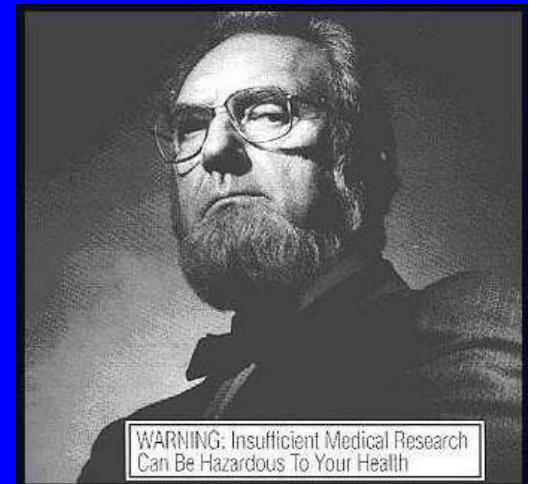


BIOE 301

Lecture Eleven



Four Questions

- What are the major health problems worldwide?
- Who pays to solve problems in health care?
- How can technology solve health care problems?
- How are health care technologies managed?

Three Case Studies

- Prevention of infectious disease
 - HIV/AIDS
- Early detection of cancer
 - Cervical Cancer
 - Ovarian Cancer
 - Prostate Cancer
- Treatment of heart disease
 - Atherosclerosis and heart attack
 - Heart failure

Outline

- The burden of cancer
- How does cancer develop?
- Why is early detection so important?
- Strategies for early detection
- Example cancers/technologies
 - Cervical cancer
 - Ovarian cancer
 - Prostate cancer

The Burden of Cancer: U.S.

■ Cancer:

- 2nd leading cause of death in US
- 1 of every 4 deaths is from cancer

■ 5-year survival rate for all cancers:

- 62%

■ Annual costs for cancer:

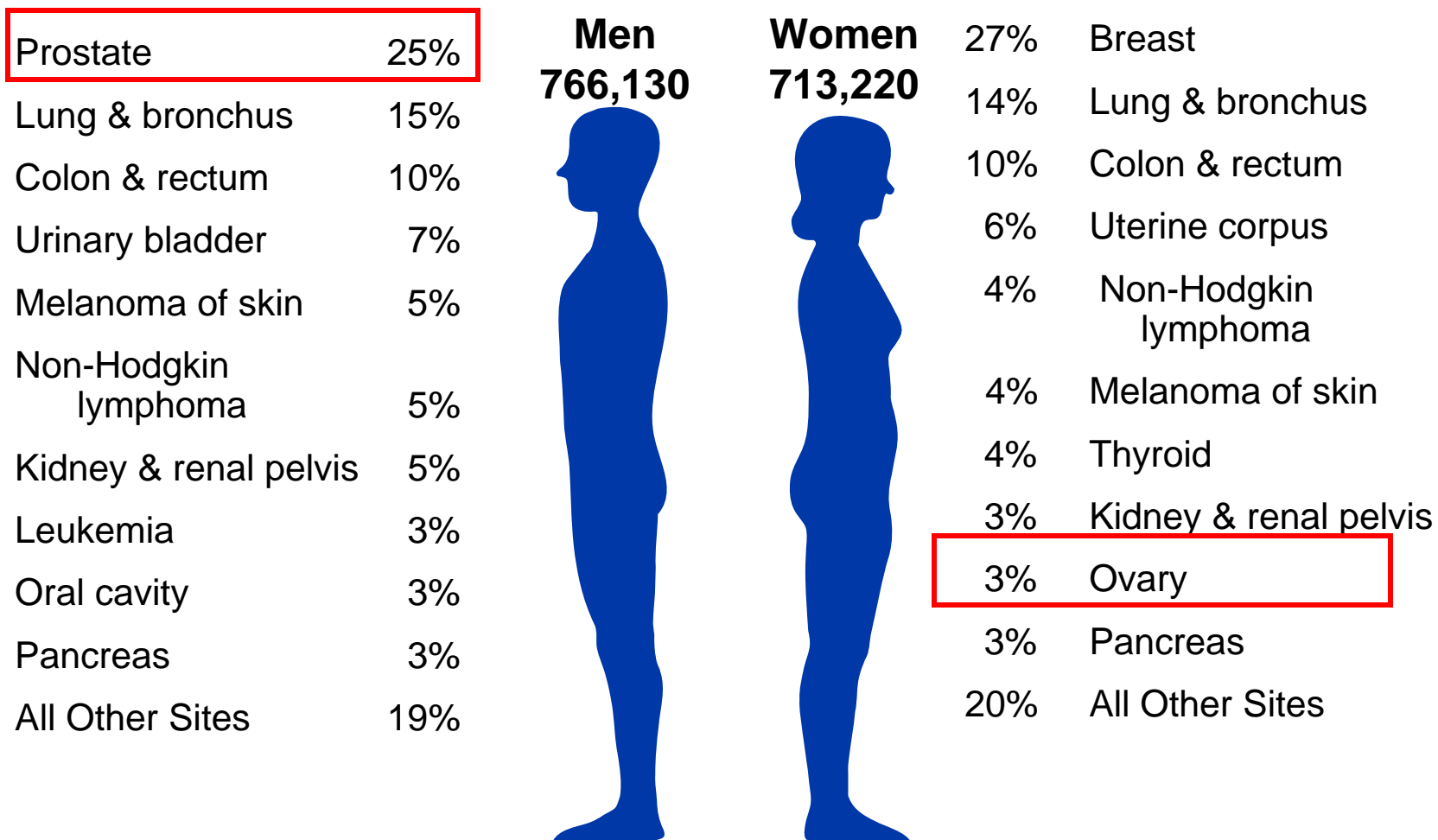
- \$172 billion
 - \$61 billion - direct medical costs
 - \$16 billion - lost productivity to illness
 - \$95 billion - lost productivity to premature death

U.S. Cancer Incidence & Mortality 2009

- New cases of cancer:
 - United States: 1,479,350
 - Texas: 98,200
- Deaths due to cancer:
 - United States: 562,340

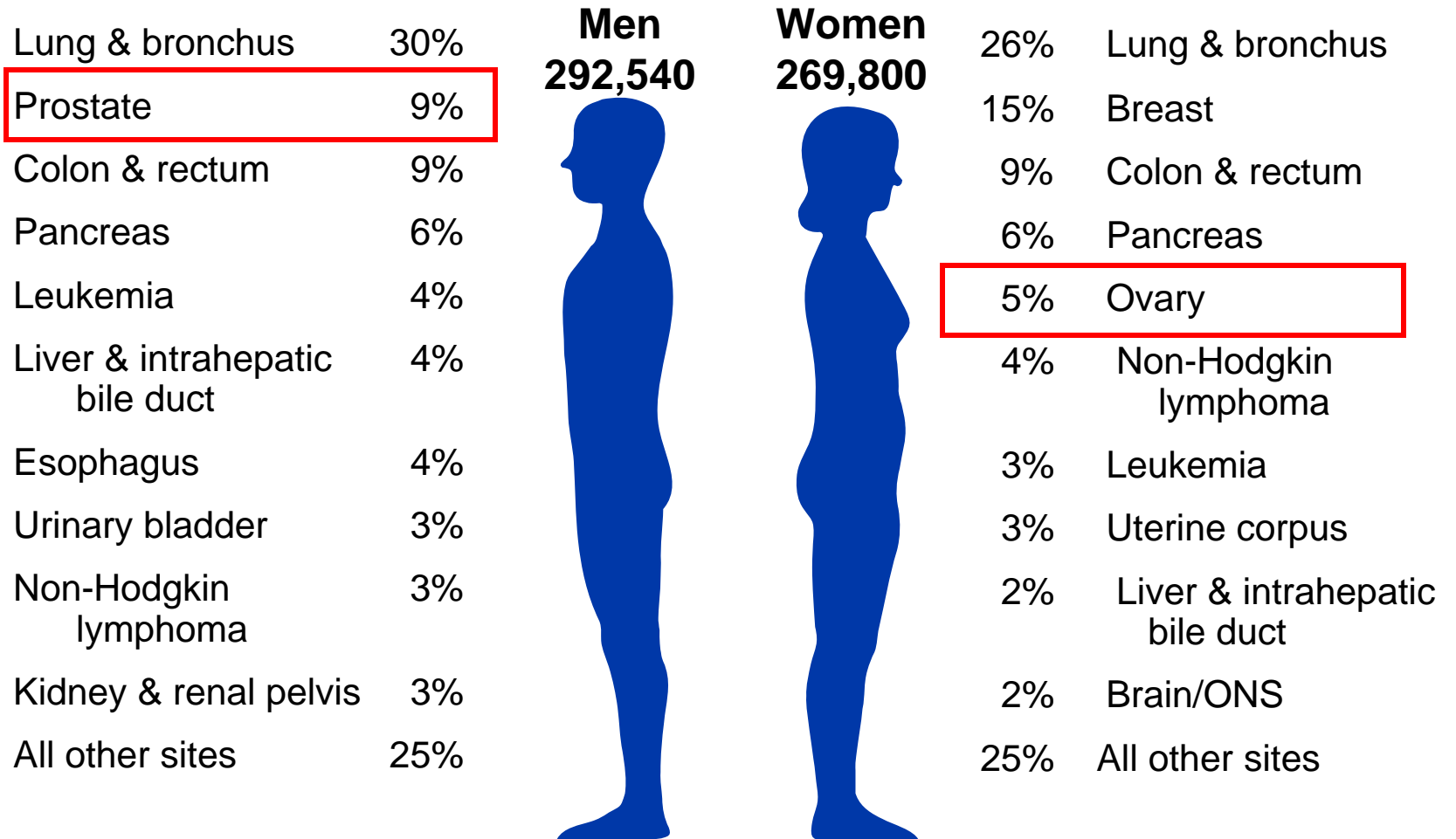
www.cancer.org, Cancer Facts & Figures

2009 Estimated US Cancer Cases*



*Excludes basal and squamous cell skin cancers and in situ carcinomas except urinary bladder.
 Source: American Cancer Society, 2009.

2009 Estimated US Cancer Deaths*



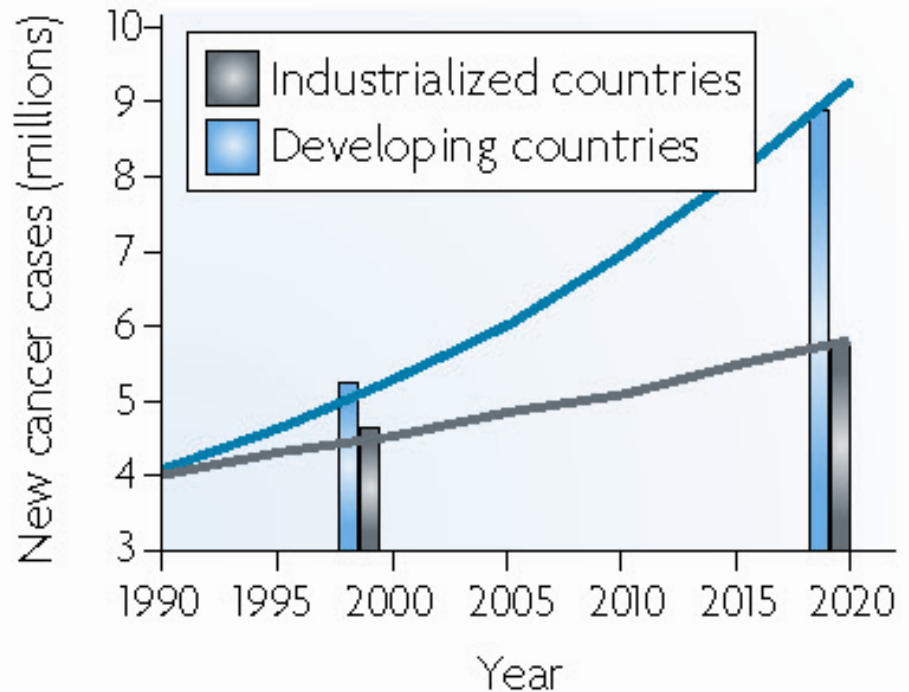
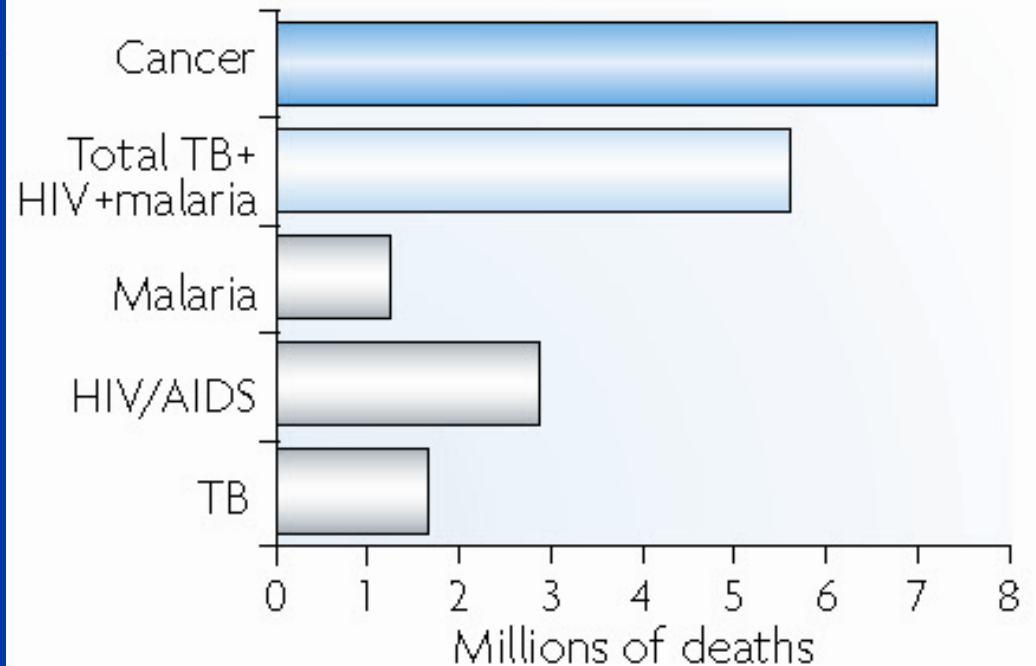
ONS=Other nervous system.
Source: American Cancer Society, 2009.

Worldwide Burden of Cancer

- Today:
 - 11 million new cases every year
 - 6.2 million deaths every year (12% of deaths)
- Can prevent 1/3 of these cases:
 - Reduce tobacco use
 - Implement existing screening techniques
 - Healthy lifestyle and diet
- In 2020:
 - 15 million new cases predicted in 2020
 - 10 million deaths predicted in 2020
 - Increase due to aging population
 - Increase in smoking

Global Cancer Trends

Lingwood, et al;
The challenge of cancer
control in Africa;
Nat Rev CA, 8:398, 2008.



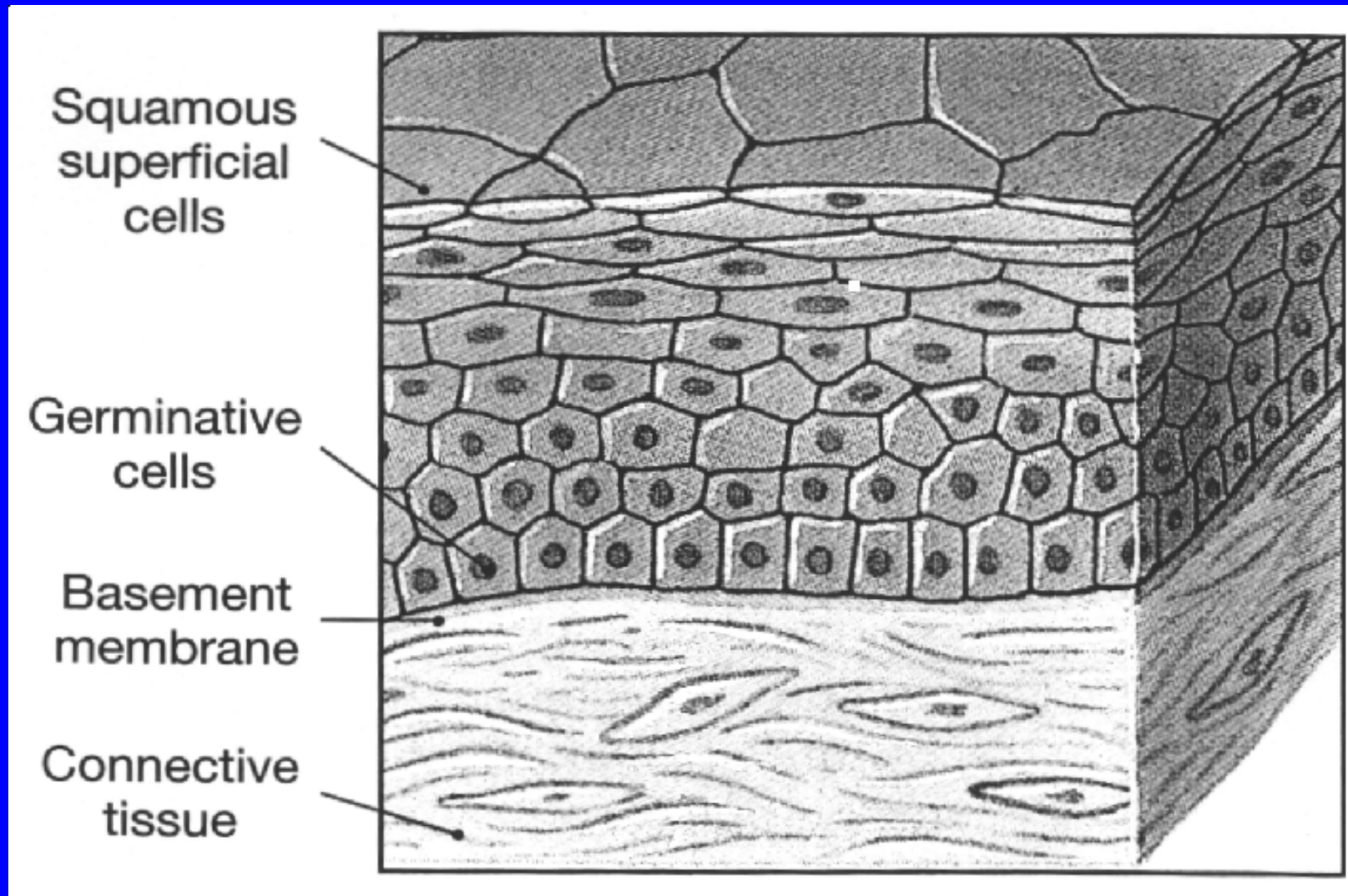
Worldwide Burden of Cancer

- 23% of cancers in developing countries caused by infectious agents
 - Hepatitis (liver)
 - HPV (cervix)
 - H. pylori (stomach)
- Vaccination could be key to preventing these cancers

