



Title 21 Vacancy Announcement
U.S. Department of Health and Human Services (HHS)
Food and Drug Administration (FDA)
Center for Drug Evaluation and Research (CDER)
Office of Surveillance and Epidemiology (OSE)

Application Period: February 28, 2023 – March 13, 2023

Area of Consideration: United States Citizenship is required. You must be a U.S. Citizen or U.S. National. Foreign nationals or legal permanent residents are not eligible for consideration.

Position: Associate Director for Oncology

Series: AD- 0601

Location(s): Silver Spring, MD

Salary: Starting at \$155,700

Work Schedule: Full Time

Cures Band(s): Band E

Full Performance Band Level: Band E

Travel Requirements: 25% or less

Bargaining Unit: 8888

Relocation Expenses Reimbursement: You may qualify for reimbursement of relocation expenses in accordance with agency policy.

This position is being filled under a stream-lined hiring authority, Title 21, Section 3072 of the 21st Century Cures Act. The candidate selected for this position will serve under a career or career-conditional appointment and be paid under the provisions of this authority.

Additional information on 21st Century Cures Act can be found here:

[**21st Century Cures Act Information**](#)

Introduction

The Food and Drug Administration (FDA) is the regulatory, scientific, public health and consumer protection agency responsible for ensuring all human and animal drugs, medical devices, cosmetics, foods, food additives, drugs and medicated feeds for food producing animals, tobacco and radiation emitting devices are safe, and effective.

The Center for Drug Evaluation and Research (CDER), is responsible for regulating prescription drugs, including new drugs, generic drugs, biological products and biosimilars as well as over-the-counter drugs (OTC). CDER's drug regulatory responsibilities include premarket review of

new drugs and generic drugs; maintenance of the OTC drug monograph system; monitoring of all marketed drug safety and promotional activities; review, monitoring, and enforcement of drug quality during the entire drug life cycle; and ensuring drug products in the market comply with the law.

The mission of the Office of Surveillance and Epidemiology (OSE) is to detect, assess, prevent, and manage the risks of medications so that they can be relied upon to treat disease and improve health using four core functions: Pharmacovigilance, Pharmacoepidemiology, Risk Management, Medication Error Prevention and Analysis.

The mission of the respective divisions in the Office of Pharmacovigilance and Epidemiology (OPE) is to protect the public using epidemiologic evidence to assess the safety and effectiveness of drugs and biologics by detecting, assessing, and evaluating the safety and effectiveness of drugs and biologics using observational methods; and conducting drug and biologic safety and effectiveness surveillance and research using the best available epidemiologic methodologies. This position is in the Division of Epidemiology I (DEPI I), Office of Pharmacovigilance and Epidemiology (OPE), Office of Surveillance and Epidemiology (OSE), Center for Drug Evaluation and Research (CDER).

Duties/Responsibilities

- As the **Associate Director of Oncology**, the incumbent applies knowledge of administrative and program management principles and skills to carrying out the mission of the Division. Seeks and develops the most cost effective and fiscally responsible methods to conduct the Division programs and to solve unusual and often precedent setting problems associated with the Division programs. Provides occupational specific technical and administrative direction and supervision 25 percent or more of the time to subordinate staff performing the work and functions of the organizational unit. Obtains resources and identifies strategic objectives for the organization.
- In collaboration with the Division Leadership, provides supervisory oversight and direction for all the Division's activities related to the safety and effectiveness (related to real-world evidence) of oncology drug and biologic products, and other therapeutic areas as needed. Initiates decision-making processes and documents, and participates in discussions and decisions concerning Division, Office, and Center plans, programs, and activities, both in strategic planning and in the actual determination, allocation, and administration of program segment(s), functions, and activities. Provides authoritative advice and assessments of the impact of actual and proposed Administration or Congressional actions on the programs, functions, and activities of the Division.

