

INFORMAȚII PERSONALE



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Sexul feminin | Naționalitatea Romana

Dana Cristina CRAIU

LOCUL DE MUNCA

1. UMF Carol Davila, Departamentul 6 – Neuroștiinte Clinice, Disciplina Neurologie Pediatrica II
2. Clinica Neurologie Pediatrica, Centru de Referinta de Boli Rare Neurologice Pediatrica, Spital Clinic Prof. Dr. Alexandru Obregia, membru in rețelele Europene de Boli Rare EpiCARE si ENDO-ERN

EXPERIENȚA PROFESIONALĂ

MEDICALA

2014 - 2021  
 2000 - 2014  
 1994 - 1999

Medic primar Neurologie Pediatrica  
 Medic specialist Neurologie Pediatrica  
 Medic rezident Neurologie Pediatrica

Spital Clinic de Psihiatrie Prof. Dr Al. Obregia, Sos. Berceni 10, sect 4, Bucuresti (<https://spital-obregia.ro/>; <https://bolirare-obregia.ro/>)

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

1993 - 1994

Medic Stagiar

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

- Ingrijire pacienti de toate varstele

ACADEMICA

2014 - 2021  
 2011 - 2014  
 2004 - 2011  
 1999 - 2004

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

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

- Activitati de predare studenti, rezidenti, cadre medicale (cursuri postuniversitare)
- Activitati cercetare clinica, genetica neurologica, neurofiziologie clinica

CONDUCERE

2021 – prez  
 2020 – prez  
 2018 – prez  
 2016 – prez  
  
 2015 – prez  
 2009 – 2017  
 2009 – 2017  
 2008 – prez  
  
 2008 - 2016

Chair al Comisiei de Ghiduri a EPNS (Soc Europeana de Neurologie Pediatrica)  
 Prodecan Cercetare UMF Carol Davila – Facultatea de Medicina  
 Presedinte al Comisiei de Neurologie Pediatrica a Ministerului Sanatatii  
 Coordonator Centru de Expertiza Boli Rare Neurologie Pediatrica Obregia, membru EpiCARE si ENDO-ERN  
  
 Presedinte SRIE (Societatea Romana Impotriva Epilepsiei, Chapter al ILAE)  
 Presedinte CNA (Comisia de Advisori Nationali) a EPNS  
 Presedinte TAB (Training Advisory Board) a EPNS  
 Sef Disciplina Neurologie Pediatrica II – Departament Neuroștiinte Clinice, UMF Carol Davila – Facultatea de Medicina  
 Sef Clinica Neurologie Pediatrica Spital Al. Obregia

## EDUCAȚIE ȘI FORMARE

2012	Medic Specialist Pediater (examen Martie 2011, OMS 740/2012)	Scrieți nivelul EQF, dacă îl cunoașteți
2005	Doctor în Medicină (sustinere Nov 2004, confirmat Ordin MEC 3956/24.04.2005)	
	Medic Primar Neurolog Pediater (examen iun 2004, confirmat Ordin MS 1067/2004)	
2004	Competența Neurofiziologie (EEG, EMG, PEC) – sesiune examen aprilie 2001 Diploma cu Seria A, Nr 1056/ 2001	
2001	Medic Specialist Neurolog Pediater (Confirmat OMS 900/1999) Dobândire titlu Doctor Medic, Specializarea Medicină (Diploma Seria L. Nr. 426) în urma absolvirii Facultății de MG (1987-1993)	
1999	UMF Carol Davila București, Str Dionisie Lupu Nr. 37, sector 2, București ( <a href="https://umfcd.ro/">https://umfcd.ro/</a> )	
1993	▪ Abilități teoretice și practice, conform curriculumelor de pregătire	

