

**REIMBURSABLE DETAIL**  
**Center for Tobacco Products**  
**Office of Science**

The Center for Tobacco Products (CTP), Office of Science, Division of Individual Health Science (DIHS) is offering a Detail opportunity for a position in one of the following series: Supervisory Pharmacologist, GS0405; Supervisory Health Scientist, GS-0601; Supervisory Physician, GP-0602; (Deputy Director) GS-15. Applicants at the GS-14 level are encouraged to apply. The Detail is available immediately for a period of 120 days. PHS Commissioned Corps Officers may apply. A temporary promotion may be considered.

**Bargaining Unit Status:** Non-Bargaining Unit Position

**Position:** Deputy Director, DIHS

**Office Location:** Food and Drug Administration  
Center for Tobacco Products  
Calverton Tower  
11785 Beltsville Rd  
Beltsville, MD 20705

**Opening Date:** **Friday October 23, 2020**

**Closing Date:** **Friday, October 30, 2020**

**Area of Consideration:** **CTP-Wide**

The Center for Tobacco Products, Office of Science offers a fast-paced, dynamic environment and an opportunity to work with dedicated, energetic people who really want to make a difference and improve public health. The position is ideal for someone who wants to have a critical role in the organization and would enjoy the challenge of handling a variety of assignments related to the regulation of tobacco products.

The incumbent serves as the Deputy Director in the Division of Individual Health Science (DIHS). The incumbent directs, manages, and plans the Division programs that ensure the goals, mission and programs of the Center for Tobacco Products are supported by an aggressive and scientifically rigorous program for targeted populations and public health. The incumbent remains cognizant of and reviews and evaluates the state of individual harm resulting from the use of tobacco and ongoing research underway throughout the scientific community, identifies emerging problems and areas of primary interest to FDA, works closely with the Division Director, Deputy Director for Regulatory Science and Office Director to develop research programs to be undertaken in CTP and establishes priorities for their implementation.

**Duties Include:**

- Supervises a staff of 10 or more health, behavioral, pharmacological and clinical scientists by providing scientific technical and administrative direction.
- Plans and schedules work to be accomplished by subordinates, sets and adjusts long and short-term priorities and prepare schedules for completion of work, when necessary.
- Develops performance standards and evaluates work performance of subordinate staff.
- Identifies ways to improve production or increase the quality of work directed.
- Serves as principal advisor and leading authority to the Office Director on tobacco medical issues.
- Maintains close personal contact with the “state of science” in order to inculcate the most advanced theories and practices the medical and scientific research matters Office/Center programs.
- Serves as senior technical expert advisor for establishing programs and policies regarding the analysis and interpretation of medical, clinical, toxicological, pharmacological, and behavioral data collected from industry, private sources and other nations.
- Contributes to the preparation of scientific papers, congressional correspondence and testimony, Federal Register documents, technical reports and private sector information programs.
- Shares policies and promotes collaboration with other Centers within the Federal establishment to obtain interest and collaborative support in proposed and ongoing research projects.
- Serves as technical expert with respect to all phases of medical, behavioral and clinical pharmacological and interpretation relating to topics under consideration.
- Writes and/or presents comprehensive technical medical, pharmacological, behavioral and clinical pharmacology reports prepared based on the data collected and provides professional advice and consultation to public health professionals, senior scientists and management officials.
- Prepares comprehensive reports of study results which include discussion of substantive health issues and research objectives; assessment of the adequacy and quality of data used in the analyses; and explanation of the methodologies, results, and relevance to health issues as they relate to the behavioral and clinical pharmacology problem under study.
- Disseminates results of research projects in a wide range of venues including publications, peer reviewed journals, summaries, manuscripts, and special reports.

**Desired Knowledge and Skills:**

- Mastery of the theories, principles and methods in tobacco-related behavioral and clinical pharmacology matters and associated scientific disciplines sufficient to review a variety of complex industry applications and to apply new scientific and technological developments to novel and critical problems which cannot be solved by the use of conventional methods.

- Knowledge of scientific developments and current practices in behavioral and clinical pharmacology through review of published literature, tobacco industry documents, and by continuing professional development through interactions with experts in the field.
- Experience preparing papers and reports for presentation to include those for professional groups and for publication in professional journals.
- Knowledge of CTP missions, programs, and organizational structures sufficient to collaborate with other CTP staff on public health issues and problems.
- Ability to communicate effectively to accurately represent the FDA/CTP and the assigned program area in dealing with representatives of other agencies and organizations.
- Excellent organizational skills.
- Excellent oral and written communication skills.
- Exceptional interpersonal relationship skills and ability to collaboratively lead teams (e.g., maximize each person's contributions, reconcile divergent viewpoints, and maintain harmonious working relationships).

**Application Procedure:**

Supervisory concurrence should be obtained before you apply to this Detail; however, is not required. The Detail opportunity is open to all qualified candidates at the GS-14 grade levels or Commissioned Corps Officers (O6).

Please enter **Detail: CTP, OS Deputy Director, DIHS GS-15 (October)** in the subject line of e-mail.

Interested applicants should submit a copy of their resume, most recent copy of SF-50, copy of unofficial transcripts, and statement of interest via email to:

R. Michele Martin  
Program Analyst  
Office of Management, Center for Tobacco Products, FDA  
[Rebecca.Martin@fda.hhs.gov](mailto:Rebecca.Martin@fda.hhs.gov)

Detail is reimbursable.  
Travel Expenses will not be paid.

**Candidates must express interest by Friday, October 30, 2020.**

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