

PROVISIONAL PROGRAMME
(contents subject to change)
Palais des Congrès de Montréal, Canada
6 to 8 December 2024
(on-demand access available until 5 March 2025)

FRIDAY 6 DECEMBER

SESSION 1 JOINT OPENING SESSION

08.30 – 08.35	Welcome – A Al-Chalabi (UK) and B Dickie/N Cole (UK)
08.35 – 08.45	Welcome from Host Association/Dignitary
08.45 – 09.25	Stephen Hawking Memorial Lecture: C1 Distinguishing Myth and Reality in Biomedical Research – T Caulfield (Canada)
09.25 – 09.35	International Alliance Forbes Norris Award
09.35 – 09.50	IPG Award and winner's research presentation
09.50 – 10.00	Longitude Prize announcement

10.00 – 10.30 COFFEE

SESSION 2A GENETICS	SESSION 2B IMPROVING CLINICAL MANAGEMENT	SESSION 2C PRESYMPTOMATIC DETECTION AND EARLY DIAGNOSIS
10.30 – 11.00 C2 Genetics and phenotypic heterogeneity of ALS/MND in Africa and people of African ancestry J Heckmann (South Africa) / D Singh Sokhi (Kenya)	10.30 – 10.50 C9 Dextromethorphan/quinidine (DMQ) in the treatment of bulbar symptoms in ALS: Patient-reported outcomes in a multicenter study S Spittel (Germany)	10.30 – 10.45 C16 Pre-symptomatic ALS/FTD: from biology to prevention? M Benatar (USA)
11.00 – 11.15 C3 IBD methods detecting distant relatedness between ALS patients facilitate new pathogenic gene discovery P Merbaum (Netherlands)	10.50 – 11.10 C10 Treatment patterns and survival benefit of edaravone-treated people with ALS in the ALS/MND Natural History Consortium S Apple (USA)	10.45 – 11.00 C17 Community support information requirements for ALS gene carriers C Haddad (USA)
11.15 – 11.30 C4 Leveraging identity-by-descent to build pedigrees for disease gene discovery L Henden (Australia)	11.10 – 11.30 C11 Development of a saliva-suppressing scopolamine patch without side effects L Ogino (Japan)	11.00 – 11.15 C18 Design of preventative trials in ALS S Fradette (USA)
11.30 – 11.45 C5 Investigating the female protective effect in ALS: Insights from TARDBP mutations carriers in Italy M Grassano (Italy)	11.30 – 11.45 C12 Longitudinal psychometric properties of the DYALS scale in monitoring swallowing function changes in ALS patients C Cattaneo (Italy)	11.15 – 11.30 C19 Pre-motor changes in PD: lessons for ALS and FTD R Postuma (Canada)
11.45 – 12.00 C6 Insights into oligogenic inheritance in familial ALS through Whole Genome Sequencing analysis M Deng (China)	11.45 – 12.00 C13 Development of optimal screening cut-off values from ALSFRS-R bulbar subscore in detecting different levels of dysphagia in a large cohort of ALS patients F Cerri (Italy)	11.30 – 11.45 C20 FTD Detection and diagnosis E Finger (Canada)

<p>12.00 – 12.15 C7 Exome-wide protein domain burden analysis identifies genetic associations with ALS risk H Marriott (UK)</p>	<p>12:00 – 12.15 C14 Self-assessment of ALS functional rating scale on the patient's smartphone proves to be non-inferior to clinic data capture L Steinfurth (Germany)</p>	<p>11.45 – 12.30 PANEL DISCUSSION / Q&A</p>
<p>12.15 – 12.30 C8 Splitformer identifies intronic splicing variants in ALS M Zhang (China)</p>	<p>12.15 – 12.30 C15 Qualitative experiences of Acceptance and Commitment Therapy from the perspectives of people living with MND and therapists: Data from an uncontrolled feasibility study and a randomised controlled trial R Gould (UK)</p>	

