Program

Vaccine Technology VII



Conference Co-Chairs

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(McGill University, Canada)

Charles Lutsch (Sanofi, France)

Tarit Mukhopadhyay

(University College London, United Kingdom)

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Vaccine Technology © Conferences History

An ECI Conference Series

Vaccine Technology I (2006)

Barry C. Buckland, John G. Aunins, Emilio A. Emini, and Jerald C. Sadoff Puerto Vallarta, Mexico

Vaccine Technology II (2008)

Barry C. Buckland, John G. Aunins, Paula Marques Alves, and Kathrin Jansen Albufeira, Algarve, Portugal

Vaccine Technology III (2010)

Barry C. Buckland, John G. Aunins, Paula Marques Alves, and Kathrin Jansen Nuevo Vallarta, Mexico

Vaccine Technology IV (2012)

Barry C. Buckland, John G. Aunins, Paula Marques Alves, and Kathrin Jansen Albufeira, Algarve, Portugal

Vaccine Technology V (2014)

Laura Palomares, Manon Cox, John Aunins and Kathrin Jansen Playa del Carmen, Mexico

Vaccine Technology VI (2016)

Laura Palomares, Tarit Mukhopadhyay, Manon Cox and Nathalie Garçon Albufeira, Portugal

Welcome from the VT VII Chairs

Welcome everyone to Mont-Tremblant, a constantly evolving destination bringing nice surprises within a Natural Park in the Laurentide's region, for the 7th Vaccine Technology (VT-VII) conference! It is enthusing that our meeting is being established as the premiere vaccine technology conference and contributes to informing on the scientific and technological trends in developing vaccines to meet the global needs of public health priorities.

Throughout this time, the VT series has been the main forum where industry, academia, public health authorities and philanthropic organizations meet to facilitate linkages and enable collaborations between all regions of the world.

With over 200 participants from 27 countries on 5 continents, this year's meeting is certainly one of the largest VT-ECI conferences ever, and one of the most diverse to date involving many students, academics, government, international bodies and industry representatives to invest in the future and sustain the growth of the vaccine field as a global priority. We will have the largest number of participants from low and medium income countries ever to attend the meeting and this is made possible by the generous sponsorship from the Bill and Melinda Gates Foundation. In addition, the kind sponsorship of our industry partners enabled the attendance a large number of academics and students.

The conference program includes more than 45 oral presentations, two keynote addresses, and six lead lectures. In addition, we have four thematic workshops. The first workshop titled: "Meet the funders for Global Health" recognize the important theme of funding for sustainability of vaccine development for global interventions. Other workshops aim to highlight technical insights in cell and process engineering, -omics tools and analytics.

As has been the tradition and a key success factor of this conference series, a significant amount of time has been allocated to poster sessions. More than 90 posters are presented, and ten short talks have been selected from these posters to underline the important topics covered. All participants are strongly encouraged to take full advantage of this opportunity to further discuss the excellent work that will be presented during the oral and poster sessions. To encourage discussion and debate, a reception will be held during the evening poster session.

Innovation in antigen design and deployment of advanced vaccine technologies are essential to strengthen the public health vaccination policies and support the preparedness plans against emerging and re-emerging infectious diseases. We find ourselves now in a position with the ability to respond to these pathogens of epidemic potential; the debate today is how best to respond – and will be a test of both the technical means and the political will.

Recently, the true potential of therapeutic vaccines was realized by the patients and health authorities are challenging the field to speed up the development of these new medical

interventions that harness the human immune system. With this in mind, we have put together a program that will present the new directions, recognize the challenges, and build on the successes in the vaccine development arena, as shared by leading academic, institutional and industrial experts.

The success of this conference is fundamentally determined by the active engagement each one of you in the dialogue that is enabled by this conference venue, sharing your thoughts and expertise with other participants as we collectively contribute to the accelerated development of the vaccine sciences and technologies and make them available globally. For those of you returning to the VT conference series – welcome back. For those first-time participants – welcome! This is a unique opportunity to rub shoulders with some of the leaders in the vaccine field – so we warmly encourage you to make a friend!

Finally, we would like to thank all the oral session chairs, workshop program session chairs, poster session chairs, and scientific committee members for their time and efforts. All have worked with remarkable dedication to put together a high-quality program. And, once again, thanks to the generous contributions from our sponsors for enabling an outstanding international attendance.

Special thanks to Barbara Hickernell and her dedicated team at ECI, particularly Kathy Chan, Kevin Korpics, and Tressa D'Ottavio for their tireless help and enormous assistance with the logistics and numerous details. Many thanks to John Aunins, for his friendly help in supporting the conference as ECI liaison for this conference. We also want to thank the many student volunteers, and Fairmont Hotel Mont-Tremblant staff, for helping make this meeting a success.

Once again, welcome to Mont-Tremblant and the Laurentide's region in Quebec. We look forward to meeting each of you personally.

