

## CALIFORNIA STATE SCIENCE FAIR 2005 PROJECT SUMMARY

Nomo(s)	Project Number
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	01210
Project Title	
RNA Alignment Scoring Based on Covariance	
Abstract	
<b>Objectives/Goals</b>	uanage. The new method
uses covariance to see how similar the 2 sequences are. I will look at the results of the tests, and other	
aspects such as time needed to make the test (It will be compared using a program I wrote) to see if this	
new method is valid.	
Methods/Materials	
The purpose of my project is to test a new method of RNA comparison to see if it#s accurate. My	
hypothesis was that it would prove to be a very effective method of comparing RNA sequences. The metorials I used for this project were: Federa Core 2 for 64 hit computers. VI (Which comes with	
Fedora Core 3) PubMed (http://www.ncbi.nlm.nib.gov/entrez/query.fogi) ClustalW	
(http://www.ebi.ac.uk/clustalw/). A program written by me for the comparison tests.	
Results	
When I ran the sequences through my program (for example, Sequence a1 and Sequence a2 I got these	
results for both comparison methods (The base comparison method results are show first, then the	
Covariance comparison results are shown):	
Sequence A 23.349056% 53.949463% Sequence P 23.141212% 50.022525%	
Sequence C 33 $163483\%$ 52 $011089\%$	
Sequence D 20.452732% Failed	
Conclusions/Discussion	
By looking at the results I have concluded that the new comparison method does not produce the same	
results and that it is not a valid form of comparison. I have reached this conclusion by observing that the	
covariance comparison results are all in the same general area when compared to the standard comparison	
results. Another factor that makes this new comparison method invalid is that it finds every spot the KNA sequence could convert with itself. So if a sequence like accession were used, it would score 100% covariance	
similarity to the sequence tata, when in reality they have nothing in common (Other than they both only	
use 2 bases#) This method is also very inefficient in comparing RNA strands because the buffer that holds	
where the RNA sequence co varies with its self grows cubically. That makes it very hard for a standard	
computer to compare long sequences (I tried a 30,000 length sequence and it failed.) This method also	
takes a very long time to compare the RNA sequences. My conclusion is that this new comparison method	
s invalid in an respects. This goes against my hypothesis which believed that it would have some validity	
I was testing a new way to compare RNA with a program I wrote	
i was testing a new way to compare KINA with a program i wrote.	
Help Received	

Inspiration for the project came from my brother Jeff.