Listă de lucrări științifice

Alina Bogliș

Teza de doctorat - in derulare:

„Studiul mutațiilor genetice în dizabilitatea intelectuală la pacienții cu anomalii congenitale multiple". Conducator stiintific Prof.univ.Dr. Banescu Claudia, Universitatea de Medicină, Farmacie, Științe și Tehnologie „George Emil Palade" din Târgu Mureș.

Lucrare de disertatie:

„Aspecte genetice privind etiopatogenia malformațiilor congenitale". Conducator stiintific Prof.univ.Dr. Banescu Claudia, Târgu Mureș, 2016.

Capitole carti de specialitate – co-autor

1. Bănescu Claudia, Duicu Carmen, Todoran Anamaria, Moldovan Valeriu, **Bogliș Alina**, Tripon Florin, Crauciuc Andrei. *Boli multifactoriale*, in *Noțiuni de genetică fundamentală și genetică medicală*. Editura University Press, Târgu Mureș, 2015, pg 209-230, ISBN 978-973-169-430-6, cod CNCSIS 210. **(2,857 pct)**
2. Robert Nussbaum, Roderick McInnes, Huntington Willard, Laurențiu Bohîlțea, Roxana Bohîlțea, etc. Thompson and Thompson. Medical Genetics Ed. 8th - Robert L. Nussbaum. Romanian Edition. Editure Hipocrate, 2018, ISBN: 9786069457504.

Articole publicate in extenso in reviste cotate ISI, cu factor de impact (FI) – autor principal/corespondent

1. **Boglis A**, Cosma SA, Tripon F, Bănescu C. *Exon 21 deletion in the OPHN1 gene in a family with syndromic X-linked intellectual disability: Case report.* Medicine (Baltimore) 2020, DOI: 10.1097/MD.0000000000021632, ISSN 0025-7974, **FI: 1.552**, Ahead of print, autor principal, corespondent. **(38,25 pct)**
2. Tripon F, **Bogliș A,** Micheu C, Streață I, Bănescu C. *Pitt-Hopkins Syndrome: Clinical and Molecular Findings of a 5-Year-Old Patient.* Genes (Basel) 2020, 28;11(6):596, DOI: 10.3390/genes11060596, ISSN 2073-4425, **FI: 3,759**, autor corespondent. **(71,385 pct)**
3. **Boglis A,** Tripon F, Bănescu C. *The utility of molecular genetic techniques in craniosynostosis cases associated with intellectual disability.* Rev Romana Med Lab, 2018;26(4):471-477, DOI: 10.2478/rrlm-2018-0033, ISSN 2284-5623, **FI: 0,8**, autor principal, corespondent. **(27 pct)**

