

# **Computational Analysis of RNA-Mediated Host-Virus interactions**

## Bridge UnderGrad Science (BUGS) Summer Research Program

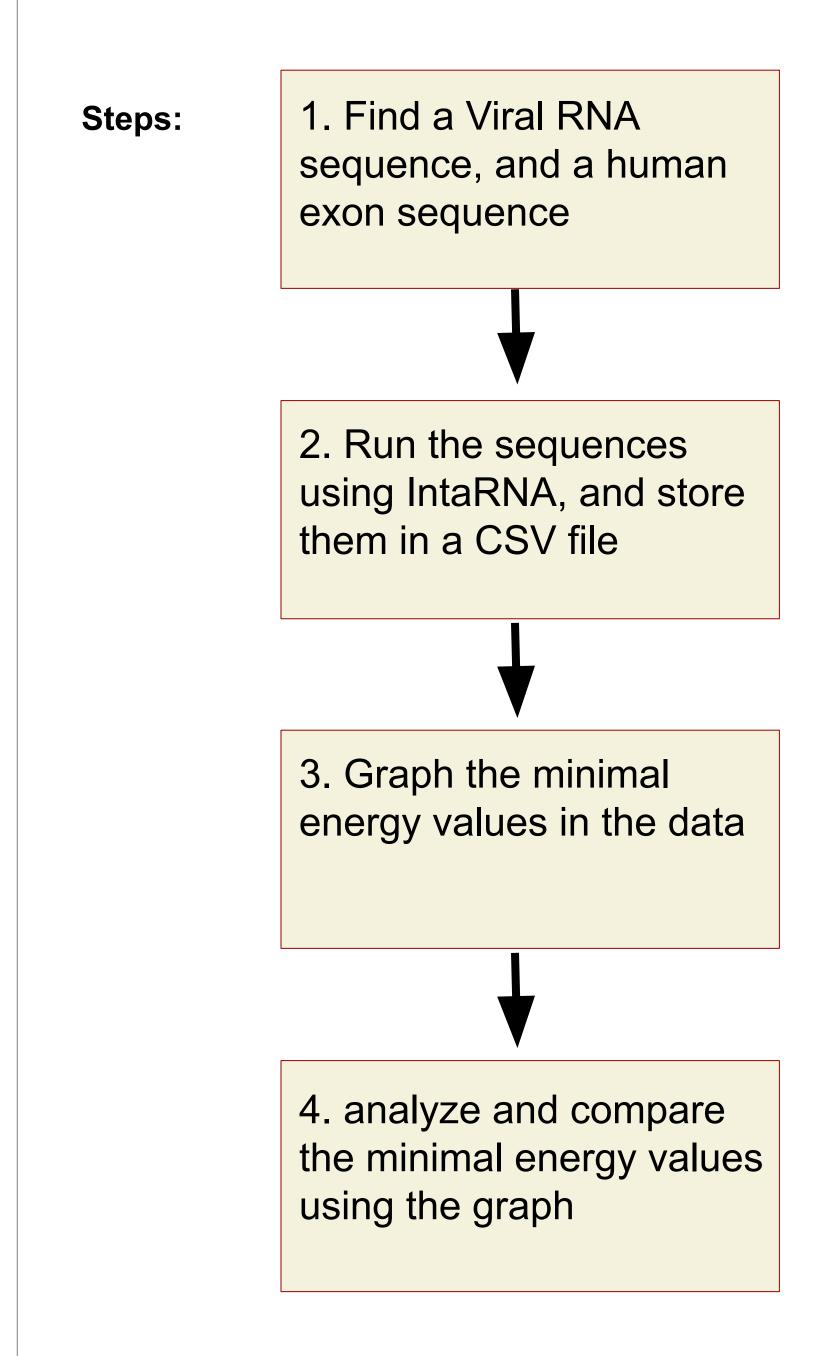
Abstract **RNA Strands** RNA-mediated host-viruses are viruses that happen from an RNA viral strand Nucleotide sequences of coding transcripts on the reference chromosom Protein-coding transcript • Transcript biotypes: protein\_coding, nonsense\_mediated\_decay, non\_stop\_decay, IG\_\*\_gene, TR\_\*\_gene, sequences polymorphic pseudogene, protein coding LoF Acidianus filamentous virus 2, complete genome NCBI Reference Sequence: NC\_009884.1 GenBank Graphics >NC\_009884.1 Acidianus filamentous virus 2, complete genome There are many difficulties that occur when studying RNA viral interactions. GACGAAAAGGAGTATTTCGCATTCCAAGAGATTAATAGATTTTTTTC ΔΑΑΤΤΑΔΑCΑΑGAGGAAAATGAAAAAGAAAAAATAGTAAAAAATTAAATGAAAGTGATGTAAAAAGCAT AACAAAAATGATAGTATTGAAGTTTTGTTGGGTAGAGTGAATTTTATTGAAGTGGAA AAATAGTCCCGATGACACGTATGGTTGTGTTGCTGAAGAGAGTTAACTATTACACGTC GTACAATGATGAATTATATGTTTTCTTCTATTCAGTCGGCTGGGTAAAATTTTCTTAGC TTTTATTATTTTTTTTTTTTTCGTTTTTTATGTTCTATTTGTTAGAATGCCTATAAA1 IntaRNA simulates RNA strand interactions and displays the energy used in rgtaatctataccgcataatgtagtagcaagtacttctttttctgtcgttacgataggatct

