EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Curriculum Vitae

Personal information

Ingeborg Barisic

Work experience

1. Employer: Polyclinic Breyer
   - Start date: Dec 2022
   - End date: present
   - Position: Genetic consultant
   - Activities:
   - Country: Croatia

1. Employer: Medical School University of Zagreb
   - Start date: 2010
   - End date: present
   - Position: Teaching Professor
   - Activities: present
   - Country: Croatia
2. Employer: Faculty of Pharmacy and Biochemistry University of Zagreb
   - Start date: 2007
   - End date: present
   - Position: Teaching professor
   - Activities:
   - Country: Croatia
3. Employer: Faculty for Educational and Rehabilitation Sciences
   - Start date: 1996
   - End date: present
   - Position: Teaching professor
   - Activities:
   - Country: Croatia
4. Employer: Croatian Paediatric Society, Croatian Medical Association
   - Start date: 1997
   - End date: present
   - Position: Editor_in_Chief of the journal Paediatria Croatica
   - Activities:
   - Country: Croatia
5. Employer: Children's University Hospital Zagreb, Medical School University of Zagreb
   - Start date: 2003
   - End date: 2019
   - Position: Head of the Referral Centre of the Croatian Ministry of Health for the Surveillance of Congenital Anomalies
   - Activities:
   - Country: Croatia
6. Employer: Childrens University Hospital Zagreb, Medical School University of Zagreb
   - Start date: 1999
   - End date: 2003
   - Position: Head of the Department of Paediatrics
   - Activities:
   - Country: Croatia
7. Employer: Childrens University Hospital Zagreb, Medical School University of Zagreb
   - Start date: 1996
   - End date: 122019
   - Position: Head of the Department of Medical Genetics and Reproductive Health
   - Activities:
   - Country: Croatia
8. Employer: Public Health Centre Trnje, Clinical Hospital Centre Zagreb, Children's University Hospital Zagreb
   - Start date: 091979
   - End date: 1995
   - Position: Medical doctor, resident in pediatrics, pediatrician
   - Activities:
   - Country: Croatia

Education and training

1. Subject: Medical School University of Zagreb
   - Start date: 2004
   - End date:
   - Qualification: 2010 Full Professor of Paediatrics
   - Organisation:
   - Country: Croatia
2. Subject: Ministry of Health of the Repubic of Croatia
   - Start date:
   - End date: 1999
   - Qualification: Sub_specialisation in Medical Genetics
   - Organisation:
   - Country: Croatia

3. Subject: Medical School University of Zagreb
    - Start date: 011987
    - End date: 1989
    - Qualification: Ph.D.
    - Organisation:
    - Country: Croatia
4. Subject: Clinical Hospital Centre Zagreb, Medical School University of Zagreb
    - Start date: 1980
    - End date: 1986
    - Qualification: Specialization in Paediatrics
    - Organisation:
    - Country: Croatia
5. Subject: Medical School University of Zagreb
    - Start date: 1985
    - End date: 1986
    - Qualification: Postgraduate studies in Medical Genetics
    - Organisation:
    - Country:
6. Subject: Medical School University of Zagreb
    - Start date: 1981
    - End date: 1982
    - Qualification: Postgraduate studies in Paediatrics
    - Organisation:
    - Country: Croatia
7. Subject: Medical School University of Zagreb
    - Start date: 1977
    - End date: 1979
    - Qualification: Postgraduate studies in Biomedicine
    - Organisation:
    - Country: Croatia
8. Subject: Medical School University of Zagreb
    - Start date: 1972
    - End date: 1977
    - Qualification: M.D.
    - Organisation:
    - Country: Croatia

