



OXFORD
GLOBAL

4th Annual Cell & Gene Therapy Congress

25-26 October 2018 | London, UK

Day 1

Stream 1 – Cell & Gene Therapy Development and Clinical Trials

- Recent Clinical advances in:
 - Off-The-Shelf Natural Killer Cells
 - Targeting Solid Tumours
 - CAR T Therapies For Head & Neck Cancer
 - Rare & Severe Childhood Diseases
 - Chemotherapy & Radiation Exposure
 - Ophthalmological Disease
- Future challenges with the development of gene and advanced therapies
- Clinical safety testing models for effective and commercially successful CAR and TCR cell immunotherapies
- Patient focussed aspects to clinical development and clinical trials
- Panel discussion:** Future challenges with the development of gene and advanced therapies

Day 1

Stream 2 – Cell and Gene Therapy Bioprocessing and Manufacturing Strategies

- CAR T Including:
 - Allogenic CAR T Performance in Manufacturing & Industrialisation
 - CAR T Process Development & Manufacturing Considerations
 - Development of Gamma-Delta Cell Therapy
- Implementation of affordable And Scalable Manufacturing Strategies
- Quality Assurance Through the Entire Supply Chain
- Regulatory Considerations for Expedited Quality Development And Bioprocessing
- Viral Safety By Design For Cell & Gene Therapy Products
- Bypassing Genome Editing Tools
- Comparability For Viral Vectors

Day 2 Stream 1

Part 1 – Cell and Gene Therapy Manufacturing

- Workshop:** Cell and Gene Therapy Manufacturing
- Scale-Up Solutions And Automation In Manufacturing of Cell Therapies

Part 2 – Commercialising Cell & Gene Treatments

- Innovative Pricing Strategies For ATMPs
- Business Transformation And Partnering Strategies To Accelerate Commercialisation
- Lessons for CAR T Commercialisation from: Bluebird Bio And Celgene
- How Do We Ensure Affordability And Accessibility To Patients
- Regulatory Challenges For The Development & Approval Of Novel Cell & Gene Therapies
- Futures Of CAR T Therapies
- Panel Discussion:** Reimbursement And Pricing Strategies for Cell And Gene Therapies

Benefits to Attending

- ✓ **Hear from pioneers and leaders in the cell and gene therapy field.** 2018 Attendees include: Senior Vice President, GSK; Senior Vice President, Cellectis; Vice President, Janssen; Director, Biogen; and Vice President, Bluebird Bio
- ✓ **Discuss collaborative solutions to cell and gene therapy development and commercialisation challenges.** This event brings together innovators addressing current trends in novel therapeutic areas, Important regulatory considerations and critical reimbursement and pricing strategies
- ✓ **Discover the latest innovations in cell therapy bioprocessing and manufacturing.** Case studies include CAR T cell immunotherapies, the application of genome editing technology, novel vector systems and viral safety for gene therapy and scalable manufacturing strategies
- ✓ **Unparalleled networking opportunities.** The two-day congress offers dedicated networking breaks creating an interactive platform for scientific discussions and 1-2-1 meetings. The exhibition hall and poster presentation spaces offer a relaxed and professional environment for discussion
- ✓ **A high-quality programme devised with the help of our esteemed advisory board.** Presentations covering updates in upstream and downstream processing, recent advances in CAR T cell therapies and the future challenges for Advanced Therapeutic Medicinal Products
- ✓ This event is part of our flagship **Cell Series**, which also includes our **7th Annual Cell Culture & Bioprocessing, 5th Annual Stem Cell Congress** and inaugural **Biobanking Congress**

2018 Speakers Include:



Axel Hoos
GSK



David Sourdiv
Cellectis



Peter Olagunju
Bluebird Bio

Meet Senior Decision Makers

Over 300 VPs, Directors & Senior Managers from leading pharmaceutical organisations, biotech companies and academic institutions will attend the event. Delegate job functions include:

Cell Therapy
Gene Therapy
Regulatory Affairs

Process Development
Biotherapeutics
Commercialisation

Stem Cell Therapy
Cell Therapy Manufacture
Regenerative Medicine

Discover New Solutions

Formal and informal meeting opportunities offer delegates the chance to discuss key solutions with leading service providers. Services to be discussed include:

Cell Activation
Flow Cytometry
Separation Reagents

CRISPR, TALEN, RNAi
Tissue Printing
Cell Storage

Assay Development
Cryopreservation
Cell Line Development

For booking details & registration fees please visit:

<https://www.oxfordglobal.co.uk/celltherapy-congress/>

COMPLIMENTARY PRE-CONGRESS WEBINARS

Strategies In CAR T Development And Targeting –30th August 2018, 10am GMT(UK). 11am CET(Europe)

Led by:

Katrien Reynders, Programme Director, Celyad

Peggy Sotiropoulou, Research & Development Manager, Celyad

COMPLIMENTARY PRE-CONGRESS Q&As

Outlook of Cell & Gene Therapies in 2018

Michael Kalos, Senior Vice President, Immunooncology & Oncology Cellular Therapies, Janssen

CAR T Therapies – Targeting Unmet Medical Needs

Paul Rennert, President & Chief Executive Officer, Aleta Biotherapeutics

Reserved and Confirmed Cell & Gene Therapy 2018 Speakers:

- Axel Hoos, Senior Vice President & Oncology Therapeutic Area Head, GSK
- Michael Kalos, Vice President, Immunooncology and Oncology Cellular Therapies, Janssen
- David Sourdivo, Executive Vice President, Technical Operations, Cellectis
- Nebojsa Milovic, Vice President, Cell & Gene Therapy Pharmaceutical Sciences, Takeda
- Stefanos Theoharis, Senior Vice President, Corporate Development, Cell Medica
- Adrien Lemoine, Senior Vice President, Business Development & Alliance Management, Orchard Therapeutics
- Suyash Prasad, Senior Vice President & Chief Medical Officer, Audentes Therapeutics
- Lothar Germeroth, Senior Vice President, Juno Therapeutics
- Peter Olagunju, Vice President, Patient Operations & Lentiglobin Program Leader, Bluebird Bio
- Nina Kotsopoulou, Vice President, Process Development, Autolus
- Aniz Girach, Chief Medical Officer, Nightstar Tx
- Jens Hasskarl, Senior Medical Director, Clinical R&D & Program Lead, CRP – Juno CAR T Therapy, Celgene
- Diane Wilkinson, Director of Regulatory & CMC, Biogen
- Jerome Zeldis, Chief Medical Officer & President of Clinical Development, Sorrento Therapeutics
- Ram Mandalam, President & Chief Executive Officer, Cellerant
- Paul Rennert, President & Chief Executive Officer, Aleta Biotherapeutics
- Mark Plavsic, Chief Technical Officer, Lysogene
- Keith Foster, Chief Scientific Officer, Sutura Therapeutics and Professor of Translational Medicine, University of Reading
- John Maher, Chief Science Officer, Leucid Therapeutics and Senior Lecturer of Immunology, King's College London
- Owain Millington, Head of New Product Development, TC BioPharm
- Jennifer Chow, Global Commercial CAR T Lead, Celgene
- Lawrence Thompson, Principal Scientist & Group Leader, Pfizer
- Sarah Snykers, Quality Control Manager, Celyad
- Ricardo Baptista, Lead Scientist, Process Development
- Katy Rezvani, Chief of Cellular Therapy, Medical Director of GMP Facility, MD Anderson Center
- Panos Kefalas, Head of Health Economics & Market Access, Cell & Gene Therapy Catapult
- Magda Papadaki, Head of Manufacturing & Innovation, Association of the British Pharmaceutical Industry
- Paolo Morgese, Director of EU Market Access & Member Relations, Alliance for Regenerative Medicine



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For more information please contact marketing@oxforglobal.co.uk

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**4rd Annual Cell and Gene Therapy Congress
Day One – 25th October 2018**

