Mark Bogs is the Senior Director of Sterility Assurance at ICU Medical and is responsible for supporting pharmaceutical moist heat terminal sterilization, medical device Gamma, E-Beam, X-Ray and Ethylene Oxide sterilization, and aseptic manufacturing across global ICU Medical sites and third party manufacturing. He has 30 years industry experience in start up aseptic filling operations, established aseptic filling operations for small volume parenterals and Blow Fill Seal, and Ethylene Oxide, Gamma and E-Beam radiation sterilization. Mark has been a board member of PDA Midwest snce 2018, serving as chapter President Elect from 2020-2021, and serving as chapter President from 2022-2023.

He has previously held Global Quality Director roles at Pfizer, Hospira, and Hollister and site QA/RA Director roles at Catalent Pharmaceutical, Dentsply Pharmaceutical, and LifeWatch. He has operations experience as the Manufacturing Manager at the start up of an aseptic pharmaceutical site at Dentsply Pharmaceutical and as the Plant Manager of an Ethylene Oxide sterilization plant for Sterigenics in the Chicagoland area. He began his pharmaceutical career at Abbott Laboratories Hospital Products Division as the Manufacturing QA Supervisor for the Small Volume Parenteral aseptic filling and finishing operation in North Chicago, IL.
Mark holds a BS in Chemistry from Purdue University and an MBA in International Business and Finance from Loyola University of Chicago.