

Advancing Development & Manufacturing

**Pharmaceutical  
Technology**  
EUROPE

**EDITORIAL  
CALENDAR**

**30<sup>th</sup>**

**A N N I V E R S A R Y**

**Covering the Pharma Industry since 1989**



UBM

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# EDITORIAL COVERAGE: EXPERT INSIGHT AND ANALYSIS

*Pharmaceutical Technology Europe* 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 Europe* 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 Europe* 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, semi-solid, 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.

### 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 column reviews legislation, court decisions, and regulatory changes in Europe and other regions.

## 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.

## CONTRIBUTION GUIDELINES

For information about contributing editorial features to *Pharmaceutical Technology Europe*, visit [www.PharmTech.com/pharmtech-author-guidelines](http://www.PharmTech.com/pharmtech-author-guidelines).

## SPECIAL ISSUES IN 2019

### August 2019

#### 30th Anniversary Issue

The editors of *Pharmaceutical Technology Europe* mark 30 years of publishing insight and analysis about bio/pharmaceutical development and manufacturing with a retrospective of industry accomplishments and a look at challenges ahead.

### March 2019

#### Outsourcing Resources:

#### Technology and Process Support for Bio/Pharma Development and Manufacturing

The editors review the market for outsourced resources, including analysis of contract services, as well as business, regulatory, and supply chain issues.

### September 2019

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

This special supplement covers developments in the synthesis of APIs and pharmaceutical intermediates, formulation, development, and finished-product manufacturing. This issue will feature a CPhI Worldwide Planning Guide.

# 2019 EDITORIAL CALENDAR

## JANUARY

Ad Close: 6 January

### Focus

Packaging Trends  
Special Report: Salary and 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

Transforming Bio/Pharma  
Development and Manufacturing

#### Outsourcing

Outsourcing Review

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

Pharmapack, 6-7 February, Paris, France

### Value-Added

Readex Ad Study or  
1/2-page profile for Pharmapack

## FEBRUARY

Ad Close: 29 January

### 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

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

Global Drug Delivery,  
12-14 March, Berlin, Germany

### Value-Added

e-news Product/Service Profile

## MARCH

Ad Close: 21 February

### 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

#### Outsourcing

Outsourcing Review

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

- Pittcon, 17-19 March, Philadelphia, PA USA
- DCAT Week, 17-23 March, New York, NY USA
- BIO Europe, 25-27 March, Vienna, Austria
- INTERPHEX, 2-4 April, New York, NY USA
- POWTECH, 9-11 April, Nuremberg, Germany

### Value-Added

½-page Product/Service Profile

### Special Themed Supplement

#### Outsourcing Resources

**Technology and Process Support for Bio/  
Pharma Development and Manufacturing**

An annual review of trends and  
developments for outsourcing services.

## APRIL

Ad Close: 28 March

### 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

#### Operations

Tech Transfer

#### Supply Chain

Anticounterfeiting

#### Outsourcing

Formulation Best Practices

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

- CPhI North America,  
30 April-2 May 2, Chicago, IL USA
- Making Pharmaceuticals UK,  
30 April-2 May 2, Coventry, UK

### Value-Added

½-page profile for CPhI North America or  
e-news Product/Service Profile

# 2019 EDITORIAL CALENDAR

## MAY

Ad Close: 23 April

### 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

#### Operations

Packaging Trends

#### Outsourcing

Outsourcing Review

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

BIO International Convention,  
3-6 June, Philadelphia, PA USA

### Value-Added

½-page Product/Service Profile

## JUNE

Ad Close: 21 May

### 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

#### Operations

Process Operations

#### Supply Chain

Track and Trace/Serialization

#### Outsourcing

Manufacturing Best Practices

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

Chemspec, 26-27 June, Basel, Switzerland

### Value-Added

e-news Product/Service Profile

## JULY

Ad Close: 25 June

### 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

#### Analytics

Endotoxin Testing

#### Operations

Packaging Trends

#### Outsourcing

Outsourcing Review

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

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

### Value-Added

· Free Profile with Full Page  
· Readex ad study

## AUGUST

Ad Close: 23 July

### Focus

**30th Anniversary Issue**

### 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

#### Analytics

Protein Characterization

#### Operations

Lab Operations

#### Supply Chain

Logistics/Shipping

#### Outsourcing

Analytical Best Practices

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Value-Added

Full Page Profile: Past, Present and Future



# 2019 EDITORIAL CALENDAR

## SEPTEMBER

Ad Close: 20 August

### 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

#### Analytics

Container Closure Testing

#### Operations

Facility Design and Engineering

#### Outsourcing

Outsourcing Review

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

Making Pharmaceuticals Europe,  
18-19 Sept., Milan, Italy

### Value-Added

Free Whitepaper Listing

### Special Themed Supplement

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

Developments in the synthesis of APIs  
and pharmaceutical intermediates,  
formulation, development, and finished-  
product manufacturing. This issue  
will feature a CPhI Planning Guide.

## OCTOBER

Ad Close: 24 September

### 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

#### Analytics

Dissolution Testing  
Statistical Solutions

#### Operations

Packaging Trends

#### Supply Chain

Raw Materials Tracking

#### Outsourcing

Formulation Best Practices

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

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

### Value-Added

½-page CPhI Profile

## NOVEMBER

Ad Close: 24 October

### 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

#### Analytics

Lyophilization

#### Operations

Tech Transfer

#### Outsourcing

Outsourcing Review

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Shows

CPhI India, Dates TBD, Delhi, India

### Value-Added

½-page Product/Service Profile

## DECEMBER

Ad Close: 19 November

### 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 Experiment

#### Analytics

Extractables and Leachables Testing

#### Operations

Process Operations

#### Supply Chain

Drug Product Tracking

#### Outsourcing

Manufacturing Best Practices

### Departments

European Regulatory Watch  
Product Spotlight  
Ask the Compliance Expert

### Value-Added

Free full-page Corporate Profile

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\*Advertising deadlines may be subject to change.

\*\*Receipt of an article or abstract does not guarantee inclusion in a particular issue of the magazine or on [www.PharmTech.com](http://www.PharmTech.com)

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