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1. Tripon F, Crauciuc GA, **Bogliş A,** Moldovan V, Sándor-Kéri J, Benedek IJ, Trifa AP, Bănescu C. *Co-occurrence of PML-RARA gene fusion, chromosome 8 trisomy, and FLT3 ITD mutation in a young female patient with de novo acute myeloid leukemia and early death: A CARE case report.* Medicine (Baltimore) 2020, 99(14):e19730. DOI: 10.1097/MD.0000000000019730, ISSN 0025-7974, **FI: 1.552**, co-autor. **(4,785 pct)**
2. Bănescu C, Tripon F, Trifa AP, Crauciuc AG, **Bogliș A,** Lazar E, Dima D, Macarie I, Duicu C, Iancu M. *Presence of copy number aberrations and clinical prognostic factors in patients with acute myeloid leukemia: an analysis of effect modification. Authors' reply.* Pol Arch Intern Med 2020, 130(4):347-348. DOI: 10.20452/pamw.15327, ISSN 0032-3772, **FI: 3,007,** co-autor. **(6,010 pct)**
3. Tripon F, Iancu M, Trifa A, Crauciuc GA, **Boglis A**, Dima D, Lazar E, Bănescu C. *Modelling the Effects of MCM7 Variants, Somatic Mutations, and Clinical Features on Acute Myeloid Leukemia Susceptibility and Prognosis.* J Clin Med 2020, 9(1):158. DOI: 10.3390/jcm9010158, ISSN 2077-0383, **FI: 3,303,** co-autor. **(8,068 pct)**
4. Tripon F, Iancu M, Trifa A, Crauciuc GA, **Boglis A,** Balla B, Cosma A, Dima D, Candea M, Lazar E, Jimbu L, Banescu C. *Association Analysis of TP53 rs1042522, MDM2 rs2279744, rs3730485, MDM4 rs4245739 Variants and Acute Myeloid Leukemia Susceptibility, Risk Stratification Scores, and Clinical Features: An Exploratory Study.* J Clin Med 2020, 9(6):1672. DOI: 10.3390/jcm9061672, ISSN 2077-0383, **FI: 3,303,** co-autor. **(5,378 pct)**
5. Bănescu C, Tripon F, Trifa AP, Crauciuc AG, Moldovan VG, **Bogliş A,** Benedek I, Dima D, Cândea M, Duicu C, Iancu M. *Cytokine rs361525, rs1800750, rs1800629, rs1800896, rs1800872, rs1800795, rs1800470, and rs2430561 SNPs in relation with prognostic factors in acute myeloid leukemia*. Cancer Med 2019, 8(12):5492-5506. DOI: 10.1002/cam4.2424, ISSN 2045-7634, **FI: 3,491**, co-autor. **(6,124 pct) ARTICOL PEREMIAT UEFISCIDI**
6. Meliţ LE, Mărginean CO, Bănescu C, **Bogliş A**, Mocan S, Iancu M. *The relationship between TLR4 rs4986790 and rs4986791 gene polymorphisms and Helicobacter pylori infection in children with gastritis.* Pathol Res Pract 2019, 215(12):152692. DOI: 10.1016/j.prp.2019.152692, ISSN 1618-0631, **FI: 2,050**, co-autor. **(7,625 pct)**
7. Tripon F, Crauciuc GA, Moldovan VG, **Bogliş A**, Benedek IJ, Lázár E, Bănescu C. *Simultaneous FLT3, NPM1 and DNMT3A mutations in adult patients with acute myeloid leukemia – case study.* Rev Romana Med Lab 2019, 27(3):245-54. DOI:10.2478/rrlm-2019-0022, ISSN 2284-5623, **FI: 0,945**, co-autor. **(4,167 pct)**
8. Crauciuc GA, Tripon F, **Bogliş A**, Făgărăşan A, Bănescu C. *Multiplex ligation dependent probe amplification - a useful, fast and cost-effective method for identification of small supernumerary marker chromosome in*

children with developmental delay and congenital heart defect. Rev Romana Med Lab 2018, 26(4):461-670. DOI:10.2478/rrlm-2018-0032, ISSN 2284-5623, **FI: 0,8,** co-autor. **(5,4 pct)**

9. Bănescu C, Iancu M, Trifa AP, Dobreanu M, Moldovan VG, Duicu C, Tripon F, Crauciuc A, Skypnyk C, **Bogliș A,** Lazar E. *Influence of XPC, XPD, XPF, and XPG gene polymorphisms on the risk and the outcome of acute myeloid leukemia in a Romanian population.* Tumour Biol 2016, 37(7):9357-66. DOI: 10.1007/s13277-016-4815-6, ISSN 1010-4283, **FI: 2.926**, co-autor. **(5,353 pct)**

Articole publicate in extenso in reviste cotate ISI, fara factor de impact (FI)

1. **Bogliș A**, Zeleniuc M, Calvente M, Tripon F, Crauciuc GA, Duicu C, Bănescu C. *A novel 2q11.2q14.3 duplication resulting from a small supernumerary marker chromosome associated with developmental delay, intellectual disability, and congenital heart malformation.* Proceedings of 5th Medical Genetics Congress with International Participation, Filodiritto Editore-Proceedings 2019, 2:53-59. ISBN 978-88-85813-54-0, fara FI, autor principal. **(15 pct)**
2. **Bogliș A**, Radu GC, Tripon F, Crauciuc GA, Demian S, Duicu C, Bănescu C.. *XRCC1 Arg194Trp and Arg399Gln polymorphisms and risk of non-Hodgkin lymphoma in a Romanian population*. Rev Med Chir Soc Med Nat Iasi 2016, 120(3):644-650, ISSN 0048-7848, fara FI, autor principal. **(15 pct)**

