



September 06, 1996 / 45(RR-12);1-35

20011029 086

Update: Vaccine Side Effects, Adverse Reactions, Contraindications, and Precautions Recommendations of the Advisory Committee on Immunization Practices (ACIP)

Summary

This report provides updated information concerning the potential adverse events associated with vaccination for hepatitis B, poliomyelitis, measles, mumps, diphtheria, tetanus, and pertussis. This information incorporates findings from a series of recent literature reviews, conducted by an expert committee at the Institute of Medicine (IOM), of all evidence regarding the possible adverse consequences of vaccines administered to children. This report contains modifications to the previously published recommendations of the Advisory Committee on Immunization Practices (ACIP) and is based on an ACIP review of the IOM findings and new research on vaccine safety. In addition, this report incorporates information contained in the "Recommendations of the Advisory Committee on Immunization Practices: Use of Vaccines and Immune Globulins in Persons with Altered Immunocompetence" (MMWR 1993;42{No. RR-4}) and the "General Recommendations on Immunization: Recommendations of the Advisory Committee on Immunization Practices (ACIP)" (MMWR 1994;43{No. RR-1}). Major changes to the previous recommendations are highlighted within the text, and specific information concerning the following vaccines and the possible adverse events associated with their administration are included: hepatitis B vaccine and anaphylaxis; measles vaccine and a) thrombocytopenia and b) possible risk for death resulting from anaphylaxis or disseminated disease in immunocompromised persons; diphtheria and tetanus toxoids and pertussis vaccine (DTP) and chronic encephalopathy; and tetanus-toxoid-containing vaccines and a) Guillain-Barre syndrome, b) brachial neuritis, and c) possible risk for death resulting from anaphylaxis. These modifications will be incorporated into more comprehensive ACIP recommendations for each vaccine when such statements are revised. Also included in this report are interim recommendations concerning the use of measles and mumps vaccines in

- a. persons who are infected with human immunodeficiency virus and

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- b. persons who are allergic to eggs; ACIP is still evaluating these recommendations.

INTRODUCTION

Immunization has enabled the global eradication of smallpox (1), the elimination of poliomyelitis from the Western hemisphere (2), and major reductions in the incidence of other vaccine-preventable diseases in the United States (Table 1). However, although immunization has successfully reduced the incidence of vaccine-preventable diseases, vaccination can cause both minor and, rarely, serious side effects. Public awareness of and controversy about vaccine safety has increased, primarily because increases in vaccine coverage resulted in an increased number of adverse events that occurred after vaccination. Such adverse events include both true reactions to vaccine and events coincidental to, but not caused by, vaccination. Despite concerns about vaccine safety, vaccination is safer than accepting the risks for the diseases these vaccines prevent. Unless a disease has been eradicated (e.g., smallpox), failure to vaccinate increases the risks to both the individual and society.

In response to concerns about vaccine safety, the National Childhood Vaccine Injury Act of 1986 established a no-fault compensation process for persons possibly injured by selected vaccines (3). The Act also mandated that the Institute of Medicine * (IOM) review scientific and other evidence regarding the possible adverse consequences of vaccines administered to children.

IOM constituted an expert committee to review all available information on these vaccine adverse events; such information included epidemiologic studies, case series, individual case reports, and testimonials. To derive their conclusions, the IOM committee members created five categories of causality to describe the relationships between the vaccines and specific adverse events. The first IOM review examined certain events occurring after administration of pertussis and rubella vaccines (Table 2) (4). The second IOM review examined events occurring after administration of all other vaccines usually administered during childhood (i.e., diphtheria and tetanus toxoids and measles, mumps, hepatitis B, Haemophilus influenzae type b {Hib}, and poliovirus vaccines) (Table 3) (5). Two other IOM committees have met since the findings of the second review were published. These two committees have published their findings concerning both the diphtheria and tetanus toxoids and pertussis vaccine (DTP) and chronic nervous system dysfunction (Figure 1) (6) and research strategies for vaccine-associated adverse events (7).

The Advisory Committee on Immunization Practices (ACIP) recently reviewed the findings of the IOM committees and modified the previously published ACIP recommendations to ensure consistency with IOM conclusions. These recommendations, which are included in this report, update all previously published ACIP recommendations pertaining to the precautions, contraindications, side effects, and adverse reactions ** associated with specific vaccines. ACIP accepted the IOM conclusions for almost all vaccine adverse events; the few exceptions generally occurred because new information

that was available to ACIP had not been available when the IOM committees published their recommendations. These exceptions included a) oral poliovirus vaccine (OPV) and Guillain-Barre syndrome (GBS), b) tetanus-toxoid- containing vaccines and GBS, and c) DTP and chronic nervous system dysfunction.

In addition, this report incorporates information contained in the "Recommendations of the Advisory Committee on Immunization Practices: Use of Vaccines and Immune Globulins in Persons with Altered Immunocompetence" (MMWR 1993; 42{No. RR-4}) and the "General Recommendations on Immunization: Recommendations of the Advisory Committee on Immunization Practices (ACIP)" (MMWR 1994;43{No. RR-1}). To facilitate recognition of the new recommendations in this report, all major changes that are being made to the previously published ACIP statements are highlighted within the text. These changes include information on the following vaccines and the possible adverse events associated with their administration:

- Hepatitis B vaccine and anaphylaxis;
- Measles vaccine and a) thrombocytopenia and b) possible risk for death resulting from anaphylaxis or disseminated disease in immunocompromised persons;
- DTP and chronic encephalopathy; and
- Tetanus-toxoid-containing vaccines and a) GBS, b) brachial neuritis, and c) possible risk for death resulting from anaphylaxis.

The modifications contained in this report, and possibly other changes as new information becomes available, will be incorporated into more comprehensive ACIP recommendations for each vaccine when such statements are revised.

HEPATITIS B VACCINE

The following recommendations concerning adverse events associated with hepatitis B vaccination update those applicable sections in "Hepatitis B Virus: A Comprehensive Strategy for Eliminating Transmission in the United States Through Universal Childhood Vaccination -- Recommendations of the Immunization Practices Advisory Committee (ACIP)" (MMWR 1991;40{No. RR-13}).

Vaccine Side Effects and Adverse Reactions

Hepatitis B vaccines are safe to administer to adults and children. More than an estimated 10 million adults and 2 million infants and children have been vaccinated in the United States, and at least 12 million children have been vaccinated worldwide.

Vaccine-Associated Side Effects

Pain at the injection site (3%-29%) and a temperature greater than 37.7 C (1%-6%) have been among the most frequently reported side effects among adults and children receiving vaccine (8-12). In placebo-controlled studies, these side effects were reported no more frequently among vaccinees than among persons receiving a placebo (11,12). Among children receiving both hepatitis B vaccine and DTP, these mild side effects have been observed no more frequently than among children receiving only DTP.

The recommendation to begin hepatitis B vaccination soon after birth has raised the concern that a substantial number of infants will require an extensive medical evaluation for elevated temperatures secondary to hepatitis B vaccination. Several population-based studies to evaluate this possibility are in progress.

Adverse Events

In the United States, surveillance of adverse reactions indicated a possible association between GBS and receipt of the first dose of plasma-derived hepatitis B vaccine (CDC, unpublished data; 13). However, an estimated 2.5 million adults received one or more doses of recombinant hepatitis B vaccine during 1986-1990, and available data concerning these vaccinees do not indicate an association between receipt of recombinant vaccine and GBS (CDC, unpublished data).

Based on reports to the Vaccine Adverse Events Reporting System (VAERS), the estimated incidence rate of anaphylaxis among vaccine recipients is low (i.e., approximately one event per 600,000 vaccine doses distributed). Two of these adverse events occurred in children (CDC, unpublished data). In addition, only one case of anaphylaxis occurred among 100,763 children ages 10-11 years who had been vaccinated with recombinant vaccine in British Columbia (D. Scheifele, unpublished data), and no adverse events were reported among 166,757 children who had been vaccinated with plasma-derived vaccine in New Zealand (5). Although none of the persons who developed anaphylaxis died, this adverse event can be fatal; in addition, hepatitis B vaccine can -- in rare instances -- cause a life-threatening hypersensitivity reaction in some persons (5). Therefore, subsequent vaccination with hepatitis B vaccine is contraindicated for persons who have previously had an anaphylactic response to a dose of this vaccine.

Large-scale hepatitis B immunization programs for infants in Alaska, New Zealand, and Taiwan have not established an association between vaccination and the occurrence of other severe adverse events, including seizures and GBS (B. McMahon and A. Milne, unpublished data; 14). However, systematic surveillance for adverse reactions in these populations has been limited, and only a minimal number of children have received recombinant vaccine. Any presumed risk for adverse events that might be causally associated with hepatitis B vaccination must be balanced with the expected risk for hepatitis B virus (HBV)-related liver disease. Currently, an estimated 2,000-5,000 persons in each U.S. birth cohort will die as a result of HBV-related liver disease because

of the 5% lifetime risk for HBV infection.

As hepatitis B vaccine is introduced for routine vaccination of infants, surveillance for vaccine-associated adverse events will continue to be an important part of the program despite the current record of safety. Any adverse event suspected to be associated with hepatitis B vaccination should be reported to VAERS. VAERS forms can be obtained by calling (800) 822-7967.

POLIOMYELITIS PREVENTION

The following recommendations concerning adverse events associated with poliomyelitis vaccination update those applicable sections in "Poliomyelitis Prevention:

Recommendation of the Immunization Practices Advisory Committee (ACIP)" (MMWR 1982;31:22-6,31-4) and "Poliomyelitis Prevention: Enhanced-Potency Inactivated Poliomyelitis Vaccine -- Supplementary Statement" (MMWR 1987;36:795-8).

Precautions and Contraindications Pregnancy

Although no conclusive evidence documents the adverse effects of OPV or inactivated poliovirus vaccine (IPV) in pregnant women and their developing fetuses, vaccination of pregnant women should be avoided. However, if immediate protection against poliomyelitis is necessary, OPV or IPV can be given.

Immunodeficiency

Persons who have congenitally acquired immune-deficiency diseases (e.g., combined immunodeficiency, hypogammaglobulinemia, and agammaglobulinemia) should not be given OPV because of their substantially increased risk for vaccine-associated disease. Furthermore, persons who have altered immune status resulting from acquired conditions (e.g., human immunodeficiency virus {HIV} infection, leukemia, lymphoma, or generalized malignancy) or who have immune systems compromised by therapy (e.g., treatment with corticosteroids, alkylating drugs, antimetabolites, or radiation) should not receive OPV because of the theoretical risk for paralytic disease.

IPV -- and not OPV -- should be used to vaccinate immunodeficient persons and their household contacts. Many immunosuppressed persons are already immune to polioviruses because of previous vaccination or exposure to wild-type virus when they were immunocompetent. Although such persons should not receive OPV, their risk for paralytic disease may be less than that of persons who have congenitally acquired immunodeficiency. Although a protective immune response to IPV in the immunodeficient patient cannot be ensured, the vaccine is safe and some protection may result from its administration. If a household contact of an immunodeficient person is vaccinated inadvertently with OPV, the OPV recipient should avoid close physical contact with the immunodeficient person for approximately 4-6 weeks after vaccination.

(i.e., during the period of maximum excretion of vaccine virus). If such contact cannot be avoided, rigorous hygiene and hand washing after contact with feces (e.g., after diaper changing) and avoidance of contact with saliva (e.g., by not sharing eating utensils or food) should be practiced. These practices are an acceptable, but probably less effective, alternative than refraining from contact. Because immunodeficiency is possible in other children born to a family in which one child is immunodeficient, OPV should not be administered to a member of such a house-hold until the immune status of the recipient and other children in the family is documented.

Adverse Reactions OPV

In rare instances, administration of OPV has been associated with paralytic poliomyelitis in healthy recipients and their contacts. Very rarely, OPV has caused fatal paralytic poliomyelitis in immunocompromised persons (5). Other than efforts for identifying persons with immune-deficiency conditions, no procedures are currently available to identify persons likely to experience such adverse reactions. Although the risk of vaccine-associated paralysis is extremely small for vaccinees and their susceptible, close, personal contacts, they should be informed of this risk.

