About Tevogen Bio

We are a clinical-stage specialty immunotherapy company harnessing the power of CD8+ cytotoxic T lymphocytes ("CD8+ CTLs"), one of nature's most powerful immunological weapons, to develop off-the-shelf, precision T cell therapies for the treatment of infectious diseases, cancers, and neurological disorders with the aim of addressing the significant unmet needs of large patient populations.

We believe that personalized immunotherapies which are accessible to the masses are the next frontier of medicine, and a disruptive business model is necessary to increase the pace of medical innovation. Core to the company's successful biopharma business model is its culture of Creativity, Optimism, Respect, Equality, and keen focus on organizational and manufacturing efficiency.

Job Description

Job Title: Senior Director, General Counsel
Work Location: Tevogen Corporate HQ, Warren, NJ
Reports to: Chief Executive Officer
Position: Full-time, Exempt

Essential Duties and Responsibilities:

As a member of the leadership team, you will serve as the principal legal officer and lead the general legal function and compliance function at Tevogen Bio. Your main responsibilities will involve providing oversight across departments, including finance and R&D departments. In addition, your regulatory experience will be leveraged to advise on the evolving regulatory and governance compliance environment.

- Advise and counsel on general securities law, NYSE and NASDAQ matters, and general corporate governance matters.
- Collaborate with internal stakeholders on SEC filings, such as 10-Ks, 10-Qs, 8-Ks, and proxy statements.
- Assist with communications related to trading windows and equity related communications.
- Assist with the preparation of materials to be provided to the Board of Directors and its committees as needed.
- Draft and review amendments to governance documents, corporate policies and procedures, and Board committee charters as needed.
- Provide leadership on cross-functional teams, identify legal issues, and provide legal options/analysis of possible legal solutions.
- Proactively address the rapidly evolving regulatory and governance compliance environment including changes in laws and regulations, regulatory focus areas, and industry best practices.
- Draft, negotiate, and review various research and development agreements, including but not limited to, clinical trial agreements, informed consent documents, clinical supply agreements, quality agreements, investigator-initiated study agreements, contract research organization (CRO) agreements, and confidentiality agreements.
- Be responsible for advising clients on U.S. and ex-U.S. regulatory matters and compliance with laws, regulations, rules, guidance, industry standards, and company policies and procedures that arise through

out the product life cycle, including in the areas of product development, regulatory submissions and strategy, social media, data, research and development, quality system compliance, etc.
- Work with colleagues to develop appropriate and compliant strategies for interacting with healthcare providers, payors, government officials, and other stakeholders.

Education and/or Work Experience Requirements:

- A Juris Doctor Degree with admission to and current good standing with the bar of at least one state or the District of Columbia is required.
- A minimum of 12 years of experience advising business partners in the life-sciences industry while working at a law firm, in a regulatory agency, or as an in-house lawyer.
- Experience in a law firm supporting public companies in the life-sciences industry on securities matters is required.
- Must have experience in a wide variety of areas of U.S. FDA regulatory law for investigational as well as marketed products.
- Demonstrate excellent strategic and analytical skills and the ability to help business partners find creative and compliant solutions to problems.
- Be able to work independently in a fast-paced environment.

Travel Requirements:

This position currently requires onsite work at Tevogen's Warren, NJ HQ. Some domestic and international travel is expected.

We are an equal opportunity employer. All applicants will be considered for employment without attention to race, color, religion, sex, sexual orientation, gender identity, national origin, veteran, or disability status.

Prospective new hires for this position must be fully vaccinated against COVID-19 as a condition of employment. Vaccine verification will be required on your start date unless an exemption has been approved by the Company as a medical or religious accommodation.

Tevogen Bio Plans to Become Publicly Listed on NYSE via Business Combination with Semper Paratus Acquisition Corporation. Learn more here: *https://www.globenewswire.com/news-release/2023/06/29/2696959/0/en/Tevogen-Bio-to-Become-Publicly-Listed-on-NYSE-via-Business-Combination-with-Semper-Paratus-Acquisition-Corporation.html*