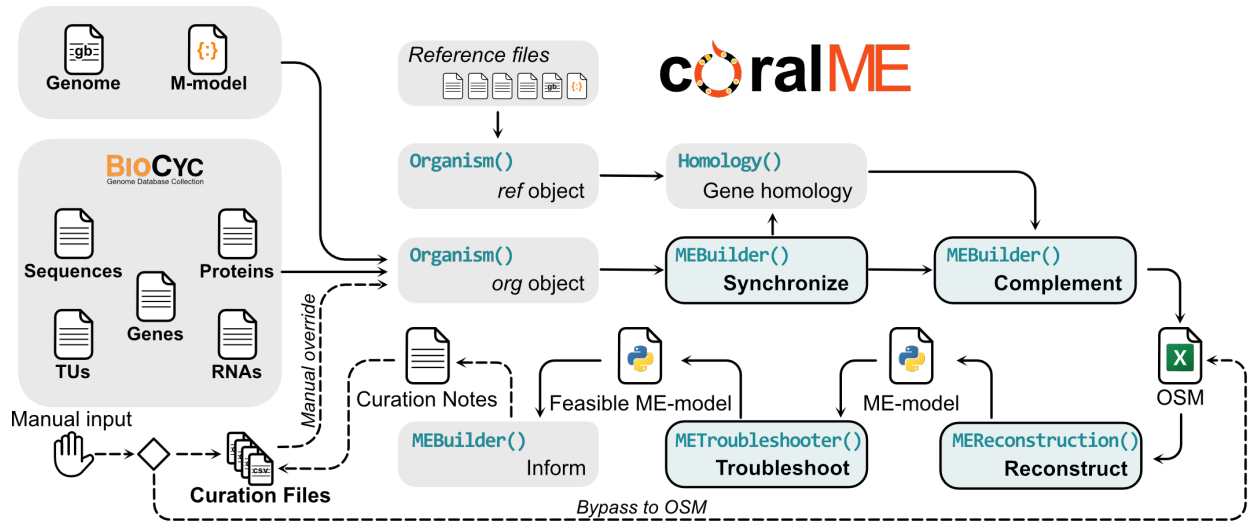
See an example of [organism.json](#)

## 2.3 Architecture of coralME

coralME is composed of 4 main classes that process and exchange organism-specific information for the reconstruction of a ME-model. The classes are:

```
class Organism()  
  
class MEBuilder()  
  
class MEREconstruction()  
  
class Homology()
```





### 2.3.1 Organism()

**class** coralme.builder.organism.Organism(*config*, *is\_reference*)

Organism class for storing information about an organism

This class acts as a database containing all necessary information to reconstruct a ME-model. It is used to retrieve and store information of the main (**org**) and the reference (**ref**) organisms. Information in Organism is read and manipulated by methods in the MEBuilders class.

#### Parameters

- **config** (*dict*) – Dictionary containing configuration and settings.
- **is\_reference** (*bool*) – If True, process as reference organism.

Role: Store information about an organism

This class acts as a database containing all necessary information to reconstruct a ME-model. It is used to retrieve and store information of the main (**org**) and the reference (**ref**) organisms. Information in Organism() is read and manipulated by methods in the MEBuilders() class. The reference can be set as any of the provided organisms in coralME, available [here](#), although we advise to choose *E. coli* and *B. subtilis* for gram-negative and gram-positive bacteria, respectively.

### 2.3.2 MEBuilders()

**class** coralme.builder.main.MEBuilders(*\*args*, *\*\*kwargs*)

MEBuilders class to coordinate the reconstruction of ME-models.

#### Parameters

- **\*args** – Positional arguments are passed as paths to JSON files that update the configuration of the parent class.
- **\*\*kwargs** – Further keyword arguments are passed on as dictionaries to update the configuration of the parent class.

Role: Coordinate the roles of other classes.

This class acts as the main coordinator between other objects, e.g. Organism, Homology, MEProcessor, and METroubleshooter. It contains methods to manipulate class Organism by using attributes in class Homology, and manually curated files in the folder containing the main organism. Moreover, it is called by objects to access stored information in other objects.

