

## FINAL program 22.3.25

	THURSDAY, MARCH 20 <sup>th</sup> ,2025	
08:00-09:40	Neuroimmunology	ALL A
Chairs:	Brian Weinshenker, USA, Avi Gadoth, Israel	
08:00-08:50		
	Capsule: Myelin oligodendrocyte glycoprotein-lgG is a biomarker of a specific neuroimmune disease characterized by optic neuritis, myelitis, acute dissemi	
	encephalomyelitis and occasionally cortical encephalitis. Rituximab, although effective, is less effective than for neuromyelitis optica spectrum disorder associat with aquaporin 4-IgG. It remains uncertain whether the disease is due to the direct effects of the antibody or whether the antibody is a marker of autoimmunity	
	that may be mediated by other effectors	
08:00-08:10	Moderator: <u>Brian Weinshenker</u> , USA Introduction and Pre-Debate Voting	
08:10-08:25	Yes: Patrick Waters, UK	
08:25-08:40	No: Thomas Berger, Austria	
08:40-08:50	Discussion, Rebuttals and Post-Debate Voting	
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08:50-09:40	Time to redefine generalized myasthenia gravis (gMG): are corticosteroids the backbone of the MG treatment?	
	Capsule: For decades, treatment of gMG consisted mainly of cholinesterase inhibitors, immunosuppresants and costicosteroids. Recently, monoclonal anti-	bodies
	have been added, but have they changed the scene?	
08:50-09:00	Moderator: Thomas Berger, Austria	
	Introduction and Pre-Debate Voting	
09:00:09:15	Yes: Hakan Cetin, Austria	
09:15-09:30	No: Anna Kostera-Pruszczyk, Poland	
09:30-09:40 <b>09:40-10:10</b>	Discussion, Rebuttals and Post-Debate Voting  Coffee Break, Exhibition & ePosters Visits	
10:10-11:10		ALL A
Chairs:	Amos Korczyn, Israel; Petr Marusic, Czech Republic, Natan Bornstein, Israel	ALL A
10:10-10:15	Welcome to CONy 2025 - Natan Bornstein, Israel; Amos Korczyn, Israel	
10:15-10:20	Welcome address - Irena Rektorova, Czech Republic	
10:20-10:25	Best e-Poster Award - Natan Bornstein, Israel	
10:25-10:30	Welcome address on behalf of the Czech Neurological Society - Petr Marusic, Czech Republic	
10:30-10:45	The contribution of Oskar Fischer and Arnold Pick to the field of dementia - <u>Irena Rektorova</u> , Czech Republic	
10:45-10:50	CONy Excellence in Neurology Award to Prof. Friedemann Paul - presented by <u>Amos Korczyn</u> , Israel	
10:50-11:10	NMOSD - an emerging spectrum - Friedemann Paul, Germany *recording	

	THURSDAY, MARCH 20 <sup>th</sup> ,2025	
11:10-12:10	Plenary Session	HALL A
Chairs:	George Chakhava, Georgia; Viktoriia Gryb, Ukraine	'
11:10-11:40	A Plan for Parkinson - Michael Okun, USA	
11:40-12:10	The impact of climate changes on neurological diseases - Jacques Reis, France	
12:10-13:10	Industry Sponsored Symposium	HALL A
13:10-14:10	Lunch Break, Exhibition & ePosters Visits	
14:10-15:50	Neuroimmunology (continued)	HALL A
Chairs:	Klaudia Duka Glavor, Croatia; Ali Hasnain, Ireland	
14:10-15:00	All patients with PML should be treated with pembrolizumab	
	Capsule: Progressive multifocal leukoencephalopathy (PML) is a devastating condition caused by JC virus reactivation observed mainly	•
	patients but also in patients with inflammatory diseases treated with various immunosuppressants. Disability and mortality of PML co	-
	reconstitution (IRIS) which is sometimes seen after the diagnosis, especially after stopping immunosuppressants. Should all patients w	vith PML receive immune
	check-point inhibitors such as pembrolizumab?	
14:10-14:20	Moderator: Avi Gadoth, Israel	
14.20 14.25	Introduction and Pre-Debate Voting	
14:20-14:35 14:35:14:50	Yes: <u>Uros Rot</u> , Slovenia	
14:50:15:00	No: Michel Toledano, USA  Discussion, Bobyttale and Bost Debate Veting	
14:50:15:00	Discussion, Rebuttals and Post-Debate Voting	
15:00-15:50	Is CAR-T cell therapy appropriate for NMOSD?	
	Capsule: Chimeric antigen receptor (CAR)-T cells are autologous T cells engineered to target a variety of antigens. Potential advantage	es of this form of treatment
	include the tissue distribution properties of T cells and self-replication. CAR-T cells have revolutionized the treatment of B-cell maligna	ncies and have recently been
	applied to autoimmune disease. There are a number of toxicities including cytokine release syndrome. Does CAR T cell therapy offer u	ınique advantages for
	NMOSD that justify its cost and toxicity?	
15:00-15:10	Moderator: <u>Joab Chapman,</u> Israel	
	Introduction and Pre-Debate Voting	
15:10-15:25	Yes: Brian Weinshenker, USA	
15:25-15:40	No: Petra Nytrova, Czech Republic	
15:40-15:50	Discussion, Rebuttals and Post-Debate Voting	
15:50-16:20	Coffee Break, Exhibition & ePosters Visits	



	THURSDAY, MARCH 20 <sup>th</sup> , 2025	
16:20-18:00	Neuroimmunology (continued)	
Chairs:	Boleslav Lichterman, Russia; Petra Nytrova, Czech Republic	
16:20-17:10	Can primary CNS vasculitis be diagnosed without biopsy?	
	Capsule: Primary central nervous system vasculitis (CNSV) is a challenging diagnosis due to its rarity and clinical variability. Traditionally, brain biopsy has recognized as a gold standard to establish definitive diagnosis. However, its invasive nature and limited sensitivity, despite being relatively high, raises a question: Can primary CNSV be diagnosed without a biopsy? In this debate we will consider alternative diagnostic methods, and their reliability comparbiopsy.	the
16:20-16:30	Moderator: <u>Michel Toledano</u> , USA Introduction and Pre-Debate Voting	
16:30-16:45	Yes: Sarlota Mesaros, Serbia	
16:45-17:00	No: Joab Chapman, Israel	
17:00-17:10	Discussion, Rebuttals and Post-Debate Voting	
17:10-18:00	Narcolepsy is an autoimmune disorder	
	Capsule: The current body of literature supports that narcolepsy is an autoimmune disorder. However, the role of autoantibodies has yet to be establish	ned.
	Moreover, reports of using immunotherapies in narcolepsy patients remain limited and inconsistent. Nonetheless, narcolepsy has been strongly linked that alleles and T-cell receptor polymorphisms. More recently, it has been argued that alterations in cytokine levels, gut microbiota, and microglial activindicate a neuro-inflammation in the disease's development, and during this debate we will discuss current evidence pro and against the immune theor as address the potential role for epigenetic silencing.	vation may
17:10-17:20	Moderator: <u>Ivana Rosenzweig</u> , UK Introduction and Pre-Debate Voting	
17:20-17:35	Yes: Roland Liblau, France	
17:35-17:50	No: Mehdi Tafti, Switzerland	
17:50-18:00	Discussion, Rebuttals and Post-Debate Voting	
18:00	Networking Reception	



