

Science Enrichment Program (SEP)

Introduction to Cancer Biology

Dr. Chakravarty

Assistant Attending, Department of Pathology and Laboratory Medicine

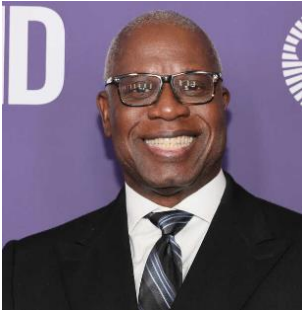
Lead Scientist, OncoKB, Center of Molecular Oncology

11 December 2024, HOPP Science Enrichment Program (SEP)



**Memorial Sloan Kettering
Cancer Center**

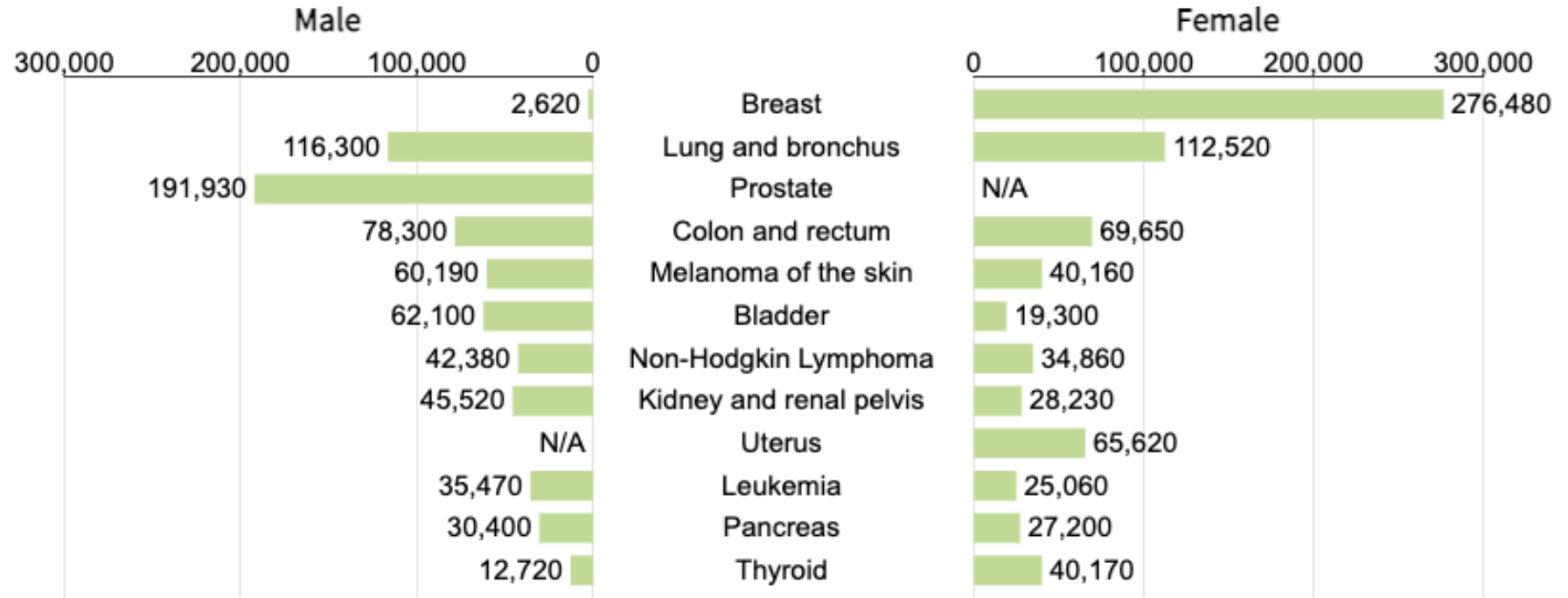
What two things do these people have in common?



What two things do these people have in common?



1 in 2 men and 1 in 3 women are at lifetime risk for developing cancer in their lifetime



Source: Cancer Facts & Figures 2020, American Cancer Society (ACS), Atlanta, Georgia, 2020.

Who gets cancer?

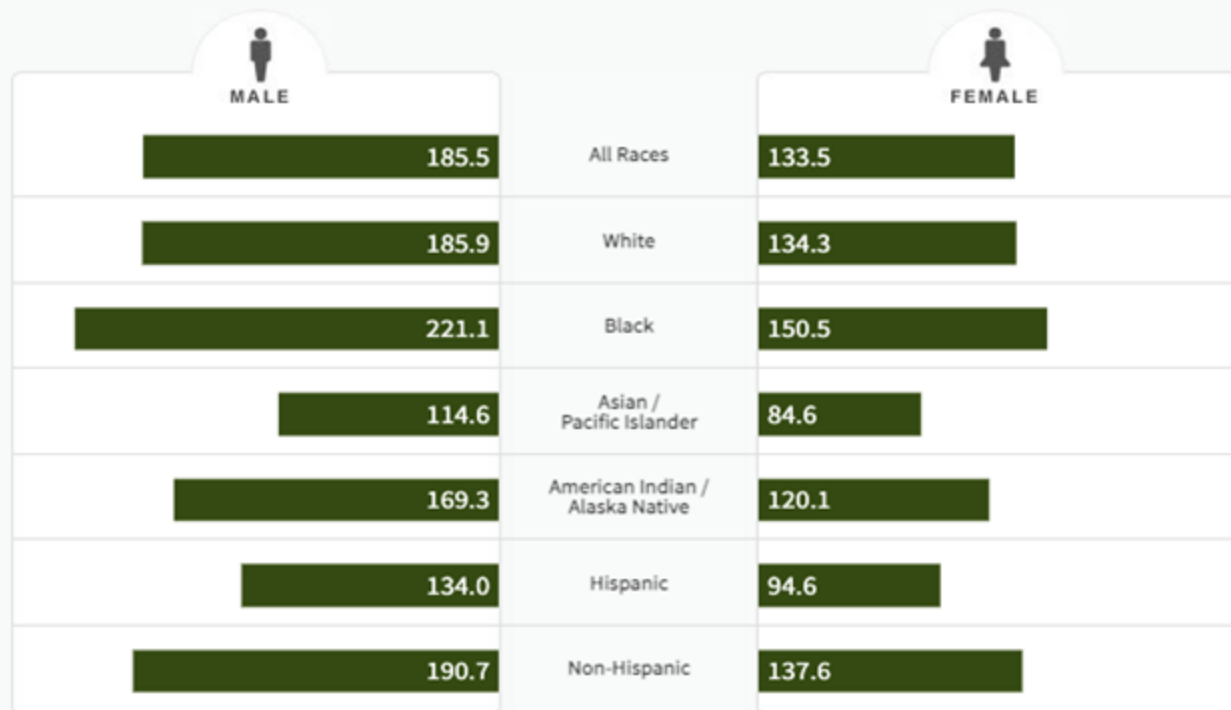
Rate of New Cases per 100,000 Persons by Race/Ethnicity & Sex: Cancer of Any Site



SEER 21 2013-2017, Age-Adjusted

Who dies from cancer?

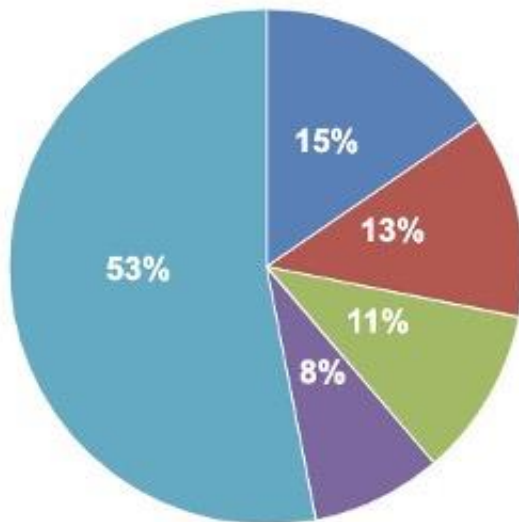
Death Rate per 100,000 Persons by Race/Ethnicity & Sex: Cancer of Any Site



U.S. 2014–2018, Age-Adjusted

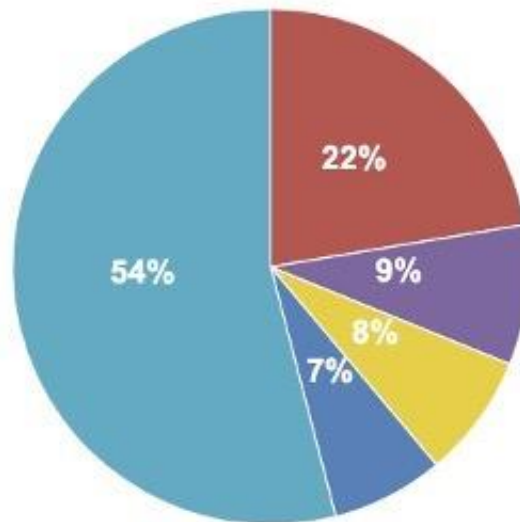
What are the most common cancers?

New Cancer Cases, 2020



- Breast: 279,100 (15%)
- Lung and bronchus: 228,820 (13%)
- Prostate: 191,930 (11%)
- Colon and rectum: 147,950 (8%)
- Other: 958,790 (53%)

Cancer Deaths, 2020



- Lung and bronchus: 135,720 (22%)
- Colon and rectum: 53,200 (9%)
- Pancreas: 47,050 (8%)
- Breast: 42,690 (7%)
- Other: 327,860 (54%)

What are the most common cancers?

In 2014, among men, Black/African American men had the highest lung cancer incidence and mortality rates in the United States.

- Smoking rates alone do not explain disparities in the development of the disease

Native Hawaiian people lung cancer rates surpass those of White people b/c of historically high smoking prevalence

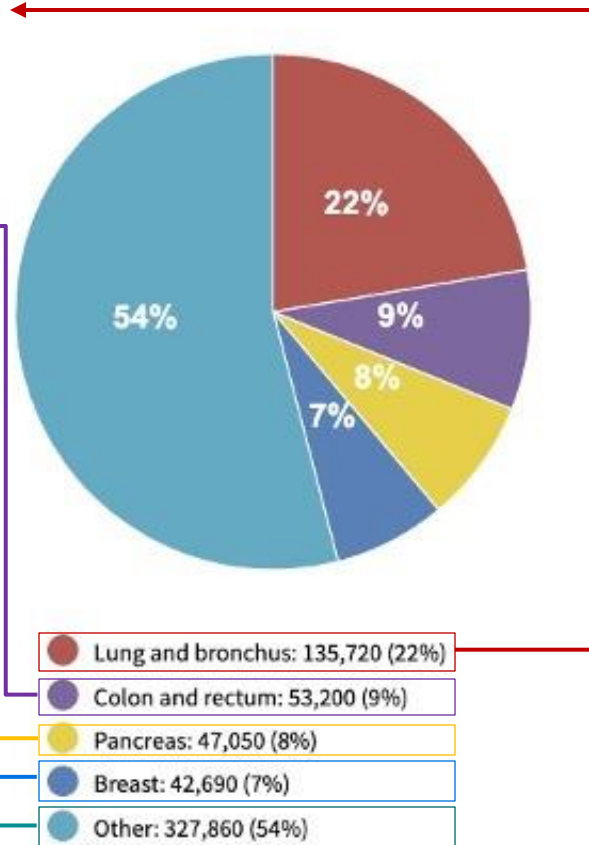
- 28% difference in incidence in the diagnosis of distant stage CRC between Black/African American and non-Hispanic whites.
- This difference accounted for more than 60% of the total mortality disparity from CRC.
- In addition, AA have an earlier median age at CRC diagnosis when compared to nHw

African Americans/Black patients (AA/B) have both higher incidence and mortality rates compared to NHW. Compared to NHW, AA/B typically with more aggressive disease, and stage-matched AA are less likely to receive systemic therapy, surgery, chemotherapy or radiotherapy and more likely to refuse tx

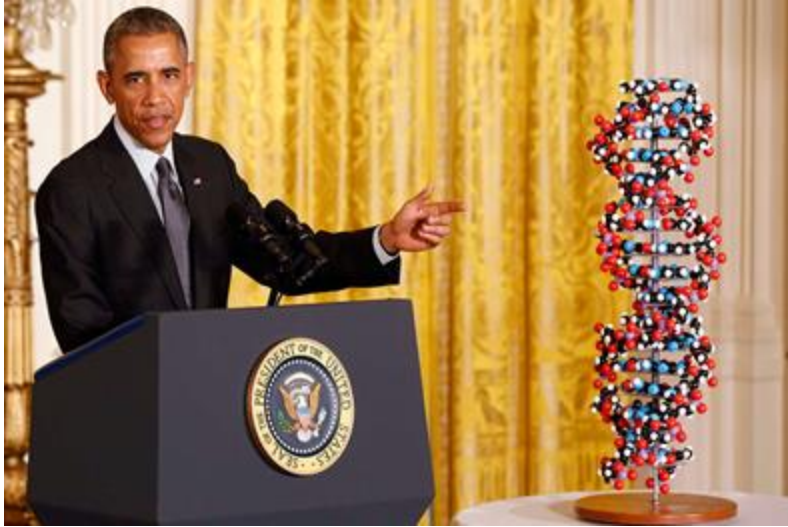
The racial disparity in deaths from breast cancer has remained at 40% or higher for a decade. Black women younger than age 50 had a death rate that was twice as high as White women that age. Plus, Black women are more likely than White women to die of breast cancer at any age.

Cervical cancer incidence is >30% higher in Hispanic women vs. white women. Liver and stomach cancer rates in Hispanic people are 2X those in White people. Prostate cancer though lower in Hispanic men is 44% higher in men living in Puerto Rico

Cancer Deaths, 2020

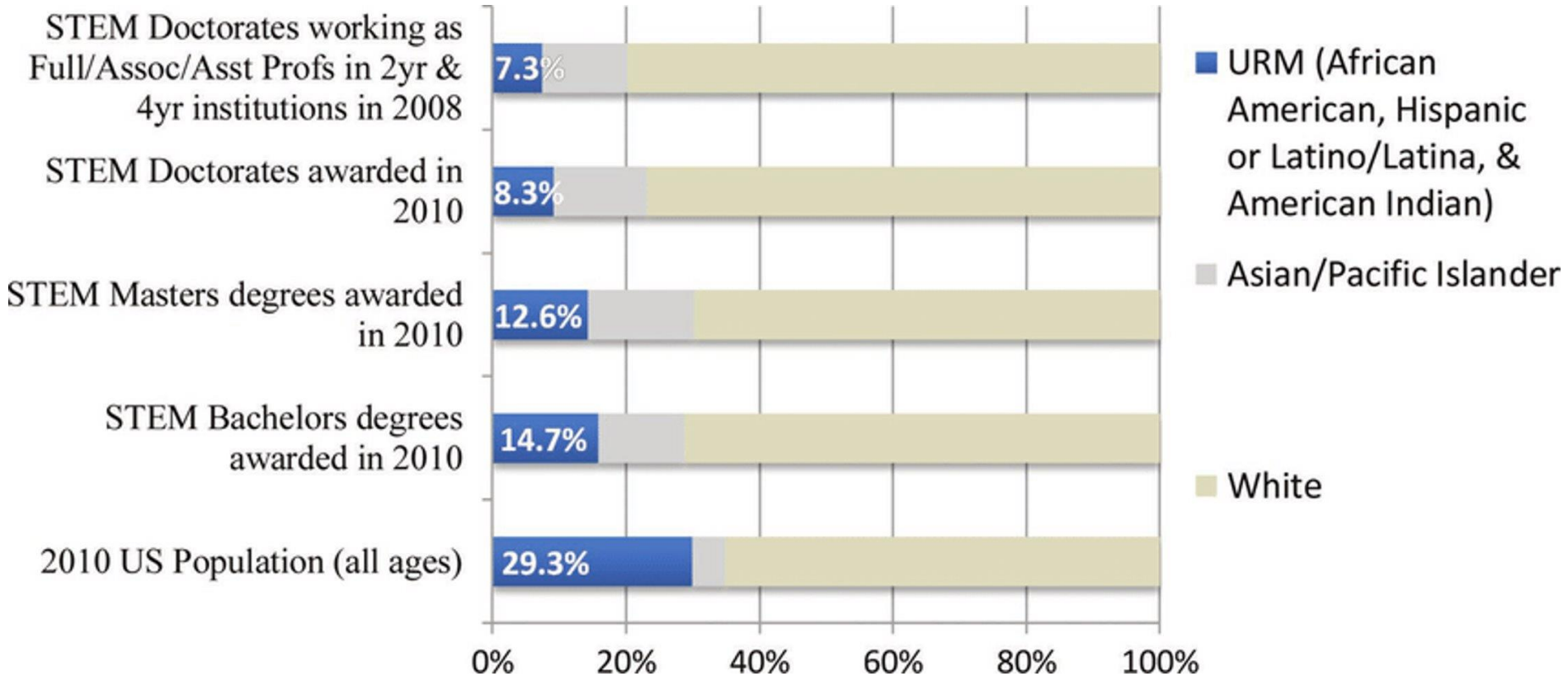


Precision Medicine “Moonshot” Initiative



“scale up efforts to identify genomic drivers in cancer and apply that knowledge in the development of more effective approaches to cancer treatment.”

And yet we're losing you...



