INFORMAŢII PERSONALE

Jurca-Simina Iulia-Elena

Str. Lt. Anca- Virgil, nr. 40, loc. Abrud, jud Alba, cod 515100, România
+40741030292
jurca.iulia@umft.ro, jurca.iulia@gmail.com

Sexul feminin | Data naşterii 16/09/1986 | Naţionalitatea română

EXPERIENŢA PROFESIONALĂ

Medic specialist Genetică medicală

01.04.2021- prezent — Societatea Civila Medicala ” DR BABEU”, Timisoara, Blv. Liviu Rebreanu nr 190

01.2021- prezent — Evaluare medicala la distanta prin platforma tehnologica Medicentrum.ro, detinuta de S.C. TRUDOC ADVICE S.R.L.., cu sediul social in Sibiu, str. Strandului, nr. 29, sc.B, ap.12, jud. Sibiu

Tipul sau sectorul de activitate

Evaluare genetica medicala fizic si la distanta, in regim ambulatoriu si de tip telemedicina

Medic specialist Pediatrie

01.01.2022- prezent — Spitalul Clinic de Urgență pentru Copii “Louis Țurcanu”, Str Iosif Nemoianu nr 2, Timișoara

03.2021- 31.12.2021 — Unitatea administrativ-teritoriala a Municipiului Timisoara, str. Bv. C.D. Loga nr.1, jud Timis

01.04.2021- prezent — Societatea Civila Medicala ” DR BABEU”, Timisoara, Blv. Liviu Rebreanu nr 190

01.2018- prezent — Policlinica Regina Maria Iulius Mall, Str. Aristide Demetriade Nr 1, in incinta Iulius Mall, etaj 1, Timișoara si Centrul Medical Sfânta Maria, Strada Romulus 62, Timișoara

Tipul sau sectorul de activitate

Medic coordonator al Centrului de vaccinare pediatrica + efectuare garzi in Unitatea de Primire a Urgentelor

Medic vaccinator Centru vaccinare Incuboxx, medic coordonator al Centrului de vaccinare din 02.08.2021 pana in 31.12.2021

Consultatii de pediatrie in regim ambulatoriu, fizice si prin telemedicina

Asistent universitar la plata cu ora, personal extern- disciplina Genetica Medicala

10.2021- 01.2022
09.2016- 01.2020 — Universitatea de Medicină şi Farmacie „Victor Babeș”, Piața Eftimie Murgu nr 2, Timișoara

Tipul sau sectorul de activitate

Activitate didactica- lucrări practice Genetica Medicala

Medic rezident Genetica medicala

01.2018- 08.2020 — Universitatea de Medicină şi Farmacie Oradea, Spitalul Municipal „Dr. Gavril Curteanu” Oradea

Detașată la:

Universitatea de Medicina şi Farmacie „Victor Babeș”, Piața Eftimie Murgu nr 2, Timișoara

Spitalul Clinic de Urgență pentru Copii “Louis Țurcanu”, Str Iosif Nemoianu nr 2, Timișoara

Tipul sau sectorul de activitate

Stagii efectuate si sectoare de activitate : genetica medicala. genetică clinică, citogenetică, genetică moleculară, informatică și biostatistică, obstetrică și ginecologie, medicină internă, neurologie, endocrinologie + 1 an, 5 luni și 2 săptămâni stagii considerate de la prima specializare în pediatrie

Medic rezident Pediatrie

01.2012- 07.2017 Spitalul Clinic de Urgență pentru Copii "Louis Țurcanu", Str Iosif Nemoianu nr 2, Timișoara

Tipul sau sectorul de activitate

Stagii efectuate si sectoare de activitate : pediatrie generala, nefrologie pediatrica, gastro-enterologie pediatrica, cardiologie pediatrica, pneumologie-ftiziologie pediatrica, boli infectioase, chirurgie si ortopedie pediatrica, toxicologie, diabet si nutritie pediatrica, neurologie pediatrica, psihiatrie pediatrica, onco-hematologie pediatrica, urgente pediatrice, terapie intensiva pediatrica, neonatologie, genetica, ecografie generala, etica cercetarii.

