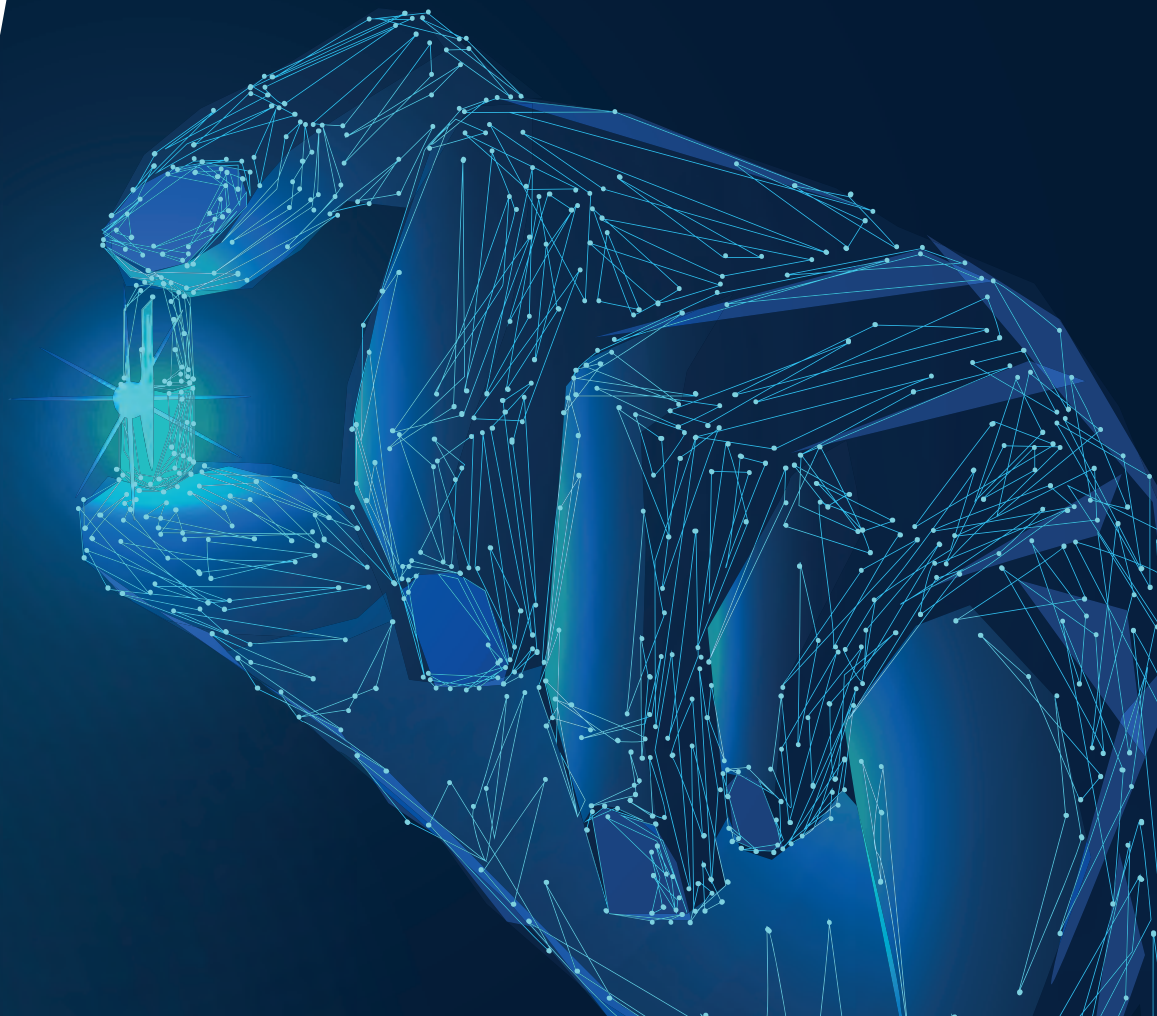


Pharmaceutical Technology[®]

2019 EDITORIAL CALENDAR

www.pharmtech.com



Covering the pharma industry since 1977

EDITORIAL COVERAGE: EXPERT INSIGHT AND ANALYSIS

**Pharmaceutical
Technology**

Pharmaceutical Technology sets the standard for publishing independent, industry-leading information on the technologies, strategies, and regulations crucial to professionals developing and manufacturing pharmaceuticals and biopharmaceuticals. The editorial mix of peer-reviewed papers, technical articles, technology reports, regulatory and business columns, and expert commentary provide comprehensive coverage of process and formulation development, manufacturing operations, drug delivery, packaging, labeling, and distribution.