Biology Fundamentals



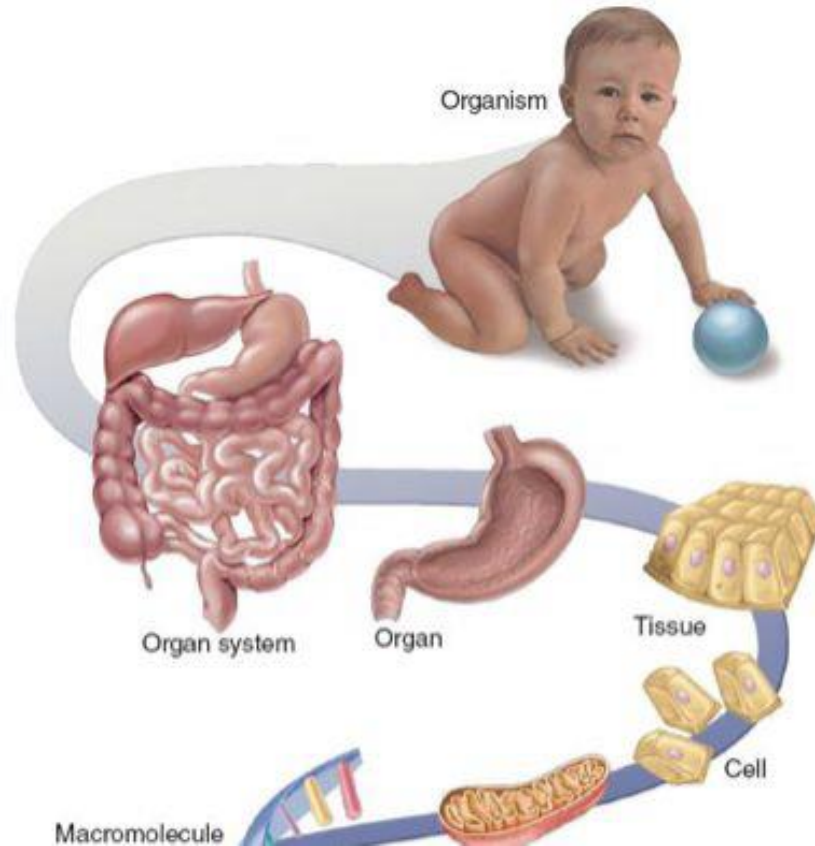
Today's Lecture Outline

- **The cell**
 - From cells to organisms
- The central dogma
- **The human genome**
 - DNA
 - Gene expression
 - Transcription
 - RNA
 - Translation
- **Mutations: Overview**
 - Oncogenes and tumor suppressor genes
 - Gene mutations
 - Chromosomal mutations

DNA → RNA → protein → organism

Blueprint in hieroglyphics → blueprint in English → construction workers scaffolding etc → building with rooms and occupants

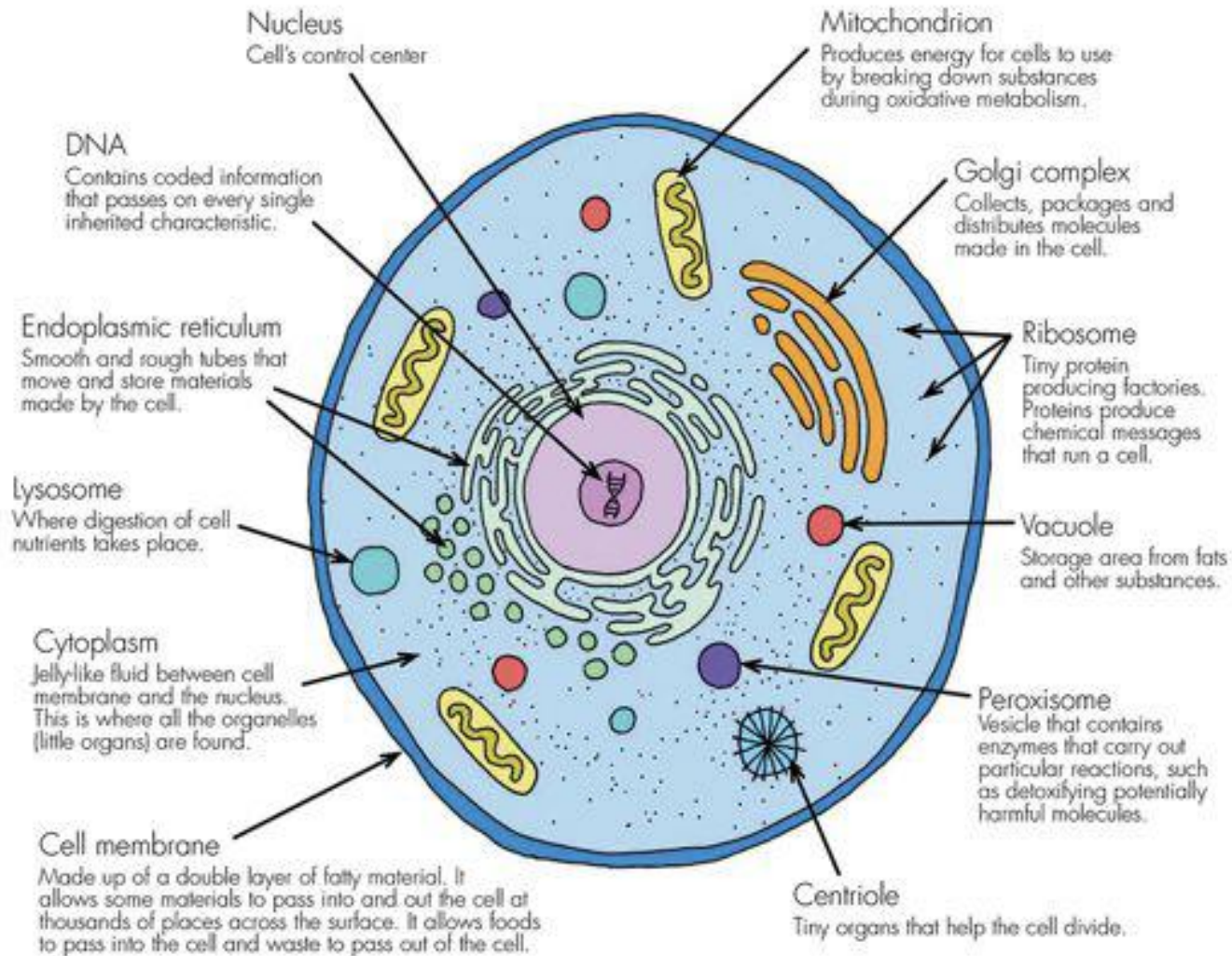
From Cells to Organisms



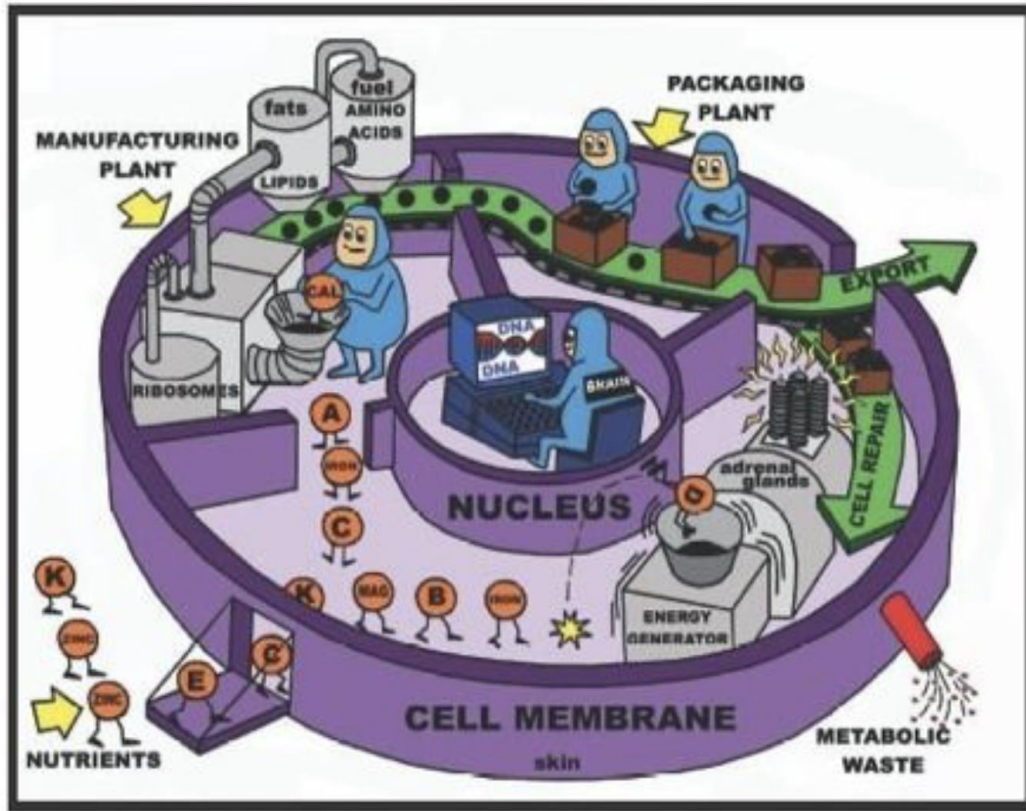
The Cell



Cells are the basic building blocks of life



Cells are like mini-factories



Nucleus = control center

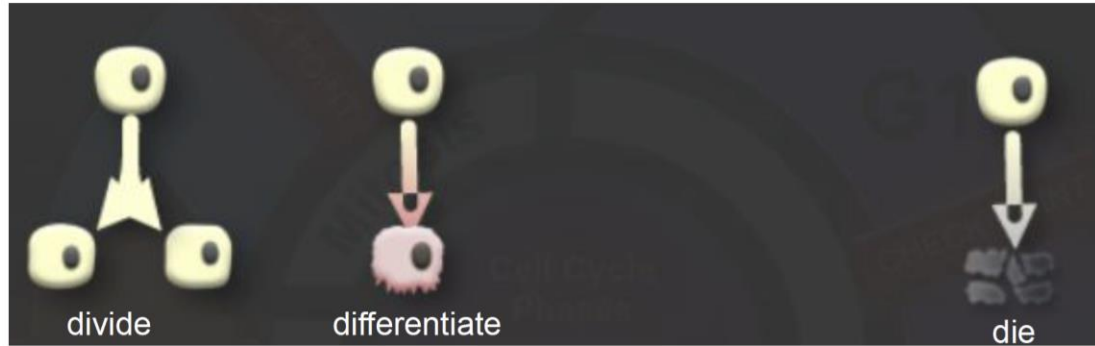
DNA = blueprint

Cell Membrane = shipping and receiving

Cytoplasm = factory floor

Proteins = workers

Cells can divide, differentiate or die



Cells divide to produce two identical daughter cells.

Cells stop dividing to specialize in structure and function, a process called **differentiation**.

Cells can also undergo programmed cell death, or **apoptosis**, a process that eliminates unnecessary cells during development and removes unhealthy or damaged cells in the mature organism.

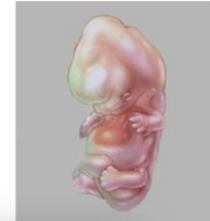
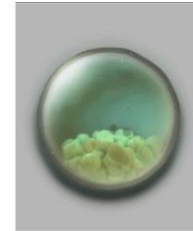
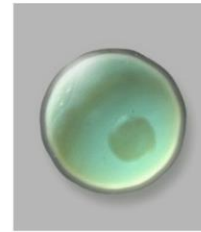
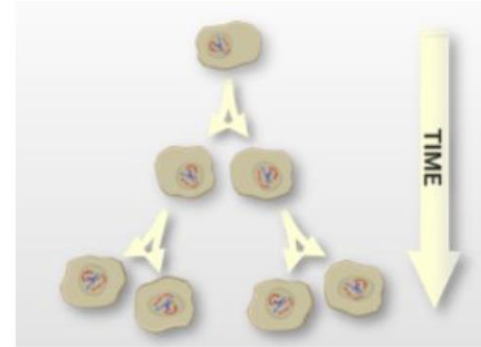
**Cell Division
(birth/reproduction)**



Cell Division

Cells division is essential to life

- Multicellular organisms need **cell division** to grow and to replace dead or damaged cells.
- In humans, many types of cells divide.
 - For example, repeated divisions allow a single fertilized egg cell to develop into an adult with more than 37 trillion cells.
- After growth, division remains important in normal cell turnover, such as in our skin and gut, where cells are continuously renewed. Other cells have to divide to heal wounds like skin cuts or broken bones.



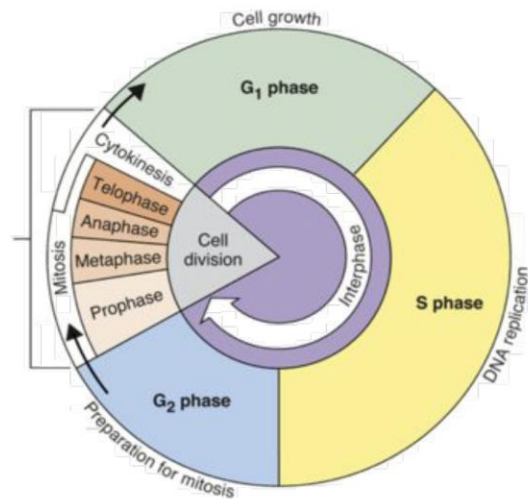
Cell Division and the Cell Cycle

The cell cycle



- Each time the cell divides into two, it goes through a sequence of events that includes
 - Growth
 - DNA replication
 - preparation to divide
 - and cell division or **mitosis**
- This is called the **cell cycle**

- *To remain healthy, it's critical for an organism to maintain the right number of cells.*
 - This is achieved primarily by regulating the cell cycle
 - **Cell cycle regulators** are molecular signals that may stimulate or halt cell division, instruct cells to differentiate, or initiate cell death.



onlinesciencenotes.com/cell-cycle-its-different-phases-and-duration/

In cancer, cell cycle regulators do not function properly leading to too many cells.

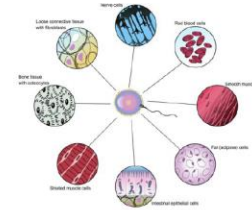
True or False

Cell Differentiation (get a job)






Cell Differentiation due to different gene expression

All cells have the same DNA, but different cells express different genes



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Cell type	Red blood	Muscle	Pancreatic
			
Gene type			
Housekeeping	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Hemoglobin	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Insulin	<input type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>
Myosin	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>

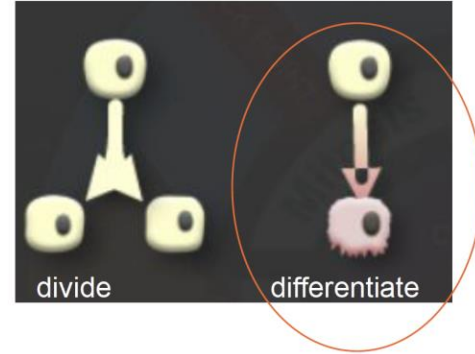
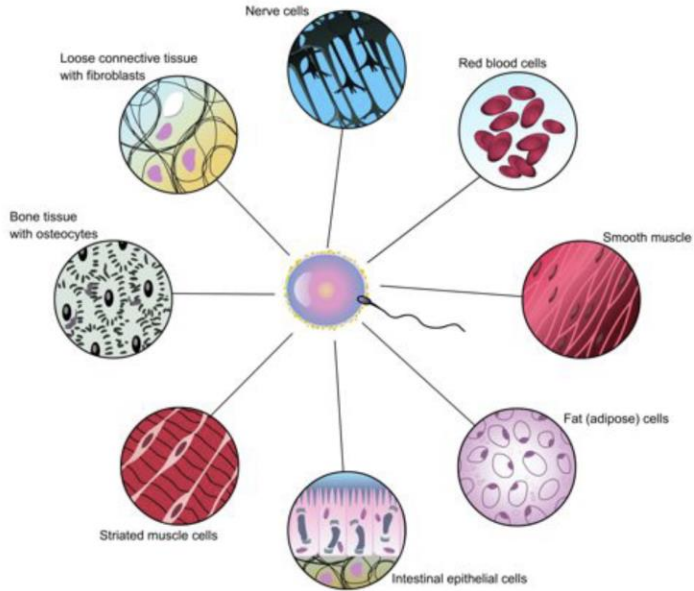
All the cells in a person's body have the **same DNA** and the same genes

However, the difference between cells in different tissues and organs is that the **"expression" of the genes** differs between cells

1. Expression means that the message from the **DNA is being copied and made into protein**

For example, liver cells express different proteins than skin cells, even though their DNA is the same

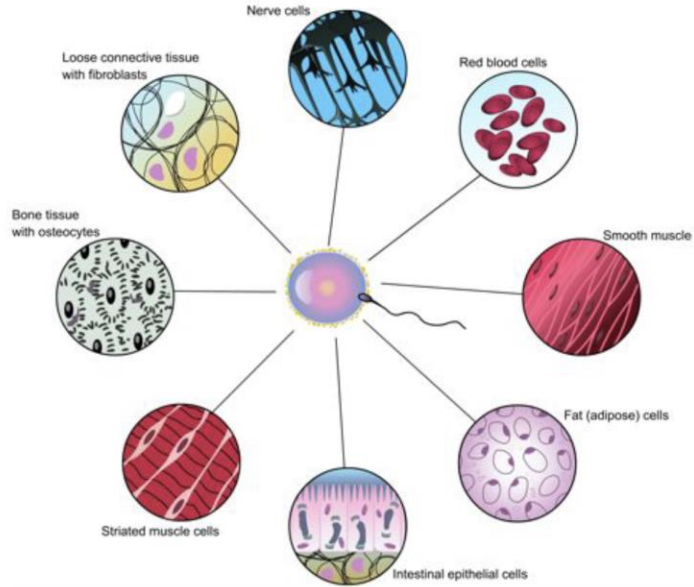
Cell Differentiation



Cells stop dividing to specialize in structure and function

Isn't it amazing that your body with all of its specialized organs developed from a single cell?

