WHAT IS ACTEMRA?
ACTEMRA is a prescription medicine that targets the interleukin-6 (IL-6) signaling pathway. ACTEMRA is used to treat adults with giant cell arteritis (GCA).

IMPORTANT SIDE EFFECT INFORMATION
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 taking ACTEMRA have died from these infections.

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
WELCOME TO ACTEMRA

Now that you have been prescribed ACTEMRA subcutaneous (SC) injections, it is normal to have questions. In this brochure, we hope to provide a better understanding of what giant cell arteritis (GCA) is and how ACTEMRA may help.

Patient support resources
Have questions? Call the ACTEMRA Help Line at 1-800-228-3672 to speak with one of our specialists.
You may qualify for financial assistance. For more information on help paying for your treatment, please see pages 22-24.

ACTEMRA can cause serious side effects
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 serious infections while taking ACTEMRA, including tuberculosis (TB), and infections caused by bacteria, fungi, or viruses that can spread throughout the body. Some people have died from these infections. Your healthcare provider should test you for TB before starting ACTEMRA.

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.

INSIDE THIS BROCHURE YOU’LL FIND:

- Understanding GCA
- How is GCA treated?
- What is ACTEMRA?
- How ACTEMRA is believed to work
- Clinical trial results
- What to expect with ACTEMRA treatment
- Important Side Effect Information
- Coverage & patient assistance

Inside this kit you’ll also find:

- A Patient’s Guide to Using the Prefilled Syringe—a helpful instructional handbook
- Magnet with important numbers
- Genentech-Access.com
- Help Line: 1-800-228-3672

IMPORTANT SIDE EFFECT INFORMATION
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 taking ACTEMRA have died from these infections.

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
What is GCA?

Giant cell arteritis, or GCA, causes inflammation in the medium and large blood vessels. These blood vessels are known as arteries. GCA most often causes problems in the arteries found in your head, specifically the temples. The aorta (the biggest artery leaving the heart) can also be affected, along with its branch arteries.

GCA typically occurs in people over 50, and most commonly in people between 70 and 80 years old. Women are 2 to 3 times more likely than men to develop GCA.

Explaining your disease to others may come as a challenge. Here’s one way you could explain GCA to friends and family:

“Some of my blood vessels are inflamed. It can cause headaches and jaw pain, and may affect my vision.”

UNDERSTANDING GCA

What are common symptoms of GCA?

- New-onset headaches
- Aches, pains, and stiffness in the shoulders, neck, or hips
- Scalp tenderness
- Jaw pain
- Vision problems
- Generally feeling unwell
- Weight loss
- Fever

[Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.]
UNDERSTANDING GCA

How is GCA diagnosed?

There are a number of factors that help your healthcare provider determine if you have GCA. Symptoms (such as the ones on page 5) may offer the first sign. There are other important ways to help diagnose GCA. These include:

• Lab testing
• Imaging tests to detect inflammation in the arteries
• A temporal artery biopsy (TAB) performed by a surgeon, in which a sample of the blood vessel is removed and tested for inflammation

Why is it important to be treated for GCA quickly?

GCA should be treated as soon as possible following diagnosis. Serious issues may occur if GCA is left untreated, including blindness, stroke, and swelling of the arteries.

You’ve taken an important step by talking with your healthcare provider about GCA. It’s important that you continue to have these conversations throughout treatment.

Living with GCA

Maintaining a healthy lifestyle is important—especially if you have GCA. Here are some tips to discuss with your healthcare provider*:

Eat a healthy diet
• Eat fresh fruits and vegetables, whole grains, lean meat, and fish
• Take calcium and vitamin D to help prevent thinning bones

Exercise, if your healthcare provider says you can
• Speak with your healthcare provider to plan an exercise routine that’s right for you

Get regular checkups

Discuss the risks of smoking and drinking alcohol with your healthcare provider

*Tips on living with GCA provided by the Mayo Clinic.

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
Steroid treatment
Fortunately, there are options to treat GCA. Treatment usually involves high doses of initial steroids to control the inflammation. Once GCA is under control, the steroid dose can slowly be tapered off. While steroids have been proven in immediately treating the disease, long-term use of steroids may result in serious side effects.

As of May 2017, you have another option: ACTEMRA—the first and only FDA-approved biologic treatment for GCA.

Important Side Effect Information | Hepatitis B Infection
If you have hepatitis B, a virus that affects the liver, or are a carrier of the virus, ACTEMRA can cause the virus to become active. Your healthcare provider should test you for hepatitis B before starting treatment. Tell your healthcare provider right away if you see any signs of these symptoms: feeling very tired, vomiting, chills, dark urine, skin or eyes look yellow, clay-colored bowel movements, stomach discomfort, skin rash, little or no appetite, fevers, or muscle aches.

A biologic is a type of medication that is developed in living systems, and made from proteins that are similar to proteins that occur naturally. These treatments can be used to target certain parts of your immune system, and to fight inflammation in certain diseases such as GCA.

What is a biologic treatment?

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
With ACTEMRA, you have an option for treating GCA that may reduce your overall exposure to steroids. ACTEMRA has been used to treat over half a million patients with other conditions since 2010. As of May 2017, it’s also a treatment option for patients with GCA.

ACTEMRA is a prescription medicine that targets the IL-6 signaling pathway and is used to treat adults with GCA.

**Important Side Effect Information | Stomach tears**

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). This happens most often in people who also take nonsteroidal anti-inflammatory drugs (NSAIDs), corticosteroids, or methotrexate.

Tell your healthcare provider right away if you see any of these side effects: fever, stomach-area pain that does not go away, or if you see a change in your bowel habits.

**How ACTEMRA is believed to work**

ACTEMRA helps block the action of interleukin-6 (IL-6), a protein in your body. Research has shown that patients with GCA may have a higher amount of IL-6 in their bodies.

**Without ACTEMRA**

IL-6 connects to the cell and tells the cell to activate. When the cell activates, it may contribute to the signs and symptoms of GCA.

**With ACTEMRA**

ACTEMRA blocks IL-6 from connecting to the cell.

**IL-6 Signal may contribute to signs and symptoms of GCA**

**IL-6**  
**Cell**  
**Receptor**

**IL-6**  
**ACTEMRA**  
**IL-6**
The GCA clinical trial included patients receiving two types of treatment:

1. ACTEMRA plus steroids that were tapered off by 6 months
   - Some patients took ACTEMRA once a week; some took ACTEMRA every 2 weeks

2. Steroids alone
   - Some patients tapered off steroids at 6 months, some at 1 year

The percentage of patients who achieved sustained remission when treated with ACTEMRA weekly plus an initial dose of steroids was 4 times larger when compared with patients taking steroids alone, tapering off at 6 months.

ACTEMRA may help achieve remission when taken with steroids that are tapered off over time

RESULTS OF THE ACTEMRA CLINICAL TRIAL IN GCA

<table>
<thead>
<tr>
<th>ACTEMRA (weekly) plus initial dose of steroids</th>
<th>steroids alone (tapering off at 6 months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACTEMRA plus steroids that were tapered off by 6 months</td>
<td>ACTEMRA may increase your risk of certain cancers by changing the way your immune system works.</td>
</tr>
</tbody>
</table>

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

Important Side Effect Information | Increased risk of cancer

ACTEMRA may help achieve remission when taken with steroids that are tapered off over time

up to 56% of patients in the ACTEMRA treatment group achieved sustained remission at 1 year

14% of patients in the steroids alone treatment group (tapering off at 6 months) achieved sustained remission at 1 year

- Results for all patient groups: 56% (ACTEMRA weekly) and 53.1% (ACTEMRA every 2 weeks) vs 14% (6-month steroid taper group) and 17.6% (1-year steroid taper group)

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
If you’ve never had injection therapy before, you may be wondering what to expect. Here are a few things that will happen before, during, and after your treatment.

1. **Blood tests:** At the start of your ACTEMRA treatment, and 4 to 8 weeks after your first injection, your healthcare provider will take blood tests. He or she will repeat the tests every 3 months to check for changes in your liver function tests, neutrophil count, and platelet count. Your healthcare provider should also do blood tests to check your cholesterol levels 4 to 8 weeks after your first ACTEMRA SC injection, and then every 24 weeks.

2. **Getting to know your treatment:** Before you start on ACTEMRA SC injections, it’s important to know all the facts. So make sure you review the ACTEMRA Medication Guide, available at ACTEMRA.com. Especially take note of the “What is the most important information I should know about ACTEMRA?” and “Before you receive ACTEMRA, tell your healthcare provider about all of your medical conditions” sections.

3. **Your first injection:** Your healthcare provider or nurse can help to train you or your caregiver in his or her office on how to properly inject ACTEMRA. During this training session, you or your caregiver should perform the first injection. You should not attempt to inject yourself before receiving this in-office training. Only patients or caregivers who have been properly trained should use the ACTEMRA prefilled syringe.

4. **Monitoring for side effects at home:** ACTEMRA may lead to serious allergic reactions, including death. So it’s important to pay close attention to how you’re feeling during and after an injection. Allergic reactions may happen during and after any injection, even if they have not happened before. Contact 911 immediately, as well as your healthcare provider, if you experience any of these reactions:
   - Shortness of breath or trouble breathing
   - Swelling of the lips, tongue, or face
   - Chest pain
   - Feeling dizzy or faint
   - Moderate or severe abdominal pain or vomiting

[Learn more about what to expect at ACTEMRA.com/GCA](#)
After reading about ACTEMRA, please talk to your healthcare provider if you have any questions.

ACTEMRA is:

• Available by medical prescription only
• For adults with giant cell arteritis (GCA)

ACTEMRA can cause serious side effects

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.

Before taking ACTEMRA, tell your healthcare provider if you have:

• An infection, think you may have an infection, are being treated for an infection, or get a lot of infections that return. Infection signs, with or without a fever, include:
  - Sweating or chills
  - Shortness of breath
  - Warm, red or painful skin or sores on your body
  - Feel very tired
  - Muscle aches
  - Blood in phlegm
  - Diarrhea or stomach pain
  - Cough
  - Weight loss
  - Burning when you urinate or urinating more often than normal

IMPORTANT SIDE EFFECT INFORMATION

• Any of the following conditions that may give you a higher chance of getting infections. These include: diabetes, HIV, or a weak immune system
• Tuberculosis (TB) or have been in close contact with someone who has TB. Your healthcare provider should test you for TB before starting ACTEMRA and during treatment with ACTEMRA
• Lived in or currently live in parts of the United States known for fungal infections. These parts include the Ohio and Mississippi River Valleys and the Southwest
• Hepatitis B or have had hepatitis B

Be sure to contact your healthcare provider or nurse if you see any signs of these side effects.

Stomach tears

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). This happens most often in people who also take nonsteroidal anti-inflammatory drugs (NSAIDs), corticosteroids, or methotrexate.

Please see full Prescribing Information and Medication Guide, including Serious Side Effects, for more Important Safety Information.
Tell your healthcare provider right away if you see any of these side effects:

- Fever
- Stomach-area pain that does not go away
- Change in your bowel habits

Before starting ACTEMRA, 4 to 8 weeks after starting treatment, and then every 3 months, your healthcare provider will regularly do blood tests to check for the following side effects:

**Changes in blood test results**

*Low neutrophil count:* Neutrophils are white blood cells that help the body fight infection.

*Low platelet count:* Platelets are blood cells that help with clotting, which stops bleeding.

*Increase in liver function test levels*

*Increase in blood cholesterol levels*

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. Your cholesterol levels should be checked 4 to 8 weeks after the start of your treatment, and then every 6 months after that.

**IMPORTANT SIDE EFFECT INFORMATION (CONT’D)**

**Increased risk of cancer**

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

**Hepatitis B infection**

If you have hepatitis B, a virus that affects the liver, or are a carrier of the virus, ACTEMRA can cause the virus to become active. Your healthcare provider should test you for hepatitis B before starting treatment. Tell your healthcare provider right away if you see any signs of these symptoms:

- Feeling very tired
- Vomiting
- Chills
- Dark urine
- Skin or eyes look yellow
- Clay-colored bowel movements
- Stomach discomfort
- Skin rash
- Little or no appetite
- FEVER
- Muscle aches

If you have hepatitis B, a virus that affects the liver, or are a carrier of the virus, ACTEMRA can cause the virus to become active. Your healthcare provider should test you for hepatitis B before starting treatment. Tell your healthcare provider right away if you see any signs of these symptoms:

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- Clay-colored bowel movements
- Stomach discomfort
- Skin rash
- Little or no appetite
- FEVER
- Muscle aches

Please see full Prescribing Information and Medication Guide, including Serious Side Effects, for more Important Safety Information.
Serious allergic reactions

Serious allergic reactions, including death, can happen with ACTEMRA infusions or injections, even if they did not occur with an earlier infusion or injection. If you had hives, a rash, or experienced flushing after injecting, you should tell your healthcare provider or nurse before your next injection.

Contact 911 immediately, as well as your healthcare provider or nurse, if you experience any of these reactions:
• Shortness of breath or trouble breathing
• Chest pain
• Feeling dizzy or faint
• Moderate or severe abdominal pain or vomiting

Nervous system problems

While rare, multiple sclerosis has been diagnosed in some people taking ACTEMRA.

Increased blood pressure (also called hypertension)

Upper respiratory tract infections (like common cold and sinus infections)

Headache

Injection site reactions

Tell your healthcare provider if you have these or any other side effect that bothers you or does not go away.

ACTEMRA & pregnancy

Tell your healthcare provider if you are planning to become pregnant, are pregnant, plan to breast-feed, or are breast-feeding. You and your healthcare provider should decide if you will take ACTEMRA or breast-feed. You should not do both. If you are pregnant and taking ACTEMRA, join the pregnancy registry. The purpose of this registry is to check the health of the pregnant mother and her baby. To learn more, call 1-877-311-8972 or talk to your healthcare provider to register.

Reporting side effects

Tell your healthcare provider right away if you are experiencing any side effects. You may report side effects to the FDA at 1-800-FDA-1088 or www.FDA.gov/medwatch. You may also call Genentech at 1-888-835-2555.

Most common side effects

Tell your healthcare provider if you have these or any other side effect that bothers you or does not go away:

While rare, multiple sclerosis has been diagnosed in some people taking ACTEMRA.

Increased blood pressure (also called hypertension)

Upper respiratory tract infections (like common cold and sinus infections)

Headache

Injection site reactions

Tell your healthcare provider if you are planning to become pregnant, are pregnant, plan to breast-feed, or are breast-feeding. You and your healthcare provider should decide if you will take ACTEMRA or breast-feed. You should not do both. If you are pregnant and taking ACTEMRA, join the pregnancy registry. The purpose of this registry is to check the health of the pregnant mother and her baby. To learn more, call 1-877-311-8972 or talk to your healthcare provider to register.

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Please see full Prescribing Information and Medication Guide, including Serious Side Effects, for more Important Safety Information.
COVERAGE & PATIENT ASSISTANCE

ACTEMRA offers a variety of assistance programs that patients may be eligible for.

TYPE OF HEALTH INSURANCE COVERAGE

Government Insurance (Medicare and Medicaid)

- Patients with Medicare Part D and limited incomes may be eligible for co-pay support through the government Low Income Subsidy (LIS) Program. Patients eligible for the full subsidy pay $8.25 per drug co-pay
- Patients who don’t qualify for the LIS Program and who can’t afford their co-payments may be eligible for free drug through the Genentech® Access to Care Foundation (GATCF)

Commercial Insurance

- Patients eligible for the ACTEMRA Co-pay Card Program pay $5 per drug co-pay. The Co-pay Card may be used to cover co-pays, co-insurance, deductibles, or other out-of-pocket costs for ACTEMRA only. Terms and conditions apply

Uninsured/Underinsured

- Uninsured patients or those rendered uninsured by payer denial or the underinsured may be eligible for free drug through the Genentech® Access to Care Foundation (GATCF)

To learn more about how we can help, contact us. Call 1-800-ACTEMRA (1-800-228-3672)

What should I know?

- Eligible commercially insured patients pay $5 per drug co-pay, which can be applied to deducibles, co-pays, co-insurance, or other out-of-pocket costs for ACTEMRA
- There are no income limits
- The Co-pay Card covers your enrollment for 12 months at a time. You must reapply at the end of 12 months
- The Co-pay Card can be used if you’re just starting ACTEMRA or if you’re already on ACTEMRA

How does it work?

- For example, Julie is charged a $100 co-pay for her ACTEMRA. She pays $5 herself, and uses her Co-pay Card to cover the rest.

Uninsured/Underinsured

- She can use her card to cover up to $15,000 for ACTEMRA in co-pays during her enrollment year

How you may be eligible for the Co-pay Card*

- Your physician recommends ACTEMRA for an FDA-approved use such as Giant Cell Arteritis (GCA), and:
  - You have a commercial (private or nongovernmental) health plan. This includes plans available through state and federal exchanges
  - You are not receiving support from the Genentech® Access to Care Foundation (GATCF) or any other charitable organization
  - You do not use a federal or state-funded health care program. This includes, but is not limited to, Medicare, Medicaid, VA, DoD or TRICARE

GCA patients not eligible for the Co-pay Card may be eligible for the Genentech® Access to Care Foundation (GATCF). See page 24.

*For detailed terms and conditions, please see page 26.

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
Some patients may be eligible for co-pay support through the government Low Income Subsidy (LIS) Program. These patients include:
- Medicare Part D patients
- Patients with limited incomes

Patients eligible for the full subsidy will pay $8.25 per drug co-pay.

Some patients may be eligible for free drug through the Genentech® Access to Care Foundation (GATCF). These patients include:
- Medicare patients who don’t qualify for the LIS Program and who can’t afford their co-payments
- Uninsured patients

Help for Medicare, Medicaid, and underinsured patients

Getting your ACTEMRA prescription

Call your specialty/mail-order pharmacy. If they are a participating pharmacy, provide the Rx BIN number and Member ID from your card. If they are not a participating pharmacy, call 1-866-681-3261 for assistance.

Contact Genentech Rheumatology Access Solutions® to learn more. We are available at 1-866-681-3261 from 6 AM-5 PM PT, Monday through Friday, or 24 hours a day at www.Genentech-Access.com/ACTEMRA.

Please see Important Side Effect Information on pages 16-21. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects.
By using the ACTEMRA Co-pay Card Program, the patient acknowledges and confirms that at the time of usage, (s)he is currently eligible and meets the criteria set forth in the terms and conditions described.

The Co-pay Card is valid ONLY for patients with commercial (private or nongovernmental) insurance. Patients using Medicare, Medicaid or any other government funded program to pay for their medications are not eligible. Patients who start utilizing their Government coverage during their enrollment period will no longer be eligible for the program.

This Co-pay Card program is not health insurance or a benefit plan. Distribution or use of the Co-pay Card does not obligate use or continuing use of any specific product or provider. Patient or guardian is responsible for reporting the receipt of all Co-pay Card program benefits or reimbursement received, to any insurer, health plan, or other third party that pays for or reimburses any part of the prescription filled using the Co-pay Card program, as may be required.

The Co-pay Card is not valid for medications the patient receives for free or that are eligible to be reimbursed by private insurance plans or other healthcare or pharmaceutical assistance programs (such as: GATCF or any other charitable organization) that reimburse the patient in part or for the entire cost of his/her medication. Patient, guardian, pharmacist, prescriber and any other person using the Co-pay Card agree not to seek reimbursement for all or any part of the benefit received by the recipient through the offer.

The Co-pay Card will be accepted by participating pharmacies, physician offices or hospitals. To qualify for the benefits of this Co-pay Card program, the patient may be required to pay out-of-pocket expenses for each treatment. Once enrolled, this Co-pay Card program will not honor claims with date of service or medication dispensing that precede program enrollment by more than 120 days. This Co-pay Card is only available with a valid prescription and cannot be combined with any other rebates, free trial, or similar offer for the specified prescription. Use of this Co-pay Card must be consistent with all relevant health insurance requirements and payer agreements. Participating patients, pharmacists, physician offices and hospitals are obligated to inform third-party payers about the use of the Co-pay Card as provided for under the applicable insurance or as otherwise required by contract or law. The Co-pay Card may be used, purchased, traded or offered for sale, purchase or trade. The Co-pay Card is Limited to 1 person per peren during this offering period and is not transferable. This program expires within 12 months from enrollment. This program is not available where prohibited by law, for Massachusetts residents, the Co-pay Card is not valid for any prescription drug that has an AB-rated generic equivalent as determined by the United States Food and Drug Administration. For Massachusetts residents, this program shall expire on or before July 1, 2019. For Massachusetts residents, this program expires on or before July 1, 2019. The patient or their guardian must be 18 years or older to receive Co-pay Card program assistance. The Co-pay Card program is (1) void if the card is reproduced, (2) void where prohibited by law, (3) only valid in the United States and Puerto Rico, and (4) only valid for Genentech products. Healthcare providers may not advertise or otherwise use the program as a means of promoting their services or Genentech’s products to patients. Genentech, Inc. reserves the right to rescind, revoke, or amend the program without notice at any time.

The Co-pay Card is limited to 1 per person during this offering period and is not transferable. This program expires within 12 months from enrollment. This program is not valid where prohibited by law. For Massachusetts residents, the Co-pay Card is not valid for any prescription drug that has an AB-rated generic equivalent as determined by the United States Food and Drug Administration. For Massachusetts residents, this program shall expire on or before July 1, 2019. For Massachusetts residents, this program expires on or before July 1, 2019. The patient or their guardian must be 18 years or older to receive Co-pay Card program assistance. The Co-pay Card program is (1) void if the card is reproduced, (2) void where prohibited by law, (3) only valid in the United States and Puerto Rico, and (4) only valid for Genentech products. Healthcare providers may not advertise or otherwise use the program as a means of promoting their services or Genentech’s products to patients. Genentech, Inc. reserves the right to rescind, revoke, or amend the program without notice at any time.

The ACT Fast Program ensures quick access to therapy
For eligible commercial and government insured patients waiting longer than 5 days for a coverage decision, the ACT Fast Program provides ACTEMRA SC at no charge for up to 6 months or until coverage is established, whichever comes first. For more information, call 1-800-ACTEMRA.
A PATIENT’S GUIDE TO USING THE PREFILLED SYRINGE

Please see full Instructions for Use before using the prefilled syringe.

WHAT IS ACTEMRA?
ACTEMRA is a prescription medicine that targets the interleukin-6 (IL-6) signaling pathway. ACTEMRA is used to treat adults with giant cell arteritis (GCA).

IMPORTANT SIDE EFFECT INFORMATION
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 taking ACTEMRA have died from these infections.

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
LEARNING TO SELF-INJECT WITH THE ACTEMRA PRE FILLED SYRINGE

By choosing ACTEMRA subcutaneous (SC) injections, you and your healthcare provider have taken an important step in treating your giant cell arteritis (GCA). This easy-to-follow injection brochure will help you or your caregiver feel comfortable and confident using the ACTEMRA pre filled syringe. If you still have questions, a team of experts are here to help. Simply call 1-800-ACTEMRA (1-800-228-3672), Monday-Friday from 9 AM to 9 PM ET, for additional support.

INSIDE YOU’LL FIND INFORMATION ABOUT:

- Getting to know the ACTEMRA pre filled syringe
- Preparing for your injection
- Steps to self-injection
- Important Side Effect Information

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
UNDERSTANDING THE ACTEMRA PREFILLED SYRINGE

**Plunger** | Do not hold or pull back on the plunger. You must press the plunger all the way down to get the full dose of medication.

