



# MMWR<sup>TM</sup>

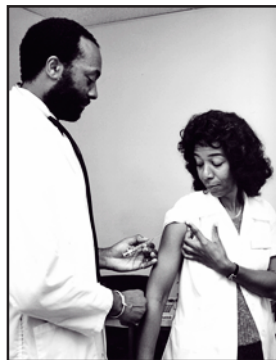
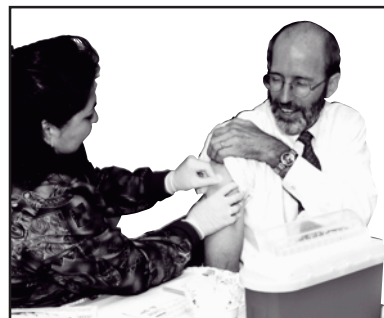
## Morbidity and Mortality Weekly Report

Early Release

February 9, 2006 / Vol. 55

### Influenza Vaccination of Health-Care Personnel

**Recommendations of the Healthcare Infection Control Practices Advisory Committee (HICPAC) and the Advisory Committee on Immunization Practices (ACIP)**



The *MMWR* series of publications is published by the Coordinating Center for Health Information and Service, Centers for Disease Control and Prevention (CDC), U.S. Department of Health and Human Services, Atlanta, GA 30333.

#### SUGGESTED CITATION

Centers for Disease Control and Prevention. Influenza vaccination of health-care personnel: recommendations of the Healthcare Infection Control Practices Advisory Committee (HICPAC) and the Advisory Committee on Immunization Practices (ACIP). Early Release 2006; 55(February 9, 2006):[inclusive page numbers].

#### Centers for Disease Control and Prevention

Julie L. Gerberding, MD, MPH  
*Director*

Dixie E. Snider, MD, MPH  
*Chief Science Officer*

Tanja Popovic, MD, PhD  
*Associate Director for Science*

#### Coordinating Center for Health Information and Service

Steven L. Solomon, MD  
*Director*

#### National Center for Health Marketing

Jay M. Bernhardt, PhD, MPH  
*Director*

#### Division of Scientific Communications

Judith R. Aguilar  
*(Acting) Director*

Mary Lou Lindegren, MD  
*Editor, MMWR Series*

Suzanne M. Hewitt, MPA  
*Managing Editor, MMWR Series*

Teresa F. Rutledge  
*Lead Technical Writer-Editor*

Jeffrey D. Sokolow, MA  
*Project Editor*

Beverly J. Holland  
*Lead Visual Information Specialist*

Lynda G. Cupell  
Malbea A. LaPete  
*Visual Information Specialists*

Quang M. Doan, MBA  
Erica R. Shaver  
*Information Technology Specialists*

## CONTENTS

Introduction .....	1
Summary Recommendations .....	2
Background .....	2
Transmission of Influenza in Health-Care Settings .....	3
Strategies for Improving HCP Vaccination Rates .....	5
Recommendations for Using Inactivated Influenza Vaccine and LAIV Among HCP .....	6
Recommendations for Prioritization of Influenza Vaccination During the 2005–06 Influenza Season .....	10
Side Effects and Adverse Reactions Associated with Vaccination .....	10
Additional Information Regarding Influenza Infection Control in Health-Care Settings .....	11
References .....	12

# Influenza Vaccination of Health-Care Personnel

## Recommendations of the Healthcare Infection Control Practices Advisory Committee (HICPAC) and the Advisory Committee on Immunization Practices (ACIP)

Prepared by

Michele L. Pearson, MD<sup>1</sup>

Carolyn B. Bridges, MD<sup>2</sup>

Scott A. Harper, MD<sup>3</sup>

<sup>1</sup>Division of Healthcare Quality Promotion, National Center for Infectious Diseases

<sup>2</sup>Epidemiology and Surveillance Division, National Immunization Program

<sup>3</sup>Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases

### Summary

*This report summarizes recommendations of the Healthcare Infection Control Practices Advisory Committee (HICPAC) and the Advisory Committee on Immunization Practices (ACIP) concerning influenza vaccination of health-care personnel (HCP) in the United States. These recommendations apply to HCP in acute care hospitals, nursing homes, skilled nursing facilities, physician's offices, urgent care centers, and outpatient clinics, and to persons who provide home health care and emergency medical services. The recommendations are targeted at health-care facility administrators, infection-control professionals, and occupational health professionals responsible for influenza vaccination programs and influenza infection-control programs in their institutions. HICPAC and ACIP recommend that all HCP be vaccinated annually against influenza. Facilities that employ HCP are strongly encouraged to provide vaccine to their staff by using evidence-based approaches that maximize vaccination rates.*

### Introduction

Influenza transmission and outbreaks in hospitals (1–8) and nursing homes (9–13) are well documented. HCP can acquire influenza from patients or transmit influenza to patients and other staff. Despite the documented benefits of HCP influenza vaccination on patient outcomes (14,15) and HCP absenteeism (16) and on reducing influenza infection among staff (16,17), vaccination coverage among HCP remain low (i.e., <50%) (18). Because HCP provide care to patients at high risk for complications of influenza, HCP should be considered a high priority for expanding influenza vaccine use. In addition, older HCP (i.e., aged  $\geq 65$  years) and those who have underlying chronic medical conditions or who might be pregnant are at increased risk for influenza-related

complications. Achieving and sustaining high vaccination coverage among HCP will protect staff and their patients, and reduce disease burden and health-care costs.

This report summarizes recommendations of the Healthcare Infection Control Practices Advisory Committee (HICPAC) and the Advisory Committee on Immunization Practices (ACIP) concerning influenza vaccination of health-care personnel (HCP)\* in the United States. These recommendations are targeted at health-care facility administrators, infection control professionals, and occupational health professionals responsible for influenza vaccination programs and influenza infection control programs in their institutions. HICPAC and ACIP recommend that all HCP be vaccinated annually against influenza. Facilities that employ HCP are strongly encouraged to provide vaccine to their staff by using

The material in this report originated in the National Center for Infectious Diseases, Rima F. Khabbaz, MD, Director; Division of Healthcare Quality Promotion, Denise M. Cardo, MD, Director; Division of Viral and Rickettsial Diseases, Steve Monroe, PhD, Acting Director; and National Immunization Program, Anne Schuchat, MD, Director; Epidemiology and Surveillance Division, Alison Mawle, PhD, Acting Director.

**Corresponding preparer:** Michele L. Pearson, MD, Division of Healthcare Quality Promotion, National Center for Infectious Diseases, 1600 Clifton Road, NE, MS A-31, Atlanta, GA 30333. Telephone: 404-639-4251; Fax: 404-639-4046; E-mail: mpearson@cdc.gov.

\* In this report, the term HCP refers to all paid and unpaid persons working in health-care settings who have the potential for exposure to infectious materials, including body substances, contaminated medical supplies and equipment, contaminated environmental surfaces, or contaminated air. HCP might include (but are not limited to) physicians, nurses, nursing assistants, therapists, technicians, emergency medical service personnel, dental personnel, pharmacists, laboratory personnel, autopsy personnel, students and trainees, contractual staff not employed by the health-care facility, and persons (e.g., clerical, dietary, housekeeping, maintenance, and volunteers) not directly involved in patient care but potentially exposed to infectious agents that can be transmitted to and from HCP. The recommendations in this report apply to HCP in acute care hospitals, nursing homes, skilled nursing facilities, physician's offices, urgent care centers, and outpatient clinics, and to persons who provide home health care and emergency medical services.

evidence-based approaches that maximize vaccination rates. This report supplements ACIP's previous statement regarding use of influenza vaccine and antiviral agents (1), which provides details regarding the epidemiology of influenza transmission in nonhealth-care settings, influenza vaccination of nonhealth-care personnel, composition of influenza vaccines, and use of antiviral medications.

## Summary Recommendations

The summary recommendations contained in this report are categorized by using the HICPAC evidence ranking system (Table 1). The recommendations were drafted after review of peer-reviewed scientific articles, and whenever possible are based on well-designed studies; certain recommendations are based on strong theoretic rationale and expert opinion. All recommendations have been approved by HICPAC and ACIP. The committees involved in drafting and reviewing these recommendations included persons with expertise in infectious diseases, infection control, pediatrics, vaccinology, internal medicine, and public health. The recommendations are as follows:

- Educate HCP regarding the benefits of influenza vaccination and the potential health consequences of influenza illness for themselves and their patients, the epidemiology and modes of transmission, diagnosis, treatment, and nonvaccine infection control strategies, in accordance with their level of responsibility in preventing health-care-associated influenza (category IB).
- Offer influenza vaccine annually to all eligible HCP to protect staff, patients, and family members and to decrease HCP absenteeism. Use of either available vaccine (inactivated and live, attenuated influenza vaccine [LAIV]) is recommended for eligible persons. During periods when inactivated vaccine is in short supply, use of LAIV is especially encouraged when feasible for eligible HCP (category IA).
- Provide influenza vaccination to HCP at the work site and at no cost as one component of employee health pro-

grams. Use strategies that have been demonstrated to increase influenza vaccine acceptance, including vaccination clinics, mobile carts, vaccination access during all work shifts, and modeling and support by institutional leaders (category IB).

- Obtain a signed declination from HCP who decline influenza vaccination for reasons other than medical contraindications (category II).
- Monitor HCP influenza vaccination coverage and declination at regular intervals during influenza season and provide feedback of ward-, unit-, and specialty-specific rates to staff and administration (category IB).
- Use the level of HCP influenza vaccination coverage as one measure of a patient safety quality program (category II).

## Background

### Influenza Among HCP

A limited number of prospective and cross-sectional studies provide estimates of incidence of influenza and influenza-like illness (ILI) among HCP (17,19,20). In one serosurvey of HCP, 23% of HCP had documented serologic evidence of influenza infection after a mild influenza season; however, of these, 59% could not recall having influenza, and 28% could not recall any respiratory infection, suggesting a high proportion of asymptomatic illness (17). In a randomized trial of influenza vaccine among HCP, 13% of placebo recipients subsequently had influenza infection (18). In a cross-sectional survey of house staff, 37% reported ILI during an 8-month period (September–April); 9% reported more than one illness. Length of illness varied (range: 1–10 days; mean: 7 days), as did the number of days of work missed (range: 0–10 days; mean: 0.7 days) (20).