Amine Kamen, Nathalie Garcon, Charles Lutsch and Tarit Mukhopadhyay Chairs, Vaccine Technology VII Conference

Conference Sponsors

Bill & Melinda Gates Foundation

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Thermo Fisher Scientific

Tunnell Consulting

Univercells

Xell AG

Conference program

Friday June 22	Breakfast		Checkout																											
Thursday June 21	Breakfast		Session 7	Therapeutic vaccines	Lead speaker: Dr. John Bell		Coffee break	Session 8	Short talks selected from	posiei absilacis	Poster session 3		Grazing lunch					Coffee break	Session 9	Bioprocessing advances in	vaccine manufacturing (III)	Networking			Closing Keynote	David Robinson	Banquet			
Wednesday June 20	Breakfast		Session 5	Formulation and delivery	Lead speaker: Dr. Jean	Haensler	Coffee break	Session 6	Capacity building and	ווופול פוונסוו אומיו (ד)			Lunch		Workshop 4	Analytical tools		Coffee break	Session 6	Capacity building and	intervention plan (II)	Lead speaker: Dr. Jerome Kim		Dinner			Poster session 2			
Tuesday June 19	Breakfast		Session 4	Bioprocessing advances in vaccine	manufacturing (II)	Lead speaker: Dr. David Kaslow	Coffee break	Session 4 cont.			Workshop 3	Genomics for vaccines		Boxed lunch		External activities		Networking								Dinner on your own				
Monday June 18	Breakfast		Session 1	Technological and clinical	advances in vaccinology (I)	Lead speaker: Dr. Barney	Coffee break	Session 2	Technological and clinical	advances III vaccinology (II)			Lunch		Workshop 2	Manufacturing equipment		Coffee break	Session 3	Bioprocessing advances in	vaccine manufacturing (I)	Lead speaker: Manon Cox		Dinner			Poster session 1			
Sunday June 17															Registration and	refreshments				Workshop 1	Innovation in global	health			Opening keynote	Dr. Alejandro Cravioto	Dinner			
Time	7:30	8:00	8:30	00:6	9:30	10:00	10:30	11:00	11:30		12:00	12:30	13:00	13:30	14:00	14:30	15:00	15:30	16:00	16:30	17:00	17:30	18:00	18:30	19:00	19:30	20:00	20:30	21:00	21:30

Room locations and notes

- General Sessions and Workshops will be held in Mali III-IV.
- Poster Sessions will be in Mali I-II.
- Meals will be in Soutana 1-2. The conference banquet location will be announced on site.
- The ECI office is the Meeting Planner Office.
- Audio, still photo and video recording by any device (e.g., cameras, cell phones, laptops, PDAs, watches) is strictly prohibited during the technical sessions, unless the author and ECI have granted prior permission.
- Speakers Please have your presentation loaded onto the conference computer prior to the session start (preferably the day before).
- Speakers Please leave discussion time as previously directed by your session chair.
- Please do not smoke at any conference functions.
- Turn your mobile telephones to vibrate or off during technical sessions.
- Please write your name on your program so that it can be returned to you if lost or misplaced.
- After the conference, ECI will send an updated participant list to all participants. Please
 check your listing now and if it needs updating, you may correct it at any time by logging
 into your ECI account.
- Emergency Contact Information: Because of privacy concerns, ECI does not collect or maintain emergency contact information for conference participants. If you would like to have this information available in case of emergency, please use the reverse side of your name badge.

Sunday, June 17, 2018

14:00 – 16:00 Conference check-in

16:30 – 18:30 Workshop 1: Meet the Funders in Global Health

Moderators: Vivian Hsu, Bill & Melinda Gates Foundation;

Torey de Rozario, Bill & Melinda Gates Foundation; Tarit Mukhopadhyay, University College London

Objectives:

- To connect scientists from academia and industry to organizations who fund innovations in Global Health, and to provide greater insight on how to gain access to those funding streams
- To hear from Bill & Melinda Gates Foundation grantees on their experience in working with a funding organization and to hear an overview of their funded technology
- Ultimately, we aim to support continued funding of innovative technology to further vaccine development and manufacturing for global needs

Opening

Vivian Hsu, Bill & Melinda Gates Foundation

Meet the funders

- 1. Bill & Melinda Gates Foundation (BMGF), Torey de Rozario
- 2. Global Health Investment Fund (GHIF), Glenn Rockman
- 3. The National Institute for Innovations in Manufacturing (NIIMBL), Chris Roberts
- 4. Coalition for Epidemic Preparedness Innovations (CEPI), Simone Blayer

BMGF Grantees

- 1. UCL- ULTRA Platform Grant, Tarit Mukhopadhyay & Lourdes Velez Suberbie
- 2. University of Kansas Center for Research- MSA, David Volkin
- 3. Vaxess- Microneedles Platform Technology Grant, Michael Schrader

Open discussion

Questions via Sli.do. Go to address www.sli.do and use event code #G330.

