

A DNA-Based Archival Storage System

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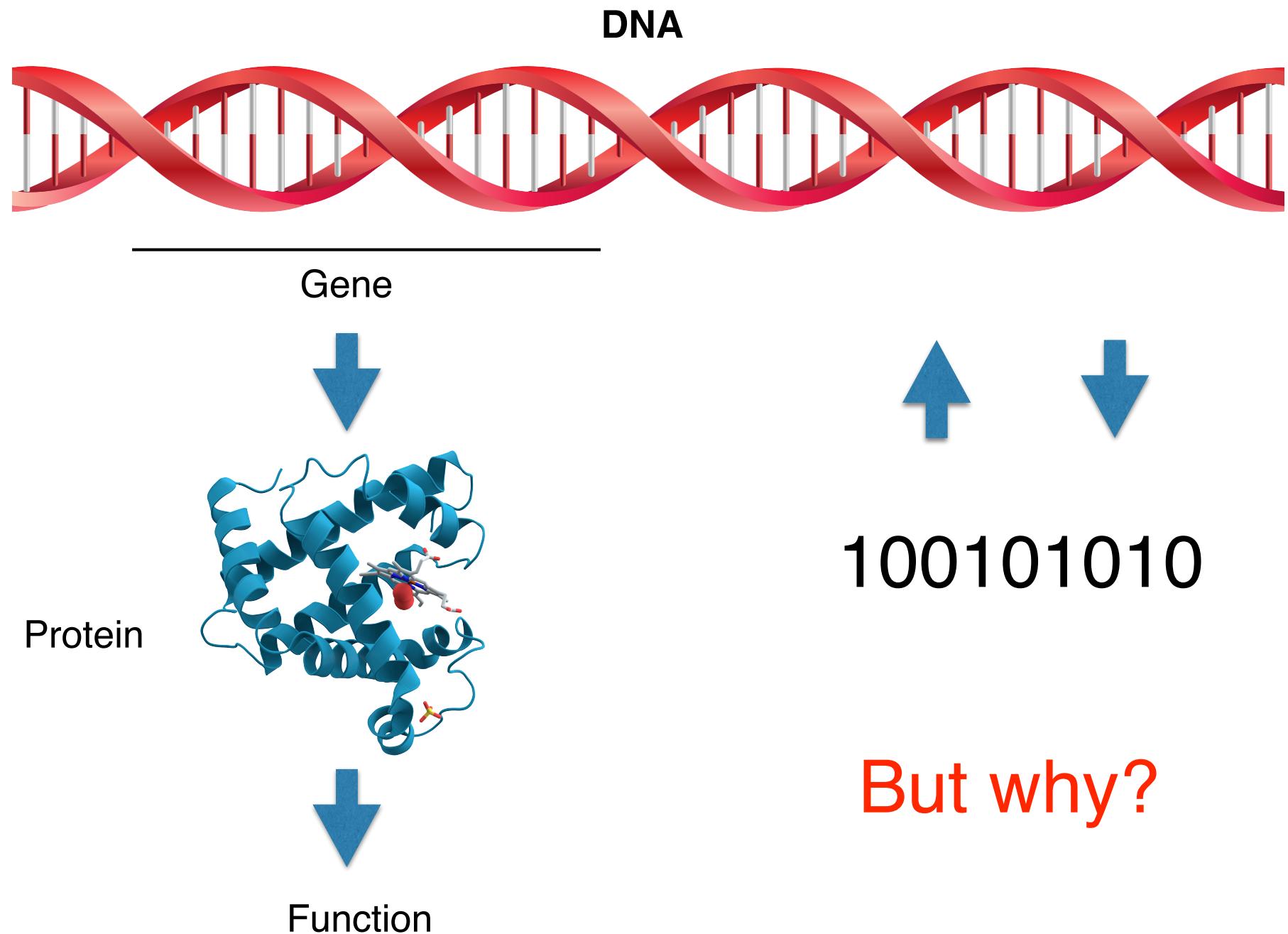


Microsoft
Research

*joint work with Karin Strauss, Doug Carmean, Georg Seelig, James Bornholt,
Randolph Lopez, Lee Organick, Yuan Chen, Chris Takahashi, Bichlien Nguyen,
Sergey Yekhanin, Siena Dumas Ang.*



Life evolved a fantastic storage medium...



DNA molecules for digital data

Extremely dense

Theory: 1 exabyte in 1 mm³



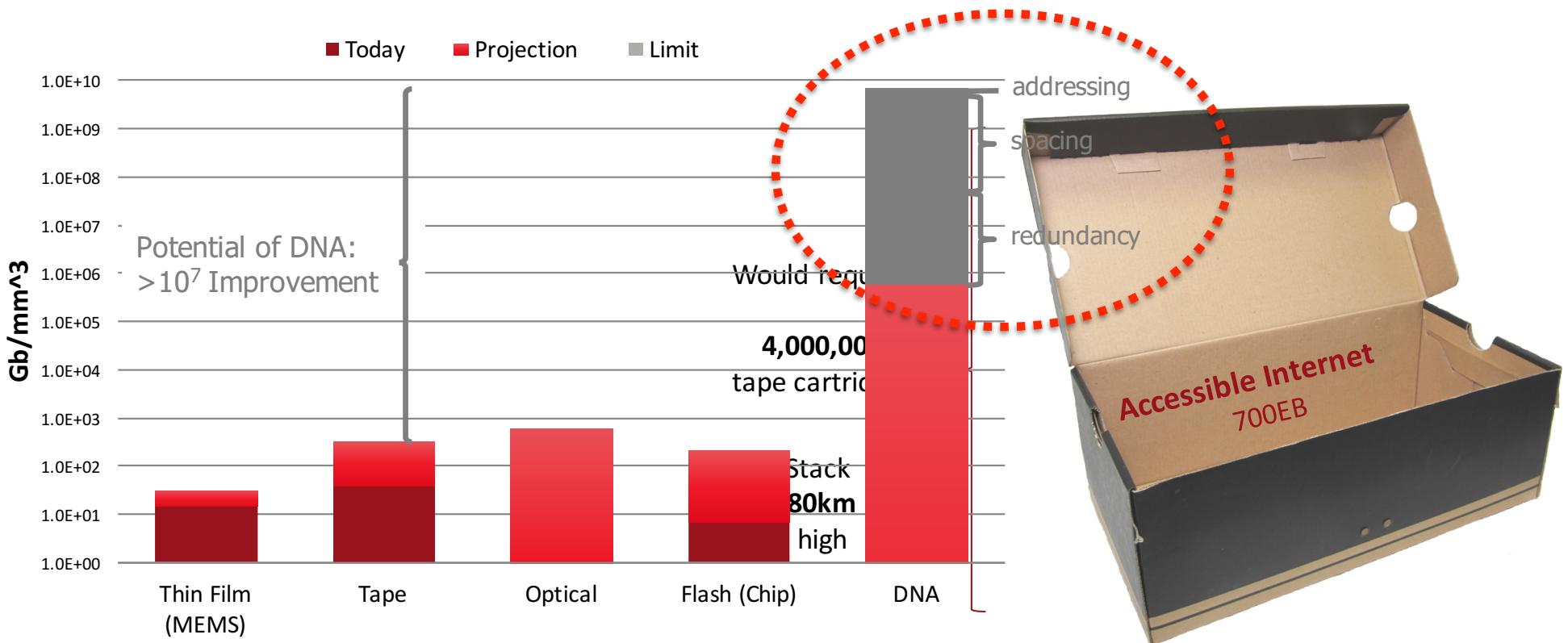
Extremely durable

Half life > 500 years

And readers never become obsolete!



Comparing Storage Technologies



The ultimate hierarchy

	<i>Access Time</i>	<i>Durability</i>
Flash	ms	~5 yrs
HDD	10s ms	~5 yrs
Tape	minutes	~15-30 yrs
DNA-based Archival	hrs	centuries

A DNA-based archival storage system

[ASPLoS'16]

11010101...



11010101...



Redundancy
and density



Write

Efficient
retrieval



Read

Wet lab
experiments

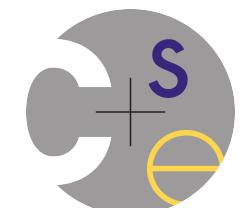


Store

ACATCG...



