

BioPharm^{INTERNATIONAL}

The Science & Business of Biopharmaceuticals

2023 MEDIA PLANNER

Covering the biopharmaceutical development and
manufacturing industry since 1988

#1 SOURCE FOR PRINT, DIGITAL, AND CONTENT MARKETING SOLUTIONS

AN **MJ** life sciences[®] BRAND

BioPharminternational.com

The **MJH** life sciences[®] Advantage

60+

Brands

3.9M+

Email Reach

7M+

Unique Visitors
per Month

16M+

Average Page Views
per Month

1.9M+

Print
Circulation

1000s

KOL & SAP Relationships

1500+

Conferences & Events

As an MJH Life Sciences[®] brand you'll be partnering with the largest privately held medical media company in North America.

Speed to Market

Driven by our flexibility and entrepreneurial spirit

Relationships

Ability to leverage the credibility and experience of our key opinion leaders (KOLs) and strategic alliance partners (SAPs)

Audience

An unmatched active audience spanning 63 brands and 17 various specialties

Partnering with us means you'll reach your audience with the right message at the right time.



CERTIFICATE OF REGISTRATION
Information Security Management System
ISO/IEC 27001:2013

This is to certify that the Information Security Management System of:
MJH Life Sciences
2 Clarke Drive, Suite 100
Cranbury, New Jersey 08512
United States

Conforms with the requirements of ISO/IEC 27001:2013 for the scope listed below:

The ISMS supporting the confidentiality, integrity, and availability of customer data, supplier information, and MJH Life Science's internal data related to all sub-LLC's under the MJH Life Sciences LLC, all sales services as provided by all MJH Life Sciences Brands, all services as provided by the Truth Serum Network (Agency), all shared services located out of headquarters, and all supporting departments: HR, IT, and Finance located at the headquarters at 2 Clarke Drive, Suite 100, Cranbury, New Jersey environment.

MJH Life Sciences takes threats to the availability, integrity, and confidentiality of our clients' information seriously. As such, MJH Life Sciences is an ISO/IEC 27001:2013 certified provider whose Information Security Management System (*ISMS*) has received third-party accreditation from the International Standards Organization.

- ISO (*International organization of Standardization*) takes a risk-based approach to information security and risk mitigation
- ISO 27001 & ISO 27701 consist of 114 controls and 10 management system clauses
- MJH and their compliance partners conduct internal and external audits annually that test the efficiency and enforcement of all ISO controls



BioPharm International[®] covers all aspects of biopharmaceutical development, analysis, processing, and packaging, as well as business strategies and regulatory issues.

Key Topics

- Accelerating Drug Development
- Automation
- Biopharma Analysis
- Drug Delivery Systems
- Emerging Therapies
- Fill/Finish
- Outsourcing Strategies
- Process Development and Control
- Quality Control

Development

- ADCs Development
- Cell Therapy Development
- Drug Delivery Technologies
- Excipients
- Formulation
- Gene Therapy Development
- Monoclonal Antibodies Development
- Next-Generation Antibody Development
- Nucleic Acid-Based Therapeutics
- Process Development
- Regenerative Medicine Development
- Vaccine Development

Upstream Processing

- Automation
- Cell Culture
- Bioreactors
- Single-Use Systems
- Media and Supplements
- Expression Systems
- Scale up
- Fermentation
- Biochemicals and Raw Materials
- Perfusion

Downstream Processing

- Automating Downstream Processes
- Cell Harvesting
- Chromatography Resins
- Process Chromatography
- Process Optimization
- QbD for Downstream
- Processing Residual Impurities
- Scale Up
- Separation and Purification
- Single-Use Systems
- Viral Clearance

Manufacturing

- Aseptic Manufacturing
- Processes Cell and Gene Therapy
- Container Closures
- Contamination Control
- Continuous Manufacturing
- Digitalization
- Facilities
- Fill/Finish
- Lyophilization
- Packaging Trends
- Primary Packaging
- Process Analytical Technology
- Process Modeling
- Process Monitoring/Controls
- Scaling Manufacturing
- Systems Single-Use
- Consumables Sterilization
- Methods

Supply Chain

- Cold Chain
- Drug Product Security
- Materials Sourcing
- Shipping/Logistics

Analytics

- Adventitious Agent Testing
- Bioassay Development
- Biosimilar Analysis
- Cleaning Validation
- Data Integrity
- Environmental Control
- Extractables and Leachables
- Testing Glycosylation
- Lab Data Management
- Lab Operations
- Protein Characterization
- Stability Testing

Quality/Regulations

- Audits and Inspections
- CMC Strategies
- Compendial Compliance Update
- Data Mining CAPA
- Final Product Inspection
- Form 483s and Warning Letters
- GMPs: Emerging Therapies
- GMPs: Sterile/Aseptic Manufacturing
- Good Distribution Practices
- IND/NDA/BLA Filings
- Supplier Oversight

Outsourcing

- Bioanalytical Studies
- Bioprocessing Contract Services
- Clinical Trial Materials
- Contract Packaging
- Contract Testing Services
- Formulation
- Method Development
- Outsourcing Trends
- Preclinical Studies
- State of Outsourcing Industry
- Tech Transfer

BioBusiness

- Business Development
- Economic Development
- Emerging Companies
- Global Biopharma Markets
- Incubators
- Industry Consortia
- Intellectual Property
- Investment Outlook
- Partnerships

[To contribute, view the submission guidelines: biopharminternational.com/editorial_info](https://www.biopharminternational.com/editorial_info)

OUR DATABASE

BioPharm^{INTERNATIONAL}[®]



CUSTOM AUDIENCE SEGMENTATION TOOL

Meet your customers where they are —
in print, online, eNewsletters or webcasts.

CAST™ is the largest pharma/science global database in the market. This propriety tool contains over **700,000 unduplicated decision-makers** from global companies in the industry, allowing you to communicate with your target audience through the information channels they are using.

