# IMPORTANT SAFETY INFORMATION

**Do not use TEZSPIRE** if you are allergic to tezepelumab-ekko or any of its ingredients.

Do not use to treat sudden breathing problems.

# TEZSPIRE may cause serious side effects, including:

* **severe allergic (hypersensitivity) reactions, such as rash or eye allergy.** Call your healthcare provider or get emergency help right away if you have any of the following symptoms of an allergic reaction:
  + rash
  + breathing problems
  + hives
  + red, itchy, swollen, or inflamed eyes

# Before using TEZSPIRE, tell your healthcare provider about all of your medical conditions, including if you:

* have ever had a severe allergic reaction
* have a parasitic (helminth) infection.
* have recently received or are scheduled to receive any vaccinations. You should not receive a “live vaccine” if you are treated with TEZSPIRE.
* are pregnant or plan to become pregnant.
* are breastfeeding or plan to breastfeed. It is not known if TEZSPIRE passes into your breast milk. Talk to your healthcare provider about the best way to feed your baby if you use TEZSPIRE.
* are taking prescription and over-the-counter medicines, vitamins, or herbal supplements.

**Do not** change orstop taking your other asthma medicines unless instructed to do so by your healthcare provider.

**The most common side effects of TEZSPIRE include:** Sore throat, joint and back pain. These are not all the possible side effects.

# [What is TEZSPIRE?] [Approved Use]

* TEZSPIRE is a prescription medicine used with other asthma medicines for the maintenance treatment of severe asthma in people 12 years of age and older whose asthma is not controlled with their current asthma medicine.
* TEZSPIRE helps prevent severe asthma attacks (exacerbations) and can improve your breathing.
* TEZSPIRE is not used to treat sudden breathing problems. Tell your healthcare provider if your asthma does not get better or if it gets worse after you start treatment with TEZSPIRE.
* It is not known if TEZSPIRE is safe and effective in children under 12 years of age.

**Print**

**[Please see accompanying full Prescribing Information, including Patient Information.]**

**Journal**

**[Please see Brief Summary of full Prescribing Information, including Patient Information.]**

**Digital**

**[Full Prescribing Information including Patient Information.] Must hyperlink to PI. Where possible, clicking Prescribing Information should go to top of PI and Patient Information should go directly to Patient Information section.**

DIGITAL ASSETS

*You may report side effects related to AstraZeneca products by clicking here.*

The word “here” must hyperlink to https://us-aereporting.astrazeneca.com/adverse-events.html. The Spanish version of the adverse event reporting language is also available at this site.

JOURNAL ADVERTISEMENTS

The following language must be used in HCP journal ads, consumer book of record print ads, and paid advertising print placements:

*You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.FDA.gov/medwatch or call 1-800-FDA-1088.*

PRINT ASSETS

*You are encouraged to report negative side effects of AstraZeneca prescription drugs by calling 1-800-236-9933. If you prefer to report these to the FDA, call 1-800-FDA-1088.*

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