presage BIOSCIENCES

a new era in cancer drug development realized

Clinical Scientist

Our Company

Presage Biosciences, Inc. is an innovative oncology company with an entirely novel platform for assessing new drugs and combinations to treat solid tumors. Our proprietary technology platform, CIVO®, simultaneously analyzes multiple cancer drug candidates and drug combinations within a single living tumor.

Presage is a growing, privately held company led by an experienced management team and is headquartered in the vibrant South Lake Union area of Seattle, WA.

The Role

The Clinical Scientist is responsible for providing scientific support to the Clinical Project Team, leading medical communications, conducting data analyses, and providing scientific input on Presage's preclinical and clinical programs.

Key Responsibilities

- Review and interpret CIVO biomarker data and contribute to data summaries for partners and publications.
- Contribute to the development of quantitative image-based assays of cellular responses within the tumor microenvironment and generate initial data and analyses of these assays.
- Develop subject-matter expertise in relevant tumor biology and pathway target engagement areas including literature review and summarization for team decision-making.
- Support the qualification of internally developed immunohistochemical assays in collaboration with Presage's Histology team and external pathologists.
- Deliver high quality presentations of scientific data to health care professionals in multiple settings, including advisory boards, investigator meetings, and other appropriate venues to enhance understanding and awareness of the CIVO platform and the Presage's Phase 0 clinical trial product.
- Participate in the design, development, and review of clinical trial protocols, amendments, informed consent forms, study guides, plans, case report forms, and any other clinical research related documents (including plans, policies, procedures, work instructions, forms, logs, etc.) and study reports.
- Participate as needed in the preparation of the clinical contribution to INDs, IBs/IFUs, regulatory briefing documents, and other relevant documentation for submission to the FDA and other regulatory authorities.
- Participate in the identification and recruitment of study sites and investigators for clinical sites.
- Manage safety communication and recording, reviewing, and reporting of safety events in concert with the study's medical monitor.

Qualifications

- Graduate level education (PhD) required with at least 3 years of experience in biotech industry.
- Oncology experience required.

- Familiarity with signal transduction and the immune oncology landscape required.
- Experience creating and fostering productive professional relationships and engaging in scientific dialogue with KOLs and other external collaborators.
- Experience compiling and delivering written and oral scientific presentations.
- Ability to develop subject-matter expertise in relevant disease areas.
- Must be able to work well independently and in groups.
- Excellent interpersonal, written, and verbal communication skills.
- Must be comfortable working in a fast-paced, competitive environment with ambiguity and complexity.
- Must be able to handle multiple, unrelated tasks simultaneously.
- Must possess strong attention to detail with excellent analytical, critical thinking and organizational skills.
- A positive attitude and willingness to learn new things.
- Experience with image analysis software (e.g., HALO) and biostatistics a plus.
- Experience with digital pathology, histology, immunohistochemistry, and digital slide scanning a plus.

Work Environment

In-office with the possibility of working remotely for a portion of that time, with prior approval. Travel may be required.

Physical Demands

The physical demands described here are representative of those that must be met by an employee to successfully perform the essential functions of this job. Reasonable accommodations may be made to enable individuals with disabilities to perform the essential functions.

While performing the duties of this job, the employee is occasionally required to stand; walk; sit; use hands to handle or feel objects, tools, or controls; reach with hands and arms; climb stairs; balance; stoop, kneel, crouch or crawl. The employee must occasionally lift and/or move up to 25 pounds. Specific vision abilities required by the job include close vision, distance vision, peripheral vision, depth perception, color vision, and the ability to adjust focus.

Join Us!

If you enjoy a stimulating office environment, are based in the Seattle area, and most importantly, have a passion for improving the lives of patients with cancer, come join us!

To apply for this position, please send your CV with cover letter to:

Presage Biosciences, Inc.
Attn: Human Resources
530 Fairview Ave N, Suite 1000
Seattle, WA 98109
E-mail: hr@presagebio.com

Presage Biosciences, Inc. is an Equal Opportunity Employer.