World Congress on Rare Diseases

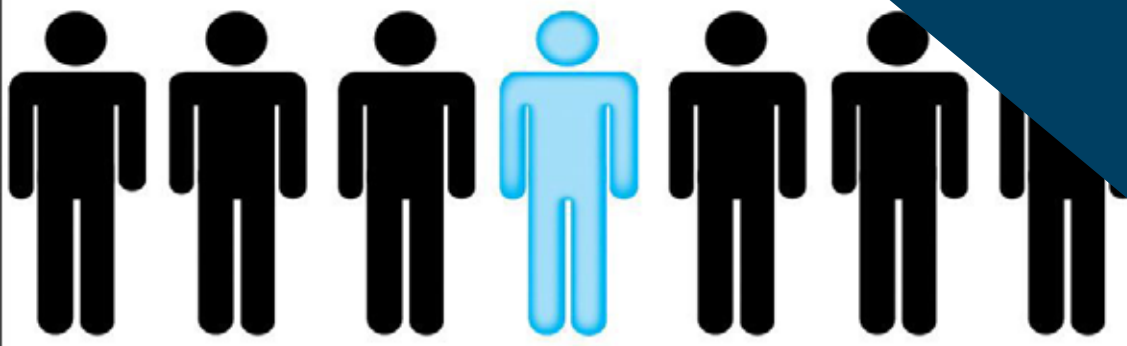
December 05-07, 2018
Dubai, UAE



TENTATIVE PROGRAM

Conferences Series llc LTD
47 Churchfield Road
London W3 6AY, UK

World Rare Diseases 2018

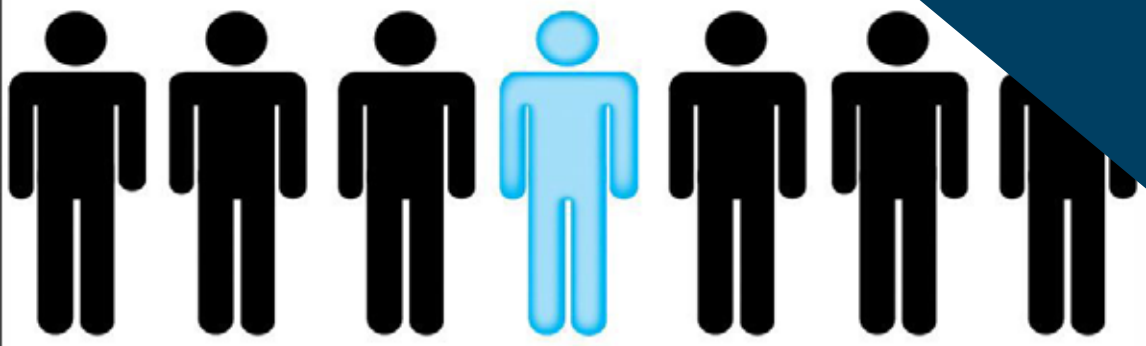


EVENT PROGRAM

December 05, 2018

Opening Ceremony	08:30-09:00
Keynote Sessions	09:00-10:30
Panel Discussion	10:30-10:40
Clinical Research & Treatment on Rare Diseases	11:00-13:00
Lunch Break	13:00-14:00
Mystery & Future of Rare Diseases, Living with a Rare Diseases	14:00-16:00
Workshop	16:00-18:00

