

PMPS

Pharmaceutical Manufacturing and Packing Sourcer



2026

Media Pack

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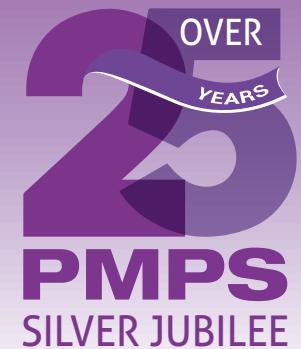
SAMEDAN
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Working With Us

Pharmaceutical Manufacturing and Packing Sourcer (PMPS) has built its reputation as an industry-leading platform over 25 years. It is a trusted medium to raise profiles plus promote products and services.

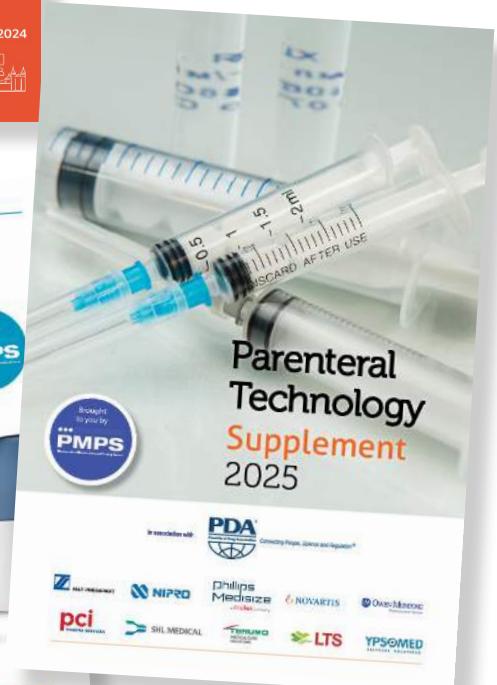
The advantages of appearing in the publication include:

1. Access to an international readership of supply chain decision makers
2. Association with strong, opinion-led editorial features on industry focus topics
3. Affiliation with a growing number of international, *PMPS*-sponsored trade events and conferences
4. Enduring advertising partnerships and a loyal affiliate network
5. Digital-first exposure to reach an even wider audience



About Us

PMPS is a specialist, B2B magazine providing in-depth coverage of topics vital to the pharmaceutical manufacturing, packaging and supply chain sectors. Published quarterly, it is a respected channel for senior executives and strategic decision makers to keep up to date with industry trends and the latest product developments. *PMPS* highlights the innovations and trends that matter through expert thought leadership and effective marketing opportunities.



Event-related Supplements

Be part of our event-related Supplements, partnered with several integral events across the pharma industry and covering key areas of drug delivery.

There are options for both editorial and promotional submissions:

SPOTLIGHT

Spotlight on emissions reductions for a lower CO₂ footprint in pharmaceutical packaging

When we are making concentrated efforts to become more sustainable, how will we approach reducing greenhouse gas emissions in its manufacturing processes?

At Christofor

Our packaging strategy, *Right packaging, Right place and time*, with a focused effort on the use of a sustainable approach, has been a reality and a success.

However, over the span of the last three years, we have dramatically chosen to approach approximately 40% lower (see Figure 1).

These statements that we will need to act swiftly and collaboratively to meet the Paris Climate Agreement of limiting global warming to 1.5°C will

carbon dioxide (CO₂) levels, matched with a significant reduction in the last two decades, CO₂ levels had remained relatively stable over the last 1,000 years. However, in the last 20 years, the CO₂ levels have increased by 300 ppm (Figure 2).

The planet is facing significant climate challenges, with the number of extreme weather events occurring more frequently. There has also been a significant increase in global average temperature from 1990 to 2005.

These statements demonstrate that we all need to act swiftly and collaboratively to meet the Paris Climate Agreement of limiting global warming to 1.5°C. The

chains they can include constraints from purchased materials for use in manufacturing, as well as constraints through the use of the company's own products by the end users.

Aptar's reduction initiative progress

Aptar has set well-defined science-based targets (2018) for Scope 1 and Scope 2 emissions, and is currently working on requirements to keep global warming at 1.5°C by 2030.