Wrap-up

19:00 – 20:00 **Opening Keynote**

The impact of vaccines worldwide and the challenges to achieve universal immunization Dr. Alejandro Cravioto, Chair of WHO Strategic Advisory Group of Experts (SAGE) and Faculty Medicine of the Universidad National Autonoma de Mexico

20:00 – 22:00 Opening Reception Dinner

Monday. June 18. 2018

07:00 – 08:30	Breakfast
08:30 – 10:30	Session 1: Technological and Clinical Avances in Vaccinology (I) Session Chairs: David Weiner, The Wistar Institute, USA; Frank Bähner, CureVac AG, Germany
08:30 – 09:10	Lead talk: Structure-based vaccines for respiratory viruses Dr. Barney Graham, Deputy Director of the Vaccine Research Center of the NIH, USA
09:10 – 09:35	mRNA Vaccines: On the progress from promise to reality Hari Pujar, Moderna, USA
09:35 – 10:00	RNActive®-An mRNA-based vaccine technology for next generation prophylactic vaccines Edith Jasny, Senior Scientist CureVac AG, Tübingen, Germany
10:00 – 10:25	Virus-like particle vaccines against BK and JC polyomaviruses Diana V. Pastrana, NCI/NIH, USA
10:25 – 10:55	Coffee break (Sponsored by GE Healthcare and GSK)
10:55 – 12:35	Session 2: Technological and Clinical Advances in Vaccinology (II) Session Chairs: Udo Reichl, Max Planck Institute, Germany; Hari Pujar, Moderna Therapeutics, USA
10:55 – 11:20	Molecular quality engineering for low cost vaccine production Kerry Routenberg Love, Koch Cancer Institute at MIT, USA
11:20 – 11:45	Single-cell analysis uncovers a novel influenza A virus-derived defective interfering particle for antiviral therapy Sascha Young Kupke, Max Planck Institute for Dynamics of Complex Technical Systems Magdeburg, Germany
11:45 – 12:10	Persistent antibody clonotypes dominate the serum response to influenza following repeated vaccination over multiple years Jiwon Lee, University of Texas at Austin, USA
12:10 – 12:35	Pan-HA antibodies confer protection in mice against influenza Aziza Manceur, National Research Council, Canada
12:35 – 14:00	Lunch
14:00 – 15:30	Workshop 2: Advanced Technologies. Equipment and Instrumentation for Vaccine Manufacturing Moderators: Charles Lutsch, Sanofi-Pasteur, France; Laura Palomares, IBT, UNAM, Mexico
	Short presentations:
	Optimisation of a flocculation step using a scale-down model with 3D-printed impellers and focused beam reflectance measurement (FBRM) particle-size monitoring. Francis DiGennaro, Merck & Co., Inc., USA
	CRISPR-dCAS9 for controlling Baculovirus replication and increasing production of

Virus-Like particles

Mark Bruder, University of Waterloo, Canada

Monday, June 18, 2018 (continued)

EXPISF - A chemically-defined Baculovirus-based expression system for enhanced protein production in SP9 Cells.

Maya Yovcheva, Thermo Fischer Scientific Inc., USA

Fully automated high-throughput process development for the novel purification of Rotavirus Vaccines.

Shaleem I. Jacob, University College London, UK

Influenza virus capture using membrane chromatography: Improving selectivity by matrix design and pseudo-affinity ligand interactions

Stefan Fischer-Frühholtz, Sartorius Stedim Biotech, Germany

A scalable adenovirus production process, from cell culture to purified bulk

Åsa Hagner-McWhirter, GE Healthcare, Sweden

Panel Discussion (all speakers)

15:30 – 16:00	Coffee break (Sponsored by McGill University - Faculty of Engineering)
16:00 – 18:20	Session 3: Bioprocessing Advances in Vaccine Manufacturing (I) (Sponsored by Sartorius Stedim Biotech GmbH) Session Chairs: Linda Lua, Queensland University, Australia; Richard Peluso, Merck and Co., USA
16:00 – 16:40	Lead talk: The story of a successful biotech (ad)venture: The development of Flublok Manon Cox, NextWaveBio, USA
16:40 – 17:05	Accelerating bioprocess development by analysis of all available data: A USP case study Diego Suarez-Zuluaga, Intravacc, Netherlands
17:05 – 17:30	Purifying viruses with a sheet of paper: Single-use steric exclusion chromatography as a capture platform for vaccine candidates Pavel Marichal-Gallardo, Max Planck Institute for Dynamics of Complex Technical Systems, Germany
17:30 – 17:55	Vero SF technology platform: Strategy for rapid and effective vaccine development; flavivirus vaccines case study Nicolas Sève, Sanofi Pasteur, France
17:55 – 18:20	Bioprocess intensification for production of a Peste des petits ruminants virus (PPRV) vaccine Manuel J.T. Carrondo, iBET, Portugal
18:30 – 20:00	Dinner
20:00 – 22:00	Poster session 1 and Social Hour

