Advancing Development & Manufacturing

Pharmaceutical Technology® EUROPE

2022 Editorial Calendar

AN MJH life sciences™ BRAND

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EDITORIAL COVERAGE

Pharmaceutical Technology EUROPE

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 provides 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 and manufacturing strategies; drug delivery trends; emerging dosage forms; vaccines and biologic-drug development; drug ingredient quality; and processing equipment.

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

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 Europe*™ Editorial Advisory Board, which comprises leading scientists, managers, directors and consultants.

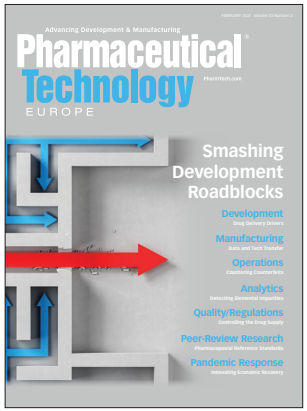
KEY TOPICS

DRUG DEVELOPMENT

Features address advances in API synthesis of small- and large-molecule drug substances and excipients, as well as formulation and drug delivery challenges. Topics including early development strategies, solubility enhancement, particle characterization, excipients, and stability are covered for traditional and emerging dosage forms.

MANUFACTURING

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.

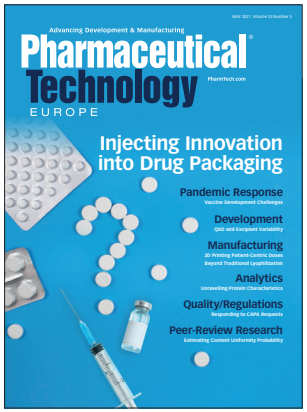


ANALYTICAL TESTING

Feature articles and case studies address vital quality and analytical practices including contamination control, dissolution, extractables and leachables, stability testing, protein characterization, cleaning validation, and more.

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.



QUALITY/REGULATIONS

Experts review current regulatory authority initiatives and offer insight on regulatory authority activities, good manufacturing practices, good laboratory practices, statistical analysis and more.

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

CONTRIBUTION GUIDELINES

For information about contributing editorial features to *Pharmaceutical Technology Europe*™, visit the Editorial Info link on www.PharmTech.com.

SPECIAL EDITORIAL COVERAGE

Pharmaceutical Technology EUROPE

January 2022 – Editors' Tech Talk:
APIs, Excipients, and Formulation Advances
The editors analyze recent new drug approvals and trends in API synthesis, formulation strategies, excipients, and process development.

February 2022 – Interactive eBook:
Bio/Pharma Outsourcing Innovation
Contract research, development, and manufacturing organizations share details on the technologies, processes, equipment, and other innovations that help accelerate drug development, manufacturing, packaging, and quality control.

March 2022 – Interactive eBook:
Quality and Regulatory Sourcebook
Stay ahead of the latest regulations, guidance documents, and compendial documents guiding drug development and manufacturing; gain insight into practical quality practices for bio/pharma organizations.

April 2022 – Editors' Tech Talk:
Emerging Therapies
The editors examine challenges associated with developing, formulating, and manufacturing new drug modalities and dosage forms.

May 2022 – Interactive eBook:
Trends in Manufacturing
New technologies and processes are accelerating drug production while reducing costs and improving quality. Learn about new strategies from process development through commercial manufacturing for a range of dosage forms.

June 2022 – Editors' Tech Talk:
Biopharmaceutical Drug Development Manufacturing
The editors and report on novel technologies for the formulation, manufacture, purification, and delivery biologic-based drugs.

August 2022 – Editors' Tech Talk:
Aseptic Processing and Manufacturing
The editors review regulatory requirements, quality challenges, and new processes and technologies to produce sterile drugs safely and economically.

September 2022 – Editors' Tech Talk:
Solid Dosage Drug Development and Manufacturing
The editors share expert insight on trends in the development of solid-dosage drug forms, including excipients, APIs, formulation, and new manufacturing processes and equipment.

October 2022 – Interactive eBook:
Trends in Formulation
Experts share new processes, novel excipients, and new methodologies to address formulation challenges associated with complex molecules, particle engineering, bioavailability limits, and demands for safer dosage forms for patients.

November 2022 – Editors' Tech Talk:
Automating Bio/Pharma Processes
The editors review how artificial intelligence, robotics, virtual reality, remote monitoring, and other automation strategies are impacting bio/pharma process development and manufacturing.

2022 EDITORIAL CALENDAR

Pharmaceutical Technology EUROPE

JANUARY
Ad Close: 7 January

FOCUS
Pharma Industry Outlook
Special Coverage: Annual Employment Survey

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Formulation Strategies for Early Drug Development
Drug Appearance and Taste
Solubility/Bioavailability

Manufacturing
Biologic-Based Drug Manufacturing
Facility Design and Engineering

Quality/Regulations
European Regulatory Update
Ask the Compliance Expert

Analytics
Drug Substance Testing

Outsourcing
State of Outsourcing Industry

VALUE-ADDED
FREE 3-Minute Podcast Posted on *www.PharmTech.com* or FREE Whitepaper Listing in the *PharmTech* Whitepapers Section
FREE Direct eResponse Ad Leads (Ask your rep for details.)