## Cursuri educaționale la care am participat

- Curs Training VNS – 2021 – 1 zi
- Cursuri SEEG – Venetia – câte 1 săptămână/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 în epilepsii - 2008 (educație la distanță – EUREPA)
- Curs: EEG Basic - 2007 (educație la distanță- EUREPA)
- San-Servolo școala de vară: Copilul cu epilepsie – 07.2007 (2 săptămâni)
- San-Servolo: școala de vară; Epilepsii remediable chirurgical – 07.2006 (2 săptămâni)
- Curs: Epilepsii genetice – 2005 (educație la distanță - EUREPA)
- Breakfast seminarii în congresele de epilepsii – annual
- Curs: 'Explorări electrice în Neurologie' - EEG, EMG, EP, 2001
- 3 luni Bursa educațională – EEG pediatric, video-EEG, evaluare prechirurgicală în Stichting Epilepsie Instellingen, Olanda, sub direcția îndrumare a Dr. Walter van Emde Boas, (sept - dec 2000)
- Curs: 'Progres în Tratatamentul Epilepsiilor', 1999 (București)
- Curs: 'Explorări electrice în Neurologie' - EEG, EMG, EP, 1997
- Curs: Progres în Tratatamentul Epilepsiilor', 1996 (București)

## COMPETENȚE PERSONALE

## Limba maternă

Romana

## Alte limbi străine cunoscute

Engleza

Germana

	INTELEGERE		VORBIRE		SCRIERE
	Ascultare	Citire	Participare la conversație	Discurs oral	
Engleza	C1	C1	C1	C1	C1
Germana	A1	A1	A1	A1	A1

## Competențe organizaționale/manageriale

1. Program de studii pentru obținerea atestatului „Managementul serviciilor de sănătate” (14.03 – 27.07.2016) – Școala Națională de Sănătate Publică, Management și Perfectionare în Domeniul Sanitar București (Seria At Nr 0285)
2. Curs de formare ca formator organizat de ILAE -EUREPA (Liga Internațională Impotriva Epilepsiei): Train the Trainees – 2008 – Malta – 1 săptămână

Bun organizator, cu spirit de echipă

Abilități de organizare cursuri, conferințe, programe de formare

Bun manager de proiecte de cercetare sau de dezvoltare activități și programe în 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 distanta si telemedicina: Zoom, Google Meets, Teams, DocBook
- folosire programe de baze de date

**INFORMATII SUPLIMENTARE**
**Alte afiliari**

- Membru Fondator SRNP (societatea Romana de Neurologie Pediatria) 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 EPNS din 2005
- Membru ICNA (International Child Neurology Association) din 2001
- Membru fondator RONEP (Fundatia Romana de Neurologie si Epileptologie) – 1998
- Membru EUREPA din 1998

**Cursuri educationale predate**
**Cursuri pentru studenti**

2000 - 2007, 2 cursuri din 10 pentru modulul de Neurologie Pediatrica:

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

2008-prezent

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

**Cursuri pentru rezidenți:**

- 2008-prezent: Epilepsii, Status epilepticus, Sindroame neurocutanate, EHIP, Hidrocefalii, Edem cerebral, HIC, EEG de baza – teoretic si practic,
- Journal club – saptamanal – prin rotatie cu colegii din colectiv
- Presentari cazuri – saptamanal impreuna cu colegii din colectiv prin rotatie

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

- Curs de EEG basic – 3 saptamani
- Curs de sindroame in epilepsie – 1 sapt
- Curs de tratament in epilepsie – 1 saptamana

**Cursuri optionale**

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

Trainer – Virepa BEEG (Basic EEG) curs de 3 saptamani – 2/an din 2014 pana in prezent

Trainer – Virepa PEEG (Pediatric EEG) curs de 3 saptamani – 1/an din 2018 pana in prezent

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

Tratatul: Afectiuni neuromusculare la sugar, copil si adolescent, sub redactia Sanda Magureanu, editura Amaltea 2004, a primit premiul Gheorghe Marinescu al Acadmiei Romane in 2006

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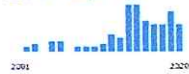
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Total Publications

100



Articles

18

Average citations per item

13,68

Sum of Times Cited

1.368

Without self citations

1.344

Citing articles

1.188

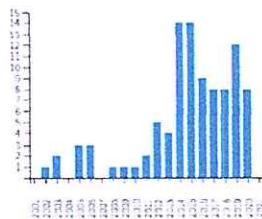
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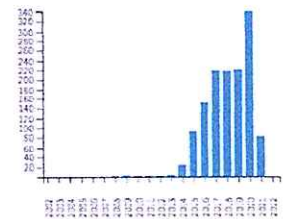
AUTHOR: (Craiu D)

Timespan=All years. Indexes=SCI-EXPANDED, SSCI, A&HCI, CPCI-S, CPCI-SSH, BKCI-S, BKCI-SSH, ESCI, CCR-EXPANDED, IC.

