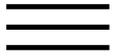


2009 monovalent vaccines reported to the Vaccine Adverse Event Reporting System, United States, October 1, 2009.

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## Vaccine

Volume 28, Issue 45, 21 October 2010, Pages 7248-7255

Adverse events following influenza A (H1N1) 2009 monovalent vaccines reported to the Vaccine Adverse Event Reporting System, United States, October 1, 2009–January 31, 2010

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<https://doi.org/10.1016/j.vaccine.2010.09.021>

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### Abstract

The United States (US) influenza A (H1N1) 2009 monovalent (2009-H1N1) vaccination program began in October 2009. Reports to the vaccine adverse event reporting system (VAERS), a US spontaneous reporting system, were reviewed to identify potential rare events or unusual adverse event (AE) patterns after 2009-H1N1 vaccination. The adverse event profile after 2009-H1N1 vaccine in VAERS ( $\approx 1/410,000$  reports) was consistent with that of seasonal influenza vaccines, although the reporting rate was higher after 2009-H1N1 than seasonal influenza vaccines, this may be, at least in part, a reflection of stimulated reporting. Death, Guillain-Barré syndrome and anaphylaxis

reports after 2009-H1N1 vaccination were rare (each <2 per million doses administered).



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## Keywords

H1N1 vaccines; Vaccine safety; Post-marketing surveillance

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<sup>†</sup> *Disclaimer:* The findings and conclusions in this article are those of the authors and do not necessarily represent the views CDC or FDA.

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