23 a 26 Jul

BH 2025

Belo Horizonte - MG

gerin

14º Encontro Nacional de Distúrbios do Movimento

gerin 2025

Organização

Egberto Reis Barbosa
Clínica Neurológica HC FMUSP

Francisco Cardoso
Professor Titular, Setor de Neurologia,
Departamento de Clínica Médica, UFMG

Hélio Teive
Serviço de Neurologia, Departamento
de Medicina Interna, HC, UFPR

Luiz Augusto F. Andrade
Hospital Israelita Albert Einstein, SP

23 julho

07h30
Registration

08h15 - 09h30
1º ENCONTRO CIENTÍFICO LARGE-PD-BRAZIL

08h15 - 08h20
Introduction
Speaker
Vitor Tumas, SP
08h20 - 08h45
LARGE-PD: the ongoing studies
Speaker
Ignacio Fernandez Mata, USA
08h45 - 09h00
Brazilian network for the validation of alpha-synuclein biomarkers in Parkinson's disease
09h00 - 09h15
Profile of patients with idiopathic Parkinson's disease carrying variants in the GBA and LRRK2 genes in Brazil
Sarah Teixeira Camargos, MG
09h15 - 09h30
Global overview of rare causal variants and common high-risk variants in Parkinson's disease in Brazil
Sarah Teixeira Camargos, MG

09h30 - 10h00
PRESENTATION OF RESEARCH PROJECTS
Speakers
Artur Schumacher Schuh, RS
Bruno Lopes dos Santos Lobato, PA
Pedro Braga Neto, CE

09h30 - 09h45
Validity of the "body-first" and "brain-first" models in the brazilian cohort of the large-pd study
Gabriela Lopes de Morais, SP
09h45 - 10h00
Development of a human cell-based model to explore the association of pesticides with Parkinson's disease
Bruno Lopes dos Santos Lobato, PA

10h00 - 10h30 Coffee Break

10h30 – 12h00
PRESENTATION OF RESEARCH PROJECTS
Speakers
Carlos Roberto de Mello Rieder, RS
Delson José da Silva, GO
Denise Hack Nicaretta, RJ
Fernanda Martins Maia Carvalho, CE
Grace H Letro, SP
Henrique Ballalai Ferraz, SP
Laura Silveira Moriyama, SP
Marcus Vinicius Della Coletta, AM
Rubens Gisbert Cury, SP
Sarah Teixeira Camargos, MG
Vanderci Borges, SP

10h30 - 10h45
Genetic factors associated with neuroinflammation and the risk of developing levodopa-induced dyskinesias in Parkinson's disease
Joyce Yamamoto, SP
10h45 - 11h00
Evaluation of genetic factors and polymorphisms in the development of sarcopenia in patients with Parkinson's disease: association with clinical phenotypes in a Brazilian cohort
Pedro Braga Neto, CE
11h00 - 11h15
Association of female health factors with the severity of Parkinson's disease in Brazilian women: translation and application of the women's Cleveland Clinic Mata questionnaire
Ângela Vieira Pimentel, SP
11h15 - 11h30
Gender differences in the presentation and progression of young-onset Parkinson's disease in Brazil
Vanderci Borges, SP
11h30 - 11h45
Pain in Parkinson's disease
Grace Helena Letro, SP
11h45 - 12h00
Assessment of people's perceptions about being informed of a possible risk of developing Parkinson's disease in the future
Rafael Pallos da Silveira, SP

MOVEMENT DISORDERS AND PARKINSON'S DISEASE SOCIETY (MDS PAS COURSES)

12h00 - 12h45
Registration

12h45 - 13h00
OPENING COMMENTS - GENETIC OF MOVEMENT DISORDERS: FROM BASIC SCIENCE TO CLINICAL PRACTICE
Course Directors
Malco Rossi, ARG
José Luiz Pedroso, SP
Chair - GERIN 2025
Francisco Eduardo Costa Cardoso, MG
President of the Brazilian Academy of Neurology
Delson José da Silva, GO
Scientific Committee
Rubens Gisbert Cury, SP
Sarah Teixeira Camargos, MG
Hélio Afonso Ghizoni Teive, PR
Emilia Gatto, ARG
Carlos Henrique Camargo, PR
Orlando Graziani Povoas Barsottini, SP

