# The Biology of HPV Infection and Cervical Cancer

Kaitlin Sundling, M.D., Ph.D.
Clinical Instructor

Faculty Director, Cytotechnology Program

Wisconsin State Laboratory of Hygiene and University of Wisconsin School of Medicine and Public Health

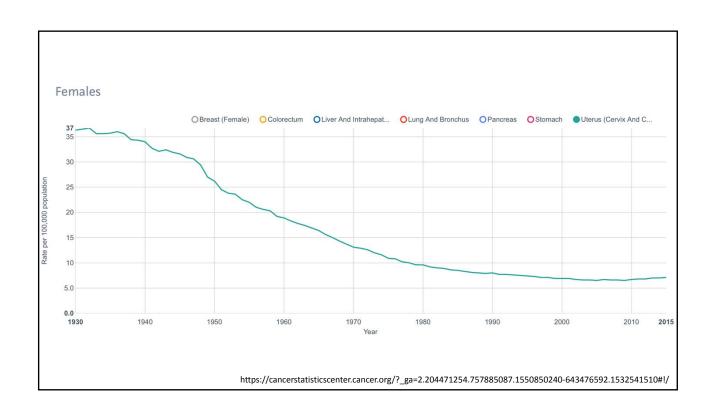
ksundling@wisc.edu @kaitlinsundling on Twitter

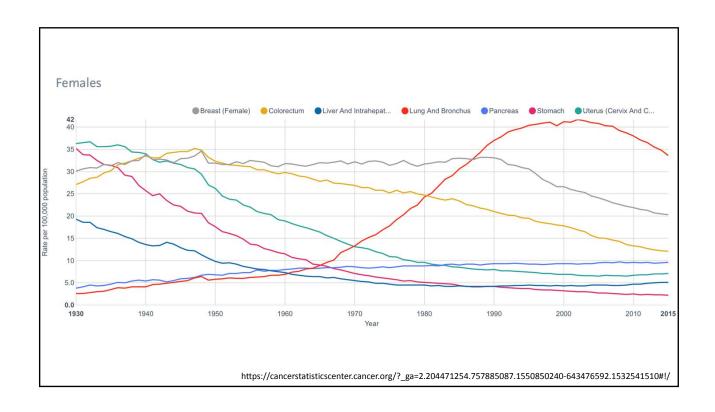
## Learning Objectives

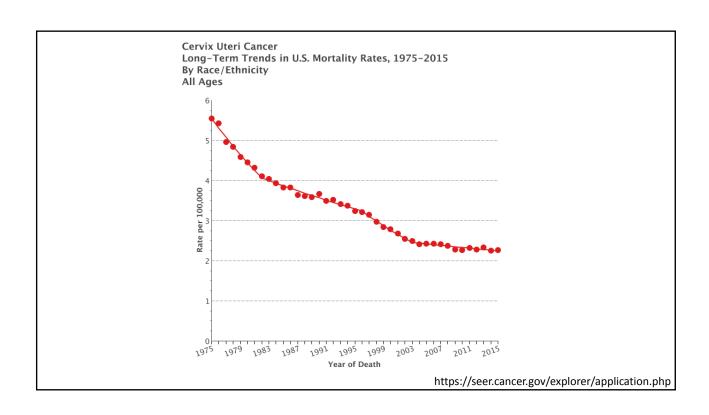
- Describe the key molecular events in HPV oncogenesis.
- Relate transient and persistent HPV infection to patient clinical history and cytomorphologic findings.
- Explain the integration of HPV testing and cytologic findings in cervical cancer screening follow up guidelines.
- Troubleshoot pitfalls in HPV testing.

