# How to Give an Oral Presentation

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## Before we begin

#### **ACKNOWLEDGEMENTS**

- David Spach
  - Slides

#### REFERENCES

Hayne & McDaniel,
 Nursing Forum 2013;
 48:4 (289-94)

# Giving an effective oral presentation

#### **PRESENTATION**

- Clear
- Concise
- Organized
- Informative
- Relevant

#### **SPEAKER**

- Energetic
- Charismatic
- Passionate
- Humorous
- Knowledgeable

#### **Outline**

- Organizing your talk
- Enhancing your presentation
- Powerpoint tips and pitfalls
- Delivery

# Types of "Fellow Talks"

- Case conference/clinical review
- Informal lab meeting/research group
- Research meeting
  - Division
  - Regional/national
- Formal venue
  - Job talk
  - Grand rounds

# Approach to organizing your talk

- Know your audience
- Know your message
- Know your material
- What is the goal?

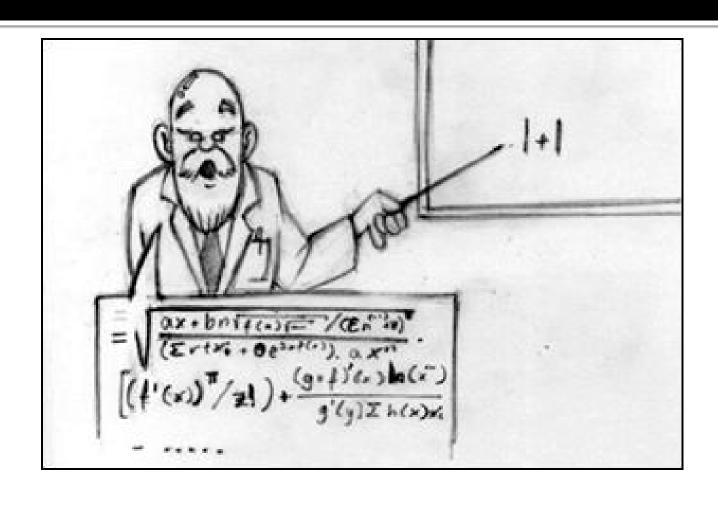
#### **Audience**

- Case conference/clinical review
  - Medical students, community providers, peers
- Informal lab meeting/research group
  - Scientists, epidemiologists, biostatisticians
- Research meeting
  - Division or Regional/national
    - Peers
- Formal venue
  - Peers, colleagues with broad expertise

# Managing a mixed audience

- Use real examples
- Focus on key information
  - Do not provide extraneous level of detail
- Present material in a step-wise level of difficulty
  - Bring people along with you

# Keep it simple



# Define your message: Research Talk

- Make it easy to understand
- Limit to 1 or 2 key messages
- Focus your content
  - Most important for background and results
- Don't overwhelm with details
  - Laser focus your presentation

# Goal of presentation

- Transfer of specific knowledge/information
- Enhance understanding of concepts
- Gain new insight
- Stimulate interest in subject
- Change behavior/practice

### "Outcomes" Related to a Presentation

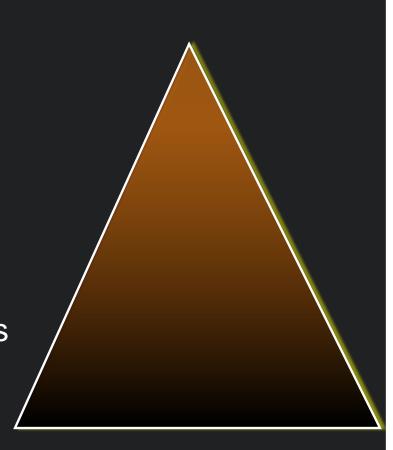
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Stimulate interest/motivate

Gain new insight

Enhance understanding of concepts

Transfer of knowledge/information



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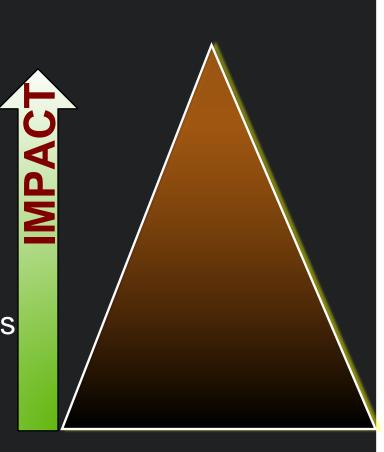
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Gain new insight

