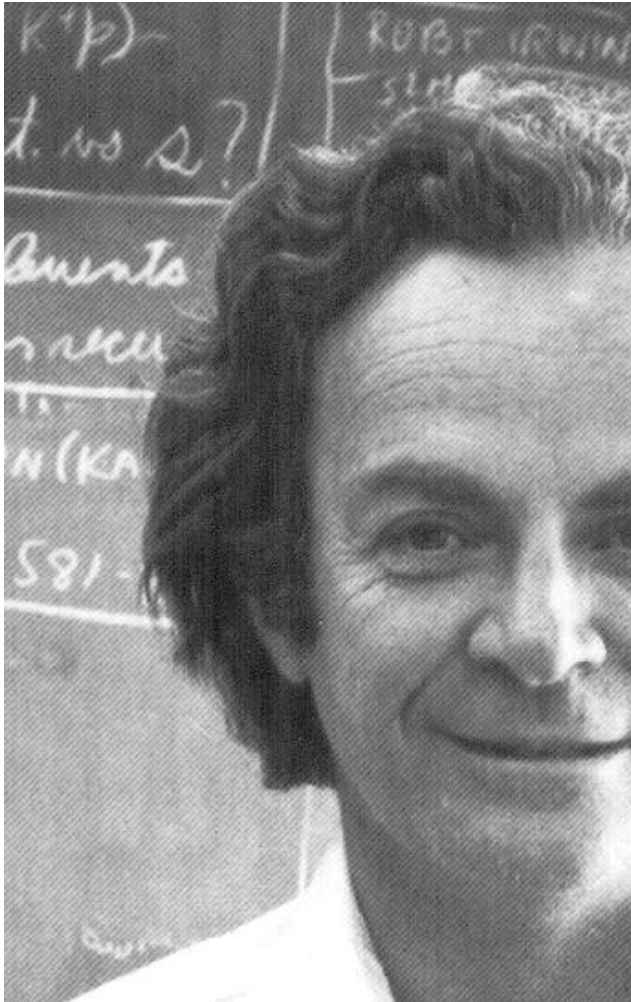


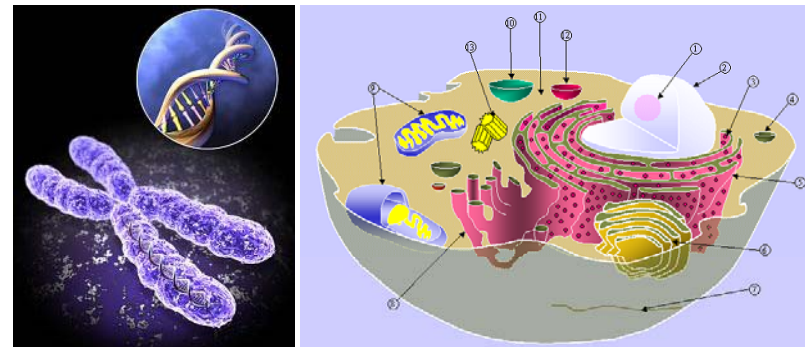


DNA Computing

Computing on molecular level



- Richard P. Feynman
- In late 1950s,
- **Biological molecules** can carry enormous amounts of information in an exceedingly small space.
 - **inborn computing power!**





Motivation of DNA Computing

- We need a totally different **new technology to overcome CMOS limitations.**
- Certain types of problems (learning, pattern recognition, large set search algorithms) are intrinsically very difficult to solve even with fast evolution of CMOS.
- It is natural **to solve biological problems** with biological tools.



Advantages of DNA Computing

- Achievement of massive **parallelism**
- Parallel molecular operation
 - Desktop: 10^9 operations/ sec
 - Supercomputer: 10^{12} operations/ sec
 - 1 mole of DNA: 10^{23} **simultaneous reactions**
- High information storage capacity
 - 6.022×10^{23} molecules/ mole \rightarrow 1 bit per cubic nanometer
- Favorable Energetics



Rise and Growth of DNA Computing

■ Adleman's work in 1994

- Hamiltonian path problem (graph problem, NP problem)
- City and road information representation using DNA sequences (indicative information)
- Solution-based DNA computing

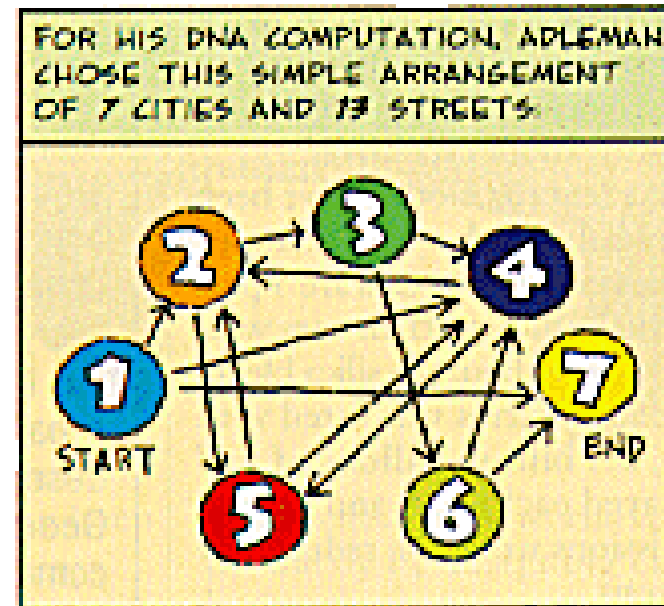
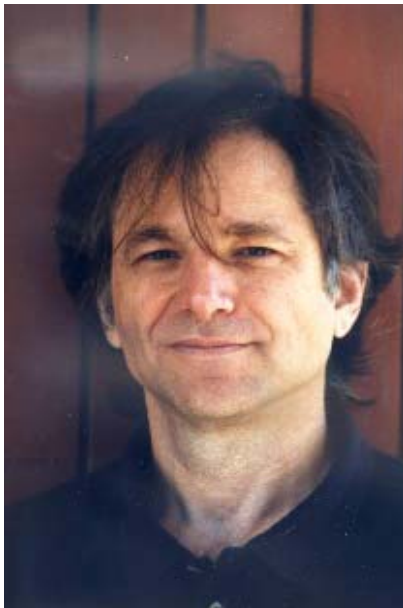
■ Liu's work in 2000