Cell Differentiation



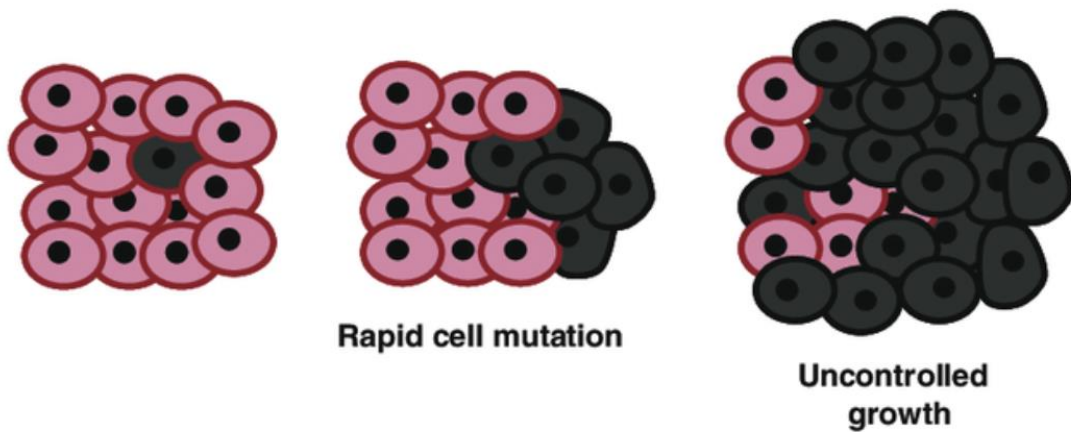
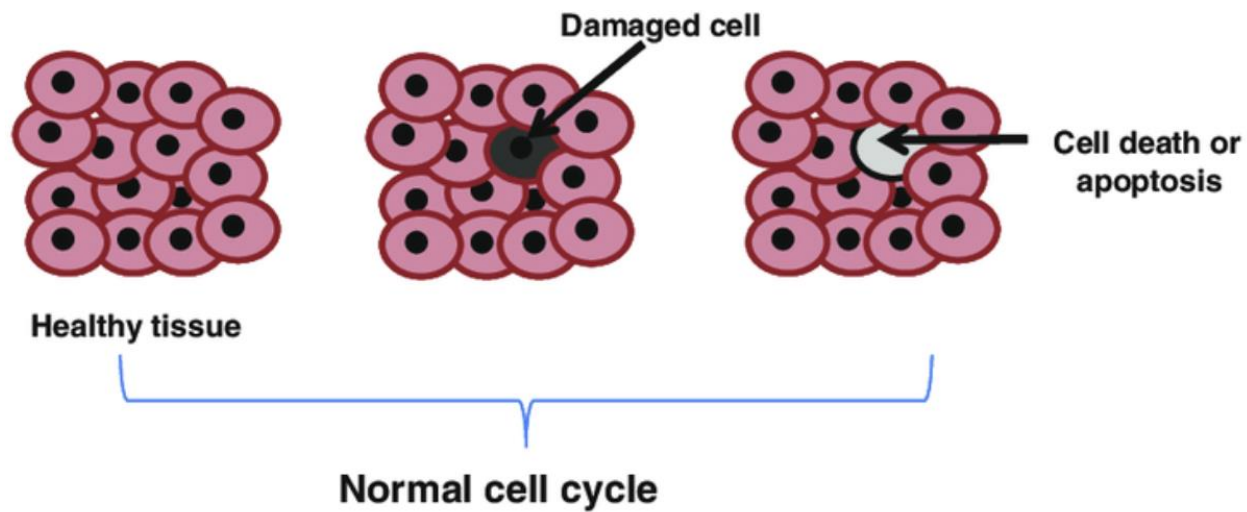
How do cells become specialized

(ie. how does a lung cell become a lung cell and a blood cell become a blood cell)?

Cell Death

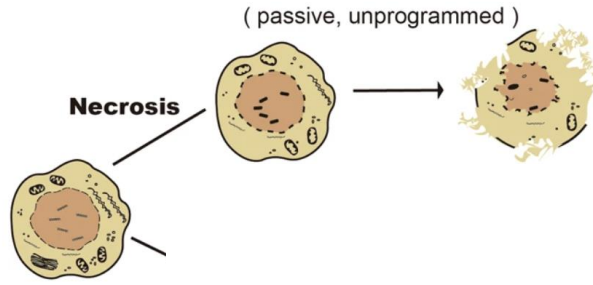
(die and leave a mess or have everything organized)





Cell Death in Normal Cells versus that in Gastric Cancer

Cell death

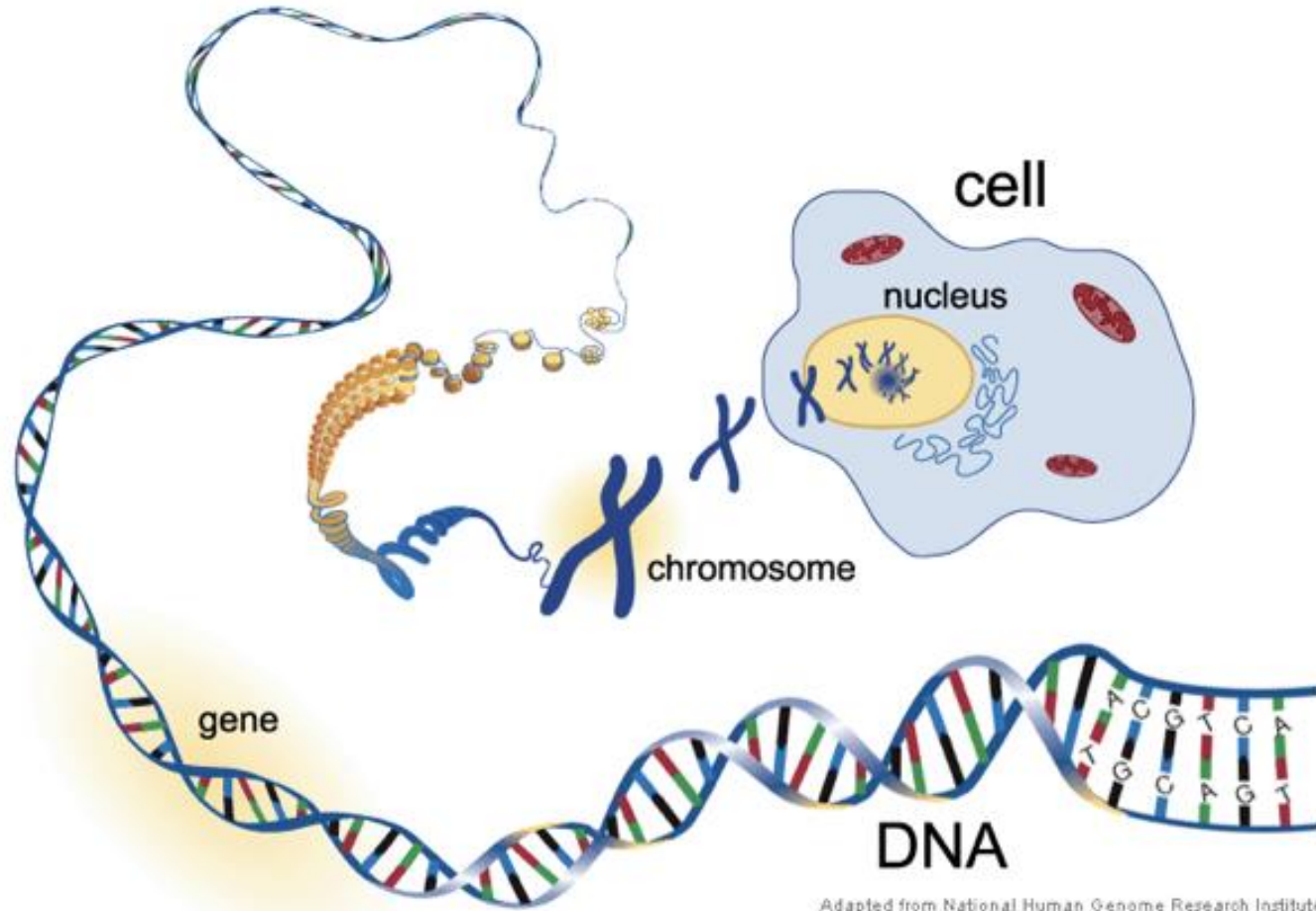


Left: The classification of cell death. Middle: Cells that undergo death in gastric cancer. Right: Effects triggered by different cell death in gastric cancer.

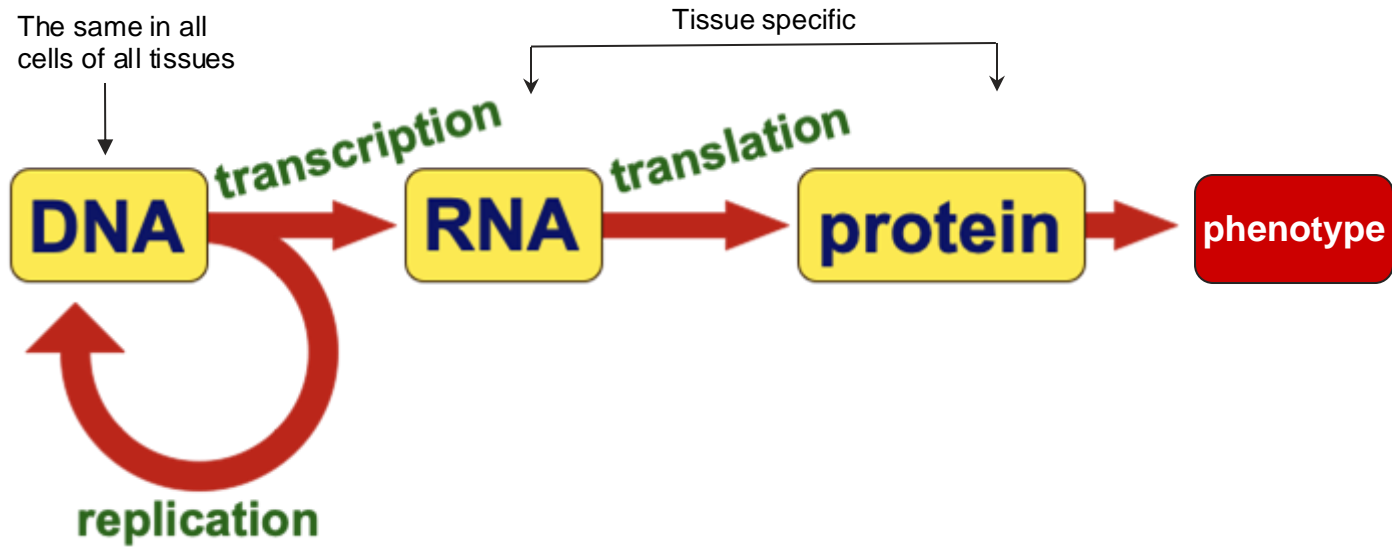
The Central Dogma
DNA → RNA → Protein



Independent of the cell type -- all cells have DNA

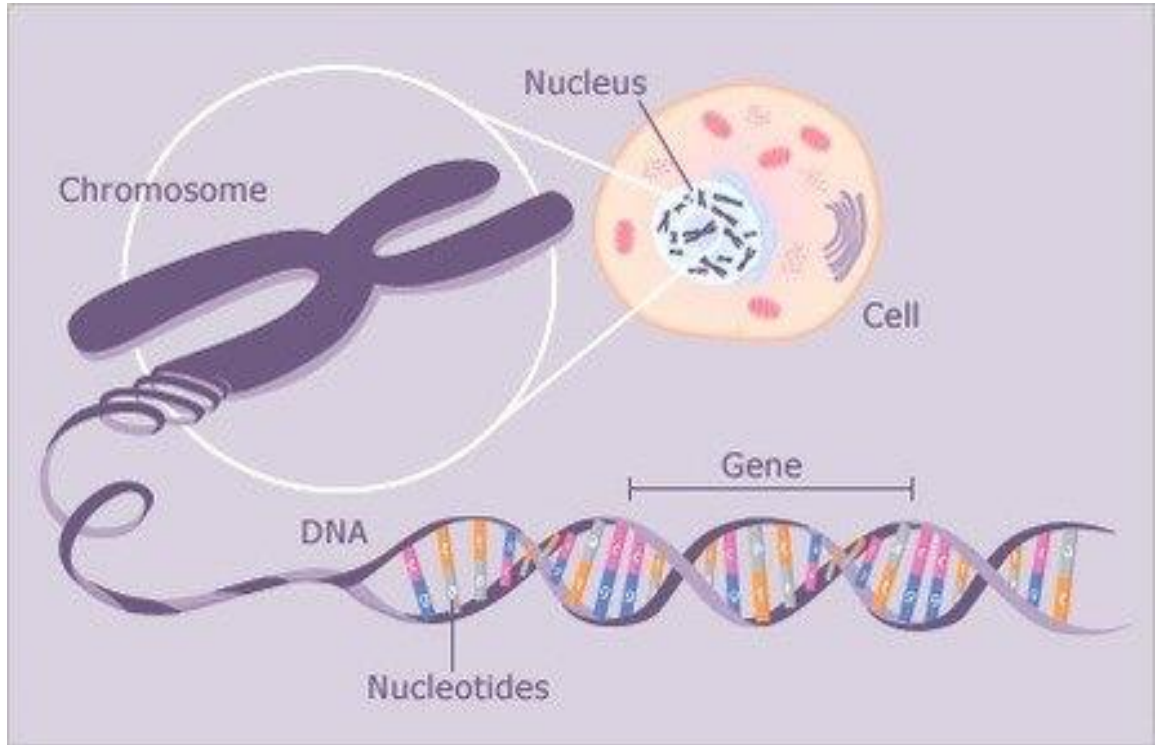


The Central Dogma




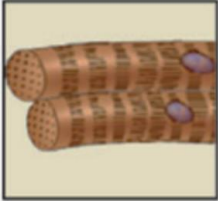
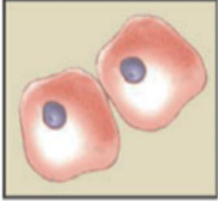












DNA and genes

- Every cell in your body has the same DNA (*consider DNA as a very thick hand-written book with multiple chapters*)
- Different genes (*words in the book*) are used in different cell types (*in the different chapters*)
- But each nucleotide in the DNA of each gene with varying base-pairs (*each letter in each word*) needs to be transcribed into mRNA (*the selected hand-written chapter needs to typed*)
- Then the mRNA needs to be translated into a functional protein (*the selected typed chapter becomes an episode of a TV show*)



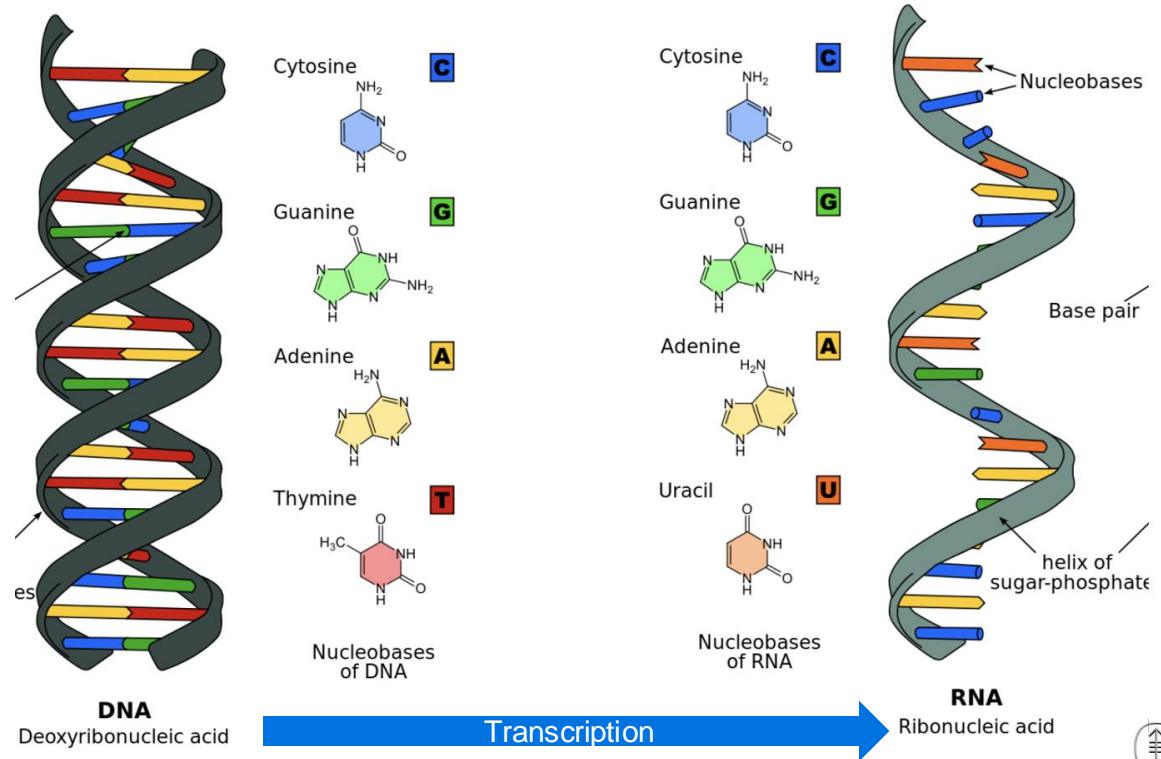
mRNA

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Cell type	Red blood	Muscle	Pancreatic X
			