Total Publications by Year



Sum of Times Cited by Year



**ACTIVITATE DE CERCETARE:**

1. Grant internațional – director proiect					
1	POSCE-Axa II/ Op.02.2.4./ INFO ACT/ 2009-2011	Informatizarea activității administrative în 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-EUROCO/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 în consorțiul European EuroEPINOMICS –RES	Fonduri de la bugetul de stat, România fiind membru ESF (European Science Foundation) care a lansat competiția de proiecte.	378.000 RON	E, Depienne C, Balling R, Barisic N, Baulac S, Caglayan HS, Craiu DC, De Jonghe P, Depienne C, Gormley P, Guerrini R, 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, Serratosa 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.011093-23/MAR-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 Universitară „Carol Davila”, București, 2013.
4	Proiect tip FP7/ European Commission- Health Programme 2/ reference No. 534055/ 2013-2016	CEC 2013: Centrul de referință de chirurgie 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 finanțat 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.

2. Grant internațional – membru în echipa de cercetare					
1	POSCE axa II – Operațiunea O2.2.1.-2009-4/ SMIS 14042/ Contract 910 – 21.12.2012/ Perioada 2012-2015	Centrul de Cercetare Translațională în Psihiatrie și Neuroștiințe	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 (Îmbunătățirea cercetării științifice a autismului precoce: O rețea pentru îmbunătățirea cercetării, serviciilor și prognosticului)	Fonduri europene	Acopera integral costurile deplasărilor la toate întâlnirile 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 deplasărilor la toate întâlnirile proiectului	Duca G.D, <b>Craiu D</b> , Boer M, Chiriac S.M, Arghir A, Tutulan-Cunța 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 deplasărilor la toate întâlnirile proiectului	
5	CA16118/2017 – 2021	European Network on Brain Malformations Neuro-MIG	Fonduri europene	Acopera integral costurile deplasărilor la toate întâlnirile proiectului	


**3. Grant național – membru în echipa de cercetare**

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 infecție HIV/ SIDA	Buget de stat	200.000.000 lei	
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	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

		somn la copil			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 in domeniile prioritare PN II – Contract 42-130/2008 /2009-2011	Cercetari multidisciplinare – clinice, citogenetice si moleculare – in sindroamele de retard mental asociate cu anomalii congenitale: contributii in cunoasterea si managementul bolilor rare (RMBGR)	Buget de stat	200.000.000 lei	Craiu D, 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, Burlău 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, 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, Guerguelcheva V, Butoiianu 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, Jeannet 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-Berrepoets C, de Coo I, Craiu D, 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. <b>Ataxia rating scales are age-dependent in healthy children.</b> <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 - <b>NEUROLOGIE PEDIATRICA - Note de curs</b>, Editura Universitara "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. <b>Functional magnetic resonance imaging contribution to language areas assessment in children with non-lesion focal epilepsy</b>, <i>Romanian Journal of</i></p>

					Neurology, Volume IX, No. 3, 2010.  <b>Craiu D</b> , Avram P, Craiu M, Cochino AV, Minciú I, Tarta-Arsene O, Butcianu N, Burlóiu 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.
8	CEEX/ 2005-2008	Integrarea tehnicilor de analiza moleculara in diagnosticarea distrofinopatiilor in perspectiva unor strategii terapeutice si profilactice	Buget de stat	40.000 lei	<b>Craiu D</b> . DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitara „Carol Davila”, Bucuresti, 2013.