Contributors from bio/pharmaceutical companies and industry supplier companies, columnists, and the editorial staff are experts, with specialized knowledge and experience in their fields.

EDITORIAL FOCUS

Each issue of *Pharmaceutical Technology* addresses a key trend in drug development and manufacturing including advances in equipment, instruments, and processes; drug formulation strategies; drug delivery trends; emerging dosage forms; biologic-drug development; CMC challenges; process development; and quality-related issues.

Through expert interviews, roundtable discussions, literature reviews, and survey analysis, the editors report on emerging trends, strategies, and best practices in these key areas.

SPECIAL COVERAGE

Throughout 2019, the editors will report on strategies bio/pharma companies can use to transform bio/pharmaceutical development and manufacturing using leading-edge technologies and methods to increase quality output and reduce costs.

PEER-REVIEWED RESEARCH

Pharmaceutical Technology publishes peer-reviewed papers in the form of data-driven research papers, literature and patent reviews, application and technical notes, and position papers on drug development topics. All papers undergo a double-blind peer-review process by the *Pharmaceutical Technology* Editorial Advisory Board, which comprises leading scientists, managers, directors, and consultants.

KEY TOPICS

Drug Development

Features address formulation and drug delivery challenges including solubility, controlled-release formulations, traditional and emerging dosage forms, and more. Advances in API synthesis of small- and large-molecule drug substances, as well as excipients to enhance drug properties, are also addressed.

MANUFACTURING, OPERATIONS, AND SUPPLY CHAIN

The editors examine problems and solutions for solid dosage, sterile, biopharmaceutical, and other drug forms. Experts share insights on manufacturing equipment, process controls, scale-up, packaging, tech transfer, supply chain, and facility and laboratory operations.



EDITORIAL COVERAGE: EXPERT INSIGHT AND ANALYSIS

**Pharmaceutical
Technology**

ANALYTICAL TESTING

Feature articles and case studies address vital quality and characterization practices including solubility, dissolution, extractables and leachables, stability testing, protein characterization, and more.

QUALITY/REGULATIONS

Experts review current regulatory authority initiatives and offer insight on quality, critical quality attributes, good manufacturing practices, statistical analysis, and more.

The **Regulatory Watch** columns review legislation, court decisions, and regulatory changes in the United States and Europe.

Ask the Compliance Expert answers reader questions about good manufacturing practices and other regulatory issues.

OUTSOURCING

Trends, partnerships, and business activities in the contract services market are described by expert columnists. Other features examine best practices for working with contract service providers for drug development, manufacturing, and laboratory studies.

OTHER EDITORIAL FEATURES

New analytical instruments, automation and process control systems, information technology tools, laboratory equipment, and manufacturing equipment are described in **Product Spotlight**.

Business developments, new facilities, and other industry supplier activities are reported in **PharmaCapsules**.

Updates on global markets, industry research, partnerships/collaborations, and the drug pipeline are also featured.

CONTRIBUTION GUIDELINES

For information about contributing editorial features to *Pharmaceutical Technology*, visit www.PharmTech.com/pharmtech-author-guidelines.



2019 EDITORIAL COVERAGE

JANUARY

Ad Close: December 7, 2018

FOCUS

Transforming Bio/Pharma Development and Manufacturing
Annual Employment Survey

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

API Development and Approval Trends
Protein/Peptide Drug Delivery

Manufacturing

Biologic-Based Drug Manufacturing
Process Development

Quality/Regulations

Regulatory Outlook

Analytics

Solubility

Operations

Packaging Trends

Outsourcing

Outsourcing Outlook

DEPARTMENTS

US Regulatory Watch
European Regulatory Watch
New Product Spotlight
PharmaCapsules
Ask the Compliance Expert

SHOWS

Pharmapack, Feb. 6-7, Paris, France

VALUE-ADDED

- FREE 3-Minute Podcast, posted on www.PharmTech.com or FREE whitepaper listing in the *PharmTech Pharma Knowledge Resources* Newsletter
- FREE Direct eResponse Ad Leads (Ask your rep for details.)

