

PLENARY SPEAKERS

LOCATION ➤ All Plenary Sessions will be held in the Plenary Hall, located in the **Grand Ballroom Centre, Lower Concourse Level.**

➤ WEDNESDAY JULY 6, 2016

PLENARY 1.0

GENETICS

➤ **CHAIR: James Dowling, CA**

08:00 – 09:00 PL 1.1 GENOMIC APPROACHES TO DIAGNOSIS OF RARE MUSCLE DISEASE

➤ **Keynote Speaker:**
Daniel MacArthur, US

09:00 – 09:30 PL 1.2 GENE DISCOVERY IN CHARCOT-MARIE-TOOTH NEUROPATHIES

➤ **Stephan Züchner, US**

09:30 – 10:00 PL 1.3 RNA SEQUENCE AND RNA ANALYSIS

➤ **James Dowling, CA**

➤ THURSDAY JULY 7, 2016

PLENARY 2.0

HOT TOPICS

➤ **CHAIRS: Vera Bril, CA & Carlos Navarrete Maldonado, CL**

08:00 – 08:15 PL 2.1 STEM CELL THERAPY IN ALS

➤ **Eva Feldman, US**

08:15 – 08:30 PL 2.2 RESULTS OF THE THYMECTOMY TRIAL IN MYASTHENIA GRAVIS

➤ **Gil Wolfe, US**

08:30 – 08:45 PL 2.3 REGAIN: A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED MULTI-CENTER PHASE 3 STUDY OF THE SAFETY AND EFFICACY OF ECULIZUMAB IN SUBJECTS WITH REFRACTORY GENERALIZED MYASTHENIA GRAVIS

➤ **James F. Howard, Jr., US**

08:45 – 09:00 PL 2.4 APPROACH TO PATIENT-CENTERED OUTCOMES RESEARCH

➤ **Richard Barohn, US**

09:00 – 09:15 PL 2.5 DO WE STILL NEED MUSCLE BIOPSY IN THE ERA OF ULTRASOUND?

➤ **Carsten Bonnemann, US**

09:15 – 09:30 PL 2.6 THERAPEUTIC APPROACHES TO INCLUSION BODY MYOSITIS

➤ **Mazen Dimachkie, US**

09:30 – 09:45 PL 2.7 TREATMENT OF AMYLOID NEUROPATHY

➤ **David Adams, France**

09:45 – 10:00 PANEL DISCUSSION

► **FRIDAY JULY 8, 2016**

► **SATURDAY JULY 9, 2016**

PLENARY 3.0 MUSCULAR DYSTROPHY

08:00 – 08:30 PL 3.1 GENE THERAPY FOR MUSCULAR DYSTROPHY

► **Keynote Speaker:**
Dongsheng Duan, US

08:30 – 09:00 PL 3.1 RNA THERAPEUTICS FOR DUCHENNE MUSCULAR DYSTROPHY

► **Keynote Speaker:**
Dana Martin, US

09:00 – 09:30 PL 3.2 ANTISENSE THERAPY FOR MYOTONIC DYSTROPHY

► **Charles Thornton, US**

09:30 – 10:00 PL 3.3 CRISPR BASED GENE EDITING FOR MUSCULAR DYSTROPHY

► **Ronald Cohn, CA**

PLENARY 4.0 MOTOR NEURON DISEASE

► **CHAIR: John Kissel, US**

08:00 – 09:00 PL 4.1 ALS THERAPY DEVELOPMENT: CHALLENGES AND OPPORTUNITIES

► **Keynote Speaker:**
Michael Benatar, US

09:00 – 09:30 PL 4.2 BIOLOGY OF C9ORF72 DISEASE

► **Leonard Petrucelli, US**

09:30 – 10:00 PL 4.3 ANTISENSE THERAPY FOR SPINAL MUSCULAR ATROPHY

► **John Kissel, US**

PROGRAM AT A GLANCE

TUESDAY
JULY 5, 2016

WEDNESDAY
JULY 6, 2016

REGISTRATION HOURS

TUESDAY, JULY 5

▶ 07:00–20:00

WEDNESDAY, JULY 6

▶ 07:00–17:00

THURSDAY, JULY 7

▶ 07:00–17:00

FRIDAY, JULY 8

▶ 07:00–19:00

SATURDAY, JULY 9

▶ 07:00–17:00

EXHIBIT HOURS

WEDNESDAY, JULY 6

▶ 10:00–16:00

THURSDAY, JULY 7

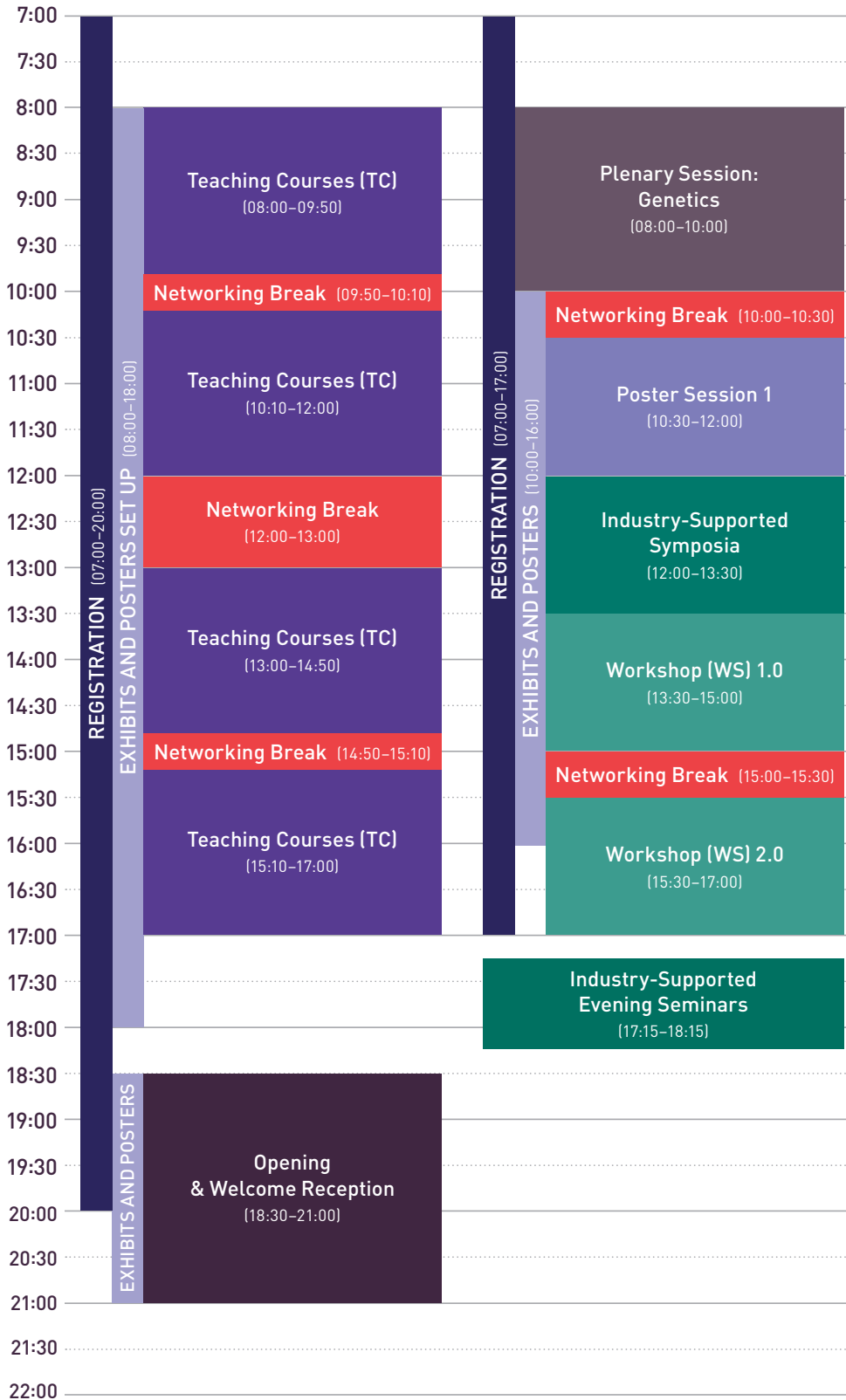
▶ 10:00–16:00

FRIDAY, JULY 8

▶ 10:00–16:00

SATURDAY, JULY 9

▶ 10:00–15:30

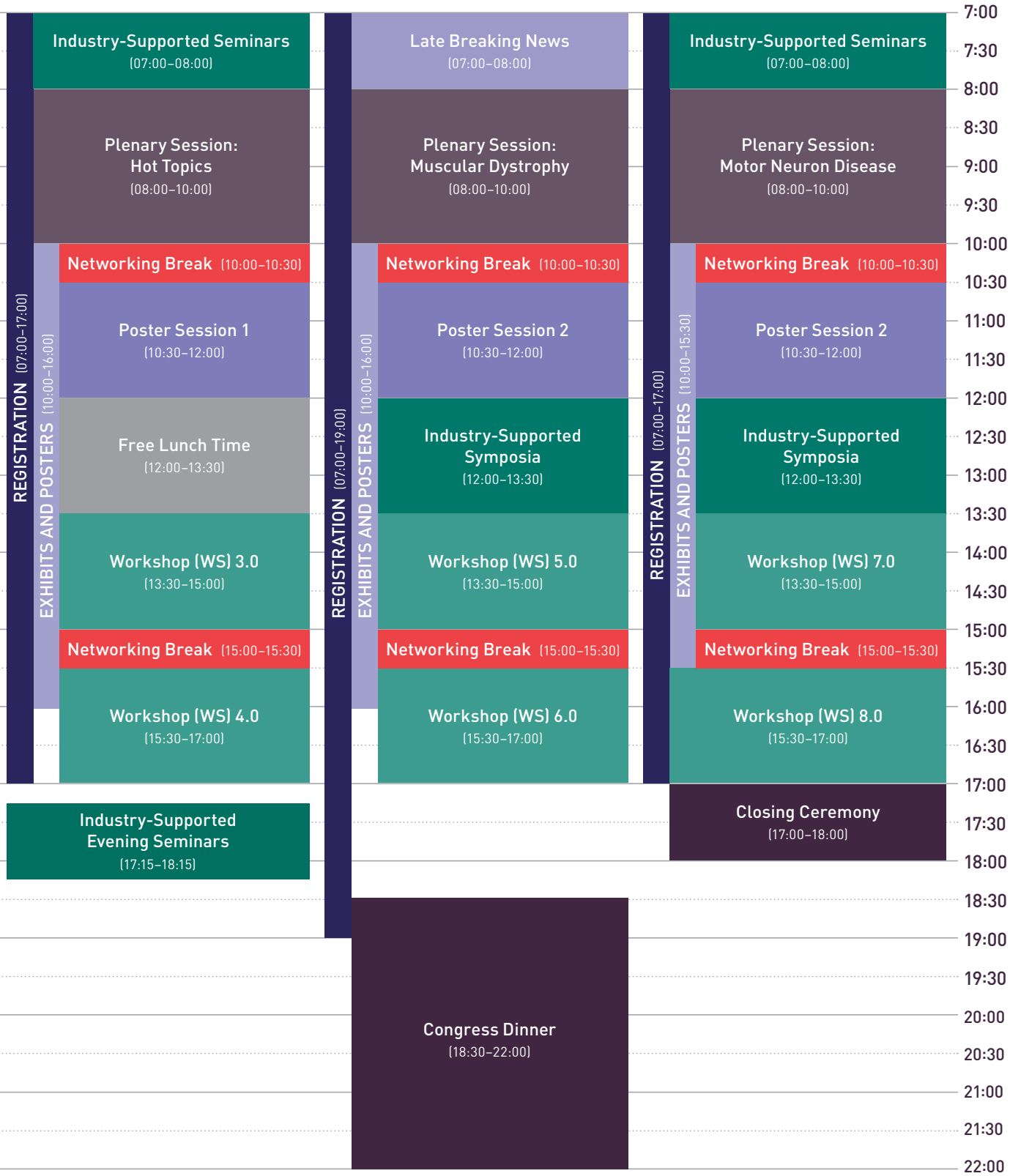


Program listings are subject to change.

THURSDAY
JULY 7, 2016

FRIDAY
JULY 8, 2016

SATURDAY
JULY 9, 2016



PRE-CONGRESS TEACHING COURSES ▶ TUESDAY JULY 5, 2016

LOCATION▶ All Teaching Course Sessions are located on the **2nd Floor of the Sheraton Centre Toronto Hotel.**

07:00–20:00 REGISTRATION OPEN

08:00–18:00 EXHIBITS AND POSTERS SET UP
LOCATION▶ Exhibit Hall, Lower Concourse

08:00–09:50 TC 1.0 - Paediatric Muscular Dystrophy

LOCATION▶ City Hall

Chairs: Kevin Flanigan, US
& Volker Straub, UK

08:00–08:55 1.1 CONGENITAL MYOPATHIES
James J. Dowling, CA

08:55–09:50 1.2 DYSTROPHINOPATHIES
Kevin Flanigan, US

09:50–10:10 NETWORKING BREAK

10:10–11:05 1.3 LIMB-GIRDLE DYSTROPHIES
Volker Straub, UK

11:05–12:00 1.4 AXIAL MYOPATHIES
Anthony A. Amato, US

08:00–09:50 TC 2.0 - Myasthenia Gravis

LOCATION▶ Dominion North

Chairs: Carolina Barnett-Tapia, CA
& Gil I. Wolfe, US

08:00–08:55 2.1 CLINICAL ASSESSMENT OF MYASTHENIA GRAVIS
Carolina Barnett-Tapia, CA

08:55–09:50 2.2 ANTIBODY TESTING IN MYASTHENIA GRAVIS
Luis Querol, ES

09:50–10:10 NETWORKING BREAK

10:10–11:05 2.3 ELECTROPHYSIOLOGICAL TESTING IN MYASTHENIA GRAVIS
Hans D. Katzberg, CA

11:05–12:00 2.4 TREATMENT OF MYASTHENIA GRAVIS
Gil I. Wolfe, US

08:00–09:50 TC 3.0 - Genetics

LOCATION▶ Churchill

Chair: Mary Reilly, UK

08:00–08:55 3.1 THE ABC'S OF GENETICS
Ronald D. Cohn, CA

08:55–09:50 3.2 EVALUATION OF VARIANTS OF UNKNOWN SIGNIFICANCE
Mary Reilly, UK

09:50–10:10 NETWORKING BREAK

10:10–11:05 3.3 THE ETHICAL IMPLEMENTATION OF GENOMIC MEDICINE
M. Stephen Meyn, US

11:05–12:00 3.4 GENETICS COUNSELLING
Jeanna McCuaig, CA

08:00–09:50 TC 4.0 - Non-immune Mediated Polyneuropathy

LOCATION▶ Simcoe & Dufferin

Chairs: James Russell, US
& P. James Dyck, US

08:00–08:55 4.1 HEREDITARY POLYNEUROPATHY
Stephan Züchner, US

08:55–09:50 4.2 ASSESSMENT OF NEUROPATHY IN DIABETES AND PRE-DIABETES
James Russell, US

09:50–10:10 NETWORKING BREAK

10:10–11:05 4.3 TREATMENT OF POEMS
P. James Dyck, US

11:05–12:00 4.4 PARANEOPLASTIC NEUROPATHY
Kristine Chapman, CA

08:00–09:50 TC5.0 - Ultrasound in Neuromuscular Disorders

LOCATION▶ Dominion South

Chairs: Francis O. Walker, US
& Ari Breiner, CA

08:00–08:25 5.1 INTRODUCTION TO ULTRASOUND
Ari Breiner, CA

08:25–08:50 5.2 ULTRASOUND OF FOCAL PERIPHERAL NEUROPATHIES
Francis O. Walker, US

08:50–09:15 5.3 ULTRASOUND OF DIFFUSE NEUROPATHIES & MOTOR NEURON DISEASE
Lisa Hobson-Webb, US

09:15–09:40 5.4 ULTRASOUND OF MUSCLE
Steven Shook, US

09:40–09:50 DISCUSSION PERIOD

09:50–10:10 NETWORKING BREAK

10:10–12:00 5.5 PRACTICAL DEMONSTRATIONS

- 10:10–10:45 Station 1**
Median and ulnar nerves
(upper limb)
- 10:45–11:20 Station 2**
Fibular and tibial nerves
(lower limb)
- 11:20–11:55 Station 3**
Muscle +/- diaphragm

Equipment for the Practical Demonstrations
generously provided by GE Healthcare and
Philips Healthcare

12:00–13:00 NETWORKING BREAK

**13:00–17:00 TC 6.0 - Inflammatory Neuropathies
co-sponsored with PNS**

LOCATION► City Hall

Chairs: Ingemar Merkies, NL
& Ivo van Schaik, NL

- 13:00–13:55 6.1 CLINICAL ASPECTS OF IMMUNE
MEDIATED NEUROPATHIES**
Jean-Marc Léger, FR
- 13:55–14:50 6.2 CLINIMETRICS OF IMMUNE MEDIATED
NEUROPATHIES**
Ingemar Merkies, NL