Praticien Attaché Associé

09.2014-01.2016 Spitalul "Raymond Poincaré", Assistance Publique – Hôpitaux de Paris
Serviciul de Genetica Medicala, Coordonator Profesor Dominique P. GERMAIN
104, bulevard Raymond Poincaré, 92380 GARCHES, Franța

Tipul sau sectorul de activitate

Genetica clinica, activități de cercetare, participare in studii clinice

Voluntar in Organizatia Salvati Copiii- Filiala Timis

10.2007-09.2014 Organizatia salvati Copiii, Filiala Timis, B-dul Republicii nr.1, Corp B, Ap 7, Timişoara, Timiș

Tipul sau sectorul de activitate

Organizare si desfasurare de programe educationale pentru sanatate si preventie, dar si drepturi ale copiilor (participant si formator)

EDUCAŢIE ŞI FORMARE

Doctor in medicina

10.2012- 24.10.2019 Universitatea de Medicină şi Farmacie „Victor Babeş" Timişoara

Confirmat prin Ordinul Ministerului Sanatatii OM-5748_13_10_2020

Teza: Strategii de evaluare ale patologiei genetice corelate progresului tehnologic si informatic

Coordonator: Prof. Dr. Puiu Maria

Tipul sau sectorul de activitate

Genetica medicala

Link sumar teza: http://www.umft.ro/2019_789

Doctor-medic

2005-2011 Universitatea de Medicină şi Farmacie „Victor Babeş" Timişoara
Facultatea de Medicina Generala

Bacalaureat

2001-2005 Liceul „Horia, Cloșca si Crișan" Abrud
Profil vocațional- pedagogic; învățător-educator

COMPETENTE PERSONALE

Limba maternă: Limba romana

Alte limbi străine cunoscute

	INTELEGERE		VORBIRE		SCRIERE
	Ascultare	Citire	Participare la conversaţie	Discurs oral	
Limba engleza	C1	C1	C1	C1	C1
Limba franceza	C1	C1	C1	C1	C1

Niveluri: A1/A2: Utilizator elementar - B1/B2: Utilizator independent - C1/C2: Utilizator experimentat
Cadrul european comun de referinţă pentru limbi străine

Competenţe de comunicare

- Facilitate si deschidere în stabilirea relațiilor interpersonale şi capacitate de a stabili bune contacte cu pacienții și familiile acestora, cu specialiștii din diverse domenii medicale.
- Adeptă a spiritului de echipă la locul de muncă.

Competenţe organizaţionale/manageriale

Cultura organizațională si abilitate in scrierea si coordonarea proiectelor adaptate ONG cu activitate in domeniul sănătății. Implicarea activă în proiecte educaționale, coordonarea lor si organizarea a diferite activități extra curriculare, de susținere directă a bolnavilor, de informare şi campanii pe tema bolilor rare și de pregătire pentru studenți în diferite arii medicale.

Competenţă digitală

AUTOEVALUARE				
Procesarea informaţiei	Comunicare	Creare de conţinut	Securitate	Rezolvarea de probleme
Utilizator experimentat	Utilizator experimentat	Utilizator independent	Utilizator experimentat	Utilizator experimentat

Niveluri: Utilizator elementar - Utilizator independent - Utilizator experimentat
Competențele digitale - Grilă de auto-evaluare

Alte competențe informatice: o bună stăpânire a suitei de programe de birou (procesor de text, calcul tabelar, software pentru prezentări), cunoștințe dobândite in cadrul procesării lucrării de licență, a editării posterelor prezentate si a prezentărilor orale, a contribuției directe la scrierea de publicații medicale; bune cunoștințe de editare foto, dobândite ca fotograf amator.

Alte competenţe

Multiplele activități de voluntariat desfășurate pe parcursul traiectoriei educaționale au contribuit la formarea mea ca personalitate activă, ambițioasa, cu spirit de inițiativă, capabilă de a se implica într-o muncă solicitantă şi de a-şi asuma un rol responsabil in activitățile întreprinse, interesata de psihologia comunicării si a copilului si adolescentului.
Temele esențiale ale cercetărilor efectuate sunt din aria geneticii, pediatriei generale si a nefrologiei.

