IMAPAC
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EVENT BROCHURE

CLINICAL TRIALS FESTIVAL ASIA 2027

Asia's Foremost Clinical Trials Conference Uniting Clinical Trials Leaders

23 - 24 March 2027 Sands Expo & Convention Centre, Singapore

PART OF BIOPHARMA APAC CONFEX
BIOPHARMA ASIA PACIFIC CONFERENCE, EXHIBITION & AWARDS
MANUFACTURING | ADC | CGT | CLINICAL TRIALS | TIDES | SUPPLY CHAIN

OFFICIAL EVENT BROCHURE

https://infohub.imapac.com/clinical-trials-festival-asia-2027

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EVENT CONTACTS

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EVENT HIGHLIGHTS

300+	100+	30+
ATTENDEES	COMPANIES	SPEAKERS

DATE & VENUE

23 - 24 March 2027

Sands Expo & Convention Centre
Singapore

CO-LOCATED EVENTS

BIOSUPPLY CHAIN ASIA 2027
BIOLOGICS MANUFACTURING ASIA 2027
ADC ASIA CONGRESS 2027
OLIGOS & PEPTIDES WORLD ASIA 2027
CELL & GENE THERAPY WORLD ASIA 2027

AWARDS

ASIA-PACIFIC BIOPHARMA EXCELLENCE AWARDS 2027

Testimonials

Innovent

I was impressed by the seamless organization and the high quality of the presentations. The topics were covered comprehensively, and the attention to detail extended even to the catering - the food was excellent.

Senior Director and ADC MedChem Group Head
Innovent Biologics
China

Dr.Reddy's

A well-organized event with excellent coordination.

Biologics CAR-T, Head Quality
Dr. Reddy's Laboratories
India

PINOTBIO

Informative - well organized, chosen speaker, full of useful information. Good opportunity to connect with the Asian colleagues, early in the year.

CEO
PinotBio
South Korea

About the Event

Join us at Clinical Trials Festival Asia 2027, the region's premier clinical research conference in its 5th year. Connect with industry leaders to explore innovations in trial design, management, and decentralized approaches, while discussing partnerships, regulations, and digital strategies. The event highlights Asia's growing role in clinical research, particularly in Singapore, South Korea, and Australia, offering valuable networking opportunities with diverse professionals to enhance trial efficiency across the region.

WHAT WE'LL COVER

Key Themes

Advancement in APAC
The Global Clinical Trials Hub

Patient-Centric Trials
Patients Recruitment, Engagement, and Outreach

Advancing Pre-Clinical Research
Bridging the Gap: From Pre-Clinical Insights to Clinical Success in ADC Development

Oncology Clinical Trials
New Modalities & Industry Innovation

Trial Logistics in APAC
Transforming Clinical Trial Logistics in APAC

APAC CRO Showcase
End-to-End Clinical Research Capabilities

Sponsors & Media Partners

EXHIBITORS

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NETWORKING PARTNER

ALMAC

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2026 SPEAKERS

Naveen Jain
VP, Pre-Clinical
Panacea Biotec
INDIA

Maneesh Ghildyal
VP & Global Head, Projects & Engineering
Biocon Biologics
INDIA

Matthew Choo, Ph.D.
Associate Director, Data Science
MSD
SINGAPORE

AGENDA AT A GLANCE

CONFERENCE DAY 1 | 23RD MARCH 2027	CONFERENCE DAY 2 | 24TH MARCH 2027
APAC Trial Harmonization & Emerging Markets Multi-country Trial Design and Next-wave Market Spotlight	**AI & Digital Transformation in Trial Execution** AI in Feasibility, eSource, Protocol Design
POSTER PRESENTATION, EXHIBITION VIEWING, & 1-1 ATTENDEE NETWORKING	
Navigating the Future of Patient-Centric Trials Recruitment, Retention, and Patient Diversity	**Transforming Clinical Trial Logistics in APAC**
LUNCH POSTER PRESENTATION, EXHIBITION VIEWING, & 1-1 ATTENDEE NETWORKING	
Advancing Pre-Clinical Research Bridging the Gap: From Pre-Clinical Insights to Clinical Success in ADC Development	**APAC Showcase** From Early-Phase to Late-Stage: Showcasing End-to-End Clinical Research Capabilities
POSTER PRESENTATION, EXHIBITION VIEWING & 1-1 ATTENDEE NETWORKING	
Decentralized Trials, Digital Endpoints, and Wearables	**Closing Panel Discussion** Rethinking Clinical Development as a Continuous Journey
END OF CONFERENCE DAY 1	END OF CONFERENCE DAY 2
AWARDS GALA DINNER 2026	

