

Janssen Sciences Ireland UC

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13th October 2021

COVID-19 Vaccine Janssen: Risk for immune thrombocytopenia (ITP) and venous thromboembolism (VTE)

Dear Healthcare Professional,

Janssen-Cilag International NV in agreement with the European Medicines Agency and the Health Products Regulatory Authority (HPRA) would like to inform you of the following:

Summary

Immune thrombocytopenia (ITP):

- Cases of ITP, some with very low platelet levels (<20,000 per μL), have been reported very rarely, usually within the first four weeks after receiving COVID-19 Vaccine Janssen. This included cases with bleeding and cases with a fatal outcome. Some of these occurred in individuals with a history of ITP.
- If an individual has a history of ITP, the risks of developing low platelet levels should be considered before vaccination, and platelet monitoring is recommended after vaccination.
- Individuals should be alert to signs and symptoms of ITP, such as spontaneous bleeding, bruising or petechiae.
- Individuals diagnosed with thrombocytopenia within 3 weeks after vaccination with COVID-19 Vaccine Janssen should be actively investigated for signs of thrombosis to assess a potential diagnosis of thrombosis with thrombocytopenia syndrome (TTS) that requires specialised clinical management.

Venous thromboembolism (VTE):

- Venous thromboembolism has been observed rarely following vaccination with COVID-19 Vaccine Janssen.
- The risk of VTE should be considered for individuals with increased risk factors for thromboembolism.
- Healthcare professionals should be alert to the signs and symptoms of VTE. Those vaccinated should be instructed to seek immediate medical attention if they develop symptoms such as shortness of breath, chest pain, leg pain, leg swelling, or persistent abdominal pain following vaccination.
- Individuals who present with thrombosis within 3 weeks of vaccination should be evaluated for thrombocytopenia to assess a potential diagnosis of thrombosis with thrombocytopenia syndrome (TTS) that requires specialised clinical management.

Background on the safety concern

COVID-19 Vaccine Janssen suspension for injection is indicated for active immunisation to prevent COVID-19 caused by SARS-CoV-2, in individuals 18 years of age and older.

Immune thrombocytopenia (ITP)

Although there was no imbalance of thrombocytopenia in clinical trials, review of post-marketing cases support ITP being an adverse drug reaction following vaccination with the COVID-19 Vaccine Janssen.

Analysis of key cases and literature suggests individuals with a medical history of ITP might be at an increased risk of decreased platelets and symptomatic ITP following vaccination with the COVID-19 Vaccine Janssen. If an individual has a history of ITP, the risks of developing low platelet levels should be considered before vaccination, and platelet monitoring is recommended after vaccination.

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Healthcare professionals should be alert to the signs and symptoms of thrombocytopenia. Those vaccinated should be instructed to seek prompt medical attention if they experience spontaneous bleeding, skin bruising (petechia) beyond the site of vaccination after a few days.

Individuals diagnosed with thrombocytopenia within 3 weeks after vaccination with COVID-19 Vaccine Janssen should be actively investigated for signs of thrombosis to assess a potential diagnosis of thrombosis with thrombocytopenia syndrome (TTS) that requires specialised clinical management.

Venous thromboembolism (VTE)

Venous thromboembolism, distinct from thrombosis with thrombocytopenia syndrome (TTS), has been observed rarely following vaccination with COVID-19 Vaccine Janssen. This should be considered for individuals at increased risk for venous thromboembolism.

During the double-blind period (median follow-up 123 days) of an ongoing phase 3 study (COV3001), venous thromboembolic events were observed in 26/21,894 (0.1%) of individuals who received COVID-19 Vaccine Janssen and 9/21,882 (0.04%) of individuals who received placebo. Of these, venous thromboembolic events were observed within 28 days in 8 individuals who received COVID-19 Vaccine Janssen and in 4 individuals who received placebo. Deep vein thrombosis and pulmonary embolism were mostly observed (21 individuals who received COVID-19 Vaccine Janssen and 8 individuals who received placebo during the entire double-blind phase). The majority of events were reported in individuals with at least one predisposing risk factor for venous thromboembolism.

In another ongoing phase 3 study (COV3009, 15,708 individuals receiving the vaccine and 15,592 placebo), there was no increase in venous thromboembolic events among individuals who received the COVID-19 Vaccine Janssen (median follow-up time 70 days).

Healthcare professionals should be alert to the signs and symptoms of VTE. Those vaccinated should be instructed to seek immediate medical attention if they develop symptoms such as shortness of breath, chest pain, leg pain, leg swelling, or persistent abdominal pain following vaccination. Individuals who present with thrombosis within 3 weeks of vaccination should be evaluated for thrombocytopenia to assess a potential diagnosis of thrombosis with thrombocytopenia syndrome (TTS) that requires specialised clinical management.

Call for reporting

Healthcare professionals should report any suspected adverse reactions associated with the use of COVID-19 Vaccine Janssen in accordance with the national spontaneous reporting system on the HPRAs website: www.hpra.ie.

▼ This product is subject to additional monitoring. This will allow quick identification of new safety information. Healthcare professionals are asked to report any suspected adverse reactions.

Company contact point

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- For further information, please visit: www.covid19vaccinejanssen.com
- COVID-19 vaccine dedicated contact centre: **+353 212 35 68 06**
- Email: JGCC_emea@its.jnj.com

Yours faithfully,

A handwritten signature in black ink, which appears to read "Brid Seoighe". The signature is fluid and cursive.

Dr Bríd Seoighe

Head of Medical Affairs

Janssen Sciences Ireland UC