CAST™ Capabilities

- Contextual data based on specific article topics and content
- Behavioral data from email engagement metrics on every contact
- On-demand ad hoc filter options to select specific titles, companies, domains and other contact information

CAST™ Flexibility

- Updated monthly, with unsubscribe list and any hard bounce emails updated nightly
- Benefit from a unique list that is fine-tuned to your targeted audience
- Multidimensional targeting and segmentation
- CAST™ is flexible enough to reach the most niche audience, based on your business needs

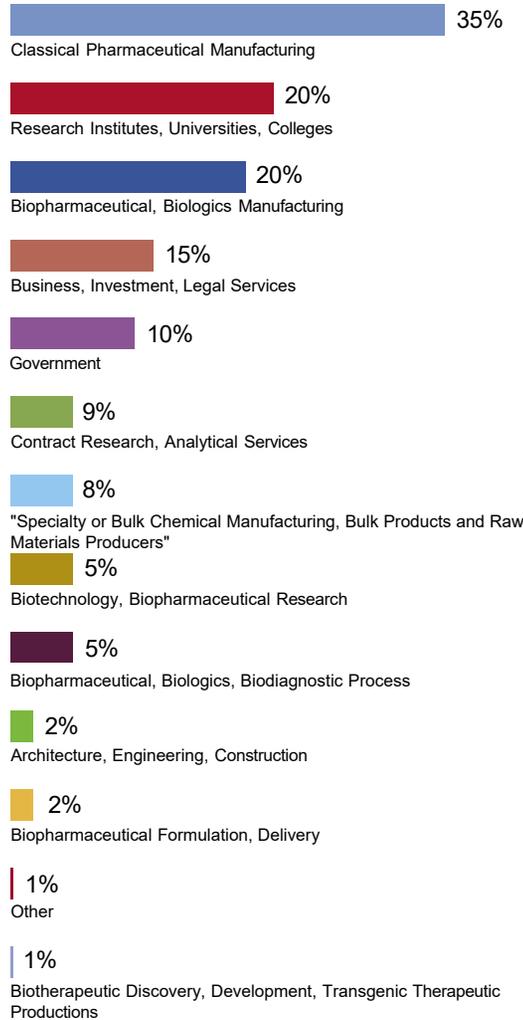


AUDIENCE – PUBLICATION

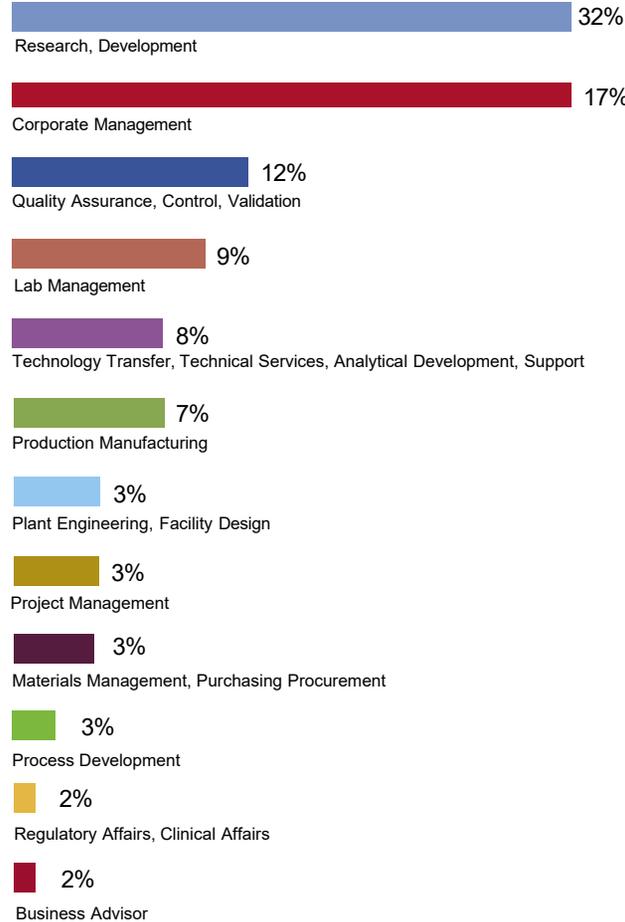
REACH 32,504 PRINT AND DIGITAL SUBSCRIBERS*

BioPharm International delivers a targeted audience of professionals involved in all stages of the development and manufacturing of biopharmaceutical therapeutics and diagnostics.

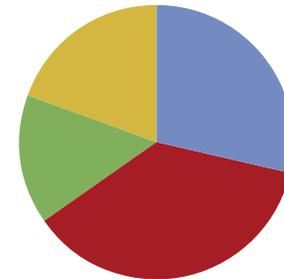
BUSINESS & INDUSTRY BREAKOUT*



TITLE & JOB FUNCTION BREAKOUT*

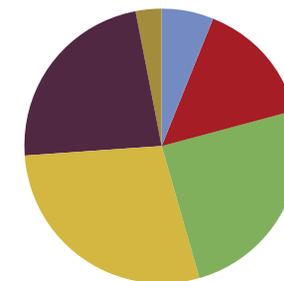


84.67% OF BIOPHARM INTERNATIONAL READERS PLAY AN ACTIVE ROLE IN THE PURCHASE OF PRODUCTS AND SERVICES IN THEIR ORGANIZATION.**



I make the decisions as part of a committee (28.67%)
 I recommend, influence or pass information to the decision-maker (36.67%)
 I am not involved (15.33%)
 I am the decision-maker (19.33%)

93.61% OF BIOPHARM INTERNATIONAL READERS HAVE MORE THAN 5 YEARS OF EXPERIENCE IN THE INDUSTRY.**



1-5 years (6.4%)
 6-10 years (15.12%)
 11-20 years (25.58%)
 21-30 years (29.07%)
 More than 30 years (23.84%)
 Less than 1 year (1.44%)

*Alliance for Audited Media December 2021 Audit Report. As filed with Alliance for Audited Media, subject to audit.

