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Simultaneous Influenza and Pneumococcal Vaccines

by Gina Pugliese, RN, MS
Medical News Editor

A recent study concluded that simultaneous administration of pneumococcal and influenza vaccines to the elderly is safe and adverse reactions are mild. This is good news, because it may provide a strategy to increase the currently poor immunization rates for two infections that are major causes of morbidity and mortality among the elderly.

The coverage of influenza vaccination is approximately 40%, and the coverage of pneumococcal vaccination is around 10%. The Immunization Practices Advisory Committee (ACIP) recommends that both vaccines be administered to persons aged 65 years or older. It is clear that there needs to be methods to improve vaccine use, and one possible improvement would be simultaneous administration of the two vaccines.

Simultaneous administration of these vaccines has been shown not to interfere with the antibody response, and the possibility of giving both vaccines at the same time is included in the ACIP recommendation. However, there are few data regarding the effects of the two vaccines when given at the same time.

Dr. Pekka Honkanen and colleagues from the National Public Health Institute in Oulu, Finland, conducted a study of 9,336 persons in Northern Finland aged 65 years or older, of whom 4,581 persons received influenza vaccine and 4,755 persons received influenza and pneumococcal vaccines. No serious reactions were observed in any vaccine recipients. The incidence of local reactions was 284 per 1,000 vaccinations in the influenza-vaccinated group and 441 per 1,000 vaccinations in the influenza-pneumococcal-vaccinated group. The incidence of fever (at least 37.5°C) was 10 and 24 per 1,000, respectively. The frequency of local reactions decreased with advancing age.

The authors conclude that simultaneous administration of pneumococcal and influenza vaccine proved to be safe, and, when indicated, the vaccines should be given at the same time.

FROM: Honkanen PO, Keistinen T, Sirkka-Liisa K. Reactions following administration of influenza vaccine alone or with pneumococcal vaccine to the elderly. *Arch Intern Med* 1996;156:205-208.

New TB Test Approved

by Gina Pugliese, RN, MS
Medical News Editor

FDA has approved a target amplification test for the direct detection of *Mycobacterium tuberculosis* in digested decontaminated respiratory specimens that are AFB-smear positive. The test is marketed by Gen-Probe Incorporated, San Diego, California, as the Gen-Probe Amplified *Mycobacterium tuberculosis* Direct (MTD) Test. It can identify *M tuberculosis* in respiratory specimens in 4 to 5 hours. The MTD test is a nucleic acid probe using target amplification and is indicated for detection of *M tuberculosis* in AFB-smear–positive sediments prepared from induced or expectorated sputum, bronchial aspirates, and specimens from bronchoalveolar lavages or tracheal aspirates. The MTD test cannot determine drug susceptibility or identify other mycobacterial species that are clinically significant.

FROM: Nightingale SL. New tuberculosis test approved. JAMA 1996;275(8):585.

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