	THURSDAY, MARCH 20 <sup>th</sup> , 2025		
08:00-09:40	Alzheimer's Disease (AD) & Dementia HALL B		
Chairs:	Marina Janelidze, Georgia; Judith Aharon Peretz, Israel		
08:00-08:50			
	<b>Capsule</b> : AD is typically perceived as a memory-predominant neurodegenerative condition. However, in ~10% of individuals non disturbances in processing of visual information, language impairment and/or behavioral/personality changes represent the coatypical clinical presentation (and associated biomarker profiles and progression rates), these individuals do not meet eligibility therefore systematically excluded from promising investigational interventions with disease modifying drugs. Here, we will discuintividuals with atypical forms of AD in clinical trials.	re cognitive complaint. Due to their criteria for clinical trials and are	
08:00-08:10	Moderator: Rik Ossenkoppele, The Netherlands Introduction and Pre-Debate Voting		
08:10-08:25	Yes: Keir Yong, UK		
08:25-08:40	No: Rosaleena Mohanty, Sweden		
08:40-08:50	Discussion, Rebuttals and Post-Debate Voting		
08:50-09:40	Are the new anti-amyloid drugs cost-effective?		
	Capsule: The cost-effectiveness of lecanemab and donanemab is being closely examined. Regulators and payors in the US, Europe, Great Britain and other jurisdictions have come to different conclusions. The usual price-point of \$100,000/QALY has been exceeded for lecanemab, and details for donanemab are available. The advent of subcutaneous formulations and stopping/maintenance rules will have to be taken into account. Competition between current manufacturers, next generation antibodies and increased efficacy with longer term administration (3-5 years) at earlier stages of AD will also change the calculations		
08:50-09:00	Moderator: Colin L. Masters, Australia Introduction and Pre-Debate Voting		
09:00:09:15	Yes: Jakub Hlavka, Czech Republic		
09:15-09:30	No: Stanislav Sutovsky, Slovakia		
09:30-09:40	Discussion, Rebuttals and Post-Debate Voting		
09:40-10:10	Coffee Break, Exhibition & ePosters Visits		
10:10-11:10	Opening Ceremony and Best e-Poster awards	HALL A	
Chairs:	Amos Korczyn, Israel; Petr Marusic, Czech Republic, Natan Bornstein, Israel		
10:10-10:15	Welcome to CONy 2025 - Amos Korczyn, Israel; Natan Bornstein, Israel		
10:15-10:20	Welcome address - Irena Rektorova, Czech Republic		
10:20-10:25	Best e-Poster Award - Natan Bornstein, Israel		
10:25-10:30	Welcome address on behalf of the Czech Neurological Society- Petr Marusic, Czech Republic		
10:30-10:45	The contribution of Oskar Fischer and Arnold Pick to the field of dementia - <u>Irena Rektorova</u> , Czech Republic		

CONy Excellence in Neurology Award to Prof. Friedemann Paul  NMOSD - an emerging spectrum - Friedemann Paul, Germany *recording  Plenary Session	
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	HALL A
George Chakhava, Georgia; Viktoriia Gryb, Ukraine	
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Industry Sponsored Symposium	HALL A
Lunch Break, Exhibition & ePosters Visits	
Alzheimer's Disease (AD) & Dementia (continued)	HALL B
Odelia Elkana, Israel; Xiao Ping Wang, China	
Should MCI patients be immunized against zoster?	
<b>Capsule</b> : Retrospective studies have suggested that immunization against herpes zoster reduces the incidence of dementia. Call i.e. should AD patients be immunized to ameliorate the disease?	n this also be of therapeutic value,
Moderator: Stanislav Sutovsky, Slovakia Introduction and Pre-Debate Voting	
Yes: <b>Lukasz Rzepiński</b> , Poland	
No: Dorota Religa, Sweden	
Discussion, Rebuttals and Post-Debate Voting	
Is AD a disease?	
<b>Capsule</b> : The definition of AD has changed several times over the years and still lacks an agreed one. Lacking understanding of the condition, it is still arguable whether it should be considered a disease or a syndrome	he causes and mechanisms of the
Moderator: Lon Schneider, USA Introduction and Pre-Debate Voting	
Yes: Colin L. Masters, Australia	
No: Amos Korczyn, Israel	
Discussion, Rebuttals and Post-Debate Voting	
Coffee Break, Exhibition & ePosters Visits	
	Industry Sponsored Symposium Lunch Break, Exhibition & ePosters Visits  Alzheimer's Disease (AD) & Dementia (continued) Odelia Elkana, Israel; Xiao Ping Wang, China Should MCI patients be immunized against zoster? Capsule: Retrospective studies have suggested that immunization against herpes zoster reduces the incidence of dementia. Ca i.e. should AD patients be immunized to ameliorate the disease? Moderator: Stanislav Sutovsky, Slovakia Introduction and Pre-Debate Voting Yes: Lukasz Rzepiński, Poland No: Dorota Religa, Sweden Discussion, Rebuttals and Post-Debate Voting  Is AD a disease? Capsule: The definition of AD has changed several times over the years and still lacks an agreed one. Lacking understanding of to condition, it is still arguable whether it should be considered a disease or a syndrome Moderator: Lon Schneider, USA Introduction and Pre-Debate Voting Yes: Colin L. Masters, Australia No: Amos Korczyn, Israel Discussion, Rebuttals and Post-Debate Voting



	THURSDAY, MARCH 20th , 2025	
16:20-18:00	Alzheimer's Disease (AD) & Dementia (continued)	HALL B
Chairs:	Yvonne Freund-Levi, Sweden; Milica G. Kramberger, Slovenia	
16:20-17:10	Monoclonal antibodies or natural products for prevention of dementia?	
	Capsule: Monoclonal antibodies and natural products are both being explored for the prevention of dementia. Monoclonal anti-	
	plaques in the brain, which are a hallmark of AD, and have been shown consistently to have positive effects on reducing amyloid levels and slowing cognitive decline. However, their high cost and potential side effects are concerns. On the other hand, natural products like dietary supplements, omega-3 fatty acids, and antioxidants may support brain health and delay cognitive decline. Both approaches have potential, but further studies are essential to determine their long-term	
	benefits and practicality	
16:20-16:30	Moderator: <u>Lon Schneider</u> , USA	
	Introduction and Pre-Debate Voting	
16:30-16:45	Natural products: <u>Magda Tsolaki</u> , Greece	
16:45-17:00	Monoclonal antibodies: <u>Jakub Hort</u> , Czech Republic	
17:00-17:10	Discussion, Rebuttals and Post-Debate Voting	
17:10-18:00	Do lifestyle factors protect against dementia by affecting amyloid metabolism?	
	Capsule: Lifestyle factors, such as physical activity, diet, and cognitive engagement, may protect against dementia by influence	
	associated with reduced amyloid plaque accumulation and improved cognitive function. Diets like the Mediterranean diet link	
	cognitive decline. Cognitive engagement through activities like reading and puzzles can delay dementia onset by reducing amy	
	predispositions and the complex nature of lifestyle adherence can limit these benefits. While promising, the relationship between lifestyle factors and amyloid	
	metabolism is not fully understood, and this debate will discuss the pros and cons of the existing evidence.	
17:10-17:20	Moderator: Milica Kramberger, Slovenia	
17.20 17.25	Introduction and Pre-Debate Voting	
17:20-17:35	Yes: Laura Bonanni, Italy *recording	
17:35-17:50	No: Giancarlo Logroscino, Italy	
17:50-18:00	Discussion, Rebuttals and Post-Debate Voting	
18:00	Networking Reception	



