GARROD SYMPOSIUM 2019
NEUROMETABOLISM

MAY 9 - 11, 2019
SHERATON CENTRE
TORONTO, ON

INTERPROFESSIONAL.UBC.CA/INITIATIVES/GARRODSYMPHOSIUM2019
LEARNING OBJECTIVES & PROGRAM

By the end of the meeting participants will be able to:

• Understand different types of inherited metabolic disorders presenting with neurological features
• Expand their knowledge for treatable neurometabolic conditions and their treatments
• Learn specific neuroimaging patterns of neurometabolic disorders
• Describe organelle specific neurometabolic disorders involving mitochondria, lysosomes and peroxisomes
• Learn the neurodevelopmental outcomes and neuropsychiatric manifestations of neurometabolic disorders

Please note: 25% of the scientific program will be interactive.

LOCATION & TRAVEL

The meeting will be held at the Sheraton Centre Toronto (123 Queen Street, Toronto, ON). A special rate of $239.00 for a standard guest room (Traditional Room: single occupancy) is available for conference delegates until April 9, 2019 or until availability. To avoid disappointment, book by calling: 1.888.627.7171, please specify that you are booking under the Garrod Symposium 2019 room block to receive these reduced rates.

REGISTRATION

ONLINE: The quickest, most secure method is registering online. Registration is available for Visa/MasterCard holders at the meeting organizer’s website: www.interprofessional.ubc.ca/initiatives/garrodsymposium2019

BY FAX: Fax your completed registration form to: 1-604-822-4835


MAIL: Send the registration form with cheque to this address:

IN 9580 REGISTRATION, Interprofessional Continuing Education, The University of British Columbia, Room 105-2194 Health Sciences Mall, Vancouver, BC, V6T 1Z3, Canada

Participants paying by credit card outside of North America: Please inform your credit card company of the transaction as some banks put a block on credit card payments made outside your country.

TUITION FEES & MEETING PROGRAM

Please see the registration form (last page) for details. The tuition fee includes: welcome reception (May 9), breakfast (May 10 & 11), lunch (May 10 & 11), webinars and workshops (May 9-11), Garrod Association membership rate (if selected) and a certificate of attendance. Registration for the dinner event on Friday will also be available at an extra cost. Pre-registration before/on March 29, 2019 is strongly recommended to take advantage of the early-bird rate. To qualify for the Garrod Association member’s rate, one must pay the fee for membership dues during the registration process.

We will be including presenter slides (if the presenter permits) in our meeting program. We will send this program to you electronically (for free) a few days prior to the meeting but if you wish to purchase the paper copy that will be distributed on-site, please select the $20 option on the registration page.

CANCELLATION & REFUND POLICY

Notifications of cancellations must be sent by email to registration.ipce@ubc.ca on or before March 29, 2019 in order for registration fees to be refunded. If you have been invoiced but not paid the fees, you are still liable to pay the cancellation charges below:

• Up to and including March 29, 2019: Registration fee will be refunded less 20% cancellation fee.
• After March 29, 2019: No refunds on registration fees will be made. Substitutions can be made.

CREDITS

This event has been approved by the Canadian Paediatric Society for a maximum of 10.75 credit hours as an Accredited Group Learning Activity (Section 1) as defined by the Maintenance of Certification program of The Royal College of Physicians and Surgeons of Canada. The specific opinions and content of this event are not necessarily those of the CPS, and are the responsibility of the organizer(s) alone.

This conference has been approved by the Canadian Association of Genetic Counsellors (CAGC) for 13.0 CECs.

CONTACT

For additional enquiries, please contact Jo at 604-822-0054 or jo.ipce@ubc.ca.
The Garrod Association and the Program Committee gratefully acknowledge the generous contributions and financial support from the following organizations:

**PLATINUM**

[Logos of the platinum sponsors]

**GOLD**

[Logos of the gold sponsors]

**BRONZE**

[Logos of the bronze sponsors]
12:00-14:00  CIMDRN Meeting (Invitation Only)  
(Room: Cedar)

13:00-16:00  Nurse/Genetic Counselor Webinar  
(Room: Maple West)

**The Hostage Syndrome: Our Relationship with Patients**

*Joe Clarke, MD, PhD, FRCPC, FCCMG, Professor Emeritus, University of Toronto, Toronto, ON*

At the end of the presentation, the listener will be able to:

1. Recount at least 4 characteristics of the “hostage syndrome” as it relates to the relationship between a health care professional and patient with a rare, inherited metabolic disease.
2. Describe how to communicate empathically with the parents of a child with a newly diagnosed inherited metabolic disease.
3. Describe what is meant by the importance of “curiosity” when managing inherited metabolic diseases.
4. Describe 3 potential obstacles to the establishment of rapport with the parents of a child with an inherited metabolic disease.