Permis de conducere

Categoria B din 2004

INFORMATII SUPLIMENTARE

Proiecte de cercetare	1. PIR16183: A Prospective and Retrospective Cohort Study to Refine and Expand the Knowledge on Patients With Chronic Forms of Acid Sphingomyelinase Deficiency (ASMD), NCT04106544, Investigational Site Number 6420001, Timisoara, Romania, 300011, Study Director: Clinical Sciences & Operations Sanofi, Study Start Date: September 27, 2019, Estimated Primary Completion Date : March 2023- **coinvestigator** incepand cu **18.02.2020- prezent** 2. Project Science and Technology in childhood Obesity Policy (STOP), Grant Agreement number 774548; Call: Horizon 2020-SFS-2016-2017; Topic: SFS-39-2017, Research Innovation Action 2018-2020. WP 8: T8.2 - Three-site RCT of an early childhood obesity intervention: design and tool development. Translation of material for intervention, Universitatea de Medicina si Farmacie Victor Babes Timisoara, Romania –***membru activ 31.08.2018- prezent*** 3. Formarea PROfesionala a personalului medical in GENetica medicala– PROGEN (PROfessional formation of medical personnel in medical GENetics) Project financed through the Social European Fund through the Operational Program Human Capital 2014-2020. Priority axe 4- Social inclusion and poverty combat. Specific objective 4.8- Improvement of the competence level of professionals' in the medical field. SMIS107623; Contract POCU: 91/4/8/107623/08.12.2017 (12.2017-12.2019)- ***formator 2017-2019*** 4. Competitiveness Operational Programme 2014-2020; priority axis 1 – Research, technological development and innovation (RD&I) to support economic competitiveness and business development action 1.1.4 Attracting high-level personnel from abroad in order to enhance the RD capacity; Title: Use of nutrigenomic models for the personalized treatment with medical foods in obese people (NutriGen) 2016-2019- ***voluntar 2017-2019*** 5. ACT1373: Evaluate the Safety, Pharmacodynamics, Pharmacokinetics, and Exploratory Efficacy of GZ/SAR402671 in Treatment-naïve Adult Male Patients With Fabry Disease- NCT02228460 - ***coinvestigator*** in perioada Decembrie 2014 - Ianuarie 2016, Garches, Franta 6. Clinical, genetic and epigenetic aspects involved in the etiology of Prader Willi/ Angelman syndromes:

	multidisciplinary approach model of rare disorders in Romania- **membru activ 2009-2011** Project CNMP Parteneriate, Contract 41113, 2009-2011
Premii	**Iulia E. Jurca-Simina**, François Vialard, Karelle Benistan, Lucia Echevarria, Philippe de Mazancourt, Alessandro P. Burlina, Dominique P. Germain, "*A 16 Mb deletion at Xp22.13 leads to highly skewed X chromosome inactivation and consequent severe phenotypic expression of Fabry disease in a heterozygous female*"- 14th European Round Table on Fabry Disease – Fabry PhD Research Initiative, **Paris, France**- Abstract, poster and oral presentation- **Fabry PhD Research Initiative- award- 13-14 March 2015**
Cursuri si conferințe	- "EUROPLAN – European Project for Rare Diseases National Plans Development", Bucharest, Romania- 18th-19th June 2010 - The course of "Interdisciplinary approach of genetic rare diseases", Timisoara, Romania- 20th- 22th September 2010 - "The Third National Medical Genetic Conference with International Participation", Timisoara, Romania- 22th-25th September 2010 - The Workshop "Gene Therapy", Timisoara, Romania- 3th November 2010 - National Conference of Pediatrics, "Emergencies and chronic diseases", Bucharest, Romania March 2012 - European Society for Pediatric Nephrology- 45th Annual Scientific Meeting, Krakow, Poland- 6th-8th September 2012 - Course "Quality Management in Medical Sector and Institutional Communication", Oradea, Romania- 25th-27th April 2013 - The 7th German- Romanian Genetics Course: "Medical genetics, today", Oradea, Romania - 30th August- 1th September 2013 - The 7th Medical Genetics National Conference with international participation, Sibiu- Paltinis, Romania- 26th- 28th September 2013 - Fabry Masterclass VI, Prague, Czech Republic - 16th-17th May 2014 - 14th European Round Table on Fabry Disease, Paris, France – 13th-14th March 2015 - European Human Genetics Conference 2015, Glasgow, England, June 2015 - European Human Genetics Conference 2017, Copenhagen, Denmark, May 2017 - „Colloque international Les Territoire de la Sante: Production agroalimentaire, Nutrition, Securite alimentaire- PaNSaTS", Timisoara, Romania, October 2017 - „Personalised genomics in Pediatric Nephrology: from the lab bench to the bedside"- The International Pediatric Nephrology Association (IPNA) teaching course, Bucharest, Romania- 17th-18th November 2017. - ICGEB Workshop on "Next Generation Diagnostics", Skopje, Republic of Macedonia, 22 -24 March 2018. - Manchester Dysmorphology Course, Nowgen Centre, Manchester, UK, 17th - 19th April 2018. - European Human Genetics Conference 2018, Milan, Italy, June 2018 - Workshop Rețele Europene de Referință, Zalau, Romania, 2018 - Evocative signs in clinical genetics Course, Gura Humorului, Romania, September 2018 - European Human Genetics Conference 2019, Gothenburg, Sweden, June 2019 - Conferinta de pediatrie Pedipractic, online 30.09-02.10.2020 - „Training on strategies to foster solutions of undiagnosed rare disease cases"- 12-14 April 2021, ISS Rome Italy - European Human Genetics Conference, August 28–31, 2021 editie online - Congresul National de Pediatrie, Craiova, 15 – 18 septembrie 2021 - Zilele OncoHelp: "Managementul multidisciplinar al patologiei oncologice- Noutăți în diagnostic și tratament" Ediția a VI-a /26 - 28 Noiembrie 2021
Membru in societăți medicale	- Societatea Romana de Genetica Medicala SRGM - European Society of Medical Genetics ESHG - Societatea Nationala Romana de Pediatrie SNRPed

