











"Beyond the Double Helix: A Journey from DNA Recombination to Unravelling Its Computational Tapestry"

Guest Lecture

Organized by

Department of Mathematics and Natural Sciences, Universitas Pendidikan Ganesha, Bali, Indonesia

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From Bali to Malaysia



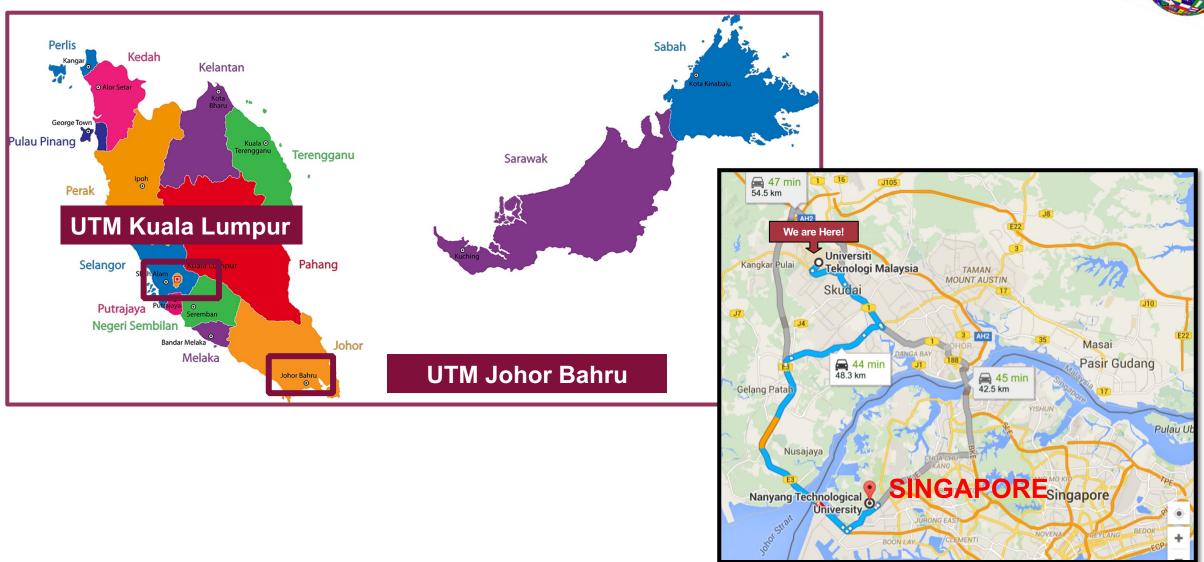
- The distance between Bali and Malaysia is estimated at 1,972.47 km or 1,225.64 miles.
- A typical flight would have an average flying time of about 3 hours.





UTM Location





12 **Faculties**

Faculties in UTM

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- Structure Materials
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- Water and Environmental Engineering

TOP 100 & #1 in MAS

Mechanical **Engineering**

- Applied Mechanics & Design
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- Materials, Manufacturing & Industrial Engineering

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• #3 in Chemistry

Biosciences

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- Management
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- Bioprocess & Polymer •#2 in Petroleum Engineering
- **Energy Engineering**



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- Electronic and Computing Engineering
- **Electrical Power** Engineering
- Control and **Mechatronics** Engineering

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- Applied Computing
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- Architecture
- **Quantity Surveying**
- **Urban and Regional** Planning
- Landscape Architecture
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Architecture/7 Built Environment **TOP 100 &** #1 in MAS

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- **Information Systems**

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- School of Education
- School of Human Resource Development & Psychology
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- Language Academy
- Centre for Advanced Studies on Islam, Science, and Civilisation
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Intelligence Informatics

Electrical and

Electronics

TOP 100

- Smart Engineering and Advanced Technology
- Business Intelligence, Humanities and Governance
- Creative Artificial Intelligence

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- **Business Administration**
- Accounting and Finance
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- Mechanical Precision Engineering
- Chemical and Environmental Engineering
- Management of Technology
- Software Engineering





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54

456

195

8

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- 1. Algebra & Analysis
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- 4. Operations Research
- 5. Statistics

Members belong to specific research groups focused on diverse studies.

(A3G) A³G

AND

Joined Department of Mathematics, Faculty of Science, UTM on 2 May 1991

MY BACKGROUND



State University of New York at Binghamton (now known as Binghamton University), New York, USA

1986 - 1989

1989 - 1990

1995 - 1998

B.Sc (Hons) Mathematics (Minor in Economics)

MA Mathematics

PhD Mathematics



"Beyond the Double Helix: A Journey from DNA Recombination to Unravelling Its Computational Tapestry"

Abstract

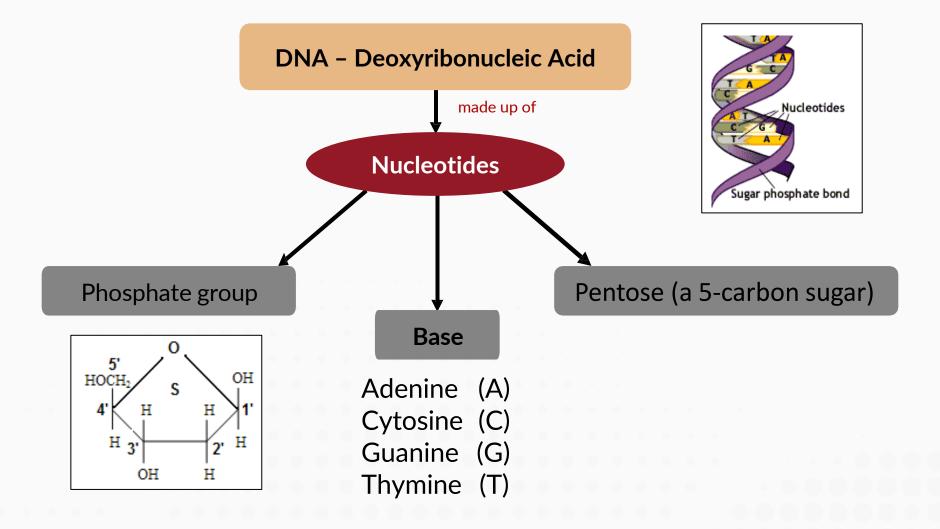
The groundbreaking work of the DNA splicing system introduced by Head in 1987 simulates the process of cutting and pasting of DNA molecules using restriction enzymes and ligases. By formal language theory, a relationship between splicing systems and molecular biology is explored. In this presentation, the mutual connections between formal language theory, biomolecular science, and the design of automated enzymatic processes and implementation within splicing systems, revealing their deterministic finite automaton structure will be shared. Additionally, experimental validation techniques are done, ensuring the theoretical results align with real-world observations.



DNA Structure



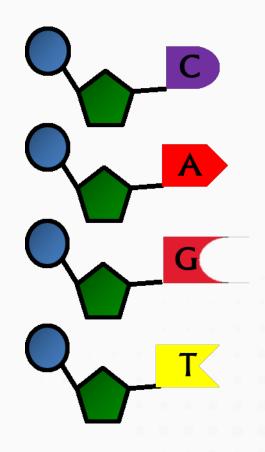
Structure of DNA

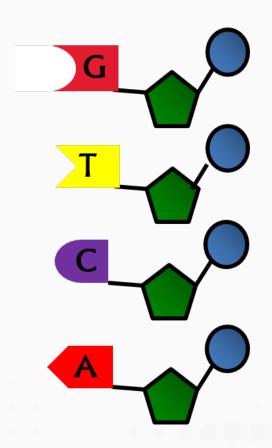




Structure of DNA (Cont.)

