

Environmental Population Analysis

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May 21, 2019

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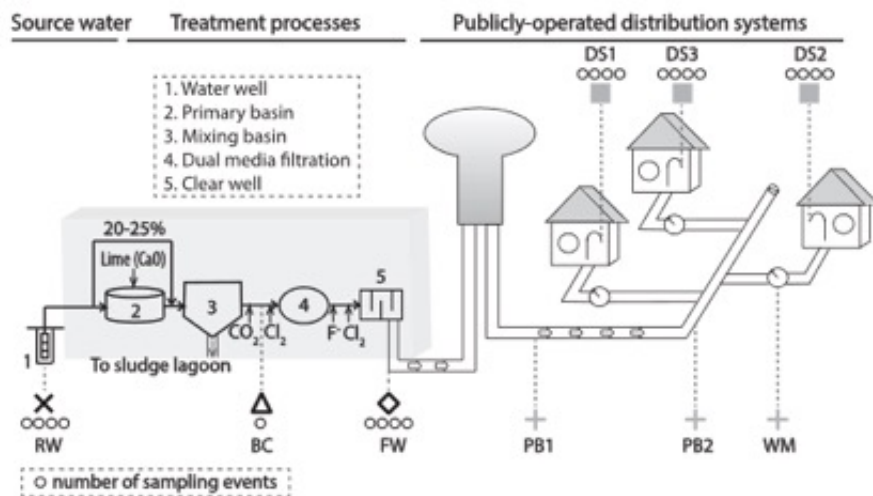


Figure 1: scheme

1 Environmental analysis

RW: Raw Water before treatment

BC: Before Chlorination

FW: Finished Water

DS1-DS3: tap water

PB1-PB2: biofilm in pipes

WM: biofilm in water meters

We are going to study the variation of microbial populations before entering the circuit (RW), at the end of the treatment (FW) and tap water samples in three houses (DS1, DS2 and DS3). We will also study samples taken in June and July.

I propose you to run the pipeline I showed you in Unit 2 whit this dataset and answer the following questions:

- 1) How many reads have we got for each sample?
- 2) Which is the trimming length you are using for the denoising step?
- 3) How many ASVs do you have before filtering deblur-table? And after filtering?
- 4) Which is the average frequency of sequences per sample before filtering deblur table?
- 5) Which is the sample with the lower number of sequences after filtering deblur table? And the one with the highest number?
- 6) Which is the most abundant phylum in each sample?
- 7) Has the study enough coverage to allow us to make any statistical inference on communities' diversity?
- 8) Studying the Unifrac Weighted PCoA plot. Is there any effect of water treatment or sampling moment on the bacteria communities?
- 9) If we compare Untreated samples (RW) vs Treated samples (FW, DS1, DS2, DS3) which are the Phyla or Classes explaining the differences among both groups? (Hint: use LEfSe)

Dataset Contents:

- 1) fastq folder: raw sample sequences
- 2) 85_otus.fasta and 85_otus_taxonomy.txt: taxonomy database
- 3) metadata.txt: sample metadata. I have included some columns useful for diversity analyses.
- 4) primers.txt: information on primers used for 16S PCR amplification
- 5) quiz.docx: this file
- 6) samplemanifest: manifest file with information of the ubication of fastq files and their corresponding tags.

1.1 Responses

- 1) How many reads have we got for each sample?

From *grep* against fastq files and verified after import on *qiime*.

File	#reads
SRR3593621_1.fastq	52945
SRR3593621_2.fastq	52945
SRR3593622_1.fastq	62218
SRR3593622_2.fastq	62218
SRR3593623_1.fastq	92740
SRR3593623_2.fastq	92740
SRR3593625_1.fastq	70366
SRR3593625_2.fastq	70366
SRR3593627_1.fastq	100615
SRR3593627_2.fastq	100615
SRR3593628_1.fastq	78495
SRR3593628_2.fastq	78495
SRR3593631_1.fastq	97332
SRR3593631_2.fastq	97332
SRR3593632_1.fastq	84361
SRR3593632_2.fastq	84361
SRR3593664_1.fastq	101827
SRR3593664_2.fastq	101827
SRR3593665_1.fastq	84850
SRR3593665_2.fastq	84850

- 2) Which is the trimming length you are using for the denoising step?

The quality of the reads is incremented substantially after the merge step by VSEARCH, consequence of a great overlap within forward and reverse sequences. In fact the filtering by means of 'qiime quality-filter q-score-joined' doesn't drop any read.

Index	Unknown	Actinobacteria	Bacteroidetes	Caldisevica	Chlorobi	Chloroflexi	Cyanobacteria	Firmicutes	Nitrospirae	OD1	OP11	OP3	Planctomycetes	Proteobacteria	Spirochaetes	WS3	Month	Group
SRR35936210.0	257.0	0.0	0.0	0.0	0.0	0.0	2847.0	0.0	0.0	0.0	0.0	0.0	183.0	29274.0	0.0	0.0	June	DS1
SRR35936210.0	65.0	0.0	0.0	0.0	0.0	0.0	5817.0	0.0	0.0	0.0	0.0	0.0	114.0	20791.0	0.0	0.0	June	DS1
SRR35936210.0	94.0	756.0	0.0	0.0	0.0	0.0	13014.0	0.0	0.0	3.0	0.0	4235.0	22161.0	0.0	0.0	June	DS2	
SRR35936213.0	27.0	96.0	0.0	0.0	0.0	0.0	5389.0	0.0	0.0	0.0	0.0	2.0	1081.0	20092.0	0.0	0.0	July	DS2
SRR35936310.0	0.0	2499.0	0.0	0.0	0.0	0.0	32169.0	0.0	0.0	0.0	0.0	0.0	662.0	6601.0	3.0	0.0	June	DS3
SRR35936312.0	0.0	51.0	0.0	0.0	0.0	0.0	21105.0	0.0	0.0	0.0	0.0	0.0	124.0	11733.0	0.0	0.0	July	DS3
SRR359366440.0	211.0	30.0	0.0	0.0	0.0	0.0	10267.0	0.0	0.0	0.0	0.0	0.0	160.0	24818.0	3.0	0.0	June	FW
SRR35936613.0	0.0	0.0	0.0	0.0	0.0	0.0	9009.0	0.0	0.0	0.0	0.0	0.0	143.0	23019.0	0.0	0.0	July	FW
SRR35936210.0	10.0	169.0	35.0	153.0	703.0	24.0	167.0	36.0	10.0	3365.0	337.0	0.0	868.0	10652.0	57.0	June	RW	
SRR35936210.0	25.0	218.0	40.0	527.0	1376.0	28.0	808.0	289.0	20.0	5803.0	879.0	0.0	4592.0	45.0	246.0	July	RW	

Figure 2: Taxonomy by phylum

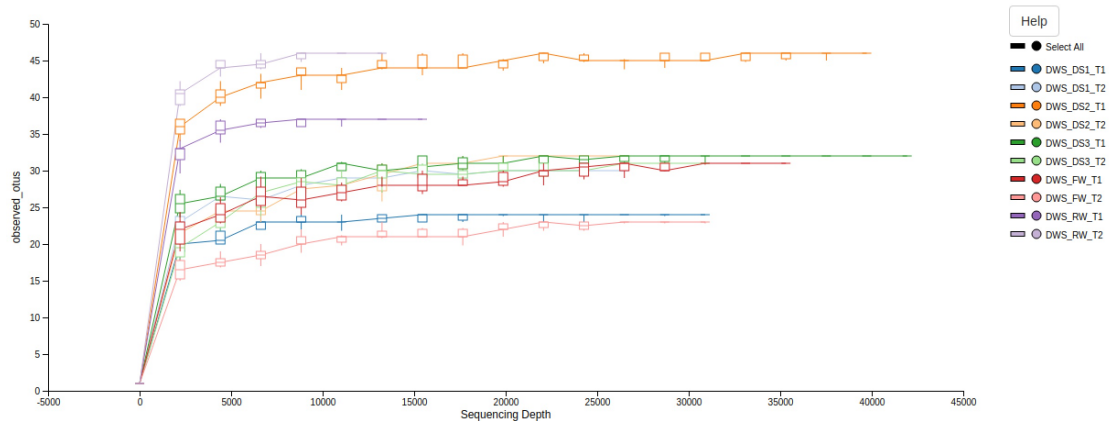


Figure 3: Rarefaction curves

I choose 372 that retains $Q \geq 38$.