	4th Annual Cell & Gene Therapy Congress	
07.30 – 08.20	Registration	
08.20 – 08.25	Oxford Global's Welcome Address	
08.25 – 08.30	Chairperson's Opening Address:	
08.30 – 09.00	Keynote Address: The Next Frontier – An Overview Of Living Therapies In Oncology Michael Kalos, Vice President, Immunooncology & Oncology Cellular Therapies, Janssen	
	Cell & Gene Therapy Development and Clinical Trials	Cell and Gene Therapy Bioprocessing and Manufacturing Strategies
	Stream Chair: To Be Confirmed	Stream Chair: Maria Rende, Senior Biotherapy Sales Specialist, Macopharma
09.00 – 09.30	Stream Keynote Address: Future Challenges With The Development Of Gene And Advanced Therapies Axel Hoos, Senior Vice President, Therapeutic Area Head of Oncology, GSK	Stream Keynote Address: Allogenic CAR T Performance In Manufacturing And Industrialisation David Sourdivo, Executive Vice President, Technical Operations, Cellectis
09.30 – 10.00	Solution Provider Presentation For Sponsorship Opportunities please contact sponsorship@oxfordglobal.co.uk	Solution Provider Presentation 
10.00 – 11.20	Morning Coffee & Refreshments, One To One Meetings x3, Poster Presentation Sessions	
11.20 – 11.50	Predicting Long Term Risk Benefit Of CAR T Cell Therapies Jens Hasskarl, Senior Medical Director, Clinical Research & Development & Program Lead CRP – Juno CAR T Therapy, Celgene	Regulatory Considerations For Expedited Quality Development & Bioprocessing Diane Wilkinson, Director, Regulatory CMC, Biogen
11.50 – 12.20	CAR T Anti-CD38 For Multiple Myeloma And Anti-CEA For Solid Tumours Jerome Zeldis, Chief Medical Officer & President of Clinical Development, Sorrento Therapeutics	Novel Process Technology For The Accelerated Manufacture Of CAR T Therapies Lothar Germeroth, Senior Vice President, Juno Therapeutics
12.20 – 12.50	Solution Provider Presentation For Sponsorship Opportunities Please Contact sponsorship@oxfordglobal.co.uk	Solution Provider Presentation For Sponsorship Opportunities Please Contact sponsorship@oxfordglobal.co.uk
12.50 – 13.50	Lunch, Poster Presentation Sessions	
13.50 – 14.20	Panel Discussion – Future Challenges With The Development Of Gene And Advanced Therapies Invited Moderator: Axel Hoos, GSK Invited Panelists: Katy Rezvani, Chief of Cellular Therapy, MD Anderson Center John Maher, Chief Scientific Officer, Leucid Bio & Senior Lecturer in Immunology, King's College London Keith Foster, Chief Science Officer, Sutura and Associate Professor In Translational Medicine, University of Reading	Viral Safety by Design for Cell And Gene Therapy Products <ul style="list-style-type: none"> • Product safety an important critical quality attribute of medicinal products. • Adventitious agents represent an undesired impurity and as such they impact product and patient safety. • ways to mitigate the risk of adventitious agents in the manufacture of cellular and gene therapy product, • looking at various interconnected links throughout the manufacturing process from starting materials and suppliers of raw materials to fill and finish. Mark Plavsic, Chief Technical Officer, Lysogene

