Update on Egg Allergy and Influenza Vaccine (Nov 2011)

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Background

Influenza vaccines are grown in eggs and there has been concern that residual egg protein (ovalbumin) could cause allergic reactions in egg-allergic recipients. However, all studies to date have suggested that this risk is very low,\(^1\)-\(^7\) and the vaccine provides substantial protection against the morbidity and mortality associated with influenza disease.\(^8\) This update reflects changes to recommendations for administration of trivalent influenza vaccine (TIV) to egg-allergic patients based on several studies, most published in the last 2 years.\(^1\)-\(^7\) New guidelines have also been issued by the Centers for Disease Control and Prevention’s (CDC) Advisory Committee on Immunization Practices (ACIP)\(^9\) (see Figure) and the American Academy of Pediatrics’ (AAP) Committee on Infectious Diseases (“Red Book” committee).\(^10\)

Conclusions:

1. The risks of vaccinating egg-allergic patients with influenza vaccine are outweighed by the risks of not vaccinating.

   In 7 published studies, over 1600 patients with egg allergy have been vaccinated without any serious reactions.\(^1\)-\(^7\) 0-6.3% of vaccinations have involved reactions confined to the skin, e.g. hives. 0-4.8% of vaccinations have involved mild respiratory or gastrointestinal symptoms. No reactions have involved symptoms of hypotension. None of these reactions required treatment with epinephrine. One study using an adjuvanted H1N1 vaccine included an additional 3,640 patients with reported, but not proven, egg allergy given influenza vaccine; 1.2% had skin reactions, 0.7% had respiratory reactions including 2 that were given epinephrine, although the authors conclude that these were not anaphylactic reactions.\(^3\) In studies that included non egg-allergic controls, similar rates of reactions are reported.\(^1,3,4,6\)

   On average 294,128 people, including 21,156 children under 5 years of age, are hospitalized each year in the United States due to influenza,\(^11\) and an average of 23,607 die.\(^12\) Much of this morbidity and mortality can be prevented by vaccination\(^13\) and it is likely that some of these preventable hospitalizations and deaths include patients not vaccinated because they are egg-allergic.

2. Patients who believe they are allergic to eggs should be evaluated by an allergist but, for those with a history of hives only after egg ingestion, this need not delay
influenza vaccination.

Persons with a history of suspected egg allergy should be evaluated by an allergist \(^ {14}\) where history and prick skin testing or specific in vitro IgE antibody testing for egg can determine whether or not they are egg allergic. Even patients with confirmed egg allergy can receive influenza vaccine. \(^ {9,10}\) For those with a history of hives only after egg ingestion, the vaccine can be administered in a primary care setting with appropriate precautions as below, and such immunization can proceed even prior to additional evaluation of the egg allergy. \(^ {9,10}\) For those with more severe reactions after egg ingestion, egg allergy evaluation and influenza vaccination can both be accomplished in the allergist's office. \(^ {9,10}\)

3. In all studies to date, even patients with a history of anaphylaxis to egg ingestion have been safely administered the influenza vaccine.

The studies on influenza vaccine in egg-allergic patients have included 185 patients with histories of anaphylaxis to egg ingestion, and all have tolerated the vaccine without serious reactions. \(^ {1,3,4,6,7}\)

4. Skin testing with the influenza vaccine itself in egg-allergic individuals does not identify patients who are at increased risk of reacting to the vaccine and is not recommended.

When influenza vaccine skin testing was done in egg-allergic patients, subjects with positive skin test results had no reactions, or no greater rate of reactions, than those with negative skin test results. \(^ {1,2,4,6,7}\) The rate (low) of reactions (minor) is the same whether skin testing is included in the protocol or not. \(^ {2}\) Skin testing has greater utility when evaluating patients with allergic reactions to influenza vaccine itself. \(^ {15}\)

5. Influenza vaccine can be administered as a single dose to egg-allergic patients.

Some studies of influenza vaccine in egg-allergic patients have given 10% of the dose and, if no reaction in 30 minutes, given the remaining 90%. The vast majority of patients ultimately tolerate the entire dose, \(^ {1-7}\) and studies giving the vaccine as a single dose also report no serious reactions. \(^ {3,4,6,7}\) The CDC and AAP have stated that persons who have experienced hives only following ingestion of egg should receive influenza vaccine and that vaccine skin testing and dividing the dose are not necessary or recommended. \(^ {9,10}\) They recommend that patients with a history of more severe reactions be referred to an allergist before receipt of vaccine. \(^ {9,10}\) Studies support a single dose even in these patients; of the 185 patients reported with a history of anaphylaxis or severe reaction to egg ingestion who have been vaccinated with TIV, 119 receive a divided dose and 66 received a single dose, all without
developing serious reactions.\textsuperscript{1, 3, 4, 6, 7}

6. Egg-allergic individuals should receive influenza vaccine in a setting where anaphylaxis can be recognized and treated and should be observed for 30 minutes after vaccination.