3) How many ASVs do you have before filtering deblur-table? And after filtering?

Before filtering I've 765 features after 104.

4) Which is the average frequency of sequences per sample before filtering deblur table? >30,070

5) Which is the sample with the lower number of sequences after filtering deblur table? And the one with the highest number?

The sample *SRR3593622* has the lower number of sequences : 15,296 The sample *SRR3593631* has the higher number of sequences : 41,934

6) Which is the most abundant phylum in each sample?

The marked in the table of figure 2.

7) Has the study enough coverage to allow us to make any statistical inference on communities diversity? Yes, the rarefaction curves reach saturation asymptotic pattern, no more reads would give us more observed OTUs (figure 3).

8) Studying the Unifrac Weighted PCoA plot. Is there any effect of water treatment or sampling moment on the bacteria communities?

The water treatment groups cluster together nearly over axis2-axis3 plane, and the not treated water groups far along the axis1. As axis1 explains 43% of differences, we conclude that treated groups are most in common between than between the not treated groups (figure 4).

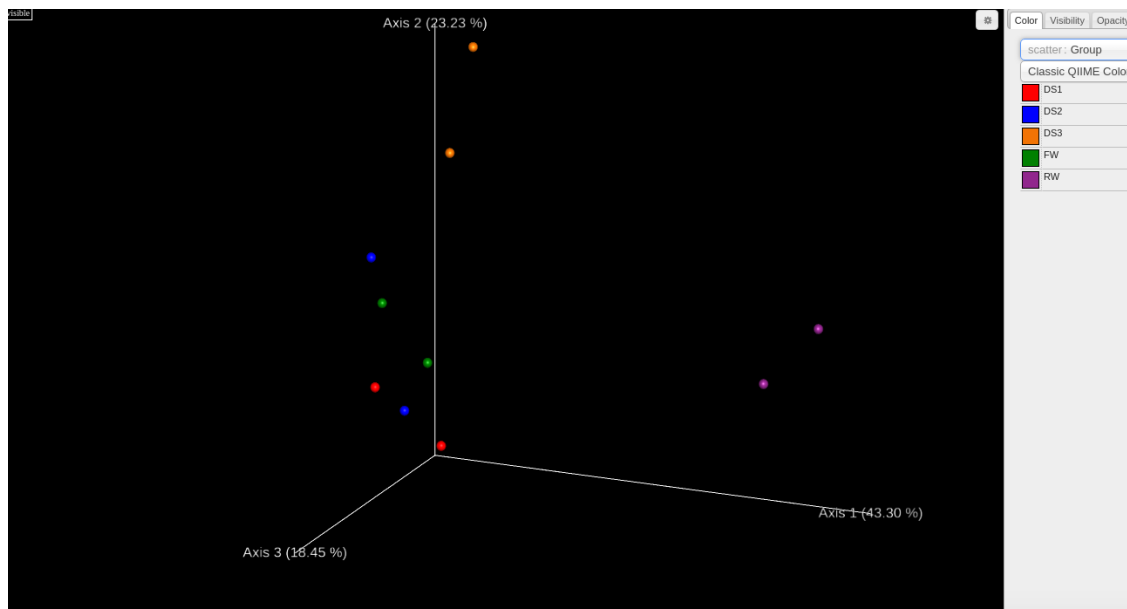


Figure 4: PCoA groups

Also the graph evidences the dissimilarity between DS3 groups and the other treated groups, above on the axis2. The biofilm presence in the distribution, surely is the explaining factor.

The difference in months are less relevant as seen in figure 5. In the case of treated groups month juny groups cluster nearer to the axis2.

- 9) If we compare Untreated samples (RW) vs Treated samples (FW, DS1, DS2, DS3) which are the Phyla or Classes explaining the differences among both groups? (Hint: use LEfSe)

The classes are drawn in the figure 6 LDA representation (TT treated and TN untrated). The most explanatory respectively are cyanobacteria and spirochaetes + OP11, coherently with the table of question 6.

Also the graph evidences the higher diversity in the not treated water, as expected.

1.2 Pipeline

1.2.1 Preprocessing and quality check

Script 1.2.1 (python)

```
1 import warnings
2 warnings.filterwarnings('ignore')
3 import pandas as pd
4 import matplotlib.pyplot as plt
5
6 FILE_ID = "SRR"
7 FASTQ_STR = "@SRR"
```

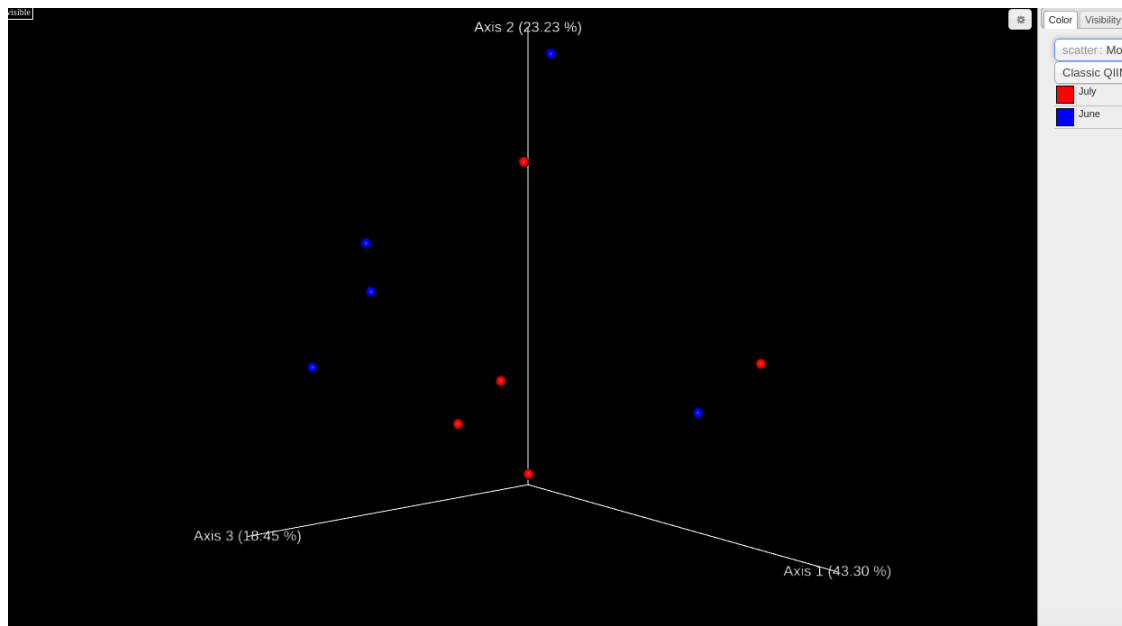


Figure 5: PCoA months

Script 1.2.2 (bash)

```
1 %bash -s "$FILE_ID" "$FASTQ_STR"
2 ssh microbiointf@192.168.56.101 env FILE_ID=$1 FASTQ_STR=$2 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 cd Documentos/Tema_2/fastq
5 head -4 ${FILE_ID}*fastq
6 grep -c $FASTQ_STR ${FILE_ID}*fastq
7 EOT
```