Gene type			
Housekeeping			
Hemoglobin			
Insulin			
Myosin			

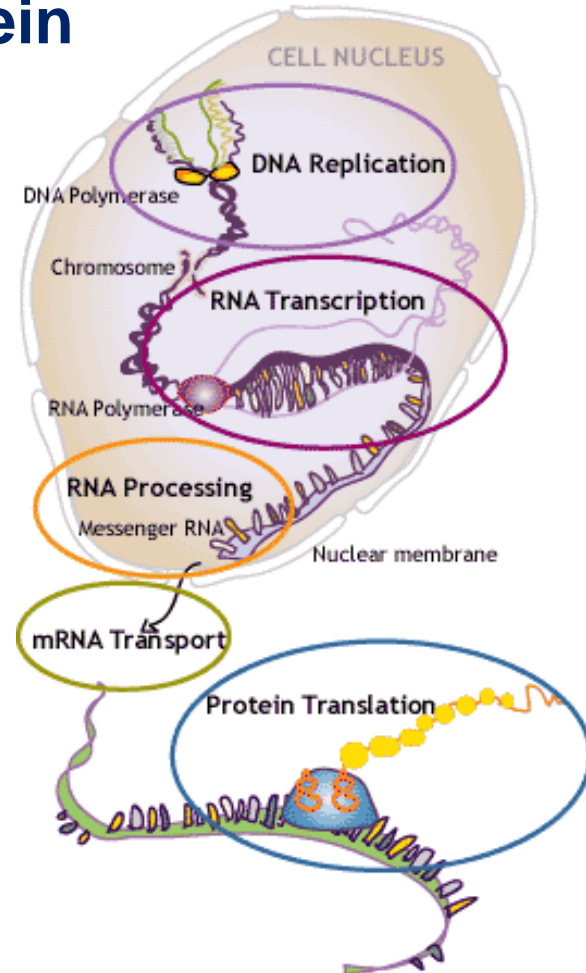
DNA – Transcription → mRNA

- Every cell in your body has the same DNA (*consider DNA as a very thick hand-written book with multiple chapters*)
- Different genes (*words in the book*) are used in different cell types (*in the different chapters*)
- But each nucleotide in the DNA of each gene with varying base-pairs (*each letter in each word*) needs to be transcribed into mRNA (*the selected hand-written chapter needs to typed*)
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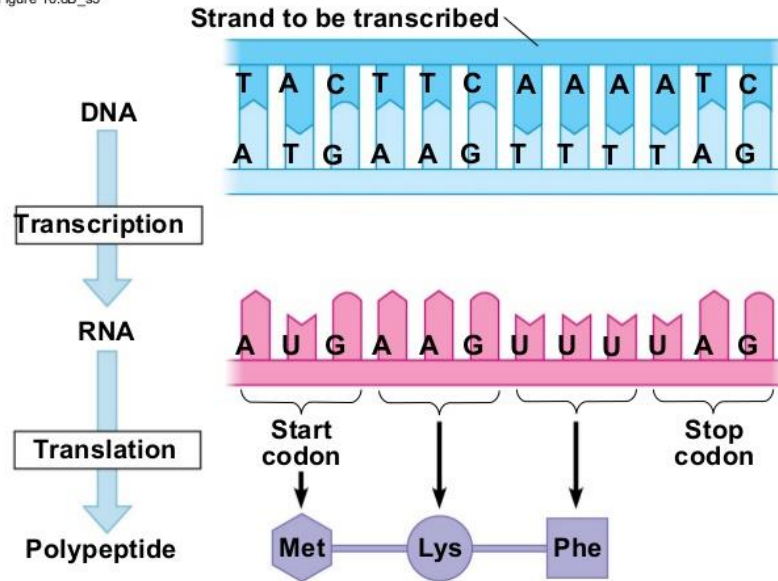
DNA-RNA – Translation → Protein

- Every cell in your body has the same DNA (*consider DNA as a very thick hand-written book with multiple chapters*)
- Different genes (*words in the book*) are used in different cell types (*in the different chapters*)
- But each nucleotide in the DNA of each gene with varying base-pairs (*each letter in each word*) needs to be transcribed into mRNA (*the selected hand-written chapter needs to typed*)
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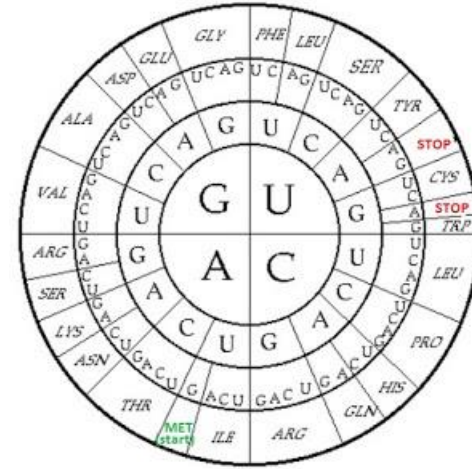
Protein Coding

Figure 10.8B_s3



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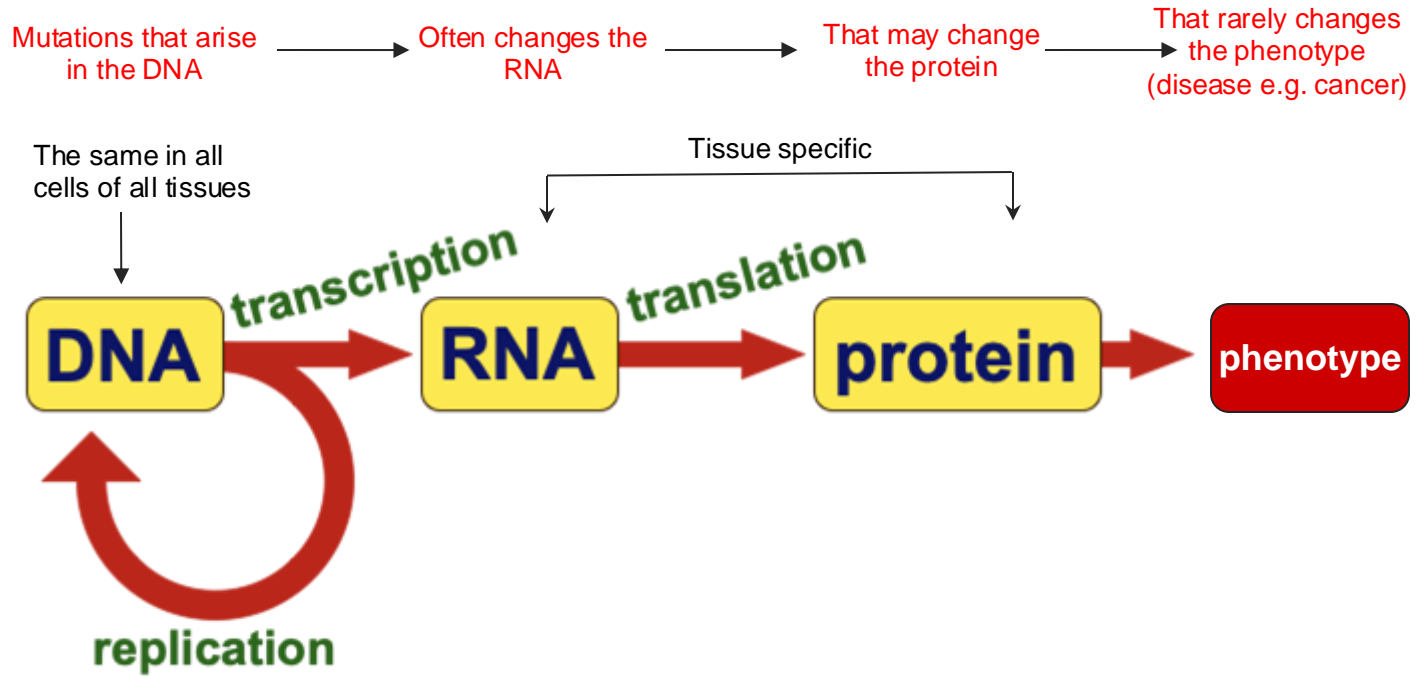
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<http://lab2webchemistry.blogspot.com/2015/08/protein-manufacturing-coding-in-our-body.html>

What is Cancer?



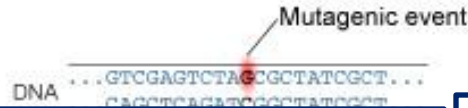


Cancer is a genetic disease meaning changes in the DNA MAY cause cancer

Types of Mutation

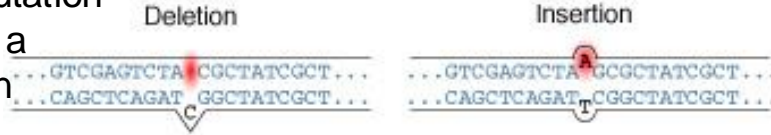
Micro

Point mutations



Silent mutation is when a base pair change in the DNA doesn't result in a change in the protein

Frameshift mutation that results in a mutant protein



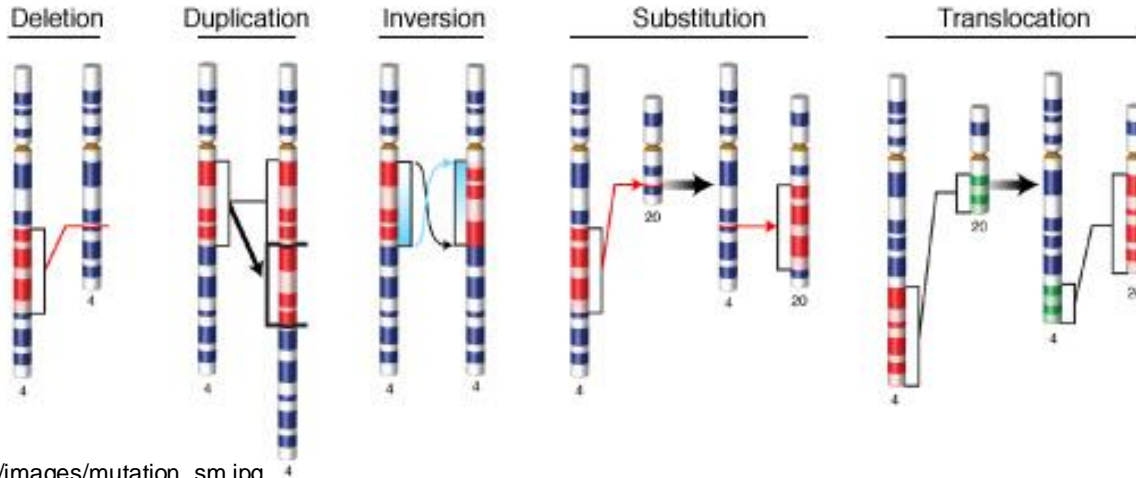
Substitution



Missense mutation that results in a mutant protein

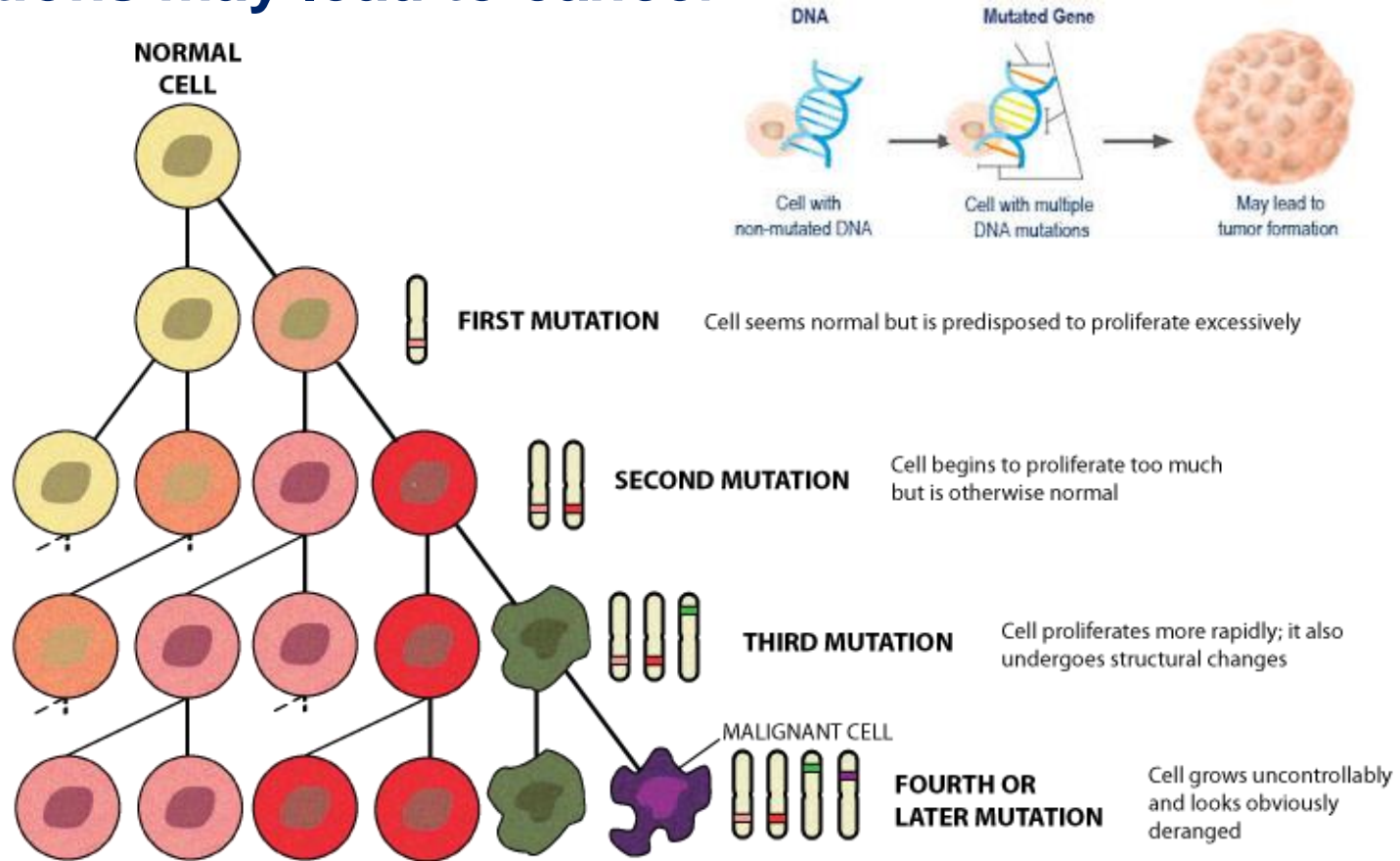
Macro

Chromosomal mutations



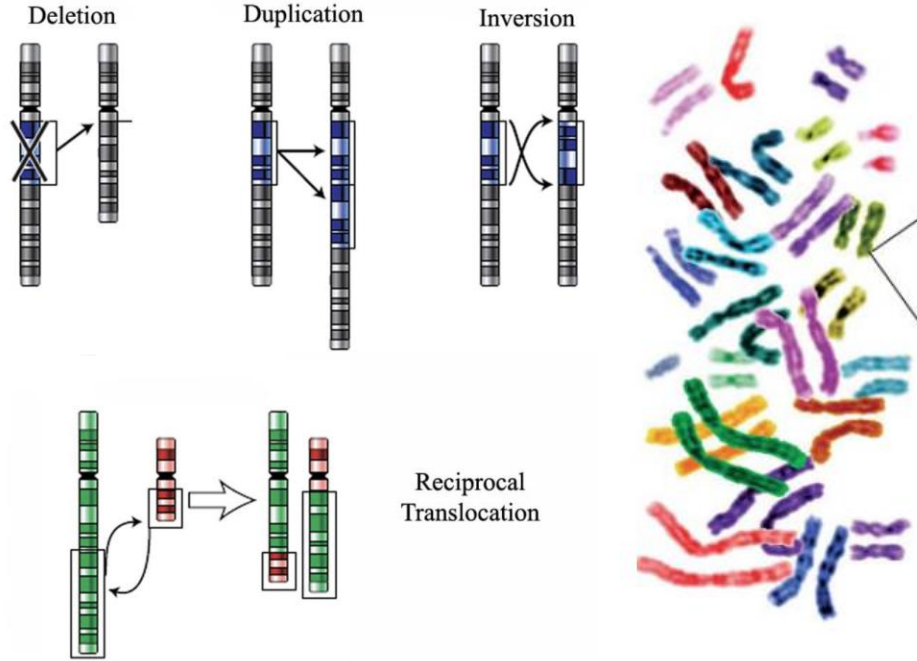
Nonsense mutations A missense mutation which changes the codon that recruits an amino acid to the mRNA into a stop codon