**August 2018 Readership Survey (multiple responses allowed).

AUDIENCE –

WEBSITE*

BioPharmInternational.com
Average Monthly Unique Browsers

61,884

Average Monthly
Page Impressions

176,027

eNEWSLETTERS*

Science & Business
eBulletin

Average Audited Distribution

26,968

Biopharma Knowledge
Resources

Average Audited Distribution

28,061

First Look**

Average
Distribution

23,189

BP Elements**

Average Distribution

29,720

BioPharm^{INTERNATIONAL}



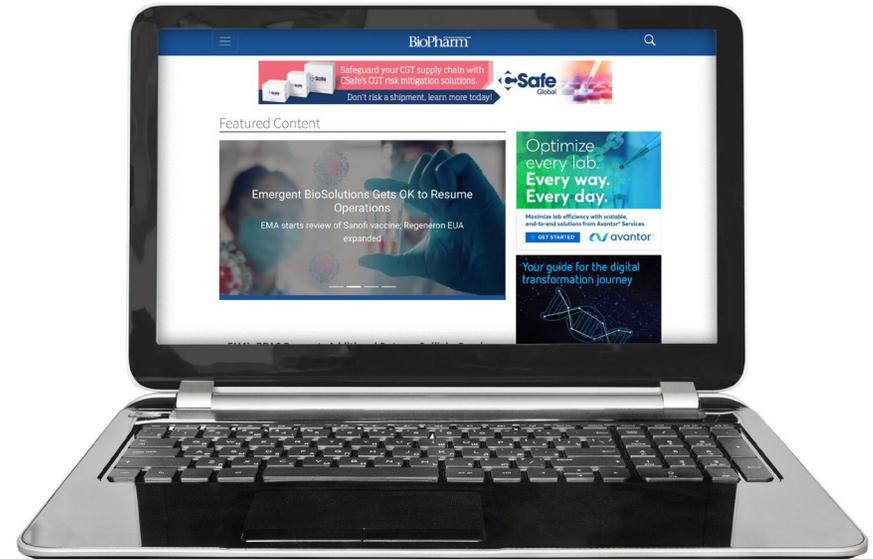
Available Opportunities

WEBSITE

- biopharminternational.com
- Banner Ads
- Expandable Video Banner Ads
- Interstitials
- Pre-Roll Videos
- Videos
- Sponsored Content
- Sponsored Link
- Ad Retargeting
- Geotargeting
- Native Advertising

eNEWSLETTERS:

- Banner ads
- Text ads



*AAM Audit, December 2021

**Publisher's Own Data

BioPharmInternational.com

BioPharmInternational.com provides comprehensive coverage for bioprocessing, quality, regulatory, and business topics including:

- Banner Ads
- Expandable Video Banner Ads
- Rich Media
- Geotargeting
- Native Advertising



Native Advertising

This program gives you the opportunity to inject thought leadership, insight, and brand awareness within the context of *BioPharm International's* trusted, editorial communities. You will receive a choice of topics, and in-article links to your gated content are served within relevant editorial content.



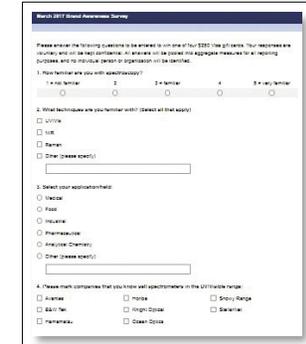
Sponsored Content Block

Exclusive sole-sponsored resource section on *BioPharm International's* website where your company can disseminate collateral, videos, whitepapers, and research, to drive website traffic, generate leads, and more. Link up to 4 assets plus your company's logo and website link. Your content block is NOT an ad unit and does not go into rotation, so it is visible 24/7 on virtually every page of *BioPharm International's* website. Each asset/link is tracked individually so you will know which assets are the most popular. For gated assets, all registration information will be provided.



Sponsored Survey Package

BioPharm International's sponsored web-based surveys can be used to better understand your client's business issues. These survey projects include a written report of findings, and can provide individual data on each survey respondent's habits and preferences.



CAST (Custom Audience Segmentation Tool)

Emails from our database that contains unduplicated decision-makers from global companies involved in industry science industries served by our leading publications.



Ad Retargeting

Once a visitor leaves *BioPharmInternational.com*, your display ad will follow them across the web, driving targeted groups to your website to learn more about your products or services.



Science & Business eBulletin

The Science & Business eBulletin is a great complement to your print and online advertising. It provides news and insights about technology and regulatory issues, the latest company changes, people moves and current conference calendar. Features include news, deals and alliances, people, products, and conferences.

First Look

BioPharm International's First Look is a monthly electronic newsletter that is sent to subscribers. It previews the latest issue of BioPharm International with links to online content and the digital edition of the magazine.

Biopharma Knowledge Resources

BioPharmInternational.com invites its audience of readers and site visitors to use the Knowledge Resources eLibrary at no charge. To download your whitepaper or application note, the viewer must complete a short response form including contact information and demographics. After the whitepaper is sent, you will receive an immediate email notification with the respondent's information. In addition, you will have access to real-time data containing all the leads generated via the password-protected website.

BP Elements

BP Elements delivers articles and news coverage on early-stage and preclinical biologics drug discovery and development, including pertinent research and cutting-edge technologies being developed in-lab for potential commercialization. Coverage will include organoids, gene-editing, data analytics, lab processes/workflows (e.g., cell counting, high-throughput screening for cell targets for potential use in biomanufacture, microscopy, fluorescent staining for protein/cell tagging), and lab technologies that specifically focus on providing deeper understanding of biological processes for the development of biologics, emerging therapies, and regenerative medicines.

VIDEO PROGRAMS AND EVENTS

VIDEO PROGRAMS

Extend your ROI at industry events with video content that can strengthen your brand reach post show with editorialized videos by BioPharm International[®] and audience engagement with the BioPharm International[®] community.

Presentation Showcase

Our Presentation Showcase program is comprised of a series of short, topic driven videos that combine our editor's interview with your speaker and outtakes from their presentation. Each package is promoted to your target audience – extending reach, expanding access and prolonging engagement well after the event has ended.

Thought Leadership Interview

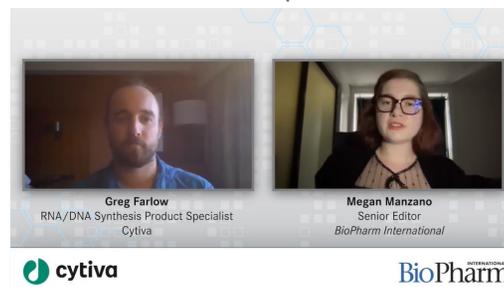
Our internal Studios team will coordinate an interview conducted by our editors and your KOLs that provides insight and digs into the key issues affecting our industry. Each video includes questions of your choice and promotion to our audience.