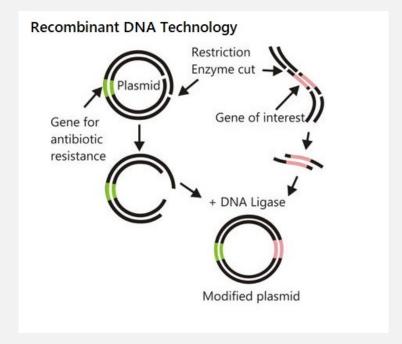
Watson-Crick Complementary







Recombinant DNA



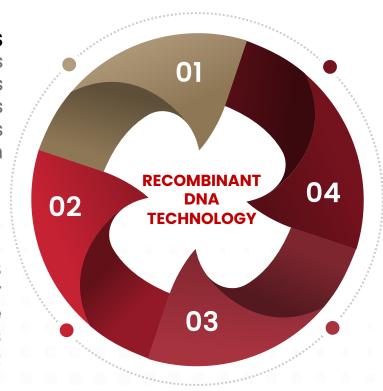
VARIOUS APPLICATIONS OF RECOMBINANT DNA TECHNOLOGY

2. THERAPEUTIC PRODUCTS

Vaccines
Growth hormones
Antibodies
Anticancer drugs
Recombinant protein

3. DIAGNOSIS

Gene therapy Monitoring device Therapeutic strategies CRISPR



1. GENETICALLY MODIFIED PRODUCTS

Fruits
Vegetables
Crops
Microbs
Animals

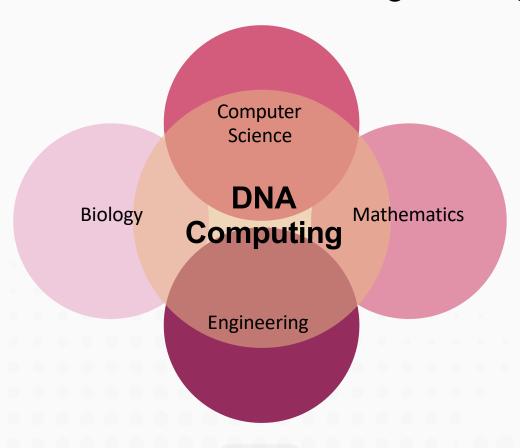
4. ENERGY APPLICATIONS

Biohydrogen Bioethanol Biomethanol Biobutanol



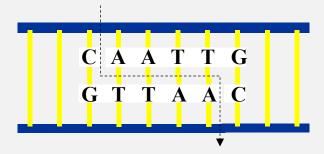
DNA Computing

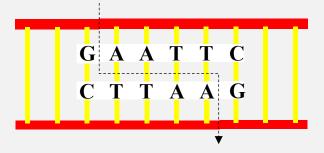
• DNA computing has emerged in the last twenty years as an exciting new research field at the intersection of Biology, Computer Science, Mathematics and Engineering.





DNA Splicing System







Mathematics and DNA Splicing System

- The mathematical modelling of splicing system was first defined by Tom Head in 1987.
- It was introduced as a mathematical model of the generative capacity of a biological system containing DNA molecules in the presence of appropriate enzymes.





Prof Head's visit to UTM, 2004

Mathematical Modelling of DNA Splicing System

How do we model it?

Mathematics and Formal Language

DNA in Mathematical Model

DNA bases

Grammar

Alphabets

h, n, i, z, a, s, m, r, o

a, c, g, t

Strings

nor & haniza & sarmin

DNA Sequence acgttgat & gcgttga

Nor + HanizaGrammar Haniza + Sarmin **DNA Splicing** acgt- -tgat & gcgt- -tga

Nor Haniza Language

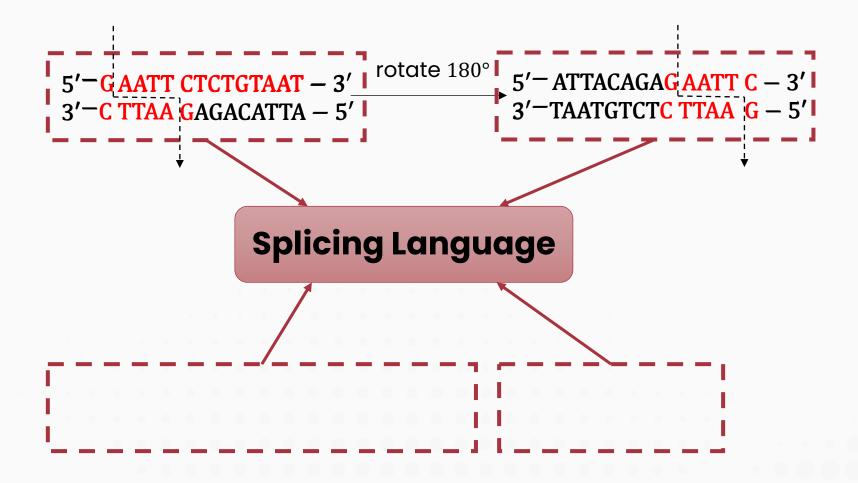
Watson-Crick

Haniza Sarmin

Language acgttga & gcgttgat



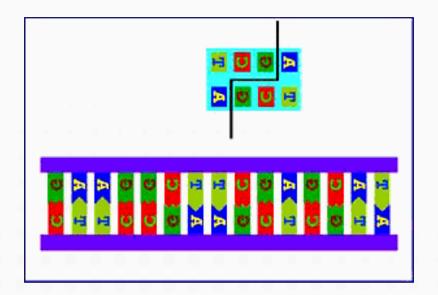
Mathematical Modelling of DNA Splicing System (Cont.)





Restriction Enzymes

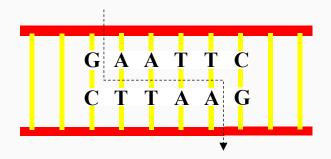
• A restriction enzyme is an enzyme that cuts double-stranded or single stranded DNA at specific recognized nucleotide sequences, known as restriction sites.



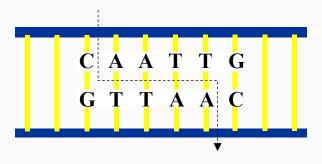


Restriction Enzymes (Cont.)

*EcoR*I: ([G/C],[A/T],[A/T],[T/A],[T/A],[C/G])



*Mfe*I: ([C/G],[A/T],[A/T],[T/A],[T/A],[G/C])





Types of DNA Splicing Systems



- Head, T. (1987). Formal Language Theory and DNA: An Analysis of the Generative Capacity of Specific Recombinant Behaviors. Bulletin of Mathematical Biology, 49(6), 737-759. doi:https://doi.org/10.1007/BF0248177
- Păun, G. (1996). On the Splicing Operation. Discrete Applied Mathematics, 70(1), 57-79. doi:https://doi.org/10.1016/0166-218X(96)00101-1 Pixton, D. (1996). Regularity of Splicing Languages. Discrete Applied Mathematics, 69(1-2), 101-124. doi:https://doi.org/10.1016/0166-218X(95)00079-7
- Goode, E., & Pixton, D. (2004). Splicing to the Limit. In N. Jonoska, G. Păun, & G. Rozenberg (Eds.), Aspects of Molecular Computing, Lecture Notes in Computer Science (pp. 189-201). Germany: Springer-Verlag.
- Yusof, Y., Sarmin, N. H., Fong, W. H., Goode, T. E., & Ahmad, M. A. (2013). *An Analysis of Four Variants of Splicing System*. Paper presented at the 20th National Symposium on Mathematical Sciences Research in Mathematical Sciences: A Catalyst for Creativity and Innovation (SKSM 2012).
- Karimi, F., Turaev, S., Sarmin, N. H., & Fong, W. H. (2014). Fuzzy Splicing Systems. In D. Hwang, J. J. Jung, & N. T. Nguyen (Eds.), Computational Collective Intelligence. Technologies and Applications, ICCCI 2014, Lecture Notes in Computer Science (pp. 20-29). Cham, Switzerland: Springer International Publishing.

Types of DNA Splicing Systems (Cont.)