12.30 – 14.00 LUNCH

SESSION 3A GLIA	SESSION 3B CLINICAL TRIALS	SESSION 3C NEUROPHYSIOLOGY
<p>14.00 – 14.15 C21 Are myelinating cells an under-appreciated therapeutic target in ALS? K Lewis (Australia)</p>	<p>14.00 – 14.20 C27 Main results from the ADORE study: An international Phase 3 trial to investigate the efficacy and safety of daily oral edaravone (FNP122) in ALS N Albareda (Spain)</p>	<p>14.00 – 14.20 C32 The novel combination of Transcranial Magnetic Stimulation and high-density EMG sensitivity captures ALS decline A Carobin (UK)</p>
<p>14.15 – 14.30 C22 Interrogating the differential toxicity of C9orf72 astrocytes against ALS motor neurons and FTD cortical neurons M Myszczyńska (UK)</p>	<p>14.20 – 14.40 C28 PrimeC, an oral candidate for ALS, demonstrates safety and efficacy in a 12-month Phase 2b trial M Cudkowicz (USA)</p>	<p>14.20 – 14.40 C33 Long intracortical inhibition is not abnormal in ALS: A threshold tracking TMS study R McMackin (Ireland)</p>
<p>14.30 – 14.45 C23 Investigation of WDR49 expression in astrocytes as a driver of ALS pathogenesis M King (UK)</p>	<p>14.40 – 15.00 C29 Safety, tolerability and efficacy of the rho kinase inhibitor fasudil in ALS (ROCK-ALS): A Phase 2, randomise, double-blind, placebo-controlled trial P Lingor (Germany)</p>	<p>14.40 – 15.00 C34 Regional cortical gamma band activity reflects motor and cognitive deficits in ALS M Trubshaw (UK)</p>
<p>14.45 – 15.00 C24 Targeting RNA binding protein SRSF3 and immune mRNAs translation restores microglia function in ALS J Kriz (Canada)</p>	<p>15.00 – 15.15 C30 HERV-K/HML-2 suppression using antiretroviral therapy in ALS D Pandya (USA)</p>	<p>15.00 – 15.15 C35 Upper limb somatosensory evoked potentials latency as a biomarker for early-stage ALS C Lunetta (Italy)</p>
<p>15.00 – 15.15 C25 Microglia-dependent synaptic dysregulation and complement activation in C9orf72 ALS iPSC-derived motor neuron-microglia co-cultures B Vahsen (UK)</p>	<p>15.15 – 15.30 C31 Long-term safety and efficacy of ultrahigh-dose methylcobalamin in early-stage ALS: JETALS interim analysis and open-label extended period R Kaji (Japan)</p>	<p>15.15 – 15.30 C36 Peripheral cause for split hand syndrome in ALS: Subclinical involvement of small hand muscles in early affected hands M de Carvalho (Portugal)</p>
<p>15.15 – 15.30 C26 VHB937, a TREM2 stabilizing and activating antibody, strongly reduces pathology after peripheral administration in a broad range of animal models for neuroinflammation and neurodegeneration D Feuerbach (Switzerland)</p>		