Tuesday. June 19, 2018

07:00 - 08:30	Breakfast
08:30 – 10:30	Session 4: Bioprocessing Advances in Vaccine Manufacturing (II) (Sponsored by Takeda Vaccines, Inc.) Session Chairs: Francesc Godia, UAB, Spain; Manon Cox, NextWaveBio, USA
08:30 – 09:10	Lead talk: Developing vaccines for low resource settings through product development partnerships (PDPs) Dr. David Kaslow, Vice President, Essential Medicines, and Director, the PATH Center for Vaccine Innovation and Access, USA
09:10 – 09:30	Continuous purification of cell culture-derived influenza A virus particles through pseudo- affinity membrane chromatography A. Raquel Fortuna, Max Planck Institute for Dynamics of Complex Technical Systems, Germany
09:30 – 09:50	Integrated scalable cyto-technology for recombinant protein bioprocessing J. Christopher Love, Koch Institute at MIT, USA
09:50 – 10:10	Low-cost cell-based production platform for seasonal and pandemic influenza vaccines Alan Yung-Chih Hu, NIIDV/NHRI, Taiwan
10:10 – 10:30	Polymer grafted chromatography media for direct capture and high-resolution purification of enveloped virus-like particles Patricia Pereira Aguilar, University of Natural Resources and Life Sciences Vienna, Austria
10:30 – 11:00	Coffee break (Sponsored by Medicago and Sanofi Pasteur)
11:00 – 11:20	Virus-like particles (VLPs) as a platform for the development of yellow fever and Zika virus vaccine candidate Renata Alvim, Federal University of Rio de Janeiro (UFRJ), Brazil
11:20 – 11:40	Manufacturing strategies for sustainable supply of ultra-low cost vaccines for global health Tania Pereira Chilima, University College London, United Kingdom
11:40 – 12:00	A tailor-made purification strategy for oncolytic measles viruses using membrane-based processes Daniel Loewe, University of Applied Sciences Mittelhessen, Germany
12:00	Boxed lunches available for pick up
12:20 – 13:40	Workshop 3: Genomics and Systems Biotechnology in Vaccine Development Moderators: Wei-Shou Hu, University of Minnesota, USA; Gautam Sanyal, Vaccine Analytics, USA
	Short presentations:
	Options and challenges for systems biology driven cell line development in virus

production

Udo Reichl, Max-Planck Institute for Dynamics of Complex Technical Systems, Germany

Acceleration and intensification of influenza pandemic seed stock candidate vaccine production from HA and NA sequence identification

Amine Kamen, McGill University, Canada

Tuesday, June 19, 2018 (continued)

FluChip-8G: Influenza Genotyping Assay for Enhanced Surveillance and Pandemic Preparedness

Kathy Rowlen, InDevR, USA

Molecular quality engineering for low cost vaccine production

Kerry Love, Massachusetts Institute of Technology, USA

High-resolution systems biology modeling of human-virus interactions Brandon Xia, McGill University, Canada

Panel Discussion (all speakers)

External activities

Networking

Dinner on your own

Wednesday, June 20, 2018

07:00 – 08:30	Breakfast
08:30 – 10:30	Session 5: Formulation and Delivering Vaccines Session Chairs: Nathalie Garcon, Bioaster, France; Lakshmi Krishnan, NRC, Canada
08:30 – 09:10	Lead talk: Formulation considerations for the development of adjuvanted vaccines Dr. Jean Haensler, Director, Antigen & Adjuvant Design, Production and Characterization Research Department, Sanofi Pasteur, Lyon, France
09:10 – 09:30	Intradermal administration of synthetic DNA vaccines induce robust cellular and humoral immune responses Jean D. Boyer, Inovio Pharmaceuticals Inc., USA
09:30 – 09:50	Thermostabilization of adenovirus-vectored vaccines, removing the need for continual cold-chain storage Alexander Douglas, Jenner Institute, University of Oxford, United Kingdom
09:50 – 10:10	Safety and biodistribution of sulfated archaeal glycolipid archaeosomes as vaccine adjuvants Mike McCluskie, National Research Council, Canada
10:10 – 10:30	Recombinant hemagglutinin proteins formulated in a novel PELC/CpG adjuvant for H7N9 subunit vaccine development Suh-Chin Wu, Institute of Biotechnology, National Tsing Hua University, Taiwan
10:30 – 11:00	Coffee break (Sponsored by Intravacc)
11:00 – 13:00	Session 6: Capacity Building and Intervention Plan for Emerging and Re-emerging Infectious Diseases (I) Session Chairs: Amadou A. Sall, Institute Pasteur Dakar, Senegal; Erin Sparrow, WHO, Geneva; Simone Blayer, CEPI, UK
11:00 – 11:20	Accelerated process development and stockpile for MERS, LASSA AND NIPAH viral vaccine Simone Blayer, CEPI, UK
11:20 – 11:40	Rapid response pipeline for stabilized subunit vaccines Keith Chappell, University of Queensland, Australia
11:40 – 12:00	Using the AdVac® vaccine manufacturing platform for rapid response to infectious disease outbreaks Guus Erkens, Janssen Vaccines, Leiden, Netherlands
12:00 – 12:20	Preclinical development of filovirus and flavivirus vaccines based on recombinant insect cell expressed subunits Axel T. Lehrer, University of Hawaii, USA
12:20 – 12:40	Development of Pan-filovirus vaccine against Ebola and Marburg virus challenges Xiangguo Qiu, Public Health Agency of Canada, Canada
12:40 – 13:00	Lightitas sanid saananaa platfasm fas anidamia psanasadnaaa
	High-titer rapid response platform for epidemic preparedness José Castillo, Univercells, Belgium