### Efficacy and Effectiveness of Influenza Vaccines Among Adults

Trivalent inactivated influenza vaccine prevents influenza illness among approximately 70%–90% of healthy adults aged <65 years when the vaccine and circulating viruses are anti-

**TABLE 1. Healthcare Infection Control Practices Advisory Committee categorization scheme for recommendations\***

Category IA	Strongly recommended for implementation and strongly supported by well-designed experimental, clinical, or epidemiologic studies.
Category IB	Strongly recommended for implementation and supported by certain experimental, clinical, or epidemiologic studies and a strong theoretic rationale.
Category IC	Required for implementation, as mandated by federal or state regulation or standard.
Category II	Suggested for implementation and supported by suggestive clinical or epidemiologic studies or a theoretic rationale.
No recommendation	Unresolved issue; practices for which insufficient evidence or no consensus regarding efficacy exist.

\*Categorized on the basis of existing scientific data, theoretic rationale, applicability, and economic impact.

genically similar (17,21–23). The effectiveness of inactivated influenza vaccine in preventing influenza illness might be lower when vaccine and circulating viruses are not well matched or among adults aged  $\geq 65$  years and persons with certain chronic conditions (e.g., diabetes, human immunodeficiency virus (HIV), or chronic obstructive pulmonary disease) (24–28). Vaccination of healthy adults also decreases work absenteeism and use of health-care resources, including antibiotics, when the vaccine and circulating viruses are well matched (17,21,23,29–31). In addition, influenza vaccine prevents secondary complications and reduces the risk for influenza-related hospitalization and death among adults aged  $\geq 65$  years with and without high-risk medical conditions (e.g., heart disease and diabetes) (32–36).

LAIV has demonstrated similar benefits in randomized controlled trials among healthy working adults aged 18–64 years. In one study, vaccination with LAIV reduced severe febrile illnesses 19% and upper respiratory tract illnesses 24%; LAIV use also was associated with fewer days of illness and of work lost, fewer health-care provider visits, and reduced use of prescription antibiotics and over-the-counter medications (37). These results were recorded during a season in which the vaccine and circulating influenza A (H3N2) strains were not well matched. In the same study, LAIV vaccination yielded similar benefits among a subset of healthy adults aged 18–49 years, and antibiotic use in this age group decreased 41%–51% (37). In one study, overall efficacy of LAIV and inactivated influenza vaccine in preventing laboratory-documented influenza was 85% and 71%, respectively (38).

## Impact of HCP Vaccination on Influenza in Health-Care Settings

Vaccination of HCP is an important component of influenza prevention programs in the United States (18). Vaccination of HCP reduces transmission of influenza in health-care settings, staff illness and absenteeism, and influenza-related morbidity and mortality among persons at increased risk for severe influenza illness (14–17). Use of antiviral drugs used for chemoprophylaxis or treatment of influenza is an adjunct to (but not a substitute for) vaccination (18).

## Transmission of Influenza in Health-Care Settings

Influenza outbreaks in hospitals (4,39) and long-term-care facilities (40) have been associated with low vaccination rates among HCP. In addition, higher vaccination levels among staff have been associated with a lower incidence of nosocomial influenza cases (14,15,39).

In one tertiary care facility in which routine surveillance for influenza was conducted, the relation between staff vaccination coverage and annual incidence of nosocomial influenza was assessed for 12 influenza seasons during 1987–2000. During this period, staff vaccination coverage increased from 4% during 1987–1988 to 67% during 1999–2000 ( $p < 0.0001$ ), and the proportion of laboratory-confirmed cases of influenza that occurred among HCP decreased from 42% during 1990–1993 to 9% during 1997–2000 ( $p < 0.0001$ ). The proportion of nosocomial cases among hospitalized patients decreased 32% to 0 ( $p < 0.0001$ ). After controlling for potential confounders by using logistic regression, a significant and inverse relationship was demonstrated between vaccination rates among HCP and the rate of nosocomial influenza among patients, suggesting that staff vaccination contributed to the observed decline in the number of nosocomial influenza cases (39).

## Staff Illness and Absenteeism

During an influenza season, HCP might acquire influenza from infected patients with resulting morbidity and absenteeism. The impact of influenza vaccination on staff illness and absenteeism has been evaluated in two randomized, placebo-controlled, double-blind trials. In one trial, HCP who received vaccine had 28% fewer documented lost work days attributable to respiratory infections (1.0 and 1.4, respectively;  $p = 0.02$ ) and 28% fewer days on which they felt unable to work, whether they were on or off duty (2.5 and 3.5, respectively;  $p = 0.02$ ). Vaccination did not reduce either the number of episodes (1.8 and 2.0, respectively) or the total number of days (13.5 and 14.6, respectively) of respiratory infection (16). In a second trial conducted in two large teaching hospitals for 3 consecutive years that measured serologically confirmed influenza, days of febrile respiratory illness, and days absent from work, HCP who received influenza vaccine had a substantially lower incidence of influenza than controls (1.7% and 13.4%, respectively) with an estimated vaccine efficacy against serologically defined influenza A and influenza B infection of 88% and 89%, respectively. HCP who received influenza vaccine also tended to have fewer total respiratory illnesses (28.7 and 40.6 per 100 persons, respectively;  $p = 0.57$ ) and days of lost work (9.9 and 21.1 per 100 persons, respectively;  $p = 0.41$ ) than did controls (17).

In a cross-sectional survey, similar reductions in staff illness episodes and days of illness were reported (20). Overall, compared with unvaccinated coworkers, vaccinated house staff reported 23% fewer ILIs (42 and 54 per 100 persons, respectively;  $p = 0.03$ ), 27% fewer days of illness (80 and 115 per 100 persons, respectively;  $p = 0.02$ ), and a 59% reduction in illness during vacation time (1.7% and 4.0% of persons,



respectively;  $p = 0.08$ ). The two groups had a similar number of lost work days attributable to ILI (18 and 21 per 100 subjects, respectively;  $p = 0.69$ ). During influenza season, vaccination was associated with reductions of 30% in ILI ( $p = 0.05$ ), 43% in the proportion of house staff reporting illnesses associated with fever and cough ( $p = 0.05$ ), and 63% in illnesses associated with fever and cough ( $p = 0.03$ ). The inability to consistently demonstrate statistically significant decreases in absenteeism among staff who received vaccination is likely attributable to the finding that HCP tend to work despite illness (17,41).

## Patient Outcomes

HCP who are clinically or subclinically infected can transmit influenza virus to other persons. Decreasing transmission of influenza from caregivers to persons at high risk might reduce influenza-related deaths among persons at high risk for complications from influenza.

Residents of long-term-care facilities are particularly vulnerable to influenza and influenza-related complications. In 1999, an estimated 1.6 million persons resided in nursing homes in the United States (42). During influenza outbreaks in long-term-care facilities, attack rates among residents have ranged as high as 25%–60%, with case-fatality rates of 10%–20% (13,43–45). When vaccine and epidemic strains are well matched, achieving increased vaccination rates among persons living in closed settings (e.g., nursing homes and other chronic-care facilities) and among staff can reduce the risk for outbreaks by inducing herd immunity (32).

Two randomized controlled trials have evaluated the impact of influenza vaccination of HCP on the outcomes of residents in nursing homes. In one study, staff vaccination was associated with a 43% decrease in incidence of ILI (odds ratio [OR] = 0.6; 95% confidence interval [CI] = 0.3–0.9) and a 44% decrease in overall mortality among facility residents, from 17% to 10% (OR = 0.6; 95% CI = 0.4–0.8) (15). No virologic data were provided in this study. In a second study, 20 long-term-care facilities were randomized to have vaccine routinely offered (intervention facilities) or not offered (control facilities) to their staff (14). Facilities were paired by number of beds and patient vaccination policies. Staff vaccination coverage was higher in intervention facilities than in control facilities (50.9% and 4.9%, respectively). Crude mortality rates were 42% lower among residents in facilities with higher staff vaccination coverage than those in control facilities (13.6% and 22.4%, respectively; OR = 0.6; 95% CI = 0.4–0.8;  $p = 0.014$ ). Incidence of laboratory-confirmed influenza did not differ between the two groups (5.4% and 6.7%, respectively), but postmortem samples from pa-

tients in control facilities were more likely to be positive for influenza by a polymerase chain reaction test than samples from patients in intervention facilities (six [20%] of 30 and none of 17, respectively;  $p = 0.055$ ), suggesting that in this study population, HCP vaccination reduced influenza-related mortality in patients despite not reducing the incidence of non-fatal influenza infection. In neither study was a significant association demonstrated between patient vaccination and mortality. Randomized trials assessing the impact of staff vaccination on patient outcomes in acute care facilities have not been conducted, but low staff vaccination coverage has been correlated with influenza outbreaks in hospitals (4,39).

## Cost-Effectiveness of Influenza Vaccine

Cost-effectiveness studies of adults aged <65 years indicate that vaccination can reduce both direct medical costs and indirect costs from work absenteeism (21,23,29,30,46,47), resulting in 13%–44% fewer health-care provider visits, 18%–45% fewer lost workdays, 18%–28% fewer days working with reduced effectiveness, and a 25% decrease in antibiotic use for ILI (21,29,48,49). Among healthy persons aged 18–64 years, vaccination can save an estimated \$60–\$4,000 per illness, depending on the cost of vaccination, the influenza attack rate, and vaccine effectiveness against ILI (23). In another economic analysis, vaccination resulted in an average annual cost savings of \$13.66 per person vaccinated (50); however, other analyses have not demonstrated cost savings (21). Among studies of healthy young adults, >70% of the costs prevented were associated with reductions in lost work productivity.

## Vaccination Coverage Levels Among HCP

During 1989–2003, HCP vaccination coverage levels in the United States increased substantially, from 10% to 40%; however, coverage levels have remained relatively constant since 1997 (18). One of the national health objectives for 2010 is to achieve HCP vaccination coverage levels of 60% (objective no. 14-29g) (51). Substantially lower vaccination rates have been reported among HCP who have contact with certain populations at high risk (12,52–54). In addition, HCP vaccination coverage varies by level and years of training, age, occupational group, and facility type (20,55,56).

## Barriers to HCP Vaccination

Reported barriers to HCP receipt or acceptance of influenza vaccination include fear of vaccine side effects (particularly ILI symptoms) (20,55,57–61), insufficient time or

inconvenience (20), perceived ineffectiveness of the vaccine (20,55,58,59), medical contraindication (55), perceived low likelihood of contracting influenza (55,60,62), reliance on treatment with homeopathic medications (55,62), avoidance of medications (57), and fear of needles (57,59). Factors facilitating vaccine acceptance include a desire for self-protection (20,58,61), previous receipt of influenza vaccine (57,58,63–65), a desire to protect patients (61), and perceived effectiveness of vaccine (20).

