COVID-19 Vaccine administration guidance

COVID-19 vaccine for the general population

The Centers for Disease Control and Prevention’s (CDC’s) Advisory Committee on Immunization Practices (ACIP) recommended the mRNA based COVID vaccine for people ages 16 and older. COVID-19 vaccine facts.

COVID-19 vaccine for patients on immunomodulatory agents

The CDC advises that immunocompromised individuals may receive COVID-19 vaccination if they have no contraindications* to vaccination. Currently, there are no data to establish the safety and efficacy of mRNA based COVID vaccines in patients taking immunosuppressive agents, including the systemic drugs and biologic agents used in dermatology. “The CDC does not consider COVID-19 vaccine to be contraindicated in persons taking biologic or traditional immunosuppressive medications, hedgehog pathway inhibitors, and PD1 inhibitors.” Therefore CDC suggests that patients should be counseled about the unknown vaccine safety profile and effectiveness in immunocompromised populations, as well as the potential for reduced immune responses and the need to continue to follow all CDC current guidance to protect themselves against COVID-19.

To access CDC interim considerations to the use of mRNA COVID vaccine review the link Interim clinical consideration to use mRNA COVID vaccine

CDC also recommends not to do antibody testing to assess for immunity to COVID-19 following mRNA COVID-19 vaccination. Shared decision making should be made between the patients and clinicians to guide discussions about the use of oral-systemic and biologic agents during the pandemic. To review the Academy’s guidance on the use of immunosuppressive agents during the pandemic review Guidance on the use of immunosuppressive agents.

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1 Based on personal communication with Sarah Schillie from the CDC Clinical Inquiries Team 01/15/2021
**Contraindications**

<table>
<thead>
<tr>
<th>CONDITIONS</th>
<th>CONTRAINDICATION TO VACCINATION</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Immunocompromising conditions</td>
<td>• None</td>
</tr>
<tr>
<td>• Pregnancy</td>
<td>• N/A</td>
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<tr>
<td>• Lactation</td>
<td>• N/A</td>
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</tbody>
</table>

**CONTRAINDICATIONS**

<table>
<thead>
<tr>
<th>CONDITIONS</th>
<th>PRECAUTION TO VACCINATION</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Moderate/severe acute illness</td>
<td>ACTIONS</td>
</tr>
<tr>
<td></td>
<td>• Risk assessment</td>
</tr>
<tr>
<td></td>
<td>• Potential deferral of vaccination</td>
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<tr>
<td></td>
<td>• 15-minute observation period if vaccinated</td>
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</tbody>
</table>

**ALLERGIES**

History of allergies that are unrelated to components of an mRNA COVID-19 vaccine, other vaccines, injectable therapies, or polysorbate, such as:

- Allergy to oral medications (including the oral equivalent of an injectable medication)
- History of food, pet, insect, venom, environmental, latex, etc., allergies
- Family history of allergies

**ALLERGIES**

History of any immediate allergic reaction* to vaccines or injectable therapies (except those related to component of mRNA COVID-19 vaccines* or polysorbate, as these are contraindicated)

**ALLERGIES**

History of the following are contraindications to receiving either of the mRNA COVID-19 vaccines:

- Severe allergic reaction (e.g., anaphylaxis) after a previous dose of an mRNA COVID-19 vaccine or any of its components
- ‘Immediate allergic reaction of any severity to a previous dose of an mRNA COVID-19 vaccine or any of its components (including polyethylene glycol [PEG])’*
- Immediate allergic reaction of any severity to polysorbate (due to potential cross-reactive hypersensitivity with the vaccine ingredient PEG)*

* These persons should not receive mRNA COVID-19 vaccination at this time unless they have been evaluated by an allergist-immunologist and it is determined that the person can safely receive the vaccine (e.g., under observation, in a setting with advanced medical care available).”
Triage of persons presenting for mRNA COVID-19 vaccination

Table Source: https://www.cdc.gov/vaccines/covid-19/info-by-product/clinical-considerations.html#Patient-counseling

† Refers only to mRNA COVID-19 vaccines currently authorized in the United States (i.e., Pfizer-BioNTech, Moderna COVID-19 vaccines)

‡ Immediate allergic reaction to a vaccine or medication is defined as any hypersensitivity-related signs or symptoms consistent with urticaria, angioedema, respiratory distress (e.g., wheezing, stridor), or anaphylaxis that occur within four hours following administration.

# These persons should not receive mRNA COVID-19 vaccination at this time unless they have been evaluated by an allergist-immunologist and it is determined that the person can safely receive the vaccine (e.g., under observation, in a setting with advanced medical care available).

COVID-19 vaccine for people with facial fillers

Two of the participants in clinical trials for Moderna’s COVID-19 vaccine who had received dermal filler injections in the cheeks were reported to have developed facial swelling after receiving the vaccination. One subject had filler injections two weeks prior to the vaccination and the other 6 months prior. A third person, who had had a history of lip injection, reportedly developed lip angioedema two days after receiving the vaccination. This participant reported a similar reaction in the past following influenza vaccination.

The localized swelling resolved itself after treatment with antihistamines or steroids in all three cases. The majority of trial participants experienced only mild side effects typical with vaccines, including headache, fatigue, and pain at the injection site. In its report on the Moderna vaccine hearing, the FDA notes that “it is possible the localized swelling in these cases is due to an inflammatory reaction from the interaction between the immune response after vaccination and the dermal filler.”

Although the data is limited, having a history of fillers should NOT prevent you from getting the COVID 19 vaccine considering the limited risks vs benefits.

To read detailed information on the Pfizer and Moderna vaccine trial click the links

FDA Pfizer Trial Summary

FDA Moderna Trial Summary
Reporting of vaccine adverse events

The Academy is closely monitoring any reports of cutaneous side effects as the vaccine rollout gets underway and asks physicians to report cases of potential dermatologic side effects in the COVID-19 Dermatology Registry.

Additional Resources on COVID 19 vaccine administration guidance from other medical societies:

1) NPF key recommendations for the management of patients with psoriatic disease during the COVID-19 pandemic: https://www.psoriasis.org/covid-19-task-force-guidance-statements/