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CONFERENCE AGENDA

Day 1

23rd March 2027

8:50

IMAPAC Representative

8:55

Chairperson Opening Remarks:

APAC Trial Harmonization & Emerging Markets: Multi-country Trial Design and Next-wave Market Spotlight

9:00

One Region, Many Rules: Designing Multi-Country Trials Across APAC's Regulatory Patchwork

- How regulatory convergence initiatives (and their limits) are shaping multi-country trial strategy today
- Practical strategies for harmonizing trial protocols, data standards, and ethics submissions across multiple countries
- Spotlight on next-wave markets and what sponsors need to know before expanding trials into them
- Lessons learned from early movers on de-risking multi-country trial design in emerging markets

9:50

One Region, Infinite Complexity: Reimagining Clinical Development Across APAC

- A big-picture reflection on the opportunities and risks of scaling clinical trials across APAC's diverse markets
- What true regulatory and operational harmonization would mean for sponsors, sites, and patients alike
- A call to action on how the industry can collectively unlock next-wave markets faster and more responsibly

Reserved for Technology/Solution Provider

10:15

The Future of Clinical Trials in APAC: Harmonization, Access, and the Next Wave of Markets

- A region-wide perspective on where APAC clinical trial harmonization stands today, and where it needs to go
- Why multi-country trial design is becoming a strategic imperative, not just an operational challenge
- A forward-looking view on which emerging markets will define the next decade of trial activity in APAC

Naveen Jain
VP, Pre-Clinical, Panacea Biotec

10:40

MORNING TEA BREAK, EXHIBITION VIEWING, 1-1 NETWORKING

Navigating the Future of Patient-Centric Trials: Recruitment, Retention, and Patient Diversity

11:15

Cracking the Code on Patient Recruitment: Strategies to Beat Enrollment Bottlenecks in APAC

- How to identify and overcome region-specific barriers to patient enrollment across diverse APAC healthcare systems
- Site selection and feasibility strategies that speed up recruitment timelines without compromising data quality
- Using digital outreach and community partnerships to widen the recruitment funnel in underserved markets

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11:40

Beyond Enrollment: Building Retention and Diversity into Trial Design

- Designing protocols that reduce patient burden and minimize dropout across long-duration, complex studies
- Practical approaches to embedding ethnic, geographic, and socioeconomic diversity into trial populations from the outset
- Case studies on retention strategies that worked (and didn't) in multi-country APAC trials

Reserved for Technology/Solution Provider

12:05

The Patient-Centric Trial: Decentralized Models, Digital Tools, and Inclusive Access

- How decentralized trial elements (telehealth, remote monitoring, e-consent) are reshaping patient experience in APAC
- Balancing technology adoption with accessibility for patients across varying digital literacy and infrastructure levels
- Measuring the real impact of patient-centric design on recruitment speed, retention, and data diversity

Reserved for Technology/Solution Provider

12:20

Rethinking Trial Access: Recruitment and Retention Models for a More Inclusive APAC

- Addressing structural and cultural barriers that limit patient access to clinical trials across APAC's varied healthcare landscapes
- Site and investigator engagement strategies that build trust and sustain patient participation through trial completion
- Frameworks for setting and measuring diversity targets that reflect real APAC disease burden and population demographics

12:35

LUNCH, EXHIBITION VIEWING, 1-1 NETWORKING

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Day 1

23rd March 2027

Advancing Pre-Clinical Research Bringing the Gap: From Pre-Clinical Insights to Clinical Success in ADC Development

2:15

Advancing Clinical Progress of Next-Generation ADC Pipelines in East Asia

- Sharing latest clinical trial updates and milestones from ongoing ADC programmes in East Asia
- Highlighting early efficacy, safety, and translational data across solid tumour and haematology indications
- Discussing regulatory pathways and strategies to accelerate ADC approvals in Asia and global markets

2:40

The Next Wave of ADC Approvals: Key Learnings from Recent FDA and EMA Submissions

- Clinical trial design strategies that accelerate ADC approval
- Overcoming safety concerns: Managing off- target toxicity
- Case studies of recent successful ADC approvals and regulatory hurdles

Reserved for Solution Provider

3:05

Translating ADC Innovation into Clinical Success Across East Asia

- Showcasing breakthrough clinical achievements and first-in-human data from East Asian biopharma companies
- Sharing lessons learned from ADC clinical development, manufacturing readiness, and global partnering efforts
- Evaluating the future landscape of ADC therapeutics including dual payloads, novel targets, and next-generation conjugation platforms

