

## INFORMAȚII PERSONALE

## Dana Cristina CRAIU



## LOCUL DE MUNCA

1. UMF Carol Davila, Departamentul 6 – Neuroștiințe Clinice, Disciplina Neurologie Pediatrică II
2. Clinica Neurologie Pediatrică, Centru de Referință de Boli Rare Neurologice Pediatrică, Spital Clinic Prof. Dr. Alexandru Obregia, membru în rețelele Europene de Boli Rare EpiCARE și ENDO-ERN

## EXPERIENȚA PROFESIONALĂ

**MEDICALA**

2004 - curent  
2000 - 2014  
1994 - 1999

Medic primar Neurologie Pediatrică  
Medic specialist Neurologie Pediatrică  
Medic rezident Neurologie Pediatrică  
Spital Clinic de Psihiatrie Prof. Dr. Al. Obregia, Sos. Berceni 10, sect 4, București (<https://spital-obregia.ro/>; <https://bolirare-obregia.ro/>)

- Ingrijire pacienti copii cu afectiuni neurologice pediatrică, predominant epilepsii, alte boli rare în domeniu, evaluare prochirurgicală pentru chirurgia epilepsiei, interpretare EEG, video-EEG
- Detaliere stagii în cadrul rezidentiatului: Spitalul de Pediatrie, V. Gomoiu (1 an - pediatrie); Psihiatrie Infanțila, Spitalul Al. Obregia (3 luni - Psihiatrie pentru copii); Neurochirurgie, Spitalul Bagdasar (3 luni - neurochirurgie); Neonatologie, Spitalul Polizu (3 luni - neonatologie); Neurologie adulți, IBCV (3 luni - neurofiziologie); Neurologie Pediatrică, Spitalul Al. Obregia (3 ani - neurologie pediatrică);

1993 - 1994

**Medic Stagiar**

Institutul National pentru Sanatatea Mamei și Copilului – Spital Polizu, Str. Gh. Polizu nr. 38-54, Sector 1, București (<https://www.insmc.ro/instituti/componenta-og-polizu/>)

- Ingrijire pacienti de toate varstele

**ACADEMICA**

2014 - 2022  
2011 - 2014  
2004 - 2011  
1999 - 2004

Prof. Univ. Neurologie Pediatrică  
Conf. Univ. Neurologie Pediatrică  
Sef Lucrari Neurologie Pediatrică  
As. Univ. per det Neurologie Pediatrică

UMF Carol Davila București, Str. Dionisie Lupu Nr. 37, sector 2, București (<https://umfd.ro/>)

- Activități de predare studenți, rezidenți, cadre medicale (cursuri postuniversitare)
- Activități cercetare clinică, genetică neurologică, neurofiziologie clinică

**CONDUCERE**

2022 - 2026  
2021 - 2026  
2020 - prez  
2018 - prez  
2016 - prez

Chair a Comisiei de Educație și training al EPNS (Soc. Europeană de Neurologie Pediatrică)  
Chair al Comisiei de Ghiduri a EPNS  
Prodecan Cercetare UMF Carol Davila – Facultatea de Medicină  
Presedinte al Comisiei de Neurologie Pediatrică a Ministerului Sănătății  
Coordonator Centru de Expertiză Boli Rare Neurologie Pediatrică Obregia, membru EpiCARE și ENDO-ERN

2015 - prez  
2009 - 2017  
2009 - 2017  
2008 - prez

Presedinte SRIE (Societatea Română Impotriva Epilepsiei, Chapter al ILAE)  
Presedinte CNA (Comisia de Advisorii Nationali) a EPNS  
Presedinte TAB (Training Advisory Board) a EPNS  
Sef Disciplina Neurologie Pediatrică II – Departament Neuroștiințe Clinice, UMF Carol Davila – Facultatea de Medicină

2008 - 2016

Sef Clinica Neurologie Pediatrică Spital Al. Obregia

**EDUCAȚIE ȘI FORMARE**

2012 Medic Specialist Pediatru (examen Martie 2011, OMS 740/2012)  
 2005 Doctor in Medicina (sustinere Nov 2004, confirmat Ordin MEC 3956/24.04.2005)  
 2004 Medic Primar Neurolog Pediatru (examen iun 2004, confirmat Ordin MS 1067/2004)  
 2001 Competenta Neurofiziologie (EEG, EMG, PEC) – sesiune examen aprilie 2001  
 Diploma cu Seria A, Nr 1056/ 2001  
 1999 Medic Specialist Neurolog Pediatru (Confirmat OMS 900/1999)  
 1993 Dobandire titlu Doctor Medic, Specializarea Medicina (Diploma Seria L. Nr. 426) in urma absolvirii Facultatii de MG (1987-1993)  
 UMF Carol Davila Bucuresti, Str Dionisie Lupu Nr. 37, sector 2, Bucuresti (<https://umfcd.ro/>)

Scrieți nivelul EQF, dacă îl cunoașteți

Cursuri educationale la care am participat

- Abilitati teoretice si practice, conform curriculumelor de pregatire
- Curs Training VNS – 2021 – 1 zi
- Cursuri SEEG – Venetia – cate 1 saptamana/an – 2018 (fizic), 2019 (fizic), 2021 (on-line)
- Curs Training VNS – 2 zile - 2018
- “Train the trainees” curs, EUREPA – ILAE, Malta 2008 – diploma de formator Eureka.
- Curs: Farmacoterapie in epilepsii - 2008 (educatie la distanta – EUREPA)
- Curs: EEG Basic - 2007 (educatie la distanta- EUREPA)
- San-Servolo scoala de vara: Copilul cu epilepsie – 07.2007 (2 saptamani)
- San-Servolo: scoala de vara; Epilepsii remediabile chirurgical – 07.2006 (2 saptamani)
- Curs: Epilepsii genetice – 2005 (educatie la distanta - EUREPA)
- Breakfast seminarii in congresele de epilepsii – annual
- Curs: ‘Explorări electrice în Neurologie ’ - EEG, EMG, EP, 2001
- 3 luni Bursa educationala – EEG pediatric, video-EEG, evaluare prechirurgicala in Stichting Epilepsie Instellingen ,Olanda, sub directa indrumare a Dr. Walter van Emde Boas, (sept - dec 2000)
- Curs: ‘Progres in Tratamentul Epilepsiilor’, 1999 (București )
- Curs: ‘Explorări electrice în Neurologie’ - EEG, EMG, EP, 1997
- Curs: Progres in Tratamentul Epilepsiilor’ , 1996 (Bucuresti)

**COMPETENTE PERSONALE**

Limba maternă

Romana

Alte limbi străine cunoscute

Engleza

Germana

	INTELEGERE		VORBIRE		SCRIERE
	Ascultare	Citare	Participare la conversatie	Discurs oral	
Engleza	C1	C1	C1	C1	C1
Germana	A1	A1	A1	A1	A1

Competențe organizaționale/manageriale

1. Program de studii pentru obtinerea atestatului „Managementul serviciilor de sanatate” (14.03 – 27.07.2016) – Scoala Nationala de Sanatate Publica, Management si Perfectionare in Domeniul Sanitar Bucuresti (Seria At Nr 0285)
2. Curs de formare ca formator organizat de ILAE -EUREPA (Liga Internationala Impotriva Epilepsiei): Train the Trainees – 2008 – Malta – 1 saptamana

Bun organizator, cu spirit de echipa

Abilitati de organizare cursuri, conferinte, programe de formare

Bun manager de proiecte de cercetare sau de dezvoltare activitati si programe in cadrul echipei UMF

Competențe digitale

**AUTOEVALUARE**

Procesarea informației	Comunicare	Creare de conținut	Securitate	Rezolvarea de probleme
utilizator independent	utilizator independent	utilizator independent	utilizator elementar	utilizator elementar

Alte competențele informatice:

- o bună stăpânire a multor programe office (word, excel, power-point)
- bune cunoștințe de editare foto, dobândite autodidact
- folosire programe de stimulare a interacțiunii – Kahoot, etc
- folosire programe comunicare la distanță și telemedicină: Zoom, Google Meets, Teams, DocBook
- folosire programe de baze de date

INFORMATII SUPLIMENTARE

Alte afilieri

- Membru Fondator SRNP (societatea Romana de Neurologie Pediatrica) 2015
- Membru Board SRNP 2015 – 2019
- Membru al Comisiei Pediatrica a ILAE 2009 - 2013
- Membru Board ILAE-Europe (fost CEA-ILAE) 2013 – 2021
- Membru al Grupului Roman de Cercetare in Genetica Epilepsiei din 2009
- Membru al Grupului Roman de Chirurgia Epilepsiei din 2007
- Membru Board EPNS 2007-prezent
- Membru EPNS din 2005
- Membru ICNA (International Child Neurology Association) din 2001
- Membru fondator RONEP (Fundatia Romana de Neurologie si Epileptologie) – 1998

Cursuri educationale predate

**Cursuri pentru studenti**

2000 - 2007, cursuri pentru modulul de Neurologie Pediatrica:

Status Epilepticus  
 Sindroame neurocutanate  
 Encefalopatia hipoxic-ischemica  
 Hipertensiunea intracraniana  
 Edem cerebral  
 Hidrocefalii

2008-prezent:

Epilepsii – definitii, clasificare, diagnostic  
 Epilepsy – sindroame la copil și adolescent, tratament  
 Status Epilepticus  
 Sindroame neurocutanate  
 Encefalopatia hipoxic-ischemica  
 Hipertensiunea intracraniana  
 Edem cerebral  
 Hidrocefalii

**Cursuri pentru rezidenti:**

- 2008-prezent: Epilepsii, Status epilepticus, Sindroame neurocutanate, EHIP, Hidrocefalii, Edem cerebral, HIC, EEG de baza – teoretic și practic,
- Journal club – săptămânal – prin rotație cu colegii din colectiv
- Prezentari cazuri – săptămânal împreună cu colegii din colectiv prin rotație

**Cursuri postuniversitare (specialisti Neurologie, NP, MF, Neonatologie, Psihiatrie adulti-copii):**

- Curs de EEG basic – 3 săptămâni
- Curs de sindroame in epilepsie – 1 sapt
- Curs de tratament in epilepsie – 1 săptămâna