In addition, Aptar has a commitment to reduce its Scope 1, 2, as well as Scope 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117, 118, 119, 120, 121, 122, 123, 124, 125, 126, 127, 128, 129, 130, 131, 132, 133, 134, 135, 136, 137, 138, 139, 140, 141, 142, 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 153, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163, 164, 165, 166, 167, 168, 169, 170, 171, 172, 173, 174, 175, 176, 177, 178, 179, 180, 181, 182, 183, 184, 185, 186, 187, 188, 189, 190, 191, 192, 193, 194, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205, 206, 207, 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Biologics: the next frontier in injectable drug delivery

What if the future of injectable drug delivery looks real just on the drug itself, but on how easily and safely we extract the drug in a patient's hand?

By Richard Clark at Biovance Design

The pharmaceutical industry made an important breakthrough in injectable drug delivery, were redefined on how to extract the drug from the healthcare professionals, is now on the way to revolutionizing the way patients with chronic diseases are managed. This is the time when the pharmaceutical industry is revolutionized by the use of art and innovative technology. You have to understand that the future of healthcare is determined, but also how we can be more efficient, streamlined, and anchored.

The consequences of change, innovation, and the need for research and advanced delivery technologies is creating both significant challenges and growing areas of opportunity. As drug approvals, and pharmaceutical companies, are looking to the future, it is clear that the need for more drug approvals, and more efficient delivery, is a priority. In addition, as of 2025, a significant portion of the pharmaceutical market will be driven by the use of art and innovative technology. This is the time when the pharmaceutical industry is revolutionized by the use of art and innovative technology. You have to understand that the future of healthcare is determined, but also how we can be more efficient, streamlined, and anchored.

As the pharmaceutical industry moves forward, it is important to understand the challenges and opportunities that lie ahead. The future of healthcare is determined, but also how we can be more efficient, streamlined, and anchored.