Available data do not indicate a measurable increased risk for GBS after receipt of OPV. Initial reports (at the time of IOM review) of two studies conducted in Finland suggested that OPV might cause GBS. These studies identified an apparent increased incidence of GBS that was temporally associated with mass OPV vaccination of children and adults who had previously received IPV (15,16). Since the IOM review, a reanalysis of the data derived from the studies conducted in Finland and an analysis of an observational study conducted in the United States have not demonstrated a causal relationship between OPV and GBS in infants (17).

Because OPV contains trace amounts of streptomycin, bacitracin, and neomycin, its use is contraindicated in persons who have previously had an anaphylactic reaction to OPV or to these antibiotics.

IPV

No serious side effects of currently available IPV have been documented. Since IPV contains trace amounts of streptomycin and neomycin, there is a possibility of hypersensitivity reactions in individuals sensitive to these antibiotics.

MEASLES PREVENTION

The following recommendations concerning adverse events associated with measles vaccination update those applicable sections in "Measles Prevention: Recommendations of the Immunization Practices Advisory Committee" (MMWR 1989; 38{No. S-9}), and they apply regardless of whether the vaccine is administered as a single antigen or as a

component of measles-rubella (MR) or measles-mumps-rubella (MMR) vaccine. Information concerning adverse events associated with the mumps component of MMR vaccine is reviewed later in this document (see Mumps Prevention), and information concerning the rubella component is located in the previously published ACIP statement for rubella (18).

Side Effects and Adverse Reactions

More than 240 million doses of measles vaccine were distributed in the United States from 1963 through 1993. The vaccine has an excellent record of safety. From 5% to 15% of vaccinees may develop a temperature of greater than or equal to 103 F (greater than or equal to 39.4 C) beginning 5-12 days after vaccination and usually lasting several days (19). Most persons with fever are otherwise asymptomatic. Transient rashes have been reported for approximately 5% of vaccinees. Central nervous system (CNS) conditions, including encephalitis and encephalopathy, have been reported with a frequency of less than one per million doses administered. The incidence of encephalitis or encephalopathy after measles vaccination of healthy children is lower than the observed incidence of encephalitis of unknown etiology. This finding suggests that the reported severe neurologic disorders temporally associated with measles vaccination were not caused by the vaccine. These adverse events should be anticipated only in susceptible vaccinees and do not appear to be age-related. After revaccination, most reactions should be expected to occur only among the small proportion of persons who failed to respond to the first dose.

Personal and Family History of Convulsions

As with the administration of any agent that can produce fever, some children may have a febrile seizure. Although children with a personal or family history of seizures are at increased risk for developing idiopathic epilepsy, febrile seizures following vaccinations do not in themselves increase the probability of subsequent epilepsy or other neurologic disorders. Most convulsions following measles vaccination are simple febrile seizures, and they affect children without known risk factors.

An increased risk of these convulsions may occur among children with a prior history of convulsions or those with a history of convulsions in first-degree family members (i.e., siblings or parents) (20). Although the precise risk cannot be determined, it appears to be low.

In developing vaccination recommendations for these children, ACIP considered a number of factors, including risks from measles disease, the large proportion (5%-7%) of children with a personal or family history of convulsions, and the fact that convulsions following measles vaccine are uncommon. Studies conducted to date have not established an association between MMR vaccination and the development of a residual seizure disorder (5). ACIP concluded that the benefits of vaccinating these children greatly outweigh the risks. They should be vaccinated just as children without such histories.

Because the period for developing vaccine-induced fever occurs approximately 5-12 days after vaccination, prevention of febrile seizures is difficult. Prophylaxis with antipyretics has been suggested as one alternative, but these agents may not be effective if given after the onset of fever. To be effective, such agents would have to be initiated before the expected onset of fever and continued for 5-7 days. However, parents should be alert to the occurrence of fever after vaccination and should treat their children appropriately.

Children who are being treated with anticonvulsants should continue to take them after measles vaccination. Because protective levels of most currently available anticonvulsant drugs (e.g., phenobarbital) are not achieved for some time after therapy is initiated, prophylactic use of these drugs does not seem feasible.

The parents of children who have either a personal or family history of seizures should be advised of the small increased risk of seizures following measles vaccination. In particular, they should be told in advance what to do in the unlikely event that a seizure occurs. The permanent medical record should document that the small risk of postimmunization seizures and the benefits of vaccination have been discussed.

Subacute Sclerosing Panencephalitis (SSPE)

Measles vaccine significantly reduces the likelihood of developing SSPE, as evidenced by the near elimination of SSPE cases after widespread measles vaccination began. SSPE has been reported rarely in children who do not have a history of natural measles infection but who have received measles vaccine. The available evidence suggests that at least some of these children may have had an unidentified measles infection before vaccination and that the SSPE probably resulted from the natural measles infection. The administration of live measles vaccine does not increase the risk for SSPE, regardless of whether the vaccinee has had measles infection or has previously received live measles vaccine (5,21).

Thrombocytopenia

Surveillance of adverse reactions in the United States and other countries indicates that MMR vaccine can, in rare circumstances, cause clinically apparent thrombocytopenia within the 2 months after vaccination. In prospective studies, the reported incidence of clinically apparent thrombocytopenia after MMR vaccination ranged from one case per 30,000 vaccinated children in Finland (22) and Great Britain (23) to one case per 40,000 in Sweden, with a temporal clustering of cases occurring 2-3 weeks after vaccination (24). With passive surveillance, the reported incidence was approximately one case per 100,000 vaccine doses distributed in Canada and France (25), and approximately one case per 1 million doses distributed in the United States (26). The clinical course of these cases was usually transient and benign, although hemorrhage occurred rarely (26). Furthermore, the risk for thrombocytopenia during rubella or measles infection is much greater than the risk after vaccination. Of 30,000 school-children in one Pennsylvania

county who had been infected with rubella during the 1963-64 measles epidemic, 10 children developed thrombocytopenic purpura (incidence: one case per 3,000 children) (27). Based on case reports, the risk for thrombocytopenia may be higher for persons who previously have had idiopathic thrombocytopenic purpura, particularly for those who had thrombocytopenic purpura after an earlier dose of MMR vaccine (5,28,29).

Revaccination Risks

There is no evidence of an increased risk for adverse reactions after administration of live measles vaccine to persons who are already immune to measles as a result of either previous vaccination or natural disease.

Precautions and Contraindications Pregnancy

Live measles vaccine, when given as a component of MR or MMR, should not be given to women known to be pregnant or who are considering becoming pregnant within the next 3 months. Women who are given monovalent measles vaccine should not become pregnant for at least 30 days after vaccination. This precaution is based on the theoretical risk of fetal infection, although no evidence substantiates this theoretical risk.

Considering the importance of protecting adolescents and young adults against measles, asking women if they are pregnant, excluding those who are, and explaining the theoretical risks to the others before vaccination are sufficient precautions.

Febrile Illness

The decision to administer or delay vaccination because of a current or recent febrile illness depends largely on the cause of the illness and the severity of symptoms. Minor illnesses, such as a mild upper-respiratory infection with or without low-grade fever, are not contraindications for vaccination. For persons whose compliance with medical care cannot be assured, every opportunity should be taken to provide appropriate vaccinations.

Children with moderate or severe febrile illnesses can be vaccinated as soon as they have recovered from the acute phase of the illness. This wait avoids superimposing adverse effects of vaccination on the underlying illness or mistakenly attributing a manifestation of the underlying illness to the vaccine. Performing routine physical examinations or measuring temperatures are not prerequisites for vaccinating infants and children who appear to be in good health. Asking the parent or guardian if the child is ill, postponing vaccination for children with moderate or severe febrile illnesses, and vaccinating those without contraindications are appropriate procedures in childhood immunization programs.

Allergic Reactions

Hypersensitivity reactions rarely occur after the administration of MMR or any of its

component vaccines. Most of these reactions are minor and consist of a wheal and flare or urticaria at the injection site. Immediate, anaphylactic reactions to MMR or its component vaccines are extremely rare. Although greater than 70 million doses of MMR vaccine have been distributed in the United States since VAERS was implemented in 1990, only 33 cases of anaphylactic reactions that occurred after MMR vaccination have been reported. Furthermore, only 11 of these cases a) occurred immediately after vaccination and b) occurred in persons who had symptoms consistent with anaphylaxis (CDC, unpublished data).

In the past, persons who had a history of anaphylactic reactions (i.e., hives, swelling of the mouth or throat, difficulty breathing, hypotension, and shock) following egg ingestion were considered to be at increased risk for serious reactions after receipt of measles-containing vaccines, which are produced in chick embryo fibroblasts. Protocols requiring caution were developed for skin testing and vaccinating persons who had had anaphylactic reactions after egg ingestion (30-34). However, the predictive value of such skin testing and the need for special protocols when vaccinating egg-allergic persons with measles-containing vaccines is uncertain. The results of recent studies suggest that anaphylactic reactions to measles-containing vaccines are not associated with hypersensitivity to egg antigens but with some other component of the vaccines. The risk for serious allergic reaction to these vaccines in egg-allergic patients is extremely low, and skin testing is not necessarily predictive of vaccine hypersensitivity (35-37). Therefore, ACIP is re-evaluating whether skin testing and the use of special protocols are routinely necessary when administering MMR or other measles-containing vaccines to persons who have a history of anaphylactic-like reactions after egg ingestion.

MMR and its component vaccines contain hydrolyzed gelatin as a stabilizer. The literature contains a single case report of a person with an anaphylactic sensitivity to gelatin who had an anaphylactic reaction after receipt of the MMR vaccine licensed in the United States (38). Similar cases have occurred in Japan (39). Therefore, ACIP is currently considering recommendations for vaccination of persons who have had an anaphylactic reaction to gelatin or gelatin-containing products. In the meantime, such persons should be vaccinated with MMR and its component vaccines with extreme caution.

MMR vaccine and its component vaccines contain trace amounts of neomycin. Although the amount present is less than that usually used for a skin test to determine hypersensitivity, persons who have experienced anaphylactic reactions to neomycin should not be given these vaccines. Most often, neomycin allergy is manifested by contact dermatitis rather than anaphylaxis. A history of contact dermatitis to neomycin is not a contraindication to receiving measles vaccine. Live measles virus vaccine does not contain penicillin.

Thrombocytopenia

Children who have a history of thrombocytopenic purpura or thrombocytopenia may be at increased risk for developing clinically significant thrombocytopenia after MMR vaccination. The decision to vaccinate should depend on the benefits of immunity to measles, mumps, and rubella and the risks for recurrence or exacerbation of thrombocytopenia after vaccination or during natural infections with measles or rubella. The benefits of immunization are usually greater than the potential risks, and administration of MMR vaccine is justified -- particularly with regard to the even greater risk for thrombocytopenia after measles or rubella disease. However, avoiding a subsequent dose might be prudent if the previous episode of thrombocytopenia occurred in close temporal proximity to (i.e., within 6 weeks after) the previous vaccination. Serologic evidence of measles immunity in such persons may be sought in lieu of MMR vaccination.

Recent Administration of Immune Globulins

Previous recommendations, based on data from persons who received low doses of immune globulin preparations, stated that MMR and its individual component vaccines could be administered as early as 6 weeks to 3 months after administration of immune globulins (40,41). However, recent evidence suggests that high doses of immune globulins can inhibit the immune response to measles vaccine for more than 3 months (42,43). Administration of immune globulins also can inhibit the response to rubella vaccine (42). The effect of immune globulin preparations on the response to mumps vaccine is unknown, but commercial immune globulin preparations contain antibodies to these viruses.

Blood (e.g., whole blood, packed red blood cells, and plasma) and other antibody-containing blood products (e.g., immune globulin; specific immune globulins; and immune globulin, intravenous {IGIV}) can diminish the immune response to MMR or its individual component vaccines. Therefore, after an immune globulin preparation is received, these vaccines should not be administered before the recommended interval (Table 4) and (Table 5). However, the postpartum vaccination of rubella-susceptible women with the rubella or MMR vaccine should not be delayed because anti-Rho(D) IG (human) or any other blood product was received during the last trimester of pregnancy or at delivery. These women should be vaccinated immediately after delivery and, if possible, tested at least 3 months later to ensure immunity to rubella and, if necessary, to measles.