Exhibit Booth Interview

Extend your ROI at industry trade shows with custom video content that can strengthen your brand reach post show. We'll conduct an interview at your booth and edit it into a 3–5-minute video with promotions to BioPharm International[®] community.

Virtual Symposiums

Virtual Conferences offer an at-home alternative to in-person events, bringing critical information directly to the screens of industry professionals — without cutting corners on attendee experience. The virtual learning environment features many of the same amenities of a live trade show or meeting, including a lobby, auditorium, exhibit hall, networking lounge, and resource center.



LIVE EVENTS

Partner with BioPharm International[®] to create your custom live events. Our editorial and events teams work with you to develop and execute best-in-class programs that meet your business and educational goals. What makes BioPharm International[®] your partner of choice?

Content Development:

The BioPharm International[®] team works with your team to produce a program based on your needs.

KOL Recruitment:

We find the speakers who will attract your target attendees at the event as well as post event.

Attendee Recruitment:

BioPharm International[®] will find and attract the people you want to attend your live event.

Post-Event Content:

Our team will create video, audio and written content based on the program. And not only do we create the content, but we also provide marketing programs to get the content out to both attendees and non-attendees.

Turnkey Logistics:

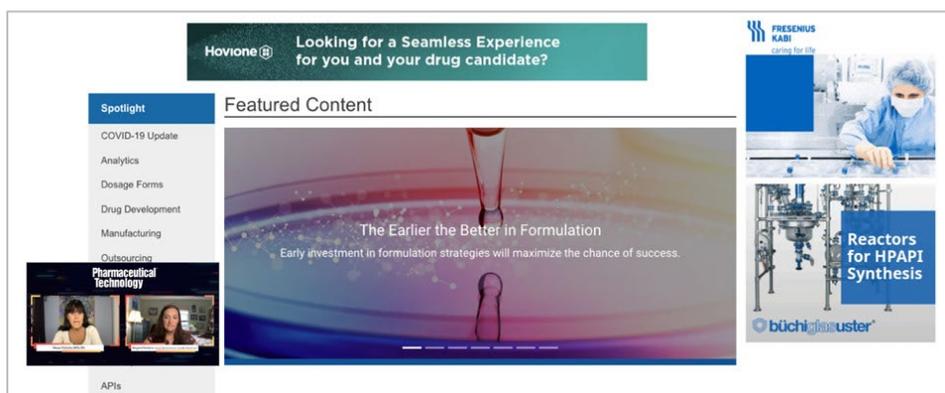
Besides the expertise of BioPharm International[®], you also get the meeting planning services of MJH Live Events to create a turnkey solution for your event.



DEDICATED VIDEO PLACEMENT

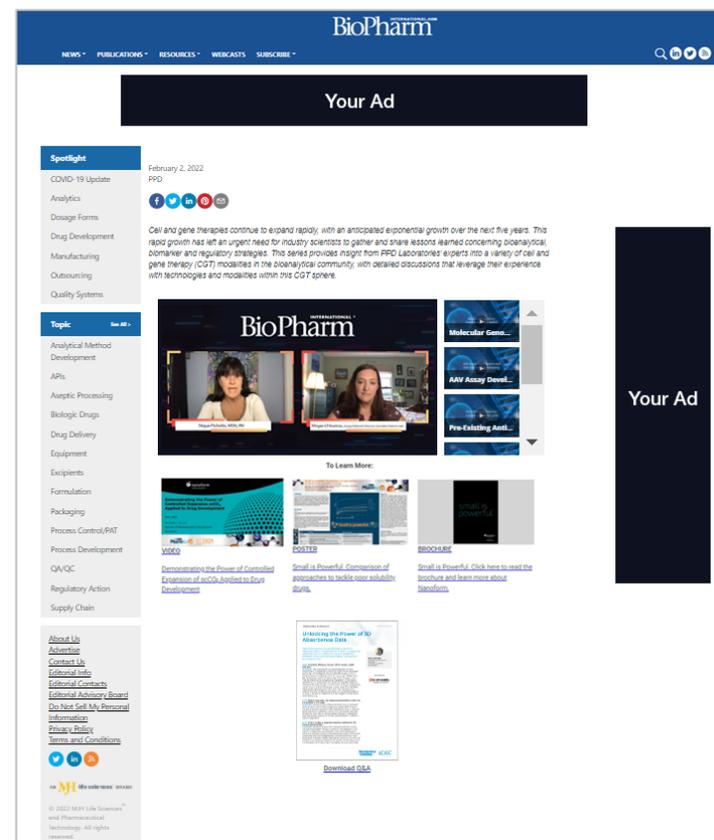
Our Dedicated Video Placement promotional program offers significant exposure to our audience right as they enter our website. This premium space places your video on our homepage as users scroll through content.

The homepage video is expandable to a larger screen and a dedicated landing page that places your video next to additional relevant content from your company.



PROGRAM FEATURES

- Your video featured on our website's home page
- Video expands to dedicated page with your branding and chosen related content
- Features your related content (whitepaper, additional videos, website, ebook, etc.)
- Exclusive branding, including your company's display ads



CONTENT MARKETING

Custom Content Creation

Our dedicated content editor will develop and write thought-provoking, insightful content about your products and services such as, but not limited to:

- Whitepapers
- Webcast summaries
- Conference presentation overviews
- Case studies
- Technical articles
- Roundtable discussions
- Infographics
- Thought Leadership interviews
- Digital primers
- Product profiles
- Market research reports
- New product write-ups
- Pharma Talks

Sponsored eBooks

A sponsored custom eBook or eBook series on topic(s) of your choice or a collaborative topic in conjunction with *BioPharm International's* editorial team. This program is designed to deliver high-quality leads.

BioPharm[®] INTERNATIONAL[®]



Lead Nurturing

Topic-driven programs that capture prospects and nurtures them by deploying high-quality content via strategically-timed communications. These programs are designed to deliver sales-ready leads.



