



14 Years of Excellence: East Asia's Biggest Biologics Manufacturing & ADC Conference

📅 17 - 18 September 2026 Songdo Convensia, Incheon, South Korea

OFFICIAL EVENT BROCHURE

<https://infohub.imapac.com/bmk-adcbio-2026-brochure>

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EVENT CONTACTS

CONFERENCE PRODUCER

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EVENT HIGHLIGHTS

600+
ATTENDEES

180+
COMPANIES

50+
SPEAKERS

DATE & VENUE

17 - 18 September 2026

Songdo Convensia
Incheon, South Korea

About the Event

South Korea has firmly established itself as a global biologics powerhouse, driven by strong R&D investment, rapid innovation, and a world-leading biopharma ecosystem. Under the Biologics World Korea 2026 umbrella, two co-located conferences Biologics Manufacturing Korea (#BMK2026), now in its 14th edition, and ADC & Bioconjugate East Asia (#ADCBio2026) bring together industry leaders from South Korea, Japan, China, and Taiwan.

Together they span the full spectrum of next-generation biologics: upstream and downstream processing, AI-driven manufacturing, regulatory pathways, and supply chain optimisation, alongside ADC development, bioconjugation strategies, CMC considerations, and commercialisation. The result is a fully integrated platform connecting biopharma, CMOs/CROs, regulators, and academia bridging innovation from discovery to market.

WHAT WE'LL COVER

Key Themes



Biomedical Advancements & Innovations

The event explores the latest breakthroughs in biomedical research and technology, highlighting how cutting-edge innovations are transforming healthcare and improving patient outcomes. Discussions will cover emerging medical technologies, advancements in diagnostics, and the future of personalized medicine.



Biopharmaceutical Manufacturing

With the growing demand for high-quality and efficient drug production, the conference delves into modern manufacturing techniques, process optimization, and the integration of automation and AI in pharmaceutical production. Experts will share insights on streamlining operations, ensuring product consistency, and scaling up production to meet global healthcare needs.



ADC & Bioconjugate

The conference will feature insights from East Asia's leading ADC biopharma and big pharma, CMOs/CROs, regulatory authorities, and academic research institutes. Discussions will cover discovery and preclinical advancements, scalable manufacturing strategies, process optimization, CMC considerations, and market access strategies.



Regulatory & Compliance Frameworks

As the biopharmaceutical industry continues to evolve, regulatory requirements remain a crucial aspect of drug development and commercialization. Sessions will address the latest policies, compliance challenges, and strategies for navigating complex regulatory landscapes across different regions.



Collaborations & Networking

The event provides a platform for key industry players, researchers, and policymakers to connect and exchange ideas. Topics will include public-private partnerships, industry collaborations, and the role of government initiatives in supporting research and development. Strengthening networks and fostering international cooperation will be key discussion points.



Sustainability & Future Trends

The industry is moving towards more sustainable and environmentally friendly practices, and this theme will highlight innovative solutions for reducing waste, improving energy efficiency, and adopting green technologies. Additionally, experts will share insights on future trends, investment opportunities, and the next big breakthroughs shaping the future of biopharmaceuticals.



Advanced Modalities

The event will spotlight cutting-edge developments in novel therapeutic modalities, RNA therapeutics, cell and gene therapies, and AI-driven bioprocessing. With participation from Korea's leading biologics manufacturers and global industry leaders, this conference is essential for staying ahead in the fast-evolving biologics landscape.

Sponsors & Media Partners

SILVER



BRONZE



EXHIBITION & NETWORKING



ASSOCIATE



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Nathan Grace Niño · nathan.grace@imapac.com · +6569836134

2026 SPEAKERS

BMK-2026



Jong Gu Kim

Director, Quality
Lotte Biologics
SOUTH KOREA



Kihyun Park, Ph.D.

Lead Scientist in AAV Process
Discovery Part
Samsung Biologics
SOUTH KOREA



Dae Won Kim, Ph.D.