3:45

AFTERNOON TEA BREAK, EXHIBITION VIEWING, 1-1 NETWORKING

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Day 1

23rd March 2027

Decentralized Trials, Digital Endpoints, and Wearables

4:15

Decentralized Trials in Practice: Scaling Digital Endpoints and Wearables Across APAC

- Practical frameworks for implementing DCT models across APAC's varied regulatory and infrastructure landscapes
- Validating digital endpoints and wearable-derived data for regulatory acceptance and clinical decision-making
- Case studies on hybrid and fully decentralized trials that improved data quality and patient convenience

4:40

From Pilot to Standard: The Maturing Role of Wearables and Digital Endpoints in Clinical Trials

- How wearables and sensor-based endpoints are moving from exploratory tools to primary and secondary trial measures
- Overcoming data integration, standardization, and interoperability challenges across devices and platforms
- Regulatory perspectives on qualifying digital endpoints for use in pivotal APAC and global trials

5:05

Building Trust in Decentralized Data: Technology, Compliance, and Patient Experience

- Ensuring data integrity, privacy, and compliance when deploying remote monitoring and digital tools across APAC markets
- Balancing sponsor technology ambitions with site and patient readiness for decentralized trial components
- What sponsors need to know about local regulatory guidance on DCT and digital endpoint adoption in APAC

5:30

END OF CONFERENCE DAY 1

6:00

CLINICAL TRIALS EXCELLENCE AWARDS 2025

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CONFERENCE AGENDA

Day 2

24th March 2027

8:50

IMAPAC Representative

8:55

Chairperson Opening Remarks:

AI & Digital Transformation in Trial Execution: AI in Feasibility, eSource, Protocol Design

9:00

The AI-Ready Trial: Rethinking Feasibility, Data Capture, and Protocol Design for Speed

- What sponsors need to build AI capability into feasibility planning and site selection processes
- Overcoming adoption barriers to AI-enabled eSource across APAC sites with varying digital maturity
- How AI and predictive analytics are being used to design smarter, more efficient trial protocols from the outset

9:25

Smarter Trials by Design: AI's Expanding Role in Feasibility and Protocol Optimization

- How AI is helping sponsors move from reactive to predictive feasibility planning
- Reducing protocol amendments by using AI to stress-test trial design before activation
- What "AI-ready" eSource infrastructure looks like, and the gaps still holding APAC sites back

Reserved for Solution Provider

Day 2

24th March 2027

9:55

Agentic and AI Transformation of Preclinical Data Analysis (tbc)

- Why real-time eSource data capture is becoming a competitive necessity, not a nice-to-have
- How AI is being layered onto existing data systems to flag risks and inefficiencies earlier
- Balancing innovation speed with data integrity and regulatory expectations across APAC markets

Matthew Choo, Ph.D.
Associate Director, Data Science, MSD

10:20

MORNING TEA BREAK, EXHIBITION VIEWING, 1-1 NETWORKING

Transforming Clinical Trial Logistics in APAC [Presentations at BSCA Room]

11:10

Engineering the Next Generation of Biopharma Manufacturing: Building Scalable, Smart & Resilient Facilities

- Connecting planning, manufacturing and logistics through integrated digital platforms
- Using real-time data and predictive intelligence to improve end-to-end decision-making
- Automating workflows and reducing operational complexity across the supply chain
- Building scalable digital capabilities that strengthen resilience, agility and performance

Maneesh Ghildyal
VP & Global Head, Projects & Engineering, Biocon Biologics

11:35

Optimising Decentralised Clinical Trials Through Connected Logistics Technologies

- Enabling direct-to-patient and site-to-patient deliveries
- Real-time shipment visibility through connected monitoring platforms
- Reducing risk with predictive temperature and lane analytics
- Supporting patient-centric clinical trials with integrated logistics solutions

Reserved for Solution Provider

Day 2

24th March 2027

12:05

Transforming Clinical Trial Supply Chains for Faster & Smarter Research

- Demand forecasting and inventory strategies
- Enhancing visibility and coordination across sponsors, depots, and logistics partners
- Digital technologies to improve trial supply agility while maintaining regulatory compliance

12:30

LUNCH, EXHIBITION VIEWING, 101 NETWORKING1

APAC CRO Showcase: From Early-Phase to Late-Stage: Showcasing End-to-End Clinical Research Capabilities