### 2.3.3 MEREconstruction()

**class** coralme.builder.main.MEREconstruction(builder)

MEREconstruction class for reconstructing a ME-model from user/automated input

#### Parameters

**MEBuilder** (coralme.builder.main.MEBuilder) –

Role: Reconstruct a ME-model from the information contained in class Organism.

This class was based almost entirely from the original ECOLme code in `build_me_model.py`. Adaptations to this code were necessary to make it applicable to other organisms.

### 2.3.4 Homology()

**class** coralme.builder.homology.Homology(org, ref, evaluate=False, verbose=False)

Homology class for storing information about homology of the main and reference organisms.

This class contains methods to predict and process homology of the main and reference organisms. Homology is inferred from the reciprocal best hits of a BLAST. The results are used to update and complement the attributes of the class Organism.

#### Parameters

- **org** (*str*) – Identifier of the main organism. Has to be the same as its containing folder name.
- **ref** (*str*) – Identifier of the reference organism. Has to be the same as its containing folder name.
- **evaluate** (*float*) – E-value cutoff to call enzyme homologs from the BLAST. Two reciprocal best hits are considered homologs if their E-value is less than this parameter.

Role: Generate and store information about homology of the main and reference organisms.

This class contains methods to predict and process homology of the main and reference organisms. Homology is inferred from the reciprocal best hits of a BLAST. The results are used to update and complement the attributes of the class Organism.

## 2.4 How to manually curate a ME-model using coralME

### 2.4.1 Manual input files

After you run coralME for the first time the following files are generated in `building_data/`. Most of them are also automatically filled by the algorithm and saved in `building_data/reference_files/`. These `_reference_files_` are meant to guide manual curation as they contain all information mapped by coralME formatted as manual input files.

`__coralME` does not overwrite any file in `building_data/`, but it will always overwrite files in `building_data/reference_files/`

- **termination\_subreactions.txt**

Input here will define translation termination subreactions and their machinery.

```
class coralme.builder.curation.TerminationSubreactions(org, id='termination_subreactions',
                                                    config={},
                                                    file='termination_subreactions.txt',
                                                    name='Translation termination
                                                    subreactions')
```

Reads manual input to define translation termination subreactions.

This class creates the property “termination\_subreactions” from the manual inputs in termination\_subreactions.txt in an instance of Organism.

Input here will define translation termination subreactions and their machinery.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

**termination\_subreactions.txt :**

subreaction enzymes PrfA\_mono\_mediated\_termination PrfA\_mono ...

- **peptide\_release\_factors.txt**

Input here will define peptide release factors.

```
class coralme.builder.curation.PeptideReleaseFactors(org, id='peptide_release_factors', config={},
                                                    file='peptide_release_factors.txt',
                                                    name='Peptide release factors')
```

Reads manual input to define peptide release factors.

This class creates the property “peptide\_release\_factors” from the manual inputs in peptide\_release\_factors.txt in an instance of Organism.

Input here will define peptide release factors.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

**peptide\_release\_factors.txt :**

release\_factor enzyme UAA generic\_RF ...

- **rna\_degradosome.txt**

Input here will define the composition of the RNA degradosome.

```
class coralme.builder.curation.RNADegradosome(org, id='rna_degradosome', config={},
                                                    file='rna_degradosome.txt', name='RNA degradosome
                                                    composition')
```

Reads manual input to add RNA degradosome composition.

This class creates the property “rna\_degradosome” from the manual inputs in rna\_degradosome.txt in an instance of Organism.

Input here will define the composition of the RNA degradosome.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### **rna\_degradosome.txt :**

enzymes Eno\_dim\_mod\_mg2(4) ...

- **special\_trna\_subreactions.txt**

Input here will define special tRNA subreactions, such as tRNA-Sec (selenocysteine) synthesis from tRNA-Ser.

```
class coralme.builder.curation.SpecialtRNASubreactions(org, id='special_trna_subreactions',
                                                    config={},
                                                    file='special_trna_subreactions.txt',
                                                    name='Special tRNA subreactions')
```

Reads manual input to define special tRNA subreactions.

This class creates the property “special\_trna\_subreactions” from the manual inputs in special\_trna\_subreactions.txt in an instance of Organism.

