Date: 09/01/2023

A Prefilled Diluent Syringe for Merck’s MMRV Family — M-M-R® II (Measles, Mumps, and Rubella Virus Vaccine Live), VARIVAX® (Varicella Virus Vaccine Live), and ProQuad® (Measles, Mumps, Rubella and Varicella Virus Vaccine Live) — is now FDA approved

Dear Valued Merck Customer,

On 08/21/2023, the FDA approved sterile diluent supplied in a prefilled syringe for vaccines in Merck’s MMRV Family. These include M-M-R® II (Measles, Mumps, and Rubella Virus Vaccine Live), VARIVAX® (Varicella Virus Vaccine Live), and ProQuad® (Measles, Mumps, Rubella and Varicella Virus Vaccine Live). When reconstituting these vaccines using the prefilled diluent syringe, the need to withdraw the sterile diluent from a vial is eliminated.

Please note that this update only impacts the sterile diluent, and there is no change to the vials containing lyophilized M-M-R® II, ProQuad, or VARIVAX (ie, packaging, NDC, etc.).

- Early in the 4th quarter of 2023, the transition from vial diluent to a prefilled diluent syringe will begin. During the transition period, you may receive either the vials or syringes of diluent with your order of M-M-R® II, ProQuad, or VARIVAX.
- The exact timing of the transition will vary based on the vaccine and the ordering channel (ie, direct from Merck, wholesaler, distributor, or CDC).
- The transition to the prefilled diluent syringes should be fully complete by the end of 2023.
- The vaccines and sterile diluent will continue to be packaged separately, allowing you to store the diluent separately from the lyophilized vaccines. Store the prefilled diluent syringes at room temperature (68°F to 77°F, 20°C to 25°C) or in the refrigerator (36°F to 46°F, 2°C to 8°C). Do not freeze the sterile diluent.
- The sterile diluent will be supplied as a carton of 10 single-dose prefilled diluent syringes (NDC: 0006-4175-88). The carton dimensions are: 4.29” x 5.236” x 1.397”.
- When shipped directly from Merck, orders of 40 doses (4 cartons) or fewer of M-M-R® II, ProQuad, and VARIVAX will be shipped together with the prefilled diluent syringes in one shipment.
- When shipped directly from Merck, orders of 50 doses (5 cartons) or more of M-M-R® II, ProQuad, and VARIVAX will be shipped in two parcels. The diluent will be shipped in an ambient temperature parcel.
- If you order vaccines from a wholesaler or distributor, please reach out to your partner directly to learn more.

With the availability of prefilled diluent syringes, Merck is proud to offer one less step for preparation of each of the vaccines in the MMRV Family (M-M-R® II, ProQuad, and VARIVAX), as compared to use of vial diluent. This change reflects our commitment to you and your patients through the evolution of the diluent presentation for these vaccines.

Indications and Usage

ProQuad is a vaccine indicated for active immunization for the prevention of measles, mumps, rubella, and varicella in children 12 months through 12 years of age.

VARIVAX is a vaccine indicated for active immunization for the prevention of varicella in individuals 12 months of age or older.

M-M-R® II is indicated for active immunization for the prevention of measles, mumps, and rubella in individuals 12 months of age or older.

Selected Safety Information

- Hypersensitivity: ProQuad, M-M-R® II, and VARIVAX are contraindicated in patients with a history of anaphylactic reaction or hypersensitivity to any component of the vaccine (including gelatin or neomycin) or to a prior dose of measles, mumps, rubella, or varicella-containing vaccine. Use caution when administering ProQuad and M-M-R® II to individuals with anaphylaxis or immediate hypersensitivity to eggs.
- ProQuad, M-M-R® II, and VARIVAX are contraindicated in certain individuals, including those with: immunodeficiency or who are immunosuppressed; an active febrile illness; untreated tuberculosis.

Selected Safety Information continues on the next page
Selected Safety Information for M-M-R*® (Measles, Mumps, and Rubella Virus Vaccine Live), VARIVAX*® (Varicella Virus Vaccine Live), and ProQuad*® (Measles, Mumps, Rubella and Varicella Virus Vaccine Live) (continued)

- Pregnancy: ProQuad, M-M-R*®, and VARIVAX are contraindicated for use in pregnant women. Do not administer ProQuad or VARIVAX to individuals who are planning to become pregnant in the next 3 months. Do not administer M-M-R*® to individuals who are planning to become pregnant in the next month.

- Febrile Seizures: Administration of ProQuad (dose 1) to children 12 to 23 months old who have not been previously vaccinated against measles, mumps, rubella, or varicella, nor had a history of the wild-type infections, is associated with higher rates of fever and febrile seizures at 5 to 12 days after vaccination when compared to children vaccinated with a first dose of both M-M-R*® and VARIVAX administered concomitantly.

- Febrile Seizures: Use caution when administering ProQuad and M-M-R*® to individuals with a history of febrile seizures.

- Family History of Immunodeficiency: Vaccination with ProQuad, M-M-R*®, and VARIVAX should be deferred in individuals with a family history of congenital or hereditary immunodeficiency until the individual’s immune status has been evaluated and the individual has been found to be immunocompetent.

- Thrombocytopenia: Transient thrombocytopenia has been reported within 4-6 weeks following vaccination with measles, mumps, and rubella vaccine. Carefully evaluate the potential risk and benefit of vaccination in children with thrombocytopenia or in those who experienced thrombocytopenia after vaccination with a previous dose of a measles, mumps, and rubella-containing vaccine.