If administration of an immune globulin preparation becomes necessary because of imminent exposure to disease, MMR or its component vaccines can be administered simultaneously with the immune globulin preparation, although vaccine-induced immunity might be compromised. The vaccine should be administered at a site remote from that chosen for the immune globulin inoculation. Unless serologic testing indicates that specific antibodies have been produced, vaccination should be repeated after the recommended interval (Table 4) and (Table 5).

If administration of an immune globulin preparation becomes necessary after MMR or its individual component vaccines have been administered, interference can occur. Usually, vaccine virus replication and stimulation of immunity will occur 1-2 weeks after vaccination. Thus, if the interval between administration of any of these vaccines and subsequent administration of an immune globulin preparation is less than 14 days, vaccination should be repeated after the recommended interval (Table 4) and (Table 5), unless serologic testing indicates that antibodies were produced.

Altered Immunocompetence

Non-HIV-Infected Persons. Replication of vaccine viruses can be enhanced in persons with immune-deficiency diseases and in persons with immunosuppression, as occurs with leukemia, lymphoma, generalized malignancy, or therapy with alkylating agents, antimetabolites, radiation, or large doses of corticosteroids. Evidence based on case reports has linked measles vaccine and measles infection to subsequent death in some severely immunocompromised children. Of the greater than 200 million doses of measles vaccine administered in the United States, fewer than five such deaths have been reported (5). Patients who have such conditions or are undergoing such therapies (excluding most HIV-infected patients) should not be given live measles virus vaccine.

Patients with leukemia in remission who have not received chemotherapy for at least 3 months may receive live-virus vaccines. The exact amount of systemically absorbed corticosteroids and the duration of administration needed to suppress the immune system of an otherwise healthy child are not well defined. Most experts agree that steroid therapy usually does not contraindicate administration of live virus vaccine when it is short term (i.e., less than 2 weeks); low to moderate dose; long-term, alternate-day treatment with short-acting preparations; maintenance physiologic doses (replacement therapy); or administered topically (skin or eyes), by aerosol, or by intra-articular, bursal, or tendon injection (44). Although of recent theoretical concern, no evidence of increased severe reactions to live vaccines has been reported among persons receiving steroid therapy by aerosol, and such therapy is not in itself a reason to delay vaccination. The immunosuppressive effects of steroid treatment vary, but many clinicians consider a dose equivalent to either 2 mg/kg of body weight or a total of 20 mg per day of prednisone as sufficiently immunosuppressive to raise concern about the safety of vaccination with live virus vaccines (44). Corticosteroids used in greater than physiologic doses also can reduce the immune response to vaccines. Physicians should wait at least 3 months after discontinuation of therapy before administering a live-virus vaccine to patients who have received high systemically absorbed doses of corticosteroids for greater than or equal to 2 weeks.

HIV-Infected Persons. Because of the increased risk for severe complications associated with measles infection and the absence of serious adverse events after measles vaccination among HIV-infected persons (41,45), ACIP has recommended that MMR vaccine be administered to all asymptomatic HIV-infected persons and that MMR

vaccine be considered for administration to all symptomatic HIV-infected persons who would otherwise be eligible for measles vaccine -- even though the immune response may be attenuated in such persons (41,44,45). There is a theoretical risk for an increase (probably transient) in HIV viral load following MMR vaccination because such effects have been observed with other vaccines (46,47).

Because of the recently reported case of pneumonitis in a measles vaccinee who had an advanced case of acquired immunodeficiency syndrome (AIDS) (48) and because of other evidence indicating a diminished antibody response to measles vaccination among severely immunocompromised persons (49), ACIP is re-evaluating the recommendations for vaccination of severely immunocompromised HIV-infected persons. In the interim, it may be prudent to withhold MMR or other measles-containing vaccines from HIV-infected persons with evidence of severe immunosuppression, defined as either a) CD4+ T-lymphocyte counts less than 750 for children ages less than 12 months, less than 500 for children ages 1-5 years, or less than 200 for persons ages greater than or equal to 6 years; or b) CD4+ T-lymphocytes constituting less than 15% of total lymphocytes for children ages less than 13 years or less than 14% for persons ages greater than or equal to 13 years (50,51).

ACIP continues to recommend MMR for HIV-infected persons without evidence of measles immunity (47) who are not severely immunocompromised (50,51). Severely immunocompromised and other symptomatic HIV-infected patients who are exposed to measles should receive immune globulin (IG), regardless of prior vaccination status (44). In addition, health-care providers should weigh the risks and benefits of measles vaccination or IG prophylaxis for severely immunocompromised HIV-infected patients who are at risk for measles exposure because of outbreaks or international travel.

Because the immunologic response to both live and killed antigen vaccines may decrease as HIV disease progresses (44,52), vaccination early in the course of HIV infection may be more likely to induce an immune response. Therefore, HIV-infected infants without severe immunosuppression should routinely receive MMR as soon as possible upon reaching their first birthday. Evaluation and testing of asymptomatic persons to identify HIV infection are not necessary before deciding to administer MMR or other measles-containing vaccine (44).

Management of Patients with Contraindications to Measles Vaccine

If immediate protection against measles is required for persons with contraindications to measles vaccination, passive immunization with IG, 0.25 mL/kg (0.11 mL/lb) of body weight (maximum dose=15 mL), should be given as soon as possible after known exposure. Exposed symptomatic HIV-infected and other immunocompromised persons should receive IG regardless of their previous vaccination status; however, IG in usual doses may not be effective in such patients. For immunocompromised persons, the recommended dose is 0.5 mL/kg of body weight if IG is administered intramuscularly

(maximum dose=15 mL). This corresponds to a dose of protein of approximately 82.5 mg/kg (maximum dose=2,475 mg). Intramuscular IG may not be needed if a patient with HIV infection is receiving 100-400 mg/kg IGIV at regular intervals and the last dose was given within 3 weeks of exposure to measles. Because the amounts of protein administered are similar, high-dose IGIV may be as effective as IG given intramuscularly. However, no data are available concerning the effectiveness of IGIV in preventing measles.

Simultaneous Administration of Vaccines

In general, simultaneous administration of the most widely used live and inactivated vaccines does not impair antibody responses or increase rates of adverse reactions (53). The administration of MMR vaccine yields results similar to the administration of individual measles, mumps, and rubella vaccines at different sites or at different times.

Vaccines recommended for administration at 12-15 months of age can be administered at either one or two visits. There are equivalent antibody responses and no clinically significant increases in the frequency of adverse events when DTP, MMR, and OPV (or IPV) vaccines and H. influenzae type b conjugate vaccine (HbCV) are administered either simultaneously at different sites or at separate times. If a child might not be brought back for future vaccinations, all vaccines (including DTP {or DTaP}, OPV {or IPV}, MMR, varicella, HbCV, and hepatitis B vaccines) may be administered simultaneously, as appropriate to the child's age and previous vaccination status.

MUMPS PREVENTION

The following recommendations concerning adverse events associated with mumps vaccination update those applicable sections in "Mumps Prevention" (MMWR 1989;38:388-92,397-400), and they apply regardless of whether the vaccine is administered as a single antigen or as a component of MR or MMR vaccine. Information concerning adverse events associated with the measles component of MMR vaccine is reviewed earlier in this document (see Measles Prevention), and information concerning the rubella component is located in the previously published ACIP statement for rubella (18).

Adverse Effects of Vaccine Use

In field trials before licensure, illnesses did not occur more often in vaccinees than in unvaccinated controls (54). Reports of illnesses following mumps vaccination have mainly been episodes of parotitis and low-grade fever. Allergic reactions including rash, pruritus, and purpura have been temporally associated with mumps vaccination but are uncommon and usually mild and of brief duration. The reported occurrence of encephalitis within 30 days of receipt of a mumps-containing vaccine (0.4 per million doses) is not greater than the observed background incidence rate of CNS dysfunction in

the normal population. Aseptic meningitis has been epidemiologically associated with receipt of the vaccine containing the Urabe strain of mumps virus, but not with the vaccine containing the Jeryl Lynn strain, the latter of which is used in vaccine distributed in the United States (5). During 1988-1992, 15 sentinel surveillance laboratories in the United Kingdom identified 13 aseptic meningitis cases that had occurred within 15-35 days after vaccination with the Urabe strain (i.e., 91 cases per 1 million doses distributed) (55). No vaccine-associated aseptic meningitis cases have been reported since 1992, when only the Jeryl Lynn strain has been used (23). Febrile seizures also have been infrequently reported. However, no evidence suggests that mumps vaccine causes residual seizure disorder (5). Although sensorineural deafness following mumps vaccination has been reported rarely, the data are inadequate to distinguish vaccine from nonvaccine causation. No association has been established between mumps vaccination and pancreatic damage or subsequent development of diabetes mellitus (5).

Contraindications to Vaccine Use Pregnancy

Although mumps vaccine virus has been shown to infect the placenta and fetus (56), there is no evidence that it causes congenital malformations in humans. However, because of the theoretical risk of fetal damage, it is prudent to avoid giving live virus vaccine to pregnant women. Live mumps vaccine, when combined with rubella vaccine, should not be administered to women known to be pregnant or who are considering becoming pregnant within the next 3 months. Women vaccinated with monovalent mumps vaccine should avoid becoming pregnant for 30 days after the vaccination. Routine precautions for vaccinating postpubertal women include asking if they are or may be pregnant, excluding those who say they are, and explaining the theoretical risk to those who plan to receive the vaccine. Vaccination during pregnancy should not be considered an indication for termination of pregnancy. However, the final decision about interruption of pregnancy must rest with the individual patient and her physician.

Severe Febrile Illness

Vaccine administration should not be postponed because of minor or intercurrent febrile illnesses, such as mild upper respiratory infections. However, vaccination of persons with severe febrile illnesses should generally be deferred until they have recovered from the acute phase of the illness.

Allergic Reactions

Hypersensitivity reactions rarely occur after the administration of MMR or any of its component vaccines. Most of these reactions are minor and consist of a wheal and flare or urticaria at the injection site. Immediate, anaphylactic reactions to MMR or its component vaccines are extremely rare. Although greater than 70 million doses of MMR vaccine have been distributed in the United States since VAERS was implemented in 1990, only 33 cases of anaphylactic reactions that occurred after MMR vaccination have

been reported. Furthermore, only 11 of these cases a) occurred immediately after vaccination and b) occurred in persons who had symptoms consistent with anaphylaxis (CDC, unpublished data).

In the past, persons who had a history of anaphylactic reactions (i.e., hives, swelling of the mouth or throat, difficulty breathing, hypotension, and shock) following egg ingestion were considered to be at increased risk for serious reactions after receipt of mumps-containing vaccines, which are produced in chick embryo fibroblasts. Protocols requiring caution were developed for skin testing and vaccinating persons who had had anaphylactic reactions after egg ingestion (30-34). However, the predictive value of such skin testing and the need for special protocols when vaccinating egg-allergic persons with mumps-containing vaccines is uncertain. The results of recent studies suggest that anaphylactic reactions to mumps-containing vaccines are not associated with hypersensitivity to egg antigens but with some other component of the vaccines. The risk for serious allergic reaction to these vaccines in egg-allergic patients is extremely low, and skin testing is not necessarily predictive of vaccine hypersensitivity (35-37). Therefore, ACIP is re-evaluating whether skin testing and the use of special protocols are routinely necessary when administering mumps-containing vaccines to persons who have a history of anaphylactic-like reactions after egg ingestion.

MMR and its component vaccines contain hydrolyzed gelatin as a stabilizer. The literature contains a single case report of a person with an anaphylactic sensitivity to gelatin who had an anaphylactic reaction after receipt of the MMR vaccine licensed in the United States (38). Similar cases have occurred in Japan (39). Therefore, ACIP is currently considering recommendations for vaccination of persons who have had an anaphylactic reaction to gelatin or gelatin-containing products. In the meantime, such persons should be vaccinated with MMR or other mumps vaccines with extreme caution.