Splicing System	Head S = (A, I, B, C)	Paun $\sigma = (A, R)$	Pixton $\zeta = (R, I)$	Goode-Pixton $(w, w') \vdash_r z$	Yusof-Goode S = (A, I, R)	Fuzzy S = (A, T, I, R, ⊙)
Initial String	u <mark>cxd</mark> v pexfq	u <mark>u₁u₂v</mark> u'u ₃ u ₄ v'	ξ <mark>α</mark> η ξ'α'η'	w = x u v y $w' = x' u' v' y'$	α <mark>uxv</mark> β γyzδ	(ua, x) (bv, y) $x, y \in [0, 1]$
Rule	(c, x, d) (e, x, f)	u ₁ # u ₂ \$u ₃ #u 4	$(\alpha, \alpha': \beta)$	$r = (\mathbf{u}, \mathbf{v}; \mathbf{u}'; \mathbf{v}')$	(u, x, v: y, x, z)	u#a\$b#v
Splicing Language	ucxfq pexdv	uu ₁ u ₄ v'	ξβη'	z = xuv'y'	α <mark>u</mark> xzδ γyx v β	$(uv, x \odot y)$ \odot is fuzzy operation

- S: splicing system
- σ and ζ : splicing schemes
- A: finite alphabet
- $T \subseteq A$: terminal alphabet

- *I*: set of initial strings
- B: set of rules with left pattern
- C: set of rules with right pattern
- $R = B \cup C$: set of rules



Theoretical vs Wet Lab





$$S = (A, I, B, C)$$



Theoretical vs Lab Results

Wet Splicing System involving CviQl and Acil

- An initial DNA molecule I used in this splicing model is a small segment taken from bacteriophage lambda between 42958 and 43117 with the length of 160 base pairs (bp).
- The initial molecule contains one cutting site each of the restriction enzymes *CviQI* and *AciI* where the genome locations for the cutting sites are found at 42992-42995 and 43036-43039 respectively.
- Five sticky ends of molecules α , β , γ , $\alpha \beta$ and $\beta \gamma$ are produced by the restriction enzymes when cutting the initial molecule. The lengths of fragments for the sticky ends are given in the following.

Fragment:
$$\alpha \frac{CviQl \text{ site}}{\beta} \frac{Acil \text{ site}}{\gamma}$$

$$|\alpha| = 35 \text{ bp}$$

$$|\beta| = 44 \text{ bp}$$

$$|\gamma| = 81 \text{ bp}$$

$$|\alpha - \beta| = 79 \text{ bp}$$

$$|\beta - \gamma| = 125 \text{ bp}$$



Theoretical vs Lab Results (Cont.) Wet Splicing System involving *CviQI* and *AciI*

Enzyme CviQI (palindromic)

5′...G▼TAC...3′

3'...CAT G...5'

• Enzyme *Aci*l (non-palindromic)

5′...C▼CGC...3′

3'...CGC \ C...5'



Theoretical vs Lab Results (Cont.) Wet Splicing System involving *CviQI* and *Aci*I

The splicing language from this splicing system S invoving one cutting site each of palindromic restriction enzyme CviQI (g, ta, c) and non-palindromic restriction enzyme Acil (c, cg, c) with different palindromic crossings is shown in the following:

$$L(S) = \{\alpha \operatorname{gtac}(\beta \operatorname{ccgg}\beta' \operatorname{gtac})^{n-1}(\alpha' + \beta \operatorname{ccgc}\gamma)\} + \{\gamma' \operatorname{gcg}(g\beta' \operatorname{gtac}\beta \operatorname{ccg})^{n-1}(\mathbf{c}\gamma + \mathbf{g}\beta' \operatorname{gtac}\alpha')\}.$$

where $n \in \mathbb{Z}^+$. The fragments of DNA strings in the splicing language L(S) are stated as follows:

$$\alpha \frac{\operatorname{gtac}}{\alpha} \left(\beta \frac{\operatorname{ccgg}}{\beta'} \beta' \frac{\operatorname{gtac}}{\beta'} \right)^{n-1} \alpha',$$

$$\alpha \frac{\operatorname{gtac}}{\alpha} \left(\beta \frac{\operatorname{ccgg}}{\beta'} \beta' \frac{\operatorname{gtac}}{\beta'} \right)^{n-1} \beta \frac{\operatorname{ccgc}}{\gamma'},$$

$$\gamma' \frac{\operatorname{gcg}}{\beta'} \left(\frac{g}{\beta'} \beta' \frac{\operatorname{gtac}}{\beta'} \beta \frac{\operatorname{ccg}}{\beta'} \right)^{n-1} \frac{c}{\gamma'} \gamma$$

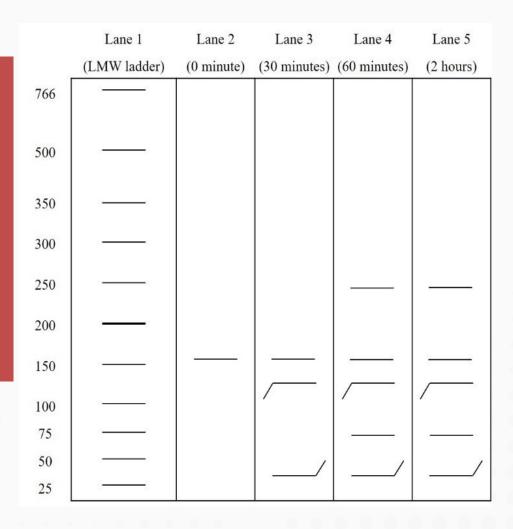
$$\gamma' \frac{\operatorname{gcg}}{\beta'} \left(\frac{g}{\beta'} \beta' \frac{\operatorname{gtac}}{\beta'} \beta \frac{\operatorname{ccg}}{\beta'} \right)^{n-1} \frac{g}{\beta'} \frac{\operatorname{gtac}}{\beta'} \alpha'$$

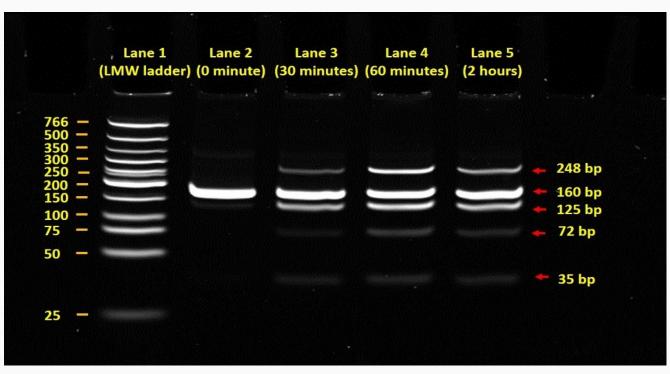
where $n \in \mathbb{Z}^+$ represents multiple copies of the specific strings.



Theoretical vs Lab Results (Cont.)

