
Regulatory Affairs

In the Center for Bio/Pharmaceutical and Biodevice Development
and the College of Sciences

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Faculty Members of the Center for Bio/Pharmaceutical and Biodevice Development

*E. Dale Sevier, Ph.D., Director of the Center for Bio/Pharmaceutical
and Biodevice Development

*Serves on the Faculty Governing Board which makes recommendations on
admissions and curriculum.

Offered by Regulatory Affairs

Master of Science degree in regulatory affairs.
Certificate in regulatory affairs, advanced
(refer to the *Graduate Bulletin*).

Course (R A)

*Refer to Curricula and Courses and University Policies sections of
this catalog for explanation of the course numbering system, unit or
credit hour, prerequisites, and related information.*

UPPER DIVISION COURSE (Intended for Undergraduates)

R A 460. Healthcare Product Regulations (1)

Healthcare product regulation of pharmaceutical, biotechnology,
and biodevice industries. Foundational knowledge of U.S. Food and
Drug Administration and laws and regulations they are responsible for
enforcing.

GRADUATE COURSES Refer to the *Graduate Bulletin*.



R A