Since mumps vaccine contains trace amounts of neomycin (25 ug), persons who have experienced anaphylactic reactions to topically or systemically administered neomycin should not receive mumps vaccine. Most often, neomycin allergy is manifested as a contact dermatitis, which is a delayed-type (cell-mediated) immune response, rather than anaphylaxis. In such persons, the adverse reaction, if any, to 25 ug of neomycin in the vaccine would be an erythematous, pruritic nodule or papule at the site of injection after 48-96 hours. A history of contact dermatitis to neomycin is not a contraindication to receiving mumps vaccine. Live mumps virus vaccine does not contain penicillin.

Recent Injection of Immune Globulin

The effect of immune globulin preparations on the response to mumps vaccine is unknown, but commercial immune globulin preparations contain mumps antibodies. Therefore, monovalent mumps or rubella-mumps vaccine should be given at least 2 weeks before the administration of an immune globulin preparation or deferred until approximately 3 months after the administration of an immune globulin preparation. For

suggested time intervals between administration of immune globulin preparations and vaccines containing live measles virus, refer to (Table 5).

Altered Immunocompetence

In theory, replication of the mumps vaccine virus may be potentiated in patients with immune deficiency diseases and by the suppressed immune responses that occur with leukemia, lymphoma, or generalized malignancy or with therapy with corticosteroids, alkylating drugs, antimetabolites, or radiation. In general, patients with such conditions should not be given live mumps virus vaccine. Because vaccinated persons do not transmit mumps vaccine virus, the risk of mumps exposure for those patients may be reduced by vaccinating their close susceptible contacts.

An exception to these general recommendations is in persons infected with HIV; asymptomatic HIV-infected children should receive MMR as soon as possible upon reaching their first birthday (44), and MMR vaccine should be considered for all symptomatic HIV-infected children who do not have evidence of severe immunosuppression (see Measles Prevention, Altered Immunocompetence).

Patients with leukemia in remission whose chemotherapy has been terminated for at least 3 months may also receive live mumps virus vaccine. Most experts agree that steroid therapy usually does not contraindicate administration of live virus vaccine when it is short term (i.e., less than 2 weeks); low to moderate dose; long-term, alternate-day treatment with short-acting preparations; maintenance physiologic doses (replacement therapy); or administered topically (skin or eyes), by aerosol, or by intraarticular, bursal, or tendon injection (44). However, mumps vaccine should be avoided if systemic immunosuppressive levels are reached by prolonged, extensive, topical application.

DTP

The following recommendations concerning adverse events associated with DTP vaccination update those applicable sections in "Diphtheria, Tetanus, and Pertussis: Recommendations for Vaccine Use and Other Preventive Measures -- Recommendations of the Immunization Practices Advisory Committee (ACIP)" (MMWR 1991;40{No. RR-10}).

Side Effects and Adverse Reactions Following DTP Vaccination

Local reactions (generally erythema and induration with or without tenderness) are common after the administration of vaccines containing diphtheria, tetanus, or pertussis antigens. Occasionally, a nodule may be palpable at the injection site of adsorbed products for several weeks. Sterile abscesses at the injection site have been reported rarely (6-10 events per million doses of DTP). Mild systemic reactions such as fever, drowsiness, fretfulness, and anorexia occur frequently. These reactions are substantially

more common following the administration of DTP than of DT, but they are self-limited and can be safely managed with symptomatic treatment.

Acetaminophen is frequently given by physicians to lessen fever and irritability associated with DTP vaccination, and it may be useful in preventing seizures among febrile-convulsion-prone children. However, fever that does not begin until greater than or equal to 24 hours after vaccination or persists for more than 24 hours after vaccination should not be assumed to be due to DTP vaccination. These new or persistent fevers should be evaluated for other causes so that treatment is not delayed for serious conditions such as otitis media or meningitis. Moderate-to-severe systemic events include high fever (i.e., temperature of greater than or equal to 40.5 C {greater than or equal to 105 F}); persistent, inconsolable crying lasting greater than or equal to 3 hours; collapse (hypotonic-hyporesponsive episode); or short-lived convulsions (usually febrile). These events occur infrequently. These events appear to be without sequelae (57-59). Other more severe neurologic events, such as a prolonged convulsion or encephalopathy, although rare, have been reported in temporal association with DTP administration.

Approximate rates for the occurrence of adverse events following receipt of DTP (regardless of dose number in the series or age of the child) are shown in (Table 6) (60,61). The frequencies of local reactions and fever are substantially higher with increasing numbers of doses of DTP, while other mild-to-moderate systemic reactions (e.g., fretfulness, vomiting) are substantially less frequent (59-61).

Concern about the possible role of pertussis vaccine in causing neurologic reactions has been present since the earliest days of vaccine use. Rare but serious acute neurologic illnesses, including encephalitis/encephalopathy and prolonged convulsions, have been anecdotally reported following receipt of whole-cell pertussis vaccine given as DTP (62,63). Whether pertussis vaccine causes or is only coincidentally related to such illnesses or reveals an inevitable event has been difficult to determine conclusively for the following reasons: a) serious acute neurologic illnesses often occur or become manifest among children during the first year of life irrespective of vaccination; b) there is no specific clinical sign, pathologic finding, or laboratory test which can determine whether the illness is caused by the DTP; c) it may be difficult to determine with certainty whether infants less than 6 months of age are neurologically normal, which complicates assessment of whether vaccinees were already neurologically impaired before receiving DTP; and d) because these events are exceedingly rare, appropriately designed large studies are needed to address the question.

Despite these methodologic difficulties, the National Childhood Encephalopathy Study (NCES) and other controlled epidemiologic studies have provided evidence that DTP can cause acute encephalopathy (64-68). This adverse event occurs rarely, with an estimated risk of zero to 10.5 episodes per million DTP vaccinations (68). A detailed follow-up of the NCES indicated that children who had had a serious acute neurologic illness after DTP administration were significantly more likely than children in the control group to

have chronic nervous system dysfunction 10 years later. These children with chronic nervous system dysfunction were more likely than children in the control group to have received DTP within 7 days of onset of the original serious acute neurologic illness (i.e., 12 {3.3%} of 367 children vs. six {0.8%} of 723 children) (69).

After reviewing the follow-up data, IOM concluded that the NCES provided evidence of an association between DTP and chronic nervous system dysfunction in children who had had a serious acute neurologic illness after vaccination with DTP. The committee proposed three possible explanations for this association. First, the acute neurologic illness and subsequent chronic nervous system dysfunction might have been caused by DTP. Second, DTP might trigger an acute neurologic illness and subsequent chronic nervous system dysfunction in children who have underlying brain or metabolic abnormalities. Such children might experience similar chronic dysfunction in the absence of DTP vaccination if other stimuli (e.g., fever or infection) are present. Third, DTP might cause an acute neurologic illness in children who have underlying brain or metabolic abnormalities that would inevitably have led to chronic nervous system dysfunction even if the acute neurologic illness had not developed (6). IOM concluded that the NCES data do not support one explanation over another.

According to IOM, the balance of evidence was consistent with a causal relationship between DTP and some forms of chronic nervous system disorders in children who had developed an acute neurologic disorder after receiving DTP. However, IOM also concluded that the results were insufficient to determine whether DTP increases the overall risk for chronic nervous system dysfunction in children.

A subcommittee of the National Vaccine Advisory Committee (NVAC) also reviewed the study and concluded that the results were insufficient to determine whether DTP administration before the acute neurologic event influenced the potential for neurologic dysfunction 10 years later (Ad hoc Subcommittee of the NVAC, unpublished data, 1994). ACIP concurs with this evaluation.

Although the NCES examined and reported risk for the 7 days after DTP vaccination, the increased risk for serious acute neurologic illness occurred primarily during the first 3 days after DTP administration (64). Thus, if an association between DTP and chronic encephalopathy exists, the risk is primarily in the first 3 days after DTP vaccination.

Among a subset of children who were participating in the NCES and who had infantile spasms, both DTP and DT vaccination appeared either to precipitate early manifestations of the condition or to lead to its identification by parents (70). IOM reviewed this and other studies and concluded that neither vaccine causes the illness (71,72).

Sudden infant death syndrome (SIDS) is listed on death certificates as the cause of death for 5,000-6,000 infants (ages 0-364 days) each year in the United States. Because the peak incidence of SIDS for infants occurs at 2-4 months of age, many instances of a close

temporal relation between SIDS and receipt of DTP are to be expected by simple chance. Only one methodologically rigorous study has suggested that DTP vaccination might cause SIDS (73). A total of four deaths were reported within 3 days of DTP vaccination, compared with 1.36 expected deaths. However, these deaths were unusual in that three of the four occurred within a 13-month interval during the 12-year study. These four children also tended to be vaccinated at older ages than their controls, suggesting that they might have had other unrecognized risk factors for SIDS independent of vaccination. In contrast, DTP vaccination was not associated with SIDS in several larger studies performed in the past decade (62,74-76). In addition, none of three studies that examined unexpected deaths among infants not classified as SIDS found an association with DTP vaccination (73,75,76). IOM reviewed these studies and concluded that the available information does not establish a causal relationship between DTP and SIDS (4).

IOM concluded recently that no available evidence indicates that DTP might cause transverse myelitis, other more subtle neurologic disorders (e.g., hyperactivity, learning disorders, and infantile autism), and progressive degenerative conditions of the CNS (4). Furthermore, one study indicated that children who received pertussis vaccine exhibited fewer school problems than those who did not, even after adjustment for socioeconomic status (77).

Recent data suggest that infants and young children who have ever had convulsions (febrile or afebrile) or who have immediate family members with such histories are more likely to have seizures following DTP vaccination than those without such histories (78,79). For those with a family history of seizures, the increased risks of seizures occurring within 3 days of receipt of DTP or 4-28 days following receipt of DTP are identical, suggesting that these histories are nonspecific risk factors and are unrelated to DTP vaccination (79).

Rarely, immediate anaphylactic reactions (i.e., swelling of the mouth, breathing difficulty, hypotension, or shock) have been reported after receipt of preparations containing diphtheria, tetanus, and/or pertussis antigens. However, no deaths caused by anaphylaxis following DTP vaccination have been reported to CDC since the inception of vaccine-adverse-events reporting in 1978, a period during which more than 80 million doses of publicly purchased DTP vaccine were administered. While substantial underreporting exists in this passive surveillance system, the severity of anaphylaxis and its immediacy following vaccination suggest that such events are likely to be reported. Although no causal relation to any specific component of DTP has been established, the occurrence of true anaphylaxis usually contraindicates further doses of any one of these components. Rashes that are macular, papular, petechial, or urticarial and appear hours or days after a dose of DTP are frequently antigen-antibody reactions of little consequence or are due to other causes, such as viral illnesses, and are unlikely to recur following subsequent injections (80,81). In addition, there is no evidence for a causal relation between DTP vaccination and hemolytic anemia or thrombocytopenic purpura.

Precautions and Contraindications General Considerations

The decision to administer or delay DTP vaccination because of a current or recent febrile illness depends largely on the severity of the symptoms and their etiology. Although a moderate or severe febrile illness is sufficient reason to postpone vaccination, minor illnesses such as mild upper-respiratory infections with or without low-grade fever are not contraindications. If ongoing medical care cannot be assured, taking every opportunity to provide appropriate vaccinations is particularly important.

Children with moderate or severe illnesses with or without fever can receive DTP as soon as they have recovered. Waiting a short period before administering DTP avoids superimposing the adverse effects of the vaccination on the underlying illness or mistakenly attributing a manifestation of the underlying illness to vaccination.

Routine physical examinations or temperature measurements are not prerequisites for vaccinating infants and children who appear to be in good health. Appropriate immunization practice includes asking the parent or guardian if the child is ill, postponing DTP vaccination for those with moderate or severe acute illnesses, and vaccinating those without contraindications or precautionary circumstances.

When an infant or child returns for the next dose of DTP, the parent should always be questioned about any adverse events that might have occurred following the previous dose.

A history of prematurity generally is not a reason to defer vaccination (82-84). Preterm infants should be vaccinated according to their chronological age from birth.