- Varicella Transmission and Precautions: Advise vaccinees administered ProQuad or VARIVAX to avoid: close contact with high-risk individuals susceptible to varicella for up to 6 weeks following vaccination since transmission of varicella vaccine virus to susceptible contacts has been reported. Varicella vaccine virus transmission may occur between vaccine recipients and contacts susceptible to varicella including healthy individuals.

- Immune Globulins and Transfusions: Immune Globulins and other blood products should not be given concomitantly with ProQuad, M-M-R*®, or VARIVAX.

- Use of Salicylates: Avoid use of salicylates in children and adolescents administered ProQuad or VARIVAX for 6 weeks following vaccination due to the association of Reye Syndrome with salicylate therapy and wild-type varicella infection.

- Adverse Events: The following adverse events have been reported for both subcutaneous and intramuscular injections of ProQuad, M-M-R*®, and VARIVAX: fever, injection-site reactions (pain/tenderness/soreness, erythema, and swelling); and rash on the body or at the injection site. Additionally, irritability has been reported for the subcutaneous injections of ProQuad, M-M-R*®, and VARIVAX.

- ProQuad Systemic Vaccine-Related Adverse Events: Systemic vaccine-related adverse events that were reported at a significantly greater rate in recipients of subcutaneous ProQuad than in recipients of the component vaccines administered concomitantly were fever and measles-like rash.

- VARIVAX Dose-related Adverse Events: In a clinical trial involving children who received 2 doses of VARIVAX 3 months apart, the incidence of injection-site clinical complaints observed in the first 4 days following vaccination was slightly higher post-dose 2 (overall incidence 25.4%) than post-dose 1 (overall incidence 21.7%), whereas the incidence of systemic clinical complaints in the 42-day follow-up period was lower post-dose 2 (66.3%) than post-dose 1 (85.8%).

- Concomitant Vaccines With ProQuad: ProQuad may be administered concomitantly with diphtheria and tetanus toxoids and acellular pertussis vaccine adsorbed (DTaP), Haemophilus influenzae type b conjugate (meningococcal protein conjugate) and hepatitis B (recombinant) vaccine. It may also be administered concomitantly with pneumococcal 7-valent conjugate vaccine and/or hepatitis A vaccine (inactivated) at separate injection sites.

- Concomitant Vaccines With VARIVAX: VARIVAX can be administered with other live viral vaccines. If not given concurrently, at least 1 month should elapse between a dose of a live attenuated measles virus-containing vaccine and a dose of VARIVAX. In children, at least 3 months should elapse between administration of 2 doses of a live attenuated varicella virus-containing vaccine.

- Tuberculin Testing: If a tuberculin test is to be done with M-M-R*® and ProQuad, it should be administered either any time before, simultaneously with, or at least 4 to 6 weeks after vaccination. With VARIVAX, tuberculin testing may be performed before the vaccine is administered or at least 4 weeks following vaccination.

- Additional M-M-R*® Precautions: Additional adverse reactions, which have been reported without regard to causality, include febrile convulsions, arthritis, thrombocytopenia, anaphylaxis, anaphylactoid reactions, arhythmia, encephalitis and encephalopathy in their diverse clinical presentations.

Selected Safety Information continues on the next page
Additional VARIVAX Precautions: It is not known if varicella vaccine virus is excreted in human milk. A boost in antibody levels has been observed in vaccinees following exposure to wild-type varicella, which could account for the apparent long-term persistence of antibody levels in studies. The duration of protection from varicella infection after vaccination is unknown.

ProQuad/VARIVAX and Herpes Zoster: The long-term effect of VARIVAX on the incidence of herpes zoster, particularly in those vaccinees exposed to wild-type varicella, is unknown at present.

Efficacy: Vaccination with ProQuad, VARIVAX, or M-M-R® may not result in protection in 100% of vaccinees.

Dosage and Administration

ProQuad:

- Each dose of ProQuad is approximately 0.5 mL and is administered intramuscularly or subcutaneously.
- At least 1 month should elapse between a dose of a measles-containing vaccine such as M-M-R® and a dose of ProQuad. At least 3 months should elapse between a dose of varicella-containing vaccine and ProQuad.

VARIVAX:

- Each dose is approximately 0.5 mL and is administered intramuscularly or subcutaneously.
  - The first dose is administered at 12 to 15 months of age.
  - The second dose is administered at 4 to 6 years of age.
  - There should be a minimum interval of 3 months between doses.
  - 12 months to 12 years of age: If a second dose is administered, there should be a minimum interval of 3 months between doses.
  - Adolescents (≥13 years of age) and Adults: 2 doses, to be administered with a minimum interval of 4 weeks between doses.

M-M-R®:

- The dose for any age is approximately 0.5 mL administered intramuscularly or subcutaneously.
  - The recommended age for primary vaccination is 12 to 15 months and the second dose should be given at 4 to 6 years of age.

Before administering VARIVAX® (Varicella Virus Vaccine Live), M-M-R® (Measles, Mumps, and Rubella Virus Vaccine Live), or ProQuad® (Measles, Mumps, Rubella and Varicella Virus Vaccine Live), please read the accompanying Prescribing Information. The Patient Information also is available for VARIVAX, M-M-R®, and ProQuad.

For additional copies of the Prescribing Information, please call 800-672-6372, visit merckvaccines.com®, or contact your Merck representative.

Sincerely,

Gabriel E. Smith
Director, Pediatric and Hepatitis Vaccines
U.S. Marketing
Merck Sharp & Dohme LLC