15.30 – 16.00 COFFEE

SESSION 4A IN VITRO MODELS	SESSION 4B EPIDEMIOLOGY	SESSION 4C MOTOR NEURON HETEROGENEITY
<p style="text-align: center;">16.00 – 16.30</p> <p>C37 ALS-in-a-dish: modelling motor neuron disease using advanced human in vitro models J Pasterkamp (Netherlands)</p>	<p style="text-align: center;">16.00 – 16.15</p> <p>C42 Association between sleep and ALS-FTSD: A prospective cohort study based on 396,918 UK Biobank participants T Yang (China)</p>	<p style="text-align: center;">16.00 – 16.30</p> <p>C48 Cellular vulnerability and disease spread in amyotrophic lateral sclerosis: Is it time to re-think upper and lower motor neurons? J Ravits (USA)</p>
<p style="text-align: center;">16.30 – 16.45</p> <p>C38 TDP-43 loss of function induces neuromuscular junction degeneration in a human stem cell derived neuromuscular assembloid model A Salzinger (UK)</p>	<p style="text-align: center;">16.15 – 16.30</p> <p>C43 A population-based mapping and ecological analysis of ALS incidence in the Republic of Ireland between 1995 and 2022 E Mac Domhnaill (Ireland)</p>	<p style="text-align: center;">16.30 – 17.00</p> <p>C49 The importance of understanding the biology and pathology of upper motor neurons for building effective treatment strategies H Ozdinler (USA)</p>
<p style="text-align: center;">16.45 – 17.00</p> <p>C39 Selective disruption of passive nucleocytoplasmic transport in C9ALS/FTD: Arginine DPRs target FG nucleoporins and disrupt protein movement in a sequence-specific manner D Solomon (UK)</p>	<p style="text-align: center;">16.30 – 16.45</p> <p>C44 Association between previous psychiatric disorders and ALS: A population-based prospective cohort study Y Tan (China)</p>	<p style="text-align: center;">17.00 – 17.15</p> <p>C50 Network spreading and local biological vulnerability in ALS A Farahani (Canada)</p>
<p style="text-align: center;">17.00 – 17.15</p> <p>C40 C210rf2 mutations point towards primary cilia dysfunction in ALS P Van Damme (Belgium)</p>	<p style="text-align: center;">16.45 – 17.00</p> <p>C45 Physical activity is associated with lower risk of ALS, including in C9orf72 expansion carriers J Gao (UK)</p>	<p style="text-align: center;">17.15 – 17.30</p> <p>C51 Somatic mosaicism in ALS and FTD reveals widespread degeneration from focal mutations Z Zhou (USA)</p>
<p style="text-align: center;">17.15 – 17.30</p> <p>C41 Embryonic motor neuron programming factors reactivate immature gene expression and suppress ALS pathologies in postnatal motor neurons E Lowry (USA)</p>	<p style="text-align: center;">17.00 – 17.15</p> <p>C46 Neighbourhood deprivation and functional impairment in ALS R Boyle (USA)</p>	
	<p style="text-align: center;">17.15 – 17.30</p> <p>C47 Epigenetic age acceleration is associated with ALS risk, survival, occupational exposures and sex X Li (USA)</p>	

17.30 – 19.00 - POSTER SESSION A

SATURDAY 7 DECEMBER

SESSION 5A CELL BIOLOGY AND PATHOLOGY	SESSION 5B RESPIRATORY MANAGEMENT	SESSION 5C NEUROIMAGING
<p>08.30 – 09.00 C52 On brains and vessels: How vascular mechanisms contribute to ALS neurodegeneration S Lewandowski (Sweden)</p>	<p>08.30 – 09.00 C57 Optimising non-invasive ventilation in ALS D McKim (Canada)</p>	<p>08.30 – 08.45 C62 Cortical and subcortical volumetric MRI biomarkers of C9orf72 repeat expansions C McMillan (USA)</p>
<p>09.00 – 09.15 C53 Sporadic ALS-TDP does not represent a single homogeneous neuropathology R Tan (Australia)</p>	<p>09.00 – 09.15 C58 Respiratory measurements, respiratory symptoms and quality of life in ALS: Results from the REVEALS study D Murray (Ireland)</p>	<p>08.45 – 09.00 C63 Regional vulnerability of the connectome topology in ALS B Kalkhoven (Netherlands)</p>
<p>09.15 – 09.30 C54 Amygdala iron changes are associated with cognitive performance, behavioural deficit and TDP-43 pathology H Spence (UK)</p>	<p>09.15 – 09.30 C59 Peak Inspiratory Flow (PIF) as a predictor of early respiratory decline in ALS U Manera (Italy)</p>	<p>09.00 – 09.15 C64 Attrition-corrected cerebral cortical thickness as a longitudinal imaging biomarker in ALS MJ Wendebourg (Netherlands)</p>
<p>09.30 – 09.45 C55 Molecular and cellular mechanisms of cognitive impairment in ALS C Gouveia Roque (USA)</p>	<p>09.30 – 09.45 C60 Respiratory quotient as independent predictor of prognosis in a large cohort of ALS patients F Cerri (Italy)</p>	<p>09.15 – 09.30 C65 Individual ALS progression markers and disease sub-trajectories encoded in multimodal neuroimaging data T Baumeister (Canada)</p>
<p>09.45 – 10.00 C56 Harnessing a key chaperone to halt TDP-43 aggregation in MND R San Gil (Australia)</p>	<p>09.45 – 10.00 C61 Monitoring the progression of hypoventilation for indicating when to use breathing support in patients with ALS/MND P Cazzolli (USA)</p>	<p>09.30 – 09.45 C66 Multimodal MRI clustering identifies three distinct neurodegeneration-based subtypes of ALS P Van Lieshout (Netherlands)</p>
		<p>09.45 – 10.00 C67 Stronger together: The effect of cognitive abilities and cortical thickness on speech timing abilities in ALS J Bradsby (Canada)</p>