Immunosuppressive therapies -- including irradiation, antimetabolites, alkylating agents, cytotoxic drugs, and corticosteroids (used in greater than physiologic doses) -- may reduce the immune response to vaccines. Short-term (less than 2-week) corticosteroid therapy or intra-articular, bursal, or tendon injections with corticosteroids should not be immunosuppressive. Although no specific studies with pertussis vaccine are available, if immunosuppressive therapy will be discontinued shortly, it is reasonable to defer vaccination until the patient has been off therapy for 1 month; otherwise, the patient should be vaccinated while still on therapy (85).

Special Considerations for Preparations Containing Pertussis Vaccine

Precautions and contraindications guidelines that were previously published regarding the use of pertussis vaccine were based on three assumptions about the risks for adverse events associated with pertussis vaccination: a) that the vaccine on rare occasions caused acute encephalopathy resulting in permanent brain damage; b) that pertussis vaccine aggravated preexisting CNS disease; and c) that certain nonencephalitic reactions are predictive of more severe reactions with subsequent doses (86). In addition, children from

whom pertussis vaccine was withheld were thought to be well protected by herd immunity, a belief that is no longer valid. The current revised ACIP recommendations reflect better understanding of the risks associated not only with pertussis vaccine but also with pertussis disease.

Contraindications

If any of the following events occur in temporal relationship to the administration of DTP, further vaccination with DTP is contraindicated (Table 7):

1. An immediate anaphylactic reaction. The rarity of such reactions to DTP is such that they have not been adequately studied. Because of uncertainty as to which component of the vaccine might be responsible, no further vaccination with any of the three antigens in DTP should be carried out. Alternatively, because of the importance of tetanus vaccination, such individuals may be referred for evaluation by an allergist and desensitized to tetanus toxoid if a specific allergy can be demonstrated (87,88).
2. Encephalopathy (not due to another identifiable cause). This is defined as an acute, severe CNS disorder occurring within 7 days following vaccination and generally consisting of major alterations in consciousness, unresponsiveness, generalized or focal seizures that persist more than a few hours, with failure to recover within 24 hours. Even though causation by DTP cannot be established, no subsequent doses of pertussis vaccine should be given. It may be desirable to delay for months before administering the balance of the doses of DT necessary to complete the primary schedule. Such a delay allows time for clarification of the child's neurologic status.

Precautions

If any of the following events occur in temporal relation to receipt of DTP, the decision to give subsequent doses of vaccine containing the pertussis component should be carefully considered (Table 7). Although these events were considered absolute contraindications in previous ACIP recommendations, there may be circumstances, such as a high incidence of pertussis, in which the potential benefits outweigh possible risks, particularly because these events are not associated with permanent sequelae (86). The following events were previously considered contraindications and are now considered precautions:

1. Temperature of greater than or equal to 40.5 C (greater than or equal to 105 F) within 48 hours not due to another identifiable cause. Such a temperature is considered a precaution because of the likelihood that fever following a subsequent dose of DTP also will be high. Because such febrile reactions are usually attributed to the pertussis component, vaccination with DT should not be discontinued.

2. Collapse or shock-like state (hypotonic-hyporesponsive episode) within 48 hours. Although these uncommon events have not been recognized to cause death nor to induce permanent neurological sequelae, it is prudent to continue vaccination with DT, omitting the pertussis component (58,89).
3. Persistent, inconsolable crying lasting greater than or equal to 3 hours, occurring within 48 hours. Follow-up of infants who have cried inconsolably following DTP vaccination has indicated that this reaction, though unpleasant, is without long-term sequelae and not associated with other reactions of greater significance (59). Inconsolable crying occurs most frequently following the first dose and is less frequently reported following subsequent doses of DTP (60). However, crying for greater than 30 minutes following DTP vaccination can be a predictor of increased likelihood of recurrence of persistent crying following subsequent doses (59). Children with persistent crying have had a higher rate of substantial local reactions than children who had other DTP-associated reactions (including high fever, seizures, and hypotonic-hyporesponsive episodes), suggesting that prolonged crying was really a pain reaction (89).
4. Convulsions with or without fever occurring within 3 days. Short-lived convulsions, with or without fever, have not been shown to cause permanent sequelae (57,90). Furthermore, the occurrence of prolonged febrile seizures (i.e., status epilepticus ***), irrespective of their cause, involving an otherwise normal child does not substantially increase the risk for subsequent febrile (brief or prolonged) or afebrile seizures. The risk is significantly increased ($p=0.018$) only among those children who are neurologically abnormal before their episode of status epilepticus (91). Accordingly, although a convulsion following DTP vaccination has previously been considered a contraindication to further doses, under certain circumstances subsequent doses may be indicated, particularly if the risk of pertussis in the community is high. If a child has a seizure following the first or second dose of DTP, it is desirable to delay subsequent doses until the child's neurologic status is better defined. By the end of the first year of life, the presence of an underlying neurologic disorder has usually been determined and appropriate treatment instituted. DT vaccine should not be administered before a decision has been made about whether to restart the DTP series. Regardless of which vaccine is given, it is prudent also to administer acetaminophen, 15 mg/kg of body weight, at the time of vaccination and every 4 hours subsequently for 24 hours (92,93).

Vaccination of infants and young children who have underlying neurologic disorders

Infants and children with recognized, possible, or potential underlying neurologic conditions present a unique problem. They seem to be at increased risk for the appearance of manifestations of the underlying neurologic disorder within 2-3 days after vaccination. However, more prolonged manifestations or increased progression of the disorder or exacerbation of the disorder as a result of DTP vaccination have not been

recognized (94). In addition, most neurologic conditions in infancy and young childhood are associated with evolving, changing neurological findings. Functional abnormalities are often unmasked by progressive neurologic development. Thus, confusion over the interpretation of progressive neurologic signs may arise when DTP vaccination or any other therapeutic or preventive measure is carried out.

Protection against diphtheria, tetanus, and pertussis is as important for children with neurologic disabilities as for other children. Such protection may be even more important for neurologically disabled children. They often receive custodial care or attend special schools where the risk of pertussis is greater because DTP vaccination is avoided for fear of adverse reactions. Also, if pertussis affects a neurologically disabled child who has difficulty in handling secretions and in cooperating with symptomatic care, it may aggravate preexisting neurologic problems because of anoxia, intracerebral hemorrhages, and other manifestations of the disease. Whether and when to administer DTP to children with proven or suspected underlying neurologic disorders must be decided on an individual basis. Important considerations include the current local incidence of pertussis, the near absence of diphtheria in the United States, and the low risk of infection with *Clostridium tetani*. On the basis of these considerations and the nature of the child's disorder, the following approaches are recommended:

1. Infants and children with previous convulsions. Infants and young children who have had prior seizures, whether febrile or afebrile, appear to be at increased risk for seizures following DTP vaccination than children and infants without these histories (79). A convulsion within 3 days of DTP vaccination in a child with a history of convulsions may be initiated by fever caused by the vaccine in a child prone to febrile seizures, may be induced by the pertussis component, or may be unrelated to the vaccination. As noted earlier, current evidence indicates that seizures following DTP vaccination do not cause permanent brain damage. Among infants and children with a history of previous seizures, it is prudent to delay DTP vaccination until the child's status has been fully assessed, a treatment regimen established, and the condition stabilized. It should be noted, however, that delaying DTP vaccination until the second 6 months of life will increase the risk of febrile seizures among persons who are predisposed. When DTP or DT is given, acetaminophen, 15 mg/kg, should also be given at the time of the vaccination and every 4 hours for the ensuing 24 hours (92,93).
2. Infants as yet unvaccinated who are suspected of having underlying neurologic disease. It is prudent to delay initiation of vaccination with DTP or DT (but not other vaccines) until further observation and study have clarified the child's neurologic status and the effect of treatment. The decision as to whether to begin vaccination with DTP or DT should be made no later than the child's first birthday.
3. Children who have not received a complete series of vaccine and who have a neurologic event occurring between doses. Infants and children who have received

one or more doses of DTP and who experience a neurologic disorder (e.g., a seizure) not temporally associated with vaccination, but before the next scheduled dose, present a special management challenge. If the seizure or other disorder occurs before the first birthday and before completion of the first three doses of the primary series of DTP, further doses of DTP or DT (but not other vaccines) should be deferred until the infant's status has been clarified. The decision whether to use DTP or DT to complete the series should be made no later than the child's first birthday and should take into consideration the nature of the child's problem and the benefits and possible risks of the vaccine. If the seizure or other disorder occurs after the first birthday, the child's neurologic status should be evaluated to ensure that the disorder is stable before a subsequent dose of DTP is given.

4. Infants and children with stable neurologic conditions. Infants and children with stable neurologic conditions, including well-controlled seizures, may be vaccinated. The occurrence of single seizures (temporally unassociated with DTP) do not contraindicate DTP vaccination, particularly if the seizures can be satisfactorily explained. Parents of infants and children with histories of convulsions should be informed of the increased risk of postvaccination seizures. Acetaminophen, 15 mg/kg, every 4 hours for 24 hours, should be given to children with such histories to reduce the possibility of postvaccination fever (92,93).
5. Children with resolved or corrected neurologic disorders. DTP vaccination is recommended for infants with certain neurologic problems, such as neonatal hypocalcemic tetany or hydrocephalus (following placement of a shunt and without seizures), that have been corrected or have clearly subsided without residua.

Vaccination of infants and young children who have a family history of convulsion or other CNS disorder

A family history of convulsions or other CNS disorder is not a contraindication to pertussis vaccination (95). Acetaminophen should be given at the time of DTP vaccination and every 4 hours for 24 hours to reduce the possibility of postvaccination fever (92,93).

Preparations Containing Diphtheria Toxoid and Tetanus Toxoid

The only contraindication to tetanus and diphtheria toxoids is a history of a neurologic or severe hypersensitivity reaction following a previous dose. Vaccination with tetanus and diphtheria toxoids is not known to be associated with an increased risk of convulsions. Local side effects alone do not preclude continued use. If an anaphylactic reaction to a previous dose of tetanus toxoid is suspected, intradermal skin testing with appropriately diluted tetanus toxoid may be useful before a decision is made to discontinue tetanus toxoid vaccination (86). In one study, 94 of 95 persons with histories of anaphylactic symptoms following a previous dose of tetanus toxoid were nonreactive following

intradermal testing and tolerated further tetanus toxoid challenge without incident (86). One person had erythema and induration immediately following skin testing, but tolerated a full IM dose without adverse effects. Mild, nonspecific skin-test reactivity to tetanus toxoid, particularly if used undiluted, appears to be fairly common. Most vaccinees develop inconsequential cutaneous delayed hypersensitivity to the toxoid. Although very rare, severe hypersensitivity reactions may occur after receipt of tetanus-toxoid-containing vaccines; these reactions can be life-threatening (5).

Persons who experienced Arthus-type hypersensitivity reactions or a temperature of greater than 103 F (greater than 39.4 C) following a prior dose of tetanus toxoid usually have high serum tetanus antitoxin levels and should not be given even emergency doses of Td more frequently than every 10 years, even if they have a wound that is neither clean nor minor.

If a contraindication to using tetanus-toxoid-containing preparations exists for a person who has not completed a primary series of tetanus toxoid immunization and that person has a wound that is neither clean nor minor, only passive immunization should be given using tetanus IG (TIG).

On the basis of a) a report of a 42-year-old man who had GBS on three separate occasions after receipt of tetanus toxoid and b) evidence that a vaccine-induced immunologic response can cause GBS, IOM concluded that tetanus-toxoid-containing vaccines can trigger the onset of GBS in adults. GBS can be a life-threatening disease. Persons who have a history of GBS associated with a particular vaccine may be at increased risk for recurrent GBS after subsequent doses of that vaccine (5). However, in a study in which an estimated 1.2 million doses of tetanus-containing toxoid were administered to persons greater than 18 years of age, two cases of GBS were expected by chance alone during the 6 weeks after vaccination, and only one case was reported (CDC, unpublished data). This finding suggests that the risk for GBS after administration of tetanus toxoid is extremely low.

No increased risk for GBS has been observed with the use of DTP in children. In a study of 0.7 million children of preschool-ages who were vaccinated with DTP during a 7-year period, three cases of GBS were expected by chance alone during the 6 weeks after vaccination, and only two cases were reported (17).

