

SCHOOL OF PUBLIC HEALTH

EPIDEMIOLOGY

UNIVERSITY *of* WASHINGTON

Session 2:

Introduction to Epidemiology

and Genetic Epidemiology

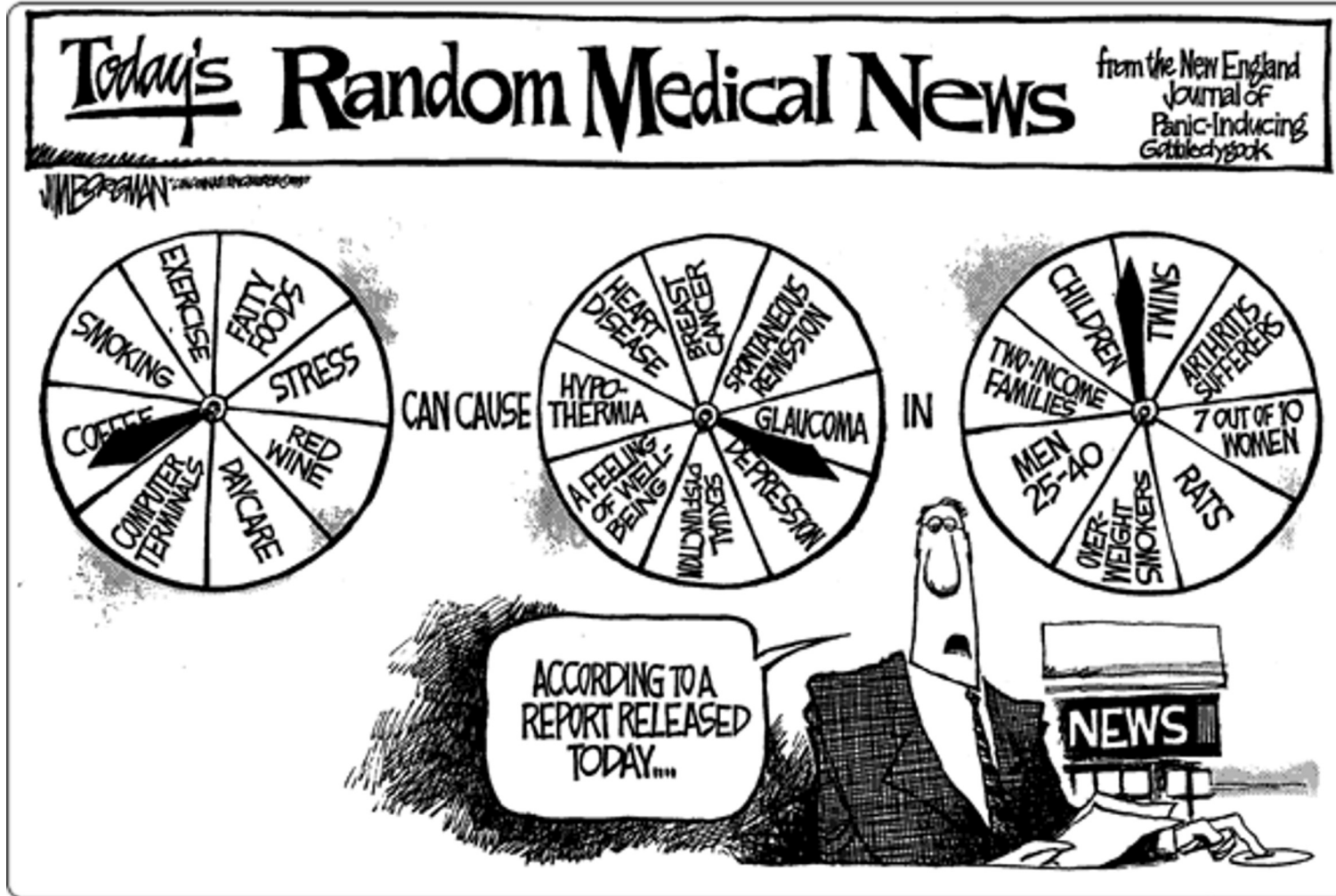


ep·i·de·mi·ol·o·gy

/,epə,dēmē'äləjē/

noun

the branch of medicine that deals with the incidence, distribution, and possible control of diseases and other factors relating to health.

