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IN THE DECEMBER 1998 ISSUE:

- Mycoplasma pneumoniae Hits Seattle School
- Wrestling with Herpes Infections
- Thinking of Influenza Vaccine? Think Pneumococcal too!
- Immune Globulin Supply Back on Track

Mycoplasma Cluster

Seattle-King County The of Public Department Health (SKCDPH) Communicable Disease Epidemiology Unit has been investigating an unusual cluster of Mvcoplasma pneumoniae infections at an elementary school in Seattle. Between September 1 and November 11 of this year, 97 persons associated with the school have had a respiratory illness characterized by fever, sore throat and prolonged cough. Fifty persons met a case definition of either one week of cough illness with a chest xray positive for pneumonia, or fever and cough of ten days or Other symptoms more duration. have included otitis media (two persons had bullous myringitis) and rash. To date, four persons have been positive for M. pneumoniaespecific IgM antibody by serologic testing and one person has tested positive for M. pneumoniae by polymerase chain reaction (PCR) on a throat swab. Serology is pending on nine additional specimens and PCR testing is pending on throat swabs from three other persons.

The majority of ill students were in the first and third grades, however cases also included family members of students and school staff. Actions taken by SKCDPH to decrease transmission of М. pneumoniae included enhancing early recognition and treatment of infection through educating parents providers regarding and the symptoms; appropriate diagnostic tests and antimicrobial treatment of М. pneumoniae infection: encouraging adherence to a four times per day handwashing regimen; reminding coughing persons to avoid spreading secretions respiratory and contaminating hands by holding the crook of the elbow over the mouth when coughing or sneezing; and reinforcing the value of maximizing

ventilation in the school and home environment.

Our investigation is continuing to better understand the epidemiology and transmission of М initial pneumoniae. An questionnaire has been administered, and a second one will be sent to homes where household transmission may have occurred. Distinguishing infection with M. pneumoniae from other causes of respiratory tract infection is difficult. Laboratory testing of new cases is important in order to confirm the diagnosis. The most useful test is M. pneumoniaespecific IgM. Culture and PCR are also useful early in disease. Please call 206-296-4774 for additional information or help in diagnosing M. pneumoniae infection.

Community-wide M. pneumoniae epidemics occur every four to eight years. Although M. pneumoniae is not reportable in the U.S., the last year widespread epidemic activity was reported appears to have been 1993. M. pneumoniae is highly transmissible. Humans are the only known source of infection. Acquisition is presumed to be by droplet spread from symptomatic patients. The incubation period is one to four weeks. The unusually long incubation period complicates epidemiologic investigations and makes effective preventive treatment and cohorting less than optimally effective at interrupting transmission. Familial spread often continues for many months. resulting in cumulative household attack rates that approach 100%. Asymptomatic carriage after infection can occur for prolonged intervals even after treatment with antibiotics. but has not been associated with transmission.

The most common clinical syndromes caused bv М. pneumoniae are acute bronchitis upper respiratory and tract infections, including pharyngitis and occasionally otitis media or Coryza, sinusitis and myringitis. croup are infrequent. Specific disease syndromes are agerelated. M. pneumoniae is the leading cause of pneumonia in school-age children and young adults but is an uncommon cause of symptomatic illness in children younger than five years of age. Initial symptoms are malaise, fever, and sometimes headache. А nonproductive cough develops within a few days and lasts for three to four weeks during which time it may become productive. Approximately 10% of children with pneumonia exhibit a rash. Chest film abnormalities are variable and characteristically are more extensive then the clinical picture would suggest. Acute bronchitis and upper respiratory tract illnesses caused by M. pneumoniae are generally mild and resolve without antibiotic therapy.

Transmissibility is reduced when appropriate therapy is used early in the course of the infection. Erythromycin, clarithromycin and azithromycin are preferred for treatment of *M. pneumoniae* infections in children younger than eight years of age. For persons eight years and older, tetracycline is equally effective and for adults, newer quinolones may be used.

Wrestling with HSV

The SKCDPH recently became aware of several cases of herpes simplex type 1 (HSV-1) infection among high school wrestlers. This infection, called herpes gladiatorum in wrestlers and scrumpox in rugby players, is thought to be one of the most common infections spread by person-to-person contact during athletic activity. HSV-1 appears to be endemic among high school wrestlers. A 1996 investigation by the Snohomish Health District and the Washington State Department of Health found that 21% of 249 high school wrestlers and coaches had either confirmed or possible herpes gladiatorum. Molecular epidemiologic analysis showed three distinct strains of HSV-1 among involved wrestlers

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suggesting that multiple discrete outbreaks occur simultaneously during the wrestling season.