Chief Executive Officer &
Founder
ICM Co., Ltd.
SOUTH KOREA



Paul Y. Song

Chief Executive Officer
NKGen Biotech
UNITED STATES



Heechung Kwon

Chief Executive Officer
GencellMed
SOUTH KOREA



Sammi Hsu

Senior Manager
Ever Supreme Biotechnology
TAIWAN



Hideto Yamaguchi

Senior Vice President, Head of
Regenerative Medicine
Technology, Astellas
CEO, **Cellafa Biosciences**
JAPAN



Cheng Yi (Jerry) Kuo

Assistant General Manager
UWell Biopharma
TAIWAN



Young Jun Seo

CEO
Elonova
SOUTH KOREA



Prof., Dr. Lee, Beom-Jin

Professor
Ajou University
SOUTH KOREA



Chen Hong Hang

Global Solution Architect, Life
Science Segment
Schneider Electric
SINGAPORE



Eric Bishop

VP of Research and
Development
Cygnus Technologies
UNITED STATES



Hemasunder Reddy

Business Development Director
Pfanstiehl, Inc.
UNITED STATES



Yo Kyung Chung

CTO and Head of R&D
Rophibio
SOUTH KOREA

ADCBIO-2026



Doo Young Jung
Chief Executive Officer
PinotBio
🇰🇷 SOUTH KOREA



Yo-Sup Rew
Chief Technology Officer
Intocell
🇰🇷 SOUTH KOREA



Wei Han Lee
Director CMC
OBI Pharma
🇹🇼 TAIWAN



Won Gyun Ahn
Associate Director
Qurient
🇰🇷 SOUTH KOREA



Chung Yi Wu
CEO
CHO Pharma
🇹🇼 TAIWAN



Dr. John Luk
CEO & Founder
ArbeleBio
🇺🇸 UNITED STATES



Wei Ting Sun
Director R&D
HoneyBear Bioscience
🇹🇼 TAIWAN



Akshaya Bansal
Executive Director, ADC
Platforms
Hummingbird Bioscience
🇸🇬 SINGAPORE



Heon Chang Lim
Director, Formulation
Samsung Biologics
🇰🇷 SOUTH KOREA



Yi Yang
CEO & Founder
Glyco-therapy Biotech
🇨🇳 CHINA



Giorgio Salciarini
Head of Sales - Technical
Business Development
BSP Pharmaceuticals



Shawn Seong
CEO & Founder
Shaperon
🇰🇷 SOUTH KOREA

Agenda At A Glance

CONFERENCE DAY 1 17 SEPTEMBER 2026		CONFERENCE DAY 2 18 SEPTEMBER 2026	
			
Leadership Panel Discussion Scaling Biologics & CGT in East Asia: Innovation, Manufacturing Challenges & Collaborative Strategies		Redefining ADC Therapeutics Addressing Toxicity, Enhancing Efficacy, and the Rise of Dual Payload Designs	
MORNING TEA, EXHIBITION VIEWING, POSTER PRESENTATIONS & 1-1 ATTENDEE NETWORKING			
Quality, Scale-Up & CMC Excellence Bridging Development and Commercialisation	Discovery Innovative Pathways in ADC Discovery: Unlocking Next-Gen Targeting Strategies	CAR-T & CAR-NK Therapies Case Studies Next-Gen Cell Therapy & Preclinical Innovation	Advancing Payloads and Linker Technologies for Next-Generation ADCs
LUNCH, EXHIBITION VIEWING, POSTER PRESENTATIONS & 1-1 ATTENDEE NETWORKING			
Case Studies in Advanced Modalities	Optimizing ADC Target Selection and Engineering for Maximum Efficacy	Advancing Gene Therapy Optimizing Vectors, Editing Technologies, and Smart Manufacturing	
AFTERNOON TEA, EXHIBITION VIEWING, POSTER PRESENTATIONS & 1-1 ATTENDEE NETWORKING			
Closing Roundtable Session From Design to Delivery: Quality, CMC, and Clinical Insights Shaping the Next Wave of Biologics		Closing Roundtable Discussion Next-Generation Modalities: Advancing Cell Therapy, Gene Therapy, and ADCs from Bench to Market	
END OF CONFERENCE DAY 1			