Output

```
==> SRR3593621_1.fastq <==
@SRR3593621.1 1 length=300
TACGTAGGGTGCAAGCGTTATCCGGATTCACTGTTCTTCCCTATTTCGTTTAGTTTTTTTTTTCCTTCTCACTTCACAGCCCTTTGCTTTACCT
  ↳ CGTCCCTCTTTTCTTCTTTACTATACTCGTTTTTATATTTGTACGTGGTTCTCCTTTTTGTTCTGTGCCTTGCGTTGTGTTCTTGTTG
  ↳ TACCCCAATTGCCATTGCTCCTTCTTCTTCATTCCCTGTCACTCTCCACGCAAGCTATCGTACTCCATCAGTTTAGTCCCCTCCTTTT
  ↳ TTCTAGCCCTCAACTCTTCCCTGCTAGTT
+SRR3593621.1 1 length=300
<6BCCGGGGGGGGGGGGGG7CEE6+,8@@,;C,,,<,<,,,,,;6,,;6,,,-,:+8+86,,,<9,:6CE,,,,,66,,,,,5:,,669
  ↳ 6,,8,89,,9,<,,9,,,:,<,95+,+++9+,9,,,,,,+,++>;74,4,+,5,,8,,8,,+66+++66,7,,,,
  ↳ ,,:5++,,+2,,,+3+74<2=2@,,+22,5,5*4+,4*/5++++3*.)*((//++(-(((*+*))3.)))+,/*/(//
  ↳ /587))6((,(*),-.)-44)43)).-
```

```
==> SRR3593621_2.fastq <==
@SRR3593621.1 1 length=300
```

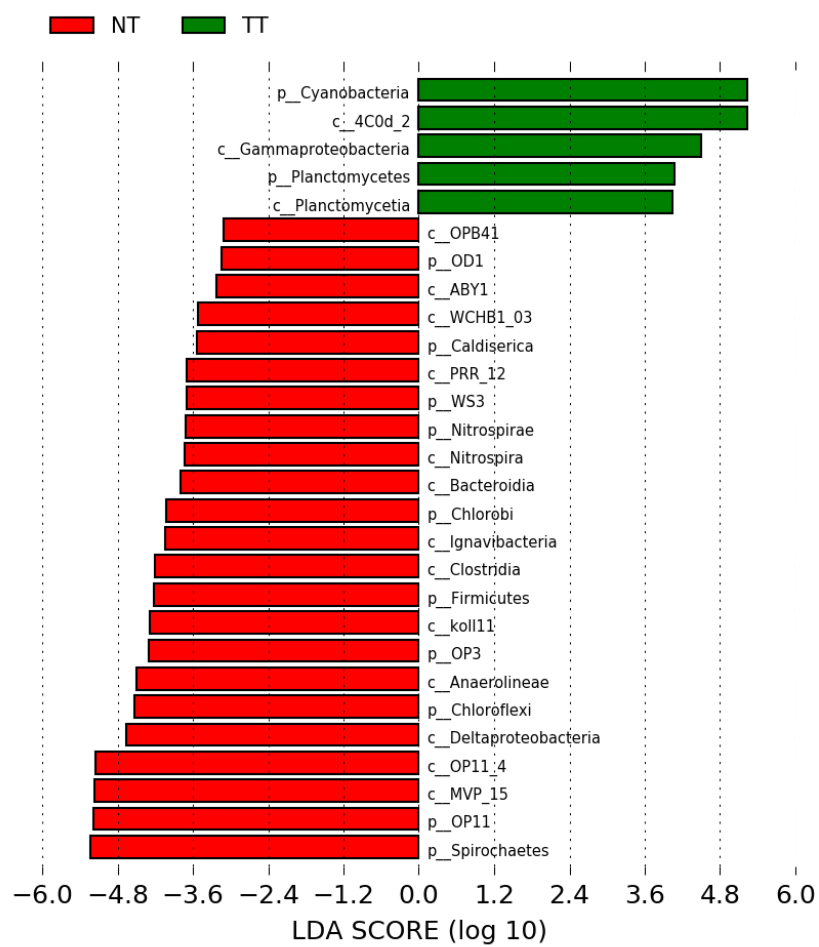


Figure 6: LDA treated/not treated

```

TTTCCTCTTTCTCTCTTCTCCTCCCCACCTTCCTCCTTCCCTCTTTCTCTTCTCCTCCTCAGATTCTCCTCCCTCCCTGCTTCCCTTTAT
↳ TTCTCCCTCTACTCCTCTCTCTCTCTGTTCTCTCCTCTACCTTTTCGTTTCTCAGTGTATCTATTTCCCTGCCGTTGCCTTCTCCT
↳ TTCTTCTTCTCCTCCTCATCTCCACTCATTTCACTTCTCCTCCATGCTTTCCCTTCTCCTCTATCACCTCCACTCTCTTCTCCTCCTCCC
↳ CCACGCCCCCTTTACGCCCCGTCCTTTCC
+SRR3593621.1 1 length=300
880--;<C@,C,-6;;6,<C@E@C,+,8,,;;,B@C,,65;,4>,,@A?,5,6;;,C6,,,,,6,,,6,,8;;,8,,,4,:5,,94,,
↳ ,,99,,,,,9,9,4,,,,,5+,3>9+934459526389+,923<9*0*09;@**3*26*++6*6++20+*)+***4)10008*03>
↳ *0*+0*3:;+:9:)*320**3*)*3;;*205:;=D*80)1*)***+001010:8;87**3*;018>5))>@E):7A*19/))*-/-
↳ -)-5(-(((((()))(((((((((//.//)

==> SRR3593622_1.fastq <==
@SRR3593622.1 1 length=300
TACGAAGGGGGCTAGCGTTGTTTCGGATTACTGGGCGTAAAGCGCACGTAGGCGGTTTTTTAAGTCAGGGGTGAAATCCCAAGGCTCAACCTT
↳ GGAAGTGCCTTTGATACTGGAAGTCTCGAGTCCGGGAGAGGTGAGTGGAATTGCTAGTGTAGAGGTGAAATTCGTAGATATTCGCAAGA
↳ ACACCACTGGCGAAGGCGGCTCACTGGCCCGGTACTGACGCTGAGGTGCGAAAGCGTGGGAGCAAACAGGATTAGATACCCTGGTAGT
↳ CCACGCCGTAAACTATGGATGCTAGCCGT
+SRR3593622.1 1 length=300
CCCC@FFGGGGGGGGGGGGGGGGGGGGFFCFFFGECFFGGGFG:+8@EFFG@FF+>BDDEE<EFC,C,,@FEGFGEE?=?,,:?DFAFFGEEE
↳ 9<,,4A?=<EFD<,A<?F9,,,:AF;?<7DF?>F+8@==FF7,,9=+F,,,:@,,8@FD9=9=:D*>D,@3=CDGGDFFGFAD>>7CC
↳ ,>,>FE,C<7<=E=@EDB75>5+2A8+*8*/211:48CE1:*775C)4>?>>B73;@DB66>>*;D5(<<07@?F66<CF86*546<A
↳ AB6<BF?0>?::<<2<62)..,3994(,,4

==> SRR3593622_2.fastq <==
@SRR3593622.1 1 length=300
TTTAACCTTGCGACCGTACTCCCCAGGCGGGATGCTTAATGCGTTAGCTGCGCCACTGAACAGCTAGCTGCCCAACGGCTAGCATCCATAGTT
↳ TACGGCGTTGACTACCACGCTATCTAATCCTCTTTTTCTCCCCACGCTTTTCGCACCTCAGCGTCAGTACCTTGCCAGTGTGCCGCTTCCG
↳ CCACTGTTGTTCTCCCGACTCTCTACGCCTTTCACCTCTCCACTCCCAGTTCCACTCTCCCCTCCCGTACTCGAGACTTCCATTTTCAA
↳ AGGCAGTTCCAAGTTTTTCGCCTTGGCATT
+SRR3593622.1 1 length=300
AC<8-;FFFFGC+7@:FFGEGEGGF@C@+++8,BFG@,,:@>CE@6EE,@76,+CC6,,,:,,4,,,:6,99C,,8=+84+,,,:94,9,,9C
↳ @,54+>68+++B;4A,,++8+<>;,+53?;5>@,24;+;@8>+3=0=0+0*0)1A)*389**53+4**/)0+***,*14)-55-7(
↳ -624.))/.;17/)),(((((/6)676((/,=2)63,4<(-343(.))5<6))66:).2(-.: (33,,(-.-1(((.4A))))))
↳ ))(((((.4))),).))),-3,()((,).

==> SRR3593623_1.fastq <==
@SRR3593623.1 1 length=300
TACGGAGGGAGCTAGCGTTGTTTCGGAATTACTGGGCGTAAAGCGCACGTAGGCGGCTTTGTGAGTTAGAGGTGAAAGCCTTGAGCTCACCTCC
↳ AGAATTGCCTTTAAGACTGCATCTCTTGAATCCAGGAGAGGTGAGTGGAATTCCTAGTGTAGAGGTGAAATTCGTAGATATTCGGATGA
↳ ACACCACTGGCGAAGGCGGCTCACTGGACTGTTATTGACGCTGCGGTGCGAAAGCTTGGGAGCAACCAGGATTAGATACCCTGGTAGT
↳ CCACGCCGTAAACCGATGATAACTAGCTGT
+SRR3593623.1 1 length=300
C@CCCGGGGGGGGGGGGGGGGDFFG7@;CE<;FC<8C@C@,CFC77+8@C:E@6@:+8=C,C,,<CC,,,,6C,,CE,,,:C,,,:@E=C,CC==
↳ ,9,,<<,<?EFA,,,<:F,,,:A,9E@,,5A@,,5,++,8:A+,,98+A,CB>=,,9CA@==,,8@+,,,@=D+@A?6@,ECC,,++53
↳ ,:,@F*4@3;55*>*9>):,4B8,*22/+314<11<*(6*)/):421*72:*15)3(413*0*)(0(,8=>?2.0<<?*/...4
↳ 9@*8(4,(42:549(3,.)).6).(46:

==> SRR3593623_2.fastq <==
@SRR3593623.1 1 length=300