13h00 - 14h40
SESSION 1
GENETIC MOVEMENT DISORDERS: FROM BASIC SCIENCE TO CLINICAL PRACTICE
Chairs
Carlos Henrique Camargo, PR
Rubens Gisbert Cury, SP

13h00 - 13h25
What movement disorders specialists should know about genetics?
Marcelo Kauffman, ARG
13h25 - 13h50
What geneticists should know about movement disorders?
Emilia Gatto, ARG
13h50 - 14h15
Genetic movement disorders in the Americas: epidemiology and genetic counseling.
Mario Cornejo-Olivas, PER
14h15 - 14h40
How to develop study groups in genetic of movement disorders. Lessons learned.
Laura Bannach Jardim, RS

14h40 - 15h55
SESSION 2
PHENOMENOLOGY OF GENETIC MOVEMENT DISORDERS
Chairs
Egberto Reis Barbosa, SP
Henrique Ballalai Ferraz, SP

14h40 - 15h05
When a genetic testing should be performed in Parkinson's disease
Ignacio Fernandez Mata, USA
15h05 - 15h30
Main genetic movement disorders in children
Laura Silveira Moriyama, SP
15h30 - 15h55
Genetic movement disorders: lessons from my patients
José Luiz Pedroso, SP

15h55 - 16h30 Coffee Break

16h30 - 18h40
SESSION 3
SPECIFIC GENETIC MOVEMENT DISORDERS
Chairs
Francisco Eduardo Costa Cardoso, MG
Pedro Braga Neto, CE

16h30 - 16h55
Genetic dystonia: guidance for the diagnosis
Sarah Teixeira Camargos, MG
16h55 - 17h20
Approach to the patient with hereditary ataxia
Malco Rossi, ARG
17h20 - 17h45
Huntington disease's phenocopies
Ricardo Maciel, MG

17h45 - 18h40
POSTER SESSION
(list at the end)
Sirius Room, Ground Floor

24 julho

07h00
Registration

07h30 - 08h30
SIMPÓSIO TEVA
CAFÉ COM CIÊNCIA: COMO DECIDIR O MELHOR CAMINHO NO MANEJO DA DOENÇA DE HUNTINGTON?
Palestrante
Gustavo Franklin, PR

08h30 - 10h10
SESSION 4
GENETIC MOVEMENT DISORDERS: MISCELLANEOUS
Chairs
Débora Palma Maia, MG
Delson José da Silva, GO

08h30 - 08h55
A journey through the differential diagnosis of neurodegeneration with brain iron accumulation (NBIA)
Orlando Graziani Povoas Barsottini, SP
08h55 - 09h20
Importance of clinical phenomenology for the interpretation of genetic data
Christos Ganos, CAN
09h20 - 09h45
How to discover new genes in movement disorders?
Stephan Zuchner, USA
09h45 - 10h10
10 genetic syndromes in childhood that movement disorders experts should prompt recognize...
Donald Gilbert, USA

10h10 -10h30 Coffee Break

10h30 - 12h00
SESSION 5
NEW TREATMENTS IN GENETIC MOVEMENT DISORDERS. REALITY AND FAILURES
Chairs
Mauro Cunningham, MG
Monica Santoro Haddad, SP

10h30 - 10h55
New Horizons in the Treatment of Genetic Parkinson's Disease
Ignacio Fernandez Mata, USA
10h55 - 11h20
Why Huntingtin-lowering therapies may need to be reconsidered
Alberto Espay, USA
11h20 - 11h45
Are we close to finding effective therapies for Hereditary ataxias?
Hélio Afonso Ghizoni Teive, PR
11h45 - 12h00
Discussion with Question and Answer

12h00 - 13h00
SIMPÓSIO BIOGEN
DO DIAGNÓSTICO À ESPERANÇA: SKYCLARYS (OMAVELOXOLONA) - O PRIMEIRO MEDICAMENTO PARA ATAXIA DE FRIEDREICH.
Palestrantes
Clécio Godeiro, RN
Marcondes C. França Junior, SP

13h00 - 13h15
ABERTURA

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14º Encontro Nacional de Distúrbios do Movimento

Comissão Organizadora
Egberto Reis Barbosa, SP
Francisco Eduardo Costa Cardoso, MG
Hélio Afonso Ghizoni Teive, PR
Luiz Augusto Franco de Andrade, SP