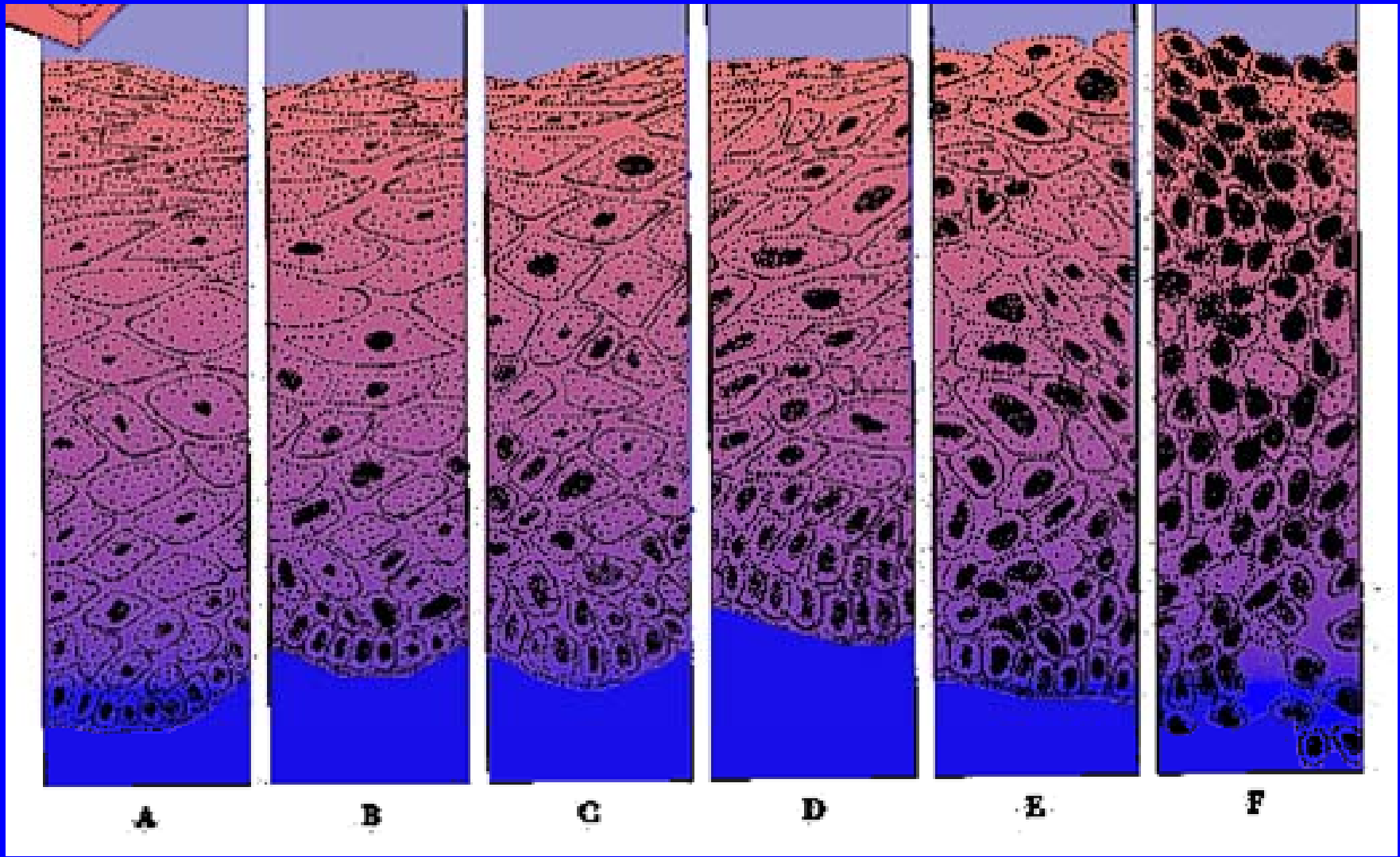
What is Cancer?

- Characterized by uncontrolled growth & spread of abnormal cells
- Can be caused by:
 - External factors:
 - Tobacco, chemicals, radiation, infectious organisms
 - Internal factors:
 - Mutations, hormones, immune conditions

Squamous Epithelial Tissue

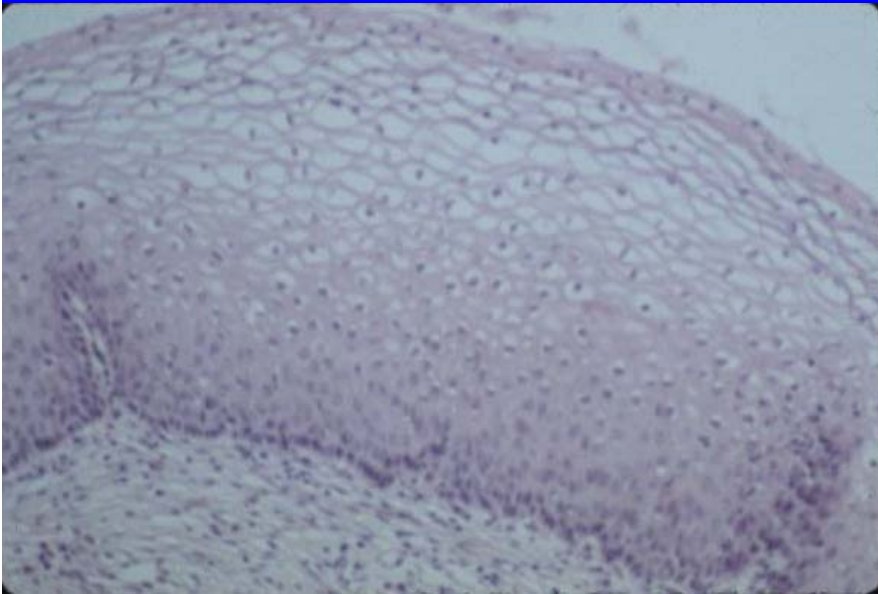


Precancer → Cancer Sequence

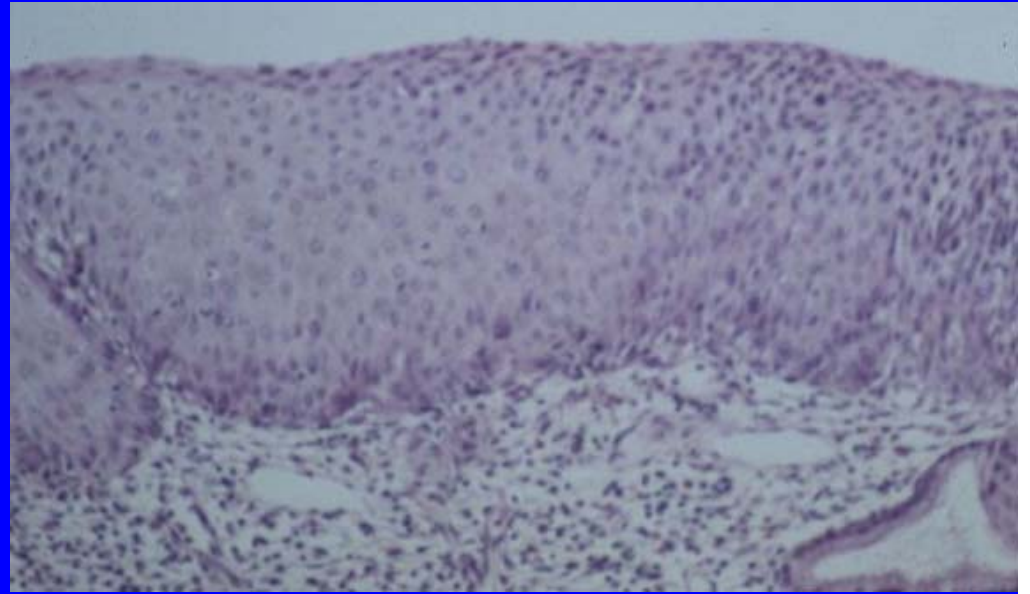


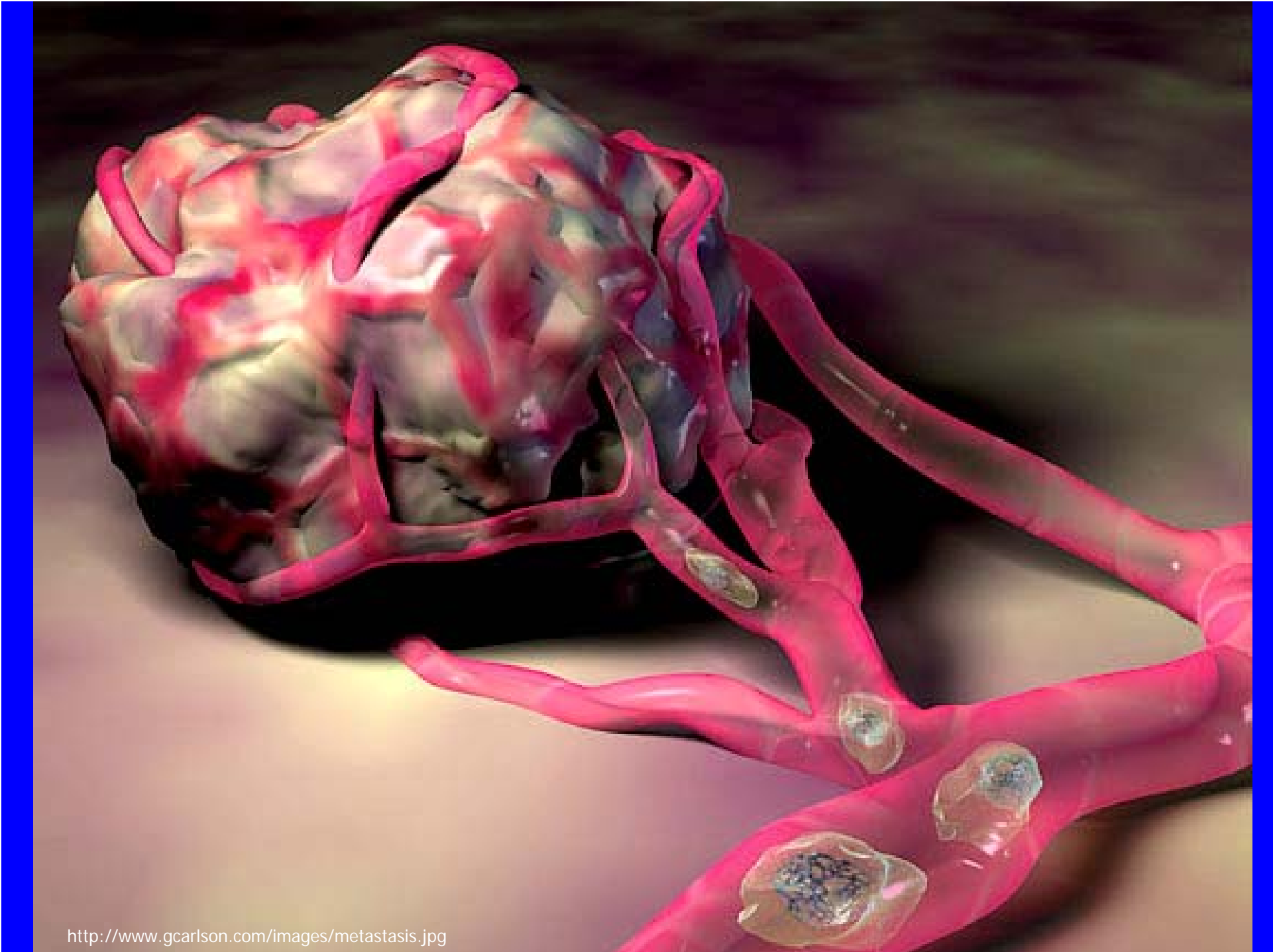
Histologic Images

Normal



Cervical Pre-Cancer





<http://www.gcarlson.com/images/metastasis.jpg>

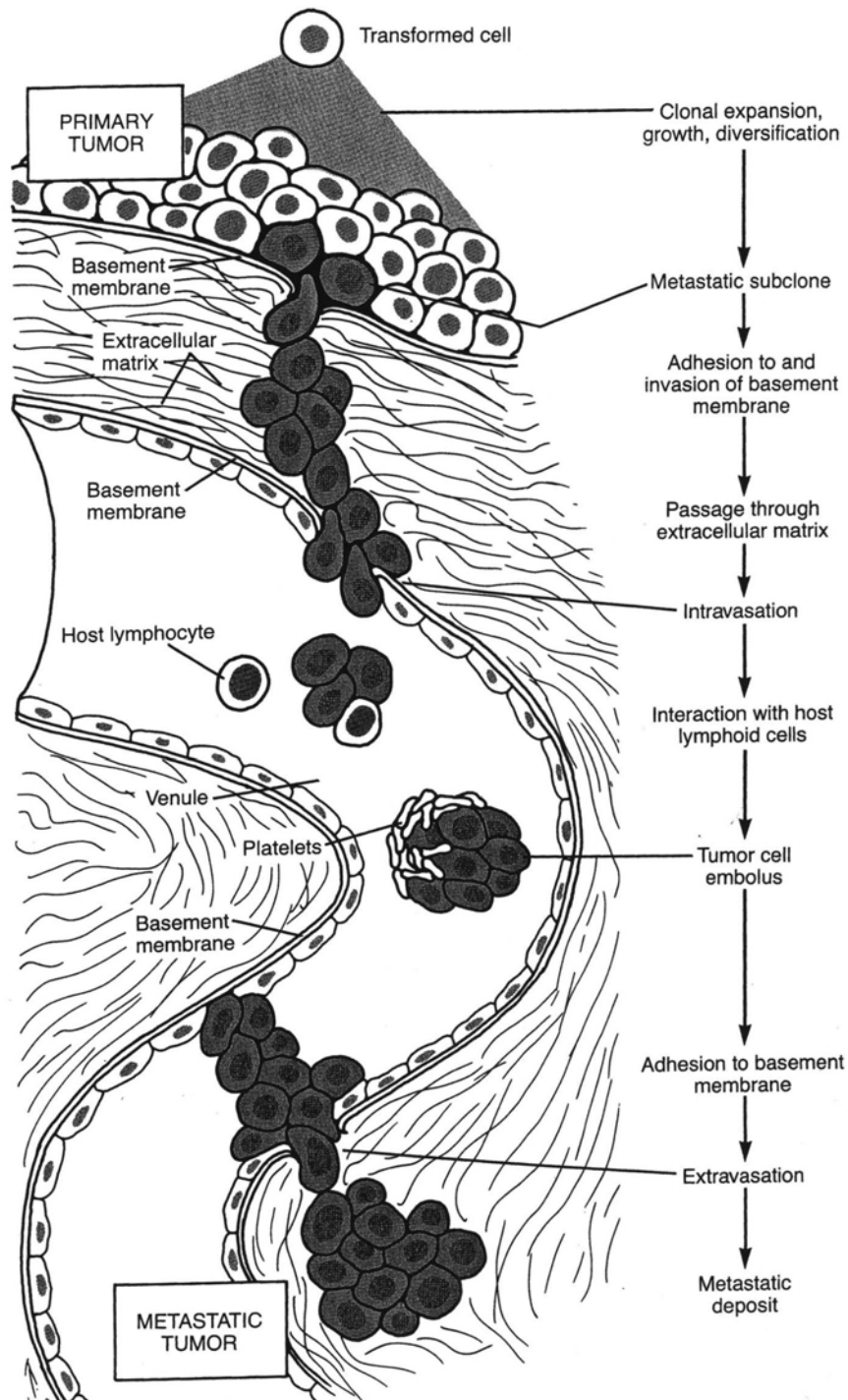
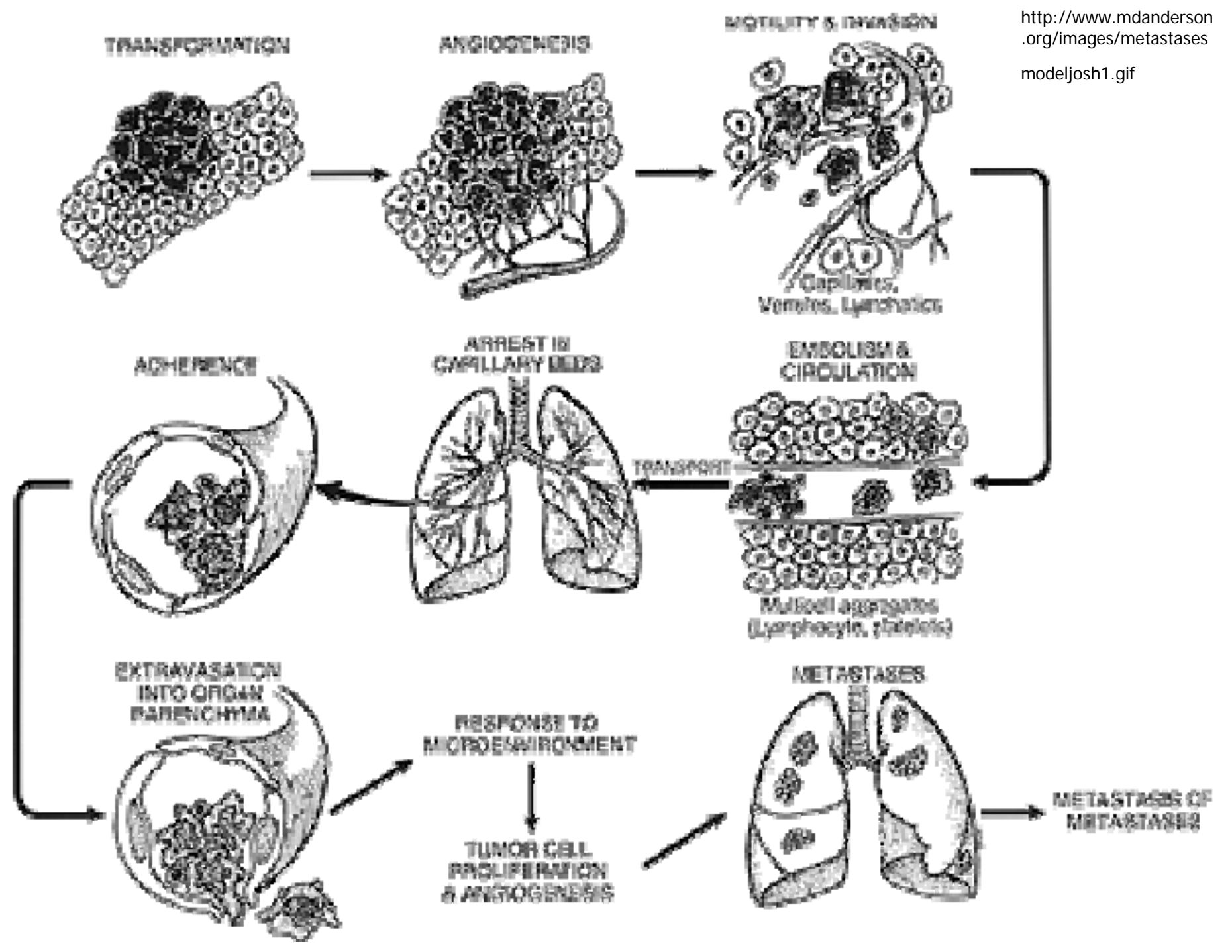


