


49 – HSV and VZV in Immuno-Competent and Immunocompromised Hosts


Speaker: Richard Whitley, MD



HSV and VZV in Immunocompetent and Immunosuppressed Patients

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7/6/2023



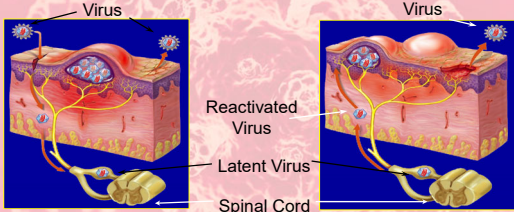
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Herpes Viruses: The Family

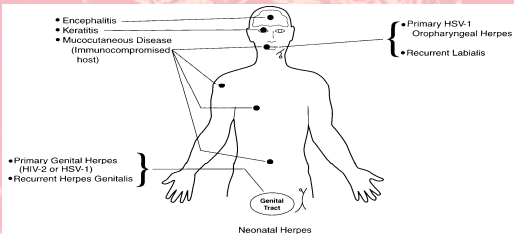
- Herpes simplex virus, type 1 (HSV-1)
- Herpes simplex virus, type 2 (HSV-2)
- Varicella zoster virus (VZV)
- Cytomegalovirus (CMV)
- Epstein Barr virus (EBV)
- Human herpesvirus 6 (HHV 6 A and B)
- Human herpesvirus 7 (HHV 7)
- Human herpesvirus 8 (HHV 8)

Viral Latency and Reactivation



Netter FH. ©2001 by Icon Learning Systems.

Clinical Manifestations of Herpes Simplex Virus Infections

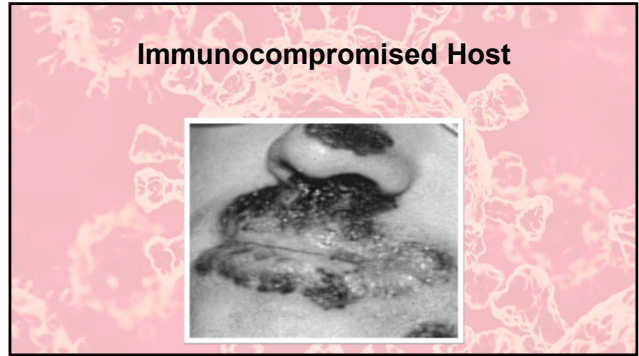
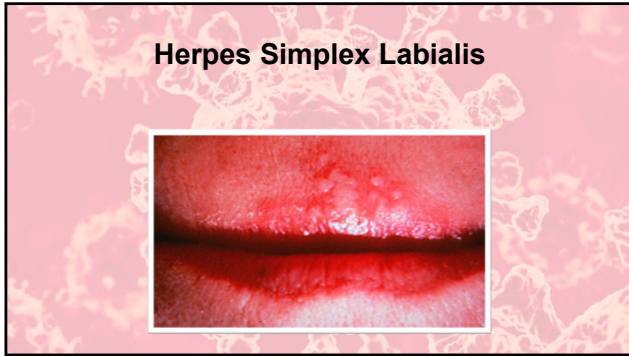


Primary Herpes Simplex Virus Infection: Cutaneous Lesions

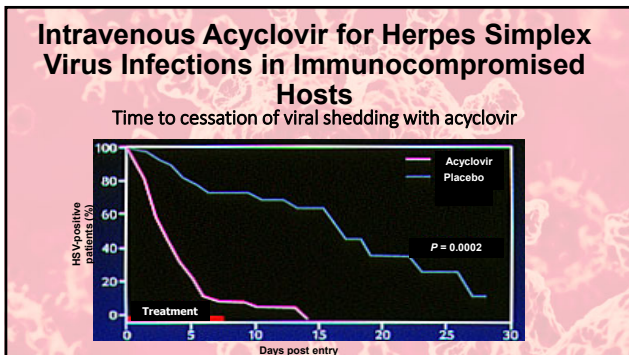
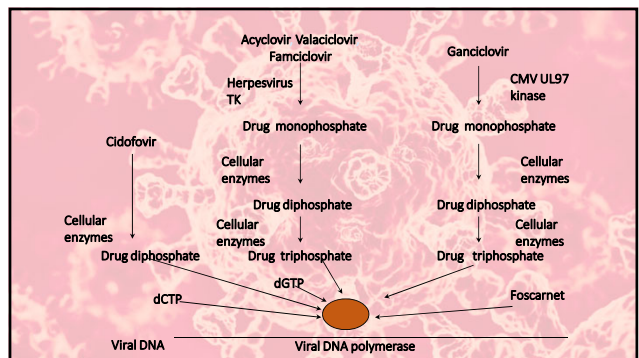