## The Pap test: A minimally invasive test for cancer and pre-cancer

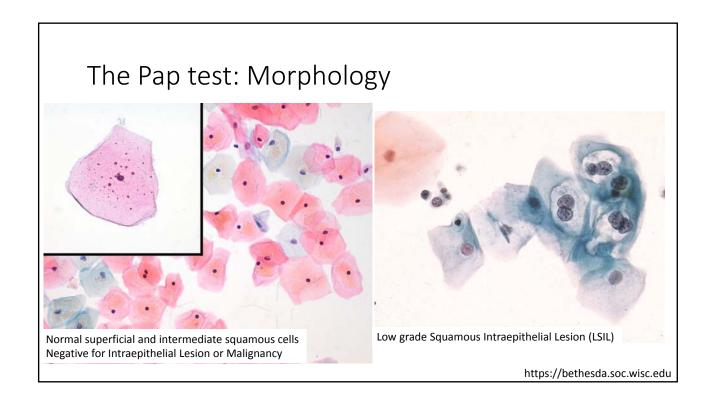
- Originally developed by Dr. George Papanicolaou, immigrant to the US from Greece
- Early scientific work used vaginal smears to study the reproductive cycles of guinea pigs
- Developed a staining method that allowed identification of benign and malignant cells under the microscope
- Original papers were published in late 1910's-1920's
- Pap test widely adopted in the 1940's

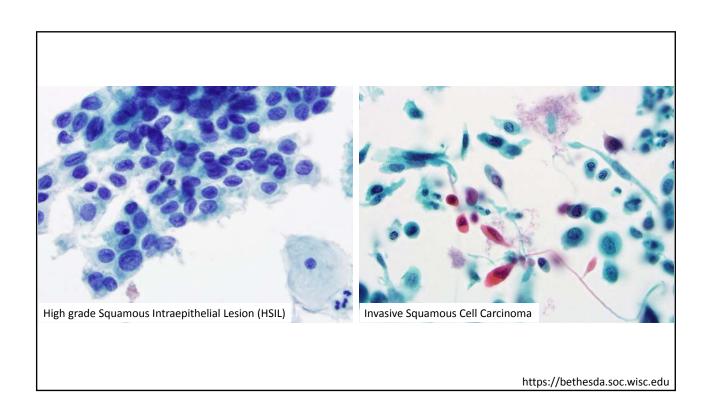








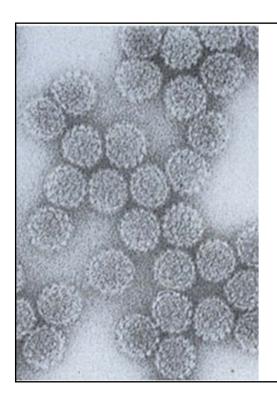






## The Pap test: A crucial component of cervical cancer prevention

- Primary prevention
  - HPV vaccination
  - Condoms (But what about areas not covered by condoms? How likely will
    patients be to use a condom for all contact, every time?)
  - Limiting sexual partners (But what about the partner's partners?)
- Secondary prevention
  - Pap test
  - Appropriate treatment and follow-up of dysplasia (precancerous lesions)



## Human papillomavirus

- Non-enveloped, circular dsDNA virus
- Early genes E6 and E7 bind p53 and Rb
- Late gene L1 makes the major coat protein
- HPV types infect birds and mammals
- Infection is ubiquitous

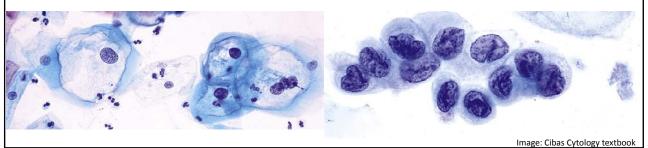
Doorbar J, et al. Rev Med Virol. 2015;25:2-23. doi:10.1002/rmv.1822

### HPV-related disease

- Anogenital tract skin and mucosa penile, vulvar, vaginal, cervical, and anal
- Oropharynx tonsils and base of tongue
- Skin most commonly low risk types, causing warts
- Papillomas of the respiratory tract and conjunctiva usually low risk types

## Low risk vs. high risk HPV infection

- Low risk HPV types can cause koilocytosis and condylomas, unlikely to cause cancer
- High risk HPV types can cause koilocytosis and condylomas, may progress to HSIL (high grade SIL) and cancer
- HPV testing almost uniformly refers to testing for **high risk HPV** types



#### **HPV** vaccination

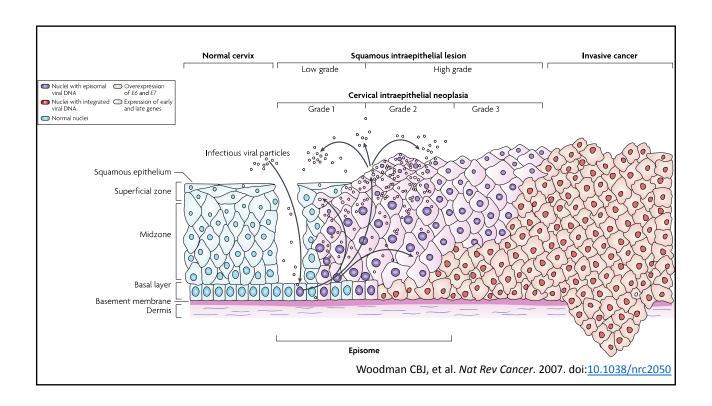
- Most effective at preventing infection when given prior to first exposure
- When given later, may still be effective in preventing infection by new HPV types

