Drug Development: How Drug Therapies are Approved for Sale in the U.S.





Join us for this exciting Event!







Gayle Freudinger, Director, Business Development, Regulatory Sciences

Rebecca Lamb-Wharton PhD, Principal Scientist, Regulatory Affairs, and Product Development

UCR's Office of Technology Partnerships will be hosting a seminar by

Cardinal Health

- Learn how new pharmaceutical and biological therapies are approved for human use by the Federal Drug Administration (FDA).
- Learn about the FDA approval process and how it requires new therapies to demonstrate safety and effectiveness over current standards of care.

This seminar will be of interest to anyone working on human health research

This presentation will be in person and on Zoom

The event is free and will include pizza