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Research

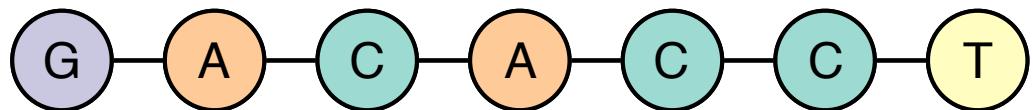


DNA molecules

Four nucleotides:

- A Adenine
- C Cytosine
- G Guanine
- T Thymine

DNA strand (oligonucleotide) is a linear sequence of these nucleotides

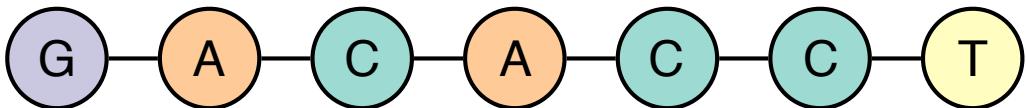


DNA molecules

Four nucleotides:

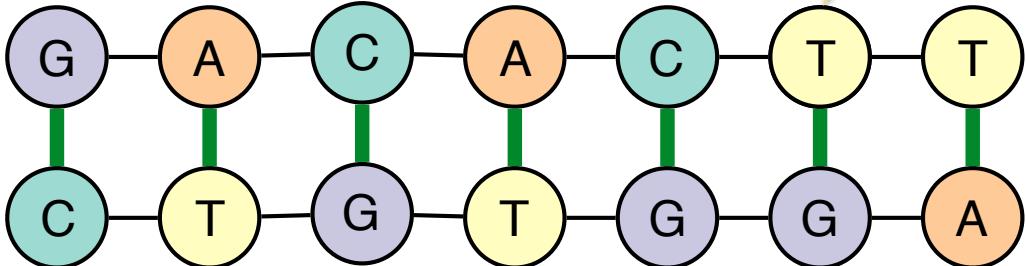
-  Adenine
-  Cytosine
-  Guanine
-  Thymine

DNA strand (oligonucleotide) is a linear sequence of these nucleotides



Two strands can bind to each other if they are complementary:

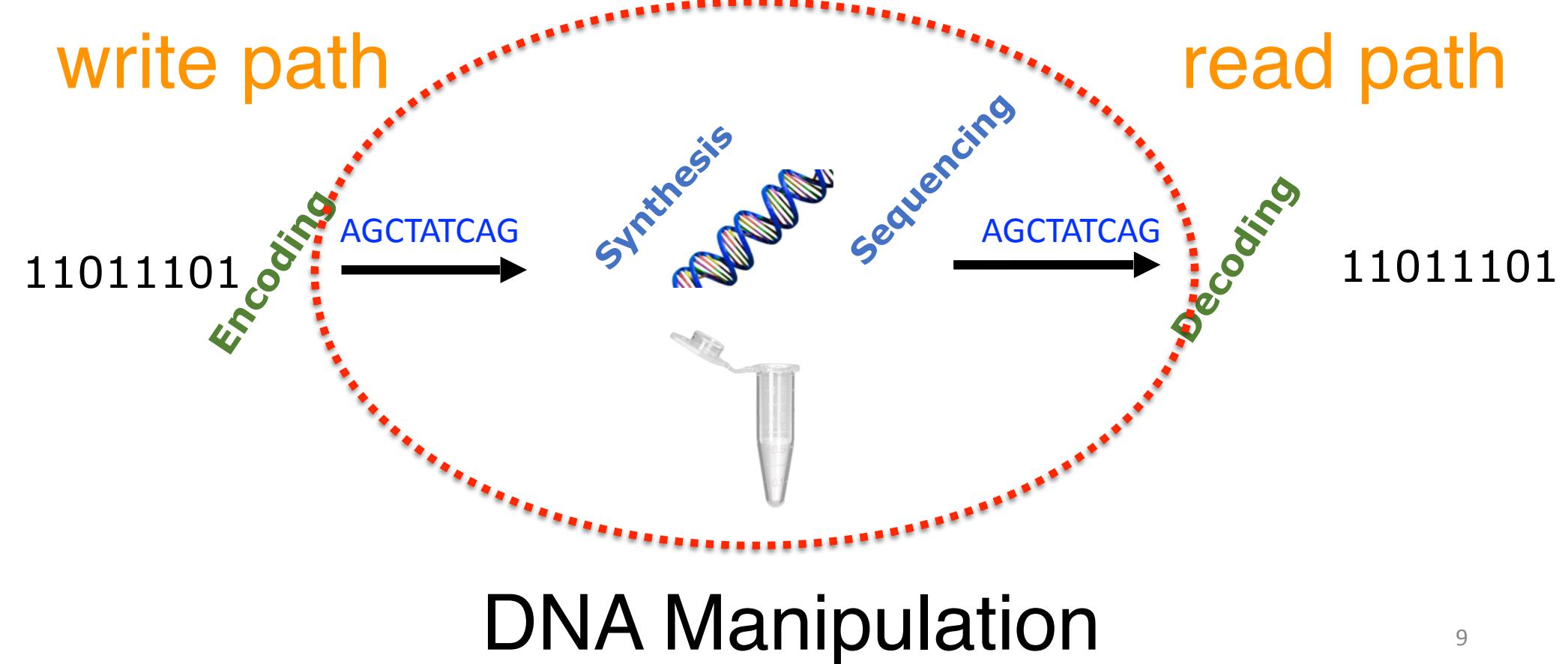
Partial errors allowed



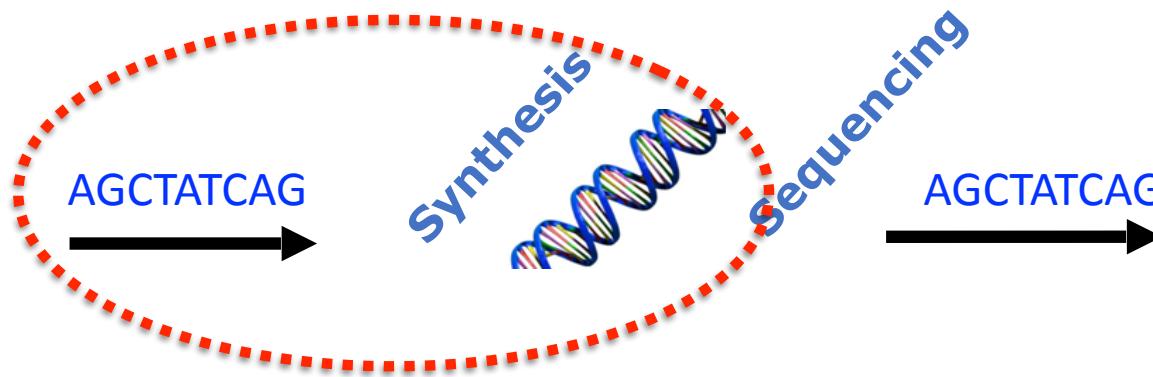
C, G are complementary

A, T are complementary

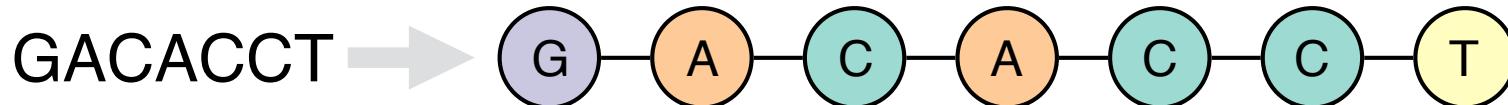
DNA data storage at 30,000 feet



DNA Manipulation: Synthesis



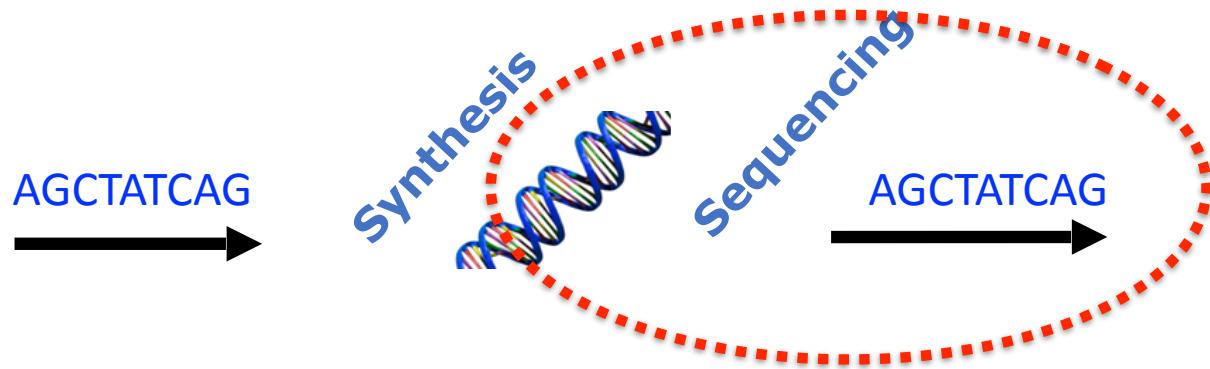
DNA Synthesis: manufacturing DNA strands



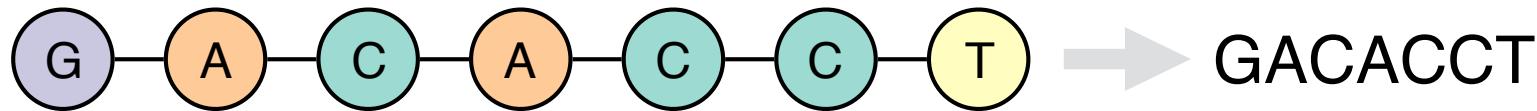
- Synthesis process appends one letter at a time
- Maximum practical sequence length of ~100s of letters
- Produces millions of copies of each sequence
- Can make many different sequences in parallel
- Normally used for genomics and genetic engineering



