

- Reviews documents submitted for regulatory action; provides secondary review of documents prepared by other OS scientists.
- Demonstrates technical leadership by: (1) interpreting complex biological, pharmacological, and toxicological aspects of reports/submissions for tobacco products; (2) reviewing study protocols and offering recommendations related to study design; (3) interpreting scientific data and if needed, performing additional analysis of data submitted; (4) preparing a comprehensive synopsis of reviews of reports/applications with recommendations for revision, acceptance or rejection, and providing specific supporting reasons for the reviewer's recommended disposition and any technical deficiencies requiring action by the sponsor; (5) implementation of policies and recommendations.
- Provides advanced expertise by keeping abreast of new findings through review of scientific literature, participation in professional meetings and undertaking independent research. Proposes areas of study for regulatory research projects in the Environmental Science Branch. Develops state-of-the-art projects in regulatory science to address gaps in scientific knowledge needed for effective regulation of tobacco products.
- Recognizes the need for, and initiation of, new and amended regulations, policies, guidances and procedures. Develops and recommends new and revised guidelines for regulated products. Provides scientific support in developing guidance and regulation.
- Provides expert advice and assistance to scientists and officials on a wide range of matters. Provides verbal and written conclusions to other federal agencies, industry, universities and state, local and foreign governments. The incumbent also compiles data to prepare presentations to support the Agency's recommendations on scientific issues.
- Ensures that the organization's strategic plan, mission, vision, and values are communicated to a team and integrated into the team's strategies, goals, objectives, work plans and work products and services.
- Articulates and communicates to the team the assignment, project, problem to be solved, actionable events, milestones, and/or program issues under review, and deadlines and time frames for completion.
- Leads a team in identifying, distributing and balancing workload and tasks among employees in accordance with established work-flow, skill level and/or occupational specialization; making adjustments to accomplish the workload in accordance with established priorities to ensure timely accomplishment of assigned team tasks; and ensuring that each employee has an integral role in developing the final team product.
- Reports to the supervisor periodically on team and individual work accomplishments, problems, progress in mastering tasks and work processes, and individual and team training needs.
- Represents the team consensus and convey the team's findings and recommendations in meetings and dealings with other supervisors, program officials, the public, and other customers on issues related to or that have an impact on the team's objectives, work products, and/or tasks.
- Supervises a team in assessing its strengths and weaknesses and provide leadership in exploring alternatives and determining what improvements can be made (e.g., in work methods, processes and procedures).

- Supervises a team of scientists. Plans and schedules work to be accomplished by subordinates, sets and adjusts long and short-term priorities and prepares schedules for completion of work, when necessary. Assigns work to subordinates based on priorities, considering difficulty and requirements of assignments as well as the capabilities of employees.
- Performs other duties as assigned.

Desired Knowledge and Skills:

- Skill in applying the theories, principles, and methods in the field of biology, pharmacology, and toxicology to provide technical expertise and leadership to a team.
- Demonstrates skill in identifying problems, gathering information, drawing conclusions, recommending solutions, preparing papers and reports for publication, providing advice to other scientists, and negotiate acceptance and implementation of recommendations.
- Knowledge of CTP missions, programs and organizations structures sufficient to collaborate with other CTP staff on public health issues and problems.

Application Procedure:

Supervisory concurrence is required to accept a detail; however, is not required to apply. The detail opportunity is open to all qualified candidates at the GS-13 and GS-14 grade levels or US PHS Commissioned Corps Officers.

Interested applicants should submit a copy of their resume, transcripts, most recent copy of SF-50 (Notification of Personnel Action) that identifies your current pay plan, series, grade, full performance level, and time in grade, and a statement indicating the interest in being considered for this detail via email to:

CTP-Recruitment@fda.hhs.gov

Please enter in the subject line:

Detail: CTP, OS, DNCS – Supervisory Interdisciplinary Scientist, GS-401/415-14.

Travel Expenses will not be paid.

Applications/resumes must be submitted by April 22, 2024.

This is not an official vacancy announcement under the Merit Promotion System.