```



```
TTTACTCTTGCCTGCGTACTCTCCCAAGCGTTCTCTCTTGCCTTAGCTTTCGCCACCACTCTTTCCACCCTCTTTTTCATCCTT
↳ TTTCCCCCTTTACTTCCCCCTTACCTCATCCTCTTTTCTCCCCCTCTTTTCGCTCCTCCTCTTCAATTTCTCTCCCGTTATCCTCCTTC
↳ GCCTCTCTTCTTCTTCCGAATATCTACCTATTTACCTCTACCCCCGTAATTCACCTCTCCTCTCCCGCTTCAAGCTCTGCACTCTTA
↳ CATGCACTTCTGCATTTTCAGCCCCATCCT
+SRR3593623.1 1 length=300
8@6--6CFC;E++7@+C6BF@EEEB,,98+++,,EE,,;66>B,,EF<@+,6,8+++,,5C<,,,,,69,,,,:,4,,99,99,4,,9
↳ ?@,9,,,+4,,,4,,9A,,,+*,)9+*4<,4,9?;+0+368++*03=<+*0+380)*+)+42*+4+20+*5:7+*3,)10**359*
↳ 15>*10)***+1*2/(*))))*2/*/***10*/00075*9*/((,0((//)6)743)),.6631(((((-())))))-)))(.(-5-
↳ )))),-4))-)))).)))-((((((,
==> SRR3593625_1.fastq <==
@SRR3593625.1 1 length=300
TACGTAGGGTCCAAGCGTTAATCGGAATTACTGGGCGTACAGCGTTTCGAGCGGTTTTTCAAGTCCGATGTGAAATCCCCTTGCTTAACTTT
↳ GGTATTGCTTTTGTACTGCTCGTCTTGAGTTTGTGAGAGGTGGCTAGAAATTCCTCGTGTAGCAGTGAATGCGTAGTGATGTGGTGGA
↳ ATACCGTTGGCGAAGGCAGCCCCCTGGGATAACACTGACGCTCATGCACGAAAGCTTGGGGAGCAACCAGGTTTAGTTACCCTGGTAGT
↳ CCTCGCCCTAAACGATGTCAACTAGTTGT
+SRR3593625.1 1 length=300
BBCCCGGGGGFSGGGGGG>,<FDGC7DAC6ECEFC@F@+@FCC@,88+88@+86@@,9,,,69,:66+88E,,69<C666,,:@F<:,CCC,
↳ ,,,,9<55,9A,,:,55C,,9++6A+,,,C,:B,4,,:+,+,8,,C,C?4@,7@F:@=,7,7C8,5,@,@+6=C+,7,7,6,+3+
↳ ,3,:>+@*,9:***13,3*1*1;C***0,4,,868>*(67)C+1*7(,*):3*2):0((.4*2*0(04)5756)4+<06**46.
↳ 6=*639((4.,+5((2(.)))62)64)6
==> SRR3593625_2.fastq <==
@SRR3593625.1 1 length=300
TTTACTCTTGCACCGTACTCCCCAGGCGGTCTACTTCTCGCGTTAGCTTCGTTACTACCCTCTTTCTCCCCACACCTACTTTTCATCCTT
↳ TTTGGCGTTGACTACCCCTCTCTCTATTCTCTTTTCTCCCCACTCTTTTCGTCATGACCGTCACTTTTATCCCACCTCGCTCCCTTCT
↳ CCATCTGTATTCTCTCTCATCTCTACGCCTTTTCTCTGCTACACCTCGTATTCTACCCCCCTCTGCCCCCTCTACCCCTTCTGTCTCCA
↳ ATTCAACTCCCTAGTTATCCTCCTCGCTT
+SRR3593625.1 1 length=300
8B<--6CDDCE@+7@+B6BF@E@C,,68+86,,CFFF,6,84>C,4EF,C496,@6,,,,,6,,,66?:++7,,89,,59,,5,4,,59
↳ C,,+,48+++9,,9,+++9,543,+2,47,5>?,33;+9@,+=+?++3*1)2+**6):9+)6+62*+303+***,0*)1))1)*
↳ :6*1:***3*2;****-76/8AE76;)/(0(4**1**0088*5(-((((/.)173(.3(63)))((,-,))6((,((.))--))
↳ (-)))))).),()((.))-.),((((,(
==> SRR3593627_1.fastq <==
@SRR3593627.1 1 length=300
TACGTAGGATGCAAGCGTTGTCCGATTATTGGGCGTAAAGAGTTTCGTAGGTGGTTTGTAAAGTTTGGTGTAAAGATTGGGGCTCAACCTT
↳ GATACTGCTCTTAATACTGTCTCAGACTCGAGTGTGGTAGAGGCTAGTGGAATTCAGTGTAGCGGTGAATTGCGTAGTTATTGGGATGA
↳ TCACCGGTGGCGTAGGCGACTTCTGGGCCATTACTGACACTGAGGAACGAAAGCCAGGGGAGCGAATGGGATTAGATACCCCTGTAGT
↳ CCTGGCCGTAACGTTGTCTACTAGTCGT
+SRR3593627.1 1 length=300
CCCCCGGDDGGGFGFFGGDDCCFEG@FE<CFECEC@F+,6C,;<CC@B,CCCCCBCC,:C9,EE9C<,:CFE,CC,:9,,,@CG<CCFC==
↳ ,,,,6<,6,<,,,<5C,,5,,,9A,4,+C+<57A;,,,=+4,,9,,C,CA>5,,8,C@=;>:C775,@=7@6C+@,@=,,3,,6,
↳ ,,:>8+@+;>@3**5***4;*4=;*/4*4,,,9+14<0>**2**055@752(0);.((2(3=*/)92(4??2.62<0)*1.)6
↳ 5=@).3<(4(-/66(41,).).6).)-4
==> SRR3593627_2.fastq <==
@SRR3593627.1 1 length=300
```

```

TTCAACCTTGC GCCCGTACTACTCAGGCGGGATACTTATCGCGTTAGCTGCGTCACATCACTGTTGATACCCCATACGCCTAGTATCCATCG
↳ TTTACGCCAGCACTACTGCGGTATCTAATCCCTTTGCTCCCGTTCGCTTTCGTTCCCTCCGTGTCAGTTATGCCCCAGCTCCTCGCCTA
↳ CGCCACCTTTGTCTTCCCCATATCTACGCATTTACCGCTACACTGGGATTTCCTAGCCTCTACCCCCCTCGATTCTGCCCGTTTC
↳ CAGTTCAGTTTCAGGTTTGACCCCAATC
+SRR3593627.1 1 length=300
@C@8-;CFFFFC+6@:F@FGAF9E766C@++8,BFGC@698BEE7AEE,C+,6,8,9C,,95,,6,86C,,4,:,,8?,,:594,9,+
↳ :A<+A,++6++94,A,,++3+3>3+,47=5,:=0::DBDC8*+=9900418ACB+0:*40++51*++3;?*10)*00)-7>(
↳ 7256(-50)+*109/96;3((6(1?557(,(247C37;22(49).)))(...<)44().44:72:)((-,(.64-5)((-.)
↳ 6)-.)6))..--))..-(-).3,,((.

==> SRR3593628_1.fastq <==
@SRR3593628.1 1 length=300
TACGTAGGATGCAAGCGTTGTCCGGATTTATTGGGCGTAAAGCGTTCGTAGGTGGTTTTTTAAGTTTGGTGTAAATCTTTGGCTCAACCTT
↳ GAAACTGCTCTTAATACTGTCTGACTCGAGTGTGGTAGAGGCTATTGGATTTCCAGTGTAGCGGTGAAATGCGTAGTTATTGGGAAGA
↳ TCACCGGTGGCGTAGGCGACTTCTGGGCCTTTACTGACACTGAGGAACGAAAGCCTGGGGAGCGAATCGGATTAGATACCCCTGTAGT
↳ CCTGGCCTTACACGTTGGCTACTAGTCGT
+SRR3593628.1 1 length=300
-CCCCGGFGGGFGGGGG>BCFCC,6CE;<FEFC@6C6,CC,888668,:CC,CBCC,:@,CC9C,,CFF66,,<9,,8@F5:,C9,6
↳ ,,,9,5,,<,,955C,,9,,4:44,+:<+,+9,,98+4,,9,9:A4=7,7A:C@;+B+,,@9=36+@+@,=,,+,+@
↳ ,,,>+@339@3*5*9**11*,3=,,**/*4,519+4<,0>*9**)/6(1)438).)2(1(:**9.(0(-8@28)6+3<+.**.6).
↳ 6=/.36)4),<.8(4((341:)5))(4

==> SRR3593628_2.fastq <==
@SRR3593628.1 1 length=300
TTCAACCTTGC GCCCGTACTACTCAGGCGGGATACTTATCGCGTTAGCTTCCCACTTCACTTGTCTATACCCGATACCCCTAGTATCCATCG
↳ TTTACGCCAGGACTACTGCGGTATCTACTCCCTTTGCTCCCGTGTCTTTCGTTCCCTCAGTGTGTCAGTTATCGCCCAGCTACTCGCCTT
↳ CCCCACCTTTGTCTTCCCTATATCTACGCATTTACCTCTACCTGGGAATTCCACTTCCCTCTTCCACACTCGAGTCTGCCAGTTTC
↳ CCGTGCACTTTTCAGGTTTCACCCCACTC
+SRR3593628.1 1 length=300
AC---;CFFFFC+6@FCEG,C,B9;66@:++8,BFFDA;9@CEG?6ED,C+,646,6:,,5,,9,<<6++68>,9:A,,5@=,9,+
↳ :A<+9,448++++99,85,,++34:><+,47@,,@3*::FEFEC*+5=CD;5681AEA+0<+5;+53*;*8;C**4)**00)/7<
↳ :09)/(/)(4*17CC?5>*.0*9@C6701(.9C))7.2)76)/)(((/7?))))) -44?2)65>4(4<?:((.4).53)).)-
↳ .)-(,(,(-.)))).).(-(,(,

==> SRR3593631_1.fastq <==
@SRR3593631.1 1 length=300
TACGGAGGGAGCTAGCGTTGTTCCGGAATTACTGGGCGTAAAGCGCCCGTAGGCGGCTTTTTAAGTCTGGTGTGACATCCCTGGGCTCAACCCC
↳ GGTACTGCCCTTGATACTGTCTATCTTGATTCTTGTAGTGGCGATTGGAATTCCTAGTGTAGAGGTGAAATTCGTAGTTATTCCGGAAGA
↳ TCACCAGTGGCGAAGGCGACTCCCTGGACCTGTCTTGCCGCTGAGGTTTCGACAGCTTGGGGAGCCACCCGTTTAGTTACCTGTAGT
↳ CCTCGCCGTACACGTTGTTACCTATCTTT
+SRR3593631.1 1 length=300
B@CCC@FGD7CFEFGFGGG>C@F@B@,CCC6EC@C@8C+,CC@6+77@68@,+8+8=CCE,,CC,,9C,9,<95,4,,4@E=:F44+
↳ ++++4<5,,=:,,5C,,,,8C,,,<7@,,,7,,,++83,3,:>=@,,3C,@3,,5@+,8,7>;35;@+7,@<=,***,
↳ ,3,8>,,67,34***8*)*4**2;:*.*5+49+0*(4))/2)4(.*,1)1)04(-3*+0.(04(45.:)+3<+1**5).
↳ <?*.,22(-4(*64(4((.))-.))6

==> SRR3593631_2.fastq <==
@SRR3593631.1 1 length=300

