

ALL CONFERENCE SESSION TIMES LISTED ARE IN CENTRAL TIME  
\*SPEAKERS LISTED IN ORDER OF PRESENTATION

## Saturday, March 15, 2025

|                 |  |                               |
|-----------------|--|-------------------------------|
| 12:00 – 6:00 PM | Registration Open for Care Center Directors        | Geppetto Foyer,<br>Tower Side |
| 12:00 – 6:00 PM | Registration Open for Large Exhibit Booth Sponsors | Trinity Exhibit Hall          |
| 5:00 – 7:00 PM  | MDA Care Center Network Happy Hour                 | Gossip Bar                    |

## Sunday, March 16, 2025

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|-------------------|---|-------------------------------|
| 7:00 AM – 7:00 PM | Registration Open   | Geppetto Foyer,<br>Tower Side |
| 7:30 AM – 3:30 PM | MDA Care Center Directors Meeting (Invitation Only)   | Chantilly East                |
| 7:30 AM – 3:30 PM | Allied Health Workshop<br><i>Shifting Care for Shifting Needs: Coordinating Care and Increasing Quality of Life via Assistive Devices</i> | Chantilly West                |
| 1:00 – 5:00 PM    | Trainee Networking Session  | Wedgewood<br>Ballroom         |
| 2:00 – 5:00 PM    | Neuromuscular Advocacy Collaborative Meeting (Invitation Only)  | Monet                         |
| 6:00 – 8:00 PM    | Welcome Reception & Posters in Exhibit Hall   | Trinity Exhibit Hall          |

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Monday, March 17, 2025

|                     |   |                            |
|---------------------|---|----------------------------|
| 6:30 AM – 6:00 PM   | Registration Open   | Geppetto Foyer, Tower Side |
| 7:00 – 8:00 AM      | <b>Industry Forum Breakfast presented by Amicus</b><br><i>Moving Forward Together: Discovering a Treatment Approach for Late-Onset Pompe Disease</i>  | Grand ABC                  |
|                     | <b>Industry Forum Breakfast presented by argenx</b><br><i>Developments in CIDP: Pathophysiology, Disease Burden, and Clinical Updates from the ADHERE/ADHERE+ Trials Investigating FcRn Inhibition</i>          | Grand DE                   |
|                     | <b>Industry Forum Breakfast presented by Avidity Biosciences</b><br><i>Advancing RNA Therapeutics: Exploring Antibody Oligonucleotide Conjugates (AOCs) for Rare Neuromuscular Diseases</i>                     | Chantilly East             |
|                     | <b>Industry Forum Breakfast presented by ITF Therapeutics</b><br><i>Learn about the Role of an HDAC inhibitor in the Treatment of Duchenne Muscular Dystrophy</i>   | Chantilly West             |
| 8:30 – 9:30 AM      | <b>Opening Remarks</b><br>Donald S. Wood, PhD<br>President and CEO of the Muscular Dystrophy Association  | Trinity Ballroom           |
|                     | <b>MDA Legacy Award</b><br>Katherine Mathews, MD  |                            |
|                     | <b>MDA Community Impact in Research Award</b><br>Donavon Decker   |                            |
| 9:30 – 10:00 AM     | Networking Break in Exhibit Hall  | Trinity Exhibit Hall       |
| 10:00 AM – 12:00 PM | <b>General Session</b><br><br><i>MDA Research: Exploring the Past, Innovating the Present, Shaping the Future</i><br>Sharon Hesterlee, PhD  | Trinity Ballroom           |
|                     | <i>The Year Ahead in MDA's Public Policy and Advocacy</i><br>Paul Melmeyer  |                            |
|                     | Keynote Address<br>Robert Califf, MD, MACC  |                            |
|                     | Keynote Panel: Looking to the Future of Neuromuscular Research  |                            |
| 12:00 – 1:30 PM     | <b>Industry Forum Lunch presented by Biogen</b><br><i>Advancing Care for Friedreich Ataxia: Clinical Insights and Patient Cases with the First Approved Treatment</i>   | Chantilly West             |
|                     | <b>Industry Forum Lunch presented by Novartis</b><br><i>Long-Term Impact of Gene Therapy: Lessons from the SMA Community</i>  | Grand ABC                  |
|                     | <b>Industry Forum Lunch presented by PPD, part of Thermo Fisher Scientific</b><br><i>The Evolution of Neuromuscular Trials: Leveraging the Challenges and Successes of the Past to Forge a Promising Future</i> | Grand DE                   |
|                     | <b>Industry Forum Lunch presented by Sarepta</b><br><i>From Assessment to Action: A Holistic Approach to Optimizing Neuromuscular Disease Care</i>  | Chantilly East             |
|                     | <b>Industry Forum Lunch presented by UCB</b><br><i>The Diagnostic Dilemma: Genetic Testing vs Muscle Biopsy in Mitochondrial Myopathies</i>   | Wedgewood Ballroom         |

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Monday, March 17, 2025 cont.

Allied Health Track

|                |  |           |
|----------------|--|-----------|
| 2:00 – 3:30 PM | <p><b>The Latest in Pulmonary and Cardiac Management in Neuromuscular Disease</b><br/> <i>(CE Accredited)</i></p> <p><i>Applying Cardiac Guidelines to the Neuromuscular Patient</i><br/>         Co-Chair – Elizabeth McNally, MD, PhD</p> <p><i>Managing Cardiac Care Before, During and After Gene Therapy</i><br/>         Beth Kaufman, MD</p> <p><i>Natural History of Pulmonary Care: How Do Lifetime Events Impact Breathing Function?</i><br/>         Co-Chair - Bethany Lussier, MD</p> <p><i>Pivotal Decision Points: Case Studies to Guide Decisions in Pulmonary Care</i><br/>         Jeanette Brown, MD, PhD</p> | Cortez AB |
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Disease Mechanism Track

|                |   |              |
|----------------|---|--------------|
| 2:00 – 3:30 PM | <p><b>Neural Regeneration – The Next Frontier</b><br/>         Chair – Kuldip Dave, PhD</p> <p><i>Is ALS a Recoverable Disease? Role of Neural Repair in the CNS and PNS</i><br/>         Brett Morrison, MD, PhD</p> <p><i>Neural Regeneration from the SCI Perspective</i><br/>         Binhai Zheng, PhD</p> <p><i>Glial Cells and Neural Plasticity: Partners in Neuroprotection and Repair</i><br/>         Isobel Scarisbrick, PhD</p> <p><i>A Novel Approach to Slowing ALS Disease Progression Using Non-Invasive Multi-site DCS</i><br/>         Nader Yaghoubi, MD, PhD</p> | Coronado BCD |
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Care Management Track