Enhance understanding of concepts

Transfer of knowledge/information



# Enhancing Presentation Delivery The Four P's

- Preparation (room and AV equipment)
- Projection
- Pauses
- Performance

#### **Presentation**

- Beware of technical issues
  - Mac: PC
    - Especially animation and video
  - Advancing slides
    - Pointer or mouse
  - Microphone
  - Webcast
    - Use mouse instead of pointer, microphone
  - Back it up
    - Email, dropbox, flash drive

# **Projection**

- Size of room and screen
- Make slides readable
- Fonts:
  - Title: 36 to 44 point size
  - Text: 28 or 32 point size
  - Nothing less than 20 point size
  - Sans-serif

## Performance

- Be organized
  - Stay within time limits

- Cut slides/concepts if needed to present in calm manner
  - ~1 minute per slide
    - Don't go over time
    - Don't rush

### Performance

- Practice
  - Enough so you are comfortable and talk flows
  - Not so much that it is memorized/over-rehearsed
  - Know what works for you
    - I usually practice 3 times
      - 1. Cut, flow
      - 2. Refine
      - 3. Optimize: timing and pacing

# Managing anxiety

- Memorize first few slides/transitions
- Bring an outline with you, but don't read it
- Consider β-blocker if very anxious
  - Try it before the meeting/presentation

# Use of the pointer

- Judicious use
- Do not allow pointer to distract
- Do not allow pointer to reveal anxiety
  - Use hand to steady —or-
  - Use mouse

# **Powerpoint**

FRIEND OR FOE?

#### **Speaker Goals for Session**

- Change your approach to preparing for a lecture
- Increase use of images to enhance presentation
- Improve presentation delivery
- Stimulate your interest to improve as a speaker

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PRESENTATION SKILLS
2013 STATE OF THE ART

Table 3. Recommendations for the Treatment of Methicillin-Resistant Staphylococcus aureus (MRSA)

Manifestation	Treatment	Adult dose	Pediatric dose	Class <sup>a</sup>	Comment
Skin and soft-tissue infection (SSTI)					
Abscess, furuncles, carbuncles	Incision and drainage			All	For simple abscesses or boils, incision and drainage is likely adequate. Please refer to Table 2 for conditions in which antimicrobial therapy is recommended after incision and drainage of an abscess due to CA-MRSA.
Purulent cellulitis (defined as cellulitis associated with purulent drainage or exudate in the absence of a drainable abscess)	Clindamycin	300–450 mg PO TID	10–13 mg/kg/dose PO every 6–8 h, not to exceed 40 mg/kg/day	All	Clostridium difficile—associated disease may occur more frequently, compared with other oral agents.
	TMP-SMX	1–2 DS tab PO BID	Trimethoprim 4–6 mg/kg/dose, sulfamethoxazole 20–30 mg/kg/dose PO every 12 h	All	TMP-SMX is pregnancy category C/D and not rec ommended for women in the third trimester of pregnancy and for children <2 months of age.
	Daxycycline	100 mg PO BID	≤45kg: 2 mg/kg/dose PO every 12 h >45kg: adult dose	All	Tetracyclines are not recommended for children under 8 years of age and are pregnancy category D.
	Minocycline	200 mg × 1, then 100 mg PO BID	4 mg/kg PO × 1, then 2 mg/kg/dose PO every 12 h	All	
	Linezolid	600 mg PO BID	10 mg/kg/dose PO every 8 h, not to exceed 600 mg/dose	All	More expensive compared with other alternatives
Nonpurulent cellulitis (defined as cellulitis with no purulent drainage or exudate and no associated abscess)	β-lactam (eg, cephalexin and dicloxacillin)	500 mg PO QID	Please refer to Red Book	All	Empirical therapy for β-hemolytic streptococci is recommended (All). Empirical coverage for CA-MRSA is rec- ommended in patients who do not respond to β-lactam ther- apy and may be considered in those with systemic toxicity.
	Clindamycin	300–450 mg PO TID	10–13 mg/kg/dose PO every 6-8 h, not to exceed 40 mg/kg/day	All	Provide coverage for both β-hemolytic streptococci and CA-MRSA
	β-lactam (eg, amoxicillin) and/or TMP-SMX or a tetracycline	Amoxicillin: 500 PO mg TID See above for TMP-SMX and tetracycline dosing	Please refer to Red Book See above for TMP- SMX and tetracycline dosing	All	Provide coverage for both β-hemolytic streptococci and CA-MRSA
	Linezolid	600 mg PO BID	10 mg/kg/dose PO every 8 h, not to exceed 600 mg/dose	All	