**Liquid** | Check the liquid. It should be clear and colorless to pale yellow. Do not inject if the liquid is cloudy, discolored, or has lumps or particles. Do not speed up the warming in any way, such as with a microwave or warm water.

**Needle Cap** | Do not remove the needle cap while allowing your ACTEMRA prefilled syringe to reach room temperature. Hold the ACTEMRA prefilled syringe with the covered needle pointed down.

**Needle** | Do not touch the needle or let it touch any surface. The syringe should be disposed of if not used within 5 minutes of needle cap removal.

**Expiration Date** | Always check date. Do not use it if the expiration date has passed.

**Trigger Fingers** | Do not touch, as this may release the needle shield early.

Image does not portray actual size

UNDERSTANDING YOUR DOSE

The recommended dose of ACTEMRA subcutaneous (SC) injection for GCA is:
- 1 prefilled syringe (162 mg) once a week, every other week, or as prescribed by your healthcare provider
- ACTEMRA can be used alone after steroids have been stopped
- ACTEMRA SC injection should never be given intravenously (into a vein)

Throughout treatment, your healthcare provider will closely monitor your lab results to see how you’re responding to ACTEMRA and check for possible side effects.

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
IMPORTANT INFORMATION ABOUT THE PREFILLED SYRINGE

- Read and follow the Instructions for Use that come with your ACTEMRA prefilled syringe before you start using it, and each time you get a prescription refill.
- Before you use the ACTEMRA prefilled syringe for the first time, make sure your healthcare provider shows you how to use it.
- Do not remove the needle cap until you’re ready to inject ACTEMRA.
- Do not try to take apart the syringe at any time.
- Do not reuse the same syringe.

If you feel your ACTEMRA prefilled syringe is damaged or have any other concerns, please call the ACTEMRA Help Line at 1-800-ACTEMRA (1-800-228-3672), Monday-Friday from 9 AM to 9 PM ET.

THE ACTEMRA PREFILLED SYRINGE

Before use

Trigger Fingers.
Do not touch, as this may release the needle shield early.

After use

Needle Shield (extended and locked)

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
Preparing for your injection

A well-lit, clean, flat surface such as a table

Alcohol pads (available at your local pharmacy)

A new ACTEMRA prefilled syringe

Cotton ball or gauze (available at your local pharmacy)

A puncture-resistant container or sharps container

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.

“If I have any questions about injecting ACTEMRA, it’s comforting to know I have the ACTEMRA Help Line.”

ACTEMRA Help Line 1-800-228-3672
Find a comfortable space with a clean, flat surface

Take the box containing the syringe out of the refrigerator and open the box. Do not touch the trigger fingers on the syringe, as this may damage the syringe

Remove one single-use ACTEMRA prefilled syringe from the box and let it sit for 30 minutes to reach room temperature. If the syringe does not reach room temperature, this could cause your injection to feel uncomfortable and make it difficult to push the plunger in

Do not speed up the warming process in any way, such as using the microwave or placing the syringe in warm water

Check the expiration date on the ACTEMRA prefilled syringe (see FIGURE A). Do not use if the expiration date has passed, because it may not be safe to use. If the expiration date has passed, safely dispose of the syringe in a sharps container and get a new one

Do not remove the needle cap while allowing your ACTEMRA prefilled syringe to reach room temperature

Keep your unused syringes in the original carton and keep in the refrigerator at 36°F-46°F (2°C-8°C). Do not freeze

Hold your ACTEMRA prefilled syringe with the covered needle pointing down (see FIGURE B)

Check the liquid in the ACTEMRA prefilled syringe. It should be clear and colorless to pale yellow. Do not inject ACTEMRA if the liquid is cloudy, discolored, or has lumps or particles in it, because it may not be safe to use. Safely dispose of the syringe in a sharps container and get a new one

Wash your hands well with soap and water

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
Step 2: Choose and prepare an injection site

Choose an injection site

- The front of your thigh and your abdomen except for the 2-inch area around your navel are the recommended injection sites (see FIGURE C).
- The outer area of the upper arms may also be used only if the injection is being given by a caregiver. Do not attempt to use the upper arm area by yourself (see FIGURE C).

Rotate injection site

- Choose a different injection site for each new injection at least 1 inch from the last area you injected.
- Do not inject into moles, scars, bruises, or areas where the skin is tender, red, hard, or not intact.

Prepare the injection site

- Wipe the injection site with an alcohol pad in a circular motion and let it air-dry to reduce the chance of getting an infection (see FIGURE D).
- Do not touch the injection site again before giving the injection.
- Do not fan or blow on the clean area.

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
Step 3: Inject ACTEMRA

- Hold the ACTEMRA prefilled syringe with one hand and pull the needle cap straight off with your other hand (see FIGURE E). Do not hold the plunger while you remove the needle cap. If you cannot remove the needle cap, you should ask a caregiver for help or contact your healthcare provider.
- Dispose of the needle cap in a sharps container or a puncture-resistant container (see FIGURE F).
- There may be a small air bubble in the ACTEMRA prefilled syringe. You do not need to remove it.
- You may see a drop of liquid at the end of the needle. This is normal and will not affect your dose.

Do not:
- Touch the needle or let it touch any surfaces.
- Use the prefilled syringe if it is dropped.
- If it is not used within 5 minutes of needle cap removal, the syringe should be disposed of in the puncture-resistant container or sharps container, and a new syringe should be used.
- Never reattach the needle cap after removal.
- Hold the ACTEMRA prefilled syringe in one hand between the thumb and index finger (see FIGURE G).

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
Step 3: Inject ACTEMRA (cont’d)

- Do not pull back on the plunger of the syringe.
- Use your other hand and gently pinch the area of skin you cleaned. Hold the pinched skin firmly. Pinching the skin is important to make sure that you inject under the skin (into fatty tissue) but not any deeper (into muscle). Injection into muscle could cause the injection to feel uncomfortable.
- Do not hold or push the plunger while inserting the needle into the skin.
- Use a quick, dart-like motion to insert the needle all the way into the pinched skin at an angle between 45° and 90° (see FIGURE H).
  It is important to use the correct angle to make sure the medicine is delivered under the skin (into fatty tissue), or the injection could be painful and the medicine may not work.
- Keep the syringe in position, and let go of the pinched skin.

FIGURE H

Slowly inject all of the medicine by gently pushing the plunger all the way down (see FIGURE I). You must press the plunger all the way down to get the full dose of medicine and to ensure the trigger fingers are completely pushed to the side.

- If the plunger is not fully depressed, the needle shield will not extend to cover the needle when it is removed.
- If the needle is not covered, carefully place the syringe into the sharps container or puncture-resistant container to avoid injury with the needle.
- Do not put the needle cap back on the needle.

Keep in mind that the ACTEMRA solution in the syringe is thicker than water. It may take some pressure to push the plunger all the way down.

FIGURE I

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
After the plunger is pushed all the way down, keep pressing down on the plunger to be sure all of the medicine is injected before taking the needle out of the skin.

Keep pressing down on the plunger while you take the needle out of the skin at the same angle as inserted (see FIGURE J).

After the needle is removed completely from the skin, release the plunger, allowing the needle shield to protect the needle (see FIGURE K).

After the injection:
- There may be a little bleeding at the injection site. You can press a cotton ball or gauze over the injection site.
- Do not rub the injection site.
- If needed, you may cover the injection site with a small bandage.

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
The ACTEMRA prefilled syringe should not be reused. Put the used syringe in your sharps container or puncture-resistant container (see FIGURE L). Do not put the needle cap back on the needle.

If your injection is given by another person, this person must also be careful when removing the needle and disposing of the syringe to prevent accidental needle stick injury and avoid passing infection.

How do I dispose of used syringes?
- Put your used needles and syringes, including ACTEMRA, in an FDA-cleared sharps disposal container or a puncture-resistant container right away after use. Do not dispose of loose needles and syringes in your household trash.
- When your sharps disposal container or puncture-resistant container is almost full, you will need to follow your community guidelines for the right way to dispose of your sharps disposal container or puncture-resistant container. There may be state or local laws about how you should dispose of used needles and syringes. For more information about safe sharps container and puncture-resistant container disposal, and for specific information about sharps container disposal and puncture-resistant container disposal in the state you live in, go to the FDA’s website at www.fda.gov/safesharpsdisposal.
- Do not dispose of your used sharps disposal container or puncture-resistant container in your household trash unless your community guidelines permit this. Do not recycle your used sharps disposal container or puncture-resistant container.
- Keep ACTEMRA prefilled syringes and the sharps disposal container or puncture-resistant container out of the reach of children.
- Record your injection: Write down the date, time, and specific part of your body where you injected yourself. It may also be helpful to write down any questions or concerns about the injection so you can ask your healthcare provider.
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. Tell your healthcare provider before your next dose if you had hives, rash, or flushing after your injection. Seek medical attention right away if you have any of the following signs of a serious allergic reaction:

- Shortness of breath or trouble breathing
- Feeling dizzy or faint
- Swelling of the lips, tongue, or face
- Chest pain
- Moderate or severe abdominal pain or vomiting
- Feeling dizzy or faint
- Chest pain
- Swelling of the lips, tongue, or face
- Moderate or severe abdominal pain or vomiting

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.

Call your healthcare provider and seek medical attention without delay if you think you might be developing an infection.

If you have any concerns or questions about your syringe, contact your healthcare provider or pharmacist for assistance.

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.
There are several resources offered to patients treating their GCA with ACTEMRA

Have questions? Call the ACTEMRA Help Line at 1-800-228-3672 to speak live with one of our specialists.
You may qualify for financial assistance. For more information on help paying for your treatment, please see pages 22-27 in the accompanying brochure.

After reading about ACTEMRA, please talk to your healthcare provider if you have any questions.

ACTEMRA is:
» Available by medical prescription only
» For adults with giant cell arteritis (GCA)

ACTEMRA can cause serious side effects
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 serious infections while taking ACTEMRA, including tuberculosis (TB), and infections caused by bacteria, fungi, or viruses that can spread throughout the body. Some people have died from these infections. Your healthcare provider should test you for TB before starting ACTEMRA.

Please see full Prescribing Information and Medication Guide, including Serious Side Effects, for more Important Safety Information, as well as full Instructions for Use.
ACTEMRA can cause other serious side effects
These include:

- Stomach tears
- Changes in blood test results, including low neutrophil and platelet counts, and increases in certain liver function test levels and blood cholesterol levels
- An increased risk of certain cancers by changing the way your immune system works

Do not take ACTEMRA if you:

- Are allergic to ACTEMRA, or any of the ingredients in ACTEMRA

Important Side Effect Information (cont’d)

ACTEMRA can cause other serious side effects
These include:

- Stomach tears
- Changes in blood test results, including low neutrophil and platelet counts, and increases in certain liver function test levels and blood cholesterol levels
- An increased risk of certain cancers by changing the way your immune system works

Do not take ACTEMRA if you:

- Are allergic to ACTEMRA, or any of the ingredients in ACTEMRA

Upper respiratory tract infections (like common cold and sinus infections)
Increased blood pressure (also called hypertension)
Headache
Injection site reactions

ACTEMRA can cause other serious side effects
These include:

- Serious allergic reactions, including death. These may happen with ACTEMRA infusions or injections, even if they did not occur with an earlier infusion or injection. If you had hives, a rash, or experienced flushing after injecting, you should tell your healthcare provider or nurse before your next injection
- Hepatitis B infection
- Nervous system problems

Tell your healthcare provider if you have these or any other side effect that bothers you or does not go away:

- Increased blood pressure (also called hypertension)
- Upper respiratory tract infections (like common cold and sinus infections)
- Headache
- Injection site reactions

Tell your healthcare provider if you are planning to become pregnant, are pregnant, plan to breast-feed, or are breast-feeding. You and your healthcare provider should decide if you will take ACTEMRA or breast-feed. You should not do both. 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.

ACTEMRA & pregnancy

Tell your healthcare provider right away if you are experiencing any side effects. You may report side effects to the FDA at 1-800-FDA-1088 or www.FDA.gov/medwatch. You may also call Genentech at 1-888-835-2555.

Please see full Prescribing Information and Medication Guide, including Serious Side Effects, for more Important Safety Information, as well as full Instructions for Use.
Visit ACTEMRA.com/GCA or call 1-800-ACTEMRA (1-800-228-3672) for more information

Please see Important Side Effect Information on pages 25-27. For more Important Safety Information, please see full Prescribing Information and Medication Guide, including Serious Side Effects, as well as full Instructions for Use.

ACTEMRA is a registered trademark of Chugai Seiyaku Kabushiki Kaisha Corp., a member of the Roche Group.
HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use ACTEMRA safely and effectively. See full prescribing information for ACTEMRA.

ACTEMRA® (tocilizumab) injection, for intravenous or subcutaneous use
Initial U.S. Approval: 2010

WARNING: RISK OF SERIOUS INFECTIONS
See full prescribing information for complete boxed warning.
- Serious infections leading to hospitalization or death including tuberculosis (TB), bacterial, invasive fungal, viral, and other opportunistic infections have occurred in patients receiving ACTEMRA. (5.1)
- If a serious infection develops, interrupt ACTEMRA until the infection is controlled. (5.1)
- Perform test for latent TB; if positive, start treatment for TB prior to starting ACTEMRA. (5.1)
- Monitor all patients for active TB during treatment, even if initial latent TB test is negative. (5.1)

Dosage and Administration (2.1, 2.5, 2.6, 2.7)
For RA, JIA and SJIA, ACTEMRA may be used alone or in combination with methotrexate: and in RA, other DMARDs may be used. (2.2)

Recommended Adult Subcutaneous Dosage:

\[
\begin{array}{|c|c|}
\hline
\text{Patients less than 100 kg weight} & 162 \text{ mg administered subcutaneously every week, followed by an increase to every week based on clinical response} \\
\hline
\text{Patients at or above 100 kg weight} & 162 \text{ mg administered subcutaneously every week} \\
\hline
\end{array}
\]

Recommended Adult Intravenous Dosage

For RA, pJIA and SJIA patients, ACTEMRA may be used alone or in combination with methotrexate and other DMARDs. (2.2)

Recommended Adult Intravenous Dosage:

\[
\begin{array}{|c|c|}
\hline
\text{Patients less than 30 kg weight} & 12 \text{ mg per kg} \\
\hline
\text{Patients at or above 30 kg weight} & 8 \text{ mg per kg} \\
\hline
\end{array}
\]

General Dosing Information (2.6)
- It is recommended that ACTEMRA not be initiated in patients with an absolute neutrophil count (ANC) below 2000 per mm³, platelet count below 100,000 per mm³, or who have ALT or AST above 1.5 times the upper limit of normal (ULN). (2.1, 5.3)
- ACTEMRA doses exceeding 800 mg per infusion are not recommended in RA or CRS patients. (2.1, 2.5, 12.3)

Dosage Modifications (2.9)
- Recommended for management of certain dose-related laboratory changes including elevated liver enzymes, neutropenia, and thrombocytopenia.

Recommended Intravenous CRS Dosage

\[
\begin{array}{|c|c|}
\hline
\text{Patients less than 30 kg weight} & 12 \text{ mg per kg} \\
\hline
\text{Patients at or above 30 kg weight} & 8 \text{ mg per kg} \\
\hline
\end{array}
\]

Dose Modifications (2.9)
- Recommended for management of certain dose-related laboratory changes including elevated liver enzymes, neutropenia, and thrombocytopenia.

Recommended Subcutaneous SJIA Dosage

\[
\begin{array}{|c|c|}
\hline
\text{Patients less than 30 kg weight} & 162 \text{ mg every two weeks} \\
\hline
\text{Patients at or above 30 kg weight} & 162 \text{ mg every two week} \\
\hline
\end{array}
\]

Recommended Subcutaneous PJIA Dosage

\[
\begin{array}{|c|c|}
\hline
\text{Patients less than 30 kg weight} & 162 \text{ mg once every two weeks} \\
\hline
\text{Patients at or above 30 kg weight} & 162 \text{ mg once every three weeks} \\
\hline
\end{array}
\]

Recommended Intravenous SJIA Dosage Every 2 Weeks

Cytokine Release Syndrome (2.5)
- Gastrointestinal (GI) perforation—use with caution in patients who may be at increased risk. (5.2)
- Laboratory monitoring—recommended due to potential consequences of treatment-related changes in neutrophils, platelets, lipids, and liver function tests. (2.8, 5.3)
- Hypersensitivity reactions, including anaphylaxis and death have occurred. (5.5)
- Live vaccines—Avoid use with ACTEMRA. (5.8, 7.3)

- -----------------------------ADVERSE REACTIONS-----------------------------

Most common adverse reactions (incidence of at least 5%): upper respiratory tract infections, nasopharyngitis, headache, hypertension, increased ALT, injection site reactions. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Genentech at 1-888-835-2555 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

- USE IN SPECIFIC POPULATIONS----------------------

- Pregnancy: Based on animal data, may cause fetal harm. (8.1)
- Lactation: Discontinue drug or nursing taking into consideration importance of drug to mother. (8.2)

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide

Revised: 11/2018
FULL PRESCRIBING INFORMATION: CONTENTS*

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*Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

WARNING: RISK OF SERIOUS INFECTIONS

Patients treated with ACTEMRA are at increased risk for developing serious infections that may lead to hospitalization or death [see Warnings and Precautions (5.1), Adverse Reactions (6.1)]. Most patients who developed these infections were taking concomitant immunosuppressants such as methotrexate or corticosteroids.

If a serious infection develops, interrupt ACTEMRA until the infection is controlled.

Reported infections include:

- Active tuberculosis, which may present with pulmonary or extrapulmonary disease. Patients should be tested for latent tuberculosis before ACTEMRA use and during therapy. Treatment for latent infection should be initiated prior to ACTEMRA use.

- Invasive fungal infections, including candidiasis, aspergillosis, and pneumocystis. Patients with invasive fungal infections may present with disseminated, rather than localized, disease.

- Bacterial, viral and other infections due to opportunistic pathogens.

The risks and benefits of treatment with ACTEMRA should be carefully considered prior to initiating therapy in patients with chronic or recurrent infection.
Patients should be closely monitored for the development of signs and symptoms of infection during and after treatment with ACTEMRA, including the possible development of tuberculosis in patients who tested negative for latent tuberculosis infection prior to initiating therapy [see Warnings and Precautions (5.1)].

1 INDICATIONS AND USAGE

1.1 Rheumatoid Arthritis (RA)
ACTEMRA® (tocilizumab) is indicated for the treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response to one or more Disease-Modifying Anti-Rheumatic Drugs (DMARDs).

1.2 Giant Cell Arteritis (GCA)
ACTEMRA® (tocilizumab) is indicated for the treatment of giant cell arteritis (GCA) in adult patients.

1.3 Polyarticular Juvenile Idiopathic Arthritis (PJIA)
ACTEMRA® (tocilizumab) is indicated for the treatment of active polyarticular juvenile idiopathic arthritis in patients 2 years of age and older.

1.4 Systemic Juvenile Idiopathic Arthritis (SJIA)
ACTEMRA® (tocilizumab) is indicated for the treatment of active systemic juvenile idiopathic arthritis in patients 2 years of age and older.

1.5 Cytokine Release Syndrome (CRS)
ACTEMRA® (tocilizumab) is indicated for the treatment of chimeric antigen receptor (CAR) T cell-induced severe or life-threatening cytokine release syndrome in adults and pediatric patients 2 years of age and older.

2 DOSAGE AND ADMINISTRATION

2.1 Rheumatoid Arthritis
ACTEMRA may be used as monotherapy or concomitantly with methotrexate or other non-biologic DMARDs as an intravenous infusion or as a subcutaneous injection.

Recommended Intravenous Dosage Regimen:
The recommended dosage of ACTEMRA for adult patients given as a 60-minute single intravenous drip infusion is 4 mg per kg every 4 weeks followed by an increase to 8 mg per kg every 4 weeks based on clinical response.

- Reduction of dose from 8 mg per kg to 4 mg per kg is recommended for management of certain dose-related laboratory changes including elevated liver enzymes, neutropenia, and thrombocytopenia [see Dosage and Administration (2.9), Warnings and Precautions (5.3), and Adverse Reactions (6.1)].
- Doses exceeding 800 mg per infusion are not recommended in RA patients [see Clinical Pharmacology (12.3)].

Recommended Subcutaneous Dosage Regimen:

<table>
<thead>
<tr>
<th>Patients less than 100 kg weight</th>
<th>162 mg administered subcutaneously every other week, followed by an increase to every week based on clinical response</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients at or above 100 kg weight</td>
<td>162 mg administered subcutaneously every week</td>
</tr>
</tbody>
</table>

When transitioning from ACTEMRA intravenous therapy to subcutaneous administration administer the first subcutaneous dose instead of the next scheduled intravenous dose.

 Interruption of dose or reduction in frequency of administration of subcutaneous dose from every week to every other week dosing is recommended for management of certain dose-related laboratory changes including elevated
liver enzymes, neutropenia, and thrombocytopenia [see Dosage and Administration (2.9), Warnings and Precautions (5.3), and Adverse Reactions (6.2)].

2.2 Giant Cell Arteritis

The recommended dose of ACTEMRA for adult patients with GCA is 162 mg given once every week as a subcutaneous injection in combination with a tapering course of glucocorticoids.

A dose of 162 mg given once every other week as a subcutaneous injection in combination with a tapering course of glucocorticoids may be prescribed based on clinical considerations.

ACTEMRA can be used alone following discontinuation of glucocorticoids.

- Interruption of dosing may be needed for management of dose-related laboratory abnormalities including elevated liver enzymes, neutropenia, and thrombocytopenia [see Dosage and Administration (2.9)].
- Intravenous administration is not approved for GCA.

2.3 Polyarticular Juvenile Idiopathic Arthritis

ACTEMRA may be used alone or in combination with methotrexate. Do not change dose based solely on a single visit body weight measurement, as weight may fluctuate.

Recommended Intravenous Dosage Regimen:

The recommended dosage of ACTEMRA for PJIA patients given once every 4 weeks as a 60-minute single intravenous drip infusion is:

<table>
<thead>
<tr>
<th>Recommended Intravenous PJIA Dosage Every 4 Weeks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients less than 30 kg weight</td>
</tr>
<tr>
<td>Patients at or above 30 kg weight</td>
</tr>
</tbody>
</table>

Recommended Subcutaneous Dosage Regimen:

<table>
<thead>
<tr>
<th>Recommended Subcutaneous PJIA Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients less than 30 kg weight</td>
</tr>
<tr>
<td>Patients at or above 30 kg weight</td>
</tr>
</tbody>
</table>

When transitioning from ACTEMRA intravenous therapy to subcutaneous administration, administer the first subcutaneous dose instead of the next scheduled intravenous dose.

Interruption of dosing may be needed for management of dose-related laboratory abnormalities including elevated liver enzymes, neutropenia, and thrombocytopenia [see Dosage and Administration (2.9)].

2.4 Systemic Juvenile Idiopathic Arthritis

ACTEMRA may be used as an intravenous infusion or as a subcutaneous injection alone or in combination with methotrexate. Do not change a dose based solely on a single visit body weight measurement, as weight may fluctuate.

Recommended Intravenous Dosage Regimen:

The recommended dose of ACTEMRA for SJIA patients given once every 2 weeks as a 60-minute single intravenous drip infusion is:
Recommended Intravenous SJIA Dosage Every 2 Weeks

| Patients less than 30 kg weight | 12 mg per kg |
| Patients at or above 30 kg weight | 8 mg per kg |

Recommended Subcutaneous Dosage Regimen:

<table>
<thead>
<tr>
<th>Recommended Subcutaneous SJIA Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients less than 30 kg weight</td>
</tr>
<tr>
<td>Patients at or above 30 kg weight</td>
</tr>
</tbody>
</table>

When transitioning from ACTEMRA intravenous therapy to subcutaneous administration, administer the first subcutaneous dose when the next scheduled intravenous dose is due.