Given the possibility of allergic reactions to any vaccine, and the theoretically increased risk for influenza vaccine in egg-allergic patients, providers who administer vaccinations should have proper resuscitative equipment available in the office to manage anaphylaxis.\textsuperscript{16} Those with a history of hives only after egg ingestion can receive the vaccine in a primary care provider's office, while those with a history of more severe reactions should receive the vaccine in an allergist’s office, where additional expertise is available should it be required.\textsuperscript{9, 10} All egg-allergic patients receiving influenza vaccine should be observed for 30 minutes after vaccination, similar to the waiting period after allergen immunotherapy.\textsuperscript{9, 10} Egg allergic individuals should not receive their influenza vaccine from a pharmacy or other non-medical office setting.

7. All influenza vaccines available in the US contain low amounts of ovalbumin.

Vaccines used in studies on administration of influenza vaccine to egg-allergic recipients have contained as much as 0.7 mcg of ovalbumin per 0.5 mL dose without serious reactions.\textsuperscript{7} It is not known if there is an amount of ovalbumin per dose that would be associated with a higher rate of reactions or more severe reactions.

All of the manufacturers of injectable influenza vaccines report the maximum amount of ovalbumin per 0.5 mL dose in their package inserts or will provide the information on request (Table). All of the claimed amounts are below 1 mcg per 0.5 mL dose. When the actual amount of ovalbumin in the vaccines has been measured in independent laboratories, the levels have been much lower than the claimed amounts.\textsuperscript{17-19}

8. All studies to date have evaluated injectable trivalent inactivated vaccine (TIV) and thus TIV, rather than intranasal live attenuated influenza vaccine (LAIV) should be used for egg-allergic patients.\textsuperscript{9, 10}

Summary

The risk of an allergic reaction to influenza vaccine in patients with egg allergy is very low, likely due to the very low amount of ovalbumin in the vaccines. Any such theoretical risk is far outweighed by the very real risk of such patients remaining unvaccinated. Thus all patients with egg allergy of any severity, including anaphylaxis, should receive influenza...
vaccine. Skin testing with the vaccine and dividing the dose are not necessary. The injectable vaccine should be administered in a medical setting where anaphylaxis can be recognized and treated should it occur. For those with a history of hives only after egg ingestion, the vaccine can be administered in the primary care provider's office. For those with more serious reactions to egg ingestion, the vaccine should be administered in an allergist's office.
REFERENCES


* Persons with egg allergy might tolerate egg in baked products (e.g., bread or cake). Tolerance to egg-containing foods does not exclude the possibility of egg allergy.
TABLE. Ovalbumin content of injectable trivalent Influenza vaccines (TIV) approved for the 2011-12 season

<table>
<thead>
<tr>
<th>Brand name</th>
<th>Manufacturer</th>
<th>Approved ages</th>
<th>Ovalbumin content (mcg per 0.5 mL dose*)†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Afluria</td>
<td>CSL Biotherapies (Merck)</td>
<td>≥ 9 years</td>
<td>&lt; 1</td>
</tr>
<tr>
<td>Fluarix</td>
<td>GlaxoSmithKline</td>
<td>≥ 3 years</td>
<td>&lt; 0.05</td>
</tr>
<tr>
<td>FluLaval</td>
<td>ID Biomedical Corporation of Quebec (GlaxoSmithKline)</td>
<td>≥ 18 years</td>
<td>&lt; 1</td>
</tr>
<tr>
<td>Fluvirin</td>
<td>Novartis</td>
<td>≥ 4 years</td>
<td>&lt; 1</td>
</tr>
<tr>
<td>Fluzone</td>
<td>Sanofi Pasteur</td>
<td>≥ 6 months</td>
<td>~0.1</td>
</tr>
<tr>
<td>Fluzone High-Dose</td>
<td>Sanofi Pasteur</td>
<td>≥ 65 years</td>
<td>~0.1</td>
</tr>
</tbody>
</table>

* Dose 0.25 mL 6-35 months, 0.5 mL ≥ 3 years

† Information in package inserts except Fluzone and Fluzone High-Dose from Sanofi Pasteur by telephone (1-800-822-2463) or e-mail (MIS.Emails@sanofipasteur.com)