Fig 7.33 – The Metastatic cascade

Neoplasia

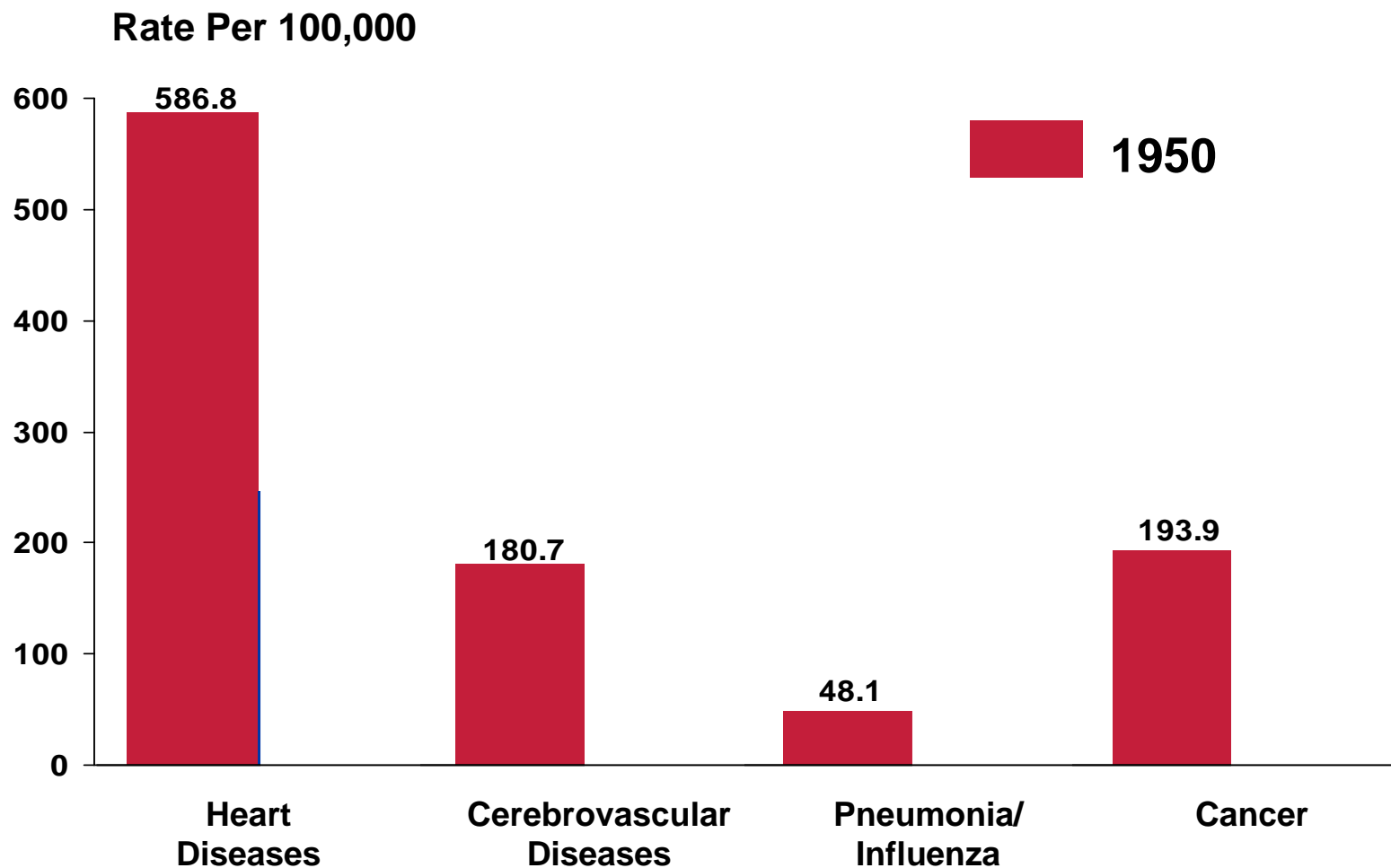


The War on Cancer

- 1971 State of Union address:
 - President Nixon requested \$100 million for cancer research
- December 23, 1971
 - Nixon signed National Cancer Act into law
 - "I hope in years ahead we will look back on this action today as the most significant action taken during my Administration."



Change in the US Death Rates* by Cause, 1950 & 2001



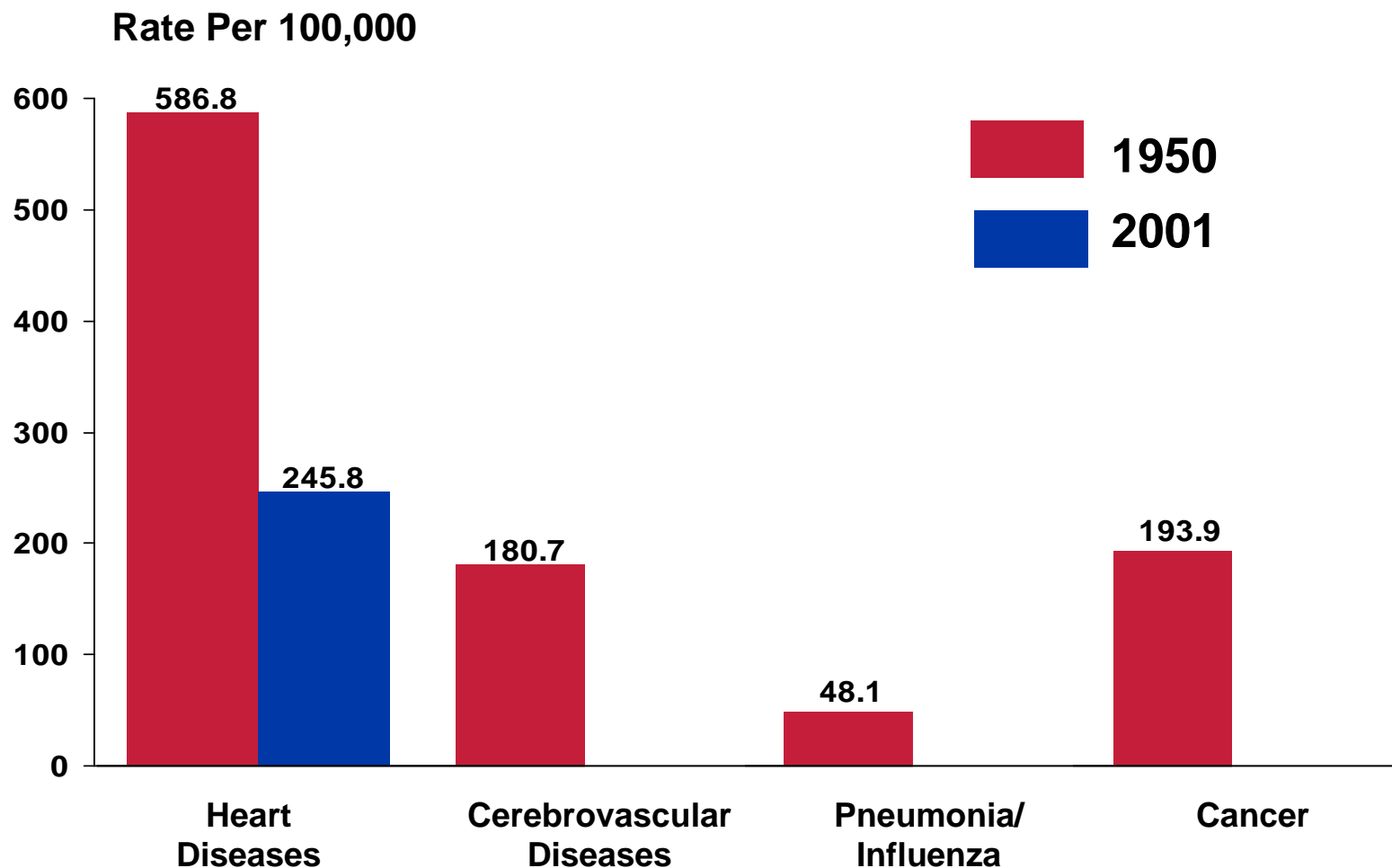
* Age-adjusted to 2000 US standard population.

Sources: 1950 Mortality Data - CDC/NCHS, NVSS, Mortality Revised.

2001 Mortality Data—NVSR-Death Final Data 2001—Volume 52, No. 3.

http://www.cdc.gov/nchs/data/nvsr/nvsr52/nvsr52_03.pdf

Change in the US Death Rates* by Cause, 1950 & 2001



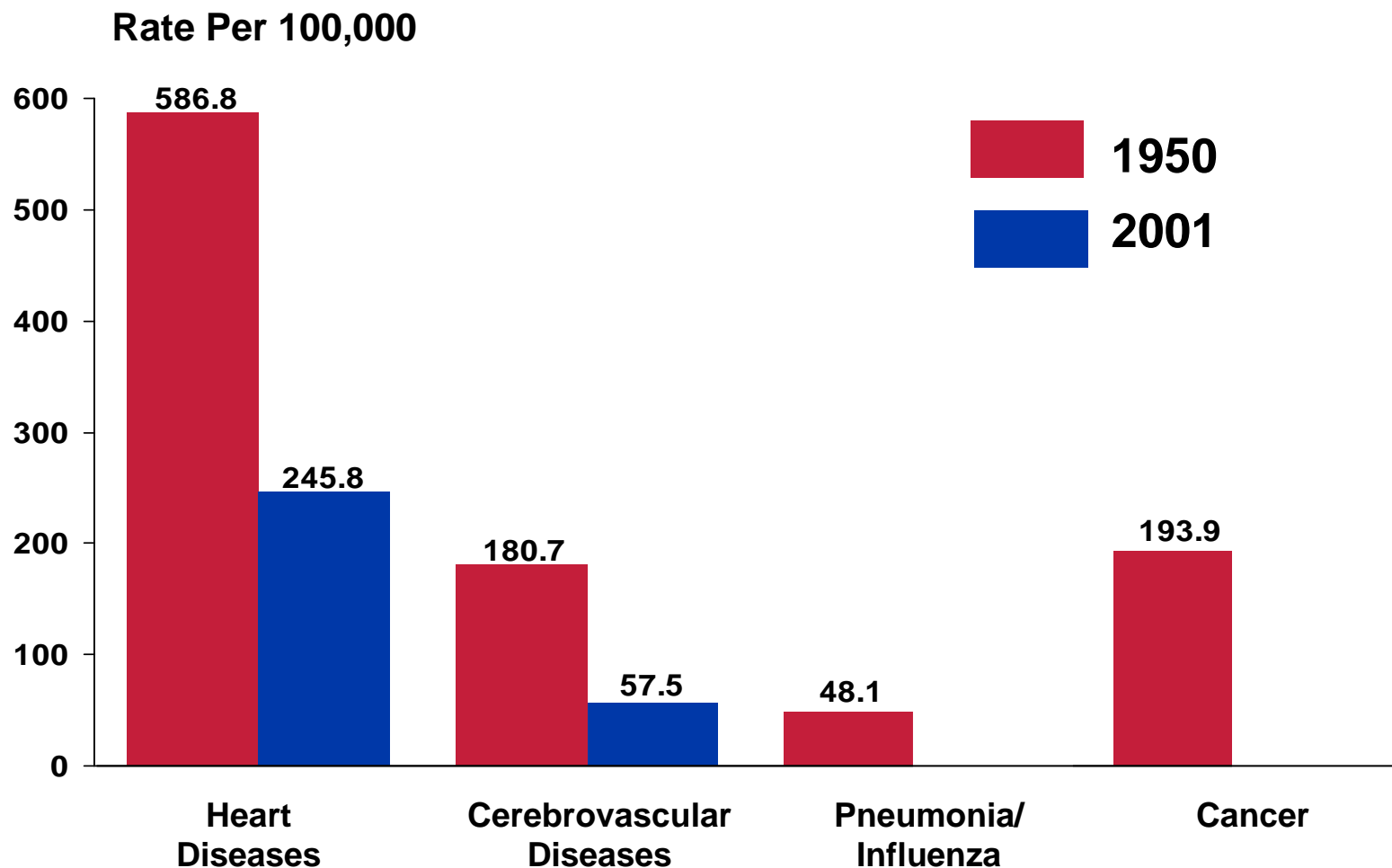
* Age-adjusted to 2000 US standard population.

Sources: 1950 Mortality Data - CDC/NCHS, NVSS, Mortality Revised.

2001 Mortality Data - NVSR-Death Final Data 2001 - Volume 52, No. 3.

http://www.cdc.gov/nchs/data/nvsr/nvsr52/nvsr52_03.pdf

Change in the US Death Rates* by Cause, 1950 & 2001



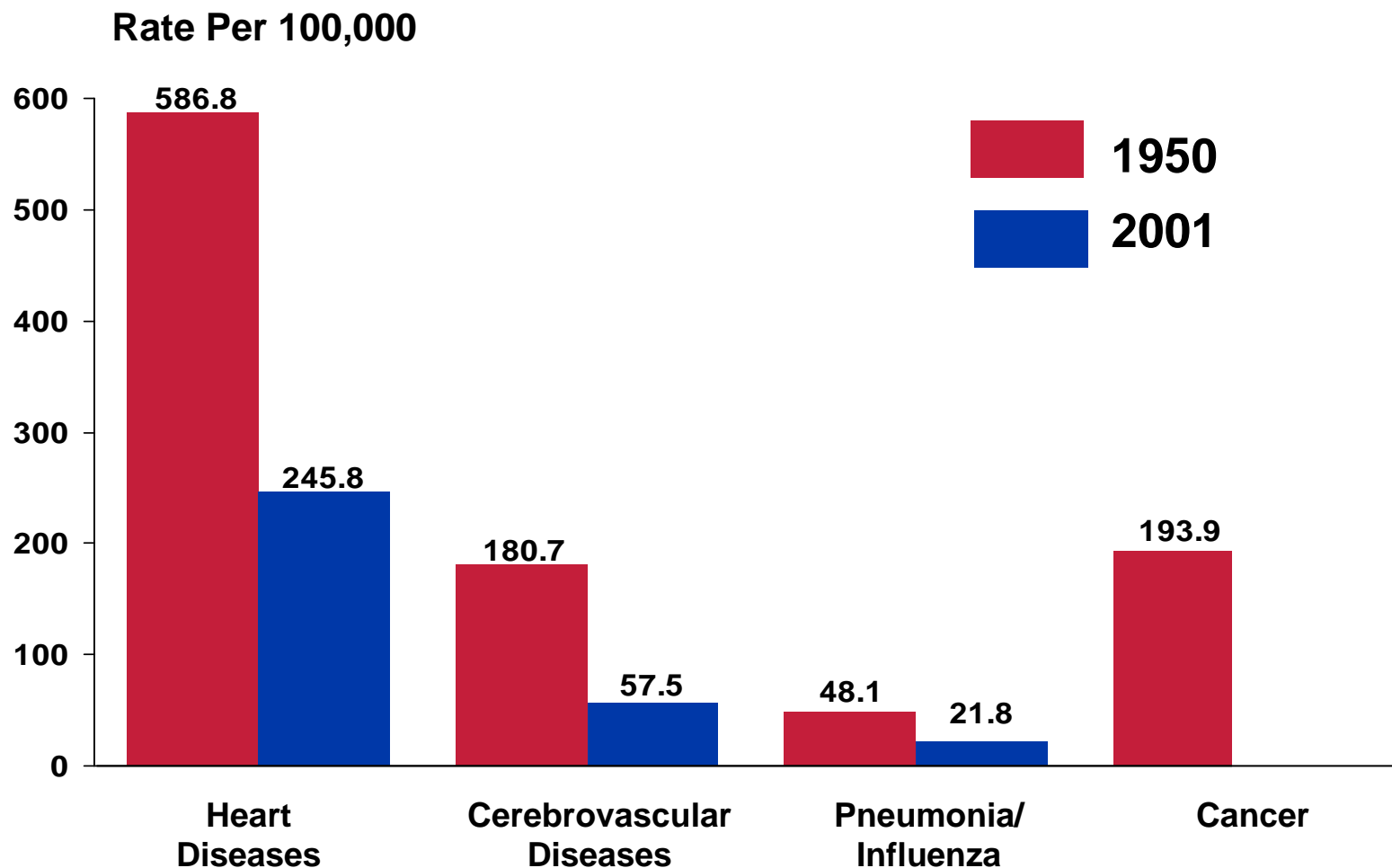
* Age-adjusted to 2000 US standard population.

Sources: 1950 Mortality Data - CDC/NCHS, NVSS, Mortality Revised.

