The 53rd Annual Meeting of the Japanese Society of Child Neurology

May 26-28, 2011 Pacifico Yokohama, Japan

PROGRAM

Wednesday, May 25, 2011

0.00	ROOM E 303	ROOM F 304
8:00		
9:00		
10:00		
11:00		
12:00		
13:00		
14:00		
15:00		
16:00		
		Practical Education Seminar 1 15:00-18:00
17:00		13.00-16.00
18:00		
	Evening Seminar	
19:00	18:00-19:00	
20:00		
20.00		Practical Education Seminar 2 19:00-22:00
21.00		
21:00		

Thursday, May 26, 2011

	ROOM A Main Hall	ROOM B 503	ROOM C 501		
8:00					
9:00	Opening Remarks 8:50-9:00				
	Presidential Lecture 9:00-9:30				
10:00		Educational Lecture 1			
	Symposium 1	9:40-10:25	Washahan 1		
	9:30-11:30	Educational Lecture 2	Workshop 1 9:40-11:30		
11:00		10:25-11:10			
12:00	Luncheon Seminar 1	Luncheon Seminar 2	Luncheon Seminar 3		
	11:45-12:45	11:45-12:45	11:45-12:45		
12.00					
13:00		English Session 1	Educational Lecture 3		
		13:00-14:00 (O-001-005)	13:00-13:45		
14:00	Symposium 2	(0-001-003)	Educational Lecture 4		
	13:00-15:00	English Session 2	13:45-14:30		
		14:00-15:00 (O-006-010)			
15:00					
	Special Lecture 1	Invited Lecture 1			
	15:10-16:00	15:10-16:00			
16:00					
	Keynote Lecture 16:00-16:50				
17.00					
17:00					
18:00					
19:00					
		Banquet Inter Continental Yokohama Grand 3F			
20:00		Inter Continental Ball Room 19:00-21:00			
		17.00-21.00			
21:00					
21.00					

	ROOM D 502	ROOM E 303	ROOM F 304	Poster Session Room 301+302/315
8:00				
9:00				
10:00	Developmental Disorder (Screening)	Genetics 1 9:40-10:40	Neonates 9:40-10:40	
	9:50-10:40 (O-011-015)	(O-021-026)	(O-043-048)	
11:00	Developmental, Psychiatric & Behavioral Disorder	Genetics 2 10:40-11:30		
	10:40-11:30 (O-016-020)	(O-027-031)		
12:00				
12.00	Luncheon Seminar 4 11:45-12:45	Luncheon Seminar 5 11:45-12:45	Luncheon Seminar 6 11:45-12:45	
13:00		Learning Disabilities 1	Epilepsy & Seizure 1	
	Symposium 3 13:00-15:00	13:00-14:00 (O-032-037)	13:00-14:00 (O-049-054)	
14:00		Learning Disabilities 2	Enilongy & Saignus 2	
		14:00-14:50 (O-038-042)	Epilepsy & Seizure 2 14:00-15:00 (O-055-060)	
15:00				
16:00				
17:00				
17.00				
				Poster Presentation $17:00 \sim$ (P-001-P-179)
18:00				(1-001-1-177)
19:00				
20:00				
21:00				

Friday, May 27, 2011

	ROOM A	ROOM B	ROOM C
8:00	Main Hall	503	501
	Morning Education Seminar 1	Morning Seminar 1 8:00-8:50	
	8:00-9:00	0.00-0.50	
9:00			
	Invited Lecture 2 9:00-9:50	Epilepsy & Seizure 3	Developmental Disorders 1 9:00-10:00
	9.00-9.30	9:10-10:00 (O-061-065)	(O-072-077)
10:00	Invited Lecture 3		
	9:50-10:40	Epilepsy & Seizure 4	Developmental Disorders (Treatment & Support)
	2011 JSCN Award for Asia Young	10:00-11:00 (O-066-071)	10:00-11:00 (O-078-083)
11:00	Investigator 10:40-11:10		(* * * * * * * * * * * * * * * * * * *
	Asian Oceanian Child Neurology International Education Program	Educational Lecture 5	Development
	11:10-11:40	11:00-11:45	11:00-12:00 (O-084-089)
12:00			
	Luncheon Seminar 7	Luncheon Seminar 8	Luncheon Seminar 9
13:00	12:15-13:15	12:15-13:15	12:15-13:15
14:00	JSCN General Assembly 13:30-14:30		
	13.30-14.30		
	Presentation of JSCN Awards to Outstanding Young Investigators	_	
15:00	14:30-15:00		
		Educational Lecture 6 15:10-15:55	
16:00	Symposium 4 15:00-17:00		B & D Seminar
	13.00-17.00	Educational Lecture 7 15:55-16:40	15:00-17:00
		10.00 10.10	
17:00			
18:00			
19:00			
20:00		Night Seminar 1	
	E E	19:00-21:00	
	Emergency Forum 19:00-22:00		
21:00			
_1.00			
6			

0.00	ROOM D 502	ROOM E 303	ROOM F 304	Poster Session Room 301+302/315
8:00		Morning Seminar 2 8:00-8:50	Morning Seminar 3 8:00-8:50	
9:00				
10:00	Peripheral Nerve 9:10-10:00 (O-090-094)	Acute Encephalitis & Encephalopathy 3 9:10-10:00 (O-117-121)		
10.00	Muscular Disease 1 10:00-10:50 (O-095-099)	Acute Encephalitis & Encephalopathy 4 10:00-11:00 (O-122-127)	Involuntary Movement & Basal Ganglia 9:50-10:50 (O-145-150)	
11:00	Muscular Disease 2 10:50-12:00 (O-100-106)	Acute Encephalitis & Encephalopathy 5 11:00-12:00 (O-128-133)	Epilepsy & Seizure 5 10:50-12:00 (O-151-157)	
12:00				
13:00	Luncheon Seminar 10 12:15-13:15	Luncheon Seminar 11 12:15-13:15	Luncheon Seminar 12 12:15-13:15	
14:00				
15:00				
16:00	Acute Encephalitis & Encephalopathy 1 15:20-16:00 (O-107-110)	Developmental Disorders 2 15:10-16:00 (O-134-138)	Congenital Anomaly & Chromosomal Abnormality 15:20-16:00 (O-158-161)	
10.00	Acute Encephalitis & Encephalopathy 2 16:00-17:00 (O-111-116)	Developmental Disorders 3 16:00-17:00 (O-139-144)	Brain Tumor & Neurosurgery 16:00-17:00 (O-162-167)	
17:00				
18:00				Poster Presentation 17:10 ~ (P-180-P-355)
19:00				
	Night Seminar 2 19:00-20:00			
20:00				
21:00				

Saturday, May 28, 2011

	ROOM A Main Hall	ROOM B 503	ROOM C 501
8:00	Morning Education Seminar 2 8:00-9:00	Morning Seminar 4 8:00-8:50	Morning Seminar 5 8:00-8:50
9:00	Special Lecture 2 9:00-9:50		Cerebral Palsy & Care 9:10-10:00
10:00		Educational Lecture 8 9:45-10:30	(O-168-172)
11:00	Workshop 2 10:00-12:00	Educational Lecture 9 10:30-11:15	Panel Discussion 10:00-12:00
12:00	Closing Remarks 12:00-12:10	Educational Lecture 10 11:15-12:00	
12.00			
13:00			
14:00			
15:00	Public Forum 13:30-16:20	Educational Seminar 13:30-16:30	The 6th Meeting of the Japanese Society for Children's Myasthenia Gravis 13:00-17:00
16:00			
17:00			
18:00			
19:00			
20:00			
21:00			

	ROOM D 502	ROOM E 303	ROOM F 304
8:00			
9:00			
7.00	Epilepsy & Seizure 6	Infection & Autoimmune Disease	Metabolic Disease 1
	9:10-10:00	9:00-9:50 (O-190-194)	9:00-10:10
10:00	(0-173-177)		(O-208-213)
	Epilepsy & Seizure 7	Imaging 9:50-11:00	
	10:00-11:00 (O-178-183)	(O-195-201)	Metabolic Disease 2 10:10-11:10
11:00			(O-214-219)
	Epilepsy & Seizure 8 11:00-12:00	Autonomic Nerves, Headache & Vascular Disease	Metabolic Disease 3
	(O-184-189)	11:00-12:00 (O-202-207)	11:10-12:00 (O-220-225)
12:00			
13:00			
14:00	The 5th Child Sleep Study Group Meeting		
14.00	13:00-15:00		
15:00			
16:00			
17:00			
18:00			
10.00			
19:00			
20:00			
21:00			