DNA Manipulation: Sequencing



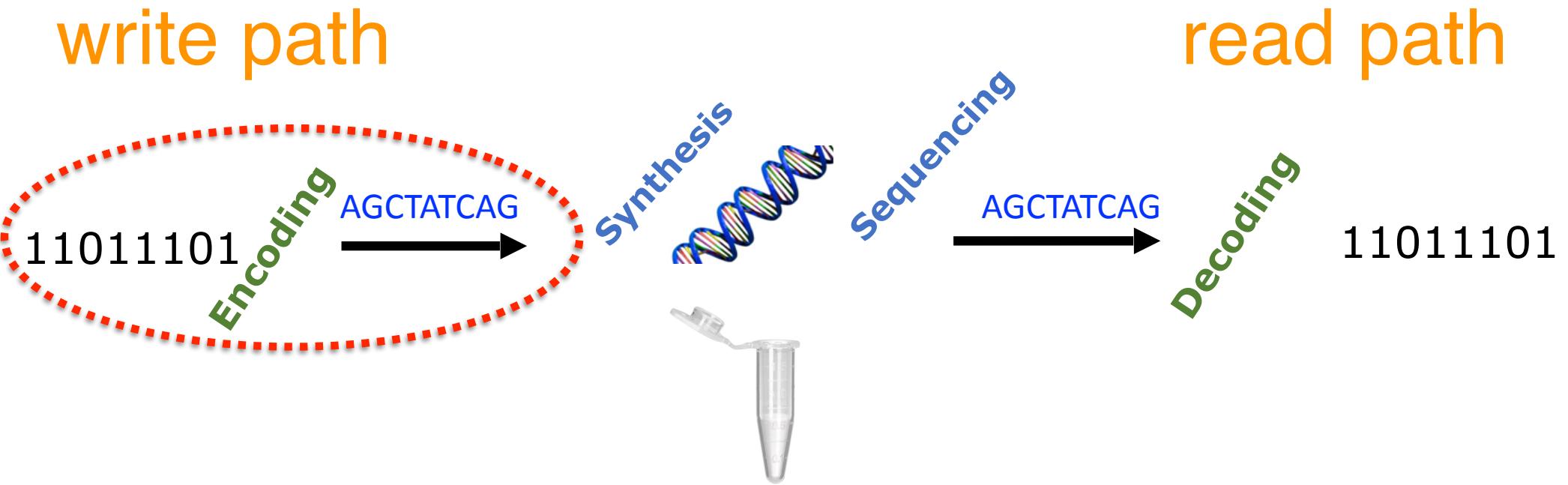
DNA Sequencing: reading DNA strands



- Produces *many* reads of a strand
- Normally used for research and diagnostics
- Currently much higher throughput than synthesis



DNA data storage at 30,000 feet



Writing digital data to DNA

The easy way: convert base 2 to base 4

10100011 10010001 11100111 11000101 10010100 1011101



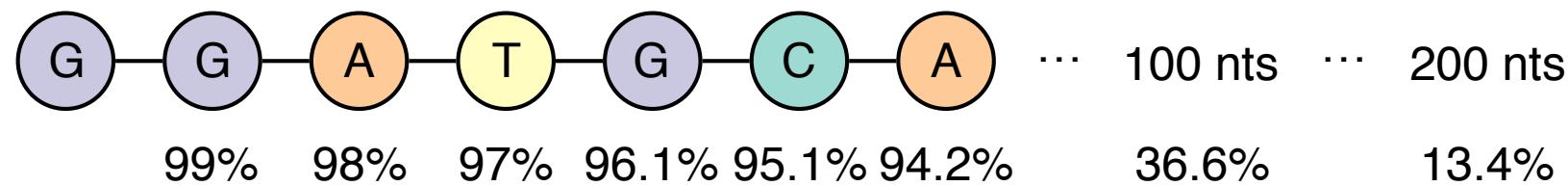
2 2 0 3 2 1 0 1 3 2 1 3 3 0 1 1 2 1 1 0 2 3 3 1



GGAT GCAC TGCT TACC GCCA GTTC

But this approach isn't feasible for more than a few bytes

$P[\text{Attach}] = 99\%$



Chunking data

Break binary data into chunks stored in separate strands

10100011 10010001 11100111 11000101 10010100 1011101



2 2 0 3 2 1 0 1 3 2 1 3 3 0 1 1 2 1 1 0 2 3 3 1



GGAT GCAC TGCT TACC GCCA GTTC

Chunking data

Break binary data into chunks stored in separate strands

10100011 10010001 11100111 11000101 10010100 1011101



2 2 0 3 2 1 0 1 3 2 1 3 3 0 1 1 2 1 1 0 2 3 3 1



ATGTT	GGAT	GCAC	AAAA	CATCC
ATGTT	TGCT	TACC	AAAC	CATCC
ATGTT	GCCAGTTC	AAAG	CATCC	

File identifiers
("primers")

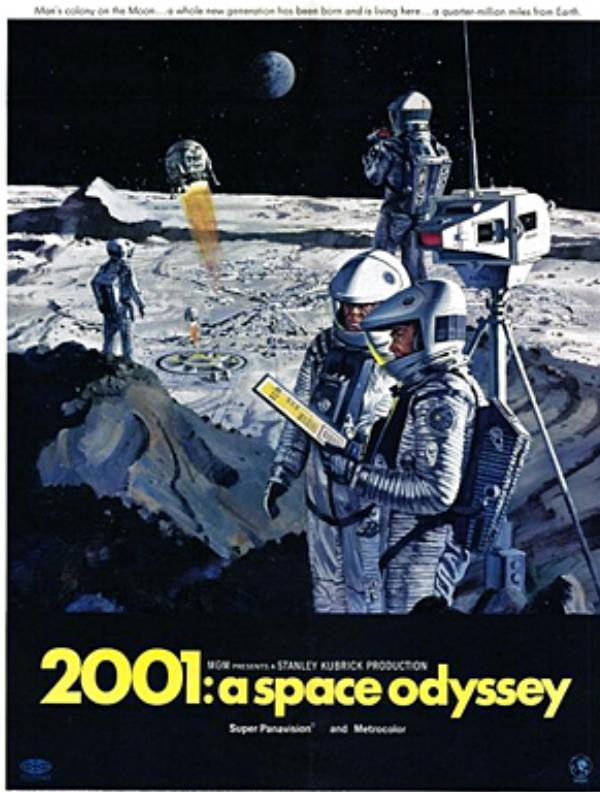
Addresses
within the file

ATGTT	GGATGCAC	AAAAA	CATCC
ATGTT	TGCTTACC	AAAC	CATCC
ATGTT	GCCAGTTC	AAAG	CATCC

File identifiers
("primers")

Addresses
within the file

An epic drama of
adventure and exploration



1 of N

2 of N

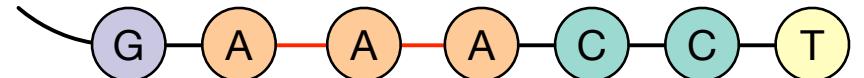
3 of N

~ 10 bytes per
DNA strand.

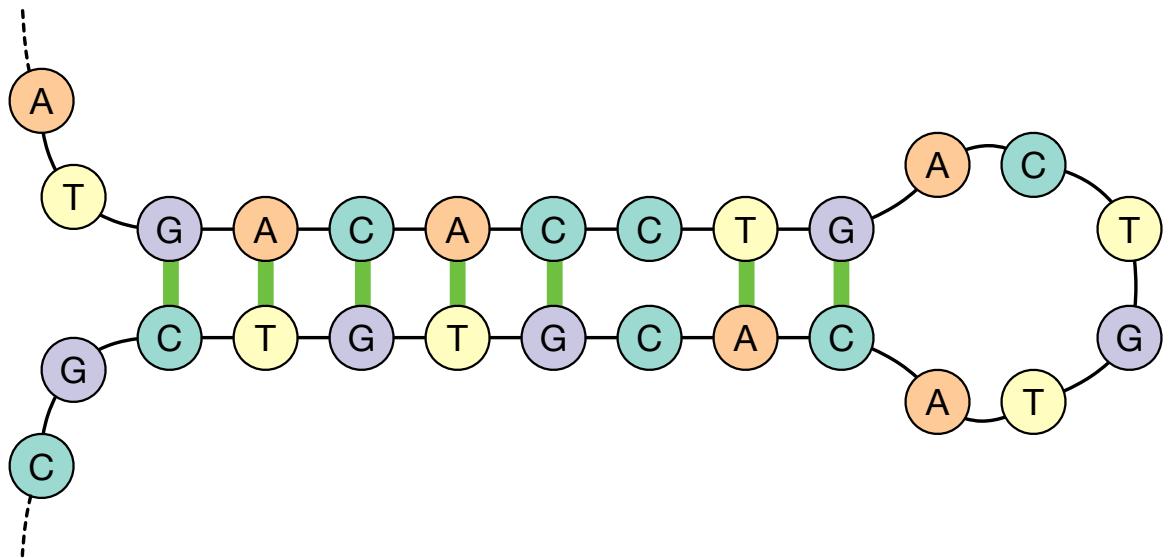
...