2001 Mortality Data - NVSR-Death Final Data 2001 - Volume 52, No. 3.

http://www.cdc.gov/nchs/data/nvsr/nvsr52/nvsr52_03.pdf

Change in the US Death Rates* by Cause, 1950 & 2001



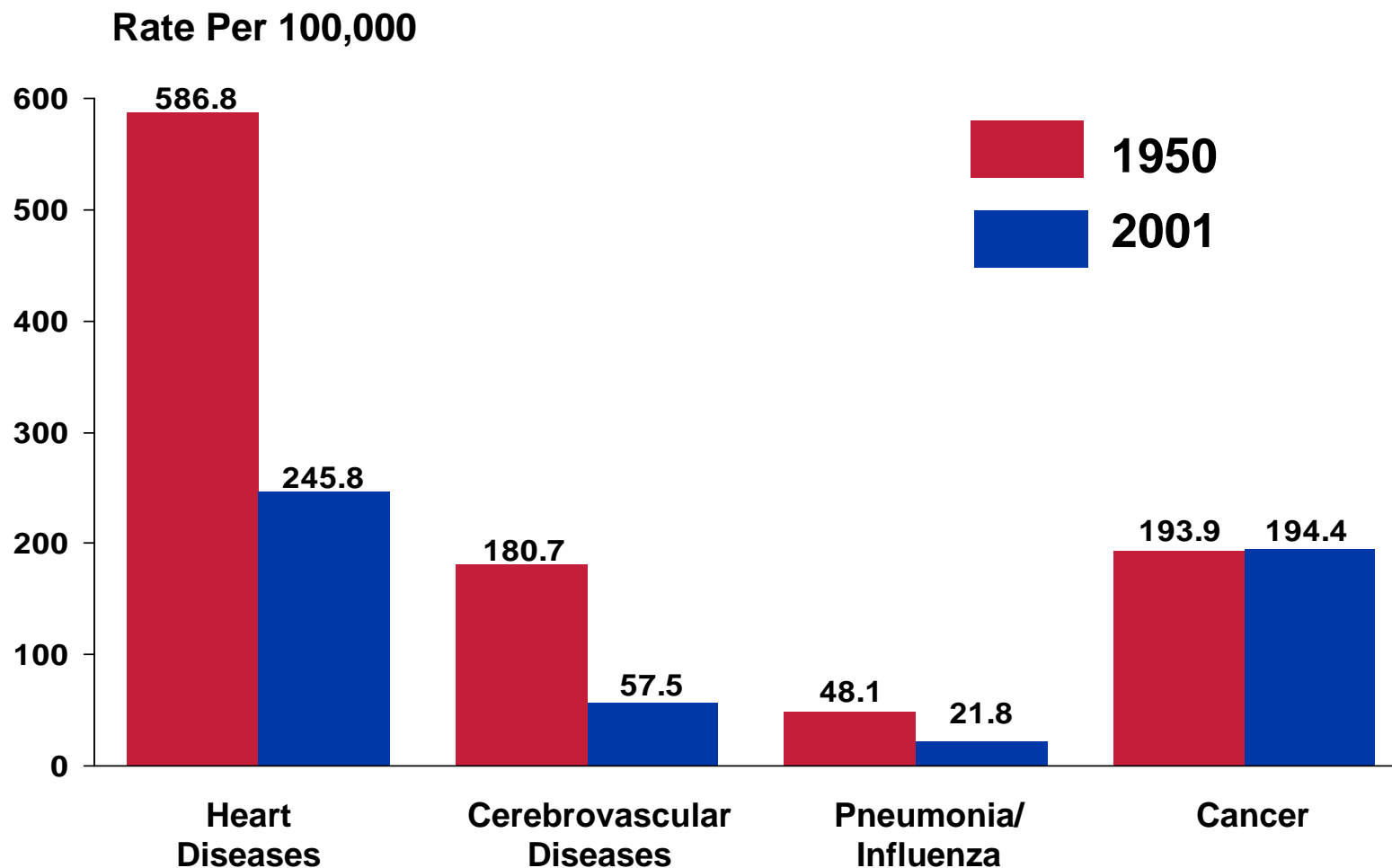
* Age-adjusted to 2000 US standard population.

Sources: 1950 Mortality Data - CDC/NCHS, NVSS, Mortality Revised.

2001 Mortality Data - NVSR-Death Final Data 2001 - Volume 52, No. 3.

http://www.cdc.gov/nchs/data/nvsr/nvsr52/nvsr52_03.pdf

Change in the US Death Rates* by Cause, 1950 & 2001



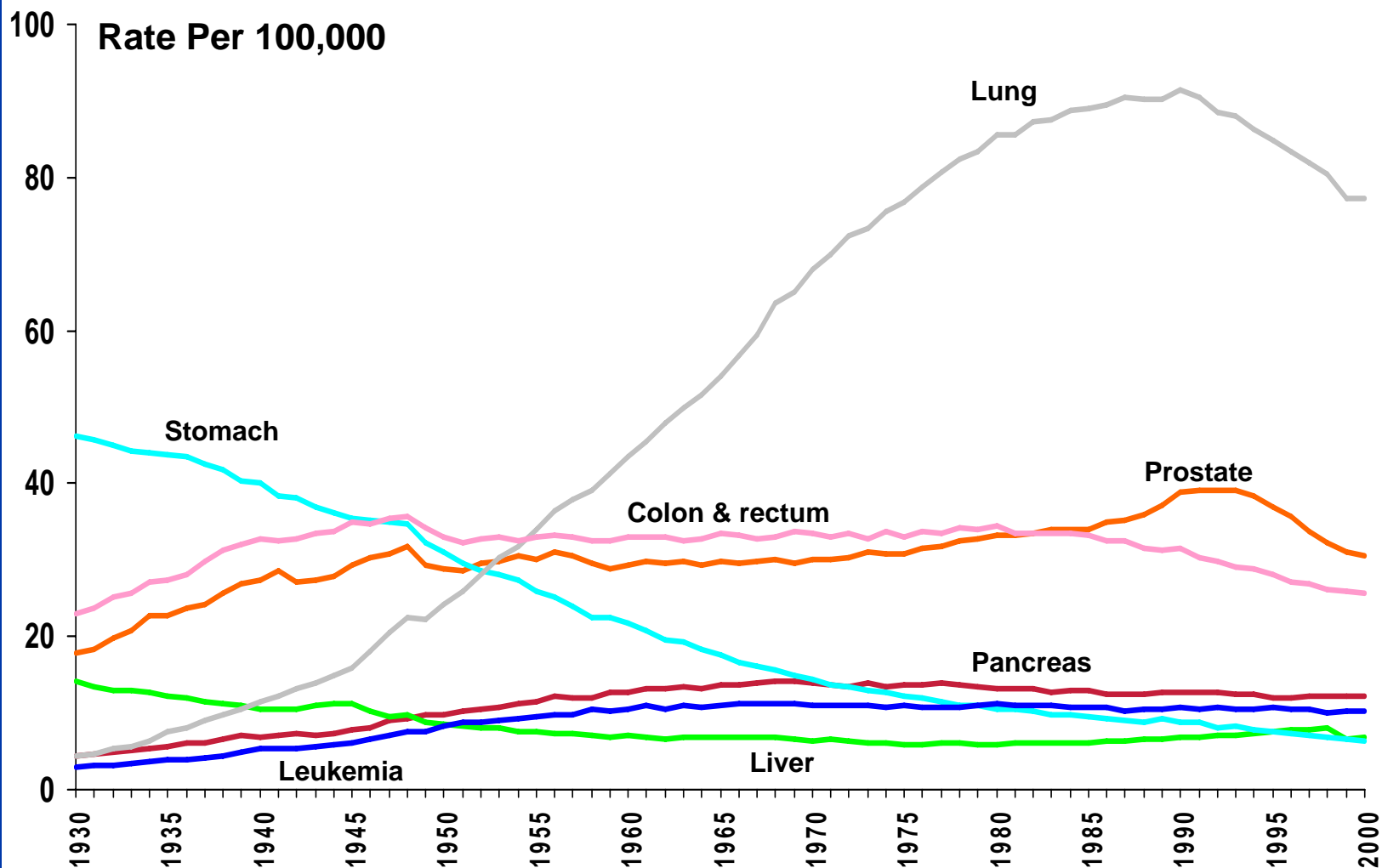
* Age-adjusted to 2000 US standard population.

Sources: 1950 Mortality Data - CDC/NCHS, NVSS, Mortality Revised.

2001 Mortality Data - NVSR-Death Final Data 2001 - Volume 52, No. 3.

http://www.cdc.gov/nchs/data/nvsr/nvsr52/nvsr52_03.pdf

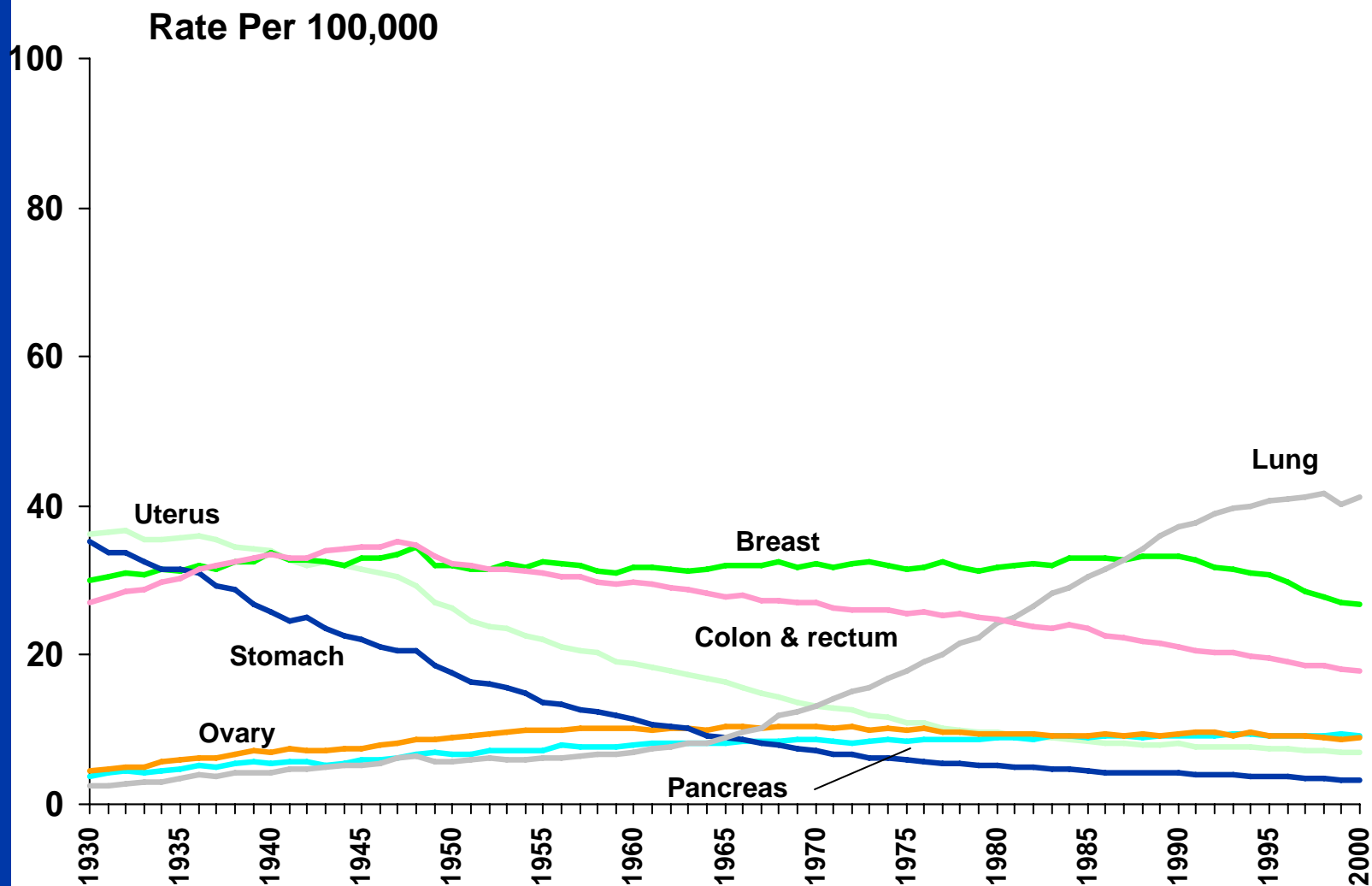
Cancer Death Rates*, for Men, US, 1930-2000



*Age-adjusted to the 2000 US standard population.

Source: US Mortality Public Use Data Tapes 1960-2000, US Mortality Volumes 1930-1959, National Center for Health Statistics, Centers for Disease Control and Prevention, 2003.

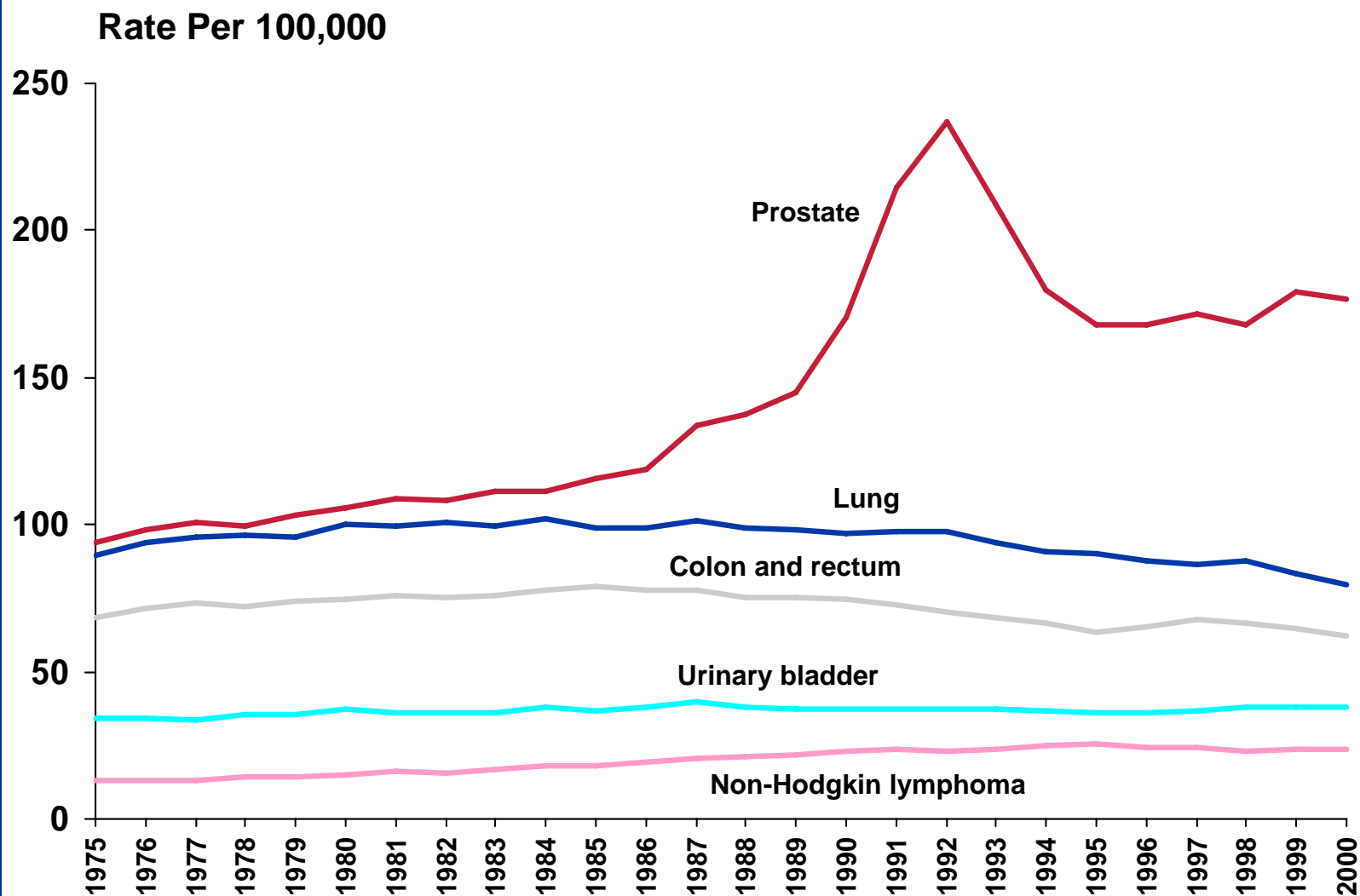
Cancer Death Rates*, for Women, US, 1930-2000



*Age-adjusted to the 2000 US standard population.

Source: US Mortality Public Use Data Tapes 1960-2000, US Mortality Volumes 1930-1959, National Center for Health Statistics, Centers for Disease Control and Prevention, 2003.

Cancer Incidence Rates* for Men, US, 1975-2000



*Age-adjusted to the 2000 US standard population.

Source: Surveillance, Epidemiology, and End Results Program, 1975-2000, Division of Cancer Control and Population Sciences, National Cancer Institute, 2003.