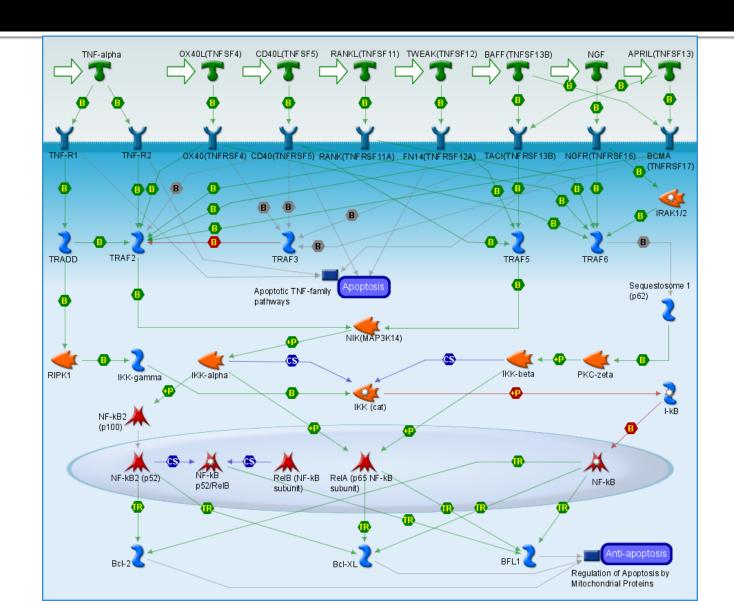
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# I know you can't read what is on this slide, but.....

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## This slide is busy but...



### Resolution

Test	Se	nsitivity	References	
	Cancer	Advanced Adenomas*		
	p	percent		
Stool-based tests				
Standard guaiac fecal occult-blood test (three stool samples)	33–50	11	Mandel et al., <sup>13</sup> Hardcastle et al., <sup>14</sup> Kronborg et al., <sup>15</sup> Imperiale et al., <sup>16</sup> Ahlquist et al. <sup>17</sup>	
Sensitive guaiac fecal occult-blood test (three stool samples)	50–75	20–25	Levin et al., <sup>2</sup> Whitlock et al., <sup>4</sup> Ahlquist et al., <sup>17</sup> Allison et al. <sup>18</sup>	
Immunochemical fecal occult-blood test (one-three stool samples)	60–85	20–50	Levin et al., <sup>2</sup> Whitlock et al. <sup>4</sup>	
Old stool DNA test (one stool sample)	51	18	Imperiale et al. <sup>16</sup>	
New stool DNA test (one stool sample)	≥80	40	Allison et al.,18 Itzkowitz et al.19	
Structural examinations of the colon				
CT colonography	Uncertain; probably >90	90 (if ≥10 mm in diameter)	Johnson et al. <sup>20</sup>	
Sigmoidoscopy	>95 (in the distal colon)	70†	Selby et al.,21 Lieberman et al.22	
Colonoscopy	>95	88–98	Lieberman et al., <sup>22</sup> Imperiale et al., <sup>23</sup> Schoenfeld et al., <sup>24</sup> Lieberman et al., <sup>21</sup> Pickhardt et al., <sup>26</sup> Cotton et al., <sup>27</sup> Rockey et al. <sup>28</sup>	

<sup>\*</sup> Advanced adenoma is defined as a tubular adenoma that is 10 mm or larger in diameter or an adenoma with villous histologic features or high-grade dysplasia.

Lieberman DA. NEJM 2009:361; 1179-1187

<sup>†</sup> If an adenoma is detected in the distal colon, the patient would undergo complete colonoscopy, which would result in the detection of some proximal advanced adenomas.