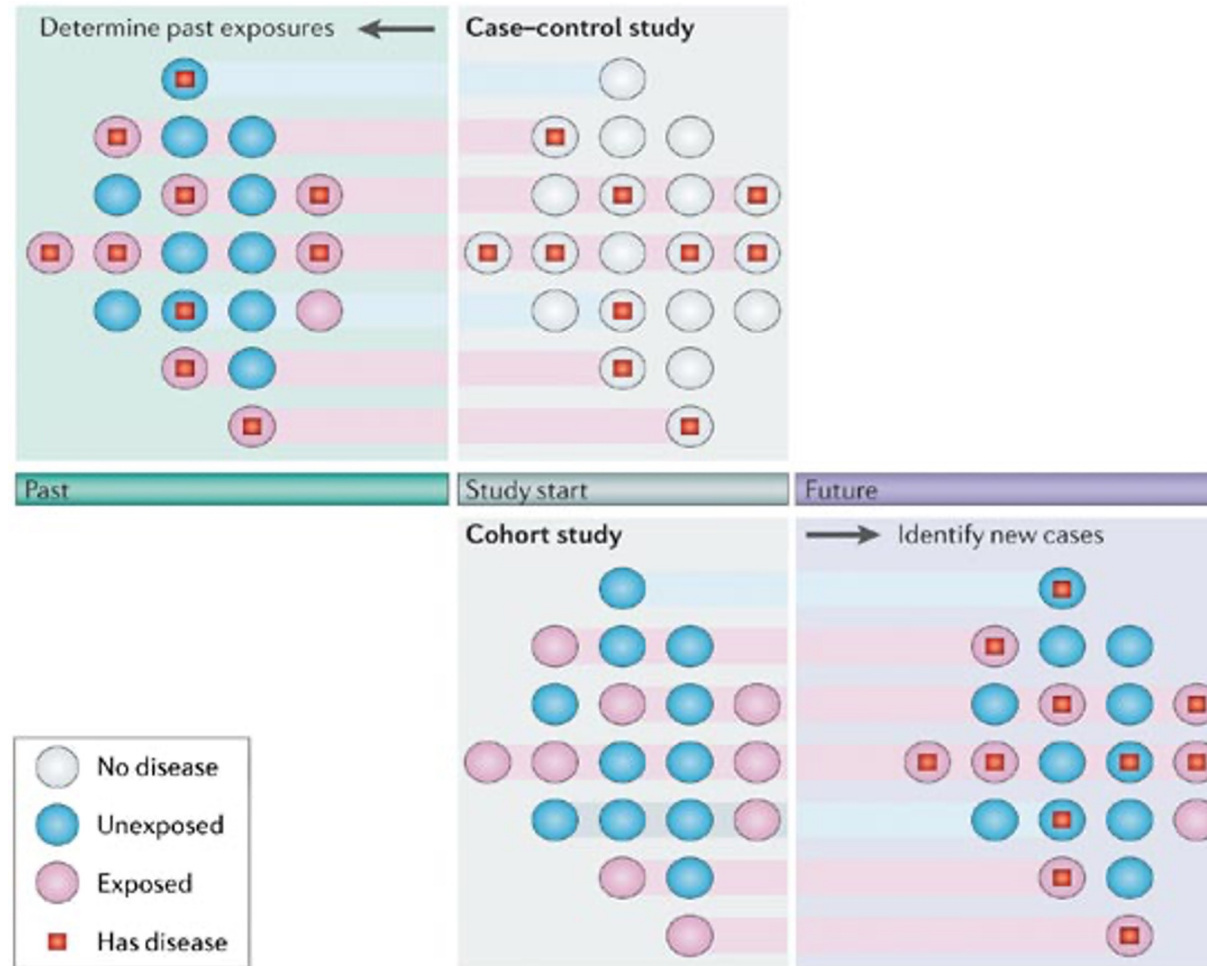


Major goals in Epidemiology

- > To obtain an *unbiased & precise* estimate of the true effect of an exposure or intervention on outcome in the population at risk
- > To use this knowledge to prevent and treat disease



Cohort vs. case-control studies



Estimated incidence rates in cohorts

Disease incidence per 100,000 per year (%)	Disease examples	Number of incident cases in 5 years for different cohort sizes		
		200,000	500,000	1,000,000
10 (0.01)	Parkinson disease, schizophrenia	91	228	457
50 (0.05)	Colorectal cancer, renal failure	456	1,141	2,282
100 (0.10)	Breast cancer, hip fracture	912	2,279	4,559
200 (0.20)	Diabetes, stroke, heart failure	1,820	4,550	9,100
500 (0.50)	Myocardial infarction, all cancers	4,524	11,309	22,618
3,000 (3.00)	Cataracts, hypertension	25,858	64,644	129,289

Estimated numbers of incident cases available after 5 years of follow-up across the entire age range in the US population are shown, assuming an attrition rate of 3% per year. Data are taken from the Incidence and Prevalence Database.

WACHOLDER, NATURE REVIEWS GENETICS 2000

Compared to cohorts, case-control studies are cheap, fast and powerful

However, case-control studies suffer from several drawbacks:

the need to identify appropriate controls

they are more sensitive to recall bias

Association and Causality

- > An exposure and outcome are associated if there is a differential distribution:
 - The prevalence of exposure differs between cases and controls.
- > An exposure is causal for the outcome if the presence (or absence) of the exposure directly or indirectly influences whether the outcome occurs.

THE FAMILY CIRCUS



"I wish they didn't turn on that seatbelt sign so much! Every time they do, it gets bumpy."

Some sources of bias in epidemiology

> **Selection Bias**

- When there is a difference between study participants and the general population (affects generalizability, e.g, the Nurses' Health Study) **OR** when cases and controls are drawn from different source populations (e.g., cases are recruited through population-based cancer registry, controls are recruited from the local gym).

> **Survival bias**

- When cases are recruited some time after they were diagnosed. Might lead to a milder form of disease. This is especially true for aggressive/fatal disease (e.g., pancreatic cancer, heart attack)

> **Diagnostic bias**

- If the investigator determining the outcome knows whether the person was exposed or not to the risk factor under study (e.g., if the radiologist knows that a potential pulmonary disease patient smokes, they may look more carefully at the x-ray).

> **Recall bias**

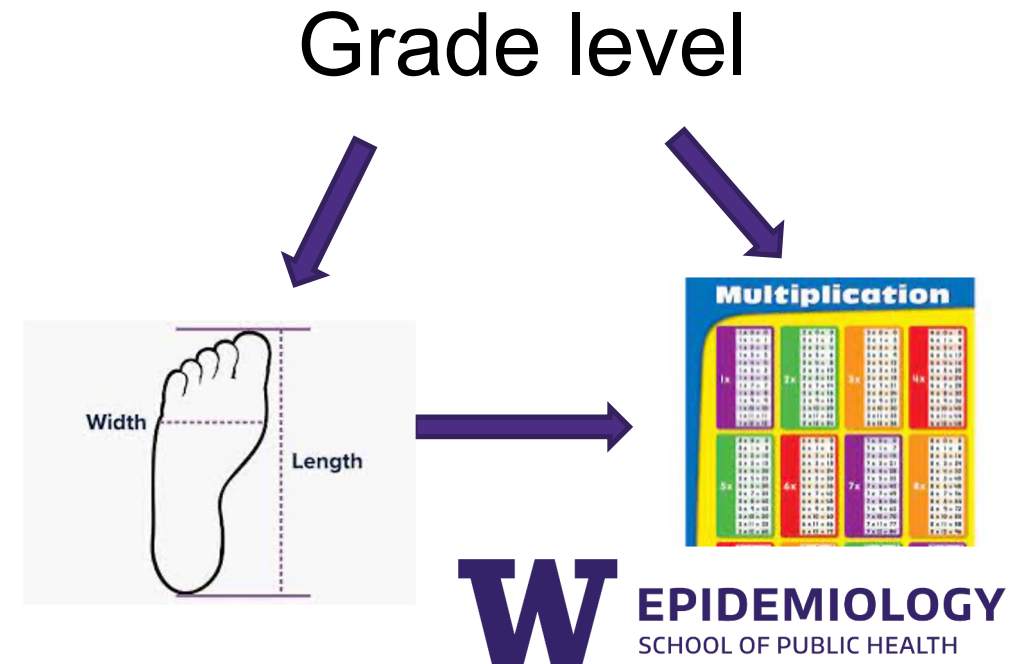
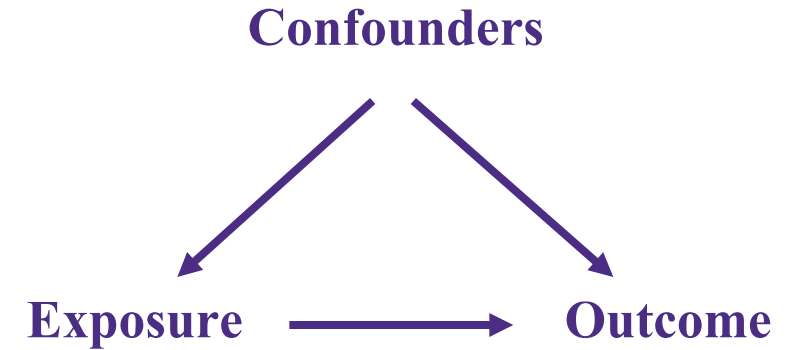
- Accuracy and completeness of exposures, lifestyle behaviors,... (e.g., cases might be more motivated to complete a questionnaire accurately).