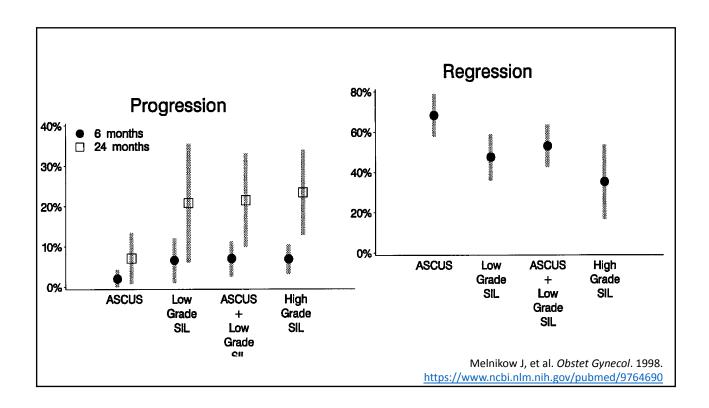
### **HPV** vaccination

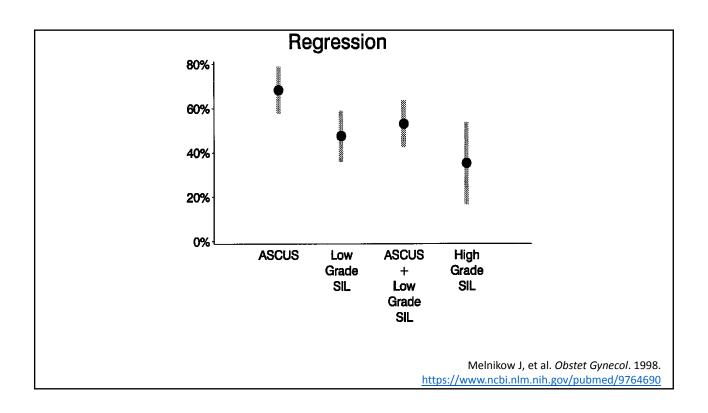
- Risks of HPV Vaccination: Allergic response to vaccine components, minor localized or febrile (fever) vaccine reactions
- Benefits of HPV Vaccination:
  - Boys: Reduced risk of genital warts, reduced risk of penile cancer, reduced risk of anal cancer, reduced risk of oropharyngeal cancer
  - Girls: Reduced risk of genital warts, reduced risk of cervical, vaginal, vulvar, and anal cancer, and reduced risk of oropharyngeal cancer
  - General public: Herd immunity
- <a href="https://www.cdc.gov/hpv/hcp/for-hcp-tipsheet-hpv.pdf">https://www.cdc.gov/hpv/hcp/for-hcp-tipsheet-hpv.pdf</a>
- <a href="https://wicancer.org/action-plans/hpv-vaccination-rates/">https://wicancer.org/action-plans/hpv-vaccination-rates/</a>

## Gardasil 9 vaccine

- L1 protein virus-like particles
- Protective agains:
  - 6, 11 low risk, causing genital warts
  - 16, 18, 31, 33, 45, 52 and 58 high risk
- Recommended for boys and girls ages 11 or 12
- May begin as early as age 9, catch up recommended up to age 26 for women, 21 for men
- FDA approval recently extended to upper limit of age 45





