

XXIII NATIONAL CONGRESS OF ITALIAN ASSOCIATION of MYOLOGY

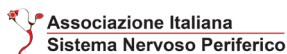
PADOVA

8th - 10th June 2023

www.congressoaim2023.it

Il XXIII Congresso Nazionale Associazione Italiana di Miologia

si svolge con i patrocini di:



Comune di Padova



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PROGRAM

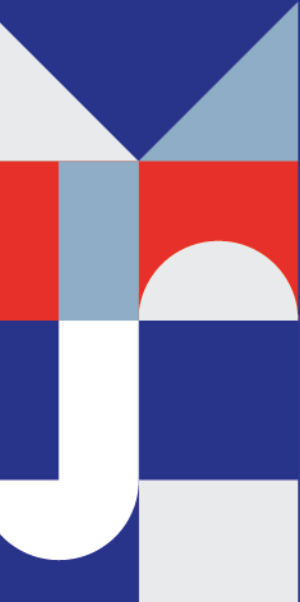
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**BOARD
OF DIRECTORS**

ITALIAN ASSOCIATION
of MYOLOGY



XXIII NATIONAL CONGRESS OF ITALIAN ASSOCIATION of MYOLOGY

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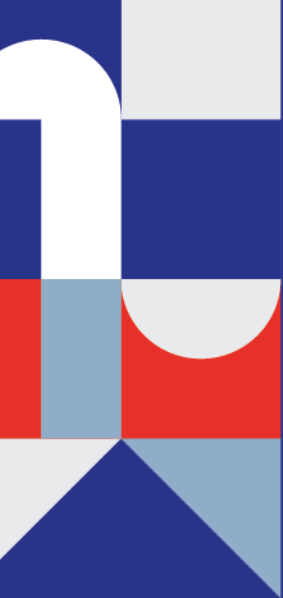
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FACULTY

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XXIII NATIONAL CONGRESS OF ITALIAN ASSOCIATION of MYOLOGY

FACULTY

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Antonio Di Muzio, Chieti	Federica Ricci, Torino
Amelia Evoli, Roma	Giulia Ricci, Pisa
Toby Ferguson, Cambridge (USA)	Carmelo Rodolico, Messina
Alessandra Ferlini, Bologna	Lucia Ruggiero, Napoli
Massimiliano Filosto, Brescia	Francesco Saccà, Napoli
Chiara Fiorillo, Genova	Dorianna Sandonà, Padova
Brian Fox, Boston (USA)	Marco Sandri, Padova
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Rocco Liguori, Bologna	Federica Trucco, Milano
Michelangelo Maestri, Pisa	Rossella Tupler, Modena
Lorenzo Maggi, Milano	Liliana Vercelli, Torino
Francesca Magri, Milano	Elisabetta Verrillo, Roma
Michelangelo Mancuso, Pisa	Carlo Viscomi, Padova
Renato Mantegazza, Milano	Massimo Domenico Zeviani, Padova
Andrea Martinuzzi, Treviso	Daniela Zuccarello, Padova

PROGRAM AT A GLANCE

ITALIAN ASSOCIATION of MYOLOGY

Thursday, 8th June 2023

	MANTEGNA HALL First Floor	PETRARCA HALL Ground Floor	PALLADIO HALL Ground Floor	POSTER AREA Third Floor	
10.10 - 10.40	LECTURE NO CME Challenges delivering ERT in lysosomal disorders: special considerations in LOPD				
10.45 - 11.00	Greetings and Introduction				
11.00 - 11.30	Main Lecture Translating knowledge on hereditary neuromuscular childhood diseases into therapies				
11.30 - 13.00	WORKSHOP Sleep disorders in Neuromuscular diseases: treatable conditions	ORAL COMMUNICATIONS SMA			
13.00 - 14.00	LUNCH			POSTER VIEWING NO CME	
14.00 - 15.00	Symposium NO CME New Frontiers in SMA				
15.00 - 16.00	Invited talks Preimplantation Genetic Test (PGT) Mechanisms regulating skeletal muscle growth and atrophy				
16.00 - 17.00	New therapeutic Approaches in Pompe Disease				
17.00 - 17.30	COFFEE BREAK				
17.30 - 18.30	WORKSHOP gender issues in Neuromuscular Disorders	ORAL COMMUNICATIONS Diagnostic and Therapeutic tools	ORAL COMMUNICATIONS Disease Mechanisms and Pathogenesis		
18.30 - 19.00	LECTURE NO CME New Born screening for SMA: the new scenarios				
19.00 - 20.00	Symposium NO CME The Motor Pool in SMA				
20.00	WELCOME COCKTAIL (Congress venue)				

Friday, 9th June 2023

	MANTEGNA HALL First Floor	PETRARCA HALL Ground Floor	PALLADIO HALL Ground Floor	POSTER AREA Third Floor		
08.00 - 08.30	MAIN LECTURE Cardiac involvement in female D/BMD-gene mutation carriers					
08.30 - 10.00	WORKSHOP MITOCHONDRIAL DISORDERS					
10.00 - 10.30	Nereo Bresolin Lecture on Mitochondrial disorders A Journey to Develop Therapy for Thymidine Kinase 2 Deficiency					
10.30 - 11.00	COFFEE BREAK					
11.00 - 12.30	ORAL COMMUNICATIONS LGMD and FSHD	ORAL COMMUNICATIONS Metabolic Myopathies	ORAL COMMUNICATIONS: Channelopathies and Inflammatory Myopathies	POSTER VIEWING NO CME		
12.30 - 13.00	MAIN Lecture Gene Therapy for Rare Neuromuscular Disorders					
13.00 - 14.00	LUNCH					
14.00 - 15.00	Symposium NO CME Myasthenia Gravis: burden of disease and innovative treatments targeting the FcRn					
15.00 - 16.00	Symposium NO CME Get off the Myasthenia Gravis rollercoaster through continuous C5 inhibition					
16.00 - 16.30	COFFEE BREAK					
16.30 - 18.30	WORKSHOP Novel therapeutic approaches in Limb Girdle Muscular Dystrophies					
18.30 - 19.30	GENERAL ASSEMBLY					
21.00	SOCIAL DINNER					

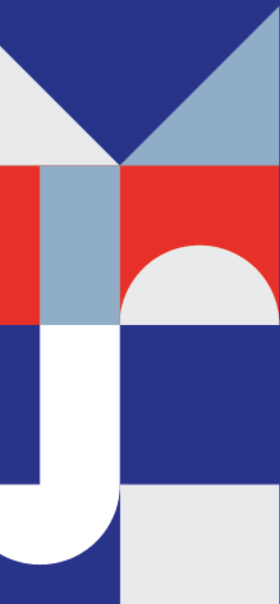
Saturday, 10th June 2023

	MANTEGNA HALL First Floor	PETRARCA HALL Ground Floor	PALLADIO HALL Ground Floor	POSTER AREA Third Floor	
08.00 - 09.00	Muscle Club				
09.00 - 09.30	MAIN Lecture Update about steroid and vamorolone therapy in DMD				
09.30 - 10.00	COFFEE BREAK				
10:00 - 11:30	ORAL COMMUNICATIONS Dystrophinopathies	ORAL COMMUNICATIONS Myopathies	ORAL COMMUNICATIONS Myasthenia Gravis		
11.30 - 13.30	ROUND TABLE MEETING with PATIENT's ASSOCIATIONS				
13.30	Awards and conclusions				



PROGRAM

ITALIAN ASSOCIATION
of MYOLOGY



Thursday, 8th June 2023

MANTEGNA HALL

10.10 - 10.40 | **LECTURE** **NO CME**

Challenges delivering ERT in lysosomal disorders: special considerations in LOPD
Brian Fox (New York, USA) introduced by Corrado Angelini (Padova)

10:45 - 11:00 GREETINGS AND INTRODUCTION

Chairpersons: *Giacomo Comi (Milano), Maurizio Corbetta (Padova), Elena Pegoraro (Padova)*

11.00 - 11.30 | **MAIN LECTURE**

Translating knowledge on hereditary neuromuscular childhood diseases into therapies
Francesco Muntoni (Londra, UK) introduced by Giacomo Comi (Milano)

11.30 - 13.00 | **WORKSHOP**

SLEEP DISORDERS IN NEUROMUSCULAR DISEASES: TREATABLE CONDITIONS

Chairpersons: *Claudio Cherchi (Roma), Antonio Trabacca (Brindisi)*

11.30 | Sleep disorders in neurological and neuromuscular conditions:
diagnostic tools and treatment approach
Michelangelo Maestri (Pisa)

11.50 | Sleep and Respiratory Disorders in Duchenne Muscular Dystrophy
Maria Grazia D'Angelo (Milano)

12.10 | Disorders of sleep in LOPD and response to ERT
Sabrina Ravaglia (Pavia)

12.30 | Sleep Disorders in SMA
Elisabetta Verrillo (Roma)

12.50 | Discussion

13.00 - 14.00 | Lunch

13.00 - 15.00 | **POSTER VIEWING NO CME**

14.00 - 15.00 | **SYMPOSIUM** **NO CME**

New frontiers in SMA

Chairperson: *Giacomo Comi (Milano)*

14.00 | What is next for SMA Management?

Stefania Corti (Milano), Eugenio Mercuri (Roma)

14.30 | How technology innovation can support the multidisciplinary approach

Davide Korn (Roma)

15.00 - 16.00 | **INVITED TALKS**

Chairpersons: *Alessandra Ferlini (Bologna), Rossella Tupler (Modena)*

15.00 | Preimplantation Genetic Test (PGT)

Daniela Zuccarello (Padova)

15.30 | Mechanisms regulating skeletal muscle growth and atrophy

Marco Sandri (Padova)

16.00 - 17.00 | **New Therapeutic Approaches in Pompe Disease**

Chairpersons: *Alberto Burlina (Padova), Gabriele Siciliano (Pisa)*

16.00 | Novelities in the treatment of infantile Pompe disease

Serena Gasperini (Monza)

16.30 | Long term clinical study data on late onset Pompe disease

Olimpia Musumeci (Messina)

17.00 - 17.30 | Coffee Break

17.30 - 18.30 | **WORKSHOP**

GENDER ISSUES IN NEUROMUSCULAR DISORDERS

Chairpersons: *Chiara Fiorillo (Genova), Paola Tonin (Verona)*

17.30 | Gender issues in SMA

Luca Bello (Padova)

17.50 | FSHD different courses in males and females

Massimiliano Filosto (Brescia)

18.10 | New avenues to treat androgen receptor' in Kennedy disease

Maria Pennuto (Padova)

18.30 - 19.00 | **LECTURE** **NO CME**

Newborn screening for SMA: the new scenarios

Federica Ricci (Torino)

19.00 - 20.00 | **SYMPOSIUM** **NO CME**

The Motor Pool in SMA

Chairperson: *Elena Pegoraro (Padova)*

19:00 | Welcome and Introduction

Elena Pegoraro (Padova)

19:05 | An introduction to motor pools and neurodegeneration

Sonia Messina (Messina)

19:20 | Tracing the motor pool-specific vulnerabilities

Toby Ferguson (Cambridge, USA)

19:35 | Clinical implication & clinical case

Valeria Sansone (Milano)

19:50 | Questions and Answers

20.00 | Welcome Cocktail

PETRARCA HALL

11.30 - 13.00 | ORAL COMMUNICATIONS | SMA

Chairpersons: Adele D'Amico (Roma), Claudio Bruno (Genova)