14:50–15:10 NETWORKING BREAK

- 15:10–16:05 6.3 PATHOGENESIS AND IMMUNOLOGY OF
IMMUNE MEDIATED NEUROPATHIES**
Hans-Peter Hartung, DE
- 16:05–17:00 6.4 IMMUNOTHERAPY: WHAT TO DO FIRST,
HOW TO START & STOP, WEAR OFF, LONG-
TERM CONSIDERATIONS, LONG-TERM
OUTCOME**
Ivo van Schaik, NL

13:00–17:00 TC 7.0 - ALS

LOCATION► Churchill

Chairs: Angela Genge, CA
& Michael Benatar, US

- 13:00–13:55 7.1 EVALUATION OF PATIENTS WITH ALS**
Angela Genge, CA
- 13:55–14:50 7.2 THE ROLE OF GENETIC TESTING IN
MOTOR NEURON DISEASE**
Matthew Harms, US

14:50–15:10 NETWORKING BREAK

- 15:10–16:05 7.3 BIOMARKERS RELEVANT TO ALS
THERAPY DEVELOPMENT**
Michael Benatar, US
- 16:05–17:00 7.4 SYMPTOMATIC MANAGEMENT OF ALS**
Stacy Rudnicki, US

13:00–17:00 TC 8.0 - Myopathy

LOCATION► Dominion North

Chair: Mark Tarnopolsky, CA

- 13:00–13:55 8.1 EVALUATION OF METABOLIC MYOPATHY**
Mark Tarnopolsky, CA
- 13:55–14:50 8.2 EXERCISE THERAPY IN MYOPATHY**
Ronni Haller, US

14:50–15:10 NETWORKING BREAK

- 15:10–16:05 8.3 TREATMENT OF MYOTONIC DYSTROPHY**
Charles Thornton, US
- 16:15–17:00 8.4 THE TREATMENT OF INFLAMMATORY
MYOPATHY**
Anthony A. Amato, US

13:00–17:00 TC 9.0 - Autonomic Neuropathy

LOCATION► Simcoe & Dufferin

Chairs: Eva L. Feldman, US
& Paola Sandroni, US

- 13:00–13:55 9.1 INTRODUCTION TO AUTONOMIC
NEUROPATHY**
Eva L. Feldman, US
- 13:55–14:50 9.2 ASSESSMENT OF AUTONOMIC
NEUROPATHY**
Pariwat Thaisetthawatkul, US

14:50–15:10 NETWORKING BREAK

- 15:10–16:05 9.3 POSTURAL ORTHOSTATIC
TACHYCARDIA SYNDROME**
Paola Sandroni, US
- 16:05–17:00 9.4 AUTOIMMUNE AUTONOMIC
NEUROPATHY**
Steven Vernino, US

**13:00–17:00 TC 10.0 - Adult Muscular Dystrophy and
Myopathy**

LOCATION► Dominion South

Chairs: Jiri Vajsar, CA
& Mazen Dimachkie, US

- 13:00–13:55 10.1 UPDATE ON FASCIO-SCAPULO-
HUMERAL DYSTROPHY (FSHD)**
Rabi Tawil, US
- 13:55–14:50 10.2 THE NEED FOR TRANSITION**
Jiri Vajsar, CA

14:50–15:10 NETWORKING BREAK

- 15:10–16:05 10.3 ADULT LIMB-GIRDLE DYSTROPHY**
Anthony A. Amato, US
- 16:05–17:00 10.4 INCLUSION BODY MYOSITIS**
Mazen Dimachkie, US

18:30–21:00 OPENING CEREMONY & WELCOME RECEPTION

LOCATION► Plenary Hall, Grand Ballroom
Centre and Exhibit Hall,
Lower Concourse

CONGRESS PROGRAM ► WEDNESDAY JULY 6, 2016

LOCATION► All Plenary & Workshop Sessions are located on the **Lower Concourse of the Sheraton Centre Toronto Hotel.**

07:00–17:00 REGISTRATION OPEN

10:00–16:00 EXHIBITS AND POSTERS OPEN
LOCATION► Exhibit Hall, Lower Concourse

08:00–10:00 Plenary Session PL 1.0
Genetics
ROOM► Plenary Hall, Grand Ballroom Centre, Lower Concourse level

Chair: James Dowling, CA

08:00–09:00 PL 1.1–GENOMIC APPROACHES TO
DIAGNOSIS OF RARE MUSCLE DISEASE
Keynote Speaker: Daniel MacArthur, US

09:00–09:30 PL 1.2–GENE DISCOVERY IN CHARCOT-
MARIE-TOOTH NEUROPATHIES
Stephan Züchner, US

09:30–10:00 PL 1.3–RNA SEQUENCE AND RNA
ANALYSIS
James Dowling, CA

10:00–10:30 NETWORKING BREAK
LOCATION► Exhibit Hall, Lower Concourse

10:30–12:00 Poster Session 1
ROOM► Exhibit Hall
See page 34 for poster information.

12:00–13:30 INDUSTRY-SUPPORTED SYMPOSIUM
See page 30 for information

13:30–15:00 Workshop WS 1.1
Imaging of Muscle
ROOM► Grand Ballroom Centre
Chair: Ari Breiner, CA

13:30–14:15 WS 1.1.1–MUSCLE ULTRASOUND
Carsten Bonnemann, US

14:15–15:00 WS 1.1.2–THE APPLICATION OF MRI IN
MUSCLE DISEASE
Volker Straub, UK

13:30–15:00 Workshop WS 1.2
Management of ALS Patients

ROOM► Osgoode Ballroom West

Chair: Hans Katzberg, CA

13:30–14:15 WS 1.2.1–SYMPTOMATIC TREATMENT
OF ALS
Stacy Rudnick, US

14:15–15:00 WS 1.2.2–END OF LIFE ISSUES IN ALS
Christen Shoesmith, US

13:30–15:00 Workshop WS 1.3
Metabolic Myopathy

ROOM► Grand Ballroom West

Chair: John Vissing, DE

13:30–14:15 WS 1.3.1–EVALUATION AND TREATMENT
OF POMPE DISEASE
Mark Tarnopolsky, US

14:15–15:00 WS 1.3.2–DIETARY AND OTHER
THERAPIES IN MUSCLE GLYCOGENOSIS
AND DISORDERS OF MUSCLE LIPID
OXIDATION
John Vissing, US

13:30–15:00 Workshop WS 1.4
Neuromuscular Databases

ROOM► Osgoode Ballroom East

Chair: Lawrence Korngut, CA

13:30–14:15 WS 1.4.1–NEUROMUSCULAR DATABASES
Lawrence Korngut, US

14:15–15:00 WS 1.4.2–TREAT NMD
Kevin Flanigan, US

13:30–15:00 **Workshop WS 1.5**
Treatment of Myasthenia Gravis

ROOM ▶ Grand Ballroom East

Chair: Susan Iannaccone, US

13:30–14:15 **WS 1.5.1–GENERAL TREATMENT**
APPROACHES

Gil Wolfe, US

14:15–15:00 **WS 1.5.2–TREATMENT OF MG IN THE**
PAEDIATRIC POPULATION

Susan Iannaccone, US

15:00–15:30 **NETWORKING BREAK**
LOCATION ▶ Exhibit Hall, Lower Concourse

15:30–17:00 **Workshop WS 2.1**
Emerging Concepts in the Pathology
and Clinical Management of
Degenerative Cervical Myelopathy
(DCM)

ROOM ▶ Osgoode Ballroom West

Chair: Michael Fehlings, CA

15:30–15:45 **WS 2.1.1–EMERGING CONCEPTS IN THE**
PATHOBIOLOGY OF DEGENERATIVE
CERVICAL MYELOPATHY, EPIDEMIOLOGY
AND CLINICAL PRESENTATION

Michael Fehlings, US

15:45–16:00 **WS 2.1.2–CLINICAL IMPLICATIONS,**
OUTCOMES AND REHABILITATION
PATHWAYS

Anthony Burns, US

16:00–16:15 **WS 2.1.3–UNDERSTANDING DISEASE**
SEVERITY THROUGH NOVEL SURROGATE
MEASUREMENT APPROACHES IN NTSCI

Sukhvinder Kalsi-Ryan, CA

16:15–16:30 **WS 2.1.4–ADVANCED TECHNIQUES IN**
IMAGING SPECIFIC TO DEGENERATIVE
MYELOPATHY

Julien Cohen-Adad, US

16:30–17:00 **PANEL DISCUSSION**

15:30–17:00 **Workshop WS 2.2**
MRI Studies in Peripheral Nerve
Disease

ROOM ▶ Grand Ballroom East

Chair: Ali Naraghi, CA

15:30–16:15 **WS 2.2.1–NOVEL PROCESSING METHODS**
FOR PERIPHERAL NERVE IMAGING

Jennifer Kollmer, DE

16:15–17:00 **WS 2.2.2–CHALLENGES IN MRI STUDIES**
OF PERIPHERAL NERVES

Ali Naraghi, US

15:30–17:00 **Workshop WS 2.3**
Outcomes in Hereditary Neuropathy

ROOM ▶ Grand Ballroom Centre

Chair: Mary Reilly, UK

15:30–16:15 **WS 2.3.1–OUTCOMES IN CMT**

Mary Reilly, UK

16:15–17:00 **WS 2.3.2–MONITORING HEREDITARY**
NEUROPATHIES IN CLINICAL TRIALS

Michael Shy, US

15:30–17:00 **Workshop WS 2.4**
Role of Skin Punch Biopsy

ROOM ▶ Osgoode Ballroom East

Chair: David Saperstein, US

15:30–16:15 **WS 2.4.1–ROLE OF SKIN PUNCH BIOPSY**
IN CLINICAL PRACTICE

David Saperstein, US

16:15–17:00 **WS 2.4.2–ROLE OF SKIN PUNCH BIOPSY**
AS A RESEARCH OUTCOME MEASURE

Michael James Polydefkis, US

15:30–17:00 **Workshop WS 2.5**
Ultrasound in Peripheral Nerve
Disease, Upper Limb

ROOM ▶ Grand Ballroom West

15:30–17:00 **WS 2.5.1–PRACTICAL DEMONSTRATION**
AND DISCUSSION

Francis O. Walker, US

Equipment for the Practical Demonstration
generously provided by GE Healthcare and
Philips Healthcare

17:15–18:15 **INDUSTRY-SUPPORTED SYMPOSIUM**
See page 30 for information

CONGRESS PROGRAM ▶ THURSDAY JULY 7, 2016

LOCATION ▶ All Plenary & Workshop Sessions are located on the **Lower Concourse of the Sheraton Centre Toronto Hotel.**

07:00–17:00 REGISTRATION OPEN

07:00–08:00 INDUSTRY-SUPPORTED SYMPOSIA
See page 31 for information

10:00–16:00 EXHIBITS AND POSTERS OPEN
LOCATION ▶ Exhibit Hall, Lower Concourse

10:30–12:00 Poster Session 1

ROOM ▶ Exhibit Hall

See page 34 for poster information.

12:00–13:30 INDUSTRY-SUPPORTED SYMPOSIUM
See page 31 for information

08:00–10:00 Plenary Session PL 2.0
Hot Topics

ROOM ▶ Plenary Hall, Grand Ballroom
Centre, Lower Concourse level

Chairs: Vera Bril, CA
& Carlos Navarrete Maldonado, CL

08:00–08:15 PL 2.1–STEM CELL THERAPY IN ALS
Eva Feldman, US

08:15–08:30 PL 2.2–RESULTS OF THE THYMECTOMY
TRIAL IN MYASTHENIA GRAVIS
Gil Wolfe, US

08:30–08:45 PL 2.3–REGAIN: A RANDOMIZED,
DOUBLE-BLIND, PLACEBO-CONTROLLED
MULTI-CENTER PHASE 3 STUDY OF THE
SAFETY AND EFFICACY OF ECULIZUMAB
IN SUBJECTS WITH REFRACTORY
GENERALIZED MYASTHENIA GRAVIS
James Howard, Jr., US

08:45–09:00 PL 2.4–APPROACH TO PATIENT-
CENTERED OUTCOMES RESEARCH
Richard Barohn, US

09:00–09:15 PL 2.5–DO WE STILL NEED MUSCLE
BIOPSY IN THE ERA OF ULTRASOUND?
Carsten Bonnemann, US

09:15–09:30 PL 2.6–THERAPEUTIC APPROACHES TO
INCLUSION BODY MYOSITIS
Mazen Dimachkie, US

09:30–09:45 PL 2.7–TREATMENT OF AMYLOID
NEUROPATHY
David Adams, FR

09:45–10:00 PANEL DISCUSSION

10:00–10:30 NETWORKING BREAK
LOCATION ▶ Exhibit Hall, Lower Concourse

13:30–15:00 Workshop WS 3.1
Challenges in Design of Investigator-
Initiated Research

ROOM ▶ Osgoode Ballroom West

Chair: Richard Barohn, US

13:30–14:15 WS 3.1.1–CHALLENGES FOR
INVESTIGATOR INITIATED TRIALS AND
FOR CONDUCTING MULTICENTER TRIALS
Richard Barohn, US

14:15–15:00 WS 3.1.2–TRANSATLANTIC CHALLENGES
Richard Barohn, US

13:30–15:00 Workshop WS 3.2
Diagnosis and Treatment of Myotonic
Dystrophy

ROOM ▶ Grand Ballroom East

Chair: Charles Thornton, US

13:30–14:15 WS 3.2.1–HOW TO TREAT MYOTONIC
DYSTROPHY
Charles Thornton, US

14:15–15:00 WS 3.2.2–CURRENT KNOWLEDGE OF
DISEASE PROGRESSION IN MYOTONIC
DYSTROPHY
Richard Moxley, III, US

13:30–15:00 Workshop WS 3.3
Modern Concepts in Genetics

ROOM ▶ Grand Ballroom Centre

Chair: Kevin Flanigan, US

13:30–14:15 WS 3.3.1–MOLECULAR DIAGNOSTICS IN
THE NEUROMUSCULAR CLINIC
Grace Yoon, CA

14:15–15:00 WS 3.3.2
Kevin Flanigan, US

- 13:30–15:00** **Workshop**
WS 3.4 – Small Fibre Neuropathy
ROOM▶ Osgoode Ballroom East
Chair: Giuseppe Lauria, IT
- 13:30–14:15** **WS 3.4.1–AMYLOID NEUROPATHY AS A MODEL OF SMALL FIBER NEUROPATHY**
David Adams, FR
- 14:15–15:00** **WS 3.4.2–DIAGNOSIS OF SMALL FIBRE NEUROPATHY**
Giuseppe Lauria, IT

- 13:30–15:00** **Workshop**
WS 3.5 – Ultrasound in Peripheral Nerve Disease, Lower Limb
ROOM▶ Grand Ballroom West
Chair: Francis O. Walker, US
- 13:30–15:00** **WS 3.5.1–PRACTICAL DEMONSTRATION AND DISCUSSION**
Francis O. Walker, USA

Equipment for the Practical Demonstration generously provided by GE Healthcare and Philips Healthcare

15:00–15:30 **NETWORKING BREAK**
LOCATION▶ Exhibit Hall, Lower Concourse

- 15:30–17:00** **Workshop WS 4.1**
Bioinformatics and Clinical Research
ROOM▶ Grand Ballroom West
Chair: Jon Katz, US
- 15:30–16:15** **WS 4.1.1–TECHNOLOGY PLATFORMS FOR COLLABORATIONS IN CLINICAL RESEARCH**
Alexander Sherman, US
- 16:15–17:00** **WS 4.1.2–USING THE ELECTRONIC MEDICAL RECORD FOR RESEARCH**
Jon Katz, US

- 15:30–17:00** **Workshop WS 4.2**
Controversies Over Large Nerve Biopsy
ROOM▶ Osgoode Ballroom West
Chair: Anthony A. Amato, US
- 15:30–16:15** **WS 4.2.1–NERVE BIOPSY ARE RARELY NEEDED**
Anthony A. Amato, US
- 16:15–17:00** **WS 4.2.2–IT IS VALUABLE**
P James Dyck, US

- 15:30–17:00** **Workshop WS 4.3**
Genetics of Hereditary Polyneuropathy
ROOM▶ Grand Ballroom East
Chair: Stephan Züchner, US
- 15:30–16:15** **WS 4.3.1–OVERVIEW OF GENETICS OF HEREDITARY POLYNEUROPATHY**
Stephan Züchner, US
- 16:15–17:00** **WS 4.3.2–CELLULAR REPROGRAMMING AND INHERITED PERIPHERAL NEUROPATHIES: PERSPECTIVES AND CHALLENGES**
Mario Saporta, US

- 15:30–17:00** **Workshop WS 4.4**
Outcome Measures in Inflammatory Neuropathy
ROOM▶ Grand Ballroom Centre
Chair: Jean-Marc Léger, FR
- 15:30–15:42** **WS 4.4.1–WHAT THE PERINOMS STUDY TAUGHT US**
Ingemar Merkies, NL
- 15:42–15:54** **WS 4.4.2–HOW WE SHOULD ASSESS INFLAMMATORY NEUROPATHY**
Jean-Marc Léger, FR

- 15:30–17:00** **Workshop WS 4.5**
Primer for Genetic Testing
ROOM▶ Osgoode Ballroom East
Chair: James Dowling, CA
- 15:30–16:15** **WS 4.5.1–EVALUATION OF VARIANTS OF UNKNOWN SIGNIFICANCE**
Mary Reilly, UK and James Dowling, CA
- 16:15–16:25** **WS 4.5.2–CLINICAL WHOLE EXOME SEQUENCING**
Livija Medne, US
- 16:25–16:45** **WS 4.5.2–GENE PANELS**
Kimberly Amburgey, CA

17:15–18:15 **INDUSTRY-SUPPORTED SYMPOSIUM**
See page 31 for information

CONGRESS PROGRAM ► FRIDAY JULY 8, 2016

LOCATION► All Plenary & Workshop Sessions are located on the **Lower Concourse of the Sheraton Centre Toronto Hotel.**

07:00–20:00 REGISTRATION OPEN

10:00–16:00 EXHIBITS AND POSTERS OPEN
LOCATION► Exhibit Hall, Lower Concourse

12:00–13:30 INDUSTRY-SUPPORTED SYMPOSIUM
See page 32 for more information

- 07:00–08:00 Late Breaking News LB 1.0**
ROOM► Plenary Hall, Grand Ballroom Centre, Lower Concourse level
Chairs: Vera Bril, CA & John England, US
- 07:00–07:20 LB 1.1–A PHASE 2 TRIAL OF RITUXIMAB IN MYASTHENIA GRAVIS: STUDY UPDATE**
Richard J. Nowak, US
- 07:20–07:40 LB 1.2–NEUROLOGICAL COMPLICATIONS ON ZIKA VIRUS**
John England, US
- 07:40–08:00 LB 1.3–GUILLAIN-BARRE SYNDROME AND VARIANTS ASSOCIATED WITH ZIKA VIRUS OUTBREAKS**
Osvaldo Nascimento, Brazil

- 08:00–10:00 Plenary Session PL 3.0 Muscular Dystrophy**
ROOM► Plenary Hall, Grand Ballroom Centre, Lower Concourse level
- 08:00–08:30 PL 3.1–GENE THERAPY FOR MUSCULAR DYSTROPHY**
Keynote Speaker: Dongsheng Duan, US
- 08:30–09:00 PL 3.1–RNA THERAPEUTICS FOR DUCHENNE MUSCULAR DYSTROPHY**
Keynote Speaker: Dana Martin, US
- 09:00–09:30 PL 3.2–ANTISENSE THERAPY FOR MYOTONIC DYSTROPHY**
Charles Thornton, US
- 09:30–10:00 PL 3.3–CRISPR BASED GENE EDITING FOR MUSCULAR DYSTROPHY**
Ronald Cohn, CA

10:00–10:30 NETWORKING BREAK
LOCATION► Exhibit Hall, Lower Concourse

- 10:30–12:00 Poster Session 2**
ROOM► Exhibit Hall
See page 34 for poster information.