Cursuri optionale

1. EEG basic – 7 zile – stud an III
2. Urgente la cam de garda

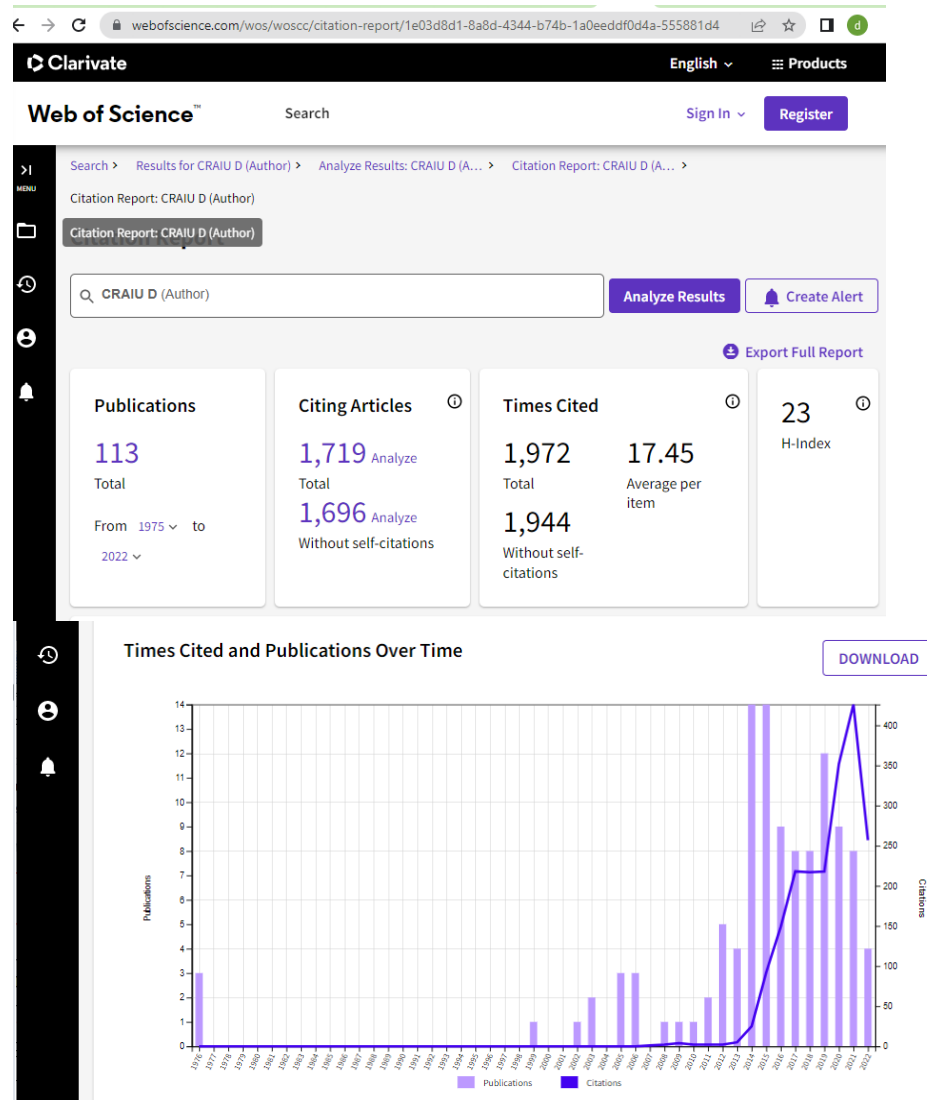
Trainer – Virepa BEEG (Basic EEG) curs de 3 săptămâni – 2/an din 2014 până în prezent

Trainer – Virepa PEEG (Pediatric EEG) curs de 3 săptămâni – 1/an din 2018 până în prezent

**LUCRARI STIINTIFICE  
PREMIATE: (in calitate de coautor)**

Tratatul: Afecțiuni neuromusculare la sugar, copil și adolescent, sub redacția Sanda Magureanu, editura Amaltea 2004, a primit premiul Gheorghe Marinescu al Academiei Române în 2006

**Citări** 1972 citări (**1944 fara autocitare**), **indice Hirsch 23**, conform Web of Science Core Collection



**ACTIVITATE DE CERCETARE:**

<b>1. Grant international – director proiect</b>					
1	POSCE-Axa II/ Op.02.2.4./ INFO ACT/ 2009-2011	Informatizarea activitatii administrative in cadrul departamentului de cercetare al Spitalului Clinic de Psihiatrie Prof. Dr. Al. Obregia	Fonduri europene	471 961 lei	Din Lista de articole ca autor principal : AP3 Coautor: CA1,3-6, 8, 11,12, 16-20, 23-25, 28, 30, 34, 40, 43, 47-52, 54, 55, 60-65
2	6-EUROCC/06.06.2011 din Programul PN II – IDEI; Proiect tip ESF-EUROCORES/ 2011-2014	IP-09: Phenotype-genotype correlations in rare epilepsy syndromes in Romania (proiect individual partener in consortiu European EuroEPINOMICS –RES	Fonduri de la bugetul de stat, Romania fiind membru ESF (European Science Foundation) care a lansat competitia de proiecte.	378.000 RON	E, Depienne C, Balling R, Barisic N, Baulac S, Caglayan HS, Helbig I, Hjalgrim H, Hoffman-Zacharska D, Jahn J, Klein KM, Koeleman BP, Komarek V, Krause R, LeGuern E, Lehesjoki AE, Lemke JR, Lerche H, Marini C, May P, Møller RS, Muhle H, Palotie A, Pal D, Rosenow F, Selmer K, Serratos JM, Sisodiya S, Stephani U, Sterbova K, Striano P, Suls A, Talvik T, von Spiczak S, Weber Y, Weckhuysen S, Zara F. <b>De novo mutations in HCN1 cause early infantile epileptic encephalopathy.</b> <i>Nat Genet.</i> 2014 Jun;46(6):640-5. doi: 10.1038/ng.2952. Epub 2014 Apr 20.
3	IBISD/ GEE006.10/ No.011093-23MAR-00-2012-2014 - terminat	Search of biomarkers for diagnosis, monitoring of disease and therapeutic response in Duchenne’s muscular dystrophy	Genethon Franta	28.000 euro	<b>Craiu D.</b> DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitara „Carol Davila”, Bucuresti, 2013.
4	Proiect tip FP7/ European Comision- Health Programme 2/ reference No. 534055/ 2013-2016	CEC 2013: Centrul de referinta de chirurgia epilepsiei la copii – proiect Colaborator al proiectului European E-Epilepsy: A European pilot network of reference centers in refractory epilepsy and epilepsy surgery	Fonduri europene	Proiect general finantat cu 1.429.420 euro	Iliescu C, <b>Craiu D</b> , Diagnostic approach of Epilepsy in Childhood and Adolescence, <i>Maedica - A Journal of Clinical Medicine</i> , Volume 8, No. 2, 2013.

<b>2. Grant international – membru in echipa de cercetare</b>					
1	POSCE axa II – Operatiunea O2.2.1.-2009-4/ SMIS 14042/ Contract 910 – 21.12.2012/ Perioada 2012-2015	Centrul de Cercetare Translationala in Psihiatrie si Neurostiinte	Fonduri europene	71.580.476 RON	NU
2	COST BM1004/ 2010-2013	Enhancing the scientific study of early autism: A network to improve research, services and outcomes (Imbunatatirea cercetarii stiintifice a autismului precoce: O retea pentru imbunatatirea cercetarii, serviciilor si prognosticului)	Fonduri europene	Acopera integral costurile deplasarilor la toate intalnirile proiectului	Grillo E1, Villard L, Clarke A, Ben Zeev B, Pineda M, Bahi-Buisson N, Hryniewiecka-Jaworska A, Bienvenu T, Armstrong J, Roche-Martinez A, Mari F, Veneselli E, Russo S, Vignoli A, Pini G, Djuric M, Bisgaard AM, Mejaški Bošnjak V, Polgár N, Cogliati F, Ravn K, Pintaudi M, Melegh B, <b>Craiu D</b> , Djukic A, Renieri A. Rett networked database: an integrated clinical and genetic network of Rett syndrome databases. <i>Hum Mutat.</i> 2012 Jul;33(7):1031-6. doi: 10.1002/humu.22072. Epub 2012 Apr 13.
3	COST BM1208/2013-2017	European Network for Human Congenital Imprinting Disorders	Fonduri europene	Acopera integral costurile deplasarilor la toate intalnirile proiectului	Duca G.D, <b>Craiu D</b> , Boer M, Chirieac S.M, Arghir A, Tutulan-Cunita A, Barca D, Iliescu C, Lungean A, Magureanu S, Budisteanu M, Diagnostic approach of Angelman Syndrome, <i>Maedica - A Journal of Clinical Medicine</i> , Volume 8, No. 4, 2013.
4	COST CA15111/2016-2019	European Network on Myalgic Encephalomyelitis/Chronic Fatigue Syndrome (EUROMENE)	Fonduri europene	Acopera integral costurile deplasarilor la toate intalnirile proiectului	
5	CA16118/2017 – 2021	European Network on Brain Malformations Neuro-MIG	Fonduri europene	Acopera integral costurile deplasarilor la toate intalnirile proiectului	

<b>3. Grant national – membru in echipa de cercetare</b>					
1	CP-D/2/ VIASAN/2001-2003	Modificările neuropsihice induse de consumul de droguri la copii și adolescenți	Buget de stat	200.000.000 lei	
2	VIASAN/2003-2004	Modificările neurologice la pacienții cu	Buget de stat	200.000.000 lei	

