

Measles

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Updated 2013 Oct 07 11:37:00 AM: Society for Healthcare Epidemiology of America (SHEA) guideline on infection prevention and control in residential facilities for pediatric patients and their families (Infect Control Hosp Epidemiol 2013 Oct) [view update](#) | [Show more updates](#)

- [General Information](#)
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Complications:

- o complications rare, occur especially in malnourished or immunocompromised children
- o thrombocytopenic purpura, 5-15% pneumonia (primary and secondary bacterial), bacterial otitis media, croup, bronchitis, bronchiolitis, diarrhea
- o severe mouth ulceration - usually due to secondary herpes infection, treat with regular mouth washes with clean water and apply 1% gentian violet to mouth lesions
- o rarely interstitial giant cell pneumonia, conjunctivitis, myocarditis, hepatitis, acute glomerulonephritis, 1/1,000 encephalomyelitis (10% mortality), pericarditis
- o subacute sclerosing panencephalitis (SSPE)
 - defective M protein spike affecting central nervous system, especially limbic system
 - incidence < 6 per million in most countries, but 56 per million in Papua New Guinea in 1990, possibly due to highly neurotropic measles virus ([Epidemiol Infect 2003 Oct;131\(2\):887](#) in [BMJ 2003 Nov 15;327\(7424\):1176](#))
 - case presentation of subacute sclerosing panencephalitis secondary to measles virus infection can be found in [N Engl J Med 2007 Aug 9;357\(6\):589](#)
- o delayed-type hypersensitivity (DTH) against tuberculin may be lost temporarily (measles can suppress T cell function)
- o most deaths follow complications and often associated with malnutrition
- o 3 cases of acute renal failure with neurologic impairment in adults have been described ([Lancet 1999 Sep 18;354\(9183\):992](#) [EBSCOhost Full Text](#))
- o measles inclusion body encephalitis is rare complication in immunocompromised host typically 1 year after measles illness or vaccination, case report in patient with stem cell transplantation ([Pediatrics 2004 Nov;114\(5\):e657](#) [EBSCOhost Full Text full-text](#))

Associated conditions:

- o atypical measles syndrome with previous exposure to killed vaccine
 - high fever, pneumonia, rash (urticarial then maculopapular with petechiae)
 - mainly in patients immunized with killed measles vaccine, given from 1963-1967
 - extremely high measles Ab titer early in illness supports diagnosis, viral culture useless
- o measles associated with increased lifetime risk for atopy (eczema, rhinitis, asthma) in cross-sectional nationwide study in Finland of 547,910 persons 14 months to 19 years ([2000 Jan 19;283\(3\):343](#)), editorial can be found in [JAMA 2000 Jan 19;283\(3\):394](#)

TOPIC OUTLINE

SUMMARY & RECOMMENDATIONS

INTRODUCTION

EPIDEMIOLOGY

- Measles infection
- Measles complications

TRANSMISSION

EFFECT OF IMMUNITY

CONTROLLING MEASLES

- The Americas
 - United States
- European Region
- African Region
- Western Pacific Region
- Eastern Mediterranean Region
- Southeast Asia Region

INFORMATION FOR PATIENTS

SUMMARY AND RECOMMENDATIONS

REFERENCES

RELATED TOPICS

Clinical presentation and diagnosis of measles

Patient information: Measles (The Basics)

Prevention and treatment of measles

a protective immune response. Travel in the developing world or contact with individuals arriving from the developing world increases the risk of exposure to measles. (See '[Effect of immunity](#)' above.)