Parenteral Technology Supplement 2025

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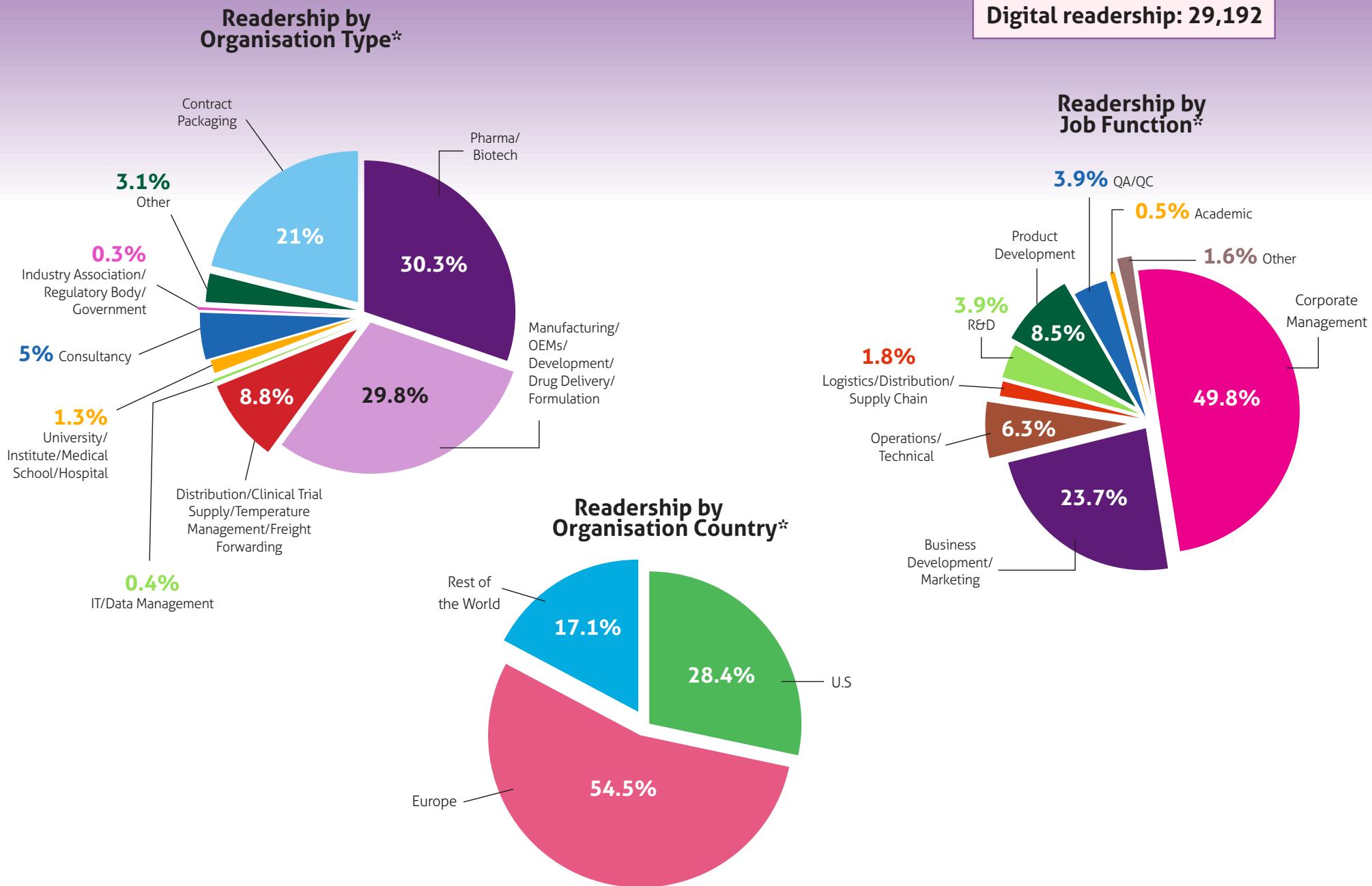
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| Issue | Winter 2026 | Spring 2026 | Summer 2026 | Autumn 2026 |
|-------------|--|--|---|--|
| Launches | January | April | June | September |
| Topics | <ul style="list-style-type: none"> Supply Chain – explore supply chain challenges and examine how they can be addressed through problem-solving in real-world scenarios Sustainability in Manufacturing and Packaging – delve into how sustainable practices in manufacturing and packaging can reduce environmental impact while maintaining efficiency and compliance Anti-Counterfeiting and Serialisation – examine strategies in anti-counterfeiting and serialisation to protect patients and ensure product integrity | <ul style="list-style-type: none"> Primary and Secondary Packaging – discuss innovations and challenges in primary (direct contact) and secondary (outer layer) packaging, from ensuring product protection to enhancing usability and compliance Tabletting and Capsuling – explore formulation and process optimisation in the making of tablets/capsules, to ensuring quality, efficiency and patient safety Cool Chain Developments – consider developments in the pharmaceutical cold chain, focusing on technologies and strategies that safeguard temperature-sensitive products in transit and storage | <ul style="list-style-type: none"> Cleanrooms and Sterilisation – give insight into maintaining controlled environments and effective sterilisation methods, ensuring safe, compliant and contamination-free pharmaceutical production New Technology in Pharma Packaging – highlight the latest developments in pharma packaging that enhance safety, efficiency, sustainability and patient engagement Innovative Drug Delivery – examine cutting-edge technologies in drug delivery, from advanced formulations to device innovations that optimise therapeutic outcomes | <ul style="list-style-type: none"> AI and Machine Learning – investigate how AI and machine learning are transforming pharma, from accelerating drug discovery to optimising manufacturing and supply chains Extractables and Leachables – discuss the impact of extractables and leachables on drug safety, packaging integrity and regulatory compliance Smart and On-Demand Packaging – discover interactive or tracking features alongside tailored, just-in-time formats in pharma packaging Medtechnology – examine how new medical technologies and digital tools are advancing drug delivery, diagnostics and patient care |
| Supplements | <ul style="list-style-type: none"> Pharmapack Europe Supplement | <ul style="list-style-type: none"> Inhalation Technology Supplement | <ul style="list-style-type: none"> Parenteral Technology Supplement | <ul style="list-style-type: none"> FlyPharma Conference Supplement CPHI WW Supplement DDL Supplement |

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Direct Marketing

Take advantage of Samedan's e-marketing service to send your own exclusive **email newsletter** to over 50,000 digital subscribers. This is a great opportunity to target the decision makers you want to reach with your news, special offer or latest product information – delivered direct to their inboxes.

Reach our database of quality-ensured and GDPR-compliant life science professionals spanning the entire pharma, biopharma and MedDev industry.

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Target specific job functions, organisation types and geographical locations

Use our e-marketing service as part of a customised multi-channel campaign
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Event Promotion

Why not promote your event with a tailored newsletter? Highlight speakers, list top exhibitors and offer an earlybird registration discount to tempt senior-level executives to attend. Let us help you design an impactful campaign – perhaps in conjunction with a dedicated event preview section in the magazine or a separate, special brochure (see page 10). Speak to our experienced team who can advise you on creating an effective omnichannel package that will get results.