Other encoding considerations

Homopolymers are bad...



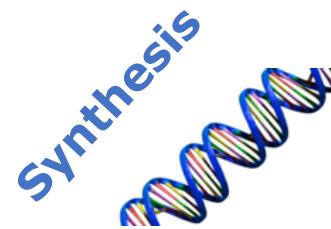
Secondary structures
are problematic ...



DNA data storage at 30,000 feet

write path

11011101
Encoding → AGCTATCAG



Synthesis
Sequencing

→ AGCTATCAG

read path

Decoding

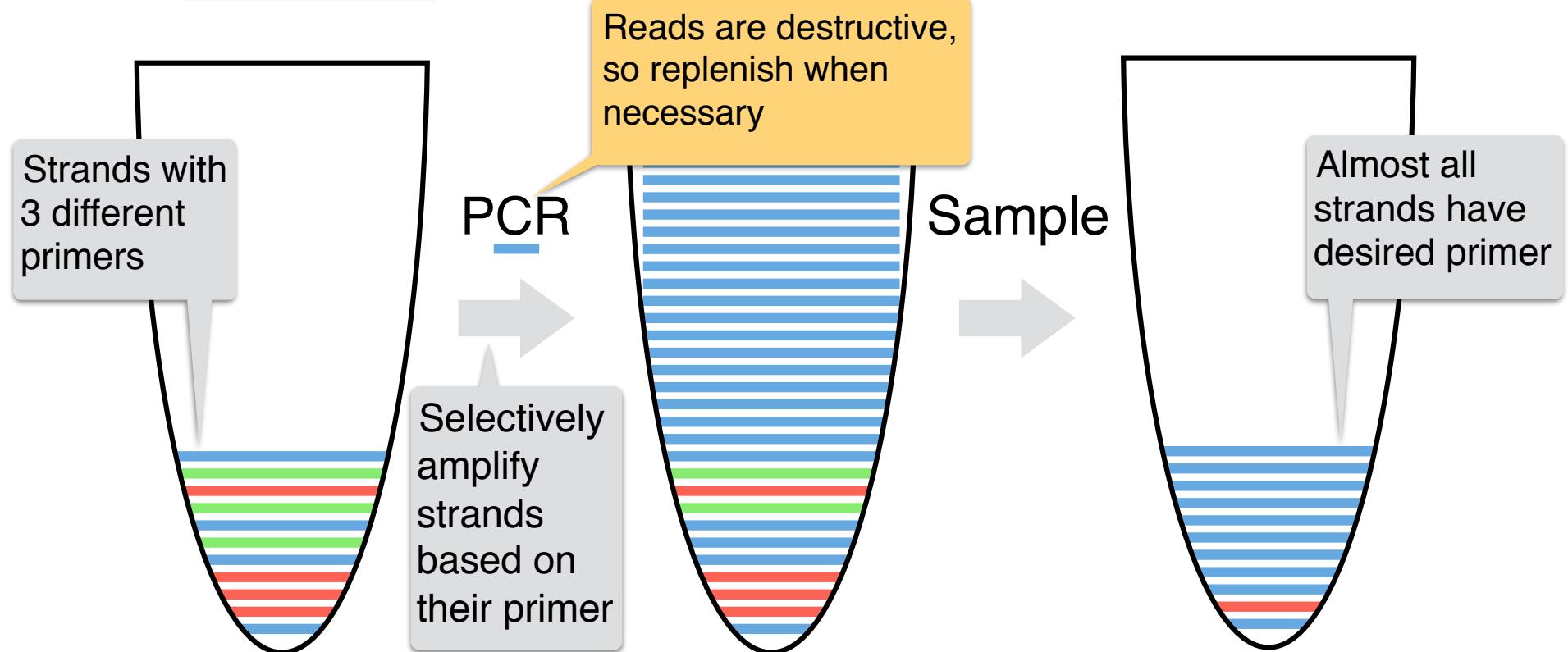
11011101

Random access?

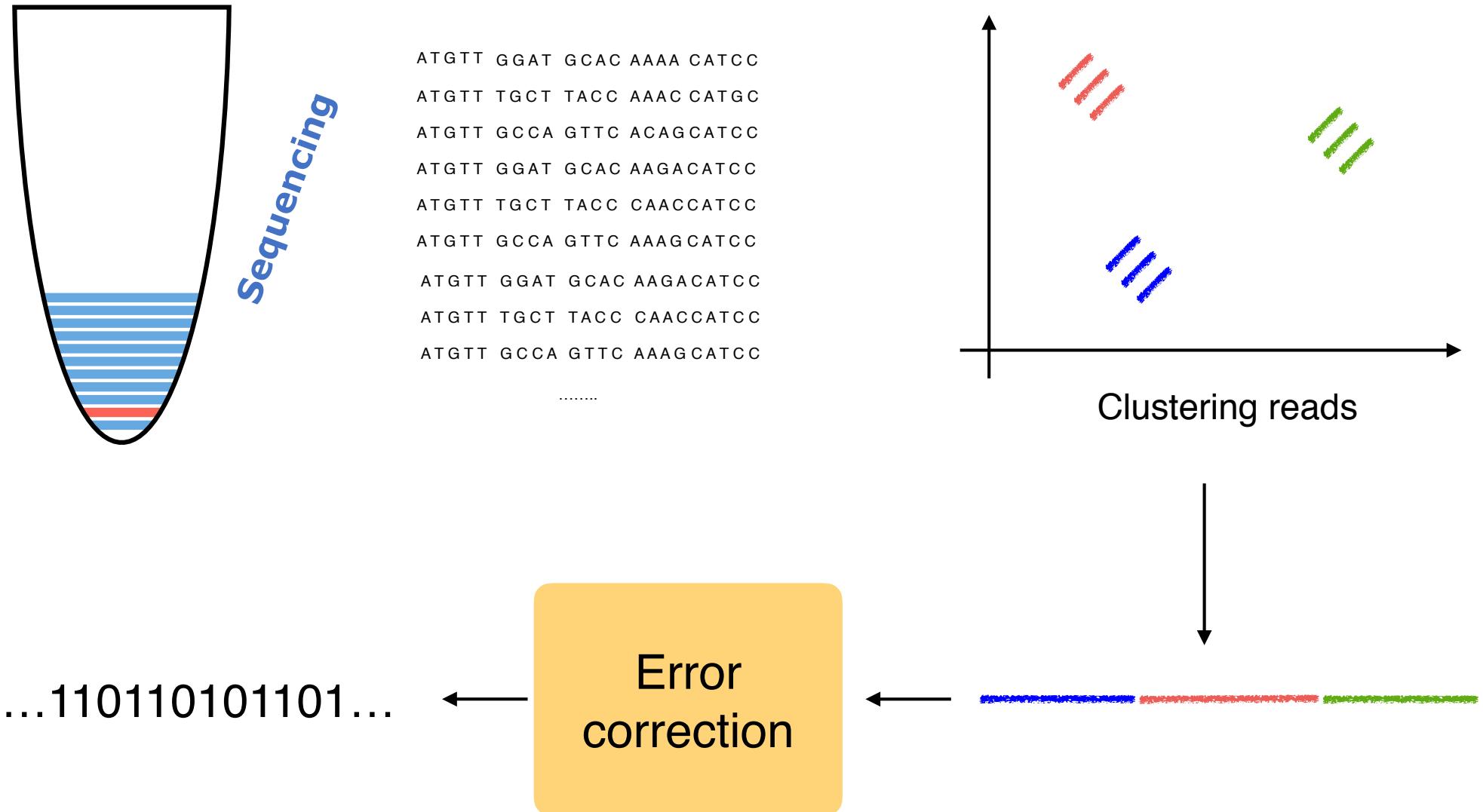


Random access!