- 11.30** | Search for new CSF biomarkers in SMA patients treated with Nusinersen
C. Panicucci, M. Bartolucci, E. Sahin, R. Russo, M. Valletta, N. Brolatti, M. Pedemonte, S. Casalini, S. Baratto, A. D'Amico, M. Pane, M. Sframeli, C. Stancanelli, S. Messina, E. Albamonte, V. Sansone, E. Mercuri, A. Chambery, U. Sezerman, A. Petretto, C. Bruno.
Genova, Istanbul, Caserta, Roma, Messina, Milano
- 11.45** | Pregnancy experience in women with spinal muscular atrophy: an overview in the cohort of the University of Naples Federico II
R. Pira Bencivenga, D. Zoppi, A. Russo, E. Cassano, S. Tozza, R. Iodice, R. Dubbioso, F. Manganelli, L. Ruggiero.
Napoli
- 12.00** | Patients and caregivers expectations on possible functional changes following disease-modifying treatment in spinal muscular atrophy
M. C. Pera, G. Coratti, J. L. Casiraghi, C. Bravetti, A. Fedeli, M. Strika, E. Albamonte, L. Antonaci, D. Rossi, M. Pane, V. Sansone, E. Mercuri.
Roma
- 12.15** | Swallowing and feeding in Spinal Muscular Atrophy (SMA) type 1: the SFERA project and the path towards a diagnostic and treatment algorithm
S. Gandolfi, C. Dosi, M.R. Scopelliti, E. Minacapilli, R. De Amicis, S. Bertoli, A. Mandelli, A. Campari, A. Schindler, R. Masson, on behalf of the SFERA working group.
Milano
- 12.30** | Facial nerve vulnerability in spinal muscular atrophy and motor unit number index (MUNIX) of the orbicularis oculi muscle
A. Barp, E. Carraro, F. Salmin, A. Lizio, M. Cheli, V. Sansone.
Milano, Valsugana
- 12.45** | Molecular and biochemical characterization of patient-derived cellular models for Spinal Muscular Atrophy with Myoclonus Epilepsy (SMA-PME)
D. Ronchi, S. Salani, M. Garbellini, M. S. Hidalgo Moreno, S. Antognozzi, M. Aureli, S. Corti, G. P. Comi.
Milano

13.00 - 14.00 | Lunch

13.00 - 15:00 | POSTER VIEWING NO CME

17:30 - 18:30 | ORAL COMMUNICATIONS | DIAGNOSTIC AND THERAPEUTIC TOOLS**Chairpersons:** *Marco Savarese (Helsinki, Finland), Giulia Ricci (Pisa)***17:30 | Engineering human stem cells for advanced neuromuscular disease and therapy modelling***V. M. Lionello, L. Pinton, S. Dastidar, D. Moore, S. Choi, H. Steele-Stallard, S. Jalal, Y. Jiang, P. Zammit, F. S. Tedesco.
Londra***17:45 | Solving the NGS negative cases:***improved WES protocols to make increasingly accurate genetic diagnoses
A. Torella, M. E. Onore, E. Picillo, A. Budillon, G. Piluso, L. Politano, V. Nigro.
Napoli***18:00 | A comparison of different NGS-based strategies for the routine clinical diagnosis of neuromuscular disorders***C. Bertolin, F. Boaretto, S. Volta, E. Pegoraro, L. Bello, G. Sorarù, C. Viscomi, M. Zeviani, L. Salvati.
Padova***18:15 | OMICS analysis to identify the genetic causes of undiagnosed neuromuscular phenotypes: the UNIFE experience within the SOLVE-RD Project***R. Selvatici, M. Neri, F. Gualandi, S. Bigoni, F. Fortunato, E. Bertini, M. Pane, E. Mercuri, A. Ferlini.
Ferrara, Roma***PALLADIO HALL****17.30 - 18.30 | ORAL COMMUNICATIONS | DISEASE MECHANISMS AND PATHOGENESIS****Chairpersons:** *Roberto Massa (Roma), Monica Sciacco (Milano)***17.30 | RNA-seq data analysis in RYR1 related congenital myopathies with cores and minicores: an Italian population study***D. Sabbatini, A. Fusto, M. Suman, D. Gorgoglione, S. Vianello, C. Romualdi, C. Marchioretto, M. Pennuto, G. Szabadkai, L. Bello, E. Pegoraro.
Padova, Londra***17.45 | Medaka fish *Oryzias latipes* a promising model to study titinopathies: the HMERF case study***V. Cetrangolo, M. Savarese, I. Conte, L. Polishchuk, V. Nigro, M. Linari, B. Udd.
Pozzuoli***18.00 | Aberrant Adenosine Triphosphate Release and Impairment of P2Y2-Mediated Signaling in Sarcoglycanopathies***S. Baratto, A. Benzi, C. Astigiano, L. Sturla, C. Panicucci, K. Mamchaoi, L. Raffaghello, S. Bruzzone, E. Gazzero, C. Bruno.
Genova, Berlino, Parigi*

18.15 | Functional characterization suggests that ACTN2 frameshift variants cause distal myopathy through protein aggregation
*J. Ranta-aho, P. H. Jonson, J. Sarparanta, B. Udd, M. Savarese.
Helsinki*

Friday, 9th June 2023

MANTEGNA HALL

08:00 - 08:30 | MAIN LECTURE

Chairpersons: *Giacomo Pietro Comi (Milano), Luisa Politano (Napoli)*

08:00 | Cardiac involvement in female D/BMD-gene mutation carriers
John Bourke (Newcastle, UK)

08.30 - 10.00 | WORKSHOP MITOCHONDRIAL DISORDERS

Chairpersons: *Michelangelo Mancuso (Pisa), Massimo Zeviani (Padova)*

08:30 | Cardiorespiratory involvement in primary mitochondrial diseases
Serenella Servidei (Roma)

08:50 | Mitochondrial optic neuropathies: differential diagnosis and therapeutic strategies
Chiara La Morgia (Bologna)

09:10 | Mitochondrial Myopathies: Models and therapies
Carlo Viscomi (Padova)

09:30 | SURF1 therapeutic development in non-human mammals
Dario Brunetti (Milano)

09:50 | Discussion

10.00 - 10.30 | NEREO BRESOLIN LECTURE ON MITOCHONDRIAL DISORDERS

A Journey to Develop Therapy for Thymidine Kinase 2 Deficiency
Michio Hirano (New York) introduced by Antonio Toscano (Messina)

10.30 - 11.00 | Coffee Break

11.00 - 12.30 | ORAL COMMUNICATIONS | LGMD AND FSHD

Chairpersons: *Francesca Magri (Milano), Giorgio Tasca (Roma)*

11.00 | Clinical and genetic analysis of the Italian cohort of LAMA2-Related Dystrophy (LAMA2-RD)

A. Zambon, S. Messina, M. Sframeli, A. Pini, M. L. Valentino, C. Fiorillo, C. Bruno, C. Panicucci, F. Ricci, T. Mongini, G. D'Angelo, R. Cima, G. Ricci, L. Politano, M. Pane, E. Mercuri, E. Pegoraro, L. Bello, E. Albamonte, F. Trucco, V. Sansone, A. Berardinelli, G. Astrea, R. Battini, M. Filosto, S. Cotti Piccinelli, F. Magri, D. Velardo, G. P. Comi, I. Moroni, F. Bruschi, L. Maggi, M. Catteruccia, E. Bertini, A. D'Amico, S. C. Previtali. Milano, Messina, Genova, Torino, Pisa, Napoli, Roma, Padova, Brescia

11.15 | The role of the NAD+/CD38 axis in the pathophysiology of alfa-sarcoglycanopathy

A. Benzi, A. Amaro, S. Pintus, F. Antonini, F. Reggiani, S. Baratto, E. Principi, G. Del Zotto, D. Cassandrini, A. D'Amico, A. Malandrini, E. Malfatti, T. Mongini, E. Pegoraro, S. Previtali, C. Rodolico, G. Tasca, F. Morandi, F. Malavasi, U. Pfeffer, S. Bruzzone, C. Bruno, L. Raffaghello. Genova, Pisa, Roma, Siena, Créteil, Torino, Padova, Milano, Messina, Newcastle

11.30 | Prenatal titinopathies are the severe end of the TTN-related myopathies spectrum: genotype- phenotype correlation on a large international cohort

M. F. Di Feo, A. Brady, F. Forzano, M. Iascone, P. D'Oria, L. Spaccini, S. Kurbatov, E. Giorgio, G. Casalis Cavalchini, A. Brusco, B. Udd, M. Savarese. Helsinki, Middlesex, Londra, Bergamo, Milano, Voronezh, Pavia, Torino

11.45 | Whole Exome Sequencing of 126 patients highlights variants in candidate genes associated with FSHD

C. Strafella, D. Megalizzi, V. Caputo, G. Trastulli, L. Colantoni, S. Bortolani, E. Torchia, M. Monforte, C. Caltagirone, E. Ricci, G. Tasca, E. Giardina, R. Cascella. Roma, Newcastle, Tirana

12.00 | When it is not FSHD: reviewing the mimics

G. Gadaleta, L. Vercelli, G. Urbano, E. Rolle, R. Tupler, T. Mongini. Torino, Modena

12.15 | New insights for the interpretation of SMCHD1 gene variants effect: a functional study on a heterogeneous cohort of patients

S. Pini, M. Francesca Di Feo, M. Chiara, M. Savarese, P. Uva, M. Iascone, F. Santorelli, M. Vandroux, J. Laporte, A. Puliti, N. Albano, M. Seri, T. Pippucci, F. Isidori, A. Cereda, B. Udd, V. Salsi, R. Tupler. Modena, Worcester, Genova, Milano, Bari, Helsinki, Bergamo, Pisa, Ilkirk, Bologna, Bergamo, Vaasa

12.30 - 13.00 | MAIN LECTURE

Chairpersons: *Adele D'Amico (Roma), Tiziana Mongini (Torino)*

Gene Therapy for Rare Neuromuscular Disorders

Stefania Corti (Milano)

13.00 - 14.00 | Lunch

13:00 - 16:30 | POSTER VIEWING NO CME

14.00 - 15.00 | **SYMPOSIUM** **NO CME**

Myasthenia Gravis: burden of disease and innovative treatments targeting the FcRn

Chairpersons: *Lorenzo Maggi (Milano), Carmelo Rodolico (Messina)*

FcRn inhibition: a novel treatment option in Myasthenia Gravis and beyond

Raffaele Iorio (Roma)

Improvement in gMG health-related quality-of-life beyond the sole relief of signs and symptoms

Francesco Saccà (Napoli)

Conclusions

Lorenzo Maggi (Milano), Carmelo Rodolico (Messina)

15.00 - 16.00 | **SYMPOSIUM** **NO CME**

Get off the Myasthenia Gravis rollercoaster through continuous C5 inhibition

Chairpersons: *Giovanni Antonini (Roma), Rocco Liguori (Bologna)*

15:00 | Innovative therapies in gMG: The role of Complement

Francesco Habetswallner (Napoli)

15:30 | A new gMG therapeutic algorithm in the precision Medicine era

Renato Mantegazza (Milano)

16.00 - 16.30 | Coffee Break

16.30 - 18.30 | **WORKSHOP**

NOVEL THERAPEUTIC APPROACHES IN LIMB GIRDLE MUSCULAR DYSTROPHIES

Chairpersons: *Carlo Minetti (Genova), Vincenzo Nigro (Napoli)*

16.30 | Future development of gene therapy in sarcoglycanopathies and FKRP-related LGMD

Sophie Olivier (Paris, France)

17.00 | Novel therapeutic approaches in sarcoglycanopathies

Dorianna Sandonà (Padova)

17.30 | Gene Therapy for Sarcoglycanopathies

Amanda Haidet-Phillips (Ohio, USA)

18.00 | Therapies in LGMD2I/R9-FKRP related Limb Girdle Muscular Dystrophy

Douglas M. Sproule (North Carolina, USA)

18.30 - 19.30 | **GENERAL ASSEMBLY**

21:00 | **SOCIAL DINNER**

ORAL COMMUNICATIONS | METABOLIC MYOPATHIES

Chairpersons: Giovanna Cenacchi (Bologna), Liliana Vercelli (Torino)