Additional information

Publications

Google Scholar H index 37 i10 index 58 Citations 5526 Publications in the last 5 years 1. Mamasoula C, Bigirumurame T, Chadwick T, Addor MC, Cavero_Carbonell C, Dias CM, Echevarría_González_de_Garibay LJ, Gatt M, Khoshnood B, Klungsoyr K, Randall K, Stoianova S, Haeusler M, Nelen V, Neville AJ, Perthus I, Pierini A, Bertaut_Nativel B, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Zymak_Zakutnia N, Barisic I, de Walle HEK, Lanzoni M, Sayers G, Mullaney C, Pennington L, Rankin J. Maternal age and the prevalence of congenital heart defects in Europe, 1995_2015: A register_based study. Birth Defects Res. 2023 Feb 3. doi: 10.1002/bdr2.2152. Epub ahead of print. PMID: 36734416. 2. Bergman, J. E. H., Barišić, I., Addor, M._C., Braz, P., Cavero_Carbonell, C., Draper, E. S., Echevarría_González_de_Garibay, L. J., Gatt, M., Haeusler, M., Khoshnood, B., Klungsøyr, K., Kurinczuk, J. J., Latos_Bielenska, A., Luyt, K., Martin, D., Mullaney, C., Nelen, V., Neville, A. J., O'Mahony, M. T., … de Walle, H. E. K. (2022). Amniotic band syndrome and limb body wall complex in Europe 1980–2019. American Journal of Medical Genetics Part A, 1– 12. https://doi.org/10.1002/ajmg.a. 3. Morris JK, Wellesley D, Limb E, Bergman JEH, Kinsner_Ovaskainen A, Addor MC, Broughan JM, Cavero_Carbonell C, Dias CM, Echevarría_González_de_Garibay LJ, Gatt M, Haeusler M, Barisic I, Klungsoyr K, Lelong N, Materna_Kiryluk A, Neville A, Nelen V, O'Mahony MT, Perthus I, Pierini A, Rankin J, Rissmann A, Rouget F, Sayers G, Stevens S, Tucker D, Garne E. Prevalence of vascular disruption anomalies and association with young maternal age: A EUROCAT study to compare the United Kingdom with other European countries. Birth Defects Res. 2022 Dec 1;114(20):1417_1426. doi: 10.1002/bdr2.2122. Epub 2022 Nov 11. PMID: 36369770. 4. Marcus E, Latos_Bielenska A, Jamry_Dziurla A, Barišić I, Cavero_Carbonell C, Den Hond E, Garne E, Genard L, Santos AJ, Lutke L, Matias Dias C, Neergaard Pedersen C, Neville AJ, Niemann A, Odak L, Pierini A, Rico J, Rissmann A, Rankin J, Morris JK. Information needs of parents of children with congenital anomalies across Europe: a EUROlinkCAT survey. BMC Pediatr. 2022 Nov 12;22(1):657. doi: 10.1186/s12887_022_03734_z. 5. Mamasoula C, Addor MC, Carbonell CC, Dias CM, Echevarría_González_de_Garibay LJ, Gatt M, Khoshnood B, Klungsoyr K, Randall K, Stoianova S, Haeusler M, Nelen V, Neville AJ, Perthus I, Pierini A, Bertaut_Nativel B, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Zymak_Zakutnia N, Barisic I, de Walle HEK, Lanzoni M, Mullaney C, Pennington L, Rankin J. Prevalence of congenital heart defects in Europe, 2008_2015: A registry_based study. Birth Defects Res. 2022 Dec 1;114(20):1404_1416. doi: 10.1002/bdr2.2117. Epub 2022 Nov 8. 6. Coi A, Barisic I, Garne E, Pierini A, Addor MC, Aizpurua Atxega A, Ballardini E, Braz P, Broughan JM, Cavero_Carbonell C, de Walle HEK, Draper ES, Gatt M, Häusler M, Kinsner_Ovaskainen A, Kurinczuk JJ, Lelong N, Luyt K, Mezzasalma L, Mullaney C, Nelen V, Odak L, O'Mahony MT, Perthus I, Randrianaivo H, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Wiśniewska K, Yevtushok L, Santoro M. Epidemiology of aplasia cutis congenita: A population_based study in Europe. J Eur Acad Dermatol Venereol. 