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- ### Most Widely Used Systemic Anti-HSV and VZV Drugs
- Acyclovir (ACV, Zovirax)
 - Famciclovir (FCV, Famvir)
 - Valacyclovir (VACV, Valtrex)
 - Foscarnet (PFA, Foscavir)
 - Ganciclovir (GCV, Cytovene)
 - Val-Ganciclovir (Valcyte)
 - Others:
 - Cidofovir



Acyclovir Prophylaxis for HSV Infection in BMT Patients

Acyclovir (250 mg iv/m2 /tid) or placebo for 18 days beginning 3 days before transplant

Group	Number of Patients	Number of HSV Infections	P
Acyclovir	10	0	~0.003
Placebo	10	7	

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Question #1

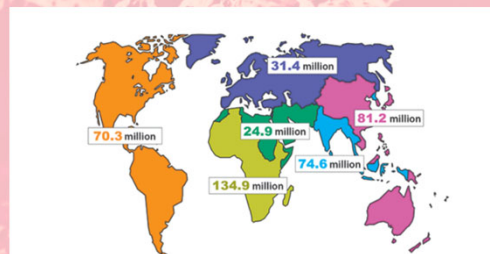
A 30 year old heart transplant has received acyclovir for the past 60 days for cutaneous HSV infection. The lesions are now progressive in spite of high-dose intravenous therapy. The most likely cause for disease progression is a deficiency or alteration of:

- A. Ribonucleotide reductase
- B. Reverse transcriptase
- C. Protease
- D. Thymidine kinase
- E. DNA polymerase

Answer #1a and b

- Three types of acyclovir resistant viruses:
 - thymidine kinase negative
 - thymidine kinase altered substrate
 - DNA polymerase mutations
- All populations of HSV contain viruses with resistant genotypes
- Progressive disease has been limited to the immunocompromised host, especially HSCT recipients and those with poorly controlled HIV
- Three normal hosts with documented ACV resistant virus had disease progression

Global Prevalence of HSV-2 Infection



Total estimated number of people (in millions) infected with HSV-2 in 2012 by WHO region, gender and age ranges. Source: WHO, as published in PLOS ONE (21 Jun 2015)

Acyclovir Therapy of Genital Herpes

Summary of clinical benefit for treatment of:

- Primary
- Recurrent
- Suppressive

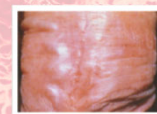
Spectrum of HSV Clinical Presentation



First infection



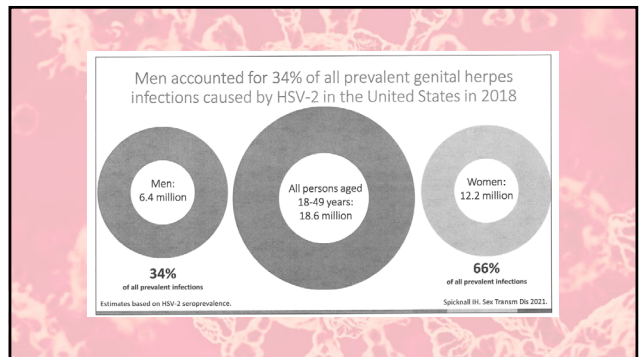
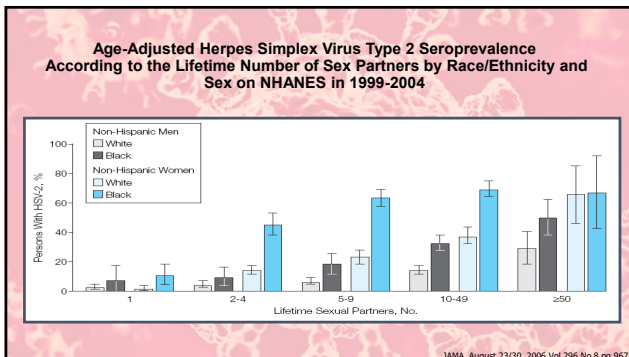
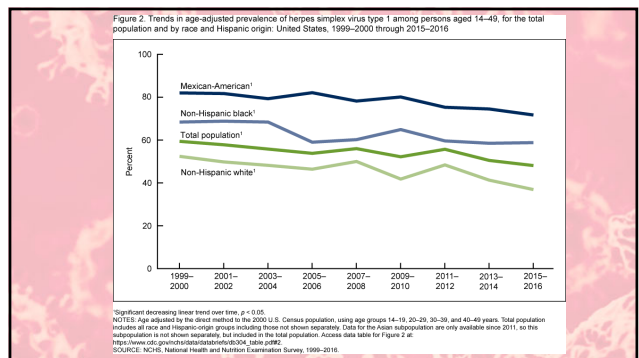
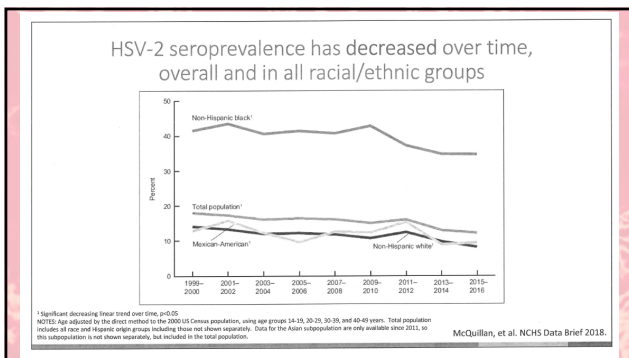
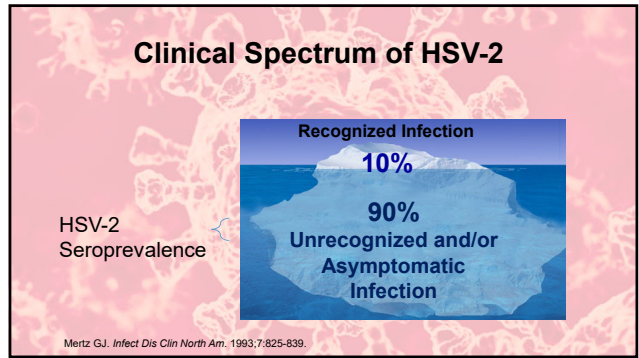
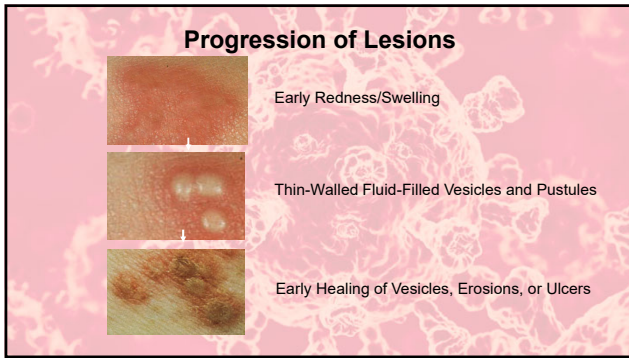
Classical recurrence