The infection is spread by contact with an opponent's cutaneous lesions or virus-bearing saliva. Transmission by fomites such as wrestling mats and equipment is thought to be insignificant. The herpetic lesions are usually found on the face, scalp and neck of wrestlers and may be accompanied by fever, headache, regional lymphadenopathy, sore throat, and itching or painful eyes. Herpes gladiatorum may be misdiagnosed as staphlococcal or streptococcal infection (impetigo), ringworm eczema, or (tinea The main serious gladiatorum). complication of herpes gladiatorum is ocular involvement, which may manifested as follicular be conjunctivitis or keratoconjunctivitis. Corneal involvement is usually characterized by pain or foreignbody sensation. Herpes keratitis is potentially vision-threatening and symptoms should trigger prompt ophthomologic evaluation. Recurrences of ocular herpes develop in 25% or more of persons with symptomatic primary infection.

Clinicians and school health practitioners should maintain a high index of suspicion for herpes gladiatorum in wrestlers with cutaneous lesions on the face, neck

or scalp with or without systemic symptoms. The diagnosis of HSV-1 is confirmed by culture. Direct fluorescent antibody (FA) is a useful and rapid diagnostic test that has lesser sensitivity than culture and does not distinguish HSV-1 from HSV-2. Specimens for both tests are obtained by vigorously swabbing the base of fresh lesions.

Persons suspected of having herpes gladiatorum should be

referred their health care to provider for evaluation. confirmation, and possibly Appropriate treatment. disease control steps to prevent transmission of HSV-1 include ongoing evaluation of wrestlers and coaches throughout the wrestling season with exclusion of persons symptomatic from wrestling until all lesions are thoroughly healed and crusted. Although usually not indicated for uncomplicated nongenital mucocutaneous herpes infections, acyclovir and related drugs may shorten the time to healing and may be useful in preventing recurrences in selected cases. gladiatorum Herpes is not reportable although the Communicable Disease Epidemiology Unit is available to answer questions on this subject at 206-296-4774.

Thanks to Chris Spitters, MD, MPH, Snohomish Health District, for providing background on the Snohomish herpes gladiatorum investigation.

Pneumovax

Pneumococcal vaccine is recommended for and given to many of the same adults who receive influenza vaccine every fall. Pneumococcal vaccine is a onetime-only dose for most adults, and the vaccine can be given any time during the year.

A second dose of pneumococcal vaccine is now recommended by ACIP (CDC's Advisory Committee on Immunization Practices) ONLY for the following groups:

- 1) Immunocompromised persons 2 to 64 years of age:
- For ages 2 to 10 years, give booster three to five years after the first dose
- For ages 11 to 64 years, give booster five years after the first dose

 Individuals over 65 years of age who were less than 65 years when they received their first dose, give booster if it has been five or more years since the first dose.

IG Supply

According to FFF Enterprises, the main distributor of the national private supply of immune globulin (IG), availability of immune globulin has improved. Health departments and private health care providers can resume ordering IG outside of an outbreak situation and without CDC approval. Providers can order IG directly from FFF Enterprises at 1-800-843-7477.

Until it is certain that the IG supply will be sustained, SKCDPH will continue to limit IG use to communicable disease exposures. Hepatitis A vaccine will continue to be recommended to travelers.

To Report:	(area code 206)		
AIDS			
Tuberculosis	731-4579		
STDs	731-3954		
Communicable	Disease 296-4774		
24-hr Report Lir	ne 296-4782		
Disease Alert:			
CD Hotline			
After hours	682-7321		
http://www.metrokc.gov/health/			

nerpes gladiatorum should be dose						
REPORTED CASES OF SELECTED DISEASES						
SEAT	FLE-KING COU					
		CASES REPORTED IN NOVEMBER		CASES REPORTED THROUGH NOVEMBER		
	1998	1997	1998	1997		
VACCINE-PREVENTABLE DISEASES	0	0	0			
Mumps	0	0	2	4		
Measles	0	0	0	1		
Pertussis	2	8	143	188		
Rubella	0	0	1	1		
SEXUALLY TRANSMITTED DISEASES						
Syphilis	3	0	34	5		
Gonorrhea	59	73	899	826		
Chlamydial infections	229	267	3171	2831		
Herpes, genital	43	57	592	626		
Pelvic Inflammatory Disease	15	13	211	266		
Syphilis, late	1	2	27	39		
ENTERIC DISEASES						
Giardiasis	19	17	244	247		
Salmonellosis	19	14	205	216		
Shigellosis	4	5	84	94		
Campylobacteriosis	14	19	216	304		
E.coli O157:H7	2	5	34	44		
HEPATITIS	_	-	• •			
Hepatitis A	10	23	381	403		
Hepatitis B	2	0	47	35		
Hepatitis C/non-A, non-B	0	0 0	1	2		
AIDS	10	32	216	298		
TUBERCULOSIS	13	5	106	126		
MENINGITIS/INVASIVE DISEASE	10	0	100	120		
Haemophilus influenzae	0	0	1	1		
Meningococcal disease	1	3	13	20		