Interruption of dosing may be needed for management of dose-related laboratory abnormalities including elevated liver enzymes, neutropenia, and thrombocytopenia [see Dosage and Administration (2.9)].

2.5 Cytokine Release Syndrome (CRS)

Use only the intravenous route for treatment of CRS. The recommended dose of ACTEMRA for treatment of CRS given as a 60-minute intravenous infusion is:

<table>
<thead>
<tr>
<th>Recommended Intravenous CRS Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients less than 30 kg weight</td>
</tr>
<tr>
<td>Patients at or above 30 kg weight</td>
</tr>
</tbody>
</table>

- Alone or in combination with corticosteroids

- If no clinical improvement in the signs and symptoms of CRS occurs after the first dose, up to 3 additional doses of ACTEMRA may be administered. The interval between consecutive doses should be at least 8 hours.
- Doses exceeding 800 mg per infusion are not recommended in CRS patients.
- Subcutaneous administration is not approved for CRS

2.6 General Considerations for Administration

- ACTEMRA has not been studied in combination with biological DMARDs such as TNF antagonists, IL-1R antagonists, anti-CD20 monoclonal antibodies and selective co-stimulation modulators because of the possibility of increased immunosuppression and increased risk of infection. Avoid using ACTEMRA with biological DMARDs.
- It is recommended that ACTEMRA not be initiated in patients with an absolute neutrophil count (ANC) below 2000 per mm³, platelet count below 100,000 per mm³, or who have ALT or AST above 1.5 times the upper limit of normal (ULN).
  - Patients with severe or life-threatening CRS frequently have cytopenias or elevated ALT or AST due to the lymphodepleting chemotherapy or the CRS. The decision to administer ACTEMRA should take into account the potential benefit of treating the CRS versus the risks of short-term treatment with ACTEMRA.

2.7 Preparation and Administration Instructions for Intravenous Infusion

ACTEMRA for intravenous infusion should be diluted by a healthcare professional using aseptic technique as follows:
Patients less than 30 kg: use a 50 mL infusion bag or bottle of 0.9% or 0.45% Sodium Chloride Injection, USP, and then follow steps 1 and 2 below.

Patients at or above 30 kg weight: use a 100 mL infusion bag or bottle, and then follow steps 1 and 2 below.

Step 1. Withdraw a volume of 0.9% or 0.45% Sodium Chloride Injection, USP, equal to the volume of the ACTEMRA injection required for the patient’s dose from the infusion bag or bottle [see Dosage and Administration (2.1, 2.3, 2.4, 2.5)].

<table>
<thead>
<tr>
<th>Dosage</th>
<th>Indication</th>
<th>Volume of ACTEMRA injection per kg of body weight</th>
</tr>
</thead>
<tbody>
<tr>
<td>4 mg/kg</td>
<td>Adult RA</td>
<td>0.2mL/kg</td>
</tr>
<tr>
<td>8 mg/kg</td>
<td>Adult RA, SJIA, PJIA and CRS (≥30 kg of body weight)</td>
<td>0.4mL/kg</td>
</tr>
<tr>
<td>10 mg/kg</td>
<td>PJIA (&lt; 30 kg of body weight)</td>
<td>0.5 mL/kg</td>
</tr>
<tr>
<td>12 mg/kg</td>
<td>SJIA and CRS (&lt; 30 kg of body weight)</td>
<td>0.6mL/kg</td>
</tr>
</tbody>
</table>

Step 2. Withdraw the amount of ACTEMRA for intravenous infusion from the vial(s) and add slowly into the 0.9% or 0.45% Sodium Chloride Injection, USP infusion bag or bottle. To mix the solution, gently invert the bag to avoid foaming.

- The fully diluted ACTEMRA solutions for infusion using 0.9% Sodium Chloride Injection, USP may be stored at 2° to 8°C (36° to 46°F) or room temperature for up to 24 hours and should be protected from light.
- The fully diluted ACTEMRA solutions for infusion using 0.45% Sodium Chloride Injection, USP may be stored at 2° to 8°C (36° to 46°F) for up to 24 hours or room temperature for up to 4 hours and should be protected from light.
- ACTEMRA solutions do not contain preservatives; therefore, unused product remaining in the vials should not be used.
- Allow the fully diluted ACTEMRA solution to reach room temperature prior to infusion.
- The infusion should be administered over 60 minutes, and must be administered with an infusion set. Do not administer as an intravenous push or bolus.
- ACTEMRA should not be infused concomitantly in the same intravenous line with other drugs. No physical or biochemical compatibility studies have been conducted to evaluate the co-administration of ACTEMRA with other drugs.
- Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. If particulates and discolorations are noted, the product should not be used.
- Fully diluted ACTEMRA solutions are compatible with polypropylene, polyethylene and polyvinyl chloride infusion bags and polypropylene, polyethylene and glass infusion bottles.

2.8 Preparation and Administration Instructions for Subcutaneous Injection

- ACTEMRA for subcutaneous injection is not intended for intravenous drip infusion.
- Assess suitability of patient for subcutaneous home use and instruct patients to inform a healthcare professional before administering the next dose if they experience any symptoms of allergic reaction. Patients should seek immediate medical attention if they develop symptoms of serious allergic reactions. ACTEMRA subcutaneous injection is intended for use under the guidance of a healthcare practitioner. After proper training in subcutaneous injection technique, a patient may self-inject ACTEMRA or the patient’s caregiver may administer ACTEMRA if a healthcare practitioner determines that it is appropriate. PJIA and SJIA patients may self-inject with ACTEMRA prefilled syringe or the patient’s caregiver may administer ACTEMRA if both the healthcare practitioner and the parent/legal guardian determines it is appropriate. The ability of pediatric patients to self-inject with the autoinjector has not been tested. Patients, or patient caregivers, should
be instructed to follow the directions provided in the Instructions for Use (IFU) for additional details on medication administration

- Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration. Do not use ACTEMRA prefilled syringes (PFS) or autoinjectors (AI) exhibiting particulate matter, cloudiness, or discoloration. ACTEMRA for subcutaneous administration should be clear and colorless to pale yellow. Do not use if any part of the PFS or AI appears to be damaged.

- Patients using ACTEMRA for subcutaneous administration should be instructed to inject the full amount in the syringe (0.9 mL) or full amount in the autoinjector (0.9 mL), which provides 162 mg of ACTEMRA, according to the directions provided in the IFU.

- Injection sites should be rotated with each injection and should never be given into moles, scars, or areas where the skin is tender, bruised, red, hard, or not intact.

2.9 **Dosage Modifications due to Serious Infections or Laboratory Abnormalities**

Hold ACTEMRA treatment if a patient develops a serious infection until the infection is controlled.

**Rheumatoid Arthritis and Giant Cell Arteritis**

<table>
<thead>
<tr>
<th>Lab Value</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Greater than 1 to 3x ULN (confirmed by repeat testing)</td>
<td>Hold ACTEMRA dosing until less than 3x ULN and follow recommendations above for greater than 1 to 3x ULN</td>
</tr>
<tr>
<td>Greater than 3 to 5x ULN (confirmed by repeat testing)</td>
<td>Dose modify concomitant DMARDs (RA) or immunomodulatory agents (GCA) if appropriate</td>
</tr>
</tbody>
</table>

For persistent increases in this range:

- For patients receiving intravenous ACTEMRA, reduce dose to 4 mg per kg or hold ACTEMRA until ALT or AST have normalized
- For patients receiving subcutaneous ACTEMRA, reduce injection frequency to every other week or hold dosing until ALT or AST have normalized. Resume ACTEMRA at every other week and increase frequency to every week as clinically appropriate.

For persistent increases greater than 3x ULN, discontinue ACTEMRA

Greater than 5x ULN | Discontinue ACTEMRA

<table>
<thead>
<tr>
<th>Lab Value</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Greater than 3 to 5x ULN (confirmed by repeat testing)</td>
<td>Hold ACTEMRA dosing until less than 3x ULN and follow recommendations above for greater than 1 to 3x ULN</td>
</tr>
<tr>
<td>Greater than 3 to 5x ULN (confirmed by repeat testing)</td>
<td>Dose modify concomitant DMARDs (RA) or immunomodulatory agents (GCA) if appropriate</td>
</tr>
</tbody>
</table>

For persistent increases greater than 3x ULN, discontinue ACTEMRA

Greater than 5x ULN | Discontinue ACTEMRA
### Low Absolute Neutrophil Count (ANC) [see Warnings and Precautions (5.3)]:

<table>
<thead>
<tr>
<th>Lab Value (cells per mm³)</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>ANC greater than 1000</td>
<td>Maintain dose</td>
</tr>
<tr>
<td>ANC 500 to 1000</td>
<td>Hold ACTEMRA dosing</td>
</tr>
</tbody>
</table>
  - When ANC greater than 1000 cells per mm³:  
    - For patients receiving intravenous ACTEMRA, resume ACTEMRA at 4 mg per kg and increase to 8 mg per kg as clinically appropriate  
    - For patients receiving subcutaneous ACTEMRA, resume ACTEMRA at every other week and increase frequency to every week as clinically appropriate  
| ANC less than 500        | Discontinue ACTEMRA |

### Low Platelet Count [see Warnings and Precautions (5.3)]:

<table>
<thead>
<tr>
<th>Lab Value (cells per mm³)</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>50,000 to 100,000</td>
<td>Hold ACTEMRA dosing</td>
</tr>
</tbody>
</table>
  - When platelet count is greater than 100,000 cells per mm³:  
    - For patients receiving intravenous ACTEMRA, resume ACTEMRA at 4 mg per kg and increase to 8 mg per kg as clinically appropriate  
    - For patients receiving subcutaneous ACTEMRA, resume ACTEMRA at every other week and increase frequency to every week as clinically appropriate  
| Less than 50,000         | Discontinue ACTEMRA |

**Polyarticular and Systemic Juvenile Idiopathic Arthritis:**

Dose reduction of ACTEMRA has not been studied in the PJIA and SJIA populations. Dose interruptions of ACTEMRA are recommended for liver enzyme abnormalities, low neutrophil counts, and low platelet counts in patients with PJIA and SJIA at levels similar to what is outlined above for patients with RA. If appropriate, dose modify or stop concomitant methotrexate and/or other medications and hold ACTEMRA dosing until the clinical situation has been evaluated. In PJIA and SJIA the decision to discontinue ACTEMRA for a laboratory abnormality should be based upon the medical assessment of the individual patient.

### 3 DOSAGE FORMS AND STRENGTHS

**Intravenous Infusion**

Injection: 80 mg/4 mL, 200 mg/10 mL, 400 mg/20 mL as a clear, colorless to pale yellow solution in 20 mg/mL single-dose vials for further dilution prior to intravenous infusion.
Subcutaneous Injection
Injection: 162 mg/0.9 mL clear, colorless to slightly yellowish solution in a single-dose prefilled syringe or autoinjector.

4 CONTRAINDICATIONS
ACTEMRA is contraindicated in patients with known hypersensitivity to ACTEMRA [see Warnings and Precautions (5.5)].

5 WARNINGS AND PRECAUTIONS
5.1 Serious Infections
Serious and sometimes fatal infections due to bacterial, mycobacterial, invasive fungal, viral, protozoal, or other opportunistic pathogens have been reported in patients receiving immunosuppressive agents including ACTEMRA. The most common serious infections included pneumonia, urinary tract infection, cellulitis, herpes zoster, gastroenteritis, diverticulitis, sepsis and bacterial arthritis [see Adverse Reactions (6.1)]. Among opportunistic infections, tuberculosis, cryptococcus, aspergillosis, candidiasis, and pneumocystosis were reported with ACTEMRA. Other serious infections, not reported in clinical studies, may also occur (e.g., histoplasmosis, coccidioidomycosis, listeriosis). Patients have presented with disseminated rather than localized disease, and were often taking concomitant immunosuppressants such as methotrexate or corticosteroids which in addition to rheumatoid arthritis may predispose them to infections.

Do not administer ACTEMRA in patients with an active infection, including localized infections. The risks and benefits of treatment should be considered prior to initiating ACTEMRA in patients:

- with chronic or recurrent infection;
- who have been exposed to tuberculosis;
- with a history of serious or an opportunistic infection;
- who have resided or traveled in areas of endemic tuberculosis or endemic mycoses; or
- with underlying conditions that may predispose them to infection.

Closely monitor patients for the development of signs and symptoms of infection during and after treatment with ACTEMRA, as signs and symptoms of acute inflammation may be lessened due to suppression of the acute phase reactants [see Dosage and Administration (2.6), Adverse Reactions (6.1), and Patient Counseling Information (17)].

Hold ACTEMRA if a patient develops a serious infection, an opportunistic infection, or sepsis. A patient who develops a new infection during treatment with ACTEMRA should undergo a prompt and complete diagnostic workup appropriate for an immunocompromised patient, initiate appropriate antimicrobial therapy, and closely monitor the patient.

Tuberculosis
Evaluate patients for tuberculosis risk factors and test for latent infection prior to initiating ACTEMRA.

Consider anti-tuberculosis therapy prior to initiation of ACTEMRA in patients with a past history of latent or active tuberculosis in whom an adequate course of treatment cannot be confirmed, and for patients with a negative test for latent tuberculosis but having risk factors for tuberculosis infection. Consultation with a physician with expertise in the treatment of tuberculosis is recommended to aid in the decision whether initiating anti-tuberculosis therapy is appropriate for an individual patient.

Closely monitor patients for the development of signs and symptoms of tuberculosis including patients who tested negative for latent tuberculosis infection prior to initiating therapy.

It is recommended that patients be screened for latent tuberculosis infection prior to starting ACTEMRA. The incidence of tuberculosis in worldwide clinical development programs is 0.1%. Patients with latent tuberculosis should be treated with standard antituberculous therapy before initiating ACTEMRA.
**Viral Reactivation**

Viral reactivation has been reported with immunosuppressive biologic therapies and cases of herpes zoster exacerbation were observed in clinical studies with ACTEMRA. No cases of Hepatitis B reactivation were observed in the trials; however patients who screened positive for hepatitis were excluded.

**5.2 Gastrointestinal Perforations**

Events of gastrointestinal perforation have been reported in clinical trials, primarily as complications of diverticulitis in patients treated with ACTEMRA. Use ACTEMRA with caution in patients who may be at increased risk for gastrointestinal perforation. Promptly evaluate patients presenting with new onset abdominal symptoms for early identification of gastrointestinal perforation [see Adverse Reactions (6.1)].

**5.3 Laboratory Parameters**

Rheumatoid Arthritis and Giant Cell Arteritis

**Neutropenia**

Treatment with ACTEMRA was associated with a higher incidence of neutropenia. Infections have been uncommonly reported in association with treatment-related neutropenia in long-term extension studies and postmarketing clinical experience.

- It is not recommended to initiate ACTEMRA treatment in patients with a low neutrophil count, i.e., absolute neutrophil count (ANC) less than 2000 per mm$^3$. In patients who develop an absolute neutrophil count less than 500 per mm$^3$ treatment is not recommended.

- Monitor neutrophils 4 to 8 weeks after start of therapy and every 3 months thereafter [see Clinical Pharmacology (12.2)]. For recommended modifications based on ANC results see Dosage and Administration (2.9).

**Thrombocytopenia**

Treatment with ACTEMRA was associated with a reduction in platelet counts. Treatment-related reduction in platelets was not associated with serious bleeding events in clinical trials [see Adverse Reactions (6.1, 6.2)].

- It is not recommended to initiate ACTEMRA treatment in patients with a platelet count below 100,000 per mm$^3$. In patients who develop a platelet count less than 50,000 per mm$^3$ treatment is not recommended.

- Monitor platelets 4 to 8 weeks after start of therapy and every 3 months thereafter. For recommended modifications based on platelet counts see Dosage and Administration (2.9).

**Elevated Liver Enzymes**

Treatment with ACTEMRA was associated with a higher incidence of transaminase elevations. These elevations did not result in apparent permanent or clinically evident hepatic injury in clinical trials [see Adverse Reactions (6.1, 6.2)]. Increased frequency and magnitude of these elevations was observed when potentially hepatotoxic drugs (e.g., MTX) were used in combination with ACTEMRA.

In one case, a patient who had received ACTEMRA 8 mg per kg monotherapy without elevations in transaminases experienced elevation in AST to above 10x ULN and elevation in ALT to above 16x ULN when MTX was initiated in combination with ACTEMRA. Transaminases normalized when both treatments were held, but elevations recurred when MTX and ACTEMRA were restarted at lower doses. Elevations resolved when MTX and ACTEMRA were discontinued.

- It is not recommended to initiate ACTEMRA treatment in patients with elevated transaminases ALT or AST greater than 1.5x ULN. In patients who develop elevated ALT or AST greater than 5x ULN treatment is not recommended.
Monitor ALT and AST levels 4 to 8 weeks after start of therapy and every 3 months thereafter. When clinically indicated, other liver function tests such as bilirubin should be considered. For recommended modifications based on transaminases [see Dosage and Administration (2.9)].

**Lipid Abnormalities**
Treatment with ACTEMRA was associated with increases in lipid parameters such as total cholesterol, triglycerides, LDL cholesterol, and/or HDL cholesterol [see Adverse Reactions (6.1, 6.2)].

- Assess lipid parameters approximately 4 to 8 weeks following initiation of ACTEMRA therapy, then at approximately 24 week intervals.

- Manage patients according to clinical guidelines [e.g., National Cholesterol Educational Program (NCEP)] for the management of hyperlipidemia.

**Polyarticular and Systemic Juvenile Idiopathic Arthritis**
A similar pattern of liver enzyme elevation, low neutrophil count, low platelet count and lipid elevations is noted with ACTEMRA treatment in the PJIA and SJIA populations. Monitor neutrophils, platelets, ALT and AST at the time of the second administration and thereafter every 4 to 8 weeks for PJIA and every 2 to 4 weeks for SJIA. Monitor lipids as above for approved adult indications [see Dosage and Administration (2.9)].

**5.4 Immunosuppression**
The impact of treatment with ACTEMRA on the development of malignancies is not known but malignancies were observed in clinical studies [see Adverse Reactions (6.1)]. ACTEMRA is an immunosuppressant, and treatment with immunosuppressants may result in an increased risk of malignancies.

**5.5 Hypersensitivity Reactions, Including Anaphylaxis**
Hypersensitivity reactions, including anaphylaxis, have been reported in association with ACTEMRA [see Adverse Reactions (6)] and anaphylactic events with a fatal outcome have been reported with intravenous infusion of ACTEMRA. Anaphylaxis and other hypersensitivity reactions that required treatment discontinuation were reported in 0.1% (3 out of 2644) of patients in the 6-month controlled trials of intravenous ACTEMRA, 0.2% (8 out of 4009) of patients in the intravenous all-exposure RA population, 0.7% (8 out of 1068) in the subcutaneous 6-month controlled RA trials, and in 0.7% (10 out of 1465) of patients in the subcutaneous all-exposure population. In the SJIA controlled trial with intravenous ACTEMRA, 1 out of 112 patients (0.9%) experienced hypersensitivity reactions that required treatment discontinuation. In the PJIA controlled trial with intravenous ACTEMRA, 0 out of 188 patients (0%) in the ACTEMRA all-exposure population experienced hypersensitivity reactions that required treatment discontinuation. Reactions that required treatment discontinuation included generalized erythema, rash, and urticaria. Injection site reactions were categorized separately [see Adverse Reactions (6)].

In the postmarketing setting, events of hypersensitivity reactions, including anaphylaxis and death have occurred in patients treated with a range of doses of intravenous ACTEMRA, with or without concomitant therapies. Events have occurred in patients who received premedication. Hypersensitivity, including anaphylaxis events, have occurred both with and without previous hypersensitivity reactions and as early as the first infusion of ACTEMRA [see Adverse Reactions (6.6)]. ACTEMRA for intravenous use should only be infused by a healthcare professional with appropriate medical support to manage anaphylaxis. For ACTEMRA subcutaneous injection, advise patients to seek immediate medical attention if they experience any symptoms of a hypersensitivity reaction. If anaphylaxis or other hypersensitivity reaction occurs, stop administration of ACTEMRA immediately and discontinue ACTEMRA permanently. Do not administer ACTEMRA to patients with known hypersensitivity to ACTEMRA [see Contraindications (4) and Adverse Reactions (6)].

**5.6 Demyelinating Disorders**
The impact of treatment with ACTEMRA on demyelinating disorders is not known, but multiple sclerosis and chronic inflammatory demyelinating polyneuropathy were reported rarely in RA clinical studies. Monitor patients
for signs and symptoms potentially indicative of demyelinating disorders. Prescribers should exercise caution in considering the use of ACTEMRA in patients with preexisting or recent onset demyelinating disorders.

5.7 Active Hepatic Disease and Hepatic Impairment
Treatment with ACTEMRA is not recommended in patients with active hepatic disease or hepatic impairment [see Adverse Reactions (6.1), Use in Specific Populations (8.6)].

5.8 Vaccinations
Avoid use of live vaccines concurrently with ACTEMRA as clinical safety has not been established. No data are available on the secondary transmission of infection from persons receiving live vaccines to patients receiving ACTEMRA.

No data are available on the effectiveness of vaccination in patients receiving ACTEMRA. Because IL-6 inhibition may interfere with the normal immune response to new antigens, it is recommended that all patients, particularly pediatric or elderly patients, if possible, be brought up to date with all immunizations in agreement with current immunization guidelines prior to initiating ACTEMRA therapy. The interval between live vaccinations and initiation of ACTEMRA therapy should be in accordance with current vaccination guidelines regarding immunosuppressive agents.

6 ADVERSE REACTIONS
The following serious adverse reactions are described elsewhere in labeling:

- Serious Infections [see Warnings and Precautions (5.1)]
- Gastrointestinal Perforations [see Warnings and Precautions (5.2)]
- Laboratory Parameters [see Warnings and Precautions (5.3)]
- Immunosuppression [see Warnings and Precautions (5.4)]
- Hypersensitivity Reactions, Including Anaphylaxis [see Warnings and Precautions (5.5)]
- Demyelinating Disorders [see Warnings and Precautions (5.6)]
- Active Hepatic Disease and Hepatic Impairment [see Warnings and Precautions (5.7)]

Because clinical studies are conducted under widely varying conditions, adverse reaction rates observed in the clinical studies of a drug cannot be directly compared to rates in the clinical studies of another drug and may not predict the rates observed in a broader patient population in clinical practice.

6.1 Clinical Trials Experience in Rheumatoid Arthritis Patients Treated with Intravenous ACTEMRA (ACTEMRA-IV)
The ACTEMRA-IV data in rheumatoid arthritis (RA) includes 5 double-blind, controlled, multicenter studies. In these studies, patients received doses of ACTEMRA-IV 8 mg per kg monotherapy (288 patients), ACTEMRA-IV 8 mg per kg in combination with DMARDs (including methotrexate) (1582 patients), or ACTEMRA-IV 4 mg per kg in combination with methotrexate (774 patients).

The all exposure population includes all patients in registration studies who received at least one dose of ACTEMRA-IV. Of the 4009 patients in this population, 3577 received treatment for at least 6 months, 3309 for at least one year; 2954 received treatment for at least 2 years and 2189 for 3 years.

All patients in these studies had moderately to severely active rheumatoid arthritis. The study population had a mean age of 52 years, 82% were female and 74% were Caucasian.

