Lab Book

Cientific Iniciation - Coral Metagenomes Letcia Costa Cavalcante

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> > 2018

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May 2018

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1.1.1 Learning LATEX

- Working folder: path

LATEX is a high-quality typesetting system, available as free software, which allows to produce scientific or technical documents [?]. I am using LATEX to create a Bioinformatics Lab Book. To compile my Lab Book, I can use command lines (pdflatex and bibtex). Afterwards I can visualise the produced .pdf file with evince or another reader. Alternatevily, I can use a Latex editor, such as TexWorks (https://www.tug.org/texworks/), which allows me to write the code and control the pdf file in the same environment (Figure 1.1).

To compile the .tex file in the command line:

\$pdflatex lab-book \$bibtex lab-book \$pdflatex lab-book \$pdflatex lab-book

To visualise the .pdf:

\$evince lab-book.pdf &

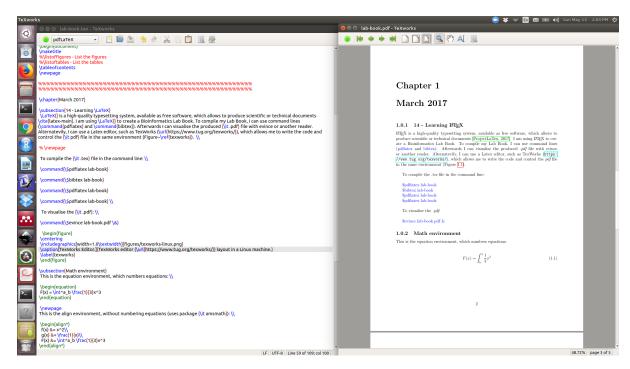


Figura 1.1: TexWorks editor (https://www.tug.org/texworks/) layout in a Linux machine.

1.1.2 Math environment

This is the equation environment, which numbers equations:

$$F(x) = \int_{b}^{a} \frac{1}{3}x^{3} \tag{1.1}$$

This is the align environment, without numbering equations (uses package amsmath):

$$f(x) = x^{2}$$

$$g(x) = \frac{1}{x}$$

$$F(x) = \int_{b}^{a} \frac{1}{3}x^{3}$$

1.1.3 15 - Short-term project proposal

Some text here. Incluing and referencing a table (table 1.1).

- $\bullet\,$ First numbered list item
- Second numbered list item

Tabela 1.1: table0

species	changes	score
Macaque	4	0.0
Human	2	14.9
Orangutan	0	0.0
Pan	0	0.0
Gorilla	0	0.0

Creation of data base of metagenomes and genomes

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2.1.1 Bibliographic search for genomes

Found a new possibility of phyla list. Because of this, there are four possibilities of list of microorganisms phyla, one of them, the SILVA database, is based in RNA sequences:

- The list of Prokariotic names with stading nomenclature http://www.bacterio.net/-classifphyla.html
- SILVA database LSU(large subunit of ribosome) https://www.arb-silva.de/browser/lsu/
- SILVA database SSU(small subunit of ribosome) https://www.arb-silva.de/browser/ssu/
- PATRIC GENOMES https://www.patricbrc.org/view/Taxonomy/2#view_tab= taxontree

The list of articles used until now is:

- 10.1038/nature14486
- 10.1038/ismej.2013.111
- 10.1038/ismej.2013.174
- 10.1038/ismej.2016.43
- 10.1038/nature12352
- 10.1038/nature14486
- 10.1038/nature21031
- 10.1038/ismej.2015.233

- 10.1038/ncomms13219
- \bullet 10.1073/pnas.0801980105
- 10.1111/1462-2920.13362
- 10.1126/science.1132690
- 10.1186/s40168-015-0077-6