**4th Annual Cell and Gene Therapy Congress
Day One – 25th October 2017**

4th Annual Cell & Gene Therapy Congress		
14.20 – 14.50	<p>Gene Therapy For Rare Diseases: Moving From Pre-Clinical Proof Of Concept To Clinical Trials</p> <ul style="list-style-type: none"> Approach to Designing Clinical Trials for Rare, Severe, Genetic Disease Informing Regulatory Strategy Clinical data from the X-linked Myotubular Myopathy and Crigler Najjar Programs <p>Suyash Prasad, Chief Medical Officer & President, Audentes Therapeutics</p>	<p>NKT Cells For The Development Of Off-The-Shelf CAR Products That Do Not Require Gene Editing</p> <ul style="list-style-type: none"> Natural Killer T-cells (NKT cells) are a highly-active subpopulation of T-cells with unique characteristics that render them ideal to off-the-shelf applications using chimeric antigen receptors (CARs). In addition, they are resistant to suppressive cells in the tumour micro-environment. NKT cells are invariant, targeting pathogen-specific glycolipids that are not present on human cells and do not therefore cause GvHD. They can therefore be used in allogeneic settings without the need for gene editing (CRISPR/TALEN/other) to remove their T-cell receptor. This reduces the complexity and cost of manufacturing CAR-NKT products and ensures all patients receive a consistent and efficacious product. Cell Medica is developing multiple products based on the CAR-NKT platform for indications of high unmet medical need, in partnership with the Baylor College of Medicine. The products are further engineered to secrete cytokines that further enhance their persistence and activity in solid tumours. <p>Stefanos Theoharis, Senior Vice President, Corporate Development, Cell Medica</p>
14.50 – 15.20	<p align="center">Solution Provider Presentation</p> <p align="center">For Sponsorship Opportunities Please Contact sponsorship@oxfordglobal.co.uk</p>	<p align="center">Solution Provider Presentation</p> <p align="center">For Sponsorship Opportunities Please Contact sponsorship@oxfordglobal.co.uk</p>
15.20 – 16.20	Afternoon Coffee & Refreshments, One To One Meetings x3, Poster Presentation Sessions	
16.20 – 16.50	<p>Optimising Delivery Of Antisense Gene Medicines</p> <ul style="list-style-type: none"> Improving Therapeutic Outcomes Decreasing Potential Toxicities Improving Cost of Goods <p>Keith Foster, Chief Science Officer, Sutura Therapeutics and Associate Professor In Translational Medicine, University of Reading</p>	<p>CAR T Process Development And Manufacturing Considerations (TBC)</p> <ul style="list-style-type: none"> CAR/TCR Approaches Assay Development Comparability <p>Nina Kotsopoulou, Director, Process Development, Autolus</p>
16.50 – 17.20	<p>Allogeneic MPC Therapy For The Treatment Of Severe Neutropenia Following Myelosuppressive Chemotherapy Or Radiation Exposure</p> <p>Ram Mandalam, President & CEO, Cellerant</p>	<p>Development Of Gamma-Delta CAR T Cell Therapies</p> <p>Owain Millington, Head of New Product Development, TC BioPharm</p>
17.20 – 17.50	<p>T4 Immunotherapy Of Head And Neck Cancer Using Pan-ErbB Retargeted CAR T Cells</p> <ul style="list-style-type: none"> A CAR has been engineered using a promiscuous ligand that engages 8 distinct ErbB dimer species Anti-tumour activity has been demonstrated in several pre-clinical solid tumour models Phase 1 evaluation is ongoing in patients with head and neck cancer, using intra-tumoural delivery and phased dose escalation to mitigate risk <p>John Maher, Chief Scientific Officer, Leucid Bio & Senior Lecturer in Immunology, King's College London</p>	<p>Quality Control and Assurance Throughout The Full Supply Chain Of Autologous CAR-T Products</p> <ul style="list-style-type: none"> Quality Control throughout manufacturing of autologous CAR-T (start + raw material, process, QC testing, product delivery) -GMP, ICH, USP & PhEur guidance <p>Sarah Snykers, Quality Control Manager, Celyad</p>

**4th Annual Cell and Gene Therapy Congress
Day One – 25th October 2017**

4th Annual Cell & Gene Therapy Congress		
17.50 – 18.20	<p>First-In-Human Trial Of Off-The-Shelf Cord Blood Derived CAR-Engineered NK Cells For The Treatment Of Cancer</p> <ul style="list-style-type: none"> • Use of human cord blood to produce NK cells engineered with a CAR to redirect specificity and enhance persistence • Expression of CARs with novel signaling domains in NK cells • Data from the first-in-human Phase I/II study of CAR-engineered NK cells to target relapsed/refractory lymphoid malignancies <p>Katy Rezvani, Chief of Cellular Therapy, MD Anderson Cancer Center</p>	<p>Comparability For Viral Vectors</p> <p>Lawrence Thompson, Principal Scientist, Pfizer</p>
18.20 – 18.50	<p>CAR19 T Cells Carrying Multi-Antigen Targeting Payloads: Tackling Persistence And Tumor Antigen Heterogeneity For The Treatment Of Diverse Tumors</p> <ul style="list-style-type: none"> • IMPACTtm technology enables robust and modular antigen targeting using a single CAR • Multiple antigens are attacked simultaneous, with low pM activity • Technology addresses critical issues of antigen loss and antigen heterogeneity <p>Paul Rennert, President & Chief Executive Officer, Aleta Biotherapeutics</p>	<p><i>Delegates are invited to attend the collocated presentations</i></p>
18.50 – 19.20	<p>Innovative Ophthalmology Endpoints</p> <ul style="list-style-type: none"> • Novel endpoints in retinal trials • Regulatory perspective of these endpoints • Future endpoints in gene therapy trials <p>Aniz Girach, Chief Medical Officer, Nightstar Tx</p>	<p><i>Delegates are invited to attend the collocated presentations</i></p>
19.20 – 19.50	Networking Drinks & End Of Day One	