Mutations may lead to cancer

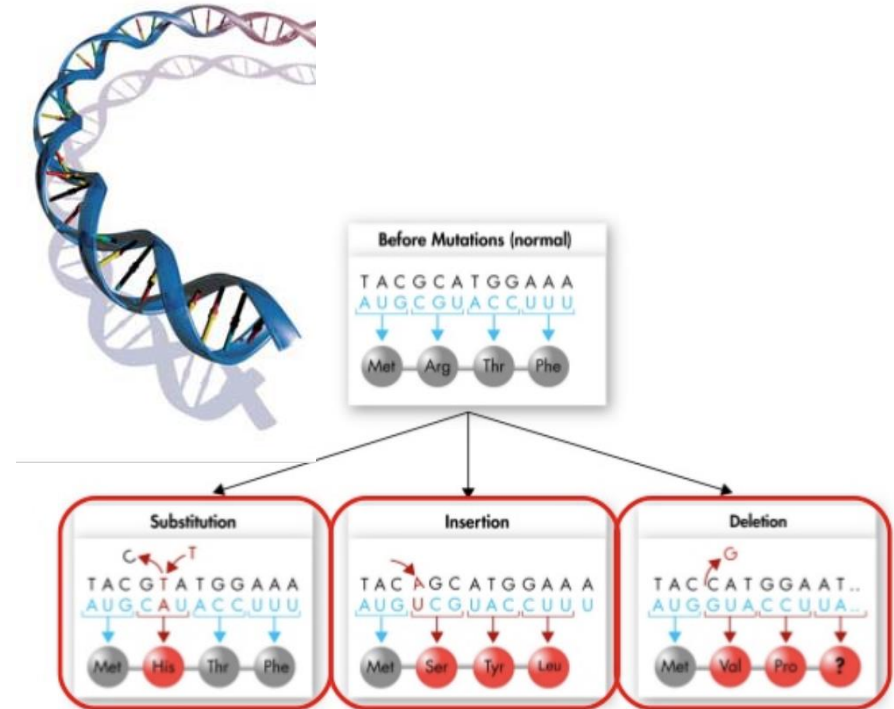


Types of Mutations

Chromosomal

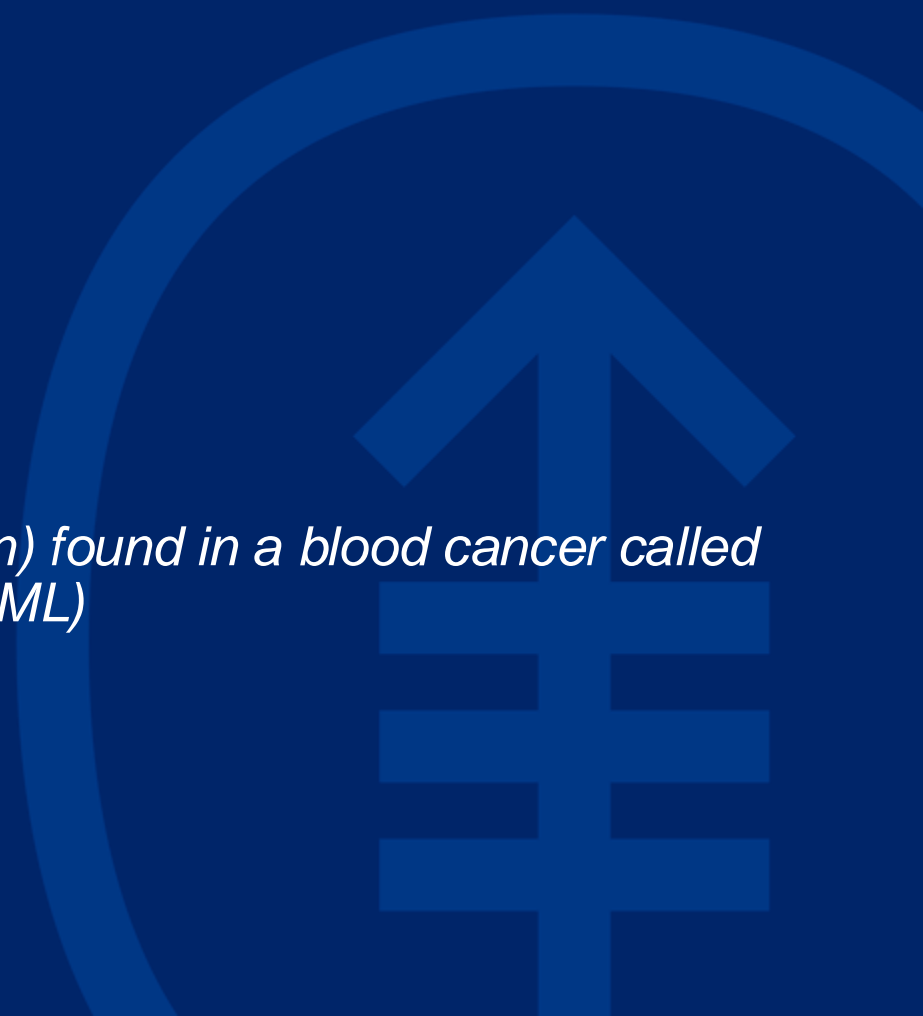


Point Mutations



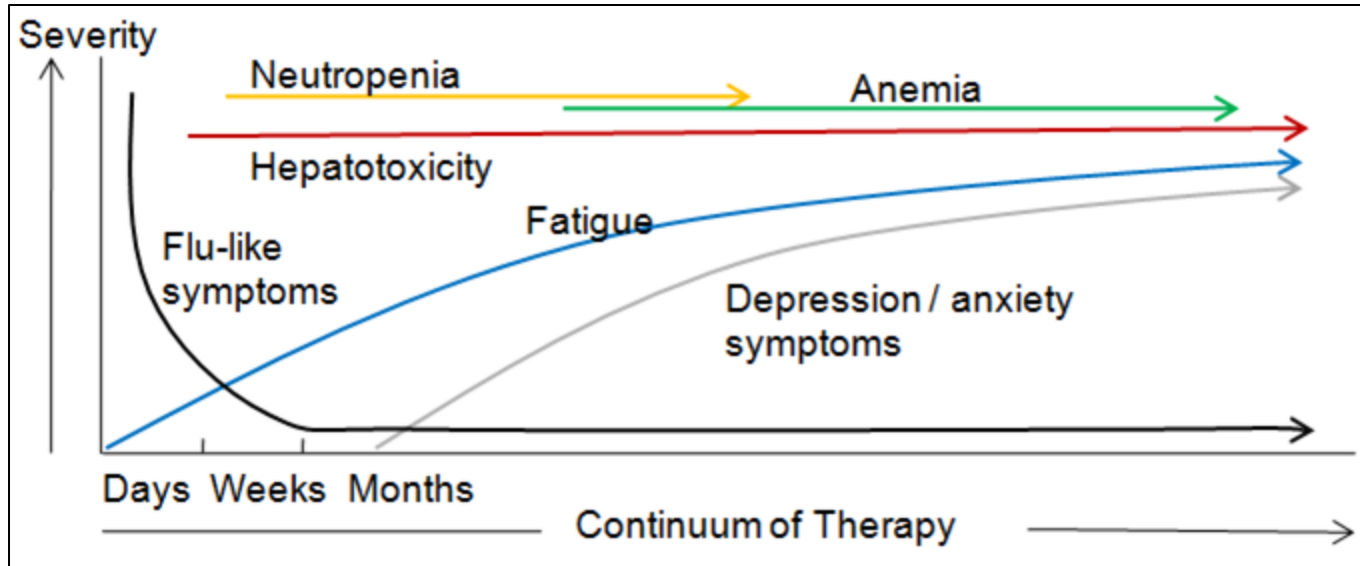
BCR-ABL1

a translocation (chromosomal mutation) found in a blood cancer called “Chronic Myelogenous Leukemia” or CML)



Before Targeted Therapies...

- Prior to 2001: <1 in 3 CML patients survived 5 years past diagnosis
- The most successful intervention at the time was Interferon a



Chromosomes 9 and 22 in blood cells of a patient with Chronic Myelogenous Leukemia

A Minute Chromosome in Human Chronic Granulocytic Leukemia

In seven cases thus far investigated (five males, two females), a minute chromosome has been observed replacing one of the four smallest autosomes in the chromosome complement of cells of chronic granulocytic leukemia cultured from peripheral blood. No abnormality was observed in the cells of four cases of acute granulocytic leukemia in adults or of six cases of acute leukemia in children. There have been several recent reports of chromosome abnormalities in a number of cases of human leukemia [including two of the seven cases reported here: Nowell and Hungerford, *J. Natl. Cancer Inst.*, **25**, 85 (1960)], but no series has appeared in which there was a consistent change typical of a particular type of leukemia.

Cells of the five new cases were obtained from peripheral blood (and bone marrow in one instance), grown in culture for 24-72 hours, and processed for cytological examination by a recently developed air-drying technique (Moorhead, *et al.*, *Exptl. Cell Research*, in press). The patients varied from asymptomatic untreated cases to extensively treated

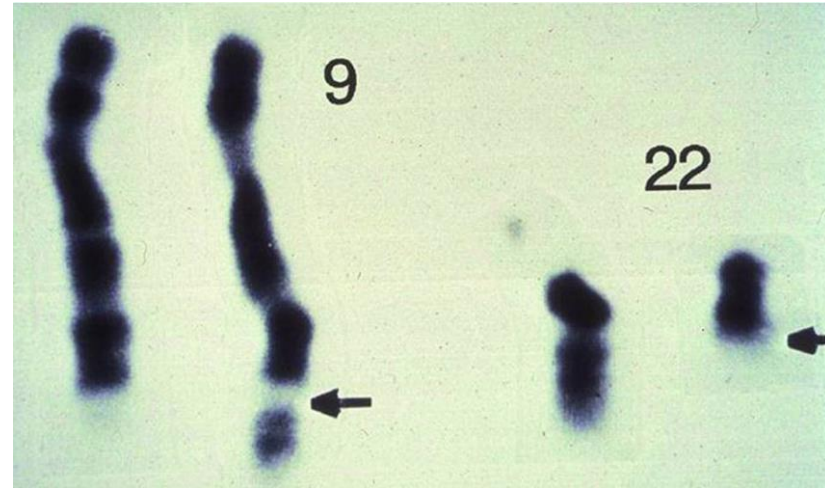
cases of several years duration in terminal myeloblastic crisis. All seven individuals showed a similar minute chromosome, and none showed any other frequent or regular chromosome change. In most of the cases, cells with normal chromosomes were also observed. Thus, the minute is not a part of the normal chromosome constitution of such individuals.

The findings suggest a causal relationship between the chromosome abnormality observed and chronic granulocytic leukemia.

PETER C. NOWELL

*School of Medicine,
University of Pennsylvania*

DAVID A. HUNGERFORD
Institute for Cancer Research



Poster Child for Targeted Therapy: Gleevec

A BCR-ABL1 in CML and ALL

BCR-ABL1 protein

BCR-ABL1 mRNA

Activation of downstream pathways

t(9;22)

TIME

THERE IS NEW AMMUNITION IN THE WAR AGAINST CANCER. THESE ARE THE BULLETS.

Revolutionary new pills like GLEEVEC combat cancer by targeting only the diseased cells. Is this the breakthrough we've been waiting for?

9 22

Gleevec™ (imatinib mesylate) Capsules 100 mg

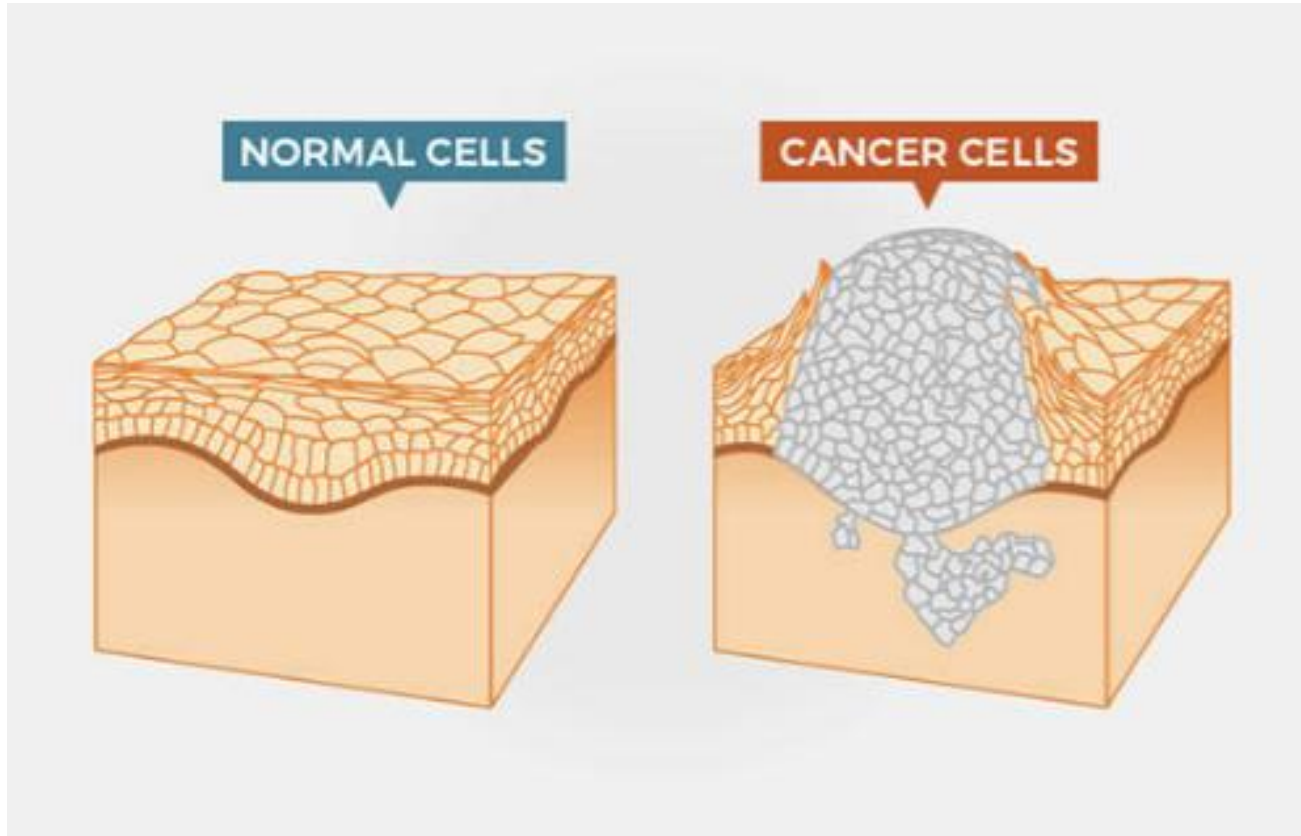
Each capsule contains 100 mg of imatinib mesylate.

120 Capsules Rx only

Cancer Fundamentals



How does cancer form?



Cancer is a disease caused when cells divide uncontrollably and spread into surrounding tissues.

“What Is Cancer?” was originally published by the National Cancer Institute.”

