

Webinar

Strategies for Foreign Biotech Companies Entering the U.S. Market



Thu, Aug 27th
20:00~ (EDT)

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Seminar Highlights

- Strategies for successful drug development in the U.S.
- Latest FDA regulatory expectations and updates
- Regulatory strategies from early development through marketing approval
- IND, NDA, and BLA submission planning
- FDA meeting preparation and regulatory communication
- Practical solutions to common regulatory challenges

Frank Li, MD, PhD, RAC, as a Featured Speaker



BLA Regulatory is pleased to invite you to this exclusive seminar hosted by R&D Support Center. Join Frank Li, Founder & Principal Consultant of BLA Regulatory, and fellow industry experts as they share practical insights into

successful U.S. drug development, FDA regulatory strategies, and the latest regulatory updates.

Whether you are planning U.S. market entry or advancing your development program, this seminar will provide valuable knowledge to help you navigate today's evolving regulatory landscape.

Meet BLA Regulatory

BLA Regulatory is a global regulatory affairs consulting firm supporting pharmaceutical, biotechnology, and advanced therapy companies throughout the product lifecycle.

Our experienced consultants provide strategic regulatory planning, FDA interactions, regulatory submissions, eCTD publishing, and global regulatory support to help clients accelerate development and achieve successful approvals.

Organizer: R&D Support Center Co., Ltd.