		infecție HIV / SIDA			
3	VIASAN/2003-2005	Depistarea precoce a bolilor neuromusculare ereditare la copil	Buget de stat	2.000.000 mii lei	
4	VIASAN/2004-2006	Diagnosticul precoce al fenomenelor paroxistice epileptice și neepileptice de somn la copil	Buget de stat	2.700.000 mii lei	<b>Craiu D</b> , Magureanu S, Van Emde Boas W. Are absences truly generalized seizures or partial seizures originating from or predominantly involving the pre-motor areas? Some clinical and theoretical observations and their implications for seizure classification. <i>Epilepsy Res.</i> 2006 Aug; 70 Suppl 1: S141-155. Epub 2006 Jul 18.
5	Parteneriate în domeniile prioritare PN II – Contract 42-130/2008 /2009-2011	Cercetări multidisciplinare – clinice, citogenetice și moleculare – în sindroamele de retard mental asociate cu anomalii congenitale: contribuții în cunoașterea și managementul bolilor rare (RMBGR)	Buget de stat	200.000.000 lei	<b>Craiu D</b> , Kaler S, Craiu M. Role of optic microscopy for early diagnosis of Menkes disease. <i>Rom J Morphol Embryol</i> 2014, 55 (3): 3-6.
6	CEEX M1/ 150/1/ 2006-2008	Abordarea integrată clinică, biochimică și citogenetică a bolilor neurogenetice pediatrice în vederea inițierii unui registru regional pentru supravegherea bolilor neurologice pediatrice	Buget de stat	1.300.000 lei	<p>Tarta-Arsene O, Barca D, Burloiu C, Craiu D, Stoian D, Leanca M, Magureanu S. Aspects of epileptic seizures in children with neurofibromatosis type 1. <i>Romanian Journal of Neurology</i>, Volume XII, No. 2, 2013.</p> <p>Bladen CL, Thompson R, Jackson JM, Garland C, Wegel C, Ambrosini A, Pisano P, Walter MC, Schreiber O, Lusakowska A, Jedrzewowska M, Kostera-Pruszyk A, van der Pol L, Wadman RI, Gredal O, Karaduman A, Topaloglu H, Yilmaz O, Matyushenko V, Rasic VM, Kosac A, Karcagi V, Garami M, Herczegfalvi A, Monges S, Moresco A, Chertkoff L, Chamova T, Guergueltcheva V, Butoianu N, Craiu D, Korngut L, Campbell C, Haberlova J, Strenkova J, Alejandro M, Jimenez A, Ortiz GG, Enriquez GV, Rodrigues M, Roxburgh R, Dawkins H, Youngs L, Lahdetie J, Angelkova N, Saugier-Verber P, Cuisset JM, Bloetzer C, Jeannot PY, Klein A, Nascimento A, Tizzano E, Salgado D, Mercuri E, Sejersen T, Kirschner J, Rafferty K, Straub V, Bushby K, Verschuuren J, Beroud C, Lochmüller H. <b>Mapping the differences in care for 5,000 spinal muscular atrophy patients, a survey of 24 national registries in North America, Australasia and Europe.</b> <i>J Neurol.</i> 2014 Jan;261(1):152-63. doi: 10.1007/s00415-013-7154-1. Epub 2013 Oct 27.</p> <p>Bosemani T, Anghelescu C, Boltshauser E, Hoon AH Jr, Pearl PL, Craiu D, Johnston MV, Huisman TA, Poretti A. <b>Subthalamic nucleus involvement in children: a neuroimaging pattern-recognition approach.</b> <i>Eur J Paediatr Neurol.</i> 2014 May;18(3):249-56. doi: 10.1016/j.ejpn.2013.09.010. Epub 2013 Oct 9.</p> <p>Todorov T, Todorova A, Motoescu C, Dimova P, Iancu D, Craiu D, Stoian D, Barbarii L, Bojinova V, Mitev V. <b>Spontaneous recurrent mutations and a complex rearrangement in the MECP2 gene in the light of current models of mutagenesis.</b> <i>Mutat Res.</i> 2012 Jun 1;734(1-2):69-72. doi: 10.1016/j.mrfmmm.2012.04.001. Epub 2012 Apr 16.</p> <p>Grillo E1, Villard L, Clarke A, Ben Zeev B, Pineda M, Bahi-Buisson N, Hryniewiecka-Jaworska A, Bienvenu T, Armstrong J, Roche-Martinez A, Mari F, Veneselli E, Russo S, Vignoli A, Pini G, Djuric M, Bisgaard AM, Mejaški Bošnjak V, Polgár N, Cogliati F, Ravn K, Pintaudi M, Melegh B, Craiu D, Djukic A, Renieri A. <b>Rett networked database: an integrated clinical and genetic network of Rett syndrome databases.</b> <i>Hum Mutat.</i> 2012 Jul;33(7):1031-6. doi: 10.1002/humu.22072. Epub 2012 Apr 13.</p>
7	CEEX M3/PNCP/ Contract Nr. 15.17./2006-2008	Promovarea dezvoltării cercetării în neurologia pediatrică în scopul participării la programele de cercetare ale uniunii europene	Buget de stat	1.000.000 lei	<p>Brandsma R, Spits AH, Kuiper MJ, Lunsing RJ, Burger H, Kremer HP, Sival DA; <b>Childhood Ataxia and Cerebellar Group.</b> Barisic N, Baxter P, Brankovic-Sreckovic V, Calabrò GE, Catsman-Berreoets C, de Coo I, <b>Craiu D</b>, Dan B, Gburek-Augustat J, Kammoun-Feki F, Kennedy C, Mancini F, Mirabelli-Badenier M, Nemeth A, Newton R, Poll-The BT, Steinlin M, Synofzik M, Topcu M, Triki C, Valente EM. Ataxia rating scales are age-dependent in healthy children. <i>Dev Med Child Neurol.</i> 2014 Jun;56(6):556-63. doi: 10.1111/dmcn.12369. Epub 2014 Jan 7.</p> <p>Craiu D, Iliescu C - NEUROLOGIE PEDIATRICA - Note de curs, Editura Universitară " Carol Davila", Bucuresti, 2013.</p> <p>Albeanu A.G, Magureanu S, Craiu D, Lagae L. Hippocampal sclerosis - cause or consequence of mesial temporal lobe epilepsy in children?, <i>Romanian Journal of Neurology</i>, Volume XI, No. 1, 2012.</p>

					<p>Oana Tarta-Arsene, Florin Preoteasa, Sanda Adriana Magureanu, Adrian Iliescu, Dana Craiu, Cristina Motoescu, Eugen Tarta-Arsene, Gabriela Ciobanu. Functional magnetic resonance imaging contribution to language areas assessment in children with non-lesion focal epilepsy, Romanian Journal of Neurology, Volume IX, No. 3, 2010.</p> <p><b>Craiu D</b>, Avram P, Craiu M, Cochino A.V, Minciu I, Tarta-Arsene O, Butoianu N, Burloiu C, Iliescu C, Magureanu S, Measles and Subacute Sclerosing Panencephalitis (SSPE) in the last 18 years in Romania, International Conference on Diagnosis and Treatment in Pediatric Neurology, , Warsaw, Poland, Medimond International Proceedings, 2008.</p>
8	CEEX/ 2005-2008	Integrarea tehnicilor de analiza moleculara in diagnosticarea distrofinoapatilor in perspectiva unor strategii terapeutice si profilactice	Buget de stat	40.000 lei	<b>Craiu D</b> . DISTROFINOPATIILE - Notiuni teoretice. Algoritmi de diagnostic si tratament. Editura Universitara „Carol Davila”, Bucuresti, 2013.

4. Studii clinice internaționale – investigator principal					
1	1042-0500/2006-004285-13/2007-2008	A double-blind, placebo-controlled, dose-ranging clinical study to evaluate the safety, tolerability, and antiepileptic activity of ganaxolone in treatment of patients with infantile spasms			
2	1042-0501/2007-2009	An open-label clinical study to evaluate the safety and antiepileptic activity of ganaxolone in treatment of patients diagnosed with infantile spasms			
3	MK-0462-082-00/	A Worldwide, Randomized, Double Blind, Placebo-Controlled, „Parallel Group Clinical Trial to Evaluate the Safety and Efficacy of Rizatriptan for the Acute Treatment of Migraine in Children and Adolescents			<p>Ho TW, Pearlman E, Lewis D, Hämäläinen M, Connor K, Michelson D, Zhang Y, Assaid C, Mozley LH, Strickler N, Bachman R, Mahoney E, Lines C, Hewitt DJ, Rizatriptan Protocol 082 Pediatric Migraine Study Group. McKhann G, Eliasziw M, Fisher PG, Tennekoon G, Hsu DT, Nassogne MC, Sekhara T, Almadani M, O'Mahony M, Richer L, Illum N, Laugaard-Jacobsen HC, Sander V, Talvik I, Kallela M, Keski-Santti P, Kiukkonen J, Nikkanen E, Nissila M, Partinen M, Peltola J, Annequin D, Cuvelier JC, Fournier-Charriere E, Laborde S, Milh M, Navez ML, Parain D, Suc A, Ebinger F, Evers S, Gaul C, Gendolla A, Jansen JP, Laengler K, Pothmann R, Schellenberg R, Agarwal S, Anand I, Chandran S, Chodhary VB, Harawat PJ, Harsha S, Jog P, Kannan A, Karardan U, Keerthi AS, Nellikunja S, Pandit L, Srinivasa R, Barbanti P, Jegere D, Strautmanis J, Mulleners W, Pop P, Van den Berg P, Bryn B, Kjaerli T, Sommerfelt K, Kochanowska I, Pietrzak M, Strzelecks J, Szatanik M, Wesolowska M, Benga I, Craiu DC, Diaconu G, Gheonea C, Popescu L, Artigas Pallares J, Blanco Barca O, Campistol Plana J, Macaya A, Mosquera Villaverde Mdel C, Reyes Martin A, San Antonio MV, Danielsson B, Ohrner Y, Arthur CP, Bala P, Gardner S, Gosalakal J, Prabhakar P, Abraham A, Adler J, Aguado M, Atalla A, Atri PB, Aurora S, Baber R, Banks JW, Bargar R, Barker J, Barrington P, Bateman L, Baur CE, Bays H, Bennett NL, Berenson FR, Berman GD, Berstein A, Bhatia P, Blumenfeld A, Bramlet D, Broker RE, Byrd S, Cady R, Calcagno J, Camacho A, Carlini W, Casadonte J, Choi S, Christensen SG, Civitarese F, Clark WD, Corder CN, David R, Dhaduk V, Duffy CA, Earle R, Edmond M, Edrozo J, Eross E, Erwin JS, Espinosa-Paccini JB, Essink B, Farmer M, Fedlman M, Fernando M, Fieve RR, Fisher M, Flitman S, Ford LB, Former SD, Fox E, Frandsen B, Fry JA, Fuller GR, Gaffney ME, Gasecki A, Gay C, Gelfand S, Giancarlo T, Giblin J, Glover MC, Goldstein G, Goodman H, Gordon G, Gorrela SV, Grainger W, Gray J, Gupta P, Halthore SN, Handal NM, Harris DJ, Harvey B, Hazan L, Hedrick J, Henry DC, Herring MO, Hines RL, Holloway W Jr, Horwitz AE, Huling R, Igleburger J, Jennings W, Jones T, Julien KA, Katie A, Kent EF, Khan A, Khan A, Kimmel MA, Klein TR, Knutson J, Krafty MB, Kratzer J, Kwentus J, Lacey D, Lane P, Lebron D, Lesh K, Ley J, Linder S, Liu E, Luber SR, Machanic B, Marcadis I, Markely HG, Markovitz PJ, Mate LJ, Mathew NT, McAllister P, McGettigan JW, Means P, Mechtler L, Mehra V, Melamed I, Miller DC, Miller JL, Miranda F, Moon M, Muhar IM, Munoz S, Murphy K, Nanavaty R, Nayak NA, Nelson J, Nett R, Neufeld N, Nussdorfer T, O'Carroll C, Oftadeh L, O'Hern R, Onder R, O'Reilly T, Palanpurwala K, Pathak L, Pearlman EM, Pendleton J, Peterson R, Poy IG, Qaqundah P, Renfro J, Richer R, Riesenberger R, Robbins L, Roberts KL, Rothner DA, Salem G, Samudrala S, Saper JR, Sarkis E, Saunders M, Schaefer F, Schreiber AO, Sebastian V, Sedill A, Silas PE, Silverboard G, Sivakumar K, Smietana S, Snell P, Sperling M, Spiegel C, Spierings E, Stedman M, Stepp WP, Strzinek R, Taber LM, Taghadosi M, Taylor L, Thurman LM, Ventre P, Wade RD, Wagner A, Weissman JD, Williams D, Wilson MC, Wisman P, Wolfson E, Woodruff BE, Wyszomierski DA, Zinn MM. <b>Efficacy and tolerability of rizatriptan in pediatric migraineurs: results from a randomized, double-blind, placebo-controlled trial using a novel adaptive enrichment design.</b> <i>Cephalalgia</i>. 2012 Jul;32(10):750-65. Epub 2012 Jun 18.</p>
4	B4Z-EW-B013/ 2008 -2011	Investigation of factors associated with changes in ADHD severity during a 2 year follow-up period in patients that are responders and stable on their first			

