



Handbook of Anticancer Pharmacokinetics and Pharmacodynamics (Cancer Drug Discovery and Development)

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There are many steps on the road from discovery of an anticancer drug to securing its final approval by the Food and Drug Administration. In this thoroughly updated and expanded second edition of the *Handbook of Anticancer Pharmacokinetics and Pharmacodynamics*, leading investigators synthesize an invaluable overview of the experimental and clinical processes of anticancer drug development, creating a single indispensable reference that covers all the steps from the identification of cancer-specific molecular targets to screening techniques and the development and validation of bioanalytical methods to clinical trial design and all phases of clinical trials. The authors have included new material on phase 0 trials in oncology, organ dysfunction trials, drug formulations and their impact on anticancer drug PK/PD including strategies to improve drug delivery, pharmacogenomics and cancer therapy, high throughput platforms in drug metabolism and transport pharmacogenetics, imaging in drug development and nanotechnology in cancer.

Authoritative and up-to-date, *Handbook of Anticancer Pharmacokinetics and Pharmacodynamics, 2nd Edition* provides in one comprehensive and highly practical volume a detailed step-by-step guide to the successful design and approval of anticancer drugs.

- Road map to anticancer drug development from discovery to NDA submission
- Discussion of molecular targets and preclinical screening
- Development and validation of bioanalytical methods
- Chapters on clinical trial design and phase 0, I, II, III clinical trials
- Pharmacokinetics, pharmacodynamics, pharmacogenomics, and pharmacogenetics of anticancer agents
- Review of the drug development process from both laboratory and clinical perspectives
- New technological advances in imaging, high throughput platforms, and nanotechnology in anticancer drug development

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Handbook of Anticancer Pharmacokinetics and Pharmacodynamics (Cancer Drug Discovery and Development) From Humana Press Bibliography

- Sales Rank: #3550554 in Books
- Published on: 2014-01-11
- Original language: English
- Number of items: 1
- Dimensions: 10.10" h x 2.00" w x 7.10" l, 4.80 pounds
- Binding: Hardcover
- 836 pages

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Editorial Review

Review

"...an invaluable overview of the experimental and clinical processes that lead to anticancer drugs, creating a single indispensable reference that covers all the steps from the identification of cancer-specific targets to phase III clinical trials." - Tumori

From the Back Cover

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Dr. Michelle Rudek received her BS in Pharmacy from the University of Pittsburgh and her dual Pharm.D., Ph.D. from Virginia Commonwealth University in a joint clinical pharmacology/oncology program with the National Cancer Institute. Dr. Rudek joined Johns Hopkins University in 2001 and is currently an Associate Professor of Oncology and Director of the Analytical Pharmacology Core Laboratory. Her research program is focused on clinical pharmacology with a focus on early phase drug development and special populations including AIDS malignancy and organ dysfunction.

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