Position: **Scientific Advisor (2, Medical Affairs)**

Reports to: **Medical Director**

Summary

The Scientific Advisor role is primarily to support the EU Medical Affairs organization in delivering on its goals through leadership and organization of key medically led activities including advisory boards, congresses, investigator meetings KOL interaction and scientific / medical communications as well as ensuring appropriate dissemination of knowhow and expertise in the company through delivery of scientific training & advice to other functions.

Specific Responsibilities:

- Management of company scientific meetings (advisory boards, symposia etc.)
- Supports Medical Affairs Lead in Congress planning – and ensuring appropriate training is provided to attendees
- Provide medical/scientific support for Aimmune medical, commercial, patient & payor advisory board meetings and other external interactions
- Supports Medical Affairs colleagues in the development and maintenance of effective working partnerships with KOLs
- Oversees the management of the KOL profiling tools and its use to support cross-functional projects (speaker selection/ trial site selection / publications / scientific updates etc.)
- Drive preparation and planning of publications relevant to the EU markets
- Prepare and review materials (including those for use by MDLs/MSLs and for congresses) both at EU and national levels ensuring alignment with applicable codes of practice, laws and regulations as well as company scientific goals
- Manage the medical components of the Value Evidence Generation Plan and EU Medical Affairs Plans
- In partnership with internal subject matter experts, develops and delivers internal training materials on Therapy Area (and Brand data) for head office and field based personnel
- Provides internal training on relevant codes of practice and internal processes to EU staff (and to non-EU staff on EU relevant topics) as required
- Supports Medical Affairs colleagues in successful development and delivery of Medical Education & Disease Awareness programs
- Provides high quality and timely briefing materials/training on new / highly impactful relevant data to the EU cross-functional team
- Provides input to and manages strategic Product Scientific Core Claims Plan
- Consults and works with colleagues from other functions in the development of core Scientific claims plans & publications plans for EU

- Conducts scientific data gap analysis to identify and optimize future marketing potential of the Brand
- Supports delivery of Brand strategies plans through defense of intercompany complaints as appropriate
- Supports Medical Affairs Lead with compilation of effective complaints against competitor promotional campaigns and defend complaints against Aimmune Brands as appropriate
- Supports and reviews, with Medical Affairs Lead, standard letters and Q&As; and provides additional information to support Medical Information enquiry handling process for Brands and Therapy Area
- Understands current and future positioning aims for the Brand and identifies, in partnership with Medical Affairs and Clinical Operations, local/regional clinical activities that support the achievement of those aims (e.g. scientific partnerships, centers of excellence etc.)
- Supports the work of Health Technology Appraisal Preparation Team
- Post-licensing: Support formulary submissions (with appropriate data requirements / and ANDs (Advanced Notification of Drugs to payor bodies)

Qualifications / Requirements:

- UK Registered Pharmacist
- 3+ years of Medical Affairs experience, preferably in an International Environment
- Fluency in English (written and oral) and preferably one other European language

Preferred Experience, Special Skills, Knowledge:

- Experience of review/sign-off of promotional materials in accordance with ABPI code of practice
- Engaged, dynamic, results-driven with ability to critically appraise scientific information to identify key insights quickly
- Strong team work skills, ability to get results in a cross-functional setting, and enjoys operating in a highly motivated team environment
- Solid understanding of the clinical and scientific aspects of therapeutic area;
- Excellent organizational, analytical and problem-solving skills
- Strong interpersonal skills, experience interacting with clinicians/KOLs
- Ability to manage multiple projects simultaneously
- Ability to travel to meetings/trainings/programs as required
- Experience in Allergy and Immunology, preferred
- Experience with scientific and medical communication required

About Aimmune Therapeutics, Inc.

Aimmune Therapeutics is a clinical-stage biopharmaceutical company founded to address the unmet medical need in food allergy, which currently has no approved treatments. Our mission is to improve the lives of people with food allergies, based on our proprietary desensitization treatments in development. Aimmune's lead investigational drug for peanut allergy, AR101, is in Phase 3 clinical testing in North America and Europe. Headquartered in the heart of San Francisco's biotechnology hub, Brisbane, California, Aimmune has additional offices in the Kings Cross area of London, and in Raleigh, North Carolina.

Qualified candidates should forward a resume and cover letter, including a statement of interest, availability, and experience to Human Resources (careers@aimmune.com) with the job title in the subject line.

Aimmune Therapeutics is an Equal Opportunity Employer.

Principals only; no recruiters please.