EDITORS' TECH TALK
APIs, Excipients, and Formulation Advances
The editors analysis recent new drug approvals and trends in API synthesis, formulation strategies, and excipient and process development.

FEBRUARY
Ad Close: 24 January

FOCUS
Bio/Pharma Analysis

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Toxicology Studies
Cannabinoid-based Drugs
Vaccine Development

Manufacturing
Aseptic/Sterile Drug Manufacturing
Process Analytical Technology
Serialization

Quality/Regulations
Computer Validation
European Regulatory Update
Ask the Compliance Expert

Analytics
Automated Analytical Workflows

Outsourcing
Method Development

SHOWS
Pittcon, 5–9 March, Atlanta

VALUE-ADDED
Product Service Profile in eNewsletter

INTERACTIVE EBOOK
Bio/Pharma Outsourcing Innovation
Contract research, development, and manufacturing organizations share details on the technologies, processes, equipment, and other innovations that help accelerate drug development, manufacturing, packaging, and quality control.

MARCH
Ad Close: 21 February

FOCUS
Drug Dosage Forms Trends

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
IND/CTA Application Process
Biopharmaceutical APIs
Accelerated Formulation Strategies

Manufacturing
Oral Solid Dose Drug Manufacturing
Biologics Drug Continuous Manufacturing
Supply Chain Continuity

Quality/Regulations
Good Distribution Practices
European Regulatory Update
Ask the Compliance Expert

Analytics
Protein Characterization

Outsourcing
Clinical Trial Materials

SHOWS
DCAT Week, 20–24 March, New York
Drug Delivery & Formulation Summit, 21–23 March, Berlin
BIO-Europe Spring, 28–30 March, Basel
ACHEMA, 4–8 April, Frankfurt

VALUE-ADDED
FREE 3-Minute Podcast Posted on *www.PharmTech.com* or FREE Whitepaper Listing in the *PharmTech* Whitepapers Section

INTERACTIVE EBOOK
Quality and Regulatory Sourcebook
Stay ahead of the latest regulations, guidances, and compendial documents guiding drug development and manufacturing, and gain insight into practical quality practices for bio/pharma organizations.

Trade show dates listed are as of 2021 September 10. Trade show dates and topics are subject to change.

2022 EDITORIAL CALENDAR

Pharmaceutical Technology EUROPE

APRIL
Ad Close: 20 March

FOCUS
Drug Manufacturing Technology

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Medicinal Chemistry
Excipient Quality
Tablet Formulation

Manufacturing
Compounded Drug Manufacturing
Fill/Finish
Packaging Trends

Quality/Regulations
Corrective and Preventive Actions
European Regulatory Update
Ask the Compliance Expert

Analytics
Cleaning Validation
Statistical Solutions

Outsourcing
Bioprocessing Contract Services

SHOWS
CPhI North America, 17–19 May, Philadelphia
Interphex, 24–26 May, New York

VALUE-ADDED
Ad Retargeting: 25,000 Impressions

EDITORS' TECH TALK
Emerging Therapies
The editors examine challenges associated with developing, formulating, and manufacturing new drug modalities and dosage forms.

MAY
Ad Close: 20 April

FOCUS
Biologic Drug Development and Manufacturing

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Pre-IND/CTA Studies
Inhalation Drug Formulation
Drug Delivery Methods

Manufacturing
Semisolid Drug Manufacturing
Lyophilization
Cold Chain

Quality/Regulations
Quality Culture
European Regulatory Update
Ask the Compliance Expert

Analytics
Dissolution Testing

Outsourcing
Formulation

SHOWS
ChemSpec Europe, 31 May–1 June, Frankfurt
BIO International Convention, June 13–16, San Diego

VALUE-ADDED
FREE Direct eResponse Ad Leads

INTERACTIVE EBOOK
Trends in Manufacturing
New technologies and processes are accelerating drug production while reducing costs and improving quality. Learn about new strategies from process development through commercial manufacturing for a range of dosage forms.

JUNE
Ad Close: 20 May

FOCUS
Aseptic Processing

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Dosing Studies
Coprocessed Excipients
Patient-Centric Formulation

Manufacturing
Biologic-Based Drug Manufacturing
Equipment Cleaning

Quality/Regulations
European Regulatory Update
Ask the Compliance Expert

Analytics
Elemental Impurities

Outsourcing
Contract Testing Services

SHOWS
Analytica, 21–24 June, Munich

VALUE-ADDED
Product Service Profile in eNewsletter

EDITORS' TECH TALK
Biopharmaceutical Drug Development Manufacturing
The editors report on novel technologies for the formulation, manufacture, purification, and delivery of biologic-based drugs.

