

INFORMAȚII PERSONALE

Dana Cristina CRAIU



LOCUL DE MUNCĂ

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 Medic primar Neurologie Pediatrică
2000 - 2014 Medic specialist Neurologie Pediatrică
1994 - 1999 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/>)
- Îngrijire pacienți copii cu afecțiuni neurologice pediatrică, predominant epilepsii, alte boli rare în domeniu, evaluare prochiurgicală pentru chirurgia epilepsiei, interpretare EEG, video-EEG
 - Detaliere stagii în cadrul rezidentiatului: Spitalul de Pediatrie, V. Gomoiu (1 an - pediatrie); Psihiatrie Infanțilă, 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 Național pentru Sănătatea Mamei și Copilului – Spital Polizu, Str. Gh. Polizu nr. 38-54, Sector 1, București (<https://www.insmc.ro/institutii/componenta-cp-polizu/>)
- Îngrijire pacienți de toate vârstele

ACADEMICA

- 2014 - 2022 Prof. Univ. Neurologie Pediatrică
2011 - 2014 Conf. Univ. Neurologie Pediatrică
2004 - 2011 Șef Lucrări Neurologie Pediatrică
1999 - 2004 As. Univ. per det Neurologie Pediatrică
UMF Carol Davila București, Str. Dionisie Lupu Nr. 37, sector 2, București (<https://umfcd.ro/>)
- Activități de predare studenți, rezidenți, cadre medicale (cursuri postuniversitare)
 - Activități cercetare clinică, genetică neurologică, neurofiziologie clinică

CONDUCERE

- 2022 - 2026 Chair a Comisiei de Educație și training al EPNS (Soc. Europeană de Neurologie Pediatrică)
2021 - 2026 Chair al Comisiei de Ghiduri a EPNS
2020 - prez Prodecan Cercetare UMF Carol Davila – Facultatea de Medicină
2018 - prez Presedinte al Comisiei de Neurologie Pediatrică a Ministerului Sănătății
2016 - prez Coordonator Centru de Expertiză Boli Rare Neurologie Pediatrică Obregia, membru EpiCARE și ENDO-ERN
- 2015 - prez Presedinte SRIE (Societatea Română Impotriva Epilepsiei, Chapter al ILAE)
2009 - 2017 Presedinte CNA (Comisia de Advisorii Naționali) a EPNS
2009 - 2017 Presedinte TAB (Training Advisory Board) a EPNS
2008 - prez Șef Disciplina Neurologie Pediatrică II – Departament Neuroștiințe Clinice, UMF Carol Davila – Facultatea de Medicină
2008 - 2016 Șef Clinica Neurologie Pediatrică Spital Al. Obregia

Alte competențe 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 studenți

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 si 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 si practic,
- Journal club – saptamanal – prin rotatie cu colegii din colectiv
- Prezentrari 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

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

ACTIVITATE DE CERCETARE:

1. Grant international – director proiect

1	POSCE-Axa II/ Op.02.2.4/ INFO ACT/ 2009-2011	Informizarea 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: CA1,3-5, 8, 11, 12, 16-20, 23-25, 28, 30, 34, 40, 43, 47-52, 54, 55, 60-65
2	6-EUROCORE/06.05.2011 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 consorziul European EuroEPIDEMICS –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, Barise N, Baulic 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, Koelman BP, Komaruk 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. De novo mutations in HCN1 cause early infantile epileptic encephalopathy. <i>Nat Genet.</i> 2014 Jun;46(6):640-5. doi: 10.1038/ng.2952. Epub 2014 Apr 20.
3	IBISD/ GEE006.10/ No.01.1093-23MAR-00-2012-2014 - terminat	Search of biomarkers for diagnosis, monitoring of disease and therapeutic response in Duchenne's muscular dystrophy	Genethon Franța	28.000 euro	Craiu D. DYSTROFIOPATIILE - 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, Craiu D. Diagnostic approach of Epilepsy in Childhood and Adolescence, <i>Maedica - A Journal of Clinical Medicine</i> , Volume 8, No. 2, 2013.

2. Grant international – membru in 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 Zeay B, Pineda M, Behi-Buisson N, Hryniewiecka-Jaworska A, Bienvenu T, Armstrong J, Roche-Martinez A, Mari F, Veneselli E, Russo S, Vignoli A, Pini G, Djuric M, Blagaard AM, Mejaki Bošnjak V, Polgar N, Cogliati F, Ravn K, Pintaudi M, Melegh B, Craiu D, 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	Duco G.D, Craiu D, Boer M, Chiriac S.M, Arghir A, Tutulan-Cunika 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 in 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 pacienți cu	Buget de stat	200.000.000 lei	

					Dana Tarta-Arsene, Florin Preoteasa, Sanda Adriana Magureanu, Adrian Ilescu, 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. Craiu D, Avram P, Craiu M, Cochino AJ, Minciul, Tarta-Arsene D, Butoiu N, Burloiu C, Ilescu C, Magureanu S, Moosles 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	Craiu D. DISTROFINOPATIILE - Notiuni teoretice, Algoritmi de diagnostic si tratament. Editura Universitara „Carol Davila”, Bucuresti, 2013.

4. Studii clinice internaționale – investigator principal

1	1042-0500/2005-004285-13/2007-2008	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			Ho TW, Pearlman E, Lewis D, Hämläinen 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, Tennelkoon G, Hsu DT, Nassogne MC, Sekhara T, Almadani M, O'Mahony M, Richer L, Illum N, Laugaard-Jacobsen HC, Sander V, Tahrik I, Kallala M, Keski-Santti P, Kuikkonen J, Nikkanen E, Nissila M, Partinen M, Pellola J, Annequin D, Cuvelier JC, Fourmier-Charriere E, Laborde S, Mili M, Navez ML, Parain D, Suc A, Ebinger F, Evers S, Gaul C, Gendolla A, Jansen JP, Laengler K, Pothmann R, Schellenberg R, Agorwot S, Anand I, Chandran S, Chodhary VB, Harawat PJ, Harsha S, Jog P, Kannan A, Karardan U, Keerthi AS, Neillikunja S, Pandit L, Srinivasa R, Barbanti P, Jegere D, Strautmanis J, Mulleners W, Pop P, Van den Berg P, Bryn B, Kjaemli T, Sommerfeldt K, Kochanowska J, Pietrak M, Strzelecks J, Szatani K M, Wesolowska M, Benga I, Craiu DC, Diaconu G, Gheonea C, Popescu L, Artigas Pallares J, Blanco Banca O, Campistol Plana J, Macaya A, Mosquera Villaverde Mdcl C, Reyes Martin A, San Antonio MV, Danielsson B, Ohmer Y, Arthur CP, Bala P, Gardner S, Gosalakkal 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 RR, Berman GD, Bernstein A, Bhaba 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, Fedman M, Ferriando M, Fieve RR, Fisher M, Ritman 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, Laocy D, Lane P, Lebron D, Lesh K, Ley J, Linder S, Liu E, Luber SR, Machanic B, Marcadis I, Markely HG, Markovitz PJ, Mate UJ, Mathew NT, McAllister P, McGettigan JW, Means P, Mechtler L, Mehra V, Melamed I, Miller DC, Miller JL, Miranda F, Moon M, Mubar IM, Munoz S, Murphy K, Naravaty 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, Riesenber R, Robbins L, Roberts KL, Rothner DA, Salem G, Samudrala S, Saper JR, Sarkis E, Saunders M, Schaeff F, Schreiber AQ, Sebastian V, Sedill A, Silas PE, Silverboard G, Sivakumar K, Smietana S, Snell P, Sperling M, Slegel C, Spierings E, Stedman M, Stepp W, Szrinczek 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, Wyssomierski DA, Zinn MM. Efficacy and tolerability of rizatriptan in pediatric migraineurs: results from a randomized, double-blind, placebo-controlled trial using a novel adaptive enrichment design. <i>Cephalalgia</i>. 2012 Jul;32(10):750-65. Epub 2012 Jun 18.
4	B4Z-EW-8013/ 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			