Tabela 2.1: table 1

DOI	DI I
DOI	Phylum
10.1038/nature14486	Candidatus Falkowbacteria
10.1038/nature14486	Candidatus Kuenenbacteria
10.1038/nature14486	Candidatus Magasanikbacteria
10.1038/nature14486	Candidatus Uhrbacteria
10.1038/nature14486	Candidatus Moranbacteria
10.1038/nature14486	Candidatus Azambacteria
10.1038/nature14486	Candidatus Yanofskybacteria
10.1038/nature14486	Candidatus Jorgensenbacteria
10.1038/nature 14486	Candidatus Wolfebacteria
10.1038/nature 14486	Candidatus Giovannonibacteria
10.1038/nature 14486	Candidatus Nomurabacteria
10.1038/nature 14486	Candidatus Campbellbacteria
10.1038/nature 14486	Candidatus Adlerbacteria
10.1038/nature 14486	Candidatus Kaiserbacteria
10.1038/nature14486	C. S. yataiensis
10.1038/nature14486	Pacebacteria
10.1038/nature14486	Candidatus Collierbacteria
10.1038/nature14486	Candidatus Beckwithbacteria
10.1038/nature14486	Candidatus Roizmanbacteria
10.1038/nature14486	Candidatus Saphirobacteria
10.1038/nature14486	Candidatus Amesbacteria
10.1038/nature 14486	Candidatus Woesebacteria
10.1038/nature14486	Candidatus Gottesmanbacteria
10.1038/nature14486	Candidatus Levybacteria
10.1038/nature14486	Candidatus Daviesbacteria
10.1038/nature14486	Candidatus Curtissbacteria
10.1038/nature 14486	WWE3
10.1038/nature14486	CPR3
10.1038/nature 14486	WS6
10.1038/nature14486	Candidatus Berkelbacteria
$10.1038^{'}$ /nature 14486	Candidatus Peregrinibacteria
10.1038/nature 14486	Candidatus Gracilibacteria
$10.1038/{ m nature} 14486$	CPR2
10.1038/nature14486	Kazan
10.1038/nature14486	Saccharibacteria (TM7)
$10.1038/{ m nature} 14486$	SR1
10.1038/ncomms13219	Candidatus Kerfeldbacteria
10.1038/ncomms13219	Candidatus Komeilibacteria
10.1038/ncomms13219	Candidatus Andersenbacteria
10.1038/ncomms13219	Candidatus Ryanbacteria
10.1038/ncomms13219	Candidatus Niyogibacteria
10.1000/110011111010210	

40.40==1	G 1.1
10.1038/ncomms13219	Candidatus Tagabacteria
10.1038/ncomms13219	Candidatus Terrybacteria
10.1038/ncomms13219	Candidatus Vogelbacteria
10.1038/ncomms13219	Candidatus Zambryskibacteria
10.1038/ncomms13219	Candidatus Taylorbacteria
10.1038/ncomms13219	Candidatus Sungbacteria
10.1038/ncomms13219	Candidatus Brennerbacteria
10.1038/ncomms13219	Candidatus Spechtbacteria
$10.1038 \rm /ncomms 13219$	Candidatus Staskawiczbacteria
10.1038/ncomms13219	Candidatus Wildermuthbacteria
10.1038/ncomms13219	Candidatus Portnoybacteria
10.1038/ncomms13219	Candidatus Woykebacteria
10.1038/ncomms13219	Candidatus Blackburnbacteria
10.1038/ncomms13219	Candidatus Chisholmbacteria
·	Candidatus Buchananbacteria
10.1038/ncomms13219	Candidatus Jacksonbacteria
10.1038/ncomms13219	
10.1038/ncomms13219	Candidatus Veblenbacteria
10.1038/ncomms13219	Candidatus Nealsonbacteria
10.1038/ncomms13219	Candidatus Colwellbacteria
10.1038/ncomms13219	Candidatus Liptonbacteria
10.1038/ncomms13219	Candidatus Harrisonbacteria
10.1038/ncomms13219	Candidatus Yonathbacteria
10.1038/ncomms13219	Candidatus Lloydbacteria
10.1038/ncomms13219	Candidatus Abawacabacteria
10.1038/ncomms13219	Candidatus Doudnabacteria
10.1038/ismej.2013.111	Candidatus Poribacteria
10.1111/1462-2920.13362	Candidatus Desantisbacteria
10.1038/nature 12352	Candidatus Omnitrophica
10.1038/nature 12352	Candidatus Aminicenantes
10.1126/science.1132690	Candidatus Micrarchaeota
$10.1038/{\rm nature} 14486$	Candidatus Magasanikbacteria
10.1073/pnas.0801980105	Candidatus Korarchaeota
10.1038/nature12352	Candidatus Fervidibacteria
10.1038/nature12352	Candidatus Aenigmarchaeota
10.1038/ismej.2016.43	Candidatus Fermentibacteria
10.1038/ismej.2013.174	Candidatus Bathyarchaeota
10.1036/j.cub.2015.01.014	Candidatus Woesearchaeota
, •	
10.1016/j.cub.2015.01.014	Candidatus Kryptonia
10.1038/nature12352	Candidatus Diapherotrites
10.1038/nature12352	Candidatus Latescibacteria
10.1038/nature21031 10.1038/ismej.2015.233	Candidatus Thorarchaeota
10.1038/ncomms13219	Candidatus Lindowbacteria
10.1038/nature 12352	Candidatus Parvarchaeota
10.1038/nature12352	Candidatus Cloacimonetes
10.1038/nature 12352	Candidatus Hydrogenedentes
10.1038/nature 12352	Candidatus Acetothermia

10.1038/nature 12352
10.1038/ncomms13219
10.1186/s40168-015-0077-6
10.1038/nature 21031
10.1038/nature 21031
10.1038/nature21031

Candidatus Nanohaloarchaeota Candidatus Eisenbacteria candidate division WOR-3 Lokiarchaeota Odinarchaeota Heimdallarchaeota

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2.2.1 Bibliographic search for metagenomes

The reserarch for coral metagenomes started last year. The actual list is:

Tabela 2.2: table 1

IDs
mgm4440319.3
mgm4440370.3
mgm4440371.3
mgm4440372.3
mgm4440373.3
mgm4440374.3
mgm4440375.3
mgm4440376.3
mgm4440377.3
mgm4440378.3
mgm4440379.3
mgm4440380.3
mgm4440381.3
mgm4445755.3
mgm4445756.3
mgm4480739.3
mgm4480740.3
mgm4480741.3
mgm4480748.3
mgm4480750.3
mgm4487909.3
mgm4487910.3
mgm4487911.3
mgm4516541.3
$\mathrm{mgm}4516694.3$
mgm4653307.3
mgm4694757.3
mgm4694758.3

mgm4694759.3mgm4694760.3SRR1275409 SRR1275449 SRR1283349 SRR1283371 SRR1283377 SRR1283433 SRR1283435 SRR1283437 SRR1286223 SRR1286225 SRR1286226 SRR1286227 SRR1286229 SRR1286232 SRR1822488 SRR1822516 SRR3499156 SRR3569370 SRR3694369 SRR3694370 SRR3694371 SRR3694372 SRR5215424 SRR5215454 SRR5215455 SRR5215456 SRR5215457 SRR5215458 SRR5215462

I found these metagenomes in the article: "Metagenomic analysis reveals a green sulfur bacterium as a potential coral symbiont"

SRR5605611

SRR2937345 SRR2937346 SRR2937347 SRR2937348 SRR2937350 SRR2937351 SRR2937351 SRR2937353 SRR2937354 SRR2937355 SRR2937356

Espcie: Platygyra carnosa Healthy

I found other metagenomes of coral from article doi 10.3389/fmars.2018.001011 uptated the file pmc_results_1.txt in the repository Lab_book. I continue to look the articles in results. Estou atualizando a lista pmc_results_2.txt Na pesquisa bibliografica olhando o ttulo ja me faz perceber se devo descartar e olhar. E olho aqueles que marquei para olhar. Ao olhar, leio o resumo procurando por metodos. E vou para os metodos do artigo para checar. Checking the sizes of metagenomes files. The mg-rast metagenomes base have 72 Gb.

The pipeline of bioinformatic is different for MGRAST and NCBI. The size of NCBI should be superestimated, because the ncbi says the file size of sra file, but most of them is paired-end metagenomes, so when we apply fastq-dump, its generate two files fastq.

Download of metagenomes

3.1 Download of mg-rast files

Espao no SDU Disponvel para o ebiodiv: 10Tb Bia: 5Tb Rilquer: 2T Remanescente: 3Tb

- Working folder: $scratch/ebiodiv/leticia.cavalcante/mg_rast$

I insert the list of metagenomes in the files before using it. After this, I used the following command line:

- Command: nohup bash download_curl_mgrast_corais.sh $_{\dot{c}}$ download_curl_mgrast_corais.nohupout &

3.2 Download of NCBI metagenomes

I use the script download_sra_wget_corais.sh, libs folder. I used the wget, because the curl is getting some problem in SDU. I noted that the size of the files is different:

Tabela 3.1: Comparing sizes of files

ID of metagenome	the size in NCBI site	size of file in SDU
SRR6785058	$317.00 \; \mathrm{Mb}$	318M
SRR6785057	$364.00 \; \text{Mb}$	365M
SRR6785056	$560.00 \; \mathrm{Mb}$	561M
SRR6785055	624.00 Mb	625M

So I checked the others files:

A Bia me informou que o SDU arrendonda os valores de tamanho dos arquivos, entao at o momento nao tive problemas com o download dos arquivos do mg_rast

