

## CONFERENCE PROGRAM

<b>Day</b>	<b>1:</b>	<b>Genomics</b>	<b>and</b>	<b>Healthcare</b>
Saturday	13.3.2010			
08.00		–		09.30
<i>Registration</i>				
09.30		–		10.20
<i>Opening Ceremony</i>				
10.20		–		10.30
<i>Exhibition Inauguration</i>				
10.30		–		11.00
<i>Coffee Break</i>				
<b>Session 1:</b>				<b>1:</b>
Governance Challenges of Genomic Applications				
	11.00	–		11.30
<b>Governance Challenges of Genomic Applications in Healthcare Systems</b>				
	<i>Mohammad</i>			<i>Afzal</i>
	<i>World Health Organization, Cairo, Egypt</i>			
	11.30	–		12.00
<b>Regulation of Genomic Research in Abu Dhabi</b>				
	<i>Wafa</i>	<i>A</i>		<i>El-Adhami</i>
	<i>Health Authority – Abu Dhabi, United Arab Emirates</i>			
12.00		–		13.30
<i>Lunch Break</i>				
<b>Session 2:</b>				<b>2:</b>
Genetic Disorders: Analysis				
	13.30	–		13.50
<b>Consanguinity and Reproductive Health among Arabs</b>				
	<i>Ghazi</i>	<i>O</i>		<i>Tadmouri</i>
	<i>Centre for Arab Genomic Studies, Dubai, United Arab Emirates</i>			
	13.50	–		14.10
<b>Population Genomics of Hearing Loss in the Palestinian Population: A Model for Genetic Heterogeneity</b>				
	<i>Moien</i>			<i>Kanaan</i>
	<i>Bethlehem University, Palestine</i>			

▪ 14.10 – 14.30  
**The EFR Project: A Collaborative Network to Establish an Arabian Biobank Resource to Identify Disease Genes of Indigenous Populations**  
**Habiba Al Safar**  
*Dubai Police General Head Quarters, Dubai, United Arab Emirates*

▪ 14.30 – 14.50  
**Cellular Organelle Disease Genomics: The ER-Associated Protein Degradation (ERAD) is a Major Mechanism Underlying Numerous Human Genetic Disorders**  
**Bassam R Ali**  
*UAE University, Al Ain, United Arab Emirates*

▪ 14.50 – 15.10  
**Genetic Disorders in Syria**  
**Rami A Jarjour**  
*Atomic Energy Commission of Syria, Damascus, Syria*

▪ 15.10 – 15.30  
**Linkage Mapping and Mutation Screening in Two Families with Autosomal Recessive Non-Syndromic Mental Retardation**  
**Suzanne Giesebrecht**  
*Institute of Human Genetics, Bonn, Germany*

15.30 – 16.00  
*Coffee Break*  
**Session 3:**  
Emerging Technologies and Integration into Healthcare

▪ 16.00 – 16.20  
**Novel Insight into Genomic Medicine: From Diagnosis to Tailored Therapy**  
**Fahd Al-Mulla**  
*Kuwait University, Kuwait*

▪ 16.20 – 16.40  
**The Application of Proteomics for Kidney Disease Diagnostics; Global Differential Protein Expression Profiling of Serum in FSGS Families**  
**Chaker Adra**

*Stem Cell Therapy Program, King Faisal Specialist Hospital and Research Centre, Riyadh, Saudi Arabia*

- 16.40 – 17.00  
**Mobile Elements Create Structural Variation: Analysis of a Complete Human Genome**  
*Abdel Halim AF Salem Deifalla*  
*Arabian Gulf University, Manama, Bahrain*

- 17.00 – 17.20  
**A Novel Immune System Mediator**  
*Safa MH Taha*  
*HH Princess Al-Jawhara Centre for Molecular Medicine, Genetics and Inherited Diseases, Manama, Bahrain*

**Day 2: Open Clinic Session / Role of Media in Genetic Literacy Sunday 14.3.2010**

**Session 4:**

Newborn Screening and Genetic Disease Control Programs

- 09.00 – 09.20  
**United Arab Emirates National Newborn Screening Program: An Evaluation (1995 – 2009)**  
*Mohamed Salah El-Den Riad*  
*Ministry of Health, Abu Dhabi, United Arab Emirates*

- 09.20 – 09.40  
**The Bahrain Program to Control Genetic Blood Diseases: 1984-2010**  
*Shaikha Al-Arrayed*  
*Salmaniya Medical Complex, Manama, Bahrain*

**Session 5:**

Open Clinic Session

- 09.40 – 10.00  
**Modern Approaches in Genetic Diagnostics in Kuwait**  
*Sadika Al-Awadi*  
*Kuwait Medical Genetics Center, Kuwait*

▪ 10.00 – 10.20  
**Childhood Short Stature: Approach to Evaluation and Diagnosis in a Genetic Counseling Clinic**  
*Hanan* *Hamamy*  
*Geneva University Hospital, Switzerland*

▪ 10.20 – 10.40  
**Mild Phenotype of Mucopolidosis Type III**  
*Fatma* *Al-Jasmi*  
*United Arab Emirates University, United Arab Emirates*

▪ 10.40 – 11.00  
**An Autosomal Recessive Syndrome of Severe Mental Retardation, Dysmorphic Facies and Skeletal Abnormalities Maps to the Long Arm of Chromosome 17**  
*Mohammed* *AbdulAziz* *AlOwain*  
*King Faisal Specialist Hospital and Research Centre, Riyadh, Saudi Arabia*

11.00 – 11.30  
*Coffee Break*

▪ 11.30 – 11.50  
**A Novel Homozygous Missense Mutation of the Leptin Gene (N103K) in an Obese Egyptian Patient**  
*Mona* *El-Gammal*  
*National Research Centre, Cairo, Egypt*

▪ 11.50 – 12.10  
**Baraitser Syndrome: A New Case**  
*Faouzi* *Maazoul*  
*Charles Nicolle Hospital, Tunis, Tunisia*

▪ 12.10 – 12.30  
**Delineation of *de novo* Copy-Number Variations in a Number of Cases with Genetic Disorders**  
*Jamil* *Al-Alami*  
*Shaffallah Medical Genetics Center, Doha, Qatar*

12.30	–	14.00
<i>Lunch Break</i>		
<b>Session</b>		<b>6:</b>
Role of Media in Genetic Literacy		
14.00	–	16.30
<i>Roundtable discussion</i>		
16.30 – 17.00		
<i>Closing Ceremony</i>		