OHOP/OND/CDER/FDA (Office of Hematology and Oncology Products/Office of New Drugs/Center for Drug Evaluation and Research/U.S. Food and Drug Administration) is seeking one clinical analyst to conduct clinical safety data analyses to support drug reviews. Key skills include SAS, R, and/or Python. CVs should be sent to Jinzhong.Liu@fda.hhs.gov.

Job Title: Clinical Analyst
Appointment Type: Full-time Employee
Job Purpose: Conduct clinical safety data analyses to support drug reviews
Location: Silver Spring, MD.
Department: Food and Drug Administration (FDA), Center for Drug Evaluation and Research (CDER), Office of New Drugs (OND), Office of Hematology and Oncology Products (OHOP)

Key Responsibilities:
1. Check clinical data sufficiency and integrity.
2. Verify clinical safety data.
3. Conduct in-depth clinical safety analyses.

Key Skills:
1. SAS
2. R and/or Python

Preferred Qualifications:
1. Master or Ph.D. degree in Biostatistics or related fields
2. U.S. permanent residency or U.S. citizen

A curriculum vitae should be sent to Dr. Liu (Jinzhong.Liu@fda.hhs.gov).