Dedicated Dialogue

BioPharm International will conduct an interview with an expert from your company (scientist, corporate manager, etc.). This interview will be marketed through a multimedia program that includes a podcast and a two-page article in an issue of *BioPharm International*.



WEBCASTS

BioPharm International educational webcasts are led by credible moderators and offer exclusive sponsorship to a qualified audience, while embracing digital engagement.

Experience — More than 500 educational webcasts produced by MJH Life Sciences™ each year

Credibility — BioPharm International has been in the industry for over 30 years

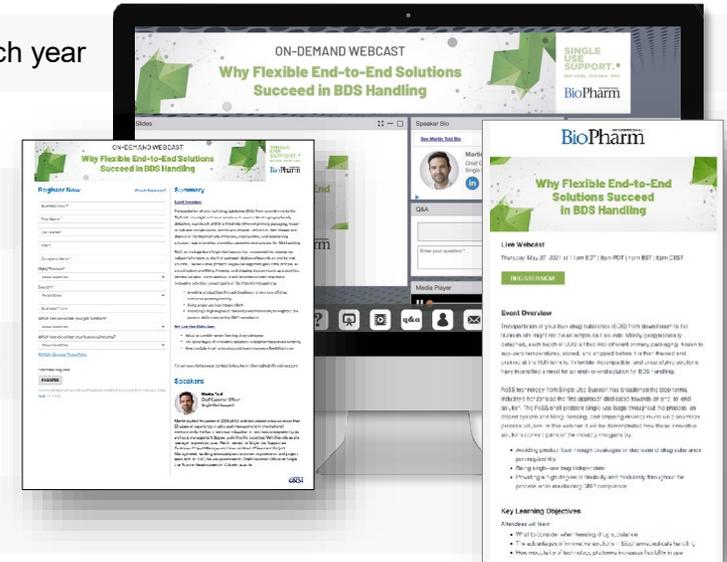
Talent — Respected speakers, producers and moderators from our editorial team

Audience/Reach — Select from 700,000+ qualified pharma/science professionals

Marketing & Promotion — Targeted audience development: print, digital and social media

Analytics — Comprehensive lead capture and data reporting for every event

Turnkey — Full-service management, marketing, training, production and hosting



Enhanced Webcasts

Cross-platform solutions that can convert a stand-alone educational webcast into an integrated content program

Utilizes social media, print and online marketing to amplify the content across the BioPharm International® community

- Repurposes webcast content cross-platform
- Delivers the content across multiple channels
- Extends reach, duration, and brand visibility
- Increases access and engagement



Turn your webcast into short-form easily digestible videos

- Post trailer on registration page
- Imbed clips into email marketing and social media posts
- Unpack long-form content into consumable moments to drive interest
- On-demand viewing

Data obtained from past MJH Life Sciences™ webcasts

PODCASTS

Podcast series regularly recorded by our editorial team that provides insights on news, new products, applications, and important trends in the industry. Our staff works with you on episodic programming to attract the most relevant listener base and to demonstrate your connection or expertise with the topic.

- Segmented by category
- Highlights important editorial topics
- New topics monthly
- Special coverage of industry events

These editorial podcasts are delivered to professionals via our brand's website and online audio channel.

BioPharm^{INTERNATIONAL}

The Science & Business of Biopharmaceuticals

2023 Editorial Calendar

Covering the biopharmaceutical development and
manufacturing industry since 1988



#1 SOURCE FOR PRINT, DIGITAL, AND CONTENT MARKETING SOLUTIONS

AN **MJ** life sciences[®] BRAND

BioPharminternational.com

SPECIAL ISSUES

March 2023

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 2023

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 2023

Emerging Therapies eBook

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

November 2023

Manufacturing and Facilities

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

SPONSORED-CONTENT EBOOKS

BioPharm International's content marketing team can develop custom publications on a range of biopharmaceutical topics for single sponsors. Contact your sales representative for details.

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 and click the "Editorial Information" link.

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.



2023 EDITORIAL CALENDAR

BioPharm^{INTERNATIONAL}

January

Ad Close: December 9, 2022

Focus

Biopharma Industry Outlook

Special Section: Employment Outlook, Training, and Survey

Special Coverage: Market Performance Measurements

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

Affinity Ligands

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

Tech Focus eNewsletter

Upstream Processing

February

Ad Close: January 13

Focus

Regenerative Medicine

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes

Biopharma Research and Technical Advances

Development

(CAR) T cell therapy

Upstream Processing

Cell Culture

Downstream Processing

Single-Use Systems

Manufacturing

Lyophilization

Facilities

FDA Voices

Quality/Regulations

CMC Strategies

Analytics

Product-specific ID and Potency (Functionality) assays

BioBusiness

Intellectual Property

Outsourcing

Method Development

Departments

Regulatory Beat

Product Spotlight

Ask the Compliance Expert

Pipeline: Protein

Therapeutics

Shows

Pittcon, March 18–22, Philadelphia

BioProcess International West, Feb, 27-March 3, San Diego

Value-Added

Supplier Spotlight Listing Newsletter

Tech Focus eNewsletter

Downstream Processing

March

Ad Close: February 10

Focus

Upstream Processing

Special Coverage: ICH9 Quality Risk Management

Introduction to series

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes

Biopharma Research and Technical Advances

Development

High-Titre Vector Producing Cells

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 20–23, New York City

PDA Annual Meeting, April 3–5, New Orleans

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.

Tech Focus eNewsletter

Analysis

Topics and trade show dates are subject to change.

2023 EDITORIAL

BioPharm^{INTERNATIONAL}

April

Ad Close: March 15

Focus

Downstream Processing
Special Coverage: ICH9 Quality Risk Management

Emerging Biopharma Topics

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

Development

Nucleic Acid-Based Therapeutics

Upstream Processing

Fixed versus Simulated Moving Bed Bioreactors

Downstream Processing

What's New in Downstream Technologies

Manufacturing

Process Monitoring/Controls

FDA Voices

Quality/Regulations

Data Mining CAPA

Analytics

Cleaning Validation

Supply Chain

Materials Sourcing

BioBusiness

Partnerships

Outsourcing

Bioprocessing Contract Services

Departments

Regulatory Beat

Product Spotlight

Ask the Compliance Expert

Pipeline: Cell Therapies

Shows

CPhI North America, April 25-27, Philadelphia
INTERPHEX, April 25-27, New York

Value-Added

Spring Conference eNewsletter Profile

Topics and trade show dates are subject to change.