	THURSDAY, MARCH 20 <sup>th</sup> , 2025	
08:00-09:40	Parkinson's Disease (PD) I	HALL C
Chairs:	Irena Rektorova, Czech Republic; Stanley Fisher, USA	
08:00-08:50		
	Capsule: Jean-Martin Charcot refined the original description of James Parkinson as disorder with characteristic motor features the clinical definition of Parkinson's disease (PD). However, we have evolved tremendously in terms of our understanding of genetic for imaging modalities, and biomarkers, supporting the vast heterogeneity observed in disease manifestation and progression. While investigate the biological underpinnings of PD, and to develop better biomarkers and imaging approaches, we are now in a positive knowledge is ready for aiding researchers classify patients in order to aid patient selection for clinical trials, in the hope that this values in developing novel therapeutic strategies for a disease that is actually a syndrome and not a single homogeneous entity.	actors, pathogenic mechanisms, e it will be essential to continue to on to debate whether the existing will increase our chance of
08:00-08:10	Moderator: Michael Okun, USA	
	Introduction and Pre-Debate Voting	
08:10-08:25	Yes: <u>Tiago Outeiro</u> , Germany	
08:25-08:40	No: Matej Skorvanek, Slovakia	
08:40-08:50	Discussion, Rebuttals and Post-Debate Voting	
08:50-09:40	The first treatment of Restless legs syndrome (RLS) should be dopamine agonists vs gabapentin and pregabalin	
	Capsule: RLS is a common neurological disorder among adult patients that often disrupts sleep and can impact activities of daily an urge to move the legs or other body parts that begin or worsen during rest or inactivity. The urge to move is typically worse in and is relieved by movement. RLS remains under-diagnosed, and many patients are not treated appropriately. The first treatment	the evening or nighttime hours
08:50-09:00	Moderator: Michal Minar, Slovakia Introduction and Pre-Debate Voting	
09:00:09:15	Dopamine agonists: Vladimira Vuletic, Croatia	
09:15-09:30	Gabapentin / pregabalin: Jarosław Slawek, Poland	
09:30-09:40	Discussion, Rebuttals and Post-Debate Voting	
09:40-10:10	Coffee Break, Exhibition & ePosters Visits	
10:10-11:10	Opening Ceremony and Best e-Poster awards	HALL A
Chairs:	Amos Korczyn, Israel; Petr Marusic, Czech Republic, Natan Bornstein, Israel	
10:10-10:15	Welcome to CONy 2025 - Amos Korczyn, Israel; Natan Bornstein, Israel	
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11:10-12:10	Plenary Session	HALL A
Chairs:	George Chakhava, Georgia; Viktoriia Gryb, Ukraine	
11:10-11:40	A Plan for Parkinson - Michael Okun, USA	
11:40-12:10	The impact of climate changes on neurological diseases - <u>Jacques Reis</u> , France	
12:10-13:10	Industry Sponsored Symposium	HALL A
13:10-14:10	Lunch Break, Exhibition & ePosters Visits	
14:10-15:50	Parkinson's Disease (PD) I (continued)	HALL C
Chairs:	Cristian Falup-Pecurariu, Romania; Magdalena Kwasniak-Butowska, Poland	
14:10-15:00	The MRI will replace molecular imaging to support the diagnosis of PD	
	Capsule: Modern MRI technology with 3T allows detection of the so-called swallow tail sign. So far, the specificity and the	e sensitivity seem to be lower thar
	<b>Capsule</b> : Modern MRI technology with 3T allows detection of the so-called swallow tail sign. So far, the specificity and the using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson	
14.10 14.20	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinso	
14:10-14:20	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinso discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting	
14:10-14:20 14:20-14:35	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinso discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic	
	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK	
14:20-14:35	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinso discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic	
14:20-14:35 14:35-14:50	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK Discussion, Rebuttals and Post-Debate Voting  GLP-1 agonists are disease modifying for PD and should be used in all patients	on's disease. The debate will
14:20-14:35 14:35-14:50 14:50-15:00	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK  Discussion, Rebuttals and Post-Debate Voting  GLP-1 agonists are disease modifying for PD and should be used in all patients  Capsule: The recent New England Journal of Medicine paper showed that GLP-1 agonists may possibly be disease modifying and	on's disease. The debate will
14:20-14:35 14:35-14:50 14:50-15:00	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK Discussion, Rebuttals and Post-Debate Voting  GLP-1 agonists are disease modifying for PD and should be used in all patients  Capsule: The recent New England Journal of Medicine paper showed that GLP-1 agonists may possibly be disease modifying ar field. Should we be giving them? What is the risk benefit ratio? Will weight loss or GI symptoms impact the decision? What other	on's disease. The debate will
14:20-14:35 14:35-14:50 14:50-15:00 <b>15:00-15:50</b>	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK Discussion, Rebuttals and Post-Debate Voting  GLP-1 agonists are disease modifying for PD and should be used in all patients  Capsule: The recent New England Journal of Medicine paper showed that GLP-1 agonists may possibly be disease modifying and field. Should we be giving them? What is the risk benefit ratio? Will weight loss or GI symptoms impact the decision? What oth Moderator: Michael Okun, USA	on's disease. The debate will
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14:20-14:35 14:35-14:50 14:50-15:00 <b>15:00-15:50</b> 15:00-15:10	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK  Discussion, Rebuttals and Post-Debate Voting  GLP-1 agonists are disease modifying for PD and should be used in all patients  Capsule: The recent New England Journal of Medicine paper showed that GLP-1 agonists may possibly be disease modifying are field. Should we be giving them? What is the risk benefit ratio? Will weight loss or GI symptoms impact the decision? What oth Moderator: Michael Okun, USA Introduction and Pre-Debate Voting  Yes: Sharon Hassin-Baer, Israel	on's disease. The debate will
14:20-14:35 14:35-14:50 14:50-15:00 <b>15:00-15:50</b>	using molecular imaging with PET or SPECT technology which are propagated in the new biological definitions of Parkinson discuss whether this can be changed  Moderator: Jarosław Slawek, Poland Introduction and Pre-Debate Voting  Yes: Irena Rektorova, Czech Republic  No: Nicola Pavese, UK Discussion, Rebuttals and Post-Debate Voting  GLP-1 agonists are disease modifying for PD and should be used in all patients  Capsule: The recent New England Journal of Medicine paper showed that GLP-1 agonists may possibly be disease modifying art field. Should we be giving them? What is the risk benefit ratio? Will weight loss or GI symptoms impact the decision? What oth Moderator: Michael Okun, USA Introduction and Pre-Debate Voting	on's disease. The debate will



THURSDAY, MARCH 20 <sup>th</sup> , 2025		
15:50-16:20	Coffee Break, Exhibition & ePosters Visits	
16:20-18:00	Parkinson's Disease (PD) I (continued) HALL C	
Chairs:	Weidong Le, China; Nana Kvirkvelia, Georgia	
16:20-17:10	Essential tremor plus (ET+) is a clinically useful concept	
	Capsule: The concept of ET+ suggests that cases of essential tremor (ET) with additional neurological symptoms form a distinct category. ET+ includes signs like	
	dystonia, cognitive changes, or gait abnormalities, broadening the understanding of tremor disorders. Proponents argue that ET+ acknowledges the complexity of	
	tremor presentations, yet critics point to the term's ambiguity and risk of diagnostic overlap. The lack of clear criteria and variable clinical relevance challenge	
	ET+'s utility. The classification remains controversial, and this debate will explore the strengths and limitations of the concept.	
16:20-16:30	Moderator: Sharon Hassin-Baer, Israel	
10.20 10.50	Introduction and Pre-Debate Voting	
16:30-16:45	Yes: Matej Skorvanek, Slovakia	
16:45-17:00	No: <u>Evzen Ruzicka</u> , Czech Republic	
17:00-17:10	Discussion, Rebuttals and Post-Debate Voting	
17:10-18:00	Focused ultrasound thalamotomy becomes the first-choice treatment for medically refractory essential tremor	
	Capsule: Medication refractory Essential tremor was in the past treated with deep brain stimulation. With the emergence of MRI guided focused ultrasound	
	thalamotomy, a non-invasive therapy that offers tremor relief, patients are referred for focused ultrasound instead of DBS. Should focused ultrasound	
	thalamotomy become the first choice of therapy in medication refractory Essential tremor?	
17:10-17:20	Moderator: Evzen Ruzicka, Czech Republic	
17.10 17.20	Introduction and Pre-Debate Voting	
17:20-17:35	Yes: <u>Ilana Schlesinger</u> , Israel	
17:35-17:50	No: Michael Okun, USA	
17:50-18:00	Discussion, Rebuttals and Post-Debate Voting	
18:00	Networking Reception	