- 13:30–15:00 Workshop WS 5.1 Approach to Muscular Dystrophies**
ROOM► Grand Ballroom Centre
Chair: Carsten Bonnemann, US
- 13:30–14:15 WS 5.1.1–THE CLINICIANS APPROACH TO LIMB GIRDLE MUSCULAR DYSTROPHY (LGMD)**
Volker Straub, UK
- 14:15–15:00 WS 5.1.2–CONGENITAL MUSCULAR DYSTROPHIES**
Carsten Bonnemann, US

- 13:30–15:00 Workshop WS 5.2 Interesting Neuromuscular Cases**
ROOM► Grand Ballroom East
Chairs: Aaron Izenberg, CA & Hans Katzberg, CA
- 13:30–13:45 WS 5.2.1–A 47-YEAR-OLD FEMALE PATIENT WITH SLOW PROGRESSIVE DISTAL AND ASYMMETRIC WEAKNESS**
Renata Andrade, BR
- 13:45–14:00 WS 5.2.2–A PATIENT WITH DISTAL WEAKNESS, CRAMPS AND FAINTING, OCULAR MOVEMENT ABNORMALITY**
Corrado Angelini, IT
- 14:00–14:15 WS 5.2.3–IN THE ERA OF EXON SEQUENCING HOW DO WE MANAGE THE PROGRESSIVE PROXIMAL, AXIAL AND FACIAL WEAKNESS LEADING TO COMPLETE BULBAR PALSY AND TONGUE FASCICULATIONS IN CHILDREN?**
Elena Gargaun, FR
- 14:15–14:30 WS 5.2.4–MYOPATHY WITH HYPERCKEMIA AND GLOBAL DEVELOPMENTAL DELAYS: THINK BEYOND THE ALPHA-DYTROGLYCONOPATHIES**
Livija Medne, US
- 14:30–14:45 WS 5.2.5–CRAMP-FASCICULATION SYNDROME - AN UNEXPECTED ETIOLOGY**
Peter Y.K. Van den Bergh, BY
- 14:45–15:00 WS 5.2.6–A CASE OF VELOPHARYNGEAL INSUFFICIENCY**
Veena Vasi, UK

13:30–15:00 **Workshop WS 5.3**
Diagnosis and Treatment of Diabetic Neuropathy

ROOM ▶ Osgoode Ballroom West

Chair: Bruce Perkins, CA

13:30–14:15 **WS 5.3.1–DIAGNOSIS AND TREATMENT OF DIABETIC NEUROPATHY**
Bruce Perkins, CA

14:15–15:00 **WS 5.3.2–THE USE OF OMEGA-3 SUPPLEMENTATION FOR MANAGING DIABETIC NEUROPATHY: RESULTS FROM A CLINICAL PILOT TRIAL**
Evan Lewis, CA

13:30–15:00 **Workshop WS 5.4**
Neuropathic Pain

ROOM ▶ Grand Ballroom West

Chair: John England, US

13:30–14:15 **WS 5.4.1–THE ROLE OF GUIDELINES IN DECISIONS ON TREATMENT**
John England, US

14:15–15:00 **WS 5.4.2–UPDATE ON TREATMENT OF NEUROPATHIC PAIN**
Jaya Trivedi, US

13:30–15:00 **Workshop WS 5.5**
Outcome Scales in Myasthenia Gravis

ROOM ▶ Osgoode Ballroom East

Chair: Ted Burns, US

13:30–14:15 **WS 5.5.1–MYASTHENIA GRAVIS IMPAIRMENT INDEX**
Carolina Barnett Tapia, CA

14:15–15:00 **WS 5.5.2–REVIEW OF CURRENT MG SCALES**
Ted Burns, US

15:00–15:30 **NETWORKING BREAK**
LOCATION ▶ Exhibit Hall, Lower Concourse

15:30–17:00 **Workshop WS 6.1**
ALS Overlap Syndromes

ROOM ▶ Grand Ballroom East

Chair: Maria Carmela Tartaglia, CA

15:30–16:15 **WS 6.1.1–GENETIC ASPECTS OF ALS OVERLAP SYNDROMES**
Ekaterina Rogaeva, CA

16:15–17:00 **WS 6.1.2–CLINICAL ASPECTS OF ALS OVERLAP SYNDROMES**
Maria Carmela Tartaglia, CA

15:30–17:00 **Workshop**
WS 6.2 – Cramps in Neuromuscular Disease

ROOM ▶ Grand Ballroom Centre

Chair: Nicholas Silvestri, US

15:30–16:15 **WS 6.2.1–TREATMENT OF MUSCLE CRAMPS**
Hans Katzberg, CA

16:15–17:00 **WS 6.2.2–ASSESSMENT OF MUSCLE CRAMPS**
Nicholas Silvestri, US

15:30–17:00 **Workshop WS 6.3**
Diabetic Neuropathy

ROOM ▶ Osgoode Ballroom East

Chair: James Russell, US

15:30–16:15 **WS 6.3.1–EPIDEMIOLOGY AND PATHOPHYSIOLOGY OF DIABETIC NEUROPATHY**
James Russell, US

16:15–17:00 **WS 6.3.2–TREATMENT OF DIABETIC NEUROPATHY**
Vera Bril, CA

15:30–17:00 **Workshop WS 6.4**
Peripheral Nerve Tumors

ROOM ▶ Osgoode Ballroom West

Chair: Wolfgang Grisold, AT

15:30–16:15 **WS 6.4.1–LYMPHOMA AND OTHER PERIPHERAL NERVE TUMORS**
Wolfgang Grisold, AT

16:15–17:00 **WS 6.4.2–NEUROFIBROMATOSIS 1 AND MALIGNANT TRANSFORMATION OF PERIPHERAL NERVE SHEATH TUMORS**
Gelareh Zadeh, US

15:30–17:00 **Workshop WS 6.5**
Ultrasound of Muscle and Nerve

ROOM ▶ Grand Ballroom West

Chair: Ari Breiner, CA

15:30–16:15 **WS 6.5.1–NEUROMUSCULAR PHYSICIANS SHOULD PERFORM NM ULTRASOUND**
Steven Shook, US

16:15–17:00 **WS 6.5.2–RADIOLOGISTS SHOULD PERFORM NM ULTRASOUND**
Linda Probyn, US

17:00–22:00 **CONGRESS DINNER**

LOCATION ▶ Willow East & Centre located at the Sheraton on the Mezzanine level

CONGRESS PROGRAM ▶ SATURDAY JULY 9, 2016

LOCATION▶ All Plenary & Workshop Sessions are located on the **Lower Concourse of the Sheraton Centre Toronto Hotel.**

07:00–19:00 REGISTRATION OPEN

07:00–08:00 INDUSTRY-SUPPORTED SYMPOSIA
See page 33 for information

10:00–15:30 EXHIBITS AND POSTERS OPEN
LOCATION▶ Exhibit Hall, Lower Concourse

08:00–10:00 Plenary Session PL 4.0
Motor Neuron Disease
ROOM▶ Plenary Hall, Grand Ballroom
Centre, Lower Concourse level

Chair: John Kissel, US

08:00–09:00 PL 4.1–ALS THERAPY DEVELOPMENT:
CHALLENGES AND OPPORTUNITIES
Keynote Speaker: Michael Benatar, US

09:00–09:30 PL 4.2–BIOLOGY OF C9ORF72 DISEASE
Leonard Petrucelli, US

09:30–10:00 PL 4.3–ANTISENSE THERAPY FOR
SPINAL MUSCULAR ATROPHY
John Kissel, US

10:00–10:30 NETWORKING BREAK
LOCATION▶ Exhibit Hall, Lower Concourse

10:30–12:00 Poster Session 2
ROOM▶ Exhibit Hall
See page 34 for poster information.

12:00–13:30 INDUSTRY-SUPPORTED SYMPOSIUM
See page 33 for information

13:30–15:00 Workshop WS 7.1
Autoantibodies in Neuromuscular
Disease
ROOM▶ Grand Ballroom Centre
Chair: Luis Querol, ES

13:30–14:15 WS 7.1.1–IGG4 AUTOANTIBODIES
RELATED TO NEUROMUSCULAR
DISEASES: THERAPEUTIC IMPLICATIONS
Luis Querol, ES

14:15–15:00 WS 7.1.2–IS IT ALL ABOUT THE
ANTIBODIES
Andrew Mammen, US

13:30–15:00 Workshop WS 7.2
Evaluation of Variants of Unknown
Significance

ROOM▶ Osgoode Ballroom East

Chair: Mary Reilly, UK

13:30–14:15 WS 7.2.1–NERVE
Mary Reilly, UK

14:15–15:00 WS 7.2.2–MUSCLE
Raveen Basran, CA

13:30–15:00 Workshop WS 7.3
Inclusion Body Myopathy

ROOM▶ Grand Ballroom East

Chair: Mazen Dimachkie, US

13:30–14:15 WS 7.3.1–GENERAL TREATMENT
APPROACHES
Mazen Dimachkie, US

14:15–15:00 WS 7.3.2–ONGOING DEVELOPMENTS IN
IBM
Anthony A. Amato, US

13:30–15:00 Workshop WS 7.4
Modern Concepts in Spinal Muscular
Atrophy

ROOM▶ Grand Ballroom West

Chair: Susan Iannaccone, US

13:30–14:15 WS 7.4.1–SMA TODAY
Susan Iannaccone, US

14:15–15:00 WS 7.4.2–UPDATE ON SPINAL
MUSCULAR ATROPHY
John Kissel, US

13:30–15:00 **Workshop WS 7.5**
Update on FSHD

ROOM ▶ Osgoode Ballroom West

Chair: Rabi Tawil, US

13:30–14:15 **WS 7.5.1–RECENT CONCEPTS IN FSHD**
Rabi Tawil, US

14:15–15:00 **WS 7.5.2–CLINICAL PRESENTATION IN FSHD**
Jeffrey Statland, US

15:00–15:30 **NETWORKING BREAK**
LOCATION ▶ Exhibit Hall, Lower Concourse

15:30–17:00 **Workshop WS 8.1**
Exercise Therapy for Metabolic Myopathies

ROOM ▶ Osgoode Ballroom East

Chair: John Vissing, DE

15:30–16:15 **WS 8.1.1–EXERCISE THERAPY IN MITOCHONDRIAL DISORDERS**
Ronni Haller, US

16:15–17:00 **WS 8.1.2–EXERCISE TRAINING AND PATHOPHYSIOLOGY OF EXERCISE IN METABOLIC MYOPATHIES**
John Vissing, DE

15:30–17:00 **Workshop WS 8.2**
How To Do Investigator-Initiated Trials; PCORI

ROOM ▶ Osgoode Ballroom West

Chair: Richard Barohn, US

15:30–16:15 **WS 8.2.1–WHY PATIENT-CENTERED OUTCOMES RESEARCH?**
Richard Barohn, US

16:15–17:00 **WS 8.2.2–INVESTIGATOR-INITIATED CLINICAL TRIALS**
Richard Barohn, US

15:30–17:00 **Workshop WS 8.3**
Metabolic Neuropathies

ROOM ▶ Grand Ballroom West

Chair: A. Gordon Smith, US

15:30–16:15 **WS 8.3.1–NEUROPATHY IN PRE-DIABETES & THE METABOLIC SYNDROME**
A. Gordon Smith, US

16:15–17:00 **WS 8.3.2–NEUROPATHY DUE TO SYSTEMIC DISEASE**
Mamatha Pasnoor, US

15:30–17:00 **Workshop WS 8.4**
Outcome Measures in Neuromuscular Disorders

ROOM ▶ Grand Ballroom Centre

Chair: Linda Lowes, US

15:30–16:15 **WS 8.4.1–BEST OUTCOME MEASURES TO USE FOR NM PATIENTS**
Linda Lowes, US

16:15–17:00 **WS 8.4.2–OUTCOME MEASURES IN MUSCULAR DYSTROPHY**
Craig McDonald, US

15:30–17:00 **Workshop WS 8.5**
Treatment of Muscular Dystrophy

ROOM ▶ Grand Ballroom East

Chair: Kevin Flanigan, US

15:30–15:42 **WS 8.5.1–GENE-DIRECTED TREATMENT OF MUSCULAR DYSTROPHY**
Kevin Flanigan, US

15:42–15:54 **WS 8.5.2–NON-GENE DIRECTED**
Craig Campbell, CA

17:00–19:00 **CLOSING CEREMONY**
LOCATION ▶ Plenary Hall,
Grand Ballroom Centre,
Lower Concourse

POSTER DISPLAY CATEGORIES

GROUP 1

Muscle Diseases of Genetic Origin: Clinical Features, Pathophysiology, Therapy

- 1.1 Dystrophinopathy
- 1.2 Muscle Dystrophies (Non-Dystrophinopathy)
- 1.3 Congenital Muscular Dystrophy
- 1.4 Congenital Myopathies / Myopathies with Prominent Muscle Contractures
- 1.5 Distal Myopathy / Myofibrillar Myopathies
- 1.6 Myotonic Myopathies
- 1.7 Facioscapulohumeral Muscular Dystrophies / Oculopharyngeal Muscular Dystrophy
- 1.8 Metabolic Myopathies / Mitochondrial Myopathies
- 1.9 Muscle Channelopathies and Related Disorders
- 1.10 Other Myopathies Including GNE – Hereditary Inclusion Body Myopathy

GROUP 2

Acquired Myopathies: Clinical Features, Pathophysiology, Therapy

- 2.1 Inflammatory / Dysimmune Myopathies
- 2.2 Inclusion Body Myositis
- 2.3 Toxic / Endocrine / Other Acquired Myopathies

GROUP 3

Diseases of Neuromuscular Junction: Clinical Features, Pathophysiology, Therapy

- 3.1 Myasthenia Gravis
- 3.2 Myasthenic Syndromes
- 3.3 Congenital Myasthenia

GROUP 4

Peripheral Neuropathy: Clinical Features, Pathophysiology, Therapy

- 4.1 Inflammatory / Dysimmune / Associated with Monoclonal Gammopathy/Paraneoplastic
- 4.2 Hereditary Peripheral Neuropathy
- 4.3 Metabolic / Toxic
- 4.4 Infectious Peripheral Neuropathy (including Leprosy, HIV)
- 4.5 Others

GROUP 5

Motor Neuron Diseases: Clinical Features, Pathophysiology, Therapy

- 5.1 Biology, Genetics
- 5.2 Biomarkers in MND
- 5.3 Epidemiology, Clinic, Treatment
- 5.4 Spinal Muscular Atrophy / Neuronopathies

GROUP 6

Novel Diagnostic Methods in Neuromuscular Diseases

- 6.1 Ultrasound
- 6.2 MRI
- 6.3 Other Biomarkers
- 6.4 Electrodiagnosis
- 6.5 Small Nerve Fibre Evaluation
- 6.6 Biochemical and Molecular Techniques