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2. **Boglis A**, Crauciuc AG, Tripon F, Radu CG, Demian S, Duicu C, Banescu C. *No association between GSTT1, GSTM1, and GSTP1 gene polymorphism and risk of non-Hodgkin lymphoma in a population from Romania.* IJIAS 2017; 19(1):1-8, ISSN 2028-9324, autor principal. **(10 pct)**
3. Crauciuc A, Tripon F, Gheorghiu A, Nemes G, **Boglis A,** Banescu C. *Development, Applications, Benefits, Challenges and Limitations of the New Genome Engineering Technique. An Update Study.* Acta Medica Marisiensis 2017, 63(1), 4-9, DOI: 10.1515/amma-2017-0007, ISSN 2668-7763, co-autor. **(1,66 pct)**
4. **Boglis A**, Rac CD, Moldovan E, Duicu C, Bănescu C. *A Rare Chromosomal Disorder – 14q Interstitial Deletion Syndrome.* Acta Medica Marisiensis 2016, 62(3):378-380, DOI: 10.1515/amma-2016-0041, ISSN 2668-7763, autor principal. **(10 pct)**
5. Bodi S, Mircea O, Irimia T, Stanca M, **Bogliș A,** Pușcașiu L. *Cesarean section and placenta previa – causation or just association*. Revista de Obstetrica si Ginecologie 2016, LXIV(2):81-85. ISSN: 1220-5532,

co-autor. **(1,66 pct)**

6. Bădescu A, Mircea O, Irimia T, Bodi S, **Bogliș A**, Pușcașiu L, Soldea V. *Deep endometriosis infiltrating the bowel – a continuing debate about the best management.* Revista de Obstetrica si Ginecologie 2016, LXIV(2):103-108. ISSN: 1220-5532, co-autor. **(1,42 pct)**

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1. **Bogliș A,** Tripon F, Crauciuc AG, Moldovan V, Duicu C, Bănescu C. *A de novo FGFR3 mutation in Crouzon syndrome: a case report and review of the literature.* European Journal of Human Genetics 2018, 26: 955-955. ISSN 1018-4813 **(4 pct)**
2. **Bogliș A,** Tripon F, Crauciuc A, Bănescu C. *MLPA versus conventional karyotyping in the investigation of the genomic rearrangements.* Rev Romana Med Lab, 2018; Supplement 26(2):87-88. ISSN 2284-5623. **(4 pct)**
3. **Bogliș A,** Tripon F, Crauciuc AG, Balla BM, Cosma AS, Bănescu C. *The use of MLPA analysis in predicting the tyrosine kinase inhibitors response in chronic myeloid leukemia patients*. Rev Romana Med Lab, 2018; Supplement 26(4):12. ISSN 2284-5623. **(4 pct)**
4. **Boglis A,** Tripon F, Moldovan V, Crauciuc A, Banescu C. *Genetic testing through multiplex ligation dependent probe amplification analysis for children with global developmental delay or intellectual disability.* European Journal of Human Genetics „The European Human Genetics Conference 2018, Milan, Italy, June 16-19, 2018". ISSN 2284-5623. **(4 pct)**
5. **Boglis A,** Tripon F, Banescu C. *Cytogenetic study in children with congenital heart defects and multiple congenital anomalies: ten years of experience.* Eur J Hum Genet 2016, Supplement 24E(1):231, ISSN 1018-4813. **(4 pct)**
6. **Bogliș A**, Moldovan V, Bănescu C. *Genetic investigations in patients with developmental delay and congenital anomalies*. Rev Romana Med Lab 2016, Supplement 24(1):69-70, ISSN 2284-5623. **(4 pct)**
7. **Bogliș A,** Kulcsar A, Moldovan V, Bănescu C. *Cytogenetic abnormalities in pediatric patients with multiple congenital anomalies.* Rev Romana Med Lab 2015, Supplement 23(1):106-107, ISSN 2284-5623. **(4 pct)**