13h15 - 15h50
Coordenador
Delson José da Silva, GO

13h15 - 13h45
Critical Appraisal of a "Biology-Based" Classification Criteria of PD
Alberto Espay, USA
13h45 - 14h10
Novidades no Manejo Sintomático Motor da DP
Henrique Ballalai Ferraz, SP
14h10 - 14h35
Estimulação Magnética em DP
Rubens Gisbert Cury, SP
14h35 - 15h00
O Que Esperar do Futuro de DBS em DP
Ricardo Maciel, MG
15h00 - 15h25
Perspectivas de Tratamentos Modificadores da DP
Francisco Eduardo Costa Cardoso, MG
15h25 - 15h50
Discussão

15h50 - 16h20 Coffee Break

16h20 - 18h00
Coordenadora
Flávia de Paiva S Rolim, CE

16h20 - 16h45
Transtornos Cognitivos e Psicose em DP – Identificação e Manejo
Pedro Renato de Paula Brandão, DF
16h45 - 17h10
Depressão e Ansiedade em DP - Identificação e Manejo
Carlos Roberto de Mello Rieder, RS
17h10 - 17h40
Dor e Hipotensão Postural em DP - Identificação e Manejo
Emilia Gatto, ARG
17h40 - 18h00
Discussão

18h00 - 19h00
SIMPÓSIO FQM
AN UPDATE ON THE ROLE OF LEVODOPA IN PARKINSON'S DISEASE
Palestrante
Oscar S. Gershanik, ARG

19h00 - 20h00
COQUETEL DE ABERTURA

25 julho

07h30 - 10h10
Coordenador
Luiz Augusto Franco de Andrade, SP

07h30 - 07h55
Tauopatias - O que há de novo
Jacy Bezerra Parmera, SP
07h55 - 08h20
Atrofia de Múltiplos Sistemas - O que há de novo
Vitor Tumas, SP
08h20 - 08h45
Interface Parkinsonismo-Ataxia: Além de AMS
Hélio Afonso Ghizoni Teive, PR
08h45 - 09h10
Parkinsonismo Atípico
Orlando Graziani Povoas Barsottini, SP
09h10 - 09h40
Discussão

10h10 - 10h40 Coffee Break

10h40 - 12h25
Coordenadora
Monica Santoro Haddad, SP

10h40 - 11h10
Genética de Parkinsonismo no Brasil
Artur Schumacher Schuh, RS
11h10 - 11h40
Coreias Adquiridas - O Que Há de Novo
Gustavo Franklin, PR
11h40 - 12h10
Doença de Huntington - Atualização
Débora Palma Maia, MG
12h10 - 12h25
Discussão

12h25 - 13h25
SIMPÓSIO TEVA
DISCINESIA TARDIA: EVIDÊNCIA, EXPERIÊNCIA E EVOLUÇÃO TERAPÊUTICA
Palestrantes
Hélio Afonso Ghizoni Teive, PR
Marcus Vinicius Della Coletta, AM

14h00 - 15h45
Coordenador
Marcus Vinicius Della Coletta, AM

14h00 - 14h25
Paralisia Cerebral - Encefalopatia Estática?
Laura Silveira Moriyama, SP
14h25 - 14h50
Tiques - O que Há de Novo
Marcelo Masruha, ES
14h50 - 15h15
Tremores - Atualização
Mariana Spitz, RJ
15h15 - 15h45
Discussão

15h45 - 16h15 Coffee Break

16h15 - 18h20
Coordenador
Roberto Cesar Pereira do Prado, SE

16h15 - 16h40
Distonia - Nova Classificação
Sarah Teixeira Camargos, MG
16h40 - 17h05
Distonia em Crianças - Universo em Expansão
Donald Gilbert, USA
17h05 - 17h30
Manejo Terapêutico das Distonias
Carlos Henrique Camargo, PR
17h30 - 17h55
Mioclonias - Atualização
Egberto Reis Barbosa, SP
17h55 - 18h20
Discussão

18h20 - 19h00
SIMPÓSIO ABBVIE
PRO CON DEBATE: É POSSIVEL ELIMINAR COMPLETAMENTE OS SINTOMAS NÃO MOTORES NA DOENÇA DE PARKINSON?
Palestrantes
Débora Palma Maia, MG
Pedro de Melo Barbosa, SP