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# **Transitions**

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# Animations

#### **Animations**

Distracting

Use appear function for simplicity

# Other Powerpoint pitfalls

- Multiple fonts
- INCONSISTENT FORMAT
  - Removes attention from the speaker and content

# "Death by Powerpoint"

Aim 1 Correlation of magnitude, severity, breadth, and phenotype of blood CD8 responses with HSV severity. We will enroll 60 HSV-2-seropositive women. Thirty will have no recognized recurrent GH and measured shedding rates of 0 of 30 days (0%) (Group 1), and 30 will have 4 or more episodes/year and shedding rates of 6 or more out of 30 days (>19%) (Group 2). The shedding rates are low and high tertiles based on 396 participants who completed 454 shedding sessions >30 days long. These assignments are relatively consistent in repeated sessions and therefore reflect a biological phenomenon within-individual (see Clinical Core). HSV-2 shedding rates will be determined by daily pooled anogenital swab PCR for HSV-2 DNA as detailed in the Cores. Subjects will be immunocompetent adults, 18-55. The age limit reduces confounding from the gradual decrease of HSV severity over time <sup>124</sup>. We will study only women to access cervical samples for Aim 2. There will be 10 HSV-1/HSV-2 (-) double negative (DN) controls. Immune seronegative (IS) DN persons occasionally 50,51 have T-cells (CD4>CD8) that react to HSV-2 peptides at low EC50. At screening, candidates will be screened for HIV-1, HBV, and HCV. These are immunomodulatory and disallow the cell sorting in Aim 1. Persons with immune suppression or who lack a cervix will be excluded. Subjects who have previously participated in daily PCR shedding studies will be eligible but will have a repeat shedding study. We will re-contact persons with group 1 or 2 phenotypes from previous studies and recruit new persons as needed. HSV-1/HSV-2 serology will be immunoblot <sup>118</sup>. Blood samples (150 ml for PBMC Ficoll/freeze) will be obtained at the beginning and end of a 30 day shedding study. As PCR specimen sets accrue, they will be analyzed to allow subject assignments to group 1, 2 or neither. Genomic DNA will be collected for HLA typing (Lab Core).

Cytobrush @ beginning and end too

Bottom line: 70 people (30 mild 30 severe 30 HSV neg), 2 bleeds for Ficoll; HLA type, 2 cervix per person

# Study design

**Design**: Prospective observational cohort study

**Population**: HSV-2 seropositive women age 18-55

HIV, Hep B, Hep C seronegative

Cohort 1 (n=30): No prior history of symptomatic genital HSV

Cohort 2 (n=30): 4 or more recurrences in past year

Control (n=10): HSV-1/HSV-2 seronegative

# Taking advantage of powerpoint

- Highlight
- Use pictures
- Use graphics

#### Physical Exam

- T=40, HR=100, BP 90/50, RR=20
- OP clear. No conjunctivitis
- Neck supple, normal ROM
- Lungs CTA B
- RRR, no m/g/r
- Abd: Soft. Guarding RUQ. No splenomegaly
- Skin: No rash
- Lymph node: No cervical, axillary LAN, shoddy inguinal LAN
- MŠ: No joint swelling/erythema

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# HSV-2 is the leading cause of

City	No. tested by M-PCR	Haemophilus ducreyi No. (%)	Treponema pallidum No. (%)	HSV No. (%)	T. pallidum/HSV No. (%)	Negative No. (%)
Chicago	49 <sup>a</sup>	6 (12)	4 (8)	24 (49)	1 (2)	14 (29)
Cincinnati	52	0	1 (2)	41 (79)	0	10 (19)
Dallas	52	0	6 (12)	35 (67)	2 (4)	9 (17)
Houston	51	0	1 (2)	38 (75)	1 (2)	11 (22)
Los Angeles	54	0	0	41 (76)	0	13 (24)
Memphis	50	10 (20) <sup>b</sup>	15 (30)	14 (28)	6 (12)	5 (10)
New York	55	0	1 (2)	36 (65)	1 (2)	17 (31)
Philadelphia	50	0	3 (6)	38 (76)	1 (2)	8 (16)
St. Louis	53	0	7 (13)	28 (53)	1 (2)	17 (32)
Total	516	16 (3) <sup>b</sup>	51 (10)	320 (62)	13 (3)	116 (22)

Similar findings in studies from around the world

a 50 patients were enrolled; 1 specimen was unsatisfactory and was not tested.
 b Includes 1 participant with both *H. ducreyi*/herpes simplex virus (HSV) detected and 2 with both *H. ducreyilT. pallidum* detected.

