

Clinical Program Leader

Teva Pharmaceuticals Industries Ltd, based in Israel, is the world's leader in generic pharmaceuticals with \$9.3 billion in total revenue and is ranked #13 in overall size in the pharmaceutical industry as a whole. Teva markets products from a wide range of therapeutic areas including respiratory, analgesic, anti-infective, cardiovascular, oncology, CNS, dermatological and anti-inflammatory. Historically, Teva has grown leveraging an aggressive acquisition strategy the result of which is a healthy blended business model of generics, innovative products and active pharmaceutical ingredients. Unlike other pharmaceutical companies today, Teva is continuing to grow at an accelerated pace with plans to be a top 10 global pharmaceutical company within the next few years. The growth plans are aggressive and we are looking for people to help execute this vision.

Teva gained a significant presence in the respiratory therapeutic area through the acquisition of IVAX and is now aggressively building an innovative Global Respiratory R&D organization for the purpose of expanding the product pipeline.

OBJECTIVES

The role of the Clinical Program Leader is critical to the program expansion and success of the clinical development programs which are currently ongoing and planned in the Global Pulmonary Clinical Research Therapeutic Area. Bringing excellent leadership skills and collaborating across teams, this individual will be charged with the responsibility of leading a clinical development program for Teva within the respiratory therapeutic area, including being the key point person and resource for matters related to a program of respiratory clinical trials. The Clinical Program Leader reports directly to the Head of the Therapeutic Area and will manage several direct reports.

RESPONSIBILITIES

- Responsible for approval and oversight of Clinical Development Plans for products in the Pulmonary Clinical Research Therapeutic Area to include:
 - Ensuring quality of the Clinical Development Program
 - Participating in the evaluation of pre-clinical development candidates.
 - Integration of all clinical activities and information within the program
 - Supporting sourcing and clinical evaluation efforts in the indication
- Develop knowledge to support Teva's position in the indication and within the respective medical communities.
- Ensure integrated view across projects, molecules and trials in the program, optimizing resource coordination and management across trials.
- Promote knowledge-sharing across trials within the program.
- Approve the design of specific studies.
- Support marketing, sourcing and clinical evaluation efforts within indications.
- Advise, guide and lead assigned Clinical Leaders.
- Interact with and maintain professional relationships with Key Opinion Leaders (KOLs), Cabinet Advisory Boards (CABs) and Data Safety Monitoring Boards (DSMBs).
- Promote clinical professionalism in the program including good interaction with sites, KOLs and other contacts, closely following GCP's and Teva's SOPs.
- Interact with regulatory authorities as needed.
- Contribute to the preparation of study results for presentation at scientific meetings and manuscripts for publication.

QUALIFICATIONS

- MS in relevant scientific, medical, or clinical discipline. Doctoral level degree (PhD, PharmD, MD or equivalent) strongly preferred.
- Must have excellent knowledge of Allergy/Immunology or Pulmonary Medicine, demonstrating medical and clinical know-how and knowledge of assigned protocols in the indications.
- Minimum of five years' experience in the pharmaceutical industry planning and managing high profile clinical trials, with proven ability to manage projects and/or lead project teams effectively.
- Comprehensive knowledge and understanding of the drug development process including clinical trial design and conduct, FDA and ICH guidelines, GCP regulations and SOPs.
- Ability to interpret study results
- Experience in preparing IND, NDA and other regulatory documents.
- Strong sense of urgency and understanding of time pressures, ability to thrive and enjoy working independently in a fast-paced, multi-tasking and innovative environment.
- High level of initiative and ownership of assigned responsibilities; ability to play a key role in troubleshooting and ensure that issues are resolved in an accurate and timely manner.
- Highly nuanced analytical capabilities; thorough, solution-oriented approach to synthesize complex and diverse information, recognizing trends and/or interrelationships.
- Hands-on approach with a strategic and operational focus; capable decision maker with high degree of flexibility to adapt to company and industry changes.
- Collaborative leader able to operate effectively in a matrix organization and to influence without authority, nurturing professional development of direct reports and creating an environment that encourages ownership, accountability, teamwork and motivation.
- High integrity with solid credibility as an individual and in representing the company.
- Ability to work effectively in a multicultural, international environment at all levels within the organization including supporting clinical development planning with other Teva groups.
- Able to develop and nurture communication between other disciplines, including promotion of knowledge sharing across trials within a program, gaining cooperation and alignment.
- Outstanding communication skills, including effective, clear, concise, well organized and accurate presentations in one-on-one, small group and large audience settings.
- Excellent written skills and ability to present technical information in a clear, understandable manner.
- Ability to be based at Teva USA in Horsham, PA and to travel approximately 30% or as needed.

SEARCH DIRECTOR:

Lucy Watson-Baker, CPC

Partner

Life Sciences Practice Group

(401) 808 8632 or (800) 405 1152 ext. 232

wb@msil.com

Roberta Packer

Director

Life Sciences Practice Group

(413) 448 6035

rpacker@msil.com

PROCESS:

If you would like to learn more about this outstanding opportunity:

1. Return a copy of your updated CV and/or resume and we will contact you to discuss next steps
2. Please send **3 references** with contact information. This will indicate your consent for MSI to conduct background reference checking