Wet Splicing System involving CviQI



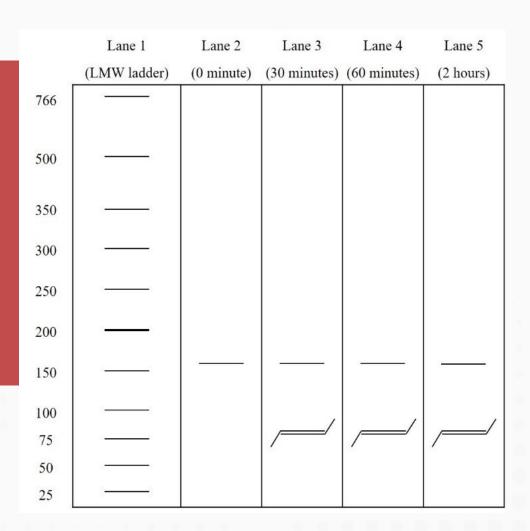


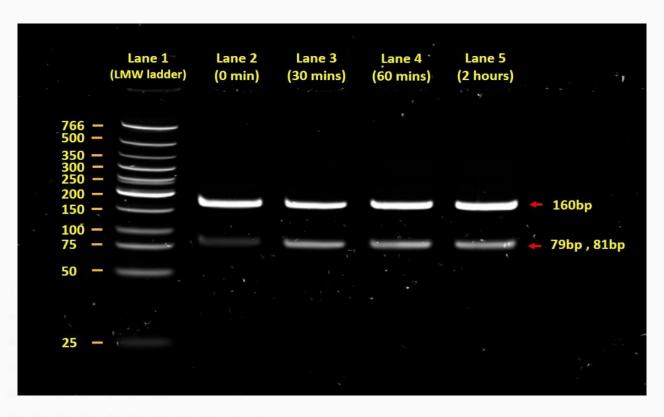
- Lane 1: LMW ladder
- Lane 2 (0 minute): 160 bp
- Lane 3 (30 minutes): 35 bp, 72 bp, 125 bp, 160 bp and 248 bp
- Lane 4 (60 minutes): 35 bp, 72 bp, 125 bp, 160 bp and 248 bp
- Lane 5 (2 hours): 35 bp, 72 bp, 125 bp, 160 bp and 248 bp



Theoretical vs Lab Results (Cont.)

Wet Splicing System involving Acil

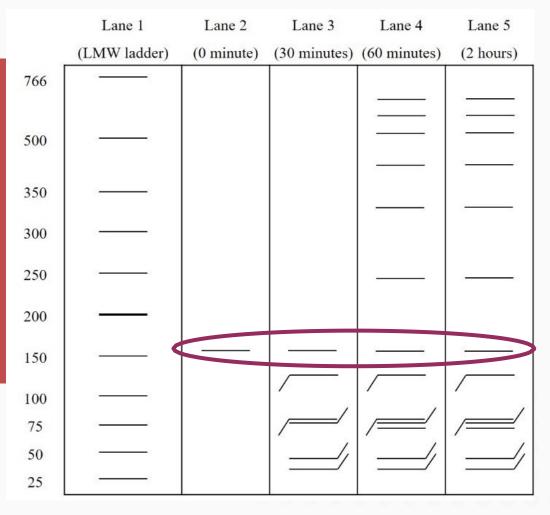


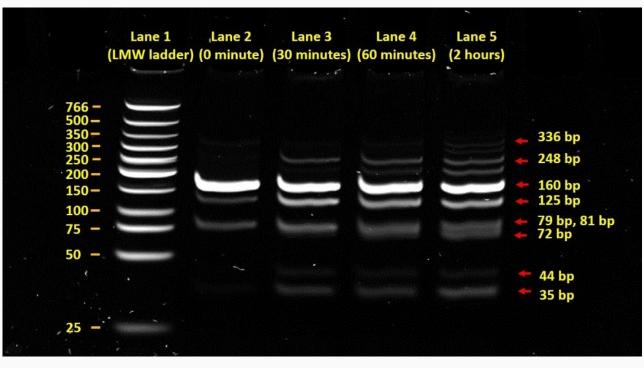


- Lane 1: LMW ladder
- Lane 2 (0 minute): 79 bp (or 81 bp) and 160 bp
- Lane 3 (30 minutes): 79 bp, 81 bp and 160 bp
- Lane 4 (60 minutes): 79 bp, 81 bp and 160 bp
- Lane 5 (2 hours): 79 bp, 81 bp and 160 bp



Theoretical vs Lab Results (Cont.) Wet Splicing System involving *CviQI* and *AciI*





- Lane 1: LMW ladder
- Lane 2 (0 minute): 79 bp (or 81 bp), 125 bp and 160 bp
- Lane 3 (30 minutes): 35 bp, 44 bp, 79 bp, 81 bp, 125 bp, 160 bp, 248 bp
- Lane 4 (60 minutes): 35 bp, 44 bp, 79 bp, 81 bp, 125 bp, 160 bp, 248 bp
- Lane 5 (2 hours): 35 bp, 44 bp, 72 bp, 79 bp, 81 bp, 125 bp, 160 bp, 248 bp, 336 bp







Universiti Teknologi Malaysia, 2007







State University of New York, Binghamton, New York, and Towson University, USA, 2010







Universiti Teknologi Malaysia, 2012







Universiti Teknologi Malaysia, 2015



Wet-lab Experiment on Splicing System





Universiti Teknologi Malaysia, 2020



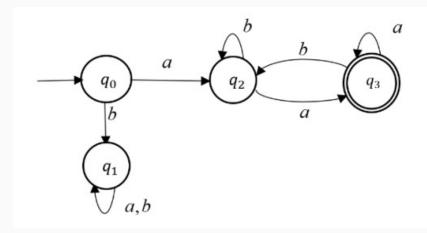
DNA Splicing in Computer Science



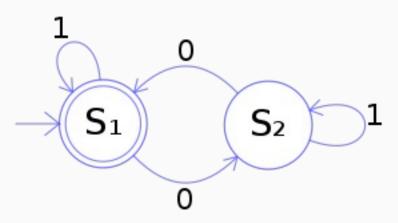


Automata of Splicing System

Automata theory is the study of <u>abstract</u> machines, as well as the <u>computational</u> <u>problems</u> that can be solved using them. It is a theory in <u>theoretical computer science</u>.



The automaton diagram for regular language



The automaton described by this <u>state</u> <u>diagram</u> starts in state S_1 , and changes states following the arrows marked 0 or 1 according to the input symbols as they arrive. The double circle marks S_1 as an accepting state. Since all paths from S_1 to itself contain an even number of arrows marked 0, this automaton accepts strings containing even numbers of 0s.

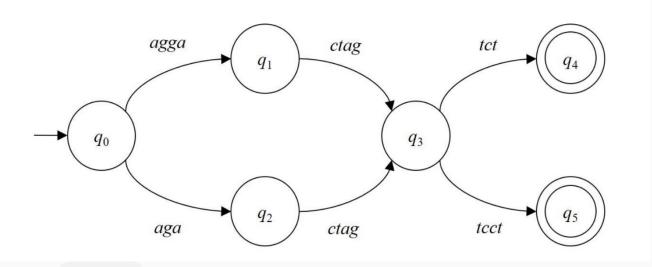


Example

Given a splicing system S = (A, I, B, C) where $I = {AGGACTAGTCT \}$ is the set of initial string, set $B = {C \setminus TA \setminus G \}$ is the set of cleavage pattern for the enzyme BfaI, and set C is the empty set.

The enzyme BfaI, $\frac{5' - CTAG - 3'}{3' - GATC - 5'}$ is a palindromic rule since the base sequence of enzyme BfaI reads the same forwards and backwards.