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REFERENCES

1. Moss WJ, Griffin DE. Measles. *Lancet* 2012; 379:153.
2. Black, FL. Measles. In: *Viral infections in humans: Epidemiology and control*, Evans, AS, Kaslow, RA (Eds), Plenum Publishing, New York 1997. p.507.
3. BABBOTT FL Jr, GORDON JE. Modern measles. *Am J Med Sci* 1954; 228:334.
4. Global measles mortality reduction and regional elimination: a status report. *J Infect Dis* 2003; 187(Suppl 1):S1.
5. LANGMUIR AD. Medical importance of measles. *Am J Dis Child* 1962; 103:224.
6. Markowitz, LE, Katz, SL. *Vaccines*, Plotkin, SA, Mortimer, EA (Eds), WB Saunders, Philadelphia 1994. p.229.
7. Arya LS, Taana I, Tahiri C, et al. Spectrum of complications of measles in Afghanistan: a study of 784 cases. *J Trop Med Hyg* 1987; 90:117.
8. Papania M, Baughman AL, Lee S, et al. Increased susceptibility to measles in infants in the United States. *Pediatrics* 1999; 104:e59.
9. Maldonado YA, Lawrence EC, DeHovitz R, et al. Early loss of passive measles antibody in infants of mothers with vaccine-induced immunity. *Pediatrics* 1995; 96:447.
10. Markowitz LE, Albrecht P, Rhodes P, et al. Changing levels of measles antibody titers in women and children in the United States: impact on response to vaccination. Kaiser Permanente Measles Vaccine Trial Team. *Pediatrics* 1996; 97:53.
11. Stein CE, Birmingham M, Kurian M, et al. The global burden of measles in the year 2000—a model that uses country-specific indicators. *J Infect Dis* 2003; 187 Suppl 1:S8.
12. Centers for Disease Control and Prevention (CDC). Global measles mortality, 2000-2008. *MMWR Morb Mortal Wkly Rep* 2009; 58:1321.
13. Pan American Health Organization. Immunization in the Americas. http://new.paho.org/sur/index.php?option=com_content&task=view&id=91&Itemid=290 (Accessed on November 08, 2010).
14. World Health Organization. Vision and Strategy 2006-2015 www.who.int/vaccines-documents/DocsPDF05/GIVS_Final_EN.pdf (Accessed on October 13, 2011).
15. World Health Organization. WHO/UNICEF joint statement -- global plan for reducing measles mortality 2006-2010. http://www.who.int/immunization/documents/WHO_IVB_05.11/en/index.html (Accessed on November 03, 2010).
16. WHO-recommended surveillance standard of measles. http://www.who.int/immunization_monitoring/diseases/measles_surveillance/en/index.html (Accessed on November 08, 2010).
17. Simons E, Ferrari M, Fricks J, et al. Assessment of the 2010 global measles mortality reduction goal: results from a model of surveillance data. *Lancet* 2012; 379:2173.
18. <http://www.measlesinitiative.org> (Accessed on November 15, 2010).

UPTODATE

for use in individuals who are immunocompromised or who have CD4 counts < 200; it is not recommended for patients who are HIV-positive or who have AIDS. However, a randomized clinical trial is currently evaluating the use of Zostavax in patients with HIV disease^(2,3,4,5,7)

What We Can Do

- Learn about HZ and HIV infection so you can accurately assess your patients' personal characteristics and health education needs; share this information with your colleagues
 - Maintain universal precautions when treating all patients with HZ
 - Clinical staff members who are susceptible to VZV should not care for patients with VZV infections
- Closely monitor patients with HZ and HIV infection or AIDS for potential complications and adverse reactions to the prescribed drug regimen
- Educate patients about prevention
 - Recommend that patients who are HIV-positive who are exposed to VZV and who have no history of chickenpox, negative anti-varicella immune globulin (IgG) blood test, or both, receive the varicella **zoster** IgG, ideally within 48 hours, but no later than 96 hours, after exposure
 - Discuss the varicella vaccine with your patients who have not had chickenpox; encourage them to talk with their treating clinician about receiving the vaccine, as appropriate; vaccinate patients, as prescribed
- Provide patients with written information on HIV and HZ—including treatment risks and benefits and potential complications—and emphasize the importance of continued medical surveillance and seeking immediate medical attention for adverse drug effects or new or worsening signs and symptoms
- Provide emotional support to patients who are HIV-positive and have HZ; request referral to a mental health clinician for counseling on coping strategies, as appropriate
- Request referral to a social worker for identification of local resources for appropriate support groups, educational programs, in-home services, or hospice
- Collaborate with your hospital's continuing education department to provide education about HZ and HIV for clinicians of all specialties