Confounding

> A confounder is often defined as a factor that is:

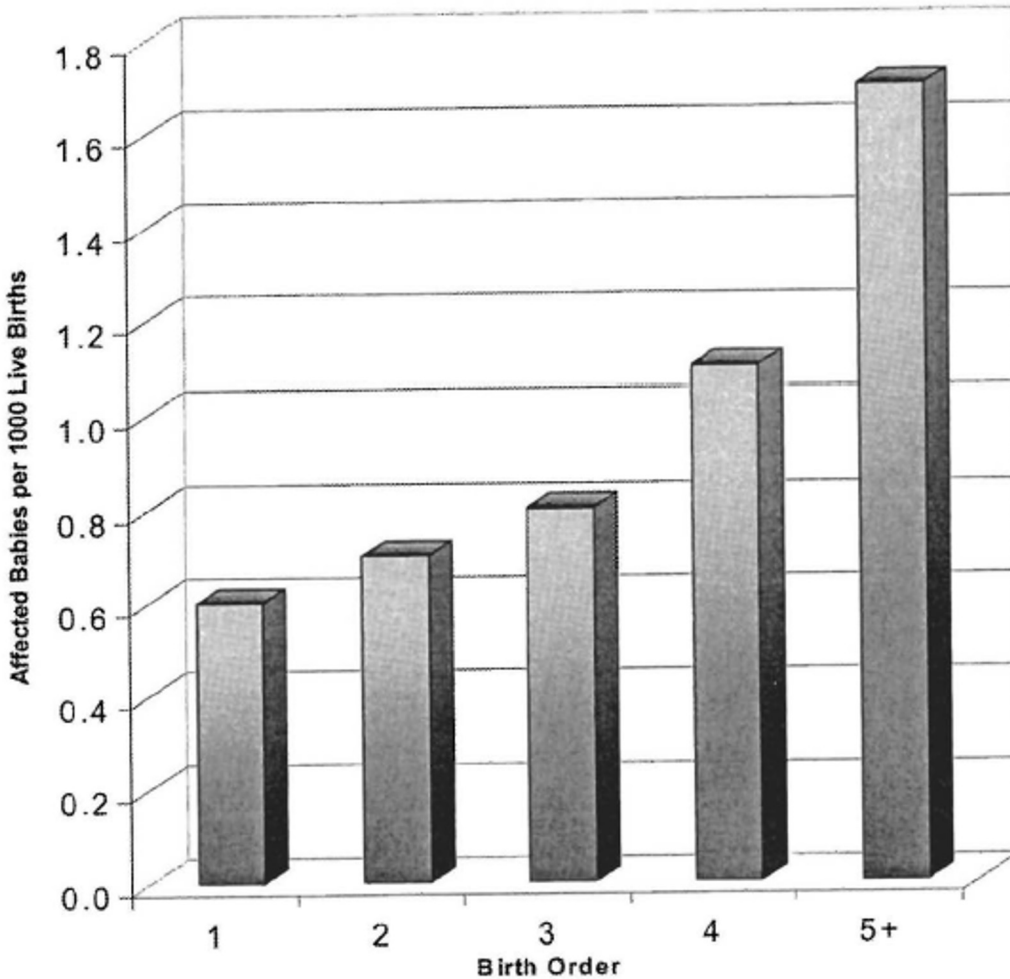
1. A risk factor for the disease
2. Associated with the exposure
3. Not a direct result of the exposure

> Confounding can lead to false positive findings.



BREAKOUT ACTIVITY

Confounding example: Birth order and Down syndrome



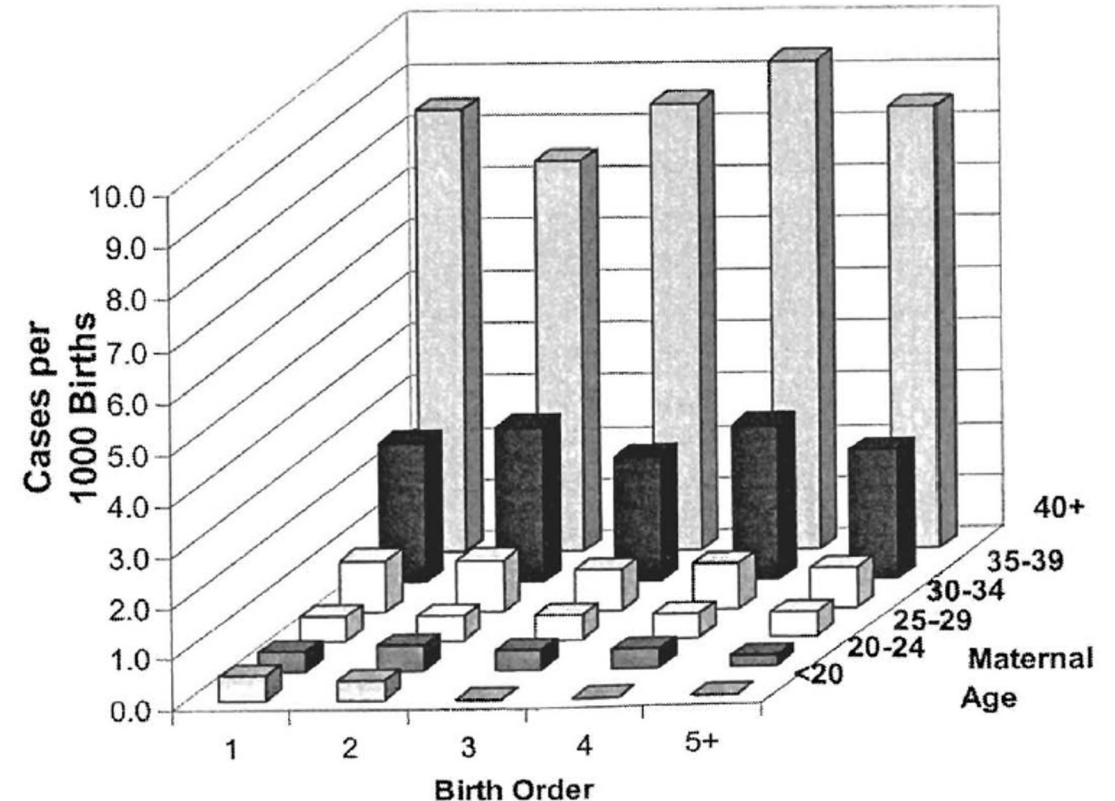
Data from Stark and Mantel (1966)

Source: Rothman 2002

1. Can you think of a factor that would confound the observed association?
1. How can you use data on your proposed confounding factor to reassess the association between birth order and Down Syndrome?
1. Can you think of potential confounders in genetic epidemiology?

Confounding example: Birth order and Down syndrome

- > Later order children have higher risk
 - Maternal age is associated with birth order
 - Maternal age is associated with Down Syndrome
- > Stratifying on maternal age, there is no longer evidence of an association between birth order and Down syndrome



Data from Stark and Mantel (1966)

Source: Rothman 2002

What is genetic epidemiology?

Genetic epidemiology is the study of the role of **genetic** factors in determining health and disease in families and in populations, and the interplay of such **genetic** factors with environmental factors.