Wednesday, June 20, 2018 (continued)

14:00 – 15:30	Workshop 4: Analytical Tools in Vaccine Development Moderators: Nathalie Garcon, Bioaster, France; Manon Cox, NextWaveBio, USA
	Short presentations:
	Structure-based vaccine design by electron microscopy Bridget Carragher, NanoImaging Services Inc., USA
	Strategies to overcome the age-old problem of immunosenescence Brian Schanen, Sanofi, USA
	VaxArray NA reagent kit Kathy Rowlen, InDevR, USA
	Laser force cytology for rapid quantification of viral infectivity Sean J. Hart, LumaCyte, USA
	Panel Discussion (all speakers)
15:30 – 16:00	Coffee break (Sponsored by LumaCyte)
16:00 – 18:30	Session 6: Capacity Building and Intervention Plan for Emerging and Re-emerging Infectious Diseases (II) Session Chairs: Amadou A. Sall, Institute Pasteur Dakar, Senegal; Erin Sparrow, WHO, Geneva; Simone Blayer, CEPI
16:00 – 16:40	Lead talk: Vaccines for poverty. Associated infection diseases: Accessing innovation Jerome Kim, Director-General International Vaccine Institute, Seoul, South Korea
16:40 – 17:00	Future prospects and application for the development of adenovirus-based vaccine Tao Zhu, Cansino Biologics, China
17:00 – 17:20	Adjuvant manufacturing scale-up and technology transfer Christopher Fox, IDRI, USA
17:20 – 17:40	A vaccine for Ebola virus – approaches and results of accelerated process development and characterization studies Randi Saunders, Merck & Co, USA
17:40 – 18:00	Product development and programmatic implementation of TYPBAR TCV® and ROTAVAC® vaccines V. K. Srinivas, Bharat Biotech International Limited, India
18:30 – 20:00	Dinner
20:00 – 22:00	Poster session 2 and Social Hour

Thursday, June 21, 2018

07:00 – 08:30	Breakfast
08:30 – 10:30	Session 7: Therapeutic Vaccines Session Chairs: Barry Buckland, BioLogicB, USA; Paula Alves, IBET, Portugal
08:30 – 09:10	Lead talk: A novel vaccinia virus backbone for the delivery of immunotherapeutic genes John Bell, Ottawa Hospital Research Institute, Ottawa, Canada
09:10 – 09:30	HER2 cancer vaccine optimization by combining <i>Drosophila</i> S2 insect cell manufacturing with a novel VLP-display technology Thomas Jørgensen, ExpreS2ion Biotechnologies, Denmark
09:30 – 09:50	Rapid design/development and clinical deployment of synthetic DNA vaccine technology for difficult immune targets David Weiner, The Wistar Institute, USA
09:50 – 10:10	Development of an analytical platform for delivery of recombinant oncolytic viruses Gautam Sanyal, Vaccine Analytics LLC, USA
10:10 – 10:30	Leveraging vectored vaccine candidates manufacturing to GMP compatible bioprocesse Cristina Peixoto, iBET, Portugal
10:30 – 11:00	Coffee break (Sponsored by Merck and Co., Inc.)
11:00 – 12:00	Session 8: Short talks selected from poster abstracts (Sponsored by Thermo Fisher Scientific) Session Chairs: Yvonne Thomassen, Intravac, Netherlands; Ernesto Chico, CIM, Cuba; Marc Aucoin, University of Waterloo, Canada; Amine Kamen, McGill University, Canada
	Purification of flavivirus VLPs by a two-step chromatographic process Matheus Souza, Federal University of Rio de Janeiro (UFRJ), Brazil
	Flavivirus production in perfusion processes using the EB66® cell line Alexander Nikolay, Max Planck Institute for Dynamics of Complex Technical Systems, Germany
	Production and characterization of HER2-displaying budded virus-like particles and their potential as cancer vaccines Lisa Nika, University of Natural Resources and Life Sciences, Vienna, Austria
	Virus-like particles: A flexible platform for universal influenza vaccine development Sarah Slack, University College London, United Kingdom
	Rapid virus titration using flow cytometry Steve George, Sanofi Pasteur, Canada

Defining the multiplicity and type of infection for the production of Zaire Ebola virus-like particles in the insect cell baculovirus expression system

Ana Ruth Pastor, Instituto de Biotecnologia-UNAM, Mexico

Combining novel and traditional approaches of vaccine development to overcome the challenges of first-in-human trial for Group A Streptococcus

Elodie Burlet, VaxForm, USA

Development of suspensions adapted Vero cell culture process for production of viruses Chun Fang Shen, National Research Council, Canada

Thursday, June 21, 2018 (continued)