- 11.00** | Exploring the role of Mobile Health Technologies in detecting subtle motor impairment in paucisymptomatic Late-onset Pompe Disease: a preliminary study
B. Labella, A. Rizzardi, S. Cotti Piccinelli, C. Zatti, C. Hansen, R. Romijnders, W. Maetzler, F. Caria, B. Risi, S. Damioli, E. Olivieri, L. Ferullo, L. Poli, A. Padovani, A. Pilotto, M. Filosto.
 Brescia, Kiel
- 11.15** | Mitochondrial involvement in Patients with Autism spectrum disorders and Intellectual disability: a histological and genetic study
C. Scuderi, S. Santa Paola, M. Lo Giudice, M. Giuliano, F. D. Di Blasi, S. Città, R. Pettinato, G. A. Vitello, C. Romano, E. Borgione.
 Troina
- 11.30** | Pregnancy in late-onset Pompe disease (LOPD) women: a multi-centre experience
O. Musumeci, L. Vercelli, M. Porcino, G. Gadaleta, I. G. Arena, T. Mongini, C. Rodolico, A. Toscano.
 Messina, Torino
- 11.45** | Prevalence of Sleep Disorders in a cohort of patients with Primary Mitochondrial Diseases
I. G. Arena, C. Consulo, M. Porcino, C. Usbergo, C. Rodolico, A. Toscano, O. Musumeci.
 Messina
- 12.00** | Clinico-pathologic features and molecular genetic spectrum in a cohort of adult patients with mtDNA maintenance disorders
M. Meneri, D. Piga, F. Magri, D. Velardo, P. Ciscato, S. Zanotti, M. Sciacco, G. P. Comi, D. Ronchi.
 Milano
- 12.15** | Deriving harmonized follow-up in hyperCKemia using a digital web tool
B. Buchignani, F. Sansone, G. Marinella, G. Astrea, S. Frosini, D. Cassandrini, A. Tonacci, R. Conte, R. Battini, A. Rubegni, F. M. Santorelli and the InGene 2.0 group.
 Pisa, Roma

ORAL COMMUNICATIONS | CHANNELOPATIES AND INFLAMMATORY MYOPATHIES

Chairpersons: *Giovanni Meola (Milano), Antonio Di Muzio (Chieti)*

- 11.00** | LAMOTRIGINE as an anti-myotonic agent in Myotonic Dystrophy typ 1:
an open-label single center pilot study
S. Cotti Piccinelli, B. Risi, F. Caria, S. Damioli, E. Bertella, F. Garofali, N. Ait Allali, B. Labella, L. Poli, V. Bonito, A. Padovani, M. Filosto.
Brescia
- 11.15** | Therapeutic approaches to adult Ab anti-NXP2+ Dermatomyositis:
a single center experience
G. Urbano, G. Gadaleta, F. Rumbolo, G. Mengozzi, S. Boschi, L. Chiadò-Piat, E. Rolle, L. Vercelli, T. Mongini.
Torino
- 11.30** | Refining the clinical spectrum and prognosis of granulomatous myositis from
a large cohort of patients
A. Lauletta, L. de le Hoye, O. Benveniste, M. Garibaldi.
Roma, Brussels, Parigi
- 11.45** | Multiple inflammatory biomarkers assay can help distinguish between idiopathic
inflammatory myopathies subtypes
M. Lucchini, V. De Arcangelis, M. Mirabella.
Roma
- 12.00** | Clinical and functional characterization of the novel Nav1.4 sodium channel mutation,
p.T592I, found in a three-generation family
C. Campanale, P. Laghetti, I. Saltarella, C. Altamura, P. Bernasconi, R. Mantegazza, R. Brugnoli, P. Tacconi, J. F. Desaphy.
Bari, Milano, Cagliari
- 12.15** | Phenotypic variability of Andersen -Tawil Syndrome in a family sharing the same
allelic KCNJ2 gene mutation
M. E. Onore, E. Picillo, A. Torella, E. Cesarone, G. Piluso, V. Nigro, L. Politano.
Napoli

MANTEGNA HALL

08.00 - 09.00 | **MUSCLE CLUB**

Chairpersons: *Serenella Servidei (Roma), Antonella Pini (Bologna)*

08.00 | Why am I cramping? Think beyond obvious

G. Tammam, E. Vegezzi, S. Gana, A. Pichiecchio, E. Alfonsi, G. Cosentino.
Pavia

08.12 | A rare case of adult-onset myopathy with facial involvement: expanding genotype-phenotype relationship and histopathological findings

S. Cotti Piccinelli, B. Risi, F. Caria, S. Damioli, E. Bertella, B. Labella, L. Poli, V. Bonito,
G. Lanzi, A. Padovani, M. Filosto.
Brescia

08.24 | A case of myalgias and hyperCKemia after exercise

M. Villa, P. Riguzzi, G. Capece, V. Zangaro, E. Sogus, M. Penzo, D. Gorgoglione,
D. Sabbatini, S. Vianello, L. Bello, E. Pegoraro.
Padova

08.36 | Diagnostic flowchart in an adult patient with asymmetric ptosis

D. Zoppi, R. Bencivenga, T. Fioretti, A. Russo, F. Masciarelli, R. Di Leo, M. Pezzella,
R. Iodice, G. Esposito, F. Manganelli, L. Ruggiero.
Napoli

08.48 | A rare cause of congenital hypotonia: clinical aspects and molecular findings

A. Decio, R. Giorda, R. Cima, S. Marelli, E. Diella, D. Sacchi, C. Legnani, F. Villa,
M.T. Bassi, M.G. D'Angelo.
Bosisio Parini

09.00 | **MAIN LECTURE**

Chairperson: *Stefano Previtali (Milano)*

Update about steroid and vamorolone therapy in DMD

Michela Guglieri (Newcastle, UK)

09.30 - 10.00 | Coffee Break

10.00 - 11.30 | **ORAL COMMUNICATIONS: DYSTROPHINOPATHIES**

Chairpersons: *Luca Bello (Padova), Riccardo Masson (Milano)*

10.00 | Dystrophin and utrophin quantitation by targeted mass spectrometry in Duchenne and Becker muscular dystrophy

P. Riguzzi, L. Bello, M. Penzo, M. Villa, G. Capece, V. Zangaro, D. Gorgoglione,
S. Vianello, D. Sabbatini, E. Canessa, Y. Hathout, E. Pegoraro.
Padova, Binghamton

- 10.15 | Natural history of Becker muscular dystrophy: a retrospective multicentre study**
D. Gorgoglione, D. Sabbatini, P. Riguzzi, A. Petrosino, M. Penzo, M. Villa, G. Capece, S. Vianello, G. S. Previtali, A. Zambon, C. Bruno, C. Panicucci, M. Traverso, A. Berardinelli, R. Nicotra, S. Parravicini, A. Gardani, A. Ferlini, F. Fortunato, E. M. Mercuri, M. Pane, C. Palermo, S. Servidei, C. Sancricca, M. Briganti, S. Messina, M. Sframeli, C. Allegra, A. D'Amico, M. Catteruccia, V. Sansone, A. Barp, M. G. D'Angelo, I. Moroni, F. Bruschi, A. Ardisson, R. Masson L. Maggi, A. Gallone, V. Nigro, E. Picillo, A. Pini, M. Giannotta, R. Battini, G. Marinella, T. Mongini, G. Gadaleta, G. Urbano, E. Rolle, F. Ricci, F. Magri, A. Lerario, G. Siciliano, G. Ricci, L. Bello, E. Pegoraro.
 Padova, Pisa, Milano, Pavia, Genova, Ferrara, Roma, Messina, Trento, Bosisio Parini, Napoli, Bologna, Torino
- 10.30 | Robust preclinical data support development of DYNE-251 as a potential treatment for individuals with DMD mutations amenable to exon 51 skipping**
E. Delage, E. Cotali, M. L. Naylor, C. A. Desjardins, R. Venkatesan, E. O'Donnell, J. Hall, R. Russo, S. Spring, K. Tang, J. W. Davis II, T. Weeden, S. Zanotti, C. Mix, B. Han, O. Beskrovnaya, A. Dugar.
 Paris
- 10.45 | Definition of diaphragmatic sleep disordered breathing and clinical meaning in Duchenne Muscular Dystrophy**
F. Trucco, M. Davies, A. A. Zambon, D. Ridout, F. Abel, F. Muntoni.
 Londra, Milano
- 11.00 | Cardiomyopathy in Duchenne muscular dystrophy: progression of myocardial involvement and pharmacologic treatment in a large cohort of patients from a single tertiary referral Centre**
M. Sframeli, L. Oreto, C. Stancanelli, C. Allegra, R. Materia, C. Zito, G. Di Bella, V. Tudisco, C. Rodolico, F. De Luca, S. Messina.
 Messina
- 11.15 | RNA in situ hybridization reveals the spatial localization of DMD isoforms in human adult brain areas**
M. S. Falzarano, M. Mietto, F. Fortunato, R. Selvatici, J. Morgan, R. Phadke, F. Catapano, F. Muntoni, A. Ferlini.
 Ferrara, Londra

11.30 - 13.30 | ROUND TABLE

Meeting with Patient's Associations

Chairpersons: Giacomo Pietro Comi (Milano), Federica Ricci (Torino)

13.30 | Awards and conclusions
 Giacomo Pietro Comi (Milano)

10.00 - 11.30 | ORAL COMMUNICATIONS | MYOPATHIES

Chairpersons: *Lucia Ruggiero (Napoli), Maurizio Moggio (Milano)*

- 10.00** | Natural history of distal and myofibrillar myopathies assessed by clinical and technological outcome measures (Dista-Myo): baseline results
S. Bortolani, M. Di Bari, E. Torchia, E. Rolle, A. Vicino, M. Rabuffetti, A. Marzegan, M. Cheli, M. Monforte, E. Ricci, J. Y. Hogrel, S. Sacconi, T. E. Mongini, L. Maggi, G. Tasca. Torino, Roma, Milano, Losanna, Parigi, Newcastle
- 10.15** | Gene editing in Emery-Dreifuss Muscular Dystrophy myoblasts
E. Cattin, E. Mattioli, E. Schena, F. Corradi, A. Pini, L. Maggi, C. Fiorillo, A. Recchia, G. Lattanzi. Bologna, Milano, Genova
- 10.30** | Human mutated MYOT and CRYAB genes cause a myopathic phenotype in Zebrafish
G. Marchetto, E. Cannone, V. Guglielmi, C. Tobia, B. Gnutti, B. Cisterna, A. Barbon P. Tonin, M. Schiavone, G. Vattemi. Verona, Brescia
- 10.45** | Myopathic Ehlers-Danlos Syndrome with Rectus femoris involvement: genetic heterogeneity behind COL12A1 dysfunction
M. Zanobio, A. Torella, P. Sabatelli, R. Zeuli, M. Scarpato, G. Piluso, L. Merlini, V. Nigro. Bologna, Napoli
- 11.00** | Myofibrillar and distal myopathies: natural history of an Italian cohort of patients
M. Cheli, S. Bortolani, E. Rolle, A. Vicino, S. Bonanno, T. Enrica Mongini, G. Tasca, L. Maggi. Milano, Torino, Roma, Newcastle
- 11.15** | Heterozygous truncating variants in DAG1 are associated with sporadic and familial isolated hyperCKemia
M. Traverso, S. Baratto, M. Di Duca, C. Panicucci, M. Iacomino, S. Casalini, M. Grandis, P. Striano, F. Zara, R. Barresi, C. Bruno, C. Fiorillo, M. Scala. Genova, Venezia