Relative Survival* (%) during Three Time Periods by Cancer Site

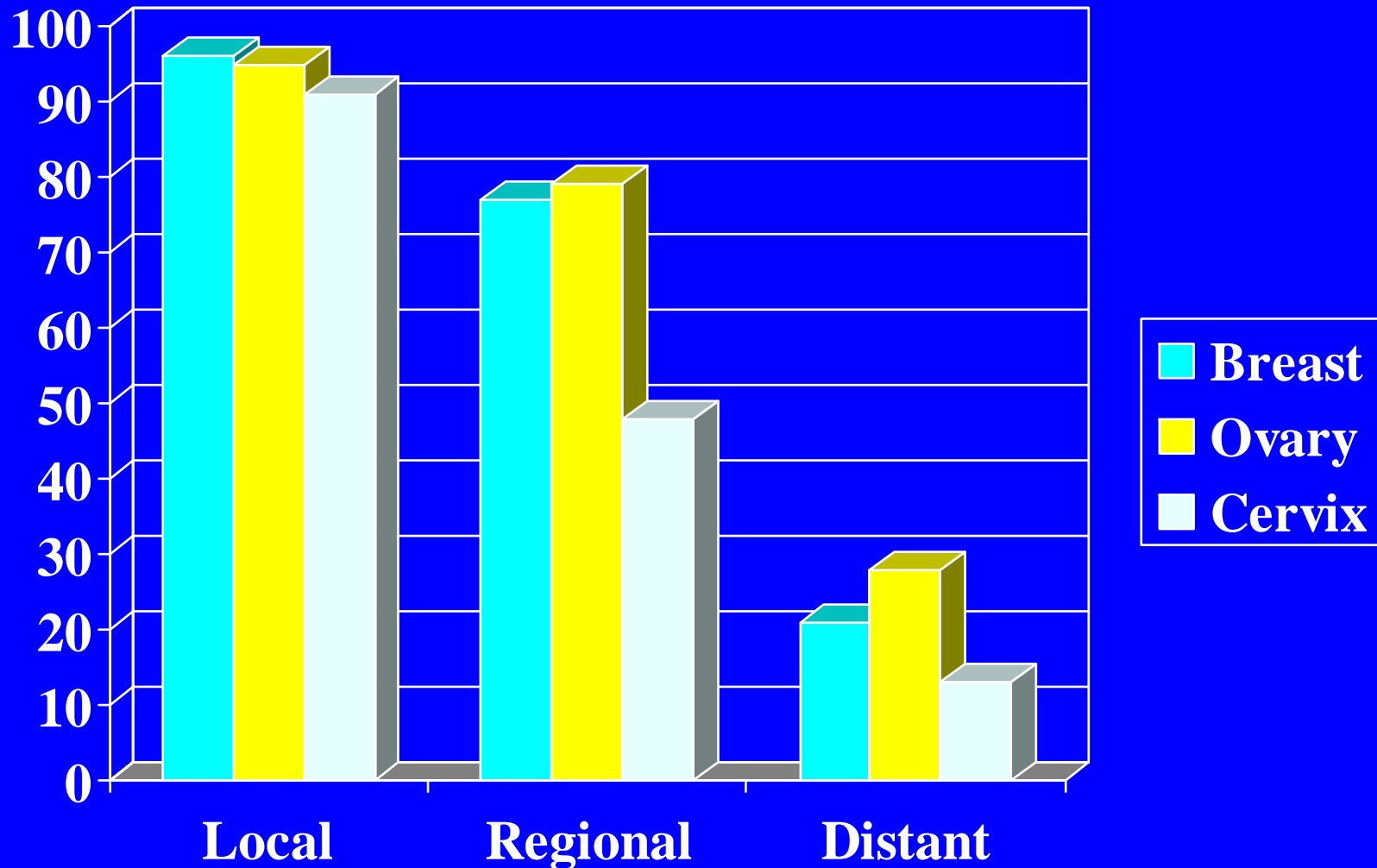
Site	1974-1976	1983-1985	1992-1999
All sites	50	52	63
Breast (female)	75	78	87
Colon & rectum	50	57	62
Leukemia	34	41	46
Lung & bronchus	12	14	15
Melanoma	80	85	90
Non-Hodgkin lymphoma	47	54	56
Ovary	37	41	53
Pancreas	3	3	4
Prostate	67	75	98
Urinary bladder	73	78	82

*5-year relative survival rates based on follow up of patients through 2000.

Source: Surveillance, Epidemiology, and End Results Program, 1975-2000, Division of Cancer Control and Population Sciences, National Cancer Institute, 2003.

Importance of Early Detection

Five Year Relative Survival Rates



Screening

- Use of simple tests in a healthy population
- Goal:
 - Identify individuals who have disease, but do not yet have symptoms
- Should be undertaken only when:
 - Effectiveness has been demonstrated
 - Resources are sufficient to cover target group
 - Facilities exist for confirming diagnoses
 - Facilities exist for treatment and follow-up
 - When disease prevalence is high enough to justify effort and costs of screening

Cancer Screening

- We routinely screen for 4 cancers:
 - Female breast cancer
 - Mammography
 - Cervical cancer
 - Pap smear
 - Prostate cancer
 - Serum PSA
 - Digital rectal examination
 - Colon and rectal cancer
 - Fecal occult blood
 - Flexible sigmoidoscopy, Colonoscopy

Screening Guidelines for the Early Detection of Breast Cancer, American Cancer Society 2009

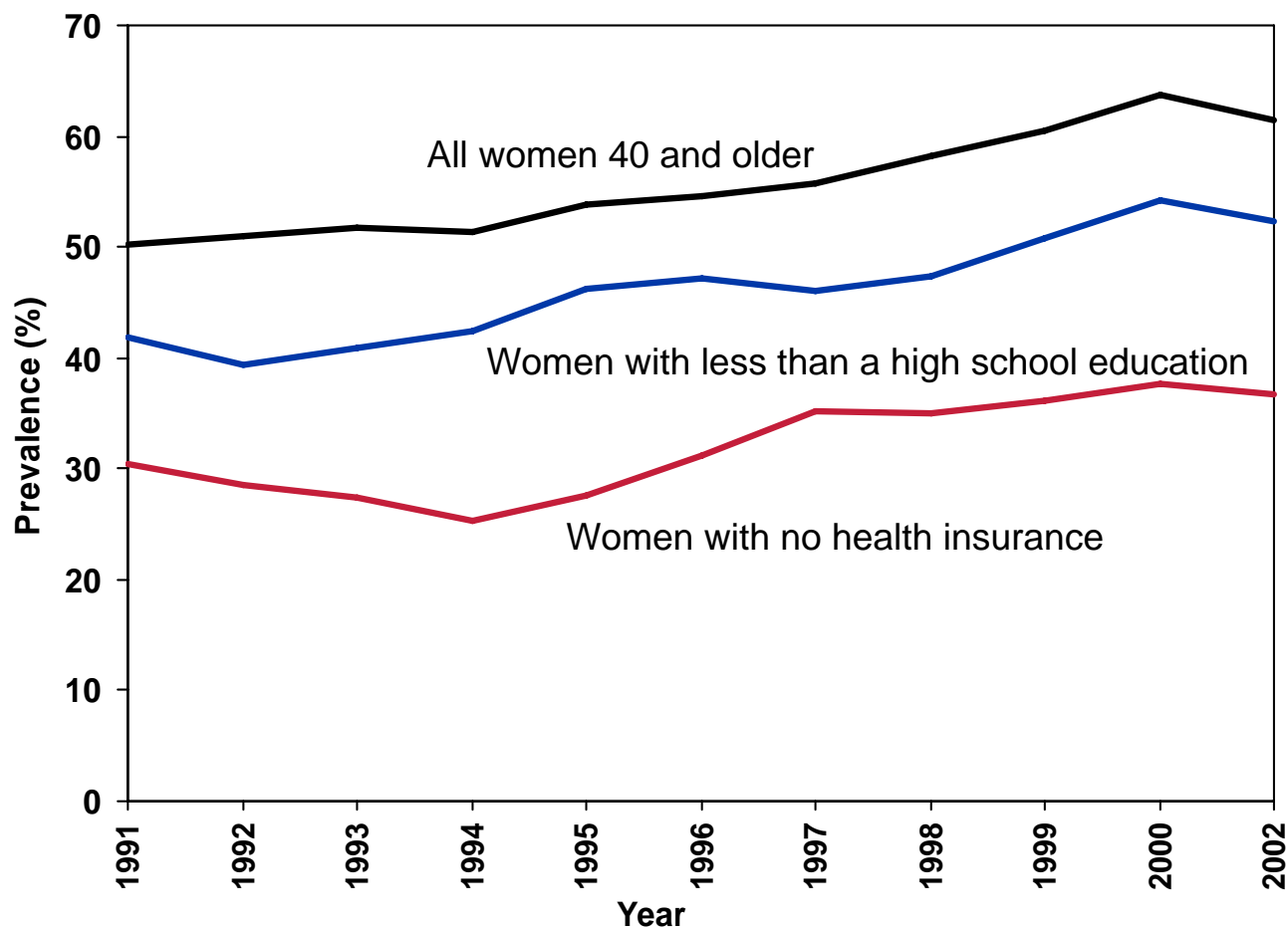
Yearly mammograms are recommended starting at age 40.

A clinical breast exam should be part of a periodic health examination, about every 3 years for women in their 20s and 30s. Asymptomatic women aged 40 and older should continue to undergo a clinical breast exam, preferably annually*.

Beginning in their early 20s, women should be told about the benefits and limitations of breast-self examination. Women should know how their breasts normally feel and report any breast changes promptly to their health care providers.

* Beginning at age 40 years, annual CBE should be performed prior to mammography.

Mammogram Prevalence (%), by Educational Attainment and Health Insurance Status, Women 40 and Older, US, 1991-2002



* A mammogram within the past year. Note: Data from participating states and the District of Columbia were aggregated to represent the United States.

Source: Behavior Risk Factor Surveillance System CD-ROM (1984-1995, 1996-1997, 1998, 1999) and Public Use Data Tape (2000, 2002), National Centers for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention 1997, 1999, 2000, 2000, 2001, 2003.

How do we judge efficacy
of a screening test?

Sensitivity/Specificity

Positive/Negative Predictive Value

Sensitivity & Specificity

■ Sensitivity

- Probability that given DISEASE, patient tests POSITIVE
- Ability to correctly detect disease
- 100% - False Negative Rate

■ Specificity

- Probability that given NO DISEASE, patient tests NEGATIVE
- Ability to avoid calling normal things disease
- 100% - False Positive Rate

Possible Test Results

	Test Positive	Test Negative	
Disease Present	TP	FN	# with Disease = TP+FN
Disease Absent	FP	TN	#without Disease = FP+TN
	# Test Pos = TP+FP	# Test Neg = FN+TN	Total Tested = TP+FN+FP+TN

$$Se = TP / (\# \text{ with disease}) = TP / (TP + FN)$$

$$Sp = TN / (\# \text{ without disease}) = TN / (TN + FP)$$

Example

- **Sputum microscopy:**
 - Procedure to detect lung cancer
- **Efficacy:**
 - 1,000 40-year-olds given the test
 - 28 people later proven to have lung cancer
 - 32 test positive, and of those 25 were truly positive
- **Calculate:**
 - Sensitivity & Specificity

Possible Test Results

	Test Positive	Test Negative	
Disease Present	25	3	# with Disease = 28
Disease Absent	7	965	#without Disease = 972
	# Test Pos = 32	# Test Neg = 968	Total Tested = 1,000

$$Se = 25/28 = 89\% \quad Sp = 965/972 = 99.3\%$$

As a patient:

What Information Do
You Want?

Predictive Value

■ Positive Predictive Value

- Probability that given a POSITIVE test result, you have DISEASE
- Ranges from 0-100%

■ Negative Predictive Value

- Probability that given a NEGATIVE test result, you do NOT HAVE DISEASE
- Ranges from 0-100%

■ Depends on the prevalence of the disease

Possible Test Results

	Test Positive	Test Negative	
Disease Present	TP	FN	# with Disease = TP+FN
Disease Absent	FP	TN	#without Disease = FP+TN
	# Test Pos = TP+FP	# Test Neg = FN+TN	Total Tested = TP+FN+FP+TN

$$PPV = TP / (\# \text{ Test Pos}) = TP / (TP + FP)$$

$$NPV = TN / (\# \text{ Test Neg}) = TN / (FN + TN)$$

Example

- **Sputum Microscopy:**
 - Procedure to detect lung cancer
- **Efficacy:**
 - 1,000 40-year-olds given the test
 - 28 people later shown to have lung cancer
 - 32 test positive, and of those 25 were truly positive
- **Calculate:**
 - Positive & Negative Predictive Value

Possible Test Results

	Test Positive	Test Negative	
Disease Present	25	3	# with Disease = 28
Disease Absent	7	965	#without Disease = 972
	# Test Pos = 32	# Test Neg = 968	Total Tested = 1,000

$$Se = 25/28 = 89\% \quad Sp = 965/972 = 99.3\%$$

$$PPV = 25/32 = 78\% \quad NPV = 965/968 = 99.7\%$$

Dependence on Prevalence

- Prevalence – is a disease common or rare?
 - $p = (\# \text{ with disease})/\text{total } \#$
 - $p = (TP+FN)/(TP+FP+TN+FN)$
- Does our test accuracy depend on p ?
 - Se/Sp do not depend on prevalence
 - PPV/NPV are highly dependent on prevalence
- $PPV = pSe/[pSe + (1-p)(1-Sp)]$
- $NPV = (1-p)Sp/[(1-p)Sp + p(1-Se)]$

Is it Hard to Screen for Rare Disease?

- **Sputum Microscopy:**
 - Procedure to detect lung cancer
- **Efficacy:**
 - 1,000 40-year-olds given the test
 - 28 people later shown to have lung cancer
 - 32 test positive, and of those 25 were truly positive
- **Calculate:**
 - Prevalence of lung cancer

Is it Hard to Screen for Rare Disease?

■ Sputum Microscopy:

- Usually offered to older smokers

■ Efficacy:

- 1,000 20-year-olds given the test
- Prevalence of lung cancer is expected to be 2.8/1000

■ Calculate:

- Sensitivity & Specificity
- Positive & Negative Predictive Value
- Suppose a 20 yo has a positive test. What is the likelihood that they have lung cancer?

Possible Test Results

	Test Positive	Test Negative	
Disease Present	2.5	.3	# with Disease = 2.8
Disease Absent	6.98	990.2	#without Disease = 997.2
	# Test Pos = 9.48	# Test Neg = 990.5	Total Tested = 1,000

$$Se = 2.5/2.8 = 89.3\% \quad Sp = 990.2/997.2 = 99.3\%$$

$$PPV = 2.5/9.48 = 26.3\% \quad NPV = 990.2/990.5 = 99.97\%$$

Cervical Cancer

Early Detection

Cervical Cancer: 2004

- 10,520 new cases in US
- 3,900 deaths in US
- Signs and symptoms:
 - Abnormal vaginal bleeding
- Risk Factors:
 - Failure to obtain regular Pap smears
 - HPV infection
 - Sex at an early age
 - Multiple sexual partners
 - Cigarette smoking

Cervical Cancer: World

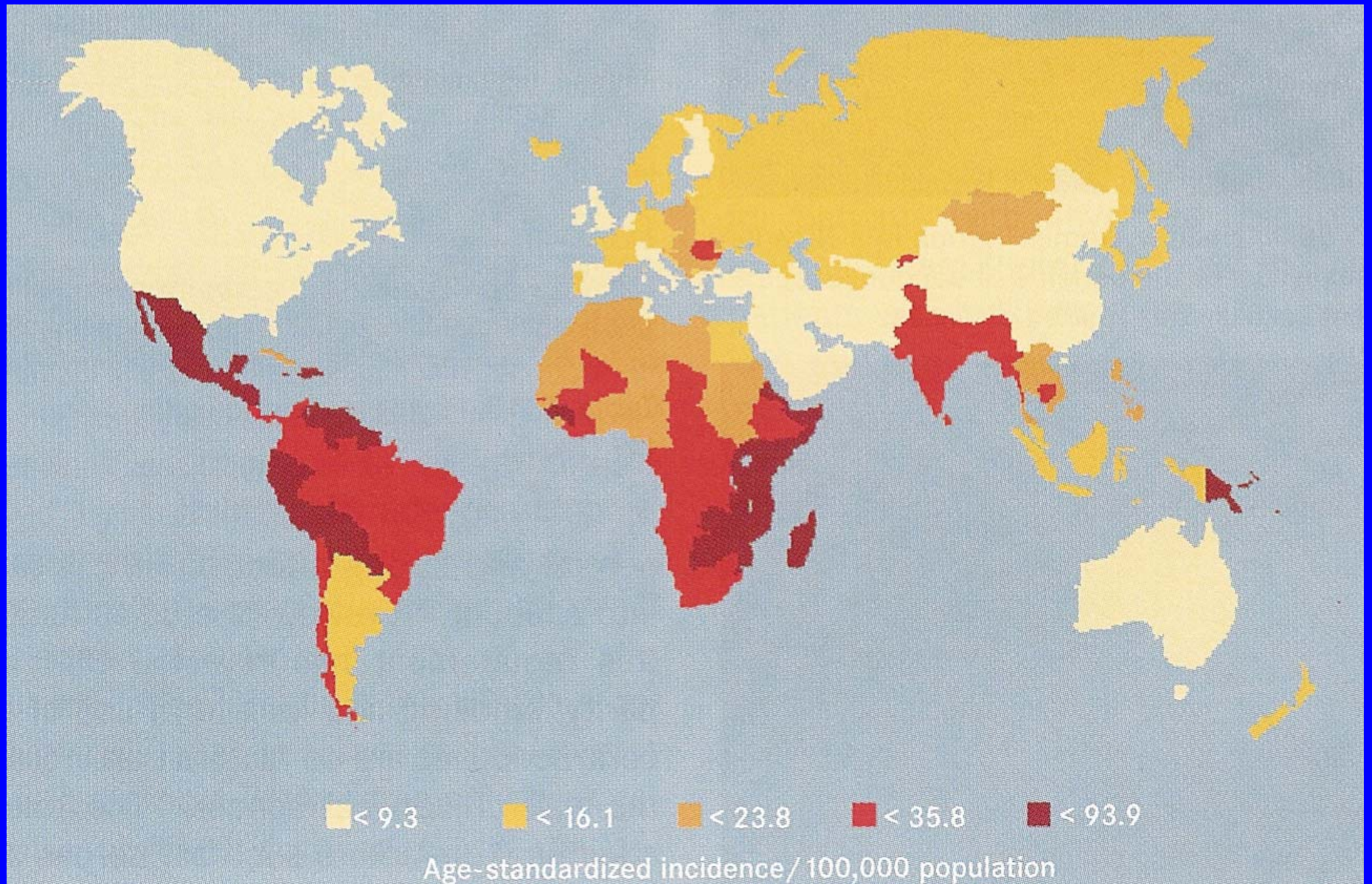
■ Incidence:

- 510,000 new cases per year worldwide
- 80% of cases occur in the developing world
- Highest incidence in:
 - Central and South America
 - Southern Africa
 - Asia