Day 1

Thursday, 17 September 2026

OPENING THEME

Scaling Biologics & CGT in East Asia

Biologics Manufacturing Korea 2026

ADC & Bioconjugate East Asia 2026

8:50

IMAPAC Opening Remarks

8:55

Chairman Opening Remarks



Yi Yang

CEO & Founder, Glyco-therapy Biotech



Fireside Chat: Scaling Biologics & ADC in East Asia: Innovation, Manufacturing Challenges & Collaborative Strategies

9:00

Fireside Chat: Scaling Biologics & ADC in East Asia: Innovation, Manufacturing Challenges & Collaborative Strategies

- Innovation across biologics and ADC pipelines: how emerging platforms are driving the next wave of development in East Asia.
- Manufacturing and formulation readiness: quality and formulation challenges specific to scaling biologics and ADC production.
- Bridging innovator and manufacturer priorities: tech transfer, quality alignment, and collaboration as ADC and biologics programs move toward commercial scale.
- Cross-border collaboration and the road ahead: positioning Korea and East Asia's ecosystem for combined biologics and ADC growth.



Shawn Seong

CEO & Founder, Shaperon



Heon Chang Lim

Director, Formulation, Samsung Biologics

9:40

Arginine-Glutamate as Ideal Excipient for High Concentration Monoclonal Antibody Formulation.

- Need for higher dose of monoclonal antibodies(mAb) formulations
- Arginine-Glutamate a promising candidate to address concerns of aggregation, viscosity, liquid-liquid phase separation and longer storage
- Mechanism driving mAb stability in formulation containing Arginine-Glutamate

**Hemasunder Reddy**

Business Development Director, Pfanstiehl, Inc.

10:05

Fatty Acid-conjugated "Fattigation" Biomaterials for Improved Physicochemical and Biological Performance

- Fattigation-Fatty acid conjugation
- Synergistic drug delivery strategies
- Redox-sensitive fattigated hyaluronic acid nanoparticles
- Receptor-oriented cancer targeting

**Prof., Dr. Lee, Beom-Jin**

Professor, Ajou University

10:30

Morning Tea Break, Exhibition Viewing & 1-1 Attendee Networking

Biologics Manufacturing Korea 2026

**Quality, Scale-Up & CMC Excellence: Bridging Development and Commercialisation**

ADC & Bioconjugate East Asia 2026

**Discovery: Innovative Pathways in ADC
Discovery: Unlocking Next-Gen Targeting Strategies**

SESSION CHAIR

**Yi Yang**

CEO & Founder, Glyco-therapy Biotech

11:15

Data Integrity 2.0 (Strategic Quality Management and Integrated Lifecycle Collaboration, MAH - CDMO - Suppliers)

- The 2026 Regulatory Paradigm Shift
- Building an Honest Organizational Foundation from Quality Culture and Metrics perspective
- Lifecycle Collaboration Model between MAH - CDMO - Supplier
- Roadmap to 2026 and beyond

**Jong Gu Kim**

Director, Quality, Lotte Biologics

11:15

Engineering Next-Generation ADCs through Glycosite-Specific Glycan Design

- Engineering glycan structures to improve ADC potency, stability, and therapeutic index.
- Modulating Fc biology to reduce off-target toxicity while preserving antitumor efficacy.
- Enabling homogeneous bispecific, dual-payload, and next-generation antibody conjugates.
- Translating glycosite-specific conjugation into scalable CMC and manufacturing advantages.

