Question for written answer
to the Commission
Rule 138
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Subject: Removal of Covid-19 vaccines side effects from EMA databases

A recent publication suggests a significant removal of side effects in databases related to Covid-19 vaccines side effects.[1] This revelation stems from a comparative analysis of EMA, VAERS, and ANSM data covering the period from 2021 to 2023 and the data available on January 1, 2024.

The publication refers to the removal of various side effects, including 4,241 reported deaths, 14,969 cases of myocarditis, 11,424 cases of pericarditis, 7,079 cases of thrombosis, 7,295 cases of embolism, 22,107 cases of amenorrhea and dysmenorrhea, 2,827 cases of deafness, and 2,282 cases of blindness.

Can the Commission verify the accuracy of this publication?

Can the Commission specifically indicate whether the figures of removed side effects are accurate? If so, what are the reasons behind the removal of these side effects?

Regardless of the publication's accuracy, it is crucial to acknowledge the valid concerns surrounding the safety and monitoring of Covid-19 vaccines sides effects. In light of these concerns, can the Commission provide a comprehensive overview of any (potential) removal and adjustment (e.g. in the form of downplaying the nature and severity) of side effects from databases related to Covid-19 vaccines side effects?

[1] https://www.covid-factuel.fr/2024/01/20/quand-lema-fait-disparaitre-les-effets-secondaires-des-vaccins/