Tabela 3.2: Comparing sizes of files 2

ID of of a reason a	the size in NCBI site	size of file in SDU	size of cleanned file
ID of metagenome	30M	29.1 MB	28M
mgm4440319.3.299.1			
mgm4440370.3.299.1	3,6M	3.5 MB	3,5M
mgm4440371.3.299.1	5,0M	4.9 MB	4,8M
mgm4440372.3.299.1	6,0M	6.0 MB	5,9M
mgm4440373.3.299.1	6,2M	6.1 MB	6,0M
mgm4440374.3.299.1	4,1M	4.1 MB	4,0M
mgm4440375.3.299.1	3.8M	3.7 MB	3.7M
mgm4440376.3.299.1	3.9M	3.9 MB	3.8M
mgm4440377.3.299.1	3,5M	3.5 MB	$3{,}4M$
mgm4440378.3.299.1	6,2M	6.2 MB	$6{,}1M$
mgm4440379.3.299.1	$7{,}0M$	$7.0~\mathrm{MB}$	$6{,}9M$
mgm4440380.3.299.1	$5{,}2M$	5.2 MB	$5{,}2M$
mgm4440381.3.299.1	$6{,}4M$	$6.4~\mathrm{MB}$	$6{,}4M$
mgm4445755.3.299.1	158M	157.0 MB	155M
mgm4445756.3.299.1	150M	149.9 MB	147M
mgm4480739.3.299.1	$8{,}0M$	7.9 MB	$7{,}9M$
mgm4480740.3.299.1	12M	11.3 MB	12M
mgm4480741.3.299.1	8,5M	8.5 MB	8,5M
mgm4480742.3.299.1	10M	12.9 MB	10M
mgm4480743.3.299.1	15M	10.0 MB	14M
mgm4484839.3.299.1	13M	14.1 MB	13M
mgm4487909.3.299.1	17M	16.5 MB	17M
mgm4487910.3.299.1	36M	35.6 MB	36M
mgm4487911.3.299.1	12M	11.4 MB	12M
mgm4516541.3.299.1	161M	$160.2~\mathrm{MB}$	163M
mgm4516694.3.299.1	193M	192.9 MB	193M
mgm4653307.3.299.1	17M	$16.0~\mathrm{MB}$	17M
mgm4694757.3.299.1	1,9G	1.8 GB	1.9G
mgm4694758.3.299.1	$2{,}2G$	$2.1~\mathrm{GB}$	$2{,}2G$
mgm4694759.3.299.1	1,7G	$1.7~\mathrm{GB}$	1,8G
mgm4694760.3.299.1	592M	1.6 GB	597M

Format Conversion of NCBI metagenomes

Adaptei o script da Bia para fazer a conversao do dos arquivos .sra Inicialmente submeti apenas um na cpu_dev para testar:

Script: teste_slurm_job_fastq_dump_corais.sh

Numero do job: *220896*

Deu certo.

O Rilquer me ajudou a criar um script que cria jobs de anotacao com o kraken2 para cada dois metagenomas.

O nome do script 'creatijobfile.sh', ele unir dois scripts: 'header' e 'ending' Adaptei para criar jobs do fastq-dump para cada 2 arquivos .sra, haja visto que no tenho uma boa ideia do quanto cada fastq-dump demorar.

Jobs submetidos dia 27/09/2018:

- job_0.sh job 221475
- job_100.sh job 221476
- job_102.sh job 221477
- job_104.sh job 221478
- job_106.sh job 221480
- job_108.sh job 221481
- job_10.sh job 221482
- job_110.sh job 221483
- job_112.sh job 221484
- job_114.sh job 221485

- job_116.sh job 221486
- job_118.sh job 221487
- job_120.sh job 221488
- job_122.sh job 221489
- job_124.sh job 221490
- job_126.sh job 221493
- job_12.sh job 221495

Jobs submetidos dia 09/10/2018:

- \bullet job_14.sh slurm 226902
- job_16.sh slurm 226903
- job_18.sh slurm 226904
- \bullet job_20.sh slurm 226905
- job_22.sh slurm 226906
- \bullet job_24.sh slurm 226907
- $job_26.sh slurm 226908$
- job_28.sh slurm 226909
- \bullet job_2.sh slurm 226910
- job_30.sh slurm 226911
- job_32.sh slurm 226912

Jobs submetidos dia 19/10/2018:

- job_34.sh slurm 231744
- $job_36.sh slurm 231745$
- \bullet job_38.sh slurm 231746
- job_40.sh slurm 231747
- job_42.sh slurm 231748
- job_44.sh slurm 231749
- $job_46.sh slurm 231750$
- job_48.sh slurm 231751

- $\bullet \ \, \text{job_4.sh}$ slurm 231752
- $\bullet \ \, \text{job_50.sh}$ slurm 231753
- $\bullet \ \, \text{job_52.sh}$ slurm 231754
- $\bullet \ \, \text{job_54.sh}$ slurm 231755

Adaptacao dos identificadores

Esse passo fez-se necessrio nas anlises anteriores, pois quando eu fazia a limpeza, os arquivos de outputs que saiam eram apenas os singletons. The output files were named as SRAXXX_good_singletons_1 and SRAXXX_good_singletons_2 and there arent any other output files beyond these and the files with bad sequences. O Pablo fez a seguinte sugesto:

```
cat SRR1275409_pass_1.fastq | sed -r 's/(SRR1275409.[0-9]+)
.([0-9]+)/\1_left/' > SRR1275409_pass_1.corr.fastq

cat SRR1275409_pass_2.fastq | sed -r 's/(SRR1275409.[0-9]+)
.([0-9]+)/\1_right/' > SRR1275409_pass_2.corr.fastq
```

Quality filter

This step is only required for NCBI metagenomes. The command line was proposed by Bia:

- trim_qual_left 25
- \bullet trim_qual_right 25

Uniformity filter (size and N bases)

7.1 Command line

Parameters:

- min_len 80
- $ns_max_p 2$
- out_format 1
- Command: nohup bash slurm_job_prinseq_single_corais_FASTA.bash & slurm_prinseq_corais.out &

