Vaccination in Adults with Chronic Inflammatory Conditions
Overview

Patients with chronic inflammatory diseases are at increased risk for infections. Immunizations are often underutilized in these patients. The objective of this activity is to promote awareness of the importance and outline best practices for pneumococcal and influenza vaccinations in these patient populations.

Target Audience

Vaccination in Adults with Chronic Inflammatory Conditions is intended for allergists/immunologists, rheumatologists, fellows, students and other medical professionals who are interested in learning about vaccination options in patients with autoimmune disease, autoimmune disease on immunosuppressive therapy, primary immunodeficiency and chronic respiratory conditions.

Learning Objectives

The information presented in this educational activity should:

- Promote awareness regarding the importance of vaccinations in patients with chronic inflammatory conditions: autoimmune disease, autoimmune disease on immunosuppressive therapy, primary immunodeficiency and chronic respiratory conditions.
- Allow the participant to be able to apply the current guidelines for pneumococcal and influenza vaccinations and to recognize clinical settings in which these vaccines are indicated or contraindicated.

Accreditation Statement

This activity has been planned and implemented in accordance with accreditation requirements and policies of the Accreditation Council for Continuing Medical Education (ACCME) through the joint providership of the American Academy of Allergy, Asthma and Immunology (AAAAI) and UNC Thurston Arthritis Research Center. The AAAAI is accredited by the ACCME to provide continuing medical education for physicians.

Designation Statement

The American Academy of Allergy, Asthma & Immunology designates this enduring material for a maximum of 1.00 AMA PRA Category 1 Credit™. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

Release Date

Vaccination in Adult Autoimmune and Immunodeficiency Conditions was originally released on October 15, 2016.

Expiration Date

This activity will expire on October 14, 2018.

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The information disclosed by the authors was reviewed in accordance with the AAAAI Disclosure Policy. No relevant financial relationships were disclosed.

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Biographical Sketches

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Dr. Saira Sheikh is board certified in Internal Medicine, Rheumatology and Allergy & Immunology with clinical interests in systemic lupus erythematosus (SLE), allergic rhinitis, urticaria and food allergy. She is director of the UNC Rheumatology Lupus Clinic and the Clinical Trials Program at the UNC Thurston Arthritis Research Center. She is Principal Investigator on several clinical trials for SLE and other rheumatologic conditions, studying new therapies for these diseases. She enjoys teaching and is actively involved in the education of medical students, residents and sub-specialty fellows. Dr. Sheikh is a member of the American Academy of Allergy, Asthma & Immunology, the American College of Allergy, Asthma & Immunology and the American College of Rheumatology.

Mildred Kwan, MD, PhD
Assistant Professor of Medicine
Dr. Mildred Kwan is board certified in Internal Medicine and Allergy & Immunology, with clinical interests in chronic urticaria, allergic and non-allergic rhinitis, and asthma. Dr. Kwan’s research interests include how immune dysregulation in disease affects the development of atopic and autoimmune conditions, the interaction of the human microbiome on immunological disease, and the effects of biologics and small molecule inhibitors on immune function. Additionally, she has a strong interest in teaching and has developed a teaching curriculum for the Allergy & Immunology fellows at both the University of Pennsylvania and the University of North Carolina. Dr. Kwan is a member of the American Academy of Allergy, Asthma & Immunology.
When is the best time to immunize patients with autoimmune disease?

- The best time to immunize patients with autoimmune disease is prior to initiating immunosuppressive therapy.

Can inactivated or killed vaccines be administered once therapy with immunosuppressive agents has been initiated?

- Yes. Inactivated or killed vaccines can be administered during the use of immunosuppressive agents such as disease-modifying antirheumatic drugs (DMARDs), long-term systemic corticosteroids, tumor necrosis factor (TNF) blocking agents and other biological agents.

Are vaccines immunogenic in patients with autoimmune disease?

- Yes. Vaccines remain immunogenic in immunosuppressed patients with autoimmune disease. Several studies have shown that while responses may be diminished compared to disease-free people, patients are able to mount protective antibody titers.

Can vaccines increase disease activity in autoimmune rheumatic diseases?

- Overall, the available evidence suggests that vaccines do not increase activity in autoimmune rheumatic diseases such as systemic lupus erythematosus or rheumatoid arthritis. Although disease flares have been reported after vaccination in controlled and uncontrolled studies, the frequency of flares was not increased in vaccinated patients compared with unvaccinated patient controls. These flares were therefore believed to represent the natural course of the disease.

Vaccination is recommended in patients with stable or low disease activity.

Is there any value in immunizing patients with primary immunodeficiency with pneumococcal or influenza vaccines?

- While the true value of immunizing patients with a primary immunodeficiency remains unknown, the standard practice is to follow usual vaccination guidelines for these patients. Injected seasonal influenza, and pneumococcal vaccines should be administered per Centers for Disease Control and Prevention (CDC) guidelines.

Why should we vaccinate against *S. pneumoniae*?