- SAT problem
- Surface-based DNA computing

■ Benenson's work in 2004

- Application to disease diagnosis and drug (antisense) administration

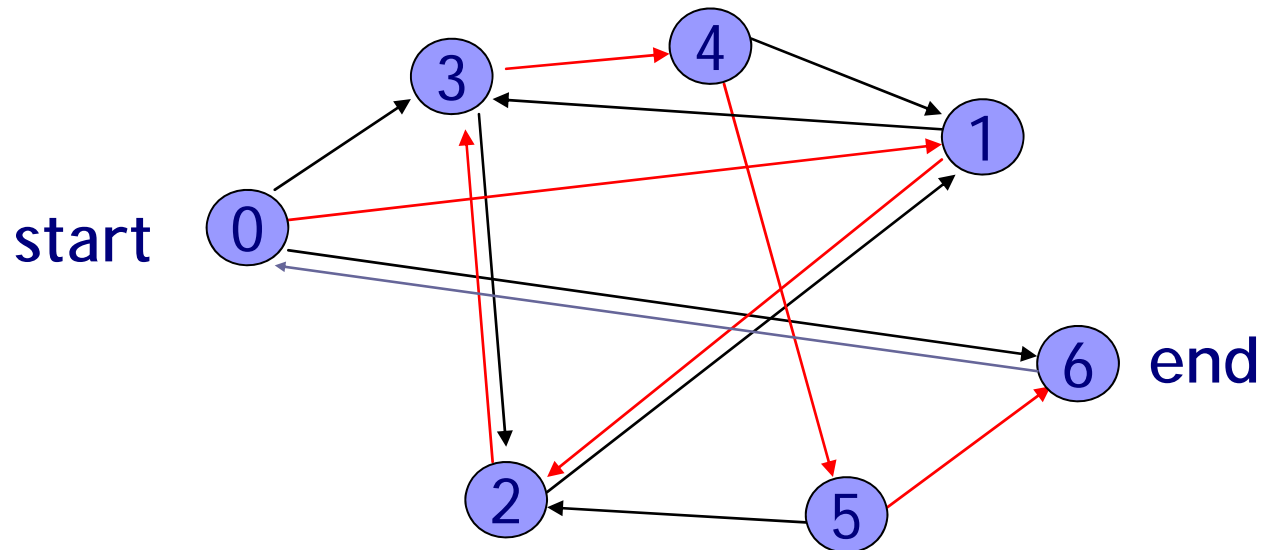
Rise of DNA computing



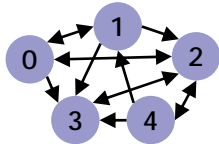
- In 1994, Adleman showed that DNA 'can compute'.
- Hamiltonian path problem (HPP)

Hamiltonian Path Problem (HPP)

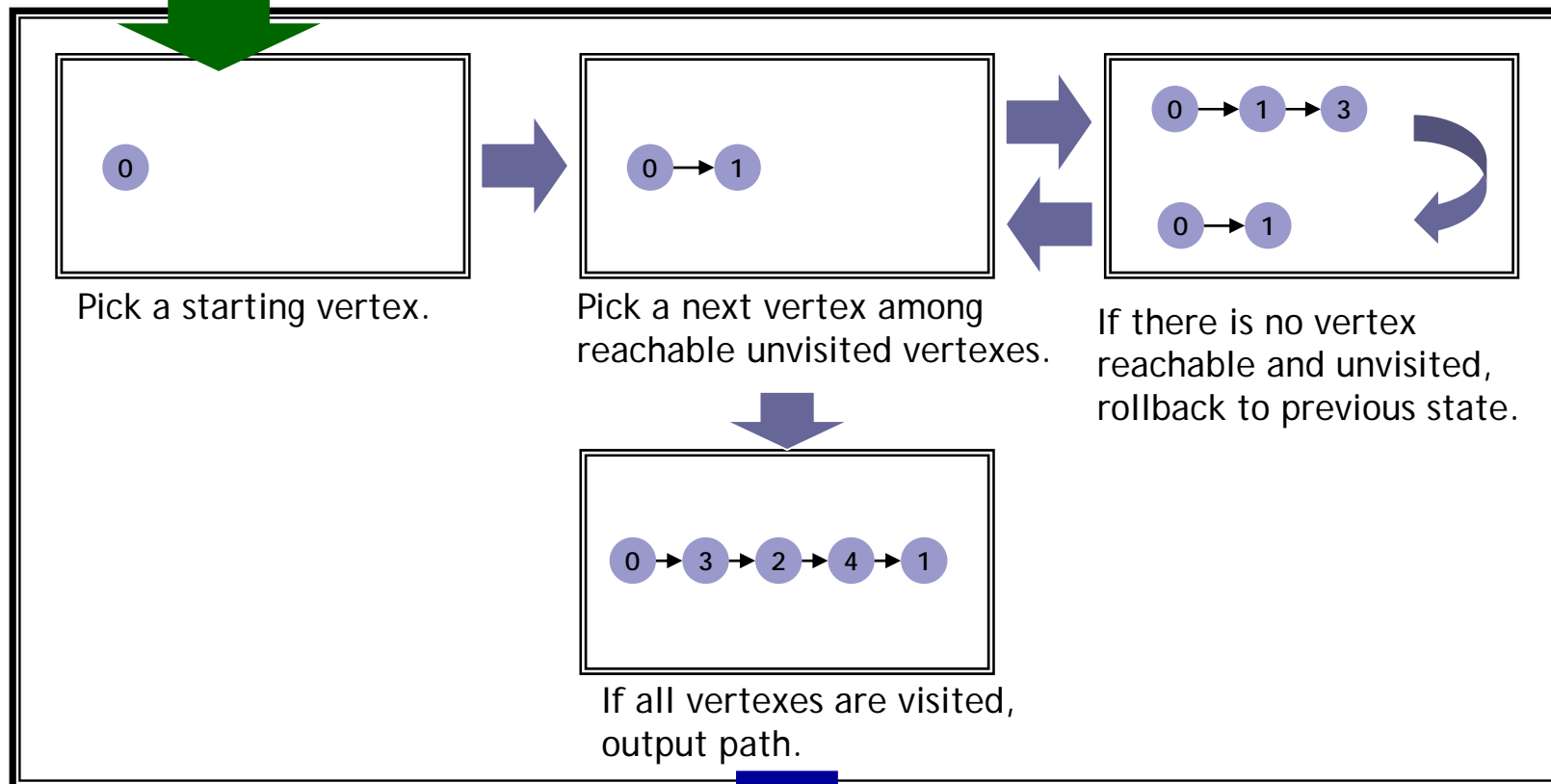
- HPP is to find a route (if it exists) that **passes through each city exactly once** with a designated start and end.