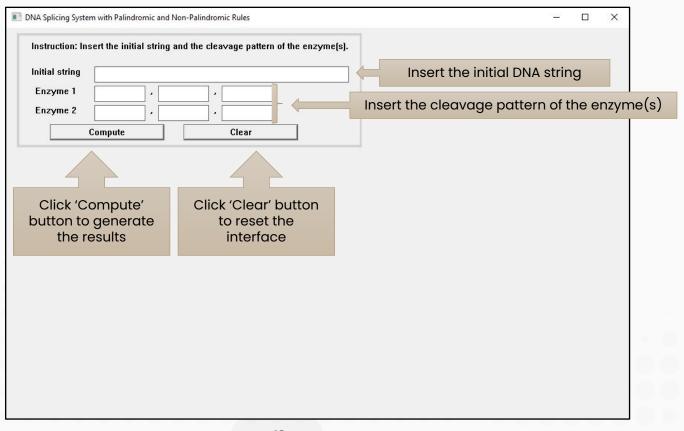
The automata for the splicing system S



Graphical User Interface (GUI) for DNA Splicing System

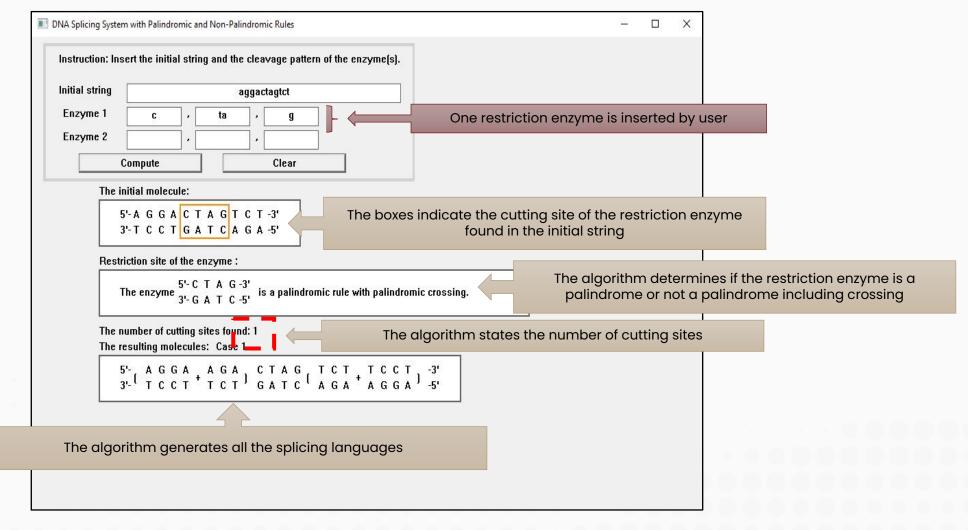
DNA Splicing Language Generator (DNASpliceGen)

A C++ program that is created in **Microsoft Visual Studio** to **develop the Graphical User Interface (GUI)** for DNA splicing systems involving palindromic and non-palindromic rules.





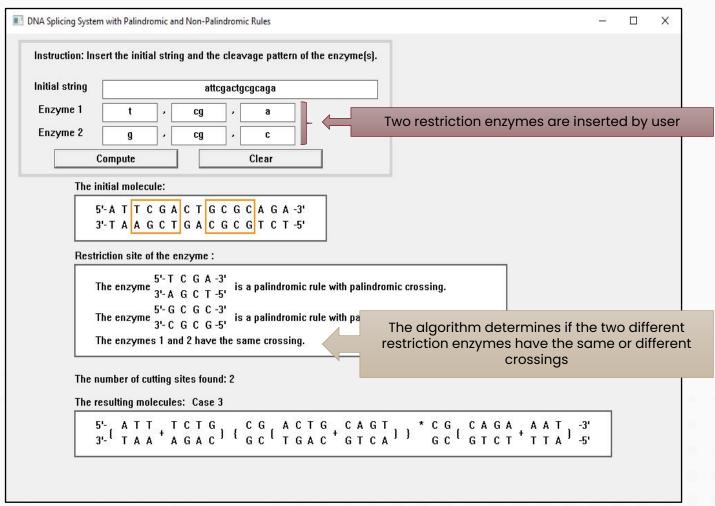
DNA Splicing Language Generator (DNASpliceGen)





DNA Splicing Language Generator (DNASpliceGen)

Output of GUI for DNA Splicing System involving Two Rules

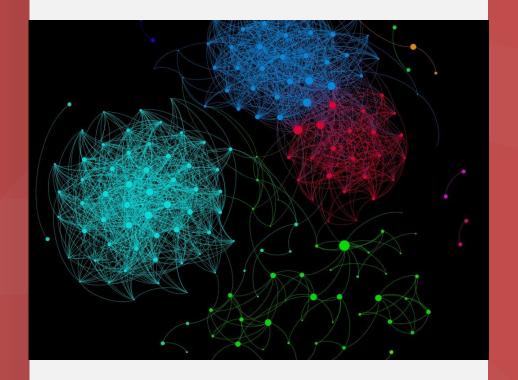


Additional features:

- Certain messages are displayed on the interface if the number of cutting sites found exceeds two
- The interface prompts the users if the cutting sites of restriction enzyme overlap
- The users will be notified if the inputs are incorrect.

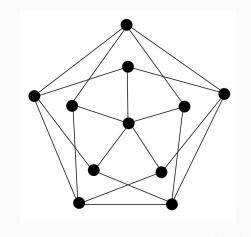


DNA Splicing in Graph Theory

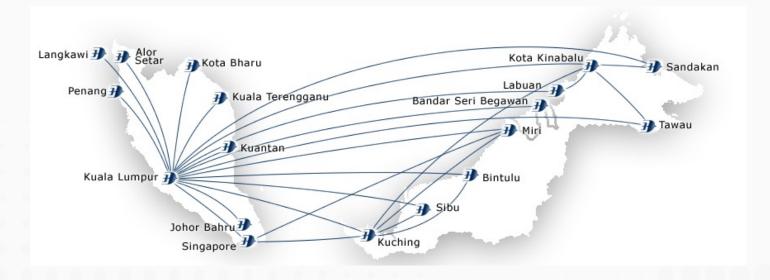


Graph Theory

A graph is a mathematical structure consists of two finite sets called the set of vertices, *V* and edges, *E*.



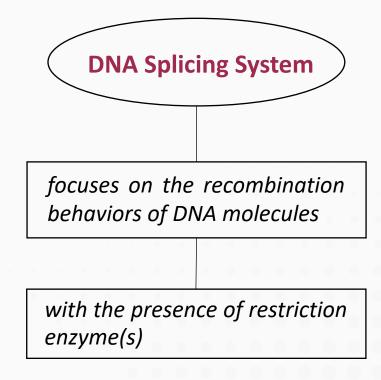


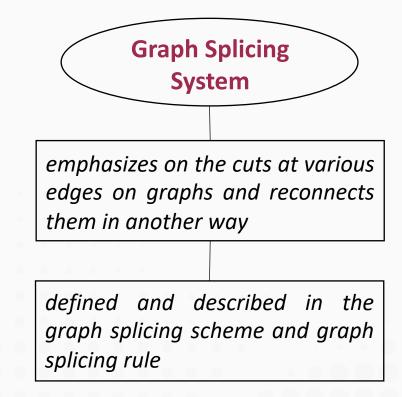


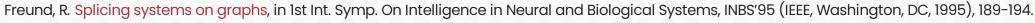


Splicing System in Graph Theory

Graph splicing system is originally introduced by Freund in 1995 to describe the DNA splicing system in the form of graphs instead of one-dimensional strings.





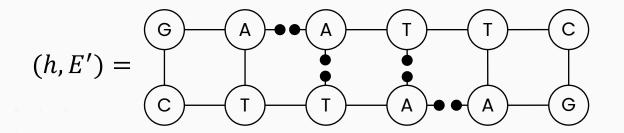


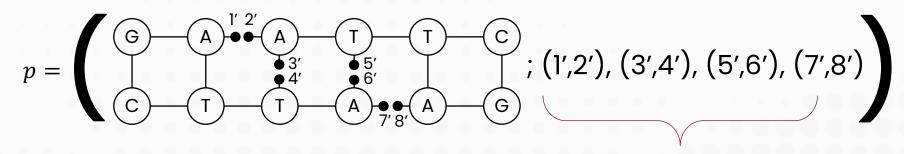
Graph Splicing Rule

• A graph splicing rule consisting the enzyme AclI can be written as follows.

$$p = ((h, E'); R)$$

 $ECORI: {AA CG TT \\ TT' GC' AA}$





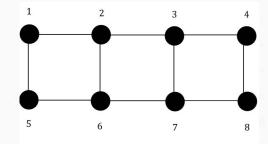


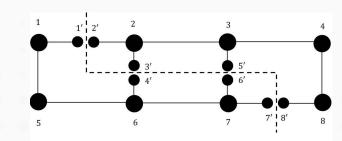
n-Cut Splicing (Cont.)