Smoking addiction gene found

Scientists say a gene makes people more likely to get hooked on tobacco, causing them to smoke more, making it harder to quit, and leading more often to deadly lung cancer. [Full story](#)

[Newsweek: Differing conclusions](#)

[Researchers make human-cow embryos](#)

[Science wishy-washy on water benefits](#) | [Vote](#)



Does aspirin prevent colorectal cancer? Depends on your DNA



Ed was unlucky enough to find the needle in the haystack!

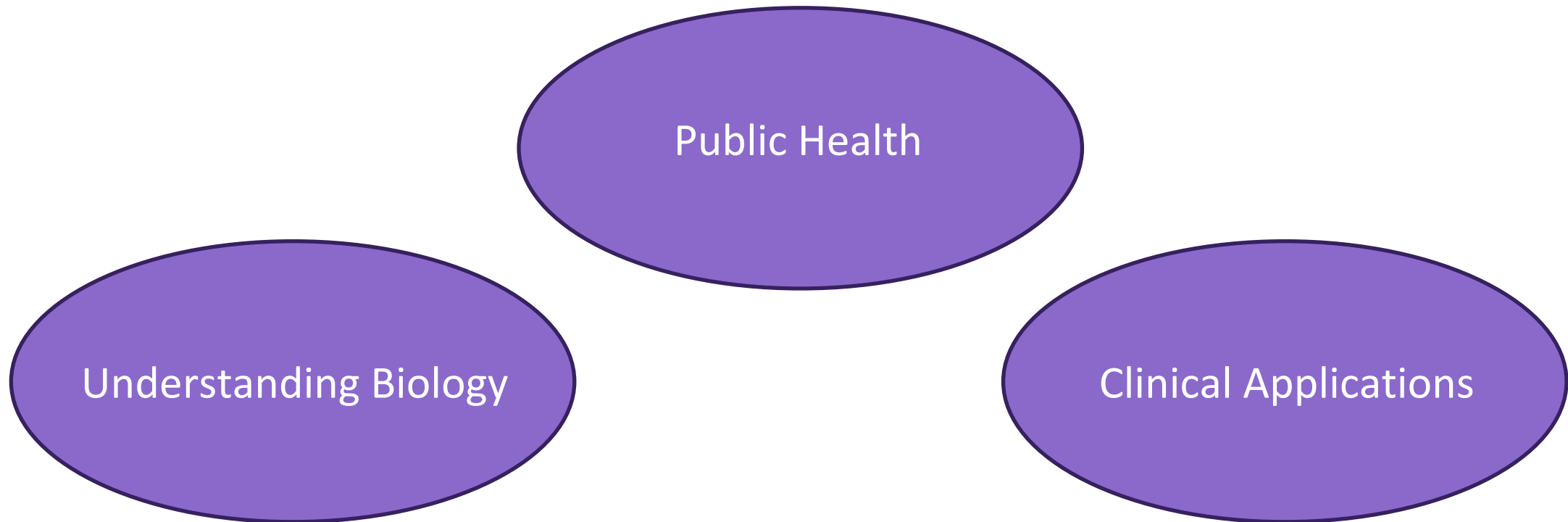
Biobanks are becoming a large part of (genetic) epi research



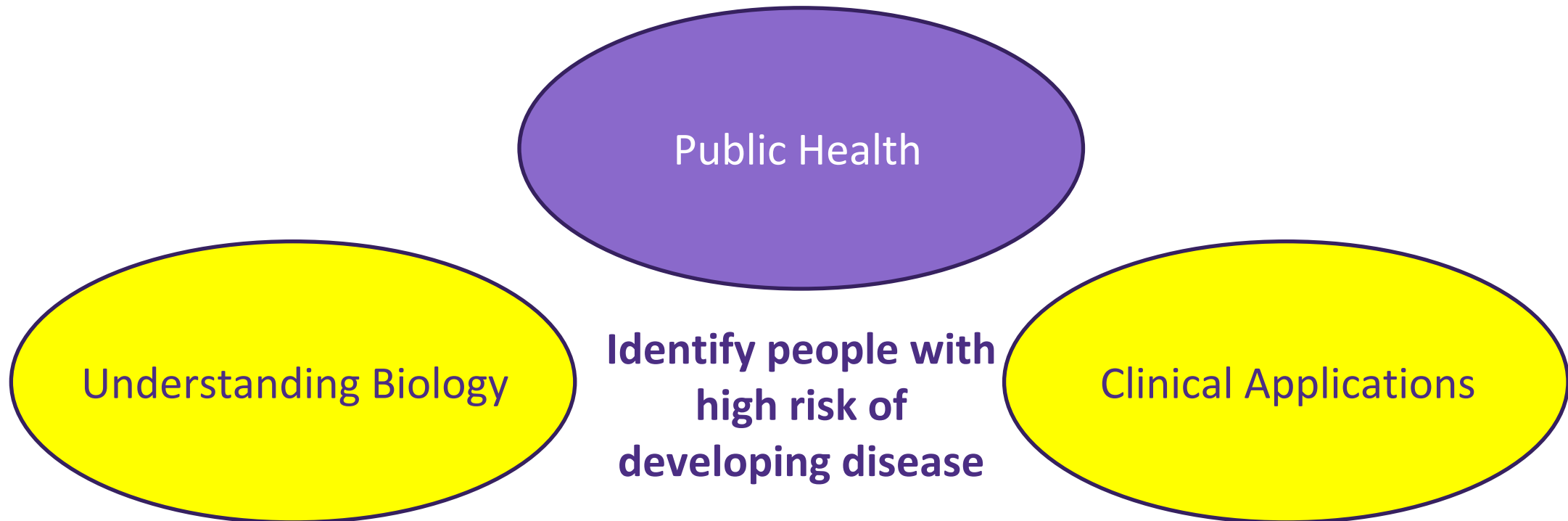
BIOBANK JAPAN

Why do we want to study how our genome is involved in disease?

Why do we want to study how our genome is involved in disease?



Why do we want to study how our genome is involved in disease?



OP-ED CONTRIBUTOR

My Medical Choice

By ANGELINA JOLIE

Published: May 14, 2013 | 1712 Comments

LOS ANGELES

[Enlarge This Image](#)



Loren Capelli

MY MOTHER fought cancer for almost a decade and died at 56. She held out long enough to meet the first of her grandchildren and to hold them in her arms. But my other children will never have the chance to know her and experience how loving and gracious she was.

We often speak of “Mommy’s mommy,” and I find myself trying to explain the illness that took her away from us. They have asked if the same could happen to me. I have always told them not to worry, but the truth is I carry a “faulty” gene, BRCA1, which sharply increases my risk of developing breast cancer and ovarian cancer.