Conventional computers

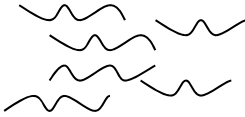


Input:
Graph information: vertexes and edges

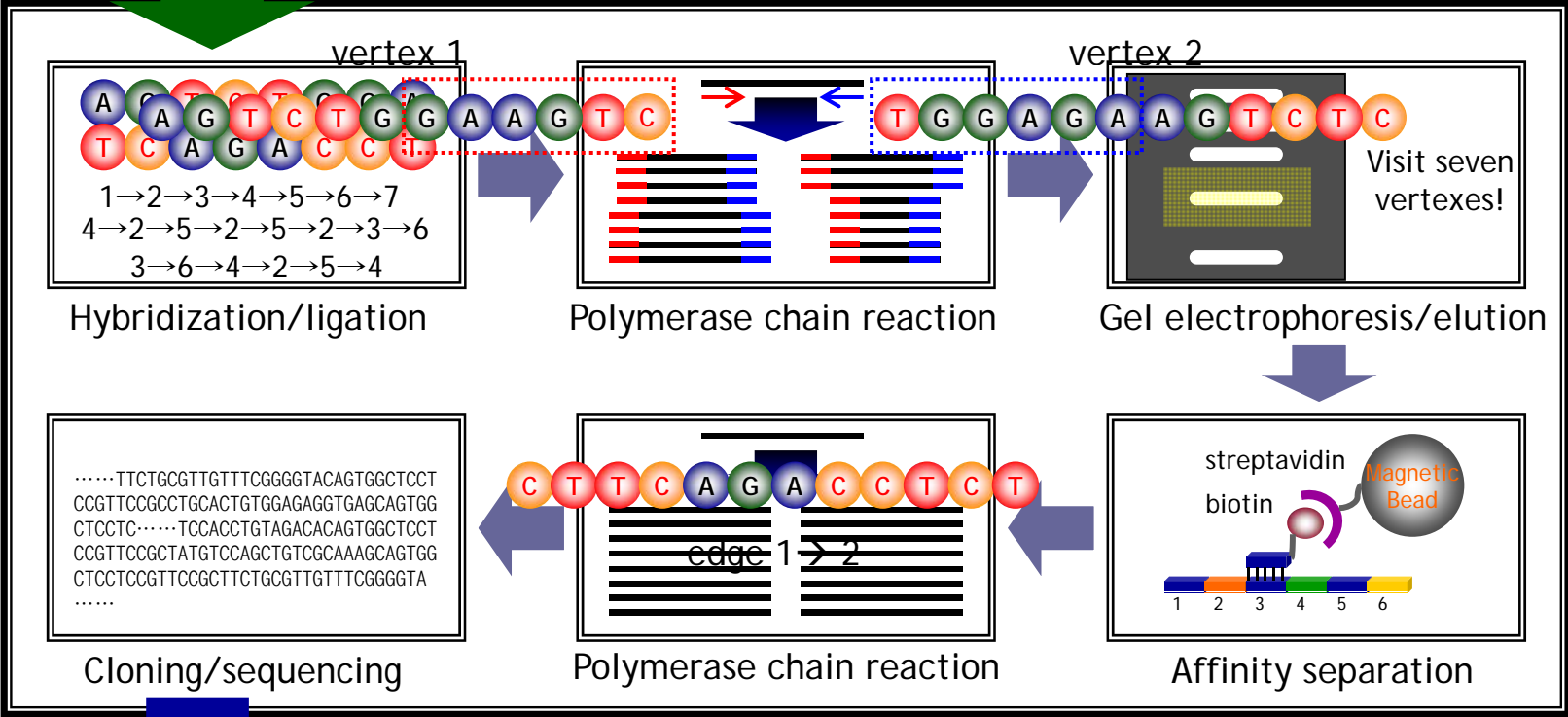


Output:
Hamiltonian path

DNA computers for HPP



Input:
Oligonucleotides mixture
(vertexes, edges)



Output:
DNA sequence
of Hamiltonian path



Experimental Implementation for HHP Conditions

- Starting with city 0 and ending with city 6
 - PCR using primers complementary to city 0 and city 6
- Visiting seven cities
 - Gel electrophoresis/elution
- Visiting every city
 - A series of affinity chromatography
 - Each affinity column contains ssDNA complementary to each city.



Traveling Salesman Problem



Traveling Salesman Problem

■ Find...

- The **cheapest way** of visiting **all the cities** and **returning to the starting point**

when a number of cities to visit and the **traveling cost between each pair of cities** are given.