	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
08:00-09:40	Multiple Sclerosis (MS)	HALL A
Chairs:	Konrad Rejdak, Poland; Jera Kruja, Albania	
08:00-08:50	European Charcot Foundation Symposium: Assessment of treatment response in progressive MS	
08.00-08.50	The Symposium is dedicated to the Memory of Prof. Giancarlo Comi	
	<b>Capsule</b> : Defining disability progression in MS remains a challenge. Universally agree upon criteria are missing. More recently the introduction of PIRA, progression independent of relapse activity, has complicated matters. Disability not related to failed recovery from relapses may conceptually allow definition with more stringency. However, it remains demanding to capture the entirety of disease activity with high granularity. Here we would like to shed light on this issue from various perspectives and discuss the use of different approaches: clinical measures, neuroimaging and functional tests. In all areas significant progress has been	
	made over recent years. This will be critically assessed.	
08:00-08:10	Moderator: Maria Trojano, Italy Introduction	
08:10-08:20	Clinical measures: Maria Trojano, Italy	
08:20:08:30	Neuroimaging: Mike Wattjes, Germany	
08:30-08:40	Functional tests: Letizia Leocani, Italy	
08:40-08:50	Discussion	
08:50-09:40	Epstein-Barr virus (EBV) is a therapeutic target in established MS  Capsule: MS is caused by an interplay between environmental and genetic factors. Infection with EBV significantly increases the risk of MS indicating that EBV ca	
	be an important factor in development of MS. Molecular mimicry between Epstein-Barr nuclear antigen 1 (EBNA1) and brain G	GlialCAM is postulated. Could we also
	treat MS by vaccinating against EBV or use antiviral drugs?	
08:50-09:00	Moderator: Jacek Losy, Poland	
00.50 05.00	Introduction and Pre-Debate Voting	
09:00:09:15	Yes: Gavin Giovannoni, UK	
09:15-09:30	No: Ron Milo, Israel	
09:30-09:40	Discussion, Rebuttals and Post-Debate Voting	
09:40-10:10	Coffee Break, Exhibition & ePosters Visits	
10:10-11:10	Plenary Session	HALL A
Chairs:	Max J. Hilz, USA; Natan Bornstein, Israel	
10:10-10:40	Neurology is psychiatry and vice versa - Adam Zeman, UK	
10:40-11:10	Apraxia - Amos Korczyn, Israel	

FRIDAY, MARCH 21 <sup>ST</sup> , 2025		
11:10-12:10	Industry Sponsored Symposium	HALL A
12:10-13:10	Lunch Break, Exhibition & ePosters Visits	
13:10-14:50	Multiple Sclerosis (continued)	HALL A
Chairs:	Jacek Losy, Poland	
13:10-14:00	Capsule: Several studies have suggested that MS diagnosis can be preceded by unspecific prodromal symptoms, months or even years before classical manifestation of the disease. Although an evident prodromal phase is associated with (among many) Parkinson's disease, Alzheimer's, rheumatoid arthritis, and	
	Crohn's disease, it is still debated whether MS is also associated with one, or whether unspecific prodromal symptoms could simply translate to early	
	manifestations of the disease itself.	
13:10-13:20	Moderator: Klaus Schmierer, UK	
12.20 12.25	Introduction and Pre-Debate Voting	
13:20-13:35	Yes: Alicja Kalinowska, Poland	
13:35-13:50 13:50-14:00	No: <u>Gavin Giovannoni</u> , UK Discussion, Rebuttals and Post-Debate Voting	
13.50-14.00	Discussion, Reputtals and Post-Departe Voting	
14:00-14:50	All patients with radiologically isolated syndrome (RIS) should be treated with disease-modifying therapies (DMT)	
	Capsule: RIS is often the first detectable manifestation of central nervous system (CNS) autoimmunity. In fact, ten years after	the diagnosis of RIS, more than 50%
	of individuals will have progressed to a formal diagnosis of clinically isolated syndrome (CIS) or multiple sclerosis (MS). There are currently over 20 approved D for patients with CIS and MS available that are effective and relatively safe. For two of these agents, namely dimethyl fumarate and fingolimod, efficacy and	
	safety were demonstrated in persons with RIS. Based on excellent biological plausibility, the early use of DMT is advocated in	•
	accumulation of neurological disability. There are emerging data to support this dogma. There is no reason to believe that a	
	would be biologically different from subsequent events that establish a diagnosis of CIS or MS. Thus, DMT should be offered to persons with RIS.	
14:00-14:10	Moderator: Joab Chapman, Israel	
	Introduction and Pre-Debate Voting	
14:10:14:25	Yes: Eva Havrdova, Czech Republic	
14:25-14:40	No: Klaus Schmierer, UK	
14:40-14:50	Discussion, Rebuttals and Post-Debate Voting	
14:50-15:20	Coffee Break, Exhibition & ePosters Visits	



	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
15:20-17:00	Multiple Sclerosis (continued)	HALL A
Chairs:	Andrijana Bogoje, Croatia; Larysa Sokolova, Ukraine	
15:20-16:10	Digital technology should replace neurological examination	
	Capsule: The neurological examination remains an important part of a patient's assessment, and its value has not been quest, students and neurology residents. A clinical provider can assess non-verbal cues, patient history, and subtle physical signs. He highly subjective and relies on a clinician's experience, intuition, and ability to observe subtle changes in a patient's behavior, a abilities. Digital technology holds the promise that it may augment neurological examination in numerous ways. Some of these reality, including advanced neuroimaging (like MRI or CT scans). Novel digital tests can track motor function, reflexes, and could and machine learning can assist in analyzing patterns in large datasets, which can enhance the accuracy of diagnoses. This debate will elucidate whether digital technology is capable of replacing the neurological examination all together by providata points.	owever, the physical examination is motor skills, speech, and cognitive se technologies are already clinical gnitive abilities. Artificial intelligence
15:20-15:30	Moderator: Larysa Sokolova, Ukraine Introduction and Pre-Debate Voting	
15:30-15:45	Yes: Letizia Leocani, Italy	
15:45-16:00	No: <u>Tjalf Ziemssen</u> , Germany	
16:00-16:10	Discussion, Rebuttals and Post-Debate Voting	
16:10-17:00	PET scanning should be a regular part of the follow up routine in patients with progressive MS	
	<b>Capsule</b> : The role of PET (Positron Emission Tomography) scanning in the routine follow-up of patients with progressive multipestablished. While PET scans can provide valuable metabolic and functional insights, their routine use in progressive MS follows:	, , ,
16:10-16:20	Moderator: Letizia Leocani, Italy Introduction and Pre-Debate Voting	
16:20-16:35	Yes: Friedemann Paul, Germany	
16:35-16:50	No: <u>Eva Havrdova</u> , Czech Republic	
16:50-17:00	Discussion, Rebuttals and Post-Debate Voting	
17:00-18:00	e-Posters Guided Tour	