				Paone L, Paul PG, Polak M, Porti F, Prosser RO, Reinauer C, Rozenkova K, Meneses TS, Simm R, Simon A, Singh Y, Spada N, van der Spek J, Stab WMA, Stoupa A, Subramanian GM, Tencat D, Tiran S, den Uijl CA, Vandermiet J, van der Wilt A, Vávrova J, Wierzbicka J, de Wit MY, Wolf NI, Wurm M, Zibordi F, Zung A, Zwaeninger-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 citare)
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 pediatric patients with multiple sclerosis		Chitris T, Arnold DL, Barwell B, Brück W, Sheng A, Giovannoni G, Greenberg B, Knapp L, Rosales K, Tardieu M, Waubant E, Wolinsky JS, Ben-Or A, Siles T, Chen Y, Putzi N, Menschikow M, Gärtner J, PARADIGMS Study Group (Andrew Kornberg, Barbara Bajen-Kornek, Sergey Lkhachev, Antonio Pereira Gomes Neto, Denise Diniz, José Pat, Rogina Alvariega, Verónica Bujnowska-Ichamova, Jean Mah, Sunita Venkateswaran, Krasanka Hufner, Kad in Gross-Paju, Bruno Brochot, Emmanuel Cheuret, François Rivier, Kumaran Deiva, Mathieu Mili, Astrid Blaschek, Regina Trollmann, Rudolf Korinthenberg, Thomas Luecke, Tjalf Ziemssen, Carlo Pozzilli, Francesco Ratti, Giancarlo Comi, Girolamo Alessandra Marife, Luigi Maria Edoardo Grimaldi, Maria Trojano, Mauro Zaffaroni, Ruggero Capra, Vincenzo Brescia Morra, Gunter Roetzheim, Juana Launay-Belene, Nerja Vaidiene-Magistris, Freddy Castro Rattan, Sandra Quinones, Barbara Steinborn, Barbara Ujma-Czapska, Mariusz Stasiulek, Mirosław Jasinski, Dana Craiu, Newey Boyle, Ekaterina Kairbekova, Rati Kheibirov, Ludmila Kuzenova, Nadezhda Malkova, Dimitria Nikolai, Jasna Janovic, Ksenia Gebeuer-Bulurov, Jaroslava Payerova, Francisco Gaxton Jimenez, Guillermo Izquierdo Ayuso, Mar Mendíbe Bilbao, Roger Hinzen, Victoria Eugenia Fernandez Sanchez, Virginia Meca-Lallana, Xavier Montalban Gamin, Karin Nordborg, Banu Ankar, Genge Yalonkaya, Kivdim Gucuyener, Murat Terzi, Serkan Ozakbas, Ursul Yilmaz, Inna Makedonska, Katerina Prokopenko, Ludmyla Tantsura, Sergii Moskova, Tetiana Kobay, Tetiana Muratova, Tetiana Nelych, Tetiana Prykhodko, Cheryl Hemingway, Evangeline Wasmer, Jay Shetty, Jay Desai, Amy Waldman, Angel Chinea Martinez, Jayne Ness, Kottli Rammoher, Michael Lloyd, Mitchell Williams, Ricardo Ayala, Ronald Davis, Vikram Bhasi). <i>Trial of Fingolimod versus Interferon Beta-1a in Pediatric Multiple Sclerosis.</i> <i>N Engl J Med.</i> 2018 Sep 13;379(11):1017-1027. doi: 10.1056/NEJMoa1802149. PMID: 30257920. Free article, 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 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-CRAD001M1C03/2013-2018	An international disease registry collecting data on manifestations, interventions and outcomes in patients with tuberous sclerosis complex- TOSCA		Cusato P, Jozwiak S, Nabout R. TSC Consensus Meeting for SEGAs and Epilepsy Management. <i>Adriaansens M, Berhouma M, Coppola G, Craiu D, Cusato P, Delellande C, De Saint Martin A, Driever PH, Fohlen M, Gajkowska W, Hertzberg C, Jansen A, Jansen S, Kotulska K, Mundera M, Moavero R, O'Callaghan E, Rajfo E, Zonnenberg BA. Management of epilepsy associated with tuberous sclerosis complex (TSC): clinical recommendations.</i> <i>Eur J Paediatr Neurol.</i> 2012 Nov;16(5):582-6. doi: 10.1016/j.ejpn.2012.05.004. Epub 2012 Jun 12. Jozwiak S, Nabout R, Cusato P. participants of the TSC Consensus Meeting for SEGAs and Epilepsy Management. <i>Management of subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC): clinical recommendations.</i> <i>Eur J Paediatr Neurol.</i> 2013 Jul;17(4):348-52. doi: 10.1016/j.ejpn.2012.12.008. Epub 2013 Feb 5. Gajkowska M, Bojnowska V, Koleva M, Dimova P, Bojnowska M, Litvinenko I, Todorov T, Ilkova E, Galusaru C, Nasag E, Craiu D, Mihov V, Todorova A. Molecular genetic diagnosis of tuberous sclerosis complex in Bulgaria: de novo mutations in the TSC1 and TSC2 genes. <i>J Genet.</i> 2018 Jun;97(2):419-427. PMID: 29912062. DOI: 10.1007/s12041-018-0927-7 (2 citare)

6. Studii clinice naționale – investigator principal

1	2008-2016	Programul național de diagnostic și tratament pentru boli rare și sepsis sever; intervenția 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, Wiegand C, Ambrosini A, Pisano P, Walter MC, Schreiber G, Lusakovska A, Jedrejewska M, Kostera-Pruszycka 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, Burciaru N, Craiu D, Komgut L, Campbell C, Haberova J, Strenkova J, Alejandro M, Jimenez A, Ortiz GG, Enriquez GV, Rodrigues M, Roxburgh R, Dawkins H, Youngs L, Lahdette J, Angelkova N, Saugier-Verbeir P, Cuisset JM, Blotzer C, Jeannot PY, Klein A, Nascimento A, Tizzano E, Selgado
---	-----------	--	--	---

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. Tarta-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. 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, Vigeveno F, Trinkla 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, Bărcă 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

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 Guernini, Ingo Helbig, Helle Hjalgrim, Dorota Hoffman-Zacharska, Johanna Jahn, Karl Martin Klein, Bobby Koeleman, Vladimir Komarek, Roland Krause, Gregor Kühlenbäumer, 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 Aldredge, Eva Andermann, Frederick Andermann, Dina Amrom, Jocelyn F Bautista, Samuel F Berkovic, Judith Bluvstein, Alex Boro, Gregory Cascino, Damian Consalvo, Patricia Crumrine, Omin 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 Esmaeili Nieh, Terence J O'Brien, Ruth Ottman, Stephen

CA8. Barba C, Parrini E, Coras R, Galuppi A, **Craiu D**, Kluger G, Parmeggiani A, Pieper T, Schmitt-Mechelke T, Striano P, Giordano F, Blumcke I, Guernini R. Co-occurring malformations of cortical development and SCN1A gene mutations

WEB OF SCIENCE

Epilepsia. 2014 Jul;55(7):1009-19. doi: 10.1111/epi.12658. Epub 2014 Jun 5. PMID: 24902755 **Free article.** (42citari)
Factor impact in 2014=5.543 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