2022 Oct 27. doi: 10.1111/jdv.18690. 7. Morris JK, Wellesley D, Limb E, Bergman JEH, Kinsner_Ovaskainen A, Addor MC, Broughan JM, Cavero_Carbonell C, Dias CM, Echevarría_González_de_Garibay LJ, Gatt M, Haeusler M, Barisic I, Klungsoyr K, Lelong N, Materna_Kiryluk A, Neville A, Nelen V, O'Mahony MT, Perthus I, Pierini A, Rankin J, Rissmann A, Rouget F, Sayers G, Stevens S, Tucker D, Garne E. Prevalence of vascular disruption anomalies and association with young maternal age: A EUROCAT study to compare the United Kingdom with other European countries Birth Defects Res. 2022 Dec 1;114(20):1404_1416. doi: 10.1002/bdr2.2117. Epub 2022 Nov 8. 8. Latos_Bielenśka A, Marcus E, Jamry_Dziurla A et al. COVID_19 and children with congenital anomalies: a European survey of parents' experiences of healthcare services. BMJ Open 2022;12:e061428. doi:10.1136/ bmjopen_2022_061428 9. Urhoj SK, Tan J, Morris JK, Given J, Astolfi G, Baldacci S, Barisic I, Brigden J, Cavero_Carbonell C, Evans H, Gissler M, Heino A, Jordan S, Lutke R, Odak L, Puccini A, Santoro M, Scanlon I, de Walle HEK, Wellesley D, Zurriaga Ó, Loane M, Garne E. Hospital length of stay among children with and without congenital anomalies across 11 European regions_A population_based data linkage study. PLoS One. 2022 Jul 22;17(7):e0269874. doi: 10.1371/journal.pone.0269874 10. Santoro M, Coi A, Pierini A, Rankin J, Glinianaia SV, Tan J, Reid A, Garne E, Loane M, Given J, Aizpurua A, Astolfi G, Barisic I, Cavero_Carbonell C, de Walle HEK, Den Hond E, García_Villodre L, Gatt M, Gissler M, Jordan S, Khoshnood B, Kiuru_Kuhlefelt S, Klungsøyr K, Lelong N, Lutke R, Mokoroa O, Nelen V, Neville AJ, Odak L, Rissmann A, Scanlon I, Urhoj SK, Wellesley D, Wertelecki W, Yevtushok L, Morris JK. Temporal and geographical variations in survival of children born with congenital anomalies in Europe: A multi_registry cohort study. Paediatr Perinat Epidemiol. 2022 Jun 8. doi: 10.1111/ppe.12884. Epub ahead of print. PMID: 35675091. 11. Tylki_Szymańska A, Almássy Z, Christophidou_Anastasiadou V, Avdjieva_Tzavella D, Barisic I, Cerkauskiene R, Cuturilo G, Djiordjevic M, Gucev Z, Hlavata A, Kieć_Wilk B, Magner M, Pecin I, Plaiasu V, Samardzic M, Zafeiriou D, Zaganas I, Lampe C. The landscape of Mucopolysaccharidosis in Southern and Eastern European countries: a survey from 19 specialistic centers. Orphanet J Rare Dis. 2022 Mar 24;17(1):136. doi: 10.1186/s13023_022_02285_x. PMID: 35331284; PMCID: PMC8943501. 12. Krnjak G, Vulin K, Pazanin L, Barisic I, Duranovic V. A case of macrophagic myofasciitis in a girl with developmental delay. Pediatr Int. 2022 Jan;64(1):e14930. doi: 10.1111/ped.14930. PMID: 35139249. 13. Loane M, Given JE, Tan J, Reid A, Akhmedzhanova D, Astolfi G, Barišić I, Bertille N, Bonet LB, Carbonell CC, Carollo OM, Coi A, Densem J, Draper E, Garne E, Gatt M, Glinianaia SV, Heino A, Hond ED, Jordan S, Khoshnood B, Kiuru_Kuhlefelt S, Klungsøyr K, Lelong N, Lutke LR, Neville AJ, Ostapchuk L, Puccini A, Rissmann A, Santoro M, Scanlon I, Thys G, Tucker D, Urhoj SK, de Walle HEK, Wellesley D, Zurriaga O, Morris JK. Linking a European cohort of children born with congenital anomalies