|                |   |           |
|----------------|---|-----------|
| 2:00 – 3:30 PM | <p><b>Learning Across Disciplines: Applying Proven Models to Neuromuscular Care</b><br/> <i>(CE Accredited)</i></p> <p><i>What Can We Learn or Model from Other Fields</i><br/>         Chair – Diana Castro, MD</p> <p><i>NMD Care: What Can We Learn from Other Fields. Successful Transition from Pediatric to Adult Congenital Heart Disease: It Takes More Than a Village</i><br/>         Kan Hor, MD</p> | Cortez CD |
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|  | <p><i>Optimizing Comprehensive Care in Neuromuscular Disease - Learning from Other Models</i><br/>         Oscar 'Hank' Mayer, MD</p>  |                      |
| <b>Monday, March 17, 2025, cont.</b>   |  |                      |
| <b>Diagnostic Rolodex Track</b>  |  |                      |
| 2:00 - 3:30 PM   | <p><b>Myasthenia Gravis: Presentation and Diagnosis; Current Therapies; and Current &amp; Future Research</b><br/> <i>(CE Accredited)</i></p> <p><i>Myasthenia Gravis: Presentation and Diagnosis; Current Therapies; and Current and Future Research</i><br/>         Chair – Sarah Heintzman, MS, APRN-CNP, FNP-C, CCRC</p> <p><i>Myasthenia Gravis: Presentation and Diagnosis; Current Therapies; and Current and Future Research</i><br/>         Mary Petrusis, MD</p> <p><i>Myasthenia Gravis Current Therapies</i><br/>         Neelam Goyal, MD</p> <p><i>Myasthenia Gravis: Current and Future Research</i><br/>         Ali Habib, MD</p>   | Coronado A           |
| <b>Catalyzing Innovation: Strategies for Streamlining Drug Development Track</b> |  |                      |
| 2:00 - 3:30 PM   | <p><b>Innovative Research Programs: Filling Gaps in Therapeutic Development</b><br/>         Chair – Laury Mignon, PhD</p> <p><i>Project PaLaDIn – Enhancing the Patient Voice in NMD</i><br/>         David Allison</p> <p><i>Treat NMD Advisory Committee for Therapeutics</i><br/>         Lindsay Alfano, PT, DPT, PCS</p> <p><i>IAMRARE Research Program: Advancing Patient-Centered Research</i><br/>         Janine Lewis</p> <p><i>MDA Kickstart Program: Advancing Gene Therapy for Ultra-Rare Neuromuscular Diseases</i><br/>         Angela Lek, PhD</p> <p><i>AFM-Telethon, A Non-Profit Association Developing Innovation for Therapeutic Applications</i><br/>         Jean-François Briand</p> <p><i>Next Generation Trial Design Today: Collaborative Data Science to Unlock Better Drugs to Patients Sooner</i><br/>         Susan Ward</p> | De Soto AB           |
| 3:30 – 3:55 PM   | Networking Break in Exhibit Hall   | Trinity Exhibit Hall |

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Monday, March 17, 2025, cont.

Allied Health Track

|                       |   |                  |
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| <p>4:00 – 6:00 PM</p> | <p><b>Optimizing Nutrition and Physical Therapy in Neuromuscular Disease: Practical Applications for Patient Care</b><br/> <i>(CE Accredited)</i><br/>         Chair – Umme Salma Vahanvaty, MS, RD, CSP</p> <p><i>Nutrition Focused Physical Exam in Patients with Neuromuscular Disease</i><br/>         Alicia Gilmore, MS, RD, CSO, LD</p> <p><i>Physical Therapy in Neuromuscular Disorders: Evidence-Based Exercise &amp; Care Recommendations</i><br/>         Constance de Monts, PT, DPT</p> <p><i>Current Trends and Challenges in Pediatric Obesity</i><br/>         Ahlee Kim, MD</p> <p><i>Constipation in Patients with Neuromuscular Disease</i><br/>         Shruti Nabar, MD</p> | <p>Cortez AB</p> |
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Disease Mechanism Track

|                       |   |                     |
|-----------------------|---|---------------------|
| <p>4:00 – 6:00 PM</p> | <p><b>Muscle Regeneration in Diseased Muscles: How Close Are We?</b></p> <p><i>Ex Vivo Gene Therapy for DMD: Safety, Efficacy and Affordability</i><br/>         Giulio Cossu, MD, FMedSci, FEAS</p> <p><i>First Clinical Trial of Myopaxon, An Allogeneic Off-The-Shelf Ipsc-Derived Myogenic Progenitor Cell Product for DMD</i><br/>         Chair – Rita Perlingeiro, PhD</p> <p><i>Primary Human Muscle Stem Cells as ATMP In Treatment of Muscular Dystrophies</i><br/>         Simone Spuler, MD</p> <p><i>Cell-Based Approaches for Treatment of Muscle Diseases: Lessons from Gene Therapy</i><br/>         Carl Morris, PhD</p> <p><i>Improving Regeneration in Aged and Dystrophic Muscle: The Role of Senescent Cells</i><br/>         Pura Muñoz-Cánoves, PhD</p> <p><i>Imaging Cell and Gene Therapies</i><br/>         Glenn Walter, PhD</p> | <p>Coronado BCD</p> |
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Monday, March 17, 2025, cont.