The most common serious adverse reactions were serious infections [see Warnings and Precautions (5.1)]. The most commonly reported adverse reactions in controlled studies up to 24 weeks (occurring in at least 5% of
patients treated with ACTEMRA-IV monotherapy or in combination with DMARDs) were upper respiratory tract infections, nasopharyngitis, headache, hypertension and increased ALT.

The proportion of patients who discontinued treatment due to any adverse reactions during the double-blind, placebo-controlled studies was 5% for patients taking ACTEMRA-IV and 3% for placebo-treated patients. The most common adverse reactions that required discontinuation of ACTEMRA-IV were increased hepatic transaminase values (per protocol requirement) and serious infections.

**Overall Infections**

In the 24 week, controlled clinical studies, the rate of infections in the ACTEMRA-IV monotherapy group was 119 events per 100 patient-years and was similar in the methotrexate monotherapy group. The rate of infections in the 4 mg per kg and 8 mg per kg ACTEMRA-IV plus DMARD group was 133 and 127 events per 100 patient-years, respectively, compared to 112 events per 100 patient-years in the placebo plus DMARD group. The most commonly reported infections (5% to 8% of patients) were upper respiratory tract infections and nasopharyngitis.

The overall rate of infections with ACTEMRA-IV in the all exposure population remained consistent with rates in the controlled periods of the studies.

**Serious Infections**

In the 24 week, controlled clinical studies, the rate of serious infections in the ACTEMRA-IV monotherapy group was 3.6 per 100 patient-years compared to 1.5 per 100 patient-years in the methotrexate group. The rate of serious infections in the 4 mg per kg and 8 mg per kg ACTEMRA-IV plus DMARD group was 4.4 and 5.3 events per 100 patient-years, respectively, compared to 3.9 events per 100 patient-years in the placebo plus DMARD group.

In the all-exposure population, the overall rate of serious infections remained consistent with rates in the controlled periods of the studies. The most common serious infections included pneumonia, urinary tract infection, cellulitis, herpes zoster, gastroenteritis, diverticulitis, sepsis and bacterial arthritis. Cases of opportunistic infections have been reported [see Warnings and Precautions (5.1)].

**Gastrointestinal Perforations**

During the 24 week, controlled clinical trials, the overall rate of gastrointestinal perforation was 0.26 events per 100 patient-years with ACTEMRA-IV therapy.

In the all-exposure population, the overall rate of gastrointestinal perforation remained consistent with rates in the controlled periods of the studies. Reports of gastrointestinal perforation were primarily reported as complications of diverticulitis including generalized purulent peritonitis, lower GI perforation, fistula and abscess. Most patients who developed gastrointestinal perforations were taking concomitant nonsteroidal anti-inflammatory medications (NSAIDs), corticosteroids, or methotrexate [see Warnings and Precautions (5.2)]. The relative contribution of these concomitant medications versus ACTEMRA-IV to the development of GI perforations is not known.

**Infusion Reactions**

In the 24 week, controlled clinical studies, adverse events associated with the infusion (occurring during or within 24 hours of the start of infusion) were reported in 8% and 7% of patients in the 4 mg per kg and 8 mg per kg ACTEMRA-IV plus DMARD group, respectively, compared to 5% of patients in the placebo plus DMARD group. The most frequently reported event on the 4 mg per kg and 8 mg per kg dose during the infusion was hypertension (1% for both doses), while the most frequently reported event occurring within 24 hours of finishing an infusion were headache (1% for both doses) and skin reactions (1% for both doses), including rash, pruritus and urticaria. These events were not treatment limiting.

**Anaphylaxis**

Hypersensitivity reactions requiring treatment discontinuation, including anaphylaxis, associated with ACTEMRA-IV were reported in 0.1% (3 out of 2644) in the 24 week, controlled trials and in 0.2% (8 out of 4009) in the all-exposure population. These reactions were generally observed during the second to fourth infusion of
ACTEMRA-IV. Appropriate medical treatment should be available for immediate use in the event of a serious hypersensitivity reaction [see Warnings and Precautions (5.5)].

**Laboratory Abnormalities**

**Neutropenia**

In the 24 week, controlled clinical studies, decreases in neutrophil counts below 1000 per mm$^3$ occurred in 1.8% and 3.4% of patients in the 4 mg per kg and 8 mg per kg ACTEMRA-IV plus DMARD group, respectively, compared to 0.1% of patients in the placebo plus DMARD group. Approximately half of the instances of ANC below 1000 per mm$^3$ occurred within 8 weeks of starting therapy. Decreases in neutrophil counts below 500 per mm$^3$ occurred in 0.4% and 0.3% of patients in the 4 mg per kg and 8 mg per kg ACTEMRA-IV plus DMARD, respectively, compared to 0.1% of patients in the placebo plus DMARD group. There was no clear relationship between decreases in neutrophils below 1000 per mm$^3$ and the occurrence of serious infections.

In the all-exposure population, the pattern and incidence of decreases in neutrophil counts remained consistent with what was seen in the 24 week controlled clinical studies [see Warnings and Precautions (5.3)].

**Thrombocytopenia**

In the 24 week, controlled clinical studies, decreases in platelet counts below 100,000 per mm$^3$ occurred in 1.3% and 1.7% of patients on 4 mg per kg and 8 mg per kg ACTEMRA-IV plus DMARD, respectively, compared to 0.5% of patients on placebo plus DMARD, without associated bleeding events.

In the all-exposure population, the pattern and incidence of decreases in platelet counts remained consistent with what was seen in the 24 week controlled clinical studies [see Warnings and Precautions (5.3)].

**Elevated Liver Enzymes**

Liver enzyme abnormalities are summarized in **Table 1**. In patients experiencing liver enzyme elevation, modification of treatment regimen, such as reduction in the dose of concomitant DMARD, interruption of ACTEMRA-IV, or reduction in ACTEMRA-IV dose, resulted in decrease or normalization of liver enzymes [see Dosage and Administration (2.6)]. These elevations were not associated with clinically relevant increases in direct bilirubin, nor were they associated with clinical evidence of hepatitis or hepatic insufficiency [see Warnings and Precautions (5.3)].

**Table 1 Incidence of Liver Enzyme Abnormalities in the 24 Week Controlled Period of Studies I to V***

<table>
<thead>
<tr>
<th></th>
<th>ACTEMRA 8 mg per kg MONOTHERAPY</th>
<th>Methotrexate</th>
<th>ACTEMRA 4 mg per kg + DMARDs</th>
<th>ACTEMRA 8 mg per kg + DMARDs</th>
<th>Placebo + DMARDs</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N = 288 (%)</td>
<td>N = 284 (%)</td>
<td>N = 774 (%)</td>
<td>N = 1582 (%)</td>
<td>N = 1170 (%)</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt; ULN to 3x ULN</td>
<td>22</td>
<td>26</td>
<td>34</td>
<td>41</td>
<td>17</td>
</tr>
<tr>
<td>&gt; 3x ULN to 5x ULN</td>
<td>0.3</td>
<td>2</td>
<td>1</td>
<td>2</td>
<td>0.3</td>
</tr>
<tr>
<td>&gt; 5x ULN</td>
<td>0.7</td>
<td>0.4</td>
<td>0.1</td>
<td>0.2</td>
<td>&lt; 0.1</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt; ULN to 3x ULN</td>
<td>36</td>
<td>33</td>
<td>45</td>
<td>48</td>
<td>23</td>
</tr>
<tr>
<td>&gt; 3x ULN to 5x ULN</td>
<td>1</td>
<td>4</td>
<td>5</td>
<td>5</td>
<td>1</td>
</tr>
<tr>
<td>&gt; 5x ULN</td>
<td>0.7</td>
<td>1</td>
<td>1.3</td>
<td>1.5</td>
<td>0.3</td>
</tr>
</tbody>
</table>

ULN = Upper Limit of Normal

*For a description of these studies, see Section 14, Clinical Studies.

In the all-exposure population, the elevations in ALT and AST remained consistent with what was seen in the 24 week, controlled clinical trials.
Lipids
Elevations in lipid parameters (total cholesterol, LDL, HDL, triglycerides) were first assessed at 6 weeks following initiation of ACTEMRA-IV in the controlled 24 week clinical trials. Increases were observed at this time point and remained stable thereafter. Increases in triglycerides to levels above 500 mg per dL were rarely observed. Changes in other lipid parameters from baseline to week 24 were evaluated and are summarized below:

- Mean LDL increased by 13 mg per dL in the ACTEMRA 4 mg per kg+DMARD arm, 20 mg per dL in the ACTEMRA 8 mg per kg+DMARD, and 25 mg per dL in ACTEMRA 8 mg per kg monotherapy.
- Mean HDL increased by 3 mg per dL in the ACTEMRA 4 mg per kg+DMARD arm, 5 mg per dL in the ACTEMRA 8 mg per kg+DMARD, and 4 mg per dL in ACTEMRA 8 mg per kg monotherapy.
- Mean LDL/HDL ratio increased by an average of 0.14 in the ACTEMRA 4 mg per kg+DMARD arm, 0.15 in the ACTEMRA 8 mg per kg+DMARD, and 0.26 in ACTEMRA 8 mg per kg monotherapy.
- ApoB/ApoA1 ratios were essentially unchanged in ACTEMRA-treated patients.

Elevated lipids responded to lipid lowering agents.

In the all-exposure population, the elevations in lipid parameters remained consistent with what was seen in the 24 week, controlled clinical trials.

Immunogenicity
As with all therapeutic proteins, there is potential for immunogenicity. The detection of antibody formation is highly dependent on the sensitivity and specificity of the assay. Additionally, the observed incidence of antibody (including neutralizing antibody) positivity in an assay may be influenced by several factors including assay methodology, sample handling, timing of sample collection, concomitant medications, and underlying disease. For these reasons, comparison of the incidence of antibodies to tocilizumab in the studies described below with the incidence of antibodies in other studies or to other products may be misleading.

In the 24 week, controlled clinical studies, a total of 2876 patients have been tested for anti-tocilizumab antibodies. Forty-six patients (2%) developed positive anti-tocilizumab antibodies, of whom 5 had an associated, medically significant, hypersensitivity reaction leading to withdrawal. Thirty patients (1%) developed neutralizing antibodies.

Malignancies
During the 24 week, controlled period of the studies, 15 malignancies were diagnosed in patients receiving ACTEMRA-IV, compared to 8 malignancies in patients in the control groups. Exposure-adjusted incidence was similar in the ACTEMRA-IV groups (1.32 events per 100 patient-years) and in the placebo plus DMARD group (1.37 events per 100 patient-years).

In the all-exposure population, the rate of malignancies remained consistent with the rate observed in the 24 week, controlled period [see Warnings and Precautions (5.4)].

Other Adverse Reactions
Adverse reactions occurring in 2% or more of patients on 4 or 8 mg per kg ACTEMRA-IV plus DMARD and at least 1% greater than that observed in patients on placebo plus DMARD are summarized in Table 2.

Table 2  Adverse Reactions Occurring in at Least 2% or More of Patients on 4 or 8 mg per kg ACTEMRA plus DMARD and at Least 1% Greater Than That Observed in Patients on Placebo plus DMARD

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>ACTEMRA</th>
<th>Methotrexate</th>
<th>ACTEMRA</th>
<th>ACTEMRA</th>
<th>Placebo + DMARDs</th>
</tr>
</thead>
<tbody>
<tr>
<td>24 Week Phase 3 Controlled Study Population</td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>
Other infrequent and medically relevant adverse reactions occurring at an incidence less than 2% in rheumatoid arthritis patients treated with ACTEMRA-IV in controlled trials were:

**Infections and Infestations:** oral herpes simplex  
**Gastrointestinal disorders:** stomatitis, gastric ulcer  
**Investigations:** weight increased, total bilirubin increased  
**Blood and lymphatic system disorders:** leukopenia  
**General disorders and administration site conditions:** edema peripheral  
**Respiratory, thoracic, and mediastinal disorders:** dyspnea, cough  
**Eye disorders:** conjunctivitis  
**Renal disorders:** nephrolithiasis  
**Endocrine disorders:** hypothyroidism

### 6.2 Clinical Trials Experience in Rheumatoid Arthritis Patients Treated with Subcutaneous ACTEMRA (ACTEMRA-SC)

The ACTEMRA-SC data in rheumatoid arthritis (RA) includes 2 double-blind, controlled, multicenter studies. Study SC-I was a non-inferiority study that compared the efficacy and safety of tocilizumab 162 mg administered every week subcutaneously and 8 mg/kg intravenously every four weeks in 1262 adult subjects with rheumatoid arthritis. Study SC-II was a placebo controlled superiority study that evaluated the safety and efficacy of tocilizumab 162 mg administered every other week subcutaneously or placebo in 656 patients. All patients in both studies received background non-biologic DMARDs.

The safety observed for ACTEMRA-SC administered subcutaneously was consistent with the known safety profile of intravenous ACTEMRA, with the exception of injection site reactions (ISRs), which were more common with ACTEMRA-SC compared with placebo SC injections (IV arm).

**Injection Site Reactions**  
In the 6-month control period, in SC-I, the frequency of ISRs was 10.1% (64/631) and 2.4% (15/631) for the weekly ACTEMRA-SC and placebo SC (IV-arm) groups, respectively. In SC-II, the frequency of ISRs was 7.1% (31/437) and 4.1% (9/218) for the every other week ACTEMRA-SC and placebo groups, respectively. These ISRs (including erythema, pruritus, pain and hematoma) were mild to moderate in severity. The majority resolved without any treatment and none necessitated drug discontinuation.

**Immunogenicity**  
In the 6-month control period in SC-I, 0.8% (5/625) in the ACTEMRA-SC arm and 0.8% (5/627) in the IV arm developed anti-tocilizumab antibodies; of these, all developed neutralizing antibodies. In SC-II, 1.6% (7/434) in
the ACTEMRA-SC arm compared with 1.4% (3/217) in the placebo arm developed anti-tocilizumab antibodies; of these, 1.4% (6/434) in the ACTEMRA-SC arm and 0.5% (1/217) in the placebo arm also developed neutralizing antibodies.

A total of 1454 (>99%) patients who received ACTEMRA-SC in the all exposure group have been tested for anti-tocilizumab antibodies. Thirteen patients (0.9%) developed anti-tocilizumab antibodies, and, of these, 12 patients (0.8%) developed neutralizing antibodies.

The rate is consistent with previous intravenous experience. No correlation of antibody development to adverse events or loss of clinical response was observed.

**Laboratory Abnormalities**

**Neutropenia**

During routine laboratory monitoring in the 6-month controlled clinical trials, a decrease in neutrophil count below 1 × 10⁹/L occurred in 2.9% and 3.7% of patients receiving ACTEMRA-SC weekly and every other week, respectively.

There was no clear relationship between decreases in neutrophils below 1 x 10⁹/L and the occurrence of serious infections.

**Thrombocytopenia**

During routine laboratory monitoring in the ACTEMRA-SC 6-month controlled clinical trials, none of the patients had a decrease in platelet count to ≤50,000/mm³.

**Elevated Liver Enzymes**

During routine laboratory monitoring in the 6-month controlled clinical trials, elevation in ALT or AST ≥3 x ULN occurred in 6.5% and 1.4% of patients, respectively, receiving ACTEMRA-SC weekly and 3.4% and 0.7% receiving ACTEMRA-SC every other week.

**Lipid Parameters Elevations**

During routine laboratory monitoring in the ACTEMRA-SC 6-month clinical trials, 19% of patients dosed weekly and 19.6% of patients dosed every other week and 10.2% of patients on placebo experienced sustained elevations in total cholesterol > 6.2 mmol/l (240 mg/dL), with 9%, 10.4% and 5.1% experiencing a sustained increase in LDL to 4.1 mmol/l (160 mg/dL) receiving ACTEMRA-SC weekly, every other week and placebo, respectively.

### 6.3 Clinical Trials Experience in Giant Cell Arteritis Patients Treated with Subcutaneous ACTEMRA (ACTEMRA-SC)

The safety of subcutaneous ACTEMRA (tocilizumab) has been studied in one Phase III study (WA28119) with 251 GCA patients. The total patient years duration in the ACTEMRA GCA all exposure population was 138.5 patient years during the 12-month double blind, placebo-controlled phase of the study. The overall safety profile observed in the ACTEMRA treatment groups was generally consistent with the known safety profile of ACTEMRA. There was an overall higher incidence of infections in GCA patients relative to RA patients. The rate of infection/serious infection events was 200.2/9.7 events per 100 patient years in the ACTEMRA weekly group and 160.2/4.4 events per 100 patient years in the ACTEMRA every other week group as compared to 156.0/4.2 events per 100 patient years in the placebo + 26 week prednisone taper and 210.2/12.5 events per 100 patient years in the placebo + 52 week taper groups.

### 6.4 Clinical Trials Experience in Polyarticular Juvenile Idiopathic Arthritis Patients Treated With Intravenous ACTEMRA (ACTEMRA-IV)

The safety of ACTEMRA-IV was studied in 188 pediatric patients 2 to 17 years of age with PJIA who had an inadequate clinical response or were intolerant to methotrexate. The total patient exposure in the ACTEMRA-IV all exposure population (defined as patients who received at least one dose of ACTEMRA-IV) was 184.4 patient years. At baseline, approximately half of the patients were taking oral corticosteroids and almost 80% were taking
methotrexate. In general, the types of adverse drug reactions in patients with PJIA were consistent with those seen in RA and SJIA patients [see Adverse Reactions (6.1 and 6.6)].

**Infections**
The rate of infections in the ACTEMRA-IV all exposure population was 163.7 per 100 patient years. The most common events observed were nasopharyngitis and upper respiratory tract infections. The rate of serious infections was numerically higher in patients weighing less than 30 kg treated with 10 mg/kg tocilizumab (12.2 per 100 patient years) compared to patients weighing at or above 30 kg, treated with 8 mg/kg tocilizumab (4.0 per 100 patient years). The incidence of infections leading to dose interruptions was also numerically higher in patients weighing less than 30 kg treated with 10 mg/kg tocilizumab (12.2 per 100 patient years) compared to patients weighing at or above 30 kg, treated with 8 mg/kg tocilizumab (4.0 per 100 patient years). The incidence of infections leading to dose interruptions was also numerically higher in patients weighing less than 30 kg treated with 10 mg/kg tocilizumab (21%) compared to patients weighing at or above 30 kg, treated with 8 mg/kg tocilizumab (8%).

**Infusion Reactions**
In PJIA patients, infusion-related reactions are defined as all events occurring during or within 24 hours of an infusion. In the ACTEMRA-IV all exposure population, 11 patients (6%) experienced an event during the infusion, and 38 patients (20.2%) experienced an event within 24 hours of an infusion. The most common events occurring during infusion were headache, nausea and hypotension, and occurring within 24 hours of infusion were dizziness and hypotension. In general, the adverse drug reactions observed during or within 24 hours of an infusion were similar in nature to those seen in RA and SJIA patients [see Adverse Reactions (6.1 and 6.6)].

No clinically significant hypersensitivity reactions associated with tocilizumab and requiring treatment discontinuation were reported.

**Immunogenicity**
One patient, in the 10 mg/kg less than 30 kg group, developed positive anti-tocilizumab antibodies without developing a hypersensitivity reaction and subsequently withdrew from the study.

**Laboratory Abnormalities**

**Neutropenia**
During routine laboratory monitoring in the ACTEMRA-IV all exposure population, a decrease in neutrophil counts below $1 \times 10^9$ per L occurred in 3.7% of patients.

There was no clear relationship between decreases in neutrophils below $1 \times 10^9$ per L and the occurrence of serious infections.

**Thrombocytopenia**
During routine laboratory monitoring in the ACTEMRA-IV all exposure population, 1% of patients had a decrease in platelet count at or less than 50,000 per mm$^3$ without associated bleeding events.

**Elevated Liver Enzymes**
During routine laboratory monitoring in the ACTEMRA-IV all exposure population, elevation in ALT or AST at or greater than 3 x ULN occurred in 4% and less than 1% of patients, respectively.

**Lipids**
During routine laboratory monitoring in the tocilizumab all exposure population, elevation in total cholesterol greater than 1.5-2 x ULN occurred in one patient (0.5%) and elevation in LDL greater than 1.5-2 x ULN occurred in one patient (0.5%).

6.5 Clinical Trials Experience in Polyarticular Juvenile Idiopathic Arthritis Patients Treated With Subcutaneous ACTEMRA (ACTEMRA-SC)
The safety of ACTEMRA-SC was studied in 52 pediatric patients 1 to 17 years of age with PJIA who had an inadequate clinical response or were intolerant to methotrexate. The total patient exposure in the PJIA ACTEMRA-SC population (defined as patients who received at least one dose of ACTEMRA-SC and accounting for treatment discontinuation) was 49.5 patient years. In general, the safety observed for ACTEMRA administered
subcutaneously was consistent with the known safety profile of intravenous ACTEMRA, with the exception of injection site reactions (ISRs), and neutropenia.

**Injection Site Reactions**

During the 1-year study, a frequency of 28.8% (15/52) ISRs was observed in ACTEMRA-SC treated PJIA patients. These ISRs occurred in a greater proportion of patients at or above 30 kg (44.0%) compared with patients below 30 kg (14.8%). All ISRs were mild in severity and none of the ISRs required patient withdrawal from treatment or dose interruption. A higher frequency of ISRs was observed in ACTEMRA-SC treated PJIA patients compared to what was seen in adult RA or GCA patients [see Adverse Reactions (6.1 and 6.3)].

**Immunogenicity**

Three patients, 1 patient below 30 kg and 2 patients at or above 30 kg, developed positive anti-tocilizumab antibodies with neutralizing potential without developing a serious or clinically significant hypersensitivity reaction. One patient subsequently withdrew from the study.

**Neutropenia**

During routine laboratory monitoring in the ACTEMRA-SC all exposure population, a decrease in neutrophil counts below $1 \times 10^9$ per L occurred in 15.4% of patients, and was more frequently observed in the patients less than 30 kg (25.9%) compared to patients at or above 30 kg (4.0%). There was no clear relationship between decreases in neutrophils below $1 \times 10^9$ per L and the occurrence of serious infections.

6.6 **Clinical Trials Experience in Systemic Juvenile Idiopathic Arthritis Patients Treated with Intravenous ACTEMRA (ACTEMRA-IV)**

The data described below reflect exposure to ACTEMRA-IV in one randomized, double-blind, placebo-controlled trial of 112 pediatric patients with SJIA 2 to 17 years of age who had an inadequate clinical response to nonsteroidal anti-inflammatory drugs (NSAIDs) or corticosteroids due to toxicity or lack of efficacy. At baseline, approximately half of the patients were taking 0.3 mg/kg/day corticosteroids or more, and almost 70% were taking methotrexate. The trial included a 12 week controlled phase followed by an open-label extension. In the 12 week double-blind, controlled portion of the clinical study 75 patients received treatment with ACTEMRA-IV (8 or 12 mg per kg based upon body weight). After 12 weeks or at the time of escape, due to disease worsening, patients were treated with ACTEMRA-IV in the open-label extension phase.

The most common adverse events (at least 5%) seen in ACTEMRA-IV treated patients in the 12 week controlled portion of the study were: upper respiratory tract infection, headache, nasopharyngitis and diarrhea.

**Infections**

In the 12 week controlled phase, the rate of all infections in the ACTEMRA-IV group was 345 per 100 patient-years and 287 per 100 patient-years in the placebo group. In the open label extension over an average duration of 73 weeks of treatment, the overall rate of infections was 304 per 100 patient-years.