**Chung Yi Wu**

CEO, CHO Pharma

Biologics Manufacturing Korea 2026

11:40

Building the Digital Foundation for Scalable Biopharma Manufacturing: Unified Operations for Resilience, Compliance, and Efficiency

- Biologics and pharmaceutical manufacturers face increasing pressure to improve operational efficiency, compliance, resilience, and scalability. However, many facilities still rely on fragmented building, utility, and power systems that create silos and limit visibility.
- This talk explores how unified, AI-powered operations can provide a smarter digital foundation for biopharma manufacturing.
- Using Schneider Electric's EcoStruxure Foresight as an example, the session will show how connected facility operations can improve uptime, streamline lifecycle management, support controlled environments, and enable more future-ready, data-driven operations.

**Chen Hong Hang**

Global Solution Architect, Life Science Segment, Schneider Electric

ADC & Bioconjugate East Asia 2026

11:40

Supporting ADC From IND To The Long Term Commercial Supply: A Step Ahead Against The Challenges Of The Bioconjugates

- Serving the Innovators from the candidate selection to the IND, from Phase I to the long term commercial supply
- Driving ADC Success with Flexible & High-Quality Solutions from Preclinical to Commercial
- Highlights on the approach to support early stage program through robust development studies.
- Ensuring right first-time-approved regulatory submission to save time and costs.

**Giorgio Salciarini**

Head of Sales - Technical Business Development, BSP Pharmaceuticals

Biologics Manufacturing Korea 2026

11:55

**Biosimilar Manufacturing:
Integrated CMC Strategies for
Global Development and Regulatory
Access**

- Evolving Biosimilar Market and Regulatory Environment
- Designing Analytical Similarity and CMC as One Integrated Strategy
- Managing Biosimilarity and Comparability from Development through Commercial Manufacturing
- Manufacturing Competitiveness: CoGS, Quality, and Supply Reliability

**Yo Kyung Chung**

CTO and Head of R&D, Rophibio

ADC & Bioconjugate East Asia 2026

12:05

**PdRaxane: A NanoMab Drug
Conjugate Addressing the
Limitations of Classical ADCs**

- Active tumor routing and retention through gp60, SPARC and PD-L1 targeting
- Non-covalent NanoMab assembly overcoming conventional conjugation and linker limitations
- Preclinical data show similar activity at 1/10 the dose — a dose-dependent reduction in toxicity that favors long-term therapy
- Integrated checkpoint blockade and nab-paclitaxel delivery remodels the immunosuppressive TME into a hot phenotype with increased effector T-cell activation

**Shawn Seong**

CEO & Founder, Shaperon

12:20

**Lunch Break, Exhibition Viewing & 1-1
Attendee Networking**

12:30

**Lunch Break, Exhibition Viewing & 1-1
Attendee Networking**

Biologics Manufacturing Korea 2026

ADC & Bioconjugate East Asia 2026

Case Studies in
Advanced ModalitiesOptimizing ADC
Target Selection
and Engineering for
Maximum Efficacy

1:45

Chemical Engineering of Next-
Generation mRNA Platforms for
RNA Therapeutics

- Innovative chemical engineering of mRNA to enhance stability, translation efficiency, and therapeutic performance
- Next-generation mRNA platform technologies beyond vaccines
- Expanding RNA therapeutics for oncology, autoimmune diseases, and other therapeutic applications
- Future opportunities for collaboration in next-generation RNA medicines across Asia

**Young Jun Seo**
CEO, Elonova

1:45

From Site-Specific Conjugation to
Dual-Payload ADCs: Enzymatic
Strategies for Next-Generation
Antibody Therapeutics

- Harnessing enzymatic glycan remodeling for precise antibody conjugation.
- Controlling conjugation sites and drug-to-antibody ratios.
- Stepwise conjugation strategies for dual-payload ADCs with tunable payload ratios.
- Expanding the design space of next-generation antibody conjugates.