Note: Event Program is subjected to change with final allotment of Speaker slots

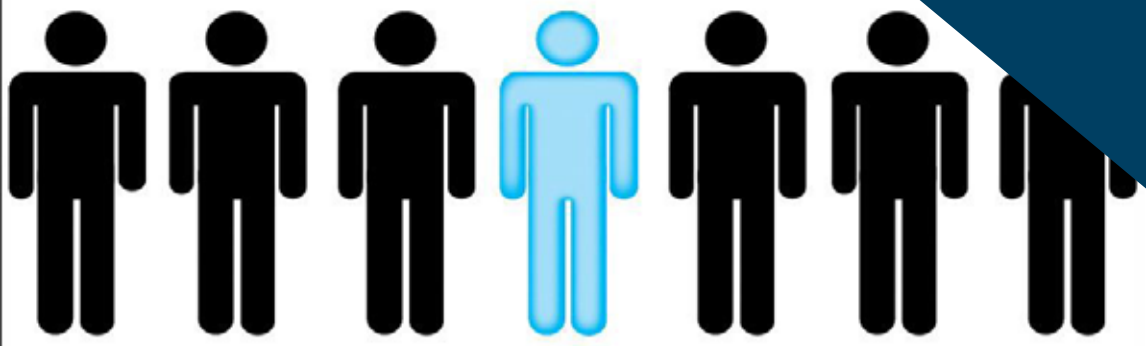


EVENT PROGRAM

December 06, 2018

Keynote Sessions	09:00-10:30
Rare Cancer Diseases, Neurology, Rare Haematology & Immune System Diseases	11:00-13:00
Lunch Break	13:00-14:00
Rare Diseases in Aging, Gastrointestinal Diseases & Pediatric Diseases	14:00-15:30
Poster Presentations	16:00-17:00
Young Research Forum	17:00-18:00

Note: Event Program is subjected to change with final allotment of Speaker slots



EVENT PROGRAM

December 07, 2018

Keynote Sessions	09:00-10:30
Rare Diseases in Endocrine System & Oncology, Oral, Eye and Ear Disease	11:00-13:00
Lunch Break	13:00-14:00
Workshop	14:00-15:00
Special Session	15:00-16:00
Networking	16:00-18:00

Note: Event Program is subjected to change with final allotment of Speaker slots



SCIENTIFIC SESSIONS

Rare Disease

Clinical Research on Rare Diseases

Treatment for Rare Diseases

Living with a Rare Diseases

Mystery Diagnosis of Rare Diseases

Future of Rare Diseases Research

Rare Diseases in Aging

Rare Gastrointestinal Diseases

Rare Paediatric Diseases

Rare Cancer Diseases

Neurology Rare Diseases

Rare Haematology & Immune System Diseases

Rare Diseases in Endocrine System

Rare Oral Diseases

Rare Diseases in Eye and Ear

Rare Oncological Diseases

Orphan Drug

Neglected Tropical Diseases

Organizing Committee Member



Scott Schliebner

Schliebner is a clinical strategist with a 25-year background in clinical drug development focused on rare diseases, orphan drugs, and advanced therapeutics. He develops innovative solutions and strategic considerations to help overcome the operational, logistical, and ethical challenges inherent to conducting research with rare disease patients. Mr. Schliebner's efforts are focused on patient-centered approaches to drug development; reducing the burden of clinical trial participation; and leveraging novel and disruptive technologies to bring new therapies to patients faster. Committed to collaborative efforts to further drug development, Mr. Schliebner has developed active partnerships with rare disease patient-focused organizations, patient advocacy groups, drug development institutes, and academic research centers.

Stef Stienstra



Strategic and creative consultant in biomedical science, with a parallel career in the Dutch Civil-Military Interaction Command in which he has responsibility for the counter measures in CBNRe threats and (medical) consequence management both in a military and a civilian (terrorism) setting. He was the director of the 2014 & 2016 World Congress of CBRNe Science & Consequence Management in Tbilisi, Georgia.



Organizing Committee Member

Nadia Bodkin

She currently holds the positions of Senior Executive Vice President of MaxifyLife, Corporate Engagement Manager of the Jansen's Foundation, Global Gene's Foundation Alliance Council Member and Founding Board Member of Rare is Not Rare. Recently, Nadia has also facilitated and coordinated the development of the Rare Advocacy Movement (RAM), a patient advocacy initiative focused on documenting the ever-evolving complex structure and unique dynamics of the rare disease patient advocacy landscape.

Leyla Camarillo

Leyla Camarillo is a knowledgeable articulate communicator with several years of experience across healthcare and, improving evaluation and training techniques in the country. Maintain strong decision-making and problem-solving abilities, as well as excellent time-management, data research, situation analysis proficiencies and, strengthening the program. Effectively train partners in network based management systems and procedures are her main goal.



SESSION TALKS - DECEMBER 05

Title: Drug Delivery by Tattooing to Treat Cutaneous Leishmaniasis

Stef Stienstra, Royal Dutch Navy, Netherlands

Title: Exploration of the impact of climate change on schistosomiasis transmission dynamics

Tayo Alex Adekiya,University of Zululand, South Africa

Title: Risk of transverse myelitis following dengue infection: A systematic review of the literature

Godwin Oligbu,St. Georges University of London, UK

Title: Diabetes and pregnancy

Dr.Mohamed Zakaria El Helf, Egyptian Ministry of Health, Egypt

Title:Perioperative Management of Paediatric Patients With Genetic Diseases

Yasser Zaghloul,Sheikh Khalifa Medical City, UAE

Title: Increasing importance of brain eating bug in Pakistan; current status and future prospects