2022 EDITORIAL CALENDAR

Pharmaceutical Technology EUROPE

JULY
Ad Close: 20 June

FOCUS
Drug Packaging Advances

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Molecule Characterization
High-Potency Drug Formulation
Biologic Drug Formulation

Manufacturing
Point-of-Use Drug Manufacturing
Automation

Quality/Regulations
European Regulatory Update
Ask the Compliance Expert

Analytics
Extractables and Leachables (raw materials)

Outsourcing
State of Outsourcing Industry

SHOWS
Controlled Release Society (TBD)

VALUE-ADDED
Case Study on an Industry Topic of Choice

AUGUST
Ad Close: 22 July

FOCUS
Ingredient Quality

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Animal Models
Cell Therapy Development
Solubility/Bioavailability

Manufacturing
Vaccine Manufacturing
Facility Design and Engineering

Quality/Regulations
GMPs for Solid-Dose Drugs
European Regulatory Update
Ask the Compliance Expert

Analytics
Automated Finished Product Inspection
Lab Data Integrity

Outsourcing
Contract Packaging

VALUE-ADDED
FREE Direct eResponse Ad Leads (Ask your rep for details.)

EDITORS' TECH TALK
Aseptic Processing and Manufacturing
The editors review regulatory requirements, quality challenges, and new processes and technologies to produce sterile drugs safely and economically.

SEPTEMBER
Ad Close: 22 August

FOCUS
Emerging Therapies

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Drug Candidate Screening
Topical Drug Development
Alternative Drug Delivery Formulation

Manufacturing
Biologic-Based Drug Manufacturing
Process Optimization
Logistics/Shipping

Quality/Regulations
Audits and Inspections
European Regulatory Update
Ask the Compliance Expert

Analytics
Environmental Monitoring

Outsourcing
Bioanalytical Studies

SHOWS
Making Pharmaceuticals, 27–28 Sept., Dublin

VALUE-ADDED
Ad Retargeting: 25,000 Impressions

EDITORS' TECH TALK
Solid Dosage Drug Development and Manufacturing
The editors share expert insight and report on trends in the development of solid-dosage drug forms, including excipients, APIs, formulation, and new manufacturing processes and equipment.

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2022 EDITORIAL CALENDAR

Pharmaceutical Technology EUROPE

OCTOBER
Ad Close: 20 September

FOCUS
Formulation Strategies

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Early Development Strategies
Advances in Small-Molecule API Synthesis
Reformulation Strategies

Manufacturing
Aseptic/Sterile Drug Manufacturing
Contamination Control
Raw Materials Traceability

Quality/Regulations
Compendial Compliance Update
European Regulatory Update
Ask the Compliance Expert

Analytics
Extractables and Leachables (processing and packaging)
Statistical Solutions

Outsourcing
Bioprocessing Contract Services

SHOWS
CPhI Worldwide, TBD

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INTERACTIVE EBOOK
Trends in Formulation
Experts share new processes, novel excipients, and new methodologies to address formulation challenges associated with complex molecules, particle engineering, bioavailability limits, and demands for safer dosage forms patients.

NOVEMBER
Ad Close: 20 October

FOCUS
Processing Equipment Trends

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
ADME Studies
Excipients for Solubility
Particle Engineering

Manufacturing
Oral Solid Dose Drug Manufacturing
Scale Up
Packaging Trends

Quality/Regulations
Supplier Oversight
European Regulatory Update
Ask the Compliance Expert

Analytics
Particle Analysis

Outsourcing
Tech Transfer

VALUE-ADDED
FREE Direct eResponse Ad Leads (Ask your rep for details.)

EDITORS' TECH TALK
Automating Bio/Pharma Processes
The editors review how artificial intelligence, robotics, virtual reality, remote monitoring, and other automation strategies are impacting bio/pharma process development and manufacturing.

DECEMBER
Ad Close: 21 November

FOCUS
Trends in Drug Development

PEER-REVIEWED RESEARCH/TECHNICAL PAPERS
Bio/Pharma Research and Technical Advances

EMERGING TOPICS IN BIO/PHARMA

TECHNICAL TOPICS

Development
Pharmacokinetics
Drug Stability
Novel Drug Forms

Manufacturing
Specialty Drug Manufacturing
Isolators and RABs

Quality/Regulations
GMPs for Sterile/Aseptic Manufacturing
European Regulatory Update
Ask the Compliance Expert

Analytics
Stability Testing

Outsourcing
Impurity Testing

VALUE-ADDED
Double Up Ad Programme

Trade show dates listed are as of 2021 September 10. Trade show dates and topics are subject to change.

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