May

Ad Close: April 14

Focus

Emerging Therapy Development and Manufacturing
Special Coverage: ICH9 Quality Risk Management
Introduction to series

Emerging Biopharma Topics

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

Development

Cell Therapy

Upstream Processing

Single-Use Systems

Downstream Processing

Cell Harvesting

Manufacturing

Process Analytical Technology
Sterilization Methods

Quality/Regulations

GMPs: Emerging Therapies

Analytics

Data Integrity

BioBusiness

Business Development

Outsourcing

Preclinical Studies

Departments

Regulatory Beat

Product Spotlight

Ask the Compliance Expert

Shows

BIO International Convention, June 13–16, San Diego
ASGCT, May 16-20, Los Angeles

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.

Tech Focus eNewsletter

Manufacturing

June

Ad Close: May 12

Focus

Accelerating Drug Development
Special Coverage: ICH9 Quality Risk Management

Emerging Biopharma Topics

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

Development

Next-Generation Antibody Development

Upstream Processing

Media and Supplements

Downstream Processing

Process Chromatography

Manufacturing

Scaling Manufacturing Systems
Packaging Trends

FDA Voices

Quality/Regulations

GMPs: Sterile/Aseptic Manufacturing

Analytics

Biosimilar Analysis

BioBusiness

Industry Consortiums

Outsourcing

Contract Testing Services

Departments

Regulatory Beat Product

Spotlight

Ask the Compliance Expert

Pipeline: Nucleic Acids

Value-Added

Supplier Spotlight Listing Newsletter

Tech Focus eNewsletter

BioBusiness Trends

Reports on investment trends, mergers, acquisitions, and business development opportunities

2023 EDITORIAL

BioPharm^{INTERNATIONAL}

July

Ad Close: June 13

Focus

Automation
Special Coverage: ICH9 Quality Risk Management

Emerging Biopharma Topics

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

Development

Cell Therapy Development

Upstream Processing

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

Departments

Regulatory Beat

Product Spotlight

Ask the Compliance Expert

Value-Added

Product/Service Profile Page (Full-page

Advertisers)

Tech Focus eNewsletter

Upstream Processing

August

Ad Close: July 14

Focus

Outsourcing Strategies
Special Coverage: ICH9 Quality Risk Management

Emerging Biopharma Topics

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

Development

Gene Therapy- Lentiviral vs AAV vectors

Upstream Processing

Scale up

Downstream Processing

Automating Downstream Processes

Manufacturing

Continuous Manufacturing

Cell Expansion

Quality/Regulations

IND/NDA/BLA Filings

Analytics

Lab Data Management

BioBusiness

Investment Outlook

Outsourcing

Contract Packaging

Departments

Regulatory Beat

Product Spotlight

Ask the Compliance Expert

Pipeline: Antibody-Drug Conjugates

Shows:

CHI Bioprocessing Summit, Aug. TBD, Boston

Value-Added

Supplier Spotlight Listing Newsletter

Tech Focus eNewsletter

Downstream Processing

September

Ad Close: August 11

Focus

Drug Delivery Systems
Special Coverage: ICH9 Quality Risk Management

Emerging Biopharma Topics

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

Development

Monoclonal Antibodies Development

Upstream Processing

Bioreactor Performance

Downstream Processing

Scale Up

Manufacturing

Process Modeling

Container Closures

Quality/Regulations

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.

Tech Focus eNewsletter

Analysis

Topics and trade show dates are subject to change.

2023 EDITORIAL

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

FDA Voices

Quality/Regulations

Compendial Compliance Update

Analytics

Glycosylation

Supply Chain

Drug Product Security

BioBusiness

Global Biopharma Markets

Outsourcing

Bioprocessing Contract Services

Departments

Regulatory Beat

Product Spotlight

Ask the Compliance Expert

Pipeline: Vaccines

Shows

CPhI Worldwide, Oct. 24-26, Barcelona

AAPS 2022 PharmSci 360, Oct. 22-26,
Orlando, FL.

Meeting on the Mesa, Oct. XX, California

Value-Added

Fall Conference eNewsletter Profile

November

Ad Close: October 12

Focus

Manufacturing at Point of Care

Emerging Biopharma Topics

Peer-Reviewed Research/Technical Notes

Biopharma Research and Technical Advances

Development

Antibody-Drug Conjugates Development

Upstream Processing

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.

Tech Focus eNewsletter

Manufacturing

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

FDA Voices

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)

Tech Focus eNewsletter

Automation

Topics and trade show dates are subject to change.