CA9. Mouthaan BE, Rados M, Barsi P, Boon P, Carmichael DW, Carrette E, **Craiu D**, Cross JH, Diehl B, Dimova P, Fabo D, Francione S, Gaskin V, Gil-Nagel A, Grigoreva E, Guekht A, Hirsch E, Hecimovic H, Helmstaedter C, Jung J, Kalviainen R, Kelemen A, Kimiskidis V, Kobulashvili T, Krsek P, Kuchukhidze G, Larsson PG, Leitinger M, Lossius MI, Luzin R, Malmgren K, Mameniskiene R, Marusic P, Metin B, Özkara C, Pecina H, Quesada CM, Rugg-Gunn F, Rydenhag B, Ryvlin P, Scholly J, Seeck M, Staack AM, Steinhoff BJ, Stepanov V, Tarta-Arsene O, Trinka E, Uzan M, Vogt VL, Vos SB, Vulliemoz S, Huiskamp G, Leijten FS, Van Eijsden P, Braun KP. **E-PILEPSY consortium. Current use of imaging and electromagnetic source localization procedures in epilepsy surgery centers across Europe**

WEB OF SCIENCE

Epilepsia. 2016 May;57(5):770-6. doi: 10.1111/epi.13347. Epub 2016 Mar 25. PMID: 27012361 **Free article** (41 citari)
Factor impact in 2016=5.699 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

CA10. Kobulashvili T, Höfler J, Dobesberger J, Ernst F, Ryvlin P, Cross JH, Braun K, Dimova P, Francione S, Hecimovic H, Helmstaedter C, Kimiskidis VK, Lossius MI, Malmgren K, Marusic P, Steinhoff BJ, Boon P, **Craiu D**, Delanty N, Fabo D, Gil-Nagel A, Guekht A, Hirsch E, Kalviainen R, Mameniskiene R, Özkara C, Seeck M, Rubboli G, Krsek P, Rheims S, Trinka E. **Current practices in long-term video-EEG monitoring services: A survey among partners of the E-PILEPSY pilot network of reference for refractory epilepsy and epilepsy surgery**

WEB OF SCIENCE

Seizure. 2016 May;38:38-45. doi: 10.1016/j.seizure.2016.03.009. Epub 2016 Apr 1. PMID: 27104922 (36 citari)
Factor impact in 2016=2.608 (<https://www.scijournal.org/impact-factor-of-seizure-eur-j-epilep.shtml>)

CA11. Thomas RH, Zhang LM, Carvill GL, Archer JS, Heavin SB, Mandelstam SA, **Craiu D**, Berkovic SF, Gill DS, Mefford HC, Scheffer IE, EuroEPINOMICS RES Consortium. **CHD2 myoclonic encephalopathy is frequently associated with self-induced seizures**

WEB OF SCIENCE

Neurology. 2015 Mar 3;84(9):951-8. doi: 10.1212/WNL.0000000000001305. Epub 2015 Feb 11. PMID: 25672921 **Free PMC article.** (34 citari)
Factor impact in 2015=7.859 (<https://www.scijournal.org/impact-factor-of-neurology.shtml>)

CA12. Hardies K, May P, Djémié T, Tarta-Arsene O, Deconinck T, **Craiu D**, AR working group of the EuroEPINOMICS RES Consortium, Helbig I, Suls A, Balling R, Weckhuysen S, De Jonghe P, Hirst J. **Recessive loss-of-function mutations in AP4S1 cause mild fever-sensitive seizures, developmental delay and spastic paraplegia through loss of AP-4 complex assembly**

WEB OF SCIENCE

Hum Mol Genet. 2015 Apr 15;24(8):2218-27. doi: 10.1093/hmg/ddu740. Epub 2014 Dec 30. **Free PMC article.** (30 citari)
Factor impact in 2015=6.387 (<https://www.scijournal.org/impact-factor-of-hum-mol-genet.shtml>)

CA13. Nissenkorn A, Levy-Drummer RS, Bondi O, Renieri A, Villard L, Mari F, Mencarelli MA, Lo Rizzo C, Meloni I, Pineda M, Armstrong J, Clarke A, Bahi-Buisson N, Mejaski BV, Djuric M, **Craiu D**, Djukic A, Pini G, Bisgaard AM, Melegh B, Vignoli A, Russo S, Angheliescu C, Veneselli E, Hayek J, Ben-Zeev B. **Epilepsy in Rett syndrome-Lessons from the Rett networked database**

WEB OF SCIENCE

Epilepsia. 2015 Apr;56(4):569-76. doi: 10.1111/epi.12941. Epub 2015 Mar 19. (22citari)
Factor impact in 2015=5.570 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

CA14. 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, **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**

WEB OF SCIENCE

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)
Factor impact in 2019=27.576 (<https://www.scijournal.org/impact-factor-of-lancet-diabetes-endocrinology.shtml>)

CA15. Beniczky S, Neufeld M, Diehl B, Dobesberger J, Trinka E, Mameniskiene R, Rheims S, Gil-Nagel A, **Craiu D**, Pressler R, Krysl D, Lebedinsky A, Tassi L, Rubboli G, Ryvlin P. **Testing patients during seizures: A European consensus procedure developed by a joint taskforce of the ILAE - Commission on European Affairs and the European Epilepsy Monitoring Unit Association**

WEB OF SCIENCE

Epilepsia. 2016 Sep;57(9):1363-8. doi: 10.1111/epi.13472. Epub 2016 Jul 21. PMID: 27440172 (19citari)
Factor impact in 2016=5.699 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

CA16. Helbig I, Lopez-Hernandez T, Shor O, Galer P, Ganesan S, Pendziwiat M, Rademacher A, Ellis CA, Hümpfer N, Schwarz N, Seiffert S, Peeden J, Shen J, Štěrbová K, Hammer TB, Möller RS, Shinde DN, Tang S, Smith L, Podun A, Krause R, Benninger F, Helbig KL, Haucke V,

effectiveness of ataluren: comparison of results from the STRIDE Registry and CINRG DMD Natural History Study

WEB OF SCIENCE

Journal of Comparative Effectiveness Research 2020; 9 (5): 341-360; DOI: 10.2217/ceer-2019-0171; Published: APR 2020; Document Type: Article (12 Citari)

Factor impact in 2020=1.458 (<https://www.scijournal.org/impact-factor-of-j-comp-eff-res.shtml>)

CA22. Grillo E, 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, Polgar N, Cogliati F, Ravn K, Pintaudi M, Melegh B, **Craiu D**, Djukic A, Renieri A. Rett networked database: An integrated clinical and genetic network of rett syndrome databases

WEB OF SCIENCE

Hum Mutat. 2012 Jul;33(7):1031-6. doi: 10.1002/humu.22072. Epub 2012 Apr 13. PMID: 22415763. (12 citari)

Factor impact in 2012=6.022 (<https://www.scijournal.org/impact-factor-of-hum-mutat.shtml>)

CA23. Siekierska A, Stamberger H, Deconinck T, Oprea SN, Partoens M, Zhang Y, Sourbron J, Adriaenssens E, Mullen P, Wiencek P, Hardies K, Lee JS, Giong HK, Distelmaier F, Elpeleg O, Helbig KL, Hersh J, Isikay S, Jordan E, Karaca E, Kecskes A, Lupski JR, Kovacs-Nagy R, May P, Narayanan V, Pendziwiat M, Ramsey K, Rangasamy S, Shinde DN, Spiegel R, Timmerman V, von Spiczak S, Helbig I; C4RCD Research Group; Chris Balak, Newell Belnap, Ana Claassen, Amanda Courtright, Matt de Both, Matthew J Huentelman, Marcus Naymik, Ryan Richholt, Ashley L Siniard, Szabolcs Szelinger, David W Craig, Isabelle Schrauwen, Zaid Afawi, Rudi Balling, Stéphanie Baulac, Nina Barišić, Hande S Caglayan, **Dana Craiu**, Rosa Guerrero-López, Renzo Guerrini, Helle Hjalgrim, Johanna Jahn, Karl Martin Klein, Eric Leguam, Johannes R Lemke, Holger Lerche, Carla Marini, Rikke S Møller, Hiltrud Muhle, Felix Rosenow, Jose Serratosa, Arvid Suls, Ulrich Stephani, Katalin Stérbová, Pasquale Striano, Federico Zara (AR working group of the EuroEPINOMICS RES Consortium), Weckhuysen S, Francklyn C, Antonellis A, de Witte P, De Jonghe P. **Biallelic VARS variants cause developmental encephalopathy with microcephaly that is recapitulated in vars knockout zebrafish**