Atypical recurrence

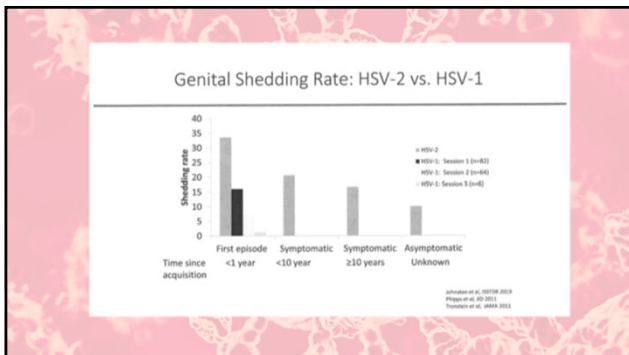
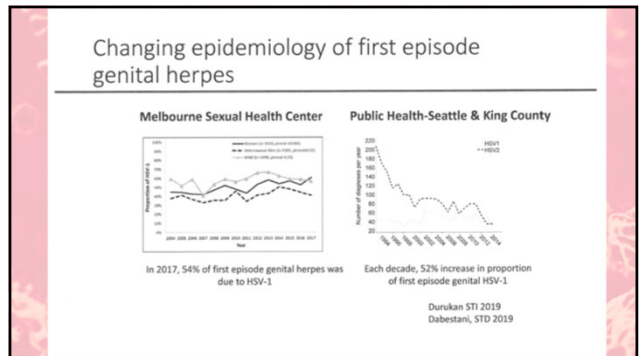
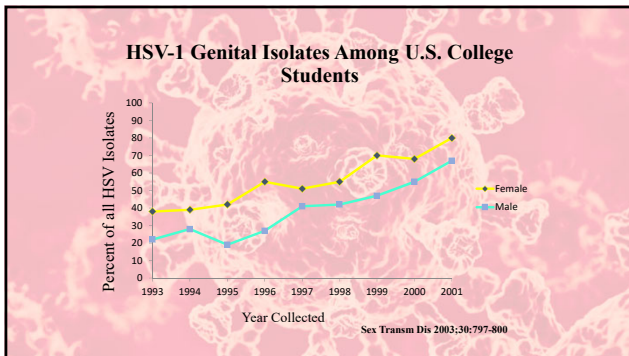
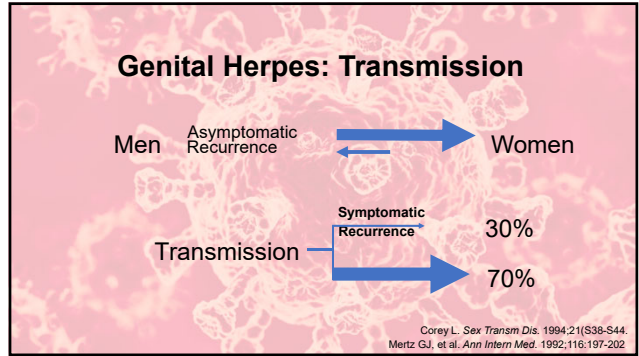
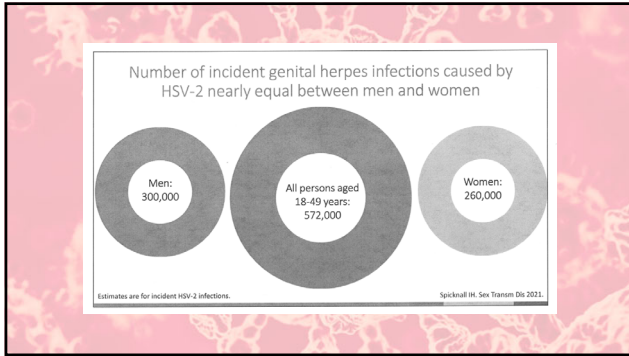
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- ### Genital Herpes: Viral Shedding
- Duration is longer in primary than in recurrent episodes
 - Higher rates in
 - People with frequent outbreaks
 - First year after acquisition
 - Primary: 12 days
 - Recurrent: 2-3 days
 - Oral antiviral suppressive therapy shortens the duration of, but does not eliminate, viral shedding
- Genital Herpes – A Clinician’s Guide to Diagnosis and Treatment, American Medical Association, 2001:1-20. Whitley RJ, et al. Clin Infect Dis 1998;26:541-555.

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Herpes Presenting as Ulceration



- The patient had been to her doctor 3 times over the past 8 months with this pruritic and mildly painful rash on her right buttock. She had been told that it was an irritation from riding a bicycle.
- What is the key to the diagnosis?
 - A. the fact that lesions recurred
 - B. site of involvement is not unusual
 - C. trauma can induce reactivation

Photo courtesy of Jeffrey Gilbert, MD

Question #2

INFECTIOUS DISEASE BOARD REVIEW 2023

PREVIEW QUESTION

An 18 year old man presents with a history of malaise, low-grade fevers, and new-onset painful genital lesions seen in the picture below. He had unprotected sexual intercourse with a female partner 2 weeks earlier. Neither he nor his partner has traveled outside the United States.



Which of the following diagnostic tests is most likely to yield the specific diagnosis?

- A. Serum RPR
- B. Serum FTA-Abs
- C. Darkfield microscopy
- D. Glycoprotein-G 1 serum antibodies
- E. PCR on lesion swab

Answer #2

INFECTIOUS DISEASE BOARD REVIEW 2023

PREVIEW QUESTION

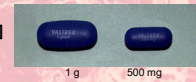
- Historically, culture of HSV was the gold standard. Using daily cultures to detect viral shedding resulted in 4-7% of all days being positive.
- Use of PCR has supplemented culture and detects shedding in up to ~25% of days. More recent data show intermittent shedding on the same day.
- A culture isolate of virus is required to test for resistance
- Serology can be used to assess prior exposure to HSV. The distinction between HSV glycoprotein 1 and 2 is diagnostic.

Oral Antiviral Therapies

- Famciclovir [Famvir®]



- Valaciclovir [Valtrex®]



- Acyclovir [Zovirax®]

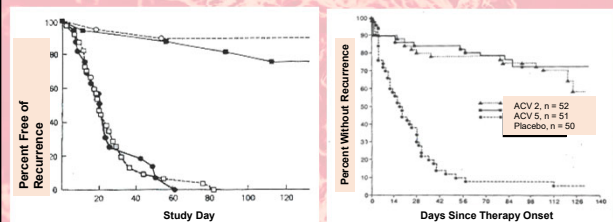


Valtrex® and Zovirax® are registered trademarks of GlaxoSmithKline.