**Wei Ting Sun**
Director R&D, HoneyBear Bioscience

2:10

Advanced Orthogonal Analysis of
HCP using Mass Spectrometry

- Overview of Mass Spectrometry in HCP Analytics
- Advantages of MS in Coverage Analysis
- Identification of HCP in DS
- Relative and Absolute Quantification of individual HCPs using MS

**Eric Bishop**
VP of Research and Development, Cygnus Technologies

2:10

Expanding the Therapeutic Index
and Overcoming Resistance via
Glycan Conjugation

- Enhanced safety & infinite creation through YTConju[®]
- Enhanced efficacy through a novel Topol
- Conquering resistance by synergistic dual-payload combinations

**Yi Yang**
CEO & Founder, Glyco-therapy Biotech

Biologics Manufacturing Korea 2026

2:25

Enhancing AAV Manufacturing Yield Through a Promoter-Optimized Mini-Helper Plasmid

- Developed a Mini-Helper plasmid - a minimized, promoter-optimized helper plasmid retaining only essential AAV helper genes.
- Boosted AAV productivity — achieved 2.0–6.5× higher AAV titers depending on serotype, with reduced cost and no compromise in particle quality.
- Validated scalability — consistent performance from flask scale up to 2 L and 200 L bioreactors, demonstrating a scalable, cost-effective AAV manufacturing solution.

**Kihyun Park, Ph.D.**

Lead Scientist in AAV Process Discovery Part, Samsung Biologics

ADC & Bioconjugate East Asia 2026

2:35

Formulation / Lyophilization Case Study for ADC Molecules: Challenges and Breakthrough

- ADC
- Formulation
- Lyophilization

**Heon Chang Lim**

Director, Formulation, Samsung Biologics

3:00

Afternoon Tea Break, Exhibition Viewing & 1-1 Attendee Networking

3:30

Closing Roundtable Session: From Design to Delivery: Quality, CMC, and Clinical Insights Shaping the Next Wave of Biologics

1. Quality by Design vs. Quality by Inspection: Closing the Gap Between Theory and Practice
2. CMC Risk Management: Identifying and De-Risking Critical Process Parameters Early
3. CDMO Selection: Beyond Capacity: What Really Predicts a Successful Partnership?
4. Resistance Mechanisms and Sequencing ADCs in Treatment Paradigms
5. Glycan Engineering for Improved ADC and Biologics Efficacy

4:30

Chairman Closing Remarks

4:35

END OF CONFERENCE DAY 1

CONFERENCE AGENDA

Day 2

Friday, 18 September 2026

OPENING THEME

Scaling Biologics & CGT in East Asia

Biologics Manufacturing Korea 2026

8:50

IMAPAC's Opening Address

8:55

Chairman's Opening Address



Wei Han Lee

Director CMC, OBI Pharma

ADC & Bioconjugate East Asia 2026

8:50

IMAPAC's Opening Address

8:55

Chairman's Opening Address



Redefining ADC Therapeutics: Addressing Toxicity, Enhancing Efficacy, and the Rise of Dual Payload Designs

Exhibition Hall, Ballroom B

9:00

Integrating Enzymatic Glycan Conjugation, Scalable One-Pot Manufacturing, and Advanced Analytical Control Strategies for Mono- and Dual-Payload ADCs

- Overcoming the limitations of traditional random conjugation through site-specific glycan remodeling and innovative linker technologies.
- Transforming enzymatic conjugation into a cost-effective, highly robust, and industrially scalable manufacturing process.
- Establishing analytical foundations, DAR characterization, and control strategies for mono- and dual-payload ADCs.



