# BioLovesData - Gene Analysis Program

#### Overview

BioLovesData is an integrated bioinformatics analysis tool using Tkinter GUI. It provides various gene and protein sequence analysis functions.

#### **Main Features**

#### 1. Transversion Ratio Calculation

- Calculates the ratio of transitions and transversions in DNA sequences
- Input: DNA sequence files in FASTA format
- Output: Number of transitions/transversions and their ratio

## 2. Consensus Symbols Analysis

- Analyzes conservation in multiple sequence alignments
- Input: FASTA file or space-separated sequences
- Output: Conservation symbols for each position (\*: identical, :: conservative, .: variable, space: gap)

#### 3. Promoter Finder

- Detects bacterial promoter regions (-10, -35 boxes)
- Input: DNA sequences (FASTA or raw sequence)
- Options: Sequence length limit, maximum ambiguity setting
- Output: Location and sequence of discovered promoter regions

#### 4. TM Domain Finder

- Predicts transmembrane domains
- Input: Protein sequences (FASTA or raw sequence)
- Options: Hydrophobicity threshold, hydrophobicity scale selection
- Output: Location, length, and hydrophobicity values of transmembrane domains

#### 5. Multiple Sequence Alignment

- Performs multiple sequence alignment
- Input: Multiple sequences in FASTA format
- Options: Gap penalty setting
- Output: Aligned sequences and gap statistics

#### How to Use

## 1. Run the Program

Execute BioLovesData.exe (Onefile program. You can delete other files if you need only this one.)

Or

```
Bash

python3 BioLovesData.py
```

#### 2. File Selection

- Click "Select File" button in the top left to choose a file for analysis
- Supported formats: FASTA (.fasta, .fa), Text (.txt)
- Use "Preview File" button to check file contents

### 3. Analysis Function Selection

- Select the desired analysis function from the left panel
- Brief descriptions are provided for each function

### 4. Option Settings

- Related options are displayed based on the selected analysis
- Adjust parameters as needed

#### 5. Run Analysis

- Click "Run Analysis" button to start the analysis
- Results are displayed in the right panel

#### 6. Save Results

• Click "Save Results" button to save analysis results as a text file

### **Test Data**

Test data provided with the program:

- consensus test1.fasta: for MSA conservation analysis testing
- multi seq align test1.fasta: for long MSA testing
- multi seq align test2.fasta: for simple MSA testing
- multi seq align test3.fasta: for aligned sequences testing
- promoter test1 SUPER BIG... .fasta: don't run it unless you have a fancy computer
- promoter test2.fasta: for finding promoter regions
- TM domain test1.fasta: for MSA transmembrain domains prediction testing

transversion ration test1.fasta: for transition/transversion ratio calculation testing

## **System Requirements**

- Python 3.11+
- tkinter (GUI library)
- numpy (numerical computation)

#### Installation

```
Bash

# Install tkinter (Ubuntu/Debian)
sudo apt-get install python3-tk

# Install numpy
pip3 install numpy
```

## File Structure

```
Plain Text
BioLovesData/
├─ BioLovesData.exe
                                       # exe file
├─ BioLovesData.py
                                       # Main GUI program
├── calculate_transversion_ratio.py # Transversion ratio analysis
consensus_symbols.py
                                     # Consensus symbols analysis
├── find_promoter.py
                                     # Promoter finder
├── find_tm_domain.py
                                      # TM domain finder
├── multi_seq_alignment.py
                                     # Multiple sequence alignment
 — Test Data
                                       # Contains test data
    — consensus test1.fasta

    BioLovesData.pdf

                                       # User guide

    □ BioLovesData.md

                                       # Same guide in markdown format
```

### **Notes**

- Large files may take time to analyze
- FASTA file headers must start with '>'
- Protein sequences should use standard amino acid codes

## **Troubleshooting**

- If the program doesn't run: Check tkinter installation
- If analysis errors occur: Check input file format
- If GUI doesn't display: Check DISPLAY environment variable setting

## **Version History**

- v2.0: Interagated into one single program and added GUI
- v1.0: A collection of standalone Python scripts