Input here will define special tRNA subreactions, such as tRNA-Sec (selenocysteine) synthesis from tRNA-Ser.

### **Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### **special\_trna\_subreactions.txt :**

subreaction enzymes PrfA\_mono\_mediated\_termination PrfA\_mono ...

- **lipoprotein\_precursors.txt**

Input here will add lipoprotein precursors.

```
class coralme.builder.curation.LipoproteinPrecursors(org, id='lipoprotein_precursors', config={},
                                                    file='lipoprotein_precursors.txt',
                                                    name='Lipoprotein precursors')
```

Reads manual input to add lipoprotein precursors.

This class creates the property “lipoprotein\_precursors” from the manual inputs in lipoprotein\_precursors.txt in an instance of Organism.

Input here will add lipoprotein precursors.

### **Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### **lipoprotein\_precursors.txt :**

name,gene AcrA,b0463 ...

- **special\_modifications.txt**

Input here will define machinery for special modifications. These modifications are a set of pre-defined modifications that are used in ME-models.

```
class coralme.builder.curation.SpecialModifications(org, id='special_modifications', config={},
                                                    file='special_modifications.txt', name='Special
                                                    protein modifications')
```

Reads manual input to define machinery for special modifications.

This class creates the property “special\_modifications” from the manual inputs in special\_modifications.txt in an instance of Organism.

Input here will define machinery for special modifications. These modifications are a set of pre-defined modifications that are used in ME-models.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

#### Examples

##### special\_modifications.txt :

modification enzymes stoich fes\_transfer CPLX0-7617,IscA\_tetra,CPLX0-7824 ...

- **excision\_machinery.txt**

Input here will define machinery for excision.

```
class coralme.builder.curation.ExcisionMachinery(org, id='excision_machinery', config={},
                                                file='excision_machinery.txt', name='Excision
                                                machinery')
```

Reads manual input to define machinery for excision.

This class creates the property “excision\_machinery” from the manual inputs in excision\_machinery.txt in an instance of Organism.

Input here will define machinery for excision.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

#### Examples

##### excision\_machinery.txt :

mechanism enzymes rRNA\_containing RNase\_E\_tetra\_mod\_zn2(2), ... ...

- **orphan\_and\_spont\_reactions.txt**

Input here will mark reactions as orphan or spontaneous. Orphan reactions will be associated with CPLX\_dummy, and spontaneous ones will not require enzymes for flux.

```
class coralme.builder.curation.OrphanSpontReactions(org, id='orphan_and_spont_reactions',
                                                    config={},
                                                    file='orphan_and_spont_reactions.txt',
                                                    name='Orphan and spontaneous reactions')
```

Reads manual input to add reactions to the ME-model.

This class creates the property “orphan\_and\_spont\_reactions” from the manual inputs in orphan\_and\_spont\_reactions.txt in an instance of Organism.

Input here will mark reactions as orphan or spontaneous. Orphan reactions will be associated with CPLX\_dummy, and spontaneous ones will not require enzymes for flux.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### orphan\_and\_spont\_reactions.txt :

name description is\_reversible is\_spontaneous subsystems CODH\_Fe\_loading Loading of Fe false true ...

- **enzyme\_reaction\_association.txt**

Input here will create the association between enzymes and reactions in the ME-model.

```
class coralme.builder.curation.EnzymeReactionAssociation(org, id='enz_rxn_assoc_df', config={},
                                                         file='enzyme_reaction_association.txt',
                                                         name='Enzyme-reaction associations')
```

Reads manual input to specify enzyme-reaction associations.

This class creates the property “enz\_rxn\_assoc\_df” from the manual inputs in enzyme\_reaction\_association.txt in an instance of Organism.

Input here will create the association between enzymes and reactions in the ME-model.

### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### enzyme\_reaction\_association.txt :

Reaction Complexes ADNt2pp NUPG-MONOMER OR NUPC-MONOMER ...

- **peptide\_compartment\_and\_pathways.txt**

Input here will modify protein locations, and translocation pathways in the ME-model.

```
class coralme.builder.curation.ProteinLocation(org, id='protein_location', config={},
                                                file='peptide_compartment_and_pathways.txt',
                                                name='Protein location')
```

Reads manual input to add protein locations.