to vital statistics and mortality records: A EUROlinkCAT study. PLoS One. 2021 Aug 27;16(8):e0256535. doi: 10.1371/journal.pone.0256535. PMID: 34449798; PMCID: PMC8396745. 14. Morris JK, Addor MC, Ballardini E, Barisic I, Barrachina_Bonet L, Braz P, Cavero_Carbonell C, Den Hond E, Garne E, Gatt M, Haeusler M, Khoshnood B, Lelong N, Kinsner_Ovaskainen A, Kiuru_Kuhlefelt S, Klungsoyr K, Latos_Bielenska A, Limb E, O'Mahony MT, Perthus I, Pierini A, Rankin J, Rissmann A, Rouget F, Sayers G, Sipek A Jr, Stevens S, Tucker D, Verellen_Dumoulin C, de Walle HEK, Wellesley D, Wertelecki W, Bermejo_Sanchez E. Prevention of Neural Tube Defects in Europe: A Public Health Failure. Front Pediatr. 2021;9:647038. doi: 10.3389/fped.2021.647038. 15. Morris JK, Garne E, Loane M, Barisic I, Densem J, Latos_Bieleńska A, Neville A, Pierini A, Rankin J, Rissmann A, de Walle H, Tan J, Given JE, Claridge H; EUROlinkCAT Consortium. EUROlinkCAT protocol for a European population_based data linkage study investigating the survival, morbidity and education of children with congenital anomalies. BMJ Open. 2021 Jun 28;11(6):e047859. 16. Santoro M, Coi A, Barišić I, Pierini A, Addor MC, Baldacci S, Ballardini E, Boban L, Braz P, Cavero_Carbonell C, de Walle HEK, Draper ES, Gatt M, Haeusler M, Klungsøyr K, Kurinczuk JJ, Materna_Kiryluk A, Lanzoni M, Lelong N, Luyt K, Mokoroa O, Mullaney C, Nelen V, O'Mahony MT, Perthus I, Randrianaivo H, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Zymak_Zakutnia N, Garne E. Epidemiology of Pierre_Robin sequence in Europe: A population_based EUROCAT study. Paediatr Perinat Epidemiol. 2021 Jun 16. doi: 10.1111/ppe.12776. Epub ahead of print. PMID: 34132407. 17. Cavadino A, Sandberg L, Öhman I, Bergvall T, Star K, Dolk H, Loane M, Addor MC, Barisic I, Cavero_Carbonell C, Garne E, Gatt M, Khoshnood B, Klungsøyr K, Latos_Bielenska A, Lelong N, Lutke R, Materna_Kiryluk A, Nelen V, Nevill A, O'Mahony M, Mokoroa O, Pierini A, Randrianaivo H, Rissmann A, Tucker D, Wiesel A, Yevtushok L, Morris JK. Signal Detection in EUROmediCAT: Identification and Evaluation of Medication_Congenital Anomaly Associations and Use of VigiBase as a Complementary Source of Reference. Drug Saf. 2021 May 9. doi: 10.1007/s40264_021_01073_z. Epub ahead of print. PMID: 33966183. 18. Dolk H, Leke AZ, Whitfield P, Moore R, Karnell K, Barišić I, Barlow_Mosha L, Botto LD, Garne E, Guatibonza P, Godfred_Cato S, Halleux CM, Holmes LB, Moore CA, Orioli I, Raina N, Valencia D. Global birth defects app: An innovative tool for describing and coding congenital anomalies at birth in low resource settings. Birth Defects Res. 2021 May 5. doi: 10.1002/bdr2.1898. Epub ahead of print. PMID: 33949803. 19. Leke AZ, Dolk H, Loane M, Casson K, Nelen V, Barišić I, Garne E, Rissman A, O'Mahony M, Neville AJ, Pierini A, Bergman JEH, Klungsøyr K, Materna_Kiryluk A, Bielenska AL, Carbonell CC, Addor MC, Tucker D. Macrolide and lincosamide antibiotic exposure in the first trimester of pregnancy and risk of congenital anomaly: A European case_control study. Reprod Toxicol. 2021 Mar;100:101_108. doi: 10.1016/j.reprotox.2021.01.006. Epub 2021 Jan 14.PMID: 33454317 20. Ana_Maria Meašić, Adriana Bobinec, Ivona Sansović Ana Močić Pavić, Silvija Pušeljić, Ingeborg Barisic. Homozygous ABCG8 mutation in a 14_year_old boy with sitosterolemia. October 2021Archives of Disease in Childhood 106(Suppl 2):A40.2_A40 DOI: 10.1136/archdischild_2021_europaediatrics.96 Conference: 10th Europaediatrics Congress, Zagreb, Croatia, 7–9 October 2021 21. Alenka Gagro, Nives Pustisek, Suzana Ozanic Bulic, Tena Nisseto, Sara Sila, Sanja Kolaček, Ingeborg Barisic. Severe presentation of Netherton syndrome: a case report. October 2021Archives of Disease in Childhood 106(Suppl 2):A67.3_A68 DOI: 10.1136/archdischild_2021_europaediatrics.159 Conference: 10th Europaediatrics Congress, Zagreb, Croatia, 7–9 October 2021 22. Ljubica Odak, Ana_Maria Meašić, Adriana Bobinec, Mijana Kero, Ivona Sansović, Katrina Vulin, Mirko Tomić, Ingeborg Barisic. Clinical exome sequencing in the diagnosis of autism spectrum disorder. October 2021Archives of Disease in Childhood 106(Suppl 2):A41.2_A41 DOI: 10.1136/archdischild_2021_europaediatrics.98 Conference: 10th Europaediatrics Congress, Zagreb, Croatia, 7–9 October 2021 23. Maša Davidović, Leona Morozin Pohovski, Nikolina Vidan Rogulj, Ivona Sansović, Adriana Bobinec, Ana_Maria Meašić, Mijana Kero, Ljubca Odak, Ivan Malčić, Ingeborg Barisic. Rare copy number variants in congenital heart defects. October 2021Archives of Disease in Childhood 106(Suppl 2):A36.1_A36 DOI: 10.1136/archdischild_2021_europaediatrics.84 Conference: 10th Europaediatrics Congress, Zagreb, Croatia, 7–9 October 2021 24. Ivona Sansović, Ljubica Odak, Adriana Bobinec, Ingeborg Barisic. A novel missense mutation in SGSH gene causing Sanfillipo type 3A mucopolysaccharidosis October 2021Archives of Disease in Childhood 106(Suppl 2):A41.1_A41 Follow journal DOI: 10.1136/archdischild_2021_europaediatrics. 