Care Management Track

|                |  |           |
|----------------|--|-----------|
| 4:00 – 6:00 PM | <p><b>Developing a Sustainable Multidisciplinary Team Model</b><br/><i>(CE Accredited)</i></p> <p>Chair – Erika Finanger, MD, MS</p> <p>Kaitlin Batley, MD</p> <p>Carolyn Kelley</p> <p>Aravindhan Veerapandiyan, MD</p> | Cortez CD |
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Diagnostic Rolodex Track

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|----------------|---|------------|
| 4:00 – 6:00 PM | <p><b>Myositis: Best Practices &amp; Updates</b></p> <p><i>Classification of Idiopathic Inflammatory Myopathies (IIM)</i><br/>Chair – Tahseen Mozaffar, MD</p> <p><i>Role of Pathology in IIM</i><br/>Suur Biliciler, MD</p> <p><i>Role of Imaging in IIM</i><br/>Namita Goyal, MD</p> <p><i>Inclusion Body Myositis: What is New?</i><br/>Bhaskar Roy, MBBS, MHS</p> | Coronado A |
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Catalyzing Innovation: Strategies for Streamlining Drug Development Track

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|-----------------|---|----------------------|
| 4:00 – 5:00 PM  | <p><b>Lessons Learned from MOVR Data</b></p> <p>Chair – Andre Paredes, PhD</p> <p>Aravindhan Veerapandiyan, MD</p> <p><i>A Reflection on Motor Neuron Disease Based on the MOVR Database</i><br/>Bhaskar Roy, MD</p> <p><i>Current State of Cardiac Therapies in Duchenne Muscular Dystrophy</i><br/>DeWayne Townsend, DVM, PhD</p> | De Soto AB           |
| 5:00 – 6:00 PM  | <p><b>MOVR Workshop (Invitation Only)</b></p>   | De Soto AB           |
| 6:00 – 8:00 PM  | <p>Poster &amp; Networking Reception</p>  | Trinity Exhibit Hall |
| 8:00 – 10:30 PM | <p>MDA 75<sup>th</sup> Celebration</p>  | Trinity Ballroom     |

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Tuesday, March 18, 2025

|                                |  |                               |
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| 6:30 AM – 4:00 PM              | Registration Open  | Geppetto Foyer,<br>Tower Side |
| 7:00 – 8:00 AM                 | <b>Industry Forum Breakfast presented by Astellas</b><br><i>Where Are We with Gene Therapies for Rare Neuromuscular Disorders?</i>   | Chantilly West                |
|                                | <b>Industry Forum Breakfast presented by Biogen</b><br><i>Pioneering Pathways: Evolving Use of Neurofilament in Neurodegenerative Disease</i>  | Grand DE                      |
|                                | <b>Industry Forum Breakfast presented by Catalyst</b><br><i>Navigating DMD Treatment: Integrating Data with Patient Stories</i>  | Grand ABC                     |
|                                | <b>Industry Forum Breakfast presented by The France Foundation</b><br><i>New Frontiers in the Treatment of DMD: Across the Age Spectrum</i>  | Chantilly East                |
| <b>ALS/FTD Track</b>           |  |                               |
| 8:30 – 9:30 AM                 | <b>Leveraging Genetics for ALS Therapeutics</b><br><br><i>Clarifying the ALS Mutome</i><br>Chair – Matt Harms, MD<br><br><i>Harnessing Genetic Data to Guide Drug Repurposing in Motor Neuron Diseases</i><br>Sara Saez Atienzar, MSc, PhD<br><br><i>Progress Update for ALS Compute: A Central Repository for Harmonized WGS Data</i><br>John Landers, PhD<br><br><i>Role of Mutant SPTLC1 in Juvenile ALS</i><br>Devesh Pant, PhD  | Coronado A                    |
| <b>Disease Mechanism Track</b> |  |                               |
| 8:30 – 9:30 AM                 | <b>Approaches to Delivering/Restoration of Large-Sized Genes</b><br>Chair – Jeffrey Chamberlain, PhD<br><br><i>SIMPLI-GT: a Novel Gene Therapy Method to Deliver and Express Large Proteins</i><br>Hichem Tassaout, PhD, PharmD<br><br><i>Recent Advances in Base Editing and Nonviral Delivery for DMD Correction</i><br>Yu-Chung (Andy) Pien<br><br><i>Engineered Lentiviral Vectors for Skeletal Muscle Targeting</i><br>Doug Millay, PhD<br><br><i>The Potential of tRNA Therapeutics to Restore Full-Length Proteins for Muscle Stop Codon Disease</i><br>Stephen Eichhorn, PhD | Coronado BCD                  |

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Tuesday, March 18, 2025

Care Management Track

|                       |  |                  |
|-----------------------|--|------------------|
| <p>8:30 – 9:30 AM</p> | <p><b>Navigating Transition: Strategies for Care Across Settings and Life Stages<br/>         (CE Accredited)</b></p> <p>Chair – Yaacov Anziska, MD</p> <p><i>Going from Pediatric to Adult Health Care When Living with Muscular Dystrophy</i><br/>         Tyus Hill</p> <p><i>Treating New Adults with Chronic Diseases</i><br/>         Christina Trout, RN, MSN</p> <p><i>Supporting Patients in the Transition to Adulthood</i><br/>         Jodi Wolff, PhD, MSSW</p> <p><i>Transition to Adult Care: More Important Now Than Ever</i><br/>         Lauren Elman, MD</p> <p>The Role of Adult Multidisciplinary Clinics in Transitioning Neuromuscular Care<br/>         Bakri Elsheikh, MBBS, FRCP, FAAN</p> | <p>Cortez CD</p> |
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Genetic Medicine Track

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| <p>8:30 – 9:30 AM</p> | <p><b>Gene Therapy Updates - Where Are We Today?<br/>         (CE Accredited)</b></p> <p><i>Genetic Medicine: Gene Therapy Updates</i><br/>         Chair – John Brandsema, MD</p> <p><i>Gene Therapy in the Clinic in 2024: Common Themes Across Neuromuscular Disorders<br/>         and Other Therapeutic Areas</i><br/>         Susan Matesanz, MD</p> <p><i>Emerging Therapies: The Research Pipeline and the Potential for Two Gene Therapies<br/>         for the Same Disease</i><br/>         Diana Bharucha-Goebel, MD</p> <p><i>Treating an Adult with Gene Therapy: They Are Not “Big Children”!</i><br/>         Emma Ciafaloni, MD, FAAN</p> | <p>Cortez AB</p> |
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Tuesday, March 18, 2025