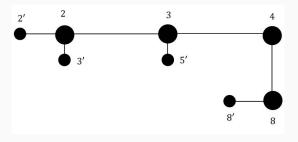
Semigraph representation of DNA molecule

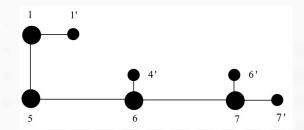
An *n*-cut splicing is applied

Two components of *n*-cut spliced semigraphs are generated





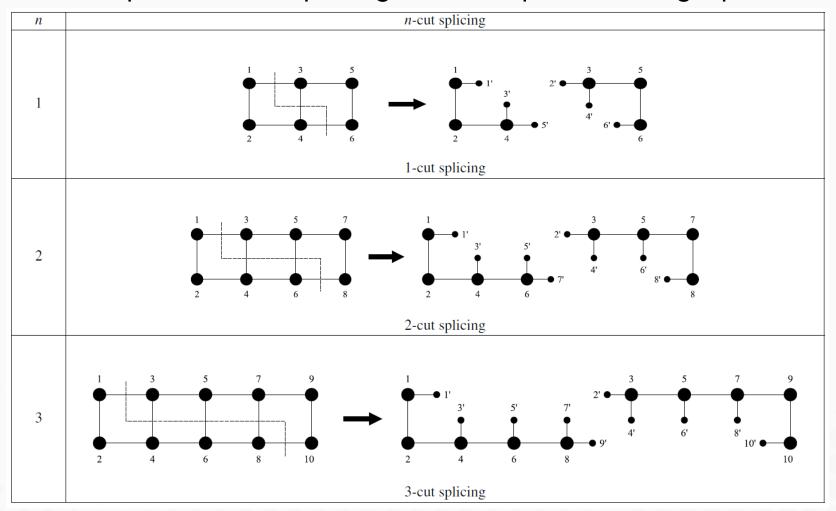






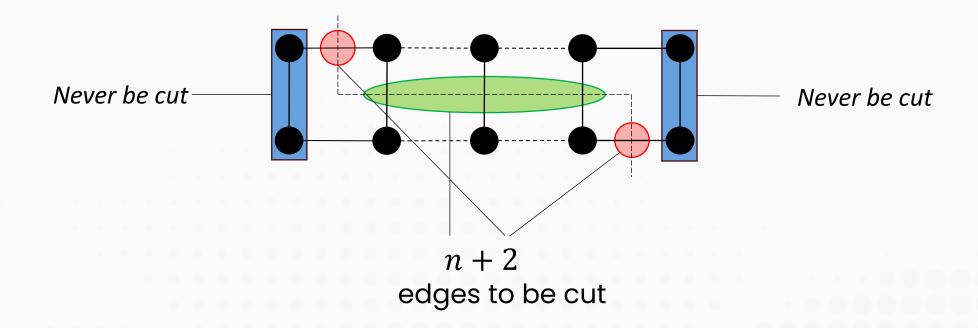
n-Cut Splicing (Cont.)

Example of *n*-cut splicing & *n*-cut spliced semigraph



n-Cut Splicing (Cont.)

An *n*-cut splicing will cut *n*+2 number of edges and the two vertices from the left most of the graphs as well as the two vertices from the right most of the graphs will never be cut.





Spectral Graph Partitioning

DNA sequence GTACCGCGTACA of length 12

The DNA sequence GTACCGCGTACA has the unit distance path graph as shown in Figure 4.1.

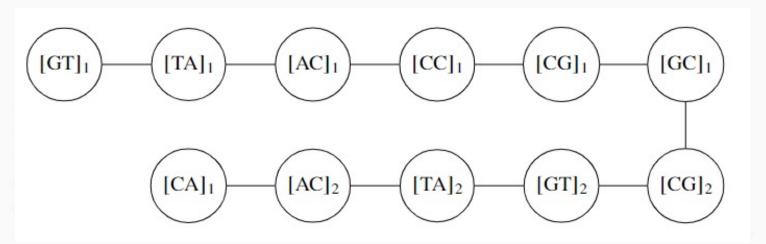


Figure 4.1: Unit distance path graph of DNA sequence GTACCGCGTACA.



DNA sequence GTACCGCGTACA of length 12 (Cont.)

The graph in Figure 4.1 is denoted as G_1 , Laplacian matrix of the graph, $L(G_1)$ is obtained.



DNA sequence GTACCGCGTACA of length 12 (Cont.)

 \succ The eigenvalues and eigenvectors of $L(G_1)$ are computed, as shown below.

Table 4.1: Set of eigenvalues and eigenvectors of $L(G_1)$.

	Eigenvalue,	ie, Eigenvector,						
ı	λ_i	v_i						
1	0	$(1 \ 1 \ 1 \ 1 \ 1 \ 1 \ 1 \ 1 \ 1)^T$						
2	0.0810	$\begin{pmatrix} -1 & -0.9190 & -0.7635 & -0.5462 & -0.2846 & 0 & 0.2846 & 0.5462 & 0.7635 & 0.9190 & 1 \end{pmatrix}^T$						
3	0.6903	$\begin{pmatrix} -1 & 0.3097 & 0.5944 & 1.0822 & 0.8308 & 0 & -0.8308 & -1.0882 & -0.5944 & 0.3097 & 1 \end{pmatrix}^T$						
4	1.7154	$\begin{pmatrix} -1 & 0.7154 & 1.2036 & -0.3728 & -1.3097 & 0 & 1.3097 & 0.3728 & -1.2036 & -0.7154 & 1 \end{pmatrix}^T$						
5	2.8308	$\begin{pmatrix} -1 & 1.8308 & -0.5211 & -0.5211 & -1.3979 & 1.6825 & 0 & -1.6825 & 1.3979 & 0.5211 & -1.8308 & 1 \end{pmatrix}^T$						
6	0.3175	$\begin{pmatrix} 1 & 0.6825 & 0.1483 & -0.4330 & -0.8768 & -1.0422 & -0.4330 & 0.1483 & 0.6825 & 1 \end{pmatrix}^T$						
7	3.6825	$\begin{pmatrix} -1 & 2.6825 & -3.5133 & 3.2287 & -1.9190 & 0 & 1.9190 & -3.2287 & 3.5133 & -2.6825 & 1 \end{pmatrix}^T$						
8	1.1692	$\begin{pmatrix} 1 & -0.1692 & -1.1406 & -0.7784 & 0.4938 & 1.1887 & 0.4938 & -0.7784 & -1.1406 & -0.1692 & 1 \end{pmatrix}^T$						
9	2.2846	$\begin{pmatrix} 1 & -1.2846 & -0.6344 & 1.4652 & 0.2173 & -1.5270 & 0.2173 & 1.4652 & -0.6344 & -1.2846 & 1 \end{pmatrix}^T$						
10	3.3097	$\begin{pmatrix} 1 & -2.3097 & 2.0251 & -0.3426 & -1.5764 & 2.4072 & -1.5764 & -0.3426 & 2.0251 & -2.3097 & 1 \end{pmatrix}^T$						
11	3.9190	$\begin{pmatrix} 1 & -2.9190 & 4.6015 & -5.9112 & 6.7420 & -7.0267 & 6.7420 & -5.9112 & 4.6015 & -2.9190 & 1 \end{pmatrix}^T$						



DNA sequence GTACCGCGTACA of length 12 (Cont.)