This class creates the property “protein\_location” from the manual inputs in peptide\_compartment\_and\_pathways.txt in an instance of Organism.

Input here will modify protein locations, and translocation pathways in the ME-model.

### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### peptide\_compartment\_and\_pathways.txt :

Complex Complex\_compartment Protein Protein\_compartment translocase\_pathway BSU02690-MONOMER Plasma\_Membrane BSU02690() Plasma\_Membrane s ...

- **translocation\_pathways.txt**

Input here will define translocation pathways and their machinery.

```
class coralme.builder.curation.TranslocationPathways(org, id='translocation_pathways', config={},
                                                       file='translocation_pathways.txt',
                                                       name='Translocation pathways')
```

Reads manual input to define translocation pathways.

This class creates the property “translocation\_pathways” from the manual inputs in translocation\_pathways.txt in an instance of Organism.

Input here will define translocation pathways and their machinery.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

#### Examples

##### **translocation\_pathways.txt :**

pathway enzyme sec BSU27650-MONOMER sec BSU35300-MONOMER sec secYEG ...

- **rna\_modification.txt**

Input here will define enzymes that perform RNA modifications for either rRNA or tRNA in the ME-model.

```
class coralme.builder.curation.RNAModificationMachinery(org, id='rna_modification_df', config={},
                                                         file='rna_modification.txt', name='RNA
                                                         Modification machinery')
```

Reads manual input to add RNA modification machinery.

This class creates the property “rna\_modification\_df” from the manual inputs in rna\_modification.txt in an instance of Organism.

Input here will define enzymes that perform RNA modifications for either rRNA or tRNA in the ME-model.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

#### Examples

##### **rna\_modification.txt :**

modification positions type enzymes source D 16,17,20,20A,21 tRNA DusB\_mono ...

- **ribosomal\_proteins.txt**

Input here will define the composition of the ribosome.

```
class coralme.builder.curation.RibosomeStoich(org, id='ribosome_stoich', config={},
                                                file='ribosomal_proteins.txt', name='Ribosomal
                                                proteins')
```

Reads manual input to define ribosome composition.

This class creates the property “ribosome\_stoich” from the manual inputs in ribosomal\_proteins.txt in an instance of Organism.

Input here will define the composition of the ribosome.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### ribosomal\_proteins.txt :

subunits proteins 30S RpsD\_mono,... 50S generic\_23s\_rRNAs, generic\_5s\_rRNAs, RplA\_mono,...

- **rho\_independent.txt**

Input here will mark genes with rho independent transcription termination.

```
class coralme.builder.curation.RhoIndependent(org, id='rho_independent', config={},
                                             file='rho_independent.txt', name='Genes with
                                             rho-independent termination')
```

Reads manual input to define genes with rho independent termination.

This class creates the property “rho\_independent” from the manual inputs in rho\_independent.txt in an instance of Organism.

Input here will mark genes with rho independent transcription termination.

### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### rho\_independent.txt :

id b0344 ...

- **sigma\_factors.txt**

Input here will mark proteins for N-terminal methionine cleavage in the ME-model.

```
class coralme.builder.curation.Sigmas(org, id='sigmas', config={}, file='sigma_factors.txt', name='Sigma
                                     factors')
```

Reads manual input to modify or add sigma factors.

This class creates the property “sigmas” from the manual inputs in sigma\_factors.txt in an instance of Organism.

Input here will mark proteins for N-terminal methionine cleavage in the ME-model.

### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### sigma\_factors.txt :

sigma,complex,genes,name RpoH\_mono,RNAP\_32H,b3461,”RNA polymerase, sigma 32 (sigma H) factor” ...

- **cleaved\_methionine.txt**

Input here will mark proteins for N-terminal methionine cleavage in the ME-model.

```
class coralme.builder.curation.CleavedMethionine(org, id='cleaved_methionine', config={},
                                                  file='cleaved_methionine.txt', name='Proteins with
                                                  N-terminal methionine cleavage')
```

Reads manual input to mark proteins that undergo N-terminal methionine cleavage.