Deu erro o job nohup: ignorando entrada

Location of PRINSEQ dir and scripts: /scratch/app/prinseq/0.20.4/bin srun Warning: can't run 1 processes on 21 nodes, setting nnodes to 1 srun Requested partition configuration not available now srun job 212425 queued and waiting for resources srun Force Terminated job 212425 srun Job has been cancelled srun error: Unable to allocate resources: No error srun Warning: can't run 1 processes on 21 nodes, setting nnodes to 1 srun Requested partition configuration not available now srun job 212428 queued and waiting for resources srun Force Terminated job 212428 srun Job has been cancelled

Ressubmeti o job com:

- Command: sbatch slurm_job_prinseq_single_corais_FASTA.bash

Figura 7.1: Erro no job no SDU

Profilling metagenomes

8.1 Mg-Rast metagenomes

I used the following script in the following folder:

- Folder: $scratch/ebiodiv/leticia.cavalcante/mg_rast/filtered_prinseq_good$
- Command: sbatch slurm_job_kraken2_corais.sh

The job doesn't work, o erro aparece na proxima figura

Ressubmeti o job, modificando a localizaca
o da DB do Kraken para a home do Rilquer. Numero do job
: $216410\,$

Esse problema foi resolvido modificando o endereco da base para o scratch do Rilquer.

8.2 Kraken-biom

Pasta onde est instalado kraken-biom:

/home/leticia/.local/bin

Para executar: python2.7 .kraken-biom

Executar o help do kraken-biom:

kraken-biom -h

Abrir no vim o arquivo .bashrc e inserir: export PATH=\$PATH:/home/leticia/.local/bin/kraken-biom

Executar o help do kraken-biom:

kraken-biom -h

Eu fiz um teste da etapa "Creation of BIOM table of abundances" da pipeline da bia com os seguintes passos: Na pasta /home/leticia/Documentos/libs/leticia_profiling_metagenomes:

- kraken-biom selected_file -o table.biom -max D -min P
- biom convert -i table.biom -o table.from_biom_with_taxonomy.txt -to-tsv -header-key taxonomy

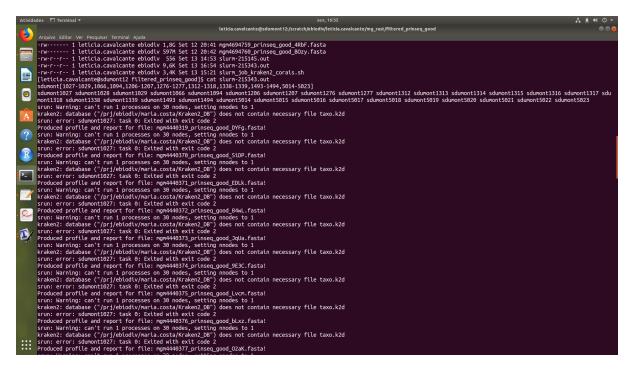


Figura 8.1: 20 erro no job no SDU

• perl filterRank.pl input table.from_biom_with_taxonomy.txt -rank p ; abundance.matrix

8.3 Teste com o kraken no scratch

Linha de teste:

perl select Groups.pl input mgm4440370_prinseq_good_SiDP.fasta_kraken.report –file_groups groups.txt $\+i$ selected_file

- First Command: $sbatch\ slurm_job_kraken2_corais.sh$
- Second Command:

 $kraken2-db\ /prj/ebiodiv/rilquer.silva/Serrapilheira \\ /Kraken2_custom_DB/\ mgm4440370_prinseq_good_SiDP.fasta$

- $-output\ mgm4440370_prinseq_good_SiDP.fasta_kraken.profiled$
- -use-names -report mgm4440370_prinseq_good_SiDP.fasta_kraken.report

Ja testei o comando acima na home do SDU e agora no scratch

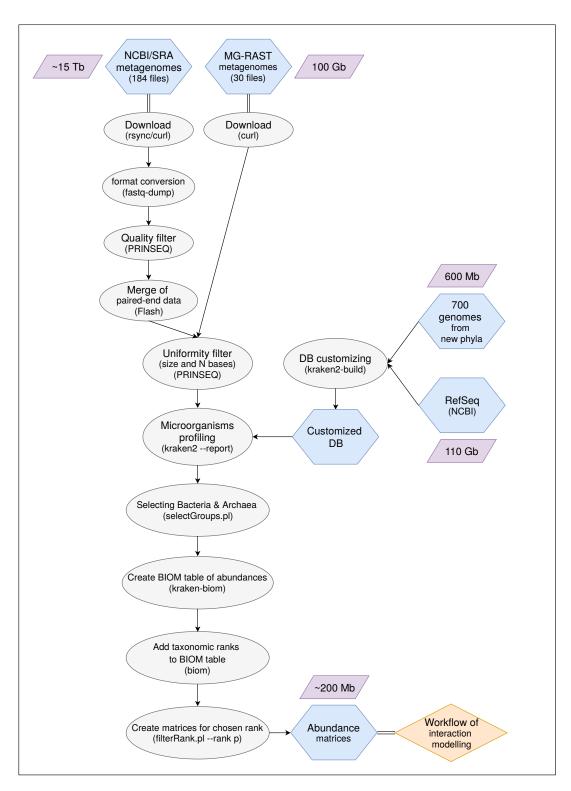


Figura 8.2: Pipeline of taxonomic annotation

8.4 Profiling no Atlantico com a ajuda do Rilquer

O Rilquer fez um script que automatiza o processo, em que a limpeza e anotacao ocorrem simultanetamente. Fiz um teste com esse script para um metagenoma com a seguinte linha:

- Comand: taxon profiling -s mgm4440378.3.299.1 -f MGRAST -k $/home/pedro/-Kraken2_custom_DB$

- Script: taxonprofiling

- Folder: /fsprofpedro/holobionts/mgrast

- Para chamar o script: taxonprofiling

Sairam 4 outputs:

- \bullet mgm4440378_kraken_class
- mgm4440378_kraken_output
- mgm4440378_kraken_report
- mgm4440378_kraken_unclass

De acordo com a pipeline da Bia, o arquivo a ser usado e o report. Teste a seguir com a pasta com os metagenomas do mg-rast inteiro:

-Comand: taxon profiling -d /fsprofpedro/holobionts/mgrast -f MGRAST -k /home/-pedro/Kraken2_custom_DB

- Folder: /fsprofpedro/holobionts

No email do Rilquer, vi que tenho que submeter o job para uma fila que no tem acesso ao fsprofpedro. Ento criei uma pasta temporaria chamada mgrast_temp, no home/pedro

- Job: profiling_metagenomes_corais_mgrast.sh
- Folder temporario em que as amostras foram copiadas: /home/pedro/m-grast_temp
- -Folder de submissao: /home/pedro/

Na pasta /fsprofpedro/holobionts, tem um exemplo de job chamado jobexample. Submissao:

-Folder de submissao: /home/pedro/mgrast_temp

- Numero: 122283.atlantico

- Command: qsub profiling_metagenomes_corais_mgrast.sh

8.5 Analises e obtencao de figuras

Apliquei o tutorial do professor para obteno de figuras no R para visualizao dos resultados.

- Script: analisys.R- Folder: /home/leticia/Documentos/libs/R

Figuras obtidas:

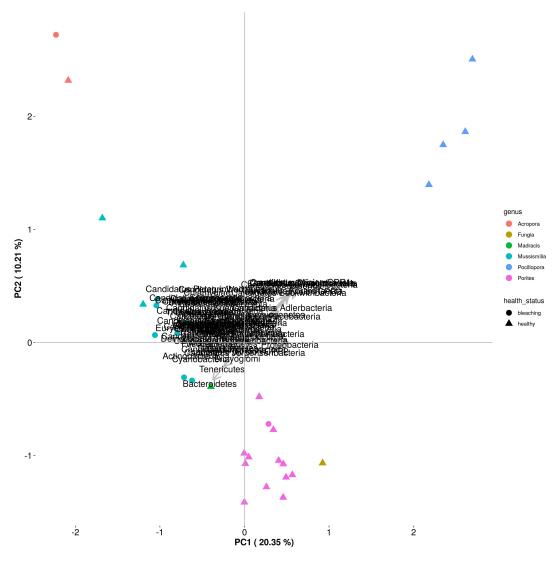


Figura 8.3: Anlise de componentes principais com todos os filos como variaveis

Na analise acima, as variaveis so muitas e ficam muito sobrepostas, fazendo com que haja grande poluicao visual. O professor recomendou em marco a utilizar uma analise de Random Forest, para que as variaveis mais importantes para os metagenomas que trabalho sejam ranqueadas. O random forest um algoritmo de machine learning que, a partir das duas categorias de sade (categorias de supervisao), elencar as variveis mais importantes para classificar as amostras nesses dois estados. o random forest abaixo su-

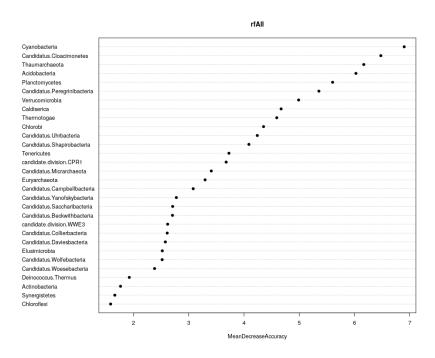


Figura 8.4: Random Forest ranqueando filos

Eu utilizei os 20 primeiros filos ranqueados para fazer o PCA. Segue esse PCA abaixo:

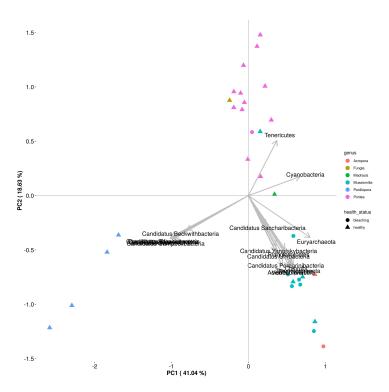


Figura 8.5: PCA com 20 filos utilizados no PCA

Uma tendencia se manteve: foi a separao das amostras por genero. Mas a poluicao visual ainda continuou, por isso fiz um random forest com os 15 primeiros filos indicados pelo random forest. Segue abaixo:

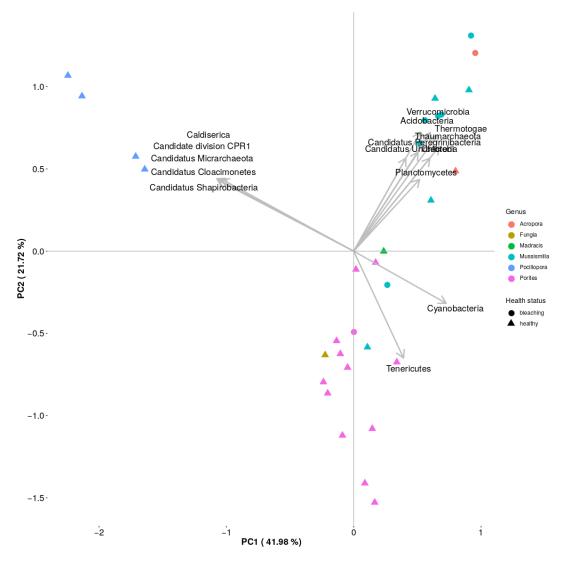


Figura 8.6: PCA com 15 filos como variaveis

Algumas tendencias tambem se mantiveram e a explicao dos eixos melhorou levemente. As amostras agrupadas no quadrante direito superior so mais diferentes das que esto no quadrante esquerdo do que das que esto no quadrante inferior direito. Na reuniao feita no dia 03/10/2018, o Amaro, o professor e Miguel me sinalizaram que existe uma separacao forte entre generos, indicando que os grupos candidatos podem ser genero específicos. Surgiu a sugesto de leitura de textos em core microbiome e específicidade de filos entre generos e o professor sugeriu fazer um random forest nao supervisionado que segue abaixo. O professor Garcia no congresso sugeriu utilizar os auto valores do PCA para ver quais podem ser mais relevantes (?).

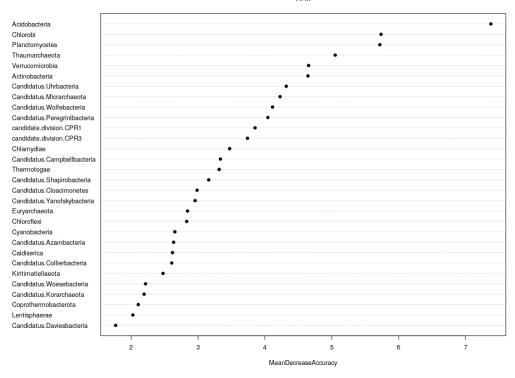


Figura 8.7: Random Forest nao supervisionado

Fiz um p
ca (libs/R/analisys.R) a partir dos primeiros 15 filos indicados no Random Forest na
o supervisionado acima. Segue abaixo:

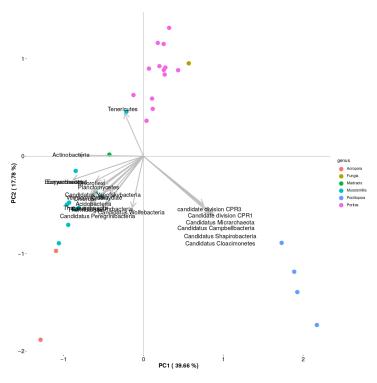


Figura 8.8: PCA a partir dos 20 filos primeiros filos que aparecem acima no random forest

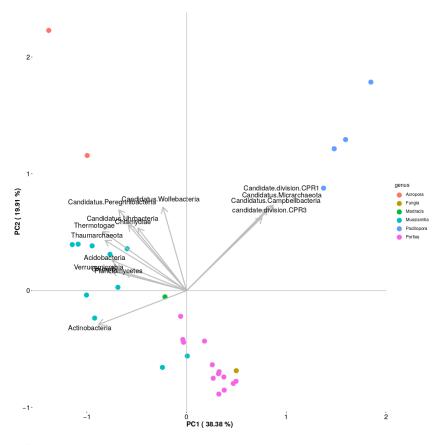


Figura 8.9: PCA a partir dos 15 filos primeiros filos que aparecem acima no random forest nao supervisionado