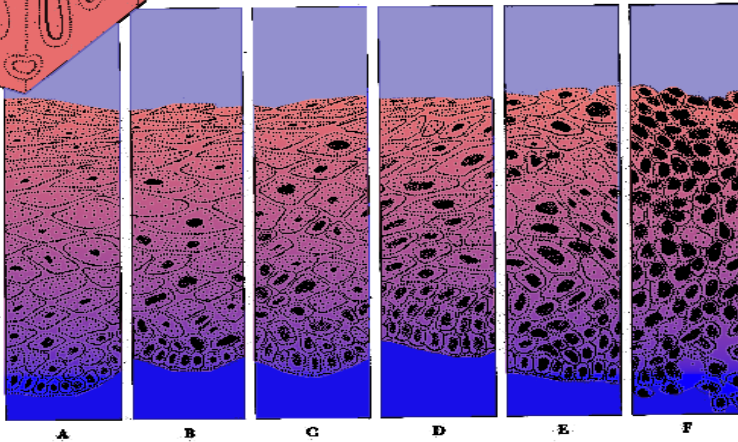
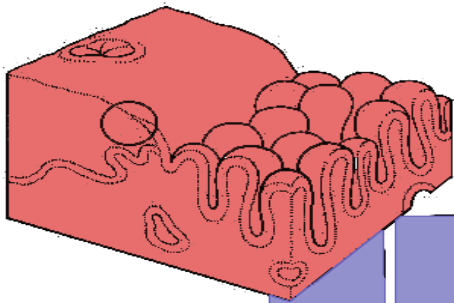
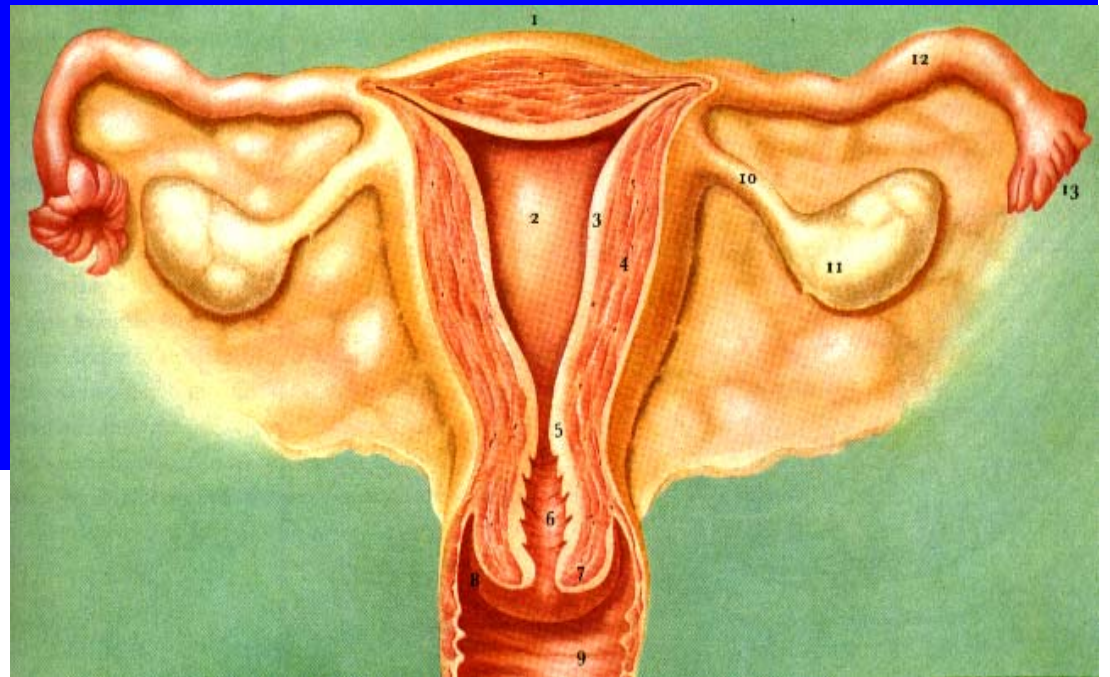
■ Mortality:

- 288,000 deaths per year worldwide
- 2nd leading cause of female cancer mortality worldwide

Global Burden of Cervical Cancer

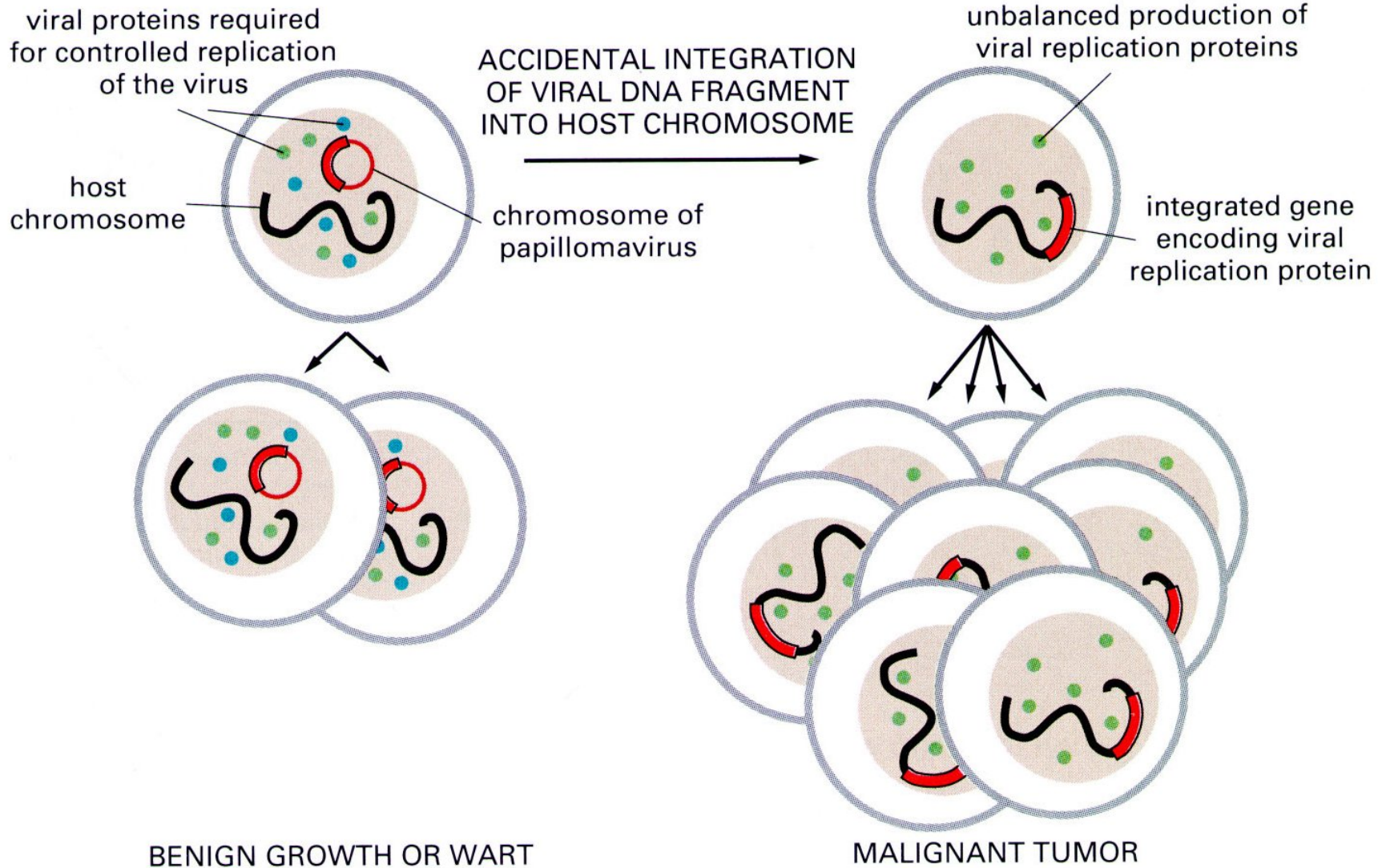


Cervical Cancer



What Initiates Transformation?

- Infection with Human Papilloma Virus (HPV)
 - Most common sexually transmitted disease
 - Asymptomatic HPV infections can be detected in 5-40% of women of reproductive age
- HPV infection is the central causative factor in squamous cell carcinoma of the cervix
 - HPV infections are transient; most young women clear them with no ill effects
 - If HPV infection persists past age 30, there is greater risk of developing cervical cancer
 - Many viral subtypes (70)
 - 13 most commonly linked to cervical cancer
 - HPV 16, 18



In a wart or benign infection, the HPV chromosomes are stably maintained in the basal epithelium as plasmids (left). Integration of viral DNA into a host chromosome alters the environment of the viral genes and disrupts control of their expression. Unregulated reproduction of viral proteins tends to drive the host cell into S phase helping to generate a cancer (right).

How Do We Detect Early Cervical Cancer?

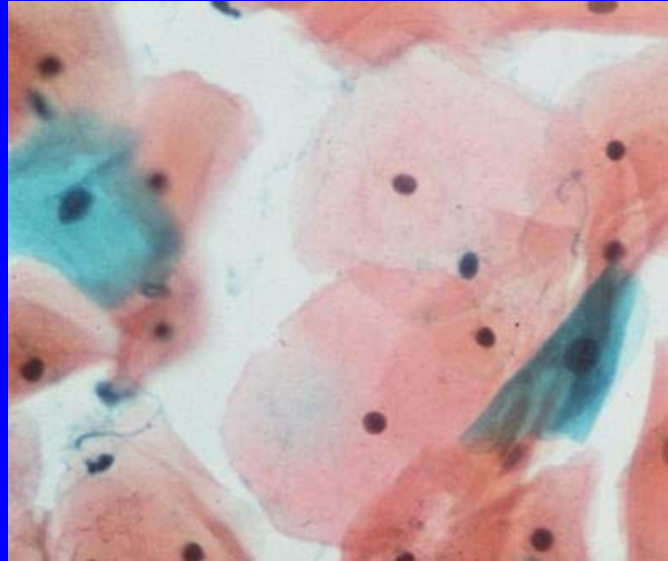
Pap Smear

(The most successful cancer-screening test in medical history)

Colposcopy + Biopsy

Screening Pap Smear

<http://www.geocities.com/HotSprings/Sauna/2329/image3.jpg>



- Each slide: 50,000-300,000 cells
- Cytotechnologists review < 100 slides per day
- 10% of "normal" slides re-screened
- Se = 62%
- Sp = 78%
- False negative smears account for 3% of U.S. Cervical Cancer cases/year

<http://www.gayfamilyoptions.org/images/hpv3.jpg>

Screening Guidelines for the Early Detection of Cervical Cancer, American Cancer Society 2009

Screening should begin approximately three years after a woman begins having vaginal intercourse, but no later than 21 years of age.

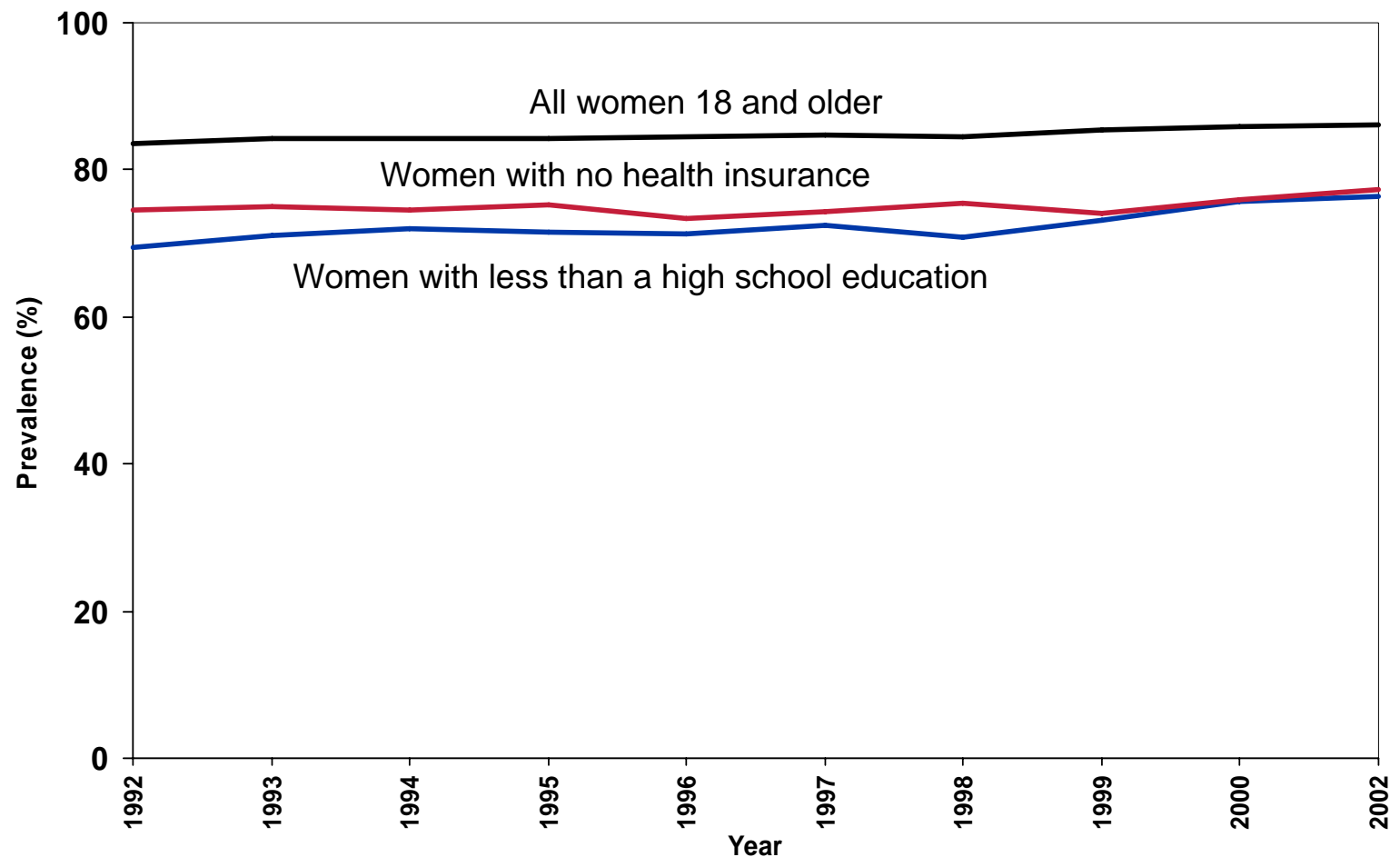
Screening should be done every year with regular Pap tests or every two years using liquid-based tests.

At or after age 30, women who have had three normal test results in a row may get screened every 2-3 years with cervical cytology (either conventional or liquid-based Pap test) alone, or every 3 years with a human papillomavirus DNA test plus cervical cytology.

Women 70 and older who have had three or more consecutive Pap tests in the last ten years may choose to stop cervical cancer screening.

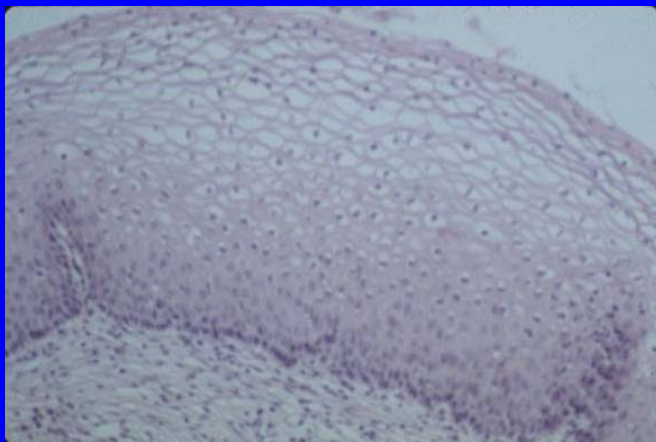
Screening after a total hysterectomy (with removal of the cervix) is not necessary unless the surgery was done as a treatment for cervical cancer.

Trends in Recent* Pap Test Prevalence (%), by Educational Attainment and Health Insurance Status, Women 18 and Older, US, 1992-2002



* A Pap test within the past three years. Note: Data from participating states and the District of Columbia were aggregated to represent the United States. Educational attainment is for women 25 and older. Source: Behavior Risk Factor Surveillance System CD-ROM (1984-1995, 1996-1997, 1998, 1999) and Public Use Data Tape (2000, 2002), National Center for Chronic Disease Prevention and Health Promotion, Center for Disease Control and Prevention, 1997, 1999, 2000, 2001, 2003.

