



تاریخ: ۱۴۰۴/۰۹/۳۰

شماره: ۲۸/۵۳/۴۸۰۳۹۶

پیوست: ۵

ساعت: ۱۳:۵۸

ریاست محترم سازمان نظام پزشکی استان قزوین

ریاست محترم انجمن داروسازان استان قزوین

موسس و مسئول فنی محترم کلیه داروخانه های استان

مدیریت محترم کلیه شرکت های پخش استان

موضوع: اطلاع رسانی در خصوص به روز رسانی نکات ایمنی واکسن MMR

با سلام و احترام

با توجه به نامه مدیر کل محترم دفتر نظارت و پایش مصرف فرآورده های سلامت به شماره ۶۵۸/۷۰۰۱۲ مورخ ۱۴۰۴/۰۹/۲۶ در خصوص به روز رسانی نکات ایمنی واکسن MMR، ممنوعیت مصرف همزمان واکسن مذکور با درمان های سرکوب کننده ایمنی شامل کورتیکواستروئید با دوز بالا، همچنین به روز رسانی یافته ها در مصرف این واکسن در دوران بارداری (به پیوست)، جهت استحضار به حضور تقدیم می گردد.

دکتر فرزانه حامدی وفا  
معاون غذا و دارو دانشگاه



Ministry of Health and Medical Educations  
Islamic Republic of Iran

09-Dec-2025

**Subject:**

The Belgium Health Authority (FAGG) posted on their website information on the contra-indication for patients receiving immunosuppressive treatment and updated pregnancy data - Measles-mumps-rubella vaccine

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Dear Sir/Madam:

The Belgium Health Authority (FAGG) posted on their website the "VIG-news" (edition November 2025) containing information on the contra-indication for patients receiving immunosuppressive treatment and updated pregnancy data

Measles-mumps-rubella vaccine and measles-mumps-rubella-varicella vaccine: new contraindication and updated pregnancy data

After analyzing the available data, the PRAC concluded that the SmPC of Priorix must be updated to include a contraindication in cases of concurrent immunosuppressive treatment (including high doses of corticosteroids). Priorix is not contraindicated for individuals receiving local or parenteral corticosteroids in low doses (for example, for asthma prophylaxis or as replacement therapy). This contraindication was already applicable to M.M.R. VaxPro and ProQuad.

Section 4.6 of the SmPC of Priorix and ProQuad, containing pregnancy data, has also been updated. It now states, as with M.M.R. VaxPro, that in a study of more than 3,500 women who were unknowingly in the early stages of pregnancy when vaccinated with a rubella-containing vaccine, no cases of congenital rubella syndrome were reported. Post-marketing research showed that congenital rubella syndrome occurred due to a rubella vaccine strain in a pregnant woman who inadvertently received a measles, mumps, and rubella vaccine. Fetal harm has not been documented when pregnant women received measles, mumps, or varicella vaccines.

**Company position:**

Belgian Health Authority website posting, containing contraindication information for patients receiving immunosuppressive therapy and during pregnancy, is consistent with the Company Core Data Sheet (CCDS) and the EU Prescribing Information (Summary of Product Characteristics, SmPC) for the measles vaccine, live attenuated virus combined with mumps and rubella vaccine, which states that M-M-R II (also known as M-M-RvaxPro in EU and UK) and ProQuad are contraindicated for patients receiving immunosuppressive therapy and for pregnant



women.

Kindly note that the labeling reference market for MMR II Vaccine for Iran is USA.

In addition, please note that ProQuad Vaccine is not registered in Iran.

**Best Regards**

*Emad Samir Naguib*  
Electronically signed by:  
Emad Samir Naguib  
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Date: Dec 9, 2025 16:25:25  
GMT+4

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# Notification regarding MMR Vaccine\_ NSS\_368

Final Audit Report

2025-12-09

Created:	2025-12-09
By:	Seba Elbarkouky (seba.elbarkouky@msd.com)
Status:	Signed
Transaction ID:	CBJCHBCAABAA5a5CqfkW5zMYtwHPJdJX373v6ZIAymOx

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