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| <p>8:30 – 10:30 AM</p>      | <p><b>Pompe Disease: Mechanisms, Therapeutic Advances, Advocacy, and Integrated Care Approaches (<i>Session in Partnership with TREAT-NMD</i>) (CE Accredited)</b></p> <p><i>Historical Learnings from Pre-ERT to Current ERT Landscape in Pompe Disease</i><br/>       Chair - Priya Kishnani, MD</p> <p><i>Understanding Molecular Challenges and Opportunities in Pompe Disease</i><br/>       Catherine Rehder, PhD</p> <p><i>Pompe Disease: Next Generation Therapies, and New Treatments on the Horizon</i><br/>       Mark Roberts</p> <p><i>Role of Biomarkers in Disease Monitoring</i><br/>       Ferdinand Knieling, PhD</p> <p><i>Multidisciplinary Approach and Rehab Services for Pompe Disease</i><br/>       Tracy Boggs, PT, MPT, NCS</p> <p><i>Diagnostic Odyssey and Living with Pompe: A Perspective from Pompe Patients</i><br/>       Tiffany House, JD</p> <p><i>Diagnostic Odyssey and Living with Pompe: A Perspective from Pompe Patients</i><br/>       Amanda Joost</p> | <p>De Soto AB</p> |
| <p><b>ALS/FTD Track</b></p> |   |                   |
| <p>9:30 – 10:30 AM</p>      | <p><b>Role of Glial Cells in Disease Pathogenesis</b><br/>       Chair – Rita Sattler, PhD</p> <p><i>Identification of a Neuroprotective Microglial State in C9ORF72 ALS/FTD</i><br/>       Justin Ichida, PhD</p> <p><i>How Astrocytes Harm Human Motor Units In ALS: a Dual Mechanism of Action</i><br/>       Ludo Van den Bosch, PhD</p> <p><i>Activation of Inflammatory Pathways in ALS</i><br/>       Allison Ebert, PhD</p> <p><i>Contribution of Astrocytic Sparc1 to Cortical Synaptic Dysfunction in C9ORF72-FTD/ALS</i><br/>       Robert Culibrk, PhD</p>  | <p>Coronado A</p> |

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Tuesday, March 18, 2025 cont.

Disease Mechanism Track

|                 |   |              |
|-----------------|---|--------------|
| 9:30 – 10:30 AM | <p><b>Targeting Signaling Pathways to Treat Muscular Dystrophies</b></p> <p><i>Preclinical Studies on the Modulation of Signaling Pathways as a Therapeutic Approach for Muscular Dystrophies</i><br/>Chair – Matthew Alexander, PhD</p> <p><i>Exploring a pathogenic role for Fibroblast Growth Factor 23 in dystrophin-deficient cardiac remodeling</i><br/>Steve Welc, PhD</p> <p><i>Development of Anti-LTBP4 Stabilizing Antibodies for the Treatment of Muscular Dystrophy</i><br/>Alexis Demonbreun, PhD</p> <p><i>CRISPR-Engineered Mutations and MERFISH Single Cell Spatial Transcriptomics Reveal Key Processes of FSHD Pathogenesis</i><br/>Kyoko Yokomori, PhD, DVM</p> <p><i>FGF21-Mediated Muscle/Bone Interactions in Duchenne Muscular Dystrophy</i><br/>Hongshuai Li, MD, PhD</p> | Coronado BCD |
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Care Management Track

|                 |   |           |
|-----------------|---|-----------|
| 9:30 – 10:30 AM | <p><b>Building Bridges: Community Partnerships for Comprehensive Multidisciplinary Care (CE Accredited)</b></p> <p><i>Community Collaboration: Collaborating with Providers in the Community to Provide Multidisciplinary Care and Bridging Unmet Needs</i><br/>Chair – Susan Apkon, MD</p> <p>Mark Terrelonge, MD, MPH</p> <p>Elizabeth 'Lynne' Wood, MD</p> | Cortez CD |
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Genetic Medicine Track

|                  |  |                      |
|------------------|--|----------------------|
| 9:30 – 10:30 AM  | <p><b>Experiences in DMD – Panel Discussion (CE Accredited)</b></p> <p>Chair – Barry J. Byrne, MD, PhD</p> <p>Julie Parsons, MD</p> <p>Kevin Flanigan, MD</p> <p>Carmen Leon Astudillo, MD</p> <p>Mindy Cameron</p> <p>Aravindhan Veerapandian, MD</p> | Cortez AB            |
| 10:30 – 10:55 AM | Networking Break in Exhibit Hall   | Trinity Exhibit Hall |

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Tuesday, March 18, 2025 cont.

Genetic Medicine Track

|                         |   |                  |
|-------------------------|---|------------------|
| <p>11:00 – 12:00 PM</p> | <p><b>New Genetic Technologies in Diagnosis and Treatment of Neuromuscular Disease</b><br/> <i>(CE Accredited)</i></p> <p><i>New Genetic Technologies in Diagnosis and Treatment of Neuromuscular Disease</i><br/>         Chair – Kelly Minks, MS CGC</p> <p>Jordan Bontrager, MS CGC</p> <p><i>It's Obviously FSHD! Or is it?</i><br/>         Natalie Katz, MD, PhD</p> <p><i>Long-Read and RNA Sequencing: Case Examples</i><br/>         Alayne Meyer, MS, CGC</p> | <p>Cortez AB</p> |
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Lab to Life Track

|                         |  |                                  |
|-------------------------|--|----------------------------------|
| <p>11:00 – 12:00 PM</p> | <p><b>Non-Muscle Organ Systems in Myotonic Dystrophy</b></p> <p><i>A Model of DM GI Dysfunction in Mice</i><br/>         Chair – Thomas Cooper, MD</p> <p><i>Altered Drug Metabolism and Increased Susceptibility to Fatty Liver Disease in Myotonic Dystrophy</i><br/>         Auinash Kalsotra, PhD</p> <p>Pradeep Mammen, MD</p> <p><i>Mechanisms of CNS Dysfunction in Myotonic Dystrophy</i><br/>         Eric Wang, PhD</p>                      | <p>Coronado<br/>         BCD</p> |
| <p>11:00 – 12:00 PM</p> | <p><b>Scientific Progress and Community Insights into Rare Myopathies</b></p> <p><i>Biology and approaches to treatment of ADSS1 myopathy</i><br/>         Chair – Alan Beggs, PhD</p> <p><i>Patient voice: ADSS1 Myopathy</i><br/>         Priyanka Kakkar &amp; Naveen Baweja</p> <p><i>Biology and approaches to treatment of GNE myopathy</i><br/>         Noah Weisleder, PhD</p> <p><i>Patient voice: GNE Myopathy</i><br/>         Al Stork</p> | <p>Cortez CD</p>                 |

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Tuesday, March 18, 2025 cont.