Detecting Cervical Pre-Cancer

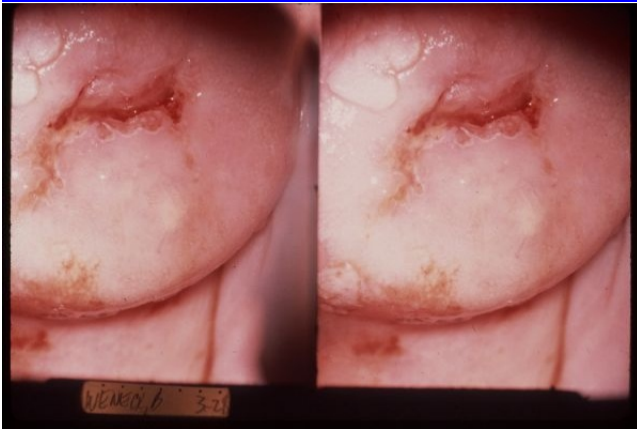


Se = 95%

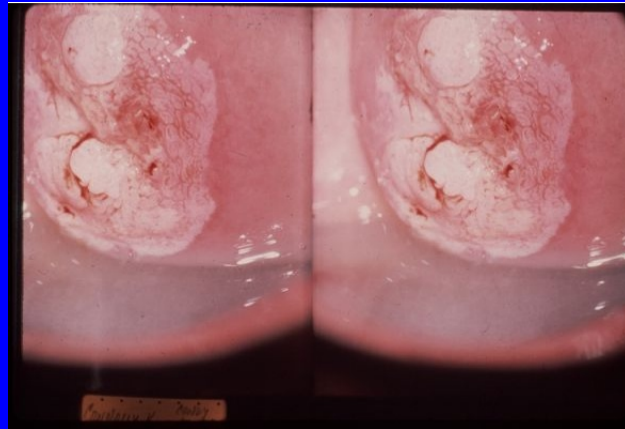
Sp = 44%

Colposcopy

CIN 1/LGSIL



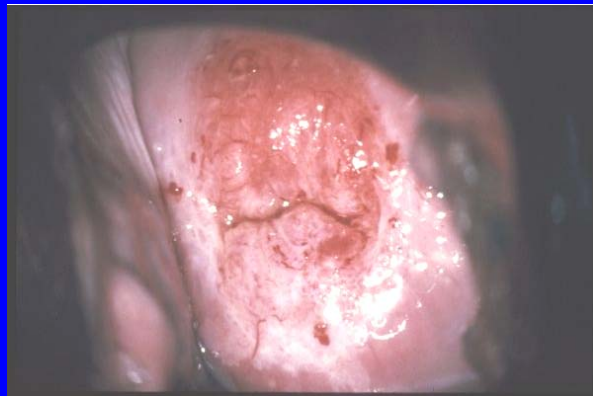
CIN 2/HGSIL



CIN 3/HGSIL



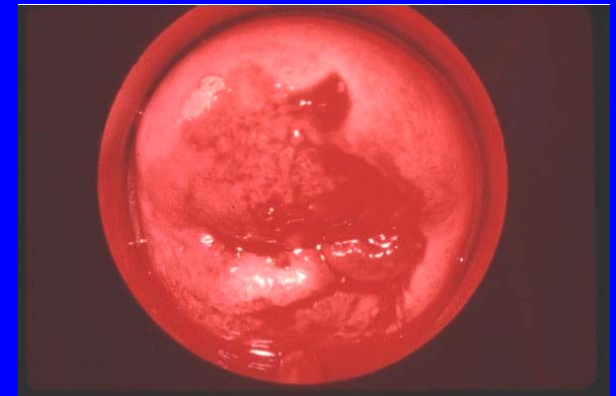
Microinvasive CA



Invasive CA



Invasive CA



Cervical Cancer

■ Screening:

- Annual Pap smear

■ Diagnosis

- Colposcopy + Biopsy

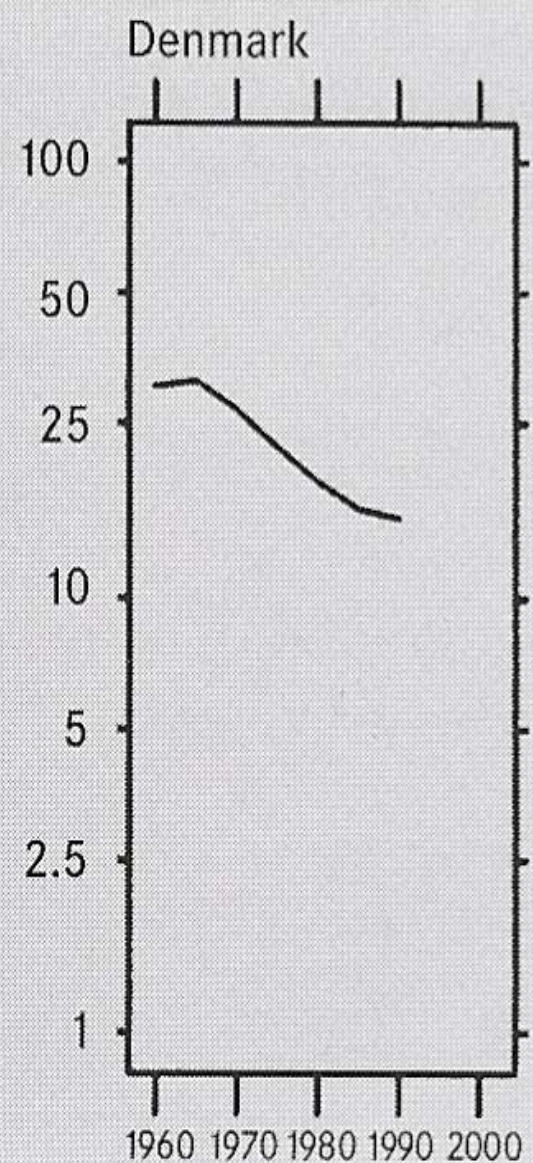
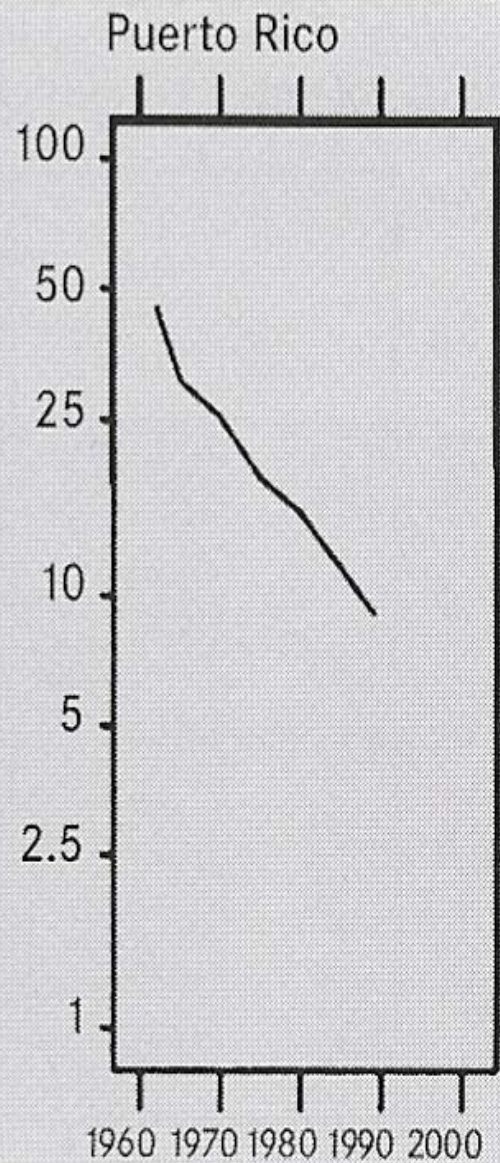
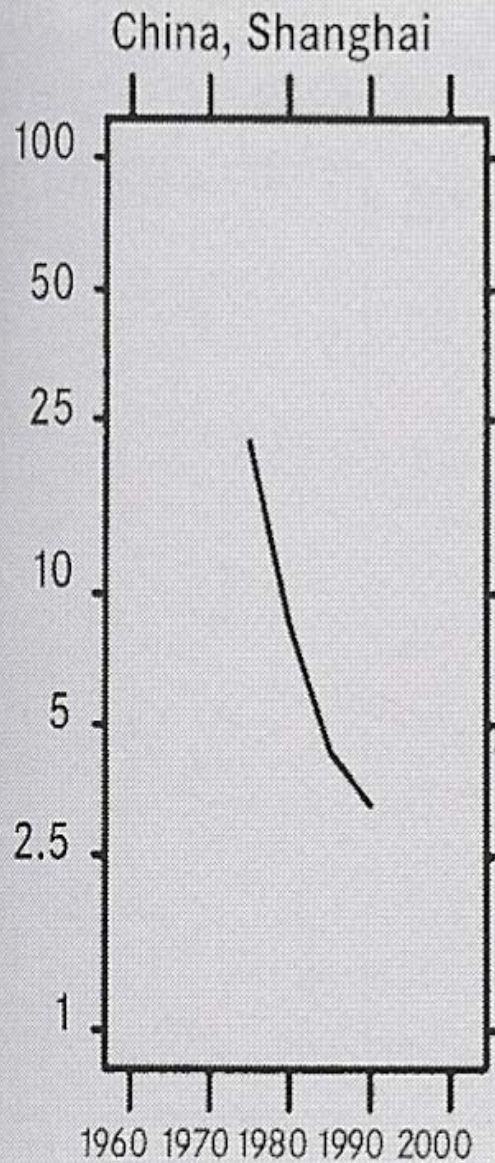
■ Treatment:

- Surgery, radiation therapy, chemotherapy

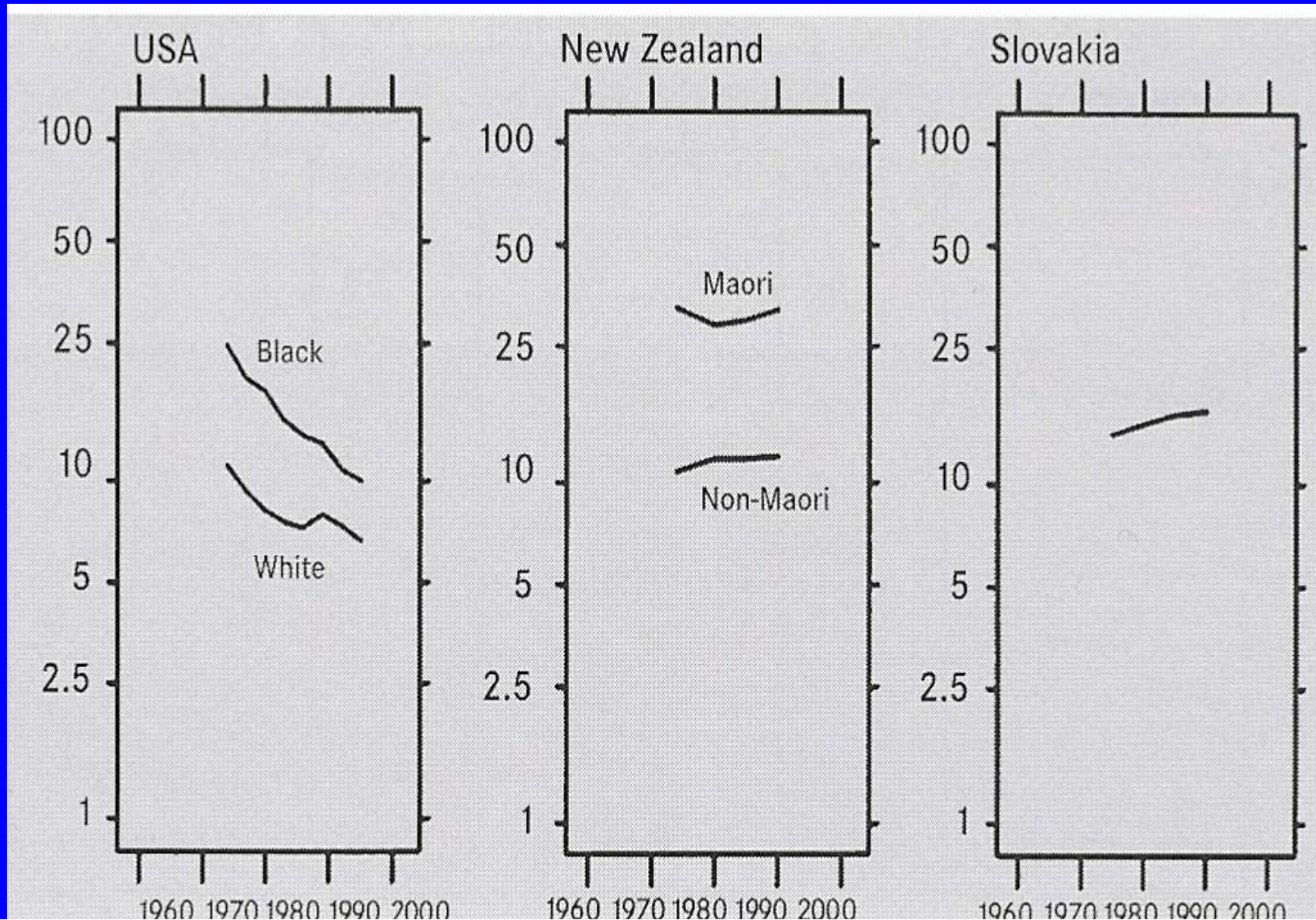
■ 5 year survival

- Localized disease: 92% (56% diagnosed at this stage)

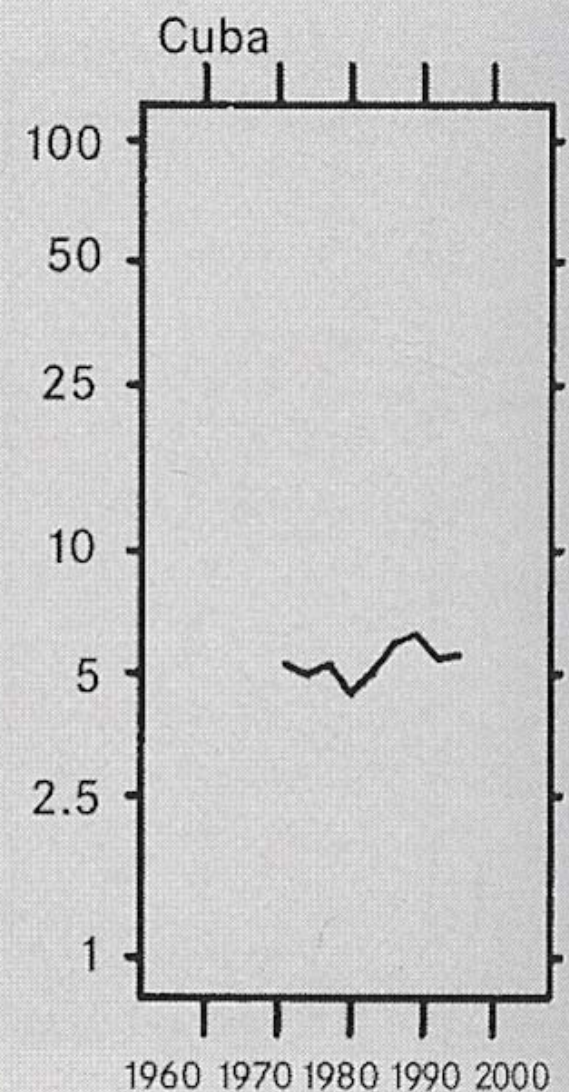
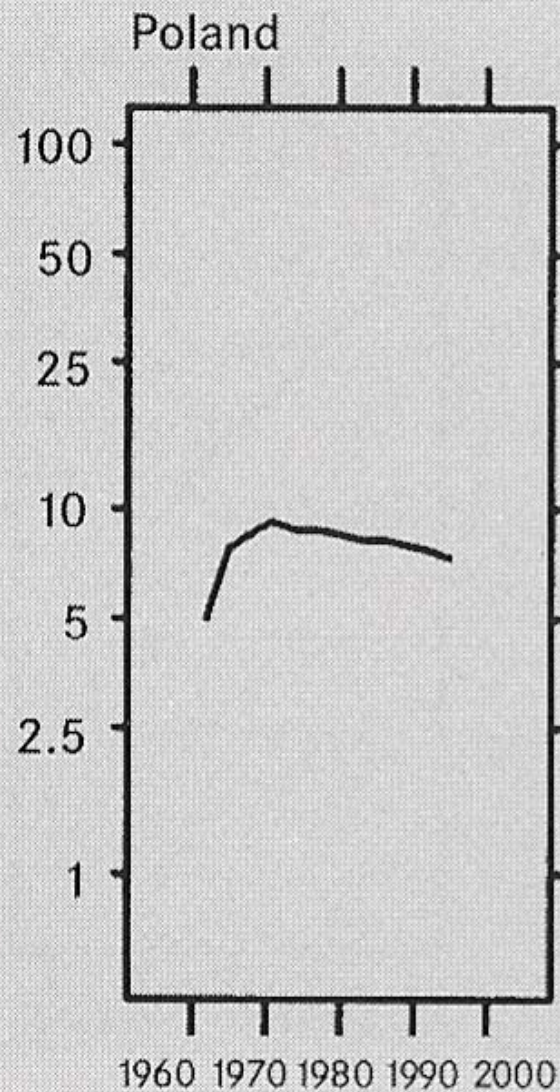
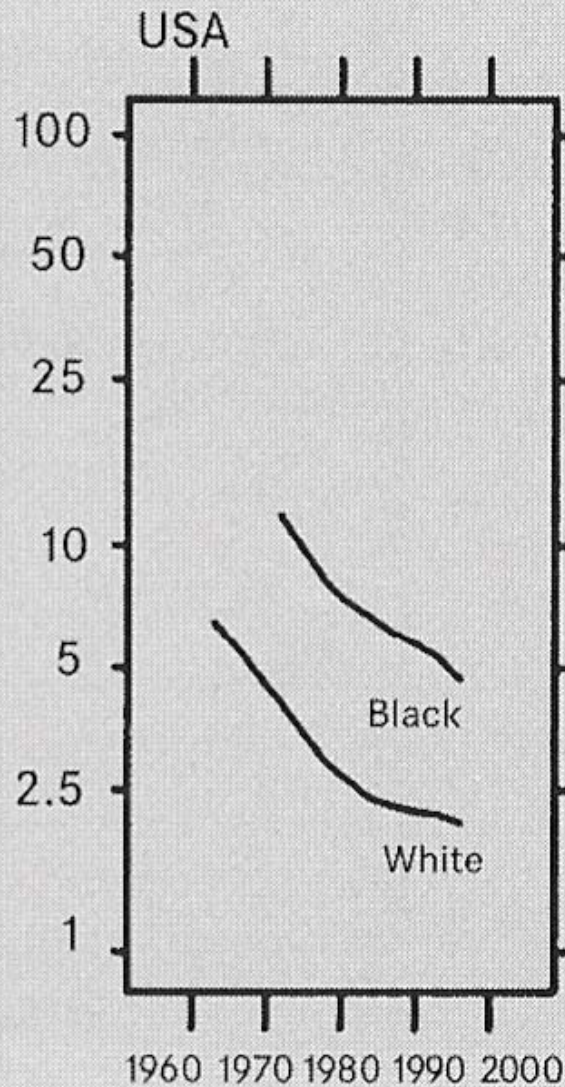
Trends in Cervical Cancer Incidence



Trends in Cervical Cancer Incidence



Trends in Cervical Cancer Mortality

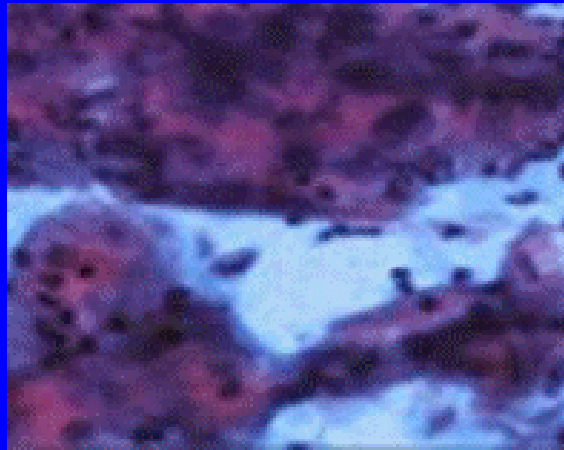


New Technologies for Cervical Cancer

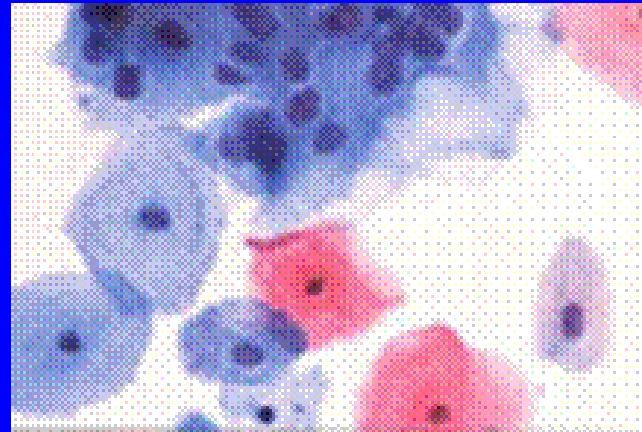
- Liquid Based Pap testing
- Automated Pap smear screening
- HPV Testing
- VIA
- HPV Vaccine

