

# CLINICAL GENOMICS

DUBAI, UNITED ARAB EMIRATES

## Conference Agenda

DAY 1 - Sunday, 19<sup>th</sup> January, 2020

| Time                 | Program  | Presenter                           |
|----------------------|--|-------------------------------------|
| 08:30 - 10:00        | Registration & Badge Collection  |                                     |
| 10:00 - 10:15        | Conference Opening   |                                     |
| <b>10:15 - 13:30</b> | <b>POPULATION GENOMICS AND HUMAN DISEASE</b>   |                                     |
| 10:15 - 10:55        | Populations and Diabetes: Gene-Environment Interaction   | Pierre Zalloua                      |
| 10:55 - 11:10        | Annotation and Visualization of the UAE-Specific Variome   | Andreas Henschel                    |
| 11:10 - 11:25        | Radiogenomic Signature helps to Achieve Personalized Therapy by Predicting Patients with Aggressive Disease    | Havjin Jacob                        |
| 11:25 - 12:05        | Coffee break and Poster Session A - Neurodevelopmental Disorders   |                                     |
| 12:05 - 12:45        | The Burden of Diabetes and Metabolic Syndrome in Africa  | Ayesha Motala                       |
| 12:45 - 13:00        | Whole Exome Sequencing for Genetic Diagnosis of Chronic Kidney Disease: Limitations and Solutions              | Hamad Ali                           |
| 13:00 - 13:15        | An Egyptian Genome Reference   | Inken Wohlers                       |
| 13:15 - 13:30        | Association Mapping of Multivariate Phenotypes in the Presence of Missing Data                                 | Saurabh Ghosh                       |
| 13:30 - 14:30        | Lunch Break  |                                     |
| <b>14:30 - 17:30</b> | <b>METABOLIC DISORDERS</b>   |                                     |
| 14:30 - 15:10        | Neurometabolic Disease Discoveries: Translating Big Data into Better Outcomes                                  | Clara Vankarnebeek                  |
| 15:10 - 15:25        | Prevalence of Fabry Disease Among Haemodialysis Patients in Saudi Arabia                                       | Salwa Alhemyadi                     |
| 15:25 - 15:40        | Integrative Genomic Approaches to Understand Complex Diseases in the UAE                                       | Youssef Idaghdour                   |
| 15:40 - 16:20        | Coffee Break and Poster Session B - Dysmorphology  |                                     |
| 16:20 - 17:00        | Next Generation Sequencing (NGS) in the Era of Inborn Errors of Metabolism (IEM): Opportunities and Challenges | Majid AlFadhel                      |
| 17:00 - 17:15        | Pre-Association Computational Analysis of Non Synonymous SNPs of Human RETN Gene                               | Imane Morjane                       |
| 17:15 - 17:30        | Metabolic Reprogramming as a Promising Approach for Everting Breast Cancer Multi-Drug Resistance               | Dana Zaher                          |
| 17:30 - 17:45        | Coffee Break   |                                     |
| <b>17:45 - 18:45</b> | <b>PUBLIC LECTURE</b>  |                                     |
| 17:45 - 18:00        | European School of Genetic Medicine and Public Awareness of Genetics: the Italian and Omani experiences.       | Giovanni Romeo and Maryam Al-Shehhi |
| 18:00 - 18:45        | To See What Everyone Sees But Think What No One Has Thought  | Rana Dajani                         |



# PAN ARAB HUMAN GENETICS CONFERENCE

19 - 20 JANUARY 2020

## Conference Agenda

### DAY 2 - Monday, 20<sup>th</sup> January, 2020

| Time                 | Program   | Presenter            |
|----------------------|---|----------------------|
| <b>10:00 - 13:00</b> | <b>NEURODEVELOPMENTAL DISORDERS</b>   |                      |
| 10:00 - 10:40        | Current State of Genomic Testing for Neurodevelopmental Disorders   | Ahmed Abou Tayoun    |
| 10:40 - 10:55        | Rapid Whole Genome Sequencing Improves Management and Cost Effectiveness of Critically Ill Patients with Neurological Presentations admitted in Neonatal and Pediatric Intensive Care Units | Shareef Nahas        |
| 10:55 - 11:35        | Coffee Break and Poster Session C - Population Genomics and Human Disease   |                      |
| 11:35 - 11:50        | Whole Exome Sequencing of Suspected Mitochondrial Disorders in Pediatric Egyptian Patients; Cairo-Newcastle Collaboration   | Laila Selim          |
| 11:50 - 12:30        | Genetics of Intellectual Disability: What have we Learnt from New Technologies?   | Alain Verloes        |
| 12:30 - 12:45        | Genomic Testing and Counseling: the Contribution of Next Generation Sequencing to Epilepsy Genetic  | Lamia Alsubaie       |
| 12:45 - 13:00        | Advances in Paediatric Genomics in the UAE; Causative Variant Discovery and Phenotype Expansion in Neurodevelopmental Disorders from the UAE.   | Hanadi Abdelrahman   |
| 13:00 - 14:00        | Lunch and Networking  |                      |
| <b>14:00 - 17:00</b> | <b>DYSMORPHOLOGY</b>  |                      |
| 14:00 - 14:40        | Dysmorphology in the Era of Clinical Genomics   | Anita Rauch          |
| 14:40 - 14:55        | Genomic Overlap Between Neurodevelopmental Disorders and Congenital Heart Defects   | Seyed Ali Shabestari |
| 14:55 - 15:10        | Recurrent Attacks of Rhabdomyolysis in a Patient with Cutis Laxa Syndrome Type IID  | Nihal Al Menabawy    |
| 15:10 - 15:50        | Coffee break and Poster Session D - Metabolic Disorders   |                      |
| 15:50 - 16:30        | Look, Listen and Learn: A Systematic Approach Relevant in an NGS Era  | Sally Ann-Lynch      |
| 16:30 - 16:45        | Case Report of the First Genetically Proven Case from Saudi Arabia with a Novel Mutation in NSD1 Gene   | Jwaher Althopaity    |
| 16:45 - 17:00        | Expanding further the phenotypic expression of Alazami syndrome   | Adila Alkindi        |
| 17:00 - 17:15        | Award presentation and closing remarks  |                      |