19:00 - 20:00

20:00 - 22:00

Reception

Banquet

HER1 therapeutic cancer vaccine: An active immunotherapy treatment for patients with tumors expressing the receptor of epidermal growth factor (EGF-R) Eduardo Suarez, CIM, Cuba 12:00 - 14:30Poster session 3 Grazing lunch Coffee break (Sponsored by NIIMBL) 14:30 - 15:0015:00 - 16:00Session 9: Bioprocessing Advances in Vaccine Manufacturing (III) Session Chairs: Linda Lua, University of Queensland, Australia; Richard Peluso, Merck and Co., USA Production of bacterial outer membrane vesicles as vaccine platform 15:00 - 15:20Matthias Gerritzen, Intravacc, Wageningen University and Research, Netherlands 15:20 - 15:40Extended gene expression for HIV-1 VLPs scale-up and production enhancement using shRNA and chemical additives Laura Cervera Gracia, Universitat Autònoma de Barcelona, Spain 15:40 - 16:00Development of scalable manufacturing process and GMP-compatible formulation for a novel recombinant schistosomiasis vaccine Damon R. Asher, MilliporeSigma, USA 16:30 - 17:30Closing Keynote Innovations in global health: What has been accomplished, what is on the horizon and where are more investments needed David Robinson, Deputy Director CMC Vaccines Development and Surveillance at the Bill & Melinda Gates Foundation, USA

highly divergent influenza A viruses

Christopher McMillan, University of Queensland, Australia

A pre-fusion, trimeric subunit influenza HA-based vaccine elicits cross-protection between

Friday. June 22, 2018

07:00 – 08:30 Breakfast

08:30 – 10:00 Check out and departures

Poster Presentations

 Stability evaluation of inactivated influenza H7N9 vaccines derived from adhesion and suspension MDCK cells

Alan Yung-Chih Hu, NIIDV/NHRI, Taiwan

2. Rapid, cost-effective and scalable gmp-compliant simian adenovirus-vectored vaccine production for early-phase clinical trials using entirely disposable product-contact components

Alexander Douglas, University of Oxford, United Kingdom

3. Flavivirus production in perfusion processes using the EB66® cell line Alexander Nikolay, Max Planck Institute for Dynamics of Complex Technical Systems, Germany

4. Latest advancements in process intensification to support global demand for affordable vaccines

Alfred Luitjens, Batavia Biosciences, Netherlands

5. Proteomic characterization of influenza H1N1 Gag virus-like particles and extracellular vesicles produced in HEK-293SF

Alina Venereo-Sanchez, McGill University, Canada

- 6. Implementation of a process-scale adenovirus purification with a single-use platform Amélie Boulais, Sartorius Stedim Biotech, France
- 7. Membrane chromatography cassettes for bind and elute applications of viruses and large proteins

Stefan Fischer-Frühholz, Sartorius Stedim Biotech, France

8. Influenza virus capture using membrane chromatography: Improving selectivity by matrix design and pseudo-affinity ligand interactions
Stefan Fischer-Frühholz, Sartorius Stedim Biotech, France

9. Defining the multiplicity and type of infection for the production of Zaire Ebola Virus-like particles in the insect cell baculovirus expression system

Ana Ruth Pastor, Instituto de Biotecnología, UNAM, Mexico

10. Structure-based vaccine design by electron microscopy

Anette Schneemann, NanoImaging Services, USA

- 11. Reducing risks with a serum-free medium for MRC-5 based vaccine production Anna-Barbara Hachmann, Thermo Fisher Scientific, Inc., USA
- 12. Process economical effects of implementation of ready-to-use micro carriers in cell-based virus vaccine production

Nicolas Seve, Anne Marie Beauchard, Sanofi Pasteur, France

- 13. **Bioprocess engineering of insect cells for accelerating vaccines development**Antonio Roldao, iBET, Portugal
- 14. **A scalable adenovirus production process, from cell culture to purified bulk** Åsa Hagner-McWhirter, GE Healthcare, Sweden

15. Vaccine protein stabilization in silica

Aswin Doekhie, University of Bath, United Kingdom

16. Experimental and computational fluid dynamics studies of adherent cells on microcarriers in an ambr® 250 bioreactor

Barney Zoro, Sartorius Stedim Biotech, United Kingdom

17. From bench scale to pilot plant operation: Business models and challenges in the biopharmaceutical industry in Brazil

Beatriz de Castro Fialho, Bio-Manguinhos/FIOCRUZ, Brazil

18. Immunization with Fc-based recombinant Epstein-Barr virus gp350 elicits potent neutralizing humoral immune response in a BALB/c mice model

Bing-chun Zhao, Sun Yat-sen University Cancer Center, China

19. Strategies to overcome the age-old problem of immunosenescence
Brian Schanen, Sanofi Pasteur, USA

20. Development and validation of a proprietary medium formulation for recombinant subunit vaccines by the Baculovirus Expression Vector System (BEVS)

Catherine Cleuziat, Boehringer-Ingelheim, France

21. Development of an animal-component free insect medium for the Baculovirus Expression Vector System (BEVS)

Catherine Nguyen, Irvine Scientific, USA

22. How to limit the use of serum in viral processes: A gibco perspective Céline Martin, Thermo Fisher Scientific, Inc., USA