Poster Session: Thursday, May 26, 2011

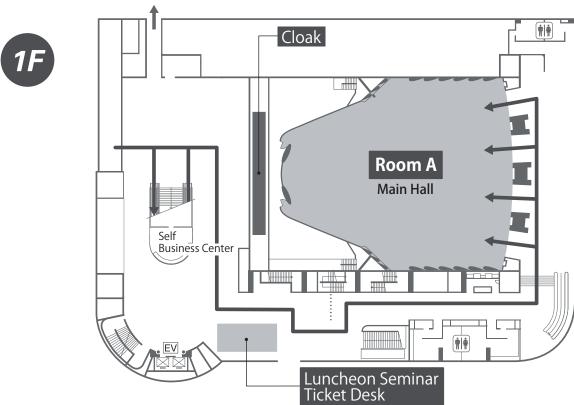
8:00	ROOM 301+302								ROOM 315		
17:00	Poster Session							Poster Session Poster Se			
77.00	Muscular Disease 1 17:00-17:40 (P-001 ~ 008)	Developmental Disorders 1 17:00-17:45 (P-017 ∼ 025)	Epilepsy & Seizure 1 17:00-17:45 (P-036 ~ 044)	Cerebral Palsy & Care (The Reality, Home & Medical Care) 17:00-17:45 (P-054 ~ 062)	Developmental Disorders (Treatment & Support) 1 17:00-17:45 (P-072 ~ 080)	Imaging 1 17:00-17:45 (P-090 ~ 098)	Acute Encephalitis & Encephalopathy 1 17:00-17:40 (P-108 ~ 115)	Infection & Autoimmune Disease I 17:00-17:45 (P-125 ~ 133)	Metabolic Degeneration 1 17:00-17:50 (P-143 ∼ 152)	Neonates 17:00-17:50 (P-162 ~ 171)	
18:00	Muscular Disease 2 17:40-18:20 (P-009 ~ 016)	Developmental Disorders 2 17:45-18:35 (P-026 ~ 035)	Epilepsy & Seizure 2 17:45-18:30 (P-045 ~ 053)	Cerebral Palsy & Care (Treatment & Gastrostoma) 17:45-18:30 (P-063 ~ 071)	Developmental Disorders (Treatment & Support) 2 17:45-18:30 (P-081 ~ 089)	Imaging 2 17:45-18:30 (P-099 ~ 107)	Acute Encephalitis & Encephalopathy 2 17:40-18:25 (P-116 ~ 124)	Infection & Autoimmune Disease 2 17:45-18:30 (P-134 ~ 142)	Metabolic Degeneration 2 17:50-18:35 (P-153 ∼ 161)	Vascular Disease 17:50-18:30 (P-172 ∼ 179)	
19:00				Poster Removal					Poster Removal		

Poster Session: Friday, May 27, 2011

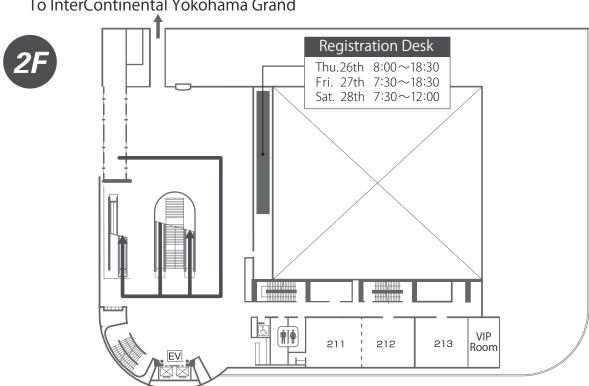
8:00	ROOM 301+302								ROOM 315	
	Poster Session								Poster Session	
17:00	Epilepsy & Seizure 4 17:10-17:55 (P-180 ~ 188)	Epilepsy & Seizure 6 17:10-18:05 (P-198 ~ 208)	Cerebral Palsy & Care 17:10-17:55 (P-216 ~ 224)	Psychiatric & Behavioral Disorder 17:10-17:55 (P-234 ~ 242)	Developmental Disorder (Screening) 17:10-17:55 (P-252 ~ 260)	Acute Encephalitis & Encephalopathy 3 17:10-17:55 (P-270 ~ 278)	Infection & Autoimmune Disease 3 17:10-17:55 (P-285 ~ 293)	Metabolic Degeneration & Muscles 17:10-17:45 (P-303 ~ 309)	Muscles 17:10-17:55 (P-321 ~ 329)	Congenital Anomaly & Chromosomal Abnormality 1 17:10-17:55 (P-338 ~ 346)
18:00	Epilepsy & Seizure 5 17:55-18:40 (P-189 ~ 197)	Learning Disabilities 18:05-18:40 (P-209 ~ 215)	Cerebral Palsy & Care (Respiratory Management & Eating) 17:55-18:20 (P-225 ~ 229) Autonomic Nerves & Headache 18:20-18:40 (P-230 ~ 233)	Development 17:55-18:40 (P-243 ~ 251)	Developmental Disorders 3 17:55-18:40 (P-261 ∼ 269)	Acute Encephalitis & Encephalopathy 4 17:55-18:25 (P-279 ~ 284)	Acute Encephalitis & Encephalopathy 5 17:55-18:40 (P-294 ~ 302)	Genetics 17:45-18:40 (P-310 ~ 320)	Peripheral Nerve 17:55-18:35 (P-330 ~ 337)	Congenital Anomaly & Chromosomal Abnormality 2 17:55-18:40 (P-347 ~ 355)
19:00				Poster Removal					Poster Removal	

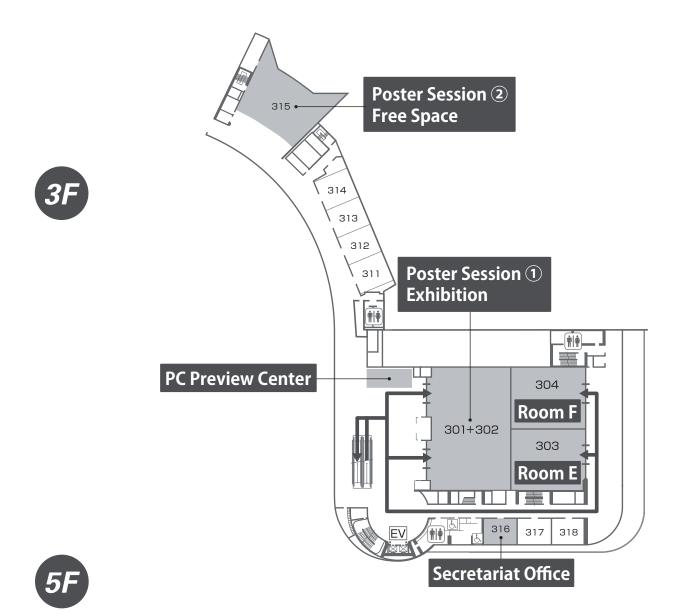
Floor Plan

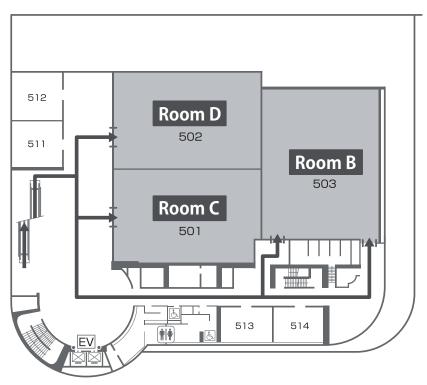
To InterContinental Yokohama Grand



To InterContinental Yokohama Grand







Presidential Lecture

Advancement and challenges in genetic disorders of copper metabolism

HIROKO KODAMA

Department of Pediatrics, Teikyo University School of Medicine, Tokyo, Japan

Keynote Lecture

On Political Measures for Intractable Diseases

ICHIRO KANAZAWA

President of the Science Council of Japan, Professor Emeritus of the University of Tokyo, Japan

Special Lecture 1

Exploring molecular etiologies of neurodegenerative diseases based on personal genome analysis

SHOJI TSUJI

Department of Neurology, The University of Tokyo Hospital, Japan

Special Lecture 2

Toward the Age of Dialogue

ORIZA HIRATA

The Osaka University Center for the Study of Communication-Design, Osaka, Japan

Invited Lecture 1

Latest Advances in the Fatty Acid Oxidation Disorders and New Therapies

INGRID TEIN

Division of Neurology, Hospital for Sick Children, Department of Pediatrics, Laboratory Medicine and Pathobiology, University of Toronto, Toronto, Ontario, Canada

Invited Lecture 2

Evidence-baced Practices for ADHD: Psychosocial, Pharmacological, and Combined Intervention

WILLIAM E. PELHAM

Center for children and Families, Florida International University, Miami, FL, USA

Invited Lecture 3

The U.S. Experience with Urea Cycle Defects: A Changing and Improving Picture

MARSHALL L. SUMMAR

Children's National Medical Center, Department of Pediatrics, The George Washington University School of Medicine and Health Sciences, Washington, NW, USA

Educational Lecture 1

Recent advances in congenital myopathy research

IKUYA NONAKA

National Center Hospital of Neurology and Psychiatry, Tokyo, Japan

Educational Lecture 2

Newborn hearing screening-Innovation of medicine for deaf infant and children

KIMITAKA KAGA

National Institute of Sensory Organs, National Tokyo Medical Center, Japan

Educational Lecture 3

Childhood epilepsy update

HIROKAZU OGUNI

Department of Pediatrics, Tokyo Women's Medical University, Tokyo, Japan

Educational Lecture 4

Recent management of status epilepticus in children

KENJI SUGAI

Department of Child Neurology, National Center Hospital, National Center of Neurology and Psychiatry, Kodaira, Japan

Educational Lecture 5

Current Directions in Early Diagnosis and Early Intervention for Autism Spectrum Disorders

YOKO KAMIO

Department of Child and Adolescent Mental Health, National Institute of Mental Health, National Center of Neurology and Psychiatry, Japan

Educational Lecture 6

Adolescence of children with developmental Disabilities

MIKIO HIRAIWA

Rabbit Developmental Research, Japan

Educational Lecture 7

Medical Treatment and Special Needs Education in Patients with Developmental Disorders

YASUSHI FUJII

Graduate School of Teacher Education, Teikyo University, Tokyo, Japan Department of Pediatrics, Faculty of Medicine, Teikyo University, Tokyo, Japan

Educational Lecture 8

Pediatric neurotransmitter disease in Japan

HARUO SHINTAKU

Department of Pediatrics, Osaka City University Graduate School of Medicine, Osaka, Japan