2:15

The Access Equation: Connecting Clinical Evidence to Market Realities in APAC

- Turning post-authorization and real-world data into evidence that supports pricing and reimbursement decisions
- The expanding role of medical affairs in shaping payer conversations and health technology assessments
- Country-specific market access hurdles across APAC and how sponsors are adapting their approach

2:40

Building Seamless Clinical Development Pathways

- The strategic advantages of consistent data, teams, and infrastructure across the entire trial lifecycle
- How early-phase insights can directly inform and de-risk late-stage trial design and execution
- Showcasing real-world examples of end-to-end clinical research driving faster, more efficient program delivery

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3:05

After the Approval: Making Post-Authorization Data Work Harder for Market Access

- Using post-authorization commitments as a strategic tool for generating real-world evidence, not just a regulatory obligation
- How medical affairs functions are being restructured to support faster, more coordinated payer engagement
- Practical approaches to closing the gap between approval and actual patient access across APAC markets

3:20 **AFTERNOON TEA BREAK, EXHIBITION VIEWING, 1-1 NETWORKING**

Closing Panel Discussion: Rethinking Clinical Development as a Continuous Journey

4:15

Panel Discussion: One Partner, Every Phase: Rethinking Clinical Development as a Continuous Journey

- Why continuity from early-phase discovery through late-stage execution is becoming a competitive differentiator in APAC
- Breaking down the operational and data silos that typically separate early and late-phase trial work
- How end-to-end capabilities support faster, more informed go/no-go decisions across the development timeline

5:00 **END OF CONFERENCE DAY 2**

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Awards & Recognition

ASIA-PACIFIC BIOPHARMA EXCELLENCE AWARDS 2027

Asia-Pacific Biopharma Excellence Awards 2027

23RD MARCH 2027SANDS EXPO & CONVENTION CENTRE, SINGAPORE

The Asia-Pacific Biopharma Excellence Awards (ABEA) 2027 aims to honour exceptional achievements in the Asian biopharma sector, recognizing excellence in bioprocessing, logistics and supply chain management, antibody-drug conjugates (ADC), and clinical trials over the past year. Celebrating the remarkable contributions of leading biopharma professionals, organizations, and technologies, the awards spotlight trailblazing leaders and trendsetters shaping the industry's future while inspiring innovation for tomorrow's biopharma landscape.

Learn More About Our Awards

Expected Audience Snapshot

85%
ATTENDANCE FROM BIOPHARMA, BIG PHARMA, CDMOS & CROS

80%
DECISION MAKERS

BY PROFILE

Big Pharma	35%
Clinical-Stage Bio Start-ups and SMEs	20%
Other Biopharm	10%
Asian Biopharma	35%

BY JOB TITLE

VPs & Directors of Clinical Operations and Research	25%
Head of Clinical Development, Patient Recruitment and Procurement	20%
Head of Regulatory/ Policy	15%
C-Level	15%
Scientific	15%
Medical Affairs	10%

BY GEOGRAPHY

15%	Korea & Japan
5%	China, Taiwan & Hong Kong
40%	Singapore
15%	Indonesia, Thailand, India & Malaysia
10%	USA & Europe
15%	Australia

Who Should Sponsor?

- CONTRACT RESEARCH ORGANIZATIONS
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- CLINICAL TRIALS COMPANIES
- CLINICAL DATA MANAGEMENT
- MEDICAL MONITORING, BIOSTATICS
- BIOMARKERS
- CELL-LINES, CELL CULTURE MEDIA/LAB EQUIPMENT
- LOGISTICS SERVICE PROVIDERS
- DIAGNOSTICS
- LAB DESIGN
- SAFETY DATA REVIEW

Why Sponsor This Event?

Thought Leadership

Present Your Expertise to the Biopharma Industry's Key Stakeholders through a Variety of Available Speaking Packages.

Host a Panel Discussion

Moderate a panel discussion including one representative from your company with 3 representatives from the end user side to share views on a current industry topic

Duration: 45 minutes

Showcase Your Solutions to Educate the Market

Join in for dedicated CMO & CRO Showcases for German Market

Branding

Emphasize on your Message by Marketing and Promoting Alongside the Event

Networking & Lead Generation

Gain Exclusive Access to Contacts of High Profile Attendees through Intimate Networking Opportunities

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Our Past Partners

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MARK YOUR CALENDAR

Save the Date

23–24 MAR 2027

Sands Expo & Convention Centre

Clinical Trials Festival Asia 2027

ALL IMAPAC EVENTS

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Discover all upcoming and future IMAPAC events across Asia-Pacific — biopharmaceutical conferences, summits, and exhibitions for the year ahead. →

OR ADD THIS EVENT TO YOUR CALENDAR

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Scan to Register

Open camera, point at code, instant registration access.