Have the latest in quality innovation, and equip your business in packaging for the future. [REGISTER NOW](#)

It is time to understand the challenges and opportunities. This conference will bring together the best minds in the pharmaceutical industry to discuss the latest in parenteral packaging. [REGISTER NOW](#)

Discover critical topics, including: Container design, Regulators, packaging, pharmaceuticals, drug delivery, storage, transportation, and more.

You won't want to miss the insightful opening plenary featuring:

• A panel discussion from the impressive Gammel Allmene, Larsen LBB Designing, Heijen, and Kastell.

• A presentation from the highly regarded Novartis.

• Keynote address from the PDA.

Check out the [REGISTER NOW](#) section.

To provide you with the most recent information, news, and opportunities in the pharmaceutical industry, we are providing you with regular news, tips, and resources. [REGISTER NOW](#)

This conference is the place to be for the latest in pharmaceutical packaging. Take our newsletter.

[REGISTER NOW](#)

Make the Most of Your Time in Copenhagen.
Take your experience to the next level by signing up for related events!

Training:

Container Cleaning, Inspection, and Basic Course

Do you have the knowledge and skills to effectively inspect and maintain pharmaceutical containers? This course is designed for you.

Electronics and Laserscanners

Electronics and Laserscanners are becoming increasingly important in pharmaceutical packaging. This course will teach you how to effectively use them.

All about Pre-Filled Syringes: The Basics Course

From glass to containers, this course will teach you how to effectively inspect and maintain pharmaceutical containers.

Container Cleaning, Inspection, and Basic Course

Do you have the knowledge and skills to effectively inspect and maintain pharmaceutical containers? This course is designed for you.

Electronics and Laserscanners

Electronics and Laserscanners are becoming increasingly important in pharmaceutical packaging. This course will teach you how to effectively use them.

Reliable Pharma Needs Reliable Thermal Protection

Topa Thermal is a company that epitomises thermal innovation. We have devoted our efforts to temperature-controlled distribution. Our commitment to cutting edge technology, sets us apart.

Why mechanical testing of thermal packaging matters – Richard Harrop

From glass to containers, this course will teach you how to effectively inspect and maintain pharmaceutical containers.

Container Cleaning, Inspection, and Basic Course

Do you have the knowledge and skills to effectively inspect and maintain pharmaceutical containers? This course is designed for you.

Electronics and Laserscanners

Electronics and Laserscanners are becoming increasingly important in pharmaceutical packaging. This course will teach you how to effectively use them.

Reliable Pharma Needs Reliable Thermal Protection



Safeguarding the Pharmaceutical Cold Chain.
Why thermal & mechanical protection is critical for pharmaceuticals. [REGISTER NOW](#)

Topa Thermal is a company that epitomises thermal innovation. We have devoted our efforts to temperature-controlled distribution. Our commitment to cutting edge technology, sets us apart.

Please enjoy this special supplement, as we discuss the importance of thermal and mechanical protection for pharmaceuticals.

As we discuss the importance of thermal and mechanical protection for pharmaceuticals, we will also cover the latest in cold chain innovations, and best practices to improve your temperature-controlled supply chain.

The evolution of global health challenges and innovation in pharmaceuticals is highlighting the importance of being on the latest developments in this sector.

Join us for our 2024 PDA Biomanufacturing Conference, taking place from 24-26 September in Gothenburg, Sweden.

The conference is the ideal forum to explore new technologies, including the application of digital twins, development approaches, CMC strategies, and quality by design. Learn how these innovations support your business to meet the needs of their customers, providing a platform for networking and collaboration.

The 2024 PDA Biomanufacturing Conference is the place to be for the latest in pharmaceuticals. You will also have the opportunity to network with industry leaders, and regulators, as well as other organisations such as the US FDA, EMA, and other product safety agencies.

Discover how to reduce costs and increase efficiency in pharmaceutical manufacturing, advanced materials, manufacturing defects, and sustainability, focusing on your industry's needs for quality by design.

Additionally, our invited guest speakers will discuss the latest in pharmaceuticals and the pharmaceutical industry, including the latest in pharmaceuticals and the pharmaceutical industry.

Throughout the conference, industry and regulatory experts will present practical insights and solutions to help attendees ensure lifecycle drug quality assurance and CGMP compliance at their companies.

The conference continues to be the essential annual CGMP event to attend and receive live and highly relevant Center for Drug Evaluation and Research (CDER), Center for Biologics Evaluation and Research (CBER), Center for Veterinary Medicine (CVM), and Office of Regulatory Affairs (ORA) updates.