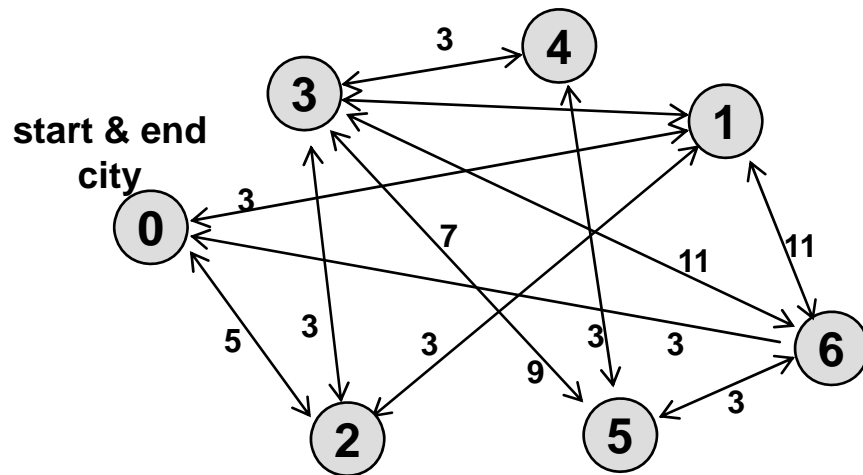
■ Previous work for weight (cost) representation

- DNA length
- DNA concentration

■ Our method for weight (cost) representation

- Thermal stability of DNA duplex
- Melting temperature (T_m), GC content

Target Problem & Encoding Method

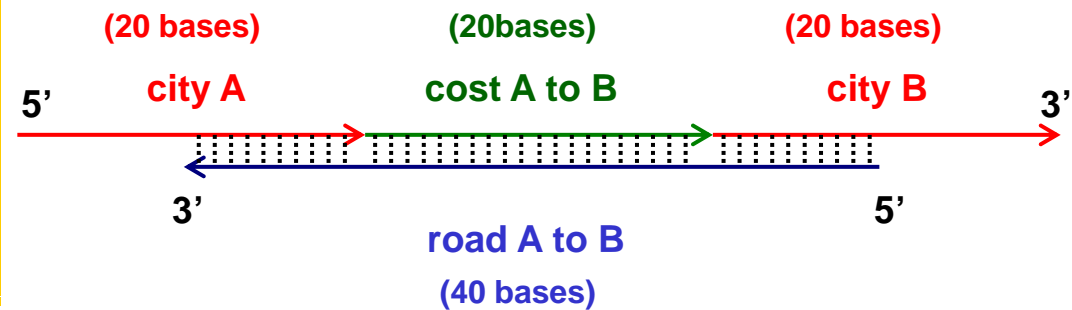


7-city traveling salesman problem

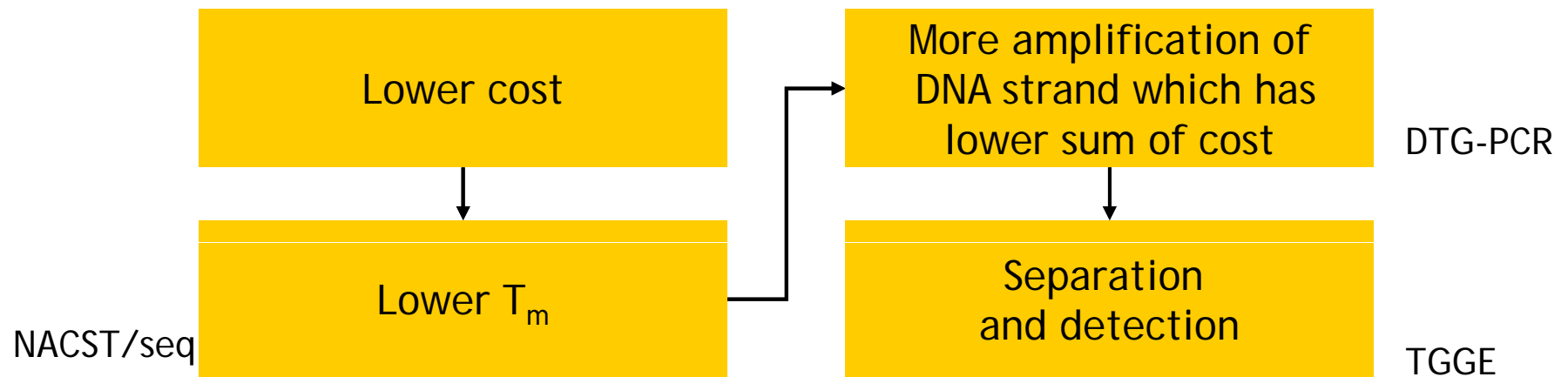
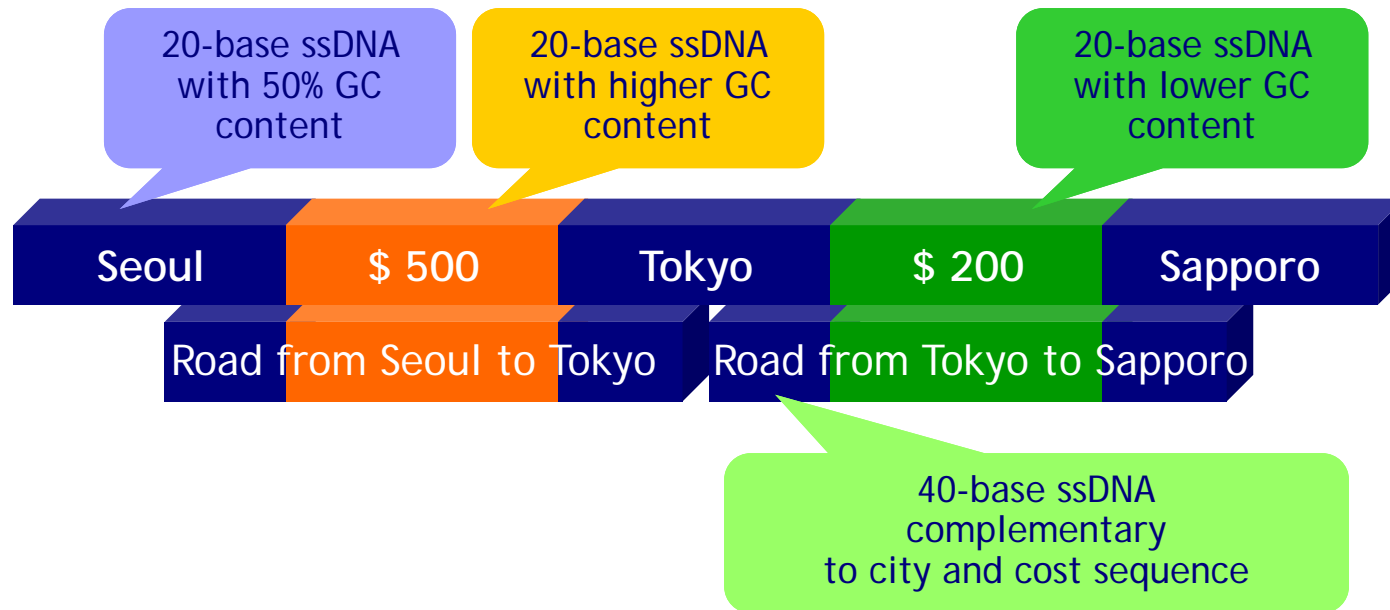
- 7 cities (0 to 6), 23 roads, 5 costs
- optimal path: '0→1→2→3→4→5→6→0'