FACEBOOK

TWITTER

GOOGLE+

SAVE

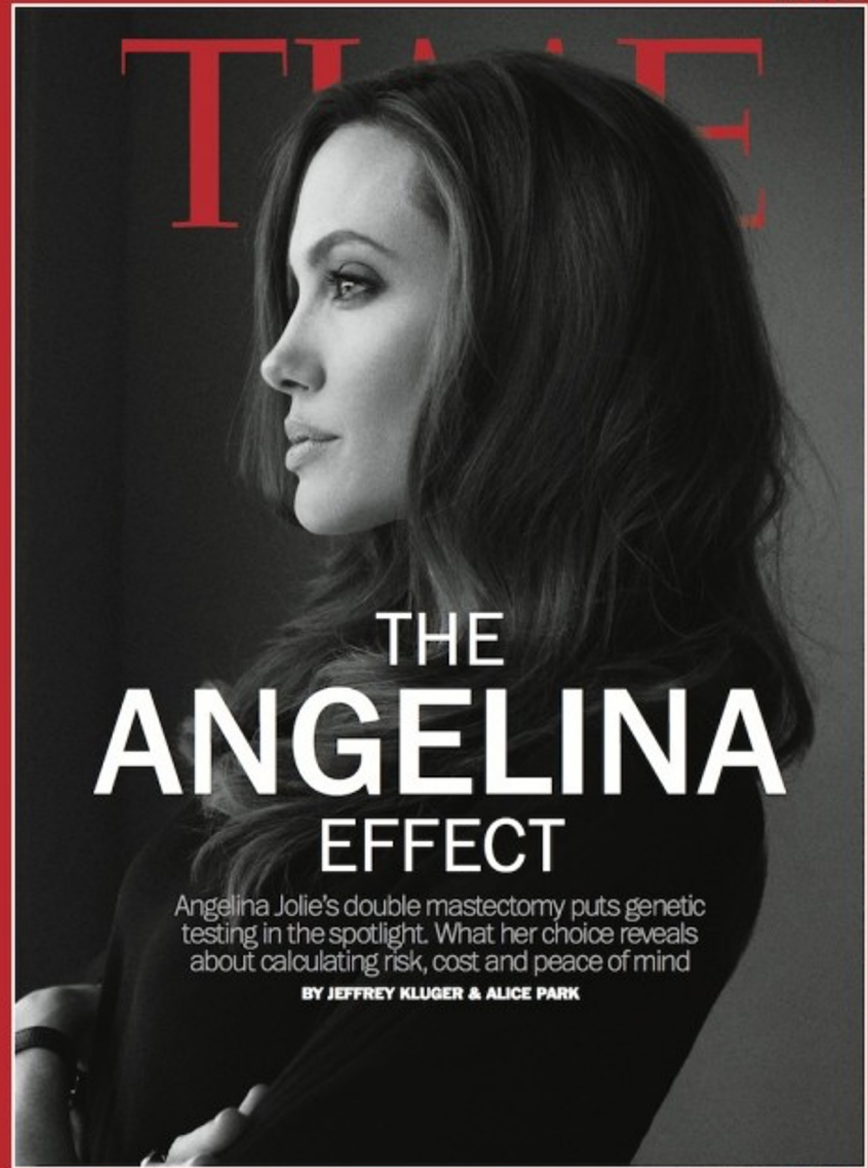
E-MAIL

SHARE

PRINT

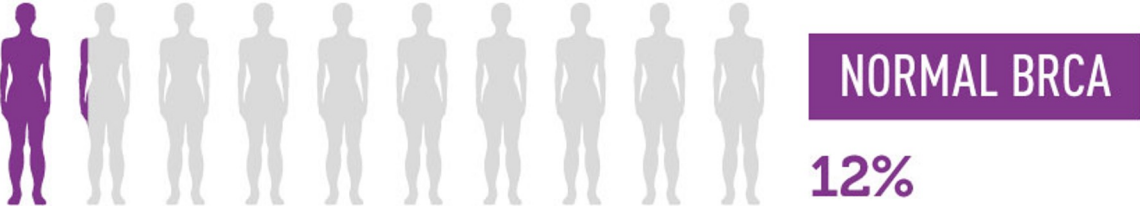
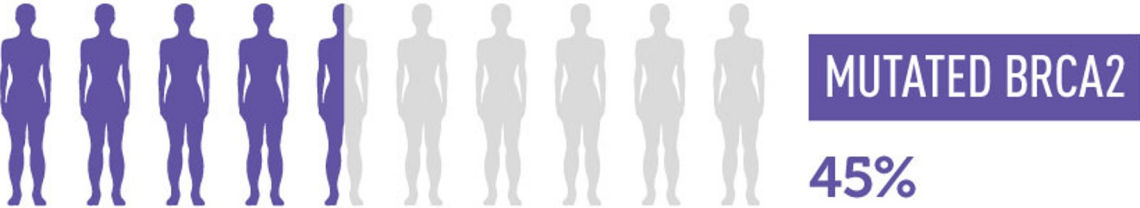
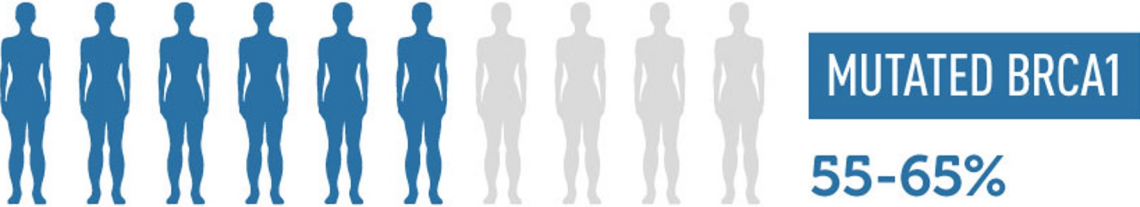
REPRINTS

