

REIMBURSABLE DETAIL
Center for Tobacco Products
Office of Science

The Center for Tobacco Products (CTP), Office of Science (OS), Division of Regulatory Science Informatics (DRSI) is offering a Detail opportunity to a Supervisory Regulatory Health Information Specialist GS-0601-14. The Detail is available immediately for a period up to 120 days. Multiple selections may be considered from this announcement. Applicants at the GS-13 and GS-14 are encouraged to apply. A temporary promotion may be considered.

Bargaining Unit Status: Non-Bargaining Unit Position

Office Location: FDA
Center for Tobacco Products
Office of Science
11785 Beltsville Drive
Beltsville, MD 20705

Opening Date: **December 17, 2021**

Closing Date: **December 23, 2021**

Area of Consideration: FDA-Wide

The CTP, OS 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. This Regulatory Health Information Specialist position is ideal for someone who wants to have a critical role in the organization and would enjoy the challenge of Information Technology (IT) software and systems development, operations and maintenance, systems design and support, data management, data analysis, and data visualization.

Duties include:

The Detail will be located in OS, DRSI, Data and Systems Branch. The primary role of the division is to strategically develop IT solutions to support the regulatory and scientific reviews of tobacco products for the OS. Duties for this position may include:

- Leads and supervises a team of individuals who manage regulatory information systems that support tobacco product review.
- Analyzes current and projected needs of regulatory reviewers, supervisors and managers to determine requirements for development and enhancement of information systems and reports generated from the systems to include directing activities with appropriate agency and contractor personnel.
- Provides management reports and presentations on various aspects of products regulated by the Office.
- Provides written analysis of the data flow and processes associated with tobacco product review and knowledge management to determine areas needing improvement and provide recommendations to management.

- Provides direction and oversight of computerized regulatory information/data pertaining to products regulated by the Office.
- Provides a thorough analysis of regulatory review and scientific health information/data needs of the Office in relation to existing capabilities for providing information to OS staff on a variety of topics associated with the regulatory review process, research and knowledge management.
- Provides advice and is responsible for coordinating the development and distribution of periodic and special reports pertaining to product review status and other matters related to various regulatory review, research and knowledge management activities handled by the Office.
- Fosters collaboration and communication within the teams, branches, DRSI, and OS.
- Performs other similar duties as assigned.

Desired Knowledge and Skills:

- Mastery knowledge management and other OS business functions that require mastery knowledge in information management, automated processes and data management.
- Mastery of a wide range of analytical and evaluative theories, database analysis, methods and procedures applicable to evaluating the effectiveness of the Office’s information management and reporting mechanisms for determining appropriate enhancements/improvements to the systems.
- Expert knowledge of the information contained in the database systems and mastery of the interrelationships between various data elements in the Office’s information systems and the ability to use that knowledge to: 1) evaluate the systems; 2) produce comprehensive reports concerning products regulated and business activities of the Office; and 3) provide quality assurance of the data and reports generated from the systems.
- Mastery of the regulatory review process for the products regulated by the Center to identify the need for specific tools or IT support mechanisms.
- Extensive knowledge of the scientific and regulatory requirements in the product submissions regulated by the Office to identify specific data needs and to identify areas where the reporting capabilities of data systems need to be enhanced or modified.
- Advanced knowledge and skill of SharePoint, or similar platforms, to facilitate collaborative efforts and knowledge management including, but not limited to, developing, managing content and testing functionality to provide direct, hands-on support of CTP’s SharePoint sites for scientific review programs and related activities.
- Knowledge of how the various scientific disciplines included in the Office’s regulatory responsibilities interrelate with each other to recognize the need to change data elements to reflect management needs.
- Excellent organizational skills, and effective communication skills both verbal and written.
- Ability to foster accountability and commitment to the mission of the Division.

Application Procedure:

This Detail opportunity is open to all qualified candidates at the GS-13 to GS-14 grade level or Commissioned Corps Officers equivalent. A temporary promotion may be available. Interested applicants should submit a copy of their resume, most recent copy of SF-50, a copy of transcripts, and statement indicating the reason for interest in being considered for this Detail

via email to:

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And

Miranda Jones
Program Analyst
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Please enter **Detail: CTP, OS, DRSI - Supervisory Regulatory Health Information Specialist GS-0601-14 (December)** in the subject line of e-mail.

Detail is reimbursable.
Travel Expenses will not be paid.

The employee will work from the CTP, OS duty station in Beltsville, MD.

Candidates must express interest by December 23, 2021.

Supervisory concurrence is required to accept a Detail; it is not required to apply.

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