This class creates the property “cleaved\_methionine” from the manual inputs in cleaved\_methionine.txt in an instance of Organism.

Input here will mark proteins for N-terminal methionine cleavage in the ME-model.



**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

**Examples****cleaved\_methionine.txt :**

cleaved\_methionine\_genes b4154 ...

- **folding\_dict.txt**

Input here will define folding pathways for proteins.

```
class coralme.builder.curation.FoldingDict(org, id='folding_dict', config={}, file='folding_dict.txt',
                                           name='Protein to folding machinery associations')
```

Reads manual input to define folding pathways for proteins.

This class creates the property “folding\_dict” from the manual inputs in folding\_dict.txt in an instance of Organism.

Input here will define folding pathways for proteins.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

**Examples****folding\_dict.txt :**

mechanism enzymes GroEL\_dependent\_folding b0014, ... ...

- **translocation\_multipliers.txt**

Input here will modify how many pores are required for the translocation of a protein.

```
class coralme.builder.curation.TranslocationMultipliers(org, id='translocation_multipliers',
                                                         config={},
                                                         file='translocation_multipliers.txt',
                                                         name='Translocation multipliers')
```

Reads manual input to define translocation multipliers.

This class creates the property “translocation\_multipliers” from the manual inputs in translocation\_multipliers.txt in an instance of Organism.

Input here will modify how many pores are required for the translocation of a protein.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

**Examples****translocation\_multipliers.txt :**

Gene,YidC\_MONOMER,TatE\_MONOMER,TatA\_MONOMER b1855,2.0,0.0,0.0 ...

- **subreaction\_matrix.txt**

Input here will define subreactions in the ME-model.

```
class coralme.builder.curation.SubreactionMatrix(org, id='subreaction_matrix', config={},
                                                  file='subreaction_matrix.txt', name='Matrix of
                                                  subreaction stoichiometries')
```

Reads manual input to add subreactions.

This class creates the property “subreaction\_matrix” from the manual inputs in subreaction\_matrix.txt in an instance of Organism.

Input here will define subreactions in the ME-model.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

#### Examples

##### subreaction\_matrix.txt :

Reaction Metabolites Stoichiometry mod\_acetyl\_c accoa\_c -1.0 mod\_acetyl\_c coa\_c +1.0 ...

- **me\_metabolites.txt**

Input here will mark metabolites in the M-model for replacement with their corrected E-matrix component.

- **elongation\_subreactions.txt**

Input here will define translation elongation subreactions and their machinery.

```
class coralme.builder.curation.MEMetabolites(org, id='me_mets', config={}, file='me_metabolites.txt',
                                             name='Metabolites to substitute from M-model')
```

Reads manual input to replace metabolites in the M-model.

This class creates the property “me\_mets” from the manual inputs in me\_metabolites.txt in an instance of Organism.

Input here will mark metabolites in the M-model for replacement with their corrected E-matrix component.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

#### Examples

##### me\_metabolites.txt :

id me\_id name formula compartment type sufbcd\_c CPLX0-1341 SufBCD complex REPLACE ...

- **subsystem\_classification.txt**

Input here will classify subsystems in umbrella classifications which are then used to set a median Keff and correct it with the complex SASA.

```
class coralme.builder.curation.SubsystemClassification(org, id='subsystem_classification',
                                                       config={}, file='subsystem_classification.txt',
                                                       name='Classification of subsystems')
```

Reads manual input to classify subsystems for Keff estimation.

This class creates the property “subsystem\_classification” from the manual inputs in subsystem\_classification.txt in an instance of Organism.

Input here will classify subsystems in umbrella classifications which are then used to set a median Keff and correct it with the complex SASA.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

subsystem\_classification.txt : subsystem central\_CE central\_AFN intermediate secondary other  
S\_Amino\_acids\_and\_related\_molecules 0 1 0 0 0 ...

- **reaction\_matrix.txt**

Input here will define reactions directly in the ME-model. Definitions here will be added to the ME-model after processing the M-model into the ME-model.

```
class coralme.builder.curation.ReactionMatrix(org, id='reaction_matrix', config={},
                                              file='reaction_matrix.txt', name='Matrix of reaction
                                              stoichiometries')
```

Reads manual input to add reactions to the ME-model.