Enough Said
Now Playing

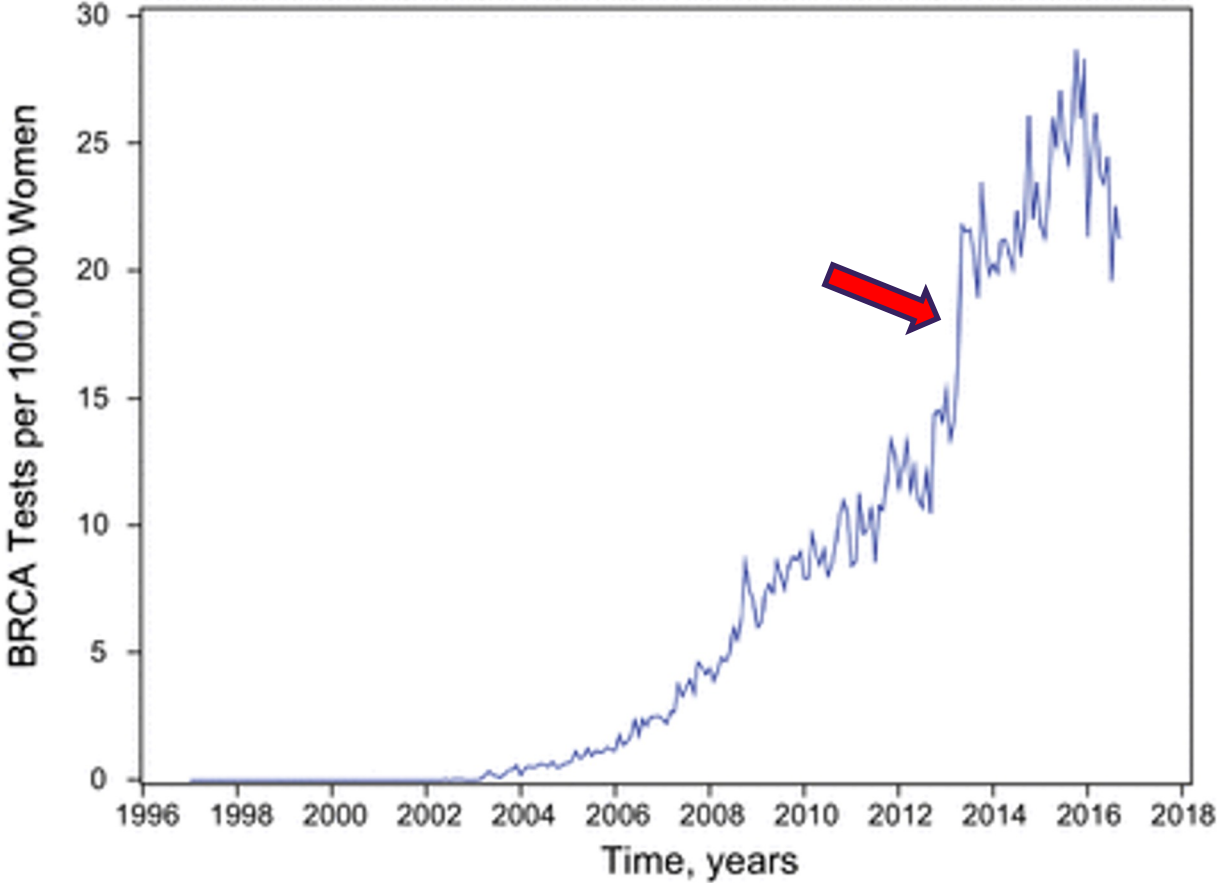


NATIONAL CANCER INSTITUTE CHANCES OF DEVELOPING BREAST CANCER BY AGE 70

Specific inherited mutations in the BRCA1 and BRCA2 genes increase the risk of breast and ovarian cancers. Testing for these mutations is usually recommended in women without breast cancer only when the person's individual or family history suggests the possible presence of a harmful mutation in BRCA1 or BRCA2. Testing is often recommended in younger women newly diagnosed with breast cancer because it can influence treatment decisions and have implications for their family members.



www.cancer.gov/brca-fact-sheet



Liede et al. BCRT 2018

Genetic testing of *BRCA1*

23andMe Granted First FDA Authorization for Direct-to-Consumer Genetic Test on Cancer Risk

March 6, 2018

Authorization allows 23andMe to report on BRCA1- and BRCA2-related genetic risk for breast, ovarian and prostate cancer

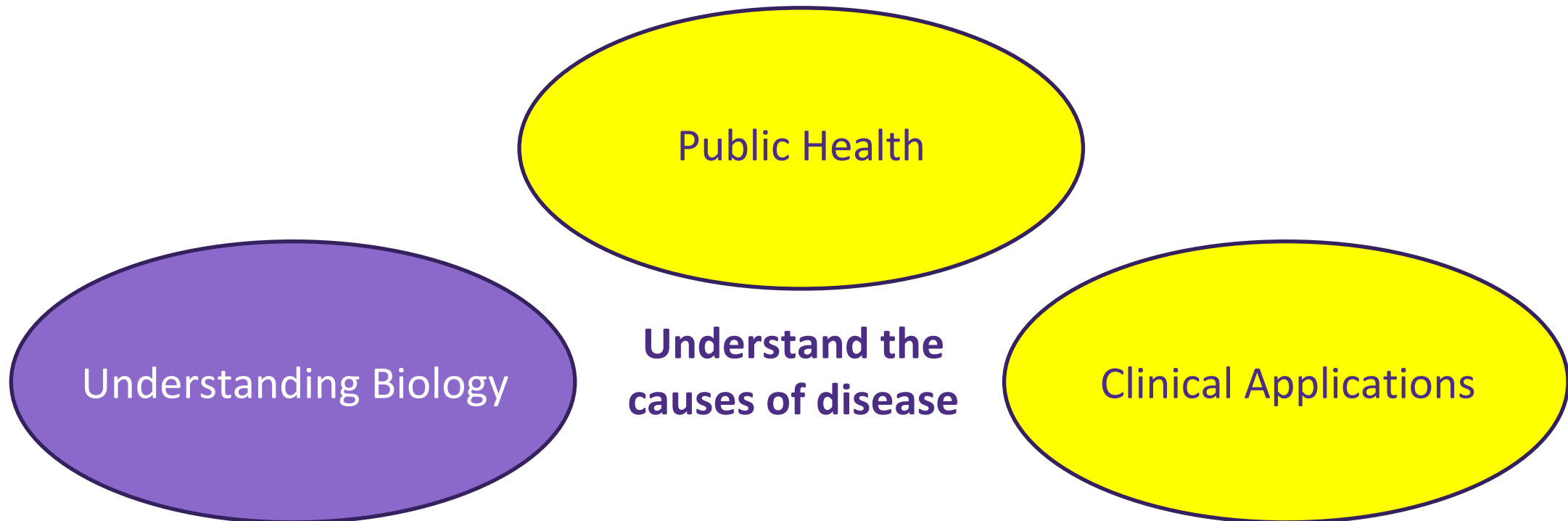
The screenshot shows a 23andMe website page with a dark purple background. At the top, there are navigation links for 'SHOP', 'LEARN', 'SIGN IN', 'REGISTER KIT', 'HELP', and a 'Shop' button. The main heading is 'DO YOU SPEAK BRCA?'. Below this, several callout boxes highlight specific BRCA variants: BRCA2 6174deIT, BRCA1 185delAG, BRCA1 5382insC, and BRCA1 185delAG. Text boxes state: 'BRCA variants affect women and men.', 'People without a BRCA variant are still at risk for cancer.', 'Specific BRCA variants are more common in certain populations.', and 'Live in the know™'. At the bottom, there is a link to 'Learn more about the 23andMe BRCA1/BRCA2 (Selected Variants) Genetic Health Risk report here' and a disclaimer: 'Have a personal or family history of cancer? Talk to your healthcare provider to determine if comprehensive genetic testing is appropriate.' The footer includes 'BRCA EDUCATION' and the text 'Let's begin at the very beginning.' followed by a detailed explanation of what BRCA stands for.

Don't Count on 23andMe to Detect Most Breast Cancer Risks, Study Warns

“In about 5,000 subjects, analysts identified at least one variant known to significantly increase an individual’s risk of breast and ovarian cancer. Among the Ashkenazi Jews in the positive group, 81 percent had one of the three founder mutations, suggesting that 23andMe’s test could be helpful for them. Among the rest, 94 percent carried variants that would have failed to be detected by 23andMe.”

New York Times, April 16, 2019

Why do we want to study how our genome is involved in disease?