GROUP 7

Basic Sciences in Neuromuscular Diseases

- 7.1 Muscle Homeostasis / Muscle Regeneration
- 7.2 Muscle Structure / Muscle Development / Muscle Growth
- 7.3 Muscle Atrophy / Degeneration
- 7.4 Nuclear Envelope / Nuclear Matrix of Muscle Cell
- 7.5 Ion Channel Function in Neuron and Muscle
- 7.6 Immune Mechanisms in Neuromuscular Diseases
- 7.7 Fundamental Approaches to Motor Neuron, Axon and Related Structures
- 7.8 Neuromuscular Junction: Basic Aspects
- 7.9 Others

GROUP 8

Miscellaneous

- 8.1 Outcome Measures in Clinical Trials
- 8.2 Biomarkers in Neuromuscular Disorders
- 8.3 Home Care / Social Programs in Neuromuscular Diseases
- 8.4 Psychological and Neuropsychological Approaches in Neuromuscular Diseases
- 8.5 Ethics in Neuromuscular Disorders
- 8.6 Rehabilitation in Neuromuscular Diseases
- 8.7 Others

POSTER SESSIONS

BEST POSTERS

Top 10 Best Posters will be displayed on the Electronic Poster Screens during the entire Congress:

SCREEN 1

PS2Group1-055 ANTISENSE TARGETING OF 3' END ELEMENTS INVOLVED IN DUX4 MRNA PROCESSING IS AN EFFICIENT THERAPEUTIC STRATEGY FOR FACIOSCAPULOHUMERAL DYSTROPHY: A NEW GENE SILENCING APPROACH
Anne-Charlotte Marsollier¹, Lucasz Ciszewski², Virginie Mariot¹, Linda Popplewell², Thomas Voit², George Dickson², Julie Dumonceaux¹; ¹Paris, FR, ²London, UK

PS1Group8-002 REDUCTION OF ISOAGGLUTININS IN IVIG BY ANTI-A DONOR SCREENING REDUCES THE RISK OF HEMOLYTIC EVENTS
Ayman Kafal, Montreal, QC, CA

SCREEN 2

PS2Group1-006 ETEPLIRSEN FOR DUCHENNE MUSCULAR DYSTROPHY (DMD): CLINICAL AND BIOCHEMICAL RESULTS WITH LONGITUDINAL COMPARISON TO EXTERNAL CONTROLS ON SIX-MINUTE WALK TEST (6MWT)
J Mendell¹, Nathalie Goemans², Louise Rodino-Klapac¹, Z Sahenk¹, Linda Lowes¹, Lindsay Alfano¹, Katherine Berry¹, E Peterson¹, S Lewis¹, K Shontz¹, P Duda³, C Donoghue⁴, J Dworzak³, B Wentworth⁴, E Kaye⁴, Eugenio Mercuri⁵, DMD Italian Network⁶; ¹Columbus, OH, US, ²Leuven, BE, ³Cambridge, MA, US, ⁴Cambridge, MA, US, ⁵Rome, IT, ⁶Milano, IT

PS2Group1-063 ENZYME REPLACEMENT THERAPY IS BENEFICIAL AFTER 5 YEARS OF TREATMENT IN A LARGE GROUP OF ADULT POMPE PATIENTS
Esther Kuperus, Michelle Kruijshaar, Stephan Wens, Juna de Vries, Marein Favejee, Chris van der Meijden, Dimitris Rizopoulos, Esther Brusse, Pieter van Doorn, Ans van der Ploeg, Nadine van der Beek; Rotterdam, NL

SCREEN 3

PS2Group1-059 FAT OXIDATION IS LIMITED IN MADD DURING EXERCISE, BUT GLUCOSE INFUSION IMPROVES EXERCISE CAPACITY
Karen Madsen¹, Nicolai Preisler¹, Astrid Emilie Buch¹, Mads Stemmerik¹, Pascal Laforêt², John Vissing¹; ¹Copenhagen, DK, ²Paris, FR

PS2Group3-004 SFEMG FINDINGS IN OCULAR COMPLICATIONS OF COSMETIC BOTOX INJECTIONS
Daniela Navarrete¹, Carlos Navarrete¹, Raul Muñoz¹, Vera Bril², Mireya Balart¹; ¹Santiago, CL, ²Toronto, ON, CA

SCREEN 4

PS2Group1-068 THE ANTI-CONVULSANTS LACOSAMIDE, LAMOTRIGINE AND RUFINAMIDE REDUCE MYOTONIA IN ISOLATED HUMAN AND RAT SKELETAL MUSCLE
Thomas Pedersen¹, Martin Skov², Ole Nielsen²; ¹Aarhus C, DK, ²Aarhus, DK

PS2Group1-064 SUGAR INFUSION IMPROVES EXERCISE CAPACITY IN PATIENTS WITH GLYCOGENIN-1 DEFICIENCY
Mads Stemmerik¹, Pascal Laforêt², Astrid Emilie Buch¹, Karen Madsen¹, John Vissing¹; ¹Copenhagen, DK, ²Paris, FR

SCREEN 5

PS2Group1-056 TREATMENT RELATED EFFECTS OF ANTI-GAA ANTIBODIES IN LATE ONSET POMPE DISEASE
Marie Wencil, Claudia Shambaugh, Namita Goyal, Virginia Kimonis, Tahseen Mozaffar; Orange, CA, US

PS2Group1-061 FATTY ACID OXIDATION DEFECTS PRESENTING AS PRIMARY MYOPATHY AND PROMINENT DROPPED HEAD SYNDROME
Seena Vengalil, Veeramani Preethish-Kumar, Kiran Polavarapu, Atchayaram Nalini, Narayanappa Gayathri, Rita Christopher, Manjunath Mahadevappa, Chandrajit Prasad; Bangalore, IN

POSTER SESSION 1WEDNESDAY, JULY 6 &
THURSDAY, JULY 7, 2016

10:30-12:00

ROOM ▶ Exhibit Hall

WEDNESDAY AND THURSDAY SESSIONS SUPPORTED BY



PS1 Group 2

PS1Group2-001 STATIN-INDUCED NECROTIZING AUTOIMMUNE MYOPATHY. RECURRENCE WITH FIBRATE USEMario Fuentealba¹, Jorge Bevilacqua²;
¹Concepcion, CL, ²Santiago, CL**PS1Group2-002 STUDY OF HYALURONIDASE-FACILITATED SCIG IN CHRONIC INFLAMMATORY DEMYELINATING POLYRADICULONEUROPATHY (CIDP)**Claudia Sommer¹, John England²,
Johannes Jakobsen³, Russell Reeve⁴,
David Gelmont⁵; ¹Wurzburg, DE, ²US,
³Copenhagen, DK, ⁴Durham, NC, US,
⁵Westlake Village, CA, US**PS1Group2-003 MECHANISM OF HYALURONIDASE-FACILITATED SCIG ALLOWING INVESTIGATIONS IN NEUROMUSCULAR DISEASE**Christopher Rabbat¹, Tobin Chettiath²,
Martin Noel³, Robert Peterman⁴, Todd
Berner⁵; ¹Kansas, AL, US, ²Westlake
Village, CA, US, ³Mississauga, ON, CA,
⁴Vienna, AT, ⁵Bannockburn, US**PS1Group2-004 RATIONALE FOR TOLL-LIKE RECEPTOR ANTAGONISM AS A POTENTIAL NOVEL THERAPEUTIC APPROACH FOR DERMATOMYOSITIS**Kirsten Gruis¹, Kanneboyina Nagaraju²,
Tahseen Mozaffar³, Anthony A. Amato⁴,
Julie Brevard¹, Lindsey Granlund¹,
Joanna Horobin¹; ¹Cambridge, MA, US,
²Washington, DC, US, ³Orange, CA, US,
⁴Boston, US**PS1Group2-005 INFLAMMATORY MYOPATHIES: NEEDLE ELECTROMYOGRAPHY CHARACTERISTICS IN A SERIE OF CASES**Rosana Scola, Paulo Lorenzoni, Claudia
Kay, Renata Ducci, Paula Rodrigues, Lineu
Werneck; Curitiba, BR**PS1Group2-006 NECROTIZING MYOPATHY ASSOCIATED TO HIV: CASE REPORT**Renata Ducci, Francisco Magalhães, Daniel
Collares, Monica Gomes-da-Silva, Paulo
Lorenzoni, Claudia Kay, Mauricio Carvalho,
Lineu Werneck, Rosana Scola; Curitiba, BR**PS1Group2-007 SRP ANTIBODY ASSOCIATED NECROTIZING MYOPATHY MIMICKED LGMD: A CASE REPORT**
Pariwat Thaiseththawatkul¹, Rodney
McComb²; ¹Omaha, US, ²Omaha, NE, US**PS1Group2-008 MYASTHENIA GRAVIS AND POLYMYOSITIS PRESENTED SIMULTANEOUSLY**
Florentina Berianu, 22, FL, US**PS1Group2-009 FOLLISTATIN GENE THERAPY IMPROVES SIX MINUTE WALK DISTANCE IN SPORADIC INCLUSION BODY MYOSITIS (SIBM)**Jerry Mendell¹, Z Sahenk¹, Mark Hogan¹,
Samiah Al-Zaidy¹, Kevin Flanigan²,
Louise Rodino-Klapac¹, Markus McColly³,
Kathleen Church¹, S Lewis¹, Linda Lowes¹,
Lindsay Alfano¹, Katherine Berry¹, Natalie
Miller¹, Igor Dvorchik¹, Melissa Moore-
Clingenpeel¹, Brian Kaspar¹; ¹Columbus,
OH, US, ²US, ³Columbus, US**PS1Group2-010 WHOLE-BODY MRI IN AMYOPLASIA CONGENITA**
Cam-Tu Emilie Nguyen¹, Sharan Goobie¹,
Craig Campbell²; ¹London, ON, CA, ²London,
ON, CA

PS1 Group 4

PS1Group4-001 RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY TO INVESTIGATE THE EFFICACY, SAFETY AND TOLERABILITY OF TWO DIFFERENT DOSES OF IGPRO20 (SUBCUTANEOUS IMMUNOGLOBULIN) FOR THE TREATMENT OF CIDP-IGG DEPENDENCY AND RESTABILIZATION PHASEIvo van Schaik¹, Vera Bril², Nan van
Geloven³, Hans-Peter Hartung⁴, Richard
Lewis⁵, G Sobue⁶, Billie Durn⁷, John-Philip
Lawo⁴, Orell Mielke⁷, David Cornblath⁸,
Ingemar Merkies⁹, On on behalf of the
PATH study group¹; ¹Amsterdam, NL,
²Toronto, ON, CA, ³Leiden, NL, ⁴Dusseldorf,
DE, ⁵Los Angeles, CA, US, ⁶Nagoya,
JP, ⁷Marburg, DE, ⁸Baltimore, MD, US,
⁹Maastricht, NL**PS1Group4-002 SWITCHING PATTERNS IN PATIENTS WITH ICD-9 DIAGNOSED CIDP INITIATING IVIG TREATMENT**Jeffrey Guptil¹, Jeffrey Allen², Micheal
Runken³, Josh Noone⁴, Emily Zacherle⁵,
Chris Blanchette⁵; ¹Durham, NC, US,
²Minneapolis, MN, US, ³Raleigh, NC, US,
⁴Charlotte, NC, US, ⁵Davidson, NC, US

- PS1Group4-003 SYSTEMIC LUPUS ERYTHEMATOSUS PRESENTING WITH AUTONOMIC AND SOMATIC SMALL FIBER NEUROPATHY**
Oscar Trujillo, Juan Idiaquez, Ricardo Fadic; Santiago, CL
- PS1Group4-004 INCIDENCE OF GUILLAIN-BARRE SYNDROME IN IRANIAN CHILDREN UNDER FIFTEEN YEARS OLD; NATIONAL AFP SURVEILLANCE REPORT (2008-2014)**
Seyed Hassan Tonekaboni, Habibeh Nejad Biglari; Tehran, IR
- PS1Group4-005 POLYNEUROPATHY IN THE LIMELIGHT: A CASE**
Sandya Tirupathi¹, Matthew Sayers², John McConville¹, K Pang¹, Marie-Louise Kane¹; ¹Btba, UK, ²Bt126ba, UK
- PS1Group4-006 VASCULITIC NEUROPATHY COMPLICATED BY ANTERIOR SPINAL ARTERY SYNDROME**
Michael Ackerl¹, Wolfgang Grisold²; ¹Vienna, AT, ²US
- PS1Group4-007 CLINICAL AND ELECTROPHYSIOLOGICAL CHARACTERISTICS OF CHRONIC INFLAMMATORY DEMYELINATING POLYNEUROPATHY IN KOREA**
Seol-Hee Baek, Jun-Soon Kim, BongJe Kim, So Hyun Ahn, Kyomin Choi, Seok-Jin Choi, Jung-Joon Sung, Yoon-Ho Hong; Seoul, KR
- PS1Group4-008 NON-TRAUMATIC PLEXOPATHIES AND RADICULOPATHIES IN CHILDREN**
Cam-Tu Emilie Nguyen¹, Craig Campbell², Hugh McMillan³, Chantal Poulin⁴, Michel Vanasse⁴, Jiri Vajsar⁵; ¹London, ON, CA, ²London, ON, CA, ³Ottawa, ON, CA, ⁴Montreal, QC, CA, ⁵Toronto, ON, CA
- PS1Group4-009 PERIPHERAL T CELL LYMPHOMA PRESENTING AS MILLER FISHER'S SYNDROME**
So Hyun Ahn, Seol-Hee Baek, Jun-Soon Kim, Kyomin Choi, Seok-Jin Choi, Yoon-Ho Hong, Jung-Joon Sung; Seoul, KR
- PS1Group4-010 A COMPARATIVE, DOUBLE-BLIND, RANDOMIZED, MULTICENTRE CLINICAL TRIAL TO ACCESS THE EFFICACY AND SAFETY OF CLAIRYG VS TEGELINE IN MAINTENANCE TREATMENT OF CHRONIC INFLAMMATORY DEMYELINATING POLYRADICULONEUROPATHY (CIDP)**
Claude Desnuelle¹, Jean Pouget², Jean Christophe Antoine³, Arnaud Lacour⁴, Jerome De Seze⁵, Christophe Vial⁶, Anne-Laure Bedat Millet⁷, Julien Cassereau⁸, David Adams⁹, Guilhem Sole¹⁰, Yann Pereon¹¹, Philippe Corcia¹², Thibault Moreau¹³, Steve Genestet¹⁴, Rabye Ouaja¹⁵, Anne Hufschmitt¹⁵, Chrystelle Mercier¹⁵; ¹Nice, FR, ²Marseille, FR, ³St Etienne, FR, ⁴Lille, FR, ⁵Strasbourg, FR, ⁶Lyon, FR, ⁷Rouen, FR, ⁸Angers, FR, ⁹Le Kremlin Bicetre, FR, ¹⁰Bordeaux, FR, ¹¹Nantes, FR, ¹²Tours, FR, ¹³Dijon, FR, ¹⁴Brest, FR, ¹⁵Les Ulis, FR
- PS1Group4-011 AN INTERNATIONAL, MULTICENTRE, EFFICACY AND SAFETY STUDY OF I10E, IQMUNE IN INITIAL AND MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC INFLAMMATORY DEMYELINATING POLYRADICULONEUROPATHY (CIDP)**
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- PS1Group4-012 NEUROMUSCULAR COMPLICATIONS ARE NOT RARE IN MIDDLE EAST RESPIRATORY SYNDROME**
Jee-Eun Kim, Su-yeon Park, Jae-Hyeok Heo, Hye-ok Kim, Sook-hee Song, Sang-Soon Park, Tai-Hwan Park, Jin-Young Ahn, Min-Ky Kim, Jae-Phil Choi; Seoul, KR
- PS1Group4-013 NOVEL ANTIGEN-SPECIFIC TREATMENT FOR ANTI-MYELIN-ASSOCIATED GLYCOPROTEIN NEUROPATHY**
Ruben Herrendorff, Pascal Haenggi, Hélène Pfister, Andreas Steck, Beat Ernst; Basel, CH
- PS1Group4-014 A EUROPEAN, RANDOMISED, DOUBLE-BLIND, CROSS-OVER STUDY OF A NEW 10% HUMAN INTRAVENOUS IMMUNOGLOBULIN VERSUS OTHER IVIG IN PATIENTS WITH MULTIFOCAL MOTOR NEUROPATHY-LIME STUDY**
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- PS1Group4-015 CORRELATION BETWEEN IGM PARAPROTEINEMIA AND MORPHOMETRIC PARAMETERS OF SURAL NERVE IN ANTI-MAG, SGGL NEUROPATHY**
Kon Ping Lin, Hua Chuan Chao, Cheng Ta Chou, Yi-Chung Lee; Taipei, TW
- PS1Group4-016 INFLAMMATORY DIABETIC NEUROPATHY: HELPFUL DIAGNOSTIC PARAMETERS**
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- PS1Group4-017 EGR2 MUTATION ENHANCE PHENOTYPE SPECTRUM OF DEJERINE-SOTTAS SYNDROME**
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- PS1Group4-018 DEMYELINATING FEATURES IN NEUROPHYSIOLOGICAL STUDY OF TRANSTHYRETIN FAMILIAL AMYLOID POLYNEUROPATHY DUE TO VAL30MET MUTATION IN A PORTUGUESE POPULATION**
Marcio Cardoso, Ana Sousa, Katia Valdrez, Teresa Coelho; Porto, PT
- PS1Group4-019 TRANSTHYRETIN-RELATED HEREDITARY AMYLOIDOSIS IN AN ARGENTINE FAMILY WITH TTR TYR114CYS MUTATION**
Marcelo Rugiero, Marcelo Chaves, Mariela Bettini, Maria Ines Araoz, Maria Saez, Patricia Sorroche, Edgardo Cristiano; Buenos Aires,AR
- PS1Group4-020 NOVEL INF2 GENE MUTATIONS IN CZECH PATIENTS WITH SPORADIC HMSN**
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- PS1Group4-021 PREDICTION OF NERVE CONDUCTION STUDIES OUTCOMES IN PATIENTS WITH FAMILIAL AMYLOIDOTIC POLYNEUROPATHY RECEIVING TAFAMIDIS THERAPY**
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- PS1Group4-022 HEREDITARY MOTOR AND SENSORY NEUROPATHY WITH PYRAMIDAL SIGNS CAUSED BY NEFL GENE MUTATION**
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- PS1Group4-023 ATYPICAL CIDP OR CMT IN THE ELDERLY? A CASE REPORT**
Min-Xia Wang¹, Penny Spring², John Pollard³, Judy Spies⁴; ¹Camperdown,NSW, AU, ²Concord,AU, ³Sydney, ACT, AU, ⁴Sydney, NSW, AU
- PS1Group4-024 ANTI-GRAVITY AEROBIC TRAINING IN PATIENTS WITH CHARCOT-MARIE-TOOTH DISEASE TYPES 1A AND 1X**
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- PS1Group4-025 COEXISTENCE OF CHARCOT MARIE TOOTH DISEASE TYPE 1A AND DIABETES: A CLINICOPATHOLOGICAL STUDY**
Kon Ping Lin, Hua Chuan Chao; Taipei, TW
- PS1Group4-026 IN VIVO FUNCTIONAL ANALYSIS OF THE NOVEL BSCL2 P.R96H MUTATION RESULTING IN HEREDITARY MOTOR NEUROPATHY**
Cheng-Tsung Hsiao, Pei-Chien Tsai, Yi-Chu Liao, Kon Ping Lin, Yi-Chung Lee; Taipei, TW
- PS1Group4-027 TWO NOVEL DE NOVO GARS MUTATIONS CAUSE EARLY-ONSET AXONAL CHARCOT-MARIE-TOOTH DISEASE**
Yi-Chu Liao, Yo-Tsen Liu, Pei-Chien Tsai, Bing-Wen Soong, Yi-Chung Lee; Taipei, TW
- PS1Group4-028 BIOPHYSICAL CHARACTERISTICS AND CLINICAL CORRELATION OF GJB1 MUTATIONS IN CHARCOT-MARIE-TOOTH DISEASE TYPE X1**
Pei-Chien Tsai, Yi-Chu Liao, Kon Ping Lin, Yo-Tsen Liu, Yi-Chung Lee; Taipei, TW
- PS1Group4-029 A NICOTINAMIDE ADENINE NUCLEOTIDE (NAD+) PRECURSOR IS A POTENTIAL THERAPY FOR DIABETIC NEUROPATHY**
Krish Chandrasekaran¹, Chen Chen¹, Avinash Sagi², James Russell³; ¹Baltimore, US, ²Baltimore, MD, US, ³US
- PS1Group4-030 FREQUENT LABORATORY TESTS ABNORMALITIES IN PERIPHERAL NEUROPATHY**
Alon Abraham, Majed Majed Alabdali, Abdulla Atsulaiman, Hana Albulaihe, Ari Breiner, Carolon Barnett, Hans Katzberg, Danah Aljaafari, Leif Lovblom, Bruce Perkins, Vera Bril; Toronto, ON, CA
- PS1Group4-031 CLINICAL AND LABORATORY FEATURES OF SMALL FIBER NEUROPATHIES (SFN) WITH IGM VS TS-HDS**
Jafar Kafaie, Minsoo Kim; Saint Louis,MO, US
- PS1Group4-032 AN INTERESTING CASE OF SCIATIC NEUROPATHY**
Jason Lazarou, Toronto, ON, CA
- PS1Group4-033 SUBACUTE BRACHIAL PLEXOPATHY ASSOCIATED WITH CYSTIC SUBCORACOID BURSTITIS**
Suk-Won Ahn, Dae-Woong Kang, Myung-Jin Kim, Jung-Joon Sung, Yoon-Ho Hong, Chang-Seop Kim; Seoul, KR
- PS1Group4-034 A CASE OF NEUROMYOTONIA ASSOCIATED WITH A CHRONIC POLYRADICULONEUROPATHY**
Anna Paula Covaleski¹, Vanessa Mota¹, Otávio Lins¹, Wilson Marques²; ¹Recife, BR, ²Ribeirão Preto, BR