		pharmacotherapy			
5	A0081041/ 2010 – 2015	Double blind, placebo controlled, parallel group, multicenter study, of the efficacy and safety of pregabalin as adjunctive therapy in children 14-16 years of age with partial onset seizures			Antinew J, Pitrosky B, Knapp L, Almas M, Pitman V, Liu J, <b>Craiu D</b> , Modequillo M, Nordli D, Farkas V, Farkas MK. Pregabalin as Adjunctive Treatment for Focal Onset Seizures in Pediatric Patients: A Randomized Controlled Trial. J Child Neurol. 2019 Apr;34(5):248-255. doi: 10.1177/0883073818821035. Epub 2019 Jan 27.PMID: 30688135 Clinical Trial. (2 citari)
6	A0081042/ 2012 – 2015	Double blind, placebo controlled, parallel group, multicenter study, of the efficacy and safety of pregabalin as adjunctive therapy in children 1 month to <4 years of age with partial onset seizures.			Mann D, Antinew J, Knapp L, Almas M, Liu J, Scavone J, Yang R, Modequillo M, Makedonska I, Ortiz M, Kyrychenko A, Nordli D, Farkas V, Farkas MK; A0081042 study group (Leanid Shalkevich, Anna Jansen, Ivan Ivanov, Vania Nedkova, Fang Fang, Yi Wang, Jean-Marc Pinard, Ulrich Brandl, Dimitrios Zafeiriou, Anna Altmann, Marianne Berenyi, Monika Bessenyei, Andras Fogaras, Geza Szabo, Aviva Fattal-Valevski, Ki Joong Kim, Ahmad Beydoun, Ghassan Hmameiss, Nor Azni Yahaya, Marissa Barlaan-Lukban, Martha Bolanos, Jo Janette De la Calzada, Maria Estrella Ibe, Maria Antonia Aurora Valencia, <b>Dana Craiu</b> , Georgeta Diaconu, Tatiana Antonova, Elena Belousova, Yulia Karakulova, Olga Khaletskaia, Olga Lvova, Maria Strachunskaya, Ruzica Kravljanc, Dimitrije Nikolic, Francisco Lopez Pison, Ying-Chao Chang, I-Ching Chou, Wang-Tso Lee, Charrin Nabangchang, Oranee Sanmaneechai, Nihal Olgac Dundar, Pinar Gencpinar, Yurii Chomolyak, Dmytro Delva, Volodymyr Martyniuk, Ronald Davis, Jose Ferreira, Jerry Tomasovic). Pregabalin adjunctive therapy for focal onset seizures in children 1 month to <4 years of age: A double-blind, placebo-controlled, video-electroencephalographic trial. Epilepsia. 2020 Apr;61(4):617-626. doi: 10.1111/epi.16466. Epub 2020 Mar 18.PMID: 32189338 Clinical Trial.
7	A0081106/ 2010-2015	A 12- month open label study to evaluate the safety and tolerability of pregabalin as adjunctive therapy in pediatric subjects 1 month to 16 years of age with partial onset seizures and pediatric and adult subject 5 to 65 years of age with primary generalised tonic clonic seizures			
8	MRZ 60201_3070_1 (2013 – 2016)	Prospective, multicenter randomized, double-blind parallel group, dose-response study of three doses Xeomin(incobotulinumtoxin A, NT 201) for the treatment of lower limb spasticity in children and adolescents (age 2-17 years) with cerebral palsy			
9	MRZ 60201_3071_1 (2013 – 2016)	Open-label, non-controlled, multicenter long-term study to investigate the safety and efficacy of Xeomin (incobotulinumtoxin A, NT 201) for the treatment of spasticity of the lower limb(s) or of combined spasticity of upper and lower limb in children and adolescents (age 2-17 years) with cerebral palsy			
10	SP0969 (din 2014 - 2017)	A multicenter, Double-Blind, Randomized, Placebo-Controlled, Parallel-Group Study to Investigate the Efficacy and Safety of Lacosamide as Adjunctive Therapy in Subjects with Epilepsy >4 Years to < years of age with partial - onset seizures			
11	EP0034 (din 2014 – 2017)	A multicenter, Open - Label, long term extension study to investigate the Efficacy and Safety of Lacosamide as Adjunctive Therapy in pediatric subjects with Epilepsy with partial - onset seizures			
12	2013-004448-45 (din 2013 – 2015)	Pilot study to Evaluate the Safety and Feasibility of Autologous Cord Blood Infusion to Children with Cerebral Palsy			
13	MCT8 – 2014-2019	Thyroid hormone analog therapy of patients with severe psychomotor retardation caused by mutation in the MCT8 thyroid hormone transporter : The Triac Trial			Groeneweg S, Peeters RP, Moran C, Stoupa A, Auriol F, Tonduti D, Dica A, Paone L, Rozenkova K, Malikova J, van der Walt A, de Coo IFM, McGowan A, Lyons G, Aarsen FK, Barca D, van Beynum IM, van der Knoop MM, Jansen J, Manshande M, Lunsing RJ, Nowak S, den Uil CA, Zillikens MC, Visser FE, Vrijmoeth P, de Wit MCY, Wolf NI, Zandstra A, Ambegaonkar G, Singh Y, de Rijke YB, Medici M, Bertini ES, Depoorter S, Lebl J, Cappa M, De Meirleir L, Krude H, <b>Craiu D</b> , Zibordi F, Oliver Petit I, Polak M, Chatterjee K, Visser TJ, Visser WE. Effectiveness and safety of the tri-iodothyronine analogue Triac in children and adults with MCT8 deficiency: an international, single-arm, open-label, phase 2 trial. Lancet Diabetes Endocrinol. 2019 Sep;7(9):695-706. doi: 10.1016/S2213-8587(19)30155-X. Epub 2019 Jul 31.PMID: 31377265 Clinical Trial. (21citari)
					Groeneweg S, van Geest FS, Abaci A, Alcantud A, Ambegaonkar GP, Armour CM, Bakhtiani P, Barca D, Bertini ES, van Beynum IM, Brunetti-Pierri N, Bugiani M, Cappa M, Cappuccio G, Castellotti B, Castiglioni C, Chatterjee K, de Coo IFM, Coutant R, <b>Craiu D</b> , Crock P, DeGoede C, Demir K, Dica A, Dimitri P, Dolcetta-Capuzzo A, Dremmen MHG, Dubey R, Enderli A, Fairchild J, Gallichan J, George B, Gevers EF, Hackenberg A, Halász Z, Heinrich B, Huynh T, Kłosowska A, van der Knaap MS, van der Knoop MM, Konrad D, Koolen DA, Krude H, Lawson-Yuen A, Lebl J, Linder-Lucht M, Lorea CF, Lourenço CM, Lunsing RJ, Lyons G, Malikova J, Mancilla EE, McGowan A, Mericq V, Lora FM, Moran C, Müller KE, Oliver-Petit I,



				Paone L, Paul PG, Polak M, Porta F, Poswar FO, Reinauer C, Rozenkova K, Meneve TS, Simm P, Simon A, Singh Y, Spada M, van der Spek J, Stals MAM, Stoupa A, Subramanian GM, Tonduti D, Turan S, den Uil CA, Vanderniet J, van der Walt A, Wémeau JL, Wierzbz J, de Wit MY, Wolf NI, Wurm M, Zibordi F, Zung A, Zwaveling-Soonawala N, Visser WE. Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. <i>Lancet Diabetes Endocrinol.</i> 2020 Jul;8(7):594-605. doi: 10.1016/S2213-8587(20)30153-4.PMID: 32559475 (1 citari)
1 4	CFTY720D2311/ 2011-00567723 (2014 – 2019)	A two-year double-blind, randomized, multicenter, active-controlled study to evaluate the safety and efficacy of fingolimod administered orally once daily versus interferon beta-1a i.m. once weekly in oediatric patients with multiple sclerosis		Chitnis T, Arnold DL, Barwell B, Brück W, Ghezzi A, Giovannoni G, Greenberg B, Krupp L, Rostásy K, Tardieu M, Waubant E, Wolinsky JS, Bar-Or A, Slites T, Chen Y, Putzki N, Merschhemke M, Gärtner J; PARADIGMS Study Group (Andrew Kornberg, Barbara Bajer-Kornek, Sergey Likhachev, Antonio Pereira Gomes Neto, Denise Diniz, José Paz, Regina Alvarenga, Veneta Bojinova-Tchamova, Jean Mah, Sunita Venkateswaran, Krasanka Hafner, Katrin Gross-Paju, Bruno Brochet, Emmanuel Cheuret, François Rivier, Kumaran Deiva, Mathieu Milh, Astrid Blaschek, Regina Trollmann, Rudolf Korinthenberg, Thomas Luecke, Tjalf Ziemssen, Carlo Pozzilli, Francesco Patti, Giancarlo Comi, Girolama Alessandra Marfia, Luigi Maria Edoardo Grimaldi, Maria Trojano, Mauro Zaffaroni, Ruggero Capra, Vincenzo Brescia Morra, Guntis Rozentals, Jurate Laurynaitiene, Nerija Vaidiene-Magistris, Freddy Castro Farfan, Sandra Quinones, Barbara Steinborn, Barbara Ujma-Czapska, Mariusz Stasiulek, Miroslaw Jasinski, Dana Craiu, Alexey Boyko, Ekaterina Kairbekova, Farit Khabirov, Liudmila Kuzenkova, Nadezhda Malkova, Dimitrije Nikolic, Jasna Jancic, Ksenija Gebauer-Bukurov, Jaroslava Payerova, Francisco Gascon Jimémez, Guillermo Izquierdo Ayuso, Mar Mendibe Bilbao, Rogier Hintzen, Victoria Eugenia Fernandez Sanchez, Virginia Meca Lallana, Xavier Montalban Gairin, Karin Nordborg, Banu Anlar, Cengiz Yalcinkaya, Kivildim Gucuyener, Murat Terzi, Serkan Ozakbas, Unsal Yilmaz, Iryna Makedonska, Kateryna Prokopenko, Liudmyla Tantsura, Sergii Moskovko, Tetiana Kobys, Tetiana Muratova, Tetiana Nehrych, Tetiana Prykhodko, Cheryl Hemingway, Evangeline Wassmer, Jay Shetty, Jay Desai, Amy Waldman, Angel Chinae Martinez, Jayne Ness, Kottil Rammohan, Michael Lloyd, Mitchel Williams, Ricardo Ayala, Ronald Davis, Vikram Bhise). <b>Trial of Fingolimod versus Interferon Beta-1a in Pediatric Multiple Sclerosis.</b> <i>N Engl J Med.</i> 2018 Sep 13;379(11):1017-1027. doi: 10.1056/NEJMoa1800149.PMID: 30207920 <b>Free article.</b> Clinical Trial
1 5	[FTY720D/fingolimod] Protocol CFTY720D2311 2016-2023 extensia)	A two-year, double-blind, randomized, multicenter, active controlled Core Phase study to evaluate the safety and efficacy of fingolimod administered orally once daily versus interferon $\beta$ -1a i.m. once weekly in pediatric patients with multiple sclerosis with five-year fingolimod Extension Phase		

### 5. Studii clinice internaționale - membru în echipa de cercetare

1	TOPMAT-MIG-3006/Phase III/2006	A Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of Topiramate for the Prophylaxis of Migraine in Pediatric Subjects 12 to 17 Years of Age		
2	UCBN01009/ 2006-2007	A Double-Blind, Randomized, Multicenter, Placebo-Controlled, Inpatient, Maximum 34 Day Study of Levetiracetam Oral Solution (20-50mg/kg/day) as Adjunctive Treatment of Refractory Partial Onset Seizures in Pediatric Epileptic Subjects Ranging in Age from 1 Month to Less Than 4 Years of Age		
3	UCB N01148 2006-2007	A Multi-Center, Open-Label, Long-Term, Follow-Up Study Of The Safety And Efficacy Of Levetiracetam In Children With Partial Onset Seizures		
4	TOSCA-CRAD001MIC03/2013-2018	An international disease registry collecting data on manifestations, interventions and outcomes in patients with tuberous sclerosis complex-TOSCA		Curatolo P, Jóźwiak S, Nabbout R; TSC Consensus Meeting for SEGA and Epilepsy Management, Adriaenssen M, Berhouma M, Coppola G, Craiu D, Cusmai R, Delalande O, De Saint Martin A, Driever PH, Fohlen M, Grajkowska W, Hertzberg C, Jansen A, Jansen F, Kotulska K, Mandera M, Moavero R, O'Callaghan F, Raffo E, Zonnenberg BA. <b>Management of epilepsy associated with tuberous sclerosis complex (TSC): clinical recommendations.</b> <i>Eur J Paediatr Neurol.</i> 2012 Nov;16(6):582-6. doi: 10.1016/j.ejpn.2012.05.004. Epub 2012 Jun 12.  Jóźwiak S, Nabbout R, Curatolo P; participants of the TSC Consensus Meeting for SEGA and Epilepsy Management. <b>Management of subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC): Clinical recommendations.</b> <i>Eur J Paediatr Neurol.</i> 2013 Jul;17(4):348-52. doi: 10.1016/j.ejpn.2012.12.008. Epub 2013 Feb 5.  Glushkova M, Bojinova V, Koleva M, Dimova P, Bojdarova M, Litvinenko I, Todorov T, Iluca E, Calusaru C, Neagu E, Craiu D, Mitev V, Todorova A. Molecular genetic diagnostics of tuberous sclerosis complex in Bulgaria: six novel mutations in the TSC1 and TSC2 genes. <i>J Genet.</i> 2018 Jun;97(2):419-427.PMID: 29932062. DOI: 10.1007/S12041-018-0927-7 (2 citari)