10.00 -11.30 | ORAL COMMUNICATIONS| MYASTHENIA GRAVIS

Chairpersons: *Amelia Evoli (Roma), Matteo Garibaldi (Roma)*

- 10.00** | Efficacy of rozanolixizumab in generalised myasthenia gravis: subgroup analyses from the randomised Phase 3 MycarinG study
G. Antonini, T. Vu, A. Drużdż, J. Grosskreutz, A. A. Habib, R. Mantegazza, K. Utsugisawa, J. Vissing, R. Beau Lejdstrom, M. Boehnlein, T. Gasalla, F. Grimson, T. Tarancón, V. Bril. Roma, Tampa, Poznań, Lübeck, Milano, Irvine, Hanamaki, Copenhagen, Monheim, Toronto, Madrid
- 10.15** | RAISE-XT: An interim analysis of safety and efficacy in an open-label extension study of zilucoplan in patients with myasthenia gravis
C. Antozzi, M. Freimer, M. Isabel Leite, A. Genge, Y. Hussain, R. Mantegazza, K. Utsugisawa, T. Vu, P. W. Duda, B. Boroojerdi, M. Vanderkelen, R. Lowcock, J. F. Howard Jr. Milano, Columbus, Oxford, Montreal, Austin, Hanamaki, Tampa, Monheim, Chapel Hill, Braine-l'Alleud
- 10.30** | Long-Term Safety, Tolerability, and Efficacy of Subcutaneous Efgartigimod PH20 in Patients with Generalized Myasthenia Gravis: Interim Results of the ADAPT-SC+ Study
J. F. Howard Jr, G. Li, T. Vu, D. Korobko, M. Smilowski, K. Banaszekiewicz, L. Liu, S. Steeland, J. Noukens, B. Van Hoorick, J. Podhorna, Y. Li, K. Utsugisawa, F. Sacca, H. Wiendl, J. L. De Bleecker, R. Mantegazza, in collaboration with the ADAPT-SC Investigator Study Group. Manchester, Chapel Hill, Tampa, Katowice, Novosibirsk, Kraków, Ghent, Napoli, Hanamaki, Münster, Milano
- 10.45** | In vitro effects of Rituximab on peripheral B cell subsets of patients with Myasthenia Gravis
J. Morroni, E. Sabatelli, S. Marini, S. Falso, R. Iorio. Roma
- 11.00** | Lambert-Eaton myasthenic syndrome in a single centre cohort of patients: clinical, neurophysiological features and therapeutic management
A. Pugliese, A. Barbaccia, F. Biasini, O. Musumeci, S. Messina, A. Toscano, C. Rodolico. Messina
- 11.15** | Early administration of Rituximab improves clinical outcome of Myasthenia Gravis: a single center cohort study
E. Latini, M. Maestri Tassoni, M. Guida, M.C. Caselli, G. Siciliano, R. Ricciardi. Pisa





POSTER SESSION

ITALIAN ASSOCIATION
of MYOLOGY

Poster Session

SMA and motor neuron disorder

Chairpersons: *Megi Meneri (Milano), Federica Trucco (Milano)*

P1 - Newborn with SMA II treated with Zolgensma: between expectations and the real life after one year of follow-up

S. Siliquini, E. Cesaroni, S. Cappanera, I. Cursio, G. Pantalone, C. Marini.
Ancona

P2 - Preliminary data on safety and efficacy of Risdiplam treatment in a small cohort of adult 5q spinal muscular atrophy

A. G. Nanni, G. Milella, G. Piccirilli, M. Ucci, S. Idrissi, A. Fraddosio, A. Introna, V. Scacco, M. Megna, I. Simone, E. D'Errico.
Bari

P3 - Profile of cognitive abilities in spinal muscular atrophy type II and III

B. Buchignani, G. Cicala, F. Moriconi, M. Ricci, A. Capasso, G. Coratti, J. Casiraghi, E. Albamonte, P. Cristofani, C. Cutrona, M. C. Pera, L. Antonaci, C. Roncoroni, D. Chieffo, V. A. Sansone, R. Battini, M. Pane, E. Mercuri.
Pisa, Roma, Milano

P4 - Nusinersen in Children with Spinal Muscular Atrophy: Experience of a Pediatric Pain Center

C. Agosto, A. Salerno, F. Benedetti, I. Maghini, A. Divisic, F. Rusalen, A. Zanin, A. Avagnina, G. Perilongo, F. Benini.
Padova

P5 - Motor unit number estimation in distal muscles of pediatric and adult patients with spinal muscular atrophy

V. Vacchiano, F. Morabito, C. Faini, R. Not, G. Scarpini, A. Pini, R. Liguori.
Bologna

P6 - Long-term comparative efficacy and safety of risdiplam versus nusinersen in children with Type 1 spinal muscular atrophy

N. Hawkins, G. Sajeev, R. Evans, A. Mahajan, D. A. Scott, J. Nam, E. Gaki, S. Sutherland, C. Kokaliaris
Oxford, Glasgow, Boston, London, Basel, Welwyn Garden City

P7 - Phenotypic spectrum of three unrelated SMA Italian patients with a compound heterozygosity for a deletion and rare missense mutation in SMN1 gene

A. Russo, D. Zoppi, T. Fioretti, R. P. Bencivenga, S. Vallone, V. Maiolo, R. Iodice, G. Esposito, L. Ruggiero.
Napoli

P8 - A novel mutation in MYH14 associated with distal spinal muscular atrophy

P. Ajdinaj, M.G. Rispoli, L. Ferri, A. Tessa, F.M. Santorelli, A. Di Muzio.
Chieti, Pisa, Pescara

P9 - New challenges in the treatment era of Spinal muscular atrophy, a single centre experience with risdiplam

C. Dosi, E. Minacapilli, R. Zanin, M.T. Arnoldi, R. Masson.

Milano

P10 - The rehabilitation project and the motor functional scales in three patients with sma type 1 treated with nusinersen: an integrated approach to better understand new phenotypes

N. Sommella, R. Not, G. Scarpini, M. Giannotta, A. Pini, A. Cersosimo.

Bologna

P11 - Using icf-based functional profile to guide development of new measures for identifying treatment effects in sma patients receiving disease-modifying therapies

M.C. Oliva, C. Ferrante, I. Gallo, T. Vespino, A. Santoro, I. Maniglio, G. Barraco, F. Distante,

A. Trabacca.

Brindisi

P12 - Preliminary results from application of EnduSMA protocol to detect endurance and fatigability in spinal muscular atrophy

R. Chiappini, G. Ricci, F. Torri, A. Govoni, L. Fontanelli, F. Magri, V. Vacchiano, M. Coccia,

G. P. Comi, R. Liguori, G. Siciliano.

Ancona, Bologna, Milano, Pisa

P13 - A potential biomarker in Spinal Muscular Atrophy: serum creatinine in nusinersen treated patients SMA

G. Placidi, C. Dosi, R. Zanin, E. Ciusani, R. Masson

Milano

P14 - A case of SMA type 0: clinical findings and literature review

M. Capelli, G. Scarpini, V. Laganà, M. Giannotta, L. Landolina, J. Sarajlija, I. Donati, C. Graziano,

G. Ancora, D. Cordelli, A. Pini.

Bologna, Cesena, Rimini

P15 - 3D-stem cell spinal cord model: a new tool to study risdiplam tool compound mechanisms and therapeutic effects for Spinal Muscular Atrophy

A. D' Angelo, P. Rinchetti, I. Faravelli, F. Beatrice, M. Nizzardo, F. Rizzo, L. Ottoboni, S. Corti.

Milano

P16 - SUNFISH Parts 1 and 2: 4-year efficacy and safety data of risdiplam in Types 2 and 3 SMA

M. Oskoui, J. W. Day, N. Deconinck, E. S. Mazzone, A. Nascimento, K. Saito, C. Vuillerot, G. Baranello, O. Boespflug-Tanguy, N. Goemans, J. Kirschner, A. Kostera-Pruszczyk, M. C. Pera, L. Servais, J. Braid, M. Gerber, K. Gorni, C. Martin, W. Y. Yeung, R. S. Scalco, E. Mercuri, on behalf of the SUNFISH Study Group.

Montreal, Palo Alto, Brussels, Ghent, Rome, Barcelona, Tokyo, Lyon, London, Milan, Paris, Leuven, Freiburg, Warsaw, Oxford, Basel, Welwyn Garden City

P17 - Risdiplam in type 2 and 3 Spinal Muscular Atrophy: results of a cohort of adult Italian patients

D. Zoppi, R. P. Bencivenga, S. Tozza, A. Russo, A. Nevano, R. Iodice, R. Dubbioso, G. Aceto, F. Manganelli, L. Ruggiero.

Napoli

P18 - Serum MyomiRs: potential biomarkers for adult SMA patients upon nusinersen treatment

C. Malacarne, F. Saraceno, R. Zanin, F. Andreetta, R. Mantegazza, L. Maggi, S. Marcuzzo, S. Bonanno.
Parma, Milano

P19 - Multidisciplinary pathway for the management of the first orally therapy for spinal muscular atrophy: a pediatric regional hospitals network experience

L. Pivato, C. Agosto, S. Paccagnella, F. Benedetti, F. Temporin, G. Perilongo, F. Venturini, F. Benini.
Padova

P20 - Clinical and demographic features of patients with SMA on treatment with risdiplam: the itaSMAC experience

E. Albamonte, L. Maggi, E. Pegoraro, S. Messina, M. Sframeli, A. Lizio, M. Pane, R. Masson, C. Dosi, A. D'Amico, A. Pini, G. Scarpini, F. Ricci, T. Mongini, A. Zanolini, G. Coratti, E. Mercuri, V. A. Sansone, on behalf of ItaSMAC.
Bologna, Messina, Milano, Padova, Roma, Torino

P21 - Pregnancy and therapy discontinuance in Spinal Muscle Atrophy: what we want to know. Discussion of a case report.

G. Ricci, L. Fontanelli, R. Chiappini, F. Torri, M. Gherardi, G. Siciliano.
Pisa

P22 - JEWELFISH: 24-month safety, pharmacodynamic and exploratory efficacy data in non-treatment-naïve patients with SMA receiving treatment with risdiplam

C. Bruno, C. A. Chiriboga, T. Duong, D. Fischer, J. Kirschner, M. Scoto, E. Mercuri, M. Gerber, K. Gorni, H. Kletzl, I. Carruthers, C. Martin, T. Gidaro, F. Muntoni, on behalf of the JEWELFISH Study Group
Genova, Palo Alto, Basel, Freiburg, London, Rome, Welwyn Garden City

MYOPATHIES

Chairperson: Andrea Barp (Trento)

P23 - A novel frameshift mutation in HSPB8 gene is responsible for a severe progressive myopathy with cardiac involvement

V. Tudisco, M. Sframeli, L. Bosco, V. Greco, R. Matera, C. Stancanelli, C. Allegra, M. Russo, I. Said, A. Lentini, S. Briuglia, F. Fattori, C. Rodolico, S. Messina.
Messina, Roma.

P24 - Associated SMN5q and MCM3AP gene mutation in a case with distal neuromuscular involvement

M. Giannotta, F. Palombo, R. Pugliano, G. Scarpini, M.L. Valentino, R. Plasmati, F. Pastorelli, M. Maffei, D.M. Cordelli, A. Pini.
Bologna

P25 - TMEM43 mutations: new variants and different phenotypes

D. Cassandrini, G. Marchetto, D. Lopercolo, A. Malandrini, C. Rodolico, P. Tonin, G. Vattermi, F.M. Santorelli.
Messina, Pisa, Siena, Verona

P26 - A novel mutation in LDB3 gene in a patient with Oculopharyngeal Muscular Dystrophy

G. Vadi, F. Torri, F. Baldinotti, G. Ali, G. Cenacchi, G. Ricci, G. Siciliano.

Bologna, Pisa

P27 - Clinical and molecular characterization of two siblings affected by Brody Myopathy

D. Velardo, S. Antognozzi, M. Rimoldi, S. Pagliarani, F. Cogiamanian, S. Barbieri, S. Corti, G. P. Comi, D. Ronchi.

Milano

P28 - Clinical and pathological findings in a couple of Italian Gypsy siblings with Charcot-Marie-Tooth type 4D and a review of the current literature

C.R. Ferrari Aggradi, S. Corti, D. Gagliardi, F. Magri, D. Velardo, G. P. Comi, E. Abati.

Milano

P29 - Centronuclear myopathy caused by a novel heterozygous *BIN1* variant: clinical, histological and MRI evaluation

F. Cortese, M. Brienza, L. De Giglio, A. Todisco, M. Garibaldi, F. Fattori, C. Bernardi, M.C. Altavista, E.M. Pennisi.

Roma

P30 - *HACD1*- Related myopathy with centronuclear myopathy and positive response to l-carnitine supplementation

G. Capece, P. Riguzzi, M. Villa, V. Zangaro, E. Sogus, M. Penzo, D. Gorgoglione, D. Sabbatini, S. Vianello, C. Bertolin, L. Bello, E. Pegoraro.