Impact of Acyclovir Therapy on Primary Genital HSV Infection

	Treatment Group (Days)		RR	P
	Acyclovir	Placebo		
Virus Shedding	2.8	16.8	6.82	0.0002
Pain	8.9	13.1	2.00	0.01
Scabbing	9.3	13.5	2.21	0.004
Healing	13.7	20.1	1.83	0.04

Effect of Acyclovir Prophylaxis on Recurrent Genital Herpes



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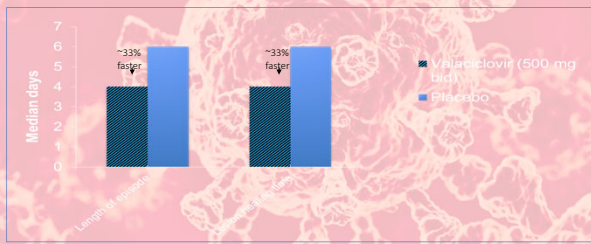
Second Generation Anti-Herpetic Medications

- Valacyclovir (prodrug of acyclovir)
- Famciclovir (prodrug of penciclovir)

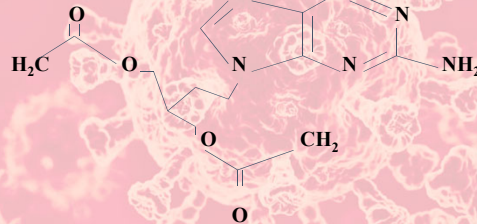
Acyclovir/Valacyclovir Kinetics

DRUG	DOSE	PHARMACOKINETICS	
		C_{max} ($\mu\text{g/mL}$)	Daily AUC ($\mu\text{g/mL}\cdot\text{h}$)
VALTRESX	1 g 3x/d	5.0	47
Oral ZOVIRAX	800 mg 5x/d	1.6	24
IV ZOVIRAX	5 mg/kg 3x/d	9.8	54
	10 mg/kg 3x/d	20.7	107

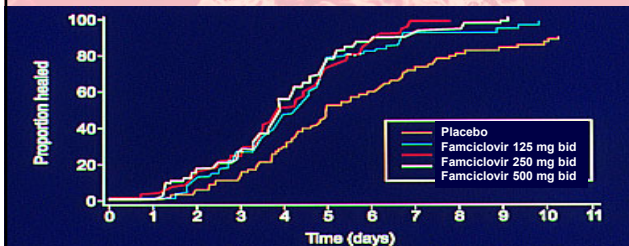
Therapy of Recurrent Genital Herpes: Duration of Disease



Famciclovir



Famciclovir Therapy of Recurrent Genital Herpes



Shorter and Shorter Therapy

- Genital Herpes
 - Valacyclovir: three days
 - Famciclovir: one day
- Labial Herpes
 - Valacyclovir: two days
 - Famciclovir: one day

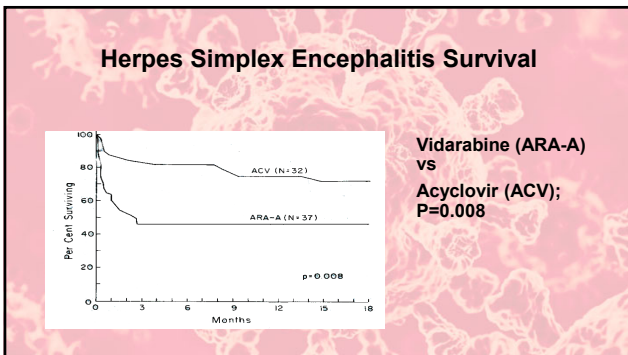
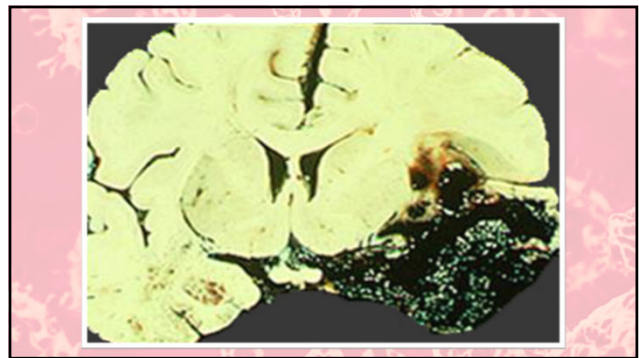
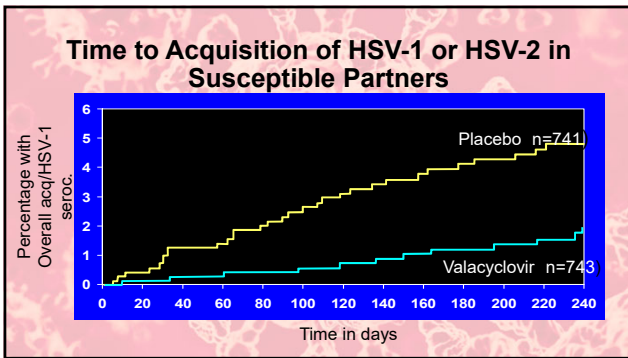
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Valacyclovir Prevention of HSV Transmission to Susceptible Partners