### 6. Studii clinice naționale – investigator principal

1	2008-2016	Programul național de diagnostic și tratament pentru boli rare și sepsis sever; Interventia pentru diagnosticul și managementul amiotrofiilor spinale și a distrofiilor musculare de tip Duchenne și Becker, precum și prevenirea transmiterii ereditare a		Bladen CL, Thompson R, Jackson JM, Garland C, Wegel C, Ambrosini A, Pisano P, Walter MC, Schreiber O, Lusakowska A, Jedrzejowska M, Kostera-Pruszyk A, van der Pol L, Wadman RI, Gredal O, Karaduman A, Topaloglu H, Yilmaz O, Matyushenko V, Rasic VM, Kosac A, Karcagi V, Garami M, Herczegfalvi A, Monges S, Moresco A, Chertkoff L, Chamova T, Guergueltcheva V, Butoianu N, Craiu D, Korngut L, Campbell C, Haberlova J, Strenkova J, Alejandro M, Jimenez A, Ortiz GG, Enriquez GV, Rodrigues M, Roxburgh R, Dawkins H, Youngs L, Lahdetie J, Angelkova N, Saugier-Verber P, Cuisset JM, Bloetzer C, Jeannot PY, Klein A, Nascimento A, Tizzano E, Salgado
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		acestora			D, Mercuri E, Sejersen T, Kirschner J, Rafferty K, Straub V, Bushby K, Verschuuren J, Beroud C, Lochmüller H. <b>Mapping the differences in care for 5,000 spinal muscular atrophy patients, a survey of 24 national registries in North America, Australasia and Europe.</b> J Neurol. 2014 Jan;261(1):152-63. doi: 10.1007/s00415-013-7154-1. Epub 2013 Oct 27.
2	2008-prezent	Program de sanatate mama si copil, subprogramul Diagnosticul si tratamentul precoce al epilepsiilor, paralizilor cerebrale si intarzierilor neuropsihomotorii la copil si prevenirea complicatiilor acestora			
3	2008-prezent	Programul national de diagnostic si tratament pentru boli rare si sepsis sever: Subprogramul de tratament al bolilor rare-scleroza multipla			
4	2008-prezent	Programul national de diagnostic si tratament pentru poliradiculonevrita acuta si alte boli autoimune acute (Boala Rasmussen, scleroza multipla la copilul <12 ani)			
5	2017 – 2018	249PED/2017 “New microarray design targeting genomic hotspots in epilepsy – a proof of concept” („Design inovativ de investigarea prin microarray a regiunilor genomice implicate in epilepsie - demonstrare experimentală a fezabilitatii”)	Bugetul de stat. Proiecte experimental-demonstrative - PN-III-CERC-CO-PED-2016	600.000 lei	
<b>7. Studii clinice naționale – subinvestigator</b>					
1	Program 3/15/7 al MSF/2002-2004	Perfecționarea tehnicilor de diagnostic si profilaxia recidivelor in epilepsia copilului			
2	Program 3/2005-2009	Prevenirea aparitiei, depistarea precoce, monitorizarea si recuperarea deficientelor senzoriale si de dezvoltare neuropsihomotorie la copil precum si profilaxia epilepsiei rezistente la tratament			
3	Proiect ERANET ERARE 2019-2022	Multi-OMICS interrogation of cerebral cortical malformations. (18-049)			

## ALTE PROIECTE SI PROGRAME

- Proiect Caritabil – Oameni mari pentru copii mici – chirurgia epilepsiei la copii – proiect de dotare a Clinicii Neurologice Pediatrica Obregia cu aparatura de video-EEG, mobilier si consumabile – in valoare de 150 000 euro - finalizat
- Programul National de Epilepsii al Ministerului Sanatatii (mama si copilul: Prevenirea reccurentelor si complicatiilor in epilepsiile copilului – participant 2001, 2002, 2003, 2004; coordonator 2005 and 2006;
- Participant - proiect: “EPILEPSY – A SIMPLE DISEASE, NOT AN UNNATURAL PHENOMENON”, sustinut de Fundatia Soros – s-au efectuat brosurile informative pentru familii si cursuri pentru parinti privind epilepsia 1999 – 2000;

## ARTICOLE

### 13 articole in reviste cu factor de impact ca autor principal:

**AP1. Craiu D, Magureanu S, van Emde Boas W.** Are absences truly generalized seizures or partial seizures originating from or predominantly involving the pre-motor areas? Some clinical and theoretical observations and their implications for seizure classification. *Epilepsy Res.* 2006 Aug;70 Suppl 1:S141-55. doi: 10.1016/j.epilepsyres.2005.11.018. Epub 2006 Jul 18. PMID: 16854562. Factor impact in 2006=2.602 (<https://www.scijournal.org/impact-factor-of-epilepsy-res.shtml>)

**AP2. Craiu D.** What is special about the adolescent (JME) brain? *Epilepsy Behav.* 2013 Jul;28 Suppl 1:S45-51. doi: 10.1016/j.yebeh.2012.12.008. PMID: 23756479. Factor impact in 2013=2.574 (<https://www.scijournal.org/impact-factor-of-epilepsy-behav.shtml>)

**AP3. Craiu D, Dragostin O, Dica A, Hoffman-Zacharska D, Gos M, Bastian AE, Gherghiceanu M, Rolfs A, Nahavandi N, Craiu M, Iliescu C.** Rett-like onset in late-infantile neuronal ceroid lipofuscinosis (CLN7) caused by compound heterozygous mutation in the MFSD8 gene and

review of the literature data on clinical onset signs. *Eur J Paediatr Neurol.* 2015 Jan;19(1):78-86. doi: 10.1016/j.ejpn.2014.07.008. Epub 2014 Aug 7. PMID: 25439737.

Factor impact in 2015=2.395 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>)

**AP4. Craiu D,** Kaler S, Craiu M. Role of optic microscopy for early diagnosis of Menkes disease. *Rom J Morphol Embryol.* 2014;55(3):953-6. PMID: 25329126; PMCID: PMC6456807.

Factor impact in 2014=0.902 (<https://www.scijournal.org/impact-factor-of-rom-j-morphol-embryo.shtml>)

**AP5. Tarța-Arsene O,** Moisa G, Bârcă DG, **Craiu D.** Neurosteroids, a new antiepileptic therapy? *Farmacia,* 2014; 62(4): 633-641.

Factor impact in 2014=0.847 (<https://www.scijournal.org/impact-factor-of-FARMACIA.shtml>)

**AP6. Craiu D.** Implications of Sex Hormones in the Treatment of Women with Epilepsy: Catamenial Epilepsy. *Acta Endo (Buc)* 2014, 10 (1): 102-117. doi: 10.4183/aeb.2014.102

Factor impact in 2014=0.313 (<https://www.scijournal.org/impact-factor-of-acta-endocrinol-buch.shtml>)

**Ap7. Craiu D,** Barborica A, Motoescu C, Donos C, Ciurea J, Mindruta I. Presurgical Evaluation and Epilepsy Surgery in MRI Negative Resistant Epilepsy of Childhood with Good Outcome. *Turk Neurosurg.* 2015;25(6):905-13. doi:10.5137/1019-5149.JTN.12093-14.0. PMID: 26617141.

Factor impact in 2015=0.672 (<https://www.scijournal.org/impact-factor-of-turk-neurosurg.shtml>)

**AP8. Iliescu C,** Tarta-Arsene O, **Craiu D.** Valproic acid, polycystic ovary syndrome and the adolescent with epilepsy. *Revista Farmacia,* 2017; 65(1):1-4

Factor impact in 2017=1.381 (<https://www.scijournal.org/impact-factor-of-FARMACIA.shtml>)

**AP9. Craiu DC.** Outpatient initiation of the ketogenic diet. *Eur J Paediatr Neurol.* 2019 Sep;23(5):672-673. doi: 10.1016/j.ejpn.2019.09.007. PMID: 31672222.

Factor impact in 2019=2.613 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>)

**AP10. Sandu C,** Magureanu SA, Iliescu C, Pomeran C, **Craiu D.** Ketogenic diet treatment for status epilepticus. *Farmacia* 2019; 67(2): 218-225.

Factor impact in 2019=1.525 (<https://www.scijournal.org/impact-factor-of-FARMACIA.shtml>)

**AP11. Craiu D,** Haataja L, Hollody K, Kršek P, Lagae L, Mall V, Parker AP, Steinlin M, Yalnizoglu D, Catsman-Berrevoets C; Committee of National Advisors in Paediatric Neurology in Europe. The training and organization of Paediatric Neurology in Europe: Special report of the European Paediatric Neurology Society & Committee of National Advisors. *Eur J Paediatr Neurol.* 2020 Sep;28:6-15. doi:10.1016/j.ejpn.2020.07.012. Epub 2020 Aug 15. PMID: 32958450.

Factor impact in 2020=2.51 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>)

**AP12. Craiu D,** Renner Primec Z, Lagae L, Vigevano F, Trinka E, Specchio N, Bakhtadze S, Cazacu C, Golli T, Zuberi SM. Vaccination and childhood epilepsies. *Eur J Paediatr Neurol.* 2022 Jan;36:57-68. doi: 10.1016/j.ejpn.2021.11.014. Epub 2021 Dec 3. PMID: 34922162.

Factor impact in 2021=2020=2.51 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>)

**AP13. Craiu DC,** Bastian AE, Zurac SA, Băilă SL, Croitoru M, Craiu M, Diaconu R, Vințan MA, Bârcă DG. Brachial and subclavian arteries aneurysms due to tuberous sclerosis complex mechanisms - case report and literature review. *Rom J Morphol Embryol.* 2022 Jan-Mar;63(1):181-189. doi: 10.47162/RJME.63.1.19. PMID: 36074682.