Following Dr. Clarke's presentation, there will be a closed session (~120mins) for case presentations/discussion.

**Growing up on Sapropterin: Experiences with young children on cofactor therapy**

*Leslie Martell, MS, RD, LDN, Metabolic Dietitian with Boston Children’s Hospital, Boston, MA, USA*

1. Describe use of Sapropterin in young children with Phenylketonuria
2. Illustrate considerations for diet liberalization in the younger population
3. Discuss strategies and approaches to working with families and young children as they grow up on Sapropterin

**“Stop, Go, Proceed with Caution”: Classification of fruit and vegetables according to protein content, using a traffic light analogy (a resource for patients with phenylketonuria)**

*Laura Nagy MSc, RD, Registered Dietitian with Metabolic Genetics and the PKU Program at the Hospital for Sick Children, Toronto, ON and Valerie Austin, BSc*

1. Understand the rationale for the development of a user-friendly resource, to support counting/controlling intake of protein from fruit and vegetables
2. Become familiar with the evidence that allows the protein from many fruits, and some vegetables, to not be counted/controlled within the prescription for natural protein
3. Understand the principles behind the development of the resource
4. Understand how the resource can be used to manage the diet for the treatment of PKU

**Transition to Adult PKU Clinic: Preliminary Findings**

*Heather Bell, RD and Sandra Tavares MSW, RSW, Registered Dietitian and Social Worker with the Fred A. Litwin Family Centre in Genetic Medicine, University Health Network/Toronto General Hospital, Toronto, ON*

1. To increase awareness of patient-identified issues relating to transition from pediatric to adult based care
**PROGRAM THURSDAY**

**May 9, 2019**

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<tr>
<th>Time</th>
<th>Event</th>
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<tr>
<td>15:00</td>
<td>Registration for Garrod Symposium Opens (Room: Willow Foyer)</td>
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<tr>
<td>16:00</td>
<td>Introduction and Welcome to the Meeting (Room: Willow)</td>
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**Saadet Andrews, MD, PhD, FCCMG, FRCPC, Biochemical Geneticist, The Hospital for Sick Children; Associate Professor, Division of Clinical and Metabolic Genetics, Department of Pediatrics, University of Toronto, Toronto, ON**

**Neal Sondheimer, MD, PhD, Associate Professor of Paediatrics and Molecular Genetics, The University of Toronto, Toronto, ON**

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<th>Time</th>
<th>Event</th>
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<tr>
<td>16:15-18:00</td>
<td>Plenary Session I: Complex Neurometabolic Disorders: phenotypes to neuroimaging (Room: Willow) Session Chair: Saadet Andrews</td>
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**Jean-Marie Saudubray, MD Senior metabolic consultant Centre de Référence Neurométabolique Adulte, Groupe Hospitalier Pitié-Salpêtrière, Paris, France**

1. Summarize the complex lipid synthesis and remodeling pathways in cell membranes and roles in the nervous system
2. List inborn errors of metabolism that impair complex lipid metabolism in the nervous system
3. Recognize the prominent motor manifestations of these disorders: Spastic paraparesis, Movement disorders, Neuropathy, Myopathy.
4. Design a tentative diagnostic algorithm

**Discussion**

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| 17:15  | Neuroimaging in Neurometabolic Disorders  
*Susan Blaser, MD, FRCPC, Pediatric Neuroradiologist, The Hospital for Sick Children; Professor of Neuroradiology, University of Toronto, Toronto, ON*  
1. Summarize van der Knaap and Valk’s concept of Selective Vulnerability  
2. Recognize that the vulnerability of white matter and grey matter evolves with brain maturation  
3. Utilize the concepts of Selective Vulnerability and MRI pattern recognition to assess imaging studies of infants and children with possible inborn errors of metabolism |