attaching to the host's exon strands. This leads to the viruses taking over the gene expression of the host. Many RNA viruses can cause death or severe symptoms. RNA viruses include Ebola, HIV, COVID-19, and many others. Studying these interactions between the RNA viral strand and the host's exons help us figure out which strands in the host's body to target in vaccines and other medicine. First, handling viruses is extremely difficult. Many research groups would not get authorization to handle a virus. Another problem is the amount of time these processes take. Studying RNA interactions manually is extremely time consuming, therefore limiting the amount of strands that can be analyzed. Because of the amount of time, and the restrictions of materials, our group used a software called IntaRNA. the interactions. This energy number can help us see which RNA strands to

target when trying to create a vaccine or cure for the disease. From our research, we found the interactions that took the least energy for a disease's interaction with a human gene. This can help us compare different RNA strands to see which is most suitable for analysis. This research contributes to the field of virology, informing targeted therapies, and vaccine design.

### Objectives

The objective of this research project is to simulate the interaction between a viral RNA strand and a human exon strand. By simulating this using IntaRNA we are trying to compare the minimal energy values to see which RNA strand to target when using vaccines.



### **Reyansh Gangal, Wilson Lee, Zhipeng Lu**

Dept of Pharmacy and Pharmaceutical Sciences, Bridge Institute, University of Southern California, Los Angeles, CA, USA

**CSV** Results

Name of Sequence 1	Start position of Strand	End position of Strand	Name of Sequence 2	Start position of Strand
id1	start1	end1	id2	start2
ENST0000641	401	764	Icl NC_009884.1	1
ENST0000641	382	1334	Icl NC_009884.1	27
ENST0000641	503	917	Icl NC_009884.1	3
ENST00000641	90	444	Icl NC_009884.1	3
ENST00000641	399	768	Icl NC_009884.1	2
ENST00000641	418	605	Icl NC_009884.1	3
ENST00000641	507	764	Icl NC_009884.1	1
ENST00000641	395	735	Icl NC_009884.1	4
ENST0000641	412	882	Icl NC_009884.1	25
ENST0000641	301	901	Icl NC_009884.1	1
ENST0000641	421	700	Icl NC_009884.1	6
ENST0000641	92	467	Icl NC_009884.1	1
ENST0000641	356	487	Icl NC_009884.1	10
ENST0000641	424	971	Icl NC_009884.1	10

TTATTTCATTTTCATTTTCCGTTAATTTCAATGAAATTTTAAATTCTTTAGGTCTTATTGGAATATAAT

TGTTACGTCATTATTTACAATTTCTATTTCTATAGTATTTTGGTAAAAACGCATAACAGGATAAAATAC TGGTCTAATATTTTCATTAGATCCCAGCACTGTATTTTTTTAGCATCGTCTAAAGTTATTTCTATTTCAT

> These are the results of running IntaRNA into a CSV file. These columns represent which parts of each strand are interacting and what their minimal energy value is.