In the 12 week controlled phase, the rate of serious infections in the ACTEMRA-IV group was 11.5 per 100 patient years. In the open label extension over an average duration of 73 weeks of treatment, the overall rate of serious infections was 11.4 per 100 patient years. The most commonly reported serious infections included pneumonia, gastroenteritis, varicella, and otitis media.

**Macrophage Activation Syndrome**

In the 12 week controlled study, no patient in any treatment group experienced macrophage activation syndrome (MAS) while on assigned treatment; 3 per 112 (3%) developed MAS during open-label treatment with ACTEMRA-IV. One patient in the placebo group escaped to ACTEMRA-IV 12 mg per kg at Week 2 due to severe disease activity, and ultimately developed MAS at Day 70. Two additional patients developed MAS during the long-term extension. All 3 patients had ACTEMRA-IV dose interrupted (2 patients) or discontinued (1 patient) for the MAS event, received treatment, and the MAS resolved without sequelae. Based on a limited number of
cases, the incidence of MAS does not appear to be elevated in the ACTEMRA-IV SJIA clinical development experience; however no definitive conclusions can be made.

**Infusion Reactions**

Patients were not premedicated, however most patients were on concomitant corticosteroids as part of their background treatment for SJIA. Infusion related reactions were defined as all events occurring during or within 24 hours after an infusion. In the 12 week controlled phase, 4% of ACTEMRA-IV and 0% of placebo treated patients experienced events occurring during infusion. One event (angioedema) was considered serious and life-threatening, and the patient was discontinued from study treatment.

Within 24 hours after infusion, 16% of patients in the ACTEMRA-IV treatment group and 5% of patients in the placebo group experienced an event. In the ACTEMRA-IV group the events included rash, urticaria, diarrhea, epigastric discomfort, arthralgia and headache. One of these events, urticaria, was considered serious.

**Anaphylaxis**

Anaphylaxis was reported in 1 out of 112 patients (less than 1%) treated with ACTEMRA-IV during the controlled and open label extension study [see Warnings and Precautions (5.5)].

**Immunogenicity**

All 112 patients were tested for anti-tocilizumab antibodies at baseline. Two patients developed positive anti-tocilizumab antibodies: one of these patients experienced serious adverse events of urticaria and angioedema consistent with an anaphylactic reaction which led to withdrawal; the other patient developed macrophage activation syndrome while on escape therapy and was discontinued from the study.

**Laboratory Abnormalities**

**Neutropenia**

During routine monitoring in the 12 week controlled phase, a decrease in neutrophil below $1 \times 10^9$ per L occurred in 7% of patients in the ACTEMRA-IV group, and in no patients in the placebo group. In the open label extension over an average duration of 73 weeks of treatment, a decreased neutrophil count occurred in 17% of the ACTEMRA-IV group. There was no clear relationship between decrease in neutrophils below $1 \times 10^9$ per L and the occurrence of serious infections.

**Thrombocytopenia**

During routine monitoring in the 12 week controlled phase, 1% of patients in the ACTEMRA-IV group and 3% in the placebo group had a decrease in platelet count to no more than 100,000 per mm$^3$.

In the open label extension over an average duration of 73 weeks of treatment, decreased platelet count occurred in 4% of patients in the ACTEMRA-IV group, with no associated bleeding.

**Elevated Liver Enzymes**

During routine laboratory monitoring in the 12 week controlled phase, elevation in ALT or AST at or above 3x ULN occurred in 5% and 3% of patients, respectively in the ACTEMRA-IV group and in 0% of placebo patients.

In the open label extension over an average duration of 73 weeks of treatment, the elevation in ALT or AST at or above 3x ULN occurred in 13% and 5% of ACTEMRA-IV treated patients, respectively.

**Lipids**

During routine laboratory monitoring in the 12 week controlled phase, elevation in total cholesterol greater than 1.5x ULN – 2x ULN occurred in 1.5% of the ACTEMRA-IV group and in 0% of placebo patients. Elevation in LDL greater than 1.5x ULN – 2x ULN occurred in 1.9% of patients in the ACTEMRA-IV group and 0% of the placebo group.

In the open label extension study over an average duration of 73 weeks of treatment, the pattern and incidence of elevations in lipid parameters remained consistent with the 12 week controlled study data.
Clinical Trials Experience in Systemic Juvenile Idiopathic Arthritis Patients Treated with Subcutaneous ACTEMRA (ACTEMRA-SC)
The safety profile of ACTEMRA-SC was studied in 51 pediatric patients 1 to 17 years of age with SJIA who had an inadequate clinical response to NSAIDs and corticosteroids. In general, the safety observed for ACTEMRA administered subcutaneously was consistent with the known safety profile of intravenous ACTEMRA, with the exception of ISRs where a higher frequency was observed in ACTEMRA-SC treated SJIA patients compared to PJIA patients and adult RA or GCA patients [see Adverse Reactions (6.2 and 6.3)].

Injection Site Reactions (ISRs)
A total of 41.2% (21/51) SJIA patients experienced ISRs to ACTEMRA-SC. The most common ISRs were erythema, pruritus, pain, and swelling at the injection site. The majority of ISRs reported were Grade 1 events and all ISRs reported were non-serious and none required patient withdrawal from treatment or dose interruption.

Immunogenicity
Forty-six of the 51 (90.2%) patients who were tested for anti-tocilizumab antibodies at baseline had at least one post-baseline screening assay result. No patient developed positive anti-tocilizumab antibodies post-baseline.

Clinical Trials Experience in Patients with Cytokine Release Syndrome Treated with Intravenous ACTEMRA (ACTEMRA-IV)
In a retrospective analysis of pooled outcome data from multiple clinical trials 45 patients were treated with tocilizumab 8 mg/kg (12 mg/kg for patients less than 30 kg) with or without additional high-dose corticosteroids for severe or life-threatening CAR T-cell-induced CRS. A median of 1 dose of tocilizumab (range, 1-4 doses) was administered. No adverse reactions related to tocilizumab were reported [see Clinical Studies (14.7)].

Postmarketing Experience
The following adverse reactions have been identified during post-approval use of ACTEMRA. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

- Fatal anaphylaxis [see Warnings and Precautions (5.5)]
- Stevens-Johnson Syndrome
- Pancreatitis

7 DRUG INTERACTIONS
7.1 Concomitant Drugs for Treatment of Adult Indications
In RA patients, population pharmacokinetic analyses did not detect any effect of methotrexate (MTX), non-steroidal anti-inflammatory drugs or corticosteroids on tocilizumab clearance. Concomitant administration of a single intravenous dose of 10 mg/kg ACTEMRA with 10-25 mg MTX once weekly had no clinically significant effect on MTX exposure. ACTEMRA has not been studied in combination with biological DMARDs such as TNF antagonists [see Dosage and Administration (2.1)].

In GCA patients, no effect of concomitant corticosteroid on tocilizumab exposure was observed.

7.2 Interactions with CYP450 Substrates
Cytochrome P450s in the liver are down-regulated by infection and inflammation stimuli including cytokines such as IL-6. Inhibition of IL-6 signaling in RA patients treated with tocilizumab may restore CYP450 activities to higher levels than those in the absence of tocilizumab leading to increased metabolism of drugs that are CYP450 substrates. In vitro studies showed that tocilizumab has the potential to affect expression of multiple CYP enzymes including CYP1A2, CYP2B6, CYP2C9, CYP2C19, CYP2D6 and CYP3A4. Its effect on CYP2C8 or transporters is unknown. In vivo studies with omeprazole, metabolized by CYP2C19 and CYP3A4, and
simvastatin, metabolized by CYP3A4, showed up to a 28% and 57% decrease in exposure one week following a single dose of ACTEMRA, respectively. The effect of tocilizumab on CYP enzymes may be clinically relevant for CYP450 substrates with narrow therapeutic index, where the dose is individually adjusted. Upon initiation or discontinuation of ACTEMRA, in patients being treated with these types of medicinal products, perform therapeutic monitoring of effect (e.g., warfarin) or drug concentration (e.g., cyclosporine or theophylline) and the individual dose of the medicinal product adjusted as needed. Exercise caution when coadministering ACTEMRA with CYP3A4 substrate drugs where decrease in effectiveness is undesirable, e.g., oral contraceptives, lovastatin, atorvastatin, etc. The effect of tocilizumab on CYP450 enzyme activity may persist for several weeks after stopping therapy [see Clinical Pharmacology (12.3)].

7.3 Live Vaccines
Avoid use of live vaccines concurrently with ACTEMRA [see Warnings and Precautions (5.8)].

8 USE IN SPECIFIC POPULATIONS
8.1 Pregnancy
Pregnancy Exposure Registry

There is a pregnancy exposure registry that monitors pregnancy outcomes in women exposed to ACTEMRA during pregnancy. Physicians are encouraged to register patients and pregnant women are encouraged to register themselves by calling 1-877-311-8972.

Risk Summary

The limited available data with ACTEMRA in pregnant women are not sufficient to determine whether there is a drug-associated risk for major birth defects and miscarriage. Monoclonal antibodies, such as tocilizumab, are actively transported across the placenta during the third trimester of pregnancy and may affect immune response in the in utero exposed infant [see Clinical Considerations]. In animal reproduction studies, intravenous administration of tocilizumab to Cynomolgus monkeys during organogenesis caused abortion/embryo-fetal death at doses 1.25 times and higher than the maximum recommended human dose by the intravenous route of 8 mg per kg every 2 to 4 weeks. The literature in animals suggests that inhibition of IL-6 signaling may interfere with cervical ripening and dilatation and myometrial contractile activity leading to potential delays of parturition [see Data]. Based on the animal data, there may be a potential risk to the fetus.

The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. All pregnancies have a background risk of birth defect, loss or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2 to 4% and 15 to 20%, respectively.

Clinical Considerations
Fetal/Neonatal adverse reactions
Monoclonal antibodies are increasingly transported across the placenta as pregnancy progresses, with the largest amount transferred during the third trimester. Risks and benefits should be considered prior to administering live or live-attenuated vaccines to infants exposed to ACTEMRA in utero [see Warnings and Precautions (5.8)]

Data
Animal Data

An embryo-fetal developmental toxicity study was performed in which pregnant Cynomolgus monkeys were treated intravenously with tocilizumab at daily doses of 2, 10, or 50 mg/kg during organogenesis from gestation day (GD) 20-50. Although there was no evidence for a teratogenic/dysmorphogenic effect at any dose, tocilizumab produced an increase in the incidence of abortion/embryo-fetal death at doses 1.25 times and higher the MRHD by the intravenous route at maternal intravenous doses of 10 and 50 mg/kg. Testing of a murine analogue of tocilizumab in mice did not yield any evidence of harm to offspring during the pre- and postnatal development phase when dosed at 50 mg/kg intravenously with treatment every three days from implantation
(GD 6) until post-partum day 21 (weaning). There was no evidence for any functional impairment of the development and behavior, learning ability, immune competence and fertility of the offspring.

Parturition is associated with significant increases of IL-6 in the cervix and myometrium. The literature suggests that inhibition of IL-6 signaling may interfere with cervical ripening and dilatation and myometrial contractile activity leading to potential delays of parturition. For mice deficient in IL-6 (Il6−/− null mice), parturition was delayed relative to wild-type (Il6+/+) mice. Administration of recombinant IL-6 to Il6−/− null mice restored the normal timing of delivery.

8.2 Lactation
Risk Summary

No information is available on the presence of tocilizumab in human milk, the effects of the drug on the breastfed infant, or the effects of the drug on milk production. Maternal immunoglobulin G (IgG) is present in human milk. If tocilizumab is transferred into human milk, the effects of local exposure in the gastrointestinal tract and potential limited systemic exposure in the infant to tocilizumab are unknown. The lack of clinical data during lactation precludes clear determination of the risk of ACTEMRA to an infant during lactation; therefore the developmental and health benefits of breastfeeding should be considered along with the mother’s clinical need for ACTEMRA and the potential adverse effects on the breastfed child from tocilizumab or from the underlying maternal condition.

8.4 Pediatric Use

ACTEMRA by intravenous use is indicated for the treatment of pediatric patients with:

- Active systemic juvenile idiopathic arthritis in patients 2 years of age and older
- Active polyarticular juvenile idiopathic arthritis in patients 2 years of age and older
- Severe or life-threatening CAR T cell-induced cytokine release syndrome (CRS) in patients 2 years of age and older.

ACTEMRA by subcutaneous use is indicated for the treatment of pediatric patients with:

- Active polyarticular juvenile idiopathic arthritis in patients 2 years of age and older
- Active systemic juvenile idiopathic arthritis in patients 2 years of age and older

The safety and effectiveness of ACTEMRA in pediatric patients with conditions other than PJIA, SJIA or CRS have not been established. The safety and effectiveness in pediatric patients below the age of 2 have not been established in PJIA, SJIA, or CRS. The ability of pediatric patients to self-inject with the autoinjector has not been tested.

Systemic Juvenile Idiopathic Arthritis – Intravenous Use

A multi-center, open-label, single arm study to evaluate the PK, safety and exploratory PD and efficacy of ACTEMRA over 12-weeks in SJIA patients (N=11) under 2 years of age was conducted. Patients received intravenous ACTEMRA 12 mg/kg every two weeks. Concurrent use of stable background treatment with corticosteroids, MTX, and/or non-steroidal anti-inflammatory drugs was permitted. Patients who completed the 12-week period could continue to the optional extension period (a total of 52-weeks or until the age of 2 years, whichever was longer).

The primary PK endpoints (C max, C min and AUC2weeks) of ACTEMRA at steady-state in this study were within the ranges of these parameters observed in patients with SJIA aged 2 to 17 years.

The safety and immunogenicity of ACTEMRA for patients with SJIA under 2 years of age was assessed descriptively. SAEs, AEs leading to discontinuation, and infectious AEs were reported by 27.3%, 36.4%, and
81.8% of patients. Six patients (54.5%) experienced hypersensitivity reactions, defined as all adverse events occurring during or within 24 hours after an infusion considered related to ACTEMRA. Three of these patients experienced serious hypersensitivity reactions and were withdrawn from the study. Three patients with hypersensitivity reactions (two with serious hypersensitivity reactions) developed treatment induced anti-tocilizumab antibodies after the event. There were no cases of MAS based on the protocol-specified criteria, but 2 cases of suspected MAS based on Ravelli criteria.¹

**Cytokine Release Syndrome – Intravenous Use**

In the retrospective analysis of pooled outcome data for patients treated with ACTEMRA for CAR T cell-induced CRS, 25 patients were children (2 years up to 12 years of age), and 17 patients were adolescents (12 years up to 18 years of age). There were no differences between the pediatric patients and the adults for safety or efficacy.

**8.5 Geriatric Use**

Of the 2644 patients who received ACTEMRA in Studies I to V [see Clinical Studies (14)], a total of 435 rheumatoid arthritis patients were 65 years of age and older, including 50 patients 75 years and older. Of the 1069 patients who received ACTEMRA-SC in studies SC-I and SC-II there were 295 patients 65 years of age and older, including 41 patients 75 years and older. The frequency of serious infection among ACTEMRA treated subjects 65 years of age and older was higher than those under the age of 65. As there is a higher incidence of infections in the elderly population in general, caution should be used when treating the elderly.

Clinical studies that included ACTEMRA for CRS did not include sufficient numbers of patients aged 65 and over to determine whether they respond differently from younger patients.

**8.6 Hepatic Impairment**

The safety and efficacy of ACTEMRA have not been studied in patients with hepatic impairment, including patients with positive HBV and HCV serology [see Warnings and Precautions (5.7)].

**8.7 Renal Impairment**

No dose adjustment is required in patients with mild or moderate renal impairment. ACTEMRA has not been studied in patients with severe renal impairment [see Clinical Pharmacology (12.3)].

**9 DRUG ABUSE AND DEPENDENCE**

No studies on the potential for ACTEMRA to cause dependence have been performed. However, there is no evidence from the available data that ACTEMRA treatment results in dependence.

**10 OVERDOSAGE**

There are limited data available on overdoses with ACTEMRA. One case of accidental overdose was reported with intravenous ACTEMRA in which a patient with multiple myeloma received a dose of 40 mg per kg. No adverse drug reactions were observed. No serious adverse drug reactions were observed in healthy volunteers who received single doses of up to 28 mg per kg, although all 5 patients at the highest dose of 28 mg per kg developed dose-limiting neutropenia.

In case of an overdose, it is recommended that the patient be monitored for signs and symptoms of adverse reactions. Patients who develop adverse reactions should receive appropriate symptomatic treatment.

11 DESCRIPTION
Tocilizumab is a recombinant humanized anti-human interleukin 6 (IL-6) receptor monoclonal antibody of the immunoglobulin IgG1κ (gamma 1, kappa) subclass with a typical H2L2 polypeptide structure. Each light chain and heavy chain consists of 214 and 448 amino acids, respectively. The four polypeptide chains are linked intra- and inter-molecularly by disulfide bonds. ACTEMRA has a molecular weight of approximately 148 kDa. The antibody is produced in mammalian (Chinese hamster ovary) cells.

Intravenous Infusion

ACTEMRA (tocilizumab) injection is a sterile, clear, colorless to pale yellow, preservative-free solution for further dilution prior to intravenous infusion with a pH of approximately 6.5. Each single-dose vial, formulated with a disodium phosphate dodecahydrate/sodium dihydrogen phosphate dehydrate buffered solution, is available at a concentration of 20 mg/mL containing 80 mg/4 mL, 200 mg/10 mL, or 400 mg/20 mL of ACTEMRA. Each mL of solution contains polysorbate 80 (0.5 mg), sucrose (50 mg), and Water for Injection, USP.

Subcutaneous Injection

ACTEMRA (tocilizumab) injection is a sterile, clear, colorless to slightly yellowish, preservative-free, histidine buffered solution for subcutaneous use with a pH of approximately 6.0.

It is supplied in a ready-to-use, single-dose 0.9 mL prefilled syringe (PFS) with a needle safety device or a ready-to-use, single-dose 0.9 mL autoinjector that delivers 162 mg tocilizumab, L-arginine (0.132 mg), L-arginine hydrochloride (18.8 mg), L-histidine (1.4 mg), L-histidine hydrochloride monohydrate (1.9 mg), L-methionine (4.03 mg), polysorbate 80 (0.18 mg), and Water for Injection, USP.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Tocilizumab binds to both soluble and membrane-bound IL-6 receptors (sIL-6R and mIL-6R), and has been shown to inhibit IL-6-mediated signaling through these receptors. IL-6 is a pleiotropic pro-inflammatory cytokine produced by a variety of cell types including T- and B-cells, lymphocytes, monocytes and fibroblasts. IL-6 has been shown to be involved in diverse physiological processes such as T-cell activation, induction of immunoglobulin secretion, initiation of hepatic acute phase protein synthesis, and stimulation of hematopoietic precursor cell proliferation and differentiation. IL-6 is also produced by synovial and endothelial cells leading to local production of IL-6 in joints affected by inflammatory processes such as rheumatoid arthritis.

12.2 Pharmacodynamics

In clinical studies in RA patients with the 4 mg per kg and 8 mg per kg intravenous doses or the 162 mg weekly and every other weekly subcutaneous doses of ACTEMRA, decreases in levels of C-reactive protein (CRP) to within normal ranges were seen as early as week 2. Changes in pharmacodynamic parameters were observed (i.e., decreases in rheumatoid factor, erythrocyte sedimentation rate (ESR), serum amyloid A and increases in hemoglobin) with doses, however the greatest improvements were observed with 8 mg per kg ACTEMRA. Pharmacodynamic changes were also observed to occur after ACTEMRA administration in GCA, PJIA, and SJIA patients (decreases in CRP, ESR, and increases in hemoglobin). The relationship between these pharmacodynamic findings and clinical efficacy is not known.

In healthy subjects administered ACTEMRA in doses from 2 to 28 mg per kg intravenously and 81 to 162 mg subcutaneously, absolute neutrophil counts decreased to the nadir 3 to 5 days following ACTEMRA administration. Thereafter, neutrophils recovered towards baseline in a dose dependent manner. Rheumatoid arthritis and GCA patients demonstrated a similar pattern of absolute neutrophil counts following ACTEMRA administration [see Warnings and Precautions (5.3)].
12.3 Pharmacokinetics

*Rheumatoid Arthritis—Intravenous Administration*

The pharmacokinetics characterized in healthy subjects and RA patients suggested that PK is similar between the two populations. The clearance (CL) of tocilizumab decreased with increased doses. At the 10 mg per kg single dose in RA patients, mean CL was 0.29 ± 0.10 mL per hr per kg and mean apparent terminal t1/2 was 151 ± 59 hours (6.3 days).

The pharmacokinetics of tocilizumab were determined using a population pharmacokinetic analysis of 1793 rheumatoid arthritis patients treated with ACTEMRA 4 and 8 mg per kg every 4 weeks for 24 weeks.

The pharmacokinetic parameters of tocilizumab did not change with time. A more than dose-proportional increase in area under the curve (AUC) and trough concentration (C_min) was observed for doses of 4 and 8 mg per kg every 4 weeks. Maximum concentration (C_max) increased dose-proportionally. At steady-state, estimated AUC and C_min were 2.7 and 6.5-fold higher at 8 mg per kg as compared to 4 mg per kg, respectively. In a long-term study with dosing for 104 weeks, observed C_min was sustained over time.

For doses of ACTEMRA 4 mg per kg given every 4 weeks, the estimated mean (± SD) steady-state AUC, C_min and C_max of tocilizumab were 13000 ± 5800 mcg•h per mL, 1.49 ± 2.13 mcg per mL, and 88.3 ± 41.4 mcg per mL, respectively. The accumulation ratios for AUC and C_max were 1.11 and 1.02, respectively. The accumulation ratio was higher for C_min (1.96). Steady-state was reached following the first administration for C_max and AUC, respectively, and after 16 weeks C_min. For doses of ACTEMRA 8 mg per kg given every 4 weeks, the estimated mean (± SD) steady-state AUC, C_min and C_max of tocilizumab were 35000 ± 15500 mcg•h per mL, 9.74 ± 10.5 mcg per mL, and 183 ± 85.6 mcg per mL, respectively. The accumulation ratios for AUC and C_max were 1.22 and 1.06, respectively. The accumulation ratio was higher for C_min (2.35). Steady-state was reached following the first administration and after 8 and 20 weeks for C_max, AUC, and C_min, respectively. Tocilizumab AUC, C_min and C_max increased with increase of body weight. At body weight at or above 100 kg, the estimated mean (± SD) steady-state AUC, C_min and C_max of tocilizumab were 55500 ± 14100 mcg•h per mL, 19.0 ± 12.0 mcg per mL, and 269 ± 57 mcg per mL, respectively, which are higher than mean exposure values for the patient population. Therefore, ACTEMRA doses exceeding 800 mg per infusion are not recommended [see Dosage and Administration (2.1)].

*Rheumatoid Arthritis—Subcutaneous Administration*

The pharmacokinetics of tocilizumab was characterized using a population pharmacokinetic analysis using a database composed of 1759 rheumatoid arthritis patients treated with 162 mg subcutaneously every week, 162 mg subcutaneously every other week, and 8 mg/kg every 4 weeks for 24 weeks.

The pharmacokinetic parameters of tocilizumab did not change with time. For the 162 mg every week dose, the estimated mean (±SD) steady-state AUC_{1week}, C_min and C_max of tocilizumab were 8200 ± 3600 mcg•h/mL, 44.6 ± 20.6 mcg/mL, and 50.9 ± 21.8 mcg/mL, respectively. The accumulation ratios for AUC, C_min, and C_max were 6.83, 6.37, and 5.47, respectively. Steady state was reached after 12 weeks for AUC, C_min, and C_max.