This class creates the property “reaction\_matrix” from the manual inputs in reaction\_matrix.txt in an instance of Organism.

Input here will define reactions directly in the ME-model. Definitions here will be added to the ME-model after processing the M-model into the ME-model.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

**reaction\_matrix.txt :**

Reaction Metabolites Stoichiometry Cs\_cyto\_import cs\_p -1.0 Cs\_cyto\_import h\_c 1.0 Cs\_cyto\_import  
cs\_c 1.0 Cs\_cyto\_import h\_p -1.0 ...

- **lipid\_modifications.txt**

Input here will define enzymes that perform lipid modifications.

- **amino\_acid\_trna\_synthetase.txt**

Input here will define amino acid tRNA ligases.

- **initiation\_subreactions.txt**

Input here will define translation initiation subreactions and their machinery.

```
class coralme.builder.curation.AminoacidtRNASynthetase(org, id='amino_acid_trna_synthetase',
                                                         config={},
                                                         file='amino_acid_trna_synthetase.txt',
                                                         name='Amino acid to tRNA synthetase
                                                         associations')
```

Reads manual input to define amino acid tRNA ligases.

This class creates the property “amino\_acid\_trna\_synthetase” from the manual inputs in amino\_acid\_trna\_synthetase.txt in an instance of Organism.

Input here will define amino acid tRNA ligases.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### **amino\_acid\_trna\_synthetase.txt :**

amino\_acid enzyme ala\_\_L\_c Ala\_RS\_tetra\_mod\_zn2(4) ...

- **post\_transcriptional\_modification\_of\_RNA.txt**

Input here will define RNA genes that undergo modifications.

```
class coralme.builder.curation.RNAModificationTargets(org, id='rna_modification_targets', config={},
                                                    file='post_transcriptional_modification_of_RNA.txt',
                                                    name='RNA modification targets')
```

Reads manual input to add RNA modification targets.

This class creates the property “rna\_modification\_targets” from the manual inputs in post\_transcriptional\_modification\_of\_RNA.txt in an instance of Organism.

Input here will define RNA genes that undergo modifications.

#### **Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### **post\_transcriptional\_modification\_of\_RNA.txt :**

bnum position modification b0202 20A D ...

- **protein\_corrections.txt**

Input here will add, modify complexes in the ME-model, as well as add, modify their modifications. You can add a complex modification ID in the replace column, which will remove that modified complex and replace it with your manually added one.

- **reaction\_median\_keffs.txt**

Input here will define median Keffs for estimation of Keffs using the SASA method.

- **transcription\_subreactions.txt**

Input here will define machinery for transcription subreactions. These subreactions are a set of pre-defined subreactions that are used in ME-models.

```
class coralme.builder.curation.TranscriptionSubreactions(org, id='transcription_subreactions',
                                                         config={},
                                                         file='transcription_subreactions.txt',
                                                         name='Transcription subreactions')
```

Reads manual input to define transcription subreactions.

This class creates the property “transcription\_subreactions” from the manual inputs in transcription\_subreactions.txt in an instance of Organism.

Input here will define machinery for transcription subreactions. These subreactions are a set of pre-defined subreactions that are used in ME-models.

#### **Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### transcription\_subreactions.txt :

mechanism enzymes Transcription\_normal\_rho\_independent Mfd\_mono\_mod\_mg2(1),NusA\_mono,NusG\_mono,GreA\_mono ...

- **generic\_dict.txt**

Input here will define generics.

```
class coralme.builder.curation.GenericDict(org, id='generic_dict', config={}, file='generic_dict.txt',
                                           name='Dictionary of generic complexes')
```

Reads manual input to define generics.

This class creates the property “generic\_dict” from the manual inputs in generic\_dict.txt in an instance of Organism.

Input here will define generics.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### generic\_dict.txt :

generic\_component enzymes generic\_16Sm4Cm1402 RsmH\_mono,RsmI\_mono ...