- In collaboration with Division leadership, develops and implements Division policies and plans, and makes critical decisions and provides expert advice and counsel concerning approaches and options that are sound and feasible in relation to Office and Center goals and objectives and Federal budgetary and economic realities. Continually evaluates budget, fiscal, and administrative controls to manage Division programs and services. Develops and makes recommendations for the enhancement and improvement of the mission and functions of the Division.
- Represents the Division and Office in dealing and negotiating with individuals representing organizations such as the Congress; other Federal agencies; State, local, and foreign governments; the regulated industry; professional and industry organizations; and public interest groups. Directs the preparations, clearance, and finalization of Division responses to inquiries covering all aspects of the program segment(s), functions, and activities of the Division, particularly as these refer to the safety and effectiveness of oncology products and other therapeutic areas as needed.
- Directs the preparation of analyses of the impact of proposed changes to Agency laws and regulations which affect the functions, program segment(s), and activities of the Division, especially with regard to the safety and effectiveness of oncology products and other therapeutic areas as needed. Decides on changes and additions to the functions, program segment(s), and activities of the Division necessary to implement new legislation or regulations and develops various scenarios for dealing with expansion or contraction of Division functions, program segment(s), and activities.

Supervisory Responsibilities: Applies a knowledge of administrative and program management principles and skills to carry out the mission of the Division as well as to address and solve unusual and often precedent setting problems associated with the Division programs. Seeks and develops the most cost effective and fiscally responsible methods to conduct these programs and to solve these problems.

Provides occupational specific technical and administrative direction and supervision to 25 percent or more of the time to subordinate staff performing the work and functions of the organizational unit. Obtains resources and identifies strategic objectives for the organization.

Conditions of Employment

- U.S. Citizenship requirement or proof of being a U.S. National must be met by closing date.
- Employment is subject to the successful completion of a background investigation, verification of qualifications, completion of onboarding forms, submission of required documents, and any other job-related requirement before or after appointment.

- Applicants must meet all qualification requirements by the closing date of this announcement.
- Direct Deposit: You will be required to have all federal salary payments electronically deposited into a bank account with a financial institution of your choice.
- FDA participates in e-Verify: All new hires must complete the I-9 form; this information will be processed through e-Verify to determine your employment eligibility. If a discrepancy arises, you must take affirmative steps to resolve the matter.
- Males born after December 31, 1959, must be registered with the Selective Service.
- One-year probationary period may be required.
- Financial Disclosure may be required.
- Ethics Clearance may be required.
- Background Investigation/Security Clearance is required. All employees must pass a security investigation. Failing to pass the background check may be grounds for removal or legal action. If hired, you may be subject to additional investigations at a later time.

Qualifications

To be placed into a Cures position, candidates must meet the following criteria:

1. Scientific, Technical, and Professional Fields
2. Qualified and Outstanding Candidates
 - a. **Qualified** applies to all candidates for Cures appointments. The FDA OTS will use the basic requirements defined in the [OPM Qualification Standards](#) as a baseline for comparing experience levels and other candidate attributes for relevant positions.
 - b. **Outstanding** candidates can be defined by existing outstanding work experience, outstanding performance rating, or both.

To qualify for this Title 21 Cures position, the candidate(s) must meet the following **required** qualifications. *Please note: Additional education and experience listed that is not indicated as required is preferable and desired. Candidates who do not meet the “desired” criteria will not be excluded from consideration for this position.*

Education Requirement:

Associate Director for Oncology, AD-0601 Series

Minimum Education Requirement: Meets the Office of Personnel Management (OPM)

Individual Requirements (IOR) for [OPM Occupational Series Qualification Requirements](#).

Desired Education:

Our ideal candidate will possess: A bachelor’s or graduate/higher level degree: major study in an academic field related to the medical field, health sciences or allied sciences appropriate to the work of the position. This degree must be from an educational program from an accrediting body recognized by the U.S. Department of Education (<https://www.ed.gov/>) at the time the degree was obtained.

Professional Experience:

Our ideal candidate will possess:

- Knowledge of the FDA Amendments Act, and the regulations and policies promulgated under that statute.
- Advanced knowledge of science (e.g., biology, regulatory science, drug safety, pharmacoepidemiology).
- Demonstrated ability to clearly communicate orally and in writing.
- Skill in working with a variety of people at all levels.
- Skill in planning, organizing, and leading the work of other professional staff.

Desired Professional Experience:

Our ideal candidate will have expertise in the use of real-world evidence for both efficacy and safety, use of real-world evidence in decision making, and experience in the oncology therapeutic area.

Education Transcripts

SUBMITTING YOUR TRANSCRIPTS: Positions which are scientific or technical in nature often have very specific educational requirements. A transcript is required to verify educational achievement. Pay careful attention to the Qualifications and Education sections to identify vacancies where a transcript is required. Even if you hold a similar position or are a current FDA employee, you are not exempt from transcript requirements.