For the 162 mg every other week dose, the estimated mean (±SD) steady-state AUC_{2week}, C_min, and C_max of tocilizumab were 3200 ± 2700 mcg•h/mL, 5.6 ± 7.0 mcg/mL, and 12.3 ± 8.7 mcg/mL, respectively. The accumulation ratios for AUC, C_min, and C_max were 2.67, 5.6, and 2.12, respectively. Steady state was reached after 12 weeks for AUC and C_min and after 10 weeks for C_max.

*Giant Cell Arteritis – Subcutaneous Administration*

The pharmacokinetics of tocilizumab in GCA patients was determined using a population pharmacokinetic analysis on a dataset composed of 149 GCA patients treated with 162 mg subcutaneously every week or with 162 mg subcutaneously every other week.

For the 162 mg every week dose, the estimated mean (±SD) steady-state C_{avg}, C_{min} and C_{max} of tocilizumab were 71.3 ± 30.1 mcg/mL, 68.1 ± 29.5 mcg/mL, and 73 ± 30.4 mcg/mL, respectively. The accumulation ratios for C_{avg} or AUC_{tau}, C_{min}, and C_{max} were 10.9, 9.6, and 8.9, respectively. Steady state was reached after 17 weeks.
For the 162 mg every other week dose, the estimated mean (±SD) steady-state $C_{avg}$, $C_{min}$, and $C_{max}$ of tocilizumab were $16.2 \pm 11.8$ mcg/mL, $11.1 \pm 10.3$ mcg/mL, and $19.3 \pm 12.8$ mcg/mL, respectively. The accumulation ratios for $C_{avg}$ or AUC$_{last}$, $C_{min}$, and $C_{max}$ were 2.8, 5.6, and 2.3 respectively. Steady-state was reached after 14 weeks.

**Polyarticular Juvenile Idiopathic Arthritis—Intravenous and Subcutaneous Administration**

The pharmacokinetics of tocilizumab (TCZ) in PJIA patients was characterized by a population pharmacokinetic analysis which included 188 patients who were treated with TCZ IV or 52 patients treated with TCZ SC.

For doses of $8 \text{ mg/kg}$ tocilizumab (patients with a body weight at or above 30 kg) given every 4 weeks intravenously, the estimated mean (±SD) $C_{max}$, $C_{trough}$, and $C_{mean}$ of tocilizumab were $183 \pm 42.3$ mcg/mL, $6.55 \pm 7.93$ mcg/mL and $42.2 \pm 13.4$ mcg/mL, respectively. For doses of $10 \text{ mg/kg}$ tocilizumab (patients with a body weight less than 30 kg) given every 4 weeks intravenously, the estimated mean (±SD) $C_{max}$, $C_{trough}$, and $C_{mean}$ of tocilizumab were $168 \pm 24.8$ mcg/mL, $1.47 \pm 2.44$ mcg/mL and $31.6 \pm 7.84$ mcg/mL, respectively.

The accumulation ratios were 1.05 and 1.16 for AUC$_{4\text{ weeks}}$, and 1.43 and 2.22 for $C_{trough}$ for $10 \text{ mg/kg}$ (BW less than 30 kg) and $8 \text{ mg/kg}$ (BW at or above 30 kg) intravenous doses, respectively. No accumulation for $C_{max}$ was observed. Following $8 \text{ mg/kg}$ every 4 weeks TCZ IV steady state peak and trough concentrations in PJIA pediatric patients (aged 2 to 17 years old) are comparable to those in adult RA patients following $8 \text{ mg/kg}$ IV every 4 weeks. Following $8\text{mg/kg}$ every 4 weeks, the average concentration ($C_{mean}$) of TCZ in PJIA patients after TCZ IV is slightly lower than in adult RA patients.

For doses of $162 \text{ mg}$ tocilizumab (patients with a body weight at or above 30 kg) given every 2 weeks subcutaneously, the estimated mean (±SD) $C_{max}$, $C_{trough}$, and $C_{mean}$ of tocilizumab were $29.4 \pm 13.5$ mcg/mL, $11.8 \pm 7.08$ mcg/mL and $21.7 \pm 10.4$ mcg/mL, respectively. For doses of $162 \text{ mg}$ tocilizumab (patients with a body weight less than 30 kg) given every 3 weeks subcutaneously, the estimated mean (±SD) $C_{max}$, $C_{trough}$, and $C_{mean}$ of tocilizumab were $75.5 \pm 24.1$ mcg/mL, $18.4 \pm 12.9$ mcg/mL and $45.5 \pm 19.8$ mcg/mL, respectively.

The accumulation ratios were 1.46 and 2.04 for AUC$_{4\text{ weeks}}$, 2.08 and 3.58 for $C_{trough}$, and 1.32 and 1.72 for $C_{max}$, for $162 \text{ mg}$ given every 3 weeks (BW less than 30 kg) and $162 \text{ mg}$ given every 2 weeks (BW at or above 30 kg) subcutaneous doses, respectively. Following subcutaneous dosing, steady state $C_{trough}$ was comparable for patients in the two body weight groups, while steady-state $C_{max}$ and $C_{mean}$ were higher for patients in the less than 30 kg group compared to the group at or above 30 kg. All patients treated with TCZ SC had steady-state $C_{trough}$ at or higher than that achieved with TCZ IV across the spectrum of body weights. The average and trough concentrations in patients after subcutaneous dosing were within the range of those achieved in adult patients with RA following the subcutaneous administration of the recommended regimens.

**Systemic Juvenile Idiopathic Arthritis—Intravenous and Subcutaneous Administration**

The pharmacokinetics of tocilizumab (TCZ) in SJIA patients was characterized by a population pharmacokinetic analysis which included 89 patients who were treated with TCZ IV or 51 patients treated with TCZ SC.

For doses of $8 \text{ mg/kg}$ tocilizumab (patients with a body weight at or above 30 kg) given every 2 weeks intravenously, the estimated mean (±SD) $C_{max}$, $C_{trough}$, and $C_{mean}$ of tocilizumab were $256 \pm 60.8$ mcg/mL, $69.7 \pm 29.1$ mcg/mL and $119 \pm 36.0$ mcg/mL, respectively. For doses of $12 \text{ mg/kg}$ tocilizumab (patients with a body weight less than 30 kg) given every 2 weeks intravenously, the estimated mean (±SD) $C_{max}$, $C_{trough}$, and $C_{mean}$ of tocilizumab were $274 \pm 63.8$ mcg/mL, $68.4 \pm 30.0$ mcg/mL and $123 \pm 36.0$ mcg/mL, respectively.

The accumulation ratios were 1.95 and 2.01 for AUC$_{4\text{ weeks}}$, and 3.41 and 3.20 for $C_{trough}$ for $12 \text{ mg/kg}$ (BW less than 30 kg) and $8 \text{ mg/kg}$ (BW at or above 30 kg) intravenous doses, respectively. Accumulation data for $C_{max}$ were 1.37 and 1.42 for $12 \text{ mg/kg}$ (BW less than 30 kg) and $8 \text{ mg/kg}$ (BW at or above 30 kg) intravenous doses, respectively. Following every other week dosing with tocilizumab IV, steady state was reached by 8 weeks for
both body weight groups. Mean estimated tocilizumab exposure parameters were similar between the two dose groups defined by body weight.

For doses of 162 mg tocilizumab (patients with a body weight at or above 30 kg) given every week subcutaneously, the estimated mean (± SD) $C_{\text{max}}$, $C_{\text{trough}}$, and $C_{\text{mean}}$ of tocilizumab were $99.8 \pm 46.2$ mcg/mL, $79.2 \pm 35.6$ mcg/mL and $91.3 \pm 40.4$ mcg/mL, respectively. For doses of 162 mg tocilizumab (patients with a body weight less than 30 kg) given every 2 weeks subcutaneously, the estimated mean (± SD) $C_{\text{max}}$, $C_{\text{trough}}$, and $C_{\text{mean}}$ of tocilizumab were $134 \pm 58.6$ mcg/mL, $65.9 \pm 31.3$ mcg/mL and $101 \pm 43.2$ mcg/mL, respectively.

The accumulation ratios were 2.27 and 4.28 for AUC$_{\text{4weeks}}$, 3.21 and 4.39 for $C_{\text{trough}}$, and 1.88 and 3.66 for $C_{\text{max}}$ for 162 mg given every 2 weeks (BW less than 30 kg) and 162 mg given every week (BW at or above 30 kg) subcutaneous doses, respectively. Following subcutaneous dosing, steady state was reached by 12 weeks for both body weight groups. All patients treated with tocilizumab SC had steady-state $C_{\text{max}}$ lower than that achieved with tocilizumab IV across the spectrum of body weights. Trough and mean concentrations in patients after SC dosing were similar to those achieved with tocilizumab IV across body weights.

**Absorption**

Following subcutaneous dosing in RA and GCA patients, the absorption half-life was around 4 days. The bioavailability for the subcutaneous formulation was 80%.

Following subcutaneous dosing in PJIA patients, the absorption half-life was around 2 days, and the bioavailability for the subcutaneous formulation in PJIA patients was 96%.

Following subcutaneous dosing in SJIA patients, the absorption half-life was around 2 days, and the bioavailability for the SC formulation in SJIA patients was 95%.

In RA patients the median values of $T_{\text{max}}$ were 2.8 days after the tocilizumab every week dose and 4.7 days after the tocilizumab every other week dose.

In GCA patients, the median values of $T_{\text{max}}$ were 3 days after the tocilizumab every week dose and 4.5 days after the tocilizumab every other week dose.

**Distribution**

Following intravenous dosing, tocilizumab undergoes biphasic elimination from the circulation. In rheumatoid arthritis patients the central volume of distribution was 3.5 L and the peripheral volume of distribution was 2.9 L, resulting in a volume of distribution at steady state of 6.4 L.

In GCA patients, the central volume of distribution was 4.09 L, the peripheral volume of distribution was 3.37 L resulting in a volume of distribution at steady state of 7.46 L.

In pediatric patients with PJIA, the central volume of distribution was 1.98 L, the peripheral volume of distribution was 2.1 L, resulting in a volume of distribution at steady state of 4.08 L.

In pediatric patients with SJIA, the central volume of distribution was 1.87 L, the peripheral volume of distribution was 2.14 L resulting in a volume of distribution at steady state of 4.01 L.

**Elimination**

ACTEMRA is eliminated by a combination of linear clearance and nonlinear elimination. The concentration-dependent nonlinear elimination plays a major role at low tocilizumab concentrations. Once the nonlinear pathway is saturated, at higher tocilizumab concentrations, clearance is mainly determined by the linear clearance. The saturation of the nonlinear elimination leads to an increase in exposure that is more than dose-proportional. The pharmacokinetic parameters of ACTEMRA do not change with time.

Population pharmacokinetic analyses in any patient population tested so far indicate no relationship between apparent clearance and the presence of anti-drug antibodies.
The linear clearance in the population pharmacokinetic analysis was estimated to be 12.5 mL per h in RA patients, 6.7 mL per h in GCA patients, 5.8 mL per h in pediatric patients with PJIA, and 5.7 mL per h in pediatric patients with SJIA.

Due to the dependence of total clearance on ACTEMRA serum concentrations, the half-life of ACTEMRA is also concentration-dependent and varies depending on the serum concentration level.

For intravenous administration in RA patients, the concentration-dependent apparent $t_{1/2}$ is up to 11 days for 4 mg per kg and up to 13 days for 8 mg per kg every 4 weeks in patients with RA at steady-state. For subcutaneous administration in RA patients, the concentration-dependent apparent $t_{1/2}$ is up to 13 days for 162 mg every week and 5 days for 162 mg every other week in patients with RA at steady-state.

In GCA patients at steady state, the effective $t_{1/2}$ of tocilizumab varied between 18.3 and 18.9 days for 162 mg subcutaneously every week dosing regimen and between 4.2 and 7.9 days for 162 mg subcutaneously every other week dosing regimen.

The $t_{1/2}$ of tocilizumab in children with PJIA is up to 17 days for the two body weight categories (8 mg/kg for body weight at or above 30 kg or 10 mg/kg for body weight below 30 kg) during a dosing interval at steady state. For subcutaneous administration, the $t_{1/2}$ of tocilizumab in PJIA patients is up to 10 days for the two body weight categories (every other week regimen for body weight at or above 30 kg or every 3 week regimen for body weight less than 30 kg) during a dosing interval at steady state.

The $t_{1/2}$ of tocilizumab intravenous in pediatric patients with SJIA is up to 16 days for the two body weight categories (8 mg/kg for body weight at or above 30 kg and 12 mg/kg for body weight below 30 kg every other week) during a dosing interval at steady-state. Following subcutaneous administration, the effective $t_{1/2}$ of tocilizumab subcutaneous in SJIA patients is up to 14 days for both the body weight categories (162 mg every week for body weight at or above 30 kg and 162 mg every two weeks for body weight below 30 kg) during a dosing interval at steady state.

**Pharmacokinetics in Special Populations**

Population pharmacokinetic analyses in adult rheumatoid arthritis patients and GCA patients showed that age, gender and race did not affect the pharmacokinetics of tocilizumab. Linear clearance was found to increase with body size. In RA patients, the body weight-based dose (8 mg per kg) resulted in approximately 86% higher exposure in patients who are greater than 100 kg in comparison to patients who are less than 60 kg. There was an inverse relationship between tocilizumab exposure and body weight for flat dose subcutaneous regimens.

In GCA patients, higher exposure was observed in patients with lower body weight. For the 162 mg every week dosing regimen, the steady-state Cavg was 51% higher in patients with body weight less than 60 kg compared to patients weighing between 60 to 100 kg. For the 162 mg every other week regimen, the steady-state Cavg was 129% higher in patients with body weight less than 60 kg compared to patients weighing between 60 to 100 kg. There is limited data for patients above 100 kg (n=7).

**Hepatic Impairment**

No formal study of the effect of hepatic impairment on the pharmacokinetics of tocilizumab was conducted.

**Renal Impairment**

No formal study of the effect of renal impairment on the pharmacokinetics of tocilizumab was conducted.

Most of the RA and GCA patients in the population pharmacokinetic analysis had normal renal function or mild renal impairment. Mild renal impairment (estimated creatinine clearance less than 80 mL per min and at or above 50 mL per min based on Cockcroft-Gault formula) did not impact the pharmacokinetics of tocilizumab.

Approximately one-third of the patients in the GCA clinical trial had moderate renal impairment at baseline (estimated creatinine clearance of 30-59 mL/min). No impact on tocilizumab exposure was noted in these patients.

No dose adjustment is required in patients with mild or moderate renal impairment.
**Drug Interactions**

In vitro data suggested that IL-6 reduced mRNA expression for several CYP450 isoenzymes including CYP1A2, CYP2B6, CYP2C9, CYP2C19, CYP2D6 and CYP3A4, and this reduced expression was reversed by co-incubation with tocilizumab at clinically relevant concentrations. Accordingly, inhibition of IL-6 signaling in RA patients treated with tocilizumab may restore CYP450 activities to higher levels than those in the absence of tocilizumab leading to increased metabolism of drugs that are CYP450 substrates. Its effect on CYP2C8 or transporters (e.g., P-gp) is unknown. This is clinically relevant for CYP450 substrates with a narrow therapeutic index, where the dose is individually adjusted. Upon initiation of ACTEMRA, in patients being treated with these types of medicinal products, therapeutic monitoring of the effect (e.g., warfarin) or drug concentration (e.g., cyclosporine or theophylline) should be performed and the individual dose of the medicinal product adjusted as needed. Caution should be exercised when ACTEMRA is coadministered with drugs where decrease in effectiveness is undesirable, e.g., oral contraceptives (CYP3A4 substrates) [see Drug Interactions (7.2)].

**Simvastatin**

Simvastatin is a CYP3A4 and OATP1B1 substrate. In 12 RA patients not treated with ACTEMRA, receiving 40 mg simvastatin, exposures of simvastatin and its metabolite, simvastatin acid, was 4- to 10-fold and 2-fold higher, respectively, than the exposures observed in healthy subjects. One week following administration of a single infusion of ACTEMRA (10 mg per kg), exposure of simvastatin and simvastatin acid decreased by 57% and 39%, respectively, to exposures that were similar or slightly higher than those observed in healthy subjects. Exposures of simvastatin and simvastatin acid increased upon withdrawal of ACTEMRA in RA patients. Selection of a particular dose of simvastatin in RA patients should take into account the potentially lower exposures that may result after initiation of ACTEMRA (due to normalization of CYP3A4) or higher exposures after discontinuation of ACTEMRA.

**Omeprazole**

Omeprazole is a CYP2C19 and CYP3A4 substrate. In RA patients receiving 10 mg omeprazole, exposure to omeprazole was approximately 2 fold higher than that observed in healthy subjects. In RA patients receiving 10 mg omeprazole, before and one week after ACTEMRA infusion (8 mg per kg), the omeprazole AUC_{inf} decreased by 12% for poor (N=5) and intermediate metabolizers (N=5) and by 28% for extensive metabolizers (N=8) and were slightly higher than those observed in healthy subjects.

**Dextromethorphan**

Dextromethorphan is a CYP2D6 and CYP3A4 substrate. In 13 RA patients receiving 30 mg dextromethorphan, exposure to dextromethorphan was comparable to that in healthy subjects. However, exposure to its metabolite, dextrorphan (a CYP3A4 substrate), was a fraction of that observed in healthy subjects. One week following administration of a single infusion of ACTEMRA (8 mg per kg), dextromethorphan exposure was decreased by approximately 5%. However, a larger decrease (29%) in dextrorphan levels was noted after ACTEMRA infusion.

**13 NONCLINICAL TOXICOLOGY**

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No long-term animal studies have been performed to establish the carcinogenicity potential of tocilizumab. Literature indicates that the IL-6 pathway can mediate anti-tumor responses by promoting increased immune cell surveillance of the tumor microenvironment. However, available published evidence also supports that IL-6 signaling through the IL-6 receptor may be involved in pathways that lead to tumorigenesis. The malignancy risk in humans from an antibody that disrupts signaling through the IL-6 receptor, such as tocilizumab, is presently unknown.

Fertility and reproductive performance were unaffected in male and female mice that received a murine analogue of tocilizumab administered by the intravenous route at a dose of 50 mg/kg every three days.
14 CLINICAL STUDIES

14.1 Rheumatoid Arthritis—Intravenous Administration

The efficacy and safety of intravenously administered ACTEMRA was assessed in five randomized, double-blind, multicenter studies in patients greater than 18 years with active rheumatoid arthritis diagnosed according to American College of Rheumatology (ACR) criteria. Patients had at least 8 tender and 6 swollen joints at baseline. ACTEMRA was given intravenously every 4 weeks as monotherapy (Study I), in combination with methotrexate (MTX) (Studies II and III) or other disease-modifying anti-rheumatic drugs (DMARDs) (Study IV) in patients with an inadequate response to those drugs, or in combination with MTX in patients with an inadequate response to TNF antagonists (Study V).

Study I evaluated patients with moderate to severe active rheumatoid arthritis who had not been treated with MTX within 24 weeks prior to randomization, or who had not discontinued previous methotrexate treatment as a result of clinically important toxic effects or lack of response. In this study, 67% of patients were MTX-naive, and over 40% of patients had rheumatoid arthritis less than 2 years. Patients received ACTEMRA 8 mg per kg monotherapy or MTX alone (dose titrated over 8 weeks from 7.5 mg to a maximum of 20 mg weekly). The primary endpoint was the proportion of ACTEMRA patients who achieved an ACR 20 response at Week 24.

Study II was a 104-week study with an ongoing optional 156-week extension phase that evaluated patients with moderate to severe active rheumatoid arthritis who had an inadequate clinical response to MTX. Patients received ACTEMRA 8 mg per kg, ACTEMRA 4 mg per kg, or placebo every four weeks, in combination with MTX (10 to 25 mg weekly). Upon completion of 52-weeks, patients received open-label treatment with ACTEMRA 8 mg per kg through 104 weeks or they had the option to continue their double-blind treatment if they maintained a greater than 70% improvement in swollen/tender joint count. Two pre-specified interim analyses at week 24 and week 52 were conducted. The primary endpoint at week 24 was the proportion of patients who achieved an ACR 20 response. At weeks 52 and 104, the primary endpoints were change from baseline in modified total Sharp-Genant score and the area under the curve (AUC) of the change from baseline in HAQ-DI score.

Study III evaluated patients with moderate to severe active rheumatoid arthritis who had an inadequate clinical response to MTX. Patients received ACTEMRA 8 mg per kg, ACTEMRA 4 mg per kg, or placebo every four weeks, in combination with MTX (10 to 25 mg weekly). The primary endpoint was the proportion of patients who achieved an ACR 20 response at week 24.

Study IV evaluated patients who had an inadequate response to their existing therapy, including one or more DMARDs. Patients received ACTEMRA 8 mg per kg or placebo every four weeks, in combination with the stable DMARDs. The primary endpoint was the proportion of patients who achieved an ACR 20 response at week 24.

Study V evaluated patients with moderate to severe active rheumatoid arthritis who had an inadequate clinical response or were intolerant to one or more TNF antagonist therapies. The TNF antagonist therapy was discontinued prior to randomization. Patients received ACTEMRA 8 mg per kg, ACTEMRA 4 mg per kg, or placebo every four weeks, in combination with MTX (10 to 25 mg weekly). The primary endpoint was the proportion of patients who achieved an ACR 20 response at week 24.

Clinical Response

The percentages of intravenous ACTEMRA-treated patients achieving ACR 20, 50 and 70 responses are shown in Table 3. In all intravenous studies, patients treated with 8 mg per kg ACTEMRA had higher ACR 20, ACR 50, and ACR 70 response rates versus MTX- or placebo-treated patients at week 24.

During the 24 week controlled portions of Studies I to V, patients treated with ACTEMRA at a dose of 4 mg per kg in patients with inadequate response to DMARDs or TNF antagonist therapy had lower response rates compared to patients treated with ACTEMRA 8 mg per kg.
<table>
<thead>
<tr>
<th>Response Rate</th>
<th>Study I</th>
<th>Study II</th>
<th>Study III</th>
<th>Study IV</th>
<th>Study V</th>
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<td></td>
<td>MTX ACTEMRA 8 mg per kg</td>
<td>Placebo + MTX 4 mg per kg + ACTEMRA 8 mg per kg</td>
<td>Placebo + MTX 4 mg per kg + ACTEMRA 8 mg per kg</td>
<td>Placebo + DMARDs + ACTEMRA 8 mg per kg</td>
<td>Placebo + MTX 4 mg per kg + ACTEMRA 8 mg per kg</td>
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<td>N=284 (95% CI)</td>
<td>N=393 (95% CI)</td>
<td>N=204 (95% CI)</td>
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<td><strong>ACR 20</strong></td>
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<tr>
<td>Week 24</td>
<td>53% (0.11, 0.27)</td>
<td>27% (0.15, 0.32)</td>
<td>27% (0.23, 0.41)</td>
<td>24% (0.30, 0.40)</td>
<td>10% (0.15, 0.36)</td>
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<td>Week 52</td>
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<td>25% (0.15, 0.28)</td>
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<td>N/A N/A</td>
<td>N/A N/A</td>
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<td><strong>ACR 50</strong></td>
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<td>Week 24</td>
<td>34% (0.04, 0.20)</td>
<td>10% (0.09, 0.20)</td>
<td>11% (0.13, 0.29)</td>
<td>9% (0.23, 0.33)</td>
<td>4% (0.05, 0.25)</td>
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<td>10% (0.14, 0.25)</td>
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<td>N/A N/A</td>
<td>N/A N/A</td>
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<td><strong>ACR 70</strong></td>
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<td>Week 24</td>
<td>15% (0.07, 0.22)</td>
<td>2% (0.03, 0.13)</td>
<td>2% (0.04, 0.18)</td>
<td>3% (0.13, 0.21)</td>
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<td>Week 52</td>
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<td>4% (0.08, 0.17)</td>
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<td>N/A N/A</td>
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<td><strong>Major Clinical Responses</strong> b</td>
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<tr>
<td>Week 52</td>
<td>N/A N/A</td>
<td>1% (0.01, 0.06)</td>
<td>N/A N/A</td>
<td>N/A N/A</td>
<td>N/A N/A</td>
</tr>
</tbody>
</table>

a CI: 95% confidence interval of the weighted difference to placebo adjusted for site (and disease duration for Study I only)  
b Major clinical response is defined as achieving an ACR 70 response for a continuous 24 week period
In study II, a greater proportion of patients treated with 4 mg per kg and 8 mg per kg ACTEMRA + MTX achieved a low level of disease activity as measured by a DAS 28-ESR less than 2.6 compared with placebo +MTX treated patients at week 52. The proportion of ACTEMRA-treated patients achieving DAS 28-ESR less than 2.6, and the number of residual active joints in these responders in Study II are shown in Table 4.