Factor impact in 2020-2021 – 1,033 (<https://www.scijournal.org/impact-factor-of-rom-j-morphol-embryo.shtml>)

## 72 articole in reviste cu factor de impact co-autor

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**CA47.** Carvill GL, McMahon JM, Schneider A, Zemel M, Myers CT, Saykally J, Nguyen J, Robbiano A, Zara F, Specchio N, Mecarelli O, Smith RL, Leventer RJ, Møller RS, Nikanorova M, Dimova P, Jordanova A, Petrou S; EuroEPINOMICS Rare Epilepsy Syndrome Myoclonic-Astatic Epilepsy & Dravet working group (Albena Jordanova, Sarah von Spiczak, Hiltrud Muhle, Hande Caglayan, Katalin Sterbova, **Dana Craiu**, Dorota Hoffman, Anna-Elina Lehesjoki, Kaja Selmer, Christel Depienne, Johannes Lemke, Carla Marini, Renzo Guerrini, Bernd Neubauer, Tiina Talvik, Arvid Suls, Eric Leguern), Helbig I, Striano P, Weckhuysen S, Berkovic SF, Scheffer IE, Mefford HC. **Mutations in the GABA Transporter SLC6A1 Cause Epilepsy with Myoclonic-Atonic Seizures.** *Am J Hum Genet.* 2015 May 7;96(5):808-15. doi: 10.1016/j.ajhg.2015.02.016. Epub 2015 Apr 9.PMID: 25865495 **Free PMC article.**

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**CA48.** Mignot C, von Stülpnagel C, Nava C, Ville D, Sanlaville D, Lesca G, Rastetter A, Gachet B, Marie Y, Korenke GC, Borggraefe I, Hoffmann-Zacharska D, Szczepanik E, Rudzka-Dybala M, Yiş U, Çağlayan H, Isapof A, Marey I, Panagiotakaki E, Korff C, Rossier E, Riess A, Beck-Woodl S, Rauch A, Zweier C, Hoyer J, Reis A, Mironov M, Bobylova M, Mukhin K, Hernandez-Hernandez L, Maher B, Sisodiya S, Kuhn M, Glaeser D, Weckhuysen S, Myers CT, Mefford HC, Hörtnagel K, Biskup S; EuroEPINOMICS-RES MAE working group (**Dana Craiu**, Peter De Jonghe, Ingo Helbig, Renzo Guerrini, Anna-Elina Lehesjoki, Carla Marini, Hiltrud Muhle, Rikke S Møller, Bernd Neubauer, Deb Pal, Kaja Selmer, Ulrich Stephani, Katalin Sterbova, Pasquale Striano, Tiina Talvik, Sarah von Spiczak), Lemke JR, Héron D, Kluger G, Depienne C. **Genetic and neurodevelopmental spectrum of SYNGAP1-associated intellectual disability and epilepsy.** *J Med Genet.* 2016 Aug;53(8):511-22. doi: 10.1136/jmedgenet-2015-103451. Epub 2016 Mar 17.PMID: 26989088

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**CA49.** Tang S, Addis L, Smith A, Topp SD, Pendziwiat M, Mei D, Parker A, Agrawal S, Hughes E, Lascelles K, Williams RE, Fallon P, Robinson R, Cross HJ, Hedderly T, Eltze C, Kerr T, Desurkar A, Hussain N, Kinali M, Bagnasco I, Vassallo G, Whitehouse W, Goyal S, Absoud M; EuroEPINOMICS-RES Consortium (**Dana Craiu**, Carol Davila, Alexandru Obregia, Peter De Jonghe, Anna-Elina Lehesjoki, Hiltrud Muhle, Bernd Neubauer, Kaja Selmer, Ulrich Stephani, Katalin Sterbova, Pasquale Striano, Tiina Talvik, Sarah von Spiczak, Sarah Weckhuysen, Hande Caglayan, Dorota Hoffman-Zacharska), Møller RS, Helbig I, Weber YG, Marini C, Guerrini R, Simpson MA, Pal DK. **Phenotypic and genetic spectrum of epilepsy with myoclonic atonic seizures.** *Epilepsia.* 2020 May;61(5):995-1007. doi: 10.1111/epi.16508. Epub 2020 May 29.PMID: 32469098

Factor impact in 2020=6.549 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

**CA50.** Carvill GL, Engel KL, Ramamurthy A, Cochran JN, Roovers J, Stamberger H, Lim N, Schneider AL, Hollingsworth G, Holder DH, Regan BM, Lawlor J, Lagae L, Ceulemans B, Bebin EM, Nguyen J; EuroEPINOMICS Rare Epilepsy Syndrome, Myoclonic-Astatic Epilepsy, and Dravet Working Group (Pasquale Striano, Federico Zara, Ingo Helbig, Rikke S Møller, Sarah von Spiczak, Hiltrud Muhle, Hande Caglayan, Katalin Sterbova, **Dana Craiu**, Dorota Hoffman, Anna-Elina Lehesjoki, Kaja Selmer, Christel Depienne, Johannes Lemke, Carla Marini, Renzo Guerrini, Bernd Neubauer, Tiina Talvik, Eric Leguern, Peter de Jonghe, Sarah Weckhuysen), Barsh GS, Weckhuysen S,

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Factor impact in 2018=10.192 (<https://www.scijournal.org/impact-factor-of-am-j-hum-genet.shtml>)

**CA51.** Ingo Helbig, Tania Lopez-Hernandez, Oded Shor, Peter Galer, Shiva Ganesan, Manuela Pendziwiat, Annika Rademacher, Colin A Ellis, Nadja Hümpfer, Niklas Schwarz, Simone Seiffert, Joseph Peeden, Joseph Shen, Katalin Štěrbová, Trine Bjørg Hammer, Rikke S Møller, Deepali N Shinde, Sha Tang, Lacey Smith, Annapurna Poduri, Roland Krause, Felix Benninger, Katherine L Helbig, Volker Haucke, Yvonne G Weber, EuroEPINOMICS-RES Consortium; GRIN Consortium (Rudi Balling, Nina Barisic, Stéphanie Baulac, Hande Caglayan, Dana Craiu, Peter De Jonghe, Christel Depienne, Renzo Guerrini, Helle Hjalgrim, Dorota Hoffman-Zacharska, Johanna Jähn, Karl Martin Klein, Bobby P C Koeleman, Vladimir Komarek, Eric Leguern, Anna-Elina Lehesjoki, Johannes R Lemke, Holger Lerche, Tarja Linnankivi, Carla Marini, Patrick May, Hiltrud Muhle, Deb K Pal, Aarno Palotie, Felix Rosenow, Susanne Schubert-Bast, Kaja Selmer, Jose M Serratos, Sanjay Sisodiya, Ulrich Stephani, Pasquale Striano, Arvid Suls, Tiina Talvik, Sarah von Spiczak, Sarah Weckhuysen, Federico Zara, Paul Avillach, Anna Bartels, Sawona Biswas, Florence Bourgeois, Batsal Devkota, Tracy Glauser, Barbara Hallinan, Allison Heath, Joel Hirschhorn, Judson Kilbourn, Sek Won Kong, Ian Krantz, In-Hee Lee, Kenneth D Mandl, Eric Marsh, Kristen Sund, Deanne Taylor, Peter White). **A Recurrent Missense Variant in AP2M1 Impairs Clathrin-Mediated Endocytosis and Causes Developmental and Epileptic Encephalopathy.** *Am J Hum Genet.* 2019 Jun 6;104(6):1060-1072. doi: 10.1016/j.ajhg.2019.04.001. Epub 2019 May 16.

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**CA52.** Eggermann T, Elbracht M, Kurth I, Juul A, Johannsen TH, Netchine I, Mastorakos G, Johannsson G, Musholt TJ, Zenker M, Prawitt D, Pereira AM, Hiort O; European Reference Network on Rare Endocrine Conditions (ENDO-ERN) (Stefan Riedl, Birgit Rami-Merhar, Greisa Vila, Sabina Baumgartner-Parzner, Walter Bonfig, Claudine Heinrichs, Dominique Maiter, Inge Gies, Martine Cools, Kristina Casteels, Albert Beckers, Sabina Zacharieva, Violeta Iotova, Tomislav Jukic, Dario Rahelic, Vassos Neodeous, Leonidas Phylactou, Michal Krsek, Jan Lebl, Claus Gravholt, Anders Juul, Vallo Tillmann, Vallo Volke, Tapani Ebeling, Thierry Brue, Patrice Rodien, Jérôme Bertherat, Christine Poitou Bernert, Philippe Touraine, Philippe Chanson, Michel Polak, Maithe Tauber, Thomas Eggermann, Joachim Spranger, Dagmar Fuhrer, Thomas Danne, Olaf Hiort, Klaus Mohnike, Dirk Prawitt, Markus Luster, Nicole Reisch, Martin Reincke, Julia Rohayem, Martin Fassnacht, Miklós Tóth, Alessandra Cassio, Sonia Toni, Csilla Krausz, Barbara Piccini, Diego Ferone, Gianni Russo, Luca Persani, Annamaria Colao, Mariacarina Salerno, Marco Boscaro, Carla Scaroni, Ferruccio Santini, Giovanni Ceccarini, Ezio Ghigo, Iveta Dzivite-Krisane, Vita Rovite, Lauma Janozola, Rasa Verkauskienė, Michael Witsch, James Clark, Johannes Romijn, Thera Links, Nienke Biermasz, Sabine Hannema, Bas Havekes, Hedi Claahsen-van der Grinten, Henri Timmers, Robin Peeters, Gerlof Valk, A A Verrijn Stuart, Harm Haak, Eystein Husebye, Jens Bollerslev, Barbara Jarzab, Agnieszka 'Szybowska, João-Filipe Raposo, Dana Craiu, Doina Piciu, Ludmila Kostalova, Jarmila Vojtková, Tadej Battelino, Roque Cardona-Hernandez, Diego Yeste, Sonia Gaztambide, Anna Nordenström, Neil Gittoes, Trevor Cole, Elizabeth Crowne, Faisal Ahmed, Mohammed Didi, Marta Korbonits, Mehel Dattani, Peter Clayton, Justin Davies). **Genetic testing in inherited endocrine disorders: joint position paper of the European reference network on rare endocrine conditions (Endo-ERN).**

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Factor impact in 2020=3.612 (<https://www.scijournal.org/impact-factor-of-orphanet-j-rare-dis.shtml>)

**CA53.** Lal D, May P, Perez-Palma E, Samocha KE, Kosmicki JA, Robinson EB, Møller RS, Krause R, Nürnberg P, Weckhuysen S, De Jonghe P, Guerrini R, Niestroj LM, Du J, Marini C; EuroEPINOMICS-RES Consortium (Rudi Balling, Nina Barisic, Stéphanie Baulac, Hande Caglayan, Dana C Craiu, Peter De Jonghe, Christel Depienne, Renzo Guerrini, Ingo Helbig, Helle Hjalgrim, Dorota Hoffman-Zacharska, Johanna Jähn, Karl M Klein, Bobby P C Koeleman, Vladimir Komarek, Roland Krause, Eric Leguern, Anna-Elina Lehesjoki, Johannes R Lemke, Holger Lerche, Tarja Linnankivi, Carla Marini, Patrick May, Hiltrud Muhle, Deb K Pal, Aarno Palotie, Felix Rosenow, Susanne Schubert-Bast, Kaia Selmer, Jose M Serratos, Ulrich Stephani, Katalin Štěrbová, Pasquale Striano, Arvid Suls, Tina Talvik, Sarah von Spiczak, Yvonne G Weber, Sarah Weckhuysen, Federico Zara), Ware JS, Kurki M, Gormley P, Tang S, Wu S, Biskup S, Poduri A, Neubauer BA, Koeleman BPC, Helbig KL, Weber YG, Helbig I, Majithia AR, Palotie A, Daly MJ. **Gene family information facilitates variant interpretation and identification of disease-associated genes in neurodevelopmental disorders.** *Genome Med.* 2020 Mar 17;12(1):28. doi: 10.1186/s13073-020-00725-6.PMID: 32183904 **Free PMC article.**