- PS1Group4-035** **QUALITY OF LIFE IN PATIENTS WITH DIABETIC PERIPHERAL NEUROPATHY: A LITERATURE REVIEW**
Semra Aciksoz, Ankara, TR
- PS1Group4-036** **SEASONAL VARIATION OF BELL'S PALSY: A HOSPITAL BASED RETROSPECTIVE STUDY OVER 9 YEARS**
Byung-Nam Yoon¹, Jung-Joon Sung², Suk-Won Ahn², Ji-Eun Kim², Yoon-Ho Hong²; ¹Incheon, KR, ²Seoul, KR
- PS1Group4-037** **THE USE OF OMEGA-3 SUPPLEMENTATION FOR MANAGING DIABETIC NEUROPATHY: RESULTS FROM A CLINICAL PILOT TRIAL**
Evan Lewis¹, Bruce Perkins², Richard Bazinet², Thomas Wolever², Vera Bril²; ¹US, ²Toronto, ON, CA
- PS1Group4-038** **B12 DEFICIENCY IS A CAUSE OF REVERSIBLE AUTONOMIC FAILURE: A CASE REPORT**
Pariwat Thaisethhawatkul, Omaha, US

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- PS1Group6-002** **UNUSUAL CAUSE OF NOCTURNAL HAND PAIN-3 CASES DIAGNOSED BY POCUS AFTER NORMAL EMG**
Abraham Chaiton, Toronto, ON, CA
- PS1Group6-003** **IMPACT OF DRISAPERSEN ON APPARENT FAT FRACTION IN DUCHENNE MUSCULAR DYSTROPHY**
Courtney Bishop¹, Rexford Newbould¹, Zhengning Lin², Robert Janiczek³, Susanne Wang²; ¹London, ON, CA, US, ²Novato, CA, US, ³Middlesex, UK
- PS1Group6-004** **USEFULNESS OF MRI IN CASES OF HYPERCKEMIA**
Pilar Marti¹, Nuria Muelas¹, Jordi Diaz-Manera², Juan J Vilchez¹; ¹Valencia, ES, ²Barcelona, ES
- PS1Group6-005** **LOWER LIMB MUSCLE VOLUME TEST, RE-TEST VARIABILITY USING MRI**
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- PS1Group6-006** **VITAMIN DEFICIENCIES IN PATIENTS WITH VARIOUS MYOPATHIES AND OTHER NEUROMUSCULAR CONDITIONS-PILOT STUDY**
Dubravka Dodig¹, Mark Tarnopolsky²; ¹Toronto, ON, CA, ²Hamilton, ON, CA
- PS1Group6-007** **POTENTIALLY CONFOUNDING VARIABLES OF GDF-15: NEW BIOMARKER OF MITOCHONDRIA DISEASES**
Akiko Ishii¹, Seitaro Nohara¹, Fumiko Yamamoto¹, Shuichi Yatsuga², Makoto Terada¹, Tetushi Aizawa¹, Tetsuto Yamaguchi¹, Kumi Yanagihara¹, Tetsuya Moriyama¹, Naoki Touzaka¹, Zenshi Miyake¹, Hiroshi Tsuji¹, Yasushi Tomidokoro¹, Kiyotaka Nakamagoe¹, Kazuhiro ISHII¹, Masahiko Watanabe¹, Yasutoshi Koga², Akira Tamaoka¹; ¹Tsukuba, JP, ²Kurume, JP
- PS1Group6-008** **A NOVEL ASSESSMENT OF BAROREFLEX ACTIVITY BY PHOTOPLETHYSMOGRAPHY AND TERNARY ARITHMETIC CODING IN A RAT MODEL**
An-Bang Liu¹, Hsien-Tsai Wu², Chun-Keng Lin²; ¹Hualien, TW, ²Shoufeng, TW
- PS1Group6-009** **AGREEMENT BETWEEN AUTOMATED AND MANUAL QUANTIFICATION OF CORNEAL NERVE FIBER LENGTH: IMPLICATIONS FOR DIABETIC NEUROPATHY RESEARCH**
Daniel Scarr, Cesar Falappa, Ilia Ostrovski, Leif Lovblom, Mohammed Farooqi, Dylan Kelly, Tong Wu, Elise Halpern, Mylan Ngo, Eduardo Ng, Andrej Orszag, Vera Bril, Bruce Perkins; Toronto, ON, CA
- PS1Group6-010** **USE OF CORNEAL NERVE FIBRE LENGTH (CNFL) FOR DIABETIC NEUROPATHY IDENTIFICATION IN OLDER PATIENTS WITH LONGSTANDING TYPE 1 DIABETES**
Mohammed Farooqi¹, Leif Lovblom¹, Daniel Scarr¹, Julie Lovshin¹, Yuliya Lytvyn¹, Genevieve Boulet¹, Alanna Weisman¹, Hillary Keenan², Michael Brent¹, Narinder Paul¹, Ilia Ostrovski¹, Vera Bril¹, David Cherney¹, Bruce Perkins¹; ¹Toronto, ON, CA, ²Boston, US
- PS1Group6-011** **VALIDITY OF AN AUTOMATED PROTOCOL OF IN VIVO CORNEAL CONFOCAL MICROSCOPY FOR DIABETIC SENSORIMOTOR POLYNEUROPATHY DETECTION IN TYPE 1 DIABETES**
Daniel Scarr, Nancy Cardinez, Ilia Ostrovski, Tong Wu, Mohammed Farooqi, Leif Lovblom, Elise Halpern, Ausma Ahmed, Mylan Ngo, Eduardo Ng, Andrej Orszag, Vera Bril, Bruce Perkins; Toronto, ON, CA
- PS1Group6-012** **VALIDATION OF COOLING DETECTION THRESHOLD AS A MARKER OF SENSORIMOTOR POLYNEUROPATHY IN TYPE 2 DIABETES**
Mohammed Farooqi, Andrej Orszag, Zoe Lysy, Leif Lovblom, Elise Halpern, Mylan Ngo, Eduardo Ng, Ari Breiner, Vera Bril, Bruce Perkins; Toronto, ON, CA

PS1Group6-013 HEREDITARY NEUROPATHIES: THE ROLE OF COPY NUMBER VARIATIONS (CNVS) IN THE NGS TARGETED GENE PANEL DIAGNOSTIC TESTING

Petra Lassuthova, Jana Neupauerová, Simona Marková, Marcela Krůtová, Radim Mazanec, Dana Brožková, Pavel Seeman; Prague ,CZ

PS1Group6-014 HOW MULTI-GENE PANELS CAN CHANGE THE LANDSCAPE OF DIAGNOSING NEUROMUSCULAR DISORDERS

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PS1Group6-015 SENSITIVITY AND SPECIFICITY OF DR1 BISULFITE SEQUENCING IN DETECTING SMCHD1 MUTATION IN A COHORT OF FSHD1 AND FSHD-LIKE PATIENTS

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PS1Group6-01 ROBUST GENOTYPING IN THE DIAGNOSTICS OF LIMB GIRDLE MUSCULAR DYSTROPHIES

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PS1Group6-017 PROFLECT: A USER-FRIENDLY TOOL TO DETECT COPY NUMBER VARIATION (CNV) AMONG AMPLICON SEQUENCING DATA

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PS1Group6-018 GENETIC SEQUENCING OF PATIENTS WITH LIMB GIRDLE MUSCLE WEAKNESS USING AN NGS PANEL

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PS1Group8-001 A PHASE III DOUBLE-BLIND, RANDOMIZED, PLACEBO-CONTROLLED STUDY (SIDEROS) ASSESSING THE EFFICACY OF IDEBENONE IN SLOWING THE RATE OF RESPIRATORY FUNCTION LOSS IN PATIENTS WITH DUCHENNE MUSCULAR DYSTROPHY RECEIVING GLUCOCORTICOID STEROIDS

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PS1Group8-003 DEVELOPMENT OF A NOVEL TOOL FOR ASSESSMENT OF CRAMP SEVERITY: THE TORONTO CLINICAL CRAMP INDEX (TCCI)

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PS1Group8-004 RELIABILITY AND VALIDITY OF THE 100 METER TIMED TEST AS AN OUTCOME MEASURE IN DUCHENNE MUSCULAR DYSTROPHY

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PS1Group8-005 INTRAVENOUS IMMUNOGLOBULIN "WEAR-OFF EFFECT" IN CIDP: STUDY DESIGN AND PROGRESS UPDATE

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PS1Group8-006 IDEBENONE REDUCES RESPIRATORY COMPLICATIONS IN PATIENTS WITH DUCHENNE MUSCULAR DYSTROPHY

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- PS1Group8-007 TREATMENT EFFECT OF IDEBENONE ON INSPIRATORY FUNCTION IN PATIENTS WITH DUCHENNE MUSCULAR DYSTROPHY**
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- PS1Group8-008 REFERENCE VALUES FOR THE THREE-MINUTE WALK TEST, NORTH STAR AMBULATORY ASSESSMENT AND TIMED TESTS IN TYPICALLY DEVELOPING BOYS AGED 2.5-5 YEARS**
Katrijn Klingels, Jasmine Hoskens, Lise Van Verdegem, Marleen Van den Hauwe, Gunnar Buyse, Nathalie Goemans; Leuven, BE
- PS1Group8-009 NEUROMUSCULAR JUNCTION IN EXPERIMENTAL AUTOIMMUNE ENCEPHALOMYELITIS: A HISTOPATHOLOGICAL ANALYSIS**
Thalita Rocha¹, Jetro Sguarezi¹, Sara Ferreira¹, Rodolfo Thomé², Liana Verinaud², Catarina Rapôso²; ¹Bragança Paulista, BR, ²Campinas, BR
- PS1Group8-010 WHEN SHOULD WE TREAT HYPERCKEMIA?**
Astrid Emilie Buch, Karen Pedersen, Sofie Ostergaard, Jesper Thomassen, Ruth Frikke-Schmidt, Nanna Witting, John Vissing; Copenhagen E, DK
- PS1Group8-011 VALIDATION OF PROTEIN BIOMARKERS FOR DUCHENNE MUSCULAR DYSTROPHY**
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- PS1Group8-012 SURVEY ON USAGE OF TELECOMMUNICATION TERMINALS IN JAPANESE PATIENTS WITH NEUROMUSCULAR DISEASES**
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- PS1Group8-013 IS HOME TREATMENT IN AUTO-IMMUNE DISEASE PATIENTS TREATED BY IVIG SAFE?**
Guilhem Sole¹, Claude Desnuelle², Jean-Philippe Azulay³, Gérard Besson⁴, Jean Christophe Antoine⁵, Françoise Buhour⁶, Alain Creange⁷, Gwendal Le Masson¹, Laurent Magy⁸, Sebastien Marcel⁹, Jean-Michel Paquet¹⁰, Francois Rouhart¹¹, Rabye Ouaja¹², Marc Gauthier-Darnis¹², Sophie Puget¹³; ¹Bordeaux, FR, ²Nice, FR, ³Marseille, FR, ⁴Grenoble, FR, ⁵St Etienne, FR, ⁶Lyon, FR, ⁷Creteil, FR, ⁸Limoges, FR, ⁹Chambery, FR, ¹⁰Laval, FR, ¹¹Brest, FR, ¹²Les Ulis, FR, ¹³Les Luis, FR
- PS1Group8-014 AT HOME VERSUS HOSPITAL IVIG FOR THE TREATMENT OF MULTIFOCAL MOTOR NEUROPATHY (MMN), CHRONIC INFLAMMATORY DEMYELINATING POLYRADICULONEUROPATHY (CIDP) AND LEWIS SUMNER SYNDROME (LSS): A COST OF ILLNESS STUDY**
Emilen Delmont¹, Claude Desnuelle², Guilhem Sole³, Isabelle Durand-Zaleski⁴, Marc Gauthier-Darnis⁵, Rabye Ouaja⁵, Sophie Puget⁶; ¹Marseille, FR, ²Nice, FR, ³Bordeaux, FR, ⁴Creteil, FR, ⁵Les Ulis, FR, ⁶Les Luis, FR
- PS1Group8-015 IMPACT OF DIABETIC NEUROPATHY ON DIABETES DISTRESS AND DEPRESSION IN LONGSTANDING T1DM: RESULTS FROM THE CANADIAN STUDY OF LONGEVITY IN TYPE 1 DIABETES**
Johnny-Wei Bai¹, Alanna Weisman¹, Mohammed Farooqi¹, Leif Lovblom¹, Elise Halpern¹, Genevieve Boulet¹, Devrim Eldelekli¹, Julie Lovshin¹, Yuliya Lytvyn¹, Hillary Keenan², Michael Brent¹, Narinder Paul¹, Vera Bril¹, David Cherney¹, Bruce Perkins¹; ¹Toronto, ON, CA, ²Boston, US
- PS1Group8-016 ORAL MOTOR COMMUNICATION INVENTORY FOR ALS: CONTENT VALIDATION**
Laura Ball¹, Gary Pattee², Jichuan Wang¹; ¹Washington, DC, US, ²Lincoln, US
- PS1Group8-017 REHABILITATION NURSING IN NEUROMUSCULAR DISEASES**
Tulay Basak, Ankara, TR
- PS1Group8-018 PROBLEMS OF FAMILIES LIVING WITH BOYS WITH DUCHENNE MUSCULAR DYSTROPHY (DMD) IN A DEVELOPING COUNTRY**
Yakup Sayın, Coşkun Özdemir; İstanbul, TR
- PS1Group8-019 REAL WORLD USE OF PRIVIGEN IN THE TREATMENT OF GBS AND CIDP: RESULTS OF A RETROSPECTIVE OBSERVATIONAL STUDY**
Ayman Kafal, Montreal, QC, CA