10.00 – 10.30 COFFEE

SESSION 6A PROTEOSTASIS & PROTEOTOXICITY	SESSION 6B EXPERIMENTAL MEDICINE AND TRIAL DESIGN	SESSION 6C COGNITIVE AND BEHAVIOURAL CHANGE
<p>10.30 – 11.00 C68 Transmission of Misfolded Proteins in Neurodegenerative Disorders: A Common Mechanism of Disease Progression V Lee (USA)</p>	<p>10.30 – 11.10 C75 Experimental Medicine to identify therapies in ALS M Turner (UK)</p>	<p>10.30 – 11.00 C81 Defining concepts and modifiers of cognition in ALS C McMillan (USA)</p>
<p>11.00 – 11.15 C69 Structures of pathological TDP-43 filaments in ALS and FTD D Arseni (UK)</p>	<p>11.10 – 11.30 C76 Evaluating and optimizing an ALS platform trial design: Insights for future directions L Chibnik (USA)</p>	<p>11.00 – 11.15 C82 Age-related incidence of neurodegenerative diseases in C9orf72 expansion carriers from a population-based cohort J Gao (UK)</p>
<p>11.15 – 11.30 C70 A stress-dependent TDP-43 SUMOylation program preserves neuronal function T Suk (Canada)</p>	<p>11.30 – 11.45 C77 Boosting clinical trial power in ALS with AI-generated digital twins C Kusiak (USA)</p>	<p>11.15 – 11.30 C83 The features of behavioural impairment in Chinese ALS patients J Tang (China)</p>

<p>11.30 – 11.45 C71 LRSAM1 mitigates neurodegeneration in ALS by regulating clearance of TDP-43 T Shirakawa (USA)</p>	<p>11.45 – 12.00 C78 Cultivating patient preferences in ALS clinical trials: The Patient-Ranked Order of Function (PROOF) R van Eijk (Netherlands)</p>	<p>11.30 – 11.45 C84 Cognitive screening in ALS in Ireland: A 10-year study of prevalence, instrument sensitivity and confounding factors E Costello (Ireland)</p>
<p>11.45 – 12.00 C72 Pre-symptomatic pathological TDP-43 aggregation is a common feature in peripheral, non-central nervous system tissues in people with ALS J Gregory (UK)</p>	<p>12.00 – 12.15 C79 Longitudinal comparison of the Italian version of the ALSFRS-R and ROADS: An Italian multicenter prospective study E Matteoni (Italy)</p>	<p>11.45 – 12.00 C85 Predicting cognitive and behavioural decline in ALS patients: a prospective pilot study A Chio (Italy)</p>
<p>12.00 – 12.15 C73 Intrinsic and TDP-43 dysfunction-driven cellular degradation loads evoke Tfe3a-associated neuroprotective autophagy in large spinal motor neurons K Asakawa (Japan)</p>	<p>12.15 – 12.30 C80 Harmonization e-learning for ALS clinical trials: ALSFRS-R harmonized online training platform for cross-continent collaboration G Kittle (USA)</p>	<p>12.00 – 12.15 C86 Development of cognitive/behavioural disturbances in motor neuron diseases: Can we predict it? P Ferraro (Italy)</p>
<p>12.15 – 12.30 C74 Enhancing proteostasis in ALS through novel brain-penetrating peptides: A therapeutic approach targeting autophagy A Amin (Australia)</p>		<p>12.15 – 12.30 C87 Prodromal mild cognitive impairment in ALS and FTD: A revised framework C McHutchison (UK)</p>