Educational Lecture 9

Recent advances on the etiology and pathogenesis of acute encephalopathy

MASASHI MIZUGUCHI

Department of Developmental Medical Sciences, the University of Tokyo, Tokyo, Japan

Educational Lecture 10

Imaging diagnosis of pediatric acute brain disorders

HIROSHI OBA

Department of Radiology, Teikyo University Hospital, Tokyo, Japan

Symposium 1

Epidemiology of Headache in children

KENTARO KUWABARA

Department of Pediatrics, Nippon Medical School, Tokyo, Japan

Diagnosis and treatment of pediatric patients with migraine and tension-type headache

KIYOSHI ARAKI

Saitama City Hospital, Saitama, Japan

Childhood periodic syndromes that are commonly precursors of migraine

TOSHIYUKI HIKITA

Department of Pediatrics, Teikyo University School of Medicine, Tokyo, Japan

Secondary headaches in children and adolescents

NAOKI ANDO

Department of Pediatrics and Neonatology, Nagoya City University, Graduates School of Medical Sciences, Nagoya, Japan

Medical treatment of headache in children and psychosomatic medicine

HIDEHIRO YASUJIMA

Department of Pediatrics, Ono Hospital, Hyogo, Japan

Symposium 2

Pediatric multiple sclerosis

RYUTARO KIRA

Department of Pediatrics, National Fukuoka-Higashi Medical Center, Japan

Clinical Features of Japanese Pediatric Patients with Anti-Aquaporin 4 Antibody, in Japan

NAOMI HINO-FUKUYO

Department of Pediatrics, University of Tohoku, Sendai, Japan

Clinical features of acute disseminated encephalomyelitis in childhood

HIROYUKI TORISU

Comprehensive Maternity and Perinatal Care Center, Kyushu University Hospital, Kyushu University, Fukuoka, Japan

Autoimmune cerebellar ataxia

TAKASHI ICHIYAMA

Department of Pediatrics, Yamaguchi University Graduate School of Medicine, Ube, Japan Japanese Study Group of Child Immune Encephalitis, Japan

Immunological mechanisms in non-herpetic acute limbic encephalitis

YUKITOSHI TAKAHASHI

National Epilepsy Center, Shizuoka, Japan

Gifu Unversity School of Medicine, Gifu, Japan.

Evaluation of clinical features and prevalence of Lyso-ganglioside antibodies of children with neuropsychiatric disorders associated with streptococcal infections

KAZUHISA HONGOU

Department of Pediatrics, University of Toyama, Toyama, Japan

Toyama Prefectural Hospital for Clippler Children, Shimoiino Toyama City, Japan

Symposium 3

The up-to-date researches of Rett syndrome

MASAYUKI ITOH

National Center of Neurology and Psychiatry, Japan

Yoshiko Nomura

Segawa Neurological Clinic for Children, Tokyo, Japan

Mutation analysis of the responsible genes for Rett syndrome

SATORU TAKAHASHI

Department of Pediatrics, Asahikawa Medical University, Asahikawa, Japan

MeCP2 and Chromatin Dynamics

SHIN-ICHI HORIKE

Kanazawa University Frontier Science Organization, Japan

Development of Rett-syndrome model cells by stem cell technologies

TOMOYUKI TAKAHASHI

Division of Gene Therapy and Regenerative Medicine, Cognitive and Molecular Research Institute of Brain Diseases, Kurume University, Japan

Therapeutic perspective for Rett syndrome patients

TAKEO KUBOTA

Department of Epigenetic Medicine, Faculty of Medicine, University of Yamanashi, Japan

Symposium 4

From a viewpoint of the medical cooperation

SUNAO TOMITA

Tokyo Metropolitan Children's Medical Center, Tokyo, Japan

From the view point of home care for adults

REIKO NAMBA

Neurology Clinic Namba, Okayama, Japan

Home Care Medicine and Respite Care for Children

AKIHIKO TAKAHASHI

Hibari Clinic, Utsunomiya-city, Japan

About the Tokyo Home Visit Service for Persons (Children) with Severe Mental - Physical Disability

NORIKO OKUYAMA

Residential and Facility Care Support, Disabled Persons Programs Division, Bureau of Social Welfare and Public Health, Tokyo Metropolitan Government, Japan

From the standpoint of care-home and at home

KEIKO SHISHIKURA

Social Welfare Corporation 'Houmonnoie' Tomo Cninic for Handicapped, Yokohama, Japan

Workshop 1

Infantile acute subdural hematoma-accident or abuse?

MAMI YAMASAKI

Department of Neurosurgery, Osaka National Hospital, National Hospital Organization, Japan

Imaging Diagnosis of Child Abuse

NORIKO AIDA

Department of Radiology, Kanagawa Children's Medical Center, Yokohama, Japan

Investigation of the mechanism of infantile acute subdural hematoma using an infant physical dummy model

YUSUKE MIYAZAKI

Institute of Science and Engineering, Kanazawa University, Ishikawa, Japan

Workshop 2

KATSUNORI FUJII

Vice Chairperson of the Council to Promote Reform of the System for Persons with Disabilities of Cabinet Office, Government Japan

Reform of Special Needs Education

NAOTO SHIMOYAMA

Ministry of Education, Culture, Sports, Science and Technology, Japan

Our action to reconstruction of support and education system for handicapped children and people

EIJI KITAZUMI

National Rehabilitation Center for Disabled Children, Tokyo, Japan

KENJI TAKAKI

Department of Health and Welfare for persons with Disabilities Ministry of Health, Labor and Welfare, Japan

Panel Discussion

abnormal movements in children with cerebral palsy

KENJI YOKOCHI

Department of Pediatric Neurology, Seirei-Mikatahara General Hospital, Hamamatsu, Shizuoka, Japan

Abnormal movements of neonate and early infancy

MASANORI KOUWAKI

Department of Neonatal Medical Center, Toyohashi Municipal Hospital, Aichi, Japan

Hypertonic state and botulinum toxin A therapy

MIZUE IAI

The Institution for Severe Intellectual and Motor Disabilities, Knagawa Children's Medical Center, Kanagawa, Japan

Deep brain stimulation for dystonia in children

SATOKO KUMADA

Department of Neuropediatrics, Tokyo Metropolitan Neurological Hospital, Tokyo, Japan

Intrathecal Bacrofen (ITB) and Functional Posterior Rhizotomy (FPR) for patients with intractable hypertonic state

MASAYA KUBOTA

Division of Neurology, NCCHD, Tokyo, Japan

Morning Education Seminar 1

The study for the diagnosis and the treatment of fragile X syndrome in Japan

EIJI NANBA

Division of Functional Genomics, Research Center for Bioscience and Technology, Tottori University, Japan

Epidemiologic study of acute encephalitis with refractory, repetitive partial seizures

HIROSHI SAKUMA

Department of Immunology, National Institute of Neuroscience, National Center of Neurology and Psychiatry, Tokyo, Japan

Establishment of medicinal treatment in infant with autism and other pervasive developmental disorders

EIJI NAKAGAWA

Department of Child Neurology, National Center Hospital, NCNP, Tokyo, Japan

Studies on prevalence, diagnosis and treatment of Menkes disease and occipital horn syndrome

HIROKO KODAMA

Department of Pediatrics, Teikyo University School of Medicine, Tokyo, Japan

Practical investigation for 1p36 deletion syndrome and establishment of the clinical guideline

TOSHIYUKI YAMAMOTO

Tokyo Women's Medical University Institute for Integrated Medical Sciences, Tokyo, Japan

Morning Education Seminar 2

Our experience of steroid therapy in patients with Duchenne muscular dystrophy

KEIKO ISHIGAKI

Department of Pediatrics, Tokyo Women's Medical University, School of Medicine, Tokyo, Japan

Steroid therapy for DMD in our hospital

TATSUYA FUJII

Department of Pediatrics, Shiga Medical Center for Children, Moriyama, Japan

Steroid therapy of Duchenne muscular dystrophy

HIROFUMI KOMAKI

Department of Child Neurology, National Center of Neurology and Pschiatry, Tokyo, Japan

Corticosteroid treatment in children with Duchenne muscular dystrophy: A view of rehabilitation medicine

YUKA ISHIKAWA

Department of pediatrics, National organization Yakumo hospital, Yakumo-cho, Futami-gun, Hokkaido, Japan

Corticosteroid treatment of Duchenne muscular dystrophy - a view of neurologist

MITSURU KAWAI

Department of Neurology, Higashisaitama National Hospital, Hasuda, Japan

Guidline for steroid therapy for DMD patients in Japan

KAZUHIRO HAGINOYA

Department of Pediatric Neurology, Takuto Rehabilitation Center for Children, FFPH, Sendai, Japan

Night Seminar 1

The current status, challenges and future perspective of "The Japan Compensation System for Cerebral Palsy"

SHIGERU UEDA

Japan Council for Quality Health Care, Tokyo, Japan

The present status, issues and wishes on the Compensation System for Birth Troubles from a standpoint of making a certificate

KIYOSHI MATSUI

Department of General Pediatrics and Neonatology, Kanagawa Children's Medical Center, Yokohama, Japan

The present status and issues of the Compensation System for Birth Troubles from the diagnostic practice

SATORU KINOSHITA

Department of Pediatrics, National Hospital Organoization Niigata Hospital, Kashiwazaki, Niigata, Japan

The present status, issues and recommendation on Compensation System for Birth Troubles from the Judging Committee

AKIRA OKA

Deaprtment of Pediatrics, Kyorin University, Tokyo, Japan

The present status and issues of the Compensation System for Birth Troubles from an obstetrician