Because tetanus vaccination has been associated rarely with recurrence of GBS, the decision to administer additional doses of tetanus-toxoid-containing vaccine to persons who have had GBS within 6 weeks after receiving tetanus toxoid should be based on the benefits of subsequent vaccination and the risk for recurrence of GBS. For example, vaccination is usually justified for children whose primary immunization schedules are incomplete (i.e., fewer than three doses have been received); but routine booster vaccination probably is not indicated for adults who have received three or more doses.

Vaccination with tetanus-toxoid-containing vaccines has been associated with brachial neuritis in adult vaccinees, with a relative risk of 5-10 in comparison with the population-based background incidence and a 1-month attributable incidence of approximately one-half to one case per 100,000 recipients of tetanus toxoid (5).

Although no evidence exists that tetanus and diphtheria toxoids are teratogenic, waiting until the second trimester of pregnancy to administer Td is a reasonable precaution for minimizing any concern about the theoretical possibility of such reactions.

Misconceptions Concerning Contraindications to DTP

Some health-care providers inappropriately consider certain conditions or circumstances as contraindications to DTP vaccination. These include the following:

1. Soreness, redness, or swelling at the DTP vaccination site or temperature of less than 40.5 C (less than 105 F).
2. Mild, acute illness with low-grade fever or mild diarrheal illness affecting an otherwise healthy child.
3. Current antimicrobial therapy or the convalescent phase of an acute illness.
4. Recent exposure to an infectious disease.
5. Prematurity. The appropriate age for initiating vaccination among the prematurely born infant is the usual chronological age from birth (82-84). Full doses (0.5 mL) of vaccine should be used.
6. History of allergies or relatives with allergies.
7. Family history of convulsions.
8. Family history of SIDS.
9. Family history of an adverse event following DTP vaccination.

References

1. World Health Organization. The global eradication of smallpox: final report of the Global Commission for the Certification of Smallpox Eradication. In: History of international public health. No. 4. Geneva, Switzerland: World Health Organization, 1980.
2. CDC. Certification of poliomyelitis eradication -- the Americas, 1994. MMWR 1994;43:720-2.

3. The National Childhood Vaccine Injury Act of 1986, section 2125 of the Public Health Service Act as codified at 42 U.S.C. 300aa-(suppl 1987).
4. Institute of Medicine, Howson CP, Howe CJ, Fineberg HV, eds. Adverse effects of pertussis and rubella vaccines. Washington, DC: National Academy Press, 1991.
5. Institute of Medicine, Stratton KR, Howe CJ, Johnston RB, eds. Adverse events associated with childhood vaccines: evidence bearing on causality. Washington, DC: National Academy Press, 1994.
6. Institute of Medicine, Stratton KR, Howe CJ, Johnston RB, eds. DPT vaccine and chronic nervous system dysfunction: a new analysis. Washington, DC: National Academy Press, 1994.
7. Institute of Medicine, Stratton KR, Howe CJ, Johnston RB, eds. Research strategies for assessing adverse events associated with vaccines: a workshop summary. Washington, DC: National Academy Press, 1994.
8. Stevens CE, Taylor PE, Tong MJ, et al. Yeast-recombinant hepatitis B vaccine: efficacy with hepatitis B immune globulin in prevention of perinatal hepatitis B virus transmission. *JAMA* 1987;257:2612-6.
9. Zajac BA, West DJ, McAleer WJ, Scolnick EM. Overview of clinical studies with hepatitis B vaccine made by recombinant DNA. *J Infect* 1986;13(suppl A):39-45.
10. Andre FE. Summary of safety and efficacy data on a yeast-derived hepatitis B vaccine. *Am J Med* 1989;87(suppl 3A): 14s-20s.
11. Szmunes W, Stevens CE, Harley EJ, et al. Hepatitis B vaccine: demonstration of efficacy in a controlled clinical trial in a high-risk population in the United States. *N Engl J Med* 1980; 303:833-41.
12. Francis DP, Hadler SC, Thompson SE, et al. The prevention of hepatitis B with vaccine: report of the Centers for Disease Control multi-center efficacy trial among homosexual men. *Ann Intern Med* 1982;97:362-6.
13. Shaw FE Jr, Graham DJ, Guess HA, et al. Postmarketing surveillance for neurologic adverse events reported after hepatitis B vaccination: experience of the first three years. *Am J Epidemiol* 1988;127:337-52.
14. Chen D-S. Control of hepatitis B in Asia: mass immunization program in Taiwan. In: Hollinger FB, Lemon SM, Margolis HS, eds. *Viral hepatitis and liver disease*. Baltimore: Williams & Wilkins, 1991:716-9.
15. Kinnunen E, Farkkila M, Hovi T, et al. Incidence of Guillain-Barre syndrome

during a nationwide oral poliovirus vaccine campaign. *Neurology* 1989;39:1034-6.

16. Uhari M, Rantala H, Niemela M, et al. Cluster of childhood Guillain-Barre cases after an oral poliovaccine campaign. *Lancet* 1989;2:440-1.
17. Rantala J, Cherry JD, Shields WD, et al. Epidemiology of Guillain-Barre syndrome in children: relationship of oral polio vaccine administration to occurrence. *J Pediatr* 1994;124:220-3.
18. ACIP. Rubella prevention: recommendations of the Immunization Practices Advisory Committee. *MMWR* 1990;39:1-18.
19. Peltola H, Heinonen O. Frequency of true adverse reactions to measles-mumps-rubella vaccine. *Lancet* 1986;1:939-42.
20. CDC. Adverse events following immunization. Atlanta: US Department of Health and Human Services, Public Health Service, CDC, 1989. (Surveillance report no. 3, 1985-1986).
21. American Academy of Pediatrics. Measles. In: Peter G, ed. 1994 Red book: report of the Committee on Infectious Diseases. 23rd ed. Elk Grove Village, IL: American Academy of Pediatrics, 1994:317-8.
22. Nieminen U, Peltola H, Syrjala MT, et al. Acute thrombocytopenic purpura following measles, mumps and rubella vaccination: a report on 23 patients. *Acta Paediatr* 1993;82:267-70.
23. Farrington P, Pugh S, Colville A, et al. A new method for active surveillance of adverse events from diphtheria/tetanus/pertussis and measles/mumps/rubella vaccines. *Lancet* 1995;345: 567-9.
24. Bottinger M, Christenson B, Romanus V, et al. Swedish experience of two dose vaccination programme aiming at eliminating measles, mumps, and rubella. *Br Med J* 1987;295:1264-7.
25. Jonville-Bera AP, Autret E, Galy-Eyraud C, Hessel L. Thrombocytopenic purpura after measles, mumps, and rubella vaccination: a retrospective survey by the French Regional Pharmacovigilance Centres and Pasteur-Merieux Serums et Vaccins. *Pediatr Infect Dis J* 1996;15:44-8.
26. Beeler J, Varricchio F, Wise R. Thrombocytopenia after immunization with measles vaccines: review of the Vaccine Adverse Events Reporting System (1990 to 1994). *Pediatr Infect Dis J* 1996;15:88-90.
27. Bayer WL, Sherman FE, Micheals RH, et al. Purpura in congenital and acquired

rubella. *N Engl J Med* 1965;273:1362-6.

28. Drachtman RA, Murphy S, Ettinger LJ, et al. Exacerbation of chronic idiopathic thrombocytopenic purpura following measles-mumps-rubella immunization. *Arch Pediatr Adolesc Med* 1994;148:326-7.
29. Vlach A, Forma EN, Miron D, Peter G. Recurrent thrombocytopenic purpura after repeated measles-mumps-rubella vaccination. *Pediatrics* 1996;97:738-9.
30. American Academy of Pediatrics. Active immunization. In: Peter G, ed. 1994 Red book: report of the Committee on Infectious Diseases. 23rd ed. Elk Grove Village, IL: American Academy of Pediatrics, 1994:36-7.
31. Lavi S, Zimmerman B, Koren G, Gold R. Administration of measles, mumps, and rubella virus vaccine (live) to egg-allergic children. *JAMA* 1990;263:269-71.
32. Greenberg MA, Birx DL. Safe administration of mumps-measles-rubella vaccine in egg-allergic children. *J Pediatr* 1988;113:504-6.
33. Herman JJ, Radin R, Schneiderman R. Allergic reactions to measles (rubeola) vaccine in patients hypersensitive to egg protein. *J Pediatr* 1983;102:196-9.
34. Stiehm ER. Skin testing prior to measles vaccination for egg-sensitive patients {Editorial}. *Am J Dis Child* 1990;144:32.
35. Kemp A, Van Asperen P, Mukhi A. Measles immunization in children with clinical reactions to egg protein. *Am J Dis Child* 1990;144:33-5.
36. Fasano MB, Wood RA, Cooke SK, et al. Egg hypersensitivity and adverse reactions to measles, mumps, and rubella vaccine. *J Pediatr* 1992;120:878-81.
37. James JM, Burks AW, Roberson PK, et al. Safe administration of the measles vaccine to children allergic to eggs. *N Engl J Med* 1995;332:1262-6.
38. Kelso JM, Jones RT, Yunginger JW, et al. Anaphylaxis to measles, mumps, and rubella vaccine mediated by IgE to gelatin. *J Allergy Clin Immunol* 1993;91:867-72.
39. Sakaguchi M, Ogura H, Inouye S. IgE antibody to gelatin in children with immediate-type reactions to measles and mumps vaccines. *J Allergy Clin Immunol* 1995;96:563-5.
40. ACIP. General recommendations on immunization: recommendations of the Immunization Practices Advisory Committee (ACIP). *MMWR* 1989;38:205-14, 219-27.

41. ACIP. Measles prevention: recommendations of the Immunization Practices Advisory Committee. MMWR 1989;38(No. S-9):1-18.
42. Siber GR, Werner BC, Halsey NA. Interference of immune globulin with measles and rubella immunization. J Pediatr 1993;122:204-11.
43. Mason W, Takahashi M, Schneider T. Persisting passively acquired measles antibody following gamma globulin therapy for Kawasaki disease and response to live virus vaccination. Presented at the 32nd meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy {Abstract 311}. Los Angeles, California, October 1992.
44. ACIP. Recommendations of the Advisory Committee on Immunization Practices (ACIP): use of vaccines and immune globulins in persons with altered immunocompetence. MMWR 1993;42(No. RR-4).
45. ACIP. General recommendations on immunization: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 1994;43(No. RR-1).
46. O'Brien WA, Grovit-Ferbas K, Namazi A, et al. Human immunodeficiency virus-type 1 replication can be increased in peripheral blood of seropositive patients after influenza vaccination. Blood 1995;86:1082-9.
47. Stanley SK, Ostrowski MA, Justement JS, et al. Effect of immunization with a common recall antigen on viral expression in patients infected with human immunodeficiency virus type 1. N Engl J Med 1996;334:1222-30.
48. CDC. Measles pneumonitis following measles-mumps-rubella vaccination of a patient with HIV infection, 1993. MMWR 1996;45:603-6.
49. Palumbo P, Hoyt L, DeMasio K, Oleske J, Connor E. Population-based study of measles and measles immunization in human immunodeficiency virus-infected children. Pediatr Infect Dis J 1992;11:1008-14.
50. CDC. 1993 Revised classification system for HIV infection and expanded surveillance case definition for AIDS among adolescents and adults. MMWR 1992;41(No. RR-17).
51. CDC. 1994 Revised classification system for human immunodeficiency virus infection in children less than 13 years of age. MMWR 1994;43(No. RR-12):1-10.
52. Arpadi SM, Markowitz LE, Baughman AL, et al. Measles antibody in vaccinated human immunodeficiency virus type 1-infected children. Pediatrics 1996;97:653-7.