12.30 – 14.00 LUNCH

SESSION 7A IMMUNITY AND NEUROINFLAMMATION	SESSION 7B AUTONOMY & DECISION- MAKING	SESSION 7C ANTISENSE AND siRNA- BASED THERAPEUTIC STRATEGIES
<p>14.00 – 14.20 C88 RIG-1 but not Sting mediate neuron specific IFN type 1 signalling in FUS-ALS induced neurodegeneration and offers new biomarker driven individualized treatment options A Herman (Germany)</p>	<p>14.00 – 14.45 C93 MAID: From legislation to practice W Johnston (Canada)</p>	<p>14.00 – 14.30 C97 Antisense-based therapies for rare neurological diseases F Bennett (USA)</p>
<p>14.20 – 14.40 C89 Investigating the metabolic crosstalk between senescent T cells and microglia from patients with ALS V Tsang (UK)</p>	<p>14.45 – 15.00 C94 The OCEAAMM: A decision support tool to evaluate competency to consent to MAID S Charbonneau (Canada)</p>	<p>14.30 – 14.45 C98 Update on Silence ALS: A platform for the discovery and development of antisense therapeutics for patients with ultra-rare forms of ALS N Shneider (USA)</p>
<p>14.40 – 15.00 C90 Functionally distinct Treg subsets exhibit unique mechanisms of dysfunction in ALS C Baecher-Allan (USA)</p>	<p>15.00 – 15.15 C95 The use of support strategies to improve mental capacity to make treatment decisions in ALS S Abrahams (UK)</p>	<p>14.45 – 15.00 C99 Antisense oligonucleotides are broadly distributed but do not provide sustained suppression of G4C2 pathology in c9ALS patients J Glass (USA)</p>
<p>15.00 – 15.15 C91 Immune checkpoint changes correlate with the progression and prognosis of ALS S Chen (China)</p>	<p>15.15 – 15.30 C96 SOD1 gene screening in ALS: Frequency of mutations, patients' attitudes to genetic information and translation to tofersen treatment in a multi-center program T Meyer (Germany)</p>	<p>15.00 – 15.15 C100 RAG-17: A promising new gene silencing therapy for SOD1 ALS. Early safety and efficacy data from a first-in-human trial J Ye (China)</p>

15.15 – 15.30 C92 Longitudinal analysis of T cell responses in ALS S Yazdani (Sweden)		15.15 – 15.30 C101 Development of an UNC13A cryptic exon skipping antisense oligonucleotide as a treatment for ALS W-H Chang (USA)
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15.30 – 16.00 COFFEE

SESSION 8A RNA BIOLOGY	SESSION 8B BIOMARKERS	SESSION 8C SPONSORED SESSION (TBC)
16.00 – 16.30 C102 Using spatial genomics to study the central nervous system in health and disease H Phatnani (USA)	16.00 – 16.20 C107 Elevated cerebrospinal fluid UCHL1 in asymptomatic C9orf72 hexanucleotide repeat expansion carriers E Dellar (UK)	
16.30 – 16.45 C103 Employing advanced long-read RNA sequencing in post mortem brain specimens to explore transcriptomic diversity associated with upper and lower motor neuron pathology A Jain (USA)	16.20 – 16.40 C108 Combination of serum neurofilament light and serum cardiac troponin T improves diagnostic accuracy in ALS P Weydt (Germany)	
16.45 – 17.00 C104 Single-nucleus transcriptome atlas of orbitofrontal cortex in ALS and FTLT with a deep learning-based decoding of alternative polyadenylation mechanisms P McKeever (Canada)	16.40 – 17.00 C109 Total tau and phosphorylated tau levels as diagnostic biomarkers for ALS T Petrozziello (USA)	
17.00 – 17.15 C105 Exploring ALS differential vulnerability using single-cell transcriptomic analysis P Alipour (Canada)	17.00 – 17.20 C110 Development of a blood test to distinguish amyotrophic lateral sclerosis from primary lateral sclerosis S Banack (USA)	
17.15 – 17.30 C106 A journey through space and time: ALS human and mouse spinal cords demonstrate altered gene expression relative to disease pathology S Howe (Australia)		