	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
08:00-09:40	Stroke	HALL B
Chairs:	Michael Teitcher, Israel; Sadagat Huseynova, Azerbaijan	
08:00-08:50	POINT(S) and COMPASS(ES). Should stroke physicians use a combination of aspirin and low dose Rivaroxaban to reduce the risk of recurrent in high-risk people with large artery disease?	
	Capsule: The COMPASS trial demonstrated that people with stable atherosclerotic vascular disease who were treated with a cand aspirin had better cardiovascular outcomes but more bleeding than people treated with aspirin alone. Most of the particular to a history of myocardial infarction or peripheral vascular disease and people with a recent stroke were excluded. However, stroke despite being treated with antiplatelets. Is this a viable treatment option for people with stroke due to large artery diseantiplatelet therapy be preferable?  Moderator: Laszlo Csiba, Hungary	ipants were enrolled into the study ver, we see many patients who suffer
08:00-08:10	Introduction and Pre-Debate Voting	
08:10-08:25	Yes: Robert Gabor Kiss, Hungary	
08:25-08:40	No: <u>Jesse Dawson</u> , UK	
08:40-08:50	Discussion, Rebuttals and Post-Debate Voting	
08:50-09:40	Is AI a useful tool for making decisions in post stroke neurorehabilitation?	
	<b>Capsule</b> : Al can collect, assemble, and process huge amounts of data. This raises the question if Al tools can also be used to en neurorehabilitation, e.g., for planning and monitoring therapeutic interventions. This could increase the quality and speed of neurorehabilitation and help overcome problems with highly trained personnel, hence increasing the availability of intellectual problems with data security and uncertainties about whether Al is helpful for focalized decisions in the rehab process. In this capture is a security and uncertainties about whether Al is helpful for focalized decisions in the rehab process. In this capture is a security and uncertainties about whether Al is helpful for focalized decisions in the rehab process.	feeding information for processes in all resources. There are, however,
08:50-09:00	Moderator: Abraham Ohry, Israel Introduction and Pre-Debate Voting	
09:00:09:15	Yes: Volker Hoemberg, Germany	
09:15-09:30	No: Dafin Muresanu, Romania	
09:30-09:40	Discussion, Rebuttals and Post-Debate Voting	
09:40-10:10	Coffee Break, Exhibition & ePosters Visits	

FRIDAY, MARCH 21 <sup>ST</sup> , 2025		
10:10-11:10	Plenary Session	HALL A
Chairs:	Max J. Hilz, USA; Natan Bornstein, Israel	
10:10-10:40	Neurology is psychiatry and vice versa - Adam Zeman, UK	
10:40-11:10	Apraxia - Amos Korczyn, Israel	
11:10-12:10	Industry Sponsored Symposium	HALL A
12:10-13:10	Lunch Break, Exhibition & ePosters Visits	
13:10:14:50	Stroke (continued)	HALL B
Chairs:	<u>Dalius Jatuzis</u> , Lithuania; <u>Peter Klivenyi</u> , Hungary	
13:10-14:00	Should we offer endovascular treatment (EVT) to patients with acute stroke and pre-stroke mRS of 3 or more?	
	<b>Capsule</b> : Randomized trials with endovascular treatment (EVT) of acute stroke have excluded patients with pre-stroke modified Rankin scale (mRS) of more than "2". Despite lacking trial data, patients with higher mRS are offered EVT. Does the lack of trial data require additional studies in high mRS patients with LVO and acute stroke? Is the data from current trials sufficient to offer treatment regardless of the pre-stroke mRS?	
13:10-13:20	Moderator: Milija Mijajlovic, Serbia Introduction and Pre-Debate Voting	
13:20-13:35	Yes: <b>Ashfaq Shuaib</b> , Canada	
13:35-13:50	No: Roman Herzig, Czech Republic	
13:50-14:00	Discussion, Rebuttals and Post-Debate Voting	
14:00-14:50	Computed tomography perfusion (CTP) is rarely needed for decision making in patients with ischemic stroke	
	Capsule: Reason is that you do not know if there is large core until you have CTP. And CTP can offer other benefits beyond indication of mechanical thrombectomy (MT)	
14:00-14:10	Moderator: Robert Mikulik, Czech Republic Introduction and Pre-Debate Voting	
14:10-14:25	Yes: Michael Teitcher, Israel	
14:25-14:40	No: Ashfaq Shuaib, Canada	
14:40-14:50	Discussion, Rebuttals and Post-Debate Voting	
14:50-15:20	Coffee Break, Exhibition & ePosters Visits	



FRIDAY, MARCH 21 <sup>ST</sup> , 2025		
15:20-17:00	Stroke (continued)	HALL B
Chairs:	Zuzana Gdovinová, Slovakia; Michal Bar, Czech Republic	
15:20-16:10	There are sufficient data to use Andexanet alpha in people with intracerebral hemorrhage (ICH) associated with factor X inhibitor use	
	Capsule: In people with ICH associated with the use of FXa inhibitors, treatment with andexanet alfa reduces anti-FXa activity and has good hemostatic efficacy.	
	There are also reports that it is associated with lower mortality and better clinical outcomes. However, there may be an increa	ased risk of thrombotic events so the
	risk benefit ratio may be hard to define. Is there sufficient data to support routine use?	
15:20-15:30	Moderator: Jesse Dawson, UK	
15.20 15.50	Introduction and Pre-Debate Voting	
15:30-15:45	Yes: Mira Katan, Switzerland	
15:45-16:00	No: Ales Tomek, Czech Republic	
16:00-16:10	Discussion, Rebuttals and Post-Debate Voting	
16:10-17:00	Time to get the gout drugs out? Colchicine for prevention of stroke. Are you CONVINCED?	
	Capsule: The use of colchicine to prevent cardiovascular events in people with atherosclerotic coronary heart disease was rece	ently approved by the FDA. At least
	some of the benefits observed are due to a reduction in stroke. The CONVINCE and CHANCE-3 trials recently assessed this in po	eople with recent ischaemic stroke.
	Should we now be using this in people with ischaemic stroke?	
16:10-16:20	Moderator: Marina Roje Bedeković, Croatia	
10.10-10.20	Introduction and Pre-Debate Voting	
16:20-16:35	Yes: <b>Ashfaq Shuaib</b> , Canada	
16:35-16:50	No: Vida Demarin, Croatia	
16:50-17:00	Discussion, Rebuttals and Post-Debate Voting	
17:00-18:00	e-Posters Guided Tour	

	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
08:00-09:40	Parkinson's Disease (PD) II - Consensus and Controversy in PD Therapeutics	HALL C
Chairs:	Stuart Isaacson, USA; Rajesh Pahwa, USA	
08:05-08:20	Co-pathologies in neurodegenerative diseases - Radoslav Matej, Czech Republic	
08:20-09:00	Should on demand use of inhaled levodopa be used first-line for off episodes?	
	Capsule: OFF episodes are a common but still underrecognized problem in PD. OFF persists despite increasing oral levodopa and	nd adjunctive dopaminergic and
	dopaminergic therapies. Should first-line therapy include on demand inhaled levodopa treatment?	
08:20-08:25	Moderator: Rajesh Pahwa, USA	
	Introduction and Pre-Debate Voting	
08:25-08:40	No: <u>Daniel Kremens</u> , USA	
08:40-08:55	Yes: Richard Dewey III, USA	
08:55-09:00	Discussion, Rebuttals and Post-Debate Voting	
09:00-09:40	Panel Discussion: Should antipsychotics be used as soon as symptoms of PDP emerge?	
09:00-09:05	Moderator: <b>Stuart Isaacson,</b> USA	
	Introduction and Pre-Panel Voting	
09:05-09:35	Capsule: Antipsychotics have established efficacy in psychosis, but D2 antagonism and of target adverse effects can limit their	
	and Clozapine on the EU have regulatory approval for PDP. Should they be prescribed for early hallucinations and/or delusions	s emerge?
	Discussion: <u>Daniel Kremens</u> , USA; <u>Rajesh Pahwa</u> , USA	
09:35-09:40	Discussion, Rebuttals and Post-Panel Voting	
09:40-10:10	Coffee Break, Exhibition & ePosters Visits	
10:10-11:10	Plenary Session	HALL A
Chairs:	Max J. Hilz, USA; Natan Bornstein, Israel	
10:10-10:40	Neurology is psychiatry and vice versa - Adam Zeman, UK	
10:40-11:10	Apraxia - Amos Korczyn, Israel	
10.40 11.10	Aproxide Anno Rotelytty 151001	