26 julho

08h30 - 10h55
Coordenador
Ylmar Correa Neto, SC

08h30 - 09h15
Best clinical papers in Movement Disorders 2024/2025
Andrew John Lees, GBR
09h15 - 09h45
Ataxias Tratáveis
Malco Rossi, ARG
09h45 - 10h10
Movimentos Anormais Psicogênicos
José Luiz Pedroso, SP
10h10 - 10h35
Discinesia Tardia
Roberta Arb Saba Rodrigues Pinto, SP
10h35 - 10h55
Discussão

10h55 - 11h15 Coffee Break

11h15 - 13h05
Coordenadores
João Carlos Papaterra Limongi, SP
Luiz Augusto Franco de Andrade, SP

11h15 - 11h40
Movimentos Anormais e Medicina Interna
Clécio Godeiro, RN
11h40 - 12h05
Interface Sono-Movimentos Anormais
Lúcio Huebra, SP

12h05 - 13h05
Sessão de Vídeos
Apresentadores
Christos Ganos, CAN
João Carlos Papaterra Limongi, SP
Luiz Augusto Franco de Andrade, SP

Caso 1
Nathalia Luisa Carlos Ferreira, Welton Cardoso dos Santos, Vitor Santos de Souza, Emilly Rennale Freitas de Melo, Sarah Teixeira Camargos, Debora Palma Maia

Caso 2
Isabela Caldeira de Oliveira, Gleica Maria de Macena Kanawa, Daniel Santos Uchoa, Andre Vinicio Cruz Messias, Igor Fortunato da Silva, Filipe Brito Porto, Julian Leticia de Freitas, Maria Sheila Guimarães Rocha

Caso 3
Lia Araujo Guabiraba, Ricardo Oliveira Horta Maciel, Francisco Eduardo Costa Cardoso

Caso 4
Leonardo Villaverde Buback Ferreira, Gabriela Rodrigues Tomaz, Renato Barradas Rodrigues, Victor Guimarães de Almeida, Larissa Braga Mendes, Enedina Maria Lobato de Oliveira, Carolina Candeias da Silva, Henrique Ballalai Ferraz

Caso 5
Victor Rebelo Procaci, Anna Maria Gomes, Thiago Yoshinaga Tonholo Silva, Raphael Pinheiro Camurugy da Hora, Jackeline Neves Pereira, Orlando Graziani Povoas Barsottini, José Luiz Pedroso

Caso 6
Tamyres Stephane Fonseca Mariano, Aracelle Victor do Carmo Faria, Lucas Wilson Matos Gomes, Delson José da Silva, Luana Mikael

Caso 7
Helton Benevides, Maria Eduarda R. Souza, Myllena Maria S. Santana, Álesson Oliveira Dutra, Jorge Dornellys, Roberto César P. do Prado, Antonio Lucas da Silva Filho

13h05 - 13h15
ENCERRAMENTO
Comissão Organizadora
Egberto Reis Barbosa, SP
Francisco Eduardo Costa Cardoso, MG
Hélio Afonso Ghizoni Teive, PR
Luiz Augusto Franco de Andrade, SP

PT.01
WHOLE GENOME SEQUENCING IN UNDETERMINED ATAXIA
Gomes AM[1], Procaci VR[1], Tonholo Silva TY[1], da Hora RPC[1], Fonseca HARD[2], Barsottini OGP[1], Pedroso JL[1] - [1]Universidade Federal de São Paulo - Setor de Neurologia Geral e Ataxias, [2]Hospital Israelita Albert Einstein - Academic Research Organization

PT.02
BRAIN STRUCTURAL IMPAIRMENT IN SPINOCEREBELLAR ATAXIA TYPE 6: NOT RESTRICTED TO THE CEREBELLUM
Massuyama BK[1], Rezende TJR[2], França Junior MC[2], Barsottini OGP[3], Pedroso JL[3] - [1]Federal University of Sao Paulo - Department of Neurology and Neurosurgery, [2]State University of Campinas - Deparment of Neurology, [3]Federal University of Sao Paulo - Department of Neurology and Neurosurgery