```

```

TTTACTCTTGCACCGTACTCCCCGGCGGTTCACTTATTGCGTTAGCTTCGCCACCCCCCTTCTTTCCACCCCGCCCCCTTTTTATCCTCTT
↳ TTTCCCCCTTTTCCTTCCCCCGTTTCTTATCCTCTTTTCTCCCCACGCTTTTCGCTCCCTCTCTTCTTTCTTTTCCCGCTATTCTCCTTC
↳ GCCCCTCTTTTCTTCCCACTTCTCCCCCTTTCCCTCTTCACTCGGACTTCCCCTCCCCCCCCCTCATTCCCGCTTCCCCCTTTCC
↳ ACCTCCCTTCCCGCTTTTCCCCCTCTCT
+SRR3593631.1 1 length=300
@B0--6CEDCE@+6@+B,BF@F@E+++77++6,, :FF,,6966>C7,EE,9+,6,@6=+++449C,5,, ,59=+++7+++4,,95,59,7,,7
↳ 8?,@,, ,+4,, ,37,9=,, +++,),++3573,7: :+2049>C,0*+033*0*11*)**0*3*++4+4/2*1:2**)*(+1*05.>*
↳ ((5)3,((+/.(.(.)(/))) )/(((,0..))4((,).).)(,).(. (,(-,(-,(((.(.))) )-,-,,(((-.)))
↳ ))((.(,.(.(((,(-()))))32((((,

==> SRR3593632_1.fastq <==
@SRR3593632.1 1 length=300
TACGTAGGATGCAAGCGTTGTCCGGATTTATTGGGCGTAAAGAGTTCGTAGGTGGTTTGTTAAGTTTGGTGTAAAGATTGGGGCTCAACCTT
↳ GATACTGCACTGAATACTGGCAGACTCGAGTGTGGTAGAGGCTAGTGGAATTCAGTGTAGCGGTGAAATGCGTAGATATTGGGAAGA
↳ ACACCGGTGGCGTAGGCGACTAGCTGGGCCATAACTGACACTGAGGAACGAAAGCCAGGGGAGCGAATGGGATTAGATACCCAGTAGT
↳ CCTGGCCGTAAACGATGGATACTAGGCGT
+SRR3593632.1 1 length=300
-ACCCFGGFFFGGFFFEFEGGFFEGEFCEFGGGGGGGFFCGEE@FCCFGEggFED,CECCFG<CFCFGGGGGGF,6C6,CCEG<CFG<8C
↳ <?, ,6C@9,,C,: ,?C,FE,,4?,9A=D<FF+BFFFFGFGG:9=,CFFG9FE=>5,EFGEEDFGGGFF,@,=E@CE+@=DA,BEE,AF
↳ ,@FGG6EEEDE:@FEG8C9;* ,0BFF*04*=>,8C@BF60509FC*):??EFFF)08>?66(. /:>5:FF??FFF2AGDC8585>44:
↳ <FFB:::2906CA:6<8?(.6:AA2029<

==> SRR3593632_2.fastq <==
@SRR3593632.1 1 length=300
TTCAACCTTGCGCCGTACTTCTCAGGCGGTATACTTATTGCGTTAGCTGCGCCACTCCACCTTTCGATCCCCACACCCCTTGTTCCCATCT
↳ TTTACCGCCCGGACTACTGCGGTATCTAACCCCTTTTCGCTCCCTCGCTTTTCGTTTCTCCTCGTCTTTTTCGCCAGCTCCCCCTTT
↳ CGCCTCCTGTTTTCTTCCCTCTATCTCCGTTTTTCCCTCTACCCGGTGATTTCCTCTCCCTCTCCCACTCGATTCTGCCATTTTA
↳ CAGTCCTTTTTCTGTTGTTACCCCCATC
+SRR3593632.1 1 length=300
AC6A- ,=CDFFF+7F:CFGG,E,CFC<68@++6,CFF@6,, ;>BE7@EF,67+,+8,, :,,, ,6,,,, ,96+++ ,6,4+, ,5,,4,,44,
↳ <A,9: ,++++++8+49,, ,+83+8>2+,+38++8@=*1* ;==A8***6*3304)1;C=+)5*3)++5+*3)*3*?45,))*) )-(-)
↳ 1((/(, (2)1( ).)/( /)16.5)((0(( (/8(- ( ).)(( ( ( ).--6).-((, ((-3( ,41((81(. ()))))).)--
↳ ..4.)).)).).-.( (-(-.-4(((4, .

==> SRR3593664_1.fastq <==
@SRR3593664.1 1 length=300
TACGTAGGATGCAAGCGTTGTCCGGATTTATTGGGCGTAAAGCGTTCGTAGGTGGTTTTTTAAGTTTGGTGTAAATCTTCCGGCTCCACCTT
↳ GATACTGCTCTTAATACTGTCAGACTCGAGTGTGGTAGAGGCTATTGGATTTCAGTGTAGCGGTGAATTGCGTAGTTTTTTGGATTA
↳ TCCCGTTGGCGTAGGCGACTTCTGGTCCTTTACTGACACTGAGGAACGAAAGCCTGGGTAGCGAATCGGATTAGTTCCCCCTTTAGT
↳ CCTGGCCGTACACGATGGTTCTTAGTCGT
+SRR3593664.1 1 length=300
CCCCGGG?FFGCGGGGG7C@FCF86CC,<ECCEC@F@,,C,88686@,:: ;,98C@,9@,, :<6@,, :EF<@9,,99,, ,8@F4,6C94=
↳ ,,, ,5<,9,,<,, ,9<9C,,5,, ,9A4+,+C+<,+995,, ,+,, ,9,,5, :A447,,8,8@,4++@++5,@, @3@+@+@, @=+++ ,4>
↳ ,7, :@**6*,3**1**3**14*,4=,, ,10,4,419+4>908*20**)/61*573)).2(( /,3/*)0(03(4;<<).+3<:)**-.)
↳ 6@6)(36(-3(*.4(3-(-)),. :))- ,4

==> SRR3593664_2.fastq <==
@SRR3593664.1 1 length=300