Cancer is a collection of different diseases

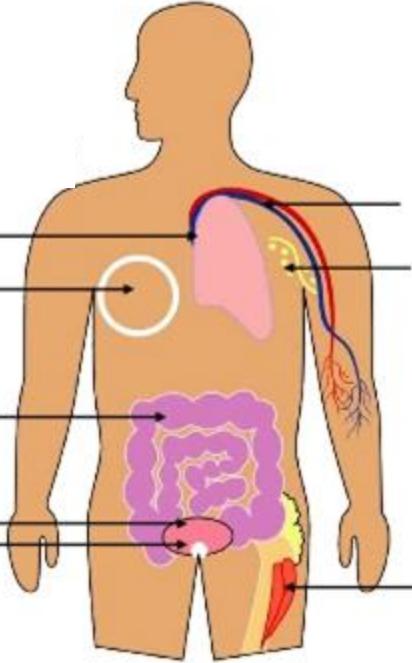
Carcinomas (cells that cover internal and external body surfaces)

Lung

Breast

Colon

Bladder
Prostate
(Men)



Leukemia
(Blood Cells)

Lymphomas
(Lymph nodes & tissues)

Sarcomas
(Cells in supportive tissues – Bone, Fat, Muscles)

Naming cancer

Cancer Prefixes Point to Location

<i>Prefix</i>	<i>Meaning</i>
---------------	----------------

adeno-	gland
---------------	--------------

chondro-	cartilage
-----------------	------------------

erythro-	red blood cell
-----------------	-----------------------

hemangio-	blood vessels
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hepato-	liver
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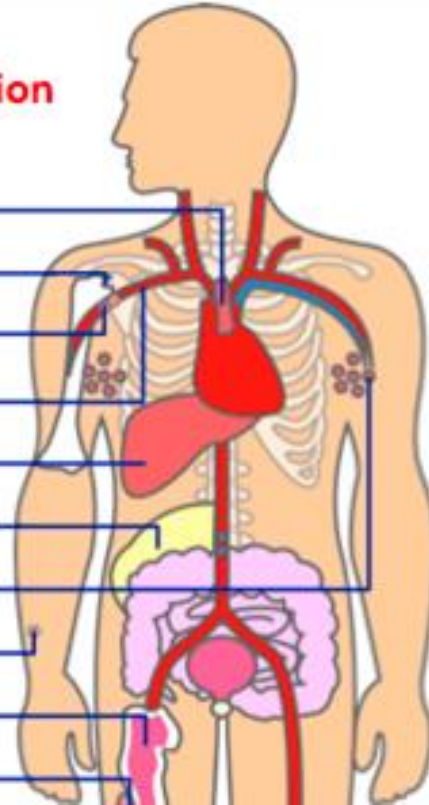
lipo-	fat
--------------	------------

lympho-	lymphocyte
----------------	-------------------

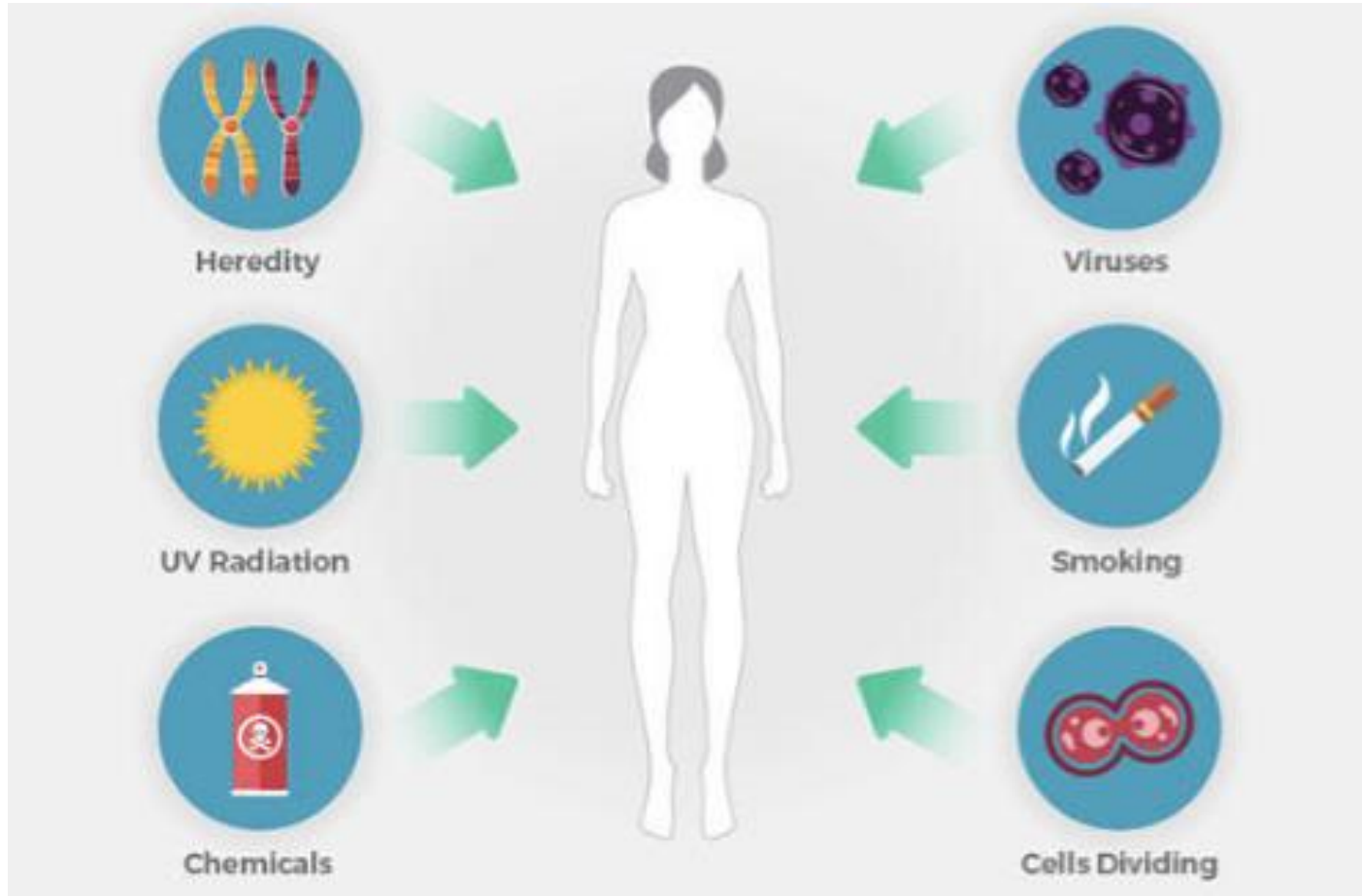
melano-	pigment cell
----------------	---------------------

myelo-	bone marrow
---------------	--------------------

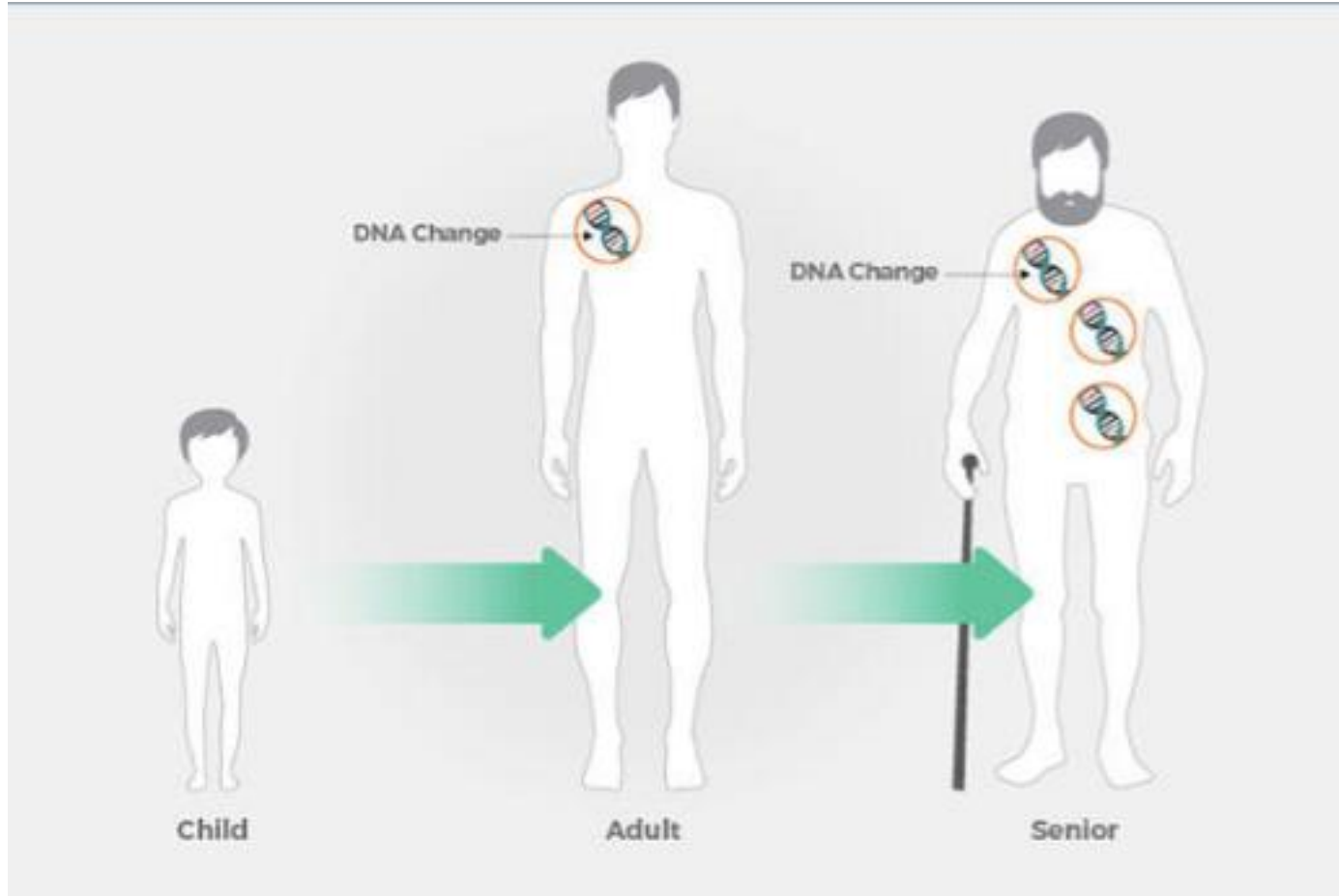
myo-	muscle
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What causes genetic changes in cancer?

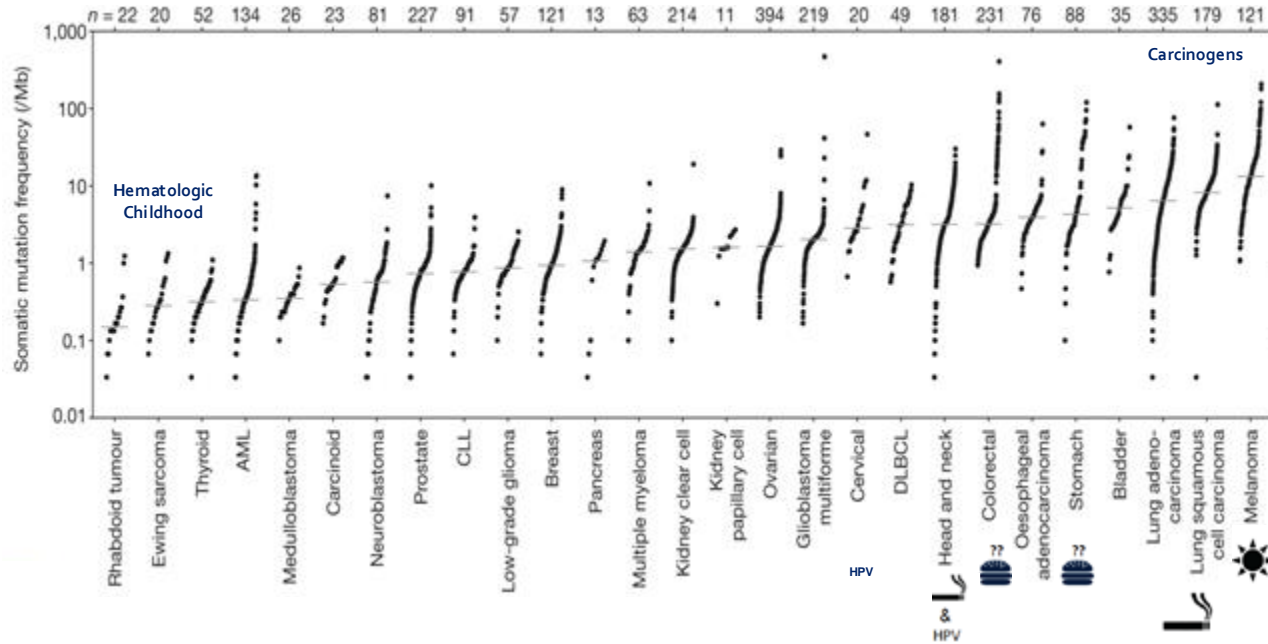


How does age relate to cancer?

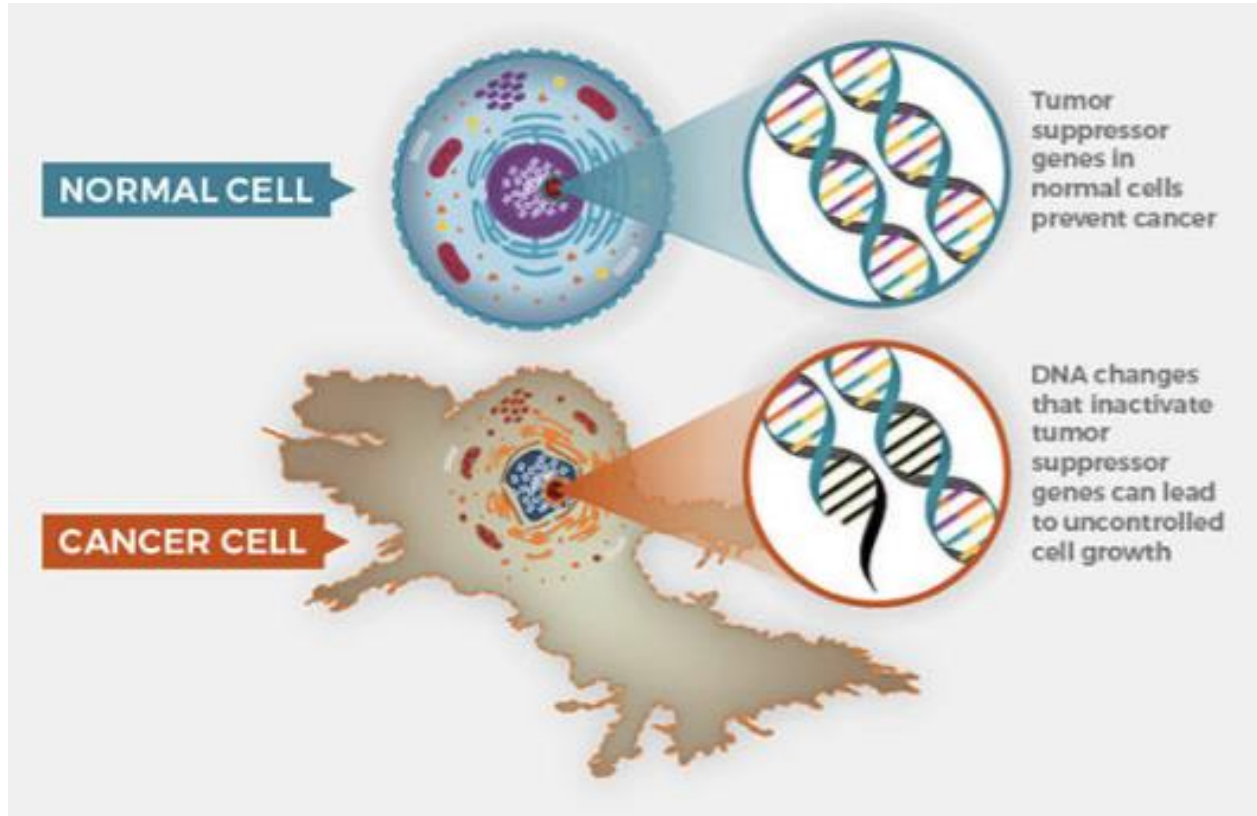


Cancer is Genetic

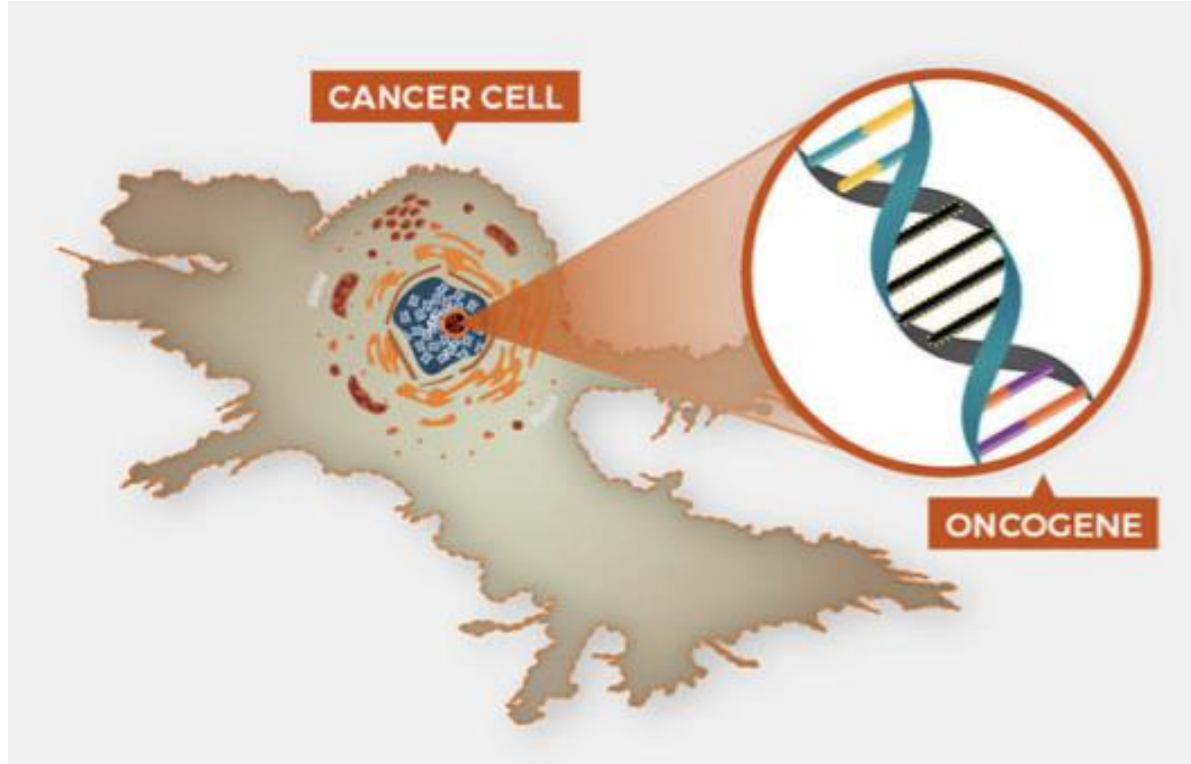
Most mutations in cancer are “passengers”