FEBRUARY

Ad Close: January 10

FOCUS

Analysis and Data Interpretation

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Biopharmaceutical APIs
Controlled-release Drug Formulation

Manufacturing

Topical Drug Manufacturing
Lyophilization

Quality/Regulations

Regulatory Focus: Sterile/Aseptic Manufacturing

Analytics

Lab Data Integrity

Operations

Facility Design and Engineering

Supply Chain

Cold Chain

Outsourcing

Analytical Best Practices

DEPARTMENTS

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SHOWS

PDA Annual Meeting, March 11-13, San Diego, CA

VALUE-ADDED

PDA News Release (Print and ePT e-newsletter)

SPECIAL THEMED SUPPLEMENT

Partnering for Bio/Pharma Success

Best practices and metrics for choosing contract service suppliers, ensuring quality control in vendor relationships, technology transfer, and intellectual property issues.

MARCH

Ad Close: February 8

FOCUS

Technology Drivers for Bio/Pharma Manufacturing Advances

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Excipients for Solubility Enhancement
Biologic Drug Formulation

Manufacturing

Solid/Semi-Solid Drug Manufacturing
Refurbished Equipment

Quality/Regulations

Regulatory Focus: Emerging Therapies

Analytics

Stability Testing

Operations

Lab Operations

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SHOWS

- Pittcon, March 17-21, Philadelphia, PA
- DCAT Week, March 19-22, New York, NY
- INTERPHEX, April 2-4, New York, NY

VALUE-ADDED

- INTERPHEX Show Guide (Print)
- INTERPHEX Show Guide (Online and special Newsletter)

2019 EDITORIAL COVERAGE

APRIL

Ad Close: March 13

FOCUS

Patient-Centric Drug Development/Formulation

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Sourcing API Development/Manufacturing
Drug Delivery Methods

Manufacturing

Biologic-Based Drug Manufacturing
Continuous Manufacturing

Quality/Regulations

Batch Records

Analytics

Particle Characterization
Statistical Solutions

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Anticounterfeiting

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Formulation Best Practices

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SHOWS

- CPhI North America, April 30–May 2, Chicago, IL
- ExcipientWorld, May 7–8, National Harbor, MD

VALUE-ADDED

- CPhI North America Exhibitor Guide (Print)
- CPhI North America Exhibitor Guide (Online and special Newsletter)

SPECIAL THEMED SUPPLEMENT

Solid Dosage Drug Development and Manufacturing featuring the CPhI North America Planning Guide

Trends in the development of solid-dosage drug forms, including excipients, API compatibility, formulation development, solubilization, and new manufacturing equipment.

MAY

Ad Close: April 9

FOCUS

Bioprocessing Productivity Enhancement

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Managing Excipient Variability
Particle Engineering

Manufacturing

Aseptic/Sterile Drug Manufacturing
Single-Use Manufacturing

Quality/Regulations

Critical Quality Attributes

Analytics

Cleaning Validation

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Packaging Trends

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SHOWS

- BIO International Convention, June 3–6, Philadelphia, PA
- DCAT Sharp Sourcing, June, Location TBD

VALUE-ADDED

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SPECIAL THEMED EBOOK

Biologics and Sterile Drug Manufacturing eBook

Novel technologies for the formulation, manufacture, purification, and delivery of sterile small- and large-molecule drugs including single-use systems, facilities and equipment, contamination issues, and process analytics.

JUNE

Ad Close: May 14

FOCUS

Advances in Drug Delivery Devices

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

High-Potency API Manufacturing
Solubility/Bioavailability

Manufacturing

Inhalation Drug Manufacturing
Scale-Up

Quality/Regulations

Good Laboratory Practices

Analytics

Glycosylation

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Track and Trace/Serialization

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Manufacturing Best Practices

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FREE 3-Minute Podcast, posted on www.PharmTech.com OR Free Whitepaper listing in the *PharmTech Pharma Knowledge Resources* Newsletter

BUYERS' GUIDE

Buyers' Guide and Best Practices

The global resource for suppliers of chemicals, raw materials, intermediates and excipients; equipment and supplies for manufacturing, packaging, and cleanrooms; laboratory equipment; and contract services. Plus, recommended best practices for laboratory, formulation, manufacturing, and packaging functions.