Wei Han Lee

Director CMC, OBI Pharma

Conference Producer:

Umairoh Nurul Hafifa · umairoh.nurul@imapac.com · +6285600287725

9:25

Beyond BsADCs: From Better Delivery to Checkmate Killing with Novel Proteasome Inhibitor Payload

- The ADC Evolution & The Lethality Gap
- The Next Wave: Overcoming Resistance
- Pioneering Novel Payloads (The QLi5 Platform)

**Won Gyun Ahn**

Associate Director, Qurient

9:50

OHPAS™, PMT™, and Nexatecan™ Technologies for Enhanced ADC Efficacy and Safety

- OHPAS™ linker design and PMT™ mechanism
- Proprietary Nexatecan™ and iso-Nexatecan™ TOP1 inhibitor payloads
- Next-generation ADC development enabled by IntoCell's platform

**Yo-Sup Rew**

Chief Technology Officer, Intocell

10:15

Morning Tea Break, Exhibition Viewing & 1-1 Attendee Networking*Exhibition Hall, Ballroom B*

Biologics Manufacturing Korea 2026

ADC & Bioconjugate East Asia 2026

**CAR-T & CAR-NK
Therapies Case
Studies: Next-Gen
Cell Therapy &
Preclinical Innovation**

SESSION CHAIR

**Hideto Yamaguchi**

Senior Vice President, Head of Regenerative
Medicine Technology, Astellas, CEO, Cella
Biosciences

**Advancing
Payloads and
Linker Technologies
for Next-
Generation ADCs**

11:00

**ICM-203 AAV Gene Therapy as a
Promising DMOAD Candidate for
OA Treatment**

- Target (Nkx3.2) discovery and validation for OA treatment
- Preclinical validation of ICM-203 for DMOAD activity
- First-in-human clinical development of ICM-203

**Dae Won Kim, Ph.D.**

Chief Executive Officer & Founder, ICM Co., Ltd.

11:00

**Dual-Payload ADC Platform:
Advancing Rational Payload
Combinations to Address Resistance**

- Addressing ADC payload resistance through rationally selected dual-payload combinations.
- Integrating payload selection, conjugation strategy, linker chemistry, and antibody engineering to optimize efficacy and minimize off-target toxicity.
- Streamlining development and manufacturing to accelerate candidate-to-IND progression.

**Akshaya Bansal**

Executive Director, ADC Platforms,
Hummingbird Bioscience

Biologics Manufacturing Korea 2026

11:25

Use of Troculeucel (Cryopreserved Enhanced Autologous NK Cell Therapy) for Neurodegenerative Diseases

- Troculeucel's mechanism of action in reducing CSF neuroinflammatory and protein biomarkers.
- Reviewing the early clinical efficacy of Troculeucel in AD, PD, and FTD patients.
- Discuss regulatory landscape in Asia to expedite development and commercialization of Troculeucel

**Paul Y. Song**

Chief Executive Officer, NKGen Biotech

ADC & Bioconjugate East Asia 2026

11:25

Combining Payload Selection, Linker Design and Site-specific Conjugation for Dual Payload ADC Development

- Importance of payload selection for the design of dual payload ADC - pharmacology and physicochemical properties perspective
- Exploitation of novel linkers to expand the applicable payload scope
- Improving stability of dual payload ADCs by applying site-specific conjugation

**Doo Young Jung**

Chief Executive Officer, PinotBio

Biologics Manufacturing Korea 2026

ADC & Bioconjugate East Asia 2026

11:50

Driving Innovation in Cell Therapy Manufacturing with AI and Robotics: From Process Optimization to Regulatory-Ready Platforms

- Feasibility study results on AI-driven process optimization for iPSC-derived NK cell production, utilizing machine learning to analyze process data generated by the dual-arm LabDroid "Maholo" automation system.
- The discussion will cover how AI-enabled data analysis and robotic automation can improve yield, enhance batch-to-batch consistency, and support robust cell quality in advanced cell therapy manufacturing.
- Outline a regulatory strategy for implementing robotic automation in GMP environments, including key considerations from interactions with regulatory agencies.