Functional annotation of metagenomes

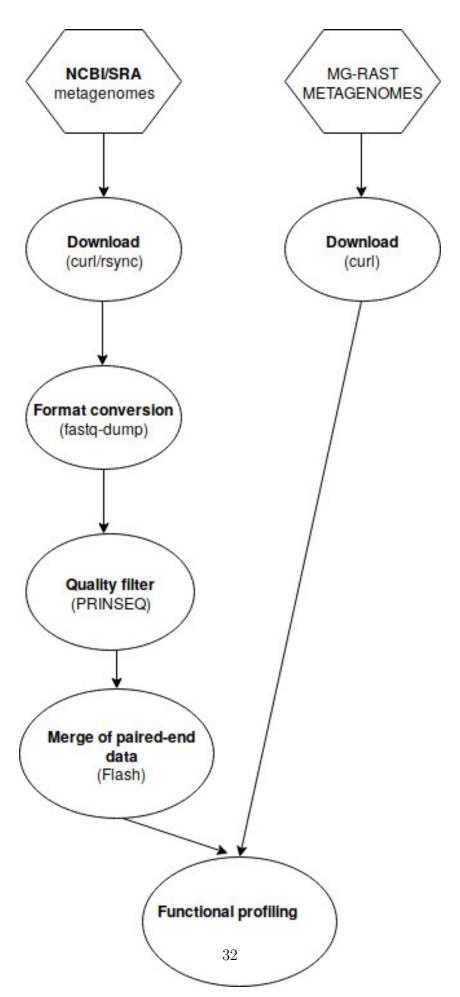


Figura 9.1: Pipeline of functional annotation

references

Articles list:

- 10.1371/journal.pone.0071301: Relata resultados que eu acreditava ter sido a primeira a encontrar
- 10.1038/nature14486: reconstruction of microorganism's genomes we use
- 10.1038/nmicrobiol.2016.48: three of life, including the Candidate Phyla Radiation
- 10.1146/annurev.micro.57.030502.090759: speaks about the uncultured majority of microorganisms
- 10.1038/ismej.2016.174: revision of rare biosphere
- 10.1038/nrmicro3400: another revision of rare biosphere
- 10.1126/science.1224041: metabolic activities of Candidatus Parcubacteria, one of super-phyla of CPR
- 10.1128/MMBR.00009-08: Revision of bioinformatic methods and steps for metagenomic
- 10.1186/s40168-018-0428-1: Sponge as holobiont. Note: This article has a important information about microbial ecology: "Network and modeling analyses aim to disentangle the strength and nature (positive, negative, or neutral) of the interactions and predict their dynamics. Bacteria-bacteria network analysis of the core microbiota in different sponge species has revealed a low connective network with very few strong and many weak unidirectional interactions (i.e., amensalism [/0] and commensalism [+/0] prevailed over cooperation [+/+] and competition [/]. These findings are consistent with mathematical models that predict that weak and non-cooperative interactions help to stabilize highly diverse microbial communities, whereas cooperation yields instability in the long term by fueling positive feedbacks"
- 10.1016/j.tim.2009.09.004: Microbial disease and the coral holobiont
- 10.3389/fmicb.2017.00618: Comparative Metagenomics of the Polymicrobial Black Band Disease of Corals

- 10.1038/nrmicro1643: The role of ecological theory in microbial ecology
- 10.1038/nrmicro3218: Explaining microbial genomic diversity in light of evolutionary ecology
- 10.1111/j.1462-2920.2009.01935.x: Metagenomic analysis of stressed coral holobionts
- 10.1038/nature06810: Functional metagenomic profiling of nine biomes
- 10.3389/fcimb.2014.00176: Microbes in the coral holobiont: partners through evolution, development, and ecological interactions
- 10.1038/ismej.2015.39: The coral core microbiome identifies rare bacterial taxa as ubiquitous endosymbionts
- 10.1111/j.1462-2920.2007.01383.x: Metagenomic analysis of the microbial community associated with the coral Porites astreoides
- 10.1038/nmicrobiol.2015.32: Metagenomics uncovers gaps in amplicon-based detection of microbial diversity
- 10.1038/ismej.2016.45: Challenges in microbial ecology: building predictive understanding of community function and dynamics
- 10.1111/j.1462-2920.2009.02113.x: Microbial functional structure of Montastraea faveolata, an important Caribbean reef-building coral, differs between healthy and yellow-band diseased colonies
- 10.1111/j.1758-2229.2010.00234.x:
- 10.1038/ismej.2011.116: Assessing the complex sponge microbiota: core, variable and species-specific bacterial communities in marine sponges

Softwares, instalacao e linhas

11.1 Profilling metagenomes

Fundamentos teoricos

meetings

Instalar o kraken-biome

Folder: /home/leticia
Command: pip install kraken-biom
Site: https://github.com/smdabdoub/kraken-biom

Para atualizacao:

git commit git push origin master

Para transferencia:

 $maquina\ remota\ para\ local:\ scp\ leticia.caval cante@login.sdumont.lncc.br:/scratch/ebiodiv/leticia.caval cantegrate.printer.gov.printer.$ /home/leticia/Documentos/dados