ATGTT	GGAT	GCAC	AAAAA	CATCC
CCACT	TGCT	TACC	AAAC	GATAC
ATGTT	GCCAG	TTC	AAAG	TATCT
TACAA				ATAGA

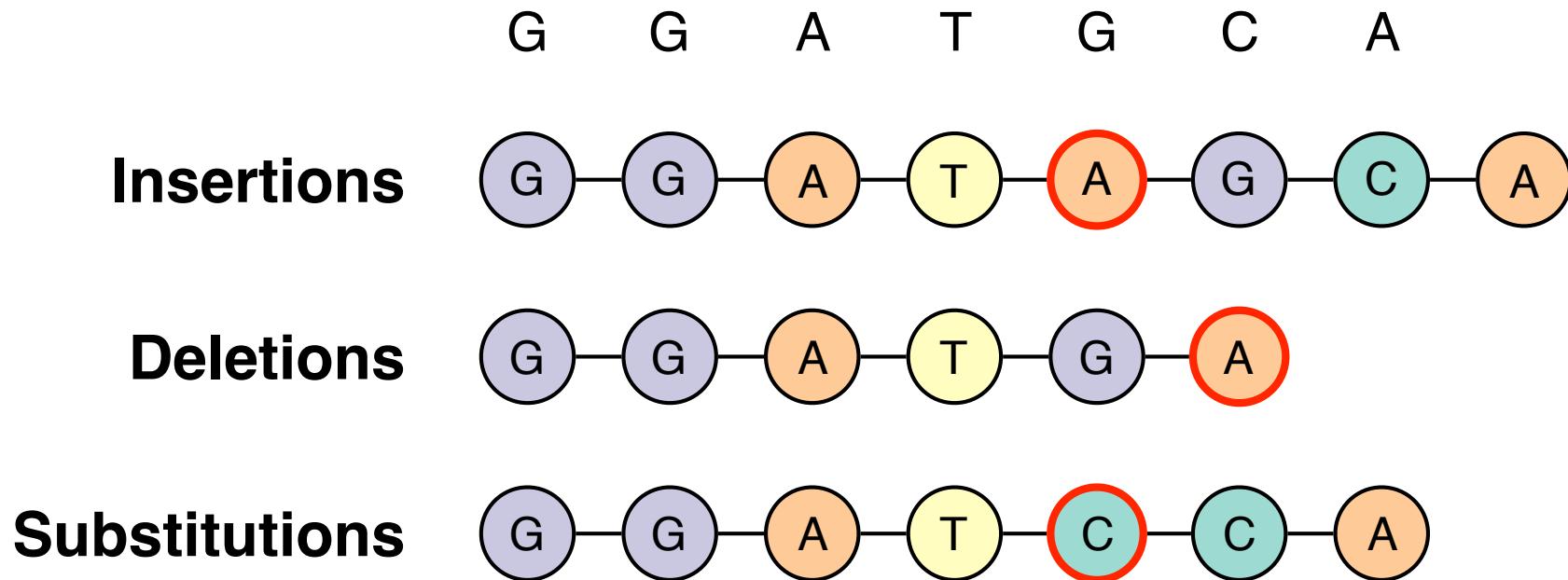


Going back to digital: Decoding



Error correction

Both synthesis and sequencing are error prone:



Aggregate error rates ~1% per position

Logical redundancy

Key identifiers
("primers")

Addresses
within the value

ATGTT	GGATGCAC	AAAA	CATCC
ATGTT	TGCTTACC	AAAC	CATCC
ATGTT	GCCAGTTC	AAAG	CATCC
ATGTT	TCGCTACG	CAAC	CATCC
ATGTT	TGCAATTG	CAAG	CATCC

Redundant data in
additional DNA strands.
Many possibilities: parity,
Reed Solomon, LDPC, ...

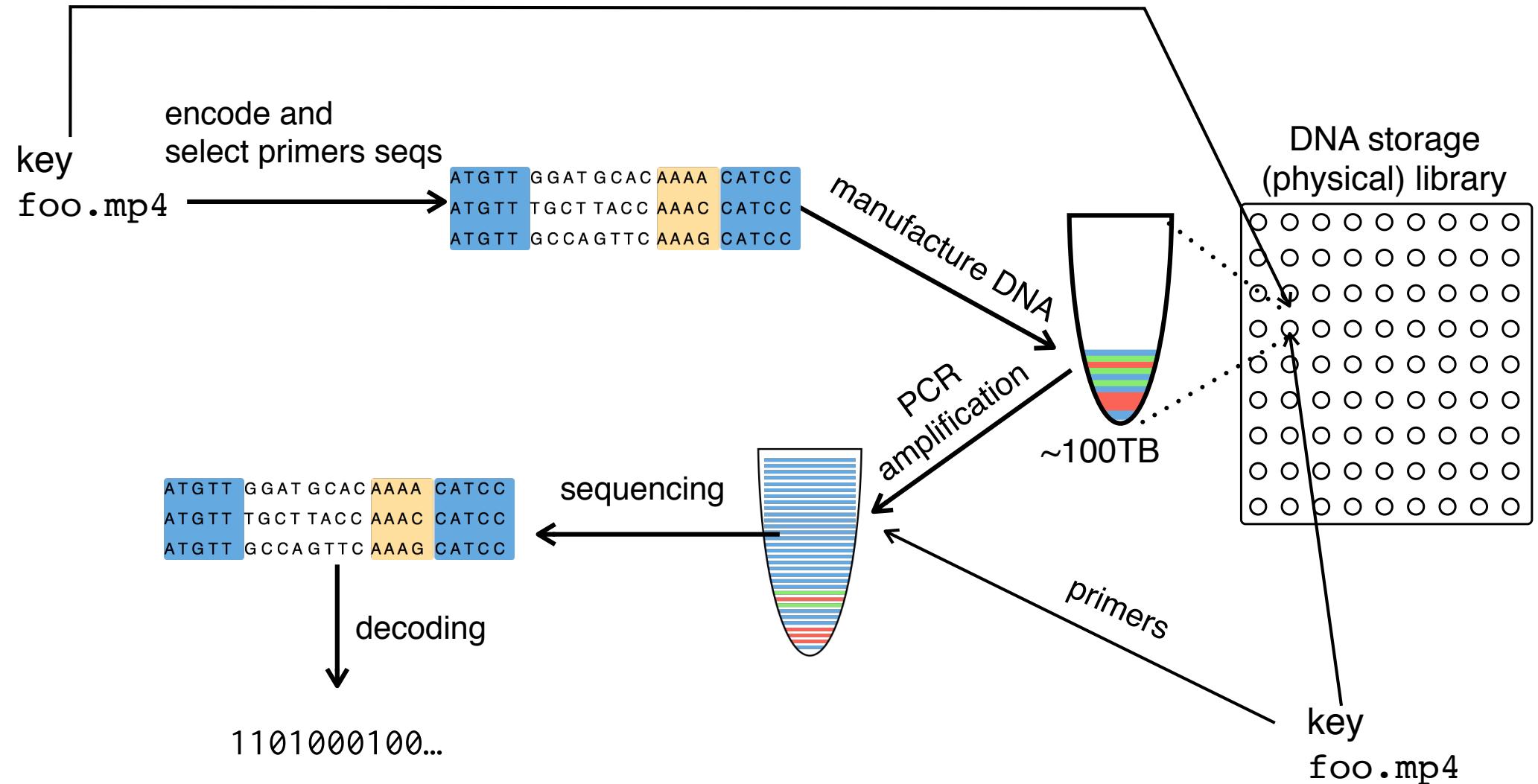
Putting it all together

SAS / SATA



Putting it all together

Data address/key specifies physical location and primer for random access.

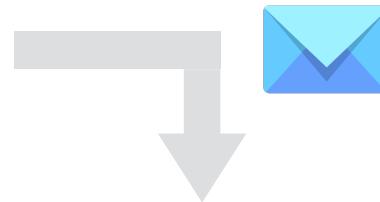


Wet lab results

Experiments



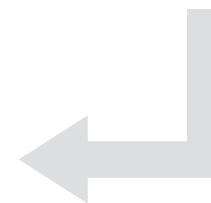
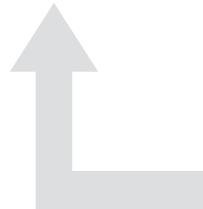
catcatgg