- PS1Group8-020 REFRACTORY POSTURAL ORTHOSTATIC TACHYCARDIA SYNDROME: EFFICACY AND SAFETY OF WEEKLY ALBUMIN INFUSIONS**
Zaeem Siddiqi, Aimee Soloway, Derrick Blackmore; Edmonton, AB, CA
- PS1Group8-021 RD-CONNECT: DATA SHARING AND ANALYSIS FOR RARE DISEASE RESEARCH WITHIN THE INTEGRATED PLATFORM AND THROUGH GA4GH BEACON AND MATCHMAKER EXCHANGE**
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- PS1Group8-022 CASE REPORT OF RECURRENT MENINGITIS SECONDARY TO CSF RHINORRHEA**
Aleena Soomro, Karachi, PK
- PS1Group8-023 CLINICAL PRESENTATION OF ANTI-NMDA ENCEPHALITIS**
Rabail Karim, Bashir Soomro; Karachi, PK
- PS1Group8-024 NUDT15 VARIANT IS THE MOST COMMON VARIANT ASSOCIATED WITH THIOPURINE-INDUCED EARLY LEUKOPENIA AND ALOPECIA IN KOREAN PATIENTS WITH VARIOUS NEUROLOGICAL DISEASES**
Sun-Young Kim¹, Dae-Seong Kim², Jin-Hong Shin², Jin-Sung Park³, Sa-Yoon Kang⁴, Ki-Jong Park⁵, Tai-Seung Nam⁶, So-Young Huh⁷, Jong-kuk Kim⁷; ¹44033, KR, ²-, KR, ³Daegu, KR, ⁴Jeju, KR, ⁵Jinju, KR, ⁶Gwangju, KR, ⁷Pusan, KR
- PS1Group8-025 DATABASE OF NEUROMUSCULAR DISEASES IN REGION OF CONCEPCION IN CHILE**
Mario Fuentealba, Concepcion, CL
- PS1Group8-026 SEROPREVALENCE OF HUMAN T-LYMPHOTROPIC VIRUS TYPE 1 IN PATIENTS WITH SURGICAL HISTORY IN KAGOSHIMA, SOUTHERN JAPAN**
Yuichi Tashiro¹, Eiji Matsuura¹, Satoshi Nozuma¹, Akihiro Hashiguchi², Osamu Watanabe², Hiroshi Takashima¹; ¹Kagoshima City, JP, ²Kagoshima, JP
- PS1Group8-027 HIGH RISK BREAST CANCER SCREENING IN WOMEN WITH NEUROFIBROMATOSIS TYPE 1**
Jeanna McCuaig, Shelley Westergard, Catherine Maurice, Paul Kongkham, Galareh Zadeh, Carolina Barnett-Tapia, Vera Bril, Raymond Kim; Toronto, ON, CA
- PS1Group8-029 UNDERSTANDING THE CANADIAN NEUROMUSCULAR DISEASE RESEARCH LANDSCAPE**
Megan Johnston¹, Christopher MacDonald¹, Jeff Dilworth², Hans Katzberg³, Jean Mah¹, Lawrence Korngut¹; ¹Calgary, AB, CA, ²Ottawa, ON, CA, ³Toronto, ON, CA
- PS1Group8-030 CANADIAN NEUROMUSCULAR DISEASES NETWORK (CAN-NMD) THE DEVELOPMENT & IMPLEMENTATION OF A WEB-BASED KNOWLEDGE SHARING AND EXCHANGE PLATFORM**
Gracia Mabaya¹, Craig Campbell¹, Cynthia Gagnon², Megan Johnston³, Laura McAdam⁴, Jeremy Dixon³, Kelvin Jones⁵, Charles Kassardjian⁴, Aneal Khan³, Jane Mitchell⁴, Annie Plourde², Maryam Oskoui⁶, Lawrence Korngut³; ¹London, ON, CA, ²Jonquière, QC, CA, ³Calgary, AB, CA, ⁴Toronto, ON, CA, ⁵Edmonton, AB, CA, ⁶Montreal, QC, CA
- PS1Group8-031 UNDERSTANDING DECISION NEEDS FOR RESPIRATORY INTERVENTIONS IN PAEDIATRIC NEUROMUSCULAR DISORDERS FROM THE PERSPECTIVE OF HEALTHCARE PROVIDERS**
Gracia Mabaya¹, Sherri Katz², Margaret Lawson², April Price¹, Dhenuka Radhakrishnan², Jean Mah³, Lawrence Korngut³, Hugh McMillan², Cheryl Scholtes¹, Allyson Shephard², Melissa Heletea², Craig Campbell¹, Lynda Hoey²; ¹London, ON, CA, ²Ottawa, ON, CA, ³Calgary, AB, CA
- PS1Group8-032 A PROPOSAL: ISAACS SYNDROME (ACQUIRED NEUROMYOTONIA) DIAGNOSTIC CRITERIA**
Osamu Watanabe¹, Kimiyoshi Arimura¹, Hiroshi Takashima²; ¹Kagoshima, JP, ²Kagoshima City, JP
- PS1Group8-033 UNDERSTANDING THE PERSPECTIVES OF YOUNG ADULTS WITH DUCHENNE MUSCULAR DYSTROPHY AS THEY TRANSITION TO ADULTHOOD AND ADULT HEALTH CARE**
Sally Lindsay, Laura McAdam, Tania Mahenderin; Toronto, ON, CA

PS1Group8-034 CHARACTERISTICS DEVELOPMENT OF A NEW IVIG (I10) THE QUALITY BY DESIGN APPROACH (QBD)

Philippe Paolantonacci, Catherine Decoupade, Philippe Appourchaux, Catherine Michalski, Rabye Ouaja, Ousmane Alfa Cisse, Ludovic Burlot; Les Ulis, FR

PS1Group8-035 PURIFICATION OF IVIG (INTRAVENOUS IMMUNOGLOBULIN) FROM IGNG MANUFACTURING PROCESS TO OPTIMIZE PRODUCT TOLERABILITY PROFILE:

EXAMPLE OF HUMAN NORMAL IMMUNOGLOBULIN (IQYMUNE® 100 MG, ML, SOLUTION FOR INFUSION)
Philippe Paolantonacci¹, Catherine Decoupade¹, Philippe Appourchaux¹, Catherine Michalski¹, Rabye Ouaja¹, Sophie Puget², Ludovic Burlot¹; ¹Les Ulis, FR, ²Les Luis, FR

PS1Group8-036 HEREDITARY MUSCLE DISORDERS IN MIDDLE EUROPE: DATA FROM HOSPITAL REGISTRY

Stan Vohanka, Josef Bednarik, Olesja Parmova, Magda Chmelikova, Lenka Fajkusova; Brno, CZ

PS1Group8-037 ANTINOCICEPTIVE AND ANTI-INFLAMMATORY EFFECTS OF COMBINED ADMINISTRATION OF VITAMIN B12 AND KETOROLAC IN RATS

MD Mizanur Rahman, Dhaka, BD

PS1Group8-038 MEASURING PRIORITIES AND GOALS OF CHILDREN WITH DUCHENNE MUSCULAR DYSTROPHY TO DEVELOP A MEANINGFUL PATIENT REPORTED OUTCOME MEASURE

Roni Propp¹, Sarah Buttle², Shannon Weir¹, Clarissa Encisa¹, Aileen Davis¹, Laura McAdam¹, Nancy Salbach¹, Unni Narayanan¹; ¹Toronto, ON, CA, ²Ottawa, ON, CA

POSTER SESSION 2

**FRIDAY, JULY 8 &
SATURDAY, JULY 9, 2016**

10:30-12:00

ROOM► Exhibit Hall

FRIDAY SESSION SUPPORTED BY

Baxalta

SATURDAY SESSION SUPPORTED BY

Biogen

PS2 Group 1

PS2Group1-001 DYSFELINOPATHIES IN BURKINA FASO: A CASE REPORT

Anselme Dabilgou, Christian Napon, Julie M A Kyelem, Alassane Drave, Anila Bhunnoo, Jean Kabore; Ouagadougou, BF

PS2Group1-002 THE QUALITY OF LIFE IN PATIENTS WITH DUCHENNE MUSCULAR DYSTROPHY, IRANIAN EXPERIENCE

Gholamreza Zamani, Morteza Heidari, Mahshid Mehdizadeh; Tehran, IR

PS2Group1-003 CURRENT STATUS OF DYSTROPHINOPATHY NATIONAL REGISTRY IN JAPAN

En Kimura¹, Madoka Mori-Yoshimura², Satomi Mitsunashi², Fumi Takeuchi³, Harumasa Nakamura³, Hirohumi Komaki², Ichizo Nishino², Mitsuru Kawai⁴, Shin'ichi Takeda²; ¹Kodaira, JP, ²Tokyo, JP, ³Kodaira, Tokyo, JP, ⁴Hasuda, Saitama, JP

PS2Group1-004 CHARACTERISTICS OF JAPANESE PATIENTS WITH BECKER MUSCULAR DYSTROPHY IN A JAPANESE NATIONAL REGISTRY OF MUSCULAR DYSTROPHY (REMUDY): HETEROGENEITY AND CLINICAL VARIATION

Madoka Mori-Yoshimura¹, Satomi Mitsunashi¹, Hirohumi Komaki¹, Naohiro Yonemoto², Harumasa Nakamura³, Fumi Takeuchi³, Yukiko Hayashi¹, Miho Murata¹, Ichizo Nishino¹, Shin'ichi Takeda¹, En Kimura²; ¹Tokyo, JP, ²Kodaira, JP, ³Kodaira, Tokyo, JP

PS2Group1-005 META-ANALYSES OF ATALUREN IN PATIENTS WITH NONSENSE MUTATION DUCHENNE MUSCULAR DYSTROPHY

Craig Campbell¹, Francesco Muntoni², Eugenio Mercuri³, Xiaohui Luo⁴, Gary Elfring⁴, Hans Kroger⁵, Peter Riebling⁵, Tuyen Ong⁵, Robert Spiegel⁵, Stuart W Peltz⁵, Craig McDonald⁶; ¹London, ON, CA,

²London, UK, ³Rome, IT, ⁴South Plainfield, AL, US, ⁵South Plainfield, NJ, US, ⁶Sacramento, CA, US

- PS2Group1-007 LONGITUDINAL EFFECT OF DRISAPERSEN VERSUS HISTORICAL CONTROLS ON AMBULATION IN DUCHENNE MUSCULAR DYSTROPHY**
Nathalie Goemans¹, Már Tulinius², Anna-Karin Kroksmark², Marleen Van den Hauwe¹, Zhengning Lin³, Susanne Wang³, Giles Champion⁴; ¹Leuven, BE, ²Gothenburg, SE, ³Novato, CA, US, ⁴Leiden, NL
- PS2Group1-008 IMPACT OF MUSCLE FUNCTION, NUTRITIONAL STATE AND SYSTEMIC INFLAMMATION, ON BONE MINERAL DENSITY IN PATIENTS WITH DUCHENNE MUSCULAR DYSTROPHY**
Oriana Cruz¹, Maricela Rodriguez-Cruz², Carlos Wong-Baeza³, Salvador Atilano-Miguel², Tomas Almeida-Becerril²; ¹Mexico, MX, ²D.f, MX, ³Mexico, D.f, MX
- PS2Group1-009 MIBG THERAPY FOR AN INOPERABLE PARAGANGLIOMA IN DUCHENNE MUSCULAR DYSTROPHY**
Denis Duboc, Marine Paul, Laurie Fanon, Karim Wahbi, Marco Alifano, Florence Tenenbaum, Laurence Guignat; Paris, FR
- PS2Group1-010 EFFECT OF METFORMIN ON IN VIVO AND EX VIVO PATHOLOGY SIGNS IN EXERCISED DYSTROPHIC MDX MICE**
Roberta Capogrosso, Anna Cozzoli, Arcangela Giustino, Paola Mantuano, Francesca Sanarica, Michela De Bellis, Annamaria De Luca; Bari, IT
- PS2Group1-011 DEVELOPMENT OF A PATIENT-REPORTED OUTCOME MEASURE FOR UPPER LIMB FUNCTION IN DUCHENNE MUSCULAR DYSTROPHY (DMD-UPPER LIMB PROM)**
Katrijn Klingels¹, Anna Mayhew², Elena Mazzone³, Michelle Eagle², Tina Duong⁴, Valérie Decostre⁵, Marion Main⁶, Marleen Van den Hauwe¹, Ulla Werlauff⁷, Imelda De Groot⁸, Sonia Messina⁹, Valeria Ricotti⁶, Giles Champion¹⁰, Laurent Servais⁵, Elizabeth Vroom¹¹, Eugenio Mercuri³, Nathalie Goemans¹; ¹Leuven, BE, ²Newcastle Upon Tyne, UK, ³Rome, IT, ⁴Stanford, US, ⁵Paris, FR, ⁶London, UK, ⁷Aarhus, DK, ⁸Nijmegen, NL, ⁹Messina, IT, ¹⁰Leiden, NL, ¹¹Veenendaal, NL
- PS2Group1-012 GENOTYPE PHENOTYPE ANALYSIS OF MULTIPLEX LIGATION DEPENDENT PROBE AMPLIFICATION (MLPA) POSITIVE DUCHENNE, BECKER MUSCULAR DYSTROPHY (DMD, BMD) PATIENTS**
Seena Vengalil, Kiran Polavarapu, Veeramani Preethish-Kumar, Atchayaram Nalini, Meera Purushottam, Deepha Sekar; Bangalore, IN
- PS2Group1-013 CLINICAL AND MUSCLE BIOPSY CHARACTERISTICS OF A COHORT OF CHILDREN UNDER TWO YEARS OF AGE WITH DUCHENNE MUSCULAR DYSTROPHY**
Ana Sousa, Elisa Costa, Ricardo Taipa, Melo Pires, Manuela Santos; Porto, PT
- PS2Group1-014 NOVEL MOUSE MODEL OF DUCHENNE MUSCULAR DYSTROPHY WITH DELETION OF EXONS 834**
Tatiana Dimitrieva, Alexey Deikin, Denis Reshetov, Dmitry Vlodayets, Eugeniia Zotova; Moscow, RU
- PS2Group1-015 THE NEED FOR TRANSITION THE NEED FOR TRANSITION. WORKSHOP TC10.2**
Jiri Vajsar, Toronto, ON, CA
- PS2Group1-016 A CASE REPORT OF A 10 YEAR OLD BOY WITH COMBINATION OF DMD AND DOWN SYNDROME**
Dmitry Vlodayets, Marina Komarova, Denis Reshetov; Moscow, RU
- PS2Group1-017 PROGNOSTIC MODEL FOR 1-YEAR CHANGE IN 6-MINUTE WALK DISTANCE (6MWD) IN PATIENTS WITH DUCHENNE MUSCULAR DYSTROPHY (DMD)**
Nathalie Goemans¹, James Signorovitch², Elyse Swallow², Jinlin Song², Susan Ward³; ¹Leuven, BE, ²Boston, MA, US, ³Cambridge, MA, US
- PS2Group1-018 DIAGNOSIS OF DUCHENNE MUSCULAR DYSTROPHY IN ITALY: CRITICAL ISSUES AND AREAS FOR IMPROVEMENTS**
Adele D'Amico¹, Michela Catteruccia¹, Marika Pane¹, Giovanni Baranello², Alessandra Govoni², Sonia Messina³, Maria Grazia D'angelo⁴, Luisa Politano⁵, Ksenjia Gorni², Stefano Carlo Previtali², Antonella Pini⁶, Roberta Battini⁷, Angela Berardinelli⁸, Federica Ricci⁹, Elena Pegoraro¹⁰, Claudio Bruno¹¹, Federica Trucco¹¹, Barbara Panasis², Giuseppe Vita³, Tiziana Mongini⁹, Maurizio Moggio², Giacomo Pietro Comi², Eugenio Mercuri¹, Enrico Bertini¹; ¹Rome, IT, ²Milan, IT, ³Messina, IT, ⁴Bosiso Parini, IT, ⁵Naples, IT, ⁶Bologna, IT, ⁷Pisa, IT, ⁸Pavia, IT, ⁹Turin, IT, ¹⁰Padua, IT, ¹¹Genoa, IT
- PS2Group1-019 THE PROPHYLACTIC USE OF PAMIDRONATE ON GLUCOCORTICOID-INDUCED BONE LOSS IN THE MDX MOUSE MODEL OF DUCHENNE MUSCULAR DYSTROPHY**
Sung-Hee Yoon, Jinghan Chen, Marc Grynepas, Jane Mitchell; Toronto, ON, CA
- PS2Group1-020-TRPV2 INHIBITION THERAPY CAN BE EFFECTIVE FOR CARDIOMYOPATHY OF MUSCULAR DYSTROPHY**
Tsuyoshi Matsumura¹, Misa Matsui¹, Yuko Iwata², Masanori Asakura², Toshio Saito¹, Harutoshi Fujimura¹, Saburo Sakoda¹; ¹Toyonaka, JP, ²Suita, JP