PT.03
BEYOND MONOGENIC PARKINSON'S DISEASE: A CASE SERIES OF PARKINSONISM WITH OTHER GENETIC CAUSES
Echeverria F[1], Quiroga SR[1], Costa G[1], Zeballos MB[2], Moron GD[2], Arakaki T[1], Garretto NS[1], Kauffman M[2] - [1]Hospital J. M. Ramos Mejia - Neurology, [2]Hospital J. M. Ramos Mejia - Neurogenetic

PT.04
EXPANSIONES REPETIDAS EN EL GEN *FGF14* COMO CAUSA DE ATAXIA HEREDITARIA DE INICIO EN LA EDAD ADULTA EN PACIENTES CHILENOS
Gassiot J[1], Pellerin D[2], Zuchner S[2], Silva C[3], Pizarro B[4], Contreras M[5], Canals F[3], Besa V[3], Miranda M[6], Bustamante ML[7] - [1]Universidad de Chile - Medical technologist undergrad, [2]University of Miami, [3]Instituto Nacional de Movimientos Anormales, [4]Universidad de Chile - PhD in Medical Sciences student, [5]Departamento de Neurología Hospital de la Serena, [6]Fundación Diagnosis y Clínica MEDS, [7]Instituto de Ciencias Biomédicas Universidad de Chile y Fundación Diagnosis.

PT.05
ATAXIA-TELANGIECTASIA (AT): A CASE SERIES
Gouvea LA[1], Novis LE[1], Silva TYT[1], Raslan IR[1], Pedroso JL[1], Barsottini OGP[1], Lago CSA[2] - [1]Universidade Federal de São Paulo, São Paulo, SP, Brazil - Department of Neurology, [2]Universidade Federal de São Paulo, São Paulo, SP, Brazil - Department of Allergy, Clinical Immunology and Rheumatology

PT.06
REPEAT EXPANSIONS IN THE *RFC1* G ENE AS A CAUSE OF ADULT-ONSET HEREDITARY ATAXIA IN CHILEAN PATIENTS
Quezada TBK[1], Houlden H[2], Dominik N[2], Pellerin D[2], Cortese A[3], Silva C[4], Yass MB[5], Cortes AM[5], Miranda M[6], Bustamante ML[7] - [1]Universidad de Chile - Medical student, [2]University College of London, [3]University College London, [4]Instituto Nacional de Movimientos Anormales (INMOV), [5]Fundación Diagnosis, [6]Fundación Diagnosis and Clínica MEDS, [7]Universidad de Chile and Fundación Diagnosis

PT.07
CASE REPORT: JUVENILE FORM OF NIEMANN-PICK TYPE C DISEASE
Belfort AC[1], Thiersch LMS[1], Camelo CCS[1], Cruz MCAOF[1], Arantes RR[1] - [1]Hospital das Clínicas da UFMG

PT.08
UNDIAGNOSED DISEASES UNIT AT THE UNIVERSITY OF CHILE: EXPERIENCE FROM 3 YEARS OF WORK
Muñoz A[1], Barreto M[2], Arancibia M[3], Miranda A[2], Pizarro B[4], Miranda M[5], Bustamante ML[6] - [1]Universidad de Chile - Medical student, [2]Fundación Diagnosis, [3]Universidad de Chile - Masters in genetics student, [4]PhD in Medical Sciences student, [5]Fundación Diagnosis and Clinica MEDS, [6]Universidad de Chile and Fundación Diagnosis

PT.09
CASE SERIES: ALTERNATING HEMIPLEGIA OF THE CHILDHOOD WITH HETEROGENEOUS MANIFESTATION
Xavier BFD[1], Thiersch LMS[1], Teixeira FL[1], Diniz SSL[1], Camelo CCS[1], Gianetti JG[1], Santos VSJ[1], Lopes DFS[1], Coelho BCL[1] - [1]UFMG - Neuropediatria

PT.10
CHOREA ASSOCIATED WITH GLUTARIC ACIDURIA TYPE 1 – DIAGNOSIS AND TREATMENT: CASE REPORT
Lamanes BF[1], Cabral JG[1], Santos IE[1] - [1]Federal University of Uberlândia - Pediatric Neurology