**Hideto Yamaguchi**

Senior Vice President, Head of Regenerative Medicine Technology, Astellas, CEO, Cellafa Biosciences

12:15

Lunch Break, Exhibition Viewing & 1-1 Attendee Networking



Advancing Gene Therapy: Optimizing Vectors, Editing Technologies, and Smart Manufacturing

Exhibition Hall, Ballroom A

SESSION CHAIR



Heechung Kwon

Chief Executive Officer, GencellMed

1:45

Advancing Allogeneic CAR-T for Solid Tumors: From Phase II Clinical Development to Next-Generation In Vivo CAR-T

- Clinical progress and Phase IIa development strategy of CAR001, an allogeneic off-the-shelf CAR-T therapy for solid tumors.
- Emerging clinical insights from recurrent glioblastoma and other solid tumor indications.
- Manufacturing and scalability considerations for next-generation allogeneic cell therapies.
- Leveraging clinical experience to develop EXO001, an in vivo CAR-T platform.



Sammi Hsu

Senior Manager, Ever Supreme Biotechnology

2:10

Clinical Development of CRA-T Cell Therapy: Manufacturing, Formulation and Administration.



Cheng Yi (Jerry) Kuo

Assistant General Manager, UWell Biopharma

2:35

Development of an Oncolytic Gene Therapy for Peritoneal Cancer Using Retargeted HSV Vectors

- To strictly redirect oncolytic herpes simplex virus (oHSV) to cancer-specific antigens, GencellMed developed the SMART™ HSV retargeting platform, a novel, double-retargeted oHSV system.
- GencellMed's lead oncology asset GCM-101: Unlike conventional oHSVs that largely retain natural viral tropism, GCM-101 is engineered to exclusively enter EpCAM-expressing cancer cells, thereby maximizing tumor selectivity and minimizing off-target infection of normal cells.
- Preclinical evaluations of GCM-101's broad therapeutic potential across solid tumors with high EpCAM expression, including ovarian, colorectal, gastric, pancreatic, and lung cancers.



Heechung Kwon

Chief Executive Officer, GencellMed

3:00

Afternoon Tea Break, Exhibition Viewing & 1-1 Attendee Networking

Premier Ballroom A

3:30

Closing Roundtable Session - Navigating Complexity: Practical Solutions for Cell, Gene, and ADC Therapeutics in East Asia**1. Cell Therapy Clinical Development: What Works and What Doesn't****Cheng Yi (Jerry) Kuo**

Assistant General Manager, UWell Biopharma

2. Next-Gen Payload and Linker Innovation: What's Actually Moving the Needle Post-ADC 1.0**Sammi Hsu**

Senior Manager, Ever Supreme Biotechnology

3. Beyond CAR-T: Overcoming Manufacturing and Access Bottlenecks in Solid Tumor Cell Therapies**Hideto Yamaguchi**

Senior Vice President, Head of Regenerative Medicine Technology, Astellas, CEO, Cellafa Biosciences