2019 EDITORIAL COVERAGE

JULY

Ad Close: June 11

FOCUS

Defining Good Manufacturing Practices

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Advances in Small-Molecule API Synthesis
Solid-Dosage Drug Forms

Manufacturing

Aseptic/Sterile Drug Manufacturing
Digital Manufacturing Connections

Quality/Regulations

Good Automation Practices

Analytcs

Endotoxin Testing

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SHOWS

Controlled Release Society, July 21–23, Valencia, Spain

VALUE-ADDED

Double Up Ad Program: Ask your Rep for details.

AUGUST

Ad Close: July 10

FOCUS

Overcoming CMC Challenges

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Excipients for Controlled-Release Formulations
Parenteral Drug Formulation

Manufacturing

Solid/Semi-Solid Drug Manufacturing
Clinical Trial Materials

Quality/Regulations

Good Manufacturing Practices

Analytcs

Protein Characterization

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Logistics/Shipping

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New Product Spotlight
PharmaCapsules
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VALUE-ADDED

FREE Product/Service Profile Posted Online and Listed in Special eNewsletter

SPECIAL THEMED SUPPLEMENT

Outsourcing Resources: Technology and Process Support for Bio/Pharma Development and Manufacturing

An annual review of the market for outsourced resources, including analysis of contract services and business, regulatory, and supply chain issues.

SEPTEMBER

Ad Close: August 9

FOCUS

Right-Scaling the Drug Manufacturing Process

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Advances in Biologics API Synthesis
Tablet Formulation

Manufacturing

Aseptic/Sterile Drug Manufacturing
Tableting

Quality/Regulations

Measuring Quality

Analytcs

Container Closure Testing

Operations

Facility Design and Engineering

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SHOWS

Pack Expo, Sept. 23–25, Las Vegas, NV
PDA/FDA Joint Regulatory Conference, Dates TBD, Washington, DC

VALUE-ADDED

Pharma Marketplace online directory program. Ask your rep for details.

SPECIAL THEMED EBOOK

PharmTech's Regulatory Sourcebook and Reference

A compilation of regulations, guidances, regulations, compendial documents, position papers, and enforcement actions from global regulatory authorities.

2019 EDITORIAL COVERAGE

OCTOBER

Ad Close: September 10

FOCUS

Drug Dosage Forms Trends

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

Drug Stability
Lipid-based Formulations

Manufacturing

Solid/Semi-Solid Drug Manufacturing
Manufacturing Data Integrity

Quality/Regulations

Inspection Readiness and Response

Analytcs

Dissolution Testing
Statistical Solutions

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Supply Chain

Raw Materials Tracking

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SHOWS

- AAPS 2019 PharmSci 360, Nov. 3-6, San Antonio, TX
- CPhI Worldwide, November 5-7, Frankfurt, Germany

VALUE-ADDED

- AAPS Exhibitor Guide (Print)
- AAPS Exhibitor Guide (Online and Special Newsletter)
- CPhI Worldwide Exhibitor Guide (Print)
- CPhI Worldwide Exhibitor Guide (Online and Special Newsletter)

SPECIAL THEMED SUPPLEMENT

APIs, Excipients, and Manufacturing featuring the CPhI/ICSE Worldwide Planning Guide

Developments in the synthesis of APIs and intermediates, plus advances in formulation, development, and finished-product manufacturing.

NOVEMBER

Ad Close: October 11

FOCUS

Best Practices for Quality Control

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

API Formulation Trends
Taste-masking

Manufacturing

Transdermal Drug Manufacturing
Process Control/Automation

Quality/Regulations

Regulatory Focus: Solid/Semi-solid Dosage Drugs

Analytcs

Lyophilization

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Tech Transfer

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DECEMBER

Ad Close: November 8

FOCUS

Manufacturing Continuity Strategies

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS

TECHNICAL TOPICS

Development

API/Excipient Interactions
Solubility/Bioavailability

Manufacturing

Biologic-Based Drug Manufacturing
Process Modeling

Quality/Regulations

Design of Experiments

Analytcs

Extractables and Leachables Testing

Operations

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Drug Product Tracking

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VALUE-ADDED

Double Up Ad Program: Ask your Rep for details.

SPECIAL SPONSORED-CONTENT ISSUE

Corporate Capabilities

Full-page descriptions of products and services from the industry's leading suppliers. Profiles are posted online for six months.

SPECIAL THEMED EBOOK

PharmTech's Regulatory Sourcebook and Reference

An updated compilation of regulations, guidances, regulations, compendial documents, position papers, and enforcement actions from global regulatory authorities.

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