



PROGRAM ANNOUNCEMENT

Wednesday, 10 October 2018

Monthly Program

Medical Affairs, Pharmacovigilance, and the Post Approval Landscape

Program Speaker:

Daniel Gharbawy, PharmD

Medical Science Liaison, Seqirus and Publicis Touchpoint Solutions

Event Agenda:

Date: Wednesday, 10 October 2018

Time: 5:15 – 6:00 PM Registration & Snacks
6:00 – 6:05 PM Welcome and Announcements
6:05 – 7:05 PM Program Presentation
7:05 – 7:30 PM Q & A

LOCATION: BD (former CareFusion) (directions on last page of this flyer)
3770 Torrey View Court
San Diego, CA 92130

Program Speaker: Daniel Gharbawy, PharmD

Program Topic:

Interactions between Medical and Regulatory Affairs are a vital part to the success of a pharmaceutical company and their products. The Medical Affairs division is typically most active following approval but can be involved in publication planning, pharmacovigilance, clinical trial design, investigator sponsored research, and observational studies in support of a product and subsequent submissions. This review will highlight the major functions of medical affairs, pharmacovigilance, and how phase IV research has shaped how the influenza vaccine is used in clinical practice.

Program Summary:

This program will focus on the structure and core responsibilities of medical affairs in support of a product pre- and post-approval.

The talk will provide updates on:

- The Medical Affairs Perspective of a Product's Lifecycle

- Responsibilities of Medical Affairs
- Influenza and Prevention
- Current Influenza Vaccines and their Clinical Importance
- FDA Label vs Advisory Committee on Immunization Practices (ACIP) Guidelines
- The Phase IV Landscape and Pharmacovigilance

Speaker Biography:

Daniel Gharbawy has been in the pharmaceutical industry in medical affairs for 9 years. He completed his PharmD at Albany College of Pharmacy and his residency and fellowship at St. Jude Children’s Research Hospital and the University of Tennessee with emphasis in Health Outcomes, Med Safety, and Policy Research. He served as an Associate Professor of Drug Information in the department of Clinical Pharmacy at the University of Tennessee Health Science Center in Memphis, TN before working in the pharmaceutical industry. He served as a Senior Medical Information Specialist and in Medical Affairs with Astra Zeneca, Horizon Pharma, Gilead, Novartis Vaccines, CSL Behring and bioCSL. His expertise has primarily focused on vaccines, but he has also participated in the approval and launch of products in the cardiovascular, rheumatology, gastroenterology and infectious disease therapeutic space. He currently serves as a Medical Science Liaison with Seqirus supporting influenza vaccines and antivirals and interacts regularly with policy and liaison members of the CDC and ACIP. He is passionate about immunizations and eradicating vaccine preventable diseases.

To register for the meeting, click the [link](#) to access the SDRAN registration system (123Signup) for this event.

Please make your reservation early. The online registration due date is Monday, 08 October 2018. Credit Cards are accepted online only. Cash and Check are accepted at the door.

Online registration (through Monday 08 October 2018):

\$15 SDRAN Member	\$25 Non-Member
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Onsite Registration:

\$25 SDRAN Member	\$35 Non-Member
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For Questions Email: programs@sdran.org

Directions to BD (former CareFusion):

From the North 5:

I-5 South toward San Diego
Exit toward Carmel Mountain Rd Left onto Carmel Mountain Rd Left onto Torrey View Court Facility is on the right

From the South 5:

I-5 North toward Del Mar
Merge onto I-5 local Bypass
Take Carmel Mountain Rd exit (32)
Turn right onto Carmel Mountain Rd
Turn left onto Torrey View Court
Facility is on the right