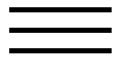


Adverse events following influenza A (H1N1)
2009 monovalent vaccines reported to the
Vaccine Adverse Event Reporting System,
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Vaccine

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 †

Claudia Vellozzi ^a ... Frank DeStefano ^a

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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



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Keywords

H1N1 vaccines; Vaccine safety; Post-marketing surveillance

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