Susceptible Partner	Val-ACV N = 743	Placebo N = 741	Total
No. acquired HSV-2	14	28	42
No. acquired HSV-1	0	4	4
No. developed clinical HSV-2	4	17	21



HSE Morbidity

Percent Patients Patient Normal / Mild Impairment

Age	Glasgow Coma Scale <6	Glasgow Coma Scale ≥6
<30	0	60
>30	0	36

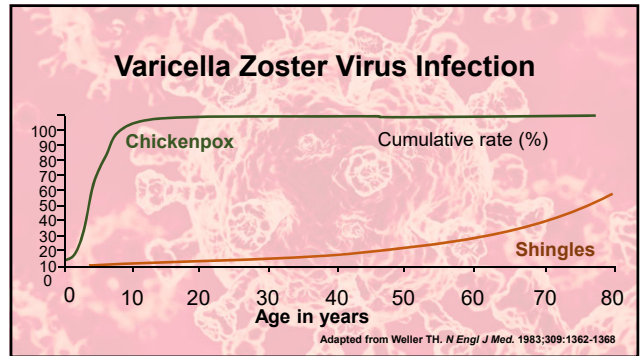
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Sensitivity and Specificity of PCR

	Biopsy Positive	Biopsy Negative
PCR Positive	53	3
PCR Negative	1	44

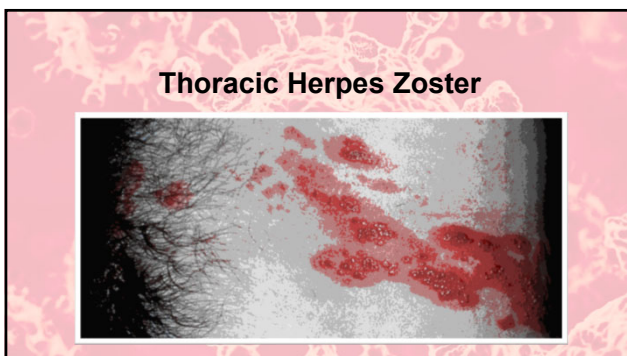
Sensitivity 98%
Specificity 94%
Positive Predictive Value 95%
Negative Predictive Value 98%



CHICKEN POX: Is Therapy of Value?


Treatment of Chicken Pox: Adults (>18 Years) < 24 Hour Duration

	Acyclovir (n=38)	Placebo (n= 38)	p
Time to maximum number of skin lesions (days)	1.5	2.1	0.002
Days of new lesion information	2.7	3.3	0.03
Time to onset of cutaneous healing (days)	2.6	3.3	<0.001
Time to 100% crusting (days)	5.6	7.4	0.001
Maximum number of lesions	268	500	0.04



Questions

1. What is the most likely diagnosis?
2. How would you prove the etiology?



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Answer

- Clinically this is herpes zoster
- The lesion shown is Tzank prep positive on skin scraping. The sensitivity of this test is only ~60% and, therefore, is not recommended
- Immunofluorescence is positive for VZV, having a sensitivity of ~80%.
- Preferably, PCR can be performed even when lesions are scabbed and has the highest sensitivity.

Question #3

What complication would you be most concerned about?

