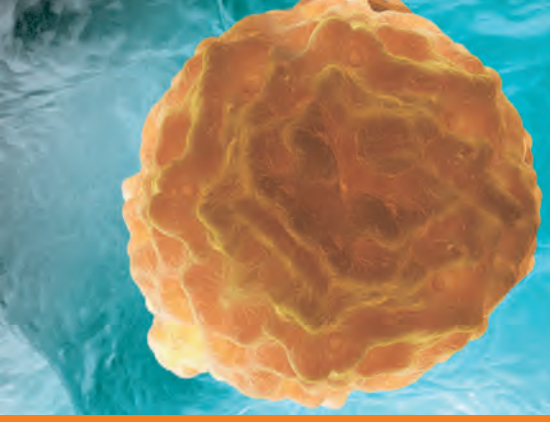

You're Invited!

Rituxan® (rituximab) for the Treatment of Granulomatosis with Polyangiitis (GPA) and Microscopic Polyangiitis (MPA)

Kidney Week 2018
Exhibitor Spotlight

Dr. Gerald Appel

Professor of Medicine
Columbia University
College of Physicians and Surgeons
New York–Presbyterian Hospital
New York, NY

Dr. Gerald Appel is a practicing nephrologist and Professor of Medicine at Columbia University, as well as a member Fellow of American Society of Nephrology. In addition to to being an active member of the nephrology community, Dr. Appel has been a recipient of numerous outstanding teaching awards and is also the author of over 300 publications. Dr. Appel received compensation from Genentech in connection with this presentation.

Date: Friday, October 26, 2018

Time: 12:00 pm - 1:00 pm

Location: *San Diego Convention Center, Exhibit Hall, San Diego, CA*

Supported by: Genentech USA, Inc.

The Exhibitor Spotlight is not a Continuing Education (CE) activity.

Please see Important Attendance Information on reverse.

INDICATION

Rituxan® (rituximab), in combination with glucocorticoids, is indicated for the treatment of adult patients with Granulomatosis with Polyangiitis (GPA) (Wegener's Granulomatosis) and Microscopic Polyangiitis (MPA).

BOXED WARNING

- **Infusion Reactions: Rituxan administration can result in serious, including fatal infusion reactions. Deaths within 24 hours of Rituxan infusion have occurred. Approximately 80% of fatal infusion reactions occurred in association with the first infusion. Monitor patients closely. Discontinue Rituxan infusion for severe reactions and provide medical treatment for Grade 3 or 4 infusion reactions.**
- **Severe Mucocutaneous Reactions: Severe, including fatal, mucocutaneous reactions can occur in patients receiving Rituxan.**
- **Hepatitis B Virus (HBV) Reactivation: HBV reactivation can occur in patients treated with Rituxan, in some cases resulting in fulminant hepatitis, hepatic failure, and death. Screen all patients for HBV infection before treatment initiation, and monitor patients during and after treatment with Rituxan. Discontinue Rituxan and concomitant medications in the event of HBV reactivation.**
- **Progressive Multifocal Leukoencephalopathy (PML), including fatal PML, can occur in patients receiving Rituxan.**

Please see next page and accompanying full Prescribing Information including **BOXED WARNING** for additional Important Safety Information.

Rituxan® Rituximab

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BOXED WARNINGS and Additional Important Safety Information

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Hepatitis B Virus (HBV) Reactivation: HBV reactivation can occur in patients treated with Rituxan, in some cases resulting in fulminant hepatitis, hepatic failure, and death. Screen all patients for HBV infection before treatment initiation, and monitor patients during and after treatment with Rituxan. Discontinue Rituxan and concomitant medications in the event of HBV reactivation.

Progressive Multifocal Leukoencephalopathy (PML), including fatal PML, can occur in patients receiving Rituxan.

Warnings and Precautions

Rituxan administration can also result in additional serious, including fatal, adverse reactions including:

- Tumor lysis syndrome (TLS): Administer aggressive intravenous hydration, anti-hyperuricemic agents, monitor renal function
- Infections: Withhold Rituxan and institute appropriate anti-infective therapy. Rituxan is not recommended for use in patients with severe, active infections
- Cardiovascular adverse reactions: Discontinue infusions in case of serious or life-threatening events
- Renal toxicity: Discontinue in patients with rising serum creatinine or oliguria
- Bowel obstruction and perforation: Consider and evaluate for abdominal pain, vomiting, or related symptoms
- Immunizations: Live virus vaccinations prior to or during Rituxan treatment are not recommended
- Embryo-Fetal toxicity: Can cause neonatal harm. Advise of potential risk to neonates and use of effective contraception
- Use of concomitant immunosuppressants other than corticosteroids has not been studied in GPA, MPA, or PV patients exhibiting peripheral B-cell depletion following treatment with Rituxan
- The safety and efficacy of retreatment with Rituxan have not been established in patients with GPA and MPA

Most Common Adverse Reactions

Granulomatosis With Polyangiitis (GPA) (Wegener's Granulomatosis) and Microscopic Polyangiitis (MPA)

Most common adverse reactions (≥15%) in the clinical study were infections, nausea, diarrhea, headache, muscle spasms, anemia, and peripheral edema. Other important adverse reactions include infusion reactions.

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.

For additional Important Safety Information, please see the Rituxan full Prescribing Information, including **BOXED WARNINGS.**

Important Attendance 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, New Jersey, Vermont, and Federal Entities (e.g., the Department of Defense and the Department of Veterans Affairs) have restrictions on receiving in-kind benefits (e.g., meals, valet 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., valet parking) in connection with 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 sunshine.gene.com).

The meal cost may vary by event location and be up to $150 per person (exceptions may apply).

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

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Rituxan®
Rituximab