**4th Annual Cell and Gene Therapy Congress
Day Two – 26th October 2017**

4th Annual Cell & Gene Therapy Congress	
08.30 – 09.30	<p>One Hour Workshop And Breakfast – Cell & Gene Therapy Manufacturing</p> <ul style="list-style-type: none"> Two Presentations from the leading representatives in cell & gene therapy <p>Operational Excellence in Cell & Gene Therapy</p> <p>Frank Thielmann, PMO & Operational Excellence Leader, Novartis</p> <p>Panel Discussion – Cost-Effectiveness in Cell & Gene Therapy Manufacture</p> <ul style="list-style-type: none"> Cost of Goods When scaling up/out automation <p>Moderator: David Sourdive, Executive Vice President, Technical Operations, Cellectis</p> <p>Invited Panellists: Nebojsa Milovic, Vice President of Cell Therapy Pharmaceutical Science, Takeda Ricardo Baptista, Lead Scientist, Process Development, Cell & Gene Therapy Catapult</p>
Cell and Gene Therapy Manufacturing	
09.30 – 10.00	<p>Solution Provider Presentation</p> <p>For Sponsorship Opportunities Please Contact sponsorship@oxfordglobal.co.uk</p>
10.00 – 11.00	Morning Coffee & Refreshments, One to One Meetings x3, Poster Presentation Sessions
11.00 – 11.30	<p>CMC For Cell Therapies – Scale-Up Solutions And Automation In Manufacturing</p> <p>Nebojsa Milovic, Vice President of Cell Therapy Pharmaceutical Science, Takeda</p>
11.30 – 12.00	<p>Scale-Up Manufacture Of Allogenic Cell Therapies Derived From Pluripotent Stem Cells</p> <p>Ricardo Baptista, Lead Scientist, Process Development, Cell & Gene Therapy Catapult</p>
Commercialising Cell & Gene Treatments	
12.00 – 12.30	<p>Innovative Pricing Schemes for ATMP</p> <ul style="list-style-type: none"> The presentation will cover the key considerations from both the payer and the manufacturer perspective in optimizing pricing and access It will concentrate on the specific challenges ATMPs present in the context of reimbursement and strategies to mitigate commercial risk <p>Panos Kefalas, Head of Health Economics & Market Access, Cell & Gene Therapy Catapult</p>
12.00 – 12.30	<p>Solution Provider Presentation</p> <p>For Sponsorship Opportunities Please Contact sponsorship@oxfordglobal.co.uk</p>
12.30 – 13.30	Lunch, Poster Presentation Sessions