DIGITAL SPECIFICATIONS

Website Ad Units						
Creative Unit Name	Initial Dimensions (WxH in pixels)	Maximum Expanded Dimensions (WxH in pixels)	Max Initial File Load Size	Host-initiated Subload	Animation/Video Guidelines	Unit-Specific Notes (See General Ad Requirements)
Leaderboard - Desktop	728 x 90	728x270	200 KB	300 KB	15 sec max animation / 30 sec max video. Scrolling ISI animation must be less than 60 sec.	Expansion must be user- initiated by click
Leaderboard - Mobile	320 x 50 or 300 x 50	320x460 (full-screen)	50 KB	100 KB	15 sec max animation / 30 sec max video. Scrolling ISI animation must be less than 60 sec.	Expansion must be user- initiated by tap
Medium Banner	300 x 250	600 x 250	150 KB	300 KB	15 sec max animation / 30 sec max video. Scrolling ISI animation must be less than 60 sec.	Expansion must be user- initiated by click
Small Banner	300 x 100	Expansion not allowed for these units	100 KB	Not allowed for this ad unit	15 sec max animation length/Video not allowed for this unit. (If using animation, expansion is not allowed.)	
Half Page	300 x 600 (desktop only)	600 x 600	200 KB	300 KB	15 sec max animation / 30 sec max video. Scrolling ISI animation must be less than 60 sec.	Expansion must be user- initiated by click
Welcome Ad	640 x 480 or 300 x 250 (desktop only)	Expansion not allowed for these units	200 KB	300 KB	15 sec max animation length/Video not allowed for this unit. (If using animation, expansion is not allowed.)	
Super Leaderboard	970 x 90	970x300	200 KB	400 KB	15 sec max animation / 30 sec max video. Scrolling ISI animation must be less than 60 sec.	Expansion must be user- initiated by click
Floating Footer	1025 x 100, 970 x 90 or 728 x 90	970x300	150 KB	300 KB	15 sec max animation length/Video not allowed for this unit. (If using animation, expansion is not allowed.)	Expansion must be user- initiated
In-Article Display Ad	300x100	Expansion not allowed for these units	100 KB	Not allowed for this ad unit	15 sec max animation length/Video not allowed for this unit. (If using animation, expansion is not allowed.)	
Wallpaper/Gutter Ads	150x1050, 160x600, 120x600	Expansion not allowed for these units	200 KB	300 KB	Animation or video is not allowed for this unit.	Must be built by third party vendor, Spotable at an additional cost
In-Banner Video	300x250, 728x90, 300x600	300x250 > 600x250 728x90 > 728x270 300x600 > 600x250	200 KB	2.2MB Total load with video	Minimum 24 fps for video / 15 sec max length /1.1 MB additional file size allowed for host-initiated video / Unlimited file size for user-initiated video	Audio and video must be user initiated.
Video Pre-Roll Ad	16:9 preferred 4:3 accepted	N/A	10 MB	N/A	Length: 15 seconds for non-skip ad, 15-60 seconds for skippable ads	n/a

DIGITAL SPECIFICATIONS

eBulletin - Email Newsletter Creative Units							
Creative Unit Name	Initial Dimensions (WxH in pixels)	Maximum Expanded Dimensions (WxH in pixels)	File Format	Max Initial File Load Size	Host-initiated Subload	Animation/Video Guidelines	Unit-Specific Notes
Medium Rectangle	300x250	Expansion not allowed for this unit	Jpg, gif, png	50 KB	Not allowed for this unit	Gif animation : 15 second max	3rd party 1x1 impression tracking pixel and click URL accepted
Leaderboard	728x90	Expansion not allowed for this unit	Jpg, gif, png	50 KB	Not allowed for this unit	Gif animation : 15 second max	3rd party 1x1 impression tracking pixel and click URL accepted
Banner	468x60	Expansion not allowed for this unit	Jpg, gif, png	50 KB	Not allowed for this unit	Gif animation : 15 second max	3rd party 1x1 impression tracking pixel and click URL accepted
Text Ad	65 Word Max 1 Click Thru URL						3rd party 1x1 impression tracking pixel and click URL accepted
First Look - Email Newsletter Creative Units							
Creative Unit Name	Initial Dimensions (WxH in pixels)	Maximum Expanded Dimensions (WxH in pixels)	File Format	Max Initial File Load Size	Host-initiated Subload	Animation/Video Guidelines	Unit-Specific Notes
Medium Rectangle	300x250	Expansion not allowed for this unit	Jpg, gif, png	50 KB	Not allowed for this unit	Gif animation : 15 second max	3rd party 1x1 impression tracking pixel and click URL accepted
Leaderboard	728x90	Expansion not allowed for this unit	Jpg, gif, png	50 KB	Not allowed for this unit	Gif animation : 15 second max	3rd party 1x1 impression tracking pixel and click URL accepted
Banner	468x60	Expansion not allowed for this unit	Jpg, gif, png	50 KB	Not allowed for this unit	Gif animation : 15 second max	3rd party 1x1 impression tracking pixel and click URL accepted
Text Ad	65 Word Max 1 Click Thru URL						3rd party 1x1 impression tracking pixel and click URL accepted
Featured Product	3-4 word title 30-word description. If the word count is exceeded, the summary will be subject to revision by our editor. One product image (120x120 pixels, format is jpg, gif, or png, Max file size of 30kb) 1 Live Click URL						3rd party 1x1 impression tracking pixel and click URL accepted

DIGITAL SPECIFICATIONS

Preroll Ad			
Video Setting	Specifications		
File Format	H.264 (mp4)		
Audio Format	MP3 or ACC (Preferred)		
Aspect Ratio	H.264		
Frame Rate	24 or 30		
Length	6 - 15 seconds for non-skippable 15 - 30 seconds for skippable		
Max File Size 10MB	Low Resolution	Medium Resolution	High Resolution
16:9 Aspect Ratio	360p or less	360p - 576p	576p - 1080p
4:3 Aspect Ratio	480p or less	480p - 576	n/a
Video Target Birate	500 kbps - 700 kbps	700 kbps - 1500 kbps	1500 kbps - 2500 kbps for 720p 2500 kbps - 3500 kbps for 1080p
Site Served	<ul style="list-style-type: none"> » Must be uploaded to YouTube (send video URL, shortened URL not allowed) Must allow embedding » Must be public or unlisted » True streaming in not allowed 		
Third-party Served	<ul style="list-style-type: none"> » Must be SSL-compliant » VAST 2.0, Vast 3.0 or VPAID (VAST 2.0 will not be accepted for skippable ads) 		
Sponsored E-blast Guidelines			
Requirements	<ul style="list-style-type: none"> » HTML creative from client » Text back up from client » (optional) Subject line and preheader » Test and final seed list* 		
Additional Needs for UNBRANDED e-blasts	<ul style="list-style-type: none"> » Opt Out link on clients creative » Suppression file from within the last 10 business days from the client From line 		
Please send the following 5 business days prior to the send date	<ul style="list-style-type: none"> » The HTML (saved as an attachment, with images hosted to your server) » Text only file (saved in Notepad- with full URLs listed for all links. The text should mirror the words in the HTML and not include coding) Your suppression file: in excel (only if sending from your company name) » Subject line: (limit to under 50 characters/including spacing) » Test seed list: email address of those to receive the test to review » Final seed list: any additional email addresses that are not on the test list but need to receive the final deployment (up to 10) 		
Timeline	<ul style="list-style-type: none"> » MJH Life Sciences™ will follow up with a proof of the e-blast at least one business day prior to the scheduled deployment to the test seed list » Please review the proof and reply to the email with approval or changes marked in a PDF. If another proof is required, a revised test will be sent » MJH Life Sciences™ will confirm that the e-blast is scheduled to deploy on the specified date » By the 15th of the following month, MJH Life Sciences™ will provide delivery metrics for all that deployed within the month 		

GENERAL NOTES

File weight calculation: All files for the ad (.html, .js, .css, images, etc.) must be included as part of the maximum file weight calculation for all file load limits. Shared libraries are also included as part of the file weight calculation unless otherwise exempted. File weights are calculated after files have been compressed into gzip format. You can use this site to check if your creative is within our specs guidelines <http://html5.iabtechlab.com/needauth?redir>.