Oligonucleotides & Encoding

- cities and costs are 20-base ssDNA
- roads are 40-base ssDNA
- 35 oligonucleotides
(7 cities, 23 roads, 5 costs)

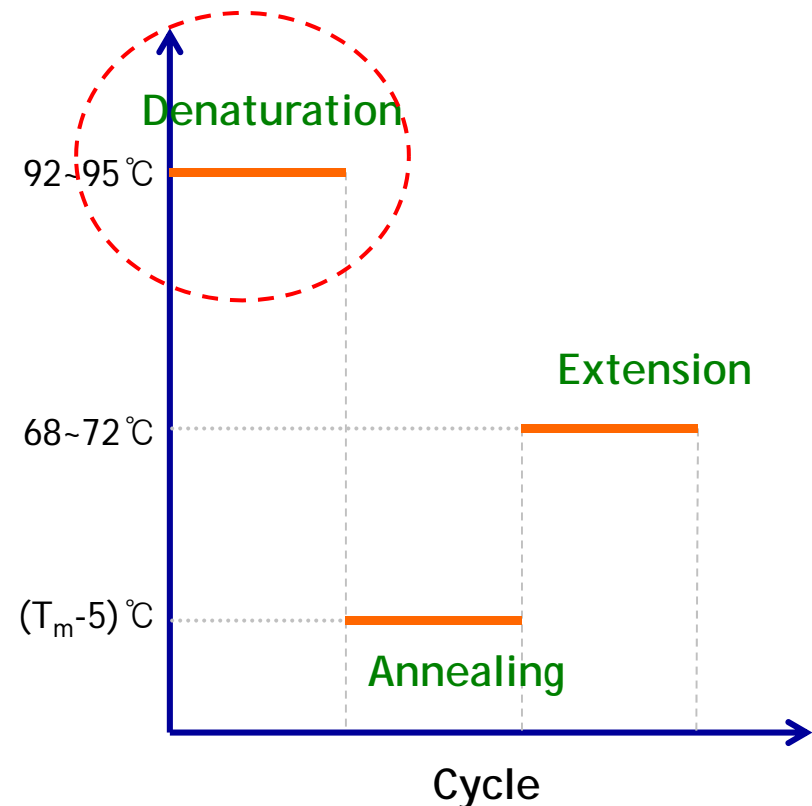


Weight (Cost) Encoding

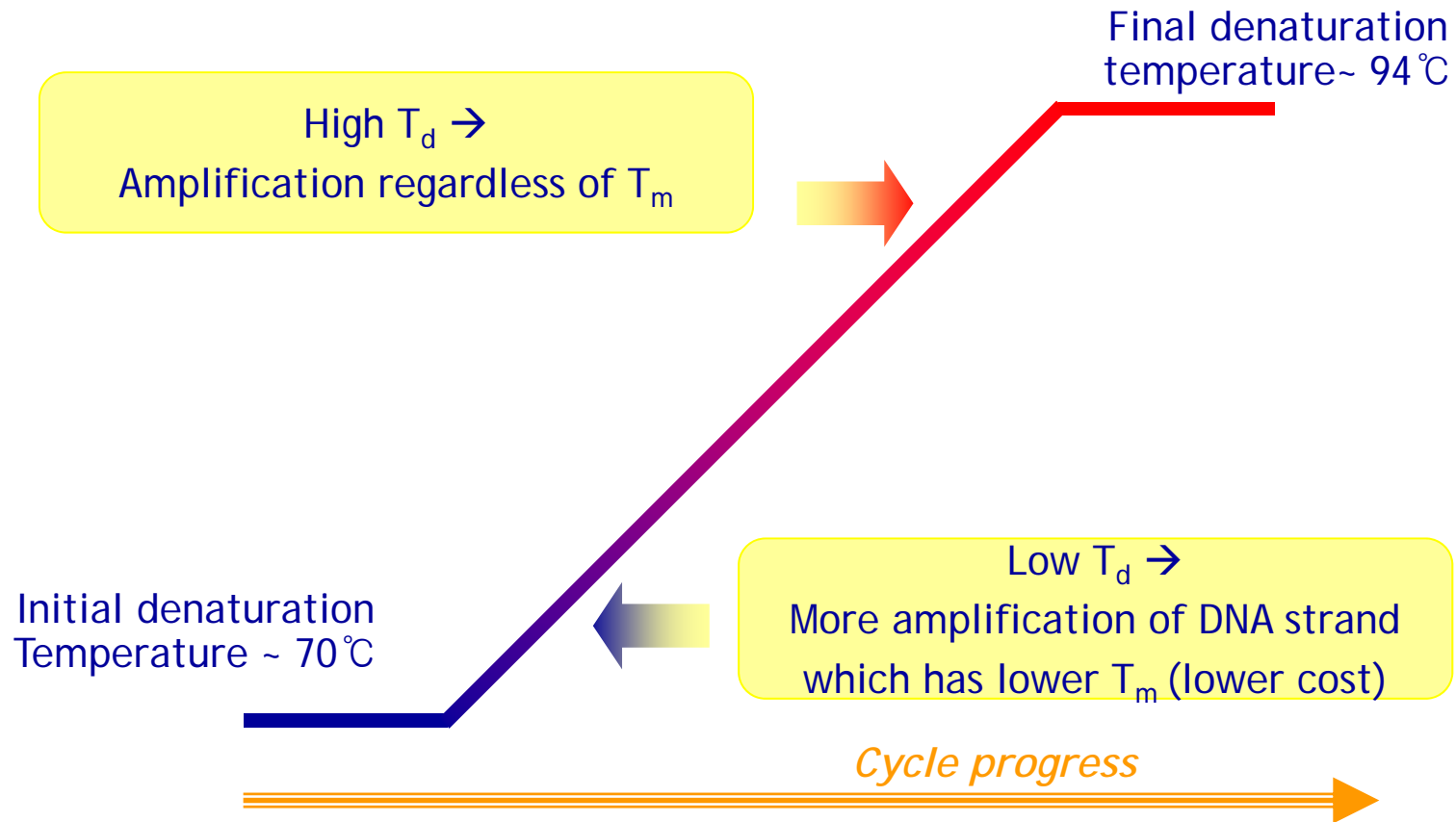


Denaturation Temperature Gradient Polymerase Chain Reaction (DTG-PCR)

- Conventional PCR
 - Denaturation (T_d)
 - Annealing (T_a)
 - Extension (T_e)
- Modification of conventional PCR protocol
 - Variation in T_d

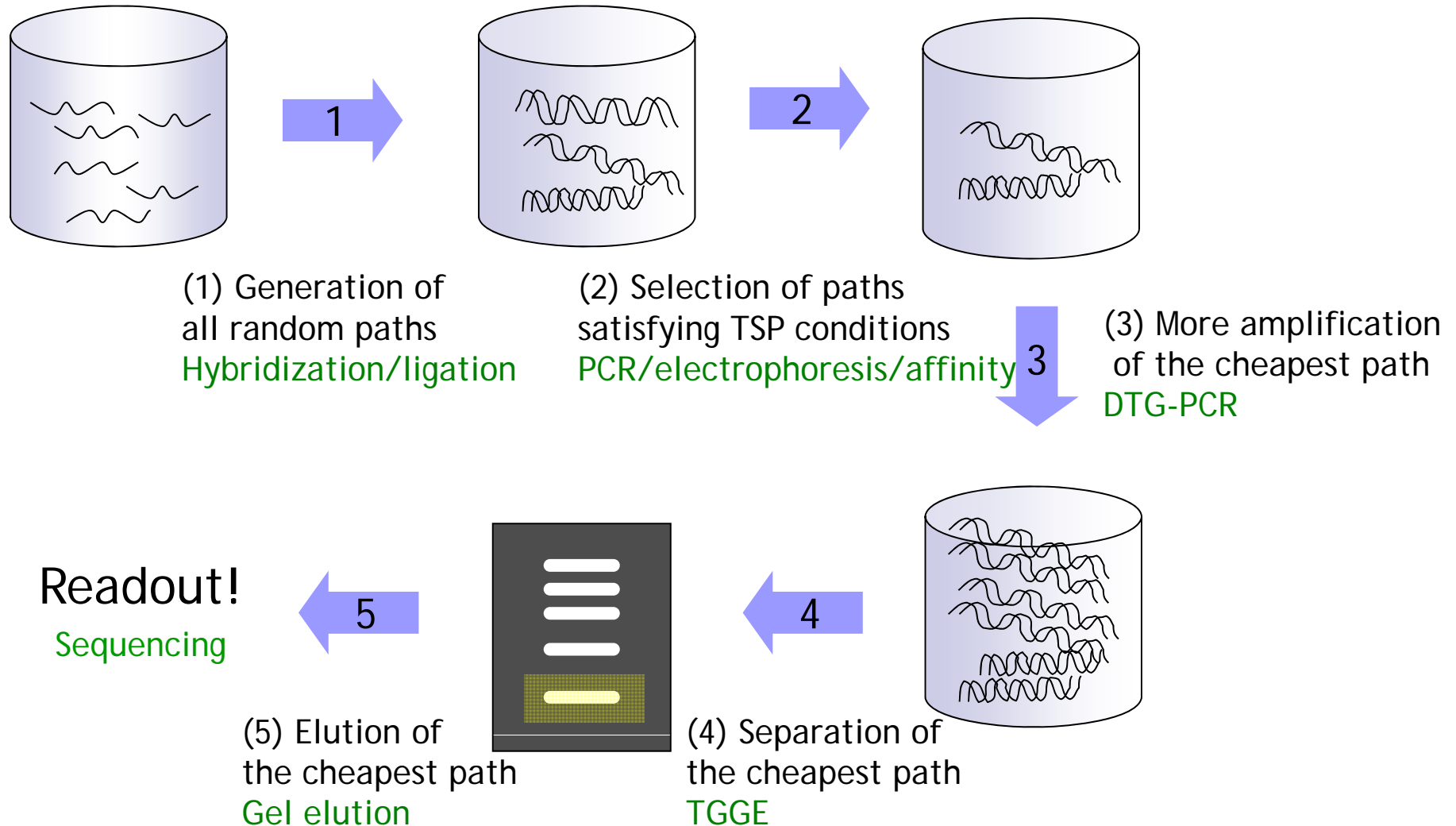


Denaturation Temperature Gradient



- **Biased operator:** more amplification of DNA strands with lower T_m
→ biased search for lower cost

Molecular Algorithm



Sequence Design for Cities and Costs

- Using NACST/seq
- Non-cross hybridization
- Similar T_m among cities
- Different T_m among costs

Vertex sequences			
No.	Sequence (5' → 3')	T_m	GC%
0	AGGCGAGTATGGGGTATATC	60.73	50
1	CCTGTCAACATTGACGCTCA	59.24	50
2	TTATGATTCCACTGGCGCTC	59.00	50
3	ATCGTACTCATGGTCCCTAC	56.81	50
4	CGCTCCATCCTTGATCGTTT	58.13	50
5	CTTCGCTGCTGATAACCTCA	59.44	50
6	GAGTTAGATGTCACGTCACG	56.97	50
Weight sequences			
Edge cost	Sequence (5' → 3')	T_m	GC%
3	ATGATAGATATGTAGATTCC	47.89	30
5	GGATGTGATATCGTTCTTGT	54.62	40
7	GGATTAGCAGTGCCTCAGTT	58.37	50
9	TGGCCACGAAGCCTTCCGTT	64.51	60
11	GAGCTGGCTCCTCATCGCGC	68.88	70