## Strategies for Improving HCP Vaccination Rates

Facilities that employ HCP are strongly encouraged to provide vaccine to staff by using evidence-based approaches that maximize vaccination rates. Successful HCP vaccination programs are multifaceted and combine publicity and education to combat fears and misconceptions about influenza and influenza vaccines, use of reminder recall systems, efforts to remove administrative and financial barriers, role modeling, and monitoring and feedback on vaccination coverage (66). In contrast, single-component interventions will likely have minimal effectiveness in achieving desired vaccination coverage levels (66,67).

### Education and Campaigns

HCP knowledge, perceptions, and attitudes regarding influenza and influenza vaccination vary (20). Basic knowledge about influenza and influenza vaccination has been associated with vaccine receipt (57,68,69), and participation in structured in-service education or conferences has been associated with improved vaccination rates (62,65). Educational programs should emphasize the benefits of HCP vaccination for staff and patients (70). Organized campaigns that promote and make vaccine accessible can improve vaccination rates among HCP (52,71).

### Role Models

Vaccination of senior medical staff or opinion leaders has been associated with higher vaccination acceptance among staff members under their leadership (55,69,72,73). For example, medical students who have contact with infectious disease specialists are more likely to be vaccinated (69).

### Improved Access

Removing administrative barriers (e.g., costs) (71) and providing vaccine in locations and at times easily accessible by HCP can substantially improve vaccine acceptance

(40,52,55,72,74,75). In one survey, 33% of HCP reported that they would reject vaccination if they were required to pay for the vaccine (76).

Making vaccine readily accessible at congregate areas (e.g., clinics), during conferences, or by use of mobile carts (40,52,55,72) has been demonstrated to improve vaccination coverage rates. Use of mobile carts has been associated with increased vaccine acceptance during outbreaks and nonoutbreak situations (75,76). In a 3-year prospective study in a 630-bed acute care hospital, a sustained four- to fivefold increase in vaccination rates was associated with using mobile carts to deliver vaccine to staff rather than requiring HCP to visit an employee health center to receive vaccine. Provision of modest incentives also has been associated with improved vaccine acceptance among HCP (77). However, the benefits of vaccine deputies or peer-vaccinators have not been consistently associated with improved HCP vaccination (52).

## Measurement and Feedback

HCP influenza vaccination coverage should be regularly measured and reported. Posting of vaccination coverage levels in different areas of the hospital is a component of successful vaccination programs (6). Monitoring vaccination coverage by facility area (e.g., ward or unit) or occupational group allows facilities to identify where vaccination levels are low and interventions should be targeted. In addition, HICPAC has recommended that HCP influenza vaccination coverage be used as a health-care quality measure in those states that mandate public reporting of health-care-associated infections (78).

The independent contribution of signed declination statements to improving HCP vaccination has not been studied. However, obtaining declination statements from HCP who refuse vaccination for reasons other than medical contraindications can assist facilities in identifying personnel who might require targeted education or other interventions to overcome barriers to vaccine acceptance. In addition, collection of such information will allow health-care facilities to determine what proportion of their staff are reached and offered vaccine.

## Legislation and Regulation

Legislative and regulatory efforts have favorably affected hepatitis B vaccination rates among HCP (79,80). As of January 2005, a total of 13 states (Alabama, Arkansas, Kentucky, Maine, Maryland, New Hampshire, New York, Oklahoma, Oregon, Pennsylvania, Rhode Island, Texas, and Utah) and the District of Columbia were reported to have enacted regulations regarding influenza vaccination of staff in long-term-care facilities (67,81). However, because only one state

(Pennsylvania) has monitored the impact of its laws on nursing home staff vaccination rates, data are insufficient to assess the overall impact of these legislative efforts on HCP influenza vaccination coverage (CDC, unpublished data, 2005).

## Recommendations for Using Inactivated Influenza Vaccine and LAIV Among HCP

All HCP should be vaccinated annually against influenza. Either inactivated influenza vaccine or LAIV can be used to reduce the risk for influenza among HCP (Table 2). LAIV is approved for use only among nonpregnant healthy persons aged 5–49 years. HCP who work with severely immunocompromised patients who require a protected environment should not receive LAIV. Inactivated influenza vaccine is approved for all persons aged >6 months who lack vaccine contraindications, including those with high-risk conditions (see Recommendations for Prioritization of Influenza Vaccine During the 2005–06 Influenza Season). Four influenza vaccines have been approved for use in the United States during the 2005–06 season (Table 3).

## Inactivated Influenza Vaccine Recommendations

### Dosage and Route

Because immunity declines during the year after vaccination, HCP eligible to receive inactivated influenza vaccine should be administered 1 dose of the current year's vaccine each year (82,83). The intramuscular route is recommended for inactivated influenza vaccine. Adults should be vaccinated in the deltoid muscle, ideally by using a needle of length >1 inch because needles of length <1 inch might not penetrate muscle tissue in certain adults (84).

### Persons Who Should Not Be Vaccinated with Inactivated Influenza Vaccine

Inactivated influenza vaccine should not be administered to persons known to have anaphylactic hypersensitivity to eggs or to other components of the influenza vaccine without first consulting a physician (see Side Effects and Adverse Reactions Associated with Vaccination). Prophylactic use of antiviral agents is an option for preventing influenza among such persons. However, persons who have a history of anaphylactic

**TABLE 2. Live, attenuated influenza vaccine (LAIV) compared with trivalent inactivated influenza vaccine**

Factor	LAIV	Trivalent inactivated influenza vaccine
Route of administration	Intranasal spray	Intramuscular injection
Type of vaccine	Live virus	Killed virus
No. of included virus strains	3 (2 influenza A, 1 influenza B)	Same as LAIV
Vaccine virus strains updated	Annually	Same as LAIV
Frequency of administration	Annually	Same as LAIV
Approved age and risk groups*	Healthy persons aged 5–49 yrs	Persons aged ≥6 mos
Can be administered to family members or close contacts of immunosuppressed persons not requiring a protected environment	Yes	Yes
Can be administered to family members or close contacts of immunosuppressed persons requiring a protected environment (e.g., hematopoietic stemcell transplant recipient)	Inactivated influenza vaccine preferred	Yes
Can be administered to family members or close contacts of persons at high risk but not severely immunosuppressed	Yes	Yes
Can be simultaneously administered with other vaccines	Yes†	Yes§
If not simultaneously administered, can be administered within 4 weeks of another live vaccine	Prudent to space 4 weeks apart	Yes
If not simultaneously administered, can be administered within 4 weeks of an inactivated vaccine	Yes	Yes

\* Populations at high risk from complications of influenza infection include persons aged ≥65 years; residents of nursing homes and other chronic-care facilities that house persons with chronic medical conditions; adults and children with chronic disorders of the pulmonary or cardiovascular systems; adults and children with chronic metabolic diseases (including diabetes mellitus), renal dysfunction, hemoglobinopathies, or immunosuppression; children and adolescents receiving long-term aspirin therapy (at risk for Reye syndrome after wild-type influenza infection); pregnant women; and children aged 6–23 months.

† No data are available regarding effect on safety or efficacy.

§ Inactivated influenza vaccine coadministration has been evaluated systematically only among adults with pneumococcal polysaccharide vaccine.



**TABLE 3. Influenza vaccine manufacturers and projected supplies for the 2005–06 influenza season**

Manufacturer	Vaccine	Formulation	Contains thimerosal as preservative	Age indication	No. of projected doses*
Sanofi Pasteur, Inc.	Fluzone <sup>®†</sup>	Multidose vial	Yes	≥6 mos	60 million <sup>§</sup>
		Single-dose prefilled 0.5-mL syringe or vial	No	≥36 mos	
		Single-dose prefilled 0.25-mL syringe	No	6–35 mos	
Chiron Corporation	Fluvirin <sup>™†</sup>	Multidose vial	Yes	≥4 yrs	18–26 million
		Single-dose prefilled 0.5-mL syringe	No <sup>¶</sup>	≥4 yrs	
GlaxoSmithKline, Inc.	Fluarix <sup>™†</sup>	Single-dose prefilled 0.5-mL syringe	No <sup>¶</sup>	≥18 yrs	8 million
MedImmune Vaccines, Inc.	FluMist <sup>™**</sup>	Single-dose nasal sprayer	No	Healthy, nonpregnant persons aged 5–49 yrs	3 million

\* As of August 2005.

† Trivalent inactivated influenza vaccine.

§ Approximately 6–8 million of the 60 million doses were projected to be distributed in single-dose prefilled syringes or vials.

¶ These preparations contain traces of thimerosal from the production process.

\*\* Live, attenuated influenza vaccine.

hypersensitivity to vaccine components but who are also at high risk for complications from influenza can benefit from vaccine after appropriate allergy evaluation and desensitization (18). Information regarding vaccine components is located in package inserts from each manufacturer. Persons with acute febrile illness typically should not be vaccinated until their symptoms have abated. However, minor illnesses with or without fever do not contraindicate use of influenza vaccine.

## LAIV Recommendations

LAIV licensed for use in the United States (FluMist<sup>™</sup> manufactured by MedImmune, Inc., Gaithersburg, Maryland [http://www.medimmune.com]) is a live, trivalent, intranasally administered vaccine that is

- attenuated, producing mild or no signs or symptoms related to influenza virus infection;
- temperature-sensitive, a property that limits the replication of the vaccine viruses at 100.4°–102.2° F (38° C–39° C) and thus restricts LAIV viruses from replicating efficiently in human lower airways; and
- cold-adapted, replicating efficiently at 77° F (25° C), a temperature that is permissive for replication of LAIV viruses but restrictive for replication of different wild-type viruses.

The immunogenicity of the approved LAIV has been assessed in multiple studies (85–91). LAIV virus strains replicate primarily in nasopharyngeal epithelial cells. The protective mechanisms induced by vaccination with LAIV are not completely understood but appear to involve both serum and nasal secretory antibodies. No single laboratory measurement closely correlates with protective immunity induced by LAIV.

## Shedding and Transmission of Vaccine Viruses

One concern regarding use of LAIV among HCP has been the potential for transmitting vaccine virus from persons receiving vaccine to nonimmune patients at high risk. Available data indicate that children and adults vaccinated with LAIV can shed vaccine viruses for >2 days after vaccination, although in lower titers than typically occur with shedding of wild-type influenza viruses. Shedding should not be equated with person-to-person transmission of vaccine viruses, although transmission of shed vaccine viruses from vaccinated persons to nonvaccinated persons has been documented in rare instances among children in a day care center (92).