**Discussion**

<table>
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| 18:00-19:30 | WELCOME RECEPTION & EXHIBITS  
(Room: Birchwood Ballroom & Foyer) |
| 19:30 - 21:00 | Garrod Executive Committee Meeting  
(Invitation Only) (Room: Maple West) |
07:00-8:00 | Registration Open & Breakfast (Provided)
(Room: Willow)

08:00-9:15 | Plenary Session II:
Treatable Neurometabolic Disorders
(Room: Willow)
Session Chair: Neal Sondheimer

08:00 | Creatine Deficiency Disorders: Present and
Future
Andreas Schulze, MD, PhD, FRCPC, Professor Paediatrics
and Biochemistry, University of Toronto; Head, Metabolic
Genetics, Medical Director, Newborn Screening Program,
Senior Associate Scientist, Research Institute, The Hospital
for Sick Children, Toronto, ON

1. Understand the function and metabolism of creatine
2. Recognize the clinical spectrum of creatine deficiency
   disorders
3. Utilize clinical observation, pathophysiological
   thinking and translational research
4. Summarize the present and future of interventions for
   patients with creatine deficiencies

Discussion

08:35 | Pyridoxine-dependent epilepsies: phenotypes
and treatment outcomes
Saadet Andrews, Saadet Andrews, MD, PhD, FCCMG,
FRCPC, Biochemical Geneticist, The Hospital for Sick
Children; Associate Professor, Division of Clinical and
Metabolic Genetics, Department of Pediatrics, University of
Toronto, ON

1. Describe clinical features of pyridoxine-dependent
   epilepsies
2. List current treatments of pyridoxine-dependent
   epilepsies
3. Discuss treatment outcomes of pyridoxine-
   dependent epilepsies

Discussion

09:45 | Plenary Session III:
Organelles and Metabolic Disease
(Room: Willow)
Session Chair: Andreas Schulze

09:45 | Mitochondrial SUMOylation in Cancer and
Metabolism
Heidi McBride, PhD, Montreal Neurological Institute, McGill
University, Montreal, QC

1. Learn how mitochondrial signaling platforms regulate
   glucose metabolism in liver
2. Explore a new mouse model of hepatocellular
   carcinoma
3. Obtain a broad view of the integration between
   mitochondrial dynamics/signaling and cell physiology

10:45 | Evaluating the role of Autophagy in Peroxisome
Biogenesis Disorders
Peter Kim, PhD, Senior Scientist, Cell Biology, Hospital
for Sick Children; Associate Professor, Department of
Biochemistry, University of Toronto, Toronto, ON

1. Explain the cellular mechanisms associated with
   peroxisome biogenesis disorder (PBD)
2. Describe the how autophagy leads to peroxisome loss
   some PBD
3. Discuss the use of autophagy inhibitors as potential
   treatment for PBD

11:20 | Zellweger Spectrum Disorder: Update and
Challenges for Therapy
Nancy Braverman, MD, MS, FACMG, Associate Professor,
Dept. of Human Genetics and Pediatrics, McGill University
Dept. of Pediatrics, McGill University Health Center
(MUHC), Scientist, Peroxisome Disease Program, Child
Health and Human Development, Research Institute of the
MUHC, Montreal, QC

1. Prepare a diagnostic model for the different
   peroxisome disorders
2. Describe new contributions to peroxisome biology
3. Discuss approaches and challenges to therapies

Discussion

12:00-12:45 | Lunch (provided)
(Room: Willow)

12:45-13:15 | Dessert & Exhibits (provided)
(Birchwood Ballroom)
13:15-14:30

Plenary Session IV:
Neurodevelopmental Outcomes and Neuropsychiatric Presentations
(Room: Willow)
Session Chair: Saadet Andrews

13:15
Neurodevelopmental Outcomes of Mucopolysaccharidosis Type I (MPS I)
Michal Inbar-Feigenberg, MD, FCCMG, The Hospital for Sick Children, Toronto, ON

1. Describe neurodevelopmental outcomes of patients with MPS type 1 post HSCT
2. Summarize the unique neurodevelopmental features of this population
3. Compare our findings with the ones described in the literature.

