Indian Institute of Technology Gandhinagar BE623 Biocomputing Sem1 2025-2026

Lab Assignment –2

Linux & Shell Scripting with Biological Data Files

Important:

- Do not copy commands from the internet your answer will be checked for correctness by running it in the lab environment.
- Some questions require looking at the actual files before deciding the correct command.
- You may combine commands using pipes ()) or use intermediate variables where needed.

Part 1 - vi Basics & File Editing

- 1. Open a new file called notes.txt in vi.
 - Insert exactly one line of text:

Have a nice day

(Make sure there is no trailing space at the end.)

- Save and exit.
- Verify that the file contains exactly one line and 15 characters.

Part 2 - Pattern Matching in FASTA Files

- 2. Display the last four lines of sequence.fasta without opening the file in an editor.
- 3. In sequence 5. fasta, print all header lines (lines starting with >).
- 4. Find all matches in sequence5.fasta where A is followed by any single character and then G.
- 5. Find all matches in sequence5.fasta where P is followed by any character except A, then L.
- 6. Print all lines in sequence 5. fasta that have exactly 2 consecutive Vs anywhere in the line.
- 7. Print all lines in sequence5.fasta that contain either AA or DD.
- 8. Print only the sequence lines (ignore headers) from sequence 5. fasta that contain the letter P.

Part 3 - Using Variables

- 9. Store the filename sequence 5. fasta in a variable called seq and print the number of sequences in it (headers count as sequences).
- 10. Store the pattern $G\setminus\{2,\setminus\}$ in a variable and search protein fasta for sequence lines (ignore headers) with 2 or more consecutive Gs.
- 11. Store "Biocomputing" in a variable, export it, and verify that it is available inside a new shell started using:

bash -c 'echo \$VARIABLE NAME'

Part 4 - File Existence & Loops

- 12. Write a shell script that checks if sequence3.fasta exists in the current folder. If yes, print the number of lines. If no, print "Missing file".
- 13. Using a for loop, go through all .fasta files in the current directory and print: filename, number of sequences, and file size in characters.
- 14. Modify the above loop so that it only prints files with more than 3 sequences.

Part 5 - Applied Data Extraction

15. From sequence5.fasta, extract only the sequence lines (no headers) that contain 3 or more cysteines (C). Save the output to a file named cys_rich.txt. Ensure the output file contains no empty lines.

Extra Challenge (Optional)

Write a single shell command that finds the file in the current directory with the largest number of sequences (by header count) and prints:

<filename> has <count> sequences

Hint: You will likely need wc, grep, sort, and head.