4. Studii clinice internaționale – investigații principale

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				Dana Cristina CRAIU
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				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, Elasziv 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, Fourmier-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 P, 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, Strzelecs 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, Ohmer Y, Arthur CP, Bala P, Gardner S, Gosalakal J, Prabhakar P, Abraham A, Adler L, 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, Forner SD, Fox E, Frandsen B, Fry JA, Fuller GR, Gaffney ME, Gasecki A, Gay C, Gelfand S, Giancarlo T, Gidlin 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 U, 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, Qaquadah P, Renfroe J, Richer R, Riesenberger R, Robbins L, Roberts KL, Rothner DA, Salem G, Samudrala S, Saper JR, Sarkis E, Saunders M, Schaerf 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.
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, Craiu D, Modequillo M, Nordli D, Farkas V, Farkas MK. Pregabalin as Adjunctive Treatment for Focal Onset Seizures in Pediatric Patients: A Randomized Controlled Trial. <i>J Child Neurol</i> . 2019 Apr;34(5):248-255. doi: 10.1177/0883073818821035. Epub 2019 Jan 27. PMID: 30688135 Clinical Trial. (2 ctan)
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, Kyrchenko A, Nordli D, Farkas V, Farkas MK; A0081042 study group (Leandri Shalkevich, Anna Jansen, Ivan Ivanov, Vania Nedkova, Fang Fang, Yi Wang, Jean-Marc Pnard, Ulrich Brandl, Dimitrios Zafeiriou, Anna Altmann, Marianne Berenyi, Monika Bessenyei, Andras Fogaras, Geza Szabo, Aviva Fattal-Valevski, Ki Joong Kim, Ahmad Beydoun, Ghassan Hmaimes, 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 Krvavjanac, Dimitrije Nikolic, Francisco Lopez Pison, Ying-Chao Chang, I-Ching Chou, Wang-Iso Lee, Charcrin Nabangchang, Oranee Sanmaneechai, Nihal Olga Dundar, Pinar Gencpinar, Yuri 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. <i>Epilepsia</i> . 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	Open-label, non-controlled, multicenter				



	(2013 – 2016)	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	EPO034 (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			<p>Groeneweg S, Peeters RP, Moran C, Stoupa A, Aurid 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, Marshande M, Lunsing RJ, Nowak S, den Uil CA, Zilikens 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 Merleir L, Krude H, Craiu D, 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. <i>Lancet Diabetes Endocrinol.</i> 2019 Sep;7(9):695-706. doi: 10.1016/S2213-8587(19)30155-X. Epub 2019 Jul 31. PMID: 31377265 Clinical Trial. (21citari)</p> <p>Groeneweg S, van Geest FS, Aboa A, Akantud 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, Craiu D, Crock P, DeGoede C, Demir K, Dica A, Dimitri P, Dolcetta-Capuzzo A, Dremmen MHG, Dubey R, Enderli A, Fairchild J, Galichan J, George B, Gevers EF, Hackenberg A, Halasz Z, Heinrich B, Huynh T, Klosowska 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, Meriq V, Lora FM, Moran C, Müller KE, Oliver-Petit I, Paone L, Paul PG, Polak M, Porta F, Poswar FO, Reinauer C, Rozenkova K, Menesse 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, Vandermiet J, van der Walt A, Wémeau JL, Wierzbka 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)</p>
14	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			<p>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, Stites T, Chen Y, Putzki N, Merschhemke M, Gärtner J; PARADIGMS Study Group (Andrew Kornberg, Barbara Bajzer-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, Tjaaf 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 Vaiciene-Magistris, Freddy Castro Farfan, Sandra Quinones, Barbara Steinborn, Barbara Ujma-Czapska, Mariusz Stasiulek, Miroslaw Jasinski, Dana Craiu, Alexey Boyko, Bateria Kairbekova, Farit Khabirov, Liudmila Kuzenkova, Nadezhda Malkova, Dimitrije Nilolic, Jasna Jancic, Ksenija Gebauer-Bukurov, Jaroslava Payerova, Francisco Gascon Jimenez, Guillermo Izquierdo Ayuso, Mar Mendibe Bilbao, Rogier Hintzen, Victoria Eugenia Fernandez Sanchez, Virginia Meca Lallana, Xavier Montalban Gairin, Karin Nordborg, Banu Anlar, Cergiz Yalçinkaya, Kivcim Gucuyener, Murat Terzi, Serkan Ozakbas, Unsal Yilmaz, Iryna Makedonska, Kateryna Prokopenko, Liudmyla Tanbura, 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, Kottli Rammoan, Michael Lloyd, Mitchel Williams, Ricardo Ayala, Ronald Davis, Vikram Bhishe). Trial of Fingolimod versus Interferon Beta-1a in Pediatric Multiple Sclerosis. <i>N Engl J Med.</i> 2018 Sep 13;379(11):1017-1027. doi: 10.1056/NEJMoa1800149. PMID: 30207920 Free article. Clinical Trial</p>
15	[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 β-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			<p>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, Mandra M, Moavero R, O'Callaghan F, Raffo E, Zonnenberg BA. Management of epilepsy associated with tuberous sclerosis complex (TSC): clinical recommendations. <i>Eur J Paediatr Neurol</i>. 2012 Nov;16(6):582-6. doi: 10.1016/j.ejpn.2012.05.004. Epub 2012 Jun 12.</p> <p>Jóźwiak S, Nabbout R, Curatolo P; participants of the TSC Consensus Meeting for SEGA and Epilepsy Management. Management of subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC): Clinical recommendations. <i>Eur J Paediatr Neurol</i>. 2013 Jul;17(4):348-52. doi: 10.1016/j.ejpn.2012.12.008. Epub 2013 Feb 5.</p> <p>Glushkova M, Bojinova V, Koleva M, Dimova P, Bojidarova 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)</p>