Abdul Matin, Majmaah University, Saudi Arabia

Title: In vitro Activity of Colistin in Combination with Doripenem against Carbapenem Non-Susceptible Acinetobacter baumannii

Reham Wasfy, Theodor Bilharz Research Institute, Egypt

Title: The burdens of Tuberous Sclerosis Complex for patients and caregivers

Emmanuelle Logette, Scientific Director -ASTB, Switzerland

Title: The Challenge of Rare Diseases in Mexico

Leyla Soraya Camarillo Blancarte, Mexican Ministry of Health, Mexico

Workshop

Title: Rare Disease PRO Registries – Empowering Patients to Advance research

Virginia Spitzer , Foundation for Sarcoidosis Research, Chicago

Title: Early occurrence and recurrence of hepatocellular carcinoma in HCV-related cirrhosis treated with direct- acting antivirals

Ali A. Ghweil, South Valley University, Egypt

SESSION TALKS - DECEMBER 06

Title: Relax, cool down and scaffold. How to rescue dopamine transporter mutants linked with Infantile/juvenile Parkinsonism-dystonia

Hafiz Muhammad Mazhar Asjad,Forman Christian College University, Pakistan

Title: Reactivation of Herpesvirus in patients with Hepatitis C treated with Direct-Acting Anti-viral agents

Ali A. Ghweil, South Valley University, Egypt

Title: Rare Bleeding Disorders in Pakistan: An Update

Munira Borhany,Pakistan

Title: Phytochemical and Pharmacological Screening of Seeds and Fruits Pulp of Cucurbita moschata Duchesne Cultivated in Egypt

Naglaa Z.H.Eleiwa, Zagazig University, Egypt

Title: Future Heredity of Rare Diseases and Orphan Drugs

Mohsen Sojoudi, Rare Diseases Foundation of Iran, Iran

Title: Lymphedema: Update on diagnosis and management

Samer Ellahham, Regional Chair, Middle East, UAE

Title: The science behind what Occipital Horn Syndrome

David Rose, Rare Revolution Magazine, UK

Title: The Rare Special Powers: Art, Science, and Patient Groups

Eleonora Passeri, Rare Special Powers community, Italy

Title: Not just one disease: Acromegaly. A plethora of ailments and conditions harshly reducing the quality of your life and wellbeing"

Ms. Maserame June Mokoka,Pituitary Community South Africa, South Africa

Workshop

Title: Clinical Studies Networks – De-Risking Trials and Studies

Noopur Singh, Foundation for Sarcoidosis Research, Chicago

Title: BESTI1 mutations in young asymptomatic subjects; OCT findings

Leyla Soraya Camarillo Blancarte, Mexican Ministry of Health, Mexico

SESSION TALKS - DECEMBER 07

Title: Liver stiffness predicts relapse after Direct acting antiviral therapy against chronic Hepatitis C Virus infection

Ali A. Ghweil, South Valley University, Egypt

Title: Evaluation of method for detection of Klebsiella pneumoniae carbapenemase (kpc) among enterobacteriaceae

Hauwa Yakubu,Bauchi state University Gadau ,Nigeria

Title: Effects of Botulinum Toxin A on Histology and Ultrastructure of Submandibular Salivary Glands.

Mohamed Yehia Abdelfattah, Assiut University, Egypt.

Title: Bridging the Information Gap – Why do we need an information Revolution

Rebecca Stewartt, Rare Revolution Magazine & The Rare Youth Project, UK

Title: Rare Diseases And Their Oral Clinical Manifestations In Two Hospital Center In Yaounde

Rose Mbede Nga Mvondo, Cameroon

Title: Universal Health In One Touch

Rose Mbede Nga Mvondo, Cameroon

Title: Fetal MRI in early diagnosis in rare disease of brain

Hamid Reza Edraki, Rare Diseases Foundation of Iran, Iran

Title: Tuberous Sclerosis Complex (TSC), the patient's side

Emmanuelle Logette, Scientific Director -ASTB, Switzerland

Workshop

Title: Molecular Diagnosis of Congenital Cataract Using Next-Generation Sequencing

Dr. Mirena C. Astiazaran,Clinical Geneticist,Mexico

Title: The RAS Pathway in Mexican patients with Oculocutaneous Syndormes: Somatic Mutations"

Dr. Oscar Chacon, Geneticist, Mexico

10 Speaker Slots Available



Venue

Dubai, UAE















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