ACTEMRA®
tocilizumab

Order Your Free ACTEMRA Travel Pack and Sharps Container

If you or your child are taking ACTEMRA SC for an FDA-approved condition, you are eligible to receive:

ACTEMRA Travel Pack



Includes:

- Prefilled syringe travel cooler
- Ice pack
- Information on treatment and the ACTEMRA Co-pay Card
- Travel mat
- TSA* card allowing you to take ACTEMRA with you when traveling

*Transportation Security Administration.

Sharps Container



A puncture-resistant container for safe disposal of needles.

To receive your complimentary ACTEMRA Travel Pack and sharps container, call **1-800-ACTEMRA (1-800-228-3672)**.

INDICATIONS AND IMPORTANT SIDE EFFECT INFORMATION

What does ACTEMRA treat?

ACTEMRA is a prescription medicine used to treat:

- Adults with moderately to severely active rheumatoid arthritis (RA) after at least one other medicine called a disease modifying antirheumatic drug (DMARD) has been used and did not work well
- Adults with giant cell arteritis (GCA)
- Patients with active polyarticular juvenile idiopathic arthritis (PJIA) 2 years of age and older

It is not known if ACTEMRA is safe and effective in children with PJIA under 2 years of age.

ACTEMRA can cause serious side effects

Serious Infections

ACTEMRA changes the way your immune system works. This can make you more likely to get infections or make any current infection worse. Some people have died from these infections. Your healthcare provider should test you for TB before starting and during treatment with ACTEMRA.

Do not take ACTEMRA if you are allergic to tocilizumab, or any of the ingredients in ACTEMRA.

ACTEMRA can cause other serious side effects. These include:

Tears (perforation) of the Stomach or Intestines

If you have diverticulitis (inflammation in parts of the large intestine), talk to your healthcare provider before taking ACTEMRA. Some people taking ACTEMRA may develop a hole in the wall of their stomach or intestines (also known as a perforation).

Changes in Blood Test Results

Your healthcare provider should do blood tests before you start receiving ACTEMRA. You should not receive ACTEMRA if your neutrophil and platelet counts are too low or your liver function test levels are too high. These may cause your healthcare provider to stop your ACTEMRA treatment for a time or change your dose.

Cancer

ACTEMRA may increase your risk of certain cancers by changing the way your immune system works.

Hepatitis B Infection

If you have or are a carrier of the hepatitis B virus (a virus that affects the liver), the virus may become active while you use ACTEMRA. Your healthcare provider may do blood tests before you start treatment with ACTEMRA and while you are using ACTEMRA.

Serious Allergic Reactions

Serious allergic reactions, including death, can happen with ACTEMRA. These reactions can happen with any infusion or injection of ACTEMRA, even if they did not occur with an earlier infusion or injection.

Nervous System Problems

While rare, Multiple Sclerosis has been diagnosed in people who take ACTEMRA.

The most common side effects of ACTEMRA include:

- upper respiratory tract infections (common cold, sinus infections)
- headache
- increased blood pressure (hypertension)
- injection site reactions

ACTEMRA & Pregnancy

Tell your healthcare provider if you are planning to become pregnant, are pregnant, plan to breast-feed, or are breast-feeding. If you are pregnant and taking ACTEMRA, join the **pregnancy registry**. To learn more, call 1-877-311-8972 or talk to your healthcare provider to register.

Tell your healthcare provider if you have any side effects. You may report side effects to FDA at 1-800-FDA-1088. You may also report side effects to Genentech at 1-888-835-2555.

For additional Important Safety Information, please see full Prescribing Information and the Medication Guide, including **Serious Side Effects**.

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