TSUGIO MAEDA

Maeda Obstetrics and Gynecology Clinic, Yaizu, Shizuoka, Japan

Night Seminar 2

Botulinum toxin in the management of upper extremities in childhood cerebral palsy

ATSUO NEZU

Clinic of Child Neurology, Yokohama Residential Care and Medical Centre for Developmentally Disabled Person, Yokohama, Japan

The experience of botulinum toxin therapy for mild lower extremity spasticity in young children with cerebral palsy

MASAO ADACHI

Department of Pediatrics, Kakogawa Municipal Hospital, Kakogawa, Japan

Efficacy of BonT injection to neck, shoulder, and upper limbs of SMID and cerebral palsy

NAOKO YOSHIDA

Department of Pediatric Neurolory, Saint Joseph Hospital for Handicapped People, Kyoyo, Japan

The 5th Child Sleep Group Meeting

A 10 year-old girl with Chiari malformation type 1 who complain snoring and apnea

KUMI KATO

Ota Memorial Sleep Center, Kanagawa, Japan

Sleep disorder from 'cyber addiction'; five school children

KYOKO HOSHINO

Minami Wakayama Medical Center, Japan

Changes of the circadian rhythm in core body temperature of adolescents with Delayed Sleep Phase Syndrome

SHIGEYUKI MATSUZAWA

Hyogo Prefectural Rehabilitation Center Hospital, Kobe, Japan

Effects of Ramelteon (Rozerem) for children with sleep disorders

SHIHOKO KIMURA-OHBA

Department of Pediatrics, Osaka University Graduate School of Medicine, Osaka, Japan

Factors affecting the nocturnal sleep development among preterm Infants

YOKO ASAKA

Konan Women's University, Japan

Kobe University Graduate School of Health Sciences, Japan

Continuance measurement of child sleep of with heart rate valiability analysis by complex demodulation method

NOBORU OHKI

NoruPro Light Systems, Inc. Tokyo, Japan

Reports on IPSA2010 and WSD

JUN KOHYAMA

Department of Pediatrics, Tokyo Bay Urayasu/Ichikawa Medical Center, Japan

B & D Seminar

How to write a case report and a clinical research paper for publication

KOUSAKU OHNO

Division of Child Neurology, Tottori University Faculty of Medicine, Japan

Writing Articles in Basic Science And in English: Is It Our Business?

TAKAO TAKAHASHI

Department of Pediatrics, Keio University School of Medicine, Japan

How to write your manuscript

MELINDA HULL

Cactus Communications K.K., Japan

2011 JSCN Award for Asia Young Investigator

The role of superior sagittal sinus compression by the occipital bone in neonatal cerebral sinovenous thrombosis

MARILYN A. TAN

Department of Pediatrics and Neurosciences, Section of Pediatric Neurology, Philippine General Hospital, Philippines

Asian Oceanian Child Neurology International Education Program

Patients with riboflavin-responsive multiple acyl-CoA dehydrogenase deficiency in Taiwan

WEN-CHEN LIANG

Departments of Pediatrics, Kaohsiung Medical University Hospital, Kaohsiung, Taiwan

Morning Seminar 1

Complications of Traumatic Brain Injury in Children: Epilepsy and Higher Cortical Dysfunction

MANA KURIHARA

Department of Pediatrics, The Kanagawa Rehabilitation Center, Japan

Hypopituitarism after traumatic brain injury

YOSHIKAZU NISHI

Department of Pediatrics, Hiroshima Red Cross Hospital, Japan

Morning Seminar 2

SHUJI WAKAI

Nakanoshima Clinic, Sapporo, Japan

TAKAMICHI YAMAMOTO

Seirei Hamamatsu General Hospital, Epilepsy Center, Shizuoka, Japan

Morning Seminar 3

YUICHI GOTO

Department of Mental Retardation and Birth Defect Research, National Institute of Neuroscience, National Center of Neurology and Psychiatry, Tokyo, Japan

YUKO SAWADA

Agilent Technologies Japan, Ltd.

Morning Seminar 4

Therapy in nocturnal enuresis according to patient's condition -DDAVP and Alarm therapy-

EIICHI HOASHI

Hoashi Clinic, Tokyo, Japan

Morning Seminar 5

L-carnitine therapy in pediatrics: Up-to-date

KIYOSHI MATSUI

Department of General Pediatrics and Neonatology, Kanagawa Children's Medical Center, Yokohama, Japan

Luncheon Seminar 1

Mental disorders related to developmental disorders

JUNZO IIDA

Faculty of Nursing, Nara Medical University, Japan

Luncheon Seminar 2

Epilepsy Management In Children - What is the evidence for what we do?-

JAMES W. WHELESS

Department of Pediatric Neurology, The University of Tennessee, USA

Luncheon Seminar 3

Antiepileptic Drug Therapy for Idiopathic Generalized Epilepsy Considering Lifestyle of Patients

MASAKI TANAKA

Tanakama Clinic, Yokohama, Japan

Practical management for children and adolescences with idiopathic generalized epilepsy from the viewpoint of long-term outcome

SHINICHIRO HAMANO

Division of Neurology, Saitama Children's Medical Center, Saitama, Japan

Luncheon Seminar 4

Can new antiepileptic drugs change epilepsy treatment?

KATSUMI IMAI

National Epilepsy Center, Shizuoka Institute for Epilepsy and Neurological Diorders, Shizuoka, Japan

Luncheon Seminar 5

Strategy of treatment for Wilson disease with neurologic symptoms

NORIKAZU SHIMIZU

Department of Pediatrics, Toho University School of Medicine, Ohashi Medical Center, Tokyo, Japan

Luncheon Seminar 6

A New Treatment Modality for Spasticity in Children with Cerebral Palsy

RYUJI KAJI

Department of Neurology, Tokushima University, Tokushima, Japan

Luncheon Seminar 7

Seeking better treatment of epilepsy: from the viewpoint of life stages and QOL of women

MAKIKO OSAWA

Department of Pediatrics, Tokyo Women's Medical University, Tokyo, Japan

Luncheon Seminar 8

Comprehensive support to AD/HD: Role of Medication

TAKASHI HAYASHI

Faculty of Nursing and Human Nutrition, Yamaguchi Prefectural University, Yamaguchi, Japan

Luncheon Seminar 9

Treatment of spasticity in children with cerebral palsy

NOBUAKI IWASAKI

Department of Pediatrics, Ibaraki Prefectural University of Health Science, Inashiki, Ibaraki, Japan

Neurosurgical management for intractable spasticity in children

SATOSHI IHARA

Department of Neurosurgery, University of Tsukuba, Ibaraki, Japan

Luncheon Seminar 10

Diagnosis and treatment for Gaucher disease and Pompe disease

HIROYUKI IDA

Department of Pediatrics, Jikei University School of Medicine, Japan

Luncheon Seminar 11

Pediatric Nutrition Management - Future Developments

YOSHIAKI TANAKA

Department of Pediatric Surgery, Kurume University School of Medicine, Japan

Luncheon Seminar 12

Management of SGA infants and children with GH and Nutrition therapy for growth and development

SHINOBU IDA

Osaka Medical Center and Research Institute for Maternal and Child Health, Osaka, Japan

Evening Seminar

Nutritional care for child neurology outpatients

DAISUKE TANAKA

Department of Pediatrics, Showa University Toyosu Hospital, Japan

Practical Education Seminar 1

MR imaging of normal and abnormal pediatric brain based on the brain development; Diagnostic clue

HIROSHI OBA

Department of Radiology, Teikyo University Hospital, Tokyo, Japan

Introduction of diffusion-weighted images and MR spectroscopy in pediatric neurology

JUN-ICHI TAKANASHI

Department of Pediatrics, Kameda Medical Center, Kamogawa, Japan

Neonatal Imaging

NORIKO AIDA

Department of Radiology, Kanagawa Childrens' Medical Center, Japan

Diagnostic imaging of brain malformations

MITSUHIRO KATO

Department of Pediatrics, Yamagata University Faculty of Medicine, Yamagata, Japan

Diagnostic imaging of congenital metabolic disorders

HARUSHI MORI

Department of Radiology, Graduate School and Faculty of Medicine, The University of Tokyo, Tokyo, Japan

Practical Education Seminar 2

EEG in the normal term infants

AKIHISA OKUMURA

Department of Pediatrics, Juntendo University School of Medicine, Japan

EEG in the normal preterm infants

TORU KATO

Department of Child Neurology, Okazaki City Hospital, Japan

Abnormal background EEG findings and ictal changes

AKIHISA OKUMURA

Department of Pediatrics, Juntendo University School of Medicine, Japan

Educational Seminar

Brain Death from A Legal Perspective

EIJI MARUYAMA

Kobe University School of Law, Japan

Japanese Criteria of Brain Death in Children

YASUKO KUSAKA

Department of Neurosurgery, The Jikei University Scool of Medicine, Tokyo, Japan

Intrahospital system construction and preparation for brain death organ transplant

MASAYA KUBOTA

Division of Neurology, NCCHD, Tokyo, Japan

Exclusion of Abused and/or Neglected Children from Donors in Brain Death

FUJIKO YAMADA

Sangenkai Yamada Clinic, Isehara-city, Kanagawa, Japan

Child Maltreatment Prevention Network, Isehara-city, Kanagawa, Japan

Japanese Medical Society on Child Abuse and Neglect, Isehara-city, Kanagawa, Japan

Neurological examination and apnea testing

IKUYA UETA

Division of Pediatric Critical Care, Shizuoka Children's Hospital, Shizuoka, Japan

Vestibular reflex in judgement of brain death

SHINICHI IWASAKI

Department of Otolaryngology, Faculty of Medicine, University of Tokyo, Tokyo, Japan

Electroencephalography

AKIHISA OKUMURA

Department of Pediatrics, Juntendo University School of Medicine, Tokyo, Japan

Public Forum

-First Section-

People inheriting the Gift of Life -Client-Centered-

YOSHIKO SHIOTANI

President of Nagasaki International University, Former Governor of Kumamoto Prefecture, Japan

-Second Section-

Raising a child with an autistic spectrum disorder. A message from a parent.