Initial file load: Includes all assets and files necessary for completing first visual display of the ad.

Host-initiated subload: Where allowed, additional files may load one second after the browser domContentLoadedEventEnd event. The ad should be able to “listen” for the browser domContentLoadedEventEnd event before subsequent files beyond the initial max file size may be loaded.

User-initiated file size: Ads that allow additional file size for host-initiated subload also allows for unlimited file load after user-initiated interaction. User initiation is the willful act of a user to engage with an ad. Users may interact by clicking or tapping the ad.

VIDEO REQUIREMENTS:

- File Format: H.264 (mp4)
- Audio Format: MP3 or ACC (Preferred).
- Aspect Ratio: 16:9 preferred, 4:3 accepted
- Frame Rate: 24 or 30
- Max File Size: 10MB
- Tags Accepted: VAST 2.0, VAST 3.0, or VPAID (VAST 2.0 will not be accepted for skippable ads). Must be SSL-Compliant.
- Video length: 15/30 sec

HTML5 NOTES:

HTML5 provides/introduces new options for developing ads. The IAB has developed “HTML5 for Digital Advertising” (<http://www.iab.com/html5>) to help ad designers provide ads in HTML5 unit that will perform more successfully across the display advertising ecosystem. Please review this document and adopt its recommendations to help improve HTML5 ad performance in the industry.

HTML5 DESIGN INDUSTRY STANDARDS INFO:

<http://www.iab.com/html5>

HTML5 REQUIREMENTS FOR GOOGLE AD MANAGER:

<https://support.google.com/admanager/answer/7046799>

GENERAL AD REQUIREMENTS (APPLY TO ALL DISPLAY ADS):

File Format - JPG, GIF, PNG, HTML5 (must be 3rd-party hosted)

Audio - Must be user-initiated. To allow for audio initiation in videos without player controls, a control may be included ofr user to initiate audio.

Max CPU - Ad not to exceed 30% CPU usage during host-initiated execution

Expansion - Must be user-initiated by click and served through a 3rd Party tag.

Hotspot - Not to exceed 1/4 size of ad. Initiated when curser rests on hotspot for atleast 1 sec. Must NOT initiate audio

Defining ad space - Ad unit content must be clearly distinguishable from normal webpage content (ad unit must have clearly defined borders and not be confused with normal page content).

Submission Lead Time - Minimum lead time for ad file submission is 5-7 business days before campaign start.

Max number of host-initiated file requests - Ad not to exceed 15 file requests during initial file load and host-initiated subload. Unlimited file requests allowed after user-interaction

PRINT SPECIFICATIONS

Print Ad Specifications

Ad Size	Bleed Ad						Non-Bleed Ad	
	Bleed Ad		Trim Size		Live Area			
	Width	Depth	Width	Depth	Width	Depth	Width	Depth
2-Page Spread	15.75"	10.75"	15.5"	10.5"	15"	10"		
Full page	8.00"	10.75"	7.75"	10.5"	7.25"	10"		
2/3 page vertical	5.25"	10.75"	5.125"	10.50"	4.625"	10"	4.5"	9.50"
1/2 page Horizontal	8.00"	5.375"	7.75"	5.25"	7.25"	4.75"	6.75"	4.625"
1/2 page Vertical	4.125"	10.75"	4.00"	10.5"	3.5"	10"	3.375"	9.50"
1/2 page Island	5.25"	7.75"	5.125"	7.625"	4.625"	7.125"	4.5"	7.00"
1/3 page Horizontal	8.00"	3.875"	7.75"	3.75"	7.25"	3.25"	6.75"	3.00"
1/3 page Vertical	2.875"	10.75"	2.75"	10.5"	2.25"	10"	2.125"	9.5"
1/3 page Square							4.5"	4.625"
1/4 page Square							3.375"	4.625"

MAGAZINE SIZE	
Bleed : 8" x 10.75"	Bleed (-) Trim = 0.125" each side
Trim : 7.75" x 10.5"	Trim (-) Live = 0.25" each side
Live Area : 7.25" x 10"	*All measurements in inches.

DIGITAL AD REQUIREMENTS

- Digital data is required for all ad submissions.** Preferred format is PDF/X-1a. Note that a standard PDF is not a preferred format: files should be a PDF/X-1a, which is a PDF subset specific to printing. Publisher shall have no obligation or liability to Advertiser of any kind (including, without limitation, the obligation to offer Advertiser make goods or any other form of compensation) if an ad is supplied to Publisher by Advertiser in any format other than our preferred formats. Non-preferred or non-acceptable formats will be charged a \$150 processing fee. All files should be built to exact ad space dimensions purchased.
- Publisher will not supply a faxed or soft proof for Advertiser-supplied files.** Advertiser is solely responsible for preflighting and proofing all advertisements prior to submission to Publisher. If Publisher detects an error before going to press, Publisher will make a reasonable effort to contact Advertiser to give Advertiser an opportunity to correct and resubmit Advertiser's file before publication.
- Ad Proofs:** To ensure that Advertiser's ad is reproduced correctly, a SWOP-certified color proof that has been made from the same file that Advertiser supplies to Publisher must be provided. Publisher cannot provide Advertiser any assurances regarding the accuracy of reproduction of any ad submitted without a SWOP proof. Publisher shall have no obligation or liability to Advertiser of any kind (including, without limitation, the obligation to offer Advertiser make goods or any other form of compensation) for any ad supplied to Publisher by Advertiser without a SWOP proof.



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knowledge that matters

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