4th Annual Cell and Gene Therapy Congress
Day Two – 26th October 2017

4th Annual Cell & Gene Therapy Congress	
13.30 – 14.00	<p>Business Transformation And Partnering Strategies To Accelerate Commercialisation</p> <ul style="list-style-type: none"> Developing a predictable route to market and patient access for CAR T therapies: Understanding Production, Clinical Trial and Value Assessment Interfaces. Using partnerships between developers, regulators and health systems to accelerate CAR T and new immunotherapy commercialization pathways. <p>Magda Papadaki, Head of Manufacturing & Innovation, Association of the British Pharmaceutical Industry</p>
14.00 – 14.30	<p>Lessons in Commercialization of Cell And Gene Therapies</p> <ul style="list-style-type: none"> Lessons in Commercialization of Cell and Gene Therapies Commercialization scorecard The importance of a streamlined patient centric approach Addressing Cost of Goods <p>Peter Olagunju, Vice President, Patient Operations & Lentiglobin Programme Leader, Bluebird Bio</p>
14.30 – 15.00	<p>Panel Discussion: Reimbursement And Pricing Strategies For Cell And Gene Therapies</p> <p>Moderator: Invitation to:</p> <p>Confirmed Panelists: Magda Papadaki, Head of Manufacturing & Innovation, The Association of the British Pharmaceutical Industry Panos Kefalas, Head of Health Economics & Market Access, Cell & Gene Therapy Catapult UK Paolo Morgese, Director of EU Market Access & Member Relations, Alliance for Regenerative Medicine</p> <p>Invitation to: Peter Olagunju, Vice President, Patient Operations & Lentiglobin Programme Leader, Bluebird Bio Adrien Lemoine, Senior Vice President, Business Development & Alliance Management, Orchard Therapeutics</p>
15.00 – 15.30	Afternoon Coffee & Refreshments, Poster Presentation Sessions
15.30 – 16.00	<p>Insights Into CAR T Commercialisation</p> <p>Jennifer Chow, Global Commercial CAR T Lead, Celgene</p>
16.00 – 16.30	<p>Academic And Manufacturing Partnerships In Cell And Gene Therapy</p> <ul style="list-style-type: none"> Creating a discovery and clinical development engine through academic alliances Partnering with CDMOs to access expertise and manufacturing capacity <p>Adrien Lemoine, Senior Vice President, Business Development & Alliance Management, Orchard Therapeutics</p>
16.30 – 17.00	<p>Areas of Strategic Importance for Cell & Gene Therapy: Going Forward</p> <p>Paolo Morgese, Director of Market Access & Member Relations, Alliance for Regenerative Medicine</p>
17.00	End of Conference

4th Annual Cell and Gene Therapy Congress

25—26 October 2018 | Novotel London West Hotel, London, UK
www.oxfordglobal.co.uk/celltherapy-congress

HOW TO REGISTER:

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I would like to attend: (Please tick as appropriate)

Industry Delegates (Biopharma, Pharma or Biotech Companies)

- Congress £899 plus VAT
1 day pass £599 plus VAT
 Day 1
 Day 2

Academic Delegates

- Congress £520 plus VAT
1 day pass £320 plus VAT
 Day 1
 Day 2

Vendor Delegates

(CROs, Consultants, Technology and Service Providers)

- Congress Only £1750 plus VAT
1 day pass £999 plus VAT
 Day 1
 Day 2

- Poster Presentation £250 plus VAT

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- Distribution of your company's promotional literature to all conference attendees £999 plus VAT

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The Delegate Booking Fee includes: lunches and refreshments throughout the Congress event, conference presentations, workshop and panel sessions, scheduled one-to-one meetings and networking/social events, conference and speaker notes. Delegates may attend, free of charge, all sessions arranged by the Organiser. An admin surcharge of £50 will be applied to payments settled following the receipt of an invoice. This charge will not be applied to payments settled online.

Vendor Delegates will not be eligible for one to one meetings unless they purchase a sponsorship meetings package. These can only be purchased directly from Oxford Global Marketing Ltd and not via the online booking facility.

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Those who have booked a poster presentation at the event must provide the poster title, abstract (200 words or less), principal author, organisation, mailing address, email, telephone, fax and additional authors, within a month of registration. All poster spaces will be for A0 (841mm x 1189mm) portrait size.

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Delegates and vendor delegates are subject to the following charges and refunds upon withdrawal or cancellation.

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Documentation

I cannot attend but would like to purchase access to the following:

- Access to the online conference presentations £499 plus VAT

VAT is charged at 20% on the attendance fees for all delegates. VAT is also charged on online and paper copy documentation and promotional literature distribution for all UK customers and for those EU customers not supplying a registration number for their own country here.

How to Pay (choose one of the following payment options)

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