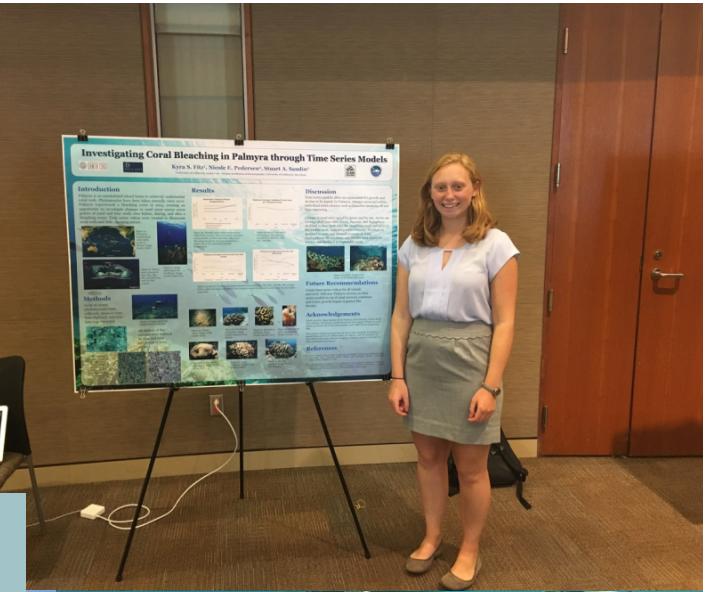


# Using ‘Omics to Answer Your Biological Questions

Slides by: Amanda S. Ackiss



**OLD DOMINION**  
UNIVERSITY

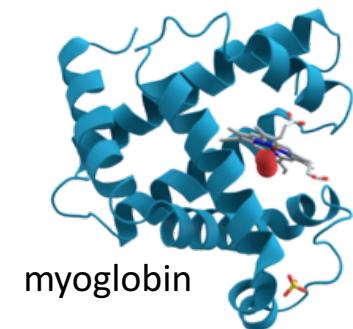
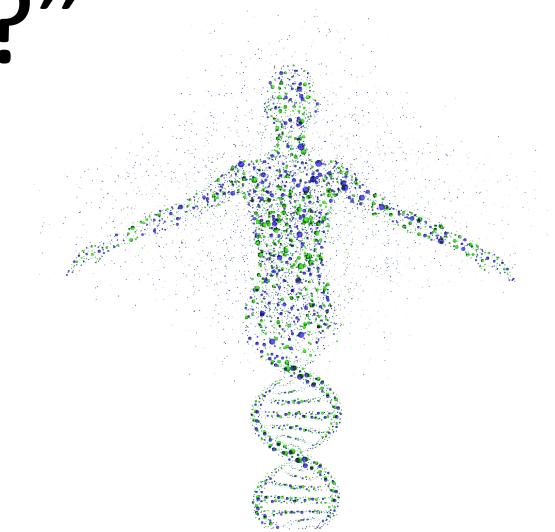


# Kyra Fitz

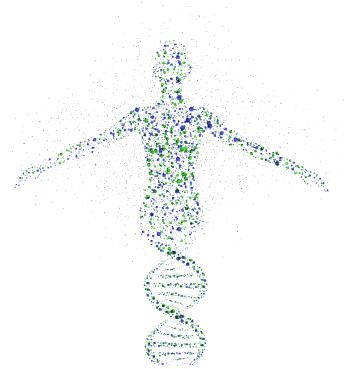


# What is “omics?”

- Genomics
  - The study of the structure, function, evolution and mapping of an organism's genes (the **genome**)
- Transcriptomics
  - The study of all the mRNA expressed (the **transcriptome**) from the genes in an organ or organism and their function
- Proteomics
  - The study of proteins that can be or are expressed by a cell, tissue, or organism (the **proteome**) and their function

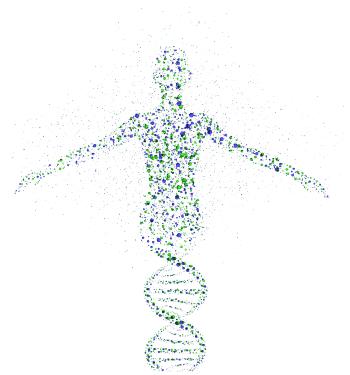


# Genomics



- Genetics vs Genomics
  - Genetics: the study of heredity, single genes, and their role in the way traits and conditions are passed from one generation to the next
  - Genomics: the study of the entirety of an organism's genes

# Genomics



- Applications:
  - Clinical and biomedical
    - Disease research & drug effects
  - Bioengineering
    - Creation of synthetic species



# Genomics



## Genomic islands of divergence and their consequences for the resolution of spatial structure in an exploited marine fish

Ian R. Bradbury,<sup>1,2,3</sup> Sophie Hubert,<sup>4</sup> Brent Higgins,<sup>4</sup> Sharen Bowman,<sup>4</sup> Tudor Borza,<sup>4,5</sup> Ian G. Paterson,<sup>2</sup> Paul V. R. Snelgrove,<sup>3</sup> Corey J. Morris,<sup>1</sup> Robert S. Gregory,<sup>1</sup> David Hardie,<sup>2</sup> Jeffrey A. Hutchings,<sup>2</sup> Daniel E. Ruzzante,<sup>2</sup> Christopher T. Taggart<sup>6</sup> and Paul Bentzen<sup>2</sup>

1 Fisheries and Oceans Canada, St. John's, NF, Canada

2 Department of Biology, Marine Gene Probe Laboratory, Dalhousie University Halifax, NS, Canada

3 Ocean Sciences Center and Biology Department, Memorial University of Newfoundland St. John's, NF, Canada

4 Atlantic Genome Center Halifax, NS, Canada

5 Department of Plant and Animal Sciences, Faculty of Agriculture, Dalhousie University Truro, NS, Canada

6 Department of Oceanography, Dalhousie University Halifax, NS, Canada

## Complete Genome Sequence of the Marine Fish Pathogen *Vibrio anguillarum* Harboring the pJM1 Virulence Plasmid and Genomic Comparison with Other Virulent Strains of *V. anguillarum* and *V. ordalii*<sup>▽†</sup>