catcatg**c**



Throughput
MBs/week



FedEx

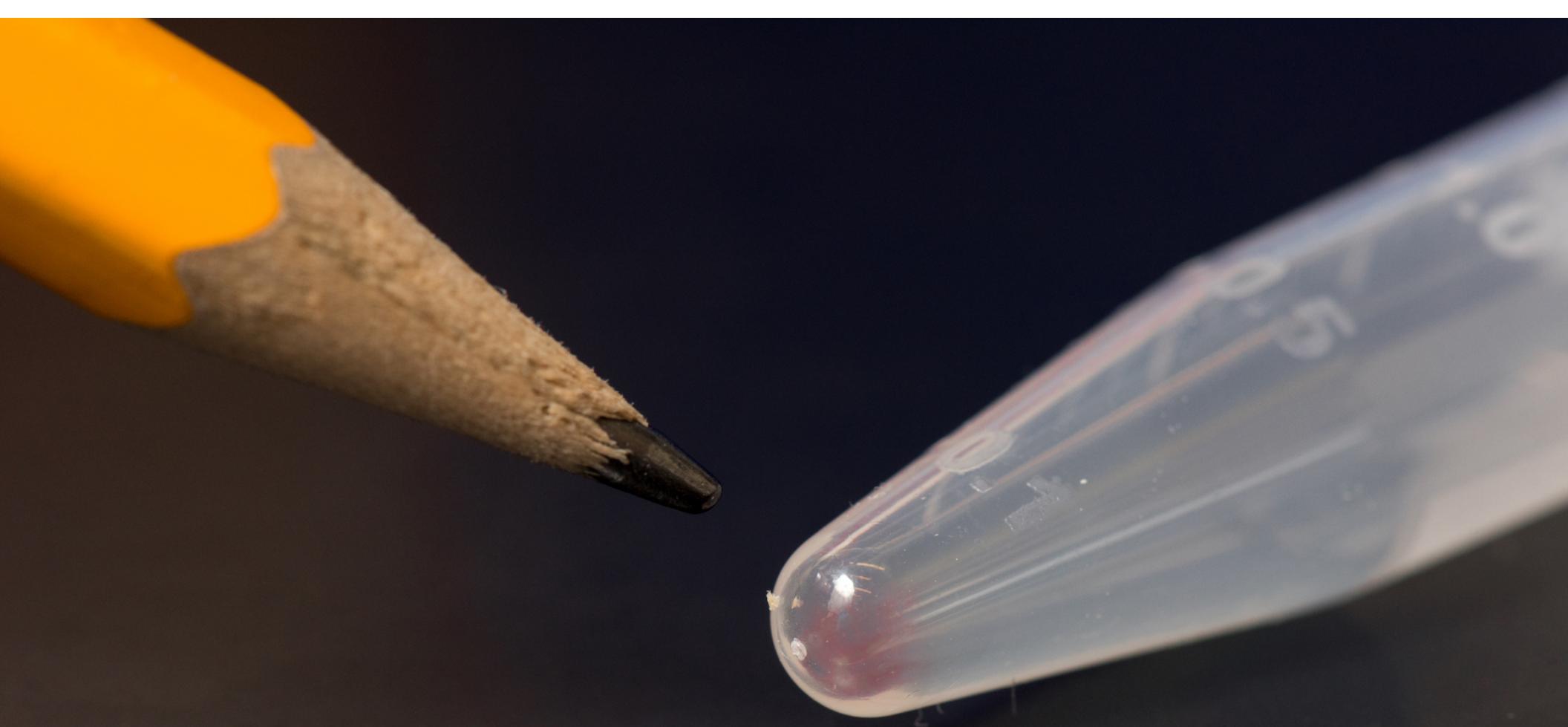


Photo: Tara Brown / UW

Decoding

Encoded and synthesized 3 files (151 kB):



Selected and PCRed one file for random access (42 kB):



Sequenced and decoded the resulting amplified pool:



Recovered *every bit* despite errors in synthesis and sequencing

The importance of redundancy

Key identifiers
("primers")

Addresses
within the value

ATGTT GGATGCACAAAAA CATCC

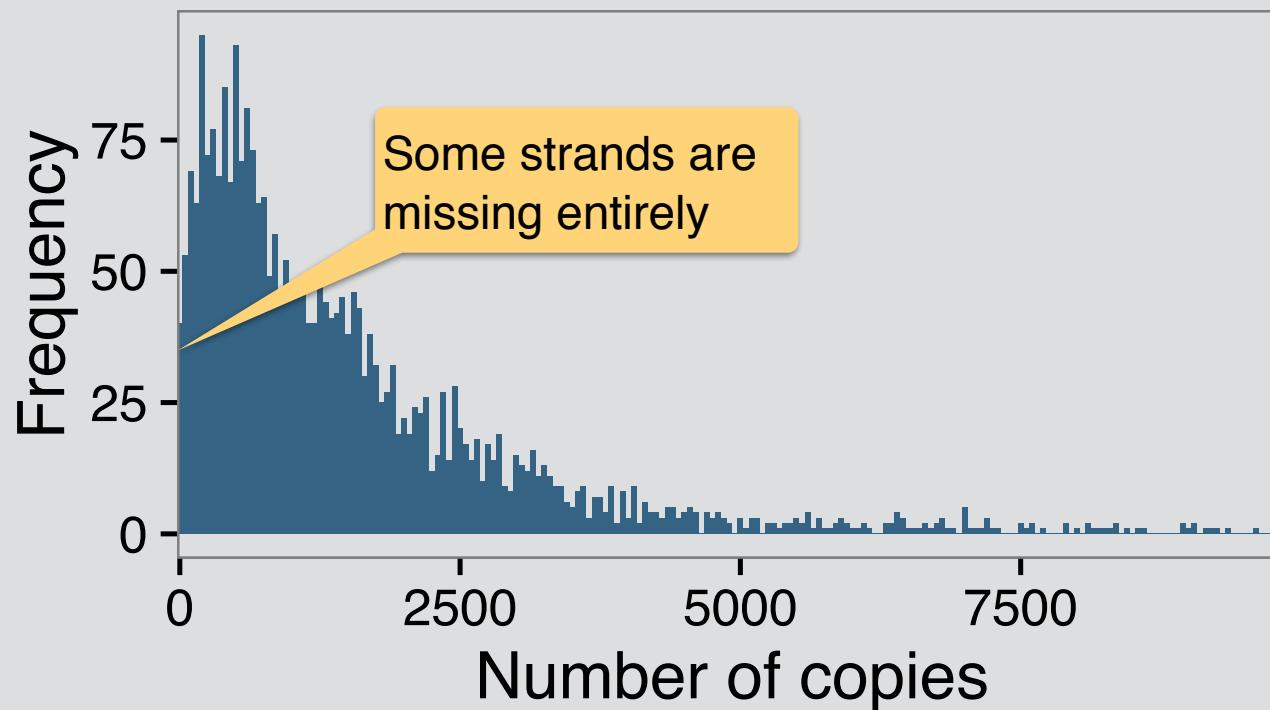
ATGTT

ATGTT

ATGTT

ATGTT

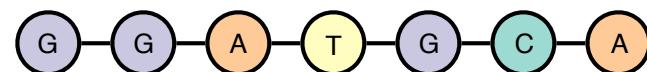
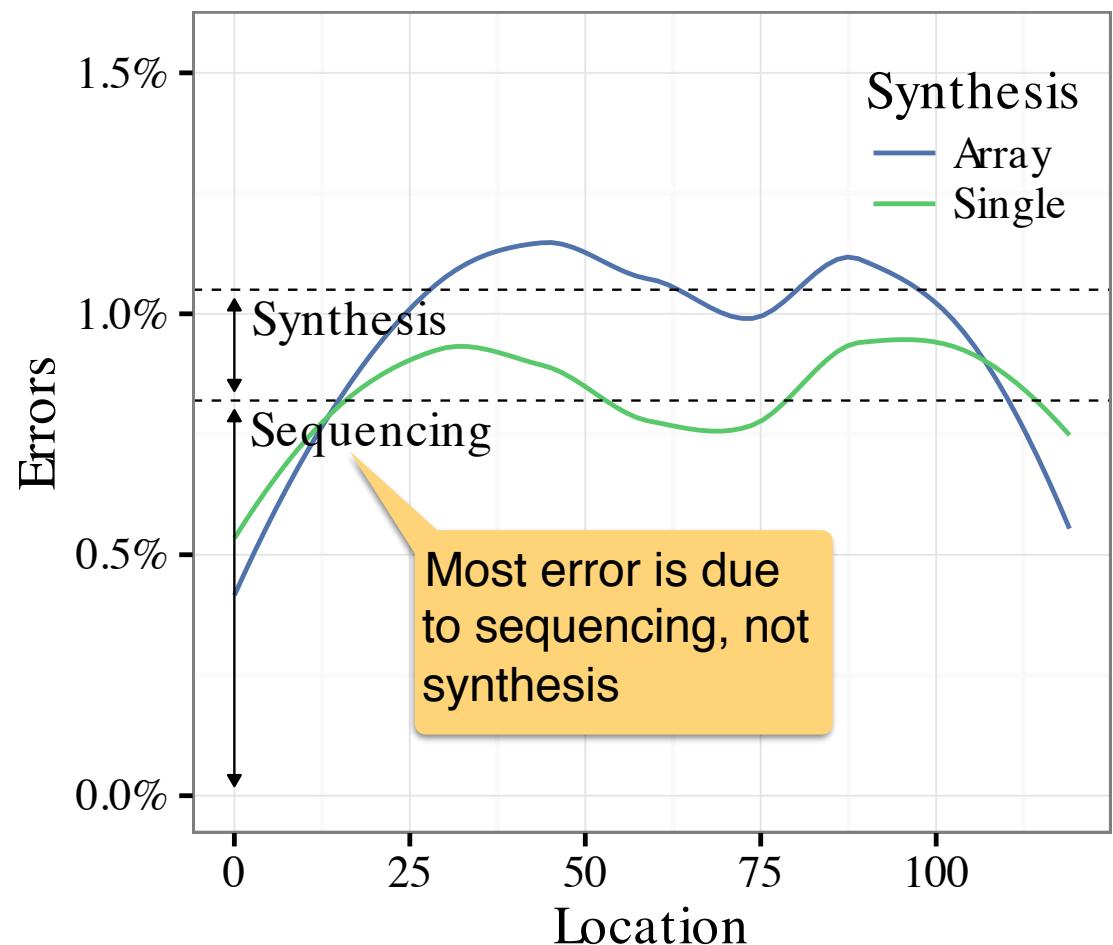
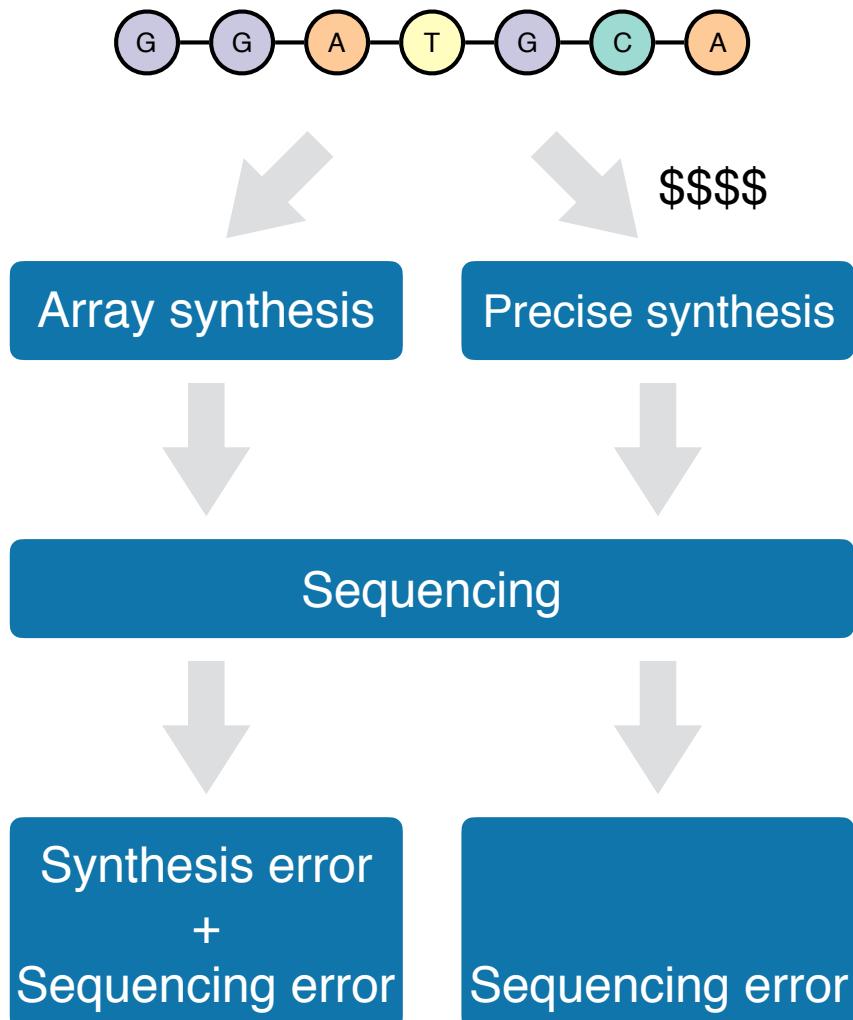
If we ignore redundancy data, we cannot recover the file.