ANEXE

CAPITOLE CARTI

1. Cristina Gug, Maria Puiu, **Iulia Jurca-Simina**. Ghiduri şi îndrumătoare de laborator: Génétique médicale- Travaux pratiques pour les étudiants en Médecine Générale, Editura Victor Babeş, Timişoara, CNCSIS: 324 © 2020, ISBN 978-606-786-212-6
2. Dominique P. Germain, **Iulia E. Jurca-Simina**. Principles of Human Genetics and Mendelian Inheritance. In A. P. Burlina (Ed.), Neurometabolic Hereditary Diseases of Adults: Diagnosis and treatment, Springer International Publishing AG, part of Springer Nature 2018(I):1-28, https://doi.org/10.1007/978-3-319-76148-0

LISTA COMPLETĂ A LUCRĂRILOR

I. Articole publicate in extenso:

a. Articole publicate în reviste cotate ISI, cu factor de impact

Prim autor	
1.	**Jurca-Simina IE,** Jugănaru I, Iurciuc MŞ, Iurciuc S, Ungureanu E, Dobrescu AI, Chiriţă-Emandi A, Voinescu OR, Olariu IC, Puiu M, Georgescu D, Borugă VM. *What if body fat percentage association with FINDRISC score leads to a better prediction of type 2 diabetes mellitus?,* **Rom J Morphol Embryol, 2019**, 60(1):205–210, PMID: 31263846, **(FI/2018 = 1,5)**
Coautor	
1.	NCD Risk Factor Collaboration (NCD-RisC, including **Jurca-Simina IE** in the writing group). *Heterogeneous contributions of change in population distribution of body mass index to change in obesity and underweight.* **Elife. 2021** Mar 9;10:e60060. doi: 10.7554/eLife.60060 **(FI/2021=8.14)**
2.	NCD Risk Factor Collaboration (NCD-RisC, including **Jurca-Simina IE** in the writing group), *Height and body-mass index trajectories of school-aged children and adolescents from 1985 to 2019 in 200 countries and territories: a pooled analysis of 2181 population-based studies with 65 million participants,* **The Lancet**, Vol. 396, Issue 10261, November, **2020**, Pages 1511-1524 **(FI/2020=60.392)**
3.	Oliveira JP, Nowak A, Barbey F, Torres M, Nunes JP, Teixeira-e-Costa F, Carvalho F, Sampaio S, Tavaresj I, Pereira O, Soares AL, Carmona C, Cardoso MT, **Jurca-Simina IE**, Spada M, Ferreirab S, Germain DP. *Fabry disease caused by the GLA p.Phe113Leu (p.F113L) variant: Natural history in males,* **Eur J Med Genet., 2020** Feb;63(2):103703. Doi: 10.1016/j.ejmg.2019.103703. Epub 2019 Jun 11. PMID: 31200018 **(FI/2020= 4.246)**
4.	Beth L. Thurberg, Dominique P. Germain, Fernando Perretta, **Iulia E. Jurca-Simina**, Juan M. Politei. *Fabry disease: Four case reports of meningioma and a review of the literature on other malignancies*, **Molecular Genetics and Metabolism Reports**, Volume 11, **2017**, Pages 75-80, ISSN 2214-4269, https://doi.org/10.1016/j.ymgmr.2016.09.005. **(FI/2018=1.354)**
5.	Gafencu M, **Jurca-Simina IE**, Costa R, Doros G. *Distal renal tubular acidosis in AIDS young woman with wasting syndrome,* **Int Urol Nephrol*, 2014,*** 46(12): 2423-2427, PMID: 25298139. DOI: 10.1007/s11255-014-0840-9 **(FI/2014=1,293)**

