You're Invited

ACTEMRA®
tocilizumab

Brittany L. Goss, D.O.
Queen Of The Valley Medical Center
Napa, CA

Wednesday, June 13, 2018
Arrival time: 6:00 pm
Presentation time: 6:30 pm

at Left Bank Restaurant
377 Santana Row
San Jose, CA 95128

ACTEMRA FOR THE TREATMENT OF RHEUMATOID ARTHRITIS (RA) AND GIANT CELL ARTERITIS (GCA)

Please register by June 8th at http://www.genersvp.com
and reference the unique meeting ID PRF84288
or RSVP to Madeline Morgan at 707-373-7279 or morgan.madeline@gene.com

Please see Important Registration Information on reverse.

INDICATION

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

ACTEMRA is indicated for the treatment of giant cell arteritis (GCA) in adult patients.

BOXED WARNING and Additional Important Safety Information

RISK OF SERIOUS INFECTIONS:

Patients treated with ACTEMRA are at increased risk for developing serious infections that may lead to hospitalization or death, including tuberculosis (TB), bacterial, invasive fungal, viral, or other opportunistic infections. 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.

ACTEMRA is contraindicated in patients with known hypersensitivity to ACTEMRA.

Please see next page and accompanying full Prescribing Information including **Boxed WARNING** for additional important safety information.

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Other serious or potentially life-threatening adverse reactions that have been reported in clinical trials with ACTEMRA include gastrointestinal perforations. Use ACTEMRA with caution in patients who may be at risk for GI perforations.

Laboratory monitoring is recommended due to potential consequences of treatment-related laboratory abnormalities in neutrophils, platelets, lipids, and liver function tests.

Hypersensitivity reactions, including anaphylaxis and death, have occurred.

- If anaphylaxis or other hypersensitivity reaction occurs, stop administration of ACTEMRA immediately and discontinue ACTEMRA permanently.

Avoid use of live vaccines concurrently with ACTEMRA, as clinical safety has not been established.

Other potential risks of ACTEMRA include demyelinating disorders and malignancies.

Treatment with ACTEMRA is not recommended in patients with active hepatic disease or hepatic impairment.

Common adverse reactions included upper respiratory tract infections, nasopharyngitis, headache, hypertension, increased ALT, and injection-site reactions (SC only).

You may report side effects to the FDA at (800) FDA-1088 or www.fda.gov/medwatch. You may also report side effects to Genentech at (888) 835-2555.

Please see accompanying full Prescribing Information, including **Boxed WARNING**, for additional important safety information.

Important Registration Information

This program is intended for Healthcare Professionals and clinical staff with significant clinical patient care responsibilities. In accordance with the PhRMA Code on Interaction with Healthcare Professionals, please understand we are unable to accommodate spouses or guests. Thank you for your cooperation.

Minnesota, Vermont, the **Department of Defense,** and the **Department of Veterans Affairs** have restrictions on receiving in-kind benefits (e.g., meals, parking) at company-sponsored events. You are accountable for understanding such restrictions and complying with them. If you are licensed in or affiliated with any of these states or federal agencies, Genentech policies may restrict you from consuming any portion of the Genentech-sponsored meal at this program or from receiving any other in-kind benefit from Genentech (e.g., parking) in connection with the program.

When you RSVP please indicate whether you will accept or opt out of Genentech's in-kind benefits (e.g., meals, valet parking) at the program. If you choose to opt out you may either pay for the meal and parking on your own, or not consume anything at the program.

For all program attendees who receive Genentech's in-kind benefits at this program, Genentech will report the attendee's name and the value received as required by federal and state disclosure laws (for more information on the federal law please visit http://sunshine.gene.com).

The meal cost may vary by event location and be up to $125 per person (exceptions may apply).

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