97 Conference: 10th Europaediatrics Congress, Zagreb, Croatia, 7–9 October 2021 25. Adriana Bobinec, Ana_Maria Meašić Ivona Sansović, Mijana Kero, Linimol Boban, Ingeborg Barisic. CDHR1 and RGR mutations in two patients with retinitis pigmentosa. October 2021Archives of Disease in Childhood 106(Suppl 2):A40.1_A40 DOI: 10.1136/archdischild_2021_europaediatrics. 95Conference: 10th Europaediatrics Congress, Zagreb, Croatia, 7–9 October 2021 26. Morozin Pohovski L, Bobinec A, Measic AM , Sansovic I, Barisic I. A new case of intragenic deletion of the TCF4 gene without features of Pitt_Hopkins syndrome. MEBM. 2020;3:56_8. https://doi.org/10.33602/mebm.3.2.8 27. van Schie JJM, Faramarz A, Balk JA, Stewart GS, Cantelli E, Oostra AB, Rooimans MA, Parish JL, de Almeida Estéves C, Dumic K, Barisic I, Diderich KEM, van Slegtenhorst MA, Mahtab M, Pisani FM, Te Riele H, Ameziane N, Wolthuis RMF, de Lange J.Warsaw Breakage Syndrome associated DDX11 helicase resolves G_quadruplex structures to support sister chromatid cohesion. Nat Commun. 2020;11(1):4287. Published 2020 Aug 27. doi:10.1038/s41467_020_18066_8 28. van de Putte R, van Rooij IALM, Haanappel CP, Marcelis CLM, Brunner HG, Addor MC, Cavero_Carbonell C, Dias CM, Draper ES, Etxebarriarteun L, Gatt M, Khoshnood B, Kinsner_Ovaskainen A, Klungsoyr K, Kurinczuk JJ, Latos_Bielenska A, Luyt K, O'Mahony MT, Miller N, Mullaney C, Nelen V, Neville AJ, Perthus I, Pierini A, Randrianaivo H, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Wiesel A, Zymak_Zakutnia N, Loane M, Barisic I, de Walle HEK, Bergman JEH, Roeleveld N. Maternal risk factors for the VACTERL association: A EUROCAT case_control study. Birth Defects Res. 2020 May 15;112(9):688_698. doi: 10.1002/bdr2.1686. Epub 2020 Apr 22. 29. van de Putte R, van Rooij IALM, Marcelis CLM, Guo M, Brunner HG, Addor MC, Cavero_Carbonell C, Dias CM, Draper ES, Etxebarriarteun L, Gatt M, Haeusler M, Khoshnood B, Klungsoyr K, Kurinczuk JJ, Lanzoni M, Latos_Bielenska A, Luyt K, O'Mahony MT, Miller N, Mullaney C, Nelen V, Neville AJ, Perthus I, Pierini A, Randrianaivo H, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Wiesel A, Zymak_Zakutnia N, Loane M, Barisic I, de Walle HEK, Roeleveld N, Bergman JEH. Spectrum of congenital anomalies among VACTERL cases: a EUROCAT population_based study. Pediatr Res. 2020 Feb;87(3):541_549. doi: 10.1038/s41390_019_0561_y. [Epub ahead of print] 30. Santoro M, Coi A, Barišić I, Garne E, Addor MC, Bergman JEH, Bianchi F, Boban L, Braz P, Cavero_Carbonell C, Gatt M, Haeusler M, Kinsner_Ovaskainen A, Klungsøyr K, Kurinczuk JJ, Lelong N, Luyt K, Materna_Kiryluk A, Mokoroa O, Mullaney C, Nelen V, Neville AJ, O'Mahony MT, Perthus I, Randrianaivo H, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Yevtushok L, Pierini A. Epidemiology of Dandy_Walker Malformation in Europe: A EUROCAT Population_Based Registry Study. Neuroepidemiology. 2019;53(3_4):169_179. doi: 10.1159/000501238. 31. Coi A, Santoro M, Garne E, Pierini A, Addor MC, Alessandri JL, Bergman JEH, Bianchi F, Boban L, Braz P, Cavero_Carbonell C, Gatt M, Haeusler M, Klungsøyr K, Kurinczuk JJ, Lanzoni M, Lelong N, Luyt K, Mokoroa O, Mullaney C, Nelen V, Neville AJ, O'Mahony MT, Perthus I, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker D, Wellesley D, Wisniewska K, Zymak_Zakutnia N, Barišić I. Epidemiology of achondroplasia: A population_based study in Europe. Am J Med Genet. 2019;179A:1791–1798. DOI: 10.1002/ajmg.a.61289. 32. Morris JK, Wellesley DG, Barisic I, Addor MC, Bergman JEH, Braz P, Cavero_Carbonell C, Draper ES, Gatt M, Haeusler M, Klungsoyr K, Kurinczuk JJ, Lelong N, Luyt K, Lynch C, O'Mahony MT, Mokoroa O, Nelen V, Neville AJ, Pierini A, Randrianaivo H, Rankin J, Rissmann A, Rouget F, Schaub B, Tucker DF, Verellen_Dumoulin C, Wiesel A, Zymak_Zakutnia N, Lanzoni M, Garne E. Epidemiology of congenital cerebral anomalies in Europe: a multicentre, population_based EUROCAT study. Arch Dis Child. 2019;104(12):1181_1187. doi: 10.1136/archdischild_2018_316733. 33. Cleary B, Loane M, Addor MC, Barisic I, de Walle HEK, Matias Dias C, Gatt M, Klungsoyr K, McDonnell B, Neville A, Pierini A, Rissmann A, Tucker DF, Zurriaga O, Dolk H. Methadone, Pierre Robin sequence and other congenital anomalies: case_control study. Arch Dis Child Fetal Neonatal Ed. 2020 Mar;105(2):151_157. doi: 10.1136/archdischild_2019_316804. Epub 2019 Jun 22.. 4 34. 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Projects