- **ribosome\_subreactions.txt**

Input here will define enzymes that perform a ribosome subreaction.

```
class coralme.builder.curation.RibosomeSubreactions(org, id='ribosome_subreactions', config={},
                                                    file='ribosome_subreactions.txt',
                                                    name='Ribosomal subreactions')
```

Reads manual input to define ribosome subreactions.

This class creates the property “ribosome\_subreactions” from the manual inputs in ribosome\_subreactions.txt in an instance of Organism.

Input here will define enzymes that perform a ribosome subreaction.

#### Parameters

**org** (*coralme.builder.organism.Organism*) – Organism object.

## Examples

### ribosome\_subreactions.txt :

subreaction enzyme gtp\_bound\_30S\_assembly\_factor\_phase1 BSU16650-MONOMER ...

- **reaction\_corrections.txt**

Input here will modify reactions at the M-model stage before ME-model building.

```
class coralme.builder.curation.ReactionCorrections(org, id='reaction_corrections', config={},
                                                    file='reaction_corrections.txt', name='Reaction
                                                    corrections')
```

Reads manual input to modify reactions in the M-model.

This class creates the property “reaction\_corrections” from the manual inputs in reaction\_corrections.txt in an instance of Organism.

Input here will modify reactions at the M-model stage before ME-model building.

**Parameters**

**org** (*coralme.builder.organism.Organism*) – Organism object.

**Examples****reaction\_corrections.txt :**

reaction\_id,name,gene\_reaction\_rule,reaction,notes COBAL2tpp,cobalt transport in via permease (no H+),BSU24740,cobalt2\_e -> cobalt2\_c,No notes ...

- **TUs\_from\_biocyc.txt**  
Input here will modify transcriptional unit information.

**2.4.2 How to curate?**

1. If you have not run coralME yet, go back to *GettingStarted.ipynb*.
2. **Copy** all of the generated *reference files* in *building\_data/reference\_files* and replace accordingly in *building\_data/*
3. **Go one by one** through the files in *building\_data/* curating as needed! Important flags are risen in *curation\_notes.json* to further guide you through curation.
4. Everytime you make a change, **run the model through the troubleshooter!** It will show you remaining gaps to look at, and the new curation notes might show new warnings.
5. **Keep iterating!** You will have finished when no gaps are present, and all remaining warnings in curation notes are irrelevant.

**2.5 Frequently Asked Questions****2.5.1 My gene identifiers are not consistent, what should I do?**

We know that consistent gene ID conventions are a problem across all platforms of bioinformatics. We tried to generalize as much as possible what the gene conventions could be, but often different genome assemblies or M-model reconstructions yield inconsistent files.

What you should do depends on your problem, so we will classify the gene ID convention issues considering there are three main sources of information that must be consistent:

- **M-model** *gene identifiers*
- **Genome** *locus\_tag*
- **Optional file** column *Accession-I*

Overall, you can assume that modifying the genome genbank file is the hardest approach and thus, the last resort.

## 1. M-model and Genbank are consistent, but they are not consistent with the BioCyc files.

Make sure that you looked for the correct BioCyc database, which corresponds to the M-model reconstruction. One quick way to ensure that is to copy one gene from your genbank or M-model and paste it in the search bar of BioCyc. Best case scenario, your microbe will appear in the list. Download the files from there and your problems are solved!

### That didn't help?

It is possible that even when ensuring the BioCyc database is correct, the **Accession-1** column of genes.txt is still not consistent. However, you can assume that the correct IDs are somewhere in the database, since you found it looking for a gene id that follows your conventions (see [Getting Started](#)).

Try:

- Adding new columns in the gene **SmartTable**, **Accession-2** or **Synonyms** could contain your IDs.
- Maybe your IDs and BioCyc's only differ by an underscore, e.g. "PP0001" and "PP\_0001". Use a text editor to change the IDs accordingly in **Accession-1** of genes.txt. Make sure not to make a mistake by editing gene IDs!

## 2. M-model and Genbank are not consistent

Make sure that you downloaded the same genbank file that was used to reconstruct the M-model, that is critical! If this is happening, you probably have the wrong genbank.

If you have a gene dictionary to convert between conventions, change the files to the IDs that are consistent with BioCyc.

## INDICES AND TABLES

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- `modindex`
- `search`



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