MASAKO SUZUKI

Itabashi Association of Parents of Children with Developmental Disabilities, Japan

Good sleep and good life (Gussuri-raifu)

MASAKO OKAWA

Department of Sleep Medicine, Shiga University of Medical Science, Otsu, Japan

The Periodontal - Systemic Disease Connection

TOSHIHIDE NOGUCHI

Aichi-gakuin University School of Dentistry Department of Periodontology, Nagoya, Japan

Educational Support in School

TOSHIHIDE KOIKE

Tokyo Gakugei University, Tokyo, Japan

The Lifestyle of Individuals with Pervasive Developmental Disorders: Live in Obscurity

HIROSHI IHARA

Department of Psychiatry, Dokkyo Medical University Koshigaya Hospital, Saitama, Japan

English Session 1

O-001 Autosomal dominant nocturnal frontal lobe epilepsy and founder effect

SU KYEONG HWANG

Department of Pediatrics, School of Medicine, Fukuoka University, Fukuoka, Japan

O-002 The analysis of surgically treated pediatric patients with epilepsy

AYATAKA FUJIMOTO

Seirei Hamamatsu General Hospital, Comprehensive Epilepsy Center, Hamamatsu, Japan

O-003 Malignant epilepsies with intractable epileptic seizures and mental retardation

MIHO FUKUDA

St. Marianna University School of Medicine, Kanagawa, Japan

O-004 Eating Behavior in Japanese and Indonesian Autistic Children

HANDAYANI MAULINA

Department of Community Health Sciences, Graduate School of Health Sciences, Kobe, Japan Nursing Study Programme, Faculty of Medicine and Health Sciences, State Islamic University Syarif Hidayatullah Jakarta, Jakarta, Indonesia

O-005 Vagus nerve stimulation (VNS) for intractable epilepsy in the pediatric population: NYU experience

TAKAMICHI YAMAMOTO

Comprehensive Epilepsy Center, Seirei Hamamatsu General Hospital, Hamamatsu, Japan

English Session 2

O-006 A case of paramyotonia congenita with a mutation in SCN4A

DIAN K. PRAMUDYA

Department of Community Medicine and Social Healthcare Science, Kobe University Graduate School of Medicine, Kobe, Japan

O-007 A SOX10 binding site mutation in GJC2 promoter causes Pelizaeus-Merzbacher-like disease

HITOSHI OSAKA

Devision of Neurology, Kanagawa Childrens Medical Centor, Yokohama, Japan

O-008 RT-multiplex PCR for detection of 10 viruses causing acute encephalopathy/encephalitis in Asia

NGAN PHAM

Department of Developmental Medical Sciences, Institute of International Health, The University of Tokyo, Tokyo, Japan

O-009 Chemical Chaperon therapy for beta-galactosidase deficiency: NOEV effect on the 88 missense mutations

EIJI NANBA

Division of Functional Genomics, Research Center for Bioscience and Technology, Tottori University, Japan

O-010 Tissue specificity of mitochondrial respiratory chain disorders with neurological symptoms

AKIRA OHTAKE

Department of Pediatrics, Saitama Medical University, Saitama, Japan

Developmental Disorder (Screening)

O-011 Psychometric Properties of the Japanese version of the Motor Observation Questionnaire for Teachers

AKIO NAKAI

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O-012 Characteristics of exploratory eye movements in patients with Asperger syndrome

TAKASHI OHYA

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O-013 Mirror neuron system in autism spectrum disorders: a near-infrared spectroscopic study

KENJI MORI

Department of Pediatrics, Institute of Health Bioscience, The University of Tokushima Graduate School, Tokushima, Japan

O-014 Functional evaluation of GABAergic neurons in autism spectrum disorders: 123 I-iomazenil SPECT study

TATSUO MORI

Department of Pediatrics, Institute of Health Bioscience, The University of Tokushima Graduate School, Tokushima, Japan

O-015 The investigation of plasma ghrelin concentration in Rett syndrome patients with MECP2 mutations

MUNETSUGU HARA

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Developmental, Psychiatric & Behavioral Disorder

O-016 Validity of impairment rating scale as a tool for functional impairment of children with ADHD

YUSHIRO YAMASHITA

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O-017 Clinical Study of administration of methylphenidate for abused children with ADHD

JUNICHI FURUSHO

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Matsudo Clinic, Chiba, Japan

O-018 The sensory integration programs improved self-efficacy and communication skills in ADHD

TAKASHI HAYASHI

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O-019 Clinical Study in childhood with eating disorder: outcome of amenorrhea

RYOICHI SAKUTA

Center for Child Development and Psychosomatic Medicine, Dokkyo Medical University, Koshigaya Hospital, Japan

O-020 Predictors of pirydoxine responsiveness in chidren with pervasive developmental disorders

SHINICHI KURIYAMA

Department of Molecular Epidemiology, Tohoku University Graduate School of Medicine, Sendai, Japan

Genetics 1

O-021 SETBP1 deletion is responsible for developmental delay distinct from Schinzel-Giedion syndrome

TOSHIYUKI YAMAMOTO

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O-022 Genetic analysis of Myotubular myopathy

EIJI UCHIDA

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School of Human Studies, Taisho University, Tokyo, Japan

O-023 Simultaneous analysis of mitochondrial DNA mutations using suspension array technology

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Nishigaki Clinic & Research Laboratory, Nagoya, Japan

Department of Genomics for Longevity and Health, Tokyo Metropolitan Institute of Gerontology, Tokyo, Japan

O-024 A novel mutation of ATP6V0A2 gene in a Japanese girl with autosomal recessive cutis laxa type 2

YOSHITO ISHIZAKI

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O-025 Autosomal dominant Aicardi-Goutieres syndrome with mild clinical phenotype

TOMONARI AWAYA

Department of Pediatrics, Kyoto University Hospital, Kyoto, Japan

O-026 The proliferative angiomatosis and genetic background in patients with tuberous sclerosis complex

HIROSHI YAMADA

Department of Pediatrics and Child Neurology, Oita University Faculty of Medicine, Oita, Japan

Genetics 2

O-027 The study for the diagnosis and the treatment of fragile X syndrome in Japan

KAORI ADACHI

Division of Child Neurology, School of Medicine, Tottori University, Yonago, Japan

Division of Functional Genomics, Center for Bioscience and Technology, Tottori University, Yonago, Japan

O-028 A new diagnostic method for Prader-Willi syndrome based on ICON probe-based DNA methylation assay

TAKEO KUBOTA

Department of Epigenetic Medicine, University of Yamanashi, Yamanashi, Japan

O-029 A proposal for diagnostic criteria and classification of congenital cerebral hypomyelination

KEN INOUE

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O-030 Estabalishment of iPS cells from patients with Pelizaeus-Merzbacher disease

KEIKO SHIMOJIMA

Tokyo Women's Medical University Institute for Integrated Medical Sciences, Tokyo, Japan

O-031 A new microdeletion syndrome at 5q31 demonstrates neonatal hypotonia and severe mental retardation

SHINJI SAITOH

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Learning Disabilities 1

O-032 Predective Factors of the Reading and Writing Abilities in Elementary School Children

KAORU HANAFUSA

Graduate School of Medicine, Denistry and Pharmaceutical Science Okayama University, Okayama-shi, Japan

O-033 Development of Visual Symptom Survey in Children

TOMOHITO OKUMURA

Osaka Medical College, LD Center, Osaka, Japan

O-034 Significance of working memory in reading ability in PDD and AD/HD

MAKIO OKA

Department of Child Neurology, Okayama University Graduate School of Medicine, Dentistry and Pharmaceutical Science, Okayama, Japan

O-035 2007-2009 year longitudinal study for the RTI model for early detection of Dyslexia

TATSUYA KOEDA

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O-036 Effects of phonological awareness and rapid naming on writing in Japanese children

EIJI WAKAMIYA

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O-037 Reading ability of junior high school students in relation to self-evaluation and depression

TOSHIYA YAMASHITA

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Learning Disabilities 2

O-038 The effect of oral reading training with personal computer for developmental dyslexia

TAKASHI TAKESHITA

Osaka Medical College LD Center, Osaka, Japan

O-039 A Training Method of Writing Kanji for a Child with Visual Information Processing Impairment(2)

MEKUMI MIZUTA

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O-040 2008-2010 year longitudinal studies for the RTI model for early intervention of Dyslexia

TATSUYA KOEDA

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O-041 Deficit of Lexical Route in Japanese Developmental Dyslexia

TOMOHITO OKUMURA

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O-042 Factors and antidote of increase in developmental disorders from the viewpoint of physicians

ASAYO ISHIZAKI

Oji Clinic; Division of Medicine The Association of Remedial Teaching for People with Developmental Handicaps, Japan

Neonates

O-043 Fetal development of the oculomotor nuclear complex in man. II. Volumetric analysis

KATSUYUKI YAMAGUCHI

Department of Pathology, Dokkyo University School of Medicine, Tochigi, Japan Department of Pediatrics, Southern Tohoku General Hospital, Koriyama, Japan