- A. Facial paralysis
- B. Keratitis
- C. Encephalitis
- D. Optic neuritis
- E. Oculomotor palsies



<http://www.itfnoroloi.org/kranjalnoropatiler/Kranjalnoropatiler.html>

Answer: #3

- This patient has Ramsay Hunt syndrome (Herpes zoster oticus), caused by VZV reactivation in the geniculate ganglion, i.e. zoster of CN VII, presenting with severe ear pain and reduced hearing or deafness. When vesicle are seen in the auditory canal, abnormalities in cranial nerves VII, and sometimes VIII, IX or X, can occur. Thus A, facial paralysis is the best answer. Acyclovir is usually recommended although its not clear if it's effective. The facial paralysis is more severe and less likely to resolve than the usual HSV related Bells Palsy.
- Keratitis would be more typical of a lesion on the tip of the nose, or zoster ophthalmicus involving the CN V ophthalmic branch.
- Encephalitis can be caused rarely by VZV and would not be the best answer. Stroke syndromes due to carotid intimal involvement are associated with zoster, and often with cranial nerve V (trigeminal involvement), but are not offered as an answer
- Optic neuritis and oculomotor paralysis would be uncommon.

Question #4 Stem

The patient has only the observed finding on his nose.

- What is your most likely diagnosis?
- What is the name of this sign?



www.medscape.com

Question #4

What complication is it most likely to be associated with this illness?

- A. Deafness
- B. Vertigo
- C. Optic neuritis
- D. Keratitis
- E. Stroke

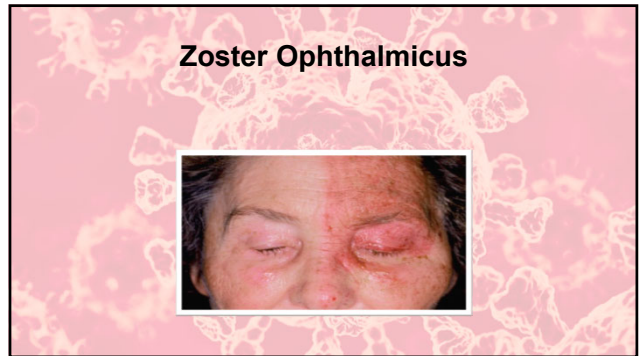
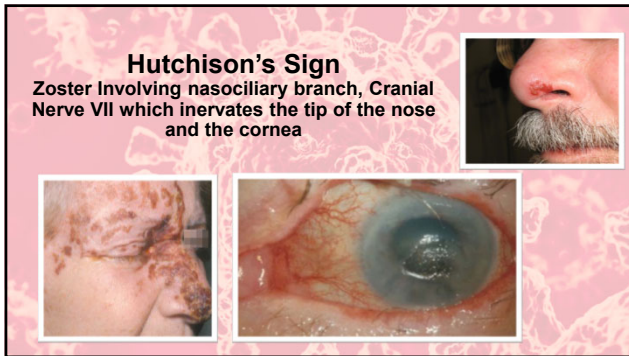
Answer: #4

This patient has Hutchinson's sign, which indicates involvement of the cranial nerve V, i.e. ophthalmic branch of the trigeminal nerve, which innervates the tip of the nose and the globe. After a prodrome of fever and headache for 1-4 days, patients develop a cutaneous rash. Days or up to 3 weeks later, the sclera and cornea can be involved. Thus, keratitis is the correct answer.

Deafness or vertigo would be more characteristic of geniculate ganglion (CN VII) involvement, i.e. Ramsay Hunt, which is a polyneuropathy involving the cranial nerve VII, and then often involves VIII, IX, X. Thus A and B are not the best answers.

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NATURAL HISTORY OF ZOSTER IN THE NORMAL HOST

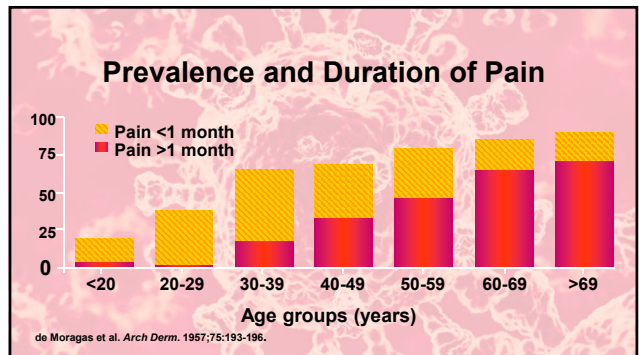
- Acute neuritis may precede rash by 48 - 72 hours
- Maculopapular eruption, followed by clusters of vesicles
- Unilateral dermatomal distribution

NATURAL HISTORY OF ZOSTER IN THE NORMAL HOST

- Events of healing:
 - Cessation of new vesicle formation: 3 - 5 days
 - Total pustulation: 4 - 6 days
 - Total scabbing: 7 - 10 days
 - Complete healing: 2 - 4 weeks
- Cutaneous dissemination can occur dissemination is extremely rare
- Postherpetic neuralgia in 10% - 40% of cases