```

```

TTCAACCTTGCGCCCGTACTACTCAGGCGGTATACTTATCGCGTTAGCTTCGCCACTTCCCCCTTCTATACCCCATACCCCTTGTATCCTTCT
↳ TTTACCGCCTGCACTACTGCGGTATCTATTCCCTTTTCGCTCCCGCTCGCTTTCGTTCCCTCAGTGCTTTTCTTCCCGGCTCCTCTCCTT
↳ CGCCTCCTGTTTTATTCCCAATATCTTCGCTTTTCCCGCTACACTTGGTTTTCTCTCGCTCTACACCCCTCGATTCTCCGATTTTC
↳ CAGTGCAGTTTCAGTTTTTCTCCCCCTT
+SRR3593664.1 1 length=300
@B6--6CFECEf+,@+@+BF9F,EF;6C8+++5EFG<@,:6@CE76ED,C+,6,?,,:,,++64,,6,669,,6,9,,?:,,:,,44,9,,
↳ <:9,A,,+4++,94,9,,++3+3>+,444,,9=338:?C=C8*++60=0*151@DD+)5+60++21*2**5=D**+))*03*308*
↳ :)85)))/))5*10/1/*/7*(7)98**1/((055@*.5(4..2)6.)+))(/,/)7))1((,4,4()6)).(--(((().))..()-).)
↳ ))...6)))--)))--)))(, )66,((-,(

==> SRR3593665_1.fastq <==
@SRR3593665.1 1 length=300
TACGGAGGGAGCTAGCGTTGTTTCGGAATTACTGGGCGTAAAGCGCACGTAGGCGGCTTTGTAAGTTAGAGGTGAAAGCCTGGAGCTCAACTCC
↳ AGAATTGCCTTTAAGACTGCATCGCTTGAATCCAGGAGAGGTGAGTGGAATTCGAGTGTAGAGGTGAAATTCGTAGATATTCGGAAGA
↳ ACACCACTGGCGAAGGCGGCTCACTGGACTGGTATTGACGCTGAGGTGCGAAAGCGTGGGAGCAAACAGGATTAGATACCCCTGGTAGT
↳ CCACGCCGTAAACGATGATAACTAGCTGT
+SRR3593665.1 1 length=300
CCCCCGGGGGGGGGGGGGGGGGGGGGGDFFEFGGGGFGGFFGFEF@@@FFGGCCG7CCC,C<<FFF,C9,CFFGEF9CAE,=FFGCFFGDFD
↳ 8=,<?=AAEFE9,,C?F@Q9EE+@FD@8=FD?EF,+?<FFF798CDFGFFGFF,+8@@EFC9EGFC:8,ECF@DFG>DFGGEDC85;D
↳ 7@>FC>5DFGGGDDGF5CCED+;CFC*2=55>;=CFCGE1?DBF:D3?B:5<9?)8>>B0<>;CC>;6F?<BF7>95+3<8F6D84<<
↳ >F4>?9:69???:6?9:365)6<A<7(64<

==> SRR3593665_2.fastq <==
@SRR3593665.1 1 length=300
TTTAATCTTGCGACCGTACTCCCCAGGCGGATAACTTAATGCGTTAGCTTCGCCACCAAAGTTCTGTGAACCCTGACAGCTAGTTATCATCGT
↳ TTTTCGCGTGGACTACCAGGTATCTTATCCTGTTTCTCCTCCCGCTTTTCGCACCCCAGCGTCAATACCAGTCCAGTTATCCGCCTTC
↳ GCCACTCTTGTTCTTCCGAATATCTACGAATTTACCTCTACACTCGGAATTCCACCCACCTCTCCTGGATTTCATGCGATCCAGTCTTA
↳ AAGGCAATTCTGATTTTAGCTCCATGCT
+SRR3593665.1 1 length=300
CCCC-CFFFFFGC@CCFFGGDFEEFDDCCC7@,5FFG<,<E<EEFEGF,C@78,78<,,9AEE,<9,,C?,4,:,,95,,CA,:@,@;A
↳ ED,9++4+8+++4,,AB,,+7439:C++7===,@EE?79??@EF8*60@F+*:+15*)**05:)+;+3)+)3:77*?,,*4;5):=<3
↳ 49A*:.)*/*18**976(---/*/*5@(((27796@3273>)67@3)(233)9)7(/462,),46.((())<<))((--(-))..642
↳ <)))((().))..4))(.).4().()

SRR3593621_1.fastq:52945
SRR3593621_2.fastq:52945
SRR3593622_1.fastq:62218
SRR3593622_2.fastq:62218
SRR3593623_1.fastq:92740
SRR3593623_2.fastq:92740
SRR3593625_1.fastq:70366
SRR3593625_2.fastq:70366
SRR3593627_1.fastq:100615
SRR3593627_2.fastq:100615
SRR3593628_1.fastq:78495
SRR3593628_2.fastq:78495
SRR3593631_1.fastq:97332
SRR3593631_2.fastq:97332
SRR3593632_1.fastq:84361
SRR3593632_2.fastq:84361
SRR3593664_1.fastq:101827
SRR3593664_2.fastq:101827