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| 11:00 – 12:00 PM | <p><b>Rational Disease-Modifying Therapies for Charcot-Marie-Tooth Disease and Related Neuropathies – Are We There Yet?</b></p> <p>Chair – Joshua Burns, PhD</p> <p><i>Genetic Discoveries and Clinical Trials in CMT</i><br/> Michael Shy, MD, FAAN, FANA</p> <p><i>Treatment Pipeline for CMT</i><br/> Mario Saporta, MD, PhD, MBA</p> <p><i>Challenges and Opportunities in Pediatric Clinical Trials</i><br/> Richard Finkel, MD</p> <p><i>Clinical Trial Readiness for TRPV4 Channelopathies</i><br/> Charlotte Sumner, MD</p>  | Coronado A         |
|                  | <p><b>Pre-Clinical and Clinical Efforts in Oculopharyngeal Muscular Dystrophy (OPMD)</b></p> <p>Chair – Mathew Wicklund, MD</p> <p><i>History of the Study of OPMD as it Sets the Stage for New Therapeutic Trials</i><br/> Co-chair – Bernard Brais, MDCM, PhD</p> <p><i>Drug Development Approaches to OPMD</i><br/> Guy Rouleau, PhD</p> <p><i>Advances in Non-Instrumental Dysphagia Assessment for Oculopharyngeal Muscular Dystrophy</i><br/> Claudia Côté, PhD</p> <p><i>Interim Clinical Data Summary: A Phase 1b/2a Open-Label, Dose Escalation Study to Evaluate the Safety and Clinical Activity of Intramuscular Doses of an AAV9-Based Gene Therapy (BB-301) Administered to Subjects with Oculopharyngeal Muscular Dystrophy (OPMD) with Dysphagia</i><br/> Jerel Banks, PhD</p> | De Soto AB         |
| 12:00 – 1:30 PM  | <p><b>Industry Forum Lunch presented by argenx</b><br/> <i>Reviewing the Evolving Treatment Approaches in Generalized Myasthenia Gravis</i></p>  | Chantilly West     |
|                  | <p><b>Industry Forum Lunch presented by Dyne Therapeutics</b><br/> <i>Harnessing the Force™ Platform to Advance Targeted Therapies for Neuromuscular Diseases</i></p>  | Grand DE           |
|                  | <p><b>Industry Forum Lunch presented by Edgewise Therapeutics</b><br/> <i>Spotlight on Becker Muscular Dystrophy: Understanding the Lived Experience of Becker and Clinical Advancements with a Novel Agent</i></p>  | Wedgewood Ballroom |
|                  | <p><b>Industry Forum Lunch presented by Genentech</b><br/> <i>Exploring Oral SMA Treatment: Discover the Latest Developments for Evrysdi</i></p>   | Grand ABC          |
|                  | <p><b>Industry Forum Lunch presented by Scholar Rock</b><br/> <i>Muscle, Myostatin, and More: Evolving Needs and Approaches in SMA</i></p>   | Chantilly East     |

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Tuesday, March 18, 2025 cont.

ALS / FTD Track

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| 2:00 – 3:00 PM | <p><b>Exploring ALS Innovations: Motor Speech, BCIs, and AI Solutions</b></p> <p><i>Where AI Can (and Can't) Help with ALS</i><br/>         Chair – Ernest Fraenkel, PhD</p> <p><i>Motor Speech Analysis and Listener Effort as ALS Outcome Measurers</i><br/>         James Berry, MD, PhD</p> <p><i>BCIs for Restoring Speech and Communication</i><br/>         Daniel Rubin, MD, PhD</p> | Coronado A |
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Disease Mechanism Track

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| 2:00 – 3:00 PM | <p><b>Fibroadipogenic Progenitors - Mechanisms and Therapeutic Opportunities</b></p> <p><i>Role of FAPs in Muscle Diseases And Its Modulation As A Therapeutic Strategy</i><br/>         Fabio Rossi, MD</p> <p><i>Uncovering the Origins of Genuine Senescent Cells to Mitigate Skeletal Muscle Damage: Are Fibroadipogenic Progenitors the Primary Source?</i><br/>         Michael Wosczyzna, PhD</p> <p><i>Fibroadipogenic Progenitors in Muscular Dystrophy: Contribution to Pathology and Therapeutic Potential</i><br/>         Chair – Marshall Hogarth, PhD</p> | Coronado BCD |
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Care Management Track

|                |  |           |
|----------------|--|-----------|
| 2:00 – 4:00 PM | <p><b>Preparing for Tomorrow's Leaders &amp; Fostering Diversity Among Providers (CE Accredited)</b></p> <p>Chair - Terry Heiman-Patterson MD<br/>         Co-Chair - Urvi Desai, MBBS, MS, MD, FAAN<br/>         Ericka Greene, MD, MACM<br/>         Vovanti Jones, MD</p> | Cortez CD |
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Genetic Medicine Track

|                |  |           |
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| 2:00 – 4:00 PM | <p><b>Gene Therapy for Intermediate/Experienced Sites (CE Accredited)</b></p> <p>Chair – Natalie Goedeker, DNP, CPNP<br/>         Craig McDonald, MD, PhD<br/>         Katherine Mathews, MD</p> | Cortez AB |
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 \*SPEAKERS LISTED IN ORDER OF PRESENTATION

Tuesday, March 18, 2025 cont.

ALS / FTD Track

|                       |   |                   |
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| <p>3:00 – 4:00 PM</p> | <p><b>Advances in Biomarker Exploration</b></p> <p>Chair – Tania Gendron, PhD</p> <p><i>HDGFL2, a Biomarker for TDP-43 Proteinopathies</i><br/>       Leonard Petrucelli, MD, PhD</p> <p><i>Multi-Domain Smartphone Assessments for Neurological and Neuromuscular Disorders: Progress in the FTD-ALS Spectrum</i><br/>       Adam Staffaroni, PhD</p> <p><i>Plasma Extracellular Vesicle Tau Isoform Ratios and TDP-43 Inform about Molecular Pathology in Frontotemporal Dementia and ALS</i><br/>       Hanna Huber, PhD</p> <p><i>Large-Scale Cerebrospinal Fluid Proteomics Identifies Molecular Signatures of C9orf72 Frontotemporal Dementia</i><br/>       Rowan Saloner, PhD</p> | <p>Coronado A</p> |
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Disease Mechanism Track

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| <p>3:00 – 4:00 PM</p> | <p><b>Genetic Therapies in the Pipeline</b></p> <p><i>AAV.U7snRNA as a Platform to Treat Neuromuscular Disorders</i><br/>       Chair – Nicolas Wein, PhD</p> <p>Melissa Spencer, PhD</p> <p><i>Gene Editing Correction of DMD Mutations in Human Cells and Humanized Mice</i><br/>       Mateusz Durbacz, MSc</p> <p><i>Advancing Duchenne Muscular Dystrophy Research: Precision Models and Therapeutic Strategies</i><br/>       Zhenya Ivakine, PhD</p> | <p>Coronado BCD</p>         |
| <p>4:00 – 4:25 PM</p> | <p>Networking Break in Exhibit Hall</p>   | <p>Trinity Exhibit Hall</p> |

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Tuesday, March 18, 2025 cont.