### 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 acestora			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, Guerguelcheva V, Butoianu N, Craiu D, Korngut L, Campbell C, Haberłova J, Strenkova J, Alejandro M, Jimenez A, Ortiz GG, Enriquez GV, Rodrigues M, Roxburgh R, Dawkins H, Youngs L, Lahdette 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. Mapping the differences in care for 5,000 spinal muscular atrophy patients, a survey of 24 national registries in North America, Australasia and Europe. <i>J Neurol</i> . 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, paraliziiilor 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")	Bu get ul de sta t. Pro iec te ex per im ent al- de mo nst rati ve - PN -III-	600. 000 lei	

			CE RC- CO - PE D- 20 16	
<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 Cartabil – 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 brosure informatice pentru familii si cursuri pentru parinti privind epilepsia 1999 – 2000;

## ARTICOLE – autor principal

### 11 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.yebbeh.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, Ilescu 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, Bărcă 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. Ilescu C, Tarța-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, Ilescu 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=2019=2.513 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>) nu este calculate inca pe 2020

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

**CA1.** 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 Koелеman, Vladimir Komarek, Roland Krause, Gregor Kuhlenbäumer, Eric Leguem, 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, Orin 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 Widdess-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. EuroEPINOMICS-RES Consortium; Epilepsy Phenome/Genome Project; Epi4K Consortium. **De Novo Mutations in Synaptic Transmission Genes Including DNM1 Cause Epileptic Encephalopathies**

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**CA47.** Larsen J, Carvill GL, Gardella E, Kluger G, Schmiedel G, Barisic N, Depienne C, Bristra E, Mang Y, Nielsen JE, Kirkpatrick M, Goudie D, Goldman R, Jahn JA, Jepsen B, Gill D, Döcker M, Biskup S, McMahon JM, Koeleman B, Harris M, Braun K, de Kovel CG, Marini C, Specchio N, Djémié T, Weckhuysen S, Tommerup N, Troncoso M, Troncoso L, Bevoit A, Wolf M, Hjalgrim H, Guerrini R, Scheffer IE, Mefford HC, Møller RS; EuroEPINOMICS RES Consortium CRP (Aarno Palotie, Anna-Elina Lehesjoki, Arvid Suls, Bobby Koeleman, Carla Marini, Christel Depienne, Dana Craiu, Deb Pal, Dorota Hoffman-Zacharska, Eric Leguerm, Federico Zara, Felix Rosenow, Hande Caglayan, Helle Hjalgrim, Hiltrud Muhle, Holger Lerche, Ingo Helbig, Johanna Jahn, Johannes Lemke, Jose M Serratosa, Kaja Selmer, Karl Martin Klein, Katalin Sterbova, Nina Barisic, Padhraig Gormley, Pasquale Striano, Patrick May, Peter De Jonghe, Renzo Guerrini, Rikke S Møller, Roland Krause, Rudi Balling, Sanjay Sisodiya, Sarah von Spiczak, Sarah Weckhuysen, Stéphanie Baulac, Tiina Talvik, Ulrich Stephani, Vladimir Komarek, Yvonne Weber). The phenotypic spectrum of SCN8A encephalopathy. Neurology. 2015 Feb 3;84(5):480-9. doi: 10.1212/WNL.0000000000001211. Epub 2015 Jan 7. PMID: 25568300 **Free PMC article.**