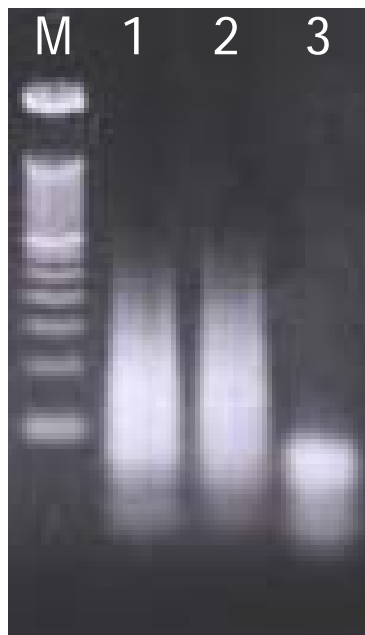


Experimental Implementation for TSP Conditions

- Starting and ending with city 0
 - PCR using primers complementary to city 0
- Visiting every city
 - A series of affinity chromatography
 - Each affinity column contains ssDNA complementary to each city.
- Cheapest path
 - DTG-PCR

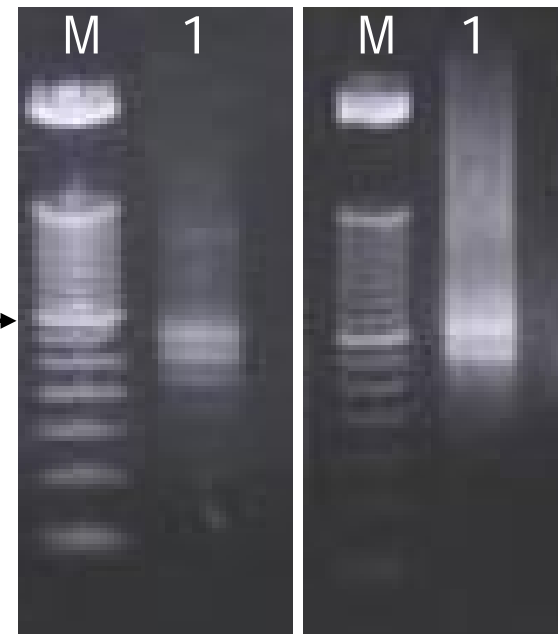
Experimental Results

- Random path generation
(by hybridization and ligation)
- Selective amplification of paths starting and ending with city 0
(by PCR using primers complementary to city 0)



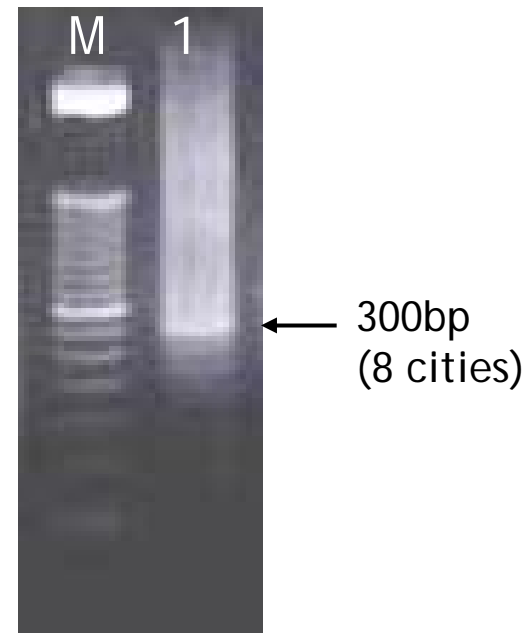
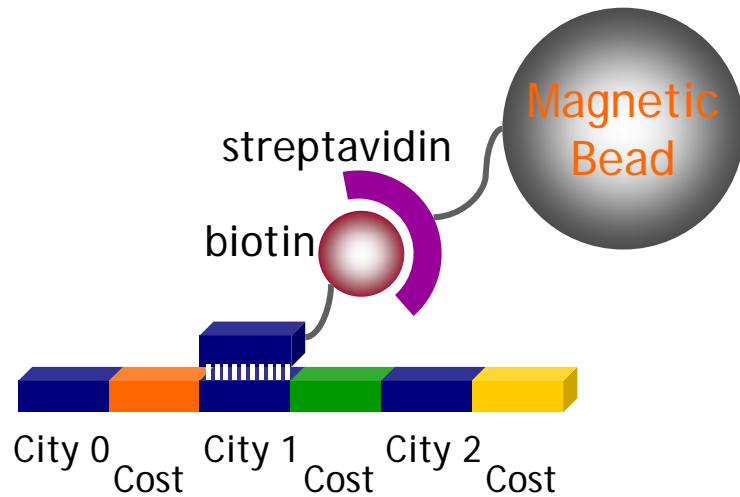
M: 50 bp ladder
lane 1,2:
after hybridization/ligation
lane 3: mixture of ssDNA

300bp →

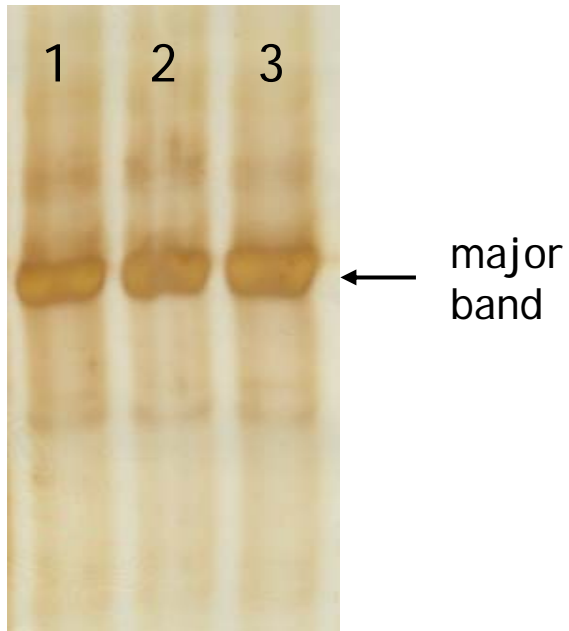


- Separation of paths containing every city
(by a series of affinity chromatography)