Table 4  Proportion of Patients with DAS28-ESR Less Than 2.6 with Number of Residual Active Joints in Trials of Intravenous ACTEMRA

<table>
<thead>
<tr>
<th>Study II</th>
<th>Placebo + MTX</th>
<th>ACTEMRA 4 mg per kg + MTX</th>
<th>ACTEMRA 8 mg per kg + MTX</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N = 393</td>
<td>N = 399</td>
<td>N = 398</td>
</tr>
<tr>
<td>DAS28-ESR less than 2.6</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Proportion of responders at week 52 (n)</td>
<td>3% (12)</td>
<td>18% (70)</td>
<td>32% (127)</td>
</tr>
<tr>
<td>95% confidence interval</td>
<td>0.10, 0.19</td>
<td>0.24, 0.34</td>
<td></td>
</tr>
<tr>
<td>Of responders, proportion with 0 active joints (n)</td>
<td>33% (4)</td>
<td>27% (19)</td>
<td>21% (27)</td>
</tr>
<tr>
<td>Of responders, proportion with 1 active joint (n)</td>
<td>8% (1)</td>
<td>19% (13)</td>
<td>13% (16)</td>
</tr>
<tr>
<td>Of responders, proportion with 2 active joints (n)</td>
<td>25% (3)</td>
<td>13% (9)</td>
<td>20% (25)</td>
</tr>
<tr>
<td>Of responders, proportion with 3 or more active joints (n)</td>
<td>33% (4)</td>
<td>41% (29)</td>
<td>47% (59)</td>
</tr>
</tbody>
</table>

*Note: *n* denotes numerator of all the percentage. Denominator is the intent-to-treat population. Not all patients received DAS28 assessments at Week 52.

The results of the components of the ACR response criteria for Studies III and V are shown in Table 5. Similar results to Study III were observed in Studies I, II and IV.

Table 5 Components of ACR Response at Week 24 in Trials of Intravenous ACTEMRA

<table>
<thead>
<tr>
<th>Component (mean)</th>
<th>Study III</th>
<th>Study IV</th>
<th>Placebo + MTX</th>
<th>Study V</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>ACTEMRA 4 mg per kg + MTX</td>
<td>ACTEMRA 8 mg per kg + MTX</td>
<td>N=205</td>
<td>ACTEMRA 4 mg per kg + MTX</td>
</tr>
<tr>
<td></td>
<td>Baseline</td>
<td>Week 24a</td>
<td>Baseline</td>
<td>Week 24a</td>
</tr>
<tr>
<td>Number of tender joints (0-68)</td>
<td>33</td>
<td>19 (-7.0, -6.4)</td>
<td>32</td>
<td>14.5 (-9.6, -6.7)</td>
</tr>
<tr>
<td>Number of swollen joints (0-66)</td>
<td>20</td>
<td>10 (-4.2, -2.3)</td>
<td>19.5</td>
<td>8 (-6.2, -4.2)</td>
</tr>
<tr>
<td>Painb</td>
<td>61</td>
<td>33 (-11.0, -5.0)</td>
<td>60</td>
<td>30 (-15.8, -9.9)</td>
</tr>
<tr>
<td>Patient global assessmentb</td>
<td>66</td>
<td>34 (-10.9, -10.9)</td>
<td>65</td>
<td>31 (-14.9, -9.9)</td>
</tr>
<tr>
<td>Physician global assessmentb</td>
<td>64</td>
<td>26 (-5.6, -0.8)</td>
<td>64</td>
<td>23 (-9.0, -4.2)</td>
</tr>
<tr>
<td>Disability index (HAQ)d</td>
<td>1.64</td>
<td>1.01 (-0.34, 0.02)</td>
<td>1.55</td>
<td>0.96 (-0.21, -0.05)</td>
</tr>
<tr>
<td>CRP (mg per dL)</td>
<td>2.79</td>
<td>1.17 (-1.30, -0.59)</td>
<td>2.61</td>
<td>0.25 (-2.156, -1.46)</td>
</tr>
</tbody>
</table>

*a* Data shown is mean at week 24, difference in adjusted mean change from baseline compared with placebo + MTX at week 24 and 95% confidence interval for that difference

*b* Visual analog scale: 0 = best, 100 = worst

*c* Health Assessment Questionnaire: 0 = best, 3 = worst; 20 questions; 8 categories: dressing and grooming, arising, eating, walking, hygiene, reach, grip, and activities
The percent of ACR 20 responders by visit for Study III is shown in Figure 1. Similar response curves were observed in studies I, II, IV, and V.

**Figure 1** Percent of ACR 20 Responders by Visit for Study III (Inadequate Response to MTX)*

*The same patients may not have responded at each timepoint.

**Radiographic Response**

In Study II, structural joint damage was assessed radiographically and expressed as change in total Sharp-Genant score and its components, the erosion score and joint space narrowing score. Radiographs of hands/wrists and forefeet were obtained at baseline, 24 weeks, 52 weeks, and 104 weeks and scored by readers unaware of treatments group and visit number. The results from baseline to week 52 are shown in Table 6. ACTEMRA 4 mg per kg slowed (less than 75% inhibition compared to the control group) and ACTEMRA 8 mg per kg inhibited (at least 75% inhibition compared to the control group) the progression of structural damage compared to placebo plus MTX at week 52.
Table 6  Mean Radiographic Change from Baseline to Week 52 in Study II

<table>
<thead>
<tr>
<th></th>
<th>Placebo + MTX N=294</th>
<th>ACTEMRA 4 mg per kg + MTX N=343</th>
<th>ACTEMRA 8 mg per kg + MTX N=353</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Week 52</strong>*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total Sharp-Genant Score, Mean (SD)</td>
<td>1.17 (3.14)</td>
<td>0.33 (1.30)</td>
<td>0.25 (0.98)</td>
</tr>
<tr>
<td>Adjusted Mean difference** (95% CI)</td>
<td>-0.83 (-1.13, -0.52)</td>
<td>-0.90 (-1.20, -0.59)</td>
<td></td>
</tr>
<tr>
<td>Erosion Score, Mean (SD)</td>
<td>0.76 (2.14)</td>
<td>0.20 (0.83)</td>
<td>0.15 (0.77)</td>
</tr>
<tr>
<td>Adjusted Mean difference** (95% CI)</td>
<td>-0.55 (-0.76, -0.34)</td>
<td>-0.60 (-0.80, -0.39)</td>
<td></td>
</tr>
<tr>
<td>Joint Space Narrowing Score, Mean (SD)</td>
<td>0.41 (1.71)</td>
<td>0.13 (0.72)</td>
<td>0.10 (0.49)</td>
</tr>
<tr>
<td>Adjusted Mean difference** (95% CI)</td>
<td>-0.28 (-0.44, -0.11)</td>
<td>-0.30 (-0.46, -0.14)</td>
<td></td>
</tr>
</tbody>
</table>

* Week 52 analysis employs linearly extrapolated data for patients after escape, withdrawal, or loss to follow up.
** Difference between the adjusted means (ACTEMRA + MTX - Placebo + MTX)
SD = standard deviation

The mean change from baseline to week 104 in Total Sharp-Genant Score for the ACTEMRA 4 mg per kg groups was 0.47 (SD = 1.47) and for the 8 mg per kg groups was 0.34 (SD = 1.24). By the week 104, most patients in the control (placebo + MTX) group had crossed over to active treatment, and results are therefore not included for comparison. Patients in the active groups may have crossed over to the alternate active dose group, and results are reported per original randomized dose group.

In the placebo group, 66% of patients experienced no radiographic progression (Total Sharp-Genant Score change ≤ 0) at week 52 compared to 78% and 83% in the ACTEMRA 4 mg per kg and 8 mg per kg, respectively. Following 104 weeks of treatment, 75% and 83% of patients initially randomized to ACTEMRA 4 mg per kg and 8 mg per kg, respectively, experienced no progression of structural damage compared to 66% of placebo treated patients.

Health Related Outcomes
In Study II, physical function and disability were assessed using the Health Assessment Questionnaire Disability Index (HAQ-DI). Both dosing groups of ACTEMRA demonstrated a greater improvement compared to the placebo group in the AUC of change from baseline in the HAQ-DI through week 52. The mean change from baseline to week 52 in HAQ-DI was 0.6, 0.5, and 0.4 for ACTEMRA 8 mg per kg, ACTEMRA 4 mg per kg, and placebo treatment groups, respectively. Sixty-three percent (63%) and sixty percent (60%) of patients in the ACTEMRA 8 mg per kg and ACTEMRA 4 mg per kg treatment groups, respectively, achieved a clinically relevant improvement in HAQ-DI (change from baseline of ≥ 0.3 units) at week 52 compared to 53% in the placebo treatment group.

Other Health-Related Outcomes
General health status was assessed by the Short Form Health Survey (SF-36) in Studies I – V. Patients receiving ACTEMRA demonstrated greater improvement from baseline compared to placebo in the Physical Component Summary (PCS), Mental Component Summary (MCS), and in all 8 domains of the SF-36.
14.2 Rheumatoid Arthritis—Subcutaneous Administration

The efficacy and safety of subcutaneously administered ACTEMRA was assessed in two double-blind, controlled, multicenter studies in patients with active RA. One study (SC-I) was a non-inferiority study that compared the efficacy and safety of ACTEMRA 162 mg administered every week subcutaneously to 8 mg per kg intravenously every four weeks. The second study (SC-II) was a placebo controlled superiority study that evaluated the safety and efficacy of ACTEMRA 162 mg administered every other week subcutaneously to placebo. Both SC-I and SC-II required patients to be >18 years of age with moderate to severe active rheumatoid arthritis diagnosed according to ACR criteria who had at least 4 tender and 4 swollen joints at baseline (SC-I) or at least 8 tender and 6 swollen joints at baseline (SC-II), and an inadequate response to their existing DMARD therapy, where approximately 20% also had a history of inadequate response to at least one TNF inhibitor. All patients in both SC studies received background non-biologic DMARD(s).

In SC-I, 1262 patients were randomized 1:1 to receive ACTEMRA-SC 162 mg every week or intravenous ACTEMRA 8 mg/kg every four weeks in combination with DMARD(s). In SC-II, 656 patients were randomized 2:1 to ACTEMRA-SC 162 mg every other week or placebo, in combination with DMARD(s). The primary endpoint in both studies was the proportion of patients who achieved an ACR20 response at Week 24.

The clinical response to 24 weeks of ACTEMRA-SC therapy is shown in Table 7. In SC-I, the primary outcome measure was ACR20 at Week 24. The pre-specified non-inferiority margin was a treatment difference of 12%. The study demonstrated non-inferiority of ACTEMRA with respect to ACR20 at Week 24; ACR50, ACR70, and DAS28 responses are also shown in Table 7. In SC-II, a greater portion of patients treated with ACTEMRA 162 mg subcutaneously every other week achieved ACR20, ACR50, and ACR70 responses compared to placebo-treated patients (Table 7). Further, a greater proportion of patients treated with ACTEMRA 162 mg subcutaneously every other week achieved a low level of disease activity as measured by a DAS28-ESR less than 2.6 at Week 24 compared to those treated with placebo (Table 7).

Table 7  Clinical Response at Week 24 in Trials of Subcutaneous ACTEMRA (Percent of Patients)

<table>
<thead>
<tr>
<th></th>
<th>SC-I</th>
<th>SC-II</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TCZ SC 162 mg every week + DMARD</td>
<td>TCZ IV 8mg/kg every four weeks + DMARD</td>
</tr>
<tr>
<td>ACR20</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Week 24</td>
<td>69%</td>
<td>73.4%</td>
</tr>
<tr>
<td>Weighted difference (95% CI)</td>
<td>-4% (-9.2, 1.2)</td>
<td>30% (22.0, 37.0)</td>
</tr>
<tr>
<td>ACR50</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Week 24</td>
<td>47%</td>
<td>49%</td>
</tr>
<tr>
<td>Weighted difference (95% CI)</td>
<td>-2% (-7.5, 4.0)</td>
<td>28% (21.5, 34.4)</td>
</tr>
<tr>
<td>ACR70</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Week 24</td>
<td>24%</td>
<td>28%</td>
</tr>
<tr>
<td>Weighted difference (95% CI)</td>
<td>-4% (-9.0, 1.3)</td>
<td>15% (9.8, 19.9)</td>
</tr>
<tr>
<td>Change in DAS28 [Adjusted mean]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Week 24</td>
<td>-3.5</td>
<td>-3.5</td>
</tr>
<tr>
<td>Adjusted mean difference (95% CI)</td>
<td>0 (-0.2, 0.1)</td>
<td>-1.4 (-1.7; -1.1)</td>
</tr>
<tr>
<td>DAS28 &lt; 2.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Week 24</td>
<td>38.4%</td>
<td>36.9%</td>
</tr>
<tr>
<td>Weighted difference (95% CI)</td>
<td>0.9 (-5.0, 6.8)</td>
<td>28.6 (22.0, 35.2)</td>
</tr>
</tbody>
</table>

TCZ = tocilizumab

a Per Protocol Population
b Intent To Treat Population
The results of the components of the ACR response criteria and the percent of ACR20 responders by visit for ACTEMRA-SC in Studies SC-I and SC-II were consistent with those observed for ACTEMRA-IV.

**Radiographic Response**

In study SC-II, the progression of structural joint damage was assessed radiographically and expressed as a change from baseline in the van der Heijde modified total Sharp score (mTSS). At week 24, significantly less radiographic progression was observed in patients receiving ACTEMRA-SC every other week plus DMARD(s) compared to placebo plus DMARD(s); mean change from baseline in mTSS of 0.62 vs. 1.23, respectively, with an adjusted mean difference of -0.60 (-1.1, -0.1). These results are consistent with those observed in patients treated with intravenous ACTEMRA.

**Health Related Outcomes**

In studies SC-I and SC-II, the mean decrease from baseline to week 24 in HAQ-DI was 0.6, 0.6, 0.4 and 0.3, and the proportion of patients who achieved a clinically relevant improvement in HAQ-DI (change from baseline of ≥ 0.3 units) was 65%, 67%, 58% and 47%, for the subcutaneous every week, intravenous 8 mg/kg, subcutaneous every other week, and placebo treatment groups, respectively.

Other Health-Related Outcomes

General health status was assessed by the SF-36 in Studies SC-I and SC-II. In Study SC-II, patients receiving ACTEMRA every other week demonstrated greater improvement from baseline compared to placebo in the PCS, MCS, and in all 8 domains of the SF-36. In Study SC-I, improvements in these scores were similar between ACTEMRA-SC every week and ACTEMRA-IV 8 mg/kg.

### 14.3 Giant Cell Arteritis—Subcutaneous Administration

The efficacy and safety of subcutaneously administered ACTEMRA was assessed in a single, randomized, double-blind, multicenter study in patients with active GCA. In Study WA28119, 251 screened patients with new-onset or relapsing GCA were randomized to one of four treatment arms. Two subcutaneous doses of ACTEMRA (162 mg every week and 162 mg every other week) were compared to two different placebo control groups (pre-specified prednisone-taper regimen over 26 weeks and 52 weeks) randomized 2:1:1:1. The study consisted of a 52-week blinded period, followed by a 104-week open-label extension.

All patients received background glucocorticoid (prednisone) therapy. Each of the ACTEMRA-treated groups and one of the placebo-treated groups followed a pre-specified prednisone-taper regimen with the aim to reach 0 mg by 26 weeks, while the second placebo-treated group followed a pre-specified prednisone-taper regimen with the aim to reach 0 mg by 52 weeks designed to be more in keeping with standard practice.

The primary efficacy endpoint was the proportion of patients achieving sustained remission from Week 12 through Week 52. Sustained remission was defined by a patient attaining a sustained (1) absence of GCA signs and symptoms from Week 12 through Week 52, (2) normalization of erythrocyte sedimentation rate (ESR) (to < 30 mm/hr without an elevation to ≥ 30 mm/hr attributable to GCA) from Week 12 through Week 52, (3) normalization of C-reactive protein (CRP) (to < 1 mg/dL, with an absence of successive elevations to ≥ 1mg/dL) from Week 12 through Week 52, and (4) successful adherence to the prednisone taper defined by not more than 100 mg of excess prednisone from Week 12 through Week 52. ACTEMRA 162 mg weekly and 162 mg every other week + 26 weeks prednisone taper both showed superiority in achieving sustained remission from Week 12 through Week 52 compared with placebo + 26 weeks prednisone taper (Table 8). Both ACTEMRA treatment arms also showed superiority compared to the placebo + 52 weeks prednisone taper (Table 8).
### Table 8  Efficacy Results from Study WA28119

<table>
<thead>
<tr>
<th></th>
<th>PBO + 26 weeks prednisone taper N=50</th>
<th>PBO + 52 weeks prednisone taper N=51</th>
<th>TCZ 162mg SC QW + 26 weeks prednisone taper N=100</th>
<th>TCZ 162 mg SC Q2W + 26 weeks prednisone taper N=49</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Sustained remission</strong>a</td>
<td>Responders, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>7 (14.0%)</td>
<td>9 (17.6%)</td>
<td>56 (56.0%)</td>
<td>26 (53.1%)</td>
</tr>
<tr>
<td>Unadjusted difference in proportions vs PBO + 26 weeks taper (99.5% CI)</td>
<td>N/A</td>
<td>N/A</td>
<td>42.0% (18.0, 66.0)</td>
<td>39.1% (12.5, 65.7)</td>
</tr>
<tr>
<td>Unadjusted difference in proportions vs PBO + 52 weeks taper (99.5% CI)</td>
<td>N/A</td>
<td>N/A</td>
<td>38.4% (14.4, 62.3)</td>
<td>35.4% (8.6, 62.2)</td>
</tr>
<tr>
<td><strong>Components of Sustained Remission</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sustained absence of GCA signs and symptomsb, n (%)</td>
<td>20 (40.0%)</td>
<td>23 (45.1%)</td>
<td>69 (69.0%)</td>
<td>28 (57.1%)</td>
</tr>
<tr>
<td>Sustained ESR&lt;30 mm/hrc, n (%)</td>
<td>20 (40.0%)</td>
<td>22 (43.1%)</td>
<td>83 (83.0%)</td>
<td>37 (75.5%)</td>
</tr>
<tr>
<td>Sustained CRP normalizationd, n (%)</td>
<td>17 (34.0%)</td>
<td>13 (25.5%)</td>
<td>72 (72.0%)</td>
<td>34 (69.4%)</td>
</tr>
<tr>
<td>Successful prednisone taperinge, n (%)</td>
<td>10 (20.0%)</td>
<td>20 (39.2%)</td>
<td>60 (60.0%)</td>
<td>28 (57.1%)</td>
</tr>
</tbody>
</table>

*a*Sustained remission was achieved by a patient meeting all of the following components: absence of GCA signs and symptomsb, normalization of ESRc, normalization of CRPd and adherence to the prednisone taper regimen e.

b Patients who did not have any signs or symptoms of GCA recorded from Week 12 up to Week 52.

c Patients who did not have an elevated ESR ≥30 mm/hr which was classified as attributed to GCA from Week 12 up to Week 52.

d Patients who did not have two or more consecutive CRP records of ≥ 1mg/dL from Week 12 up to Week 52.

e Patients who did not enter escape therapy and received ≤ 100mg of additional concomitant prednisone from Week 12 up to Week 52.

Patients not completing the study to week 52 were classified as non-responders in the primary and secondary analysis: PBO+26: 6 (12.0%), PBO+52: 5 (9.8%), TCZ QW: 15 (15.0%), TCZ Q2W: 9 (18.4%).

CRP = C-reactive protein
ESR = erythrocyte sedimentation rate
PBO = placebo
Q2W = every other week dose
QW = every week dose
TCZ = tocilizumab

The estimated annual cumulative prednisone dose was lower in the two ACTEMRA dose groups (medians of 1887 mg and 2207 mg on ACTEMRA QW and Q2W, respectively) relative to the placebo arms (medians of 3804 mg and 3902 mg on placebo + 26 weeks prednisone and placebo + 52 weeks prednisone taper, respectively).

### 14.4 Polyarticular Juvenile Idiopathic Arthritis—Intravenous Administration

The efficacy of ACTEMRA was assessed in a three-part study including an open-label extension in children 2 to 17 years of age with active polyarticular juvenile idiopathic arthritis (PJIA), who had an inadequate response to methotrexate or inability to tolerate methotrexate. Patients had at least 6 months of active disease (mean disease duration of 4.2 ± 3.7 years), with at least five joints with active arthritis (swollen or limitation of movement accompanied by pain and/or tenderness) and/or at least 3 active joints having limitation of motion (mean, 20 ± 14 active joints). The patients treated had subtypes of JIA that at disease onset included Rheumatoid Factor Positive or Negative Polyarticular JIA, or Extended Oligoarticular JIA. Treatment with a stable dose of methotrexate was permitted but was not required during the study. Concurrent use of disease modifying antirheumatic drugs (DMARDs), other than methotrexate, or other biologics (e.g., TNF antagonists or T cell costimulation modulator) were not permitted in the study.
Part I consisted of a 16-week active ACTEMRA treatment lead-in period (n=188) followed by Part II, a 24-week randomized double-blind placebo-controlled withdrawal period, followed by Part III, a 64-week open-label period. Eligible patients weighing at or above 30 kg received ACTEMRA at 8 mg/kg intravenously once every four weeks. Patients weighing less than 30 kg were randomized 1:1 to receive either ACTEMRA 8 mg/kg or 10 mg/kg intravenously every four weeks. At the conclusion of the open-label Part I, 91% of patients taking background MTX in addition to tocilizumab and 83% of patients on tocilizumab monotherapy achieved an ACR 30 response at week 16 compared to baseline and entered the blinded withdrawal period (Part II) of the study. The proportions of patients with JIA ACR 50/70 responses in Part I were 84.0%, and 64%, respectively for patients taking background MTX in addition to tocilizumab and 80% and 55% respectively for patients on tocilizumab monotherapy.

In Part II, patients (ITT, n=163) were randomized to ACTEMRA (same dose received in Part I) or placebo in a 1:1 ratio that was stratified by concurrent methotrexate use and concurrent corticosteroid use. Each patient continued in Part II of the study until Week 40 or until the patient satisfied JIA ACR 30 flare criteria (relative to Week 16) and qualified for escape.