Hiroaki Naka,<sup>1</sup> Graciela M. Dias,<sup>2</sup> Cristiane C. Thompson,<sup>3</sup> Christopher Dubay,<sup>1</sup> Fabiano L. Thompson,<sup>2</sup> and Jorge H. Crosa<sup>1\*</sup>

Department of Molecular Microbiology and Immunology, Oregon Health and Science University, Portland, Oregon 97239<sup>1</sup>; Institute of Biology, Federal University of Rio de Janeiro, UFRJ, Brazil<sup>2</sup>; and Laboratory of Molecular Genetics of Microorganisms, Oswaldo Cruz Institute, FIOCRUZ, Rio de Janeiro, Brazil<sup>3</sup>

# Transcriptomics



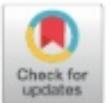
- ‘Functional’ genomic approach
- Connects the genome to gene function
- Typically represents less than 5% of the **genome**
- The **transcriptome** is the precursor to the **proteome**



# Transcriptomics

- Applications:
  - Clinical and biomedical
    - Examining the processes of cellular differentiation and carcinogenesis
    - Embryology and development
  - Biological and conservation sciences
    - Inferring phylogenetic relationships
    - Examining functional responses relative to biotic and abiotic stressors

# Transcriptomics



Transcriptomics reveals tissue/organ-specific differences in gene expression  
in the starfish *Patiria pectinifera*

Chan-Hee Kim<sup>a</sup>, Hye-Jin Go<sup>a</sup>, Hye Young Oh<sup>a</sup>, Yong Hun Jo<sup>b</sup>, Maurice R. Elphick<sup>c</sup>,  
Nam Gyu Park<sup>a,\*</sup>

<sup>a</sup> Department of Biotechnology, College of Fisheries Sciences, Pukyong National University, 45 Yongso-ro, Nam-gu, Busan 48513, Republic of Korea

<sup>b</sup> Division of Plant Biotechnology, Institute of Environmentally-Friendly Agriculture (IEFA), College of Agriculture and Life Sciences, Chonnam National University, Gwangju 500-757, Republic of Korea

<sup>c</sup> School of Biological and Chemical Sciences, Queen Mary University of London, London E1 4NS, UK

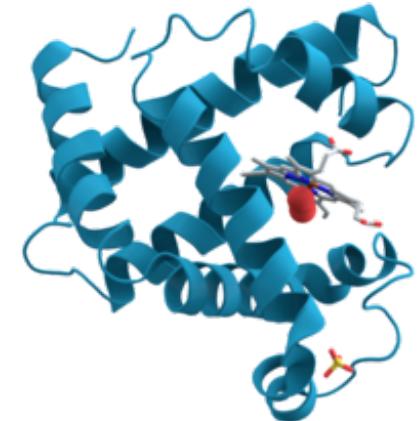
**Cellular stress responses to chronic heat shock and shell damage  
in temperate *Mya truncata***

Victoria A. Sleight<sup>1,2</sup> • Lloyd S. Peck<sup>2</sup> • Elisabeth A. Dyrynda<sup>3</sup> • Valerie J. Smith<sup>4</sup> • Melody S. Clark<sup>2</sup>

Received: 31 January 2018 / Revised: 6 April 2018 / Accepted: 1 May 2018

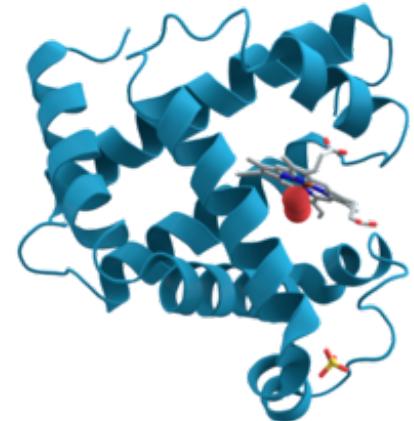
© The Author(s) 2018

# Proteomics



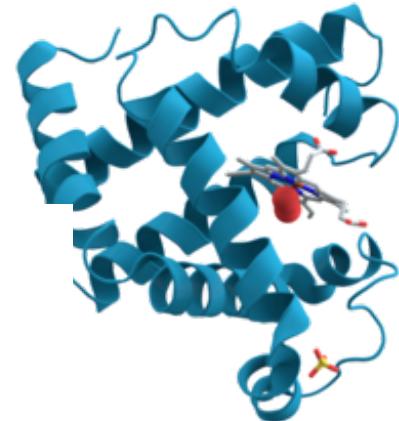
- protein + genome = *proteome*
- Why study the proteome?
  - mRNA is not always translated into protein
  - Relatively small changes in mRNA expression can produce large changes in the total amount of the corresponding protein present in the cell
- Proteomics confirms the presence of the protein & provides a direct measure of the quantity present!

# Proteomics



- Applications:
  - Clinical and biomedical
    - Biomarkers for diagnosing disease
  - Systems biology
    - Analysis of cellular systems
    - Cell-signaling cascades and gene regulatory networks
  - Conservation sciences
    - Examining functional responses relative to biotic and abiotic stressors

# Proteomics



## Exploring the Environmental Physiology of the Indo-Pacific Reef Coral *Seriatopora hystrix* with Differential Proteomics

Anderson B. Mayfield<sup>1,2\*</sup>, Yi-Jyun Chen<sup>2</sup>, Chi-Yu Lu<sup>3,4</sup>, Chii-Shiarng Chen<sup>2,5,6</sup>

<sup>1</sup>Khaled bin Sultan Living Oceans Foundation, USA

<sup>2</sup>Taiwan Coral Research Center, Taiwan

<sup>3</sup>Department of Biochemistry, College of Medicine, Kaohsiung Medical University, Taiwan

<sup>4</sup>Center for Research Resources and Development, Kaohsiung Medical University, Taiwan

<sup>5</sup>Department of Marine Biotechnology and Resources, National Sun Yat-Sen University, Taiwan

<sup>6</sup>Graduate Institute of Marine Biotechnology, National Dong-Hwa University, Taiwan

Email: \*andersonblairmayfield@gmail.com

## From Marine Venoms to Drugs: Efficiently Supported by a Combination of Transcriptomics and Proteomics

Bing Xie<sup>1,†</sup>, Yu Huang<sup>2,†</sup>, Kate Baumann<sup>3,†</sup>, Bryan Grieg Fry<sup>3,\*</sup> and Qiong Shi<sup>2,4,\*</sup>

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<sup>4</sup> BGI Shenzhen Academy of Marine Sciences, BGI Fisheries, BGI, Shenzhen 518083, China

\* Correspondence: bgfry@uq.edu.au (B.G.F.); shiqiong@genomics.cn (Q.S.);

Tel.: +61-4-0019-3182 (B.G.F.); +86-755-3630-7807 (Q.S.)

