



Job Title: Associate Medical Director or Medical Director, Clinical Science - Europe
Reports to: VP Clinical Development

ABOUT THE COMPANY

Aimmune Therapeutics is a clinical-stage biopharmaceutical company headquartered in Brisbane, California, and with its European office in central London UK. The company is developing desensitization treatments for food allergies and its lead product for peanut allergy is currently in Phase 3 development with MAA and BLA planned for the end of 2018.

Aimmune Therapeutics offers a competitive compensation and benefits package.

JOB SUMMARY

The Associate Medical Director/Medical Director of Clinical Scientist will be responsible for working as part of the medical team to design, execute and report clinical trials in food allergy.

ESSENTIAL DUTIES:

- Design and execute clinical development plans and protocols for clinical trials in food allergy
- Provide clinical trials expertise to support to study teams and participate in multidisciplinary meetings
- Medical monitoring of clinical trials and provision of medical expertise to the team
- Contribute to the analysis and discussion of study results
- Author key clinical documents, including protocols and study reports
- Contribute to the preparation of documents required for regulatory submissions
- Contribute to scientific publications
- Ensuring compliance with GCP across clinical trials
- Work closely with Regulatory, CMC, and Clinical Operations counterparts
- Develop relationships with investigators and key opinion leaders

QUALIFICATIONS & REQUIREMENTS:

- Level and title for the position will be determined by the qualifications and experience of the candidate.
- A degree combined with industry experience is required. Candidates with an MD should have 5+ years of directly related experience that includes experience in the execution, analysis and reporting of Phase 2/3 clinical trials

- Experience of contributing to the clinical sections of a marketing authorization application
- Experience in allergy, asthma, or immunology preferred
- Thorough understanding of GCP and familiarity with relevant FDA and ICH guidance
- Working knowledge of statistics as applied to clinical trial design and analysis
- Experience in writing and editing scientific research reports

SKILLS & KNOWLEDGE:

- Dedication to the conduct of clinical trials that generate high quality data and ensure patient safety
- Collaborative and accessible
- Strong commitment to goals and timelines
- Ability to absorb new information quickly and gain command of relevant literature
- Preference for working in a fast-paced, team-oriented, environment
- Desire for new professional challenges
- Possessing excellent problem-solving & decision-making skills; strives for appropriate balance between taking initiative and seeking input from others
- Ability to analyze and summarize complex data and information concisely
- Excellent written and verbal communication skills, including fluency in English

Aimmune Therapeutics is an Equal Opportunity Employer.

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.

For more information, please visit www.aimmune.com.