Padova

P31- Muscle MRI findings in a pediatric case of congenital fiber type disproportion associated with mutation in *TPM3* gene

S. Fusco, S. Parravicini, M. Paoletti, A. Iosca, M.I. Dainesi, C.A. Quaranta, R. Nicotra, V. Vacchini, A. Gardani, S. Gana, A. Berardinelli, A. Pichiecchio.

Pavia

P32 - X-linked Myopathy with Excessive Autophagy: Natural History of a French case series

G. Alfieri, G. Fernandez Eulate, I. Ackermann-Bonan, M. Spinazzi, F. Duval, G. Sole, A. Magot, F. Caillon, A. Pegat, E. Salort-Campana, A. Behin, T. Stojkovic.

Paris, Roma, Angers, Bordeaux, Nantes, Lyon, Marseille

P33 - The Italian myotilinopathy population

A. Lupi, P. Riguzzi, S. Bortolani, M. Filosto, R. Liguori, S. Ravaglia, G. Ricci, F.M. Santorelli, P. Tonin, G.N. Vattermi, L. Maggi, G. Tasca, L. Bello, E. Pegoraro.

Bologna, Brescia, Crotone, Milano, Pavia, Padova, Pisa, Verona

P34 - Clinical and histological spectrum of axial myopathy: A retrospective study

G. U. Borin, F. Lanzafame, G. Marchetto, M. Tosi, S. Piffer, S. Gibertini, D. Cassandrini, F. Santorelli, L. Maggi, G. Vattermi, P. Tonin.

Milano, Verona, Pisa

P35 - Hyperckemia and CNS involvement in a 2 yrs old child: when the things get harder.

A. Iosca, S. Fusco, M.I. Dainesi, R. Nicotra, A. Gardani, S. Dogliani, S. Parravicini, V. De Giorgis, C. Varesio, S. Orcesi, A.L. Berardinelli.

Pavia

P36 - Primary triadopathies: the *STIM1*, *Orai1* and *STAC3* players.

P. Riguzzi, L. Bello, A. Fusto, M. Villa, V. Zangaro, G. Capece, S. Tripodi, G. Minervini, M. Suman, S. Vianello, C. Agosto, G. Sorarù, C. Bertolin, V. Nigro, S. Tosatto, G. Cenacchi, R. Stramare, C. Reggiani, E. Pegoraro.
Bologna, Padova, Napoli



Myasthenia Gravis and Congenital Myasthenic Syndromes

Chairperson: *Giovanni Antonini (Roma)*

P37- Co/Q-Related Congenital Myasthenic Syndrome: A Clinical And Genetic Report

E. Olivieri, B. Risi, S. Cotti Piccinelli, F. Caria, S. Damioli, B. Labella, L. Ferullo, L. Poli, A. Padovani, M. Filosto.
Brescia

P38 - Long-Term Safety, Tolerability, and Efficacy of Efgartigimod in Patients with Generalized Myasthenia Gravis: Concluding Analyses from the ADAPT+ Study

M. Pasnoor, V. Bril, C. Karam, S. Peric, J. L. De Bleecker, H. Murai, A. Meisel, S. Beydoun, T. Vu, P. Ulrichs, B. Van Hoorick, C. T'joen, K. Utsugisawa, J. Verschuuren, R. Mantegazza, J. F. Howard Jr, in collaboration with the ADAPT Investigator Study Group.
Kansas, Toronto, Philadelphia, Belgrade, Ghent, Tokyo, Berlin, Los Angeles, Hanamaki, Leiden, Milan, Chape Hill

P39 - Usefulness and reliability of a self-monitoring diary for the management of patients with myasthenia gravis

F. Stragliati, A. Nuredini, P. Anceschi, S. Romano, S. Tinchelli, E. Chierici, E. Saccani.
Parma

P40 - Immune checkpoint inhibitors-induced Myasthenia Gravis: the need to stratify patients

E. Scarsi, S. Massucco, E. Faedo, C. Gemelli, M. Garnero, C. Genova, E. T. Tanda, C. Dellepiane, L. Benedetti, A. Schenone, M. Grandis.
Genova

P41 - Cancer frequency in muscle-specific tyrosine kinase (MuSK) myasthenia gravis

S. Falso, S. Marini, E. Sabatelli, A. Evoli, R. Iorio.
Roma

P42- Eculizumab for Immune Checkpoint Inhibitors (ICIs)-related Myasthenia Gravis

E. Rossini, A. Lauletta, P. Marchetti, G. Antonini, S. Morino, M. Garibaldi, L. Fionda.
Roma

P43 - Clinical and genetic heterogeneity in Congenital Myasthenic Syndromes: a pediatric case series

F. Trentin, B. Salce, G. Scarpini, M. Giannotta, G.M. Nocera, F. Pastorelli, F.M. Santorelli, F. Palombo, M.L. Valentino, D. Frattini, C. Fusco, V. Di Pisa, D.M. Cordelli, A. Pini.
Pisa, Bologna, Reggio Emilia

P44 - Lambert-Eaton myasthenic syndrome (LEMS) following SARS-CoV2 infection.

A. Barbaccia, A. Pugliese, F. Biasini, F. Andreetta, A. Migliorato, C. Rodolico.
Messina, Milano

P45 - Ravulizumab for the treatment of generalized myasthenia gravis: timing of response

R. E. Mantegazza, A. A. Habib, M. Benatar, T. Vu, A. Meisel, S. Attarian, M. Katsuno, S. Liao, K. N. Beasley, J. F. Howard Jr.
Irvine, Miami, Tampa, Berlin, Marseille, Nagoya, Boston, Chapel Hill, Milan

P46 - Clinical characterization in juvenile myasthenia gravis: a single centre experience.

F. Bruschi, A. Ardisson, C. Ciano, C. Antozzi, R. Mantegazza, I. Moroni.
Milano

P47 - A triple seronegative, refractory patient with Generalized Myasthenia Gravis successfully managed with Efgartigimod

Y.M. Falzone, B. Sorrenti, C. Laurini, C. Strano, L. Bosco, P. Schito, M. Scarlato, R. Fazio, M. Filippi, S. Previtali.
Milano

P48 - Long-term safety and efficacy of Eculizumab in generalized Myasthenia gravis: the 7 years, experience of one patient at Sant'Andrea Hospital of Rome

A. Lauletta, L. Tufano, E. Rossini, S. Morino, M. Garibaldi, L. Fionda.
Roma

P49 - A novel mutation of ColQ in a case of Congenital Myasthenic Syndrome

A. Lupica, P. Alonge, A. Torrente, E. Borgione, C. Scuderi, G. Crescimanno, F. Brighina.
Palermo, Troina

P50 - Light sensitivity in myasthenia gravis: clinical characteristics, pupillometry findings and impact on quality of life

E. Sabatelli, L. Bonagura, S. Falso, S. Marini, G. Della Marca, R. Iorio.
Roma

P51 - Anti-FcRn treatment with Efgartigimod in generalized Myasthenia Gravis

C. Antozzi, R. Frangiamore, E. Rinaldi, F. Vanoli, F. Andreetta, E. Ciusani, S. Bonanno, L. Maggi, A. Gallone, A. Pinna, R. Mantegazza.
Milano

Metabolic Myopathies

Chairperson: Andrea Martinuzzi (Treviso)

P52 - A NOVEL MISSENSE VARIANT IN THE ACADVL GENE IN A PATIENT WITH VLCAD DEFICIENCY: A CLINICAL AND GENETIC STUDY

F. Caria, S. Cotti Piccinelli, S. Damioli, B. Risi, E. Bertella, V. Bonito, B. Labella, L. Poli, A. Padovani, M. Filosto.
Brescia

P53 - Cardiopulmonary exercise testing as outcome measure in primary mitochondrial myopathy: comparison with other functional measures

I. G. Arena, C. Usbergo, C. Consulo, C. Rodolico, A. Toscano, O. Musumeci.
Messina

P54 - A novel compound homozygous ACO2 mutation in an infant with fatal progressive encephalopathy: biochemical characterization on muscle tissue

A. Somà, F. S. Ricci, S. Stanga, M. Mezzanotte, C. Marinaccio, R. D'Alessandro, S. Sottemano, R. Morana, M. Spada, M. Boido, T. E. Mongini.

Torino

P55 - A longitudinal analysis in a cohort of juvenile patients with late onset Pompe disease (LOPD)

M. Porcini, O. Musumeci, I. G. Arena, C. Consulo, C. Usbergo, C. Rodolico, A. Toscano
Messina

P56 - The ER-MITO (Emilia Romagna-Mitochondrial) project: prevalence and genetics of Chronic Progressive External Ophthalmoplegia (CPEO) in an Italian region

M. L. Valentino, L. Caporali, C. La Morgia, F. Palombo, M. Romagnoli, C. Fonti, A. Maresca, R. Liguori, C. Zenesini, R. D'Alessandro, V. Carelli, on behalf of the ER-MITO study group.

Bologna

P57 - Accidental LOPD diagnosis in a young man with no clinical and instrumental evidence of neuromuscular dysfunction

M. Sciacco, S. Lucchiari, M. Garbellini, G. P. Comi, D. Ronchi.

Milano

P58 - Expanding the spectrum of clinical presentations associated with COA8 pathogenic variants

S. Antognozzi, F. Magri, M. Meneri, M. Garbellini, S. Salani, F. Fortunato, M. Ripolone, S. Zanotti, P. Ciscato, M. Sciacco, V. Parente, S. Corti, G. P. Comi, D. Ronchi.

Milano

P59 - A primary cardiological phenotype caused by an inherited mtDNA single deletion: a case report from an Italian pedigree

P. Lopriore, C. Neuhofer, V. Montano, A. Meli, A. Lo Gerfo, M. A. Caligo, R. Berutti, R. Kopajtich, G. Siciliano, H. Prokisch, M. Mancuso.

Pisa, Munich, Neuherberg

P60 - Sudoscan in Pompe disease: a preliminary study

A. Lupica, P. Alonge, A. Torrente, V. Di Stefano, F. Brighina, G. Crescimanno

Palermo

LGMD and FSHD

Chairperson: *Giovanna Lattanzi (Bologna)*

P61 - Observational study: the quality of life in patients with alpha-sarcoglycan, beta-sarcoglycan and gamma-sarcoglycan gene mutation

*B. Vola, M. Cerletti, Y. Torrente, C. Sanchez Riera, M. Bianchi.
London, Milano, Roma*

P62 - A potential treatment for sarcoglycanopathy: pharmacological profile of C17 CFTR corrector and its efficacy with a new regimen of administration

*A. Benetollo, M. Scano, S. Parrasia, L. Biasutto, F. Dalla Barba, M. Carotti, L. Nogara, B. Blaauw, D. Sandonà .
Padova*

P63 - Validation of a blood-based assay for dysferlinopathy in a large Indian cohort

*D. Cox, M. Henderson, Y. Parkhurst, P. Gaitonde, R. Dastur, S.S. Emmons, R. Barresi.
Newcastle, Seattle, Venice*

P64 - A novel c.1478G>A calpain 3 gene variant associated to autosomal dominant calpainopathy

*E. Faedo, S. Massucco, C. Gemelli, C. Fiorillo, M. Moggio, M. Ripolone, E. Bellone, S. Patrone, A. Mammi, A. Geroldi, A. Gaudio, P. Mandich, F. Gotta, E. Scarsi, A. Schenone, M. Grandis.
Genova, Milano*

P65 - Facioscapulohumeral dystrophy: an epidemiologic investigation in Abruzzo

*G. Polito, C. Ciprietti, S. Melchiorre, P. Adjnaj, A. Di Muzio.
Chieti*

P66 - A four-years clinical follow-up and muscle MRI study in FSHD towards trials readiness

*F. Torri, G. Aringhieri, L. Caldararo, S. Ortore, G. Siciliano, G. Ricci.
Pisa*

P67 - A novel LMNA mutation causing atypical early-onset EDMD, with Achilles tendon contractures and limb-girdle weakness: diagnostic and therapeutic considerations

*E. Minacapilli, C. Dosi, R. Zanin, F. Ricci, S. Gibertini, L. Maggi, R. Masson.
Milano, Torino*

P68 - A novel case of TRIM32-related Limb Girdle Muscular Dystrophy

*M. Rimoldi, F. Magri, S. Antognozzi, G. Romagnoli, D. Velardo, V. Parente, C. Cinnante, S. Corti, G. P. Comi, D. Ronchi.
Milano*

P69 - A retrospective analysis of motor and respiratory progression in a population of FSHD1 patients: how fast does the disease go?

