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EISAI TO PRESENT LATEST DATA ON PERAMPANEL AND RUFINAMIDE AT 70TH AMERICAN EPILEPSY SOCIETY ANNUAL MEETING

Eisai Co., Ltd. (Headquarters: Tokyo, CEO: Haruo Naito, "Eisai") announced today that the latest data on its antiepileptic drugs (AED) perampanel (product name: Fycompa[®]) and rufinamide (product name: Inovelon[®], U.S. product name: BANZEL[®]) will be presented at the 70th American Epilepsy Society (AES) Annual Meeting to be held from December 2 to 6 in Houston in the United States.

For this year's AES meeting, poster presentations will be given on eight abstracts for perampanel which include the results of Phase II trials (Study 231, Study 233) of adjunctive perampanel in Japanese patients with refractory partial-onset seizures. Regarding rufinamide, a poster presentation will be given on the safety and cognitive development effects of adjunctive rufinamide in pediatric subjects with inadequately controlled Lennox-Gastaut Syndrome (LGS) from the final results of Study 303. Along with three poster presentations on health economics and outcome research, a total of 12 poster presentations will be given.

Perampanel is a first-in-class AED discovered and developed by Eisai. With epileptic seizures being mediated by the neurotransmitter glutamate, the agent is a highly selective, noncompetitive AMPA receptor antagonist that reduces neuronal hyperexcitation associated with seizures by targeting glutamate activity at postsynaptic AMPA receptors. The agent is currently approved in more than 50 countries and territories as an adjunctive treatment of partial-onset seizures (with or without secondarily generalized seizures) in adult and adolescent patients with epilepsy 12 years of age and older. In addition, perampanel is approved in more than 40 countries as an adjunctive treatment of primary generalized tonic-clonic (PGTC) seizures in patients with epilepsy 12 years of age and older.

Furthermore, Eisai has submitted a supplemental application to the U.S. Food and Drug Administration for a partial label change for perampanel as monotherapy for treatment of partial-onset seizures in patients with epilepsy 12 years of age and older.

Rufinamide is believed to exert its antiepileptic effects by regulating activity of voltage-gated sodium channels in the brain involved in the overexcitement of neurons that potentially causes seizures, so as to prolong their inactive state. The agent is approved as an adjunctive therapy to other AEDs in the treatment of seizures associated with LGS in Europe and the United States. In Japan, the agent is approved as an adjunctive therapy to other AEDs in the treatment of tonic and atonic seizures associated with LGS when therapy with other AEDs is considered inadequate. Rufinamide is currently approved in more than 30 countries worldwide.

Eisai considers epilepsy a therapeutic area of focus and by providing multiple treatment options in addition to perampanel and rufinamide as part of an extensive epilepsy product portfolio, Eisai seeks to make continued contributions to address the diverse needs of, as well as increasing the benefits provided to, patients with epilepsy and their families.

Major Poster Presentations for perampanel:

Abstract number	Abstract title
Abstract number: #2.189	
Sunday December 4	Evaluation of perampanel as monotherapy for focal seizures: experience from
Poster presentation: 10:00-16:00	open-label extension studies
Poster discussion: 12:00-14:00	·
Abstract number: #2.190	
Sunday December 4	Adjunctive perampanel (PER) in patients (pts) with partial seizures or primary
Poster presentation: 10:00-16:00	generalized tonic-clonic seizures (PGTCS): effect of age at diagnosis
Poster discussion: 12:00-14:00	, , , ,
Abstract number: #2.222	
Sunday December 4	A Cystomatic Paviany of Paul Warld Pavarra and Treatment Outcomes
Poster presentation: 10:00-16:00	A Systematic Review of Real World Perampanel Treatment Outcomes
Poster discussion: 12:00-14:00	
Abstract number: #2.225	An Indirect Treatment Comparison (ITC) of Perampanel versus Brivaracetam
Sunday December 4	
Poster presentation: 10:00-16:00	in Patients with Partial-Onset Seizures (POS) With or Without Secondary
Poster discussion: 12:00-14:00	Generalization
Abstract number: #2.193	
Sunday December 4	Long-term efficacy and safety of adjunctive perampanel: pooled analyses of
Poster presentation: 10:00-16:00	the open-label extension (OLE) studies
Poster discussion: 12:00-14:00	
Abstract number: #2.191	Phase II trials of adjunctive perampanel in Japanese patients with refractory
Sunday December 4	
Poster presentation:10:00-16:00	partial-onset seizures, an open-label, ascending-high-dose study (study 231)
Poster discussion: 12:00-14:00	and long-term extension study (study 233)
Abstract number: #3.238	
Monday December 5	Adjunctive Perampanel in Patients With Drug-Resistant Partial Seizures With
Poster presentation: 8:00-14:00	and Without Concurrent VNS Therapy in Phase III Studies
Poster discussion: 12:00-14:00	.,
Abstract number: #3.255	
Monday December 5	Health Care Resource Utilization Before and After Perampanel Initiation for
Poster presentation: 8:00-14:00	the Treatment of Epilepsy in the United States
Poster discussion: 12:00-14:00	

Major Poster Presentations for rufinamide:

Abstract number	Abstract title
Abstract number: #3.363 Monday December 5 Poster presentation: 12:00-18:00 Poster discussion: 12:00-14:00	Safety and Cognitive Development Effects of Adjunctive Rufunamide in Pediatric Subjects With Inadequately Controlled Lennox-Gastaut Syndrome(LGS): Final Results From Study 303

Major Poster Presentations on Health Economics and Outcome Research:

Abstract number	Abstract title
Abstract number: #2.332	
Sunday December 4	Racial Differences in the Incidence and Prevalence of Epilepsy in United
Poster presentation: 10:00-16:00	States Veteran Population
Poster discussion: 12:00-14:00	· ·

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Abstract number	Abstract title
Abstract number: #2.335	
Sunday December 4	Estimating the Economic Burden of Caregiving in Epilepsy
Poster presentation: 10:00-16:00	
Poster discussion: 12:00-14:00	
Abstract number: #2.338	The Caregiver's Direct Medical Costs in Epilepsy
Sunday December 4	
Poster presentation: 10:00-16:00	
Poster discussion: 12:00-14:00	

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