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## Impact of Genital Herpes

- Leading cause of genital ulcer disease
- Causes neonatal herpes
- Increases risk of HIV infection 2-3 fold

## Impact of Genital Herpes

#### Genital ulcer disease

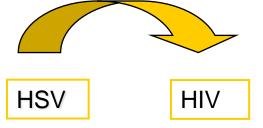




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#### **Neonatal Herpes**





Increases the risk of HIV-1 acquisition 2-3 fold

Genital ulcer disease increases risk of HIV-1 transmission



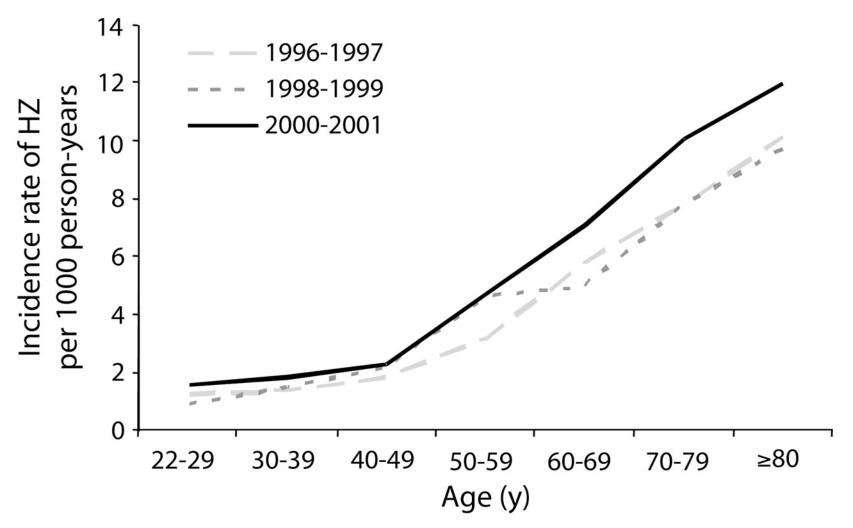
25%-50% of HIV infections attributable to HSV-2 in high prevalence settings

## Herpes zoster epidemiology

- Incidence increases with age
  - 2 cases per 1,000 person-years in <40 yrs</p>
  - 12 cases per 1,000 person-years in >80 yrs

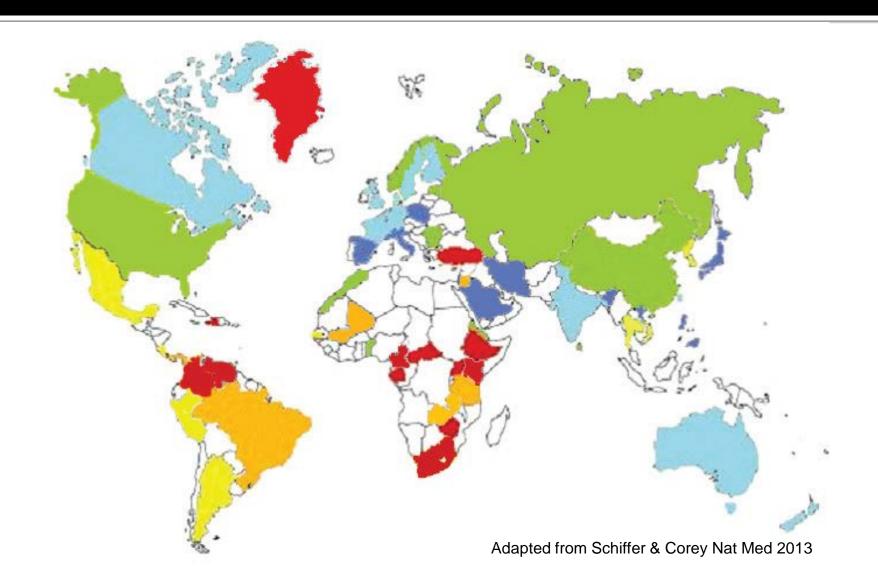
Incidence has been increasing in elderly population over past 7 years

Incidence rates of herpes zoster (HZ) in Olmsted County, MN, adults from 1996 to 2001, in 2-y increments.