In one study of 20 healthy vaccinated adults aged 18–49 years, the majority of vaccine virus shedding occurred within the first 3 days after vaccination, although in one vaccinated person, viral shedding was detected on day 7 after vaccination (93). No vaccine viruses were shed >10 days after vaccination, and duration or type of symptoms associated with receipt of LAIV did not correlate with duration of shedding of vaccine viruses (93). In another study of 14 healthy adults aged 18–49 years, 50% of vaccinated persons had viral antigen detected by direct immunofluorescence or rapid antigen tests within 7 days of vaccination; the majority of viral shedding was detected on day 2 or 3 (94). Person-to-person transmission of vaccine viruses was not assessed in either of these studies.

One study conducted in a child care center assessed transmissibility of vaccine viruses from 98 vaccinated persons to 99 unvaccinated controls aged 8–36 months; 80% of vaccine recipients shed one or more virus strains (mean duration: 7.6 days). One influenza type B isolate was recovered from a placebo recipient and confirmed to be vaccine-type virus; the

isolate retained the cold-adapted, temperature-sensitive, attenuated phenotype and possessed the same genetic sequence as a virus shed from a vaccine recipient in the same children's play group. The placebo recipient from whom the influenza type B vaccine virus was isolated exhibited symptoms that were similar to those experienced by vaccine recipients. The estimated probability of acquiring vaccine virus after close contact with a single LAIV recipient in this child care population was 0.6%–2.4% (92).

### Using LAIV for HCP

LAIV may be used for vaccination of healthy, nonpregnant persons aged 5–49 years, including HCP. When feasible, use of LAIV for vaccination of eligible HCP is especially encouraged during periods of limited supply of inactivated influenza vaccine because use of LAIV for HCP might increase availability of inactivated influenza vaccine for persons at high risk. Use of LAIV also provides an alternative vaccine strategy for HCP who avoid influenza vaccination because of an aversion to intramuscular injections.

### Persons Who Should Not Receive LAIV

The following populations should not receive LAIV:

- persons aged <5 years or >50 years;<sup>†</sup>
- persons with asthma, reactive airways disease or other chronic disorders of the pulmonary or cardiovascular systems; persons with other underlying medical conditions, including metabolic diseases such as diabetes, renal dysfunction, and hemoglobinopathies; or persons with known or suspected immunodeficiency diseases or who are receiving immunosuppressive therapies;<sup>†</sup>
- children or adolescents receiving aspirin or other salicylates (because of the association of Reye syndrome with wild-type influenza infection);<sup>†</sup>
- persons with a history of Guillain-Barré syndrome (GBS);
- pregnant women;<sup>†</sup>
- persons who have close contact with severely immunosuppressed persons (e.g., patients with hematopoietic stem cell transplants) during those periods in which the immunosuppressed person requires care in a protective environment; or
- persons with a history of hypersensitivity, including anaphylaxis, to any of the components of LAIV or to eggs.

### LAIV Dosage and Administration

Eligible HCP should receive 1 dose of LAIV. LAIV is intended only for intranasal administration and should not be administered by the intramuscular, intradermal, or intra-

venous route. Administration can be accomplished by holding an individual sprayer in the palm of the hand until thawed, with subsequent immediate administration. Alternatively, the vaccine can be thawed in a refrigerator and stored at 35.6° F–46.4° F (2° C–8° C) for ≤60 hours before use. Vaccine should not be refrozen after thawing. LAIV is supplied in a prefilled single-use sprayer containing 0.5 mL of vaccine. Approximately 0.25 mL is sprayed into the first nostril while the recipient is in the upright position. An attached dose-divider clip is removed from the sprayer to administer the second half of the dose into the other nostril. If the vaccine recipient sneezes after administration, the dose should not be repeated.

LAIV may be administered to persons with minor acute illnesses (e.g., diarrhea or mild upper respiratory tract infection, with or without fever). However, if clinical judgment indicates the presence of nasal congestion that might impede delivery of vaccine to the nasopharyngeal mucosa, deferral of administration should be considered until resolution of the illness.

Whether concurrent administration of LAIV with other vaccines affects the safety or efficacy of either LAIV or the simultaneously administered vaccine is unknown. In the absence of specific data indicating interference, adherence to ACIP's general recommendations for vaccination is prudent (95). Inactivated vaccines do not interfere with the immune response to other inactivated vaccines or to live vaccines. An inactivated vaccine can be administered either simultaneously or at any time before or after LAIV. Whenever possible, two live vaccines not administered on the same day should be administered >4 weeks apart.

### Recommended Vaccines for HCP Who Have Close Contact with Severely Immunosuppressed Persons

Inactivated influenza vaccine is the preferred vaccine for use among HCP who have close contact with severely immunosuppressed persons (e.g., patients with hematopoietic stem cell transplants) during those periods in which the immunosuppressed person requires care in a protective environment. The rationale for not using LAIV among HCP caring for such patients is the theoretic risk that a live, attenuated vaccine virus could be transmitted to the severely immunosuppressed person. HCP who receive LAIV should refrain from contact with severely immunosuppressed patients for 7 days after vaccine receipt. In addition, visitors who have received LAIV should refrain from contact with severely immunosuppressed persons for 7 days after vaccination; however, such persons need not be excluded from visitation of patients who are not

<sup>†</sup> These persons should receive inactivated influenza vaccine.

severely immunosuppressed. Either inactivated influenza vaccine or LAIV can be used to vaccinate HCP who have close contact with persons with lesser degrees of immunosuppression (e.g., persons with diabetes, persons with asthma taking corticosteroids, or persons infected with human immunodeficiency virus) or who are in close contact with all other persons at high risk.

### **Personnel Who May Administer LAIV**

The risk of acquiring vaccine viruses from the environment is unknown but likely small. Nevertheless, severely immunosuppressed persons should not administer LAIV because introduction of low levels of vaccine virus into the environment probably cannot be avoided when administering LAIV. However, other persons with conditions placing them at high risk for influenza complications (e.g., pregnant women, persons with asthma, and persons aged >50 years) may administer LAIV.

### **LAIV and Use of Influenza Antiviral Medications**

How LAIV coadministration with influenza antiviral medications affects safety and efficacy has not been studied. However, because influenza antivirals reduce replication of influenza viruses, LAIV should not be administered until 48 hours after cessation of influenza antiviral therapy, and influenza antiviral medications should not be administered for 2 weeks after receipt of LAIV.

### **LAIV Storage**

LAIV must be stored at -59° F (-15° C) or colder. LAIV may be stored in frost-free freezers without using a freezer-box. LAIV can be thawed in a refrigerator and stored at 35.6° F–46.4° F (2° C–8° C) for ≤60 hours before use. It should not be refrozen after thawing. Additional information regarding LAIV storage is available at <http://www.FluMist.com>.

## **Vaccination of Specific HCP Populations**

### **Pregnant Women**

Pregnant women are at increased risk for influenza-related complications (96–103) and hospitalizations (104). Therefore, all HCP who are pregnant during the influenza season should be vaccinated against influenza. However, pregnant women should receive only inactivated influenza vaccine; LAIV is not recommended for use during pregnancy. Inactivated influenza vaccine may be administered in any trimester. One study of influenza vaccination of approximately 2,000 pregnant women demonstrated no adverse fetal effects associated with receipt of inactivated influenza vaccine (105).

### **Breastfeeding Mothers**

Influenza vaccine does not affect the safety of mothers who are breastfeeding or their infants. Breastfeeding does not adversely affect the immune response and is not a contraindication for vaccination.

### **Persons Infected with HIV**

Detailed information on the use of influenza vaccine among persons infected with HIV has been published previously (18). Because influenza can result in serious illness and influenza vaccination can result in the production of protective antibody titers, vaccination with inactivated vaccine will benefit HIV-infected persons, including those that are pregnant.

## **Timing of Annual Influenza Vaccination of HCP**

### **Timing of Organized Vaccination Campaigns**

Planning for influenza campaigns should begin as early as February or March (106). The optimal time to vaccinate HCP is during October–November. Beginning in October each year, health-care facilities should offer influenza vaccinations to all full- and part-time staff. Particular emphasis should be placed on vaccinating HCP who care for persons at high risk. Vaccination programs should educate HCP regarding the benefits of vaccination and the potential health consequences of influenza illness for themselves and their patients. As part of employee health programs, all HCP should be provided convenient access to free influenza vaccine at the work site (107).

### **Vaccination in December and Later**

To improve vaccine coverage among HCP, influenza vaccine should continue to be offered in December and throughout the influenza season as long as vaccine supplies are available, even after influenza activity has been documented in the community. In the United States, seasonal influenza activity can increase as early as October or November, but influenza activity has not reached peak levels in the majority of recent seasons until late December–early March. Therefore, although the timing of influenza activity can vary by region, vaccine administered after November is likely to be beneficial in the majority of influenza seasons. Adults achieve peak antibody protection against influenza infection 2 weeks after vaccination (108,109).

## Recommendations for Prioritization of Influenza Vaccination During the 2005–06 Influenza Season

As a result of influenza vaccine distribution delays or supply shortages in the United States during recent influenza seasons (110,111), in September 2005, CDC issued recommendations for prioritizing the use of inactivated vaccine during the 2005–06 influenza season to ensure that early vaccine is available for those at the highest risk for complications from influenza (112). On the basis of uncertainties in doses and distribution, CDC recommended that the following groups receive priority for inactivated influenza vaccine until October 24, 2005:

- persons aged  $\geq 65$  years with and without comorbid conditions,
- residents of long-term-care facilities,
- persons aged 2–64 years with comorbid conditions,
- children aged 6–23 months,
- pregnant women,
- HCP who provide direct patient care, and
- household contacts and out-of-home caregivers of children aged  $< 6$  months (112).

These groups correspond to tiers 1A–1C in the table of inactivated influenza vaccine priority groups in the event of vaccination supply disruption that was published previously (113). After October 24, 2005, all persons were eligible for vaccination.

Tiered use of prioritization was not recommended for LAIV administration. LAIV may be administered at any time for vaccination of nonpregnant healthy persons aged 5–49 years, including the majority of HCP, other persons in close contact with persons at high risk for influenza-related complications, and others desiring protection against influenza (18).

## Side Effects and Adverse Reactions Associated with Vaccination

### Inactivated Influenza Vaccine

When educating HCP regarding potential side effects, providers should emphasize that 1) inactivated influenza vaccine contains noninfectious killed viruses and cannot cause influenza; and 2) coincidental respiratory disease unrelated to influenza vaccination can occur after vaccination. The occurrence of vaccine-related side effects has had limited to no impact on rates of absenteeism among HCP (16,17).