WEB OF SCIENCE

Nat Commun. 2019 Feb 12;10(1):708. doi: 10.1038/s41467-018-07953-w. PMID: 30755616 **Free PMC article** (11 citari)

Factor impact in 2019=12.298 (<https://www.scijournal.org/impact-factor-of-nat-commun.shtml>)

CA24. Muir AM, Myers CT, Nguyen NT, Saykally J, **Craiu D**, De Jonghe P, Helbig I, Hoffman-Zacharska D, Guerini R, Lehesjoki AE, Marini C, Møller RS, Serratosa J, Štěrbová K, Striano P, von Spiczak S, Weckhuysen S, Mefford HC, EuroEPINOMICS-RES NLES working group, Sarah Weckhuysen. Genetic heterogeneity in infantile spasms

WEB OF SCIENCE

Epilepsy Res. 2019 Oct;156:106181. doi: 10.1016/j.epilepsyres.2019.106181. Epub 2019 Jul 29. PMID: 31394400 **Free PMC article**. (9 citari)

Factor impact in 2019=2.368 (<https://www.scijournal.org/impact-factor-of-epilepsy-res.shtml>)

CA25. Rudolf G, Lesca G, Mehriouy MM, Labalme A, Salmi M, Bache I, Bruneau N, Pendziwiat M, Fluss J, de Bellescize J, Scholly J, Møller RS, **Craiu D**, Tommerup N, Valenti-Hirsch MP, Schluth-Bolard C, Sloan-Béna F, Helbig KL, Weckhuysen S, Edery P, Coulbaut S, Abbas M, Scheffer IE, Tang S, Myers CT, Stamberger H, Carvill GL, Shinde DN, Mefford HC, Neagu E, Huether R, Lu HM, Dica A, Cohen JS, Iliescu C, Pomeroy C, Rubenstein J, Helbig I, Sanlaville D, Hirsch E, Szepietowski P. Loss of function of the retinoid-related nuclear receptor (RORB) gene and epilepsy

WEB OF SCIENCE

Eur J Hum Genet. 2016 Dec;24(12):1761-1770. doi: 10.1038/ejhg.2016.80. Epub 2016 Jun 29. PMID: 27352968 **Free PMC article**. (8 citari)

Factor impact in 2016=4.172 (<https://www.scijournal.org/impact-factor-of-eur-j-hum-genet.shtml>)

CA26. Bosemani T, Anghelescu C, Boltshauser E, Hoon AH Jr, Pearl PL, **Craiu D**, Johnston MV, Huisman TA, Poretti A. Subthalamic nucleus involvement in children: A neuroimaging pattern-recognition approach

WEB OF SCIENCE

Eur J Paediatr Neurol. 2014 May;18(3):249-56. doi: 10.1016/j.ejpn.2013.09.010. Epub 2013 Oct 9. PMID: 24149100 Review (8 citari)

Factor impact in 2014=3.000 (<https://www.scijournal.org/impact-factor-of-eur-j-paediatr-neuro.shtml>)

CA27. Frullanti E, Papa FT, Grillo E, Clarke A, Ben-Zeev B, Pineda M, Bahi-Buisson N, Bienvenu T, Armstrong J, Roche-Martinez A, Mari F, Nissenkorn A, Lo Rizzo C, Veneselli E, Russo S, Vignoli A, Pini G, Djuric M, Bisgaard AM, Ravn K, Bosnjak VM, Hayek J, Khajuria R, Montomali B, Cogliati F, Pintaudi M, Hadziszewski K, **Craiu D**, Voinova V, Djukic A, Villard L, Renieri A. Analysis of the Phenotypes in the Rett Networked Database

WEB OF SCIENCE

Int J Genomics. 2019 Mar 27;2019:6956934. doi: 10.1155/2019/6956934. eCollection 2019. PMID: 31049350 **Free PMC article**. (7 citari)

Factor impact in 2019=2.336 (<https://www.scijournal.org/impact-factor-of-int-j-genomics.shtml>)

CA28. Mulhern MS, Stumpel C, Stong N, Brunner HG, Bier L, Lippa N, Riviello J, Rouhl RPW, Kempers M, Pfundt R, Stegmann APA, Kulkolich MK, Telegraf A, Lehman A; CAUSES study, Lopez-Rangel E, Houdinat N, Barth M, den Hollander N, Hoffer MJV, Weckhuysen S; EuroEPINOMICS-RES-MAE working group, Roovers J, Djamie T, Barca D, Ceulemans B, **Craiu D**, Lemke JR, Korff C, Mefford HC, Meyers CT, Siegler Z, Hiatt SM, Cooper GM, Bebin EM, Snijders Blok L, Veenstra-Knol HE, Baugh EH, Brista EH, Volker-Touw CML, van Binsbergen E, Revah-Politi A, Pereira E, McBrien D, Pacault M, Isidor B, Le Caignec C, Gilbert-Dussardier B, Bilan F, Heinzen EL, Goldstein DB, Stevens SJC, Sands TT. **NBEA: Developmental disease gene with early generalized epilepsy phenotypes**

WEB OF SCIENCE

Adjunctive Treatment for Focal Onset Seizures in Pediatric Patients: A Randomized Controlled Trial

WEB OF SCIENCE

J Child Neurol. 2019 Apr;34(5):248-255. doi: 10.1177/0883073818821035. Epub 2019 Jan 27 PMID: 30688135 Clinical Trial. (2 citari)
Factor impact in 2019=1.833 (<https://www.scijournal.org/impact-factor-of-J-CHILD-NEUROL.shtml>)

CA36. 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

WEB OF SCIENCE

J Genet. 2018 Jun;97(2):419-427 PMID: 29932062. DOI: 10.1007/S12041-018-0927-7 (2 citari)
Factor impact in 2018=1.006= FI pe 5 ani in 2019 (<https://www.springer.com/journal/12041>)

CA37. 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 Coe 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, Gallichan J, George B, Gevers EF, Hackenberg A, Halász 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, 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, Menevse 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, Wèmeau JL, Wierzbica 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

WEB OF SCIENCE

Lancet Diabetes Endocrinol. 2020 Jul;8(7):594-605. doi: 10.1016/S2213-8587(20)30153-4.PMID: 32559475 (1 citari)
Factor impact in 2020=27.576 (<https://www.scijournal.org/impact-factor-of-Lancet-diabetes-endocrinology.shtml>)

CA38. Todorov T, Todorova A, Motoescu C, Dimova P, Iancu D, Craiu D, Stoian D, Barbarii L, Bojinova V, Mitev V. Spontaneous recurrent mutations and a complex rearrangement in the MECP2 gene in the light of current models of mutagenesis

WEB OF SCIENCE

Mutat Res. 2012 Jun 1;734(1-2):69-72. doi: 10.1016/j.mrfmmm.2012.04.001. Epub 2012 Apr 16.PMID: 22525432 (1 citation)
Factor impact in 2012=1.143 (<https://www.scijournal.org/impact-factor-of-mutat-res-fund-mol-m.shtml>)

