

**Question for written answer E-003388/2025
to the Commission**
Rule 144
Gerald Hauser (PfE)

Subject: US finally rescinds authorisation for insufficiently tested COVID-19 vaccines

In its answer to question E-002126/2025¹, the Commission recently confirmed that ‘a comprehensive data dossier is not yet available’ for the COVID-19 vaccines. In the 20 November 2020 purchase agreement concluded with Pfizer, the Commission and all EU Member States signed to explicitly say that they ‘acknowledge that the long-term effects and efficacy of the Vaccine are not currently known and that there may be adverse effects of the Vaccine that are not currently known.’ As a result, every vaccinated person in the EU has been relegated to the role of experimental guinea pig. On 27 August 2025, US Secretary of Health Robert Kennedy Jr said that the emergency use authorisation for COVID-19 vaccines had been rescinded. Meanwhile, in the EU, these vaccinations are still offered and recommended unconditionally to the entire population, including pregnant women, babies and children.

1. Will the Commission now also rescind its COVID-19 vaccine authorisations for the population as a whole?
2. Who in the Commission is now responsible for vaccine damage, especially among healthy people with no pre-existing conditions?
3. What does the Commission have to say about the continuing lack of substantial data on the safety and effectiveness of COVID-19 vaccines, as the FDA has now officially confirmed?

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¹ https://www.europarl.europa.eu/doceo/document/E-10-2025-002126-ASW_EN.html