b. Articole publicate în reviste cotate ISI, fără factor de impact

Coautor	
1.	Roman Deiana, Gug Miruna, Gliga Petra, Chircă Corina, **Jurca-Simina Iulia**, Jurca Maria Claudia, Vaida Monica. *Monogenic Cause For Renal Tubulopathies -Considerations Regarding Four Cases in Fertile Women.* **Proceedings of the 4 th Congress of the Romanian Society for Minimal Invasive Surgery in Ginecology**, pages 513-517. ISBN 978-88-85813-48-9

c. Articole publicate în reviste indexate BDI

Prim autor	
1.	**Jurca-Simina IE**, Chirita-Emandi A, Andreescu N, Farcaș S, Mihailescu A, Popa AM, Tutac P, Zimbru C, Dobrescu AI, Perva IT, Murariu A, Puiu M. *Burden of rare genetic diseases –experience of Timis Regional Centre of Medical Genetics, Romania,* **Jurnalul pediatrului, 2019**, XXII (85-86): 56-65. ISSN 2065 – 4855, **Index Copernicus since 2010, CNCSIS B+**
2.	**Jurca-Simina IE**, Chirita Emandi A, Perva IT, Uhrová Mészárosová A, Corches A, Doros G, Puiu M. *Think about the founder effect in endogamous population - Congenital cataracts, Facial dysmorphism, and Neuropathy (CCFDN) Syndrome - two cases,* **Jurnalul pediatrului, 2018**, XXI(81-82): 19-25. ISSN 2065 – 4855, **Index Copernicus since 2010, CNCSIS B+**

Coautor	
1.	Sabau, I. M., Andreescu, N. I., Chiriță-Emandi, A., **Jurca-Simina, I.**, Bugi, M. A., & Puiu, M. *Genetics in anorexia nervosa*, **Jurnalul pediatrului, 2021,** XXIV (93-94): 23-27. ISSN 2065 – 4855 https://doi.org/10.37224/JP.2021.9394.05 **, Index Copernicus since 2010, CNCSIS B+**
2.	Sabau, I. M., Andreescu, N. I., Chiriță-Emandi, A., **Jurca-Simina, I.**, Bugi, M. A., & Puiu, M. *Ketogenic diet and genetic disorders*, **Jurnalul pediatrului**, **2021**, XXIV (93-94): 28-33. ISSN 2065 – 4855 https://doi.org/10.37224/JP.2021.9394.06 , **Index Copernicus since 2010, CNCSIS B+**
3.	Adela Chirita-Emandi, Gabriela Doros, **Iulia Jurca Simina**, Mihai Gafencu, Maria Puiu, *Head circumference references for school age children in western Romania*, **Rev. Med. Chir. Soc.Med. Nat., Iasi, 2015,** 119 (4): 1083-1091, **Index Copernicus since 2010, CNCSIS B+**
4.	Corina Pienar, Maria Puiu, Adela Chirita-Emandi, Simona Dumitriu, Cristina Popa, **Iulia Jurca-Simina**, Ioana Micle, Smaranda Arghirescu; *Childhood obesity: between nature and nurture*; **Jurnalul Pediatrului, 2013**, XVI (61-62): 3-8, ISSN 1221-7212, **Index Copernicus since 2010, CNCSIS B+**
5.	Mihai Gafencu, **Iulia Simina Jurca**, Laura Leahu, Andra Mitoceanu, Otilia Marginean, Gabriela Doroș, Bogdan Korbuly. *Overweight pathology in children form Timis County.* **Jurnalul Pediatrului, 2013**, XVI (63): 27-31, ISSN 2065 – 4855, **Index Copernicus since 2010, CNCSIS B+**
6.	Puiu M., **Jurca Simina I**., Dumitriu S., Arghirescu S., Chirita-Emandi A. *Multiple hereditary exostoses-Clinical features and management.* **Jurnalul Pediatrului**, **2012**, XV (57-58): 64-9, **Index Copernicus since 2010, CNCSIS B+**