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**CA48.** 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 Leguerm), Helbig I, Striano P, Weckhuysen S, Berkovic SF, Scheffer IE, Mefford HC. Mutations in the GABA Transporter SLC6A1 Cause Epilepsy with Myoclonic-Astatic 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.**

Factor impact in 2015=12.090 (<https://www.scijournal.org/impact-factor-of-am-j-hum-genet.shtml>)

- CA49.** 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-Weedl 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  
Factor impact in 2016=5.901 (<https://www.scijournal.org/impact-factor-of-j-med-genet.shtml>)
- CA50.** 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>)
- CA51.** 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, Meisler M, Berkovic SF, De Jonghe P, Scheffer IE, Myers RM, Cooper GM, Mefford HC. **Aberrant Inclusion of a Poison Exon Causes Dravet Syndrome and Related SCN1A-Associated Genetic Epilepsies.** *Am J Hum Genet.* 2018 Dec 6;103(6):1022-1029. doi: 10.1016/j.ajhg.2018.10.023. PMID: 30526861 **Free PMC article.**  
Factor impact in 2018=10.192 (<https://www.scijournal.org/impact-factor-of-am-j-hum-genet.shtml>)
- CA52.** 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 Hauke, 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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- CA53.** Eggemann 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 Maier, Inge Gies, Martine Cools, Kristina Casteels, Albert Beckers, Sabina Zachariva, Violeta Iotova, Tomislav Jukic, Dario Rahelic, Vassos Neocleous, 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, Matthe Tauber, Thomas Eggemann, 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, Mariacarla 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 Biemasz, Sabine Hannema, Bas Havekes, Hedi Claahsen-van der Grinten, Henri Timmers, Robin Peeters, Gerlof Valk, AA 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, Mehul 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).** *Orphanet J Rare Dis.* 2020 Jun 8;15(1):144. doi: 10.1186/s13023-020-01420-w. PMID: 32513286 **Free PMC article.**  
Factor impact in 2020=3.612 (<https://www.scijournal.org/impact-factor-of-orphanet-j-rare-dis.shtml>)
- CA54.** 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 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>)
- CA55.** 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, Stephanie Baulac, Nina Barisic, Rudi Balling, Hande Caglayan, Dana Craiu, Renzo Guerrini, Karl Martin Klein, Carla Marini, Hiltrud Muhle, Felix Rosenow, Jose M Serratosa, Katalin Sterbova, 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>)

**CA56.** 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 Bessenyi, 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>)

**CA57.** van den Munkhof 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>)

**CA58.** Curatolo P, Jóźwiak S, Nabbout R; TSC Consensus Meeting for SEGA and Epilepsy Management (Paolo Curatolo, Sergiusz Jóźwiak, Rima Nabbout, Miraude Adriaensen, Moncef Berhouma, Giangennaro 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>)

**CA59.** 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/dmcn.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>)

**CA60.** Santiago-Sim T, Burrage LC, Ebstein F, Tokita MJ, Miller M, Bi W, Braxton AA, Rosenfeld JA, Shahrouh 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 Jahn, Karl Martin Klein, Eric Leguern, Holger Lerche, Carla Marini, Hiltrud Muhle, Felix Rosenow, José Serratosa, 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, Kloeitzel 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>)