4:30

Chairman Closing Remarks

4:35

END OF CONFERENCE

Expected Audience Snapshot

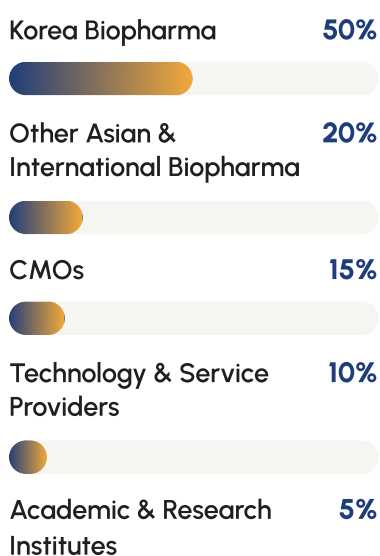
70%

ATTENDANCE FROM KOREAN BIOPHARMAS

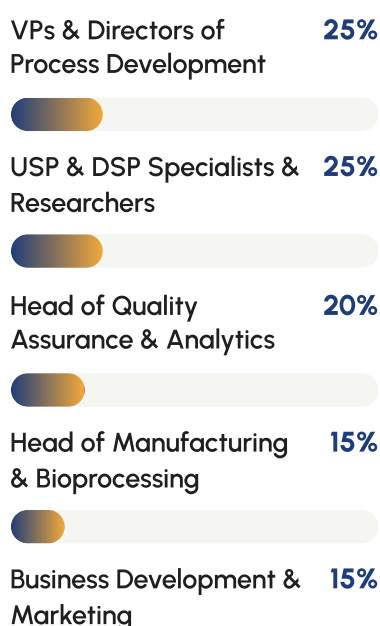
80%

DECISION MAKERS

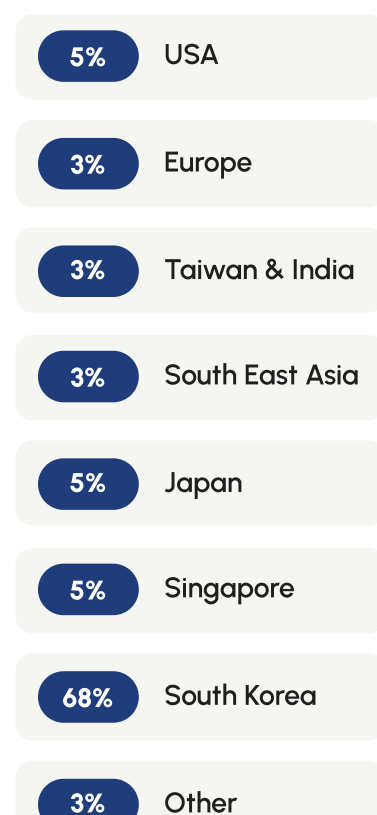
BY PROFILE



BY JOB TITLE



BY GEOGRAPHY



Who Should Sponsor?



Why Sponsor

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



Business Development Manager:

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MARK YOUR CALENDAR

Save the Date

17–18 **SEP**
2026 Songdo Convensia

BMK & ADCBio 2026 Brochure

ALL IMAPAC EVENTS



Explore Our Full Event Calendar

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

**Google Calendar**

One-click add to Gmail

**Outlook Calendar**

Add to Microsoft Outlook

**Scan to Register**

Open camera, point at code, instant registration access.

Or click the QR to open registration directly.

CATEGORY	EARLY BIRD (By 27 March 2026)	REGISTRATION (By 29 May 2026)	STANDARD (By 31 July 2026)	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
BRING YOUR TEAM - Group Discount				
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



Songdo Convensia, Incheon, South Korea

17 - 18 September 2026

3 EASY WAYS TO REGISTER

Online

<https://www.imapac.com/events/biologics-manufacturing-korea-2026#register>

Telephone

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Mail

Cheque payable to IMAPAC Pte Ltd,
36 Robinson Road, #02-01
068877
Singapore

Cheque payable to IMAPAC UK Ltd,
77 Farringdon Road, Farringdon, London
EC1M 3JU
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/biologics-manufacturing-korea-2026#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.

CATEGORY	EARLY BIRD (By 27 March 2026)	REGISTRATION (By 29 May 2026)	STANDARD (By 31 July 2026)	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
BRING YOUR TEAM - Group Discount				
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



Songdo Convensia, Incheon, South Korea

17 - 18 September 2026

3 EASY WAYS TO REGISTER

Online

<https://www.imapac.com/events/adc-bioconjugate-east-asia-2026#register>

Telephone

+65 6983 6140 (SG) / +44 161 552 8507 (UK)

Mail

Cheque payable to IMAPAC Pte Ltd, 36 Robinson Road, #02-01 068877 Singapore

Cheque payable to IMAPAC UK Ltd, 77 Farringdon Road, Farringdon, London EC1M 3JU United Kingdom

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Email	
Your primary business	
Authorising Manager	Authorising manager signature

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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 →](#)



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.

WEBSITE

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