- PS2Group1-021 A MISSENSE MUTATION IN THE PUTATIVE SARCOPLASMIC RETICULUM TRANSMEMBRANE PROTEIN DCST2 CAUSES THE STRONGMAN SYNDROME**
Talita Conte¹, Martine Tétreault^{1,2}, Marie-Josée Dicaire³, Sylvie Provost⁴, Najwa Al-Bustani⁵, Marie-Pierre Dubé⁵, Véronique Bolduc⁵, Myriam Srour⁶, Erin O’Ferrall⁵, Jean-Pierre Bouchard⁷, Gina Ravenscroft⁸, Russell Hepple⁵, Tanja Taivassalo⁵, Nigel Laing⁹, Phillipa Lamont¹⁰, Jean Mathieu¹¹, Bernard Brais⁵; ¹2b4, QC, CA, ²B, QC, CA, ³B, CA, ⁴C, QC, CA, ⁵Montreal, QC, CA, ⁶Montreal, QC, CA, ⁷Quebec, QC, CA, ⁸Nedlands,AU, ⁹Nedlands, WA, AU, ¹⁰Perth, WA, AU, ¹¹Jonquiere, QC, CA
- PS2Group1-022 HEREDITARY MYOPATHY WITH EARLY RESPIRATORY FAILURE**
Sandra Sousa¹, Jorge Oliveira², Emília Vieira², Teresa Coelho², Manuela Santos², Marcio Cardoso², Ricardo Taipa², Melo Pires², Rosário Santos²; ¹Cascais, PT, ²Porto, PT
- PS2Group1-023 GNE MYOPATHY: MILESTONES AND DISEASE PROGRESSION BASED ON PATIENT SELF-REPORTED DATA COLLECTED THROUGH THE GLOBAL PATIENT REGISTRY**
Oksana Pogoryelova¹, Phillip Cammish², Supriya Rao³, Ed Conner³, Alison Skrinar³, Hanns Lochmüller⁴; ¹Newcastle Upon Tyne, UK, ²Newcastle Upon Tyne, UK, ³Novato, US, ⁴Newcastle, UK
- PS2Group1-024 MULTISYSTEM PROTEINOPATHY WITH MOTOR NERVE CONDUCTION BLOCKS**
Oscar Trujillo, Juan Casar, Roger Gejman, Ricardo Fadic; Santiago, CL
- PS2Group1-025 PHENOTYPIC CHARACTERIZATION AND PATTERN OF MUSCLE INVOLVEMENT IN GNE MYOPATHY**
Veeramani Preethish-Kumar¹, Oksana Pogoryelova², Kiran Polavarapu¹, Narayanappa Gayathri¹, Seena Vengalil¹, Judith Hudson², Chandrajit Prasad¹, Hanns Lochmüller³, Atchayaram Nalini¹; ¹Bangalore, IN, ²Newcastle Upon Tyne, UK, ³Newcastle, UK
- PS2Group1-026 CALPAINOPATHIES IN CHILE**
Jorge Bevilacqua¹, Yves Mathieu², Martin Krahn², Marc Bartoli², Claudia Castiglioni¹, Karin Kleinsteuber¹, Jorge Díaz¹, Francesca Puppo², Mathieu Cerino², Sebastien Courrier², Svetlana Gorokhova², Alejandra Tringulao¹, Natalia Miranda¹, Patricio Gonzalez-Hormazabal¹, María De Los Ángeles Avaria¹, J Urtizberea³, Pablo Caviedes¹, Lilian Jara¹, Nicolas Levy²; ¹Santiago, CL, ²Marseille, FR, ³Hendaye, FR
- PS2Group1-027 THE FIRST FUNCTIONALLY MATURE HUMAN PRIMARY IN VITRO MUSCLE MODEL: A NEW PARADIGM TO EXPLORE MUSCLE PHYSIOPATHOLOGY AND ACCELERATE DRUG DISCOVERY FOR MUSCLE DISORDERS**
Joris Michaud¹, Mathieu Fernandes¹, Eve Duchemin-Pelletier¹, Pauline Poydenot¹, Pauline Menager²; ¹Grenoble, FR, ²Bethesda, US
- PS2Group1-028 DOMINANT TRUNCATING MUTATIONS IN THE A-BAND OF TTN ARE A CAUSE OF LIMB-GIRDLE MUSCULAR DYSTROPHY WITH CARDIOMYOPATHY**
Jennifer Roggenbuck, Ana Morales, Ray Hershberger, John Kissel; Columbus, OH, US
- PS2Group1-029 RESCUE OF FOLDING DEFECTIVE ALPHA-SARCOGLYCAN MUTANTS BY MEANS OF PROTEIN FOLDING CORRECTORS**
Chiara Fecchio¹, Marcello Carotti¹, Elisa Bianchini¹, Romeo Betto¹, Roberta Sacchetto², Dorianna Sandona¹; ¹Padova, IT, ²Legnaro (pd), IT
- PS2Group1-030 CLINICAL OUTCOME STUDY FOR DYSFERLINOPATHY: ONE-YEAR FOLLOW-UP**
Meredith James¹, Ursula Moore¹, Anna Mayhew¹, Michelle Eagle¹, Karen Bettinson¹, Elena Pegoraro², Kate Bushby¹; ¹Newcastle Upon Tyne, UK, ²Padova, IT
- PS2Group1-031 HISTOPATHOLOGICAL AND CLINICAL CHARACTERIZATION OF A SPORADIC TNPO3-MUTATED PATIENT**
Alessandra Ruggieri¹, Sara Gibertini¹, Barbara Pasanisi¹, Vincenzo Nigro², Marco Savarese², Maurizio Moggio¹, Corrado Angelini³, Renato Mantegazza¹, Lorenzo Maggi⁴, Lucia Morandi¹, Marina Mora¹; ¹Milan, IT, ²Naples, IT, ³Venezia Lido, IT, ⁴Milano, IT
- PS2Group1-032 ACE-083, A LOCALLY-ACTING MUSCLE AGENT, INCREASES MUSCLE VOLUME IN HEALTHY VOLUNTEERS**
Kenneth Attie¹, Chad Glasser¹, Michael Gartner², Brian Boes², R Pearsall¹, Xiaosha Zhang¹, Jade Sun¹, Brian Vidal¹, Ashley Bellevue¹, Monty Hankin¹, Matthew Sherman¹; ¹Cambridge, MA, US, ²Lincoln,NE, US
- PS2Group1-033 CLINICAL OUTCOME STUDY FOR DYSFERLINOPATHY: CLINICAL DATA FROM BASELINE ASSESSMENTS**
Elizabeth Harris¹, Ursula Moore¹, Catherine Bladen¹, Anna Mayhew¹, Meredith James¹, Karen Bettinson¹, Heather Hilsden¹, Hillarie Windish², The Jain Foundation COS Consortium¹, Kate Bushby¹; ¹Newcastle Upon Tyne, UK, ²Seattle, WA, US

- PS2Group1-034 MOLECULAR PATHOGENESIS OF CAVEOLIN-3-RELATED LIMB-GIRDLE-MUSCULAR-DYSTROPHY**
José Andrés González Coraspe¹, Denisa Hathazi², Hanns Lochmüller³, René Zahedi², Joachim Weis¹, Andreas Roos³; ¹Aachen, DE, ²Dortmund, DE, ³Newcastle, UK
- PS2Group1-035 TRIM32 GENE MUTATIONS DETECTED BY NEXT GENERATION SEQUENCING**
Judith Hudson¹, Eileen Graham¹, Chiara Marini Bettolo¹, Teresinha Evangelista¹, Volker Straub¹, Fiona Norwood², Kate Bushby¹, Rita Barresi¹; ¹Newcastle Upon Tyne, UK, ²London, UK
- PS2Group1-036 MUSCLE INVOLVEMENT IN LIMB GIRDLE MUSCULAR DYSTROPHY WITH GMPBP DEFICIENCY (LGMD2T)**
Sofie Østergaard, Copenhagen, DK
- PS2Group1-037 MUSCLE MRI CAN BE A POWERFUL TOOL TO DIAGNOSE LIMB GIRDLE MUSCULAR DYSTROPHY 2L**
Maria Elena Farrugia¹, Cheryl Longman¹, William Stewart¹, Volker Straub², Richard Petty¹; ¹Glasgow, UK, ²Newcastle Upon Tyne, UK
- PS2Group1-038 MOLECULAR DIAGNOSIS OF BETA-SARCOGLYCANOPATHY: REPORTING TWO NOVEL MUTATIONS IN IRAN**
Marzieh Mojbafan¹, Sirous Zeinali¹, Seyed Hasan Tonekaboni¹, Abdolazim Nejati Zadeh², Yalda Nili Pour¹; ¹Tehran, IR, ²Bandar Abbas, IR
- PS2Group1-040 A NOVEL PATHOGENIC MUTATION IN TPM3 GENE IN A 5 YEARS OLD IRANIAN PATIENT WITH AUTOSOMAL RECESSIVE NEMALINE MYOPATHY-1**
Saeid Morovvati, Yashar Morovvati; Tehran, IR
- PS2Group1-041 MTM1-RELATED MYOPATHY CARRIER FEMALES MANIFEST SIGNIFICANT ASYMMETRIES AND A SPECTRUM OF MUSCLE INVOLVEMENT**
Benjamin Cocanougher¹, Pomi Yun², Lauren Flynn³, Mina Jain², Melissa Waite², Ruhi Vasavada², Jason Wittenbach⁴, Sabine de Chastonay⁵, Sandra Donkervoort⁶, A Foley², Carsten Bonnemann²; ¹Ashburn, US, ²Bethesda, US, ³Bethesda, MD, US, ⁴Ashburn, VA, US, ⁵Torrance, CA, US, ⁶1477, MD, US
- PS2Group1-042 P4HA1 MUTATIONS CAUSE A UNIQUE CONGENITAL DISORDER OF CONNECTIVE TISSUE INVOLVING TENDON, BONE, MUSCLE AND THE EYE**
Sandra Donkervoort¹, Yaqun Zou², Antti Salo³, Aileen Barnes⁴, Ying Hu², A Foley², Elena Makareeva², Meganne Leach², Wendy DiNonno⁴, Jahannaz Dastgir², Ronald Cohn⁵, Sergey Leikin², Joan Marini², Johanna Myllyharju³, Carsten Bonnemann²; ¹1477, MD, US, ²Bethesda, MD, US, ³Oulu, FI, ⁴Newport News, VA, US, ⁵Toronto, ON, CA
- PS2Group1-043 DE NOVO DOMINANT MOSAIC MUTATIONS IN COLLAGEN 6 GENES: UNCOMMON CAUSE OF BETHLEM AND ULLRICH MYOPATHIES THAT MAY BE MISSED BY SANGER SEQUENCING**
Adele D'Amico¹, Fabiana Fattori¹, Francesca Gualandi², Stefania Petrini¹, Valentina Doria¹, Giorgio Tasca¹, Michela Catteruccia¹, Marcello Niceta¹, Marco Tartaglia¹, Alessandra Ferlini², Enrico Bertini¹; ¹Rome, IT, ²Ferrara, IT
- PS2Group1-044 A PROSPECTIVE STUDY OF HISTOLOGY, PHENOTYPES AND GENETICS IN CONGENITAL MYOPATHY PATIENTS ABOVE 5 YEARS OF AGE IN DENMARK**
Nanna Witting¹, Ulla Werlauff², Morten Duno¹, John Vissing¹; ¹Copenhagen, DK, ²Aarhus, DK
- PS2Group1-045 IDENTIFYING AND CHARACTERIZING NOVEL MUTATIONS CAUSING ARTHROGRYPOSIS**
Forough Noohi, Martine Tétreault, Jacek Majewski, Bernard Brais; Montreal, QC, CA
- PS2Group1-046 DESMINOPATHY IN CHILE, TWO FIRST CASES REPORTED**
Jorge Bevilacqua¹, Lidia González-Quereda², Ivonne Zamorano³, Claudia Castiglioni¹, Lorena Acevedo¹, Jorge Díaz¹, María José Rodríguez², Alejandra Trangulao¹, Mario Rivera¹, Pia Gallano²; ¹Santiago, CL, ²Barcelona, ES, ³Puerto Montt, CL
- PS2Group1-047 MULTIPLE DELETIONS IN MITOCHONDRIAL DNA IN MYOFIBRILLAR MYOPATHY AND CENTRONUCLEAR MYOPATHY**
Jochen Schaefer, Heinz Reichmann, Sandra Jackson; Dresden, DE
- PS2Group1-048 THE UK MYOTONIC DYSTROPHY PATIENT REGISTRY**
Nikoletta Nikolenko¹, Libby Wood², Chris Turner³, David Hilton-Jones⁴, Antonio Atalaia¹, Chiara Marini-Bettolo¹, Paul Maddison⁵, Margaret Philips⁶, Mark Roberts⁷, Mark Rogers⁸, Volker Straub¹, Simon Hammans⁹, Hanns Lochmüller¹; ¹Newcastle Upon Tyne, UK, ²Bz, UK, ³London, UK, ⁴Oxford, UK, ⁵Nottingham, UK, ⁶Derby, UK, ⁷Salford, UK, ⁸Cardiff, UK, ⁹Southampton, UK
- PS2Group1-049 18F-FDG-PET STUDY IN PATIENTS WITH MYOTONIC DYSTROPHY TYPE 1 AND 2**
Vidosava Rakocevic Stojanovic, Stojan Peric, Vera Ilic, Aleksandra Parojcic, Jovan Pesovic, Dusanka Savic-Pavicevic, Leposava Brajkovic; Belgrade, RS

- PS2Group1-050 ELEVATED PLASMA LEVELS OF CARDIAC TROPONIN-I PREDICT LEFT VENTRICULAR SYSTOLIC DYSFUNCTION IN PATIENTS WITH MYOTONIC DYSTROPHY TYPE 1: A COHORT FOLLOW-UP STUDY**
Mark Hamilton¹, Yvonne Robb², Helen Gregory³, Sarah Cumming¹, Berit Adam¹, Josephine McGhie¹, Anneli Cooper¹, Jillian Couto¹, Alexis Duncan¹, Monika Rahman¹, Anne McKeown¹, Alison Wilcox¹, Catherine McWilliam⁴, Maria Elena Farrugia¹, Richard Petty¹, Cheryl Longman¹, Iain Findlay¹, Alan Japp², Darren Monckton¹, Martin Denvir²; ¹Glasgow, UK, ²Edinburgh, UK, ³Aberdeen, UK, ⁴Dundee, UK
- PS2Group1-051 ONE-YEAR MRI-FOLLOW-UP IN 45 PATIENTS WITH FACIOSCAPULOHUMERAL MUSCULAR DYSTROPHY**
Grete Andersen¹, Julia Dahlqvist¹, Christoffer Vissing¹, Karen Pedersen², Carsten Thomsen¹, John Vissing¹; ¹Copenhagen, DK, ²Copenhagen E, DK
- PS2Group1-052 OCULOPHARYNGODISTAL MYOPATHY, A CASE REPORT FROM INDIA**
Seena Vengalil, Atchayaram Nalini, Veeramani Preethish-Kumar, Kiran Polavarapu, Narayanappa Gayathri, Anita Mahadevan; Bangalore, IN
- PS2Group1-053 NEUROMUSCULAR ELECTRICAL STIMULATION TRAINING OF THE TIBIALIS ANTERIOR MUSCLE IN FSHD1 PATIENTS**
Jeremy Garcia, Aude Doix, Pauline Lahaut, Véronique Tanant, Manuella Fournier-Mehouas, Serge Colson, Claude Desnuelle, Sabrina Sacconi; Nice, FR
- PS2Group1-054 SPEECH IMPAIRMENT IS COMMON IN EARLY ONSET FACIOSCAPULOHUMERAL DYSTROPHY**
Megan Hodge¹, Jia Feng², Jean Mah³, Cooperative International Neuromuscular Research Group Investigators²; ¹Edmonton, AB, CA, ²Washington, DC, US, ³Calgary, AB, CA
- PS2Group1-057 POMPE DISEASE IN AUSTRIA**
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- PS2Group1-058 PHASE 1 SAFETY, PHARMACOKINETICS, AND EXPLORATORY EFFICACY OF NEOGAA, A NOVEL ENZYME REPLACEMENT THERAPY, IN TREATMENT-NAIVE AND ALGLUCOSIDASE ALFA-TREATED LATE-ONSET POMPE DISEASE PATIENTS**
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- PS2Group1-060 REGULATION OF LIPOPHAGY AND LIPOLYSIS IN LIPID STORAGE MYOPATHIES AND CPT DEFICIENCY**
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- PS2Group1-062 ENERGY DEFICIT IN THE MCARDLE MOUSE MODEL AFFECTS CALCIUM HOMEOSTASIS AND FORCE GENERATION**
Thomas Krag¹, Tomas Pinos², John Vissing¹; ¹Copenhagen, DK, ²Barcelona, ES
- PS2Group1-065 EFFECT OF SENSORY INTEGRATION THERAPY ON FINE MOTOR FUNCTION IN MITOCHONDRIAL MYOPATHY**
Berkan Torpil, Semin Akel, Sedef Şahin; Ankara, TR
- PS2Group1-066 PREVALENCE STUDY OF MUSCLE CHANNELOPATHIES IN ITALY**
Lorenzo Maggi¹, Mauro Lo Monaco², Simona Portaro³, Giovanni Meola¹, Jean Francois Desaphy⁴, Sabrina Lucchiarini¹, Serena Pagliarani¹, Raffaele Dubbioso⁵, Pia Bernasconi¹, Raffaella Brugnoli¹, Paola Imbrici¹, Giacomo Pietro Comi⁶, Renato Mantegazza⁶, Maria Trojano⁴, Adele D'amico⁷, Lucio Santoro⁵, Elena Pegoraro⁸, Luisa Politano⁹, Tiziana Mongini¹⁰, Liliana Vercelli¹¹, Gabriele Siciliano¹², Giulia Ricci¹², Diana Conte-Camerino⁴, Antonio Toscano³, Valeria Sansone¹; ¹Milano, IT, ²Roma, IT, ³Messina, IT, ⁴Bari, IT, ⁵Napoli, IT, ⁶Milan, IT, ⁷Rome, IT, ⁸Padua, IT, ⁹Naples, IT, ¹⁰Turin, IT, ¹¹Torino, IT, ¹²Pisa, IT
- PS2Group1-067 A COHORT OF PEDIATRIC AGE PATIENTS WITH NON-DYSTROPHIC MYOTONIA**
Joel Freitas, Ana Sousa, Teresa Coelho, Manuela Santos; Porto, PT