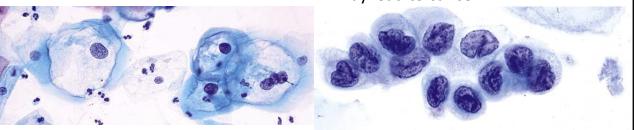


#### **Transient HPV infection**

- Usually LSIL/koilocytic changes
- Common in women in their 20s
- Regresses

#### **Persistent HPV infection**

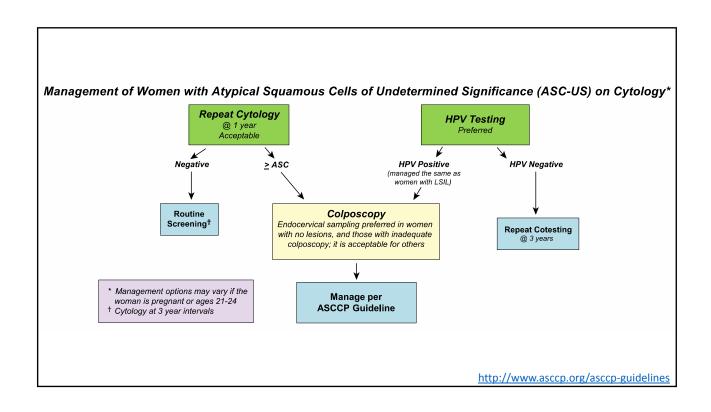
- HSIL, koilocytes less likely
- More common in women in their 30s and up
- May lead to cancer

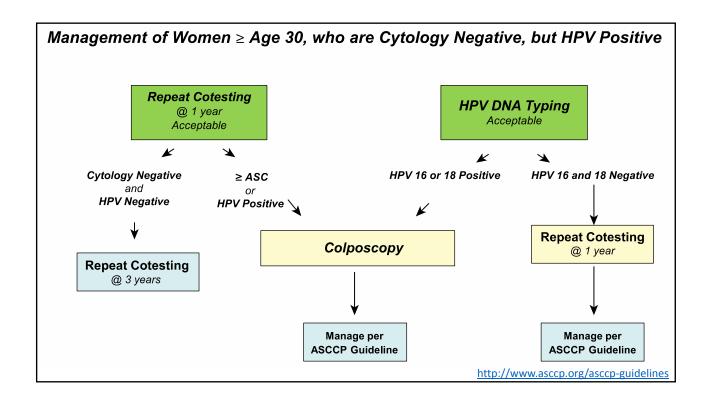


## Routine Cervical Cancer Screening Guidelines

Age group	ASCCP 2012	USPSTF 2018			
Under 21	No screening	No screening			
Age 21-29	Cytology alone every 3 years	Cytology alone every 3 years			
Age 30-65	Cytology alone every 3 years OR Cotesting with cytology and HPV testing every 5 years	Cytology alone every 3 years OR Cotesting with cytology and HPV testing every 5 years OR Primary screening with HPV testing along every 5 years			
Over 65	Discontinue screening if adequately screening and not at high risk for cervical cancer	Discontinue screening if adequately screening and not at high risk for cervical cancer			

 $\frac{https://www.uspreventiveservicestask force.org/Page/Document/UpdateSummaryFinal/cervical-cancer-screening2}{http://www.asccp.org/Assets/fcd6fdab-0325-466b-a5cd-3c1c06cf0e66/635912171989730000/asccp-guidelines-pdf}$ 





Test	Target Gene(s)	Target Biomolecule	Internal Control	Technology	HPV Types Detected	Fixative
Abbott RealTime High Risk HPV Assay	L1	DNA	Beta globin	PCR	<b>16, 18</b> , 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68	ThinPrep
APTIMA	E6 and E7	mRNA	Spiked in	Transcription mediated amplification	<b>16, 18/45</b> , 31, 33, 35, 39, 51, 52, 56, 58, 59, 66, 68	ThinPrep
Cervista	L1	DNA	HIST2H2BE	Isothermal DNA amplification, Invader FRET	<b>16, 18</b> , 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68	ThinPrep
Cobas 4800 HPV Test	L1	DNA	Beta globin	PCR	<b>16, 18</b> , 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68	ThinPrep
Hybrid Capture 2	L1	DNA	None	RNA probes and antibody detection of RNA:DNA hybrids	16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68	ThinPrep
BD Onclarity	E6 and E7	DNA	Beta globin	PCR	<b>16, 18</b> , 31, 45, 51, 52, and 59, (33, 56, 58, 66), (35, 39, 68)	SurePath

Adapted from Bibbo Comprehensive Cytopathology

## Risks of screening

- False positives
  - Unnecessary biopsies and loop electrosurgical excision procedures
    - May lead to shortened cervix or cervical stenosis
    - Reduced fertility or incompetent cervix
- False negatives
  - Lost opportunity for early treatment
  - Lesions may present when already invasive or even metastatic, requiring more invasive treatment, impact of quality of life and survival
- Direct costs to patients
- Public health impacts
- Utilization of healthcare resources