Discussion

13:50
Neuropsychiatric manifestations of neurometabolic disorders
Chantal Morel, MD, FRCP, FCCMG, Clinical and Metabolic Geneticist, Fred A. Litwin Family Centre in Genetic Medicine, University Health Network and Mount Sinai Hospital; Assistant Professor, Department of Medicine, University of Toronto, Toronto, ON

1. Recognize potential CNS manifestations of inborn errors of metabolism
2. Learn when to consider an inborn error of metabolism in a patient with neuropsychiatric symptoms
3. Be able to name groups of inborn errors of metabolism where neuropsychiatric presentations can occur, and what tests to arrange to assess for these

Discussion

14:30-15:00
Refreshment Break & Exhibitors (Birchwood Ballroom) & Posters (Pine)

15:00-16:30
Plenary Session V:
Selected Abstracts
(Room: Willow)
Session Chair: Neal Sondheimer

Platform Presentations (Selected Abstracts)
Six abstracts (15 minutes each)
**PROGRAM SATURDAY**

**08:00-09:00**
Breakfast (Provided)
(Room: Willow)

**09:00-11:00**
Plenary Session VI: Mitochondrial Neurometabolics
(Room: Willow)
Session Chair: Lauren MacNeil

**09:00**
**Exercise and Nutraceutical Approached for the Treatment of Mitochondrial Cytopathies and Diseases Associated with Mitochondrial Dysfunction**
Mark Tarnopolsky, MD, PhD, FRCP(C), Professor of Pediatrics and Medicine, CEO and CSO, Exerkine Corporation, Director of Neuromuscular and Neurometabolic Clinic, McMaster University Medical Center, Hamilton, ON

1. Summarize the pathophysiological consequences of mitochondrial dysfunction.
2. Understand the theory behind mitochondrial therapies.
3. Describe the mechanism of action of mitochondrial therapies.
4. Review the mitochondrial cocktail studies.
5. Review the exercise intervention studies.
6. Understand the basis of novel therapies for mitochondrial disease.

**09:50**
**Laboratory Diagnosis of Mitochondrial Diseases**
Tony Rupar, PhD, FCCMG, Professor in the Depts. of Pathology & Laboratory Medicine, Pediatrics and Biochemistry at Western University and the London Health Sciences Centre in London Ontario. Head, Biochemical Genetics Laboratory

1. Recognize the roles of biochemical, genetic and pathological testing
2. Describe the significance of heteroplasmy in diagnosing mitochondrial diseases
3. Plan a diagnostic strategy to diagnose mitochondrial disease

**10:25**
**A small-molecule approach to heteroplasmy shifting**
Neal Sondheimer, MD, PhD, Associate Professor of Pediatrics and Molecular Genetics, The University of Toronto, Toronto, ON

1. Analyze the impact of heteroplasmy on the phenotype of mitochondrial diseases caused by mtDNA mutation
2. Critique various approaches to heteroplasmy shifting
3. Assess the impact of G-quadruplex formation on DNA replication

**Discussion**

**11:00-11:30**
Refreshment Break & Exhibitors (Birchwood Ballroom) & Posters (Pine)

**11:30-13:00**
Garrod Association Membership Meeting (Room: Willow)

*Please see www.garrod.ca for other meetings after the official program adjourns.*
GARROD SYMPOSIUM
2019 COMMITTEES

Garrod 2019 Symposium Presidents

Saadet Andrews, MD, PhD, FCCMG, FRCPC, Biochemical Geneticist, The Hospital for Sick Children; Associate Professor, Division of Clinical and Metabolic Genetics, Department of Pediatrics, University of Toronto, Toronto, ON

Neal Sondheimer, MD, PhD, Associate Professor of Paediatrics and Molecular Genetics, The University of Toronto, Toronto, ON

Garrod 2019 Symposium Scientific Committee

Saadet Andrews, MD, PhD, FCCMG, FRCPC, Biochemical Geneticist, The Hospital for Sick Children; Associate Professor, Division of Clinical and Metabolic Genetics, Department of Pediatrics, University of Toronto, Toronto, ON

Michal Inbar-Feigenberg, MD, FCCM, Assistant Professor Division of Clinical and Metabolic Genetics, Department of Pediatrics, University of Toronto, The Hospital for Sick Children, Toronto, ON

Jonathan Kronick, MD, PhD, FRCPC, Professor Division of Clinical and Metabolic Genetics, Department of Pediatrics, University of Toronto, The Hospital for Sick Children, Toronto, ON

Lauren MacNeil, PhD, FCCMG, Metabolic Disease Laboratory, Department of Pediatric Laboratory Medicine, University of Toronto, The Hospital for Sick Children, Toronto, ON