- PS2Group1-069** **THYROTOXIC PERIODIC PARALYSIS: DOES MUSCLE MEMBRANE DYSFUNCTION UNDERLIE DISEASE PATHOGENESIS?**
Nam-Hee Kim¹, Joong-Yang Cho¹, Kyung Seok Park²; ¹Goyangsi, KR, ²Seongnamsi, KR
- PS2Group1-070** **THE CULLIN 4A, B-DDB1-CEREBLON E3 UBIQUITIN LIGASE COMPLEX MEDIATES THE DEGRADATION OF CLC-1 CHANNELS RESULTING IN MYOTONIA CONGENITA**
Ssu-Ju Fu, Yi-An Chen, Yi-Jheng Peng, Chih-Yung Tang; Taipei, TW
- PS2Group1-071** **ELECTROMYOGRAPHY AND THE RISK OF PNEUMOTHORAX**
Charles Kassardjian¹, Cullen O’Gorman², Eric Sorenson³; ¹Toronto, ON, CA, ²Brisbane, QLD, AU, ³Rochester, MN, US
- PS2Group3-009** **LATE ONSET MYASTHENIA GRAVIS (LOMG): WHEN DOES IT START?**
Nestor Genco, Mariela Bettini, Maria Ines Araoz, Felipe Peralta Calderon, Edgardo Cristiano, Hernan Gonorazky, Marcelo Rugiero; Buenos Aires, AR
- PS2Group3-010** **ROLE OF THYMECTOMY IN PREVENTING MYASTHENIC CRISIS IN GENERALIZED MYASTHENIA GRAVIS**
Gillella Pavan, A Meena, R Kumar, S Jabeen, K Rukmini, Rupam Borgohain; Hyderabad, IN
- PS2Group3-011** **OCULAR VERSUS GENERALIZED MYASTHENIA GRAVIS: 17 YEARS’ EXPERIENCE OF A TURKISH NEUROMUSCULAR CLINIC**
Belgin Mutluay, Aysun Soysal, Fikret Aysal, Ayhan Koksall, Mesude Ozerden, Ayten Dirican, Dilek Atakli, Sevim Baybas; Istanbul, TR
- PS2 Group 3**
- PS2Group3-001** **CLINICAL AND EPIDEMIOLOGICAL FEATURE OF MYASTHENIA GRAVIS IN CHILEAN POPULATION**
Gabriel Cea, David Martinez, Radrigo Salinas, Andres Stuardo; Santiago, CL
- PS2Group3-002** **EXOME SEQUENCING IDENTIFIES TARGETS IN OPHTHALMOPLEGIC MYASTHENIA GRAVIS**
Melissa Nel, Mahjoubah Jalali Sefid Dashti, Junaid Gamiieldien, Jeannine Heckmann; Cape Town, ZA
- PS2Group3-005** **A PHASE II TRIAL TO ASSESS THE EFFICACY, SAFETY AND FEASIBILITY OF 20% SUBCUTANEOUS IMMUNOGLOBULIN IN PATIENTS WITH MYASTHENIA GRAVIS EXACERBATION-SAFETY AND FEASIBILITY**
Zaeem Siddiqi, Derrick Blackmore, Ashley Mallon; Edmonton, AB, CA
- PS2Group3-006** **A PHASE II TRIAL TO ASSESS THE EFFICACY, SAFETY AND FEASIBILITY OF 20% SUBCUTANEOUS IMMUNOGLOBULIN IN PATIENTS WITH MYASTHENIA GRAVIS EXACERBATION-INTERIM ANALYSIS**
Zaeem Siddiqi, Derrick Blackmore, Ashley Mallon, Cecile Phan; Edmonton, AB, CA
- PS2Group3-007** **SUBCUTANEOUS IMMUNOGLOBULIN IN MYASTHENIA GRAVIS: TRIAL DESIGN AND PROGRESS UPDATE**
Zaeem Siddiqi, Derrick Blackmore, Ashley Mallon; Edmonton, AB, CA
- PS2Group3-008** **DISTAL MYASTHENIA GRAVIS: AN UNUSUAL PRESENTATION WITH SLOWLY PROGRESSIVE DISTAL WEAKNESS AND ATROPHY**
Renata Andrade, Alexandre Januario, João Magalhaes; Recife, BR
- PS2Group3-012** **MYCOPHENOLATE MOTETILN MYASTHENIA. EXPERIENCES AN ACADEMIC NEUROMUSCULAR CLINIC**
Ratna Bhavaraju-Sanka¹, Mithila Fadia², Alejandro Tobon², Carlayne Jackson²; ¹San Antonio, TX, US, ²San Antonio, US
- PS2Group3-013** **BULBAR MYASTHENIA PRESENTING AS UNILATERAL PALATAL PALSY**
Veena Vasi, Andrew Thompson, Keith Trimble, Sandya Tirupathi; Belfast, UK
- PS2Group3-014** **PREPURTAL MYASTHENIA - IS THERE A POSTINFECTIOUS ENTITY?**
Sandya Tirupathi, Clare Loughran; Belfast, UK
- PS2Group3-015** **ANTIGEN-SPECIFIC DEPLETION OF PLASMA CELLS IN MYASTHENIA GRAVIS**
Andreas Pelz¹, Qingyu Cheng¹, Adriano Taddeo², Andreas Radbruch¹, Andreas Meisel¹, Falk Hiepe¹, Siegfried Kohler¹; ¹Berlin, DE, ²Bern, CH
- PS2Group3-016** **CLINICAL FOLLOW-UP OF THE PREGNANCY IN MYASTHENIA GRAVIS PATIENTS**
Renata Ducci, Paulo Lorenzoni, Claudia Kay, Lineu Werneck, Rosana Scola; Curitiba, BR
- PS2Group3-017** **CHOLINERGIC TRANSMISSION OF OUTER HAIR CELL IMPAIRED IN MYASTHENIA GRAVIS**
Nam-Hee Kim¹, Joong-Yang Cho¹, Kyung Seok Park²; ¹Goyangsi, KR, ²Seongnamsi, KR
- PS2Group3-018** **RITUXIMAB FOR THE TREATMENT OF PEDIATRIC AUTOIMMUNE NEUROMUSCULAR DISORDERS**
Cam-Tu Emilie Nguyen¹, Elie Haddad², Guy D’Anjou², Jean Mathieu³, Michel Vanasse²; ¹London, ON, CA, ²Montreal, QC, CA, ³Jonquiere, QC, CA

- PS2Group3-019 EPIDEMIOLOGICAL AND CLINICAL CHARACTERISTICS OF ELDERLY MG**
Jiwon Yang¹, Byung-Nam Yoon¹, Seol-Hee Baek², Jung-Joon Sung²; ¹Incheon, KR, ²Seoul, KR
- PS2Group3-020 IMPACT OF REFRACTORY MYASTHENIA GRAVIS ON EMPLOYMENT**
Audra Boscoe¹, Gary Cutter², Haichang Xin²; ¹Lexington, MA, US, ²Birmingham, AL, US
- PS2Group3-021 IMPACT OF REFRACTORY MYASTHENIA GRAVIS ON QUALITY OF LIFE**
Audra Boscoe¹, Gary Cutter², Haichang Xin²; ¹Lexington, MA, US, ²Birmingham, AL, US
- PS2Group3-022 CLINICAL AND ECONOMIC BURDEN OF REFRACTORY GENERALIZED MYASTHENIA GRAVIS IN THE UNITED STATES**
Nicole Engel-Nitz¹, Audra Boscoe², Ryan Wolbeck¹, Jonathan Johnson¹, Nicholas Silvestri³; ¹Eden Prairie, MN, US, ²Lexington, MA, US, ³US
- PS2Group3-023 MYASTHENIA GRAVIS INDUCED BY A MEK INHIBITOR: FIRST CASE REPORTED**
Majed Majed Alabdali¹, Vera Bril², Catherine Maurice²; ¹Toronto, ON, CA, ²Toronto, ON, CA
- PS2Group3-024 INCIDENCE AND PREVALENCE OF MYASTHENIA GRAVIS IN KOREA: A POPULATION-BASED STUDY USING THE NATIONAL HEALTH INSURANCE CLAIMS DATABASE**
Su-yeon Park¹, Jee-Eun Kim¹, Jin Yong Lee¹, Nam Gu Lim², Yoon-Ho Hong¹; ¹Seoul, KR, ²Daejeon, KR
- PS2Group3-025 CLINICAL CHARACTERISTICS OF MUSK-MG IN KOREA: COMPARISON WITH DOUBLE SERONEGATIVE MG**
Kee Hong Park¹, Jung-Joon Sung², Suk-Won Ahn², Byung-Nam Yoon³, Ji-Eun Kim², Seol-Hee Baek², So Hyun Ahn², Yoon-Ho Hong²; ¹Jinju, KR, ²Seoul, KR, ³Incheon, KR
- PS2Group3-026 LATE-ONSET MYASTHENIA GRAVIS: COMPARISON WITH EARLY-ONSET AND VERY LATE-ONSET MYASTHENIA GRAVIS**
Eun Bin Cho¹, Ju-Hong Min², Sujin Lee³, Cindy W Yoon³, Hye-Jin Cho², Jin Myoung Seok², Hye Lim Lee², Byoung Joon Kim²; ¹Changwon, KR, ²Seoul, KR, ³Incheon, KR
- PS2Group3-027 DISEASES OF NEUROMUSCULAR JUNCTION, CLINICAL FEATURES, PATHOPHYSIOLOGY, THERAPY**
Varleen kaur, Patiala, IN
- PS2Group3-028 FETAL AND ADULT ACETYLCHOLINE RECEPTOR FUNCTION IN MYASTHENIC SYNDROMES**
Hakan Cetin, David Beeson, Richard Webster, Angela Vincent; Oxford, UK

- PS2Group3-029 MILD LIMB GIRDLE CONGENITAL MYASTHENIC SYNDROME CAUSED BY HOMOZYGOUS C.686-2A>G MUTATION IN GFPT-1**
Hacer Durmus¹, Serdar Ceylaner², Yesim Parman¹, Feza Deymeer¹, Piraye Oflazer-Serdaroglu¹; ¹Istanbul, TR, ²Ankara, TR
- PS2Group3-030 CONGENITAL MYASTHENIC SYNDROMES: REPORT ON 8 CASES FROM INDIA**
Atchayaram Nalini¹, Veeramani Preethish-Kumar¹, Kiran Polavarapu¹, Seena Vengalil¹, Xin-Ming Shen², Andrew Engel²; ¹Bangalore, IN, ²Minnesota, US
- PS2Group3-031 PERSISTENT DIFFUSE DEEP T WAVE INVERSION: AN ECG MANIFESTATION OF MYASTHENIA GRAVIS IN CRISIS**
Jose Eduardo Duya, Michael Joseph Agbayani; Manila, PH

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- PS2Group5-001-PRESERVATION OF MOTOR NEURON EXCITABILITY DURING THE CUTANEOUS SILENT PERIOD IN AMYOTROPHIC LATERAL SCLEROSIS**
Woogyung Kim, Kee Duk Park; Seoul, KR
- PS2Group5-002 PARKINSON DISEASE IS NOT ASSOCIATED WITH C9ORF72 REPEAT EXPANSIONS AND C19ORF12 MUTATIONS IN IRANIAN PATIENTS**
Afagh Alavi, Elahe Elahi, Maryam Malakouti Nejad, Gholamali Shahidi; Tehran, IR
- PS2Group5-003 MULTI-MODALITY, CERVICAL SPINAL CORD MRI IN ALS: A VALIDATION STUDY**
Barry Bedell¹, Thanh Nguyen¹, Vincent Auclair¹, Julien Cohen-Adad¹, Maxime Descoteaux², Felix Morency², Angela Genge¹, Jim Paskavitz³, Guillaume Gilbert¹, Donald McLaren¹, Alex Zijdenbos¹; ¹Montreal, QC, CA, ²Sherbrooke, QC, CA, ³Cambridge, MA, US
- PS2Group5-004 IMMUNOHISTOCHEMICAL STUDIES OF VALOCIN-CONTAINING PROTEIN IN THE SKIN OF PATIENTS WITH AMYOTROPHIC LATERAL SCLEROSIS**
Seiitsu Ono, Ichihara, JP
- PS2Group5-005 LOWER MOTOR NEURONE DISEASE WITH BICLONAL IGM-PARAPROTEINAEMIA**
Jochen Schaefer, Heike Kostka, Lisa Klingelhofer, Karsten Conrad; Dresden, DE

PS2Group5-007 FUNCTIONAL OUTCOMES IN HEREDITARY SPASTIC PARAPLEGIA: A PROSPECTIVE COHORT STUDY
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PS2Group5-008 GENE THERAPY FOR SPINAL MUSCULAR ATROPHY TYPE 1 SHOWS POTENTIAL TO IMPROVE SURVIVAL AND MOTOR FUNCTIONAL OUTCOMES
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PS2Group7-001-BIOENERGETIC IMPAIRMENT IN CONGENITAL MUSCULAR DYSTROPHY TYPE 1A AND LEIGH SYNDROME
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PS2Group7-002 TRANSFORMING GROWTH FACTOR- β INHIBITS ADIPOGENESIS IN REGENERATING GLYCEROL-INJURED MUSCLE
Mohamed Mahdy, Katsuhiko Warita, Yoshinao Hosaka; Tottori, JP

PS2Group7-003 ROLE OF INTERLEUKIN-6 ON UPREGULATION OF MYOSIN HEAVY CHAIN TYPE IIB MESSENGER RNA IN MOUSE MYOCYTES
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PS2Group7-004 UPREGULATION OF MYOSIN HEAVY CHAIN TYPE I AND INTERLEUKIN-6 MESSENGER RNA LEVELS BY EXOGENOUS APPLICATION OF CALCINEURIN ACTIVATORS IN C2C12 SKELETAL MYOCYTES
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PS2Group7-005 CYTOPLASMIC NOTCH AND MEMBRANAL BETA-CATENIN LINK CELL FATE CHOICE TO EMT DURING MYOGENESIS
Christophe Marcelle, Lyon, FR

PS2Group7-006 FUNCTIONS OF THE SIL1-BIP CHAPERONE SYSTEM IN MAINTAINING MUSCLE FIBER INTEGRITY
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PS2Group7-007 ANTI-INFLAMMATORY EFFECT OF ORAL-FORMULATED TACROLIMUS IN EXPERIMENTAL AUTOIMMUNE ENCEPHALOMYELITIS (EAE) MICE
Suk-Won Ahn, Myung-Jin Kim, Dae-Woong Kang, Chang-Seop Kim, Jung-Joon Sung, Yoon-Ho Hong; Seoul, KR

PS2Group7-008 MUSCLE ADAPTATIONS AND MOTOR DYSFUNCTION IN PERIPHERAL NEURODEGENERATIVE DISEASES MODELS
Ole Nielsen, Jeppe Bayley, Martin Broch-Lips, Anders Riisager, Thomas Pedersen; Aarhus, DK

PS2Group7-009 NORMAL MITOCHONDRIAL TRANSPLANTATION MAY BE USEFUL FOR THE TREATMENT OF MITOCHONDRIA-ASSOCIATED NEUROMUSCULAR DISEASES: EVIDENCES IN VITRO
Xianpeng Jiang, Robert Elliott, Jonathan Head; Baton Rouge, LA, US