Factor impact in 2020=10.506 (<https://www.scijournal.org/impact-factor-of-genome-med.shtml>)

**CA54.** Chatron N, Becker F, Morsy H, Schmidts M, Hardies K, Tuysuz B, Roselli S, Najafi M, Alkaya DU, Ashrafzadeh F, Nabil A, Omar T, Maroofian R, Karimiani EG, Hussien H, Kok F, Ramos L, Gunes N, Bilguvar K, Labalme A, Alix E, Sanlaville D, de Bellescize J, Poulat AL; EuroEpinomics-RES consortium AR working group (Ingo Helbig, Sarah von Spiczak, Stéphanie Baulac, Nina Barisic, Rudi Balling, Hande Caglayan, Dana Craiu, Renzo Guerrini, Karl Martin Klein, Carla Marini, Hiltrud Muhle, Felix Rosenow, Jose M Serratos, Katalin Štěrbová, Yvonne Weber), Moslemi AR, Lerche H, May P, Lesca G, Weckhuysen S, Tajsharghi H. **Bi-allelic GAD1 variants cause a neonatal onset syndromic developmental and epileptic encephalopathy.** *Brain.* 2020 May 1;143(5):1447-1461. doi: 10.1093/brain/awaa085.PMID: 32282878 **Free PMC article.**

Factor impact in 2020=10.750 (<https://www.scijournal.org/impact-factor-of-brain.shtml>)

**CA55.** Mann D, Antinew J, Knapp L, Almas M, Liu J, Scavone J, Yang R, Modequillo M, Makedonska I, Ortiz M, Kyrychenko A, Nordli D, Farkas V, Farkas MK; A0081042 study group (Leanid Shalkevich, Anna Jansen, Ivan Ivanov, Vania Nedkova, Fang Fang, Yi Wang, Jean-Marc Pinard, Ulrich Brandl, Dimitrios Zafeiriou, Anna Altmann, Marianne Berenyi, Monika Bessenyeyi, Andras Fogaras, Geza Szabo, Aviva Fattal-Valevski, Ki Joong Kim, Ahmad Beydoun, Ghassan Hmaimess, Nor Azni Yahaya, Marissa Barlaan-Lukban, Martha Bolanos, Jo Janette De la Calzada, Maria Estrella Ibe, Maria Antonia Aurora Valencia, Dana Craiu, Georgeta Diaconu, Tatiana Antonova, Elena Belousova, Yulia Karakulova, Olga Khaletskaya, Olga Lvova, Maria Strachunskaya, Ruzica Kravljanc, Dimitrije Nikolic, Francisco Lopez

Pison, Ying-Chao Chang, I-Ching Chou, Wang-Tso Lee, Charcrin Nabangchang, Oranee Sanmaneechai, Nihal Olgac Dundar, Pinar Gencpinar, Yurii Chomolyak, Dmytro Delva, Volodymyr Martyniuk, Ronald Davis, Jose Ferreira, Jerry Tomasovic). **Pregabalin adjunctive therapy for focal onset seizures in children 1 month to <4 years of age: A double-blind, placebo-controlled, video-electroencephalographic trial.** *Epilepsia.* 2020 Apr;61(4):617-626. doi: 10.1111/epi.16466. Epub 2020 Mar 18. PMID: 32189338 Clinical Trial.

Factor impact in 2020=6.549 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

**CA56.** van den Munckhof B, Arzimanoglou A, Perucca E, van Teeseling HC, Leijten FSS, Braun KPJ, Jansen FE; RESCUE ESES study group (Anna Jansen, Patrick van Bogaert, Lieven Lagae, Guido Rubboli, Eija Gaily, Pierangelo Veggiotti, Gaetano Cantalupo, Giuseppe Gobbi, Dana Craiu, Petia Dimova, Thomas Bast, Julia Jacobs, Sarah von Spiczak, Anja Lübbig, Stéphane Auvin, Anne de Saint-Martin, J Helen Cross, Richard Chin, Sameer Zuberi, Irene Garcia Morales, Georgia Ramantani). **Corticosteroids versus clobazam in epileptic encephalopathy with ESES: a European multicentre randomised controlled clinical trial (RESCUE ESES\*).** *Trials.* 2020 Nov 23;21(1):957. doi: 10.1186/s13063-020-04874-2. PMID: 33228736 **Free PMC article.**

Factor impact in 2020=2.063 (<https://www.scijournal.org/impact-factor-of-trials.shtml>)

**CA57.** Curatolo P, Jóźwiak S, Nabbout R; TSC Consensus Meeting for SEGA and Epilepsy Management (Paolo Curatolo, Sergiusz Jóźwiak, Rima Nabbout, Miraude Adriaensen, Moncef Berhouma, Giannennaro Coppola, **Dana Craiu**, Raffaella Cusmai, Olivier Delalande, Anne De Saint Martin, Pablo Hernáiz Driever, Martine Fohlen, Wiesława Grajkowska, Christoph Hertzberg, Anna Jansen, Floor Jansen, Katarzyna Kotulska, Marek Mandra, Romina Moavero, Finbar O'Callaghan, Emmanuel Raffo, Bernard A Zonnenberg). **Management of epilepsy associated with tuberous sclerosis complex (TSC): clinical recommendations.** *Eur J Paediatr Neurol.* 2012 Nov;16(6):582-6. doi: 10.1016/j.ejpn.2012.05.004. Epub 2012 Jun 12. PMID: 22695035 **Review.**

Factor impact in 2012=2.512 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>)

**CA58.** Brandsma R, Spits AH, Kuiper MJ, Lunsing RJ, Burger H, Kremer HP, Sival DA; Childhood Ataxia and Cerebellar Group (N Barisic, P Baxter, V Brankovic-Sreckovic, G E Calabrò, C Catsman-Berrevoets, Ifm de Co, D Craiu, B Dan, J Gburek-Augustat, F Kammoun-Feki, C Kennedy, F Mancini, M Mirabelli-Badenier, A Nemeth, R Newton, B T Poll-The, M Steinlin, M Synofzik, M Topcu, C Triki, E M Valente). **Ataxia rating scales are age-dependent in healthy children.** *Dev Med Child Neurol.* 2014 Jun;56(6):556-63. doi: 10.1111/dmnc.12369. Epub 2014 Jan 7. PMID: 24392880 **Free article.**

Factor impact in 2014=4.170 (<https://www.scijournal.org/impact-factor-of-dev-med-child-neurol.shtml>)

**CA59.** Santiago-Sim T, Burrage LC, Ebstein F, Tokita MJ, Miller M, Bi W, Braxton AA, Rosenfeld JA, Shahrour M, Lehmann A, Cogné B, Küry S, Besnard T, Isidor B, Bézieau S, Hazart I, Nagakura H, Immken LL, Littlejohn RO, Roeder E; EuroEPINOMICS RES Consortium Autosomal Recessive working group (Zaid Afawi, Rudi Balling, Nina Barisic, Stéphanie Baulac, **Dana Craiu**, Peter De Jonghe, Rosa Guerrero-Lopez, Renzo Guerrini, Ingo Helbig, Helle Hjalgrim, Johanna Jähn, Karl Martin Klein, Eric Leguern, Holger Lerche, Carla Marini, Hiltrud Muhle, Felix Rosenow, José Serratos, Katalin Sterbová, Arvid Suls, Rikke S Moller, Pasquale Striano, Yvonne Weber, Federico Zara), S. Hande Caglayan, Kara B, Hardies K, Weckhuysen S, May P, Lemke JR, Elpeleg O, Abu-Libdeh B, James KN, Silhavy JL, Issa MY, Zaki MS, Gleeson JG, Seavitt JR, Dickinson ME, Ljungberg MC, Wells S, Johnson SJ, Teboul L, Eng CM, Yang Y, Kloetzel PM, Heaney JD, Walkiewicz MA. **Biallelic Variants in OTUD6B Cause an Intellectual Disability Syndrome Associated with Seizures and Dysmorphic Features.** *Am J Hum Genet.* 2017 Apr 6;100(4):676-688. doi: 10.1016/j.ajhg.2017.03.001. Epub 2017 Mar 23. PMID: 28343629 **Free PMC article.**

Factor impact in 2017=9.358 (<https://www.scijournal.org/impact-factor-of-am-j-hum-genet.shtml>)

**CA60.** Nava C, Dalle C, Rastetter A, Striano P, de Kovel CG, Nabbout R, Cancès C, Ville D, Brilstra EH, Gobbi G, Raffo E, Bouteiller D, Marie Y, Trouillard O, Robbiano A, Keren B, Agher D, Roze E, Lesage S, Nicolas A, Brice A, Baulac M, Vogt C, El Hajj N, Schneider E, Suls A, Weckhuysen S, Gormley P, Lehesjoki AE, De Jonghe P, Helbig I, Baulac S, Zara F, Koeleman BP; EuroEPINOMICS RES Consortium (Rudi Balling, Nina Barisic, Stéphanie Baulac, Hande S Caglayan, **Dana C Craiu**, Peter De Jonghe, Christel Depienne, Padhraig Gormley, Renzo Guerrini, Ingo Helbig, Helle Hjalgrim, Dorota Hoffman-Zacharska, Johanna Jähn, Karl Martin Klein, Bobby P C Koeleman, Vladimir Komarek, Roland Krause, Eric LeGuern, Anna-Elina Lehesjoki, Johannes R Lemke, Holger Lerche, Carla Marini, Patrick May, Rikke S Møller, Hiltrud Muhle, Aarno Palotie, Deb Pal, Felix Rosenow, Kaja Selmer, José M Serratos, Sanjay Sisodiya, Ulrich Stephani, Katalin Sterbova, Pasquale Striano, Arvid Suls, Tiina Talvik, Sarah von Spiczak, Yvonne Weber, Sarah Weckhuysen, Federico Zara), Haaf T, LeGuern E, Depienne C. **De novo mutations in HCN1 cause early infantile epileptic encephalopathy.** *Nat Genet.* 2014 Jun;46(6):640-5. doi: 10.1038/ng.2952. Epub 2014 Apr 20. PMID: 24747641

Factor impact in 2014=28.317 (<https://www.scijournal.org/impact-factor-of-nat-genet.shtml>)

**CA61.** Epilepsy Phenome/Genome Project; Epi4K Consortium; EuroEPINOMICS-RES Consortium (Silke Appenzeller, Rudi Balling, Nina Barisic, Stéphanie Baulac, Hande Caglayan, **Dana Craiu**, Peter De Jonghe, Christel Depienne, Petia Dimova, Tania Djémié, Padhraig Gormley, Renzo Guerrini, Ingo Helbig, Helle Hjalgrim, Dorota Hoffman-Zacharska, Johanna Jähn, Karl Martin Klein, Bobby Koeleman, Vladimir Komarek, Roland Krause, Gregor Kuhlenbäumer, Eric Leguern, Anna-Elina Lehesjoki, Johannes R Lemke, Holger Lerche, Tarja Linnankivi, Carla Marini, Patrick May, Rikke S Møller, Hiltrud Muhle, Deb Pal, Aarno Palotie, Manuela Pendziwiat, Angela Robbiano, Filip Roelens, Felix Rosenow, Kaja Selmer, Jose M Serratos, Sanjay Sisodiya, Ulrich Stephani, Katalin Sterbova, Pasquale Striano, Arvid Suls, Tiina Talvik, Sarah von Spiczak, Yvonne Weber, Sarah Weckhuysen, Federico Zara, Bassel Abou-Khalil, Brian K Alldredge, Eva Andermann, Frederick Andermann, Dina Amrom, Jocelyn F Bautista, Samuel F Berkovic, Judith Bluvstein, Alex Boro, Gregory Cascino, Damian Consalvo, Patricia Crumrine, Orrin Devinsky, Dennis Dlugos, Michael P Epstein, Miguel Fiol, Nathan B Fountain, Jacqueline French, Daniel Friedman, Eric B Geller, Tracy Glauser, Simon Glynn, Kevin Haas, Sheryl R Haut, Jean Hayward, Sandra L Helmers, Sucheta Joshi, Andres Kanner, Heidi E Kirsch, Robert C Knowlton, Eric H Kossoff, Rachel Kuperman, Ruben Kuzniecky, Daniel H Lowenstein, Shannon M McGuire, Paul V Motika, Edward J Novotny, Ruth Ottman, Juliann M Paolicchi, Jack

