Vaccine Formulation	Contraindications and Precautions
All vaccines	Contraindication
	Severe allergic reaction (e.g., anaphylaxis) after a previous vaccine dose or to a vaccine component
	Precaution
	Moderate or severe acute illness with or without fever. Defer vaccination until illness resolves.
Td	Precautions
	GBS within 6 weeks after a previous dose of TT-containing vaccine
	History of arthus-type hypersensitivity reactions after a previous dose of TD- or DT-containing vaccines (including MCV4). Defer vaccination until at least 10 years have elapsed since the last dose.
Tdap	Contraindication
	History of encephalopathy (e.g., coma or prolonged seizures) not attributable to another identifiable cause within 7 days of administration of a vaccine with pertussis components, such as DTaP or Tdap
	Precautions
	GBS within 6 weeks after a previous dose of TT-containing vaccine
	Progressive or unstable neurologic disorder, uncontrolled seizures, or progressive encephalopathy. Defer vaccination until a treatment regimen has been established and the condition has stabilized.
	History of arthus-type hypersensitivity reactions after a previous dose of TT- or DT-containing vaccines (including MCV4). Defer vaccination until at least 10 years have elapsed since the last dose.
HPV	Contraindication
	History of immediate hypersensitivity to yeast (for Gardasil)
	Precaution
	Pregnancy. If a woman is found to be pregnant after initiation of the vaccination series, the remainder of the 3-dose regimen should be delayed until after completion of the pregnancy. If a vaccine dose has been administered during pregnancy, no intervention is needed. Exposure to Gardasil during pregnancy should be reported to Merck (800-986-8999); exposure to Cervarix during pregnancy should be reported to GlaxoSmithKline (888-452-9622).
MMR	Contraindications
	History of immediate hypersensitivity reaction to gelatin ^a or neomycin
	Pregnancy
	Known severe immunodeficiency (e.g., hematologic and solid tumors; chemotherapy; congenital immunodeficiency; long-term immunosuppressive therapy; severe immunocompromise due to HIV infection)
	Precautions
	Recent receipt (within 11 months) of antibody-containing blood product
	History of thrombocytopenia or thrombocytopenic purpura
Varicella	Contraindications
	Pregnancy
	Known severe immunodeficiency
	History of immediate hypersensitivity reaction to gelatin ^a or neomycin
	Precaution
	Recent receipt (within 11 months) of antibody-containing blood product
Influenza, inactivated,	Precautions
injectable	History of severe allergic reaction (e.g., anaphylaxis) to egg protein ^b (note: not a precaution for Flublok recombinant influenza vaccine, which is approved for persons 18–49 years of age and is manufactured without the use of eggs)
	History of GBS within 6 weeks after a previous influenza vaccine dose
Influenza, live	Contraindications
attenuated nasal spray	History of severe allergic reaction (e.g., anaphylaxis) to egg protein ^b
	Age ≥50 years
	Pregnancy
	Immunosuppression, including that caused by medications or by HIV infection; known severe immunodeficiency (e.g., hematologic and solid tumors; chemotherapy; congenital immunodeficiency; long-term immunosuppressive therapy; severe immunocompromise due to HIV infection)
	Certain chronic medical conditions, such as diabetes mellitus; chronic pulmonary disease (including asthma); chronic cardiovascular disease (except hypertension); renal, hepatic, neurologic/neuromuscular, hematologic, or metabolic disorders
	Close contact with severely immunosuppressed persons who require a protected environment, such as isolation in a bone marrow transplantation unit
	Close contact with persons with lesser degrees of immunosuppression (e.g., persons receiving chemotherapy or radiation therapy who are not being cared for in a protective environment; persons with HIV infection) is <i>not</i> a contraindication or a precaution. Health care personnel in neonatal intensive care units or oncology clinics may receive live attenuated influenza vaccine.
	Precautions

History of GBS within 6 weeks of a previous influenza vaccine dose

Receipt of specific antiviral agents (i.e., amantadine, rimantadine, zanamivir, or oseltamivir) with 48 h before vaccination