- > From Table 4.1, the **Fiedler value** obtained is 0.0810.
- ➤ The Fiedler vector obtained is
 (-1 -0.9190 -0.7635 -0.5462 -0.2846 0 0.2846 0.5462 0.7635 0.9190 1)^T.
- For **bisection cut**, the splitting value s is the median of the entries of Fiedler vector, which is s=0 in this case. Hence, the partitions of DNA sequence GTACCGCGTACA obtained are **GTACCG** and **CGTACA** with equal length of 6.
- For **gap cut**, the splitting value s is the largest gap of the sorted entries of Fiedler vector, which is 0.2846 in this case. Hence, the partitions of DNA sequence GTACCGCGTACA are **GTACCGC** (length 7) and **GTACA** (length 5), which are of unequal lengths.
- Therefore, bisection cut is the balanced cut for GTACCGCGTACA.



Shortest Path Problem and Minimized DNA String

Case of Initial Base A

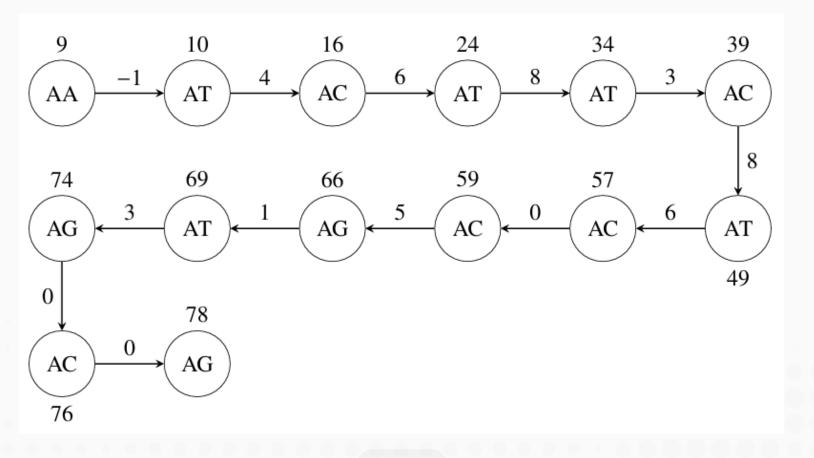
- > The string taken is from the 12841 base to the 12920 base of Lambda.
- \succ The string obtained is: α =GCGTGGGGAA TCTTTACCGG CTGATGCGCG GCTATGCCAC CGGCGGTTAT GTCGGTACAC CGGGCAGCAT GGCAGACAGC.

>Lambda_NEB

 GGATTTGTGGAAGGCGGAGAGTCAGTTCGCGGTACTGGAGGAGGCGCGCAACGTCGCCAGCTGTCTGCA ACAAGGTTACGTATCAGGAGCGCCTGAACGCGCTGGCGCAGCAGGCGGATAAATTCGCACAGCAGCAACG GAACAGCGCCTGAAGGAACAGTATGGCGATAATCCGCTGGCGCTGAATAACGTCATGTCAGAGCAGAAAA GGAAGAGAGCGCCACGGACAGTATGTCGCAGGTAAAAAGTGCAGCCACGCAGACCTTTGATGGTATTGCA <u>CAGAATATGGCGGCGATGCTGACCGGCAGTGAGCAGAACTGGCGCAGCTTCACCCGTTCCGTGCTGTCCA</u> TGATGACAGAAATTCTGCTTAAGCAGGCAATGGTGGGGATTGTCGGGAGTATCGGCAGCGCCATTGGCGG GGCTGTTGGTGGCGCCATCCGCGTCAGGCGGTACAGCCATTCAGGCCGCTGCGGCGAAATTCCATT GCAACCGGAGGATTTACGGGAACCGGCGGCAAATATGAGCCAGCGGGGATTGTTCACCGTGGTGAGTTTG TCTTCACGAAGGAGGCAACCAGCCGGATTGGCGTGGGGAATCTTTACCGGCTGATGCGCGGCTATGCCAC CGGCGGTTATGTCGGTACACCGGGCAGCATGGCAGACAGCCGGTCGCAGGCGTCCGGGACGTTTGAGCAG AATAACCATGTGGTGATTAACAACGACGGCACGAACGGGCAGATAGGTCCGGCTGCTCTGAAGGCGGTGT ATGACATGGCCCGCAAGGGTGCCCGTGATGAAATTCAGACACAGATGCGTGATGGTGGCCTGTTCTCCGG



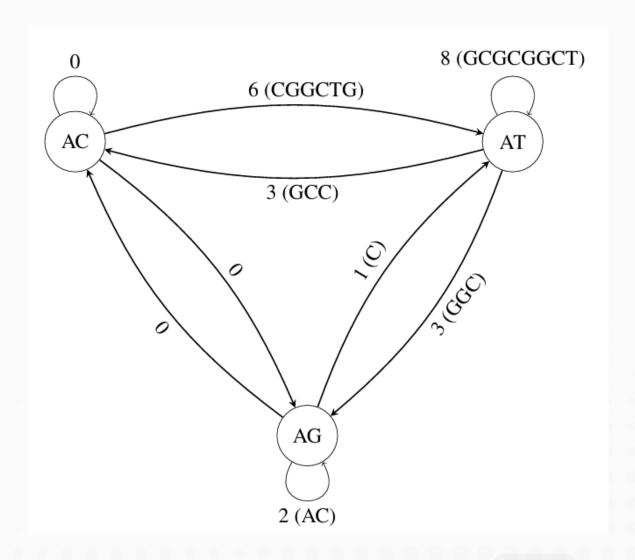
> GCGTGGGGAA TCTTTACCGG CTGATGCGCG GCTATGCCAC CGGCGGTTAT GTCGGTACAC CGGGCAGCAT GGCAGACAGC.





End vertex	Position	AA	AT	AC	AT	AT	AC	AT	AC	AC	AG	AT	AG	AC	AG
Start vertex	1 OSITION	9	10	16	24	34	39	49	57	59	66	69	74	76	78
AA	9	-	-1	5	13	23	28	38	46	48	55	58	63	65	67
AT	10	-	-	4	12	22	27	37	45	47	54	57	62	64	66
AC	16	-	-	-	6	16	21	31	39	41	48	51	56	58	60
AT	24	-	-	-	-	8	13	23	31	33	40	43	48	50	52
AT	34	-	-	-	-	-	3	13	21	23	30	33	38	40	42
AC	39	-	-	-	-	-	-	8	16	18	25	28	33	35	37
AT	49	-	-	-	-	-	-	-	6	8	15	19	23	25	27
AC	57	-	-	-	-	-	-	-	-	0	7	10	15	17	19
AC	59	-	-	-	-	-	-	-	-	-	5	8	13	15	17
AG	66	-	-	-	-	-	-	-	-	-	-	1	6	8	10
AT	69	-	-	-	-	-	-	-	-	-	-	-	3	5	7
AG	74	-	-	-	-	-	-	-	-	-	-	-	-	0	2
AC	76	-	-	-	-	-	-	-	-	-	-	-	-	-	0
AG	78	-	-	-	-	-	-	-	-	-	-	-	-	-	-





> Reduced graph with vertex set V'_a for α .