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1. Tripon F, Crauciuc AG, Duicu C, **Bogliș A,** Moldovan V, Demian S, Bănescu C. *No association between TNF alpha A308G gene polymorphism and FLT3 genes mutation in patients with acute myeloid leukemia.* European Journal of Human Genetics 2018, 26: 972-973, ISSN 1018-4813, co-autor. **(0,57 pct)**
2. Bănescu C , Crauciuc AG, Moldovan V, **Bogliș A,** Tripon F, Lazar Benedek E, Duicu C. *No influence of XPC gene polymorphisms on FLT3 or DNMT3Amutations in acute myeloid leukemia patients.* European Journal of Human Genetics 2018, 26: 611-611, ISSN 1018-4813, co-autor. **(0,57 pct)**

3. Crauciuc AG, Tripon F, **Bogliș A**, Moldovan V, Duicu C, Candea M, Bănescu C. *Role of XPC Ala499Val polymorphisms in chronic myeloid leukemia in Romanian patients.* European Journal of Human Genetics 2018, 26: 977-978, ISSN 1018-4813, co-autor. **(0,57 pct)**
4. Banescu C, Crauciuc A, Moldovan V, **Boglis A**, Lazar E, Tripon F. *No association between TERT rs2853669 polymorphism and NPM1, DNMT3A gene mutations and acute myeloid leukemia risk.* European Journal of Human Genetics „The European Human Genetics Conference 2018, Milan, Italy, June 16-19, 2018". ISSN 1018-4813. **(0,66 pct)**
5. Duicu C, Tripon F, Crauciuc AG, Aldea C, **Bogliș A,** Moldovan V, Bănescu C. *Investigarea genetică a polimorfismelor genenelor MDR1 C3435T, MDR1 C 1236T şi NPHS2 R229Q la copiii cu sindrom nefrotic.* Rev Romana Med Lab 2017; Supliment 25(2):50-51. ISSN 2284-5623. **(0,57 pct)**
6. Duicu C, Tripon F, Crauciuc AG, Aldea C, **Bogliș A,** Moldovan V, Bănescu C. *Investigarea polimorfismelor genelor TNF alpha şi IL-6 la copiii cu sindrom nefrotic.* Rev Romana Med Lab 2017; Supliment 25(2):52-53. ISSN 2284-5623. **(0,57 pct)**
7. Tripon F, Crauciuc AG, Aldea C, **Bogliș A,** Moldovan V, Bănescu C, Duicu C. *Asocierea polimorfismului VEGF +936C/T cu susceptibilitatea pentru sindromul nefrotic la copii.* Rev Romana Med Lab 2017; Supliment 25(2):53-54. ISSN 2284-5623. **(0,57 pct)**
8. Bănescu C, Crauciuc AG, Moldovan V, **Bogliș A,** Tripon F, Demian S, Duicu C. *Polimorfismul TGF-β1 T869C este asociat cu mutația FLT3 la pacienții cu LAM.* Rev Romana Med Lab 2017; Supliment 25(2):30-31. ISSN 2284-5623. **(0,57 pct)**
9. Crauciuc G, Tripon F, **Bogliș A,** Duicu C, Bănescu C. *Association between Glutathione S-transferase P1 Ile105Val gene polymorphisms and Ann Arbor stage in Lymphoma*, European Human Genetics Conference Barcelona, May 2016, European Journal of Human Genetics, 2016; 24 (E Sup. 1):468. ISSN 1018-4813. **(0,8 pct)**
10. Bănescu C, Moldovan VG, Tripon F, Crauciuc G, Duicu C**, Bogliș A**, Dobreanu M. *XPF -673C>T and XPF 11985A>G gene polymorphisms and risk of chronic myeloid leukemia,* European Human Genetics Conference Barcelona, May 2016, European Journal of Human Genetics, 2016; 24 (E Sup. 1):472. ISSN 1018-4813. **(0,57 pct)**
11. Bănescu C, Moldovan GV, Tripon F, Crauciuc A, Duicu C, **Bogliș A**, Dobreanu M. *Single nucleotide polymorphisms XPC LYS939GLN and XPG ASP1104HIS in romanian patients with chronic myeloid leukemia*, Rev Romana Med Lab 2016; 24 (Supplement 1): S57. ISSN 2284-5623. **(0,57 pct)**
12. Moldovan GV, Tripon F, Crauciuc A, **Bogliș A**, Duicu C, Dobreanu M, Bănescu C. *Short overview on MLPA*. Rev Romana Med Lab 2016; 24 (Sup. 1):S70. ISSN 2284-5623. **(0,57 pct)**
13. Moldovan GV, Duicu C, **Bogliș A,** Crauciuc A, Tripon F, Bănescu C. *Tumor necrosis factor-α, a risck factor for acute myeloid leukemia development.* Rev Romana Med Lab 2015, Supplement 23(1):100-101. ISSN 2284-5623. **(0,66 pct)**