O-044 Preterm babies born from mothers diagnosed as abruptio placenta have good neurological prognosis

SAYAKA HARADA

Department of Neonatology, Osaka City General Hospital, Osaka, Japan

O-045 The examination of the factor that influence early neonatal electroencephalogram

YOSHINOBU OYAZATO

Kakogawa Municipal Hospital, Japan

O-046 A Nationwide Survey on Therapeutic Hypothermia for Neonatal Hypoxic-Ischemic Encephalopathy in Japan

TOSHIKI TAKENOUCHI

Department of Pedaitrics, Keio University School of Medicine, Japan

O-047 A study as to school age of children with premature birth

HIDEO NAGAO

Division of Medicine for Special Support, Faculty of Education, Ehime University, Matsuyama, Japan

O-048 Hypothermia for neonatal asphyxia can not prevent brainstem dysfunction

YUKIHIRO KITAI

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Epilepsy & Seizure 1

O-049 Prediction and prevention of early relapse after ACTH therapy in West syndrome

YUMIKO ISHIZAKI

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O-050 Effect of topiramate in pediatric patients with epileptic spasms

FUMIKA ENDOH

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O-051 Absence seizure of Dravet syndrome

YUKO TSUDA

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O-052 EEG of those who were sufferd from seizure or loss of conscious without fever

TAKESHI INOUE

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O-053 Contribution of excess kurtosis of interictal EEG to seizure outcome after epilepsy surgery

TOMOYUKI AKIYAMA

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O-054 Diffusion tensor imaging in cryptogenic West syndrome: TBSS analysis

JUN NATSUME

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Epilepsy & Seizure 2

O-055 MRCP and CNV analysis in patients with PKC

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O-056 Gamma-oscillations modulated by eye movements

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O-057 Scalp-recorded high-frequency oscillations in benign partial epilepsy of childhood

KATSUHIRO KOBAYASHI

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O-058 Language lateralization in children with epilepsy using near infrared optical topography

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O-059 Pre and post hemispherotomy brain blood flow reaction using NIRS in patients with hemimegalencephaly

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O-060 Time dependent change of cognitive function at the epilepsy onset

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Epilepsy & Seizure 3

O-061 Longitudinal examination of the effect of topiramate on the weight and body composition

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O-062 Keisikasyakuyaku-to suppress hyperthermia-induced seizures via the control of intracellular Ca

HITOMI HINO

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O-063 Pharmacotherapy of chronic non-idiopathic partial epilepsies based on semiology: a prospective study

KENJI SUGAI

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O-064 Efficacy of lamotrigine for intaractable epilepsy with tuberous sclerosis

HISASHI KAWAWAKI

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O-065 A multicenter study to assess safety, efficacy and pharmacokinetics of fosphenytoin in pediatrics

EIJI NAKAGAWA

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Epilepsy & Seizure 4

O-066 Spectrum of fever sensitive epilepsy syndrome from investigation into mild cases of Dravet syndrome

MASAKO SAKAUCHI

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O-067 Diagnostic clues for early diagnosis of epileptic encephalopathy with CDKL-5 mutation/deletion

KATSUMI IMAI

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O-068 PCDH19 mutation in Japanese females with epilepsy

NORIMICHI HIGURASHI

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O-069 Quantitative analysis of antibodies to NMDA-type glutamate receptor in Rasmussen syndrome

TETSUHIRO FUKUYAMA

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O-070 Cerebrospinal Fluid Levels of Amino Acids and Ammonia in Children with Status Epilepticus

HIROKI HASUMI

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O-071 Increased level of serum IL-1 RA subsequent to resolution of clinical symptoms in West syndrome

GAKU YAMANAKA

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Developmental Disorders 1

O-072 A retrospective study of medical therapy in children with attention deficit hyperactivity disorder

RYOKO OTANI

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Center for Child Development and Psychosomatic Medicine, Dokkyo Medical University, Koshigaya Hospital, Japan

O-073 Clinical Assessment of the effect of Atomoxetine in 32 Patients with AD/HD or PDD

SHINICHIRO TAKIGUCHI

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O-074 Consideration of using properly choice MPH or ATX of 80 cases with ADHD and PDD with ADHDsympton

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O-075 Behavioral value between mothers and teachers after the medication of atomoxetine in ADHD with ASDs

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Department of Pediatrics, Tsukuba Municipal Hospital, Japan

O-076 Effect of tracheostomy and imipramine on respiratory dysrhythmia in Rett syndrome

KAORI SASAKI

Shinko Kakogawa Hospital, Hyogo, Japan

O-077 Treatment of selective bronchial air injection for atelectasis

SOICHIRO TANAKA

Takuto Rehabilitation Center, Sendai, Miyagi, Japan

Developmental Disorders (Treatment & Support)

O-078 The trial approach of the course of children's care support for mothers

KEIKO MAEDA

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O-079 5 years checkup giving priority to support

KASUMI NAGASAWA

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O-080 Maternal role consciousness scales and Parent Training

HIROYUKI YOKOYAMA

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O-081 A study on the utilization of support books at admission into the primary school

HIROMI TSUSHIMA

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O-082 The evaluation for increase of development disorders - when, where, and what do they decide to go? -

KAKUROU AOYAGI

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O-083 Motor and intellectual development of the patient with maternal uniparental disomy of chromosome 14

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Development

O-084 Follow-up study at 3 years-old on very low birth weight infants at our center

EIKO TAKADA

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O-085 The evaluation and support of the development of the 6 years old of VLBW babies

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O-086 Dynamic ground force in human crawling

ARITO YOZU

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O-087 Eating and swallowing rehabilitation for a patient with Prader-Willi syndrome

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O-088 Developmental change of visual behavior in infancy

KONISHI YUKIHIKO

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O-089 The Developmental Change of Infant's Attention by Mother-Infant Interaction -An Eye Tracking Study-

AKIO NAKAI

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Peripheral Nerve

O-090 Analysis of peripheral nerve abnormalities in pediatric patients with spinal muscular atrophy

TAKAHIRO YONEKAWA

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O-091 An infantile case suspected of vaccine associated paralytic poliomyelitis

KOTOKO SUGAYA

Saitama Children's Medical Center, Saitama, Japan

O-092 A case of GBS with enhanced Caude equina in subacte phase MRI without any abnormallity in acte phase

SAYAKA HAMAGUCHI

Tokyo Metoropolitan Hiroo Hospital, Japan

O-093 Efficacy of cyclosporin therapy in pediatric CIDP patient

KEIKO YANAGIHARA

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O-094 Successful treatment with steroid therapy for benign recurrent sixth nerve palsy

KEITARO YAMADA

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Muscular Disease 1

O-095 Diagnostic approach of metabolic myopathies using muscle metabolome profiling

TOKIKO FUKUDA

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O-096 Desirable disease explanations for patients with muscular dystrophy and their parents

OSAMU IMURA

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O-097 Survey about child neurologist trends in informing DMD boys about their disease

HARUO FUJINO

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O-098 The marker as a treatment index of muscle disease using actigraph system

SHIGEMI KIMURA

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O-099 Dysphagia in children with spinal muscular atrophy

NOZOMI SANO

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Muscular Disease 2

O-100 WITHDRAWN

O-101 Insomnia in patients with Fukuyama congenital muscular dystrophy

TAKATOSHI SATOU

Department of Pediatrics, Tokyo Wemen's Medical University, Tokyo, Japan

O-102 Fractional shorting can detect early cardiac dysfunction in Duchenne muscular dystrophy

HIROYUKI AWANO

Department of Pediatrics, Kobe University Graduate School of Medicine, Kobe, Hyogo, Japan

O-103 The effect of steroid therapy for scoliosis in patients with Duchenne muscular dystrophy

TERUMI MURAKAMI

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O-104 Review of steroid treatment in Duchenne Muscular Dystrophy

MASAHIDE GOTO

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O-105 Chemical treatment of muscular dystrophy that enhances skipping of the exon in the dystrophin gene.

MARIKO YAGI

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O-106 Extramuscular manifestations in congenital myopathy due to ACTA1 gene mutation

YOSHIAKI SAITO

Department of Child Neurology, National Center of Neurolog and Psychiatry, Japan

Acute Encephalitis & Encephalopathy 1

O-107 An analysis of serum cytokine and CPT2 gene in three cases of severe HHV-6 encephalopathy

HIROSHI MATSUMOTO

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O-108 New Treatment for mitochondrial fatty acid oxidation disorders with bezafibrate

SEIJI YAMAGUCHI

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O-109 Release of protein S-100B in serum and CSF in pediatric patients with acute encephalopathy

REIKO KOICHIHARA

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O-110 Investigation of genetic background in Japanese patients with Acute encephalopathy

MAYU SHINOHARA

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Acute Encephalitis & Encephalopathy 2

O-111 Serum and cerebrospinal fluid levels of HMGB1 in pandemic H1N1 influenza encephalopathy

TAKESHI MATSUSHIGE

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O-112 Clinical aspects of inpatient children for neurological symptoms by pandemic influenza A/H1N1 2009

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O-113 Analysis of cytokine levels in cerebrospinal fluid and serum in rotavirus encephalopathy

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O-114 Immunological assessment of acute encephalitis with refractory, repetitive partial seizures

HIROYUKI WAKAMOTO

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O-115 A research of urinary beta2-microglobulin at early phase of acute encephalopathy in children

HIROYUKI TORISU

Department of Pediatrics, Graduate School of Medicine, Kyushu University, Fukuoka, Japan Early Diagnosis of Virus Associated Encephalopathy Study Group

O-116 Clinical evaluation of acute encephalopathy with biphasic seizures and late reduced diffusion (AESD)

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O-118 Anti-N-methyl-D-aspartate-receptor encephalitis in a three-year-old boy.