- *S. pneumoniae* can cause pneumonia, bacteremia and meningitis. There are 400,000 hospitalizations for pneumococcal pneumonia annually and approximately 12,000 cases of pneumococcal bacteremia with a fatality rate of 20%. Invasive pneumococcal disease causes significant morbidity and mortality particularly in the elderly and in persons with immunocompromising conditions (e.g. primary immunodeficiency, iatrogenic immunosuppression and chronic inflammatory conditions).
Why should we vaccinate against influenza virus?
Influenza virus causes significant morbidity and mortality with average annual deaths caused by influenza measured at 23,607. In the 2009-2010 H1N1 pandemic there were 60 million Americans infected, 270,000 hospitalized and 12,500 deaths.

Persons who are considered at high risk for complications with influenza infection are those with immunodeficiency, on immunosuppressive therapy, chronic respiratory conditions such as asthma and pregnant women.

Influenza vaccines that are recommended for vaccination are inactivated influenza vaccines (IIVs) which are given by either intramuscular or intradermal routes.

What are the recent recommendations for the administration of influenza vaccine?
All persons ≥ 6 months should be vaccinated annually.

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Note: Those who received PPSV23 before age 65 years should receive another dose of the vaccine at age 65 years, or later if at least five years have elapsed since their previous PPSV23 dose.
PCV13 now because one year has passed since patient received PPSV23

Second dose of PPSV23 at age 27 (≥ 5 years since previous PPSV23 and ≥ 8 weeks since PCV13 dose)

Third dose of PPSV23 at age 65. No further PCV13 doses need to be administered, as the patient received one lifetime dose of PCV13

Annual influenza inactivated vaccine

Inactivated influenza vaccine should be used and administered by clinicians experienced in recognizing and treating anaphylaxis.

Patient should be observed for 30 minutes in office following the injection.

Both a single dose or a two step graded challenge are appropriate (10% of the initial dose followed by observation for 30 minutes then the remaining 90% of the dose and a second 30-60 minute observation period).

**Case Scenarios for Review:**

**Case 1:** A 32 year old woman has asthma and has not previously received any pneumococcal vaccines. What pneumococcal and influenza vaccines would you consider?

- PPSV23 vaccine now, at age 32
- PCV13 at age 65 followed by a dose of PPSV23 with a minimum of eight weeks from PCV13 administration
- Annual influenza inactivated vaccine

Asthma is not an indication for adult administration of PCV13 < 65 years of age unless the patient has received long-term systemic corticosteroids.

**Case 2:** A 24 year old woman with common variable immune deficiency received one dose of PPSV23 two years ago. What pneumococcal and influenza vaccines could you consider?

- PCV13 now because one year has passed since patient received PPSV23
- Second dose of PPSV23 at age 27 (≥ 5 years since previous PPSV23 and ≥ 8 weeks since PCV13 dose)
- Third dose of PPSV23 at age 65
- Annual influenza inactivated vaccine

**Case 3:** A 60 year old man with psoriatic arthritis who has not previously received any pneumococcal vaccines and is currently on systemic corticosteroids and will be starting anti-IL-17 therapy. What pneumococcal and influenza vaccines would you consider?

- PCV13 now, at age 60
- PPSV23 at least 8 weeks after the dose of PCV13
- Second dose of PPSV23 at age 65. No further PCV13 doses need to be administered, as the patient received one lifetime dose of PCV13
- Annual influenza inactivated vaccine

**Case 4:** 24 year old male with ankylosing spondylitis is currently on anti-tumor necrosis factor therapy and asks about getting a flu shot. He reports an “egg allergy” and had hives with egg as a child. Should he receive the influenza vaccine?

- Inactivated influenza vaccine should be used and administered by clinicians experienced in recognizing and treating anaphylaxis.
- Patient should be observed for 30 minutes in office following the injection.
- Both a single dose or a two step graded challenge are appropriate (10% of the initial dose followed by observation for 30 minutes then the remaining 90% of the dose and a second 30-60 minute observation period).
For reporting adverse events related to vaccinations:
http://vaers.hhs.gov/index

References:
7. CDC Pinkbook (http://www.cdc.gov/vaccines/pubs/pinkbook/index.html),
8. CDC Pneumococcal Vaccination website (http://www.cdc.gov/vaccines/vpd-vac/pneumo/default.htm)
9. CDC Influenza vaccination website (http://www.cdc.gov/flu/professionals/vaccination/index.htm)

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To receive CME credit for this activity, you must read the entire monograph, register at the American Academy of Allergy, Asthma and Immunology Society (AAAAI) website, complete and pass the post-test and complete the evaluation.

To register on the AAAAI website, please follow these directions.
■ Navigate to https://education.aaaai.org/UNCVAC
■ Click ‘Log in’ at top right corner of page
■ For returning users:
  ■ Click ‘I am a member or I have claimed credit through the AAAAI’
  ■ Enter your AAAAI username and password
  ■ If you have forgotten your login information, click ‘Forgot username or password?’
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  ■ You should be redirected to the Education site -- if not, after logging in navigate to https://education.aaaai.org/UNCVAC
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■ Once you are logged in, click the red begin button in the summary box on the right side of the page
■ Read the Instructions page and click the red continue button
■ Complete all of the steps, in order, to complete the activity and claim your credits

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"The most significant barrier to vaccination coverage identified by the CDC is lack of knowledge about vaccines among adult patients and providers ..."

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