

Should this influenza vaccine reaction be reported as an adverse event?



Injection site reactions involving pain, redness and swelling at or near the injection site occur in 10 to 64 % of influenza vaccine recipients.¹ They are inflammatory responses to the foreign material in the vaccine and the tissue irritation caused by the injection process.² These reactions occur within 48 hours of vaccination,² are usually mild and resolve spontaneously in two to three days.¹

Symptoms can be managed with cold compresses, acetaminophen or ibuprofen if required for discomfort and reassurance that this is a common occurrence with no serious consequences.^{2,3} Pressure should not be applied on the injection site.²

An injection site reaction is not a contraindication to future influenza vaccinations.² Note, it is not

recommended to take acetaminophen or ibuprofen prophylactically as there is no evidence that this is of benefit³ and some evidence it might interfere the therapeutic immune response to the vaccine.⁴

Patients should be referred to their doctors or nurse practitioners and reactions reported as adverse events following immunization (AEFI) in the following situations:

- Redness, or swelling, or pain extends past the nearest joint; AND/OR
- Redness, or swelling, or pain persists for 10 days or more²

Should the reaction pictured above be reported?

Only if the symptoms last longer than 10 days. It would be important to follow-up with the child's caregiver to find out if the symptoms have resolved.

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References:

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4. Kelso J. Allergic reactions to vaccines. In: UpToDate, Basow, DS (Ed), Waltham MA, 2017. Cited 28 Nov 2017. Available from www.uptodate.com. Subscription and login required