Complications of Zoster

Common	Uncommon
• Postherpetic neuralgia	• Cutaneous dissemination
• Ocular complications	• Herpes gangrenosum
• Ophthalmic zoster	• Hepatitis
• (uveitis, keratitis, scleritis, optic neuritis)	• Encephalitis
• Pneumonitis	• Motor neuropathies
• Scarring	• Myelitis
• Bacterial superinfection	• Hemiparesis (granulomatous CNS vasculitis)



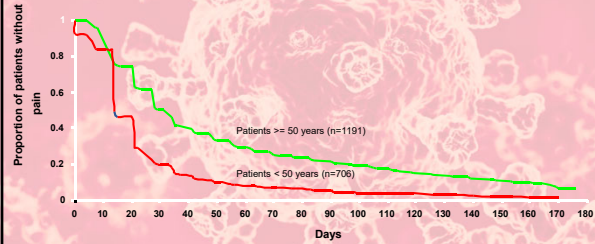
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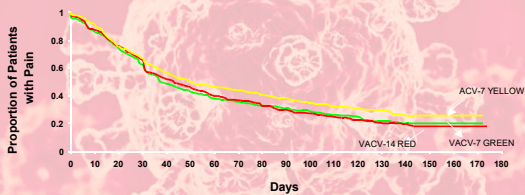
Goals of Therapy

- Accelerate cutaneous healing
- Accelerate loss of pain acute / chronic
- Prevent complications

Time to Cessation of Zoster-Associated Pain

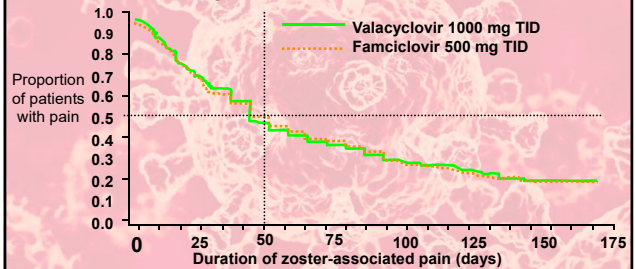


Time to Cessation of Zoster Associated Pain n = 1141



* Beutner, et al. Acyclovir versus Valacyclovir in the treatment of herpes zoster in patients > 50 years old.

Resolution of Pain in Herpes Zoster With Valacyclovir and Famciclovir



Summary of Efficacy of Concomitant Steroid Therapy with Acyclovir

- Accelerates resolution of acute neuritis
- Accelerates:
 - Return to usual activity P<0.001
 - Unaroused sleep P<0.0001
 - Cessation of analgesic use P<0.001
- Effect on chronic pain P=0.06

Question #5

What is the most likely etiologic agent?



- A. HSV
- B. VZV
- C. CMV
- D. EBV
- E. HHV6

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Answer #5

- This patient has facial palsy, also known as Bells palsy. The most likely cause of this lesion is HSV. HIV and Lyme disease are less common causes. Answers d and e are not the best answer. Of note, Lyme is rarely the cause of Bells palsy unless there are other manifestations of Lyme disease.
- For typical facial palsy, prednisone is the preferred therapy, optimally given within 3 days of onset, for one week (prednisone 60-80mg qd). Acyclovir alone is not better than placebo, although there might be some rational (unproven) to add acyclovir to prednisone.
- Ganciclovir would be a therapy for CMV, a rare cause of facial paralysis and thus not the best answer.

METHODS OF PREVENTING / MODIFYING VARICELLA

- Pre-exposure: Oka varicella vaccine
- Post-exposure: VZIG (now available in US)
- Oka varicella vaccine (<3 days after exposure)
- Acyclovir (7-14 days after exposure)

Shingles Prevention Trial: Zostavax

Attenuated, live virus (approved 2006)

- Efficacy but waning of immunity with time
 - Burden Of Illness 61.1% (51.1 – 69.1%)
 - Post-Herpetic Neuralgia 66.5% (47.5 – 79%)
 - Incidence of Herpes Zoster 51.3% (44.2 – 57.6%)

Second Generation Vaccine: Shingrix

- Recombinant adjuvanted vaccine
 - Two shots
 - > 50 years of age
- Efficacy
 - Both PHN and incidence of shingles
 - >90% for >4 years
- Adverse events
 - Local reactogenicity: redness and pain ~ 50-70%
 - Systemic malaise/fever: ~30%

Thank You
rwhitley@uab.edu