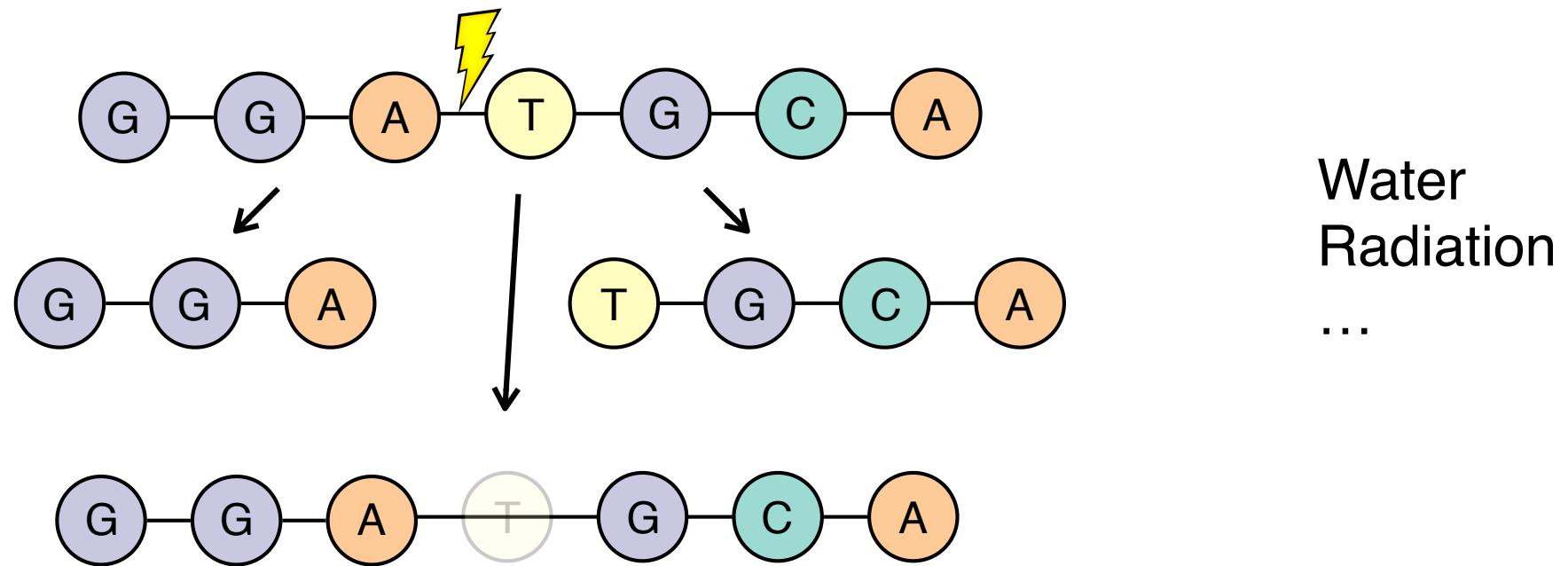
Redundant data in
multiple DNA strands.
Possible possibilities: parity,
Hamming, Reed-Solomon, LDPC, ...

Error Analysis: Synthesis and Sequencing

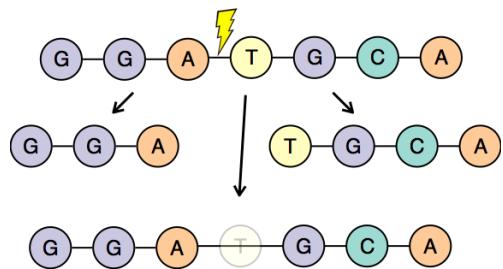
Both synthesis and sequencing are error prone, but how much?



Durability



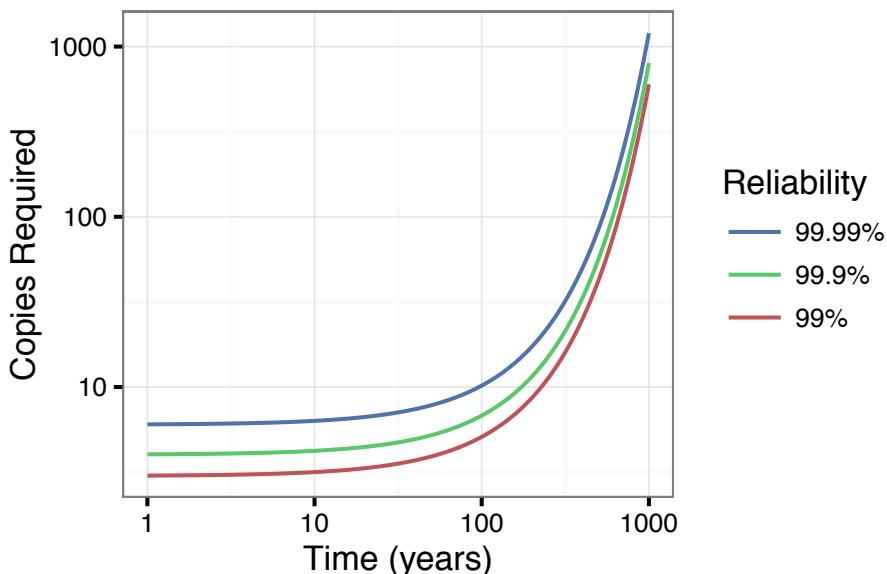
Durability



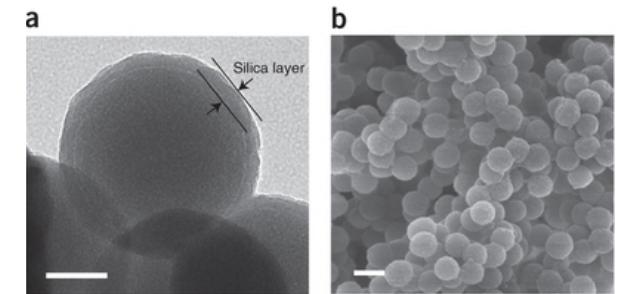
The half-life of DNA in bone: measuring decay kinetics in 158 dated fossils

Morten E. Allentoft, Matthew Collins, David Harker, James Haile, Charlotte L. Oskam, Marie L. Hale, Paula F. Campos, Jose A. Samaniego, M. Thomas P. Gilbert, Eske Willerslev, Guojie Zhang, R. Paul Scofield, Richard N. Holdaway, Michael Bunce

Half-life in bone: ~521 years.