CA39. De Liso P, Pironi V, Mastrangelo M, Battaglia D, Craiu D, Trivisano M, Specchio N, Nabbout R, Vigevano F. Fatal Status Epilepticus in Dravet Syndrome

WEB OF SCIENCE

Brain Sci. 2020 Nov 23;10(11):889. doi: 10.3390/brainsci10110889.PMID: 33238377 Free PMC article.
Factor impact in 2020=3.332 (<https://www.mdpi.com/journal/brainsci>)

CA40. Matricardi S, De Liso P, Freri E, Costa P, Castellotti B, Magri S, Gellera C, Granata T, Musante L, Lesca G, Oertel J, Craiu D, Hammer TB, Möller RS, Barisic N, Abou Jamra R, Polster T, Vigevano F, Marini C. Neonatal developmental and epileptic encephalopathy due to autosomal recessive variants inSLC13A5gene

WEB OF SCIENCE

Epilepsia. 2020 Nov;61(11):2474-2485. doi: 10.1111/epi.16699. Epub 2020 Oct 16.PMID: 33063863
Factor impact in 2020=6.549 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

CA41. Sutton F, Barca D, Komoltsev I, Craiu D, Guekht A, von Oertzen T, Cock HR. Testing blood and CSF in people with epilepsy: a practical guide

WEB OF SCIENCE

Epileptic Disord. 2020 Aug 1;22(4):381-398. doi: 10.1684/epd.2020.1191.PMID: 32782232
Factor impact in 2020=1.632 (<https://www.scijournal.org/impact-factor-of-epileptic-disord.shtml>)

CA42. Heyne HO, Singh T, Stamberger H, Abou Jamra R, Caglayan H, Craiu D, De Jonghe P, Guernini R, Helbig KL, Koelman BPC, Kosmicki JA, Linnankivi T, May P, Muhle H, Möller RS, Neubauer BA, Palotie A, Pendziwiat M, Striano P, Tang S, Wu S, EuroEPINOMICS RES Consortium, Poduri A, Weber YG, Weckhuysen S, Sisodiya SM, Daly MJ, Helbig I, Lal D, Lemke JR. De Novo Variants In Neurodevelopmental Disorders With Epilepsy

WEB OF SCIENCE

Nat Genet. 2018 Jul;50(7):1048-1053. doi: 10.1038/s41588-018-0143-7. Epub 2018 Jun 25.PMID: 29942082
Factor impact in 2018=21.691 (<https://www.scijournal.org/impact-factor-of-nat-genet.shtml>)

CA43. Cardas R, Iliescu C, Butoianu N, Seferian A, Gataullina S, Gargaun E, Nectoux J, Bienvenu T, Craiu D, Gidaro T, Servais L. DMD and West syndrome

WEB OF SCIENCE

Neuromuscul Disord. 2017 Oct;27(10):911-913. doi: 10.1016/j.nmd.2017.07.008. Epub 2017 Jul 19.PMID: 28802771
Factor impact in 2017=2.540 (<https://www.scijournal.org/impact-factor-of-neuromuscular-disord.shtml>)

CA44. Moavero R, Mühlebner A, Luinburg MJ, Craiu D, Aronica E, Curatolo P. Genetic pathogenesis of the epileptogenic lesions in Tuberous Sclerosis Complex: Therapeutic targeting of the mTOR pathway. Epilepsy Behav. 2021 Jan 8;107713. doi:

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>)

CA51. Ingo Helbig, Tania Lopez-Hernandez, Oded Shor, Peter Galer, Shiva Ganesan, Manuela Pendziwiat, Annika Rademacher, Colin A Ellis, Nadia Hümpfer, Niklas Schwarz, Simone Seifert, 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 Jahn, Karl Martin Klein, Bobby P C Koeleman, Vladimir Komarek, Eric Leguern, Anna-Elina Lehesjoki, Johannes R Lemke, Holger Lerche, Taria Linnankivi, Carla Marini, Patrick May, Hiltrud Muhle, Deb K Pal, Aarno Palotie, Felix Rosenow, Susanne Schubert-Bast, Kaia 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.

Factor impact in 2019=10.669 (<https://www.scijournal.org/impact-factor-of-am-j-hum-genet.shtml>)

CA52. 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 Maiter, Inge Gies, Martine Cools, Kristina Casteels, Albert Beckers, Sabina Zacharieva, 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 Poirou, Bernert, Philippe Touraine, Philippe Chanson, Michel Polak, Maïthe 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, Laura Janozola, Rasa Verkauskienė, Michael Witsch, James Clark, Johannes Romijn, Thera Links, Nienke Biemass, Sabine Hannema, Bas Havekes, Hedi Claahsen-van der Grinten, Henri Timmers, Robin Peeters, Gerlof Valk, A A Verjyn Stuart, Harm Haak, Eystein Husebye, Jens Bollerslev, Barbara Jarzab, Agnieszka 'Szypowska, João-Filipe Raposo, Dana Craiu, Doina Piciu, Ludmila Kostalova, Jarmila Vojtková, Tadej Battelino, Roque Cardona-Hernandez, Diego Yeste, Sonia Gaztambide, Anna Nordenström, Neil Gittos, 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>)

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 Jahn, Karl M Klein, Bobby P C Koeleman, Vladimir Komarek, Roland Krause, Eric Leguern, Anna-Elina Lehesjoki, Johannes R Lemke, Holger Lerche, Taria 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 (Leand 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 Fattai-Valevski, Ki Joong Kim, Ahmad Beydoun, Ghassan Hmaïmess, 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 Kravijanc, Dimitrije Nikolic, Francisco Lopez

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 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>)

CA62. de Kovel CG, Bristra EH, van Kempen MJ, Van't Slot R, Nijman IJ, Afawi Z, De Jonghe P, Djémié T, Guernini R, Hardies K, Helbig I, Hendrickx R, Kanaan M, Kramer U, Lehesjoki AE, Lemke JR, Marini C, Mei D, Möller 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, E LeGuern, H Lerche, P May, H Muhle, D Pal, A Palotie, F Rosenow, K Salmér, J M Serratosa, S M Sisodiya, U Stephani, K Sterbova, P Striano, T Talvik, M van Haelst, N Verbeek, S von Spiczak, Y G Weber), 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):588-80. doi: 10.1002/mgg3.235. eCollection 2016 Sep.PMID: 27652284 **Free PMC article.**
Factor impact in 2016=1.995 (<https://www.wiley.com/en-us/Molecular+Genetics+%26+Genomic+Medicine-p-9780JRN75484>) in 2018

CA63. Hardies K, Cai Y, Jardel C, Jansen AC, Cao M, May P, Djémié T, Hachon Le Camus C, Keymolen K, Deconinck T, Bhamhani V, Long C, Sajjan SA, Helbig KL. AR working group of the EuroEPINOMICS RES Consortium (Zaid Afawi, Stéphanie Baulac, Nina Barisic, Hande Caglayan, Dana Craiu, Carolien G De Kovel, Rosa Guerrero Lopez, Renzo Guernini, Helle Hjalgrim, Holger Lerche, Johanna Jahn, Karl Martin Klein, Bobby C Koeleman, Eric Leguenn, Johannes Lemke, Carla Marini, Hiltrud Muhle, Felix Rosenow, José M Serratosa, Katalin S Stérbová, Rikke S Möller, Aarno Palotie, Pasquale Striano, Yvonne Weber, Federico Zara), Suls A, Balling R, Helbig I, De Jonghe P, Depienne C, De Camilli P, Weckhuysen S. **Loss of SYNJ1 dual phosphatase activity leads to early onset refractory seizures and progressive neurological decline.** *Brain.* 2016 Sep;139(Pt 9):2420-30. doi: 10.1093/brain/aww180. Epub 2016 Jul 19.PMID: 27435091 **Free PMC article.**
Factor impact in 2016=11.075 (<https://www.scijournal.org/impact-factor-of-brain.shtml>)