Some mutations arise in Tumor Suppressor Genes == DRIVERS



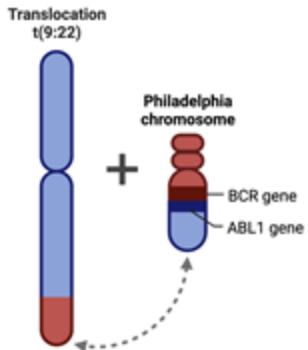
Some mutations arise in Oncogenes == DRIVERS



Passengers

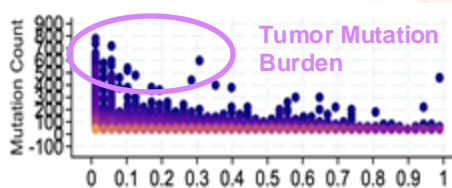
Alterations which do not alter cancer cell fitness

Translocations



MSI Status

5'—CACACACACACACA—3'
5'—CACACACA**C**ACACACA—3'



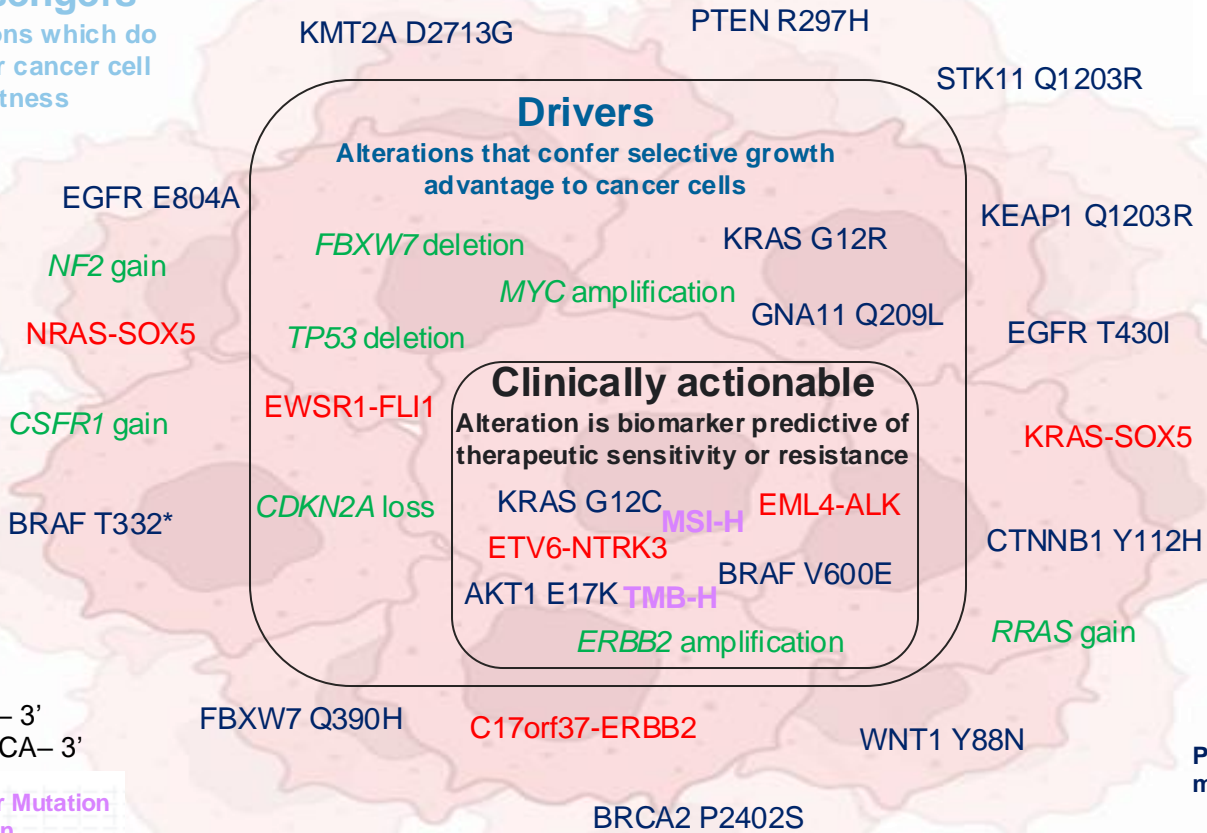
CA

Drivers

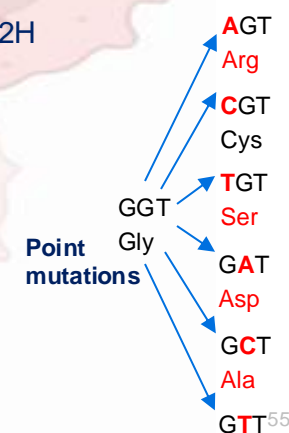
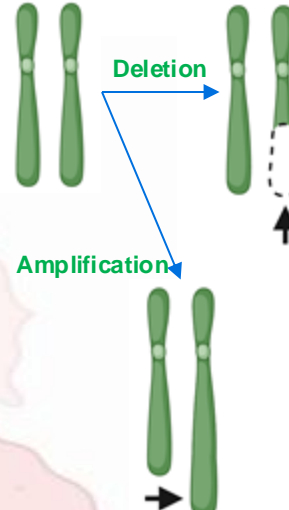
Alterations that confer selective growth advantage to cancer cells

Clinically actionable

Alteration is biomarker predictive of therapeutic sensitivity or resistance

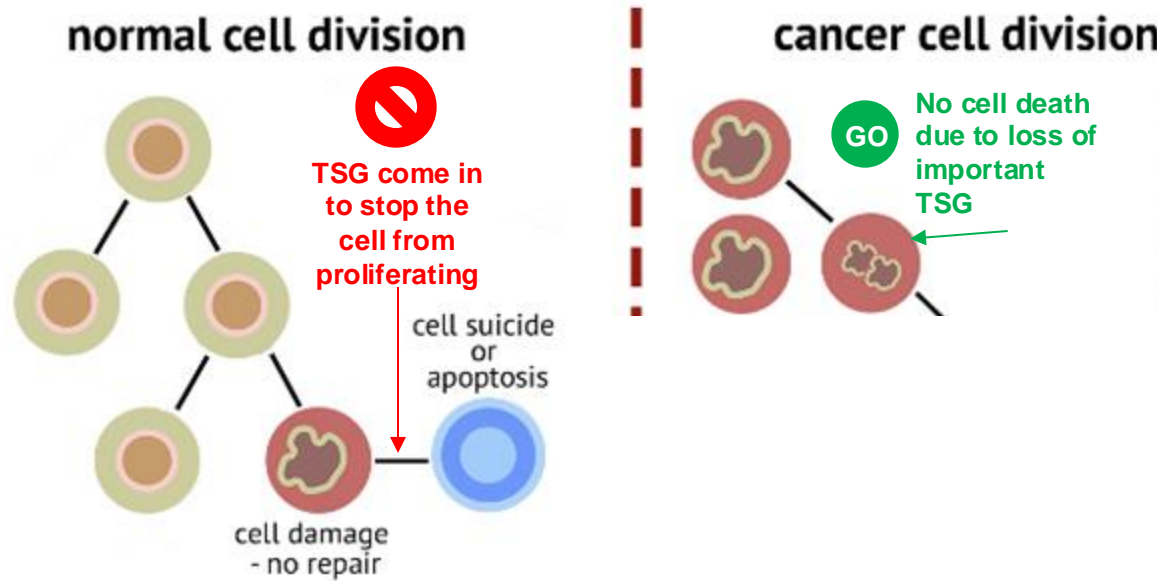


Copy Number Alterations

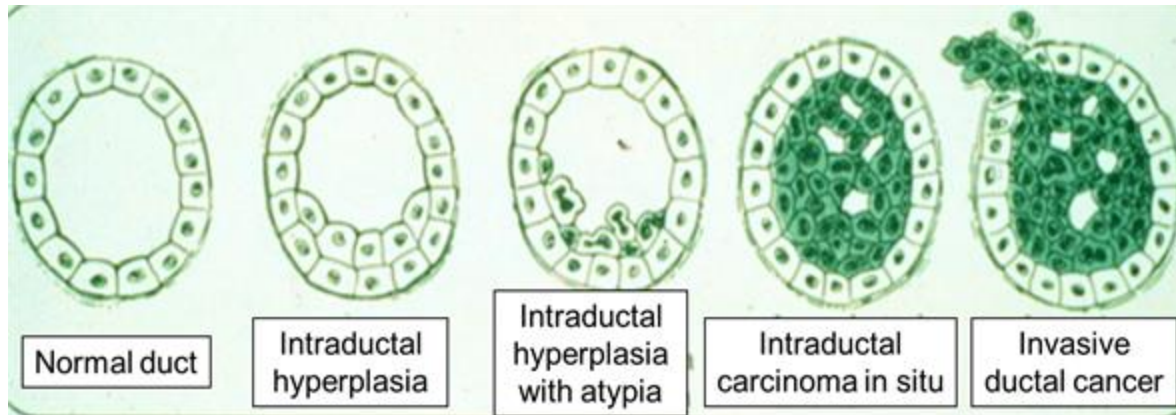
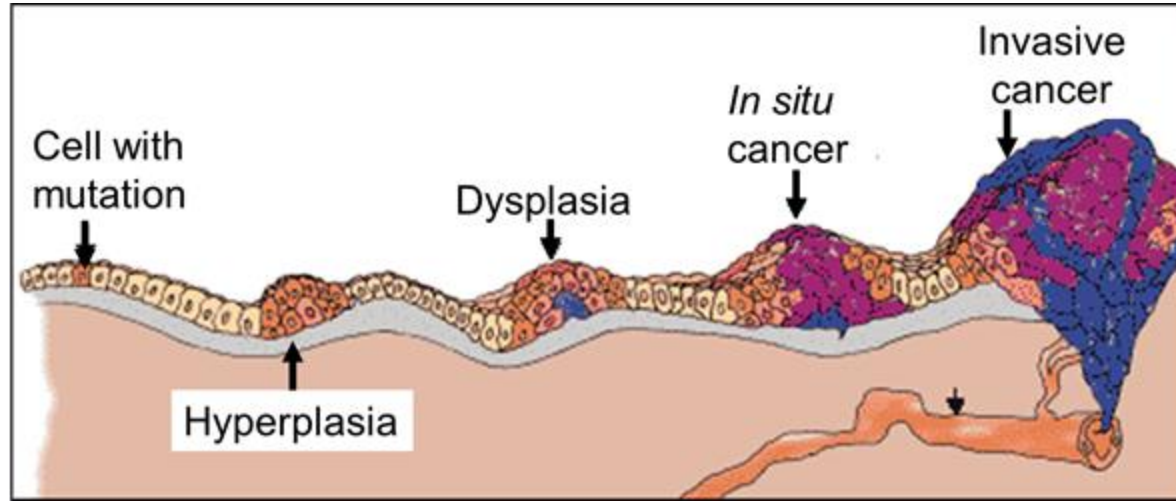


Processes are in place such that a cell with a driver mutation dies

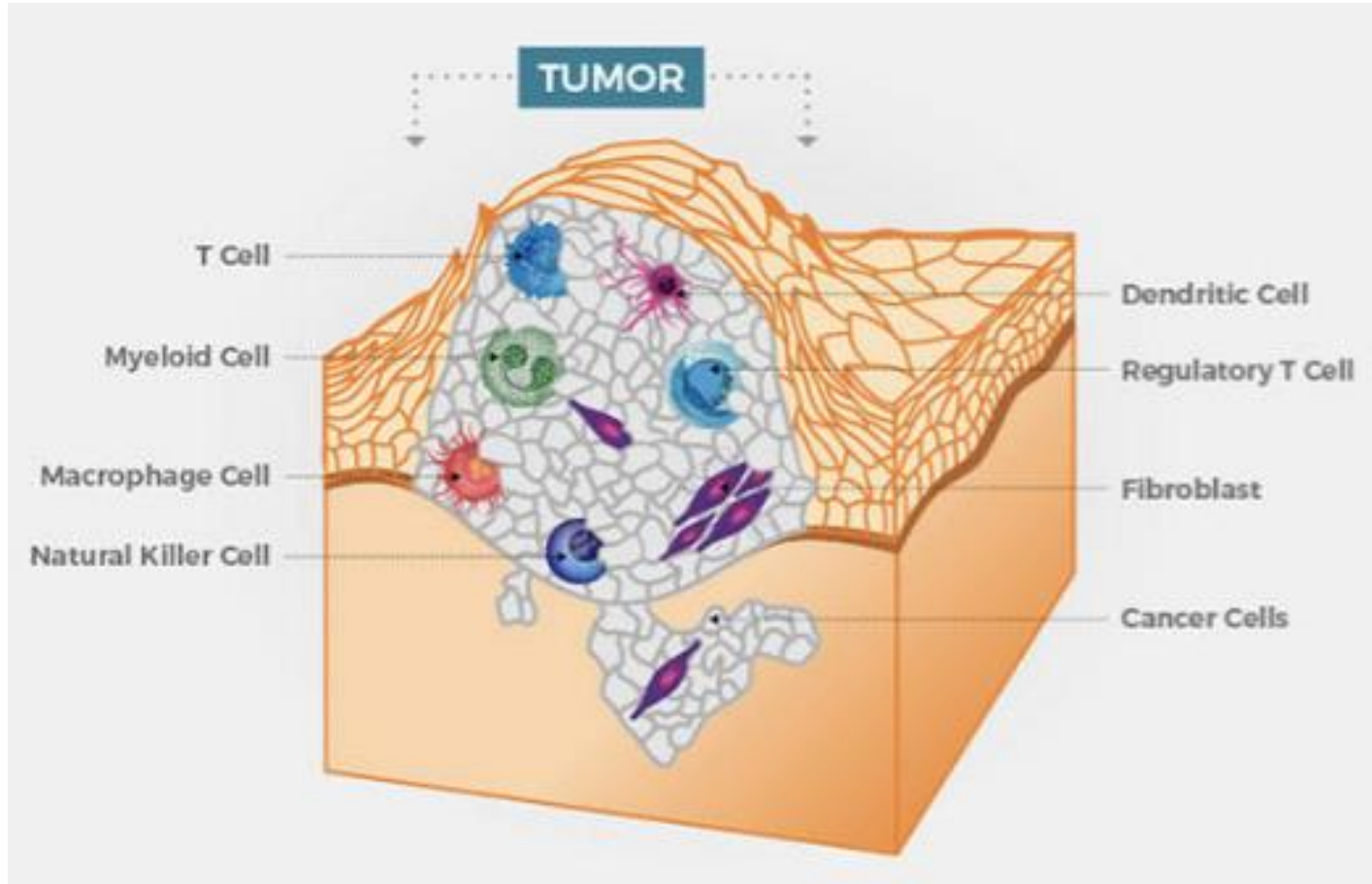
However, if the cell accumulates enough mutations → the cell with all its DNA mutations keeps making replicates of itself



Normal cells “transform” into cancer cells

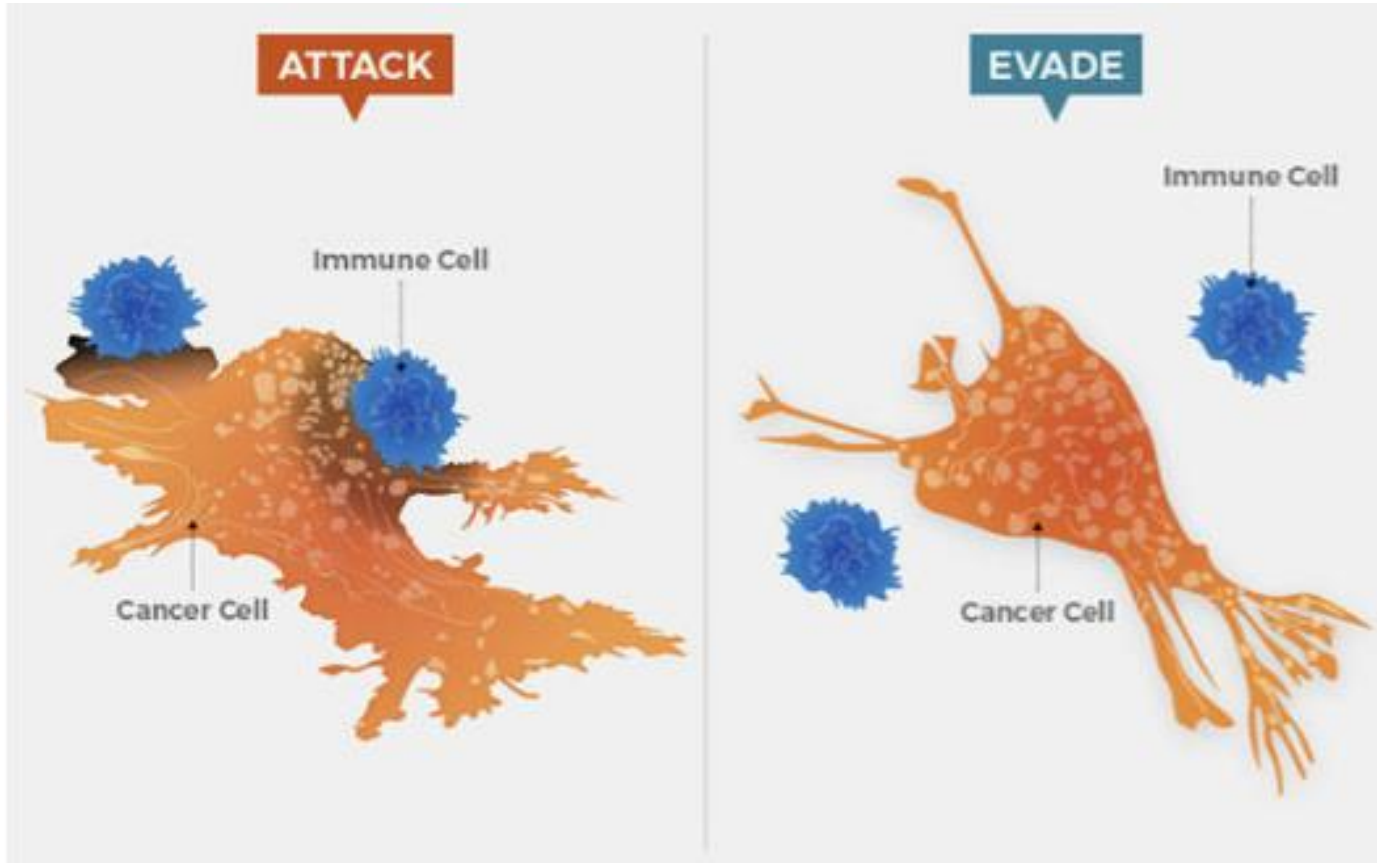


Cancer cells arise in a tissue surrounded by different types of cells → Tumor Microenvironment