Horizon 2020 Grant Agreement_821520_ConcePTION (2019_) Horizon 2020 project Grant Agreement No: 831390 — ONW — HP_PJ_03_2018, and Grant Agreement (GA) No: 825575 Acronym: EJP RD, leading partner INSERM Orphanet (2018_2022) Evaluation of the computer algorithm for the classification of congenital anomalies and coding of isolated anomalies subgroups in the JRC_EUROCAT Central Database" European Commission Joint Research Centre, Contract No. CCR.F.C793427.X0 (2018) Horizon 2020 project EUROlinkCAT: Establishing a linked European Cohort of Children with Congenital Anomalies (733001), leading partner: St George University of London (2017_2022) COST Action OC_2016_1_208_62 _ Neuro_MIG (European Network on Brain Malformations) (CA16118), leading partner Erasmus University Medical Centre (2017_2022) Reproductive and regenerative medicine – research into new platforms and potentials, KK..01.1.1.01.0008, Operational Program Competitiveness and Cohesion, European Regional Development Fund (2017_) Horizon 2020 ZIKAPLAN Project. International Committee for Birth Defect Surveillance Tools for Infectious Disease Preparedness (734584) (2017_2022) Evaluation of EUROCAT coding of minor anomalies based on a retrospective analysis of selected multiple anomaly cases, European Commission Joint Research Centre, Contract CCR.F.C791623.X0 (2016) Hunter Outcome Survey _ A Global, Multi_Centre, Long_Term, Observational Registry of Patients with Hunter syndrome, Shire Project (2016_2021) Improving EUROCAT database capacity to serve as a source for studies of rare genetic diseases, EU Commission Joint Research Centre, Contract No CCR.IHCPC443727.X0 (2015) JRC_EUROCAT_ European network of population_based registries for the epidemiological surveillance of congenital anomalies, Joint Research Centre, European Platform on Rare Disease Registration (EU RD Platform) (2015_) FP7 project _EUROmediCAT _ Safety of Medication Use in Pregnancy in Relation to Risk of Congenital Malformations, Grant Agreement no 2605982013_ 2015 JA _ EUROCAT (European Registration of Congenital Anomalies) – contract with EC DGF4 od 1.01.2011: Grant Agreement no 20102204 (2011_2013) EUROPLAN _ European Project for Rare Diseases National Plans Development, funded by the European Commission within the EU Program of Community Action in the field of Public Health (2008_2011) EPIRARE (European Platform for Rare Disease Registries) co_funded by the European Commission within the EU Program of Community Action in the field of Public Health. (2011_2013) "Epidemiology and Genetic Basis of Congenital Anomalies", Croatian Ministry of Science and Technology, project. No. 072_183107_0365 (2007_2014) „Genetic basis of Diseases in Children"_ Croatian Ministry of Science Program (2007_2014) "Epidemiological, Clinical and Biological Aspects of Birth Defects", Croatian Ministry of Science and Technology (project No. 077_1605) (2002_2007) "Molecular Cytogenetics in Diagnostic of Mental Retardation", Technological developmental_research project (program TEST) Croatian Ministry of Science and Technology (TP_01/072_01) (2001_2003) " Epidemiological, Clinical and Biological Aspects of Birth Defects", Croatian Ministry of Science and Technology (project No. 072_777) (1999_2002) „Genetic, Morphological, Immunological and Clinical Research in Children with Solid Tumors", Croatian Ministry of Science and Technology (project No. 072_333) (1997_2000) BIOMED2 "Evaluation of Prenatal Diagnosis of Congenital Anomalies by Fetal Ultrasonographic Examination" (V Contract No BMH4 CT960539) (1996_2002) „Medical and Social Evaluation of Prenatal Diagnosis ", Croatian Ministry of Science and Technology (Project No. 301_324) (1992_1997)