*G. Colacicco, E. Carraro, L. Martinelli, E. De Mattia, E. Roma, M. Croci, A. Lizio, V.A. Sansone.
Milano*

P70 - LGMDR19: A MILD GMPPB-LINKED DISEASE PHENOTYPE

*L. Ferullo, S. Cotti Piccinelli, F. Caria, B. Risi, S. Damioli, B. Labella, E. Olivieri, L. Poli, A. Padovani, M. Filosto.
Brescia*

P71 - Calpain and Becker dystrophy: a double trouble

A. Lupica, P. Alonge, A. Torrente, E. Borgione, C. Scuderi, G. Crescimanno, F. Brighina.
Palermo, Troina

P72 - LMGD-D2 TNPO3 related: A quality of life study

C. Angelini, A. A. Rodríguez.
Padova, Bilbao

P73 - Muscular and peripheral immunological profiles of patients affected by LGMDR3

C. Panicucci, A. Amaro, S. Pintus, F. Antonini, F. Reggiani, S. Baratto, E. Principi, C. Fiorillo, G. Del Zotto, D. Cassandrini, A. D'Amico, D. Lopercolo, A. Malandrini, E. Malfatti, G. Urbano, T. Mongini, A. Fusto, E. Pegoraro, S. Previtali, C. Rodolico, G. Tasca, U. Pfeffer, C. Bruno, L. Raffaghello.
Genova, Pisa, Roma, Siena, Créteil, Torino, Padova, Messina, Newcastle, Milano



Dystrophinopathies

Chairperson: Emilio Albamonte (Milano)

P74 - Feasibility of switch from prednisone to vamorolone in patients with DMD in VBP15-004 study

G. J. Van Daal, P. K. Clemens, E. C. Smith, I. Horrocks, R. Finkel, J. Mah, N. Deconinck, L. De Waele, V. Straub, G. Baranello, S. Spinty, A.M. Childs, J. Vilchez-Padilla, A. Nascimento-Osorio, E. Niks, I. de Groot, M. Katsaloulis, E. Hoffman, M. Guglieri on behalf of the VISION-DMD.
Pratteln, Glasgow, Orlando, Calgary, Brussels, Newcastle, London, Leeds, Valencia, Barcelona, Nijmegen, Rockville, Leuven, Leiden

P75 - Efficacy and safety of vamorolone during 48-week treatment in patients with Duchenne Muscular Dystrophy (DMD) in VBP15-004 study

G. J. Van Daal, E. P. Hoffman, P.R. Clemens, S.J. Perlman, E.C. Smith, I. Horrocks, R.S. Finkel, J.K. Mah, N. Deconinck, N. Goemans, J. Haberlova, V. Straub, M.M. Ryan, M. Tulinus, R. Webster, H.J. McMillan, G. Baranello, S. Spinty, A.M. Childs, K.A. Selby, J.J. Vilchez-Padilla, A. Nascimento-Osorio, E.H. Niks, I.J.M. de Groot, M. Katsaloulis, M. Guglieri on behalf of the VISION-DMD.
Pratteln, Pittsburgh, Durham, Orlando, Calgary, Brussels, Prague, Newcastle, Gothenburg, Ottawa, London, Liverpool, Vancouver, Valencia, Leiden, Athens, Sydney, Leuven, Barcelona, Nijmegen

P76 - Linkage analysis in a family with recurrence of duchenne muscular dystrophy defined the at risk haplotype and identified a clue of a more complex genomic rearrangement

M. Neri, L. Fiocco, F. Fortunato, M. Fabris, A. Margutti, C. TrabANELLI, P. Rimessi, R. Selvatici, F. Gualandi, A. Ferlini.
Ferrara

P77 - Ergoreflex Sensitivity is Associated with Cardiac Involvement in Becker Muscular Dystrophy

V. Castiglione, P. Sciarone, F. Torri, C. Borrelli, A. Barison, G. Todiere, C. Grigoratos, G. Siciliano, C. Passino, M. Emdin, G. Ricci, A. Giannoni.
Pisa

P78 - The spine fracture burden in boys with DMD treated with the novel dissociative steroid vamorolone versus deflazacort and prednisone

R. Rooman, M. Guglieri, S. Jackowski, U. Dang, M. Scharke, J. L. Jaremko, K. Koujok, M. Matzinger, N. Shenouda, K. Siminoski, M. Leinonen, S. Hasham, P. Clemens, M. McDermott, R. Griggs, E. Hoffman, L. Ward on behalf of the Ottawa Pediatric Bone Health Research Group, the FOR DMD Investigators of the Muscle Study Group and the CINRG VBP-/LTE Investigators.
Pratteln, Ottawa, New York, Newcastle, Rockville

P79 - Daily regimens of prednisone, deflazacort and vamorolone improve motor function similarly in patients with Duchenne Muscular Dystrophy

M. Guglieri, G.J. Van Daal, R. Rooman, C. McDonald, E. Henricson, A. Linden, P. Clemens, R. Griggs, P. Shieh, I. Horrocks, J. Mah, R. Finkel, N. Goemans, V. Straub, M. Ryan, H. McMillan, S. Spinty, E. Hoffman, on behalf of the VISION-DMD, FOR-DMD and CINRG investigators.
Newcastle, Pratteln, Sacramento, Pittsburgh, Los Angeles, Calgary, Memphis, Melbourne, Ottawa, Liverpool, Binghamton, New York, Leuven

P80 - Longitudinal forced vital capacity in Becker muscular dystrophy patients from the Padova cohort

M. Penzo, L. Bello, P. Riguzzi, M. Villa, G. Capece, D. Sabbatini, D. Gorgoglione, S. Vianello, A. Vianello, E. Pegoraro.
Padova

P81 - Retrospective study on the trend of NSAA and 6MWT scores of a cohort of DMD patients treated with cortisone according to age of initiation of therapy

M. I. Dainesi, A. Gardani, A. Iosca, S. Fusco, C.A. Quaranta, G. Catalano, R. Nicotra, L. Carraro, S. Parravicini, A. L. Berardinelli.
Pavia

P82 - DMD in females: a case series of eleven symptomatic carriers

G. D'Amario, I. Venezia, G. Cordaro, C. Guidetti, V. Napoli, A. L. Frongia, M. Pane, E. Mercuri.
Roma

P83 - Motor Functional findings and muscle MRI patterns of DMD patients with small mutations

C. Brogna, L. Cristiano, T. Verdolotti, G. Tasca, L. Ficociello, A. Frongia, N. Forcina, R. Ferrante, G. Stanca, M. Pane, E. Mercuri.
Roma

P84 - Genome-Wide Association Study for the identification of genetic modifiers of cardiomyopathy in Duchenne muscular dystrophy

D. Sabbatini, D. Gorgoglione, M. Penzo, S. Vianello, P. Riguzzi, M. Villa, A. Fusto, G. Capece, A. Berardinelli, C. Bruno, C. Panicucci, G. P. Comi, F. Magri, A. D'Amico, M. Catteruccia, L. Travaglioni, G. D'Angelo, V. Sansone, E. Albamonte, A. Di Bari, T. Mongini, C. Brusa, L. Maggi, E. Canioni, A. Gallone, M. Pane, D. Leone, L. Politano, E. Picillo, V. Nigro, S. Messina, M. Sframelì, G. Vita, R. Masson, S. Parravicini, G. Sorarù, L. Bello, E. Pegoraro.
Padova, Genova, Milano, Bosisio Parini, Torino, Roma, Napoli, Messina, Pavia

P85 - Poor Bone Health in Patients with Duchenne Muscular Dystrophy

E. Carraro, L. Martinelli, M. Pane, C. Palermo, A. Lizio, F. Trucco, E. Albamonte, E. Mercuri, V. A. Sansone.
Milano, Roma



Disease Mechanisms and Pathogenesis

Chairperson: *Linda Ottoboni (Milano)*

P86- Defining molecular and functional consequences of titin mutations in muscle progenitors from affected patients

C. Fiorillo, S. Baratto, S. Baldassari, E. Sondo, M. Traverso, G. Astrea, A. Biquand, E. Malfatti, M. Savarese, C. Bruno, N. Pedemonte.
Genova, Pisa, Parigi, Helsinki

P87 - Zebrafish models as new tools to study Sarcoglycanopathies

F. Dalla Barba, M. Carotti, A. Benetollo, M. Scano, P. Caccin, F. Argenton, D. Sandonà.
Padova

P88 - Circulating muscle-derived miR-206 links skeletal muscle dysfunction to heart sympathetic denervation

M. Ronfini, V. Di Mauro, V. Prando, A. Franco-Romero, A. Mazzaro, G. Favaro, F. Lo Verso, L. Dokshokova, A. Armani, G. Sorarù, M. Guescini, D. Catalucci, M. Mongillo, M. Sandri, T. Zaglia.
Padova, Milano, Urbino

P89 - Spastic paraparesis and myopathic signs associated to *POLRMT* mutations

F. Magri, S. Antognozzi, M. Meneri, F. Fortunato, M. Garbellini, M. Ripolone, A. Giacobbe, L. Napoli, G. P. Comi, D. Ronchi.
Milano

P90 - Whole miRNome profiling in myasthenia gravis thymus: novel insights into the molecular events underlying autoimmunity

M. C. Tarasco, N. Iacomino, L. Scandiffio, F. Bortone, S. Marcuzzo, S. Bonanno, R. Frangiamore, D. Pistillo, E. Voulaz, M. Alloisio, L. Maggi, C. Antozzi, R. Mantegazza, P. Cavalcante
Milano, Monza

P91 - MicroRNAs as biomarkers of responsiveness to immunosuppressive drugs in myasthenia gravis: towards personalized medicine

N. Iacomino, M. C. Tarasco, M. Ballardini, R. Frangiamore, F. Vanoli, L. Maggi, C. Antozzi, M. Foti, R. Mantegazza, P. Cavalcante.
Milano, Monza

P92 - Evaluation of aggrephagy markers in genetically defined myofibrillar myopathies' patients

G. Riolo, E. Iannibelli, S. Gibertini, A. Ruggieri, A. Carnazzi, M. Cheli, G. Tasca, S. Bortolani, L. Maggi.
Milano, Roma

P93 - Nuclear platform reorganization and nuclei orientation are lost in EDMD2 human myoblasts subjected to mechanical stimulation

V. Cenni, C. Evangelisti, S. Neri, P. Sabatelli, M. Cavallo, G. Lattanzi, E. Mattioli.
Bologna

P94 - Differential dysferlin expression in human and rat fast and slow skeletal muscle suggests different role

M. Meznaric, C. Angelini.
Padova



P95 - Transcriptomic and morphological alterations in hTNPO3 MUT-microinjected Zebrafish embryos modelling Limb Girdle Muscular Dystrophy D2 in vivo

M. Bergonzoni, M.T. Rodia, R. Costa, S. Pacilio, M. Fazzina, R. Casadei, F. Frabetti, G. Cenacchi. Bologna, Rimini

P96 - Emerging role of myomiRs as biomarkers and therapeutic targets in Emery-Dreifuss muscular dystrophies

S. Bonanno, C. Malacarne, E. Mattioli, F. Saraceno, P. Cavalcante, L. Santovito Spiro, M. Cheli, C. Fiorillo, F. Andreetta, R. Mantegazza, S. Marcuzzo, L. Maggi, G. Lattanzi. Bologna, Chicago, Genova, Parma

