

**Activity Outline**  
**FDA Grand Rounds: A Pandemic and a Call to Action for One Health: The FDA One Health Initiative**  
**June 11, 2020**  
**Webcast**

**Activity Coordinator:**

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**Series Description**

The FDA Grand Rounds is webcast every other month to highlight cutting-edge research underway across the agency and its impact on protecting and advancing public health. Each session features an FDA scientist presenting on a key public health challenge and how FDA is applying science to its regulatory activities.

**Lecture Description**

The Food and Drug Administration (FDA) is responsible for protecting the public health by ensuring the safety, efficacy, and security of human and veterinary drugs, biological products, and medical devices; and by ensuring the safety of our nation's food supply, cosmetics, and products that emit radiation. There are many factors that pose risks or threaten public health. FDA's efforts to improve health goes beyond recognizing disease transmission in an isolated lens of human health, but also acknowledges human-animal interactions and associated environmental factors such as socioeconomic status, behavior, and other social determinants. Addressing human and animal intersectoral factors that are both biological and environmental is a complex endeavor requiring a great deal of knowledge, skills, and resources from various disciplines. However, as health disciplines become progressively specialized, they also become more siloed. One Health is a concept that embraces a multisectoral and transdisciplinary approach to solving health problems by recognizing the interconnection between humans, animals, and their shared environment. Many global changes and activities associated with increased human-animal interactions are enabling disease transmission that become epidemics or pandemics that adversely impact public health. This presentation will explain the One Health Concept and the FDA One Health Initiative. It will also highlight the benefits of One Health and how FDA is operationalizing One Health actions agency wide.

**References**

- Will COVID-19 generate global preparedness?  
[https://www.thelancet.com/journals/lancet/article/PIIS0140-6736\(20\)30559-6/fulltext](https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30559-6/fulltext)
- World Bank's One Health Framework: Operational framework for strengthening human, animal and environmental public health systems at their interface.  
<http://documents.worldbank.org/curated/en/703711517234402168/Operational-framework-for-strengthening-human-animal-and-environmental-public-health-systems-at-their-interface>
- The Need for National Strategy to Address Vector-Borne Disease Threats in the United States.  
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7058377/>
- One Health AIDS to ZIKA. [https://www.youtube.com/watch?v=KXjXUMzsQ\\_s](https://www.youtube.com/watch?v=KXjXUMzsQ_s)
- One Health proof of concept: Bringing a transdisciplinary approach to surveillance for zoonotic viruses at the human-wild animal interface. <https://www.ncbi.nlm.nih.gov/pubmed/28034593>
- The Three One Health Websites: 1. One Health Initiative Website: <http://www.onehealthinitiative.com/mission.php> 2. One Health Commission Website: <https://www.onehealthcommission.org/> 3 One Health Platform website: <http://onehealthplatform.com/>

**Series Objectives**

- Discuss the research conducted at the FDA
- Explain how FDA science impacts public health

**Learning Objectives** After completion of this activity, the participant will be able to:

- Explain the mission and goals of the FDA One Health Initiative.
- Identify recent FDA One Health Initiative related activities.
- Discuss why a One Health approach to pandemics is the new norm.

**Target Audience**

This activity is intended for physicians, pharmacists, nurses, and other scientists within the agency external scientific communities.

**Agenda**

**Lecture 1 June 11, 2020**

Time	Topic	Speaker
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12:00 - 1:00 PM	A Pandemic and a Call to Action for One Health: The FDA One Health Initiative	Brianna Skinner, DVM Bernadette Dunham, DVM, Ph.D.
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## Continuing Education Accreditation



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In support of improving patient care, FDA Center for Drug Evaluation and Research is jointly accredited by the Accreditation Council for Continuing Medical Education (ACCME), the Accreditation Council for Pharmacy Education (ACPE), and the American Nurses Credentialing Center (ANCC) to provide continuing education for the healthcare team.



IPCE CREDIT™

This activity was planned by and for the healthcare team, and learners will receive 1 Interprofessional Continuing Education (IPCE) credit(s) for learning and change.

## CME

FDA Center for Drug Evaluation and Research designates this live activity for a maximum of 1.00 *AMA PRA Category 1 Credit(s)*™. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

## CPE

This knowledge-based activity has been assigned ACPE Universal Activity Number JA0002895-0000-20-022-L04-P for 1.00 contact hour(s).

## CNE

FDA Center for Drug Evaluation and Research designates this activity for 1.00 contact hour(s).

## Requirements for Receiving CE Credit

**Physicians, pharmacists, nurses, and those claiming non-physician CME:** participants must attest to their attendance and complete the final activity evaluation via the CE Portal ([ceportal.fda.gov](http://ceportal.fda.gov)). For multi-day activities, participants must attest to their attendance and complete the faculty evaluation each day. Final activity evaluations must be completed within two weeks after the activity - no exceptions.

Pharmacists will need their NABP e-profile ID number as well as their DOB in MMDD format in order to claim CE credit.

## Important Note regarding completion of evaluations and receiving credit

Attendees have 14 days from the last day of the activity to log in, complete the required evaluation(s) and attest to your attendance to claim credit. Physicians and nurses may then view/print statement of credit. Pharmacists should log into the CPE monitor 10 weeks after the last session of the activity to obtain their CE credit.

## Disclosure

### Faculty

- Dunham, DVM, Ph.D., Bernadette, Professorial Lecturer, The George Washington University - nothing to disclose
- Skinner, Brianna, DVM, Chief Regulatory Veterinarian, Office of Counter-Terrorism & Emerging Threats - nothing to disclose

### Planning Committee

- Dinatale, Miriam, Team Leader, Food and Drug Administration - nothing to disclose
- Pfundt, Tiffany, PharmD, Pharmacist, FDA - nothing to disclose
- Thomas, Devin, LCDR, MPH, CHES, Health Promotions Specialist, FDA/OC/OCS/OSPD - nothing to disclose

- Wheelock, Leslie, MS, RN, Director, OSPD, FDA, OC, OCS, OSPD - nothing to disclose

**CE Consultation and Accreditation Team**

- Bryant, Traci, M.A.T., CE Consultant, FDA/CDER/OEP/DLOD - nothing to disclose
- Zawalick, Karen, CE Team Leader, FDA/CDER/OEP/DLOD - nothing to disclose

**Registration Fee and Refunds**

Registration is complimentary, therefore refunds are not applicable.