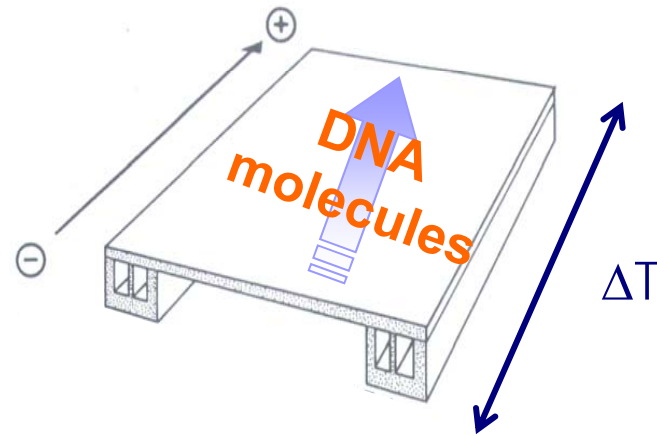
- More amplification of paths with lower costs
(by DTG-PCR)



- Separation of the path with lowest cost
(by TGGE)



- TGGE
(Temperature Gradient - Gel Electrophoresis)



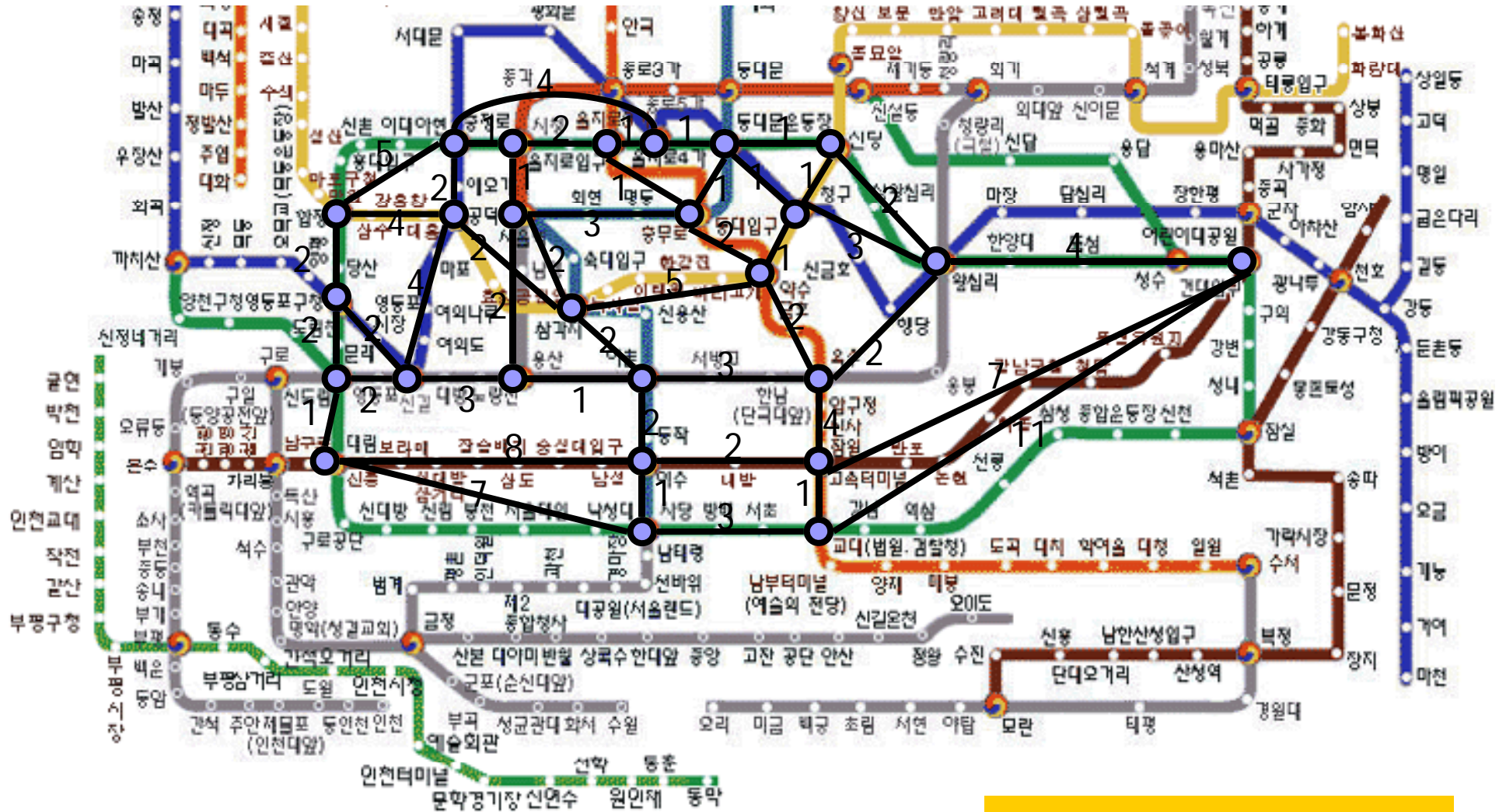
■ Readout

(by cloning and sequencing)

```
.....TTCTGCGTTGTTTCGGGGTACAGTGGCTCCTCCGTT  
CCGCCTGCACTGTGGAGAGGTGAGCAGTGGCTCCTCCGTT  
CCGCGTGGATTACAAGGCCATCGCAGTGGCTCCTCCGTT  
CCGCATACGGCGTGGTTTTTCGGGCAGTGGCTCCTCCGTT  
CCGCAAACGGTCGTAAGTGATGAACAGTGGCTCCTCCGTT  
CCGCGCACAGTCCACCTGTAGACACAGTGGCTCCTCCGTT  
CCGCTATGTCCAGCTGTCGCAAAGCAGTGGCTCCTCCGTT  
CCGCTTCTGCGTTGTTTCGGGGTA.....
```



Toward Larger Problems



Subway routes in Seoul