ALS / FTD Track

|                |   |            |
|----------------|---|------------|
| 4:30 – 6:00 PM | <p><b>Newest Therapeutic Target Discoveries</b></p> <p>Chair – Nicholas Maragakis, MD</p> <p><i>SARM1 Inhibition Confers Neuroprotection in Preclinical Models of Neurological Injury and Disease</i><br/>         Shilpa Sambashivan, PhD</p> <p><i>Validation of an AAV Gene Therapy for Amyotrophic Lateral Sclerosis</i><br/>         Philip Wong, PhD</p> <p><i>Applying Advanced AAV Strategies for Sporadic ALS</i><br/>         Defne Amado, MD, PhD</p> <p><i>Repeat Associated Non-AUG (RAN) Proteins as Therapeutic Targets in C9orf72 ALS and Other Neurodegenerative Diseases</i><br/>         Laura Ranum, MD, PhD</p> <p><i>Therapeutic Development in Neurodegenerative Diseases with TDP-43 Proteinopathy</i><br/>         Clotilde Lagier-Tourenne, MD, PhD</p> | Coronado A |
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Disease Mechanism Track

|                |   |              |
|----------------|---|--------------|
| 4:30 – 6:00 PM | <p><b>Genetic Modifiers – New Targets for Muscular Dystrophies</b></p> <p><i>Alpha7 Integrin Enhancing Small Molecule for the Treatment of Duchenne Muscular Dystrophy</i><br/>         Chair – Dean Burkin, PhD</p> <p><i>Genetic Modifiers in Congenital Myopathy</i><br/>         Vandana Gupta, PhD</p> <p><i>Modifiers of LMNA-Associated Muscular Dystrophy</i><br/>         Lori Wallrath, PhD</p> <p><i>Galectin-3 Inhibition Reduces Inflammation and Shifts Macrophage Profiles in LAMA2-CMD</i><br/>         Yonne Tenorio de Menezes, PhD</p> <p><i>Targeting Cell-Matrix Interactions for the Muscular Dystrophies</i><br/>         Rachelle Crosbie, PhD</p> <p><i>Genetic Modifiers in LGMD</i><br/>         Elizabeth McNally, MD</p> | Coronado BCD |
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Tuesday, March 18, 2025 cont.

Care Management Track

|                |  |           |
|----------------|--|-----------|
| 4:30 – 6:00 PM | <p><b>Changing Policies and Pushing Boundaries</b><br/> <i>(CE Accredited)</i></p> <p><i>Changing Policies and Pushing Boundaries</i><br/>         Chair – Leigh Maria Ramos-Platt, MD</p> <p><i>Getting the Genie Out of the Bottle</i><br/>         Lamar Davis II, MD</p> <p><i>Enterprise Partnership: Collaborating in High-Cost Drug Program Development</i><br/>         Malika Maddison, MHA, MBA, RT(T)</p> <p><i>Collaborating Across 50 States &amp; Within Your Own</i><br/>         Chamindra Laverty, MD</p> <p><i>Leading for Your Patients at the National Level</i><br/>         Nicholas Johnson, MD, MSCI, FAAN</p> | Cortez CD |
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Genetic Medicine Track

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|----------------|---|----------------------|
| 4:30 – 6:00 PM | <p><b>Gene Therapy Funding Strategies: Workshop for MDA Care Centers</b><br/> <i>(CE Accredited)</i></p> <p>Chair – Hoda Abdel-Hamid, MD, MSc, FAAN<br/>         Co-Chair – Barry Byrne, MD, PhD<br/>         Ashutosh Kumar, MD<br/>         Matthew Ginsberg, MD<br/>         Crystal Proud, MD</p> | Cortez AB            |
| 6:00 – 8:00 PM | Poster & Networking Reception   | Trinity Exhibit Hall |



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## Wednesday, March 19, 2025

|                   |   |                               |
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| 8:00 – 11:00 AM   | Registration Open   | Geppetto Foyer,<br>Tower Side |
| 8:00 AM – 2:30 PM | <p><b>New, Novel, Noteworthy &amp; Late-Breaking: NMD Highlights</b></p> <p>We are thrilled to have had an overwhelming number of incredible abstract submissions as we reconvene in 2025. To that end, we have redesigned our Clinical Trial Presentations to include a wider range of topics which also allows for even more live presentations selected directly from the abstract submissions.</p> <p>This is a final day not to be missed!</p> |                               |
| 8:00 – 2:30 PM    | <b>Clinical Trial Updates</b>   | Coronado<br>ABCD              |
| 8:00 – 8:15 AM    | <i>First-In-Human Phase 1 Study of Orally Administered SAT-3247 In Healthy Volunteers and Adult Participants with Duchenne Muscular Dystrophy (DMD)</i><br>Phil Lambert, PhD  |                               |
| 8:15 – 8:30 AM    | <i>RGX-202, An Investigational Gene Therapy for The Treatment of Duchenne Muscular Dystrophy: Interim Clinical Data</i><br>Carolina Tesi Rocha, MD  |                               |
| 8:30 – 8:45 AM    | <i>Safety and Efficacy from The Ongoing Phase 1/2 DELIVER Trial of DYNE-251 In Males with DMD Mutations Amenable to Exon 51 Skipping</i><br>Kevin Flanigan, MD  |                               |
| 8:45 – 9:00 AM    | <i>CONNECT1-EDO51: A 12-Week Open-Label Phase 2 Study to Evaluate PGN-EDO51 Safety and Efficacy in People with Duchenne Amenable to Exon 51 Skipping</i><br>Paul Streck, MD   |                               |
| 9:00 – 9:15 AM    | <i>Del-Zota Produced Statistically Significant Increases in Exon Skipping and Dystrophin Levels in EXPLORE44, A Phase 1/2 Study in Patients with DMD44</i><br>Aravindhan Veerapandiyam, MD  |                               |
| 9:15 – 9:30 AM    | <i>Long-term functional outcomes and safety of delandistrogene moxeparvovec in DMD: EMBARK 2-year and Pooled 3-year analyses</i><br>Crystal Proud, MD   |                               |
| 9:30 – 9:45 AM    | <i>Muscle MRI Outcomes in Patients With Duchenne Muscular Dystrophy Treated with Delandistrogene Moxeparvovec: Findings from EMBARK Part 1</i><br>Krista Vandenborne, PhD   |                               |
| 9:45 – 10:00 AM   | <b>BREAK</b>  |                               |
| 10:00 – 10:15 AM  | <i>Safety and Efficacy of DT-DEC01 Therapy in Non-Ambulatory Duchenne Muscular Dystrophy Patients Up to 24 Months After Systemic Administration</i><br>Maria Siemionow, MD, PhD, DSc  |                               |
| 10:15 – 10:30 AM  | <i>Interim Data Following 24 Weeks of Treatment With WVE-N531 In the Phase 2 Open-Label FORWARD-53 Study</i><br>Li-Jung Tai, MD, PhD  |                               |
| 10:30 – 10:45 AM  | <i>CANYON Trial Results: Sevasetmen, An Investigational Fast Skeletal Myosin Inhibitor, Reduced Muscle Damage Biomarkers and Stabilized Function in BMD</i><br>Craig McDonald, MD   |                               |
| 10:45 – 11:00 AM  | <i>Efficacy and Safety of Apitegromab in Individuals with Type 2 and Type 3 Spinal Muscular Atrophy Evaluated in The Phase 3 SAPPHIRE Trial</i><br>Thomas Crawford, MD  |                               |
| 11:00 – 11:15 AM  | <i>Long-Term Vatiquinone Treatment Slows FA Disease Progression Relative to FACOMS Natural History)</i><br>Jonathan Cherry, PhD   |                               |