View the agenda.

Save up to 20% during our early registration period (ending 17 July).

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24-26 SEPTEMBER 2024
GOTHENBURG, SWEDEN

Shaping the Future of BioManufacturing

The evolution of global health challenges and innovation in pharmaceuticals is highlighting the importance of being on the latest developments in this sector.

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View the agenda.

Save up to 20% during our early registration period (ending 17 July).

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09-11 SEPTEMBER | WASHINGTON, DC

Joint Regulatory Conference 2024

CGMP: Leading with Quality and Integrity

The PDA/FDA Joint Regulatory Conference 2024, the annual FDA co-sponsored Current Good Manufacturing Practice (CGMP) conference, is taking place from 9 to 11 September in Washington, D.C.

The conference will offer a comprehensive agenda with substantive sessions and practical case studies that will provide insight into the everyday application of Current Good Manufacturing Practice (CGMP) and methodologies.

Attend to hear directly from Peter Marks, MD, PhD, Director, Center for Biologics Evaluation and Research (CBER), US FDA, and Patricia Cavazzoni, MD, Director, Center for Drug Evaluation and Research (CDER), US FDA.

Concurrent sessions will focus on topics related to practices and innovations, quality systems, and supply chains. FDA regulators continue to serve as session speakers and panelists.

Throughout the conference, industry and regulatory experts will present practical insights and solutions to help attendees ensure lifecycle drug quality assurance and CGMP compliance at their companies.

The conference continues to be the essential annual CGMP event to attend and receive live and highly relevant Center for Drug Evaluation and Research (CDER), Center for Biologics Evaluation and Research (CBER), Center for Veterinary Medicine (CVM), and Office of Regulatory Affairs (ORA) updates.

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Save up to 20% during our early registration period (ending 17 July).

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285

Registrations

1959

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Development of drug delivery devices goes through numerous critical stages. Of these stages, design transfer plays a significant role as part of the product lifecycle to assure the final device design is properly translated into production specifications. Given the complexity of a drug delivery device project, a sound design transfer process is necessary to meet the project timeline and cost.

Pharma and biotech companies then have the option to establish a final assembly and packaging infrastructure internally or they can partner up with third party contract manufacturers, or they can work with end-to-end solution providers like SHL to execute the final stage of their combination product projects.

SHL's Taras and Amy will explore the key differences, advantages, and potential project risks associated with each route. The speakers will also discuss the role of an integrated process development strategy – referencing SHL as an example – in developing an optimal design transfer and manufacturing solution to transform engineering designs into manufactured drug delivery devices successfully and consistently.

Key topics that will be covered in this webinar include:

- Design transfer in the drug delivery device industry
- Critical factors to consider in the design transfer processes
- Comparison of different final assembly and packaging options
- SHL Medical's process development strategy and capabilities

[Click here to book your place today.](#) If you require further information, please do not hesitate to contact us.

Not going to be able to make it? [Register now](#) and we'll send you a link to the on demand version after the live webinar.

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Company Profile

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Product Profile

£2,150

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- Highly configurable
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Product:

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Nemera's **discrete** system.

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COMPANY PROFILE

H&T Presspart

Bringing your drug delivery devices to market



IBT Preempt is a global leader in manufacturing drug delivery systems. With over 80 years of experience partnering with leading pharmaceutical companies, we are recognized for precision, reliability and quality – a critical concern for patients. We depend on our products.

Our corporate offices drug delivery systems for the pharmaceutical and high-volume markets, including oncology, cardiology, and

Pharming and biologics. We create custom samples. Through our extensive network of

As an efficient, lean, accountable
partner, we can help you
achieve your
manufacturing
and distribution
objectives. With manufacturing at
our business, we specialize
in manufacturing and design
engineering, efficiency and
cost reduction.

of programmes. 

centrifuge tubes
In France, we established
whole-cell productive cells,
with the best supplies
of chosen technologies,
and biocatalysis tooling,
and especially complete

For full-scale production, no
expensive mould tooling is
needed.

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A person is holding a blue sign with white text that reads "the heart of Pharma". The sign is partially visible, showing the top part of the text. The background shows an office environment with desks and papers.

A person is holding a teal ribbon banner that reads "Celebrate 30!" in white text. The banner is being held up, and the background shows a hallway with other people.