```

```
SRR3593665_1.fastq:84850
SRR3593665_2.fastq:84850
```

Script 1.2.3 (bash)

```
1 %%bash
2 ssh microbiointf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6 qiime tools import --type 'SampleData[PairedEndSequencesWithQuality]' \
7     --input-path samplemanifest \
8     --output-path paired-end-demux.qza \
9     --input-format PairedEndFastqManifestPhred33
10 EOT
```

Output

Imported samplemanifest as PairedEndFastqManifestPhred33 to paired-end-demux.qza

Script 1.2.4 (bash)

```
1 %%bash
2 ssh microbiointf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6 qiime demux summarize --i-data paired-end-demux.qza --o-visualization paired-end-demux.qzv
7 EOT
```

Output

Saved Visualization to: paired-end-demux.qzv

1.2.2 Determination of ASV using Deblur

Sample pre-processing

Script 1.2.5 (bash)

```
1 %%bash
2 ssh microbiointf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6
7 # Deblur does not currently support unpaired paired-end readings,
8 #so we have to use the VSEARCH algorithm to merge the readings:
9 qiime vsearch join-pairs --i-demultiplexed-seqs paired-end-demux.qza \
```

```

10         --o-joined-sequences joined-reads.qza
11 qiime demux summarize --i-data joined-reads.qza --o-visualization joined-reads.qzv
12
13 # Filtering of the readings according to their quality.
14 qiime quality-filter q-score-joined --i-demux joined-reads.qza \
15         --o-filter-stats filt_stats.qza \
16         --o-filtered-sequences joined-filt-reads.qza
17
18 qiime demux summarize --i-data joined-filt-reads.qza --o-visualization joined-filt-reads.qzv
19
20 EOT

```

Output

```

Saved SampleData[JoinedSequencesWithQuality] to: joined-reads.qza
Saved Visualization to: joined-reads.qzv
Saved SampleData[JoinedSequencesWithQuality] to: joined-filt-reads.qza
Saved QualityFilterStats to: filt_stats.qza
Saved Visualization to: joined-filt-reads.qzv

```

Determination of ASV/features

Script 1.2.6 (bash)

```

1  %%bash
2  ssh microbioinf@192.168.56.101 /bin/bash <<"EOT"
3  export PATH=$PATH:/home/microbioinf/miniconda3/bin
4  source activate qiime2-2018.11
5  cd Documentos/Tema_2
6
7  rm -rf deblur_output
8  qiime deblur denoise-16S --i-demultiplexed-seqs joined-filt-reads.qza \
9         --p-trim-length 372 \
10         --p-sample-stats \
11         --p-jobs-to-start 2 \
12         --p-min-reads 1 \
13         --output-dir deblur_output
14
15  qiime feature-table summarize --i-table deblur_output/table.qza --o-visualization
16  ↪ deblur_output/deblur_table_summary.qzv
17
18  EOT

```

Output

```

Saved FeatureTable[Frequency] to: deblur_output/table.qza
Saved FeatureData[Sequence] to: deblur_output/representative_sequences.qza
Saved DeblurStats to: deblur_output/stats.qza
Saved Visualization to: deblur_output/deblur_table_summary.qzv

```

Script 1.2.7 (bash)

```
1 %%bash
2 ssh microbiointf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6
7 # Exclude sOTUS that have a frequency below 0.1% of the mean depth.
8 # This threshold would exclude those sOTUs that are due to Illumina sequencing errors (0.1%
  → of total reads).
9
10 qiime feature-table filter-features --i-table deblur_output/table.qza \
11                                   --p-min-frequency 30 \
12                                   --p-min-samples 1 \
13                                   --o-filtered-table deblur_output/deblur_table_filt.qza
14
15 # Exclude low frequency sOTUS
16 qiime feature-table filter-seqs --i-data deblur_output/representative_sequences.qza \
17                                --i-table deblur_output/deblur_table_filt.qza \
18                                --o-filtered-data deblur_output/rep_seqs_filt.qza
19
20
21 # Summarize
22 qiime feature-table summarize --i-table deblur_output/deblur_table_filt.qza
  → --o-visualization deblur_output/deblur_table_filt_summary.qzv
23
24 EOT
```

Output

```
Saved FeatureTable[Frequency] to: deblur_output/deblur_table_filt.qza
Saved FeatureData[Sequence] to: deblur_output/rep_seqs_filt.qza
Saved Visualization to: deblur_output/deblur_table_filt_summary.qzv
```

1.2.3 Phylogenetic distances determination using FastTree

Script 1.2.8 (bash)

```
1 %%bash
2 ssh microbiointf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6 rm -rf tree_out
7 mkdir tree_out
8 #Sequence alignment
9 qiime alignment mafft --i-sequences deblur_output/rep_seqs_filt.qza \
10                       --p-n-threads 3 \
11                       --o-alignment tree_out/rep_seqs_filt_aligned.qza
```

```

12
13 #Mask hypervariable regions
14 qiime alignment mask --i-alignment tree_out/rep_seqs_filt_aligned.qza \
15                       --o-masked-alignment tree_out/rep_seqs_filt_aligned_masked.qza
16
17 #Calculate phylogenie
18
19 qiime phylogeny fasttree --i-alignment tree_out/rep_seqs_filt_aligned_masked.qza \
20                           --p-n-threads 2 \
21                           --o-tree tree_out/rep_seqs_filt_aligned_masked_tree
22
23 #Root the tree
24
25 qiime phylogeny midpoint-root --i-tree tree_out/rep_seqs_filt_aligned_masked_tree.qza \
26                               --o-rooted-tree
27                               → tree_out/rep_seqs_filt_aligned_masked_tree_rooted.qza
28 EOT