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Wednesday, March 19, 2025 cont.

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| 11:15 – 11:30 AM    | <i>Topline Efficacy and Safety Results from REACH: Phase 3 Placebo-Controlled Trial of Losmapimod for Facioscapulohumeral Muscular Dystrophy (FSHD)</i><br>Jeffrey Statland, MD     |           |
| 11:30 – 11:45 AM    | <i>Miglustat: A First-in-Class Enzyme Stabilizer for Late-Onset Pompe Disease</i><br>Jon Brudvig, PhD   |           |
| 11:45 AM – 12:00 PM | <b>BREAK</b>  |           |
| 12:00 – 12:15 PM    | <i>RAINBOWFISH: 2-year Efficacy and Safety Data in Risdiplam-Treated Infants with Presymptomatic Spinal Muscular Atrophy (SMA)</i><br>Richard Finkel, MD                            |           |
| 12:15 – 12:30 PM    | <i>Exploring Higher Doses of Nusinersen in Spinal Muscular Atrophy (SMA): Final Results from Parts B and C of the 3-part DEVOTE Study</i><br>Thomas Crawford, MD                    |           |
| 12:30 – 12:45 PM    | <i>Safety and Efficacy of DYNE-101 in Adults with DM1: Phase 1/2 ACHIEVE Trial Data</i><br>James Lilleker, MD, MBChB, MRCP, PhD   |           |
| 12:45 – 1:00 PM     | <i>FREEDOM-DM1: A Phase 1, Placebo-Controlled Single Ascending Dose Study to Evaluate PGN-EDODM1 in People with Myotonic Dystrophy Type 1 (DM1)</i><br>Jane Larkindale, PhD         |           |
| 1:00 – 1:15 PM      | <i>Long-Term Safety, Tolerability and Efficacy of AMO-02 in Children, Adolescents and Adults with Congenital and Childhood Myotonic Dystrophy</i><br>Emily Fantelli                 |           |
| 1:15 – 1:30 PM      | <i>Interim Study Update for the BB-301 Gene Therapy Phase 1b/2a First in Human Trial in Subjects with Oculopharyngeal Muscular Dystrophy with Dysphagia</i><br>Jerel Banks, MD, PhD |           |
| 1:30 – 1:45 PM      | <i>A Clinical DMD Cytosine Base Editing Drug</i><br>Chunyan He, PhD   |           |
| 1:45 – 2:00 PM      | <i>Initial Experience From the INSPIRE DUCHENNE Phase I/II Study of SGT-003 Microdystrophin Gene Therapy for Duchenne Muscular Dystrophy</i><br>Kevin Flanigan, MD                  |           |
| 2:00 – 2:15 PM      | <i>Oral Thromboxane Receptor Antagonist Demonstrates Significant Cardiac Benefit in 12-Month Phase 2 Trial in Duchenne Muscular Dystrophy Patients</i><br>John Jerry Parent         |           |
| 2:15 – 2:30 PM      | <i>Intrathecal Onasemnogene Abeparvovec For Patients with Spinal Muscular Atrophy: Phase 3, Randomized, Sham-Controlled, Double-Blind STEER Study</i><br>Crystal Proud, MD          |           |
| 8:00 – 9:45 AM      | <b>Preclinical Gene Correction Strategies</b>   | Cortez CD |
| 8:00 - 8:15 AM      | <i>ARCUS-Mediated Excision of Exons 45-55 Leads to Functional Del45-55 Dystrophin and Restoration of Skeletal Muscle-Function for the Treatment of DMD</i><br>Adam Mischler, PhD    |           |
| 8:15 – 8:30 AM      | <i>Splicing is Improved Using a Novel AAV-Microrna Delivery Platform As A Treatment For Myotonic Dystrophy Type 1</i><br>Martin Goulet, PhD   |           |
| 8:30 – 8:45 AM      | <i>AAV Gene Therapy For LMNA-Associated Laminopathies</i><br>Monique Otero  |           |
| 8:45 – 9:00 AM      | <i>A Survival-Enhanced R155H Homozygote VCP Mouse Model: A Platform for Testing AAV Gene Therapy in Multisystem Proteinopathy</i><br>Lan Weiss, PhD                                 |           |

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## Wednesday, March 19, 2025 cont.