	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
11:10-12:10	Industry Sponsored Symposium	HALL A
12:10-13:10	Lunch Break, Exhibition & ePosters Visits	
13:10-15:10	Parkinson's Disease (PD) II - Consensus and Controversy in PD Therapeutics (continued)	HALL C
Chairs:	Stuart Isaacson, USA; Daniel Kremens, USA	
13:10-13:50	Nondopaminergic mechanisms should routinely be added to levodopa when OFF fluctuations occur	
	<b>Capsule</b> : Despite increasing levodopa and adjunctive dopaminergic therapies, OFF time often persists. This may indicate the liderapies of the dopaminergic pathways to fully resolve OFF episodes. Striatal adenosine receptors are overactive in PD, and impact direct and nondopaminergic receptor antagonists be added to levodopa as soon as OFF fluctuations emerge?	
13:10-13:15	Moderator: Daniel Kremens, USA Introduction and Pre-Debate Voting	
13:15-13:30	Yes: Peter Jenner, UK	
13:30-13:45	No: Rajesh Pahwa, USA	
13:45-13:50	Discussion, Rebuttals and Post-Debate Voting	
13:50-14:30	Optimal PD clinical care should always include Wearables + AI	
	<b>Capsule</b> : Clinical recognition of OFF fluctuations and dyskinesia can be difficult in routine practice. The emergence of wearable and report these motor states, and combined with emerging AI will continue to improve recognition. Should wearable be used history or examination is unclear?	
13:50-13:55	Moderator: <u>Stuart Isaacson</u> , USA Introduction and Pre-Debate Voting	
13:55-14:10	Yes: Rajesh Pahwa, USA	
14:10-14:25	No: Richard Dewey III, USA	
14:25-14:30	Discussion, Rebuttals and Post-Debate Voting	



	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
14:30-15:10	Parkinson's Disease (PD) II - Consensus and Controversy in PD Therapeutics (continued)	HALL C
14:30-15:10	Dopamine agonists therapy on PD should avoid predominant D2-family receptor affinity	
	Capsule: Dopamine agonists emerged in the early levodopa era and were an important treatment option for decades. These in dopamine agonists. Their use has been associated with D2 associated side effects. Other dopamine agonists have D1- and D2 activity (i.e. apomorphine) or selective D1-family dopamine agonists (i.e. tavapadon) and avoid D2-family predominant side ef agonists be avoided?	-family ("dopamine-like") receptor
14:30-14:35	Moderator: Richard Dewey III, USA Introduction and Pre-Debate Voting	
14:35-14:50	Yes: Peter Jenner, UK	
14:50-15:05	No: <u>Stuart Isaacson</u> , USA	
15:05-15:10	Discussion, Rebuttals and Post-Debate Voting	
14:50-15:20	Coffee Break, Exhibition & ePosters Visits	

15:10-17:15	Parkinson's Disease (PD) II Consensus and Controversy in PD Therapeutics (continued)	HALL C
Chairs:	Avner Thaler, Israel	
15:10-15:50	Immediate-release CD/LD should always be replaced with extended-release CD/LD whenever OFF fluctuations emerge	
	<b>Capsule</b> : Immediate release levodopa/carbidopa is a foundational therapy but is limited by its short plasma half-life and varia onset and shortened duration of benefit. Should the extended release with mucoadhesive polymer formulation be used instead as soon as off fluctuations emerge?	
15:10-15:15	Moderator: Martin Bareš, Czech Republic Introduction and Pre-Debate Voting	
15:15-15:30	Yes: Richard Dewey III, USA	
15:30-15:45	No: <u>Daniel Kremens</u> , USA	
15:45-15:50	Discussion, Rebuttals and Post-Debate Voting	



	FRIDAY, MARCH 21 <sup>ST</sup> , 2025	
15:50-17:15	Parkinson's Disease (PD) II Consensus and Controversy in PD Therapeutics (continued)	HALL C
15:50-16:30	Adjunctive continuous subcutaneous apomorphine infusion should be considered as an early add-on therapy to baseline oral/transdermal therapies in all patients with OFF fluctuations	
	<b>Capsule</b> : Apomorphine has dopamine-like postsynaptic receptor activity and dopamine-like robust efficacy. Conversion of exogits subsequent release from presynaptic striatal nerve terminals is compromised with progression of PD neurodegeneration. Stapomorphine infusion be added as soon as levodopa fails to maintain good-ON time?	
15:50-15:55	Moderator: Rajesh Pahwa, USA Introduction and Pre-Debate Voting	
15:55-16:10	Yes: <b>Daniel Kremens</b> , USA	
16:10-16:25	No: Avner Thaler, Israel	
16:25-16:30	Discussion, Rebuttals and Post-Debate Voting	
16:30-17:10	Subcutaneous delivery replacement of oral levodopa should always be used before surgical options when motor fluctuations persist despite optimized oral therapy	
	<b>Capsule</b> : New treatments have recently emerged to treat PD, such as subcutaneous infusion of foslevodopa-foscarbidopa. Subcoral levodopa has been demonstrated to improve motor fluctuations, dyskinesia, morning and nocturnal akinesia, sleep, and of these therapies are minimally invasive and easy to implement, should they be considered as the first option before surgical op	quality of life in PD patients. Since
16:30-16:35	Moderator: <u>Diego Santos-Garcia</u> , Spain Introduction and Pre-Debate Voting	
16:35-16:50	Yes: Stuart Isaacson, USA	
16:50-17:05	No: Rajesh Pahwa, USA	
17:05-17:10	Discussion, Rebuttals and Post-Debate Voting	
17:10-17:15	Recap of Parkinson's Disease (PD) II and Closing Remarks  Rajesh Pahwa, USA; Stuart Isaacson, USA	

SATURDAY, MARCH 22 <sup>ND</sup> , 2025		
08:00-09:00	e-Posters Guided Tour	
09:00-10:40	Headache	HALL A
Chairs:	Magdalena Wysocka-Bakowska, Poland; Vlasta Vukovic Cvetkovic, Croatia	
09:00-09:50	anti-CGRP therapies should be first line for migraine prevention	
	Capsule: Insurance companies in the US and elsewhere make physicians use older preventive medications, in spite of poor efficacy and significant adverse effects.  New guidelines state that the anti- CGRP medications are effective and safer than older medications and should be used first line, even though they are more	
	expensive.  Moderator: <b>Tomas Nežádal,</b> Czech Republic	
09:00-09:10	Introduction and Pre-Debate Voting	
09:10-09:25	Yes: <b>Antoinette Maassen van den Brink</b> , The Netherlands	
09:25-09:40	No: <b>Gisela M. Terwindt</b> , The Netherlands	
09:40-09:50	Discussion, Rebuttals and Post-Debate Voting	
09:50-10:40	There is a need for a newer botulinum neurotoxins for prevention of chronic migraine	
	Capsule: OnabotulinumtoxinA is well established as a preventive treatment for chronic migraine. Is there a need for other similar biologics to be available for	
	migraine prevention which are more efficacious and act longer?	
09:50-10:00	Moderator: Alan Rapoport, USA	
09.30-10.00	Introduction and Pre-Debate Voting	
10:00-10:15	Yes: Peter McAllister, USA	
10:15-10:30	No: <b>Christian Lampl</b> , Austria	
10:30-10:40	Discussion, Rebuttals and Post-Debate Voting	
10:40-11:10	Coffee Break, Exhibition & ePosters Visits	
11:10-12:10	Plenary session	HALL A
Chairs:	Zvezdan Pirtošek, Slovenia, Andriy Dubenko, Ukraine	
11:10-11:40	What can neuropathology teach us in the era of biomarkers - Lea Grinberg, Brazil/USA	
11:40-12:10	Czech physicians and authors: their gifts to world medicine and culture - Abraham Ohry, Israel	
12:10-13:10	Lunch Break, Exhibition & ePosters Visits	