**CA61.** 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, Suis 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 Jahn, 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 Serratosa, 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>)

**CA62.** 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 Jahn, 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 Serratosa, 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 Widdess-Walsh, Melodie R Vinawer, 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 Esmaeili 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>)

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## PARTICIPARI LA CONFERINTE CU LUCRARE

1. À XXVIII-A Conferinta Nationala a Societatii Romane Impotriva Epilepsiei, Bucuresti, 4-7 noiembrie 2020
2. Conferinta Nationala de Pediatrie, Bucuresti, 2020
3. Scoala Neuro-MIG, Editia 3, Bucuresti, 5-6 martie 2020
4. Conferinta Excelenta in pediatrie, Copenhaga, 5-7 decembrie 2019
5. Conferinta Psihiatria romaneasca intre stiinta si practica, Bucuresti, 20-23 noiembrie 2019
6. À XXVII-A Conferinta Nationala a Societatii Roamane Impotriva Epilepsiei, Bucuresti, 14-16 noiembrie 2019
7. Conferinta Nationala de Medicina de Familie, Bucuresti, 23-26 octombrie 2019
8. A IV-a Conferinta Nationala a Societatii Romane de Neurologie Pediatrica, Bucuresti, 3-5 oct 2019
9. Masterclass de epilepsii, Bucuresti, 2 octombrie 2019
10. 9th Migrating Course on Epilepsy, 19-22 septembrie 2019, Vrdnik, Serbia
11. 13th EPNS, 17-21 septembrie 2019
12. Al 14 lea congres National de PEDIATRIE, Cluj Napoca, 11-14 septembrie 2019
13. Scoala de vara de epilepsie, Sucevita, Suceava, 11-13 iulie 2019
14. 33rd International Epilepsy Congress, Bangkok, Thailanda, 22-26 iunie 2019
15. Adriatic Neurology Forum, Puglia, Italia, 23-26 mai 2019
16. Conferinta Internationala a Studentilor, Bucuresti, 11-14 aprilie 2019
17. Conferinta Nationala de Pediatrie, Bucuresti, 3-5 Aprilie 2019
18. 10th Anniversary of the European SEEG Course, Venice, Italia, 12-16 februarie 2019
19. 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
20. À XXVI-A Conferinta Nationala a Societatii Roamane Impotriva Epilepsiei, Bucuresti, 4-6 octombrie 2018