† These authors contributed equally to this work.

# Sequencing Methods

## First and Next Generation



by Dr. Eric Garcia

# Outline

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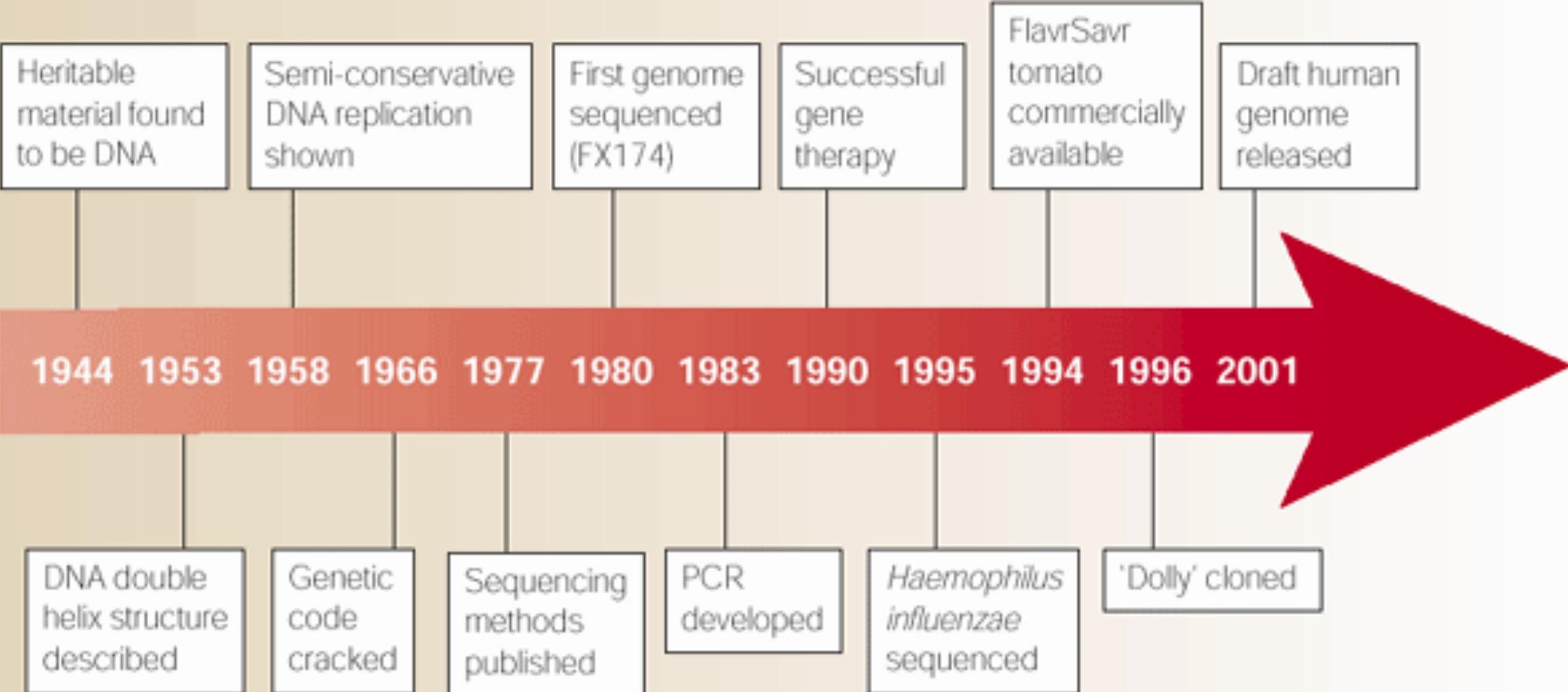
## INTRO

- Timeline of technological advancements in genetics
  - Sanger sequencing
  - PCR
  - Next Gen