23. A pre-fusion, trimeric subunit influenza HA-based vaccine elicits cross-protection between highly divergent influenza A viruses

Christopher McMillan, The University of Queensland, Australia

24. Superinfection arising in stable lentiviral vector producer cell lines bearing Cocal-G envelope proteins

Christopher Perry, University College London, United Kingdom

25. Development of suspensions adapted Vero cell culture process for production of viruses

Chun Fang Shen, National Research Council of Canada, Canada

26. Addressing the challenges of influenza virus-like particles purification Cristina Peixoto, iBET, Portugal

27. A genome-wide CRISPR screen to generate high-yield cell lines for pandemic influenza vaccine production

David M. Sharon, McGill University, Canada

28. Process development for a flexible vaccine vector platform based on recombinant life virus

Dieter Palmberger, ACIB - Austrian Center of Industrial Biotechnology, Austria

29. HER1 therapeutic cancer vaccine: An active immunotherapy treatment for patients with tumors expressing the receptor of epidermal growth factor (EGF-R)
Eduardo Suarez, CIM, Cuba

30. Combining novel and traditional approaches of vaccine development to overcome the challenges of first-in-human trial for Group A Streptococcus

Elodie Burlet, VaxForm, USA

31. Accelerated mass production of influenza virus seed stocks in HEK-293 suspension cell cultures by reverse genetics

Ernest Milián, VCN Biosciences, Spain

32. Implementation of a strategy to produce a broadly neutralizing monoclonal antibody against Zika and dengue viruses

Esmeralda Cuevas-Juárez, Instituto de Biotecnología, UNAM, Mexico

- 33. Optimization of a flocculation step using a scale-down model with 3D-printed impellers and Focused Beam Reflectance Measurement (FBRM) particle-size monitoring Francis DiGennaro, Merck & Co., Inc., USA
- 34. Vaccination with viral vectors expressing NP, M1 and chimeric hemagglutinin induces broad protection against influenza virus challenge in mice Guha Asthagiri Arunkumar, Icahn School of Medicine at Mount Sinai, USA
- 35. A new porcine suspension cell line (PBG.PK-21) provides efficient production for influenza and yellow fever vaccine viruses Gwendal Gränicher, Max Planck Institute for Dynamics of Complex Technical Systems, Germany
- 36. **HILIC-LC/MS** method for non-derivatized amino acid analysis in spent media Hari Kosanam, Merck, USA
- 37. Process development of chromatography-based purification on pandemic influenza virus-like particle based vaccines
 Hiachun Lai, NHRI, NTHU, Taiwan
- 38. Δπ=0 reverse osmosis enriches a high osmotic pressure solution from a low-titre fermentation broth to a saturated solution or salt form using RO and NF membranes Ho Nam Chang, Korea Advanced Institute of Science and Technology, South Korea
- 39. **Upstream process intensification using Viral Sensitizer technology** Jean-Simon Diallo, Ottawa Hospital Research Institute, Canada
- 40. Molecular understanding of the serum antibody repertoires after seasonal influenza vaccination among different age cohorts

 Jiwon Jung, The University of Texas at Austin, USA
- 41. Formulation development of a recombinant protein based non-replicating rotavirus (NRRV) vaccine candidate: Antigen-adjuvant-preservative interactions

 John M. Hickey, University of Kansas, USA
- 42. **Disruptive micro-facility for affordable vaccine manufacturing** José Castillo, Univercells, Belgium
- 43. Orbital shaken bioreactor for influenza A virus production in high cell density cultivations
 Juliana Coronel, Max Planck Institute for Dynamics of Complex Technical Systems, Germany
- 44. Vaxarray for hemagglutinin and neuraminidase potency testing of influenza vaccines Kathy Rowlen, InDevR, Inc., USA

- 45. Vaxarray potency assay for rapid assessment of "pandemic" flu vaccines Kathy Rowlen, InDevR, Inc., USA
- 46. Application of analytical characterization tools in process and formulation development of low cost vaccines using the ULTRA manufacturing platform Kawaljit Kaur, University of Kansas, USA
- 47. Comparaison of rabies virus purification using different methods Khaled Trabelsi, Pasteur Institute of Tunisia, Tunisia
- 48. Comparative transcriptome analysis of a Trichoplusia ni cell line reveals distinct host responses to intracellular and secreted protein products expressed by recombinant baculoviruses

Krisztina Koczka, ACIB - Austrian Center of Industrial Biotechnology, Austria

49. Generation and efficacy assessment of a chimeric antigen E2-CD154 as a marker Classical Swine Fever Virus subunit vaccine produced in HEK 293 and CHO K1 mammalian cells

Lidice Méndez-Pérez, Center for Genetic Engineering and Biotechnology, Cuba

50. Production and characterization of HER2-displaying budded virus-like particles and their potential as cancer vaccines