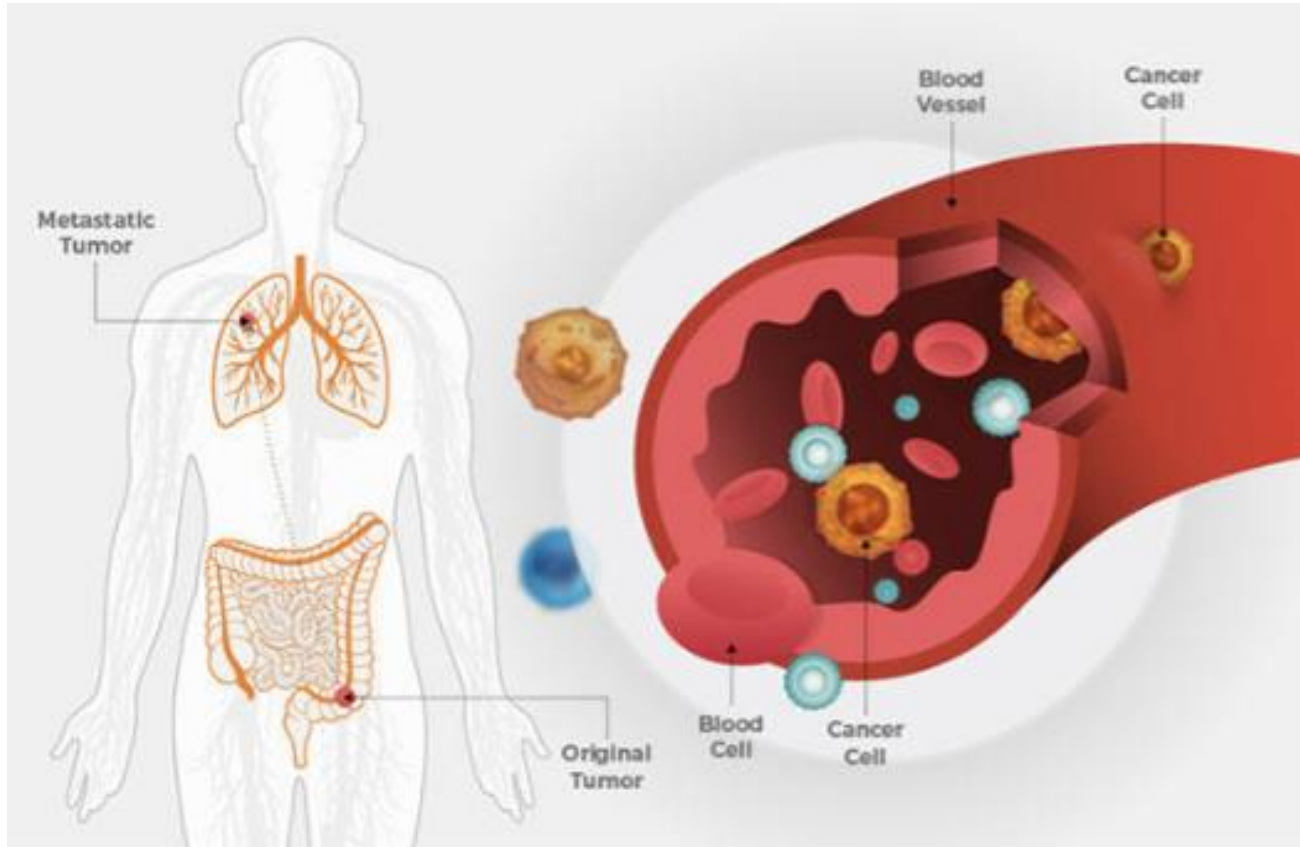


Interaction between cancer cells and its microenvironment affects their ability to grow and spread

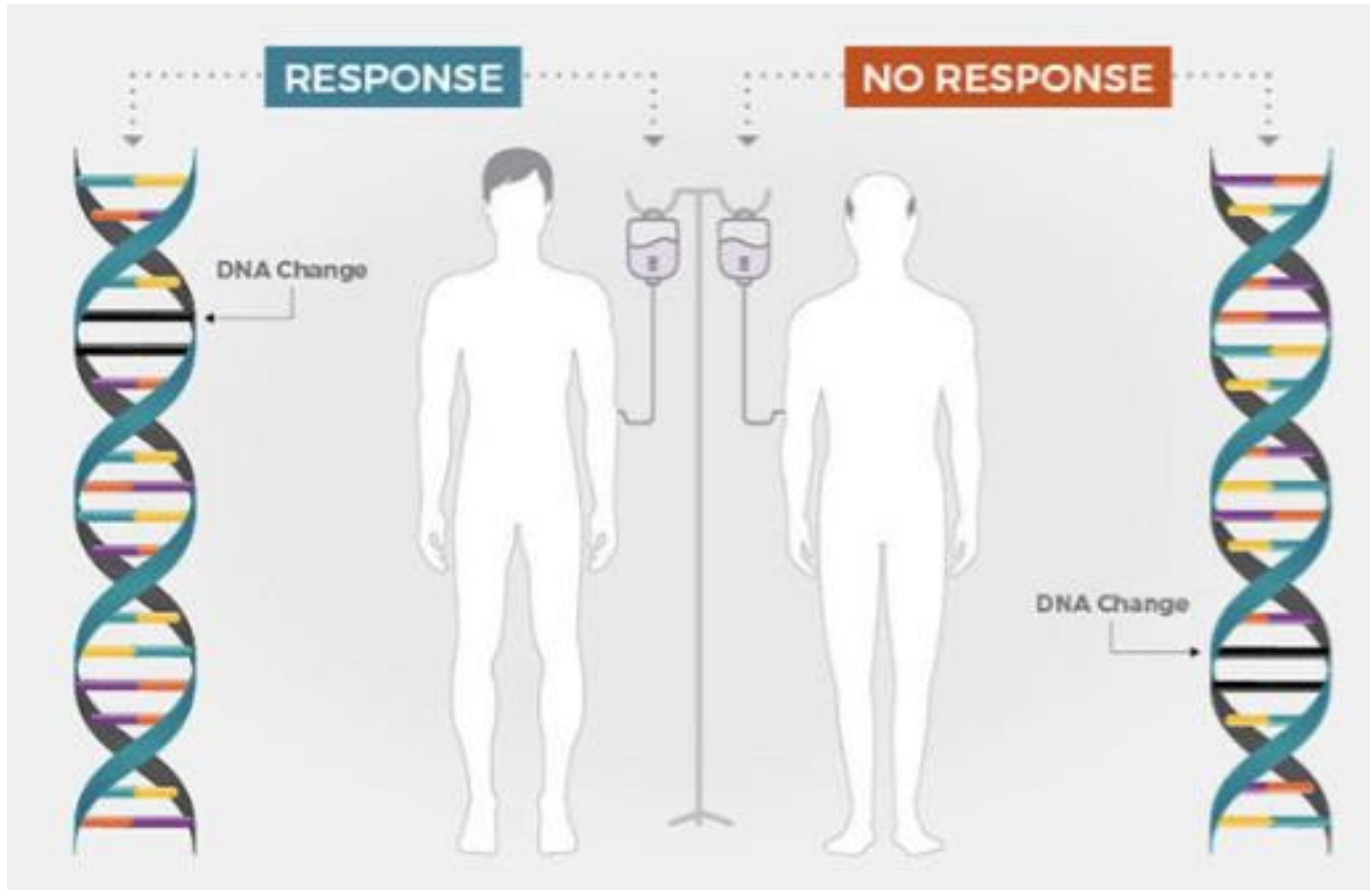
Particularly a cancer cell's interaction with the immune cells in its microenvironment



The cancer cell microenvironment also affects its ability to spread to distant organs (metastasize)

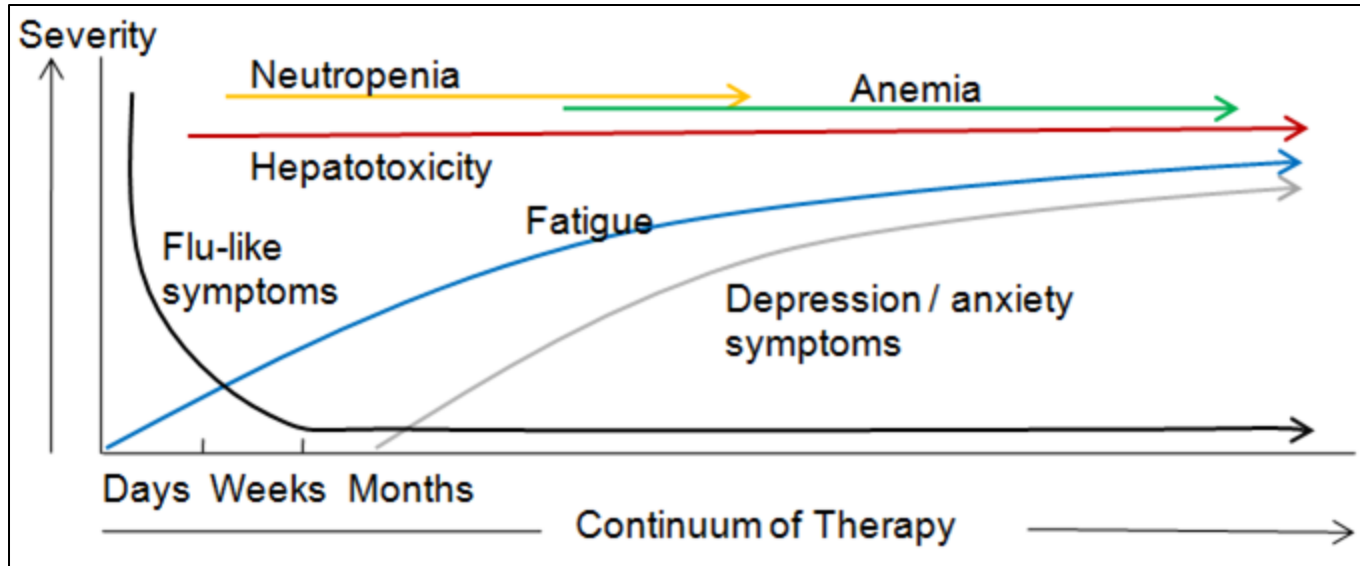


Genetic mutations can predict a patient's response to certain cancer treatments

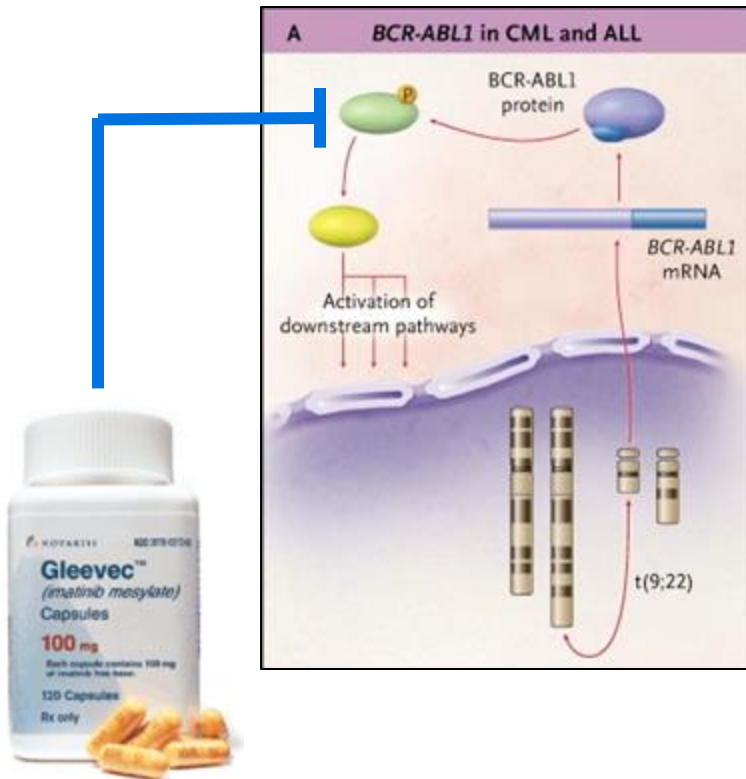


Before Targeted Therapies...

- Prior to 2001: <1 in 3 CML patients survived 5 years past diagnosis
- The most successful intervention at the time was Interferon a

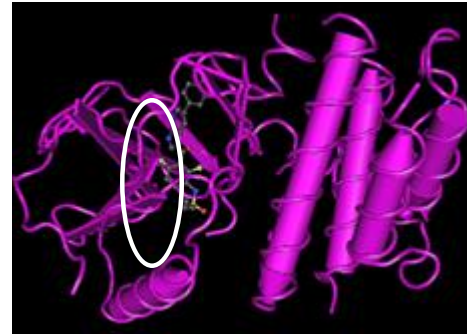
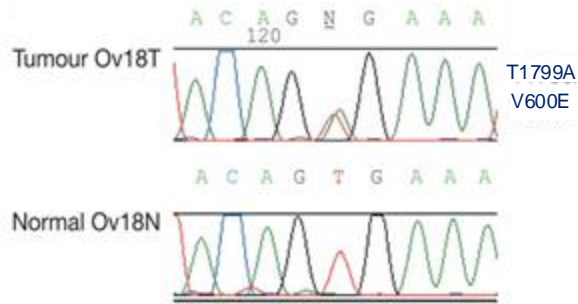


Poster Child for Targeted Therapy: Gleevec

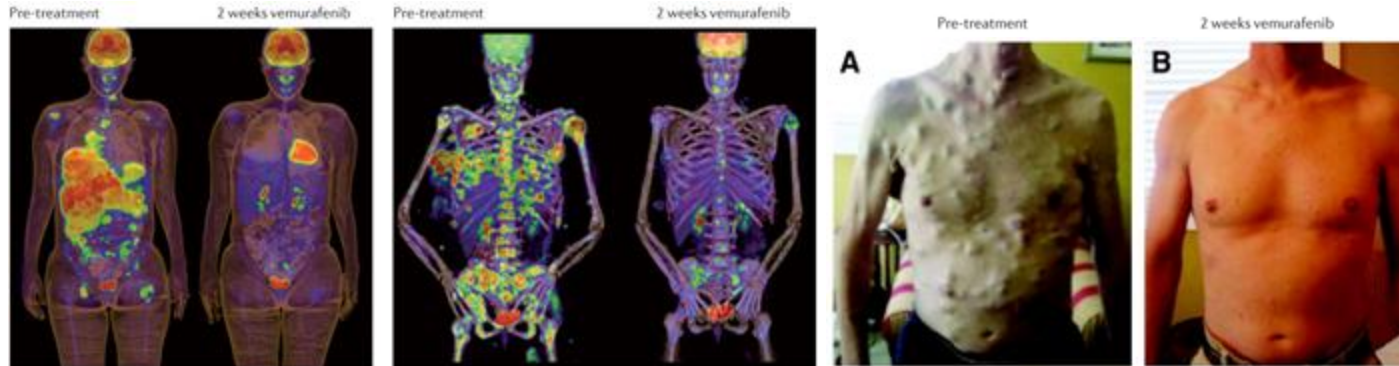


From sequencing to drug to treatment?

DNA Sequencing □ Discovery of a driver mutation **Drug** □ Inhibits the encoded protein
BRAF V600E **Vemurafenib**



Changes in Clinical Practice



Hallmarks of Cancer Cells



Questions??