Andreas Schulze, MD, PhD, FRCPC, Professor Paediatrics and Biochemistry, University of Toronto, Toronto, ON

Neal Sondheimer, MD, PhD, Associate Professor of Paediatrics and Molecular Genetics, The University of Toronto, Toronto, ON

Garrod 2019 Symposium Organizer

Jo Nam, Senior Education Manager, Interprofessional Continuing Education, University of British Columbia, Vancouver, BC

Garrod Association Executive Committee

Pranesh Chakraborty, President of Garrod Association, MD, FRCPC, FCCMG, Associate Professor, Pediatrics and Biochemistry/Microbiology/Immunology, University of Ottawa; Medical Director, Newborn Screening Ontario; Metabolic Physician, Department of Pediatrics, Children’s Hospital of Eastern Ontario, Ottawa, ON

Andreas Schulze, Vice President of Garrod Association, MD, PhD, FRCPC, Professor Paediatrics and Biochemistry, University of Toronto, Toronto, ON

Jagdeep S Walia, Secretary/Treasurer of Garrod Association, MBBS, FRCPC, FCCMG, Clinical Geneticist and Associate Professor, Head, Division of Medical Genetics (Department of Pediatrics), Director of Research (Department of Pediatrics), Kingston Health Sciences Centre and Queen’s University, Kingston, ON
AFFILIATION / PROFESSION
(please select only one)
☐ Physician  ☐ Registered Nurse  ☐ Dietitian  ☐ Genetic Counsellor  ☐ Lab
☐ Other: __________________________________________

DELEGATE INFORMATION
Please write in block letters and use one registration form per person. Please photocopy the form if needed.
☐ Dr.  ☐ Mr.  ☐ Ms.

Last Name ___________________________ First Name ___________________________ Initials __________

Institution / Organization
________________________________________________________________________

Mailing Address
________________________________________________________________________

City ___________________________ Prov/State ___________________________ Postal Code __________

Daytime Telephone Number / Local ___________________________ Email Address for Meeting Correspondence ___________________________

(If applicable) Please inform us of any dietary requirements for the meeting:
________________________________________________________________________

REGISTRATION RATES
(Rates include all applicable taxes)

Non-Members
☐ $450 On or before March 29, 2019
☐ $430 After March 29, 2019 or Onsite

Members**
☐ $330 On or before March 29, 2019
☐ $310 After March 29, 2019 or Onsite

(This rate includes the $30 Garrod Association membership dues)

Student/ Trainee
☐ $100 On or before March 29, 2019
☐ $150 After March 29, 2019 or Onsite

Allied Health Professional
☐ $100 On or before March 29, 2019
☐ $150 After March 29, 2019 or Onsite

SOCIAL EVENTS
Additional* Guest for Welcome Reception (Thursday)
*The welcome reception is included for Garrod registrants.

Gala Dinner at Hockey Hall of Fame (Friday)

Additional guest(s)

Regular Garrod Attendee
☐ $80 On or before March 29, 2019
☐ $80 After March 29, 2019 or Onsite

Additional guest(s)
☐ $120 On or before March 29, 2019
☐ $120 After March 29, 2019 or Onsite

(Multiple rates include all applicable taxes)

METHOD OF PAYMENT

Payment by Credit Card Most secure and easy method
1. Complete the full registration online at www.interprofessional.ubc.ca with your Visa or MasterCard.
2. Fax the registration form to +1-604-822-4835 and indicate that you would like to pay with Visa or MasterCard. We will send you the secure online link to enter your credit card information.
*Please do not fax your credit card information

Payment by Cheque
Please make your cheque payable to the University of British Columbia and send it along with this form to: Interprofessional Continuing Education, The University of British Columbia, Room 105–2194 Health Sciences Mall, Vancouver, BC V6T 1Z3.

Alternative Payment Methods
Mail or fax complete registration form along with one of the following:
1. Signed purchase order (PO)
2. Letter of Authorization (LOA) from the manager on the organization’s letterhead stating that they will be paying the registration fees. Must include the amount of registration fees, name and contact information of the manager
3. Signed cheque requisition form (ChReq)

Please indicate below how you would like to pay:
☐ Cheque: Payment is enclosed with mailed registration form
☐ Credit Card: Please email me a secure link to enter my credit card number
☐ PO/LOA/ChReq: Purchase order/letter of authorization/cheque requisition form is enclosed with faxed/mailed registration form