The primary endpoint was the proportion of patients with a JIA ACR 30 flare at week 40 relative to week 16. JIA ACR 30 flare was defined as 3 or more of the 6 core outcome variables worsening by at least 30% with no more than 1 of the remaining variables improving by more than 30% relative to Week 16.

ACTEMRA treated patients experienced significantly fewer disease flares compared to placebo-treated patients (26% [21/82] versus 48% [39/81]; adjusted difference in proportions -21%, 95% CI: -35%, -8%).

During the withdrawal phase (Part II), more patients treated with ACTEMRA showed JIA ACR 30/50/70 responses at Week 40 compared to patients withdrawn to placebo.

14.5 Polyarticular Juvenile Idiopathic Arthritis—Subcutaneous Administration

Subcutaneously administered ACTEMRA in pediatric patients with polyarticular juvenile idiopathic arthritis (PJIA) was assessed in a 52-week, open-label, multicenter, PK-PD and safety study to determine the appropriate subcutaneous dose of ACTEMRA that achieved comparable PK/PD profiles to the ACTEMRA-IV regimen. PJIA patients aged 1 to 17 years with an inadequate response or inability to tolerate MTX, including patients with well-controlled disease on treatment with ACTEMRA-IV and ACTEMRA-naive patients with active disease, were treated with subcutaneous ACTEMRA based on body weight.

Patients weighing at or above 30 kg (n = 25) were treated with 162 mg of ACTEMRA-SC every 2 weeks and patients weighing less than 30 kg (n = 27) received 162 mg of ACTEMRA-SC every 3 weeks for 52 weeks. Of these 52 patients, 37 (71%) were naive to ACTEMRA and 15 (29%) had been receiving ACTEMRA-IV and switched to ACTEMRA-SC at baseline.

The efficacy of subcutaneous ACTEMRA in children 2 to 17 years of age is based on pharmacokinetic exposure and extrapolation of the established efficacy of intravenous ACTEMRA in polyarticular JIA patients and subcutaneous ACTEMRA in patients with RA [see Clinical Pharmacology (12.3) and Clinical Studies (14.2 and 14.4)].

14.6 Systemic Juvenile Idiopathic Arthritis—Intravenous Administration

The efficacy of ACTEMRA for the treatment of active SJIA was assessed in a 12-week randomized, double blind, placebo-controlled, parallel group, 2-arm study. Patients treated with or without MTX, were randomized (ACTEMRA:placebo = 2:1) to one of two treatment groups: 75 patients received ACTEMRA infusions every two weeks at either 8 mg per kg for patients at or above 30 kg or 12 mg per kg for patients less than 30 kg and 37 were randomized to receive placebo infusions every two weeks. Corticosteroid tapering could occur from week six for patients who achieved a JIA ACR 70 response. After 12 weeks or at the time of escape, due to disease worsening, patients were treated with ACTEMRA in the open-label extension phase at weight appropriate dosing.

The primary endpoint was the proportion of patients with at least 30% improvement in JIA ACR core set (JIA ACR 30 response) at Week 12 and absence of fever (no temperature at or above 37.5°C in the preceding 7 days).
JIA ACR (American College of Rheumatology) responses are defined as the percentage improvement (e.g., 30%, 50%, 70%) in 3 of any 6 core outcome variables compared to baseline, with worsening in no more than 1 of the remaining variables by 30% or more. Core outcome variables consist of physician global assessment, parent per patient global assessment, number of joints with active arthritis, number of joints with limitation of movement, erythrocyte sedimentation rate (ESR), and functional ability (childhood health assessment questionnaire-CHAQ).

Primary endpoint result and JIA ACR response rates at Week 12 are shown in Table 9.

Table 9  Efficacy Findings at Week 12

<table>
<thead>
<tr>
<th></th>
<th>ACTEMRA N=75</th>
<th>Placebo N=37</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Primary Endpoint: JIA ACR 30 response + absence of fever</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Responders</td>
<td>85%</td>
<td>24%</td>
</tr>
<tr>
<td>Weighted difference (95% CI)</td>
<td>62 (45, 78)</td>
<td>-</td>
</tr>
</tbody>
</table>

JIA ACR Response Rates at Week 12

<table>
<thead>
<tr>
<th></th>
<th>ACTEMRA N=75</th>
<th>Placebo N=37</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>JIA ACR 30</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Responders</td>
<td>91%</td>
<td>24%</td>
</tr>
<tr>
<td>Weighted difference (95% CI)</td>
<td>67 (51, 83)</td>
<td>-</td>
</tr>
</tbody>
</table>

| **JIA ACR 50**   |              |              |
| Responders       | 85%          | 11%          |
| Weighted difference (95% CI) | 74 (58, 90) | -            |

| **JIA ACR 70**   |              |              |
| Responders       | 71%          | 8%           |
| Weighted difference (95% CI) | 63 (46, 80) | -            |

*The weighted difference is the difference between the ACTEMRA and Placebo response rates, adjusted for the stratification factors (weight, disease duration, background oral corticosteroid dose and background methotrexate use).

b CI: confidence interval of the weighted difference.

The treatment effect of ACTEMRA was consistent across all components of the JIA ACR response core variables. JIA ACR scores and absence of fever responses in the open label extension were consistent with the controlled portion of the study (data available through 44 weeks).

**Systemic Features**

Of patients with fever or rash at baseline, those treated with ACTEMRA had fewer systemic features; 35 out of 41 (85%) became fever free (no temperature recording at or above 37.5°C in the preceding 14 days) compared to 5 out of 24 (21%) of placebo-treated patients, and 14 out of 22 (64%) became free of rash compared to 2 out of 18 (11%) of placebo-treated patients. Responses were consistent in the open label extension (data available through 44 weeks).

**Corticosteroid Tapering**

Of the patients receiving oral corticosteroids at baseline, 8 out of 31 (26%) placebo and 48 out of 70 (69%), ACTEMRA patients achieved a JIA ACR 70 response at week 6 or 8 enabling corticosteroid dose reduction. Seventeen (24%) ACTEMRA patients versus 1 (3%) placebo patient were able to reduce the dose of corticosteroid by at least 20% without experiencing a subsequent JIA ACR 30 flare or occurrence of systemic symptoms to week 12. In the open label portion of the study, by week 44, there were 44 out of 103 (43%) ACTEMRA patients off oral corticosteroids. Of these 44 patients 50% were off corticosteroids 18 weeks or more.
**Health Related Outcomes**

Physical function and disability were assessed using the Childhood Health Assessment Questionnaire Disability Index (CHAQ-DI). Seventy-seven percent (58 out of 75) of patients in the ACTEMRA treatment group achieved a minimal clinically important improvement in CHAQ-DI (change from baseline of $\geq 0.13$ units) at week 12 compared to 19% (7 out of 37) in the placebo treatment group.

### 14.7 Systemic Juvenile Idiopathic Arthritis—Subcutaneous Administration

Subcutaneously administered ACTEMRA in pediatric patients with systemic juvenile idiopathic arthritis (SJIA) was assessed in a 52-week, open-label, multicenter, PK-PD and safety study to determine the appropriate subcutaneous dose of ACTEMRA that achieved comparable PK/PD profiles to the ACTEMRA-IV regimen.

Eligible patients received ACTEMRA subcutaneously dosed according to body weight, with patients weighing at or above 30 kg (n = 26) dosed with 162 mg of ACTEMRA every week and patients weighing below 30 kg (n = 25) dosed with 162 mg of ACTEMRA every 10 days (n=8) or every 2 weeks (n=17) for 52 weeks. Of these 51 patients, 26 (51%) were naive to subcutaneous ACTEMRA and 25 (49%) had been receiving ACTEMRA intravenously and switched to subcutaneous ACTEMRA at baseline.

The efficacy of subcutaneous ACTEMRA in children 2 to 17 years of age is based on pharmacokinetic exposure and extrapolation of the established efficacy of intravenous ACTEMRA in systemic JIA patients [see Clinical Pharmacology (12.3) and Clinical Studies (14.2 and 14.6)].

### 14.8 Cytokine Release Syndrome—Intravenous Administration

The efficacy of ACTEMRA for the treatment of CRS was assessed in a retrospective analysis of pooled outcome data from clinical trials of CAR T-cell therapies for hematological malignancies. Evaluable patients had been treated with tocilizumab 8 mg/kg (12 mg/kg for patients < 30 kg) with or without additional high-dose corticosteroids for severe or life-threatening CRS; only the first episode of CRS was included in the analysis. The study population included 24 males and 21 females (total 45 patients) of median age 12 years (range, 3–23 years); 82% were Caucasian. The median time from start of CRS to first dose of tocilizumab was 4 days (range, 0-18 days). Resolution of CRS was defined as lack of fever and off vasopressors for at least 24 hours. Patients were considered responders if CRS resolved within 14 days of the first dose of tocilizumab, if no more than 2 doses of tocilizumab were needed, and if no drugs other than tocilizumab and corticosteroids were used for treatment. Thirty-one patients (69%; 95% CI: 53%–82%) achieved a response. Achievement of resolution of CRS within 14 days was confirmed in a second study using an independent cohort that included 15 patients (range: 9–75 years old) with CAR T cell-induced CRS.

### 16 HOW SUPPLIED/STORAGE AND HANDLING

**For Intravenous Infusion**

ACTEMRA (tocilizumab) injection is a preservative-free, sterile clear, colorless to pale yellow solution. ACTEMRA is supplied as 80 mg/4 mL (NDC 50242-135-01), 200 mg/10 mL (NDC 50242-136-01), and 400 mg/20 mL (NDC 50242-137-01) individually packaged 20 mg/mL single-dose vials for further dilution prior to intravenous infusion.

**For Subcutaneous Injection**

ACTEMRA (tocilizumab) injection is supplied as a preservative-free, sterile, clear, colorless to slightly yellowish solution for subcutaneous administration. The following packaging configurations are available:

- Each single-dose prefilled syringe delivers 162 mg/0.9 mL (NDC 50242-138-01).
- Each single-dose autoinjector (ACTPen™) delivers 162 mg/0.9 mL (NDC 50242-143-01).

**Storage and Stability:** Do not use beyond expiration date on the container, package, prefilled syringe, or autoinjector. ACTEMRA must be refrigerated at 2°C to 8°C (36°F to 46°F). Do not freeze. Protect the vials, syringes, and autoinjectors from light by storage in the original package until time of use, and keep syringes and autoinjectors dry.
17 PATIENT COUNSELING INFORMATION
Advise the patient to read the FDA-approved patient labeling (Medication Guide).

Patient Counseling
Advise patients and parents or guardians of minors with PJIA, SJIA, or CRS of the potential benefits and risks of ACTEMRA.

- Infections:
  Inform patients that ACTEMRA may lower their resistance to infections. Instruct the patient of the importance of contacting their doctor immediately when symptoms suggesting infection appear in order to assure rapid evaluation and appropriate treatment.

- Gastrointestinal Perforation:
  Inform patients that some patients who have been treated with ACTEMRA have had serious side effects in the stomach and intestines. Instruct the patient of the importance of contacting their doctor immediately when symptoms of severe, persistent abdominal pain appear to assure rapid evaluation and appropriate treatment.

- Hypersensitivity and Serious Allergic Reactions
  Assess patient suitability for home use for subcutaneous injection. Inform patients that some patients who have been treated with ACTEMRA have developed serious allergic reactions, including anaphylaxis. Advise patients to seek immediate medical attention if they experience any symptom of serious allergic reactions.

Instruction on Injection Technique
Perform the first injection under the supervision of a qualified healthcare professional. If a patient or caregiver is to administer subcutaneous ACTEMRA, instruct him/her in injection techniques and assess his/her ability to inject subcutaneously to ensure proper administration of subcutaneous ACTEMRA and the suitability for home use [See Patient Instructions for Use].

Prior to use, remove the prefilled syringe (PFS) or autoinjector from the refrigerator and allow to sit at room temperature outside of the carton for 30 minutes (PFS) or 45 minutes (autoinjector), out of the reach of children. Do not warm ACTEMRA in any other way.

Advise patients to consult their healthcare provider if the full dose is not received.

A puncture-resistant container for disposal of needles, syringes and autoinjectors should be used and should be kept out of the reach of children. Instruct patients or caregivers in the technique as well as proper needle, syringe and autoinjector disposal, and caution against reuse of these items.

Pregnancy Exposure Registry
Inform patients that there is a pregnancy registry to monitor fetal outcomes of pregnant women exposed to ACTEMRA [see Use in Specific Populations (8.1)].

Pregnancy
Inform female patients of reproductive potential that ACTEMRA may cause fetal harm and to inform their prescriber of a known or suspected pregnancy [see Use in Specific Populations (8.1)].

ACTEMRA® (tocilizumab)
Manufactured by:
Genentech, Inc.
A Member of the Roche Group
1 DNA Way
South San Francisco, CA 94080-4990
US License No. 1048
What is the most important information I should know about ACTEMRA?

**ACTEMRA can cause serious side effects including:**

1. **Serious Infections.** ACTEMRA is a medicine that affects your immune system. ACTEMRA can lower the ability of your immune system to fight infections. Some people have serious infections while taking ACTEMRA, including tuberculosis (TB), and infections caused by bacteria, fungi, or viruses that can spread throughout the body. Some people have died from these infections. Your healthcare provider should test you for TB before starting ACTEMRA.

Your healthcare provider should monitor you closely for signs and symptoms of TB during treatment with ACTEMRA.

- You should not start taking ACTEMRA if you have any kind of infection unless your healthcare provider says it is okay.

Before starting ACTEMRA, tell your healthcare provider if you:

- think you have an infection or have symptoms of an infection, with or without a fever, such as:
  - sweating or chills
  - feel very tired
  - cough
  - shortness of breath
  - muscle aches
  - weight loss
  - warm, red, or painful skin or sores on your body
  - blood in phlegm
  - muscle aches
  - weight loss
  - diarrhea or stomach pain
  - burning when you urinate or urinating more often than normal
  - are being treated for an infection.
  - get a lot of infections or have infections that keep coming back.
  - have diabetes, HIV, or a weak immune system. People with these conditions have a higher chance for infections.
  - have TB, or have been in close contact with someone with TB.
  - live or have lived, or have traveled to certain parts of the country (such as the Ohio and Mississippi River valleys and the Southwest) where there is an increased chance for getting certain kinds of fungal infections (histoplasmosis, coccidiodomycosis, or blastomycosis). These infections may happen or become more severe if you use ACTEMRA. Ask your healthcare provider, if you do not know if you have lived in an area where these infections are common.
  - have or have had hepatitis B.

After starting ACTEMRA, call your healthcare provider right away if you have any symptoms of an infection. ACTEMRA can make you more likely to get infections or make worse any infection that you have.

2. **Tears (perforation) of the stomach or intestines.**

Tell your healthcare provider if you have had diverticulitis (inflammation in parts of the large intestine) or ulcers in your stomach or intestines. Some people taking ACTEMRA get tears in their stomach or intestine. This happens most often in people who also take nonsteroidal anti-inflammatory drugs (NSAIDs), corticosteroids, or methotrexate.

Tell your healthcare provider right away if you have fever and stomach-area pain that does not go away, and a change in your bowel habits.

3. **Changes in certain laboratory test results.** Your healthcare provider should do blood tests before you start receiving ACTEMRA. If you have rheumatoid arthritis (RA) or giant cell arteritis (GCA) your healthcare provider should do blood tests 4 to 8 weeks after you start receiving ACTEMRA and then every 3 months after that. If you have polyarticular juvenile idiopathic arthritis (PJIA) you will have blood tests done every 4 to 8 weeks during treatment. If you have systemic juvenile idiopathic arthritis (SJIA) you will have blood tests done every 2 to 4 weeks during treatment. These blood test are to check for the following side effects of ACTEMRA:

- low neutrophil count. Neutrophils are white blood cells that help the body fight off bacterial infections.
- low platelet count. Platelets are blood cells that help with blood clotting and stop bleeding.
- increase in certain liver function tests.
- increase in blood cholesterol levels. You may also have changes in other laboratory tests, such as your blood cholesterol levels. Your healthcare provider should do blood tests to check your cholesterol levels 4 to 8 weeks after you start receiving ACTEMRA, and then every 6 months after that.
You should not receive ACTEMRA if your neutrophil or platelet counts are too low or your liver function tests are too high. Your healthcare provider may stop your ACTEMRA treatment for a period of time or change your dose of medicine if needed because of changes in these blood test results.

4. **Cancer.** ACTEMRA may increase your risk of certain cancers by changing the way your immune system works. Tell your healthcare provider if you have ever had any type of cancer.

See “What are the possible side effects with ACTEMRA?” for more information about side effect.

---

**What is ACTEMRA?**

ACTEMRA is a prescription medicine called an Interleukin-6 (IL-6) receptor antagonist. ACTEMRA is used to treat:

- Adults with moderately to severely active rheumatoid arthritis (RA), after at least one other medicine called a Disease-Modifying Anti-Rheumatic Drug (DMARD) has been used and did not work well.
- Adults with giant cell arteritis (GCA).
- People with active PJIA ages 2 and above.
- People with active SJIA ages 2 and above.
- People age 2 years and above who experience severe or life-threatening Cytokine Release Syndrome (CRS) following chimeric antigen receptor (CAR) T cell treatment
- ACTEMRA is not approved for subcutaneous use in people with CRS.

It is not known if ACTEMRA is safe and effective in children with PJIA, SJIA, or CRS under 2 years of age or in children with conditions other than PJIA, SJIA or CRS.

**Do not take ACTEMRA:** if you are allergic to tocilizumab, or any of the ingredients in ACTEMRA. See the end of this Medication Guide for a complete list of ingredients in ACTEMRA.

**Before you receive ACTEMRA, tell your healthcare provider about all of your medical conditions, including if you:**

- have an infection. See “What is the most important information I should know about ACTEMRA?”
- have liver problems.
- have any stomach-area (abdominal) pain or been diagnosed with diverticulitis or ulcers in your stomach or intestines.
- have had a reaction to tocilizumab or any of the ingredients in ACTEMRA before.
- have or had a condition that affects your nervous system, such as multiple sclerosis.
- have recently received or are scheduled to receive a vaccine:
  - All vaccines should be brought up-to-date before starting ACTEMRA.
  - People who take ACTEMRA should not receive live vaccines.
  - People taking ACTEMRA can receive non-live vaccines.
- plan to have surgery or a medical procedure.
- plan to become pregnant or are pregnant. It is not known if ACTEMRA will harm your unborn baby.
- **Pregnancy Registry:** Genentech has a registry for pregnant women who take ACTEMRA. The purpose of this registry is to check the health of the pregnant mother and her baby. If you are pregnant or become pregnant while taking ACTEMRA, talk to your healthcare provider about how you can join this pregnancy registry or you may contact the registry at 1-877-311-8972 to enroll.
- plan to breastfeed or are breastfeeding. You and your healthcare provider should decide if you will take ACTEMRA or breast-feed. You should not do both.

Tell your healthcare provider about all of the medicines you take, including prescription, over-the-counter medicines, vitamins and herbal supplements. ACTEMRA and other medicines may affect each other causing side effects. Especially tell your healthcare provider if you take:

- any other medicines to treat your RA. You should not take etanercept (Enbrel®), adalimumab (Humira®), infliximab (Remicade®), rituximab (Rituxan®), abatacept (Orencia®), anakinra (Kineret®), certolizumab (Cimzia®), or golimumab (Simponi®), while you are taking ACTEMRA. Taking ACTEMRA with these medicines may increase your risk of infection.
- medicines that affect the way certain liver enzymes work. Ask your healthcare provider if you are not sure if your medicine is one of these.

Know the medicines you take. Keep a list of them to show to your healthcare provider and pharmacist when you get a new medicine.

**How will I receive ACTEMRA?**

Into a vein (IV or intravenous infusion) for Rheumatoid Arthritis, PJIA, SJIA, or CRS:

- If your healthcare provider prescribes ACTEMRA as an IV infusion, you will receive ACTEMRA from a healthcare provider through a needle placed in a vein in your arm. The infusion will take about 1 hour to give you the full dose of medicine.
- For rheumatoid arthritis or PJIA you will receive a dose of ACTEMRA about every 4 weeks.
For SJIA you will receive a dose of ACTEMRA about every 2 weeks.
For CRS you will receive a single dose of ACTEMRA, and if needed additional doses.
While taking ACTEMRA, you may continue to use other medicines that help treat your rheumatoid arthritis, PJIA, or SJIA such as methotrexate, non-steroidal anti-inflammatory drugs (NSAIDs) and prescription steroids, as instructed by your healthcare provider.
Keep all of your follow-up appointments and get your blood tests as ordered by your healthcare provider.

Under the skin (SC or subcutaneous injection) for Rheumatoid Arthritis or Giant Cell Arteritis or PJIA or SJIA:
See the Instructions for Use at the end of this Medication Guide for instructions about the right way to prepare and give your ACTEMRA injections at home.
ACTEMRA is available as a single-dose Prefilled Syringe or single-dose Autoinjector (ACTPen™)
You may also receive ACTEMRA as injection under your skin (subcutaneous). If your healthcare provider decides that you or a caregiver can give your injections of ACTEMRA at home, you or your caregiver should receive training on the right way to prepare and inject ACTEMRA. Do not try to inject ACTEMRA until you have been shown the right way to give the injections by your healthcare provider.
For PJIA, you may self-inject with the Prefilled Syringe or your caregiver can give you ACTEMRA if both your healthcare provider and parent/legal guardian find it appropriate.
For SJIA, you may self-inject with the Prefilled Syringe or your caregiver can give you ACTEMRA if both your healthcare provider and parent/legal guardian find it appropriate.
Your healthcare provider will tell you how much ACTEMRA to use and when to use it.

What are the possible side effects with ACTEMRA?
ACTEMRA can cause serious side effects, including:
See “What is the most important information I should know about ACTEMRA?”
Hepatitis B infection in people who carry the virus in their blood. If you 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. Tell your healthcare provider if you have any of the following symptoms of a possible hepatitis B infection:
- feel very tired
- skin or eyes look yellow
- little or no appetite
- vomiting
- clay-colored bowel movements
- fevers
- chills
- stomach discomfort
- muscle aches
- dark urine
- skin rash

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. Tell your healthcare provider before your next dose if you had hives, rash or flushing after your injection. Seek medical attention right away if you have any of the following signs of a serious allergic reaction:
- shortness of breath or trouble breathing
- swelling of the lips, tongue, or face
- chest pain
- feeling dizzy or faint
- moderate or severe abdominal pain or vomiting

Nervous system problems. While rare, Multiple Sclerosis has been diagnosed in people who take ACTEMRA. It is not known what effect ACTEMRA may have on some nervous system disorders.

The most common side effects of ACTEMRA include:
- upper respiratory tract infections (common cold, sinus infections)
- headache
- increased blood pressure (hypertension)
- injection site reactions

Call your doctor for medical advice about 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.

General information about the safe and effective use of ACTEMRA.
Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not give ACTEMRA to other people, even if they have the same symptoms that you have. It may harm them. You can ask your pharmacist or healthcare provider for information about ACTEMRA that is written for health professionals.

What are the ingredients in ACTEMRA?
Active ingredient: tocilizumab
Inactive ingredients of Intravenous ACTEMRA: disodium phosphate dodecahydrate, polysorbate 80, sodium dihydrogen phosphate dihydrate, sucrose, and water for injection.
Inactive ingredients of Subcutaneous ACTEMRA: L-arginine, L-arginine hydrochloride, L-methionine, L-histidine, L-histidine hydrochloride monohydrate, polysorbate 80, and water for injection.