References

1. DynaMed. (2012, December 3). **Zoster**. Ipswich, MA: EBSCO Publishing. Retrieved from <http://search.ebscohost.com/login.aspx?direct=true&db=dme&AN=113997> (GI)
2. Erbeling, E. J., & Ghanem, K. G. (2011). Johns Hopkins HIV guide: **Herpes zoster**. Johns Hopkins POC-IT Guides. Retrieved from http://www.hopkinsguides.com/hopkins/sub/view/Johns_Hopkins_HIV_Guide/545093/all/Herpes_zoster (GI)

NURSING REFERENCE CENTER

SHINGLES

EVIDENCE BASED CARE SHEETS

Food for Thought

- The benefit of vaccinating patients who have already experienced a bout of HZ has been unclear; now, researchers who conducted a recent retrospective study of 1,036 immunocompetent patients 60 years of age or older found that varicella vaccination may not reduce the short-term risk of recurrent HZ (Tseng et al., 2012)

Red Flags



Zostavax should not be given to patients who have had anaphylactic reaction to gelatin, neomycin, or other components of the vaccine; have a weakened immune system (e.g., due to blood or bone cancer or to HIV/AIDS with T-cell count < 200); or are receiving medical treatments that suppress the immune system



Varivax should not be given to women for 3 months prior to pregnancy or during pregnancy

What Do I Need to Tell the Patient/Patient's Family?

- Review with patient/family the signs and symptoms of HZ, its complications, and adverse effects of medications; advise patients to seek immediate medical attention for new or worsening signs or symptoms; emphasize the importance of regular medical surveillance
- Advise patients to get plenty of rest and to avoid strenuous activities while recuperating; take a cool bath or use wet compresses on blisters to relieve itching and pain; and take nonprescription analgesics (e.g., ibuprofen, naproxen), use calamine lotion, and oral antihistamines (diphenhydramine)

References

1. DynaMed. (2012). *Zoster*. Ipswich, MA: EBSCO Publishing.

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- [+] Rifampin
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cyclosporine, digoxin, doxycycline, fluoroquinolones, nifedipine, zolpidem, quinidine, corticosteroids, hormonal contraceptives, methadone, oral anticoagulants, oral sulfonylureas, theophyllines, phenytoin, cyclosporine, ketoconazole, verapamil

★ **Drug-lab test** • Rifampin inhibits standard assays for serum folate and vitamin B₁₂

■ **Nursing Considerations** 

Assessment

- **History:** Allergy to any rifamycin, acute hepatic disease, pregnancy, lactation
- **Physical:** Skin color, lesions; T; gait, muscle strength; orientation, reflexes, ophthalmologic examination; liver evaluation; CBC, LFTs, renal function tests, urinalysis

Interventions

- Administer on an empty stomach, 1 hr before or 2 hr after meals.
- Administer in a single daily dose.
- Consult pharmacist for rifampin suspension for patients unable to swallow capsules.
- Prepare patient for the reddish-orange coloring of body fluids (urine, sweat, sputum, tears, feces, saliva); soft contact lenses may be permanently stained; advise patients not to wear them during therapy.

⊗ **Warning** Arrange for follow-up visits for liver and renal function tests, CBC, and ophthalmologic examinations.

Teaching Points

- Take drug in a single daily dose. Take on an empty stomach, 1 hour before or 2 hours after meals.
- Take this drug regularly; avoid missing any doses; do not discontinue this drug without consulting your health care provider.
- Have periodic medical checkups, including eye examinations and blood tests, to evaluate the drug effects.
- You may experience these side effects: Reddish orange coloring of body fluids (tears, sweat, saliva, urine, feces, sputum; stain will wash out of clothing, but soft contact lenses may be permanently stained; do not wear them); nausea, vomiting, epigastric distress; skin rashes or lesions; numbness, tingling, drowsiness, fatigue (use caution if driving or operating dangerous machinery; use precautions to avoid injury).
- Report fever, chills, muscle and bone pain, excessive tiredness or weakness, loss of appetite, nausea, vomiting, yellowing of skin or eyes, unusual bleeding or bruising, skin rash or itching.

LIPPINCOTT'S NURSING DRUG GUIDE

RIFAMPIN