

HANDBOOK OF PHARMACEUTICAL GENERIC DEVELOPMENT

ORAL Liquids

VOLUME **IV** – Part **ONE**
Drug Development-Oral Liquid Dosage Forms

BLOCK J D & BELLE D

GENERIC DEVELOPMENT
Handbook of Pharmaceutical
Generic Development *Series*

Handbook of Pharmaceutical Generic Development Series

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Handbook of Pharmaceutical Generic Development

Part I (Development) & Part II (Model ANDA or EU Dossier)

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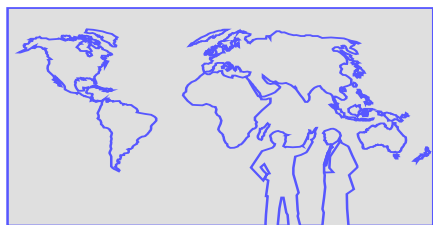
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Part One

Oral Liquids

A circular inset image showing a clear glass bottle tilted to pour a thick red liquid into a silver spoon. The liquid is captured mid-pour, creating a smooth arc from the bottle's spout to the spoon's bowl.

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Handbook of Generic Drug Development Oral Liquid Dosage Forms

This handbook represents the current International Edition of the ongoing 24 volume *series* of Generic Drug Development and appears under the cumulative title of the Handbook *series* of Generic Drug Development. The ongoing series is updated annually at the end of each year. This is an ongoing process as new data, specifications and process techniques are added on a continual and expanding basis. This handbook is fact, never fully complete, as each new annual edition brings an enlarged and extended profile in the drug development process, as well as new agency rules, guidelines and guidance to industry which continues to be added year by year as the global product data base expands. Currently over 150 scientific publications and drug development conferences are annually referenced in the 24 volume Handbook *series* of Generic Drug Development.

This mammoth task presents a continual ongoing commitment by the scientific review committee to the improvement of the technical databases and the product specific drug development requirements and know-how technology accessed through the world wide IAGIM joint ventures and know-how projects currently active in over 15 countries.

The Handbook is available in electronic format (Online and CD ROM) and the e-format is up-dated annually to association members of IAGIM.

This Handbook edition includes additional data on analytical method validation has been redesigned to meet the Guidance for Industry - Organization of an Abbreviated New Drug Application and an Abbreviated Antibiotic Application as well as all FDA guideline and requirements of the Center of Drug Evaluation and Research (CDER) to date of publishing.

Editor-in-chief.

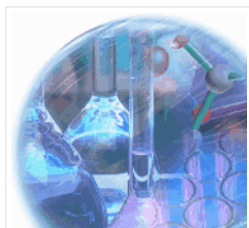


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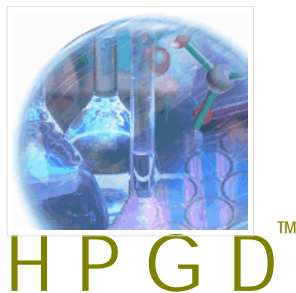
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