BioPharm INTERNATIONAL®

The Science & Business of Biopharmaceuticals

2022 Editorial Calendar

Covering the biopharmaceutical development and manufacturing industry since 1988

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EDITORIAL COVERAGE

BioPharm INTERNATIONAL®

SPECIAL ISSUES

March 2022

Regulatory Sourcebook
The editors present a compilation of news, trends, and strategies related to regulations, guidance documents, compendial documents, and enforcement actions from global regulatory authorities in an interactive eBook format.

May 2022

Partnerships for Outsourcing eBook
The editors review best practices and metrics for choosing contract service suppliers, ensuring quality control in vendor relationships, technology transfer, and intellectual property issues in an interactive eBook format.

September 2022

Emerging Therapies ebook
Innovative process development and manufacturing practices to accelerate new biopharmaceutical modalities are explored in an interactive eBook format.

November 2022

Manufacturing and Facilities
Strategies and technologies to address limited production capacity for biopharmaceutical are covered in this interactive eBook.

SPONSORED-CONTENT eBOOKS

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EXPERT INSIGHT AND ANALYSIS

BioPharm International provides the biopharmaceutical industry with comprehensive coverage of key scientific, technology, regulatory, and business topics, as well as issues related to bio/pharma's response to the global pandemic.

The editorial mix of peer-reviewed papers, practical advice on managing bioprocessing and technology, regulatory and business columns, and expert commentary provides comprehensive coverage of upstream and downstream processing, manufacturing operations, regulations, formulation, scale up, technology transfer, drug delivery, analytical testing, and more.

The print and digital editorial coverage provides technical and business insight and analysis for all biologic-based therapies including monoclonal antibodies, vaccines, biosimilars, protein therapeutics, cell therapies, gene therapies, antibody-drug conjugates, and other emerging therapies.

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

To contribute, view the submission guidelines:
biopharminternational.com/editorial_info

EDITORIAL FEATURES

Peer-Reviewed Research Papers
BioPharm International publishes peer-reviewed papers in the form of technical case studies/application notes; topical literature or patent reviews; novel research; or science-based opinion papers. All papers undergo a double-blind peer-review process by *BioPharm International*'s Editorial Advisory Board of leading scientists, managers, directors, and consultants.

Technical Articles
Feature articles in *BioPharm International* offer timely technical and scientific discussions of drug development challenges and solutions. Articles are authored by industry experts and the magazine's editorial team.

Topics cover the full spectrum of biopharmaceutical development and manufacturing including upstream processing, downstream processing, manufacturing, quality and regulations, analytics, facilities and equipment, laboratory operations, packaging, logistics, supply chain, and business issues including intellectual property, market research, and funding.

Regulatory Beat
The latest developments, guidance documents, and enforcement action from international regulatory authorities, as well as expert analysis, are addressed in this monthly feature.

Ask the Compliance Expert
Questions about enforcement, standard operating procedures, working with FDA, and other compliance issues are answered by regulatory experts.

2022 EDITORIAL CALENDAR

BioPharm INTERNATIONAL®

January

Ad Close: December 10, 2021

Focus
Biopharma Industry Outlook
Special Section: Employment Outlook, Training, and Survey

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

Development
Vaccine Development

Upstream Processing
Automating Upstream Processes

Downstream Processing
Chromatography Resins

Manufacturing
Process Monitoring/Controls

Quality/Regulations
Form 483s and Warning Letters

Analytics
Bioassay Development

Supply Chain
Cold Chain

BioBusiness
Investment Outlook

Outsourcing
State of Outsourcing Industry

Departments
Regulatory Beat
Product Spotlight
Ask the Compliance Expert

Value-Added
Ad Retargeting: Up to 10,000 Impressions

February

Ad Close: January 13

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Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Regenerative Medicine Development

Upstream Processing
Cell Culture

Downstream Processing
Single-Use Systems

Manufacturing
Lyophilization
Facilities

Quality/Regulations
CMC Strategies

Analytics
Lab Operations

BioBusiness
Intellectual Property

Outsourcing
Method Development

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Product Spotlight
Ask the Compliance Expert
Pipeline: Protein Therapeutics

Shows
Pittcon, March 5–9, Atlanta
BioProcess International West, March 14–17, San Diego

Value-Added
Supplier Spotlight Listing Newsletter

March

Ad Close: February 11

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Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Excipients

Upstream Processing
What's New in Upstream Technologies

Downstream Processing
Separation and Purification

Manufacturing
Cell and Gene Therapy
Fill/Finish

Quality/Regulations
Good Distribution Practices

Analytics
Protein Characterization

BioBusiness
Global Biopharma Markets

Outsourcing
Clinical Trial Materials

Departments
Regulatory Beat
Product Spotlight
Ask the Compliance Expert

Shows
DCAT Week, March 20–24, New York City
PDA Annual Meeting, April 4–6, Dallas

Value-Added
Button Ad (220x75 px) in *BioPharm International*'s *eBulletin* Newsletter

Interactive eBook
Quality and Regulatory Sourcebook
A compilation of news, trends, and strategies related to regulations, guidances, compendial documents, and enforcement actions from global regulatory authorities in an updated eBook.

Topics and trade show dates are subject to change.