14. Banescu C, Tripon F, Crauciuc A, Moldovan V, **Boglis A**, Demian S, Lazar E, Duicu C, Trifa PA. *Investigation of microdeletions and somatic mutations with prognostic impact in patients with acute myeloid leukemia.* Rev Romana Med Lab, 2018; Supplement 26(2):61-62. ISSN 2284-5623. **(0,44 pct)**
15. Cosma AS, Tripon F, Crauciuc A, Moldovan GV, **Bogliș A**, Demian S, Lazar E, Duicu C, Trifa PA, Bănescu C. *Investigation of XPC Lys939Gln gene polymorphism in chronic myeloproliferative neoplasms: a case-control study.* Rev Romana Med Lab, 2018; Supplement 26(2):83-84. ISSN 2284-5623. **(0,4 pct)**
16. Crauciuc A, Tripon F, **Bogliș A**, Muntean I, Bănescu C. *Investigation of VEGF -2549 I/D gene polymorphism in congenital heart defects.* Rev Romana Med Lab, 2018; Supplement 26(2):63-64. ISSN 2284-5623. **(0,8 pct)**

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1. **Boglis A,** Tripon F, Cosma AS, Banescu C. *Subtelomeric copy number variations in patients with intellectual disability/developmental delay detected by multiplex ligation-dependent probe amplification analysis.* Book of Abstracts 2019, 2:117, ISSN 2602-1609, The 12th Conference of Ph.D. Students and Young Doctors, December 11th, Targu Mures. **(2 pct)**
2. **Bogliș A,** Zeleniuc M, Tripon F, Crauciuc AG, Duicu C, Bănescu C. *Duplication of 2q11.2q14.3 resulting from a small supernumerary marker chromosome in a patient with developmental delay, intellectual disability and congenital heart malformation.* Romanian Journal of Rare Diseases, 2018; Supplement 1:67-68. ISSN 2068–5882. **(2 pct)**
3. **Bogliș A,** Crauciuc AG, Tripon F, Cosma AS, Balla BM, Bănescu C. *CHARGE syndrome due to a de novo 13q deletion – a case report*. Book of abstracts, Nr. 1/2018, ISSN 2602-1609, The 11th Conference of Ph.D. Students and Young Doctors, December 11th, 2018. **(2 pct)**
4. **Bogliș A,** Tripon F, Crauciuc AG, Moldovan V, Trifa A, Lazar E, Bănescu C. *Prognostic value of ABCG2 gene Q141K polymorphism in Romanian patients with acute myeloid leukemia with FLT3 mutation.* Romanian Journal of Rare Diseases 2017; Supplement 2:38. ISSN 2068–5882. **(2 pct)**
5. **Bogliș A,** Tripon F, Moldovan V, Crauciuc AG, Bănescu C. *Detection of genomic imbalances in children with intellectual disability using MLPA analysis in Tîrgu Mureș.* Acta Medica Marisiensis 2017; Supplement 63(4):108. ISSN 2068-3324 **(2 pct)**
6. **Boglis A**, Banescu C. *Genotype-phenotype correlations in structural abnormalities of chromosome 18.* Acta Medica Marisiensis 2016; 62(Supplement 8):25-26. ISSN 2068-3324 **(2 pct)**
7. **Boglis A,** Banescu C. *Genetic abnormalities in the congenital long QT syndrome.* Romanian Journal of Rare Diseases 2016; Supplement 1:30. ISSN 2068 – 5882 **(2 pct)**
8. **Boglis A**, Banescu C. - *18q21 deletion syndrome-a case report and clinical review of the literature.* Romanian Journal of Rare Diseases 2016; Supplement 1:31. ISSN 2068 – 5882 **(2 pct)**