## NEXT GENERATION

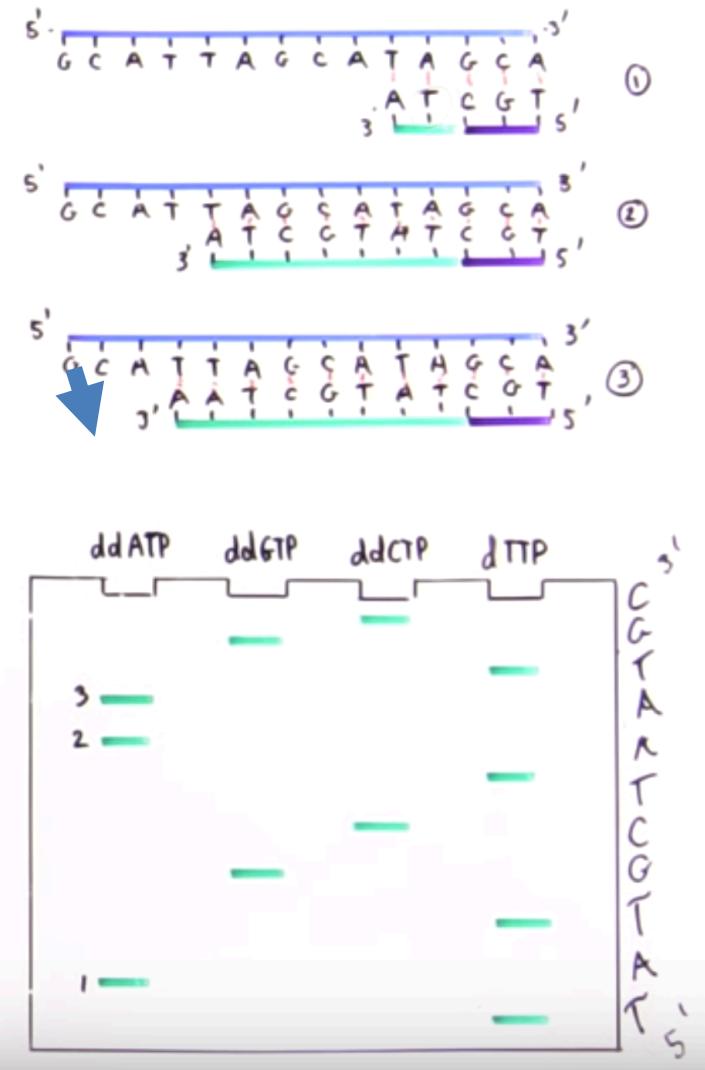
- Template preparation
- Sequencing

## Timeline | DNA milestones



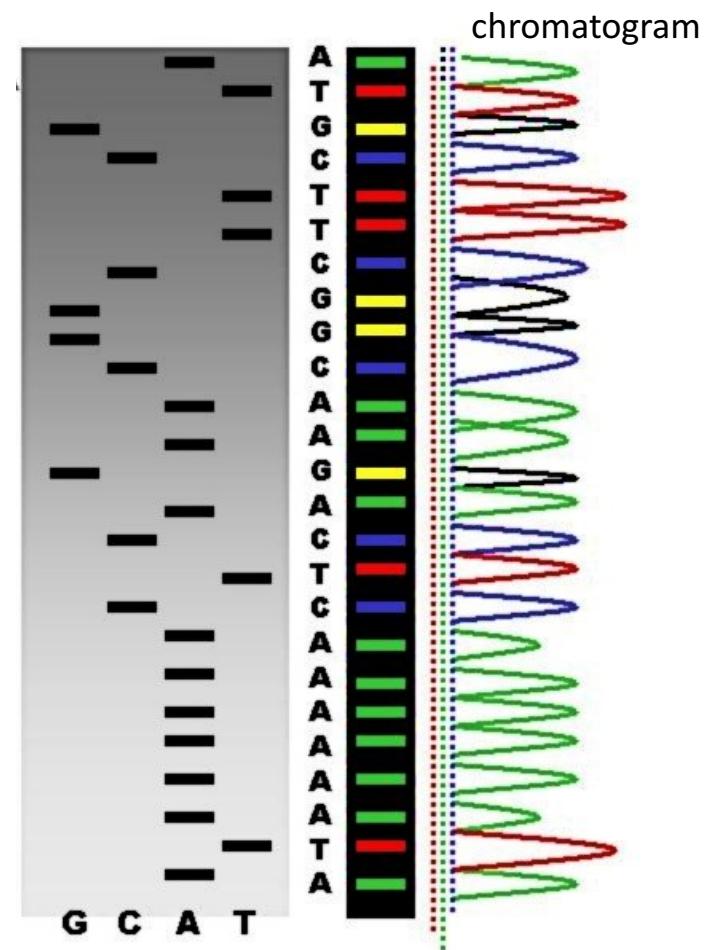
# Sanger Sequencing

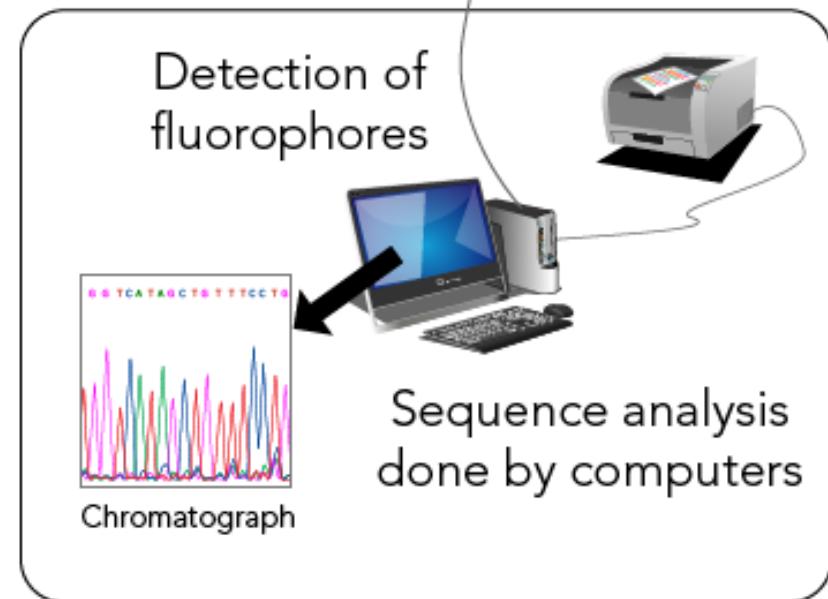
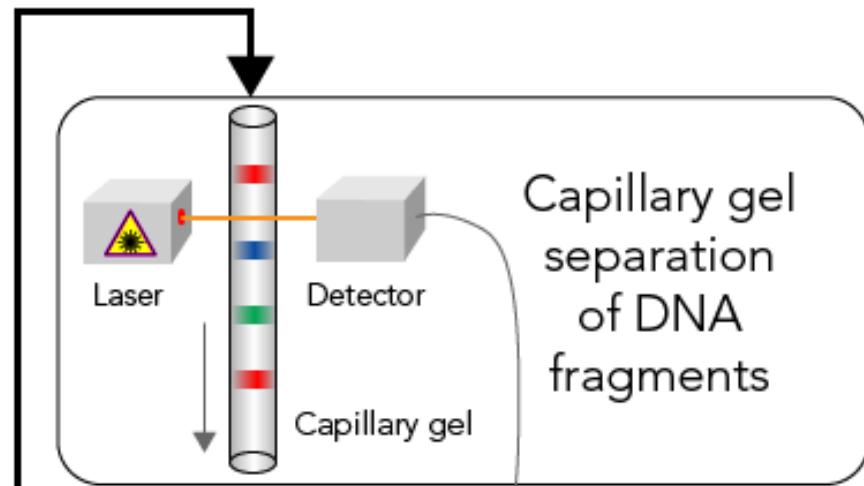
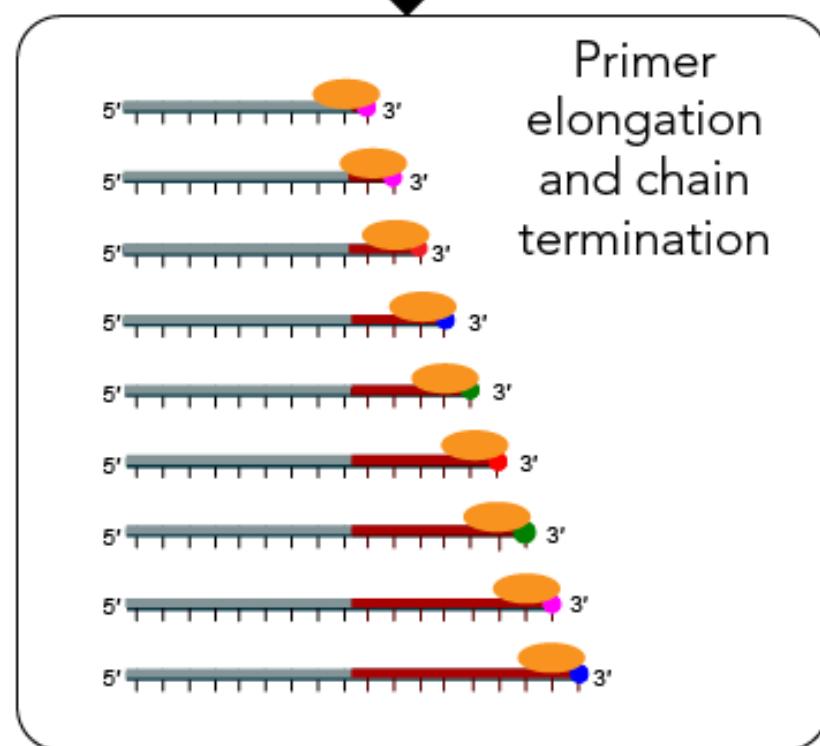
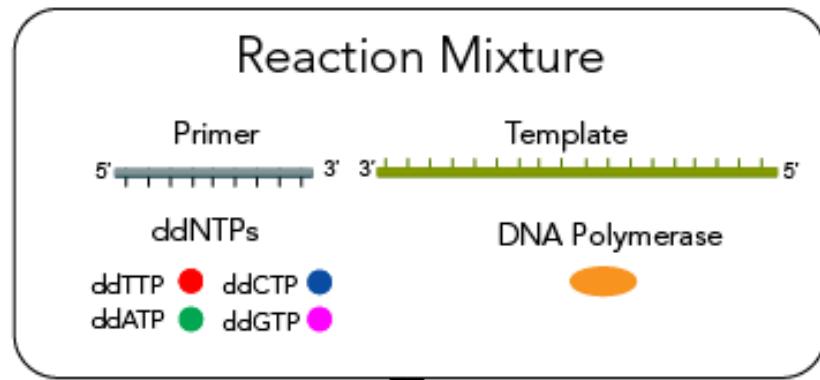
- Developed in 1977 by Frederick Sanger
- 1. Denature DNA
- 2. Mix template with:
  - Primer
  - Polymerase
  - All 4 dNTP
  - Then add only **ONE type of ddNTP**
- 3. Repeat this with the other 3 ddNTP
- 4. Gel reactions in diff. lanes