PT.12
HUNTINGTON'S DISEASE-LIKE 2 PATIENT'S PROFILE IN A BRAZILIAN COHORT
Silva CC[1], Boone DL[1], Tumas V[2], Vilela G[2], Cavalcanti M[1], Saba R[1], Borges V[1], Ferraz H[1] - [1]Universidade Federal de São Paulo - Department of Neurology, [2]Faculdade de Medicina de Ribeirão Preto - Department of Neuroscience and Behavior Science

PT.13
NEURODEGENERATION WITH BRAIN IRON ACCUMULATION (NBIA) IN A PEDIATRIC PATIENT WITH SUSPECTED KUFOR-RAKEB SYNDROME
Lopes DFS[1], Xavier BFD[1], Coelho BCL[1], Camelo CCS[1], Teixeira FL[1], Gianetti JG[1], Thiersch LMS[1], Diniz SSL[1], Santos VSJ[1] - [1]Hospital das Clínicas of the Federal University of Minas Gerais

PT.14
TWNK MUTATION AND ITS INTERPLAY WITH PARKINSONISM: A CASE REPORT.
Silva EAR[1], Maciel R, Rosa N, Cunningham M, Palma DM, Camargos S[2], Cardoso FEC[2] - [1]Universidade Federal de Minas Gerais - assistent neurologist, [2]Universidade Federal de Minas Gerais - Professor of neurology

PT.15
OMPOUND HETEROZYGOUS *VPS13A* VARIANTS IN A CHILEAN CHOREA-ACANTHOCYTOSIS PATIENT.
Fernandez-Toledo E[1,2], Meza P[1], Saffie-Awad P[3] - [1]University of Concepcion - Specialties, [2]Universidade Federal de Minas Gerais - Pós-Graduação em Ciências Aplicadas à Saúde do Adulto,, [3]Clinica Santa María - Neurology

PT.16
TREATABLE CHILDHOOD-ONSET GENETIC DYSTONIA-ATAXIA SYNDROME
Wainberg F[1], Merello M[1], Rossi M[1] - [1]FLENI - Movement disorders section, Neurology department.

PT.17
CEREBRAL CREATINE DEFICIENCY DUE TO A HOMOZYGOUS NONSENSE MUTATION IN THE *GAMT* GENE (P.W174*): A GENETICALLY CONFIRMED CASE REPORT
Cordeiro JC[1], Furlam TO[1], Guabiraba LA[1], Souza VS[1], Machado GCOG[1], Santos WC[1], Carmargos ST[1], Arantes RR[2] - [1]Hospital das Clínicas da Universidade Federal de Minas Gerais - Neurologia, [2]Hospital das Clínicas da Universidade Federal de Minas Gerais - Genética

PT.18
SAFETY AND EFFICACY OF VATIQUINONE TREATMENT IN FRIEDREICH ATAXIA PATIENTS FROM MOVE-FA: A PHASE 3, DOUBLE-BLIND, PLACEBO-CONTROLLED TRIAL
Ferreira JJG[1], Lynch D[2], Duquette A[3], França Junior MC[4], Perlman S[5], Durr A[6], Machado DS[7], Federhen A[7], Golden L[7], Zesiewicz T[8] - [1]PTC Therapeutics - Medical Affairs, [2]Children's Hospital of Philadelphia, [3]Centre Hospitalier de l'Universite de Montreal, [4]Universidade Estadual de Campinas, [5]University of California, [6]Sorbonne Université, [7]PTC Therapeutics, [8]University of South Florida

PT.19
IMPROVEMENT IN UPRIGHT STABILITY SUBSCALE OF MFARS WITH VATIQUINONE TREATMENT IN MOVE-FA: A PHASE 3, DOUBLE-BLIND, PLACEBO-CONTROLLED TRIAL
Ferreira JJG[1], Lynch D[2], Duquette A[3], França Junior MC[4], Perlman S[5], Durr A[6], Machado DS[1], Federhen A[1], Golden L[1], Zesiewicz T[7] - [1]PTC Therapeutics, [2]Children's Hospital of Philadelphia, [3]Centre Hospitalier de l'Universite de Montreal, [4]Universidade Estadual de Campinas, [5]University of California, [6]Sorbonne Université, [7]University of South Florida