9. **Boglis A,** Banescu C. *Chromosomal abnormalities in patients with sexual developmental disorders: a retrospective study*. Acta Medica Marisiensis 2016; Supplement 61(1):17. ISSN 2068-3324 **(2 pct)**
10. **Boglis A**, Banescu C. *A rare structural chromosomal abnormality detected by cytogenetic investigations in patients with congenital anomalies: the isochromosome*. Acta Medica Marisiensis 2016; Supplement 61(1):120. ISSN 2068-3324 **(2 pct)**
11. **Boglis A**, Banescu C. *Prenatal diagnosis of fetal cytogenetic abnormalities: a retrospective study.* Asklepsios 9th International Medical Congress for Students and Young Doctors, Sibiu, 2016, pg. 9. ISSN 1843-0406 **(2 pct)**
12. **Boglis A**, Moldovan GV, Banescu C. *Genetics in cystic fibrosis: review of clinical, molecular and therapeutic aspects.* Asklepsios 9th International Medical Congress for Students and Young Doctors, Sibiu, 2016, pg 30. ISSN 1843-0406 **(2 pct)**
13. **Boglis A**, Banescu C. *Clinical and cytogenetic study in patients with Turner syndrome.* Asklepsios 9th International Medical Congress for Students and Young Doctors, Sibiu, 2016, pg 100. ISSN 1843-0406 **(2 pct)**
14. **Boglis A,** Tripon F, Banescu C. *Genetic investigations in children with intellectual disability and congenital anomalies.* Orvos Kepzes. A gradualis es posztgradualis kepzes folyoirata 2016; XCI(1):17. ISSN 0030-6037 **(2 pct)**
15. **Boglis A**, Moldovan GV, Duicu C, Banescu C. *Diagnosis aspects in Beckwith-Widemann syndrome – genetic department experience from Tirgu Mures.* Romanian Journal of Rare Diseases 2015; Supplement 1:14. ISSN 2068-5882 **(2 pct)**
16. **Bogliș A**, Rac C, Mărginean O, Bănescu C. *Sindromul de deleție interstițială a cromozomului 14q- tulburare cromozomială rară.* Al XI-lea Congres Național de Pediatrie cu participare internațională, Târgu Mureș, 2013. ISSN 2344-3324 **(2 pct)**
17. **Bogliș A**, Mirică MN, Jankus HS, Gliga M. *Sindromul nefrotic la vârstnic-particularități etiologic*. Revista de Medicină și Farmacie Târgu Mureș 2010; Supliment 56(2):92-93. ISSN 1221-2229 **(2 pct)**

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1. Bănescu C, **Bogliș A**, Crauciuc A, Moldovan Valeriu, Tripon F. *The role of genetic testing in te managemnet of acute myeloid leukemia: from cytogenetic to molecular genetic*. Romanian Journal of Rare Diseases, 2018; Supplement 1:40. ISSN 2068 – 5882 **(0,4 pct)**
2. Anciuc M, Bănescu C, Tatar D, Cucerea MC, **Bogliș A.** *From theory to practice. A congenital heart disease and ambiguous genitalia case and literature review.* Romanian Journal of Rare Diseases 2017; Supplement 2:35. ISSN 2068–5882 **(0,4 pct)**
3. Crauciuc AG, Tripon F, Moldovan V, **Bogliș A**, Macarie I, Bănescu C. *TNF - A 238A>G gene*

polymorphisms are not associated with FLT3 ITD gene mutation and the risk of developing AML. Romanian Journal of Rare Diseases 2017; Supplement 2:18. ISSN 2068–5882 **(0,33 pct)**

