SCIENTIFIC PROGRAM

Wednesday **21 Nov**ember 2007

08.00 - 09.30	Registration
Session 1	Leaders in Genomics and Healthcare
09.30 – 10.00	The Human Variome Project and Pilot Projects Prof. Richard Cotton, The Human Variome Project
10.00 – 10.30	Integration of Genomic Sciences and Genomic Medicine Prof. Edison Liu, The Human Genome Organisation
10.30 – 11.00	Newborn Screening in the Region Dr. Danuta Krotoski, National Institutes of Health, USA
11.00 – 11.30	Priorities, Publication and Credit Dr. Myles Axton, Nature Genetics, USA
11.30 – 12.00	Coffee Break
Session 2	Complex and Recessive Disorders in the Arab World
12.00 – 12.30	Genetic Disorders in Arab Populations Dr. Ghazi O. Tadmouri, Centre for Arab Genomic Studies, UAE
12.30 – 13.00	Advances in the Genetics of Type 2 Diabetes Mellitus Prof. Riad Bayoumi, Sultan Qaboos University, Oman
13.00 – 13.30	Delineation of the Clinical and Molecular Basis Underlying Several New Recessive Disorders in the UAE Prof. Lihadh Al-Gazali, UAE University, UAE
13.30 – 14.30	Lunch Break
Session 3	From Phenotypes to Genotypes
14.30 – 14.50	Mutation in WNT10A is Associated with an Autosomal Recessive Ectodermal Dysplasia: The Odonto-Onycho- Dermal Dysplasia Prof. Andre Megarbane, Saint Joseph University, Lebanon
14.50 – 15.10	Anhydrotic Ectodermal Dysplasia in Omani Families Dr. Anna Rajab, Ministry of Health, Oman
15.10 – 15.30	A Novel Approach for the Study of the Genetics of Hypertension in Arabs: Heritability and Linkage Analysis Results of "Oman Family Study" Prof. Mohammad O. Hassan, Sultan Qaboos University, Oman



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15.30 – 15.50

A Whole-Genome Scan in a Large Family with Leukodystrophy and Oligodontia Reveals Linkage to 10q22

Dr. Eliane Chouery, Saint Joseph University, Lebanon

15.50 – 16.10

Coffee Break

16.10 - 16.30

Microarray as an Efficient Modern Molecular Tool in the Detection of Submicroscopic Genetic Abnormalities in Multiple Congenital Anomalies: Diagnosis and Prevention

Dr. Sabitha Murthy, Al Wasl Hospital, United Arab Emirates

16.30 – 16.50

Dominant Effect of Smoking on Airway Epithelium Gene Expression Profiles Among Individuals of European, African, Southeast Asian and Arabian Ancestries

Prof. Lotfi Chouchane, Weill Cornell Medical College, Qatar

16.50 – 17.10

Independent Introduction of Two Lactase Persistence Alleles into Human Populations Reflect Different History of Adaptation to Milk Culture

Dr. Nabil S. Enattah, University of Helsinki, Finland

^{*} The 2nd Pan Arab Human Genetics Conference is accredited by the Faculty of Medicine and Health Sciences, UAE University, as the equivalent of 11 CME hours:



21 November 2007

- Session 1: 2 CME hrsSession 2: 1.30 CME hrs
- Session 3: 2.20 CME hrs

22 November 2007

- Session 4: 2.10 CME hrs
- Session 5: 1.40 CME hrs
- Session 6: 1.15 CME hrs



SCIENTIFIC PROGRAM

Thursday **22 Nov**ember 2007

Session 4	Bioethics and Human Genetics
09.30 – 10.00	International Bioethics and Human Genetics. The activities of UNESCO Prof. Henk ten Have, UNESCO
10.00 – 10.20	Ethics of Mutation Databases: Correctness in Reporting Genetic Variation and its Effects Prof. Richard Cotton, The Human Variome Project
10.20 – 10.40	Genetic Counseling in the Muslim World: The Challenges Dr. Aida Al Aqeel, King Faisal Specialist Hospital and Research Centre, Saudi Arabia
10.40 – 11.00	Ethical Concerns to the use of Pre-implantation Genetic Diagnosis in the Gulf Cooperative Council States Dr. Hamza A. Eskandarani, King Faisal University, Saudi Arabia
11.00 – 11.20	What is the Impact of Genetic Counseling and Prenatal Diagnosis in Genetic Diseases Prevention in an Arab Muslim Population? Prof. Habiba Bouhamed Chaabouni, Charles Nicolle Hospital, Tunisia
11.20 – 11.40	Prevention of Genetic Diseases: Understanding Families and Communities hold the Key for Success Dr. Anna Rajab, Ministry of Health, Oman
11.40 – 12.00	Coffee Break
Session 5	National Startegies for the Prevention of Genetic Disorders in the Region
12.00 – 12.20	Bahrain National Hereditary Diseases Strategy (1984-2007) Dr. Shaikha Al-Arrayed, Salmaniya Medical Complex, Bahrain
12.20 – 12.40	National Strategy for the Prevention of Genetic and Congenital Disorders in Jordan Prof. Hanan Hamamy, National Center for Diabetes, Endocrinology and Genetics, Jordan
12.40 – 13.00	Newborn Screening for Inborn Errors of Metabolism-Combining Molecular and Biochemical Testing: A Saudi Perspective Dr. Moeen Al-Sayed, King Faisal Specialist Hospital and Research Center, Saudi Arabia
13.00 – 13.20	Newborn Screening in Lebanon: 12 Years Experience Dr. Issam Khneisser, Saint Joseph University, Lebanon



Thursday **22** November 2007

13.20 – 13.40	Genotype-Phenotype Correlation in Gaucher Disease: Strategy for Prevention and Therapy in Egypt Prof. Rabah Shawky, Ain Shams University, Egypt

13.40 - 14.45 Lunch Break

Session 6 **Fast Track Reports**

14.45 – 15.00 **ER Retention and Degradation of Mutated Proteins is** a Common Mechanism in Numerous Loss-of-Function **Recessive Diseases**

Dr. Bassam R. Ali, UAE University, United Arab Emirates

15.00 – 15.15 Succesful Preimplantation Diagnosis in a Family Affected with Zellweger Syndrome

Dr. Moeen Al-Sayed, King Faisal Specialist Hospital and

Research Center, Saudi Arabia

15.15 – 15.30 **Evaluation of Cell Free Fetal DNA in Maternal Plasma of**

Early and Late Onset Pre-Eclampsia Patients

Dr. Maha A. El Bassuoni, Menoufiyia University, Egypt

15.30 – 15.45 Molecular Analysis of F8 in Lebanese Hemophilia A Patients: Report on 23 Novel Mutations and Phenotype-

Genotype Correlation

Dr. Claudia D. Khayat, Hotel Dieu de France Hospital,

Lebanon

15.45 – 16.00 Most Encountered Genetic Disorders in Egypt: **Classification and Registry**

Prof. Mona El-Ruby, National Research Centre, Egypt

16.00 – 16.30 Coffee Break

Closure

16.30 – 17.00 **Announcing the Dubai Declaration**

^{*} Certificates of Attendance and Accreditation will be distributed towards the end of Session 6 to attendees who submit the completed conference questionnaires.