# Sanger Sequencing

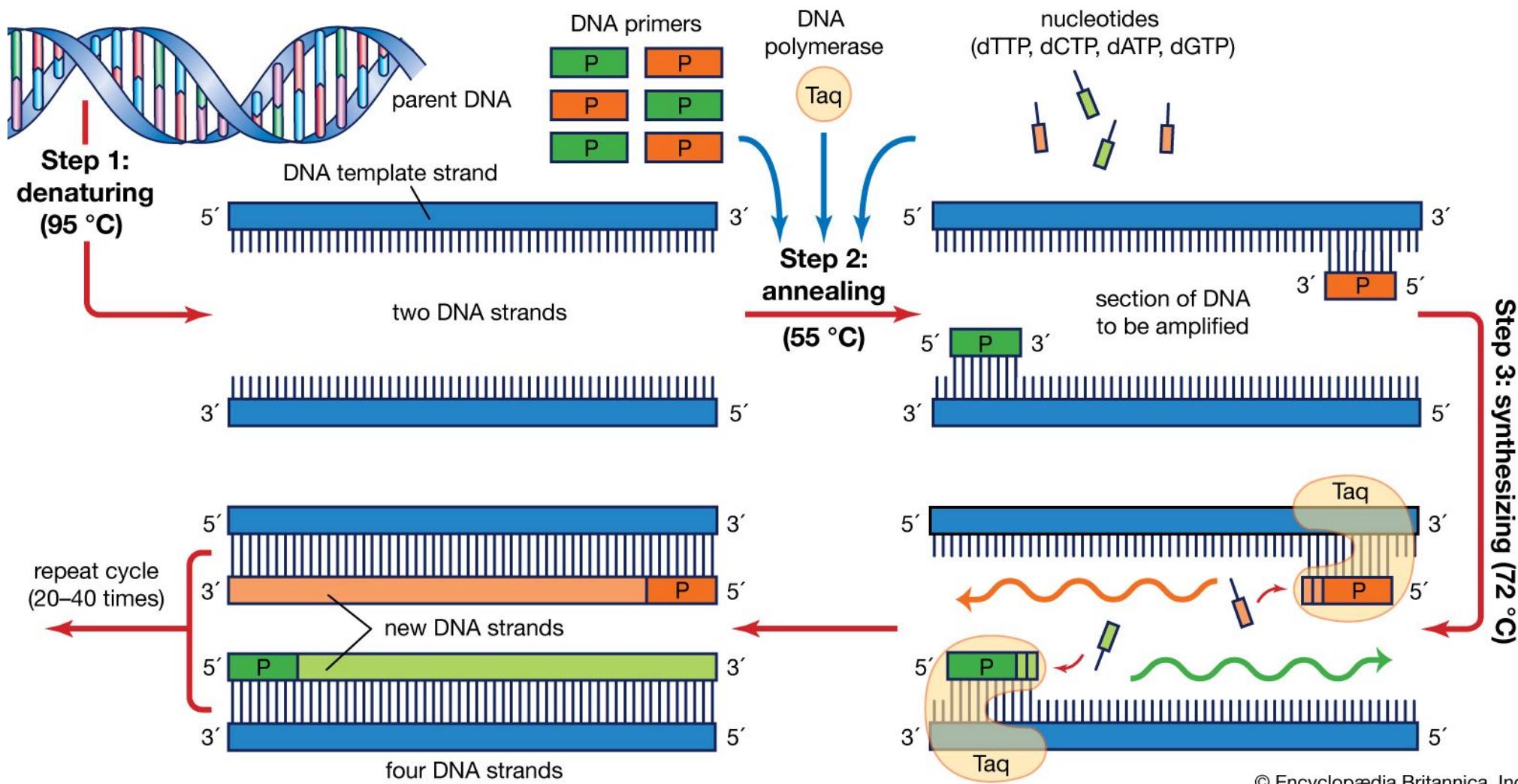
- Developed in 1977 by Frederick Sanger
1. Denature DNA
  2. Mix template with:
    - Primer
    - Polymerase
    - All 4 dNTP
    - Then add only **ONE type of ddNTP**
  3. Repeat this with the other 3 ddNTP
  4. Gel reactions in diff. lanes
  5. Expose to X-ray





