



Introduction to Transcriptomics Analysis

Class 03 - Basic File Navigation with Linux



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Outline of Topics

- Exercise 1: Creating and moving into directories.
- Exercise 2: Absolute vs relative paths.
- Exercise 3: Moving, renaming and copying files.
- Exercise 4: Taking a look into the files.
- Exercise 5: Organising directories



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- Exercise 1: Creating and moving into directories.

1. Print working directory with **pwd**.
2. Using **mkdir** create a directory called `Linux_exercises`
3. Change the working directory to `Linux_exercises` with **cd**
4. Create five new directories called `exercise01`, `exercise02`, `exercise03`, `exercise04` and `exercise05`
5. List all the directories with **ls** and **ls -lh**. Discuss the differences.
6. Print the working directory.
7. Change the working directory to `exercise01`
8. Create two new directories called `test01` and `test02`
9. Eliminate the `test01` directory with **rmdir**
10. Change the working directory to home typing **cd** with no arguments.



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- **Exercise 2: Absolute vs relative paths.**

1. Change the working directory to `Linux_exercises` and then to `exercise02`.
2. Create two directories, `sample01` and `sample02`.
3. Change the directory to `sample02`.
4. Specify the absolute and relative paths of `sample01` from your current working directory.
5. Change the working directory to `home`.
6. Specify the absolute and relative path of `sample02` from your current working directory.
7. List the items for the directory `Linux_exercises` and inside this one `exercise02` using `ls` and as argument the absolute path.
8. List the items for the directory `Linux_exercises` and inside this one `exercise01` using `ls` and as argument the relative path.



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- Exercise 3: Moving, renaming and copying files.

1. Change the working directory to `Linux_exercises` and then to `exercise03`.
2. Copy the directories `sample01` and `sample02` from the `exercise02` directory using **cp**.
3. List the items from the `exercise02` directory.
4. Move the directory `test02` from the `exercise01` directory to your current working directory using **mv**.
5. List the items from the `exercise01` directory.
6. Change the name of the `test02` directory to `sample03` using the command **mv**.
7. List the items for the directory `exercise03`
8. Change your working directory to your home.
9. Create a new directory called `RNASeq`



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- Exercise 4: Taking a look into the files.

1. Copy the files “Araport11_GFF3_genes_transposons.201606.gff” and “Artha_TAIR10_genome.fasta” from the directory /storage/users/RNASEq2020UNIMI/00_DATA/00_reference into the RNASEq directory.
2. Change the working directory to RNASEq.
3. Print the first ten lines of the “Artha_TAIR10_genome.fasta” file using **head**.
4. Print the last twenty lines of the “Artha_TAIR10_genome.fasta” file using **tail -n 20**.
5. Open the file “Araport11_GFF3_genes_transposons.201606.gff” with **less**. Use the arrows up and down to navigate the file. Exit from the less screen pushing the “q” key.
6. Search the lines with the character “>” in the “Artha_TAIR10_genome.fasta” file using the command **grep “>”**.
7. Redirect the output of the command **grep “>”** on “Artha_TAIR10_genome.fasta” using the symbol “>” again (no quotes this time) into a file called “ID_list.txt”.



- Exercise 4: Taking a look into the files.
 8. Check the content of the “ID_list.txt” file using less.
 9. Ignore all the lines with the character “#” using **grep -v “#”** on the file “Araport11_GFF3_genes_transposons.201606.gff”. To avoid to print the whole file in your screen pipe the output of **grep -v “#”** Araport11_GFF3_genes_transposons.201606.gff into a **head** command using the symbol “|”.
 10. Cut the second column in the file “Araport11_GFF3_genes_transposons.201606.gff” using the command **cut -f 2**.
 11. Change the working directory to home.



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• Exercise 5: Organising directories

1. Change the working directory to RNASeq.
2. Create a directory called 00_references
3. Move the files “Araport11_GFF3_genes_transposons.201606.gff” and “Artha_TAIR10_genome.fasta” into 00_references.
4. Delete the file “ID_list.txt”.
5. Create a directory called 01_raw_reads
6. Copy all the files from the directory /storage/users/RNASeq2020UNIMI/00_DATA/01_transcriptome into the directory 01_raw_reads.
7. Use **ls -lh** to check the size that it is being use in the directory 01_raw_reads. Alternatively use **df -lh** to check the space used in the hard drive by the directory 01_raw_reads.
8. Create the directories: 02_processed_reads, 03_mapped_reads, 04_transcript_models and 05_quantification.