“Association does not imply causation”



HDL (“Good”) Cholesterol and Myocardial Infarction (MI)

↑ HDL -> ↓ MI risk

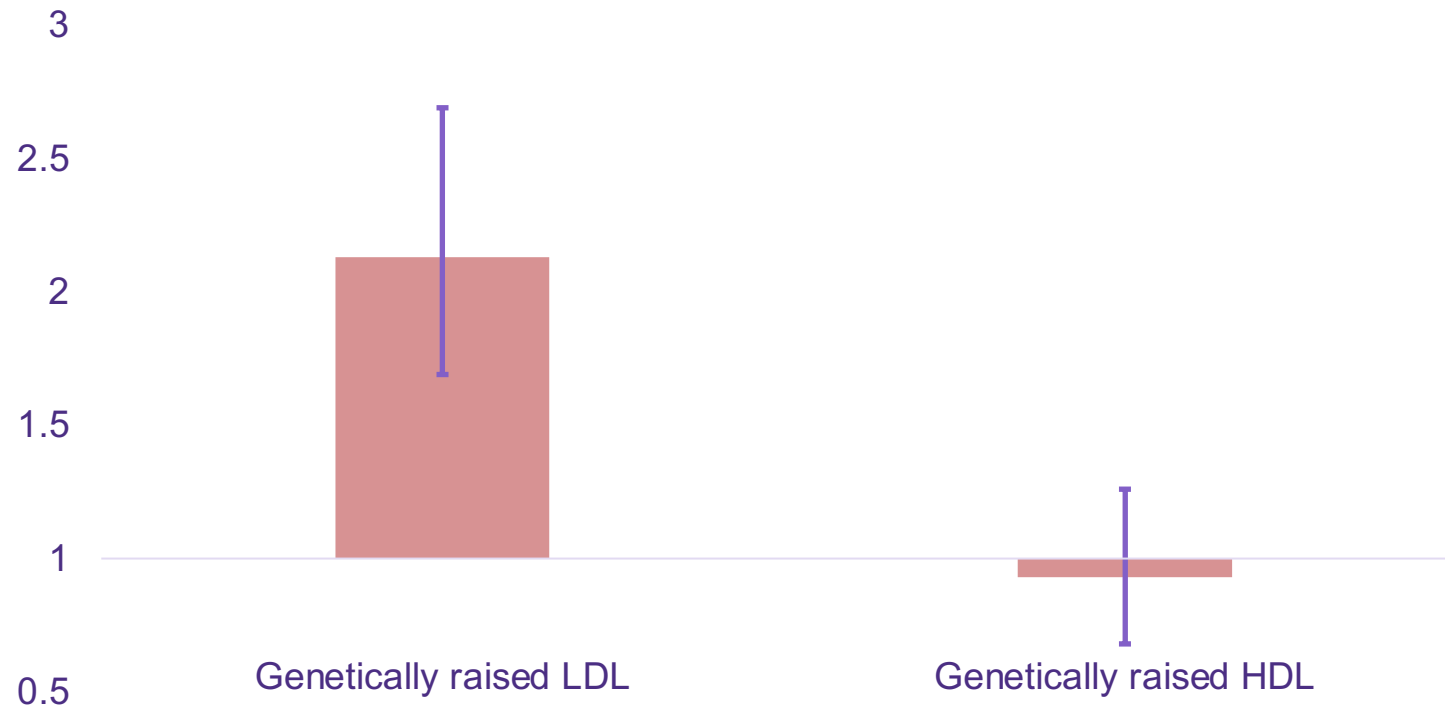


Increasing HDL concentrations might help decrease cardiovascular disease risk.

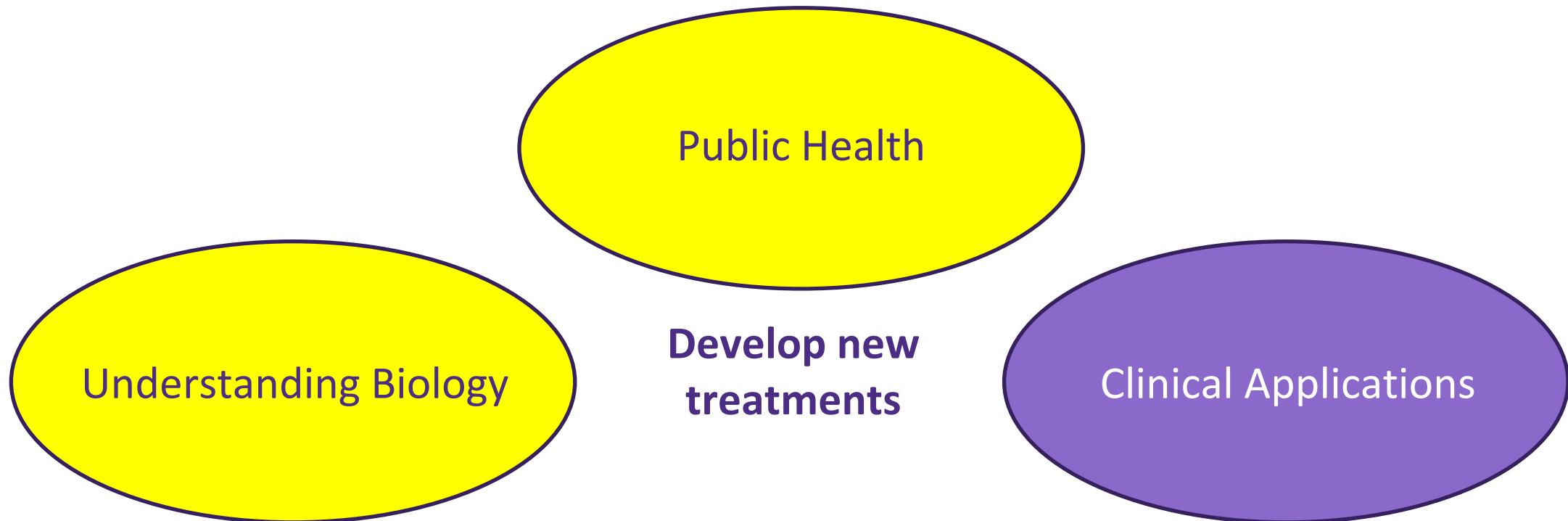


People who carry genetic variants that increase HDL do not have a lower risk of MI

Since HDL is correlated with exercise, weight loss, diet (nuts, fish) it is likely that these lower your risk for MI rather than HDL itself

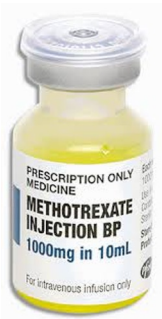


Why do we want to study how our genome is involved in disease?