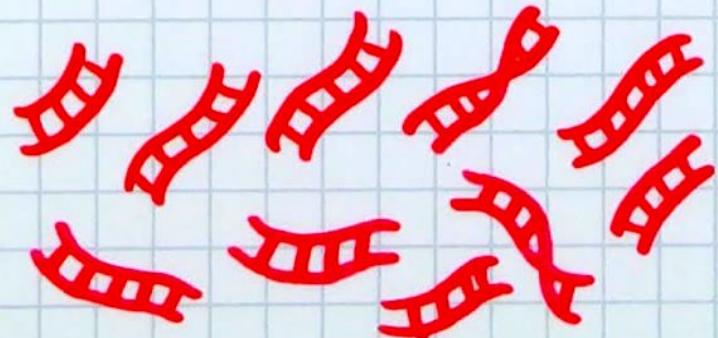
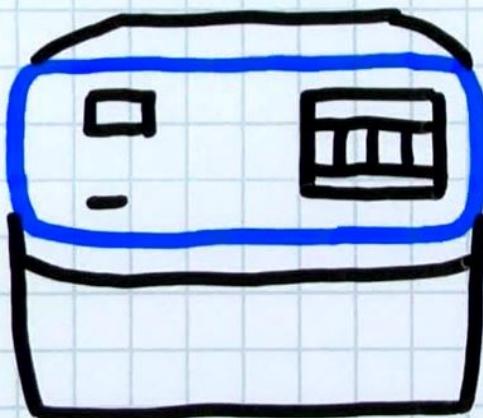
# PCR

Developed by Mullis 1985

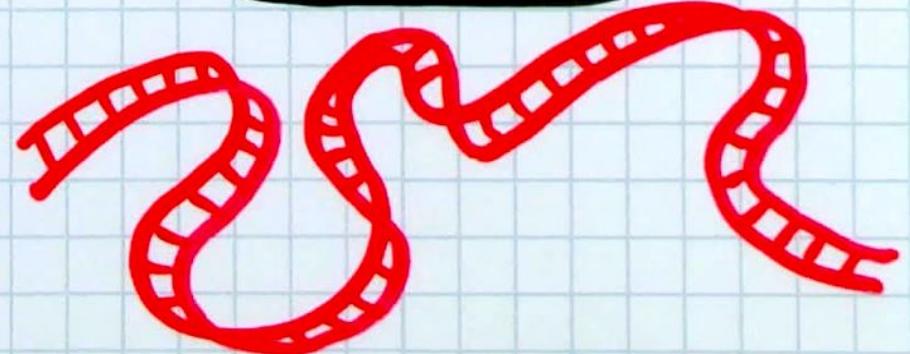
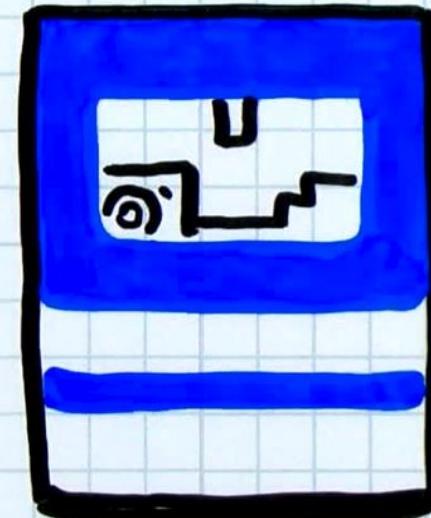


# Next Generation Sequencing

NGS  
MASSIVELY  
PARALLEL



SANGER



Fragmented  
input DNA



End Repair



DNA Library  
Prep

dA Tailing



Adaptor Ligation



Size Selection



Amplification



# Sequencing technologies – the next generation

---

- Template preparation
  - Different types of preparations
  - Different types of loading platforms
- Sequencing
  - By CTR
  - By ligation
  - By Single-nucleotide addition
  - Real Time Sequencing

# Template Preparation

- Clonally amplified templates

Requires  DNA (20x more)

PCR

Immobilizing:

- 1.- BEADS, chemically crosslinked to glass surface
- 2.- CLUSTERS, covalently attached (adaptors)