Lisa Nika, University of Natural Resources and Life Sciences, Vienna, Austria

- 51. Rapid fermentation optimization for vaccine development Lourdes Velez, University College London, United Kingdom
- 52. CRISPR-dCAS9 for controlling baculovirus replication and increasing production of virus-like particles

Mark Bruder, University of Waterloo, Canada

- 53. **Purification of flavivirus VLPs by a two-step chromatographic process**Matheus Souza, Federal University of Rio de Janeiro (UFRJ), Brazil
- 54. ExpiSf™: A chemically-defined baculovirus-based expression system for enhanced Protein production in Sf9 cells

Maya Yovcheva, Thermo Fisher Scientific, Inc., USA

- 55. **Media formulation to support the growth of Vero cells in suspension**Megan Logan, University of Waterloo, Canada
- 56. WITHDRAWN
- 57. Accelerating the manufacture of glycoconjugate vaccines for pneumococcal disease Neha Patel, University College London, United Kingdom
- 58. WITHDRAWN
- 59. Development of a vaccine production platform for poultry diseases in Africa: Newcastle Disease Virus non-replicative adenovirus-vectored vaccine
 Omar Farnos, McGill University, Canada
- 60. Formulation and stabilization of a recombinant human Cytomegalovirus vector for use as a candidate vaccine for HIV-1
 Ozan S. Kumru, University of Kansas, USA

61. Formulation development of a stable, orally delivered live human neonatal rotavirus (rv3-bb) vaccine candidate

Prashant Kumar, University of Kansas, USA

62. Case study: Single-use platform for complete process development and scale-up of an Adenovirus

Rachel Legmann, Pall Life Sciences, USA

63. Development and characterization of a murine hepatoma model expressing hepatitis C virus (HCV) non-structural antigens for evaluating HCV vaccines

Kamran Haq, National Research Council Canada, Canada

64. Case studies: Raman implementation for process lifecycle management in fermentation based processes

Roberto Ortiz, Merck &Co., USA

65. Adaptation of Vero cells to suspension culture and rabies virus production on different SERUM free media

Samia Rourou, Institut Pasteur de Tunis, Tunisia

- 66. **Virus-like particles: A flexible platform for universal influenza vaccine development** Sarah Slack, University College London, United Kingdom
- 67. Characteristics of rVSV-ZEBOV production kinetics in HEK293 and Vero cells Sascha Kiesslich, McGill University, Canada
- 68. Laser force cytology for rapid quantification of viral infectivity Sean J. Hart, LumaCyte, USA
- 69. Fully automated high-throughput process development for the novel purification of rotavirus vaccines

Shaleem I. Jacob, University College London, United Kingdom

- 70. **Affinity resin screening for optimal DSP application to rotavirus vaccine production.** Stephen A. Morris, University College London, United Kingdom
- 71. Rapid virus titration using flow cytometry

Steve George, Sanofi Pasteur, Canada

- 72. **Highly efficient influenza virus production: A MDCK-based high-cell-density process**Thomas Bissinger, Max Planck Institute for Dynamics of Complex Technical Systems,
 Germany
- 73. A clinically validated Drosophila S2 based vaccine platform for production of malaria vaccines

Thomas Jørgensen, ExpreS2ion Biotechnologies, Denmark

74. Technology transfer for local vaccine production in Argentina

Valeria Brizzio, Sinergium Biotech, Argentina

75. Real-time stability of a hepatitis E vaccine (Hecolin®) demonstrated with potency assays and multifaceted physicochemical methods

Xiao Zhang, Sun Yat-Sen University Cancer Center, China

76. Evaluating the effect of formulation on the uptake of a ZIKA subunit vaccine candidate by antigen-presenting cells

Xiaoling Li, Envigo, United Kingdom

- 77. Implications of MDCK cell heterogeneity in cell-based influenza vaccine production Xuping Liu, East China University of Science and Technology, China
- 78. An outbreed mouse model of yellow fever for study of pathogenesis and development of vaccines and therapeutics

Yakhya Dieye, Institut Pasteur de Dakar, Senegal

- 79. Development of stabilizing formulations of a trivalent inactivated poliovirus vaccine in a dried state for delivery in the Nanopatch™ microprojection array Ying Wan, University of Kansas, USA
- 80. Efficient influenza vaccine manufacturing: Single MDCK suspension cells in chemically defined medium

Yixiao Wu, East China University of Science and Technology, China

- 81. Analytical characterization and formulation assessment of model secretoryimmunoglobulin-A (slgAs) for their potential use as low cost, orally delivered slgAs Yue (Martin) Hu, University of Kansas, USA
- 82. **Steric exclusion chromatography for the purification of recombinant baculovirus**Daniel Loewe, University of Applied Sciences Mittelhessen, Germany
- 83. **Production and purification of Zika virus for an inactivated virus vaccine candidate**Renato Mancini Astray, Instituto Butantan, Brazil
- 84. Co-formulation of broadly neutralizing antibodies 3BNC117 and PGT121:Analytical challenges during pre-formulation characterization and storage stability studies Vineet Gupta, University of Kansas, USA