## Potential biological pitfalls of HPV testing

- 10% of invasive carcinomas may be HPV negative<sup>1</sup>
  - Presumed loss of HPV viral DNA in the tumor after acquisition of other mutations, such as DNA repair defects
  - L1 gene may be lost<sup>2</sup>

Flanagan MB. Archives of Pathology & Laboratory Medicine. 2018. doi: 10.5858/arpa.2018-0001-RA
 Burd EM. Clinical Microbiology Reviews. 2016. doi: 10.1128/CMR.00013-15

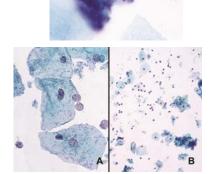
**Table 2** Secondary genetic typing for 205 HSIL cases of found to be either totally negative for HPV or positive for other high-risk HPV genotypes other than HPV 16/18 on initial testing.

HPV type	Total					Single				ixed	Risk		
	N	Bx Conf	% HSIL	Bx Conf	N	Bx Conf	% HSIL	Bx Conf	N	Bx Conf	% HSIL	Bx Conf	
			cases				cases				cases		
31	30	12	14.6	17.6	23	10	11.2	14.7	7	2	3.4	2.9	High
52	13	5	6.3	7.4	10	5	4.9	7.4	3	0	1.5	0.0	
58	13	6	6.3	8.8	11	4	5.4	5.9	2	2	1.0	2.9	
35	9	7	4.4	10.3	8	6	3.9	8.8	1	1	0.5	1.5	
45	7	3	3.4	4.4	1	1	0.5	1.5	6	2	2.9	2.9	
33	6	3	2.9	4.4	4	2	2.0	2.9	2	1	1.0	1.5	
59	6	2	2.9	2.9	4	2	2.0	2.9	2	0	1.0	0.0	
16	3	1	1.5	1.5	1	1	0.5	1.5	2	0	1.0	0.0	
56	1	0	0.5	0.0	1	0	0.5	0.0	0	0	0.0	0.0	
Negative	76	18	37.1	26.5									

McCarthy E, Ye C, Smith M, Kurtycz DFI. *Journal of the American Society of Cytopathology*. 2016. doi:10.1016/j.jasc.2016.08.001

## Potential biological pitfalls of HPV testing

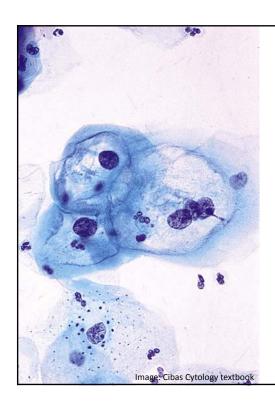
- Endogenous flora (coccobacilli including *Gardnerella sp.*, lactobacilli) and cytolysis may lead to interference
- Shifts in high risk HPV types with increasing HPV vaccination
  - Gardasil 9: 16, 18, 31, 33, 45, 52, 58
  - Most HPV tests: 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68
- Selective pressure on the L1 gene due to HPV vaccination



https://bethesda.soc.wisc.edu

## Technical pitfalls in HPV testing

- SurePath vs. ThinPrep
  - SurePath vials should contain the collection device, while ThinPrep should not
  - SurePath fixative contains a small amount of formaldehyde
  - Some laboratories have validated a boiling pre-processing step
    - Burd EM. Clinical Microbiology Reviews. 2016. doi:10.1128/CMR.00013-15
- Alternate collection methods/sources
  - Vaginal self-collection
  - Urine



## On the horizon

- · HPV primary screening
  - Australia and the Netherlands
  - Potential reflex to cytology
  - Concerns: PPV, NPV, colposcopy infrastructure
- HPV testing combined with other tests to improve specificity for precancerous lesions
  - DNA methylation
  - Gene expression
  - IHC staining
- CDC Grand Rounds: Preventing Cervical Cancer in the 21<sup>st</sup> century
  - https://www.cdc.gov/grand-rounds/pp/2019/20190125cervical-cancer.html

## Thank you!

Questions, comments, suggestions, or potential collaborations?

ksundling@wisc.edu @kaitlinsundling on Twitter