d. Lucrări publicate în rezumat

i. Lucrări publicate în rezumat la manifestări ştiinţifice internaţionale

Prim autor	
1.	**Jurca-Simina, I. E**.; Chirita-Emandi, A.; Andreescu, N.; Serban, C. L.; Zimbru, C.; Puiu, M. *Molecular genetic diagnostic in skeletal disorders - a Western Romanian delineation.* **European Journal of Human Genetics**, Volume 28, Issue SUPPL, Page 840-841, Supplement 1, Meeting Abstract E-P04.49, Published DEC **2020**, Indexed 2021-01-19, ISSN 1018-4813, eISSN 1476-5438
2.	**Jurca-Simina, I.;** Chirita-Emandi, A.; Andreescu, N.; Olariu, N.; Isac, R.; Farkas, F.; Andrei, Z.; Gafencu, M.; Puiu, M. *Bilateral Multicystic Dysplastic Kidney in a three-generation family.* **European Journal of Human Genetics**, Volume 27, Page 905-905, Supplement 1, Meeting Abstract E-P03.27, Published JUL **2019**, Indexed 2019-10-23, ISSN 1018-4813, eISSN 1476-5438
3.	**Jurca-Simina, I. E.;** Rabes, J.; Richard, P. A.; Jauny, C.; Koraichi, F.; Carlier, R.; Hagege, A. A.; de Mazancourt, P.; Puiu, M.; Germain, D. P. *Pitfalls in the diagnosis of Fabry disease: further evidence that p.Asp313Tyr is a non-pathogenic polymorphism.* **European Journal of Human Genetics**, Volume 26, Page 291-291, Supplement S, Meeting Abstract P06.09A, Published OCT **2018**, Indexed 2018-10-01, 50th European-Society-of-Human-Genetics (ESHG) Conference, Copenhagen, DENMARK, ISSN 1018-4813, eISSN 1476-5438
4.	**Jurca-Simina Iulia-Eena,** Chirita-Emandi Adela, Andreescu Nicoleta, Cristian Zimbru, Urtila Patricia, Ioana Micle, Puiu Maria. *Heterozygous known mutation in LPL gene causing Lipoprotein Lipase Deficiency with severe Hypertriclyceridemia in a child.* **Balkan Journal of Medical Genetics**, vol 21, **2018**, supplement 1, ISSN1311-0160, p57 (ICGEB Workshop "Next Generation Diagnostics", Skopje, March 22-24,2018)
5.	**I. E. Jurca- Simina**, R. M. Jurac, M. Cucuruz, C. Jinca, E. Boeriu, C. Popa, S. Arghirescu, M. Puiu. *Particularities of ATRA therapy in pediatric patients with acute promyelocytic leukemia.* **European Journal of Human Genetics**, Volume 22, Supplement 1, May **2014**, Milan, page 493, J15.17
6.	**I. E. Jurca- Simina,** M. Puiu, M. Gafencu. *Renal disease's genetic counseling- a must for an affected family*, **European Journal of Human Genetics**, Volume 21 Supplement 2, p 424, ISSN1018-4813, **2012** Impact Factor-4.319.pag 576
Coautor	
1.	Serban, C. L.; Andreescu, N.; **Jurca-Simina, I**.; Corches, A.; Emandi, A. Chirita; Puiu, M.. *Elucidating myopathies with high creatine-kinase- from unsolved cases to common diagnosis.* **European Journal of Human Genetics**, Volume 28, Issue SUPPL, Page 430-430, Supplement 1, Meeting Abstract P10.17.C, Published DEC **2020**, Indexed 2021-01-19, ISSN 1018-4813, eISSN 1476-5438
2.	Marcovici, T.; Puiu, M.; Bacos, C.; **Jurca-Simina, I**.; Belei, O.; Marginean, O.; Grozavu, A. *Classic Dravet Syndrome in an adolescent male - case report.* **European Journal of Human Genetics**, Volume 27, Page 963-963, Supplement 1, Meeting Abstract E-P09.18, Published JUL **2019**, Indexed 2019-10-23, ISSN 1018-4813, eISSN 1476-5438
3.	Manea, A.; **Jurca-Simina, I**.; Cioboata, D.; Costescu, O.; Doandes, F.; Lungu, N.; Boia, M. *Rapid and optimal diagnosis in malformative syndromes at newborns*. **European Journal of Human Genetics**, Volume