|                     |   |           |
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| 9:00 – 9:15 AM      | <i>Novel CMT2E Mouse Models with NEFL E397K Mutation Reveal Early Chronic Axonal Neuropathy and Phenotype Rescue via AAV Knockdown-and-Replace Therapy</i><br>Christian Lorson, PhD   |           |
| 9:15 – 9:30 AM      | <i>Systemic AAV Gene Therapy with Next Generation Engineered Capsids for Treatment of CNS and Cardiac Symptoms in Friedreich's Ataxia</i><br>Ryan Kast, PhD                           |           |
| 9:30 – 9:45 AM      | <i>CSF Delivery of INS1202 AAV9-SOD1-shRNA Rescues Muscle Function and Hallmarks of Neurodegeneration in a Disease Model Of ALS</i><br>Laura Ferraiuolo, PhD                          |           |
| 9:45 – 10:00 AM     | <b>BREAK</b>  |           |
| 10:00 – 1:00 PM     | <b>Preclinical Studies in Dystrophies and Myopathies</b>  | Cortez CD |
| 10:00 – 10:15 AM    | <i>Identification of Enzymatically Modified Isoquercitrin (EMIQ) as a Therapeutic Candidate for Myotonic Dystrophy tTpe 1</i><br>Subodh Mishra, PhD                                   |           |
| 10:15 – 10:30 AM    | <i>Discovery of Small Molecules that Bind CUG Repeats, Displace Muscleblind Protein, and Improve Pathogenesis of Myotonic Dystrophy Type 1</i><br>Domi Stickens, PhD                  |           |
| 10:30 – 10:45 AM    | <i>Evaluation of an Adiponectin Receptor Agonist, ALY688, In The Mdx Model Of DMD</i><br>Henry Hsu, MD  |           |
| 10:45 – 11:00 AM    | <i>Novel Inhibitor PK007 Reduces Muscle Inflammation and Myonecrosis in Mdx Mouse Models</i><br>Sai Yarlagadda, PhD   |           |
| 11:00 – 11:15 AM    | <i>Irodanoprost, A Tissue-Targeted EP4 Receptor Agonist, Improves Muscle Histology And Function in a Rat Model Of Severe Duchenne Muscular Dystrophy</i><br>Paul Kostenuik, PhD       |           |
| 11:15 – 11:30 AM    | <i>Novel Small Molecule Mitochondriotropics Reverse Skeletal Muscle Pathology and Improve Resilience in The D2.Mdx Model Of Duchenne Muscular Dystrophy</i><br>Matt Whiteman, MD, PhD |           |
| 11:30 – 11:45 AM    | <i>Non-Viral Gene Therapy for DMD Allowing Full-Length Dystrophin Delivery to Skeletal, Cardiac, and Diaphragm Muscles</i><br>Ivan Krivega, PhD                                       |           |
| 11:45 AM – 12:00 PM | <b>BREAK</b>  |           |
| 12:00 – 12:15 PM    | <i>Small Non-Coding tRNA Derivative Stabilizes Heart Function in Mice with Duchenne Muscular Dystrophy</i><br>Russell Rogers, PhD   |           |
| 12:15 – 12:30 PM    | <i>MLAB-001: A Novel Muscle-Targeted Notch Agonist as a Potential New Treatment for Duchenne Muscular Dystrophy</i><br>Yunzhang Zhu, PhD  |           |
| 12:30 – 12:45 PM    | <i>Extracellular Vesicles Ameliorate Innate Immune Responses to AAV Gene Therapy</i><br>Bradley Hamilton, PhD   |           |
| 8:00 - 9:45 AM      | <b>Advancing NMD Research with Digital Tools and Patient Data</b>   | Cortez AB |
| 8:00 - 8:15 AM      | <i>De-Risking Duchenne Muscular Dystrophy Drug Development Through Collaboration, Data Sharing and Understanding of Disease Stages and Progression</i><br>Paige Martin, PhD           |           |

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Wednesday, March 19, 2025 cont.

|                  |   |
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| 8:15 – 8:30 AM   | <i>BetterLife FSHD and FSHD Navigator: Harnessing Patient Data and Lived Experience to Improve Health and Drive Research</i><br>Amanda Hill, MBA & Anna Gilmore               |
| 8:30 – 8:45 AM   | <i>Feasibility of Using a Wearable Digital Health Technology Sensor to Assess Ambulation in DM1 and CMT</i><br>Laurent Servais, MD, PhD                                       |
| 8:45 – 9:00 AM   | <i>Remote Monitoring of Physical Activity and Upper Limb Function in Adults with Charcot-Marie-Tooth Disease</i><br>Kayla Cornett, PhD  |
| 9:00 – 9:15 AM   | <i>Digital Health Technology for Remote Symptoms Monitoring in Myasthenia Gravis</i><br>Ashkan Vaziri, PhD  |
| 9:15 – 9:30 AM   | <i>Reliability and Sensitivity of a Home-Based Video Assessment for Patients with Multisystem Proteinopathy</i><br>Lindsay Alfano, PT, DPT, PCS                               |
| 9:30 – 9:45 AM   | <i>Reliability of Remote Assessments in Myotonic Dystrophy Type 1</i><br>Laura Tufano   |
| 9:45 – 10:00 AM  | <b>BREAK</b>  |
| 10:00 – 12:00 PM | <b>Molecular Insights into NMDs</b>   |
|                  | Cortez AB   |
| 10:00 – 10:15 AM | <i>A Large, Real-World Study of Serum NfL from People with ALS Enrolled in the ARC Study</i><br>Fernando Vieira, MD   |
| 10:15 – 10:30 AM | <i>Low-Density Lipoprotein Receptor is Critical for Sporadic ALS CSF-Induced Neurotoxicity</i><br>Jamie Kay Wong, PhD   |
| 10:30 – 10:45 AM | <i>Skeletal Muscle Targeted CIC-1 Ion Channel Inhibitor Improves Skeletal Muscle Function and Respiratory Function in A Rat Model of MuSK-MG</i><br>Martin Brandhøj Skov, PhD |
| 10:45 – 11:00 AM | <i>Restoration of Motor Neuron Function Via Pharmacological in Vivo Regulation of a Potassium Channel in SMA Mice</i><br>Nandhini Sivakumar, PhD                              |
| 11:00 – 11:15 AM | <i>Spinal and Bulbar Muscular Atrophy Cardiomyocytes Exhibit Arrhythmia</i><br>Asuka Eguchi, PhD  |
| 11:15 – 11:30 AM | <i>Long-Read Whole Genomic Sequencing Reveals Novel Pathogenic Alleles in Friedreich Ataxia</i><br>Sanjay Bidichandani, MBBS, PhD   |
| 11:30 – 11:45 AM | <i>Spatial Transcriptomics of Motor Neuron Subcellular Compartments Reveals an Axon-Specific RNA Signature and Local Translation Defects in FUS-ALS</i><br>Diana Piol, PhD    |