Start vertex	End vertex	Path taken	Path length		
		$AC \rightarrow AC$	0		
	AC	$AC \rightarrow AG \rightarrow AC$	0 + 0 = 0		
		$AC \rightarrow AG \rightarrow AT \rightarrow AC$	0 + 1 + 3 = 4		
		$AC \rightarrow AT \rightarrow AC$	6 + 3 = 9		
		$AC \rightarrow AT \rightarrow AG \rightarrow AC$	6 + 3 + 0 = 9		
AC	AG	$AC \rightarrow AC \rightarrow AG$	0 + 0 = 0		
		AC→AG	0		
		$AC \rightarrow AT \rightarrow AG$	6 + 3 = 9		
	AT	$AC \rightarrow AC \rightarrow AT$	0 + 6 = 6		
		$AC \rightarrow AG \rightarrow AT$	0 + 1 = 1		
		AC→AT	6		
		$AG \rightarrow AC$	0		
	AC	$AG \rightarrow AG \rightarrow AC$	2 + 0 = 2		
		$AG \rightarrow AT \rightarrow AC$	1 + 3 = 4		
		$AG \rightarrow AC \rightarrow AG$	0 + 0 = 0		
	AG	$AG \rightarrow AC \rightarrow AT \rightarrow AG$	0 + 6 + 3 = 9		
AG		$AG \rightarrow AG$	2		
		$AG \rightarrow AT \rightarrow AG$	1 + 3 = 4		
		$AG \rightarrow AT \rightarrow AC \rightarrow AG$	1 + 3 + 0 = 4		
		$AG \rightarrow AC \rightarrow AT$	0 + 6 = 6		
	AT	$AG \rightarrow AG \rightarrow AT$	2 + 1 = 3		
		AG→AT	1		

Start vertex	End vertex	Path taken	Path length		
		$AT \rightarrow AC$	3		
	AC	$AT \rightarrow AG \rightarrow AC$	3 + 0 = 3		
		$AT \rightarrow AT \rightarrow AC$	8 + 0 = 8		
	AG	$AT \rightarrow AC \rightarrow AG$	3 + 0 = 3		
		AT→AG	3		
AT		$AT \rightarrow AT \rightarrow AG$	8 + 3 = 11		
		$AT \rightarrow AC \rightarrow AT$	3 + 6 = 9		
	AT	$AT \rightarrow AC \rightarrow AG \rightarrow AT$	3 + 0 + 1 = 4		
		$AT \rightarrow AG \rightarrow AT$	3 + 1 = 4		
		$AT \rightarrow AG \rightarrow AC \rightarrow AT$	3 + 0 + 6 = 9		
		$AT \rightarrow AT$	8		

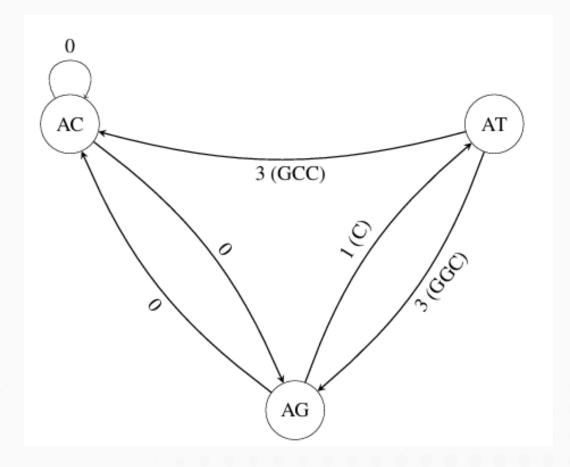
Calculation of the shortest path



> Shortest path and the minimized DNA string for each path

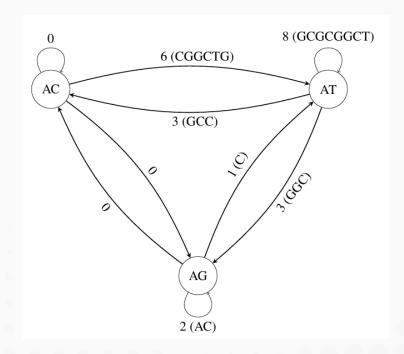
Start vertex	End vertex	Path taken	Path length	Minimized DNA string
	AC	$AC \rightarrow AC$	0	ACAC
AC	AG	$AC \rightarrow AG$	0	ACAG
	AT	$AC \rightarrow AG \rightarrow AT$	0 + 1 = 1	ACAGCAT
	AC	$AG \rightarrow AC$	0	AGAC
AG	AG	$AG \rightarrow AC \rightarrow AG$	0 + 0 = 0	AGACAG
	AT	$AG \rightarrow AT$	1	AGCAT
	AC	$AT \rightarrow AC$	3	ATGCCAC
AT	AG	$AT \rightarrow AG$	3	ATGGCAG
	AT	$AT \rightarrow AG \rightarrow AT$	3 + 1 = 4	ATGGCAGCAT





> Simplified graph with vertex set V'_a for α .

> The minimized DNA string is ATGCCACACAGCATGCCAGAC.



> Reduced graph form



Ongoing Research

A Theoretical DNA Based Computer Model for Food Authentification Process

The authentication of food is an urgent concern owing to the increasing population and direct consequences of food on public health. Food authentication using DNA and omics-based methods is gaining ground due to critical advantages notably in the areas of food adulteration in plant and animal-based food and feed products and in determining the quality of food and food spoilage. Besides, there is greater demand for the detection of genetically modified foods (GMOs) and the detection of allergens, toxins, and carcinogens like tobacco in the food. Advanced DNA and omics-based methods (genomics, metabolomics, and proteomics) have been used in the food industry including DNA-based methods that rely on specific markers known as reference genes for food authentication. Novel methods like CRISPR-Cas have been recently introduced for the management of beneficial microorganisms relevant to food like probiotics.

Publication

https://people.utm.my/nizasarmin/journal-papers/



International Journal of Applied Mathematics and Statistics

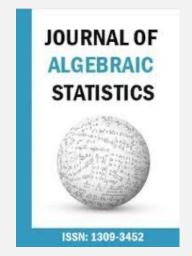


Year: 2018

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Issue

Issue Number: 3











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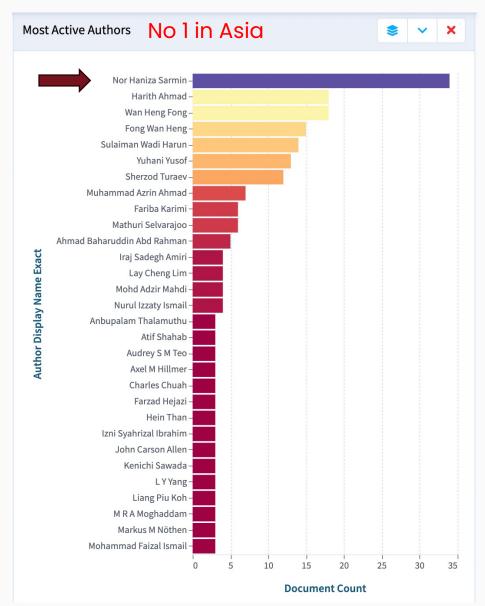
Fuzzy Splicing Systems

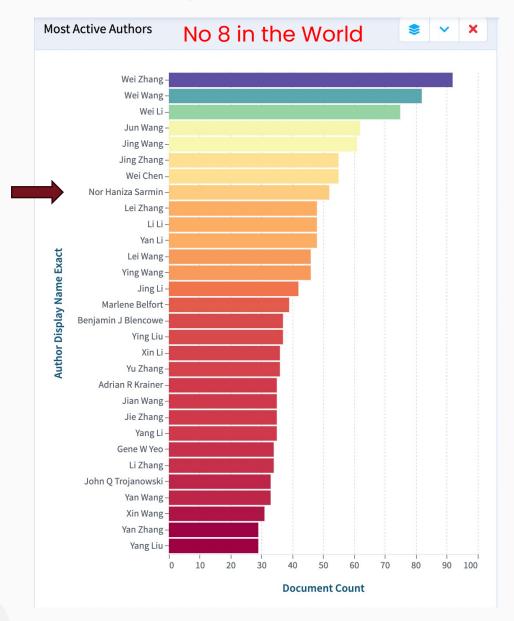
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Ranking in Splicing Systems (Lens.org)





Active Collaborator on DNA Splicing System



Specialization:

- Formal Languages and Automata
- 2. DNA Computing
- 3. Artificial Intelligence
- 4. Cryptography

Ass. Prof. Dr. Sherzod Turaev
College of Information Technology, United Arab Emirates University

Co-supervisor for some PhD students

Active Collaborator on DNA Splicing System



Specialization:

- Spectral of Laplacian Hypergraph and Graph
- Distance Matrices and Quadratic Embedding of Graphs
- 3. Machine Learning
- 4. DNA Sequencing

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In the Name of God for Mankind