	27, Page 1882-1883, Supplement 2, Meeting Abstract E-P11.38, Published JUL **2019**, Indexed 2019-10-23, ISSN 1018-4813, eISSN 1476-5438
4.	D.P. Germain, J.-B. Riviere, I. Dabaj, J. Bataille , C. Jauny, **I.E. Jurca-Simina**, L. Faivre And I. Haegy. *Clove syndrome: a case report*, **Twenty-sixth European Meeting on Dysmorphology**, 9 – 11 September **2015**, Le Bischenberg, France
5.	M. Gafencu, G. Doros, D. Dan, **I. Jurca Simina**, L. N. Bogdan, M. Puiu. *Rare Diseases week in Timisoara - a campaign with a good start*. **European Journal of Human Genetics**, Volume 22, Supplement 1, May **2014**, Milan, page 346, P18.40-M
6.	Gafencu, Mihai; Doros, Gabriela; Costa, Rodica; Schiller, Adalbert; Kundani, Nilima; **Jurca-Simina, Iulia Elena**. *Renal involvement in HIV infected Romanian children.* **Pediatric Nephrology**, Volume 27, Issue 9, Page 1690-1691, Published SEP **2012**, Indexed 2012-09-12, Meeting Abstract, ISSN 0931-041X
7.	M. Gafencu, R. Costa, G. Doros, K. Nilima, A. Schiller, **I. Jurca- Simina.** *Renal involvement in HIV infected children*, **Pediatric Nephrology**, vol 27 issue 9, pp 1605-1829(**2012**), The 45th Annual Meeting on September 6th – 8th 2012, Krakow, Poland, ISSN online 1432-198X
8.	Doros G., Popoiu A., Gafencu M., **Jurca-Simina I.E**., Leahu L., But A. *Risk factors for cardiovascular disease in school age children and teenagers,* **46th Annual Meeting of the AEPC**, Istanbul, Turkey, 23-26 May **2012**, **Cardiology in the Young**, Vol 22, Suppl. 1, p.111-112, ISSN 1047-9511, Impact factor 2012 0.948

ii. Lucrări publicate în rezumat la manifestări științifice naționale

Prim autor	
1.	**Jurca-Simina Iulia-Elena,** Chirita-Emandi Adela, Nicoleta Andreescu, Olariu Nicu, Isac Raluca, Jurca-Simina Florin-Ioan, Gafencu Mihai, Puiu Maria. *Autosomal Dominant Multicystic Dysplastic Kidney Phenotype – No Genotype Identified (Yet)*, **Abstract book of „Personalised genomics in Pediatric Nephrology: from the lab bench to the bedside"- The International Pediatric Nephrology Association (IPNA) teaching course and the VIIth Meeting of the Southeastern Europe Pediatric Nephrology Working Group (VIIth SEPNWG)**, Bucharest, Romania- 17th-18th of November **2017**, ISBN 978-973-0-25841-7.
2.	**Iulia Jurca-Simina,** Alexandra Mihăilescu, Iulia Perva, Alexandra Sima, Iulian Puiu Velea, Adela Chiriță-Emandi, Nicoleta Andreescu, Mihai Niculescu, Maria Puiu. *NutriGen. Utilisation de modèles nutrigénomiques pour personnaliser les traitements diététiques dans l'obésité*, **Colloque international Les Territoire de la Sante: Production agroalimentaire, Nutrition, Securite alimentaire- PaNSaTS**, **2017**, Timisoara, Romania
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