53. Deforest A, Long SS, Lischner HW, et al. Simultaneous administration of measles-mumps-rubella vaccine with booster doses of diphtheria-tetanus-pertussis and poliovirus vaccines. *Pediatrics* 1988;81:237-46.
54. Hilleman MR, Buynak EB, Weibel RE, et al. Live, attenuated mumps-virus vaccine. *N Engl J Med* 1968;278:227-32.
55. Miller E, Goldacre M, Pugh S, et al. Risk of aseptic meningitis after measles, mumps, and rubella vaccine in UK children. *Lancet* 1993;341:979-95.
56. Yamauchi T, Wilson C, St. Geme JW Jr. Transmission of live, attenuated mumps virus to the human placenta. *N Engl J Med* 1974;290:710-2.
57. Hirtz DG, Nelson KB, Ellenberg JH. Seizures following childhood immunizations. *J Pediatr* 1983;102:14-8.
58. Baraff LJ, Shields WD, Beckwith L, et al. Infants and children with convulsions and hypotonic-hyporesponsive episodes following diphtheria-tetanus-pertussis immunization: follow-up evaluation. *Pediatrics* 1988;81:789-94.
59. Long SS, DeForest A, Pennridge Pediatric Associates, Smith DG, Lazaro C, Wassilak SGF. Longitudinal study of adverse reactions following diphtheria-tetanus-pertussis vaccine in infancy. *Pediatrics* 1990;85:294-302.
60. Cody CL, Baraff LJ, Cherry JD, Marcy SM, Manclark CR. The nature and rate of adverse reactions associated with DTP and DT immunization in infants and children. *Pediatrics* 1981; 68:650-60.
61. Baraff LJ, Cody CL, Cherry JD. DTP-associated reactions: an analysis by injection site, manufacturer, prior reactions and dose. *Pediatrics* 1984;73:31-6.
62. Cherry JD, Brunell PA, Golden GS, Karzon DT. Report of the Task Force on Pertussis and Pertussis Immunization -- 1988. *Pediatrics* 1988;81(suppl):939-84.
63. Kulenkampff M, Schwartzman JS, Wilson J. Neurological complications of pertussis inoculation. *Arch Dis Child* 1974;49:46-9.
64. Alderslade R, Bellman MH, Rawson NSB, et al. The National Childhood Encephalopathy Study: a report on 1000 cases of serious neurological disorders in infants and young children from the NCES research team. In: Department of Health and Social Security. Whooping cough: reports from the Committee on the Safety of Medicines and the Joint Committee on Vaccination and Immunization. London: Her Majesty's Stationery Office, 1981.
65. Walker AM, Jick H, Perera DR, et al. Neurologic events following diphtheria-

tetanus-pertussis immunization. *Pediatrics* 1988;81:345-9.

66. Gale JL, Thapa PB, Wassilak SGF, et al. Risk of serious acute neurological illness after immunization with diphtheria-tetanus-pertussis vaccine. *JAMA* 1994;271:37-41.
67. Griffin MR, Ray WA, Mortimer EA, et al. Risk of seizures and encephalopathy after immunization with the diphtheria-tetanus-pertussis vaccine. *JAMA* 1990;263:1641-5.
68. Institute of Medicine, Howson CP, Howe CJ, Fineberg HV, eds. Adverse effects of pertussis and rubella vaccines. Washington, DC: National Academy Press, 1991:86-124.
69. Miller DL, Madge N, Diamond J, et al. Pertussis immunization and serious acute neurological illnesses in children. *Br Med J* 1993;307:1171-6.
70. Bellman MH, Ross EM, Miller DL. Infantile spasms and pertussis immunization. *Lancet* 1983; 1:1031-4.
71. Shields WD, Nielsen C, Buch D, et al. Relationship of pertussis immunization to the onset of neurologic disorders: a retrospective epidemiologic study. *J Pediatr* 1988;113:801-5.
72. Melchior JC. Infantile spasms and early immunization against whooping cough: Danish survey from 1970 to 1975. *Arch Dis Child* 1977;52:134-7.
73. Walker AM, Jick H, Perera DR, Thompson RS, Knause TA. Diphtheria-tetanus-pertussis immunization and sudden infant death syndrome. *Am J Public Health* 1987;77:945-51.
74. Hoffman HS, Hunter JC, Damus K, et al. Diphtheria-tetanus-pertussis immunization and sudden infant death: results of the National Institute of Child Health and Human Development Cooperative Epidemiological Study of Sudden Infant Death Syndrome Risk Factors. *Pediatrics* 1987;79:598-611.
75. Griffin MR, Ray WA, Livengood JR, Schaffner W. Risk of sudden infant death syndrome (SIDS) after immunization with the diphtheria-tetanus-pertussis vaccine. *N Engl J Med* 1988;319: 618-23.
76. Bouvier-Colle MH, Flahaut A, Messiah A, Jouglu E, Hatton F. Sudden infant death and immunization: an extensive epidemiological approach to the problem in France -- Winter 1986. *Int J Epidemiol* 1989;18:121-6.
77. Butler NR, Haslum M, Golding J, Stewart-Brown S. Recent findings from the 1970

Child Health and Education Study: preliminary communication. J R Soc Med 1982;75:781-4.

78. Stetler HC, Orenstein WA, Bart KJ, Brink EW, Brennan J-P, Hinman AT. History of convulsions and use of pertussis vaccine. J Pediatr 1985;107:175-9.
79. Livengood JR, Mullen JR, White JW, Brink EW, Orenstein WA. Family history of convulsions and use of pertussis vaccine. J Pediatr 1989;115:527-31.
80. Mortimer EA Jr, Sorensen RU. Urticaria following administration of diphtheria-tetanus toxoids-pertussis vaccine. Pediatr Infect Dis 1987;6:876-7.
81. Lewis K, Jordan SC, Cherry JD, Sakai RS, Le CT. Petechiae and urticaria after DTP vaccination: detection of circulating immune complexes containing vaccine-specific antigens. J Pediatr 1986;109:1009-12.
82. Bernbaum J, Daft A, Samuelson J, Polin RA. Half-dose immunization for diphtheria, tetanus, pertussis: response of preterm infants. Pediatrics 1989;83:471-6.
83. Bernbaum J, Anolik R, Polin RA, Douglas SD. Development of the premature infants host defense and its relationship to routine immunizations. Clin Perinatol 1984;11:73-84.
84. Koblin BA, Townsend TR, Munoz A, Onorato I, Wilson M, Polk BF. Response of preterm infants to diphtheria-tetanus-pertussis vaccine. Pediatr Infect Dis J 1988;7:704-11.
85. Gross PA, Lee H, Wolff JA, Hall CB, Minnefore AB, Lazicki ME. Influenza immunization in immunosuppressed children. J Pediatr 1978;92:30-5.
86. ACIP. Diphtheria, tetanus, and pertussis: guidelines for vaccine prophylaxis and other preventive measures -- recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1985;34:405-14,419-26.
87. Jacobs RL, Lowe RS, Lanier BQ. Adverse reactions to tetanus toxoid. JAMA 1982;247:40-2.
88. Mansfield LE, Ting S, Rawls DO, Frederick R. Systemic reactions during cutaneous testing for tetanus toxoid hypersensitivity. Ann Allergy 1986;57:135-7.
89. Blumberg DA, Mink CM, Lewis K, et al. Severe DTP-associated reactions {Abstract}. In: Manclark CR, ed. The Sixth International Symposium on Pertussis, Abstracts. Bethesda, MD: US Department of Health and Human Services, Public Health Service, Food and Drug Administration, 1990:223-4; DHHS publication no. (FDA)90-1162.

90. Ellenberg JH, Hirtz DG, Nelson KB. Do seizures in children cause intellectual deterioration? N Engl J Med 1986;314:1085-8.
91. Maytal J, Shinnar S. Febrile status epilepticus. Pediatrics 1990;86:611-6.
92. Ipp MM, Gold R, Greenberg S, et al. Acetaminophen prophylaxis of adverse reactions following vaccination of infants with diphtheria-pertussis-tetanus toxoids-polio vaccine. Pediatr Infect Dis J 1987;6:721-5.
93. Lewis K, Cherry JD, Sachs MH, et al. The effect of prophylactic acetaminophen administration on reactions to DTP vaccination. Am J Dis Child 1988;142:62-5.
94. Livingston S. Comprehensive management of epilepsy in infancy. Springfield, IL: Charles C. Thomas, 1972:159-66.
95. ACIP. Pertussis immunization: family history of convulsions and use of antipyretics -- supplementary ACIP statement: recommendations of the Immunization Practices Advisory Committee (ACIP). MMWR 1987;36:281-2.
 - o An independent research organization chartered by the National Academy of Sciences. ** In this publication, the terms "side effects" and "adverse reactions" are used interchangeably to denote the undesirable secondary effects resulting from vaccination. *** Any seizure lasting greater than 30 minutes or recurrent seizures lasting a total of 30 minutes without the child regaining full consciousness.

+----- --+ | Erratum: Vol. 45, No. RR-12 ||
 || ===== || SOURCE: MMWR 46(10) DATE: Mar 14, 1997 ||
 || || In the MMWR Recommendations and Reports entitled "Update: ||
 Vaccine Side Effects, Adverse Reactions, Contraindications, and || Precautions --
 Recommendations of the Advisory Committee on || Immunization Practices (ACIP)," the
 first complete paragraph on || page 10 should read: "OPV should not be administered to
 persons || who have experienced an anaphylactic reaction to a previous dose || of OPV."
 The next paragraph should read: "IPV should not be || administered to persons who have
 experienced a) an anaphylactic || reaction following a previous dose of IPV or b) an
 anaphylactic || reaction to streptomycin, polymyxin B, or neomycin." | +-----
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Table_1

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 1. The maximum number of cases of specified vaccine-preventable diseases ever reported for a calendar year compared with the number of cases of disease and vaccine adverse events reported for 1995 -- United States

Category	Maximum no. reported cases during prevaccine era	Year(s) maximum no. cases reported	Reported no. cases during 1995 *
Disease			
Congenital rubella syndrome	20,000 +	1964-65	7
Diphtheria	206,939	1921	0
Invasive Haemophilus influenzae	20,000 +	1984	1,164
Measles	894,134	1941	309
Mumps	152,209	1968	840
Pertussis	265,269	1934	4,315
Poliomyelitis (wild)	21,269	1952	0
Rubella	57,686	1969	146
Tetanus	601	1948	34
Vaccine adverse events &	0		10,594

* Provisional totals.

+ Estimated because national reporting did not exist in the prevaccine era.

& Total number reported to the Vaccine Adverse Events Reporting System (VAERS).

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Table_2

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 2. Summarized conclusions of evidence regarding the possible association between specific adverse effects and receipt of diphtheria and tetanus toxoids and pertussis vaccine (DTP) * and RA 27/3 measles-mumps-rubella (MMR) + vaccine, by determination of causality -- Institute of Medicine, 1991 &

Conclusion, by determination of causality	Adverse event reviewed	
	DTP vaccine	RA 27/3
1. No evidence was available to establish a causal relationship	Autism	None
2. Inadequate evidence to accept or reject a causal relationship	Aseptic meningitis Chronic neurologic damage @ Erythema multiforme or other rash Guillain-Barre syndrome Hemolytic anemia Juvenile diabetes Learning disabilities and attention-deficit disorder	Radiculo other Thromboc purpur

	Peripheral mononeuropathy Thrombocytopenia	
3. Evidence favored rejection of a causal relationship	Infantile spasms Hypsarrythmia Reye syndrome Sudden infant death syndrome	None
4. Evidence favored acceptance of a causal relationship	Acute encephalopathy ** Shock and unusual shock-like state	Chronic
5. Evidence established a causal relationship	Anaphylaxis Protracted, inconsolable crying	Acute ar

* The evidence only differentiated between components of DTP in the event of protra inconsolable crying, for which the evidence specifically implicated the pertussis component.

+ Trivalent MMR vaccine containing the RA 27/3 rubella strain.

& This table is an adaptation of a table published previously by the Institute of M (4), an independent research organization chartered by the National Academy of Sc The National Childhood Vaccine Injury Act of 1986 mandated that IOM review scient other evidence (e.g., epidemiologic studies, case series, individual case reports testimonials) regarding the possible adverse consequences of vaccines administere children. IOM constituted an expert committee to review and summarize all availab information; this committee created five categories of causality to describe the between the vaccines and specific adverse events.

@ IOM reviewed this adverse event again in 1994 (5).