CA64. Larsen J, Johannesen KM, Ek J, Tang S, Marini C, Blichfeldt S, Kibaek M, von Spiczak S, Weckhuysen S, Frangu M, Neubauer BA, Uldall P, Striano P, Zara F; MAE working group of EuroEPINOMICS RES Consortium (D C Craiu, H S Caglayan, T Talvik, Y G Weber, N Barisic), Kleiss R, Simpson M, Muhle H, Nikanorova M, Jepsen B, Tommerup N, Stephani U, Guernini R, Duno M, Hjalgrim H, Pal D, Helbig I, Möller RS. **The role of SLC2A1 mutations in myoclonic astatic epilepsy and absence epilepsy, and the estimated frequency of GLUT1 deficiency syndrome.** *Epilepsia.* 2015 Dec;56(12):e203-8. doi: 10.1111/epi.13222. Epub 2015 Nov 5.PMID: 26537434.
Factor impact in 2015=5.570 (<https://www.scijournal.org/impact-factor-of-epilepsia.shtml>)

CA65. 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 (Guy McKhann, Michael Eliasziw, Paul Graham Fisher, Gihan Tennekoon, Daphne T Hsu, Marie-Cécile Nassogne, Tayeb Sekhara, Mahmud Almadani, Michael O'Mahony, Lawrence Richer, Niels Illum, Hans-Christian Laugaard-Jacobsen, Valentin Sander, Inga Talvik, Mikko Kallela, Petra Keski-Santti, Jarmo Kuikkonen, Eija Nikkanen, Markku Nissila, Markku Partinen, Jukka Peltola, Daniel Annequin, Jean-Christophe Cuvelier, Elisabeth Fournier-Charrière, Sylvie Laborde, Mathieu Milh, Marie-Louise Navez, Dominique Parain, Agnes Suc, Friedrich Ebinger, Stefan Evers, Charly Gaul, Astrid Gendolla, Jan-Peter Jansen, Klaus Laengler, Raymond Pothmann, Ruediger Schellenberg, Sandeep Aganwal, Ish Anand, Sathish Chandran, V B Chodhary, Praveen Jain Harawat, Shaneur Harsha Pramod Jog, Alagarsamy Kannan, Ummer Karandan, Anthathi Soma Keerthi, Shankara Nellikunja, Lekha Pandit, R Srinivasa, Piero Barbanti, Daina Jegere, Jurgis Strautmanis, Wim Mulleners, Paul Pop, Peter Van den Berg, Besna Bryn, Thorbjorn Kjaemli, Kristian Sommerfelt, Iwona Kochanowska, Marek Pietrzak, Jolanta Strzelecks, Marek Szatanik, Marzena Wesolowska, Ileana Benga, Dana-Cristina Craiu, Georgeta Diaconu, Cristian Gheonea, Laura Popescu, Josep Artigas Pallares, Oscar Blanco Barca, Jaume Campistol Plana, Alfons Macaya, Maria del Carmen Mosquera Villaverde, Alejandro Reyes Martin, Maria Victoria San Antonio Bernt Danielsson, Ylva Ohmer, Charles Peter Arthur, Pronab Bala, Sharyn Gardner, Jayaprakash Gosalakkal, Ponnudas Prabhakar, Alber Abraham, Lawrence Adler, Mario Aguado, Ashraf Atalla, Padmini B Atri, Sheena Aurora, Riaz Baber, James W Banks, Robert Bargar, James Barker, Patricia Barrington, Lucinda Bateman, Christopher E Baur, Harold Bays, Nathan L Bennett, Frank R Berenson, Gary D Berman, Alan Bernstein, Perminder Bhatia, Andrew Blumenfeld, Dale Bramlet, Robert E Broker, Sheri Byrd, Roger Cady, John Calcagno, Alvaro Camacho, Walter Carlini, Joseph Casadonte, Steve Choi, Shane G Christensen, Frank Civitarese, William D Clark, Clinton N Corder, Ronald David, Vital Dhaduk, Cheryl A Duffy, Robert Earle, Michael Edmond, Johnny Edrozo, Eric Eross, John S Erwin, Juan B Espinosa-Paccini, Beal Essink, Mildred Farmer, Michael Fedlman, Marylou Fernando, Ronald R Fieve, Mark Fisher, Stephen Flitman, Linda B Ford, Stephen D Forner, Eileen Fox, Brad Frandsen, James A Fry, Gene R Fuller, Mary Elizabeth Gaffney, Andrew Gasecki, Charles Gay, Stephen Gelfand, Thomas Giancarlo, John Giblin, Martin C Glover, Gary Goldstein, Herbert Goodman, Glenn Gordon, Sushma V Gorrela, William Grainger, James Gray, Piyush Gupta, Srinivas N Halothore, Nelson M Handal, Duane J Harris, Bryan Harvey, Lydie Hazan, James Hedrick, Dan C Henry, Mark O Herring, Richard L Hines, Willis Holloway Jr, Alexander E Horwitz, Randall Huling, James Igleburger, William Jennings, Thomas Jones, Katie A Julien, Alan Katie, Edward F Kent, Antaram Khan, Arifulla Khan, Murray A Kimmel, Tracy R Klein, James Knutson, Mary Beth Krafty, James Kratzer, Joseph Kwentus, Daniel Lacey, Paula Lane, Diana Lebron, Kurt Lesh, Joseph Ley, Steve Linder, Edwin Liu, Steven R Lubet, Bennett Mechanic, Isaac Marcadis, Herbert G Markely, Paul J Markovitz, Laszlo J Mate, Ninan T Mathew, Peter McAllister, John W McGettigan, Paul Means, Laszlo Mechtler, Vishal Mehra, Isaac Melamed, David C Miller, Janice L Miller, Fernando Miranda, Mia Moon, Ivy M Muhar, Shanan Munoz, Kevin Murphy, Rajiv Nanavaty, Nicholas A Nayak, Jeffrey Nelson, Robert

1. Sandu C, Burloiu CM, Barca DG, Magureanu SA, Craiu DC. Ketogenic Diet in Patients with GLUT1 Deficiency Syndrome. *Maedica (Bucur)*. 2019 Jun;14(2):93-97. doi: 10.26574/maedica.2019.14.2.93. PMID: 31523287; PMCID: PMC6709387
2. Tarta-Arsene O, Leanca M, Gandeia L, Craiu D. Is there an encephalographic trait to septo-optic dysplasia? (de Morsier Syndrome), *Romanian Journal of Neurology*, Volume XIII, No. 2, 2014.
3. Barca D, Tarta-Arsene O, Dica A, Iliescu C, Budisteanu M, Motoescu C, Butoianu N, Craiu D. Intellectual disability and epilepsy in down syndrome. *Maedica (Bucur)*. 2014 Dec;9(4):344-50. PMID: 25705303; PMCID: PMC4316878.
4. 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, *Romanian Journal of Neurology*, Volume XII, No. 2, 2013.
5. Duca DG, Craiu D, Boer M, Chiriac SM, Arghir A, Tutulan-Cunita A, Barca D, Iliescu C, Lungeanu A, Magureanu S, Budisteanu M. Diagnostic approach of angelman syndrome. *Maedica (Bucur)*. 2013 Sep;8(4):321-7. PMID: 24790661; PMCID: PMC3968465.
6. Iliescu C, Craiu D. Diagnostic approach of epilepsy in childhood and adolescence. *Maedica (Bucur)*. 2013 Jun;8(2):195-9. PMID: 24371485; PMCID: PMC3865130.
7. Albeanu A.G, Magureanu S, Craiu D, Lagae L, Hippocampal sclerosis - cause or consequence of mesial temporal lobe epilepsy in children? *Romanian Journal of Neurology*, Volume XI, No. 1, 2012.
8. Tarta-Arsene O., Preoteasa F., Magureanu S.A., Agavriloiu M., Motoescu C.H., Tarta-Arsene E. Utilitatea imaginii prin rezonanță magnetică funcțională în evaluarea copiilor cu epilepsie. *Romanian Journal of Neurology*, vol 10, nr 1/2011
9. Dana Craiu. Dieta cetogenică – din nou în actualitate. *Medica Academica*, 2011
10. Tarta-Arsene O, Preoteasa F, Magureanu S, Iliescu A, Craiu D, Motoescu C, Tarta-Arsene E, Ciobanu. Functional magnetic resonance imaging contribution to language areas assessment in children with non-lesion focal epilepsy, *Romanian Journal of Neurology*. 2010; 9(3):130-136; DOI:10.37897/RJN.2010.3.4
11. Tarta-Arsene O, Craiu D, Ciobanu G, Barca D.G, Motoescu C, Varsatura si clipitul unilateral ictal asociat cu crize epileptice focale temporale drepte, *Revista de Neurologie si Psihiatrie a Copilului si Adolescentului din Romania*, Volumul 13, Nr. 2, 2010.

