

**Opportunity Title:** FDA NCTR Summer 2025 Student Research Participation Program **\*\*CLOSED\*\***  
**Opportunity Reference Code:** FDA-NCTR-2025-0001

**Organization** U.S. Food and Drug Administration (FDA)

**Reference Code** FDA-NCTR-2025-0001

**How to Apply** *Connect with ORISE...on the GO!* Download the new ORISE GO mobile app in the Apple App Store or Google Play Store to help you stay engaged, connected, and informed during your ORISE experience and beyond!

A complete application consists of:

- An application
- Transcripts – [Click here for detailed information about acceptable transcripts](#)
- A current resume/CV, including academic history, employment history, relevant experiences, and publication list
- One educational or professional recommendation. Your application will be considered incomplete, and will not be reviewed until one recommendation is submitted.

All documents must be in English or include an official English translation.

If you have questions, send an email to [ORISE\\_FDA.NCTR@orau.org](mailto:ORISE_FDA.NCTR@orau.org). Please include the reference code for this opportunity in your email.

**Application Deadline** 2/15/2025 11:59:00 PM Eastern Time Zone

**Description** \*Applications will be reviewed on a rolling-basis.

**FDA Office and Location:** Summer research opportunities are available at the National Center for Toxicological Research (NCTR), U.S. Food and Drug Administration (FDA) in Jefferson, Arkansas. Selected individuals will train alongside NCTR investigators who conduct FDA mission-critical, translational research that provides a scientifically sound basis for FDA regulatory decisions, and that reduces risks associated with FDA-regulated products. For additional information about this opportunity, NCTR or FDA, please visit [www.FDA.gov](http://www.FDA.gov).

**Anticipated Appointment Start Date:** May 27th, 2025 and end date August 1st, 2025. Start date is flexible and will depend on a variety of factors.

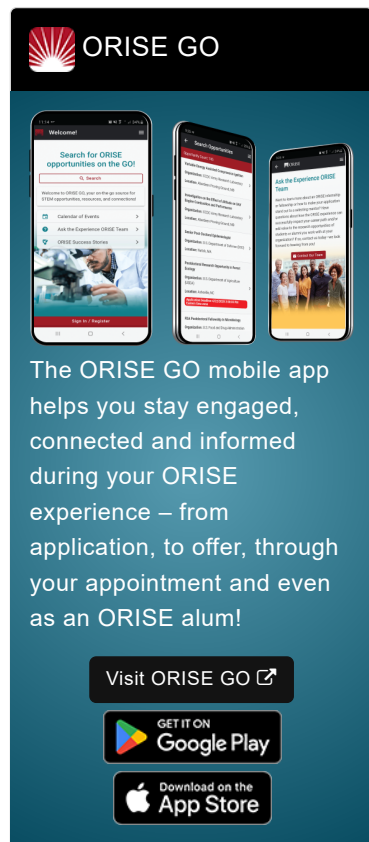
**Appointment Length:** The appointment will initially be for ten weeks, but may be renewed upon recommendation of FDA and is contingent on the availability of funds.

**Level of Participation:** The appointment is full time.

**Citizenship Requirements:** This opportunity is available to U.S. citizens only.


This program, administered by ORAU through its contract with the U.S. Department of Energy to manage the Oak Ridge Institute for Science and Education, was established through an interagency agreement between DOE and FDA. The participant will receive a monthly stipend commensurate with educational level and experience. Proof of health insurance is required for participation in this program. Participants do not become employees of FDA, DOE or the program administrator, and there are no employment-related benefits.


Completion of a successful background investigation by the Office of Personnel Management is required for an applicant to be on-boarded at FDA. OPM can complete a background investigation only for individuals, including non-US Citizens, who have resided in the US for a total of three of




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the past five years.

**FDA Ethics Requirements**

If an ORISE Fellow, to include their spouse and minor children, reports what is identified as a Significantly Regulated Organization (SRO) or prohibited investment fund financial interest in any amount, or a relationship with an SRO, except for spousal employment with an SRO, and the individual will not voluntarily divest the financial interest or terminate the relationship, then the individual is not placed at FDA. For additional requirements, see [FDA Ethics for Nonemployee Scientists](#).

FDA requires ORISE participants to read and sign their FDA Education and Training Agreement within 30 days of his/her start date, setting forth the conditions and expectations for his/her educational appointment at the agency. This agreement covers such topics as the following:

- Non-employee nature of the ORISE appointment;
- Prohibition on ORISE Fellows performing inherently governmental functions;
- Obligation of ORISE Fellows to convey all necessary rights to the FDA regarding intellectual property conceived or first reduced to practice during their fellowship;
- The fact that research materials and laboratory notebooks are the property of the FDA;
- ORISE fellow's obligation to protect and not to further disclose or use non-public information.

**Qualifications** The qualified candidate should be currently pursuing or have received an associate's, bachelor's, master's, or doctoral degree in the one of the relevant fields.

Applicants must be U.S. citizens and at least 18 years of age at the time of the appointment. Applicants **must be a student** in good standing at an accredited U.S. college or university, or accepted as an entering graduate or professional student at an accredited college or university, with a cumulative GPA of 2.50 or higher (based on a 4.0 scale).

**Point of Contact** [Sherry](#)

- Eligibility Requirements**
- **Citizenship:** U.S. Citizen Only
  - **Degree:** Associate's Degree, Bachelor's Degree, Master's Degree, or Doctoral Degree.
  - **Minimum Overall GPA:** 2.50
  - **Discipline(s):**
    - **Chemistry and Materials Sciences** ([12](#) 👁)
    - **Computer, Information, and Data Sciences** ([2](#) 👁)
    - **Engineering** ([3](#) 👁)
    - **Environmental and Marine Sciences** ([1](#) 👁)
    - **Life Health and Medical Sciences** ([45](#) 👁)
    - **Mathematics and Statistics** ([10](#) 👁)
    - **Physics** ([16](#) 👁)
    - **Science & Engineering-related** ([1](#) 👁)
  - **Age:** Must be 18 years of age

**Affirmation** I certify that I will be at least 18 years of age and a U.S. citizen at the time of the appointment. I am a student in good standing at an accredited U.S. college or university, or I have been accepted as an entering graduate or

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professional student at an accredited college or university, with a cumulative GPA of 2.50 or higher (based on a 4.0 scale).

AND

I am a U.S. citizen, or I have lived in the United States for at least 36 out of the past 60 months. (36 months do not have to be consecutive.)

AND

I have read the FDA Ethics Requirements.