P97 - Microbiota Dysbiosis influences Immune System and Muscle Pathophysiology of Dystrophin Deficient Mice

A. Farini, L. Tripodi, C. Villa, F. Strati, B. Cassani, F. Caprioli, F. Facciotti, M. Quattrocelli, Y. Torrente. Milano, Cincinnati

P98 - Efficient decellularization method for the accomplishment of engineered human skeletal muscle

S. D'Agostino, F. Magarotto, G. Tafuro, G. Germano, C. Biz, A. Semenzato, P. Gamba, M. Muraca, M. Pozzobon. Padova



Diagnostic and Therapeutic tools

Chairperson: Rita Barresi (Venezia)

P99 - Attenuated total reflection-Fourier transform infrared spectroscopy (ATR-FTIR) in muscle diseases

G. Primiano, C. Sancricca, A. Primiano, J. Gervasoni, A. Urbani, F. Marini, A. Sabino, R. Calvani, S. Servidei.

Roma

P100 - Eye tracking communication device opens world of communication for a non-verbal child affected by SMARD 1

A. Ferrero, N. Gironella, F. Re, A. Ubaldi, M. Coccia.

Ancona

P101 - Sympathetic neuropathology is revealed in muscles affected by Amyotrophic Lateral Sclerosis

A. Mazzaro, V. Vita, M. Ronfini, I. Casola, A. Klein, G. Dobrowolny, G. Sorarù, A. Musarò, M. Mongillo, T. Zaglia.

Padova, Roma

P102 - A new 6MWT web-tool to facilitate clinical monitoring of neuromuscular patients

G. Marinella, B. Buchignani, A. Rubegni, S. Frosini, E. Dati, A. Tonacci, R. Conte, F. Sansone, G. Diodato, M.C. Scudellari, F. Torri, G. Ricci, G. Siciliano, R. Battini, G. Astrea, F. M. Santorelli.

Pisa, Roma

P103 - Proposal for a model of care in a pediatric neuromuscular clinic: experience in Trento area

V. Fiorito, A. Iodice, G. Zappa, U. Pradal, R. Zuccarino.

Trento

P104 - Challenging diagnosis between an atypical phenotype of motor neuron disease and myositis: a case report.

L. Becattini, G. Vadi, F. Bianchi, L. Fontanelli, B. Giovannini, C. Meoni, G. Ricci, G. Siciliano.

Pisa

P105 - High plasma p-tau181 levels in amyotrophic lateral sclerosis: clinical-electrophysiological correlations and longitudinal trajectories

V. Vacchiano, A. Mastrangelo, C. Zenesini, S. Baiardi, P. Avoni, B. Polisch, S. Capellari, F. Salvi, R. Liguori, P. Parchi.

Bologna

P106 - An exploratory study on the use of a high-content image analysis system in myopathies

F. Torri, C. Filipponi, B. Ciurli, M. Lai, G. Alì, G. Vadi, M. Pistello, G. Siciliano, G. Ricci.

Pisa

P107 - Exploratory study of a passive wearable device (Exoband) as walking aid in neuromuscular patients

C. Semplicini, M. Agostini, C. Andrigo, V. Notararigo, S. Masiero, F. Piccione, G. Sorarù.

Padova



P108 - Diaphragmatic ultrasound: a promising technique for respiratory assessment of patients with facio-scapulo-humeral muscular dystrophy (FSHD)

E. Torchia, S. Bortolani, R. Inchingolo, A. Smargiassi, M. Monforte, G. Tasca, M. Bonini, L. Richeldi, E. Ricci.
Roma

P109 - Different approaches to the genetic diagnosis of hereditary peripheral neuropathies: a 2 years multicenter study

L. Bosco, M. Catteruccia, D. Diodato, M. Tosi, E. Pennisi, M. Garibaldi, S. Pro, E. S. Bertini, A. D'Amico, F. Fattori.
Roma

P110 - Reduction of Body Cellular Mass and Body Cellular Mass Index as an early marker of sarcopenia in Amyotrophic Lateral Sclerosis patients

G. Greco, L. Boffa, M. Goglia, S. Seraceno, A. M. Ruggieri, G. Merra, L. Romano, E. Frezza, G. Vietri, F. Gruosso, G. Nardino, I. Petitta, P. Leuratti, A. De Lorenzo, R. Massa.
Roma

P111 - Unravelling combined RNA interference and gene therapy in vitro and in vivo disease models as a potential therapeutic strategy for CMT2A

F. Rizzo, S. Bono, M.D. Ruepp, S. Salani, L. Ottoboni, E. Abati, V. Melzi, C. Cordiglieri, S. Pagliarini, R. De Gioia, A. Anastasia, M. Taiana, M. Garbellini, S. Lodato, P. Kunderfranco, D. Cazzato, D. Cartelli, C. Lonati, N. Bresolin, G. P. Comi, M. Nizzardo, S. Corti.
Milan, London

P112 - D4Z4 methylation analysis combined with machine learning pipelines: a novel tool for identifying FSHD subjects

V. Caputo, C. Strafella, G. Trastulli, D. Megalizzi, C. Fabrizio, A. Termine, L. Colantoni, J. Gimenez, M. Monforte, C. Caltagirone, E. Ricci, G. Tasca, R. Cascella, E. Giardina.
Rome, Newcastle

P113 - MethylSeq based assay to assess the epigenetic setting of D4Z4 repetitive elements in facioscapulohumeral muscular dystrophy

V. Salsi, M. Chiara, S. Pini, L. Ruggiero, F. Santorelli, S. C. Previtali, M. G. D'Angelo, C. Rodolico, S. Bonanno, L. Maggi, R. Tupler.
Modena, Worcester, Bari, Napoli, Pisa, Milano, Messina, Bosisio Parini

P114 - The case of Friedreich Ataxia as an example to exploit the pleiotropic effects of nanoparticles

C. Villa, M. Mistretta, A. Farini, Y. Torrente.
Milano

P115 - Pediatric Palliative Care is the answer for met and unmet needs of children with neuromuscular diseases

C. Agosto, F. Benedetti, I. Maghini, P. Lazzarin, F. Benini.
Padova

P116 - IN VITRO ELECTROSTIMULATION ASSAY FOR FUNCTIONAL EVALUATION OF ENGINEERED SKELETAL MUSCLE TISSUE

S. Pacilio, T. Cramer, R. Costa, V. Papa, M. T. Rodia, B. Fraboni, M. L. Focarete, G. Cenacchi.
Bologna

P117 - Unraveling the role of high-resolution ultrasonography of peripheral nerves in adult spinal muscular atrophy

S. Bonanno, G. Devigili, R. Togni, M. Corradi, P. Lanteri, R. Mantegazza, R. Eleopra, L. Maggi. Milano

P118 - Health360 Trials: a software module to manage clinical trials and patients from different clinical institutions

F. Sansone, G. Diodato, M. C. Scudellari, F. M. Santorelli, G. Siciliano, A. Malandrini, S. Matà, C. Angelini, A. Tonacci, The InGene. Project Group, R. Conte. Pisa, Siena, Firenze, Roma, Padova

Congenital Myopathies

Chairperson: *Caterina Agosto (Padova)*

P119 - Novel *PIEZO2* variants are responsible for a very suggestive clinical phenotype in early childhood

D. Diodato, M. Catteruccia, A. Sancesario, M. Tosi, L. Bosco, F. Fattori, G. De Luca, I. Mizzoni, A. Carlesi, E. Bertini, A. D'Amico. Roma

P120 - Cognitive, adaptive and perseverative aspects characterization of children with XLMTM: an explorative study

F. Cumbo, M. Tosi, A. Carlesi, I. Mizzoni, M. Catteruccia, E. S. Bertini, A. D'Amico. Roma

P121 - Collagen VI-related myopathies with onset in pediatric age: the value of clinical findings when not all the diagnostic pieces fit in

G. Guardi, M. Giannotta, G. Scarpini, M. L. Valentino, M. Maffei, F. Palombo, A. Ferlini, D.M. Cordelli, F. Gualandi, A. Pini. Bologna, Ferrara

P122 - Congenital myopathy as a phenotypic expression of *CACNA1S* gene mutation: Case Report and literature systematic review

G. Marinella, A. Orsini, M. Scacciati, E. Costa, G. Astrea, S. Frosini, R. Pasquariello, A. Rubegni, G. Sgherri, M. Corsi, A. Bonuccelli, D. Peroni, R. Battini. Pisa

P123 - Causative variants profile in a large Italian cohort of Collagen VI , related dystrophy

L. Fiocco, M. Neri, A. Margutti, F. Fortunato, A. D'Amico, A. L. Berardinelli, A. Pini, C. Fusco, C. Rodolico, C. Fiorillo, C. Bruno, C. Cereda, E. Pegoraro, E. Bertini, E. Ricci, E. Picillo, E. Mercuri, F. Vercellino, F. Massaro, G. Ciana, I. Moroni, L. Boccone, L. Maggi, L. Morandi, L. Merlini, L. Politano, M. Scutifero, M. Pane, M. Pedemonte, M. Filosto, M. Del Mastio, M. Traverso, P. Gasparini, P. Boffi, R. Massa, S. Messina, T. E. Mongini, R. Selvatici, A. Ferlini, F. Gualandi. Ferrara, Roma, Pavia, Reggio Emilia, Messina, Genova, Padova, Napoli, Prato, Udine, Milano, Cagliari, Bologna, Firenze, Trieste, Torino

P124 - Novel RYR1 pathological variants discovered in Italian patients with dusty core disease

D. Piga, S. Zanotti, F. Magri, S. Salani, L. Napoli, M. Ripolone, F. Fortunato, D. Cassandrini, F. Fattori, M. G. D'Angelo, E. Albamonte, V. Nigro, S. Corti, G. P. Comi.
Milano, Pisa, Roma, Lecco, Napoli

Channelopathies and Inflammatory Myopathies

Chairperson: *Michela Coccia (Ancona)*

P125 - High dose intravenous corticosteroids in refractory Immune Checkpoint Inhibitor-related Myocarditis/Myositis/Myasthenia overlap syndrome

L. Tufano, A. Lauletta, S. Morino, B. Musumeci, G. Antonini, M. Garibaldi.
Roma

P126 - Cognitive function assessment in a cohort of patients with Myotonic Dystrophy Type 1: a single-center experience

B. Risi, S. Cotti Piccinelli, E. Ferrari, F. Caria, S. Damioli, E. Bertella, B. Labella, L. Poli, V. Bonito, A. Padovani, M. Filosto.
Brescia

P128 - Myotonic Dystrophy type 1: motor and cardio-respiratory evolution trajectories

E. Diella, A. LoMauro, M. Delle Fave, G. Canella, G. Meola, R. Cima, M. G. D'Angelo.
Milano, Bosisio Parini

P129 - A novel hypokalemic periodic paralysis mutation in the SCN4A gene shared by two families and description of a homozygous patient

S. Pagliarini, V. Sansone, F. Gerardi, M. Scarlato, S. C. Previtali, S. Lucchiari, G. Meola, G. P. Comi.
Milano, Bergamo

P130 - Beyond diagnostic criteria: the pitfalls in the diagnosis of inflammatory myopathies, a case series

L. Bosco, Y.M. Falzone, V. Canti, P. Rovere Querini, U. Del Carro, M. Filippi, S.C. Previtali.
Milano

P131 - Compound CACNA1S heterozygosity resulting in a novel phenotype of congenital myopathy and early onset periodic paralysis: report of two probands

S. K. Aburahma, M. Shboul, L. A. Rousan, S. Lucchiari, F. Biella, G. P. Comi, G. Meola, S. Pagliarini.
Irbid, Milano

P132 - A case of myasthenia and myositis after Covid vaccine

G. Vadi, F. Torri, M. Maestri, G. Ali, M. Lencioni, G. Siciliano, G. Ricci.
Pisa

P133 - Novel HSPG2 gene mutation causing Schwartz Jampel syndrome in a Moroccan family.