PT.20
LONG-TERM VATIQUINONE TREATMENT SLOWS FA DISEASE PROGRESSION RELATIVE TO FACOMS NATURAL HISTORY
Ferreira JJG[1], Cherry JJ[1], Duquette A[2], França Junior MC[3], Perlman S[4], Durr A[5], Machado DS[1], Federhen A[1], Golden L[1], Lynch D[6], Zesiewicz T[7] - [1]PTC Therapeutics, [2]Centre Hospitalier de l'Universite de Montreal, [3]Universidade Estadual de Campinas, [4]University of California, [5]Sorbonne Université, [6]Children's Hospital of Philadelphia, [7]University of South Florida

PT.21
DENTATORUBRAL-PALLIDOLUYSIAN ATROPHY (DRPLA): A CASE REPORT
Alves KN[1], Perini LB[1], Fernandez AL[1], Silva SMCA[1], Pinto RASR[1] - [1]Hospital do Servidor Público Estadual de São Paulo - Neurologia

PT.22
MANIFESTAÇÕES CLÍNICAS EM INDIVÍDUOS COM ALELOS INTERMEDIÁRIOS NO GENE DA DOENÇA DE HUNTINGTON: UMA REVISÃO SISTEMÁTICA DE RELATOS DE CASO E SÉRIE DE CASOS
Rodrigues KL[1], Boone DL[1], Fernandes MM[2], Corso AMS[3], Santos FM[2], Pinto RASR[4], Borges V[1], Pinto RASR[1], Ferraz HB[1] - [1]Federal University of São Paulo, [2]State University of Rio Grande do Norte, [3]Federal University of Paraná, [4]Santa Marcelina Faculty of Medicine

PT.23
MOVEMENT DISORDERS IN HEREDITARY PRIONOPATHY
Machado LS[1], Gomes FLC[1], Ferreira LVB[1], Silva TYT[1], Pedroso JL[1], Barsottini OGP[1] - [1]Universidade Federal de São Paulo - Department of Neurology

PT.24
AUTOSSOMAL-DOMINANT STRIATAL DEGENERATION WITH TREMORS: AN UNIQUE ASSOCIATION OF PDE8B AND GBA1 MUTATIONS
Salanti LG[1], Bonilha PAAM[1] - [1]Hospital Universitário Cajuru - Neurologia

PT.25
UNMASKING MIMICS IN CEREBRAL PALSY: A CASE OF ATP1A3 MUTATION WITH MYOPATHIC FEATURES
Quintero-Giraldo LP[1], Bobadilla-Quesada EJ[2] - [1]SES Hospital de Caldas - Neurología, [2]Hospital Universitario San Ignacio - Unidad de Neurociencias

PT.26
EXPANDING THE PHENOTYPIC SPECTRUM OF VPS16-RELATED DYSTONIA: A CASE REPORT
Carrá MA[1], Sanguinetti A[1], Ziegler G[1], Povedano GP[1], Tschopp AL[1] - [1]Hospital Churruca Visca - Neurology

PT.27
INTRAFAMILIAL PHENOTYPIC VARIABILITY IN SPG4-ASSOCIATED HEREDITARY SPASTIC PARAPLEGIA: A CASE REPORT
Carrá MA[1], Sanguinetti A[1], Tschopp AL[1], Ziegler G[1], Povedano GP[1] - [1]Hospital Churruca Visca - Neurology

PT.29
RAPID-ONSET GENERALIZED DYSTONIA CAUSED BY A MUTATION IN THE *MT-ND6 GENE*
Costa MA[1], Boone DL[1], Silva CC[1], Pinto RASR[1], Aguiar PC[1], Tengan CH[1], Borges V[1], Ferraz HB[1] - [1]Federal University of São Paulo - Department of Neurology and Neurosurgery

PT.30
FAHR DISEASE IN YOUNG ADULT AGE: A CASE REPORT OF A 22 YEARS-OLD BOY WITH IDIOPATIC BASAL GANGLIA CALCIFICATION
Ferreira NLC[1], Mendes PFB[1], Oliveira LA[1], Ruela VP[1], Machado GCOG[1], Souza VS[1], Camargos ST[1], Fernandes BFS[1] - [1]Hospital das Clínicas da UFMG - Neurology

PT.31
CASE REPORT OF A PROGRESSIVE NEURODEGENERATIVE DISEASE AFTER HEAD TRAUMA: A PROBABLE LEUKOENCEPHALOPATHY
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