Or click the QR to open registration directly.

Registration Form

CATEGORY	EARLY BIRD (By 3 Oct 2026)	REGISTRATION (By 5 Dec 2026)	STANDARD (By 6 Feb 2027)	FINAL CALL
BIOPHARMA, ACADEMIC & RESEARCH — BASED IN ASIA (APAC)				
2-Day Conference Pass **Job Profiles:** Manufacturing, Bioprocessing, Production, USP, DSP, Operations, Site Heads, Drug Substance and Product Manufacturing, Process Development, Quality, Analytics and other related technical designations	SGD 1,495	SGD 1,695	SGD 1,895	SGD 2,095
BIOPHARMA, ACADEMIC & RESEARCH — BASED OUTSIDE OF ASIA				
2-Day Conference Pass **Job Profiles:** Manufacturing, Bioprocessing, Production, USP, DSP, Operations, Site Heads, Drug Substance and Product Manufacturing, Process Development, Quality, Analytics and other related technical designations	SGD 1,995	SGD 2,195	SGD 2,395	SGD 2,595
BIG PHARMA, SOLUTION / SERVICE / TECHNOLOGY PROVIDERS & CDMO/CMO				
2-Day Conference Pass Business Development, Marketing & Sales profiles	SGD 2,995	SGD 3,195	SGD 3,395	SGD 3,595

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Asia & International Biopharma

☐ Buy 2, Get 3 passes
☐ Buy 3, Get 5 passes

Limited to professionals in the "BUYER" profile working in Process Development, R&D, QC & Assurance, Manufacturing Heads, Engineering, Upstream, Downstream or Analytical in Big Pharmas, International / Asian Biopharmas, and Academic & Research Institutes.

INNER CIRCLE Speaker referrals and professionals working in the same organisation as our speakers are entitled to a 15% discount.

- Access to all conference proceedings for up to 30 days.
- Site Tour Package is applicable only with the purchase of a conference pass.

Standard Package Group Discount

3 Delegates Save 10% ☐ 5 Delegates Save 20% ☐ 6+ Delegates Save 25% ☐

EVENT VENUE

Sands Expo & Convention Centre
Singapore
23 - 24 March 2027

3 EASY WAYS TO REGISTER

Online
Visit:
https://www.imapac.com/events/clinical-trials-festival-asia-2027#register

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United Kingdom

YOUR DETAILS

Name _____
Job Title _____ Organization _____
Address _____
Postcode _____ Country _____
Tel _____ Fax _____
Email _____
Your primary business _____
Authorising Manager _____ Authorising manager signature _____

PAYMENT DETAILS

Payment is due immediately upon receipt of this registration form if you pay by credit card or 3 working days after your receipt of payment notification. Non-credit card payment will incur an additional US$100 (Bank handling fee)

I want to pay: ☐ Credit Card ☐ Bank Transfer ☐ Cheque

Offline credit card: ☐ Visa ☐ MasterCard ☐ American Express

Online credit card payment: Please register at https://www.imapac.com/events/clinical-trials-festival-asia-2027#register

Name on Card _____ Card Number _____
Expiry Date (MM/YY) _____ CVC Code _____ Signature _____

TERMS & CONDITIONS

Payment & Registration
Full payment is due upon registration if you pay by credit card, or 3 working days after receipt of invoice. Registration will not be confirmed until full payment has been received.
Unpaid registration will be billed 50% of the registration fee if you do not attend the event. A copy of the conference documentation will be mailed to you. Staff at the event will request a credit card guarantee for delegates without proof of payment.

Ticket Classification
All registrations are reviewed to ensure correct classification. Biopharma tickets are reserved for professionals employed by pharmaceutical or biopharmaceutical companies that develop, own or commercialise therapeutic drug products. Service, Solution and Technology Provider tickets apply to organisations offering services, platforms, technologies, equipment, consultancy or manufacturing services to the industry (including CROs, CDMOs, vendors, consultants and suppliers).
Purchasing an incorrect ticket does not guarantee entry. IMAPAC retains full discretion to deny access, reclassify registrations and issue an invoice for the price difference. Incorrect ticket purchases under an inappropriate classification are not eligible for refunds under any circumstances.