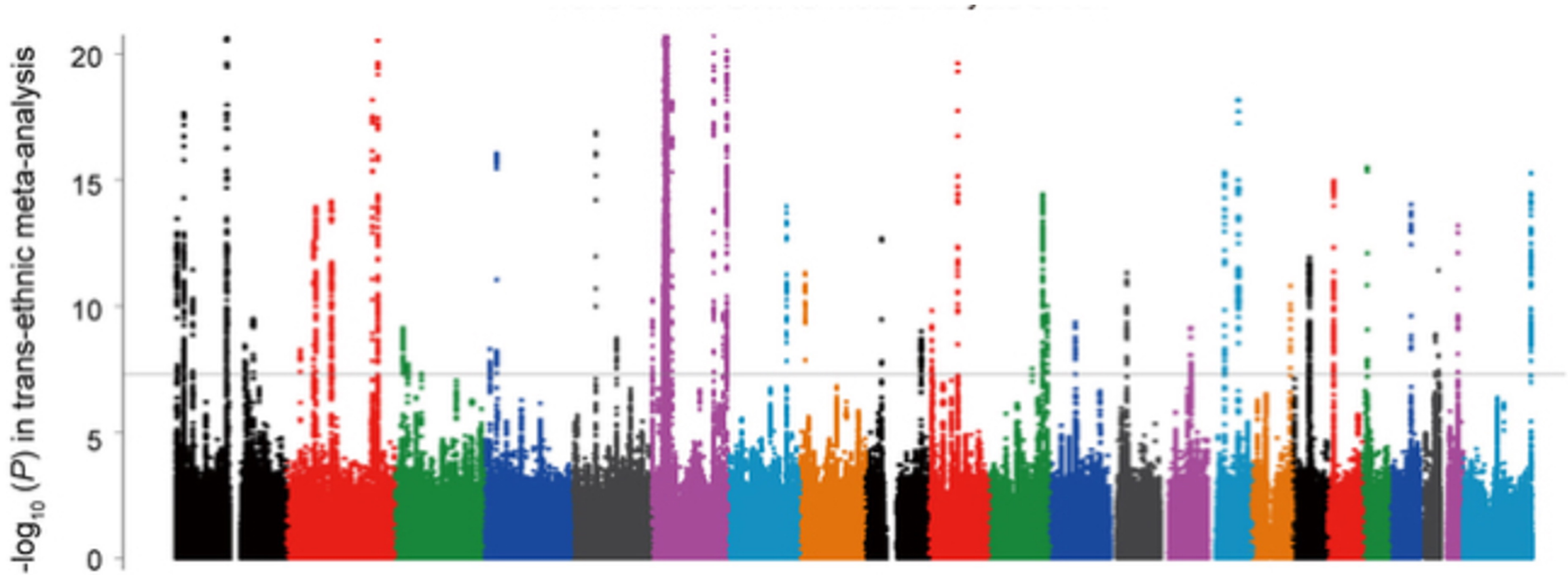


Rheumatoid Arthritis – an inflammatory, crippling, incurable disease

> In 2014, an estimated 0.5% of US adults age ≥ 18 had RA (~1.3M people).



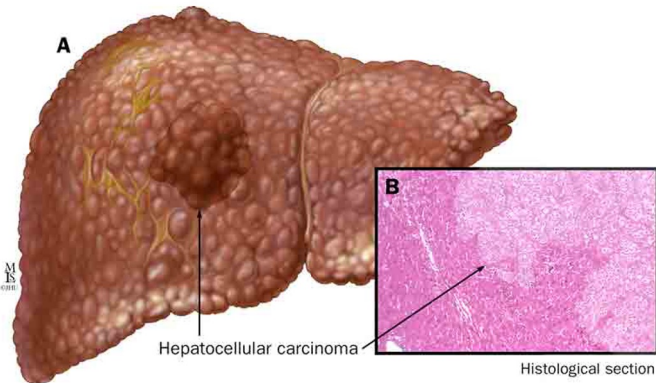
A study of 10 million genetic variants in 29,880 RA cases and 73,758 controls



Okada, Nature, 2014

Identified genes are targets of approved therapies for RA, and further suggest that drugs approved for other diseases may be repurposed for the treatment of RA

flavopiridol



CAPRIDINE-β
(c-1748)

Novel Chemotherapies for Prostate Cancer Patients Throughout Multiple Stages and Clinical States of Treatment

AV Therapeutics, Inc.
Advanced Cancer Chemotherapies

The advertisement features a photograph of a smiling elderly man in a white shirt being touched on the shoulder by a doctor in a white coat. The text is overlaid on the right side of the image.

Cancers/psoriasis

Breast cancer

NDC 0069-0189-21

Ibrance™
(palbociclib)
capsules

The Pfizer logo is a green oval with the word 'Pfizer' in white. Below it, the text 'Ibrance™ (palbociclib) capsules' is displayed in a bold, black, sans-serif font.

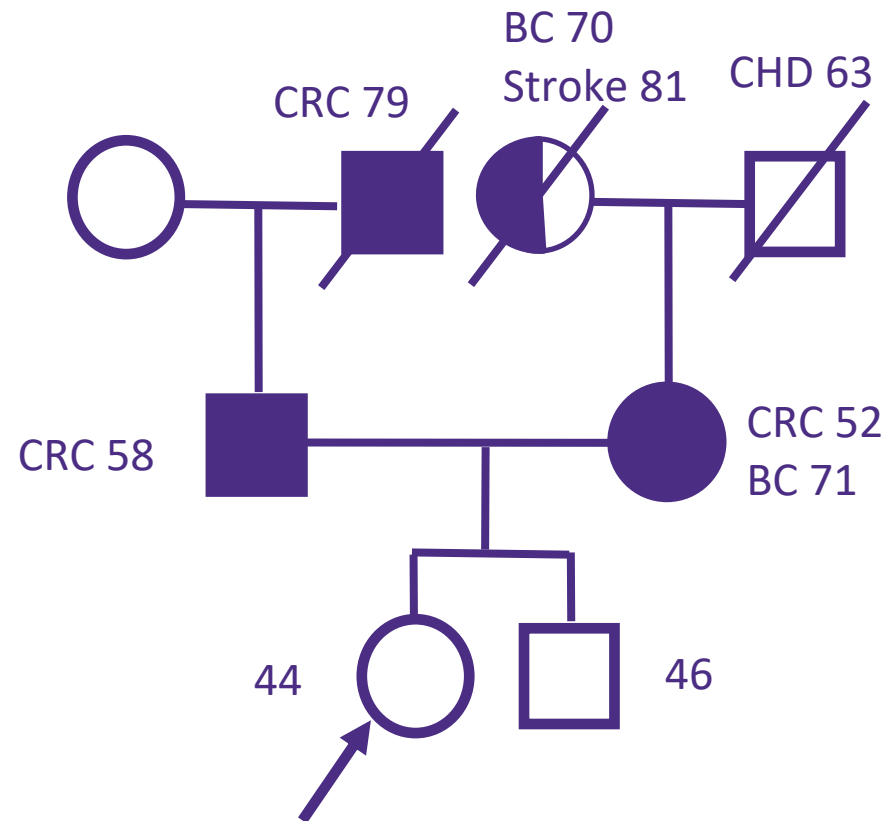
125 mg

For Oncology Use Only
21 Capsules Rx only

Lymphoma/Leukemia/Liver cancer

Okada, Nature, 2014

What would you say to this patient?



Key

Male



Affected male



Deceased male



Female



Affected Female



Deceased Female



Affected female with different cancer



CRC – colorectal cancer

BC – breast cancer