JOB TITLE: Senior Scientist, CMC Analytical

DEPARTMENT: Technical Operations (On-site)

REPORTING TO: Senior Director, Analytical Science and Quality Control

DATE: September 2023

The Company:

Cassava Sciences, Inc. located in Austin, Texas (USA) is a pharmaceutical company focused on neuroscience. Our small molecule, solid oral dosage form product is in late Phase 3 clinical stage development, targeting Alzheimer's disease.

The Position:

This senior role within Technical Operations is an on-site, office-based position and requires strong skills in analytical chemistry, ability to author technical and regulatory filing documents and demonstrated practice of the cGMP and other applicable FDA regulations.

Responsibilities:

• Subject matter expert in various analytical techniques required in support of drug substance and drug product development.
• Actively participate in the validation, transfer and management of early to late-phase QC methods.
• Works independently or collaboratively with cross-functional teams to develop and meet goals and timelines associated with departmental deliverables.
• Ensures work is compliant with cGMP and relevant regulatory requirements.
• Prepare and review Technical Reports, presentations, SOPs and regulatory dossier sections pertaining to CMC Analytical Chemistry.
• Makes sound scientific decisions and serves as back-up to management.

Requirements:

M.Sc. in chemistry or a related scientific field is required.

A minimum of eight years of analytical development and/or quality control experience within the pharmaceutical industry or other relevant experience is required for this role.

Excellent verbal, written and presentation skills.
Contact:

jstroe@cassavasciences.com
No consultants, no recruiters please.