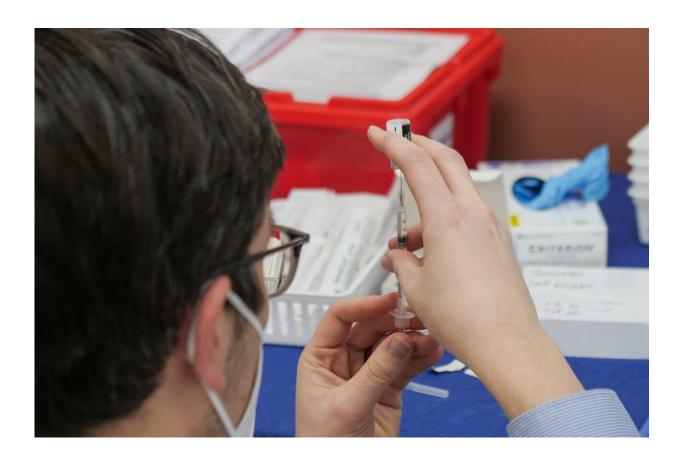


Clot questions over AstraZeneca and J&J vaccine

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The Johnson & Johnson and AstraZeneca vaccines have come under suspicion of causing very rare but serious blood clots in a handful of cases among the millions vaccinated in the drive to bring the pandemic



under control.

Here is what we know about the two vaccines, which are based on the same technology.

What happened?

The European Medicines Agency (EMA) said on Tuesday that <u>blood</u> <u>clots</u> should be listed as a "very rare" side effect of the Johnson & Johnson <u>coronavirus vaccine</u>.

As of April 13, 2021, eight cases of <u>blood</u> clots were reported in the United States—all in patients under 60, mostly women—out of seven million people vaccinated.

Earlier this month, the EMA made the same recommendation for the vaccine made by Anglo-Swedish pharmaceutical company AstraZeneca.

For both vaccines, however, the EU medicines regulator stressed that overall benefits in preventing COVID-19 clearly outweighed the risks.

The clots occur in veins in the brain—called cerebral venous sinus thrombosis (CVST)—as well as the abdomen and arteries.

They are uncommon both for where they appear in the body and because they occur together with low levels of blood platelets, which help the blood to <u>clot</u>.

Paradoxically, this can mean the patient has both haemorrhages as well as blood clots.

There have been 287 cases of these atypical thromboses worldwide following AstraZeneca jabs, including 142 in the European Economic



Area (the EU, Iceland, Norway and Liechtenstein), the EMA told AFP on Tuesday.

As with the J&J jab, most of these cases were in women under 60 and occurred within two weeks of vaccination.

US regulators have suspended the J&J vaccine and are set to announce a decision on Friday.

Suspected causes

Both of these vaccines—as well as the Russian Sputnik V jab—are based on a technique known as "viral vector".

The vaccines use a common-cold causing adenovirus, modified so it can't replicate, as a "vector" to shuttle genetic instructions into human cells, telling them to create a protein of the coronavirus.

This trains the <u>immune system</u> to be ready for the live coronavirus.

AstraZeneca opted for a chimpanzee adenovirus, J&J for a human adenovirus.

"Whilst a causal link between certain COVID-19 vaccinations, platelet abnormalities and blood clots has not, so far, been confirmed, the index of suspicion is rising that these rare cases may be triggered by the adenovirus component of the AstraZeneca and J&J vaccines," said Eleanor Riley, an immunology professor at Edinburgh University.

What mechanism?

Although nothing has been proven, Peter Marks of the US Food and



Drug Administration has said the "probable cause" of the clots could be an unusual reaction to the vaccines.

The FDA has also said that while in normal circumstances an anticoagulant drug called heparin is used to treat these blood clots, in this case administering heparin may be dangerous.

That might be because the disorder resembles a rare reaction to heparin itself, in which the body forms antibodies in response to the thinner, and in turn triggers platelets to form dangerous clots.

A team of German and Austrian scientists made the connection between this previously-known <u>immune response</u> to heparin and the AstraZeneca vaccine in March.

And the EMA said this was "one plausible explanation", calling for more research.

Balancing risk

The European agency has said no specific risk factors for this type of reaction have been confirmed for the AstraZeneca vaccine, but different countries have opted to restrict access to the jab by age.

The reasoning is that the older people get, the more they risk developing a serious form of COVID, and the more it is in their interest to be vaccinated, despite potential side effects.

For example, Britain has said it will not administer AstraZeneca to the under 30s, while France is not using it on under 55s.

In the US, health authorities said the suspension of the J&J vaccine was "out of an abundance of caution".



People who have had the shot within the past three weeks were asked to report to their doctors if they experience severe headaches, <u>abdominal pain</u>, leg pain, or shortness of breath.

Ian Douglas, professor of pharmacoepidemiology at the London School of Hygiene & Tropical Medicine, said it was important to underscore how rare these events appear to be.

"To put this into perspective, it's similar to the chance of being struck by lightning in any given year in the UK," he told the Science Media Centre in response to the US announcement.

"On the other hand, the risks from COVID-19 are substantial."

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