

Opportunity Title: FDA Renal Transporter Biomarkers Fellowship

Opportunity Reference Code: FDA-CDER-2021-0632

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

Reference Code FDA-CDER-2021-0632

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

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

If you have questions, send an email to ORISE.FDA.CDER@orau.org_ Please include the reference code for this opportunity in your email.

Application Deadline 6/30/2021 3:00:00 PM Eastern Time Zone

Description

*Applications will be reviewed on a rolling-basis.

A research opportunity is available in the Office of Translational Sciences/ Office of Clinical Pharmacology (OCP), Center for Drug Evaluation and Research (CDER), Food and Drug Administration (FDA) in Silver Spring, Maryland.

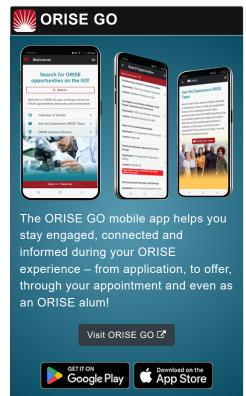
The Center for Drug Evaluation and Research (CDER) performs an essential public health task by making sure that safe and effective drugs are available to improve the health of people in the United States. As part of the U.S. Food and Drug Administration (FDA), CDER regulates over-the-counter and prescription drugs, including biological therapeutics and generic drugs. This work covers more than just medicines. The project will investigate the roles of renal drug transporters (OCT2, MATE1 and MATE2/K) in pharmacokinetics of drugs and/or endogenous biomarkers, and the impact on primary efficacy endpoint measurement in the CKD clinical trials.

Under the guidance of the mentor, the participant will learn about the following:

- Literature review and in-house database analysis.
- Data analysis of PK, biomarker, preclinical and/or clinical data, trial design,
- PBPK model building and analyses.
- Development of quantitative methodologies within the modeling and simulation framework.

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 initial appointment is for six four months, but may be renewed upon recommendation of FDA contingent on the availability of funds. The participant will receive a monthly stipend commensurate with educational level and experience. Proof of health





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insurance is required for participation in this program. The appointment is full-time at FDA in the Silver Spring, Maryland, area. 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 the past five years.

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 EDA.
- ORISE fellow's obligation to protect and not to further disclose or use nonpublic information.

Qualifications

The qualified candidate should be currently pursuing or have received a doctoral degree in one of the relevant fields. Degree must have been received within five years of the appointment start date.

Knowledge of drug development and PBPK modeling is preferred.

Eligibility Requirements

- Degree: Doctoral Degree received within the last 60 months or currently pursuing.
- Discipline(s):
 - Life Health and Medical Sciences (46 ●)
 - Mathematics and Statistics (1 ●)

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