- Single DNA molecule Template

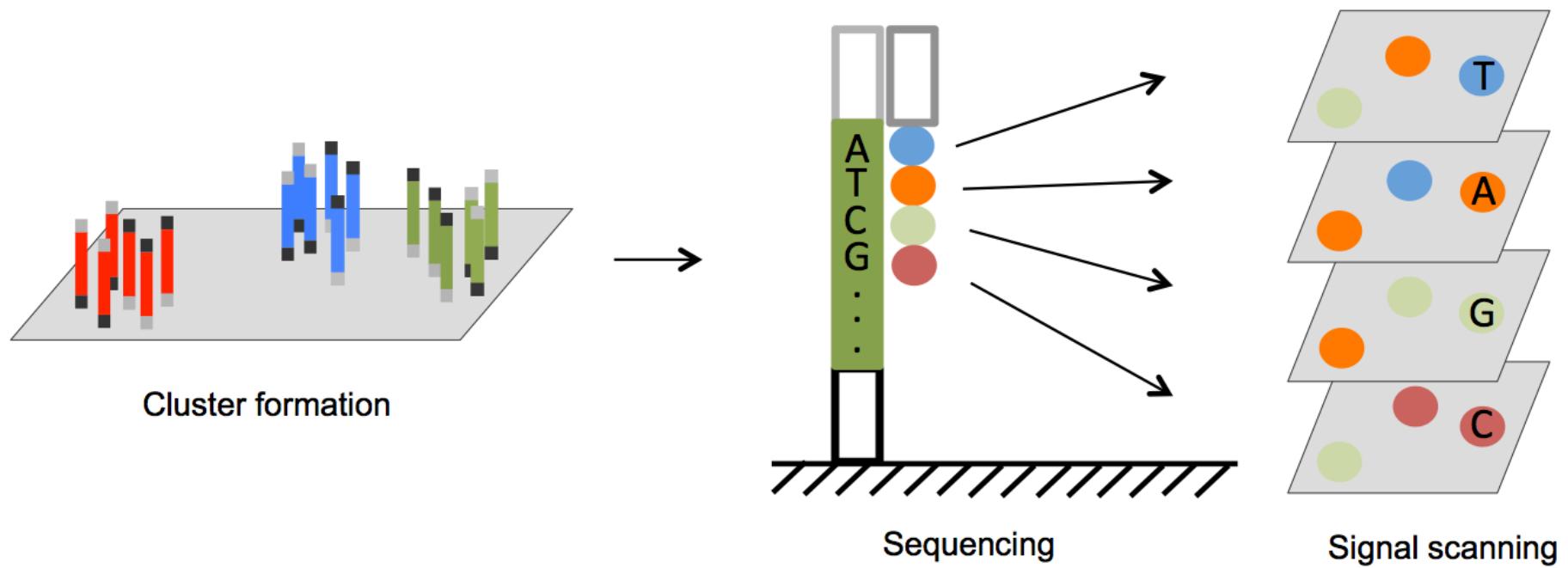
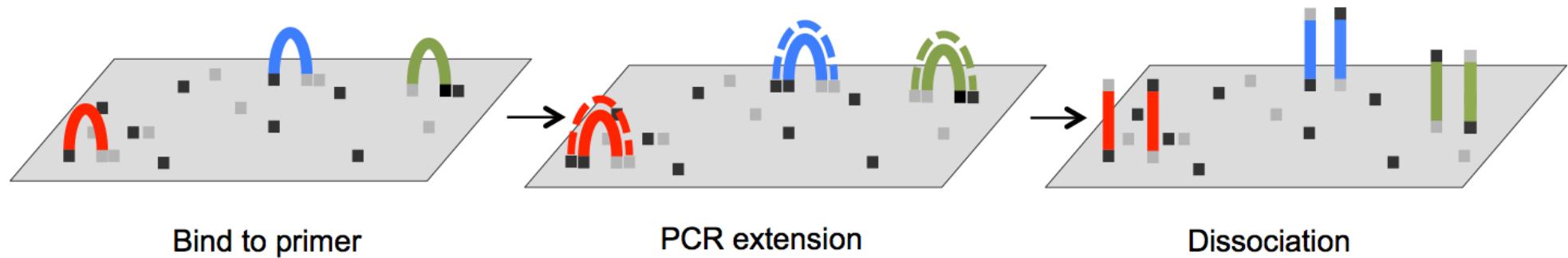
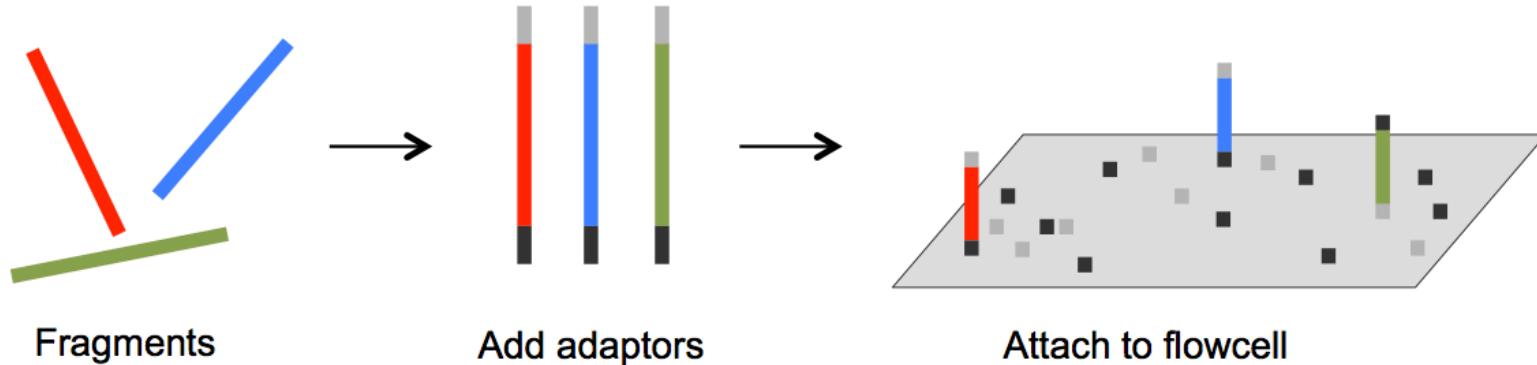