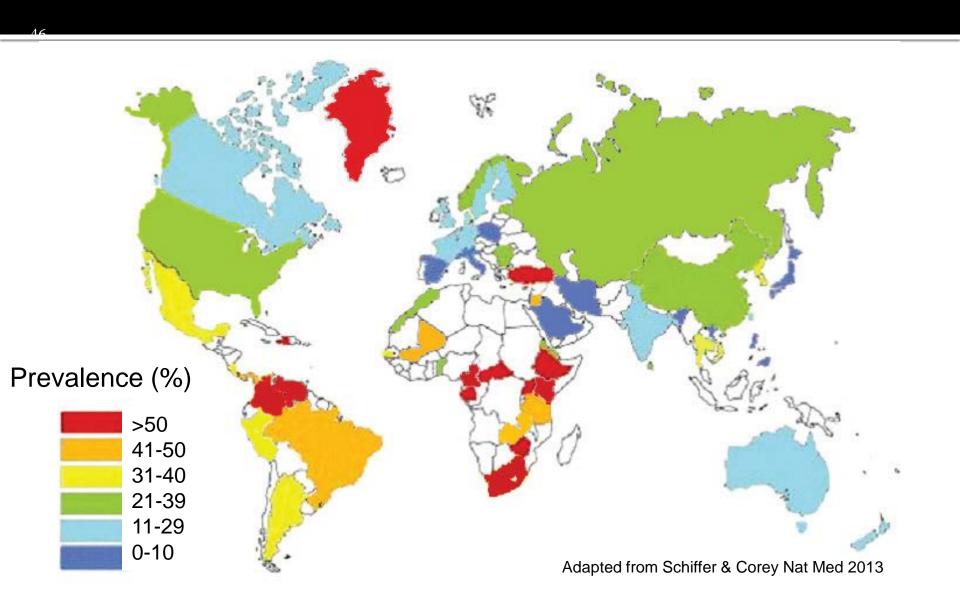


Yawn B P et al. Mayo Clin Proc. 2007;82:1341-1349





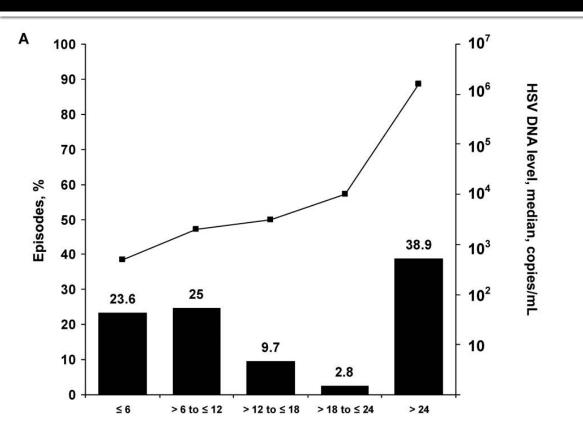
#### HSV-2 seroprevalence in women



## Tips: use of graphics

- Remember figure legends
- Explain axes
  - Take time to orient people to slide so they can interpret findings

#### HSV-2 shedding episodes are short



Median episode duration=13 hrs

HSV-2 reactivation occurs 3 times more often than previously thought

HSV-2 is leaked from neurons at a near constant rate

Mark et al, J Infec Dis 2008; 198:1141 Schiffer et al, Sci Transl Med 2009; 1(7)

## Delivery

- Appearance
  - Dress appropriately for talk
- Mechanics
  - Aviod spelng mistaeks
  - Cite references appropriately
    - At beginning or end of talk, or on each slide

#### Delivery: 3 "E"s

- Eye contact
  - Goal: Maintain with minimal use of notes
- Engagement
  - Goal: Involve audience with different strategies
- Elocution
  - Goal: Clear, articulate presentation
    - Speak loudly
    - Avoid monotone
    - Avoid verbal clutter
      - Um, ah, you know

## Delivery

- Movement
  - Avoid excessive movement
- Posture
  - Relaxed, facing audience
    - Do not face screen

## Evolving in public speaking

- Read your reviews, and learn from them
- Ask for honest evaluation from mentors, peers
- Video and review your presentation:
  - Identify verbal clutter
  - Identify nervous behaviors
  - Worth the pain

# Giving an effective oral presentation

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- Informative
- Relevant

#### **SPEAKER**

- Energetic
- Charismatic
- Passionate
- Humorous
- Knowledgeable

#### Conclusions

- Many opportunities to hone oral presentation skills during fellowship
  - Oral presentation skills are essential for academic research career
- It will take time to learn what strategies work best for you
  - Learn from each talk