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O-119 Study of differential diagnosis between Anti-NMDA receptor encephalitis and AERRPS

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O-120 A girl case of anti-NMDA receptor encephalitis who is doing rapid recovery.

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O-121 The side effect of antiepileptic drug for postencephalitis (second report:sleepiness)

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O-123 Management for pediatiric febrile refractory status epilepticus affects occurence of AEFCSE

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O-124 Findings of electroencephalography monitoring in AESD spectrum

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O-125 Analysis of 43 cases of influenza-associated encephalopathy

MASAYO HANABUSA

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MOMOKO OKA

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O-127 Four cases of AESD with mildly abnormal MRI

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O-128 Deep Gray Matter Lesions in Acute Encephalopathy with Biphasic Seizures and Late Reduced Diffusion

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O-129 The study of 17 cases with delirium in influenza about Imaging and electroencephalographic findings

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O-130 Pediatric Death due to 2009 Pandemic Flu Associated Encephalopathy

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O-131 Correlation between central sparing and clinical course in acute encephalopathy

TATSUYA FUKASAWA

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O-132 Study of acute encephalopathy without abnormality of magnetic resonance imaging

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O-133 Severity of type A acute encephalopathy in the ABC-classification

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O-134 Early detection of autism in ethnic minorities at the routine infant health checkups

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O-135 High Functioning Pervasive Developmental Disorders with Motor Developmental Delay

KEIKO KOTERAZAWA

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O-136 Cognitive functions in patients with 22q11.2 deletion syndrome -comparison with Williams syndrome-

MIHO NAKAMURA

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O-137 Study about non-social problems for adult Asperger's syndrome

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O-138 Melatonin treatment of sleep-wake cycle disorders in children with brain damage

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O-139 Clinical Study of early methylphenidate treatment response cases for children with AD/HD

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O-140 Effects of atomoxetine in patients with pervasive developmental disorder associated with AD/HD

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O-141 Usefulness of administration of methylphenidate for AD/HD patients with epilepsy

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O-142 A study of psychiatric symptoms and treatment of pervasive developmental disorder

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O-143 Advanced parental age and behavior in offspring of autism spectrum disorder

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O-144 Behavioral Problems of Pervasive Developmental Disorder using SDQ

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Involuntary Movement & Basal Ganglia

O-145 A family of Paroxysmal kinesigenic dyskinesia

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O-146 Clinical Characteristic of Pediatric Restless Legs Syndrome (RLS)

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O-147 Low dosis l-Dopa therapy for Gilles de la Tourette syndrome

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O-148 Case report - Paroxysmal tremulous movement followed by postural impairment in Angelman syndrome

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O-149 The experience of intrathecal baclofen therapy to two cases of intractable dystonia

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O-150 Clinical features of 12 cases with Hyperekplexia having glycinergic neurotransmissive gene mutations

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O-151 Epileptic characterization and their basic pathogenesis in 4 infants with cerebrovascular thrombosis

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O-152 Surgical indication in epilepsy with porencephaly due to perinatal middle cerebral artery infarction

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O-153 Efficacy and tolerability of lamotrigine in children with epilepsy

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O-154 An effect of lamotrigine for various epileptic seizure types

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O-155 Clinical Investigation of Lamotrigine for Patients with Intractable Childhood Onset Epilepsy

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O-156 Efficacy of lamotrigine and ACTH combined therapy against West syndrome

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O-157 The evaluation of oxidative stress during ACTH therapy in West syndrome

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O-159 A case of neurofibromatosis type 1 with glaucoma in infancy

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O-160 Crouzon syndrome, Apert syndrome, Pfeiffer syndrome

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O-161 In utero exposure to dioxin causes neocortical dysgenesis through the actions of p27^{Kip1}

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HIDENORI SUGANO

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O-164 Two cases of intramedullary spinal astrocytoma manifesting mild symptoms at the first examination

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O-165 A case of syringomyelia presenting as acute progressive gait disturbance

YUJI HASHIMOTO

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O-166 Treatment strategy for renal angiomyolipoma in patients with Tuberous Sclerosis

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O-167 A short-term effect of the levetiracetam use example for the intractable epilepsy

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Cerebral Palsy & Care

O-168 Organic diseases and epilepsies in patients with extreme severe motor and intellectual disabilities

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O-169 Long term prognosis of SMID patients required endotracheal intubation

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O-170 Investigation of gastroesophageal reflux in cases of sevearly disabled children

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O-171 Energy expenditure by indirect calorimetry in children with severe motor and intellectual disability

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O-172 The Factors to Acquire to Walk for Motor Handicapped Children from the Viewpoint of Foot Stamp

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O-173 Clinical characteristics of West syndrome with choreic movement

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O-174 Childhood absence epilepsy associated with diurnal enuresis

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O-175 High coexistence of neurally mediated syncope in patients with juvenile myoclonic epilepsy

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O-176 The investigation for the role of serum prolactin in pseudoseizure

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O-177 Treatment of Status Epilepticus

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O-178 Clinical characteristics of fourteen infants who had got West syndrome after neonatal asphyxia

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O-179 Spontaneous remission of West syndrome after RS virus infection

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O-180 Discontinuation of vitamin B6 and antiepileptic drugs after seizure control in West syndrome

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O-181 A good response to antiepileptic drugs for West syndrome with preterm birth

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O-182 ineffective cases of West syndrome with tuberous sclerosis for vigabatrin therapy

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O-183 efficacy and cerebrospinal fluid finding of West syndrome by intravenous immunoglobulin therapy

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O-184 Assessment of thalamic blood flow in intractable epilepsy with callosotomy using SPECT

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O-185 Characteristics of epilepsy in focal cortical dysplasia in childhood

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O-186 Epidemiology of status epilepticus in Tottori prefecture

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O-187 Cognitive outocome of 62 epileptic patients with forcal cortical dysplasia

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O-188 Effect of antiepileptic drugs to abnormal movements without EEG findings

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O-189 Analysis of the Intelligence with Panayiotopoulos syndrome(PS) using the Wechsler Intelligence Scale

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Infection & Autoimmune Disease

O-190 Clinical course toward definite diagnosis and brain images of congenital cytomegalovirus infection

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O-191 Clinical feature and characteristic head MRI findings of congenital cytomegalovirus infection

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O-192 Comparison between symptomatic and asymptomatic cases with congenital cytomegalovirus infection

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O-193 Clinical characteristic of pediatric multiple sclerosis in Japan

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O-194 TNF alpha may induce marked polymorphonuclear cell predominance in the CSF with bacterial meningitis

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O-195 Preliminary analysis of anomalies in pyramidal tract of the brainstem

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O-196 Association between callosal injury assessed by tractography and neurodevelopmental outcome

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O-197 Hippocampal abnormality on diffusion-weighted images in prolonged or clusterd febrile seizures

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O-198 Acute encephalopathy with reduced diffusion in diffuse subcortical white matter at acute phase

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O-199 Differences in the time course of splenial and white matter lesions in MERS

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O-200 Age specific benzodiazepine receptor ditstribution in children by using 123I iomazenil SPECT

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O-201 Diagnosis and surgical treatment of severe stenosis of foramen magnum associated with achondroplasia

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Autonomic Nerves, Headache & Vascular Disease

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O-203 Profiles of blood biomarkers in alternating hemiplegia of childhood

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O-204 Highschool students suffering from SIH cannnot attend the entrance examinations

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O-205 Identification of susceptibility gene for Moyamoya disease

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O-206 Basic pathogenesis of proliferative vascular anomalies in 9 patients with Klippel-Trenaunay syndrome

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O-207 Analysis of a new model mouse of hypoxic ischemic brain injury for extremely premature infants

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O-208 A case of Adrenoleukodystrophy with pyramidal tract lesion

AKIHIKO MIYAUCHI

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O-209 Epidemiologic study of Congenital dysmyelinating disorders of central nervous systems

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O-210 Ten cases with early onset Dentatorubral-Pallidoluysian Atrophy of childhood

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O-211 Nation-wide survey of long-term clinical corse and management in patients with Cockayne syndrome

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O-212 A long-term clinical course of childhood cerebral adrenoleukodystrophy

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O-213 Lesions of monoaminergic neuron system in patients with xeroderma pigmentosum group A

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Metabolic Disease 2

O-214 Chemical chaperone therapy for Gaucher disease type3

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O-215 Clinical and genetic study of 42 Japanese patients with Type2 Gaucher disease

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O-216 Fabry database: study on molecular pathology and prediction of clinical phenotype of Fabry disease

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O-217 A baby with mucopolysaccharidosis type II found by newborn screening for lysosomal storage diseases

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O-218 The Effiacacy on the Brain of Enzyme Replacement Therapy for Mucopolysaccharidosis II

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O-219 Nationwide survey of Alexander disease in Japan

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Metabolic Disease 3

O-220 Study of the diagnosis for atypical cases with Wilson disease

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O-221 A case of neurological Wilson disease diagnosed by molecular genetic analysis

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O-222 Present report of mitochondrial respiratory chain disorders in Japan

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O-223 High dose riboflavin therapy for a case of L-2-hydroxyglutaric aciduria

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O-224 Transiently high serum lactic acid in patients with monocarboxylate transporter 8

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O-225 A case of CPT 2 deficiency with acute encephalopathy related to HHV-6 infection

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P-001 Haplotype-phenotype correlation and survival analysis in Fukuyama congenital muscular dystrophy

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P-002 First trimester chorionic villi have a myogenic potential for muscular dystrophy therapy

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P-003 Survey about child neurologist trends in informing boys with DMD of their disease

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P-004 Genetic Testing of spinal muscular atrophy and SMARD1

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P-005 A case of spinal muscular atrophy 3 with a missense mutation of SMA1 with rapid motor deterioration

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P-006 Spine surgery for neuromuscular scoliosis in Japan

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P-007 The 11 Cases of Myasthenia Gravis

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P-008 Three cases of pneumonia and atelectasis treated by intrapulmonary percussive ventilation

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Muscular Disease 2

P-009 Childhood onset of dysferlinopathy with calf hypertrophy

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P-010 CAV gene mutation with rippling muscle disease

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P-011 X-Linked Myotubular Myopathy; a case report with fatal hepatic hemorrhage.

