



GARROD ASSOCIATION ANNUAL SYMPOSIUM - 2010



Agenda

Location: Sheraton Hotel Newfoundland
St. John's, NL

Friday, June 25, 2010

Time	Speaker	Topic
08:00 - 09:00	<i>Breakfast / Registration</i>	
09:00 - 09:05	Dr. Pranesh Chakraborty	Welcome and opening remarks
09:05 - 09:50	Keynote Speaker Dr. Ingrid Tein	Overview of Muscle Metabolism and New Clinical Developments in Metabolic Myopathies.
09:50 -10:35	Keynote Speaker Dr. Mark Tarnopolsky	Exercise is the only effective therapy for mitochondrial disease and aging.
10:35 - 11:00	<i>Coffee Break and Poster viewing</i>	
11:00 -11:45	Keynote Speaker Dr. Jean Michaud	Pathology of Metabolic Myopathies.
11:45 -12:00	Questions for the invited speakers	
12:00-1:00	<i>Lunch</i>	
1:15 - 2:00	Keynote Speaker Dr. Brian Robinson	The Laboratory Diagnosis of Mitochondrial Myopathies
2:00 – 3:00	ORAL PRESENTARION SESSION 1 – Chair, Dr. Lesley Turner	
2:00 - 2:20	Dr. Chitra Prasad	Spectrum of Pompe Disease in South Western Ontario: Role of Enzyme Replacement Therapy.
2:20-2:40	Dr. Sarah Dyack	L-Tyrosine reverses significant symptoms of NemaLine Rod Myopathy in Adult with Childhood onset form: Our 5 year experience.
2:40-3:00	Dr. Anne-Marie Lamhonwah	CFTR Expression in Human Muscle: Implications for Exercise Intolerance in Cystic Fibrosis.
3:00 - 3:30	<i>Coffee Break and Poster viewing</i>	
3:30 - 5:00	ORAL PRESENTATION SESSION 2 – Chair, Dr. John Mitchell	
3:30- 3:50	Dr. Lesley Turner	Case Presentation of Neutral Lipid Storage Disease with Myopathy
3:50 – 4:10	Dr. Lorne Seargeant	Fabry Disease Identified after Renal Biopsy for Proteinuria
4:10 – 4:30	Dr. Andre Mattman	Emphysema in a 40 year old male with late infantile galactosialidosis
4:30 – 4:50	Dr. Gregory Pastores	Safety and efficiency of velaglucerase alfa in patients with type 1 Gaucher disease who were treatment naïve or previously treated with imiglucerase
6:30-	<i>Dinner</i>	Johnson Geo Centre



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Saturday, June 26, 2010

Time	Speaker	Topic
08:00 - 09:00	<i>Breakfast / Registration</i>	
	ORAL PRESENTATION SESSION 3 – Chair, Dr. Murray Potter	
9:00 – 9:20	Dr. Pranesh Chakraborty	The first three years of screening for medium chain acyl-CoA dehydrogenase deficiency (MCADD) in Ontario.
9:20 – 9:40	Dr. Annette Feigenbaum	Severe Neonatal Presentation of Arginase 1 deficiency.
9:40 – 10:00	Dr. Pierre Allard	Argininosuccinic acid lyase deficiency nearly missed by amino acid chromatography.
10:00- 10:20	<i>Coffee Break</i>	
10:20 -11:00	Garrod Council Meeting	
11:15 -12:00	Annual General Meeting	
12:00-	Adjournment	
6:30-	<i>PUB CRAWL</i>	<i>Meet in the lobby of the Sheraton</i>

Note:

- A booklet of presentation abstracts will be distributed at the meeting.