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2022 EDITORIAL CALENDAR

BioPharm INTERNATIONAL®

April

Ad Close: March 15

Focus
Downstream Processing

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Nucleic Acid-Based Therapeutics

Upstream Processing
Bioreactors

Downstream Processing
What's New in Downstream Technologies

Manufacturing
Process Monitoring/Controls

Quality/Regulations
Data Mining CAPA

Analytics
Cleaning Validation

Supply Chain
Materials Sourcing

BioBusiness
Partnerships

Outsourcing
Bioprocessing Contract Services

Departments
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Ask the Compliance Expert
Pipeline: Cell Therapies

Shows
CPhI North America, May 17–18, Philadelphia
INTERPHEX, May 24–26, New York

Value-Added
Spring Conference eNewsletter Profile

May

Ad Close: April 14

Focus
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Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Upstream Processing
Single-Use Systems

Downstream Processing
Cell Harvesting

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Process Analytical Technology
Sterilization Methods

Quality/Regulations
GMPs: Emerging Therapies

Analytics
Data Integrity

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Shows
BIO International Convention, June 13–16, San Diego

Value-Added
Ad Retargeting: Up to 10,000 Impressions

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

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Ad Close: May 12

Focus
Accelerating Drug Development

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Next-Generation Antibody Development

Upstream Processing
Media and Supplements

Downstream Processing
Process Chromatography

Manufacturing
Scaling Manufacturing Systems
Packaging Trends

Quality/Regulations
GMPs: Sterile/Aseptic Manufacturing

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Biosimilar Analysis

BioBusiness
Industry Consortiums

Outsourcing
Contract Testing Services

Departments
Regulatory Beat
Product Spotlight
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Pipeline: Nucleic Acids

Value-Added
Supplier Spotlight Listing Newsletter

Topics and trade show dates are subject to change.

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2022 EDITORIAL CALENDAR

BioPharm INTERNATIONAL®

July

Ad Close: June 13

Focus
Automation

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Cell Therapy Development

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Expression Systems

Downstream Processing
QbD for Downstream Processing

Manufacturing
Single-Use Consumables

Quality/Regulations
Form 483s and Warning Letters

Analytics
Extractables and Leachables Testing

Supply Chain
Shipping/Logistics

BioBusiness
Incubators

Outsourcing
Formulation

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Ask the Compliance Expert

Value-Added
Product/Service Profile Page (Full-page Advertisers)

August

Ad Close: July 15

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Outsourcing Strategies

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Scale up

Downstream Processing
Automating Downstream Processes

Manufacturing
Continuous Manufacturing
Facilities

Quality/Regulations
IND/NDA/BLA Filings

Analytics
Lab Data Management

BioBusiness
Investment Outlook

Outsourcing
Contract Packaging

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Product Spotlight
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Pipeline: Antibody-Drug Conjugates

Value-Added
Supplier Spotlight Listing Newsletter

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Ad Close: August 12

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Biopharma Research and Technical Advances

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Bioreactor Performance

Downstream Processing
Scale Up

Manufacturing
Process Modeling
Container Closures

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Audits and Inspections

Analytics
Adventitious Agent Testing

BioBusiness
Intellectual Property

Outsourcing
Bioanalytical Studies

Departments
Regulatory Beat
Product Spotlight
Ask the Compliance Expert

Shows
PDA/FDA Joint Regulatory Conference, TBD
BPI East TBD

Value-Added
Ad Retargeting: Up to 10,000 Impressions

Interactive eBook
Emerging Therapies ebook
Innovative process development and manufacturing practices to accelerate new biopharmaceutical modalities are explored.

Topics and trade show dates are subject to change.

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2022 EDITORIAL CALENDAR

BioPharm INTERNATIONAL®

October

Ad Close: September 14

Focus
Fill/Finish

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

Development
Formulation

Upstream Processing
Fermentation

Downstream Processing
Process Optimization

Manufacturing
Aseptic Manufacturing Processes

Quality/Regulations
Compendial Compliance Update

Analytics
Glycosylation

Supply Chain
Drug Product Security

BioBusiness
Global Biopharma Markets

Outsourcing
Bioprocessing Contract Services

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Regulatory Beat
Product Spotlight
Ask the Compliance Expert
Pipeline: Vaccines

Shows
CPhI Worldwide, TBD
AAPS 2022 PharmSci 360, Oct. 16–19, Boston

Value-Added
Fall Conference eNewsletter Profile

November

Ad Close: October 12

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Process Development and Control

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

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Antibody-Drug Conjugates Development

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Biochemicals and Raw Materials

Downstream Processing
Residual Impurities

Manufacturing
Contamination Control
Primary Packaging

Quality/Regulations
Supplier Oversight

Analytics
Environmental Control

BioBusiness
Economic Development

Outsourcing
Tech Transfer

Departments
Regulatory Beat
Product Spotlight
Ask the Compliance Expert

Value-Added
Button Ad (220x75 px) in BioPharm International's eBulletin Newsletter

Interactive eBook
Manufacturing and Facilities
Strategies and technologies to address limited production capacity for biopharmaceutical are covered in this interactive eBook.

December

Ad Close: November 10

Focus
Quality Control

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes
Biopharma Research and Technical Advances

Development
Process Development

Upstream Processing
Perfusion

Downstream Processing
Viral Clearance

Manufacturing
Digitalization of Manufacturing Systems
Facilities

Quality/Regulations
Final Product Inspection

Analytics
Stability Testing

BioBusiness
Emerging Companies

Outsourcing
Outsourcing Trends

Departments
Regulatory Beat
Product Spotlight
Ask the Compliance Expert
Pipeline: Gene Therapies

Value-Added
Corporate Capabilities Profile (Full-page Advertisers)

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