AUTOR CARTI

1. Craiu D, Iliescu I, Vintan M.A., Mindruta I, *EPILEPTOLOGY – BASIC COURSE (East European Course Of Epilepsy)*, Editura Medbook Publishing, Bucuresti, 2014.
2. Craiu D, Iliescu C - *NEUROLOGIE PEDIATRICA - Note de curs*, Editura Universitara "Carol Davila", Bucuresti, 2013.
3. Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013.
4. Craiu D. Cum prezentăm un caz neuropediatric la examen?. Editura Universitara „Carol Davila”, Bucuresti, 2007.
5. Magureanu S (Craiu D - coautor), *Elemente de Neurologie Pediatrica*, Editura Universitara Carol Davila, Bucuresti 1999

AUTOR CAPITOLE CARTE:

Capitole carti - autor unic

1. Craiu D. Diagnostic: Algoritm de diagnostic. In: Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013: 107-110.
2. Craiu D. Tratamentul: Tratamentul medicamentos corticoterapic. In: Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013: 116-121.
3. Craiu D. Tratamentul: Îmbunătățirea homeostazei calciului. In: Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013: 174.
4. Craiu D. Tratamentul: Abordări terapeutice în curs de cercetare. Ameliorarea calitatii, functiei si masei musculare. Creșterea fortei musculare. In: Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013: 183-184.
5. Craiu D. Tratamentul: Măsuri suportive. In: Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013: 184-185.
6. Craiu D. Evoluție, Prognostic. In: Craiu D. *DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament*. Editura Universitara „Carol Davila”, Bucuresti, 2013: 197-198.
7. Craiu D. Compliance to antiepileptic treatment. In: Craiu D, Iliescu C, Vintan M, Mindruta I, *EPILEPTOLOGY – BASIC COURSE, EAST EUROPEAN COURSE OF EPILEPSY*, Editura MedBook, Cheile Gradistei, Romania, 2014: 149-153.
8. Craiu D. Drug treatment discontinuation when, how and whom to stop? In: Craiu D, Iliescu C, Vintan M, Mindruta I, *EPILEPTOLOGY – BASIC COURSE, EAST EUROPEAN COURSE OF EPILEPSY*, Editura MedBook, Cheile Gradistei, Romania, 2014: 177-182.

14. Butoianu N, Magureanu S, Craiu M, Craiu D. Aspectul clinic al distrofinopatiilor: Femeile purtatoare simptomatice. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 44-45.
15. Butoianu N, Magureanu S, Craiu M, Craiu D. Aspectul clinic al distrofinopatiilor: Pacienti asimptomatici sau minim simptomatice. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 45.
16. Robanescu N, Craiu D. Investigatii pentru diagnostic și urmarire: Evaluarea musculo-articulara (bilantul de recuperare fizica si kinetoterapie). In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 97-102.
17. Butoianu N, Minciu I, Iliescu C, Craiu D. Investigatii pentru diagnostic și urmarire: Diagnosticul diferential al distrofinopatiilor. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 111-113.
18. Robanescu L, Craiu D. Tratamentul: Tratamentul afectarii musculo-articulare. Tratamentul de recuperare motorie a pacientului DMD. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 121-124.
19. Robanescu L, Thiery A, Craiu D. Tratamentul: Ortezarea. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 124-129.
20. Craiu D, Cinteza E. Tratamentul: Tratamentul afectarii cardiace. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 154-155.
21. Sandu C, Craiu D. Tratamentul: Consideratii privind ingrijirea de urgenta. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 169-170.
22. Verhaart EC I, Aartsma-Rus A, Craiu D. Tratamentul: Abordari terapeutice in curs de cercetare. Terapia genica. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 170-172.
23. Verhaart EC I, Aartsma-Rus A, Craiu D. Tratamentul: Abordari terapeutice in curs de cercetare. Terapia celulara. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 172-173.
24. Verhaart EC I, Aartsma-Rus A, Craiu D. Tratamentul: Abordari terapeutice in curs de cercetare. Cresterea sintezei proteinelor compensatorii (Utrofina). In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 175-176.
25. Verhaart EC I, Aartsma-Rus A, Craiu D. Tratamentul: Abordari terapeutice in curs de cercetare. Tratamente dedicate tipului de mutatie. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 176-181.
26. Verhaart EC I, Aartsma-Rus A, Craiu D. Tratamentul: Abordari terapeutice in curs de cercetare. Ameliorarea calitatii, functiei si masei musculare. Reducerea stresului oxidativ. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 181-182.
27. Craiu D, Verhaart EC I, Aartsma-Rus A. Tratamentul: Abordari terapeutice in curs de cercetare. Ameliorarea calitatii, functiei si masei musculare. Cresterea masei musculare. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 182-183.
28. Craiu D, Thiery A, Budisteanu M. Abordarea multidisciplinara a pacientului cu distrofinopatie. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 187-189.
29. Butoianu N, Iancu D, Craiu D. Prezenta cazuri clinice. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 199-215.
30. Craiu D. Tratamentul: Tratamentul medicamentos corticoterapie. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 116-121.
31. Craiu D. Tratamentul: Imbunatatirea homeostazei calciului. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 174.
32. Craiu D. Tratamentul: Abordari terapeutice in curs de cercetare. Ameliorarea calitatii, functiei si masei musculare. Cresterea fortei musculare. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 183-184.
33. Craiu D. Tratamentul: Masuri suportive. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 184-185.
34. Craiu D. Evolutie, Prognostic. In: Craiu D. DISTROFINOPATIILE - Noțiuni teoretice. Algoritmi de diagnostic și tratament. Editura Universitară „Carol Davila”, București, 2013: 197-198.
35. Minciu I, Pomoran C, Craiu D, Morfogeneza sistemului nervos, NEUROLOGIE PEDIATRICA - NOTE DE CURS, Editura Universitară "Carol Davila", București, 2013: 11-18.