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P-012 A case of mitochondrial disease with hyperlactosis in infancy

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P-013 Reconsideration of juvenile polymyositis

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P-014 Five cases of autoimmune myopathies

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P-015 Hypo-globulinemia in patients with congenital myotonic dystrophy

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P-016 Paramyotonia congenita in a Japanese boy.

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P-017 An effect and a side effect of Methylphenidate and Atomoxetine for ADHD

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P-020 Fourteen cases of ADHD in our hospital

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P-021 The effect of medication on continuous performance test in children with AD/HD

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P-022 A 20 year-old women with ADHD who complain various sleep problem for stimulants.

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P-023 Effectiveness of aripiprazole in a clinic - Clinical experience of 70 patients -

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P-024 The effect of Bio-Active Therapy to autism

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P-025 Investigation of medication in children with autism and other pervasive developmental disorders

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Developmental Disorders 2

P-026 Clinical research of "Attention-Deficit Hyperactivity" in the infancy

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P-027 Brief checklist of developmental disorders at the age from 4 to 6 years for general pediatricians

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P-028 The child why other diseases were discovered with the main complaint of the developmental disability

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P-029 Clinical studies of head circumference and higher brain function in children with PDD

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P-030 WITHDRAWN

P-031 Eye movement of reading in developmental dyslexia

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P-032 Clinical findings in patients diagnosed as developmental disorders after their health check-ups

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P-033 Asperger's disorder of female

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P-034 Generation and Characterization of the CDKL5 Knockout Mouse

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P-035 Identification and characterization of interacting proteins of CDKL5

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Epilepsy & Seizure 1

P-036 Cytokine and chemokine levels in the cerebrospinal fluid of patients with West syndrome

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P-037 Intravenous injection of phenobarbital for benign convulsions with mild gastroenteritis

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P-038 A case of repeated afebrile convulsion with mild gastroenteritis

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P-039 Convulsive Diseases in Osaka Infant Home

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P-040 A case of febrile myoclonus

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P-041 Establishment of the standardized diagnostic and therapeutic system for breath holding spell

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P-042 A case of a boy who has become restless after the repetitive complex partial seizures with IgG-GluR

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P-043 A case of severe myoclonic epilepsy in infancy (autopsy case)

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P-044 Acute encephalopathy with severe myoclonic epilepsy in infancy.

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P-045 Early-onset absence epilepsy with onset in early infancy

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P-046 A case of symptomatic partial epilepsy presenting epileptic negative myoclonus

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P-047 A case of four year old autistic boy with pattern-sensitive epilepsy

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P-048 Two cases of nocturnal frontal lobe epilepsy diagnosed by video-monitering electroencephalogram

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P-049 Two cases of EEG exacerbation during treatment with CBZ

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P-050 EEG patterns of nonconvulsive seizures in pediatric neuro-intensive care unit

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P-211 Problems of Reading and Writing in Japanese School Children (Part 2): Relationship with the ADHD-RS TOMOKA KOBAYASHI

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P-269 Neural correlates for adaptation to environmental change: An fMRI study

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P-270 Acute encephalopathy presenting with transient hypotonia and reversible reduced diffusion; 3 cases

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P-271 Changes in abnormalities on MR images with a case of acute encephalitis

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P-272 MR spectroscopy in acute encephalopathy with biphasic seizures and late reduced diffusion[AESD]

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P-273 Acute encephalopathy with biphasic seizures and late reduced diffusion on severe cortical dysplasia

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P-274 A case of acute encephalopathy with the features of both ANE and AESD

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P-276 Clinical investigation of acute encephalopathy with prolonged convulsions

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P-277 Five cases of acute encephalopathy with genetic polymorphism

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P-278 Developmental Outcome of Children with AESD

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P-280 Autoimmune-mediated encephalitis manifesting with SLE-like profiles: A case report

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P-283 A case of anti-GluR antibody-positive encephalopathy showing the basal ganglia involvement

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P-287 Difficulty in Immunization for Children with Development Disorder

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P-288 An infant with Acute Flaccid Paralysis

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P-293 A case of infectious pachymeningitis with left blepharoptosis

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P-295 Serum and CSF biomarker in childhood neurological disorders; 4th report

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P-297 A case of Dravet syndrome with acute encephalopathy

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P-298 Genetic seizure susceptibility and a SCN1A mutation in pateints with acute encephalopathy

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P-300 Urinary retention secondary to clinically mild encephalitis with a reversible splenial lesion

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P-301 A case of acute encephalitis with right frontal dysfunction detected by NIRS during recovery phase

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P-302 Three Cases of HHV-6 Encephalopathy Showing Hyperperfusion in Cerebral Blood Flow SPECT

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P-305 Three cases of adults having Pelizaeus-Merzbacher disease with PLP1 gene duplication

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P-306 A case of neonate Alexander disease identified a heterozygous missense mutation, L352R

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P-307 A case of early diagnosed infantile Alexander disease with a novel GFAP gene mutation

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P-308 Adrenoleukodystrophy treated as an ACTH insensitivity syndrome in a pre-pubescent male

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P-309 Management of cerebral salt wasting syndrome using ANP and BNP

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P-310 Characteristics of 157 Japanese patients with Gorlin Syndrome

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P-311 A mother and a child of Gorlin syndrome with Alu-mediated PTCH1 gene

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P-312 Gorlin syndrome with nasal glioma

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P-313 Bone density of patient with Gorlin syndrome

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P-314 A case of neonatal seizure with SCN2A mutation which progress is not benign

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P-315 A boy of Subcortical band heterotopia with DCX mutation

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P-316 Clinical Application of Multiplex ligation-dependent probe amplification (MLPA)

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P-317 Deletions of SCN1A 5'genomic region with promoter activity in Dravet syndrome

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P-318 MECP2 gene analysis of patients with mental retardation

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P-319 TSC mutation analysis of Japanese patients with tuberous sclerosis

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P-320 Comprehensive genetic analysis of overlapping syndromes of RAS/RAF/MEK/ERK pathway

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P-321 A new form of childhood-onset autophagic vacuolar myopathy

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P-322 Two patients with Ullrich congenital muscular dystrophy confirmed by RT-PCR from lymphocyte mRNA

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P-323 Clinical course of severe infantile congenital myopathy

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P-324 Clinical features of congenital myasthenic syndrome in Japan

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P-325 A case report of End-Plate Acetylcholinesterase Deficiency

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P-326 Two patients with childhood-onset Pompe disease presented contrasting effect of ERT

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P-327 The progress of brotherly childhood-onset Pompe disease of Enzyme Replacement Therapy

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P-328 Pulmonary rehabilitation program for the children with neurological or neuromuscular diseases

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P-329 Preliminary report: The different metamorphosis of mitochondria in BBB consisted cells in MELAS

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Peripheral Nerve

P-330 Extremely low birth weight infant with uncertain auditory neuropathy

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P-331 A sibling case of distal Hereditary Motor Neuropathy type VI

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P-332 Case of a two-year old boy suspected of Dejerine Sottas syndrome with PMP22 gene mutation

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P-333 The family of Charcot-Marie-Tooth disease with pyramidal features

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P-334 A female case of MADSAM neuropathy-subtype classification of CIDP and the treatment-

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P-335 A atypical GBS patient presenting as enhancement of cauda equina on MRI resembling mononeuritis

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P-336 Twenty-four Japanese cases with Restless legs syndrome (RLS)

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P-337 Clinical study of five infants that show inconstant abnormal oculoar motor

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P-339 2 cases of Pitt-Hopkins syndrome

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P-340 A missense mutation in a family with CRASH syndrome

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P-341 Delayed progressive hydrocephalus in an infancy with MPPH-CM syndrome

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P-342 A patient with MECP2 duplication syndrome

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P-343 Two cases of 1p36 deletion syndrome with sleep apnea syndrome

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P-344 Intractable epilepsy in children with 2q23.1 microdeletion syndrome

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P-345 Microdeletion of 2q23.1 associated with Angelman-like phenotype

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P-347 Five cases of congenital absence of the portal vein undergoing liver transplantation

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P-348 Effective treatment with NPPV for prolonged respiratory abnormalities in Joubert syndrome

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P-349 Two infants of Septo-Optic dysplasia found with ophthalmologic abnormality

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P-350 The case of Leucodysplasia, microcephaly, cerebral malformation

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P-351 A case of Young-Simpson syndrome

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P-352 Clinical study for screening system of Sotos syndrome in Japan

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P-353 A Clinical feature with trisomy 13 providing under intensive treatment

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P-354 Chronic manifestations and therapeutic interventions for the long-term survivors with trisomy 13

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P-355 Two cases of the RAS/MARK syndrome with refractory epilepsy and severe developmental delay

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