FOREIGN EDUCATION: If you are using education completed in foreign colleges or universities to meet the qualification requirements, you must show that the education credentials have been evaluated by a private organization that specializes in interpretation of foreign education programs and such education has been deemed equivalent to that gained in an accredited U.S. education program; or full credit has been given for the courses at a U.S. accredited college or university. For more information about this requirement, please visit the [U.S. Department of Education website for Foreign Education Evaluation](#).

Security Clearance Requirements

Background Investigation/Security Clearance Requirements: Non-Sensitive/High Risk

If not previously completed, a background security investigation will be required for all appointees. Appointment will be subject to the applicant's successful completion of a background security investigation and favorable adjudication. Failure to successfully meet these requirements may be grounds for appropriate personnel action. In addition, if hired, a background security investigation or supplemental investigation may be required later.

Vaccination Requirements

To ensure compliance with an applicable preliminary nationwide injunction, which may be supplemented, modified, or vacated, depending on the course of ongoing litigation, the Federal Government will take no action to implement or enforce the COVID-19 vaccination requirement pursuant to Executive Order 14043 on Requiring Coronavirus Disease 2019 Vaccination for Federal Employees. Therefore, to the extent a federal job announcement includes the requirement that applicants must be fully vaccinated against COVID-19 pursuant to Executive Order 14043, that requirement does not currently apply. Federal agencies may request information regarding the vaccination status of selected applicants for the purposes of implementing other workplace safety protocols, such as protocols related to masking, physical distancing, testing, travel, and quarantine.

Ethics Clearance Requirements

This position may require financial disclosure reporting and will be subject to FDA's prohibited financial interest regulation. If you are hired, you may be required to divest of certain financial interests. You are advised to seek additional information on this requirement from the hiring official before accepting any job offers. For more information please visit the FDA Ethics web page: <https://www.fda.gov/about-fda/jobs-and-training-fda/ethics>.

Equal Employment Opportunity

Equal Employment Opportunity Policy

The United States Government does not discriminate in employment on the basis of race, color, religion, sex (including pregnancy and gender identity), national origin, political affiliation, sexual orientation, marital status, disability, genetic information, age, membership in an employee organization, retaliation, parental status, military service, or other non-merit factor.

[Equal Employment Opportunity \(EEO\) for federal employees & job applicants](#)

Reasonable Accommodation

Reasonable Accommodation Policy

Federal agencies must provide reasonable accommodation to applicants with disabilities where appropriate. Applicants requiring reasonable accommodation for any part of the application process should follow the instructions in the job opportunity announcement. For any part of the remaining hiring process, applicants should contact the hiring agency directly.

Determinations on requests for reasonable accommodation will be made on a case-by-case basis. A reasonable accommodation is any change to a job, the work environment, or the way things are usually done that enables an individual with a disability to apply for a job, perform job duties or receive equal access to job benefits.

Under the Rehabilitation Act of 1973, federal agencies must provide reasonable accommodations when: An applicant with a disability needs an accommodation to have an equal opportunity to apply for a job. An employee with a disability needs an accommodation to

perform the essential job duties or to gain access to the workplace. An employee with a disability needs an accommodation to receive equal access to benefits, such as details, training, and office-sponsored events. You can request a reasonable accommodation at any time during the application or hiring process or while on the job. Requests are considered on a case-by-case basis. Learn more about [disability employment and reasonable accommodations](#) or [how to contact an agency](#).

E-Verify

The Food and Drug Administration participates in the USCIS Electronic Employment Eligibility Verification Program (E-Verify). E-Verify helps employers determine employment eligibility of new hires and the validity of their Social Security numbers.

How to Apply

How to Apply: Submit resume with cover by **March 13, 2023**, to: OSE-PMAS-Admin-Team@fda.hhs.gov. Candidate resumes may be shared with hiring official within CDER with a similar job vacancy. Candidates can opt out of this process by annotating resume with “do not share”. Job Reference ID: **ADO-DEPIIMar2023**.

Announcement Contact

For questions regarding this Cures position, please email OSE-PMAS-Admin-Team@fda.hhs.gov.

The U.S. Department of Health and Human Services is an equal opportunity employer with a smoke free environment.

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