** Defined in the controlled studies that were reviewed as encephalopathy, encephali encephalomyelitis.

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Table_3

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 3. Summarized conclusions of evidence regarding the possible association betwe receipt of childhood vaccines, by determination of causality -- Institute of Medicin

DT/Td/Tetanus toxoid +	Measles vaccine &	Mumps vaccine &	OPV/IPV @
1. No evidence was available to establish a causal relationship			
None	None	Neuropathy Residual seizure disorder	Transverse (IPV) Thrombocyt (IPV) Anaphylaxi
2. Inadequate evidence to accept or reject a causal relationship			
Residual seizure	Encephalopathy	Encephalopathy	Transverse

disorder other than infantile spasms	Subacute sclerosing panencephalitis	Aseptic meningitis	(OPV)
Demyelinating	Residual seizure	Sensorineural	Guillain-B
diseases of the	disorder	deafness (MMR)	syndrome
central nervous	Sensorineural	Insulin-dependent	Death from
system	deafness (MMR)	diabetes mellitus	
Mononeuropathy	Optic neuritis	Sterility	
Arthritis	Transverse myelitis	Thrombocytopenia	
Erythema multiforme	Guillain-Barre	Anaphylaxis **	
	syndrome		
	Thrombocytopenia		
	Insulin-dependent		
	diabetes mellitus		

3. Evidence favored rejection of a causal relationship

Encephalopathy &&	None	None	None
Infantile spasms			
(DT only) @@			
Death from SIDS			
(DT only) @@ ***			

4. Evidence favored acceptance of a causal relationship

Guillain-Barre	Anaphylaxis **	None	Guillain-B
syndrome +++ &&&			syndrome
Brachial neuritis +++			

5. Evidence established a causal relationship

Anaphylaxis +++	Thrombocytopenia	None	Poliomyeli
	(MMR)		recipien
	Anaphylaxis (MMR) **		(OPV)
	Death from measles-		Death from
	vaccine-strain viral		vaccine-
	infection ++ @@@		infectio

* This table is an adaptation of a table published previously by the Institute of the National Academy of Sciences. The National Childhood Vaccine Injury Act of 1986 (Public Law 99-361) requires the Institute of Medicine (IOM) to conduct epidemiologic studies, case series, individual case reports, and testimonials) r to children. IOM constituted an expert committee to review and summarize all ava to describe the relationships between the vaccines and specific adverse events.

+ DT=diphtheria and tetanus toxoids for pediatric use; Td=diphtheria and tetanus t
& If the data derived from studies of a monovalent preparation, then the causal re exclusively from studies of the measles-mumps-rubella (MMR) vaccine, the vaccine the monovalent preparation, the causal relationship determined for the multivale

@ For some adverse events, the IOM committee was charged with assessing the causal vaccine (OPV) (i.e., for poliomyelitis) or only inactivated poliovirus vaccine (two vaccines differed for the other adverse events, the vaccine to which the adv

** The evidence used to establish a causal relationship for anaphylaxis applies to favored acceptance of a causal relationship, but this evidence was less convinci of symptoms or the possible attenuation of symptoms by medical intervention.

++ This table lists weight-of-evidence determinations only for deaths that were cla a consequence of vaccine-strain viral infection. However, if the evidence favore

vaccine and a possibly fatal adverse event, then the evidence also favored the a and death from the adverse event. Direct evidence regarding death in association Guillain-Barre syndrome, b) tetanus toxoid and anaphylaxis, and c) OPV and polio

&& The evidence derived from studies of DT. If the evidence favored rejection of a also favored rejection of a causal relationship between Td and tetanus toxoid an

@@ Infantile spasms and SIDS occur only in an age group that is administered DT but

*** The evidence derived primarily from studies of DTP, although the evidence also f

+++ The evidence derived from studies of tetanus toxoid. If the evidence favored acc and an adverse event, then the evidence also favored acceptance of (or establish

&&& This conclusion differs from the information contained in the ACIP recommendatio published this table.

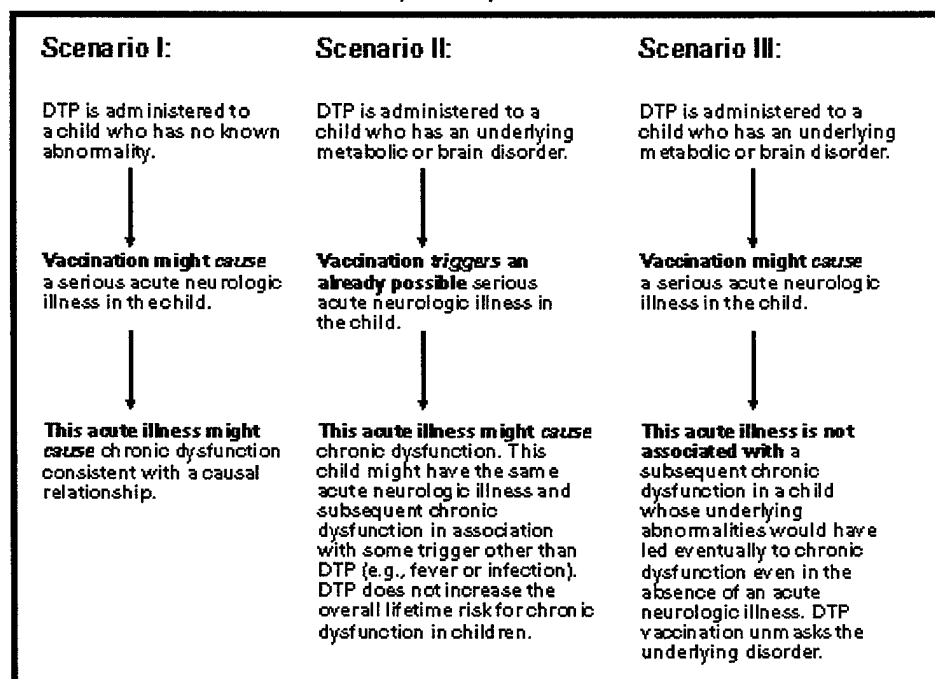
@@@ Deaths occurred primarily among persons known to be immunocompromised.

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Figure_1

FIGURE 1. Scenarios in which acute neurologic illnesses that occur after vaccination with diphtheria and tetanus toxoids and pertussis vaccine (DTP) might be associated with subsequent chronic nervous system dysfunction — Institute of Medicine, * 1994



*This information is an adaptation of information published previously by the Institute of Medicine (IOM) (6), an independent research organization chartered by the National Academy of Sciences. The National Childhood Vaccine Injury Act of 1986 mandated that IOM review scientific and other evidence (e.g., epidemiologic studies, case series, individual case reports, and testimonials) regarding the possible adverse consequences of vaccines administered to children.

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Table_4

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 4. Guidelines for spacing the administration of immune globulin preparations *
mumps, or rubella virus

Simultaneous administration		No
Immunobiologic combination	Recommended minimum interval between doses	Immunobio
		First
		Immune globu
		Vaccine
Immune globulin and vaccine	Should generally not be administered simultaneously. + If simultaneous administration of measles-mumps-rubella {MMR}, measles-rubella, and monovalent measles vaccine is unavoidable, administer at different sites and revaccinate or test for seroconversion after the recommended interval (Table 5).	
* Blood products containing large amounts of immune globulin (such as serum immune g and HBIG), intravenous immune globulin {IGIV}, whole blood, packed red cells, plas		
+ The duration of interference of immune globulin preparations with the immune respo measles-rubella, and monovalent measles vaccine is dose-related (Table 5).		

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Table_5

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 5. Suggested intervals between administration of immune globulin preparations for various indications and vaccines containing live measles virus *

Indication	Dose (including mg IgG/kg)	Time inte (mos) bef measles vaccinat
Tetanus (TIG) prophylaxis	250 units (10 mg IgG/kg) IM	3
Hepatitis A (IG) prophylaxis		
Contact prophylaxis	0.02 mL/kg (3.3 mg IgG/kg) IM	3
International travel	0.06 mL/kg (10 mg IgG/kg) IM	3
Hepatitis B prophylaxis (HBIG)	0.06 mL/kg (10 mg IgG/kg) IM	3
Rabies immune globulin (HRIG)	20 IU/kg (22 mg IgG/kg) IM	4
Varicella prophylaxis (VZIG)	125 units/10 kg (20-40 mg IgG/kg) IM (maximum 625 units)	5
Measles prophylaxis (IG)		
Standard (i.e., nonimmunocompromised contact)	0.25 mL/kg (40 mg IgG/kg) IM	5

Immunocompromised contact	0.50 mL/kg (80 mg IgG/kg) IM	6
Blood transfusion:		
Red blood cells (RBCs), washed	10 mL/kg (negligible IgG/kg) IV	0
RBCs, adenine-saline added	10 mL/kg (10 mg IgG/kg) IV	3
Packed RBCs (Hct 65%) +	10 mL/kg (60 mg IgG/kg) IV	6
Whole blood cells (Hct 35%-50%) +	10 mL/kg (80-100 mg IgG/kg) IV	6
Plasma/platelet products	10 mL/kg (160 mg IgG/kg) IV	7
Replacement therapy for immune deficiencies		
	300-400 mg/kg IV & (as IGIV)	8
Treatment of:		
Immune thrombocytopenic purpura @	400 mg/kg IV (as IGIV)	8
Immune thrombocytopenic purpura @	1,000 mg/kg IV (as IGIV)	10
Kawasaki disease	2 g/kg IV (as IGIV)	11

* This table is not intended for determining the correct indications and dosage for immune globulin preparations. Unvaccinated persons may not be fully protected against measles during the entire suggested time interval, and additional doses of immune and/or measles vaccine may be indicated after measles exposure. The concentration of antibody in a particular immune globulin preparation can vary by lot. The rate of clearance after receipt of an immune globulin preparation also can vary. The recommended time intervals are extrapolated from an estimated half-life of 30 days for passive antibody and an observed interference with the immune response to measles vaccine 5 months after a dose of 80 mg IgG/kg (42).

+ Assumes a serum IgG concentration of 16 mg/mL.

& Measles vaccination is recommended for most HIV-infected children who do not have severe immunosuppression, but it is contraindicated for patients who have congenital disorders of the immune system.

@ Formerly referred to as idiopathic thrombocytopenic purpura.

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Table_6

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 6. Adverse events * occurring within 48 hours after vaccination with diphtheria and tetanus toxoids and pertussis vaccine (DTP)

Event	Frequency of event +
Local reaction	
- Pain	1 per 2 doses
- Swelling	2 per 5 doses
- Redness	1 per 3 doses
Systemic reaction	
- Fever ≥ 100.4 F (≥ 38 C)	1 per 2 doses
- Fretfulness	1 per 2 doses
- Drowsiness	1 per 3 doses
- Anorexia	1 per 5 doses
- Vomiting	1 per 15 doses

- Persistent, inconsolable crying (i.e., for ≥ 3 hrs) 1 per 100 doses
- Fever ≥ 105 F (≥ 40.5 C) 1 per 330 doses
- Collapse (hypotonic-hyporesponsive episode) 1 per 1,750 doses
- Convulsions (with or without fever) 1 per 1,750 doses

* Adapted from Cody CL, Baraff LJ, Cherry JD, et al., 1981 (60).

+ Rate per total number of doses, regardless of dose number in DTP series.

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Table_7

Note: To print large tables and graphs users may have to change their printer settings to landscape and use a small font size.

TABLE 7. Contraindications and precautions to subsequent vaccination with diphtheria and tetanus toxoids and pertussis vaccine (DTP)

Classification/Response to DTP vaccination

Contraindications

- An immediate anaphylactic reaction.
- Encephalopathy occurring within 7 days after vaccination.

Precautions

- Fever ≥ 105 F (≥ 40.5 C) that is not attributed to another identifiable cause occurring within 48 hours after vaccination.
- Collapse or shock-like state (i.e., a hypotonic-hyporesponsive episode) occurring within 48 hours after vaccination.
- Persistent, inconsolable crying lasting ≥ 3 hours and occurring within 48 hours after vaccination.
- Convulsions with or without fever occurring within 3 days after vaccination.

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Page converted: 09/19/98

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D. Currently Applicable Classification Level: Unclassified

E. Distribution Statement A: Approved for Public Release

F. The foregoing information was compiled and provided by:
DTIC-OCA, Initials: __VM__ **Preparation Date** 10/25/01

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