D. Marelli, R. Brugnoli, C. Cappelletti, N. Iacomino, E. Canioni, L. Maggi, A. Ardisson.
Milano

P134 - A case of Amyotrophic Lateral Sclerosis initially misdiagnosed for myositis*C. Meoni, F. Bianchi, L. Becattini, L. Fontanelli, B. Giovannini, G. Siciliano.**Pisa***P135 - Long-term treatment and follow up in adult patients with anti-HMGRC immune-mediated myopathy***L. Vercelli, G. Gadaleta, G. Urbano, G. Brodini, L. Chiadò-Piat, F. Rumbolo, T. Manetta, T. Mongini.*
*Torino***P136 - Pre- and post-natal outcomes in congenital and childhood onset DM1 - the impact of parental diagnostic delay***F. Trucco, A. di Bari, F. Salmin, A. Lizio, E. Albamonte, A. Zanolini, M. Beretta, J. Casiraghi, L. Antonaci, R. de Sanctis, A. Salvalaggio, M. Catteruccia, M. Tosi, G. Marinella, R. Danti, F. Bruschi, A. Ferrero, B. Risi, A. Barp, M. Veneruso, C. Fiorillo, A. Berardinelli, A. Pini, R. Zuccarino, M. Filosto, M. Coccia, I. Moroni, G. Astrea, A. D'Amico, F. Ricci, M. Pane, E. Mercuri, V. Sansone.*
*Milano, Torino, Roma, Pisa, Ancona, Brescia, Trento, Genova, Pavia, Bologna***P138 - Myopathy in the oncologic patient: move beyond muscle inflammation***R. Costa, V. Di Pisa, R. D'Angelo, S. Rossi, R. Rinaldi, G. Cenacchi.**Bologna***P139 - Inclusion body myositis functional rating scale in IBM patients: a cross-sectional study***A. Vicino, M. Cheli, S. Bonanno, L. Maggi.**Milano, Lausanne***P140 - Preclinical Data Support the Initiation of the ACHIEVE Trial of DYNE-101 in Individuals with Myotonic Dystrophy Type 1 (DM1)***E. Delage, E. Cotali, D. Wolf, T. Picariello, L. Schlaefke, R. Russo, A. Chang, S. Hildebrand, J. Najim, Q. Qiu, T. Weeden, J. W. Davis II, C. Mix, B. Han, S. Zanotti, O. Beskrovnaya, W. Farwell.**Paris, Waltham*

GENERAL INFORMATION

CONGRESS VENUE

PADOVA CONGRESS

Via Goldoni 8 - Cancelli C
Padova

PARKING

Parcheggio Nord - Cancelli E
Via Goldoni 6, Padova
€5 per day - Open H24

ORGANIZING SECRETARIAT



First Class - Meetings and Conferences
Sede Milano

Via Vittoria Colonna 40 - 20149 Milano

Ph. +39 02 30066329

aim2023@fclassevents.com

www.fclassevents.com

The Organizing Secretariat will be available to the participants at the congress venue during the following times:

- Thursday, 8th June from 09.00 to the end of the works
- Friday, 9th June from 07.15 to the end of the works
- Saturday, 10th June from 07.30 to the end of the works

AIM SECRETARIAT

The AIM Secretariat Desk will be present at the congress venue (next to the registration area) and will follow the following times

- June 8th from 10:00 to the end of the works
- June 9th from 08:30 to the end of the works

REGISTRATION

The meeting is reserved for a maximum number of 450 participants.

Participation in the event is subject to online registration by Monday 29 May 2023.

After the deadline, it will be possible to register directly at the congress venue.

	By 8 th May 2023	From 9 th May 2023
AIM MEMBER	234,00 €	275,00 €
AIM MEMBERS UNDER 35	152,00 €	193,00 €
NOT AIM MEMBER	328,00 €	394,00 €

VAT (22%) excluded

XXIII NATIONAL CONGRESS OF ITALIAN ASSOCIATION of MYOLOGY

The registration fee includes:

Admission to scientific sessions

Congress Kit

Lunches and coffee breaks

Welcome Cocktail | 8th June 2023

Social Dinner | 9th June 2023

Certificate of attendance

CME certificate (it will be sent after the conclusion of the event after verifying the attendance and learning requirements)

PAYMENT METHOD

Online - Bank transfer and credit card

Onsite - ATM and credit card

CANCELLATIONS AND REFUNDS

Cancellations received by 21 April 2023 entitle to a refund of the amount paid, less 30% for administrative costs.

Cancellations received after this date will not give the right to any refund.

All refunds will in any case be made after the Congress has taken place.

BADGE

At the time of registration, each registered participant will be given a name badge which must always be shown in order to access the classrooms where the scientific sessions are held, the exhibition area, the catering services and the Welcome Cocktail on Wednesday 8th June.

CERTIFICATE OF ATTENDANCE

The certificate of participation can be downloaded online after completing the evaluation questionnaire. The Organizing Secretariat will send an e-mail to all participants on 10th June containing useful information to download it.

CLOAKROOM AND LUGGAGE STORAGE

A cloakroom and left-luggage office will be made available.

WELCOME COCKTAIL

The Welcome Cocktail will be held on Wednesday 8 June at 20:00 on the third floor of the Padova Congress. Please wear your personal Congress badge to access the restricted area.

SOCIAL DINNER

The social dinner will be held on Friday 9th June at 21:00 at:

VILLA MOLIN

Via Ponte della Cagna 106 - Padova

Entry will be permitted only upon presentation of the relative invitation received at the time of registration for the Congress.

Accreditamento ECM

Il **XXIII Congresso Nazionale Associazione Italiana di Miologia**, Padova 8-10 giugno 2023 è inserito nel Piano Formativo anno 2023 First Class - Provider nr. 362 ed è stato accreditato presso la Commissione Nazionale per la Formazione Continua con **Obiettivo Formativo**:

18 - Contenuti tecnico-professionali (conoscenze e competenze) specifici di ciascuna professione, di ciascuna specializzazione e di ciascuna attività ultra-specialistica, ivi incluse le malattie rare e la medicina di genere per le seguenti figure professionali:

PROFESSIONE

Farmacista

Biologo

Logopedista

Terapista Occupazionale

Psicologo

Medico Chirurgo

Infermiere Pediatrico

Fisioterapista

Infermiere

Dietista

Tecnico di Neurofisiopatologia

Tecnico Sanitario di Laboratorio
Biomedico

Terapista della Neuro e Psicomotricità
dell'età Evolutiva

DISCIPLINE

Farmacia Ospedaliera; Farmacia Territoriale;

Biologo

Logopedista

Terapista Occupazionale

Psicoterapia; Psicologia;

*Cardiologia; Genetica Medica; Malattie Metaboliche e Diabetologia; Malattie Dell'apparato Respiratorio; Medicina Fisica e Riabilitazione; Neonatologia; Neurologia; Neuropsichiatria Infantile; Pediatria; Chirurgia Pediatrica; Neurochirurgia; Ortopedia e Traumatologia; Anestesia e Rianimazione; Laboratorio di Genetica Medica; Neurofisiopatologia, Neuroradiologia
Pediatria (pediatri di libera scelta);
Scienza dell'alimentazione e dietetica; Medicina Interna*

Infermiere Pediatrico

Fisioterapista

Infermiere

Dietista

Tecnico di Neurofisiopatologia

Tecnico Sanitario di Laboratorio Biomedico

Terapista della Neuro e Psicomotricità dell'età Evolutiva

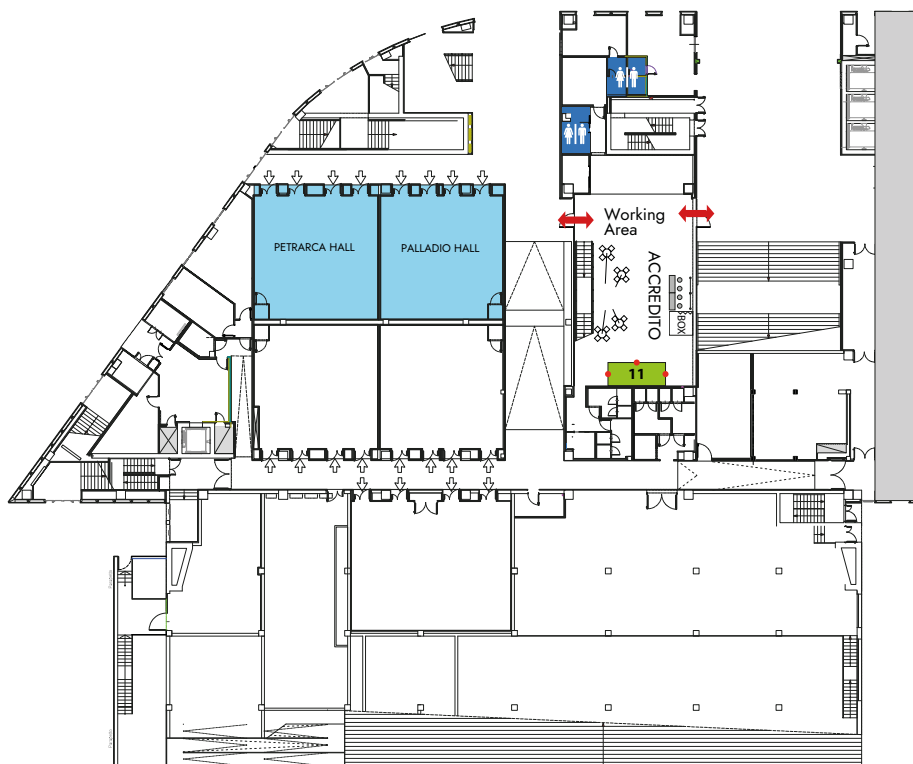
Il congresso è stato accreditato per un numero massimo di n. 450 partecipanti. Oltre tale numero e per professioni/ discipline differenti da quelle accreditate non sarà possibile rilasciare i crediti formativi. Si precisa che i crediti verranno erogati a fronte di una partecipazione del 90% ai lavori scientifici (farà fede esclusivamente l'orario di ingresso e uscita riportato nel foglio firme) e del superamento della prova di apprendimento con almeno il 75% delle risposte corrette.

L'evento, ID ECM 362-383225, ha ottenuto 4,8 crediti formativi.

Exhibition area

Padova Congress

GROUND FLOOR



Exhibition area

Padova Congress

FIRST FLOOR



AIM SECRETARIAT 11

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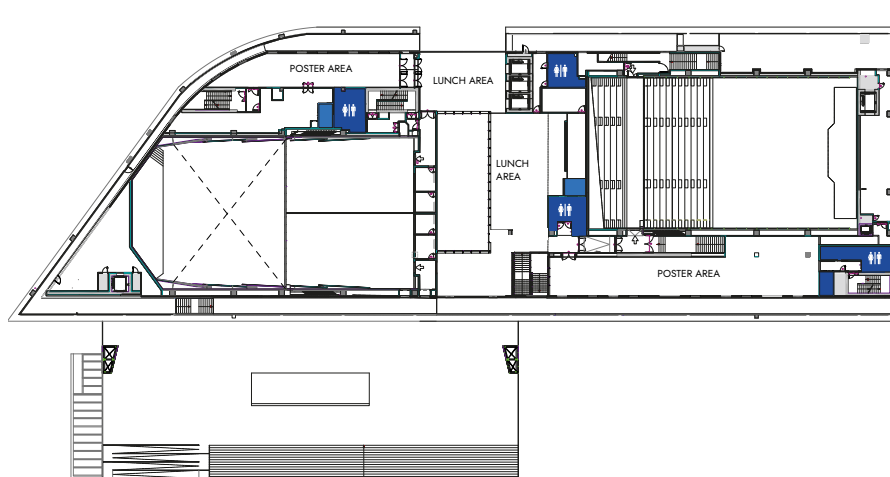
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Exhibition area

Padova Congress

THIRD FLOOR





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XXIII NATIONAL CONGRESS OF ITALIAN ASSOCIATION of MYOLOGY

PADOVA
8th - 10th June 2023

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ORGANIZING SECRETARIAT



FIRST CLASS
EVENTS & CONFERENCES

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