ETH zürich



DNA synthetic fossils survive:

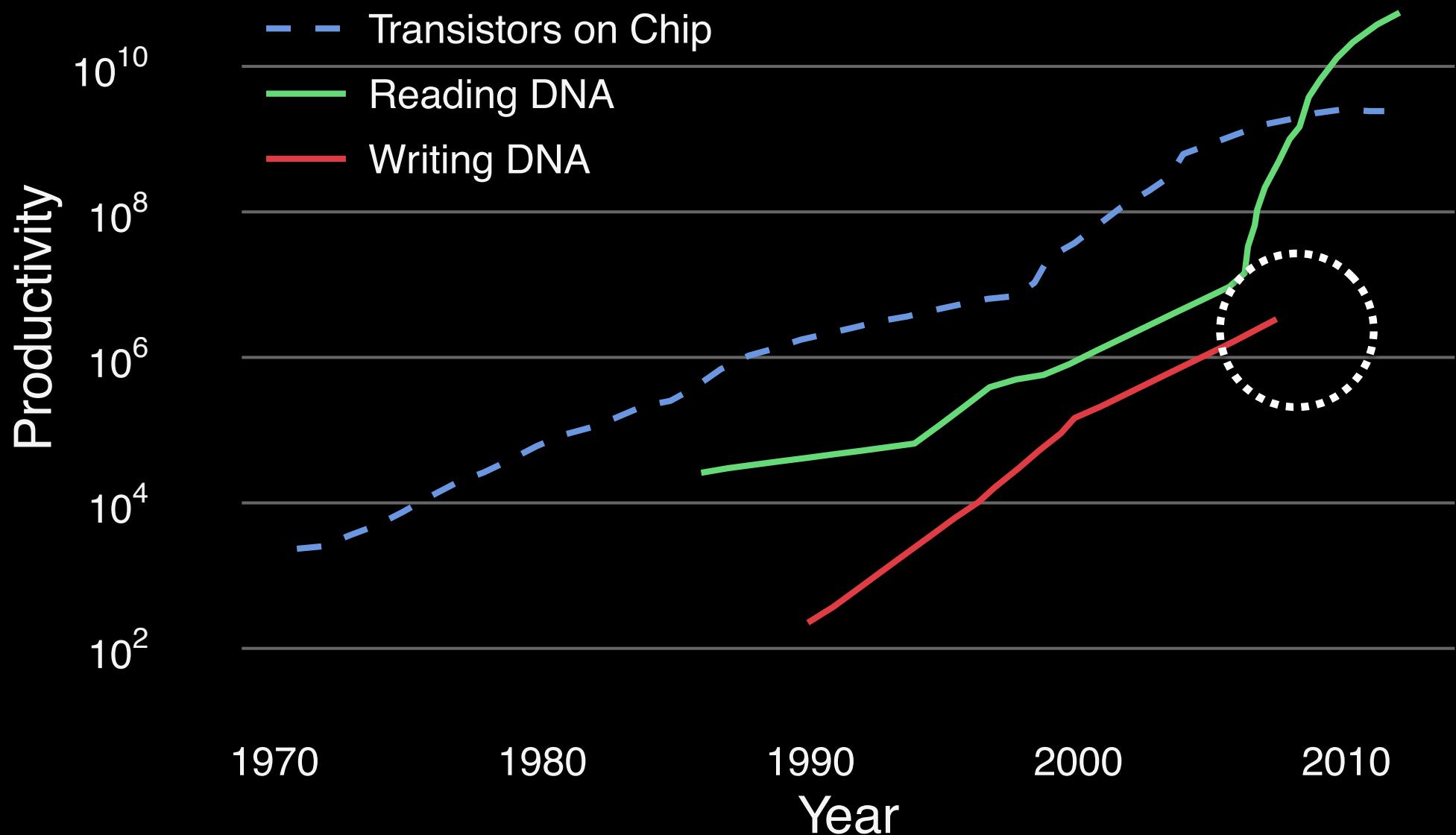
Time	Temperature
1 week	70°C
2,000 years	10°C
2,000,000 years	-18°C

Source: Grass et al. Robust Chemical Preservation of Digital Information on DNA in Silica with Error-Correcting Codes

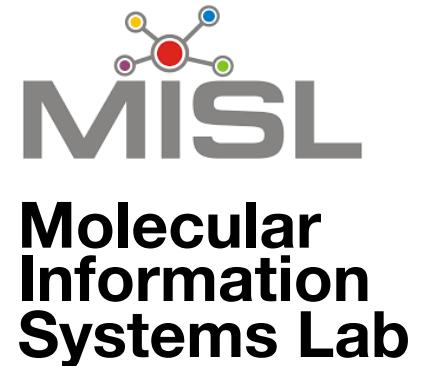
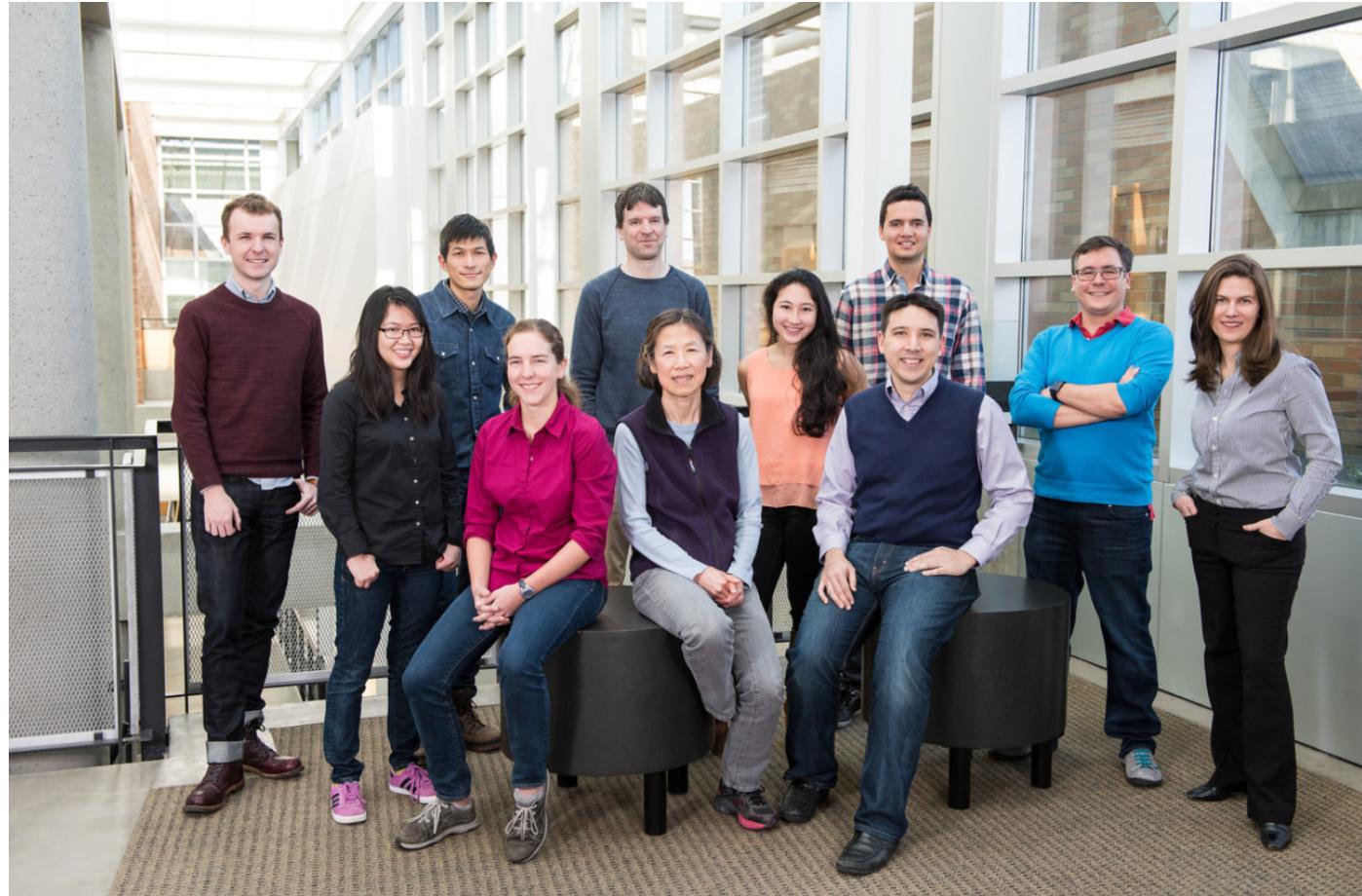
MBs/week → GBs/second

DNA manipulation productivity is growing

And cost is decreasing...



Thanks!



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