20. Sandu, C.; Fitzsimmons, G.; Sewell, M.; et al. KETOGENIC DIET SERVICE-SPECTRUM OF REFERRED CASES ACCORDING TO AGE. *EPILEPSIA* Volume: 58 Special Issue: SI Supplement: 5 Pages: S192-S192 Meeting Abstract: p1076 Published: DEC 2017. Conference: 32nd International Epilepsy Congress Location: Barcelona, SPAIN Date: SEP 02-06, 2017
21. Papuc, Sorina Mihaela; Tutulan-Cunita, Andreea-Cristina; Budisteanu, Magdalena; et al. Rare genomic imbalances in patients with neurodevelopmental disorders and complex phenotypes. *MOLECULAR CYTOGENETICS* Volume: 10 Supplement: 1 Meeting Abstract: 1.P69 Published: JUN 29 2017
22. Rudolf, G.; Lesca, G.; Mehrjouy, M. M.; Rudolf G, Lesca G, Mehrjouy M, Labalme A, Salmi, Bache I, Bruneau N, Pendziwiat M, Fluss J, de Bellescize J, Scholty J, Moller R, Craiu D, Tommerup N, et al. LOSS-OF-FUNCTION OF THE RETINOID-RELATED NUCLEAR RECEPTOR (RORB) GENE AND EPILEPSY. *EPILEPSIA* Volume: 57 Special Issue: SI Supplement: 2 Pages: 121-121 Meeting Abstract: P388 Published: DEC 2016; WOS:000425754400348; ISSN: 0013-9580; eISSN: 1528-1167
23. Lawerman, T. F.; Brandsma, R.; Lunsing, R. J.; et al. Group Author(s): European Pediat Neurology Soc. European pediatric normative values for the scale for assessment and rating of ataxia (SARA). *MOVEMENT DISORDERS* Volume: 31 Supplement: 2 Pages: S347-S347 Meeting Abstract: 1080 Published: JUN 2016. Conference: 20th International Congress of Parkinson's Disease and Movement Disorders Location: Berlin, GERMANY Date: JUN 19-23, 2016
24. Arbune, A. A.; Tarta-Arsene, O.; Craiu, D. Seizure triggers in focal and generalised epilepsies. *EUROPEAN JOURNAL OF NEUROLOGY* Volume: 23 Special Issue: SI Supplement: 2 Pages: 167-167 Meeting Abstract: P11120 Published: JUN 2016
25. Kobulashvili, T.; Dobesberger, J.; Hoefler, J.; et al. Current practices in long-term video-EEG monitoring services: a survey among partners of the E-PILEPSY pilot network of reference for refractory epilepsy and epilepsy surgery. *EUROPEAN JOURNAL OF NEUROLOGY* Volume: 23 Special Issue: SI Supplement: 2 Pages: 767-767 Meeting Abstract: P32041 Published: JUN 2016
26. Bastian, A. E.; Manole, E.; Mageriu, V.; et al. The importance of a fast and accurate diagnosis in neonatal onset hypotonia: A Merosin-deficient Congenital muscular dystrophy type 1A (MDC1A) case presentation. *VIRCHOWS ARCHIV* Volume: 467 Supplement: 1 Pages: S104-S104 Meeting Abstract: PS-05-033 Published: SEP 2015
27. Lawerman, T. F.; Brandsma, R.; Barisic, N.; et al. European SARA age validation trial in children -Preliminary results-. *MOVEMENT DISORDERS* Volume: 30 Supplement: 1 Pages: S360-S360 Meeting Abstract: 925 Published: JUN 2015. Conference: 19th International Congress of Parkinson's Disease and Movement Disorders Location: San Diego, CA Date: JUN 14-18, 2015 Sponsor(s): Int Parkinson Movement Disorders Soc
28. Syrbe, S.; Hedrich, U.; Riesch, E.; et al. Group Author(s): EuroEPINOMICS RES Consortium. DE NOVO LOSS-OR GAIN-OF-FUNCTION MUTATIONS IN KCNA2 CAUSE EPILEPTIC ENCEPHALOPATHY. *EPILEPSIA* Volume: 56 Special Issue: SI Supplement: 1 Pages: 77-77 Meeting Abstract: p0280 Published: FEB 2015. Conference: 31st International Epilepsy Congress Location: Istanbul, TURKEY Date: SEP 05-09, 2015
29. Djemie, T.; Dejanovic, B.; Suls, A.; et al. Group Author(s): EuroEPINOMICS Dravet Working Grp. SIMULTANEOUS IMPAIRMENT OF NEURONAL AND METABOLIC FUNCTION OF GEPHYRIN IN A PATIENT WITH EPILEPTIC ENCEPHALOPATHY. *EPILEPSIA* Volume: 56 Special Issue: SI Supplement: 1 Pages: 223-224 Meeting Abstract: p0909 Published: FEB 2015. Conference: 31st International Epilepsy Congress Location: Istanbul, TURKEY Date: SEP 05-09, 2015
30. Budisteanu, M.; Tutulan-Cunita, A.; Papuc, S. M.; et al. INVESTIGATION OF GENETIC CAUSES OF EPILEPSY IN CHILDREN WITH NON-SPECIFIC INTELLECTUAL DISABILITY USING ACGH-BASED TECHNOLOGY. *EPILEPSIA* Volume: 55 Special Issue: SI Supplement: 2 Pages: 37-37 Meeting Abstract: 095 Published: JUN 2014. Conference: 11th European Congress on Epileptology Location: Stockholm, SWEDEN Date: JUN 29-JUL 03, 2014
31. Suls, A.; Jaehn, J. J.; Kecskes, A.; et al. Group Author(s): EuroEPINOMICS RES Consortium. DE NOVO LOSS-OF-FUNCTION MUTATIONS IN CHD2 CAUSE A FEVER-SENSITIVE MYOCLONIC EPILEPTIC ENCEPHALOPATHY SHARING FEATURES WITH DRAVET SYNDROME. *EPILEPSIA* Volume: 55 Special Issue: SI Supplement: 2 Pages: 79-80 Meeting Abstract: p235 Published: JUN 2014. Conference: 11th European Congress on Epileptology Location: Stockholm, SWEDEN Date: JUN 29-JUL 03, 2014
32. Nissenkorn, A.; Bondi, O.; Drumer, Levi R.; et al. EPILEPSY IN RETT SYNDROME: LESSONS FROM THE RETT NETWORKED DATABASE. *EPILEPSIA* Volume: 55 Special Issue: SI Supplement: 2 Pages: 145-145 Meeting Abstract: p446 Published: JUN 2014. Conference: 11th European Congress on Epileptology Location: Stockholm, SWEDEN Date: JUN 29-JUL 03, 2014
33. Thomas, R. H.; Zhang, L. M.; Carvill, G. L.; et al. Group Author(s): euroEPINOMICS-RES. CHD2 MUTATIONS PRODUCE AN EARLY CHILDHOOD ENCEPHALOPATHY WITH PROMINENT PHOTOSENSITIVE

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, Bangkok, 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 Romane Impotriva Epilepsiei, Bucuresti, 4-6 octombrie 2018
27. 19th International Symposium on Severe Infantile Epilepsies Old and New Treatments, 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, 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. EEG, scleroza tuberoasa.

ROMÂNIA
MINISTERUL SĂNĂTĂȚII

CERTIFICAT

DE MEDIC PRIMAR

Se certifică prin prezentul că:

CRAIU DANA CRISTINA

este confirmat(ă) **MEDIC** primar în specialitatea **NEUROLOGIE PEDIATRICA**
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

Seria SI Nr. 014087

ROMÂNIA
MINISTERUL SĂNĂTĂȚII

CERTIFICAT

DE MEDIC SPECIALIST

Se certifică prin prezentul că:

CRAIU P. DANA- CRISTINA

este confirmat(ă) **MEDIC** specialist în specialitatea **PEDIATRIE**
prin Ordinul Ministrului Sănătății nr. **740** din **2012**
pe baza examenului susținut în sesiunea **MARTIE 2011** și promovat cu media generală **9.13**
Eliberat la data de **24.07.2012** cu numărul **13999**
Prezentul certificat s-a eliberat în conformitate cu prevederile legale.

DIRECTOR GENERAL,
IC. ADRIAN COCOS

CONSILIER,
GABRIELA CHELBA

Seria SI Nr. 014318



18.12.2023