### Local Reactions

The most frequent side effect of vaccination (affecting 10%–64% of patients) is soreness at the vaccination site, typically lasting  $< 2$  days (21,114–116). Local reactions typically are mild and rarely interfere with a person's ability to conduct everyday activities. In a controlled trial, only body aches (25.1%) were reported more frequently after inactivated influenza vaccine than placebo-injection (20.8%) (117).

### Systemic Reactions

Fever, malaise, myalgia, and other systemic symptoms can occur after vaccination with inactivated vaccine and most often affect persons (e.g., infants) with no previous exposure to the influenza virus antigens in the vaccine (118,119). Such reactions typically begin 6–12 hours after vaccination and can persist for 1–2 days. Recent placebo-controlled trials demonstrate that among older persons and healthy young adults, administration of split-virus (i.e., detergent-disrupted virion) influenza vaccine is not associated with higher rates of systemic symptoms (e.g., fever, malaise, myalgia, and headache) compared with placebo injections (21,114–116). No increase in asthma exacerbations has been documented in association with receipt of influenza vaccine (117).

### Severe Adverse Events

Immediate and presumably allergic reactions (e.g., hives, angioedema, allergic asthma, and systemic anaphylaxis) rarely occur after influenza vaccination (120). These reactions probably result from hypersensitivity to certain vaccine components; the majority of reactions probably are caused by residual egg protein. Although current influenza vaccines contain only a limited quantity of egg protein, this protein can induce immediate hypersensitivity reactions among persons who have severe egg allergy. Persons who have had hives or swelling of the lips or tongue, or who have experienced acute respiratory distress or collapse after eating eggs should consult a physician for appropriate evaluation to help determine if vaccine should be administered. Persons who have documented immunoglobulin E (IgE)-mediated hypersensitivity to eggs, including those who have had occupational asthma or other allergic responses to egg protein, might also be at increased risk for allergic reactions to influenza vaccine, and consultation with a physician should be considered. Protocols have been published for administering influenza vaccine safely to persons with egg allergies (121–123).

Hypersensitivity reactions to any vaccine component can occur. Although exposure to vaccines containing thimerosal can



lead to induction of hypersensitivity, the majority of patients do not have reactions to thimerosal when it is administered as a component of vaccines, even when patch or intradermal tests for thimerosal allergy indicate hypersensitivity (124,125). When reported, hypersensitivity to thimerosal typically has consisted of local, delayed hypersensitivity reactions (124).

## GBS

Investigations to date indicate no substantial increase in GBS associated with influenza vaccines (other than the 1976 swine influenza vaccine) (126–130). If current influenza vaccines pose a risk for GBS, the estimated risk is approximately one additional case per million persons vaccinated, with the total combined number of GBS cases peaking 2 weeks after vaccination (131). This estimated risk for GBS is substantially less than the risk for severe influenza, which can be prevented by vaccination among all age groups, especially persons aged  $\geq 65$  years and those who have medical indications for influenza vaccination. The potential benefits of influenza vaccination in preventing serious illness, hospitalization, and death substantially outweigh the possible risks for experiencing vaccine-associated GBS. The average case-fatality ratio for GBS is 6% and increases with age (132,133). No evidence indicates that the case-fatality ratio for GBS differs among vaccinated persons and those not vaccinated.

Incidence of GBS among the general population is low, but persons with a history of GBS have a substantially greater likelihood of subsequently experiencing GBS than persons without such a history (128,134). Whether influenza vaccination might increase the risk for recurrence of GBS is unknown; for this reason, persons who are not at high risk for severe influenza complications and who are known to have experienced GBS within 6 weeks after a previous influenza vaccination should not receive vaccine. Chemoprophylaxis using influenza antivirals might be an alternative for such persons. Although data are limited, for the majority of persons who have a history of GBS and who are at high risk for severe complications from influenza, the established benefits of influenza vaccination justify yearly vaccination. Health-care professionals should promptly report all clinically significant adverse events after influenza vaccination to the Vaccine Adverse Event Reporting System (VAERS), even if evidence is lacking that the vaccine caused the event.

## LAIV

Until additional data are available, persons at high risk for experiencing complications from influenza infection (e.g.,

immunocompromised patients; patients with asthma, cystic fibrosis, or chronic obstructive pulmonary disease; or persons aged  $\geq 65$  years) should not be vaccinated with LAIV. Protection from influenza among these groups should be accomplished by using inactivated influenza vaccine.

Among adults, runny nose or nasal congestion (28%–78%), headache (16%–44%), and sore throat (15%–27%) have been reported more often among vaccine recipients than placebo recipients (37,135,136). In one clinical trial among a subset of healthy adults aged 18–49 years, signs and symptoms reported more frequently among LAIV recipients ( $n = 2,548$ ) than placebo recipients ( $n = 1,290$ ) within 7 days after each dose included cough (13.9% and 10.8%, respectively); runny nose (44.5% and 27.1%, respectively); sore throat (27.8% and 17.1%, respectively); chills (8.6% and 6.0%, respectively); and tiredness or weakness (25.7% and 21.6%, respectively) (37). Pneumonia, bronchitis, bronchiolitis, or central nervous system events have not been observed more frequently among LAIV than among placebo recipients.

## Severe Adverse Events

Serious adverse events associated with receipt of LAIV among healthy adults aged 18–49 years occur at a rate of  $< 1\%$  (137). However, surveillance should continue for adverse events that might not have been detected in previous studies. Health-care professionals should promptly report to VAERS all clinically significant adverse events after LAIV administration, even if evidence is lacking that the vaccine caused the event.

## Additional Information Regarding Influenza Infection Control in Health-Care Settings

Additional information on controlling and preventing influenza in health-care settings is available in the following publications:

- CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2005;54(No. RR-8):1–40.
- Garner JS, Hospital Infection Control Practices Advisory Committee. Guideline for isolation precautions in hospitals. Infect Control Hosp Epidemiol 1996;17:53–80.
- CDC. Guidelines for preventing health-care-associated pneumonia, 2003: recommendations of CDC and the Healthcare Infection Control Practices Advisory Committee. MMWR 2003;53(No. RR-3):1–36.
- CDC. Respiratory hygiene/cough etiquette in health-care settings. Atlanta, GA: US Department of Health and

Human Services, CDC; 2003. Available at <http://www.cdc.gov/flu/professionals/infectioncontrol/resphygiene.htm>.

- Bradley SF. The Long-Term-Care Committee of the Society for Healthcare Epidemiology of America. Prevention of influenza in long-term care facilities. *Infect Control Hosp Epidemiol* 1999;20:629–37.
- Sneller V-P, Izurieta H, Bridges C, et al. Prevention and control of vaccine-preventable diseases in long-term care facilities. *Journal of the American Medical Directors Association* 2000;1(Suppl):S2–37.
- Bridges CB, Kuehnert MJ, Hall CB. Transmission of influenza: implications for control in healthcare settings. *Clin Infect Dis* 2003;37:1094–101.
- CDC. Detection and control of influenza outbreaks in acute care facilities. Atlanta, GA: US Department of Health and Human Services, CDC; 2001. Available at <http://www.cdc.gov/ncidod/hip/INFECTFluBook2001.pdf>.
- Talbot TR, Bradley SE, Cosgrove SE, Ruef C, Siegel JD, Weber DJ. Influenza vaccination of healthcare workers and vaccine allocation for healthcare workers during vaccine shortages. *Infect Control Hosp Epidemiol* 2005;26:882–90.

## References

1. Malavaud S, Malavaud B, Sandres K, et al. Nosocomial outbreak of influenza virus A (H3N2) infection in a solid organ transplant department. *Transplantation* 2001;72:535–7.
2. Maltezos HC, Drancourt M. Nosocomial influenza in children. *J Hosp Infect* 2003;55:83–91.
3. Weinstock DM, Eagan J, Malak SA, et al. Control of influenza A on a bone marrow transplant unit. *Infect Control Hosp Epidemiol* 2000;21:730–2.
4. Cunney RJ, Bialachowski A, Thornley D, Smaill FM, Pennie RA. An outbreak of influenza A in a neonatal intensive care unit. *Infect Control Hosp Epidemiol* 2000;21:449–54.
5. Hall CB, Douglas RG Jr. Nosocomial influenza infection as a cause of intercurrent fevers in infants. *Pediatrics* 1975;55:673–7.
6. Salgado CD, Farr BM, Hall KK, Hayden FG. Influenza in the acute hospital setting. *Lancet Infect Dis* 2002;2:145–55.
7. Kapila R, Lintz DI, Tecson FT, Ziskin L, Louria DB. A nosocomial outbreak of influenza A. *Chest* 1977;71:576–9.
8. Sartor C, Zandotti C, Romain F, et al. Disruption of services in an internal medicine unit due to a nosocomial influenza outbreak. *Infect Control Hosp Epidemiol* 2002;23:615–9.
9. Mast EE, Harmon MW, Gravenstein S, et al. Emergence and possible transmission of amantadine-resistant viruses during nursing home outbreaks of influenza A (H3N2). *Am J Epidemiol* 1991;134:988–97.
10. Drinka PJ, Krause P, Schilling M, Miller BA, Shult PA, Gravenstein S. Report of an outbreak: nursing home architecture and influenza-A attack rates. *J Am Geriatr Soc* 1996;44:910–3.
11. Degelau J, Somani SK, Cooper SL, Guay DRP, Crossley KB. Amantadine-resistant influenza in a nursing facility. *Arch Intern Med* 1992;152:390–2.
12. Coles FB, Balzano GJ, Morse DL. An outbreak of influenza A (H3N2) in a well immunized nursing home population. *J Am Geriatr Soc* 1989;40:589–92.
13. Morens DM, Rash VM. Lessons from a nursing home outbreak of influenza A. *Infect Control Hosp Epidemiol* 1995;16:275–80.
14. Carman WF, Elder AG, Wallace LA, et al. Effects of influenza vaccination of health-care workers on mortality of elderly people in long-term care: a randomised controlled trial. *Lancet* 2000;355(9198):93–7.
15. Potter J, Stott DJ, Roberts MA, et al. Influenza vaccination of health care workers in long-term-care hospitals reduces the mortality of elderly patients. *J Infect Dis* 1997;175:1–6.
16. Saxen H, Virtanen M. Randomized, placebo-controlled double blind study on the efficacy of influenza immunization on absenteeism of health care workers. *Pediatr Infect Dis J* 1999;18:779–83.
17. Wilde JA, McMillan JA, Serwint J, Butta J, O’Riordan MA, Steinhoff MC. Effectiveness of influenza vaccine in health care professionals: a randomized trial. *JAMA* 1999;281:908–13.
18. CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). *MMWR* 2005;54(No. RR-8):1–40.
19. Elder AG, O’Donnell B, McCrudden EA, Symington IS, Carman WF. Incidence and recall of influenza in a cohort of Glasgow healthcare workers during the 1993–4 epidemic: results of serum testing and questionnaire. *BMJ* 1996;313(7067):1241–2.
20. Lester RT, McGeer A, Tomlinson G, Detsky AS. Use of, effectiveness of, and attitudes regarding influenza vaccine among house staff. *Infect Control Hosp Epidemiol* 2003;24:839–44.
21. Bridges CB, Thompson WW, Meltzer MI, et al. Effectiveness and cost-benefit of influenza vaccination of healthy working adults: a randomized controlled trial. *JAMA* 2000;284:1655–63.
22. Palache AM. Influenza vaccines. A reappraisal of their use. *Drugs* 1997;54:841–56.
23. Demicheli V, Jefferson T, Rivetti D, Deeks J. Prevention and early treatment of influenza in healthy adults. *Vaccine* 2000;18:957–1030.
24. Blumberg EA, Albano C, Pruett T, et al. The immunogenicity of influenza virus vaccine in solid organ transplant recipients. *Clin Infect Dis* 1996;22:295–302.
25. Dorrell L, Hassan I, Marshall S, Chakraverty P, Ong E. Clinical and serological responses to an inactivated influenza vaccine in adults with HIV infection, diabetes, obstructive airways disease, elderly adults and healthy volunteers. *Int J STD AIDS* 1997;8:776–9.
26. Govaert TM, Thijs CT, Masurel N, Sprenger MJ, Dinant GJ, Knottnerus JA. The efficacy of influenza vaccination in elderly individuals. A randomized double-blind placebo-controlled trial. *JAMA* 1994;272:1661–5.
27. McElhane JE, Beattie BL, Devine R, Grynock R, Toth EL, Bleackley RC. Age-related decline in interleukin 2 production in response to influenza vaccine. *J Am Geriatr Soc* 1990;38:652–8.
28. CDC. Assessment of the effectiveness of the 2003–04 influenza vaccine among children and adults—Colorado, 2003. *MMWR* 2003;53:707–10.
29. Nichol KL, Lind A, Margolis KL, et al. The effectiveness of vaccination against influenza in healthy, working adults. *N Engl J Med* 1995;333:889–93.
30. Campbell DS, Rumley MH. Cost-effectiveness of the influenza vaccine in a healthy, working-age population. *J Occup Environ Med* 1997;39:408–14.