```

Output

```

Saved FeatureData[AlignedSequence] to: tree_out/rep_seqs_filt_aligned.qza
Saved FeatureData[AlignedSequence] to: tree_out/rep_seqs_filt_aligned_masked.qza
Saved Phylogeny[Unrooted] to: tree_out/rep_seqs_filt_aligned_masked_tree.qza
Saved Phylogeny[Rooted] to: tree_out/rep_seqs_filt_aligned_masked_tree_rooted.qza

```

1.2.4 Taxonomic assignment

Assignment database training

Script 1.2.9 (bash)

```

1 %%bash
2 ssh microbiointf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbiointf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6
7 #Import reference sequences
8 qiime tools import \
9   --type 'FeatureData[Sequence]' \
10   --input-path 85_otus.fasta \
11   --output-path 85_otus.qza
12
13 #Import reference taxonomy
14 qiime tools import \
15   --type 'FeatureData[Taxonomy]' \
16   --input-format HeaderlessTSVTaxonomyFormat \
17   --input-path 85_otu_taxonomy.txt \
18   --output-path ref-taxonomy.qza

```



```
19
20 EOT
```

Output

```
Imported 85_otus.fasta as DNASequencesDirectoryFormat to 85_otus.qza
Imported 85_otu_taxonomy.txt as HeaderlessTSVTaxonomyFormat to ref-taxonomy.qza
```

Script 1.2.10 (bash)

```
1 %%bash
2 ssh microbioinf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbioinf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6
7 #Trim reference sequences to the region intra-primers according to primers.txt
8 qiime feature-classifier extract-reads \
9   --i-sequences 85_otus.qza \
10  --p-f-primer GTGCCAGCMGCCGCGGTAA \
11  --p-r-primer CCGTCAATTCMTTTRAGTTT \
12  --p-min-length 100 \
13  --p-max-length 400 \
14  --o-reads ref-seqs.qza
15
16 #Generate classifier naive-bayes
17 qiime feature-classifier fit-classifier-naive-bayes \
18   --i-reference-reads ref-seqs.qza \
19   --i-reference-taxonomy ref-taxonomy.qza \
20   --o-classifier classifier.qza
21
22 EOT
```

Output

```
Saved FeatureData[Sequence] to: ref-seqs.qza
Saved TaxonomicClassifier to: classifier.qza
```

Taxonomic assignment of representative sequences

Script 1.2.11 (bash)

```
1 %%bash
2 ssh microbioinf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbioinf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6
7 rm -rf taxa
```

```

8  #Taxonomic assignment of the representative sequences of each sOTU
9  qiime feature-classifier classify-sklearn --i-reads deblur_output/rep_seqs_filt.qza \
10 --i-classifier classifier.qza \
11 --p-n-jobs 2 \
12 --output-dir taxa
13
14 #Export to tabular file
15 qiime tools export --input-path taxa/classification.qza --output-path taxa
16
17 #Obtain interactive graph to visualize the abundance in each sOTU per sample
18 qiime taxa barplot --i-table deblur_output/deblur_table_filt.qza \
19 --i-taxonomy taxa/classification.qza \
20 --m-metadata-file metadata.txt \
21 --o-visualization taxa/taxa_barplot.qzv
22
23 EOT

```

Output

Saved FeatureData[Taxonomy] to: taxa/classification.qza
 Exported taxa/classification.qza as TSVTaxonomyDirectoryFormat to directory taxa
 Saved Visualization to: taxa/taxa_barplot.qzv

1.2.5 Diversity analysis

Alpha diversity

Script 1.2.12 (bash)

```

1  %bash
2  ssh microbioinf@192.168.56.101 /bin/bash <<"EOT"
3  export PATH=$PATH:/home/microbioinf/miniconda3/bin
4  source activate qiime2-2018.11
5  cd Documentos/Tema_2
6
7  # Obtain rarefaction curves with max-depht = num of reads of the richest sample
8  qiime diversity alpha-rarefaction --i-table deblur_output/deblur_table_filt.qza \
9                                     --p-max-depth 41934 \
10                                    --p-steps 20 \
11                                    --i-phylogeny
12                                     ↪ tree_out/rep_seqs_filt_aligned_masked_tree_rooted.qza \
13                                    --m-metadata-file metadata.txt \
14                                    --o-visualization rarefaction_curves.qzv
15
16 EOT

```

Output

Saved Visualization to: rarefaction_curves.qzv

Beta diversity

Script 1.2.13 (bash)

```
1 %%bash
2 ssh microbioinf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbioinf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
6 rm -rf diversity
7 # Obtain metrics curves with sampling-depth 19845 = num of reads of the poorest sample
8 qiime diversity core-metrics-phylogenetic --i-table deblur_output/deblur_table_filt.qza \
9     --i-phylogeny tree_out/rep_seqs_filt_aligned_maske
10     ↪ d_tree_rooted.qza
11     ↪ \
12     --p-sampling-depth 15296 \
13     --m-metadata-file metadata.txt \
14     --p-n-jobs 2 \
15     --output-dir diversity
16 EOT
```

Output

```
Saved FeatureTable[Frequency] to: diversity/rarefied_table.qza
Saved SampleData[AlphaDiversity] % Properties(['phylogenetic']) to:
  ↪ diversity/faith_pd_vector.qza
Saved SampleData[AlphaDiversity] to: diversity/observed_otus_vector.qza
Saved SampleData[AlphaDiversity] to: diversity/shannon_vector.qza
Saved SampleData[AlphaDiversity] to: diversity/evenness_vector.qza
Saved DistanceMatrix % Properties(['phylogenetic']) to:
  ↪ diversity/unweighted_unifrac_distance_matrix.qza
Saved DistanceMatrix % Properties(['phylogenetic']) to:
  ↪ diversity/weighted_unifrac_distance_matrix.qza
Saved DistanceMatrix to: diversity/jaccard_distance_matrix.qza
Saved DistanceMatrix to: diversity/bray_curtis_distance_matrix.qza
Saved PCoAResults to: diversity/unweighted_unifrac_pcoa_results.qza
Saved PCoAResults to: diversity/weighted_unifrac_pcoa_results.qza
Saved PCoAResults to: diversity/jaccard_pcoa_results.qza
Saved PCoAResults to: diversity/bray_curtis_pcoa_results.qza
Saved Visualization to: diversity/unweighted_unifrac_emperor.qzv
Saved Visualization to: diversity/weighted_unifrac_emperor.qzv
Saved Visualization to: diversity/jaccard_emperor.qzv
Saved Visualization to: diversity/bray_curtis_emperor.qzv
```

Script 1.2.14 (bash)

```
1 %%bash
2 ssh microbioinf@192.168.56.101 /bin/bash <<"EOT"
3 export PATH=$PATH:/home/microbioinf/miniconda3/bin
4 source activate qiime2-2018.11
5 cd Documentos/Tema_2
```

```
6 qiime taxa collapse --i-table deblur_output/deblur_table_filt.qza \  
7                       --o-collapsed-table deblur_output/L3_collapse_table.qza \  
8                       --p-level 3 \  
9                       --i-taxonomy taxa/classification.qza  
10  
11 qiime tools export --input-path deblur_output/L3_collapse_table.qza \  
12                       --output-path lefse_table/  
13  
14 biom convert -i lefse_table/feature-table.biom \  
15               -o lefse_table/feature-table.txt \  
16               --header-key "taxonomy" --to-tsv  
17  
18 EOT
```

Output

Saved FeatureTable[Frequency] to: deblur_output/L3_collapse_table.qza
Exported deblur_output/L3_collapse_table.qza as BIOMV210DirFmt to directory lefse_table/