21. 19th International Symposium on Severe Infantile Epilepsies Old and New Treatments, Roma, 20-22 septembrie 2018
22. Universitatea de vara a SAMF, Brasov, 2-6 sept 2018
23. 13th European Congress on Epileptology, Viena, 26-30 august 2018
24. Scoala de vara de Dermatologie, Bucuresti, 24-26 august 2018
25. Scoala de vara de epilepsie SRIE, Sucevita, Suceava, 11-13 iulie 2018
26. Curs National URGEMED X, Bucuresti, 14-15 iunie 2018
27. IV East Course of Epilepsy, Chernihiv, Ucraina, 13-15 iunie 2018
28. Forum Perinatologia - Abordare Multidisciplinara in Perinatologie, Bucuresti, 24 martie 2018
29. A XXV-A Conferinta Nationala a Societatii Romane Impotriva Epilepsiei, Bucuresti, 14-18 noiembrie 2017
30. Forum Pediatric, Bucuresti, 9-10 noiembrie 2017
31. Simpozion Zilele Synevo, Bucuresti, 9 noiembrie 2017
32. A III-a Conferinta Nationala a Societatii Romane de Neurologie Pediatrica, Bucuresti, 26-28 oct 2017
33. Al XXI -lea Simpozion National de PsihoNeuroendocrinologie, Iasi, 4-6 oct 2017
34. Scoala de vara Noi Orizonturi in Medicina, Bucuresti, 27-30 iunie 2017
35. Simpozion Impreuna pentru viata - o sansa la normalitate, Bucuresti, 17 iunie 2017
36. Al 13 lea Congres National al Societatii de Pediatrie, Bucuresti, 7-10 iunie 2017
37. EPICARE, Bucuresti, 3 iunie 2017
38. Congresul UMF, Bucuresti, 29-31 mai 2017
39. Congresul de Epilepsie, Grecia, 26-28 mai 2017
40. Conferinta Internationala a Studentilor, Bucuresti, 30 martie - 2 aprilie 2017
41. Simpozion Duchenne Expert Academy, Bucuresti, 31 martie 2017
42. Workshop de boli neuromusculare, Bucuresti, 21-22 martie 2017
43. EPNS Research Meeting, Essen, Germania, 28-29 octombrie 2016
44. Conferinta Nationala Interdisciplinata, Cum Diagnosticam si cum tratam bolile renourinare la copii, Bucuresti, 20-22 octombrie 2016
45. 12th European Congress on Epileptology, Praga, 11-15 septembrie 2016
46. Cursul European de Epilepsie, Cheile Gradistei, 15-17 iunie 2016
47. Congresul UMF, Bucuresti, 2-4 iunie 2016
48. U-Task European Taskforce Childhood Epilepsy Surgery, Antiparos, Grecia, 23-27 mai 2016
49. Conferinta Internationala a Studentilor, Bucuresti, 14-17 aprilie 2016
50. A XXIII-A Conferinta Nationala a Societatii Romane Impotriva Epilepsiei, Bucuresti, 19-21 noiembrie 2015
51. Al XVI lea Congres SNPCAR, Sibiu, 23-26 septembrie 2015
52. Conferinta Nationala a Societatii de Pediatrie din Rep Moldova, Chisinau, 8-9 iunie 2015
53. Conferinta Nationala de Pediatrie 1-2 aprilie 2015
54. Conferinta Regionala de Medicina Familiei, Bucuresti, 11-13 dec 2014
55. A IX Conferinta Nationala de Scleroza Multipla, Iasi, 23-24 octombrie 2014
56. EPNS Research Meeting, Bucuresti, 12-13 septembrie 2014

**INTERES SPECIAL:** Epilepsie, chirurgia epilepsiei, EEG, scleroza tuberoasa.

ANEXE: Copii diplome si atestate

1. Diploma Medic Specialist Neurologie Pediatrica
2. Diploma Medic Primar Neurologie Pediatrica
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



**ROMÂNIA**  
**MINISTERUL SĂNĂTĂȚII**

**CERTIFICAT**  
**DE MEDIC SPECIALIST**

Se certifică prin prezentul că:

**CRAIU P. DANA CRISTINA**

este confirmată) **MEDIC** specialist în specialitatea **NEUROLOGIE PEDIATRICA**  
 prin Ordinul Ministrului Sănătății nr. **908** din **1999**  
 pe baza examenului susținut în sesiunea **OCTOMBRIE 1999** și promovat cu media generală **9,79**  
 Eliberat la data de **16.12.2016** cu numărul **31629**

Prezentul certificat s-a eliberat în conformitate cu prevederile legale.

**DIRECTOR GENERAL,**  
**PETRU ȚAGOREAN**

**CONSILIER,**  
**GABRIELA CHELBA**

TS

Seria S1 Nr. 032313

**ROMÂNIA**  
**MINISTERUL SĂNĂTĂȚII**

**CERTIFICAT**  
**DE MEDIC PRIMAR**

Se certifică prin prezentul că:

**CRAIU DANA CRISTINA**

este confirmat(ă) **MEDIC** primar în specialitatea **NEUROLOGIE PEDIATRICĂ**  
 prin Ordinul Ministrului Sănătății nr. **1067** din **2004**  
 pe baza examenului susținut în sesiunea **IUNIE 2004** și promovat cu media generală **.10,00**  
 Eliberat la data de **.09.09.2016** cu numărul **12931**

Prezentul certificat s-a eliberat în conformitate cu prevederile legale.

**DIRECTOR GENERAL,**  
**IONUȚ SEBASTIAN IAVOR**

**CONSILIER,**  
**IOANA MIHALEA**

TS

Seria P1 Nr. 014007