31. Smith JW, Pollard R. Vaccination against influenza: a five-year study in the Post Office. *J Hyg (Lond)* 1979;83:157-70.
32. Patriarca PA, Weber JA, Parker RA, et al. Risk factors for outbreaks of influenza in nursing homes. A case-control study. *Am J Epidemiol* 1986;124:114-9.
33. Gross PA, Hermogenes AW, Sacks HS, Lau J, Levandowski RA. The efficacy of influenza vaccine in elderly persons. A meta-analysis and review of the literature. *Ann Intern Med* 1995;123:518-27.
34. Mullooly JP, Bennett MD, Hornbrook MC, et al. Influenza vaccination programs for elderly persons: cost-effectiveness in a health maintenance organization. *Ann Intern Med* 1994;121:947-52.
35. Nordin J, Mullooly J, Poblete S, et al. Influenza vaccine effectiveness in preventing hospitalizations and deaths in persons 65 years or older in Minnesota, New York, and Oregon: data from 3 health plans. *J Infect Dis* 2001;184:665-70.
36. Hak E, Nordin J, Wei F, et al. Influence of high-risk medical conditions on the effectiveness of influenza vaccination among elderly members of 3 large managed-care organizations. *Clin Infect Dis* 2002;35:370-7.
37. Nichol KL, Mendelman PM, Mallon KP, et al. Effectiveness of live, attenuated intranasal influenza virus vaccine in healthy, working adults: a randomized controlled trial. *JAMA* 1999;282:137-44.
38. Treanor JJ, Kotloff K, Betts RF, et al. Evaluation of trivalent, live, cold-adapted (CAIV-T) and inactivated (TIV) influenza vaccines in prevention of virus infection and illness following challenge of adults with wild-type influenza A (H1N1), A (H3N2), and B viruses. *Vaccine* 1999;18:899-906.
39. Salgado CD, Giannetta ET, Hayden FG, Farr BM. Preventing nosocomial influenza by improving the vaccine acceptance rate of clinicians. *Infect Control Hosp Epidemiol* 2004;25:923-8.
40. Saito R, Suzuki H, Oshitani H, Sakai T, Seki N, Tanabe N. The effectiveness of influenza vaccine against influenza A (H3N2) virus infections in nursing homes in Niigata, Japan, during the 1998-1999 and 1999-2000 seasons. *Infect Control Hosp Epidemiol* 2002;23:82-6.
41. Nguyen-Van-Tam J, Granfield R, Pearson J, Fleming D, Keating N. Do influenza epidemics affect patterns of sickness absence among British hospital staff? *Infect Control Hosp Epidemiol* 1999;20:691-4.
42. Jones A. The National Nursing Home Survey: 1999 summary. National Center for Health Statistics. *Vital Health Stat* 2002;13(152):1-125.
43. Horman JT, Stetler HC, Israel E, Sorley D, Schipper MT, Joseph JM. An outbreak of influenza A in a nursing home. *Am J Public Health* 1986;76:501-4.
44. Meiklejohn G, Hall H. Unusual outbreak of influenza A in a Wyoming nursing home. *J Am Geriatr Soc* 1987;35:742-6.
45. Goodman RA, Orenstein WA, Munro TF, Smith SC, Sikes RK. Impact of influenza A in a nursing home. *JAMA* 1982;247:1451-3.
46. Office of Technology Assessment. Cost effectiveness of influenza vaccination. Washington, DC: US Government Printing Office; 1981. Available at [http://www.wws.princeton.edu/ota/disk3/1981/8112\\_n.html](http://www.wws.princeton.edu/ota/disk3/1981/8112_n.html).
47. Mixeu MA, Vespa GN, Forleo-Neto E, Toniolo-Neto J, Alves PM. Impact of influenza vaccination on civilian aircrew illness and absenteeism. *Aviat Space Environ Med* 2002;73:876-80.
48. Nichol KL, Mallon KP, Mendelman PM. Cost benefit of influenza vaccination in healthy, working adults: an economic analysis based on the results of a clinical trial of trivalent live attenuated influenza virus vaccine. *Vaccine* 2003;21:2207-17.
49. Nichol KL, Mendelman P. Influence of clinical case definitions with differing levels of sensitivity and specificity on estimates of the relative and absolute health benefits of influenza vaccination among healthy working adults and implications for economic analyses. *Virus Res* 2004;103:3-8.
50. Nichol KL. Cost-benefit analysis of a strategy to vaccinate healthy working adults against influenza. *Arch Intern Med* 2001;161:749-59.
51. US Department of Health and Human Services. Healthy people 2010 (conference ed, in 2 vols). Washington, DC: US Department of Health and Human Services; 2000. Available at <http://www.health.gov/healthypeople>.
52. Bryant KA, Stover B, Cain L, Levine GL, Siegel J, Jarvis WR. Improving influenza immunization rates among healthcare workers caring for high-risk pediatric patients. *Infect Control Hosp Epidemiol* 2004;25:912-7.
53. Ikeda RM, Drabkin PD. Influenza A outbreaks in nursing homes [Comment]. *J Am Geriatr Soc* 1992;40:1288.
54. Odelin MF, Pozzetto B, Aymard M, Defayolle M, Jolly-Million J. Role of influenza vaccination in the elderly during an epidemic of A/H1N1 virus in 1988-1989: clinical and serological data. *Gerontology* 1993;39:109-16.
55. Sartor C, Tissot-Dupont H, Zandotti C, Martin F, Roques P, Drancourt M. Use of a mobile cart influenza program for vaccination of hospital employees. *Infect Control Hosp Epidemiol* 2004;25:918-22.
56. Russell DW, Cameron DJ, Lockey RF, Behnke RH, Sinnott JT, Ganguly R. Influenza vaccination acceptance among health care professionals. *Vaccine* 1991;9:691-2.
57. Heimberger T, Chang HG, Shaikh M, Crotty L, Morse D, Birkhead G. Knowledge and attitudes of healthcare workers about influenza: why are they not getting vaccinated? *Infect Control Hosp Epidemiol* 1995;16:412-5.
58. Ong AK, Srimanunthiphol J, Frankel RI. Influenza vaccination status of healthcare workers and the extent of their domestic contact with individuals at high risk for influenza-related complications. *Infect Control Hosp Epidemiol* 2000;21:735-7.
59. Goldstein AO, Kincade JE, Gamble G, Bearman RS. Policies and practices for improving influenza immunization rates among healthcare workers. *Infect Control Hosp Epidemiol* 2004;25:908-11.
60. Begue RE, Gee SQ. Improving influenza immunization among healthcare workers. *Infect Control Hosp Epidemiol* 1998;19:518-20.
61. LaVela SL, Smith B, Weaver FM, Legro MW, Goldstein B, Nichol K. Attitudes and practices regarding influenza vaccination among healthcare workers providing services to individuals with spinal cord injuries and disorders. *Infect Control Hosp Epidemiol* 2004;25:933-40.
62. Harbarth S, Siegrist CA, Schira JC, Wunderli W, Pittet D. Influenza immunization: improving compliance of healthcare workers. *Infect Control Hosp Epidemiol* 1998;19:337-42.
63. Weingarten S, Riedinger M, Bolton LB, Miles P, Ault M. Barriers to influenza vaccine acceptance. A survey of physicians and nurses. *Am J Infect Control* 1989;17:202-7.
64. Nichol KL, Hauge M. Influenza vaccination of healthcare workers. *Infect Control Hosp Epidemiol* 1997;18:189-94.
65. Watanakunakorn C, Ellis G, Gemmel D. Attitude of healthcare personnel regarding influenza immunization. *Infect Control Hosp Epidemiol* 1993;14:17-20.