Memberships

2019 _ Representative of Croatia to the Board of Member States on European Reference Networks (ERNs) 2018 _ Member of the Policy Board of the European Joint Program on Rare Diseases (EJP), 2018 _ 2021 Representative of Croatia to the group of countries signatories to the Declaration "Towards access to at least 1 million sequenced genomes in the European Union by 2022", 2013 _ 2022 Member of the Orphanet Board and National representative of Orphanet Croatia 2013 _ Member of COMP 2014 _ Member of Croatian Academy of Medical Sciences 2013 _ 2017 Representative of the Republic of Croatia at Commission's Expert Group on Rare Diseases 2011 _ 2016 President of EUROCAT (European Network of Population_Based Registries for the Surveillance of Congenital Anomalies) Association 2010 _ 2019 Chair of the Commission for the Adoption and Implementation of the National Plan for Rare Diseases of the Republic of Croatia. 2009 _ Representative of the National Society of Human Genetics at ESHG 2009 _ 2018 President of Croatian Society of Human Genetics of the Croatian Medical Association 2008 _ 2018 President of Croatian Society of Rare Diseases of the Croatian Medical Association 2009 _ 2011 Member of the American Academy of Medical Genetics 2008 _ 2017 Member of European Association of Science Editors 1999 _ 2019 Member of the Board of Croatian Pediatric Society of the Croatian Medical Association 1999 _ 2011 Member of European Cytogeneticists Association 1998 _ 2020 Member of European Society of Human Genetics

Other Relevant Information

Awards: 2017_ Honorary award of the Croatian Medical Association 2017_ Best medical doctor award, Croatian Alliance of Patients with Rare Diseases 2014 and 2016 _Ernst Mayerhofer Award of the Croatian Pediatric Society for the best work of Croatian pediatricians 2009 _ Main Board Diploma of Croatian Medical Association 2005 _ "Ladislav Rakovac" Award _ Croatian Medical Association