Parent, Kristen Park, Annapurna Poduri, Lynette Sadleir, Ingrid E Scheffer, Renée A Shellhaas, Elliott Sherr, Jerry J Shih, Rani Singh, Joseph Sirven, Michael C Smith, Joe Sullivan, Liu Lin Thio, Anu Venkat, Eileen P G Vining, Gretchen K Von Allmen, Judith L Weisenberg, Peter Widess-Walsh, Melodie R Winawer, Andrew S Allen, Samuel F Berkovic, Patrick Cossette, Norman Delanty, Dennis Dlugos, Evan E Eichler, Michael P Epstein, Tracy Glauser, David B Goldstein, Yujun Han, Erin L Heinzen, Michael R Johnson, Ruben Kuzniecky, Daniel H Lowenstein, Anthony G Marson, Heather C Mefford, Sahar Esmaeeli Nieh, Terence J O'Brien, Ruth Ottman, Stephen Petrou, Slavé Petrovski, Annapurna Poduri, Elizabeth K Ruzzo, Ingrid E Scheffer, Elliott Sherr). **De Novo Mutations in Synaptic Transmission Genes Including DNM1 Cause Epileptic Encephalopathies.** *Am J Hum Genet.* 2017 Jan 5;100(1):179. doi: 10.1016/j.ajhg.2016.12.012.PMID: 28061363 **Free PMC article.**

Factor impact in 2017=9.358 (<https://www.scijournal.org/impact-factor-of-am-j-hum-genet.shtml>)

**CA62.** de Kovel CG, Brilstra EH, van Kempen MJ, Van't Slot R, Nijman IJ, Afawi Z, De Jonghe P, Djémié T, Guerrini R, Hardies K, Helbig I, Hendrickx R, Kanaan M, Kramer U, Lehesjoki AE, Lemke JR, Marini C, Mei D, Møller RS, Pendziwiat M, Stamberger H, Suls A, Weckhuysen S; EuroEPINOMICS RES Consortium (R Balling, N Barisic, S Baulac, H S Caglayan, D C Craiu, C Depienne, p Gormley, H Hjalgrim, D Hoffman-Zacharska, J Jähn, K M Klein, V Komarek<sup>1</sup>, E LeGuern<sup>1</sup>, H Lerche<sup>1</sup>, P May, H Muhle, D Pal<sup>1</sup>, A Palotie<sup>1</sup>, F Rosenow<sup>1</sup>, K Selmer<sup>1</sup>, J M Serratosa<sup>1</sup>, S M Sisodiya<sup>1</sup>, U Stephani<sup>1</sup>, K Sterbova<sup>1</sup>, P Striano<sup>1</sup>, T Talvik<sup>1</sup>, M van Haelst<sup>1</sup>, N Verbeek<sup>1</sup>, S von Spiczak<sup>1</sup>, Y G Weber<sup>1</sup>), Koeleman BP. **Targeted sequencing of 351 candidate genes for epileptic encephalopathy in a large cohort of patients.** *Mol Genet Genomic Med.* 2016 Jul 30;4(5):568-80. doi: 10.1002/mgg3.235. eCollection 2016 Sep.PMID: 27652284 **Free PMC article.**

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## PARTICIPARI LA CONFERINTE CU LUCRARE

1. À XXXI-A Conferința Națională a Societății Române Impotriva Epilepsiei, București, 22-25 noiembrie 2023
2. Școala de Vara de Epilepsie, Sinaia, 19-21 mai 2023
3. À XXX-A Conferința Națională a Societății Române Impotriva Epilepsiei, București, 23-26 noiembrie 2022
4. Școala de Vara de Epilepsie, Slanic-Ph, 1-3 septembrie, 2022
5. 34th International Epilepsy Congress (IEC 2021 virtual) 28 Aug-1 Sep 2021
6. À XXIX-A Conferința Națională a Societății Române Impotriva Epilepsiei, București, 25-27 noiembrie 2021
7. À XXVIII-A Conferința Națională a Societății Române Impotriva Epilepsiei, București, 4-7 noiembrie 2020
8. Conferința Națională de Pediatrie, București, 2020
9. Școala Neuro-MIG, Ediția 3, București, 5-6 martie 2020
10. Conferința Excelența în pediatrie, Copenhaga, 5-7 decembrie 2019
11. Conferința Psihiatria românească între știință și practică, București, 20-23 noiembrie 2019
12. À XXVII-A Conferința Națională a Societății Române Impotriva Epilepsiei, București, 14-16 noiembrie 2019

13. Conferinta Nationala de Medicina de Familie, Bucuresti, 23-26 octombrie 2019
14. A IV-a Conferinta Nationala a Societatii Romane de Neurologie Pediatrica, Bucuresti, 3-5 oct 2019
15. Masterclass de epilepsii, Bucuresti, 2 octombrie 2019
16. 9th Migrating Course on Epilepsy, 19-22 septembrie 2019, Vrdnik, Serbia
17. 13th EPNS, 17-21 septembrie 2019
18. Al 14 lea congres National de PEDIATRIE, Cluj Napoca, 11-14 septembrie 2019
19. Scoala de vara de epilepsie, Sucevita, Suceava, 11-13 iulie 2019
20. 33rd International Epilepsy Congress, Bamgkok, Thailanda, 22-26 iunie 2019
21. Adriatic Neurology Forum, Puglia, Italia, 23-26 mai 2019
22. Conferinta Internationala a Studentilor, Bucuresti, 11-14 aprilie 2019
23. Conferinta Nationala de Pediatrie, Bucuresti, 3-6 Aprilie 2019
24. 10th Anniversary of the European SEEG Course, Venice, Italia, 12-16 februarie 2019
25. Simpozion Aniversar la 50 ani de existenta a serviciului de recuperare copii, Centrul National Clinic de Recuperare Neuropsihomotorie Copii Dr N. Robanescu, Bucuresti, 13 decembrie 2018
26. A XXVI-A Conferinta Nationala a Societatii Roamane Impotriva Epilepsiei, Bucuresti, 4-6 octombrie 2018
27. 19th International Symposium on Severe Infantile Epilepsies Old and New Rreatments, Roma, 20-22 septembrie 2018
28. Universitatea de vara a SAMF, Brasov, 2-6 sept 2018
29. 13th European Congress on Epileptology, Viena, 26-30 august 2018
30. Scoala de vara de Dermatologie, Bucuresti, 24-26 august 2018
31. Scoala de vara de epilepsie SRIE, Sucevita, Suceava, 11-13 iulie 2018
32. Curs National URGEMED X, Bucuresti, 14-15 iunie 2018
33. IV East Course of Epilepsy, Chernihiv, Ucraina, 13-15 iunie 2018
34. Forum Perinatologia - Abordare Multidisciplinara in Perinatologie, Bucuresti, 24 martie 2018
35. A XXV-A Conferinta Nationala a Societatii Romane Impotriva Epilepsiei, Bucuresti, 14-18 noiembrie 2017
36. Forum Pediatric, Bucuresti, 9-10 noiembrie 2017
37. Simpozion Zilele Synevo, Bucuresti, 9 noiembrie 2017
38. A III-a Conferinta Nationala a Societatii Romane de Neurologie Pediatrica, Bucuresti, 26-28 oct 2017
39. Al XXI -lea Simpozion National de PsihoNeuroendocrinologie, Iasi, 4-6 oct 2017
40. Scoala de vara Noi Orizonturi in Medicina, Bucuresti, 27-30 iunie 2017
41. Simpozion Impreuna pentru viata - o sansa la normalitate, Bucuresti, 17 iunie 2017
42. Al 13 lea Congres National al Societatii de Pediatrie, Bucuresti, 7-10 iunie 2017
43. EPICARE, Bucuresti, 3 iunie 2017
44. Congresul UMF, Bucuresti, 29-31 mai 2017
45. Congresul de Epilepsie, Grecia, 26-28 mai 2017
46. Conferinta Internationala a Studentilor, Bucuresti, 30 martie - 2 aprilie 2017
47. Simpozion Duchenne Expert Academy, Bucuresti, 31 martie 2017
48. Workshop de boli neuromusculare, Bucuresti, 21-22 martie 2017
49. EPNS Research Meeting, Essen, Germania, 28-29 octombrie 2016
50. Conferinta Nationala Interdisciplinata, Cum Diagnosticam si cum tratam bolile renourinare la copil, Bucuresti, 20-22 octombrie 2016
51. 12th European Congress on Epileptology, Praga, 11-15 septembrie 2016
52. Cursul European de Epilepsie, Cheile Gradistei, 15-17 iunie 2016
53. Congresul UMF, Bucuresti, 2-4 iunie 2016
54. U-Task European Taskforce Childhood Epilepsy Surgery, Antiparos, Grecia, 23-27 mai 2016
55. Conferinta Internationala a Studentilor, Bucuresti, 14-17 aprilie 2016
56. A XXIII-A Conferinta Nationala a Societatii Romane Impotriva Epilepsiei, Bucuresti, 19-21 noiembrie 2015
57. Al XVI lea Congres SNPCAR, Sibiu, 23-26 septembrie 2015
58. Conferinta Nationala a Societatii de Pediatrie din Rep Moldova, Chisinau, 8-9 iunie 2015
59. Conferinta Nationala de Pediatrie 1-2 aprilie 2015
60. Conferinta Regionala de Medicina Familiei, Bucuresti, 11-13 dec 2014
61. A IX Conferinta Nationala de Scleroza Multipla, Iasi, 23-24 octombrie 2014
62. EPNS Research Meeting, Bucuresti, 12-13 septembrie 2014

**INTERES SPECIAL:** Epilepsie, chirurgia epilepsiei, genetica epilepsiei, Craiu D, Rener Primec Z, Lagae L, Vigevano F, Trinko E, Specchio N, Bakhtadze S, Cazacu C, Golli T, Zuberi SM. Vaccination and childhood epilepsies. Eur J Paediatr Neurol. 2022 Jan;36:57-68. doi: 10.1016/j.ejpn.2021.11.014. Epub 2021 Dec 3. PMID: 34922162. EEG, scleroza tuberoasa.

ANEXE: Copii diplome si atestate

1. Diploma Medic Specialist Neurologie Pediatria
2. Diploma Medic Primar Neurologie Pediatria
3. Diploma Medic Specialist Pediatru
4. Atestat Competenta Neurofiziologie (EEG, EMG, PEC)
5. Diploma de doctor in medicina
6. Diploma absolvire curs de Management Sanitar











18.12.2023