66. Briss PA, Rodewald LE, Hinman AR, et al. Reviews of evidence regarding interventions to improve vaccination coverage in children, adolescents, and adults. The Task Force on Community Preventive Services. *Am J Prev Med* 2000;18(1 Suppl):97-140.
67. Poland GA, Tosh P, Jacobson RM. Requiring influenza vaccination for health care workers: seven truths we must accept. *Vaccine* 2005;23:2251-5.
68. Martinello RA, Jones L, Topal JE. Correlation between healthcare workers' knowledge of influenza vaccine and vaccine receipt. *Infect Control Hosp Epidemiol* 2003;24:845-7.
69. Nafziger DA, Herwaldt LA. Attitudes of internal medicine residents regarding influenza vaccination. *Infect Control Hosp Epidemiol* 1994;15:32-5.
70. Manuel DG, Henry B, Hockin J, Naus M. Health behavior associated with influenza vaccination among healthcare workers in long-term-care facilities. *Infect Control Hosp Epidemiol* 2002;23:609-14.
71. McArthur MA, Simor AE, Campbell B, McGeer A. Influenza vaccination in long-term-care facilities. *Infect Control Hosp Epidemiol* 1999;20:499-503.
72. Ohrt CK, McKinney WP. Achieving compliance with influenza immunization of medical house staff and students. A randomized controlled trial. *JAMA* 1992;267:1377-80.
73. Fedson DS, Houck P, Bratzler D. Hospital-based influenza and pneumococcal vaccination: Sutton's Law applied to prevention. *Infect Control Hosp Epidemiol* 2000;21:692-9.
74. Pachucki CT, Pappas SA, Fuller GF, Krause SL, Lentino JR, Schaaff DM. Influenza A among hospital personnel and patients. Implications for recognition, prevention, and control. *Arch Intern Med* 1989;149:77-80.
75. Adal KA, Flowers RH, Anglim AM, et al. Prevention of nosocomial influenza. *Infect Control Hosp Epidemiol* 1996;17:641-8.
76. Steiner M, Vermeulen LC, Mullahy J, Hayney MS. Factors influencing decisions regarding influenza vaccination and treatment: a survey of healthcare workers. *Infect Control Hosp Epidemiol* 2002;23:625-7.
77. Shannon SC. Community hospitals can increase staff influenza vaccination rates. *Am J Public Health* 1993;83:1174-5.
78. McKibben L, Horan TC, Tokars JI, et al. Guidance on public reporting of healthcare-associated infections: recommendations of the Healthcare Infection Control Practices Advisory Committee. *Infect Control Hosp Epidemiol* 2005;26:580-7.
79. Agerton TB, Mahoney FJ, Polish LB, Shapiro CN. Impact of the bloodborne pathogens standard on vaccination of healthcare workers with hepatitis B vaccine. *Infect Control Hosp Epidemiol* 1995;16:287-91.
80. US Department of Labor, Occupational Safety and Health Administration. 2a CFR Part 1910.1030. Occupational exposure to bloodborne pathogens: final rule. *Federal Register* 1991;56:64004-182.
81. Stewart A, Cox M, Rosenbaum S. The epidemiology of U.S. immunization law: immunization requirements for staff and residents of long-term care facilities under state laws/regulations. Washington, DC: George Washington University School of Public Health and Health Services; 2005. Available at <http://www.gwumc.edu/sphhs/healthpolicy/immunization/EUSIL-LTC-report.pdf>.
82. Cate TR, Couch RB, Parker D, Baxter B. Reactogenicity, immunogenicity, and antibody persistence in adults given inactivated influenza virus vaccines—1978. *Rev Infect Dis* 1983;5:737-47.
83. Kunzel W, Glathe H, Engelmann H, Van Hoecke C. Kinetics of humoral antibody response to trivalent inactivated split influenza vaccine in subjects previously vaccinated or vaccinated for the first time. *Vaccine* 1996;14:1108-10.
84. Poland GA, Borrud A, Jacobson RM, et al. Determination of deltoid fat pad thickness. Implications for needle length in adult immunization. *JAMA* 1997;277:1709-11.
85. Riddiough MA, Sisk JE, Bell JC. Influenza vaccination. *JAMA* 1983;249:3189-95.
86. King JC Jr, Lagos R, Bernstein DI, et al. Safety and immunogenicity of low and high doses of trivalent live cold-adapted influenza vaccine administered intranasally as drops or spray to healthy children. *J Infect Dis* 1998;177:1394-7.
87. Belshe RB, Gruber WC, Mendelman PM, et al. Correlates of immune protection induced by live, attenuated, cold-adapted, trivalent, intranasal influenza virus vaccine. *J Infect Dis* 2000;181:1133-7.
88. Boyce TG, Gruber WC, Coleman-Dockery SD, et al. Mucosal immune response to trivalent live attenuated intranasal influenza vaccine in children. *Vaccine* 1999;18:82-8.
89. Zangwill KM, Droge J, Mendelman P, et al. Prospective, randomized, placebo-controlled evaluation of the safety and immunogenicity of three lots of intranasal trivalent influenza vaccine among young children. *Pediatr Infect Dis J* 2001;20:740-6.
90. Bernstein DI, Yan L, Treanor J, Mendelman PM, Belshe R. Effect of yearly vaccinations with live, attenuated, cold-adapted, trivalent, intranasal influenza vaccines on antibody responses in children. *Pediatr Infect Dis J* 2003;22:28-34.
91. Nolan T, Lee MS, Cordova JM, et al. Safety and immunogenicity of a live-attenuated influenza vaccine blended and filled at two manufacturing facilities. *Vaccine* 2003;21:1224-31.
92. Vesikari T. Randomized, double-blind, placebo-controlled trial of the safety and transmissibility and phenotypic stability of a live, attenuated, cold-adapted influenza virus vaccine (CAIV-T) in children attending day care [Abstract G-450]. Presented at the 41st Annual Interscience Conference on Antimicrobial Agents and Chemotherapy (ICAAC); Chicago, Illinois, December 16-19, 2001.
93. Talbot TR, Crocker DD, Peters J, et al. Duration of virus shedding after trivalent intranasal live attenuated influenza vaccination in adults. *Infect Control Hosp Epidemiol* 2005;26:494-500.
94. Ali T, Scott N, Kallas W, et al. Detection of influenza antigen with rapid antibody-based tests after intranasal influenza vaccination (FluMist). *Clin Infect Dis* 2004;38:760-2.
95. CDC. General recommendations on immunization: recommendations of the Advisory Committee on Immunization Practices (ACIP) and the American Academy of Family Physicians (AAFP). *MMWR* 2002;51(No. RR-2):1-36.
96. Freeman DW, Barno A. Deaths from Asian influenza associated with pregnancy. *Am J Obstet Gynecol* 1959;78:1172-5.
97. Widelock D, Csizmas L, Klein S. Influenza, pregnancy, and fetal outcome. *Public Health Rep* 1963;78:1-11.
98. Harris JW. Influenza occurring in pregnant women: a statistical study of thirteen hundred and fifty cases. *JAMA* 1919;72:978-80.
99. Irving WL, James DK, Stephenson T, et al. Influenza virus infection in the second and third trimesters of pregnancy: a clinical and seroepidemiological study. *BJOG* 2000;107:1282-9.
100. Kort BA, Cefalo RC, Baker VV. Fatal influenza A pneumonia in pregnancy. *Am J Perinatol* 1986;3:179-82.



101. Kirshon B, Faro S, Zurawin RK, Samo TC, Carpenter RJ. Favorable outcome after treatment with amantadine and ribavirin in a pregnancy complicated by influenza pneumonia. A case report. *J Reprod Med* 1988;33:399–401.
102. Schoenbaum SC, Weinstein L. Respiratory infection in pregnancy. *Clin Obstet Gynecol* 1979;22:293–300.
103. Shahab SZ, Glezen WP. Influenza viruses. In: Gonik B, ed. *Viral diseases in pregnancy*. New York, NY: Springer-Verlag; 1994.
104. Neuzil KM, Reed GW, Mitchel EF, Simonsen L, Griffin MR. Impact of influenza on acute cardiopulmonary hospitalizations in pregnant women. *Am J Epidemiol* 1998;148:1094–102.
105. Heinonen OP, Shapiro S, Monson RR, Hartz SC, Rosenberg L, Slone D. Immunization during pregnancy against poliomyelitis and influenza in relation to childhood malignancy. *Int J Epidemiol* 1973;2:229–35.
106. National Foundation for Infectious Diseases. Call to action: influenza immunization among health-care personnel, 2003. Bethesda, MD: National Foundation for Infectious Diseases; 2003. Available at <http://www.nfid.org/publications/calltoaction.pdf>.
107. CDC. Interventions to increase influenza vaccination of health-care workers—California and Minnesota. *MMWR* 2005;54:196–9.
108. Gross PA, Russo C, Dran S, Cataruozolo P, Munk G, Lancey SC. Time to earliest peak serum antibody response to influenza vaccine in the elderly. *Clin Diagn Lab Immunol* 1997;4:491–2.
109. Brokstad KA, Cox RJ, Olofsson J, Jonsson R, Haaheim LR. Parenteral influenza vaccination induces a rapid systemic and local immune response. *J Infect Dis* 1995;171:198–203.
110. CDC. Updated recommendations from the Advisory Committee on Immunization Practices in response to delays in supply of influenza vaccine for the 2000–2001 season. *MMWR* 2000;49:888–92.
111. CDC. Delayed influenza vaccine availability for 2001–02 season and supplemental recommendations of the Advisory Committee on Immunization Practices. *MMWR* 2001;50:582–5.
112. CDC. Update: influenza vaccine supply and recommendations for prioritization during the 2005–06 influenza season. *MMWR* 2005;54:850.
113. CDC. Tiered use of inactivated influenza vaccine in the event of a vaccine shortage. *MMWR* 2005;54:749–50.
114. Govaert TM, Dinant GJ, Aretz K, Masurel N, Sprenger MJ, Knottnerus JA. Adverse reactions to influenza vaccine in elderly people: randomised double blind placebo controlled trial. *BMJ* 1993;307(6910):988–90.
115. Margolis KL, Nichol KL, Poland GA, Pluhar RE. Frequency of adverse reactions to influenza vaccine in the elderly. A randomized, placebo-controlled trial. *JAMA* 1990;264:1139–41.
116. Nichol KL, Margolis KL, Lind A, et al. Side effects associated with influenza vaccination in healthy working adults. A randomized, placebo-controlled trial. *Arch Intern Med* 1996;156:1546–50.
117. American Lung Association Asthma Clinical Research Centers. The safety of inactivated influenza vaccine in adults and children with asthma. *N Engl J Med* 2001;345:1529–36.
118. Scheifele DW, Bjornson G, Johnston J. Evaluation of adverse events after influenza vaccination in hospital personnel. *CMAJ* 1990;142:127–30.
119. Barry DW, Mayner RE, Hochstein HD, et al. Comparative trial of influenza vaccines. II. Adverse reactions in children and adults. *Am J Epidemiol* 1976;104:47–59.
120. Bierman CW, Shapiro GG, Pierson WE, Taylor JW, Foy HM, Fox JP. Safety of influenza vaccination in allergic children. *J Infect Dis* 1977;136(Suppl):S652–5.
121. James JM, Zeiger RS, Lester MR, et al. Safe administration of influenza vaccine to patients with egg allergy. *J Pediatr* 1998;133:624–8.
122. Murphy KR, Strunk RC. Safe administration of influenza vaccine in asthmatic children hypersensitive to egg proteins. *J Pediatr* 1985;106:931–3.
123. Zeiger RS. Current issues with influenza vaccination in egg allergy. *J Allergy Clin Immunol* 2002;110:834–40.
124. Aberer W. Vaccination despite thimerosal sensitivity. *Contact Dermatitis* 1991;24:6–10.
125. Kirkland LR. Ocular sensitivity to thimerosal: a problem with hepatitis B vaccine? *South Med J* 1990;83:497–9.
126. Schonberger LB, Bregman DJ, Sullivan-Bolyai JZ, et al. Guillain-Barré syndrome following vaccination in the National Influenza Immunization Program, United States, 1976–1977. *Am J Epidemiol* 1979;110:105–23.
127. Safranek TJ, Lawrence DN, Kurland LT, et al. Reassessment of the association between Guillain-Barré syndrome and receipt of swine influenza vaccine in 1976–1977: results of a two-state study. *Expert Neurology Group. Am J Epidemiol* 1991;133:940–51.
128. Hurwitz ES, Schonberger LB, Nelson DB, Holman RC. Guillain-Barré syndrome and the 1978–1979 influenza vaccine. *N Engl J Med* 1981;304:1557–61.
129. Kaplan JE, Katona P, Hurwitz ES, Schonberger LB. Guillain-Barré syndrome in the United States, 1979–1980 and 1980–1981. Lack of an association with influenza vaccination. *JAMA* 1982;248:698–700.
130. Chen R, Kent J, Rhodes PH, et al. Investigation of a possible association between influenza vaccination and Guillain-Barré syndrome in the United States, 1990–1991 [Abstract 40]. *Post Marketing Surveillance* 1992;6:5–6.
131. Lasky T, Terracciano GJ, Magder L, et al. The Guillain-Barré syndrome and the 1992–1993 and 1993–1994 influenza vaccines. *N Engl J Med* 1998;339:1797–802.
132. Ropper AH. The Guillain-Barré syndrome. *N Engl J Med* 1992;326:1130–6.
133. Prevots DR, Sutter RW. Assessment of Guillain-Barré syndrome mortality and morbidity in the United States: implications for acute flaccid paralysis surveillance. *J Infect Dis* 1997;175 (Suppl 1):S151–5.
134. Barohn RJ, Saperstein DS. Guillain-Barré syndrome and chronic inflammatory demyelinating polyneuropathy. *Semin Neurol* 1998;18:49–61.
135. Jackson LA, Holmes SJ, Mendelman PM, Huggins L, Cho I, Rhorer J. Safety of a trivalent live attenuated intranasal influenza vaccine, FluMist, administered in addition to parenteral trivalent inactivated influenza vaccine to seniors with chronic medical conditions. *Vaccine* 1999;17:1905–9.
136. King JC Jr, Treanor J, Fast PE, et al. Comparison of the safety, vaccine virus shedding, and immunogenicity of influenza virus vaccine, trivalent, types A and B, live cold-adapted, administered to human immunodeficiency virus (HIV)-infected and non-HIV-infected adults. *J Infect Dis* 2000;181:725–8.
137. Izurieta HS, Haber P, Ball R, et al. Post-licensure surveillance of the first live, cold-adapted influenza vaccine in the U.S. [Abstract]. *Pharmacoepidemiol Drug Saf* 2004;13(Suppl):S145.