	SATURDAY, MARCH 22 <sup>ND</sup> , 2025
13:10-14:50	Headache (continued) HALL A
Chairs	<u>Ivan Milanov</u> , Bulgaria; <u>Natan Bornstein</u> , Israel
13:10-14:00	Psychedelics such as psilocybin and ketamine are reasonable treatment choices for both migraine and cluster headache
	Capsule: Psychedelic drugs such as psilocybin and ketamine are reasonably effective treatments for migraine and cluster headache in spite of strong adverse
	events. Should they be approved by the FDA and European authorities for these indications?
13:10-13:20	Moderator: Licia Grazzi, Italy
15.10-15.20	Introduction and Pre-Debate Voting
13:20-13:35	Yes: Peter McAllister, USA
13:35-13:50	No: <b>Christian Lampl</b> , Austria
13:50-14:00	Discussion, Rebuttals and Post-Debate Voting
14:00-14:50	Neurostimulation/modulation is as effective as pharmacotherapy for acute and preventive migraine treatment
	<b>Capsule</b> : Several electrical stimulation devices have been cleared by the FDA, as they appear to be effective and safe for migraine therapy. One is cleared for the acute and preventive treatment of cluster headache. Do they work as well as medications, are they safe and should they be used more often?
44.00.44.40	Moderator: <u>Tomas Nežádal</u> , Czech Republic
14:00-14:10	Introduction and Pre-Debate Voting
14:10-14:25	Yes: Miguel Lainez, Spain
14:25-14:40	No: Licia Grazzi, Italy
14:40-14:50	Discussion, Rebuttals and Post-Debate Voting
14:50-15:20	Coffee Break, Exhibition & ePosters Visits



	SATURDAY, MARCH 22 <sup>ND</sup> , 2025	
15:20-17:00	Headache (continued)	HALL A
Chairs	Marcin Kopka, Poland; Elsa Parreira, Portugal	
15:20-16:10	Even though migraine pathophysiology begins several days before symptom onset, treating acutely during prodrome/aura can be an effective strategy	
	Capsule: Migraine pathophysiology may begin several hours or days before the pain and disability start. Is it appropriate to treat patients during the prodrome	
	stage to prevent subsequent painful headaches and disability?	
15.20 15.20	Moderator: <u>Messoud Ashina</u> , Denmark	
15:20-15:30	Introduction and Pre-Debate Voting	
15:30-15:45	Yes: Gisela M. Terwindt, The Netherlands	
15:45-16:00	No: <u>Dimos D. Mitsikostas</u> , Greece	
16:00-16:10	Discussion, Rebuttals and Post-Debate Voting	
16:10-17:00	Medication underuse headache is a helpful concept which can prevent chronification and MOH	
	Capsule: Medication underuse headache is defined as a headache which begins when patients with severe and frequent attacks of	migraine do not get started on
	effective migraine preventives when they are eligible to do so and do not take rapid acting and effective medications as soon as the	headache begins to stop a
	migraine attack quickly. This results in medication underuse headache and causes both chronification and medication overuse head	lache (MOH), with significant
	consequences.	
16:10-16:20	Moderator: Alan Rapoport, USA	
10.10-10.20	Introduction and Pre-Debate Voting	
16:20-16:35	Yes: Wanakorn Rattanawong, Thailand	
16:35-16:50	No: Dimos D. Mitsikostas, Greece	
16:50-17:00	Discussion, Rebuttals and Post-Debate Voting	
17:00	Closing ceremony & Invitation to Krakow – Prof. Konrad Rejdak	



SATURDAY, MARCH 22 <sup>ND</sup> , 2025		
08:00-09:00	e-Posters Guided Tour	
09:00-10:40	Epilepsy	HALL B
Chairs:	<u>Lilach Goldstein</u> , Israel; <u>Ivan Rektor</u> , Czech Republic	
09:00-09:50	Are the newest drugs for epilepsy, cenobamate and fenfluramine better than the older drugs?	
	<b>Capsule</b> : Drugs introduced to treat epilepsy in the 1990's and 2000's did not produce seizure freedom at greater rates than old better?	er drugs. Are the newest drugs
09:00-09:10	Moderator: Maria Mazurkiewicz-Bełdzinska, Poland Introduction and Pre-Debate Voting	
09:10-09:25	Yes: Michael Sperling, USA	
09:25-09:40	No: <u>Andreja Bujan Kovač</u> , Croatia	
09:40-09:50	Discussion, Rebuttals and Post-Debate Voting	
09:50-10:40	Should we still use therapeutic drug monitoring when treating our patients with epilepsy?	
	Capsule: Does therapeutic drug monitoring really lead to better outcomes and seizure control or is management using clinical p	parameters adequate?
09:50-10:00	Moderator: Ruta Mameniskiene, Lithuania	
09.30-10.00	Introduction and Pre-Debate Voting	
10:00-10:15	Yes: <u>Ilan Blatt</u> , Israel	
10:15-10:30	No: Manjari Tripathi, India	
10:30-10:40	Discussion, Rebuttals and Post-Debate Voting	
10:40-11:10	Coffee Break, Exhibition & ePosters Visits	
11:10-12:10	Plenary session	
Chairs:	<u>Zvezdan Pirtošek</u> , Slovenia, <u>Andriy Dubenko</u> , Ukraine	HALL A
11:10-11:40	What can neuropathology teach us in the era of biomarkers - Lea Grinberg, Brazil/USA	
11:40-12:10	Czech physicians and authors: their gifts to world medicine and culture - Abraham Ohry, Israel	
12:10-13:10	Lunch Break, Exhibition & ePosters Visits	

SATURDAY, MARCH 22 <sup>ND</sup> , 2025			
13:10-14:50	Epilepsy (continued)	HALL B	
Chairs	Hadassa Goldberg-Stern, Israel; Nandan Yardi, India		
13:10-14:00	Should we use add-on therapy or substitution therapy for epilepsy when the first drug does not work?		
	Capsule: For most patients, is add-on and substitution of a new drug best when the first drug fails to control seizures? What is the evidence?		
13:10-13:20	Moderator: Elinor Ben Menachem, Sweden		
15.10-15.20	Introduction and Pre-Debate Voting		
13:20-13:35	Add on: Alla Guekht, Russia		
13:35-13:50	Substitution: Andreas Schulze-Bonhage, Germany		
13:50-14:00	Discussion, Rebuttals and Post-Debate Voting		
14:00-14:50	Case studies. Michael Sperling, USA		
14:00 14:40	Case Discussion: intractable epilepsy and seizure clusters. Established and novel therapies, and administration methods, inclu	ding trans-nasal.	
14:00-14:40	Michael Sperling, USA & Faculty		
14:40-14:50	Discussion		
14:50-15:20	Coffee Break, Exhibition & ePosters Visits		



	SATURDAY, MARCH 22 <sup>nd</sup> , 2025		
15:20-17:00	Epilepsy (continued)	HALL B	
Chairs	Andreja Bujan Kovač, Croatia; Tetyana Litovchenko, Ukraine		
15:20-16:10	Should we be targeting nuclei for deep brain stimulation other than the anterior thalamic nucleus for drug-resistant focal epilepsy?		
	Capsule: Stimulation of the anterior nucleus of the thalamus has been shown to reduce seizure frequency in a randomized controlled trial. Multiple sub thalamicnuclei, including pulvinar and centromedian, are being now stimulated instead in clinical practice. Is this justified?		
15:20-15:30	Moderator: Irena Dolezalova, Czech Republic Introduction and Pre-Debate Voting		
15:30-15:45	Yes: Elinor Ben-Menachem, Sweden		
15:45-16:00	No: Martin Holtkamp, Germany		
16:00-16:10	Discussion, Rebuttals and Post-Debate Voting		
16:10-17:00	Should we treat seizures that we see in the subclinical electrographic seizures in EEG in status epilepticus when clinical seiz	ures have stopped?	
	<b>Capsule</b> : It is common to see electrographic seizures after cessation of status epilepticus in the intensive care unit. Can we just aggressive therapy? Is there evidence to support improved outcome with or without treatment?	tify treating these with continued	
16:10-16:20	Moderator: Vladimir Komarek, Czech Republic Introduction and Pre-Debate Voting		
16:20-16:35	Yes: <u>Ilan Blatt</u> , Israel		
16:35-16:50	No: <u>Irena Dolezalova</u> , Czech Republic		
16:50-17:00	Discussion, Rebuttals and Post-Debate Voting		
17:00	Closing ceremony & Invitation to Krakow – Prof. Konrad Rejdak		