17.30 – 19.00 POSTER SESSION B

08.30 – 10.00 POSTER SESSION C

10.00 – 10.30 COFFEE

SESSION 9A MURINE MODELS	SESSION 9B INTERDISCIPLINARY CARE: POLICY AND PRACTICE	SESSION 9C BIOMARKERS (II)
<p>10.30 – 10.50 C111 A TBK1 variant causes autophagolysosomal and motoneuron pathology without neuroinflammation in mice D Brenner (Germany)</p>	<p>10.30 – 10.50 C118 Clinicians' Perceptions of Palliative Care for Amyotrophic Lateral Sclerosis (ALS): Results of National Surveys of Interdisciplinary ALS and Palliative Care Clinicians in the United States K Bischoff (USA)</p>	<p>10.30 – 10.50 C124 Novel prognostic protein discovery using Olink proximity extension assay in a large longitudinal ALS cohort D Lester (UK)</p>
<p>10.50 – 11.10 C112 Developing an inducible muscle-specific TDP-43 mouse model to investigate ALS A Russell (Australia)</p>	<p>10.50 – 11.10 C119 How do multidisciplinary teams support people with MND to make decisions about gastrostomy placement: a qualitative multiple case study S White (UK)</p>	<p>10.50 – 11.10 C125 Detection and Discovery of Urinary Biomarkers of Immune Dysfunction for Amyotrophic Lateral Sclerosis V Karnaros (Australia)</p>
<p>11.10 – 11.30 C113 TDP-43 cytoplasmic mislocalisation drives synaptic dysfunction and persistent non-motor behavioural changes in the rNLS8 mouse model of ALS/FTD W Luan (Australia)</p>	<p>11.10 – 11.30 C120 <i>How do people living with MND and their carers experience specialized care? Development and validation of a patient reported questionnaire</i> AK Schmidt (Netherlands)</p>	<p>11.10 – 11.30 C126 Targeted proteomics upon Tofersen treatment highlights candidate therapy-responsive markers for SOD1-linked ALS A Catanese (Germany)</p>
<p>11.30 – 11.45 C114 Spatial transcriptomic profiling of the lower motor neuron circuitry reveals an axonal-specific RNA signature and local translation defects in mutant FUS-mediated ALS D Piol (Belgium)</p>	<p>11.30 – 11.50 C121 Virtual care delivery models and team functioning in ALS – a pan-Canadian experience with interdisciplinary virtual care in ALS B Ritsma (Canada)</p>	<p>11.30 – 11.50 C127 Longitudinal analysis of immune cell changes in FUS-ALS patients treated with a FUS antisense oligonucleotide O Rifai (USA)</p>
<p>11.45 – 12.00 C115 Unveiling a novel mechanistic link between oxidative stress and pathophysiology of Amyotrophic Lateral Sclerosis E Eftekharpour (Canada)</p>	<p>11.50 – 12.10 C122 Proposal for a U.S National Integrated ALS Care and Research Network System- Excerpts from the National Academies of Medicine Report on ALS S Babu (USA)</p>	<p>11.50 – 12.10 C128 Corneal small fiber neuropathy in ALS patients: a corneal confocal microscopy study Z Zou (China)</p>
<p>12.00 – 12.15 C116 Comparison of the motor neuron transcriptome in mouse models of ALS and SMA reveals defects specific to protein translation. H Smith (UK)</p>	<p>12.10 – 12.30 C123 The EU ALS Coalition call for action: Roadmap of the policy changes needed across Europe to improve the lives of PLWALS and their carers across Europe J Grosskreutz (Germany)</p>	<p>12.10 – 12.30 C129 Seed-amplification assays: Windows into ALS and kindred diseases R Smith (USA)</p>
<p>12.15 – 12.30 C117 Assessing in vivo axonal transport of mitochondria and signalling endosomes in distinct α-motor neuron subtypes in SOD1G93A mice A Tosolini (Australia)</p>		

12.30 – 13.30 LUNCH

SESSION 10 JOINT CLOSING SESSION	
	Invitation to San Diego 2025
13.40 – 13.50	Poster Prize Awards
	Healey Center Prize
	Lalji Family Award
14.10 – 14.30	C130 Early onset sleep alterations in ALS and their mechanisms – L Dupuis (France)
14.30 – 15.00	C131 Outcome measures in ALS clinical trials: progress to date and future directions – A Genge (Canada)
15.00 – 15.15	Late breaking news