## Healthcare Infection Control Practices Committee Membership List, June 2005

**Chairman:** Patrick J. Brennan, MD, University of Pennsylvania School of Medicine, Philadelphia, Pennsylvania.

**Executive Secretary:** Michele L. Pearson, MD, CDC, Atlanta, Georgia.

**Members:** Vicki L. Brinsko, Vanderbilt University Medical Center, Nashville, Tennessee; Raymond Y. W. Chinn, MD, Sharp Memorial Hospital, San Diego, California; E. Patchen Dellinger, MD, University of Washington School of Medicine, Seattle, Washington; Nancy E. Foster, American Hospital Association, District of Columbia; Steven M. Gordon, MD, Cleveland Clinic Foundation, Cleveland, Ohio; Lizzie J. Harrell, PhD, Duke University Medical Center, Durham, North Carolina; Carol O'Boyle, PhD, University of Minnesota, Minneapolis, Minnesota; Dennis M. Perrotta, PhD, Texas Department of Health, Austin, Texas; Harriett M. Pitt, MS, Long Beach Memorial Medical Center, Long Beach, California; Robert J. Sherertz, MD, Wake Forest University School of Medicine, Wake Forest, North Carolina; Nalini Singh, MD, Children's National Medical Center, District of Columbia; Kurt B. Stevenson, MD, Qualis Health, Boise, Idaho; Philip W. Smith, MD, University of Nebraska Medical Center, Omaha, Nebraska.

**Liaison Representatives:** William Baine, MD, Agency for Healthcare Research and Quality, District of Columbia; Joan Blanchard, MSS, Association of periOperative Registered Nurses, Denver, Colorado; Georgia Dash, MS, Association for Professionals of Infection Control and Epidemiology, Inc., Philadelphia, Pennsylvania; Sandra L. Fitzler, American Healthcare Association, District of Columbia; David Henderson, MD, National Institutes of Health, Bethesda, Maryland; Lorine Jay, Health Services Resources Administration, Atlanta, Georgia; Stephen F. Jencks, MD, Center for Medicare and Medicaid Services, Baltimore, Maryland; Chiu S. Lin, PhD, Food and Drug Administration, Rockville, Maryland; Mark Russi, MD, American College of Occupational and Environmental Medicine, New Haven, Connecticut; Rachel Stricoff, MPH, Advisory Committee for the Elimination of Tuberculosis, New York, New York; Michael Tapper, MD, Society for Healthcare Epidemiology of America, Inc., New York, New York; and Robert Wise, MD, Joint Commission on the Accreditation of Healthcare Organizations, Oakbrook, Illinois.

## Advisory Committee on Immunization Practices Membership List, June 2005

**Chairman:** Myron J. Levin, MD, Professor of Pediatrics and Medicine, University of Colorado Health Sciences Center, Denver, Colorado.

**Executive Secretary:** Larry Pickering, MD, National Immunization Program, CDC, Atlanta, Georgia.

**Members:** Jon S. Abramson, MD, Wake Forest University School of Medicine, Winston-Salem, North Carolina; Ban Mishu Allos, MD, Vanderbilt University School of Medicine, Nashville, Tennessee; Guthrie S. Birkhead, MD, New York State Department of Health, Albany, New York; Judith Campbell, MD, Baylor College of Medicine, Houston, Texas; Reginald Finger, MD, Focus on the Family, Colorado Springs, Colorado; Janet Gildsdorf, MD, University of Michigan, Ann Arbor, Michigan; Tracy Lieu, MD, Harvard Pilgrim Health Care and Harvard Medical School, Boston, Massachusetts; Edgar Marcuse, MD, Children's Hospital and Regional Medical Center, Seattle, Washington; Julia Morita, MD, Chicago Department of Health, Chicago, Illinois; Gregory Poland, MD, Mayo Clinic College of Medicine, Rochester, Minnesota; John B. Salamone, National Italian American Foundation, District of Columbia; Patricia Stinchfield, Children's Hospital and Clinics, St. Paul, Minnesota; John J. Treanor, MD, University of Rochester School of Medicine and Dentistry, Rochester, New York; Robin Womeodu, MD, University of Tennessee Health Sciences Center, Memphis, Tennessee.

**Ex-Officio Members:** James E. Cheek, MD, Indian Health Service, Albuquerque, New Mexico; Stephen Phillips, DO, Department of Defense, Falls Church, Virginia; Geoffrey S. Evans, MD, Health Resources and Services Administration, Rockville, Maryland; Bruce Gellin, MD, National Vaccine Program Office, District of Columbia; Linda Murphy, Centers for Medicare and Medicaid Services, Baltimore, Maryland; George T. Curlin, MD, National Institutes of Health, Bethesda, Maryland; Norman Baylor, MD, Food and Drug Administration, Bethesda, Maryland; Kristin Lee Nichol, MD, Department of Veterans Affairs, Minneapolis, Minnesota.

**Liaison Representatives:** American Academy of Family Physicians, Jonathan Temte, MD, Clarence, New York, and Richard Clover, MD, Louisville, Kentucky; American Academy of Pediatrics, Margaret Rennels, MD, Baltimore, Maryland, and Carol Baker, MD, Houston, Texas; American Association of Health Plans, Robert Scalettar, MD, North Haven, Connecticut; American College Health Association, James C. Turner, MD, Charlottesville, Virginia; American College of Obstetricians and Gynecologists, Stanley Gall, MD, Louisville, Kentucky; American College of Physicians, Kathleen Neuzil, MD, Seattle, Washington; American Medical Association, Litjen Tan, PhD, Chicago, Illinois; American Pharmacists Association, Stephan L. Foster, PharmD, Memphis, Tennessee; Association of Teachers of Preventive Medicine, W. Paul McKinney, MD, Louisville, Kentucky; Biotechnology Industry Organization, Clement Lewin, PhD, Cambridge, Massachusetts; Canadian National Advisory Committee on Immunization, Monica Naus, MD, Vancouver, British Columbia; Healthcare Infection Control Practices Advisory Committee, Steve Gordon, MD, Cleveland, Ohio; Infectious Diseases Society of America, Samuel L. Katz, MD, Durham, North Carolina, and William Schaffner, MD, Nashville, Tennessee; London Department of Health, David M. Salisbury, MD, London, United Kingdom; National Association of County and City Health Officials, Nancy Bennett, MD, Rochester, New York; National Coalition for Adult Immunization, David A. Neumann, PhD, Bethesda, Maryland; National Immunization Council and Child Health Program, Mexico, Romeo Rodriguez, Mexico City, Mexico; National Medical Association, Dennis A. Brooks, MD, Baltimore, Maryland; National Vaccine Advisory Committee, Charles Helms, MD, PhD, Iowa City, Iowa; Pharmaceutical Research and Manufacturers of America, Damian A. Braga, Swiftwater, Pennsylvania, Peter Paradiso, PhD, Collegeville, Pennsylvania; and Society for Adolescent Medicine, Amy Middleman, MD, Houston, Texas.