Requires  DNA

No PCR

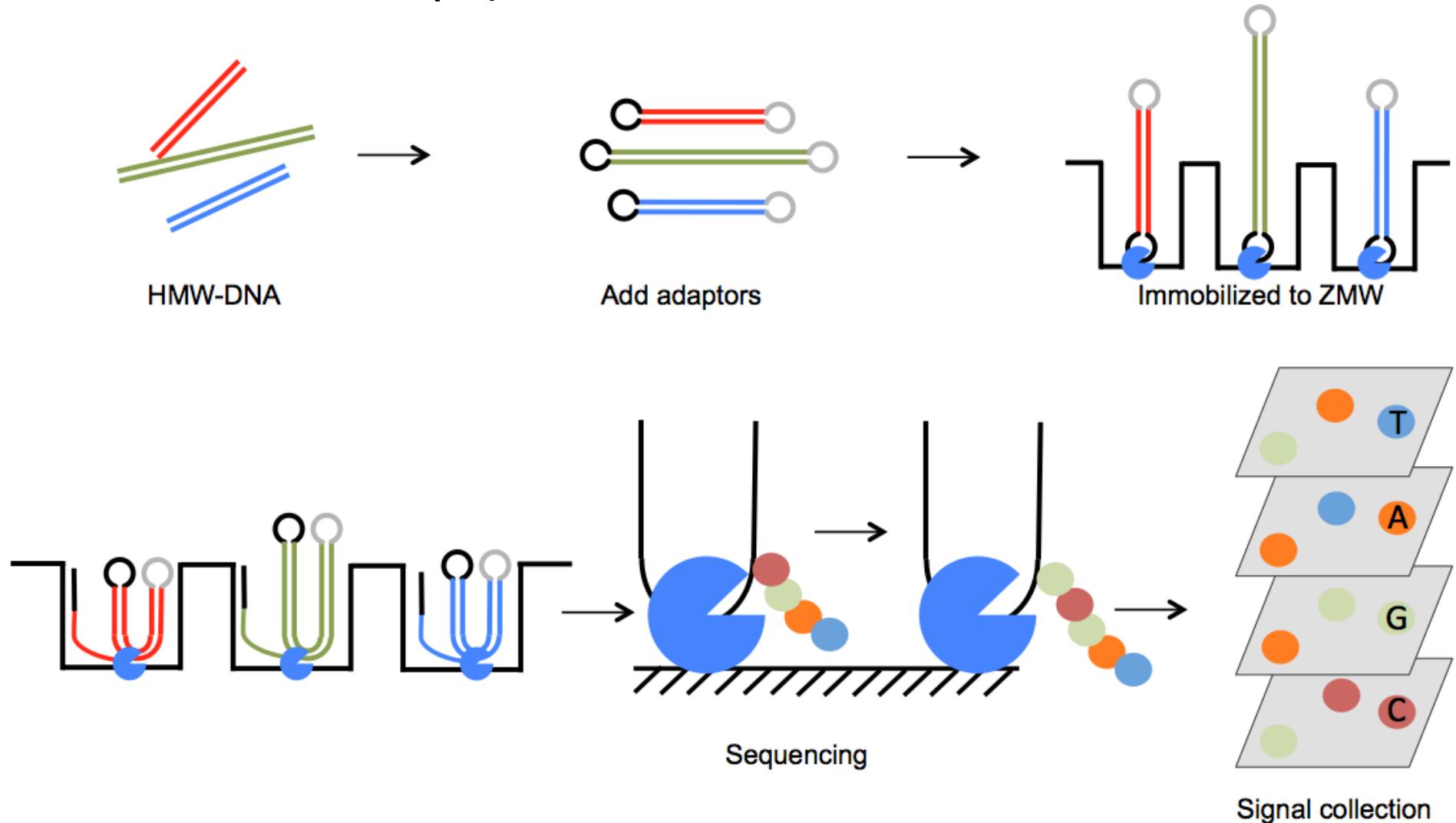
Immobilizing:

- 1.- templates
- 2.- primer
- 3.- polymerase

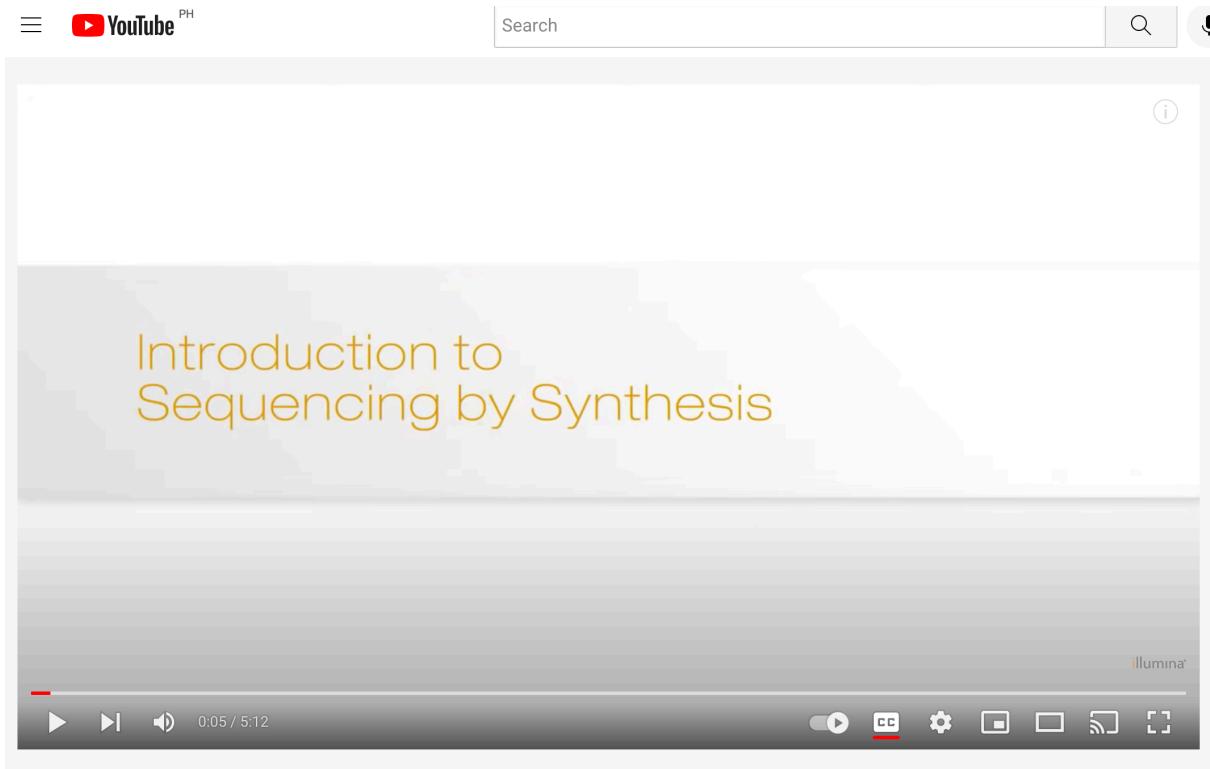


# Single DNA Molecule Template- PacBio

Immobilizing polymerases = longer reads (1000s to 10,000s bps)



# Illumina Online Resources



<https://www.youtube.com/watch?v=fCd6B5HRaZ8&t=>