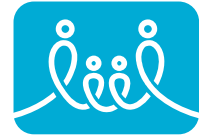




CONGRESS AGENDA

Day 1 | Saturday | September 19
Le Meridien Dubai Hotel



TIME	TOPICS	SPEAKER
08:00-08:50	Registration التسجيل	
08:20-08:30	Welcome & Introduction الترحيب والمقدمة	
Session 1: Premarital Genetic Screening (Great Ballroom 1) الفحص الجيني ما قبل الزواج Chair: Dr. Fatima Abdulaziz & Dr. Mariam Alshehhi		
08:30-08:50	Inborn error of metabolism in Adults	Dr. Ali Al-Beshri
08:55-09:40	جلسة حوارية مشتركة عن الفحص الجيني ما قبل الزواج في دولة الامارات العربية المتحدة Premarital Genetic Screening in the UAE	د. علي البشري - استشاري علم الوراثة السريرية وعلم الجينوم - مستشفى توام د. سعاد العور - وزارة الصحة ووقاية المجتمع سعادة القاضي د. علي الذيابي - قاضي تمييز أول - محاكم دبي Dr. Ali Al-Beshri - Consultant Clinical Genetics & Genomics- Tawam Hospital Dr. Suad Alawar- MOHAP - His Excellency Judge / Ali Elthabahi - Senior Judge in Cassation - Dubai Court
	Q & A / Panel Discussion أسئلة وأجوبة	
09:40-10:20	Official Inauguration الافتتاح الرسمي للمؤتمر وتكريم العائلات والبراعة	
10:20-10:35	Coffee Break استراحة	
MAIN HALL (Great Ballroom 1) Session 2: Innovation in Rare Disease Chair: Dr. Amal Al Tenaiji & Dr. Noor Almeheri		
10:35 - 10:55	Porphyria, insights on new treatment	Dr. Nadia Alhashmi, Oman
10:55 - 11:15	Novel Diagnostic / Laboratory Methods	Dr. Osama Aldirbashi, Qatar
11:15 - 11:35	Novel Therapeutic Approach to Metabolic Genetic Disorders	Dr. Amal Mohammed Alhashem, UAE
11:35 - 12:05	Symposium 1 Sanofi Beyond the Spleen: The Neurological Face of Gaucher Disease	Dr. Amal Alteneijji, UAE
12:05 - 12:15	Q & A / Panel Discussion قلاصلا ةطارتسا	
12:15 - 13:15	Lunch Break & Prayer	
MAIN HALL (Great Ballroom 1) Session 3: Metabolism and Biochemical Genetics Chair: Dr. Ame! Aljasmi Almaeenei & Dr. Noura Aldhaheri		
13:15 - 13:35	New and rare Inherited bone marrow failure syndromes	Dr. Leslie Lehmann, USA
13:35 - 13:55	UNICAS: Sant Joan de Deu Hospital's Innovative Strategy to Address Rare Diseases	Dr. Encarna Guillen, Spain
13:55 - 14:15	Insights on Lysosomal Storage Disease	Prof. Fatma Aljasmi, UAE
14:15 - 14:35	Updates in MPS II Management	Dr. Hind Alsharhan, Kuwait
14:35 - 14:45	Q & A / Panel Discussion ةبوجأو ةللسأ	
14:45 - 15:00	Coffee Break ةطارتسا	
15:00 - 15:20	Symposium 2 GenPharm - FAODs: When Fat Can't Fuel	Dr. Amal Altenaiji, UAE



CONGRESS AGENDA

Day 1 | Saturday | September 19
Le Meridien Dubai Hotel



TIME	TOPICS	SPEAKER
MAIN HALL (Great Ballroom 1) Session 4: Genomic Insights and Emerging Precision Therapies Chair: Dr. Faizah Hashimy & Dr. Hafima Aldhanhani		
15:20 - 15:40	NFIX missense variants that disrupt the β -hairpin loop result in a severe form of Malan syndrome in adolescence with rapidly evolving scoliosis and muscle wasting	Dr. Benjamin Cocanougher, USA
15:40 - 16:00	Advanced Therapies Program of Sant Joan de Deu hospital	Dr. Fatima Nuñez, Spain
16:00 - 16:20	Autism, Insight about Genetics	Dr. Maryam Alshehhi, UAE
16:20 - 16:40	Decoding PACS1 Syndrome: From Gene Discovery to Precision Therapy	Dr. Gary Thomas, USA
16:40 - 16:50	Q & A / Panel Discussion الأسئلة والإجابات	
16:50 - 17:00	Sponsors Awarding and Closing & End of Day	





CONGRESS AGENDA

Day 2 | Sunday | September 20
Le Meridien Dubai Hotel



TIME	TOPICS	SPEAKER
08:00-08:50	Registration	
08:50 - 09:00	Welcome & Introduction	
Session 5: Metabolic Emergencies (MAIN HALL-Great Ballroom 1) Session Chair: Dr. Mahra Alfattan & Dr. Ali Al-Beshri		
08:30 - 08:50	Beyond the hospital: Taking rare disease care home	Dr. Hamda Abulhoul, UAE
08:50 - 09:10	Insights on Urea Cycle Defects	Dr. Noura Aldhaheeri, UAE
09:10 - 09:30	Genetics Role in Intensive Care Management	Dr. Fatma Ali Rabea, UAE
09:30 - 09:50	Symposium 3 Biomarin - Taiba Access Rare Symposium Achondroplasia: From Pathophysiology to Targeted Therapy	Dr. Sareea Al Remeithi, UAE
09:50 - 10:00	Q & A / Panel Discussion	
10:00 - 10:15	Coffee Break	
10:15 - 10:35	Symposium 4 - Immedica Unlocking the Mystery: A Rare Disorder as a Treatable Cause of Spastic Diplegia	Dr. Amal Altenajji, UAE
Session 6: Treatable Metabolic Disorders (MAIN HALL-Great Ballroom 1) Chair: Dr. Dalia Alraei Said & Dr. Heba Elabd		
10:35 - 10:55	Vitamin Responsive Disorders	Dr. Tawfeg Ben Omran, Qatar
10:55 - 11:15	CTX, Rare Cause of Adult Neurological Disease	Dr. Arif Khan, UAE
11:15 - 11:35	The New Landscape for Precision Medicine	Dr. Theodore W. Laetsch, USA
11:35 - 11:55	From Genetics to Hemodynamics: Hereditary Pediatric Pulmonary Hypertension from Genetic Perspective	Dr. Ahmed Adel Hassan, UAE
11:55 - 12:15	Symposium 5 - Biologix Recognize Early, Act Early: Transforming Alpha-Mannosidosis Care	Dr. Fatma Al Jasmi, UAE
12:15 - 12:30	Q & A / Panel Discussion	
12:30 - 13:30	Lunch Break & Prayer	
13:30 - 14:00	Symposium 6 - AstraZeneca HPP Patient Experience Moderator: Dr. Maryam Al Shehhi, UAE	