Liquid Based Pap Smear

- Rinse collection device in preservative fluid
- Process suspension of cells to deposit a monolayer of cells on a microscope slide



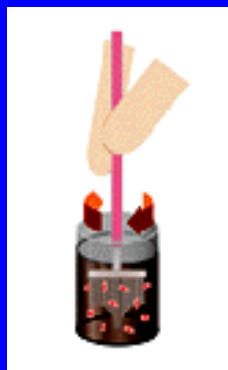
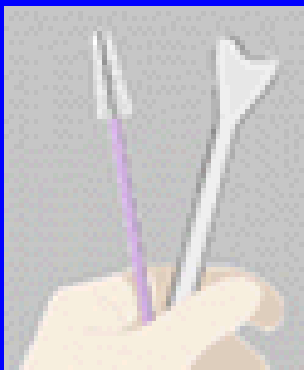
Conventional Pap



Liquid Based Pap

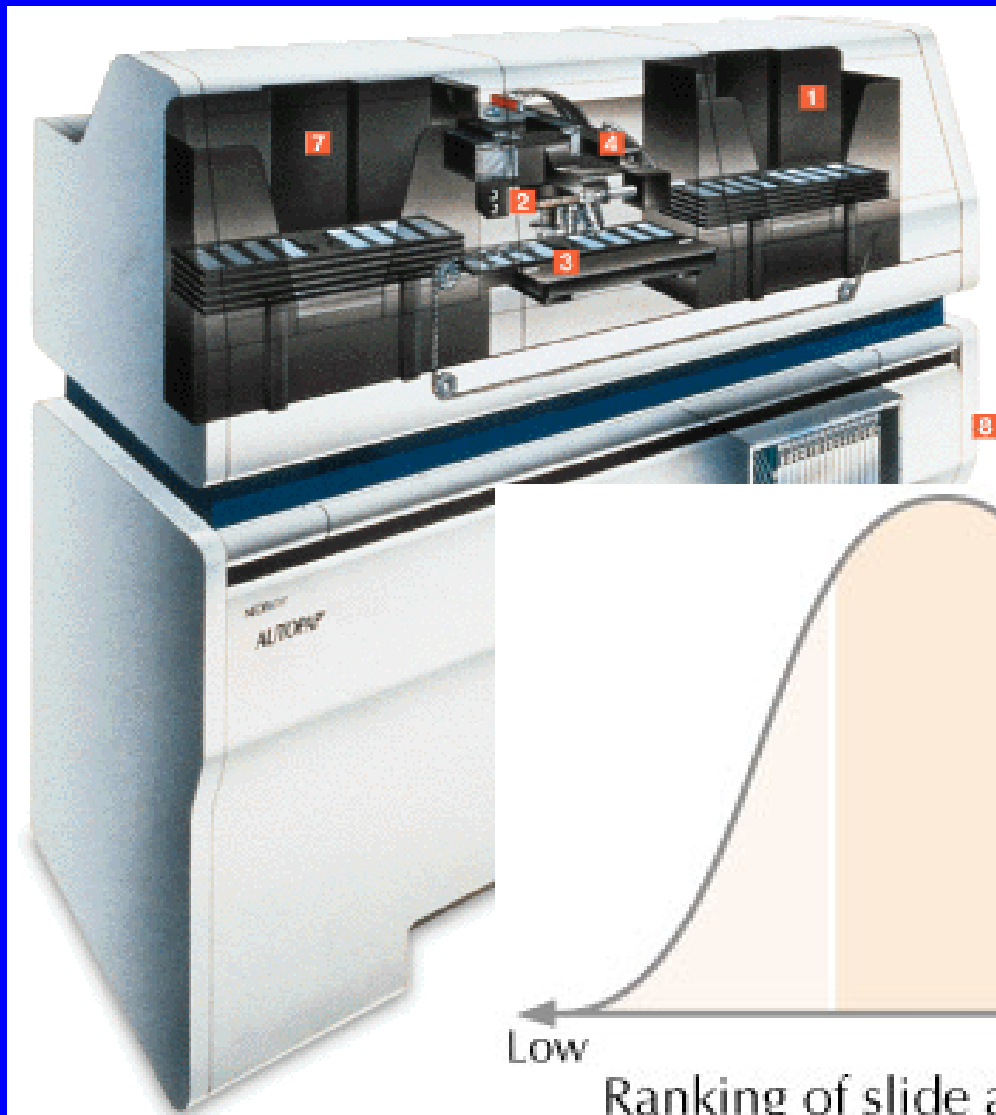
Liquid Based Pap Smear

- Gentle dispersion breaks up blood, mucous, non-diagnostic debris, and mixes sample
- Negative pressure pulse draws fluid through filter to collect a thin, even layer of cells
- Monitor flow through filter during collection to prevent cells from being too scant or too dense
- Cells then transferred to a glass slide



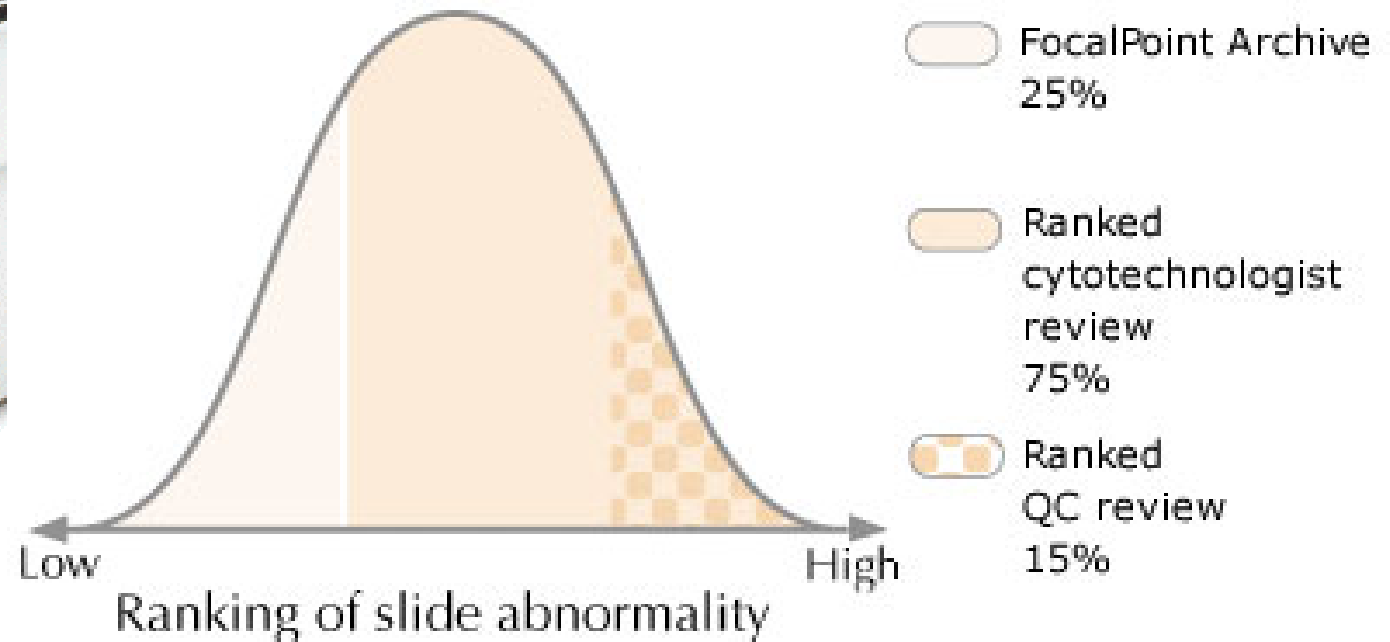
Automated Pap Smear Screening

<http://www.tripathimaging.com/images/cutaway.gif>



- TriPath Care Technologies

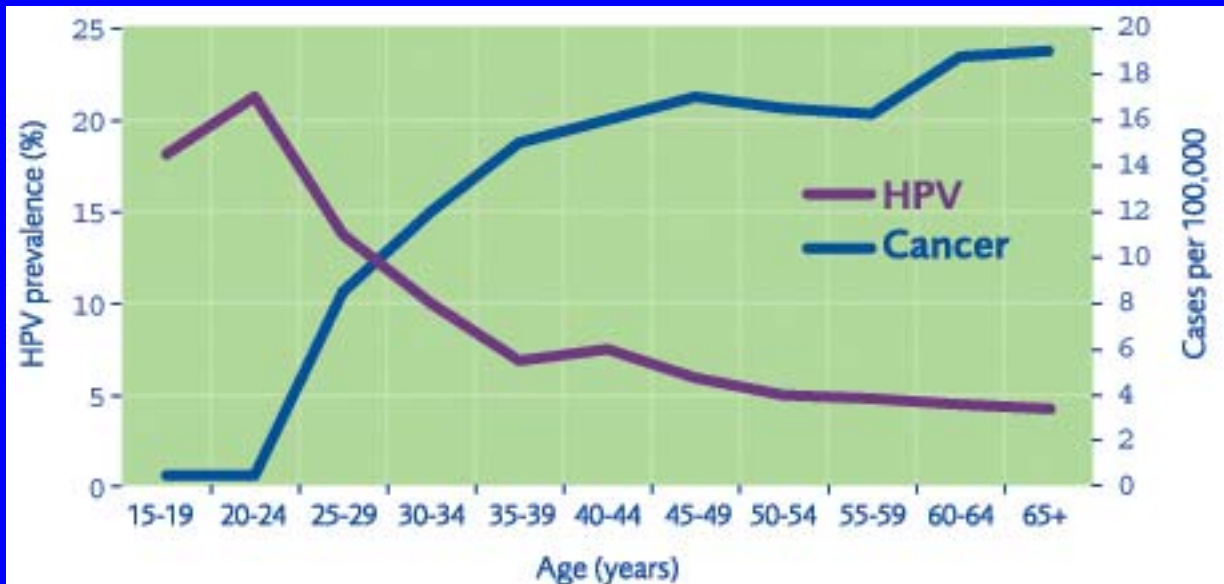
- <http://www.tripathimaging.com/usproducts/index.htm>



HPV Testing

- The DNAwithPap Test is FDA-approved for routine adjunctive screening with a Pap test for women age 30 and older.
- Digene
 - <http://www.digene.com>

<http://www.digene.com/PapX/YLC-5301-30%20VER%20X.mpg>





1. Release Nucleic Acids

Clinical specimens are combined with a base solution which disrupts the virus or bacteria and releases target DNA. No special specimen preparation is necessary.



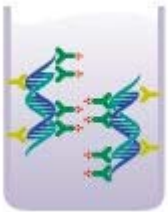
2. Hybridize RNA Probe with Target DNA

Target DNA combines with specific RNA probes creating RNA:DNA hybrids.



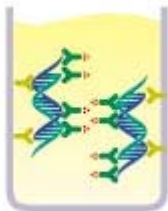
3. Capture Hybrids

Multiple RNA:DNA hybrids are captured onto a solid phase coated with universal capture antibodies specific for RNA:DNA hybrids.



4. Label for Detection

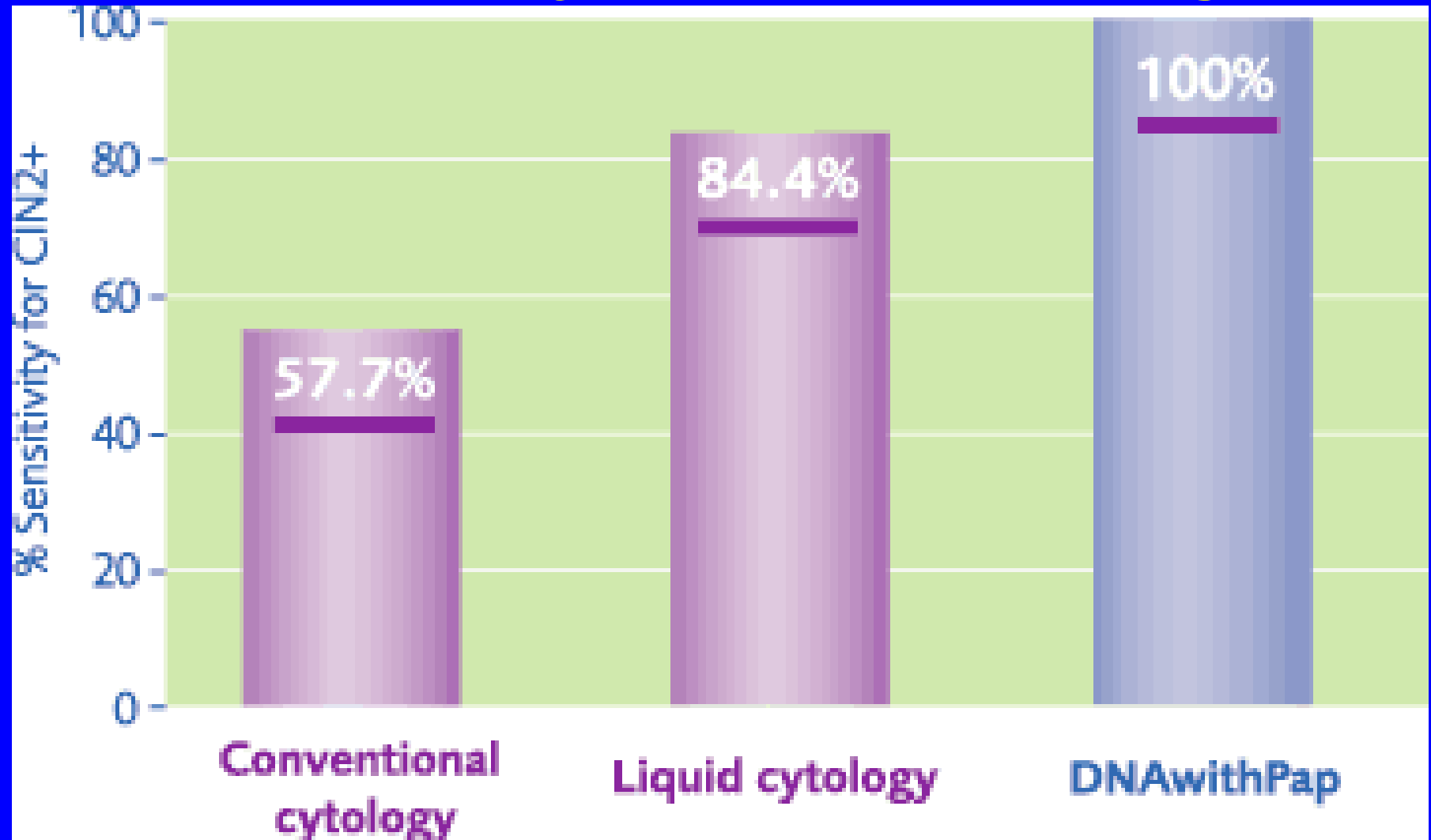
Captured RNA:DNA hybrids are detected with multiple antibodies conjugated to alkaline phosphatase. Resulting signal can be amplified to at least 3000-fold.



5. Detect, Read and Interpret Results

The bound alkaline phosphatase is detected with a chemiluminescent dioxetane substrate. Upon cleavage by alkaline phosphatase, the substrate produces light that is measured on a luminometer in Relative Light Units (RLUs).

Sensitivity of HPV Testing



<http://www.digene.com/images/s>

ens.gif

Study of 5,671 women age >30 years

Comparison of Various Techniques

	Sensitivity	Specificity
Pap smear	60-80%	45-70%
Colposcopy	90-100%	20-50%
Digene HPV Test	80-90%	57-89%
VIA	67-79%	49-86%

HPV Vaccine

■ 2006:

- Gardasil vaccine to prevent HPV infection was licensed for use in girls & women ages 9-26 in USA and 48 other countries
- Protects against 2 strains of HPV responsible for 70% of cervical cancers

■ Non-infectious vaccine

- Made by inserting gene for protein found in the HPV capsid into a different virus or yeast. Recombinantly produced HPV capsid protein self-assembles into virus like particles (VLPs).

HPV Vaccine

■ Gardasil

- Protects against new HPV infections
- Not effective for women who have already been exposed to HPV
- Given as a series of 3 shots over a 6 months
- Cost: \$360
- This cost is a barrier even in developed countries, and is likely to limit its immediate impact in developing countries

HPV Vaccine Efficacy Trials

Manufacturer	Vaccine	Location	Participants	Projected End
Merck	VLPs of L1 protein from HPV 6/11/16/18, made in yeast, aluminum adjuvant	U.S., S. America, Europe	17,800 women, 16 to 26 years old	2007
		U.S., S. America, Europe, Asia	3800 women, 24 to 45 years old	2008
		U.S., S. America, Europe, Asia, Africa	3700 men, 16 to 24 years old	2008
GSK	VLPs of L1 protein from HPV 16/18, made in baculovirus, AS04 adjuvant	U.S., S. America, Europe, Asia Pacific	18,000 women, 15 to 25 years old	2010
		Costa Rica (run by NCI)	12,000 women, 18 to 25 years old	2010

HPV & Cervical Cancer

- Do condoms prevent HPV?
- Do we still need to screen women who have been vaccinated?

Summary of Cancer

- The burden of cancer
 - Contrasts between developed/developing world
- How does cancer develop?
 - Cell transformation → Angiogenesis → Motility
→ Microinvasion → Embolism → Extravasation
- Why is early detection so important?
 - Treat before cancer develops → Prevention
- Accuracy of screening/detection tests
 - Se, Sp, PPV, NPV

Summary of Cervical Cancer

■ Cervical cancer

- 2nd Leading cause of cancer death in women in world
- Caused by infection with HPV
- Precancer → cancer sequence
- Precancer is very common

■ Screening & Detection

- Pap smear; colposcopy + biopsy
- Reduces incidence and mortality of cervical cancer
- Insufficient resources to screen in developing countries

■ New technologies

- Automated reading of Pap smears → reduce FN rate
- HPV testing
- VIA