Why is the J&J vaccine on pause



On April 13, 2021, the FDA and CDC called for a pause on the use of the Johnson & Johnson (J&J) COVID-19 vaccine after 6 recipients in the U.S. developed a **rare disorder involving blood clots** within about 2 weeks of vaccination.

Waiting for a CDC review

- This pause is being recommended out of an abundance of caution, as these adverse events appear to be extremely rare.
- More than 6.8 million people in the U.S. have received J&J vaccines and 6 recipients are known to have developed a type of blood clot called cerebral venous sinus thrombosis (CVST) in combination with low levels of blood platelets (thrombocytopenia).
- All 6 cases occurred among women between the ages of 18 and 48, and symptoms occurred 6 to 13 days after vaccination.
- CDC will review these cases and assess their potential significance.

What about vaccines in Virginia?

- Virginia will stop using the J&J vaccine while safety issues are being examined.
- Pfizer and Moderna will be used as a substitute and appointments rescheduled as needed.
- This will not impact Virginia's move to Phase 2
 by April 18; however, it could impact the availability
 of appointments going forward.
 - People who have received the J&J vaccine and develop severe headache, abdominal pain, leg pain, or shortness of breath within
 3 weeks should contact their health care provider or call 911 if it's a medical emergency.
 - People who received the J&J vaccine more than a month ago are at very low risk.

Common side effects after any vaccine include lightheadedness, nausea, or fainting. Individuals should be observed for a minimum of 15 minutes following a COVID -19 vaccination.



Sign up for **v-safe**, a **smartphone-based tool** that provides health check-ins after a COVID-19 vaccine. Through v-safe, you can tell CDC about any **side effects**.