4. Bănescu C, Crauciuc AG, Moldovan V, **Bogliș A**, Lazar Benedek E, Tripon F. *No association between copy number changes and FLT3 or DNMT3A gene mutations in acute myeloid leukemia*. Romanian Journal of Rare Diseases 2017; Supplement 2:37-38. ISSN 2068–5882 **(0,33 pct)**
5. Moldovan V, **Bogliș A,** Crauciuc AG, Tripon F, Bănescu C. *Influence of TNF - α rs1800750 gene polymorphism on the risk and relationship with prognostic factors (DNMT3A, FLT3 mutations) in acute myeloid leukemia.* Romanian Journal of Rare Diseases 2017; Supplement 2:49-50. ISSN 2068–5882 **(0,33 pct)**
6. Radu CG, Tripon F, Crauciuc AG, **Bogliș A,** Demian S, Bănescu C. *Oxidative stress gene GPX1 Pro197Leu polymorphism. a risk factor for non - Hodgkin lymphoma.* Romanian Journal of Rare Diseases 2017; Supplement 2:57-58. ISSN 2068–5882 **(0,33 pct) PREMIU CEL MAI BUN POSTER**
7. Radu CG, Tripon F, Crauciuc AG, **Bogliș A,** Demian S, Bănescu C. *CAT C262T and MNSOD Ala16Val polymorphisms and 5 years survival rates among non - Hodgkin lymphoma patients.* Romanian Journal of Rare Diseases 2017; Supplement 2:58-59. ISSN 2068–5882 **(0,33 pct)**
8. Tripon F, Crauciuc AG, **Bogliș A,** Moldovan V, Duicu C, Lazar E, Trifa PA, Bănescu C. *No association between TERT RS2736100 A>C SNP, FLT3 ITD and FLT3 D835 gene mutations and acute myeloid leukemia.* Romanian Journal of Rare Diseases 2017; Supplement 2:63-64. ISSN 2068–5882 **(0,25 pct)**
9. Bănescu C, Tripon F, Moldovan V, **Bogliș A**, Crauciuc AG, Trifa PA. *Investigation of NPM1 type a mutation by HRM, CASTPCR and allele specific RT-PCR methods in acute myeloid leukemia.* Acta Medica Marisiensis 2017; Supplement 63(4):11. ISSN 2068-3324 **(0,33 pct)**
10. Tripon F, Crauciuc AG, **Bogliș A**, Duicu C. Bănescu C. *MDR1 3435T> C gene polymorphisms and risk of nephrotic syndrome in children.* Acta Medica Marisiensis 2016, 62:25, ISSN 2668-7763. **(0,4 pct)**
11. Bănescu C, Tripon F, Moldovan V, Cândea M, **Bogliș A,** Crauciuc A. *Cytokine gene polymorphisms and risk of chronic myeloid leukemia in a Romanian population: a case-control study.* 14th National Symposium Of Microscopic Morphology With International Participation, Tîrgu-Mureş 4-6 May 2016, Acta Medica Marisiensis, Vol 61. ISSN 2068-3324 **(0,33 pct)**
12. Bănescu C, Tripon F, Cândea M, **Bogliș A,** Crauciuc A, Moldovan. *TNF-α rs 1800629 and TGF -β rs1982073 polymorphisms and risk of chronic myeloid leukemia in a Romanian population from Transylvania*. Romanian Journal of Rare Diseases. 2016 (Sup.1):30. ISSN 2068–5882 **(0,33 pct)**
13. Moldovan GV, Duicu C, **Bogliș A,** Bănescu C. *Genetic aspects in familial hypercholesterolemia.* Romanian Journal of Rare Diseases 2015; Supplement 1:47. ISSN 2068-5882 **(0,5 pct)**
14. Tripon F, Crauciuc G, **Boglis A**, Radu C, Stefan A, Duicu C, Banescu C. *The role of reactive oxygen stress and proinflammatory cytokine genes in Hodgkin lymphoma*. Romanian Journal of Rare Diseases 2015; Supplement 1:29. ISSN 2068-5882 **(0,28 pct)**

15. Bănescu C, **Boglis A,** Moldovan V, Mărginean O, Grama A, Duicu C. *Genotype-phenotype correlations in patients with Beckwith-Wideman syndrome*. Jurnalul Pediatrului 2015; Supplement 18(2): 67. ISSN 2360-4557 **(0,33 pct)**
16. Jankus HS, Ilie AB, **Bogliș A,** Mirică MN. *Importanța remediilor homeopate Cantharis CH7 și Staphysagria CH7 în tratamentul nocicepției produsă experimental.* Revista de Medicină și Farmacie Târgu Mureș 2010, Supliment 56(2):30. ISSN 1221-2229. **(0,5 pct)**

Data **Semnatura**

04.08.2020