SPOTLIGHT

Championing a safe transition to low carbon inhalers

Edmunds with Prof. Balazs at OHF inhalers, Mark Althaus at Biogas, and Amanda Grace at GSK & Energy Materials to discuss the green transition and the challenges offering

II forward

Edmunds continues: "We are actively developing a low carbon alternative to the current product, which is a drug delivery system, known as a nebulizer, that disperses liquid medication into a fine mist that can be inhaled directly with a single breath. This is a much more efficient way of dispersing medication than using a metered dose inhaler (MDI), which is a device that releases a fixed amount of medication into the air when the user inhales into it, which requires a deliberate and controlled breath."

Edmunds is a cardiologist, with key leadership experience at Biogen, and at Celsius and Climate & Change, guiding their respective research and advocacy movements. He has also been a member of the World Health Organization's Task Force on Climate Change and its subcommittee on health. To this end, OHF has been working with the WHO to develop a framework for the safe and effective use of inhaled corticosteroids (ICS) in the tropics. As a speaker at the recent *ICM2021* conference, Edmunds highlighted the importance of the WHO's *ICS* guidance, which is designed to support the safe and effective use of ICS in low- and medium-HDI countries, as well as the WHO's *ICS* guidance for the safe and effective use of ICS in high-HDI countries.

He is optimistic in his transition to low carbon and is aiming to keep it up.

EDMUND: "Major pharmaceutical companies are making great strides in transitioning the way they do business. GSK has committed to becoming net zero by 2050 and is working to support its partners to do the same. AstraZeneca, Cipla, Novartis, and others are also making significant progress. There are many others that are working to develop regulatory pathways to demonstrate that the low carbon products are safe and effective. However, there are still many others, but they may have a smaller budget and less overall strength."

MARK ALTHAUS: "From a technology perspective, we are looking at a range of different approaches. We are looking at a range of next-generation products and we have many choices. The next-generation products are designed to reduce the energy consumption of the device, increase its lifetime, and reduce the ongoing costs of the device. However, the main driver is sustainability. Patients need to use

their inhalers more frequently. There is a technology perspective, and there is a business perspective. We are looking at a range of next-generation products and we have many choices. The next-generation products are designed to reduce the energy consumption of the device, increase its lifetime, and reduce the ongoing costs of the device. However, the main driver is sustainability. Patients need to use

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their inhalers more frequently. Patients need to use

"We are pleased to continue our partnership with PMPS, a publication that shares our passion for driving the pharmaceutical sector forward through groundbreaking innovation and meaningful connections"

– Tara Dougal, brand director for CPHI Worldwide

Promotional Opportunities with FlyPharma



Join us for the FlyPharma Conference and Exhibition

Each year Samedan's **FlyPharma Conference** series gets bigger and better. Hosted in cities with superior pharma cargo and logistics facilities across Europe – and farther afield – these events draw in the cream of the industry's air transport and supply chain executives. Alongside a speaker programme featuring senior executives discussing the latest industry trends and challenges, there is an outstanding exhibition of state-of-the-art equipment and services.

Prior to the event, *PMPS* publishes a comprehensive preview in its **FlyPharma Supplement**, showcasing the companies taking part and the highlights of the show. This is distributed in advance to our extensive readership via the digital edition, as well as physically in our top quality, glossy print magazine at the meeting, giving excellent publicity to participants. From advertorials like Spotlights, Company Profiles and Q&As, to pages of advertising, this is the perfect vehicle to show your expertise to your target market.

For more information, contact **Simon Caplan**
(simon@samedanltd.com)

FlyPharma Amsterdam 2025 touches down for pharma logistics excellence

What's in store for pharma logistics leaders in Amsterdam this year?

From 8-9 October, the FlyPharma Conference and Exhibition will be held at the RAI Amsterdam, the largest and most modern exhibition and conference centre in the Netherlands. Hosted by Samedan, the event will bring together over 200 pharmaceutical and logistics industry experts from around the world to discuss the latest trends and challenges in the industry.

The FlyPharma Conference will feature a range of speakers, including senior executives from pharmaceutical companies, logistics providers, and industry associations. The exhibition will feature over 50 exhibitors showcasing the latest equipment and services for the pharmaceutical supply chain.

As well as the conference and exhibition, the event will also include networking opportunities, including a welcome reception and a closing dinner.

SPOTLIGHT

Trends and expectations in the pharmaceutical industry: a DHL perspective

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FlyPharma Supplement
Uncover the potential in your pharma supply chain
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