Teen With Rabies Recovering

CLINICAL

FluMist Found Safe and Effective

The live, attenuated, cold-adapted influenza vaccine, also known as CAIV-T (FluMist), was safe and effective in children aged 60 months to 17 years in the second year of its use for prevention of flu. The CAIV-T vaccine was first available for use during the 2003-2004 season and was designed to contain three flu strains that matched those recommended by the FDA for the annual trivalent inactivated vaccine. Robert B. Belshe, M.D., of St. Louis University and colleagues reviewed a safety trial that included 6,657

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children aged 5-17 years and an efficacy trial including 312 children aged 60-71 months. (Clin. Infect. Dis. 2004;39;920-7). The safety trial evaluated medically attended events within 42 days of vaccine administration. Overall, the frequency of events in four categories—acute respiratory events, acute gastrointestinal events, systemic bacterial infections, and rare events possibly associated with wild-type influenza—was not significantly different between the 4,452 children in the vaccine group and the 2,205 children in a placebo group. Results from the efficacy study showed an efficacy rate of 87% for children aged 60 months and older and no significant increase in the frequency of fevers greater than 37.8° C. In addition, no significant increase in frequencies of runny nose, nasal congestion, vomiting, or muscle aches was noted among the vaccine recipients compared with the placebo group. The data confirm that the efficacy of CAIV-T extends to the youngest children in the age range for which it is currently recommended, Peter F. Wright, M.D., of Vanderbilt University, Nashville, Tenn., said in an accompanying editorial (Clin. Infect. Dis. 2004;39:928-9).

Meningococcal Disease: It's Not Just a College Problem

Adolescents are also at increased risk

With the spectacular success of Hinfluenzae type b (Hib) and pneumococcal conjugate vaccines, N meningitidis is now the leading cause of bacterial meningitis in children and young adults.1-3 Although meningococcal disease is often thought to be associated with college students, ongoing surveillance shows that adolescents and young adults are also at increased risk for contracting this devastating and potentially life-threatening disease.4-6 Data from the Centers for Disease Control and Prevention (CDC) show that following infancy, meningococcal disease rates begin to rise again in early adolescence and peak between the ages of 15 to 24 years.⁵ Young people in this age group share certain

Mean rates of meningococcal disease by age group in the United States, 1996–2002.⁵

risk factors that may account for the

increased risk of infection.4,7



Adapted from Centers for Disease Control and Prevention (CDC). *MMWR Morb Mortal Wkly Rep.* 1996–2002.⁵

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Case in Point

Susan T, a 15-yearold high school student, was taken to the emergency room by her parents because of fever, vomiting, and an abrupt onset of unresponsiveness.

Physical findings included a temperature of 104°F and purpuric lesions on her arms and legs. Blood and CSF specimens were taken and appropriate antibiotics were started. A few hours later she became lethargic and comatose. She developed meningococcal septicemia, which subsequently caused gangrene in her arms and legs. To save her life, she needed bilateral below-the-knee amputation. Because laboratory diagnosis confirmed N meningitidis serogroup C infection, her case was potentially vaccine preventable.



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The consequences of infection

Meningococcal disease in adolescents

progressive and often fatal — even with

compared with younger populations.6,11

Of the survivors, up to 1 in 5 will suffer

from permanent sequelae including

amputation, severe scarring, hearing

the best of care. According to recent

data, case fatality rates are 5 times

higher among 15- to 24-year-olds

can often be devastating

and young adults can be rapidly

An opportunity for improved control of meningococcal disease

Up to 83% of the cases in adolescents and young adults are caused by potentially vaccine-preventable serogroups of *N meningitidis* (C, Y, and W-135).^{46,12} Therefore, routine vaccination of these higher-risk populations might be a reasonable strategy for controlling this devastating disease. However, the shorter duration of protection and lack of booster response after multiple immunizations limits the use of the currently licensed polysac-charide meningococcal vaccine to "peak risk" populations (eg, college students and military recruits).¹³

Building on the success of other conjugate vaccines, the introduction of conjugate meningococcal vaccines may give health-care providers the opportunity to protect adolescents and young adults through their entire period of increased risk, which in turn, could dramatically reduce the incidence of meningococcal disease in the United States.¹⁴

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A teenaged girl who contracted rabies of from a bat and received an experimental

treatment has been upgraded to fair condition at the Children's Hospital of Wisconsin (Wauwatosa), a hospital spokesperson said in an interview at press time. The girl is the first known person to survive rabies without receiving a vaccine. The bat bit the girl on Sept. 12, 2004. She reportedly thought that the bite was just a scratch, and she and those with her assumed, incorrectly, that only healthy bats could fly, so she did not see a doctor for a vaccine. She presented to Children's Hospital on Oct. 18 with symptoms of rabies, including slurred speech and fluctuating consciousness. The doctors induced a temporary coma and treated her with antiviral drugs to boost her immune system and allow her natural immunity to fight the virus. The details of the treatment and the specifics of the drugs used are under wraps until the doctors publish their findings in a medical journal. A rabies vaccine will prevent the disease only if given within days of exposure; it is useless in saving the patient's life in advanced cases.

Neonatal Infections Limit Growth

Extremely low-birth-weight infants (401-1,000 g) who developed neonatal infections were significantly more likely to have neurodevelopmental problems in early childhood, compared with noninfected infants in a cohort study of 6,093 children, said Barbara J. Stoll, M.D., of Emory University, Atlanta, and her colleagues. The infants were assessed at 18-22 months' corrected gestational age and classified as uninfected (2,161 infants), clinical infection only (1,538 infants), sepsis (1,922 infants), sepsis and necrotizing enterocolitis (279 infants), or meningitis, either with or without sepsis (193 infants). At follow-up, 41% of the children had at least one neurodevelopmental problem (JAMA 2004; 292:2357-65). Scores of less than 70 on the Mental Development Index and the Psychomotor Development Index were significantly more common among children with any of the previously mentioned infections, compared with uninfected children. In addition, children in any of the infection groups were significantly more likely to have cerebral palsy, vision impairment, and neurodevelopmental impairment, and to have a head circumference in less than the 10th percentile, compared with uninfected children.

11-Valent Vaccine Shows Promise

A new 11-valent pneumococcal conjugate vaccine (Pn-PD) was safe and effective in a randomized, single-blind study of 154 infants who received the vaccine at ages 2, 4, 6, and 12-15 months, reported Anu Nurkka of the National Public Health Institute in Helsinki, Finland, and colleagues The vaccine used Haemophilus influenzae protein D as a carrier and contained pneumococcal capsular polysaccharides of serotypes 1, 3, 4, 5, 6B, 7F, 9V, 14, 18C, 19F, and 23F. Overall, three doses of Pn-PD provoked a strong antibody response, compared with a control vaccine, with a significant booster response after the fourth dose (Pediatr. Infect. Dis. J. 2004;23:1008-14). Mild local skin reactions were common.

—Heidi Splete