Cancellation & Substitution Policy
Cancellations must be received in writing. If the cancellation is received more than 14 days before the conference, attendees will receive a full credit note towards a future conference. Cancellations received 14 days or less (including the fourteenth day) prior to the conference will be liable for the full fee.
A substitution from the same organisation can be made at any time in writing at no extra charge. In the event that IMAPAC cancels a conference, payments received at the cancellation date will be credited towards attendance at a future conference, or in the event of postponement by IMAPAC, towards the rescheduled date. Credit notes remain valid for twelve months.

Changes to Conference & Agenda
IMAPAC reserves the right to postpone or cancel an event, to change the location or alter the advertised speakers for an event. IMAPAC is not responsible for any loss or damage as a result of substitution, alteration, postponement or cancellation of an event due to causes beyond its control including, without limitation, acts of God, natural disasters, sabotage, accident, trade or industrial disputes, terrorism or hostilities. In the event that a conference is cancelled, IMAPAC is not liable for any costs incurred by delegates in connection with their attendance. Changes to the agenda will be updated on our website as soon as possible.

Data Protection
The personal information shown and/or provided by you will be held in a database. It may be used to keep you up to date with developments in your industry. Sometimes your details may be obtained or made available to reputable third parties for marketing purposes. If you do not wish your details to be used for this purpose, please write to: data@imapac.com.

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BIOPHARMA APAC CONFEX
BIOPHARMA ASIA PACIFIC CONFERENCE, EXHIBITION & AWARDS

MANUFACTURING | ADC | CGT | CLINICAL TRIALS | TIDES | SUPPLY CHAIN

THIS EVENT IS PART OF

Biopharma APAC Confex 2027

Connecting Leaders. Anchoring Partnerships.

Biopharma APAC Confex is Asia-Pacific's leading biopharma gathering, bringing together 2,000+ attendees and 700+ companies across co-located conferences spanning biologics manufacturing, clinical trials, ADCs, cell & gene therapy, TIDES, and supply chain innovation — supporting collaboration, knowledge sharing, and strategic partnerships across the region.

2000+	700+	250+	72+
ATTENDEES	COMPANIES	SPEAKERS	NETWORKING HOURS

CO-LOCATED CONFERENCES

- CELL & GENE THERAPY WORLD ASIA 2027
- OLIGONUCLEOTIDES & PEPTIDES WORLD ASIA CONGRESS 2027
- CLINICAL TRIALS FESTIVAL ASIA 2027
- ADC ASIA CONGRESS 2027
- BIOSUPPLY CHAIN ASIA 2027
- BIOLOGICS MANUFACTURING ASIA 2027

A VARIETY OF PRODUCTS TO CATER TO GLOBAL BIOPHARMACEUTICAL INDUSTRY'S MARKET INTELLIGENCE AND BUSINESS CONNECTIONS NEEDS.

01 B2B EVENTS

Thought-leadership, industry knowledge-sharing and quality networking are at the core of IMAPAC's B2B events.

Create an engaging experience for your audiences, explore our B2B Events Solutions.

02 360 MARKETING SOLUTIONS

Expand your reach and influence! Be it with customized events, digital marketing solutions, content creation or thought-leadership.

Tap into our pool of audience that spans across various sub-sectors and regions critical in the burgeoning APAC biopharmaceutical sector.

03 DASHBOARDS

A quick-view analytics tool to power your decisions. Access region or country-specific data to uncover pipeline information, product focus.

Customize the dashboard based on your needs. Drill down to the data you need using custom views.

04 CUSTOM RESEARCH

Based on your objectives and budget, you can customize a package to one that best meets your intelligence needs. This allows you to maximize your ROI without spending on items that are not in line with your strategies.

With access to a database of 60,000+ life science professionals. We conduct research with a highly-qualified audience.

05 REPORTS

Discover in-depth market information of the growing biopharmaceutical sector.

Learn More →

IMAPAC
IMAGINE YOUR IMPACT

HELPING THE GLOBAL BIOPHARMACEUTICAL INDUSTRY

Generate Tangible Growth

IMAPAC is the one-stop shop of market intelligence and business connections for the global biopharmaceutical industry. We help companies expand their influence, make data driven decisions through our industry intelligence, and link up with other biopharmaceutical professionals to exchange knowledge and engage in quality networking. At the core of IMAPAC is our mission to help biopharmaceutical businesses globally generate tangible growth. At the same time, our heart lies in giving back to society. We empower the next generation by equipping underprivileged children with an education.

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