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Medicine

JOIN LEADERS

IN REMS

AbbVie Alexion - Astra Zeneca Rare Disease Unit Amneal Pharmaceuticals Apcer Life Sciences Apellis Pharmaceuticals, Inc. Apotex Ascend Laboratories LLC, a subsidiary of Alkem Laboratories Ltd Avadel Pharmaceuticals Axian Consulting Ltd BioMarin Bristol Myers Squibb Chinook Therapeutices, Inc. Compass Pathways Compliance Architects LLC ConnectiveRx Endo Pharmaceuticals Gilead Sciences GSK

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Venebio Group, LLC

Viatris



REMS Industry Consortium

Innovating Patient Safety. Mitigating Risk.

BECOME A LEADER IN REMS

Join the REMS Industry Consortium today!



WHY RIC?

RIC stands as a trusted and reliable source, offering the latest insights into cutting-edge REMS (Risk Evaluation and Mitigation Strategies) issues. Our consortium membership is comprised of drug and biopharmaceutical companies, as well as REMS program service providers, that are building a bridge for open communication and collaboration among many stakeholders, including our membership, healthcare community members, and FDA.

We unite industry stakeholders in a collective effort to address challenges, sharing invaluable experiences and best practices to guide your REMS programs. By joining RIC, you gain the opportunity to network and build lasting relationships with REMS industry professionals. Most importantly, you become an active contributor to advancing the REMS industry, making a significant impact on the field.

BENEFITS AS A RIC MEMBER



Interact directly with the FDA to find ways to cut down the approval time for REMS innovation as the industry's unified voice



Contribute to REMS industry innovation



Network with REMS industry professionals at programs like the RIC Annual Meeting



Exclusive knowledge-sharing opportunities



Opportunity to help develop leading-edge REMS education



Generate cost savings with hours of free consultation

HAVE YOUR VOICE HEARD THROUGH RIC WORKING GROUPS

BEST PRACTICES

Leverage industry experience to suggest ways to streamline REMS program elements and enhance efficiencies for stakeholders.

COMMUNICATIONS & MARKETING

Conducts qualitative review and feedback for materials produced by all working groups. Develops RIC communications plan, oversight of RIC branding/messaging, and identifies topics for white papers and supports publication.

EDUCATION/CERTIFICATION

Develops educational resources and RIC credentialing programs for various stakeholders, which includes the development of core curriculum and ancillary training modules.

FDA INTERACTIONS

Seeks to engage with FDA by responding to FDA outreach on REMS-related initiatives and preparing recommendations for FDA on ideas generated by working groups and task forces.

INNOVATION & TECHNOLOGY

Explores ways to increase vendor capabilities. Identifies tactics to spur innovation and technology improvement.