SATURDAY, MARCH 22 <sup>nd</sup> , 2025		
08:00-09:00	e-Posters Guided Tour	
09:00-10:40	Sleep	HALL C
Chairs:	Natan Bornstein, Israel; Elsa Parreira, Portugal	
09:00-09:50	Sleep enhances brain clearance of amyloid and other neurotoxic substances	
	Capsule: The hypothesis that sleep facilitates brain clearance of amyloid-6, tau, and other neurotoxic waste via the glymphatic system has gained significant traction over the last decade. Several preclinical studies demonstrated that slow-wave sleep may promote cerebrospinal fluid influx, enhancing perivascular clearance of metabolic byproducts; perhaps in keeping, other studies showed, and that sleep deprivation may accelerate A6 deposition. These findings have been taken to support sleep-based interventions as a potential neuroprotective strategy against AD. Nonetheless, direct clinical support for this process is still limited. Recent experimental observations have challenged the initial observations. To-date neither the glymphatic hypothesis nor the earlier classical hypothesis adequately explain how solutes and fluid move into, through and out of the brain parenchyma. We will revisit all the current evidence of mechanisms for extravascular transport into and out of the brain of hydrophilic solutes unable to cross the blood—brain barrier.	
09:00-09:10	Moderator: Claudio Bassetti, Switzerland Introduction and Pre-Debate Voting	
09:10-09:25	Yes: Lea Grinberg, Brazil/USA	
09:25-09:40	No: Ivana Rosenzweig, UK	
09:40-09:50	Discussion, Rebuttals and Post-Debate Voting	
09:50-10:40	Is sleep assessment essential in general neurology practice?	
	Capsule: Sleep is essential for brain, mental, physical and societal health. Brain integrity is on the other hand essential for a normal sleep-wake-circadian cycle Although the bidirectional relationship between sleep and neurological health and disorders is undeniable, sleep-wake circadian disturbances are often overlooked in neurology. Emerging evidence suggests that sleep loss/disturbances are not only a consequence but can also be a risk factor as well as a modul of neurological disorders. Insomnia, sleepiness/hypersomnia, sleep disordered breathing and parasomnias are prevalent in conditions such as stroke, dement epilepsy, movement disorders, MS, and headache syndromes, yet sleep history is rarely incorporated into standard neurological practice. As a consequence, integrating sleep-wake-circadian assessments may have a tremendous impact on the overall care of neurological patients, while ignoring can have negative effects such as increasing the risk of seizure or stroke recurrence, cognitive decline, mortality. Integrating sleep-wake circadian assessment in general neurolog practice is challenging because tools are not always validated and diagnostic approaches, such as polysomnography, can be resource-intensive and still not readily available in all settings. We challenge the audience to consider whether (and how) sleep-wake circadian assessment could become a standard compos of neurological evaluation or remain a specialized field for dedicated sleep medicine experts.	
09:50-10:00	Moderator: <u>Diego García-Borreguero</u> , Spain Introduction and Pre-Debate Voting	
10:00-10:15	Yes: Claudio Bassetti, Switzerland	
10:15-10:30	No: Ivana Rosenzweig, UK	

SATURDAY, MARCH 22 <sup>nd</sup> , 2025			
10:30-10:40	Discussion, Rebuttals and Post-Debate Voting		
10:40-11:10	Coffee Break, Exhibition & ePosters Visits		
11:10-12:10	Plenary session	HALL A	
Chairs:	<u>Zvezdan Pirtošek</u> , Slovenia, <u>Andriy Dubenko</u> , Ukraine		
11:10-11:40	What can neuropathology teach us in the era of biomarkers - Lea Grinberg, Brazil/USA		
11:40-12:10	Czech physicians and authors: their gifts to world medicine and culture - Abraham Ohry, Israel		
12:10-13:10	Lunch Break, Exhibition & ePosters Visits		
13:10-14:50	ALS	HALL C	
Chairs	Ervin Jancic, Croatia; Stanislav Sutovsky, Slovakia		
13:10-14:00	Physiological stress, as derived from smoking and extreme exercise, is a risk factor for Amyotrophic Lateral Sclerosis (ALS)		
	Capsule: Extreme physical activity and smoking have been linked to an increased risk of developing ALS. Physiological stress, when ongoing, extreme or		
	uncontrolled, may thus result in neurodegeneration, particularly with ALS		
13:10-13:20	Moderator: Pamela Shaw, UK		
	Introduction and Pre-Debate Voting		
13:20-13:35	Yes: <u>Amir Dori</u> , Israel		
13:35-13:50	No: Osman Sinanovic, Bosnia and Herzegovina		
13:50-14:00	Discussion, Rebuttals and Post-Debate Voting		
14:00-14:50	0 For neuroprotection in ALS - targeted therapies represent a better approach than therapeutic cocktails		
	Capsule: Function of the nervous system is largely dependent on energy supply, provided by oxygen, glucose and lipids. Interventions can target such less specific		
	factors (and others), but can also interfere with specific factors, such as disease-causing genes. Recently, specific treatment strategies – represented by tofersen		
	and nusinersen – were shown to be extremely successful, should these or non-specific cocktails be preferred?		
14:00-14:10	Moderator: <u>Amir Dori</u> , Israel		
14.00-14.10	Introduction and Pre-Debate Voting		
14:10-14:25	Yes: Pamela Shaw, UK		
14:25-14:40	No: <u>Albert Ludolph</u> , Germany		
14:40-14:50	Discussion, Rebuttals and Post-Debate Voting		
14:50-15:20	Coffee Break, Exhibition & ePosters Visits		



SATURDAY, MARCH 22 <sup>ND</sup> , 2025			
15:20-17:00	Neurodegenerative Diseases	HALL C	
Chairs	Ornit Chiba-Falek, USA; Radoslav Matej, Czech Republic		
15:20-16:10	The age-dependent decrease of brain clearing mechanisms is responsible for late-onset neurodegenerative diseases		
	<b>Capsule</b> : One of the main common features of neurodegenerative disorders is abnormal protein aggregation. This so-called 'proteinopathy' triggers different pathogenic events, such as alteration of axonal transport, loss of synapses and eventually cell loss in the brain. At the cellular and tissular levels, the brain possesses molecular debris clearing mechanisms. Is age-dependent decay of these clearing mechanisms responsible for proteinopathy in late-onset neurodegenerative diseases?		
15:20-15:30	Moderator: <u>Johannes Attems</u> , UK Introduction and Pre-Debate Voting		
15:30-15:45	Yes: <u>Bogdan Popescu</u> , Romania		
15:45-16:00	No: Radoslav Matej, Czech Republic		
16:00-16:10	Discussion, Rebuttals and Post-Debate Voting		
16:10-17:00	Palliative care should be discussed with people with progressive neurological disease early in the disease progression		
	<b>Capsule</b> : Unfortunately, there still are neurological disorders which cannot be healed or slowed down in their progression, such neurodegenerative pathogenic background. Once diagnosed, the prognosis is estimated, including a time frame of neurological devastating conditions, is it important to inform patients about palliative care options and procedures in the early disease progression.	I function deterioration. For these	
16:10-16:20	Moderator: Bogdan Popescu, Romania Introduction and Pre-Debate Voting		
16:20-16:35	Yes: Robert Rusina, Czech Republic		
16:35-16:50	No: Vladimira Vuletic, Croatia		
16:50-17:00	Discussion, Rebuttals and Post-Debate Voting		
17:00	Closing ceremony & Invitation to Krakow – Prof. Konrad Rejdak		