

# Adrianna Mostowska

## List of Publications by Year in Descending Order

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**Version:** 2024-04-26

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

95  
papers

1,455  
citations

21  
h-index

34  
g-index

111  
ext. papers

1,651  
ext. citations

3  
avg, IF

4.4  
L-index

#	Paper	IF	Citations
95	Paraoxonase 1 concerning dyslipidaemia, cardiovascular diseases, and mortality in haemodialysis patients. <i>Scientific Reports</i> , <b>2021</b> , 11, 6773	4.9	5
94	Polymorphism rs368234815 of interferon lambda 4 gene and spontaneous clearance of hepatitis C virus in haemodialysis patients: a case-control study. <i>BMC Infectious Diseases</i> , <b>2021</b> , 21, 102	4	1
93	The Prevalence and Morphology of Supernumerary Teeth in Children With Nonsyndromic Cleft Lip and Palate. <i>Cleft Palate-Craniofacial Journal</i> , <b>2021</b> , 10556656211027750	1.9	1
92	Paraoxonase 1 gene (PON1) variants concerning hepatitis C virus (HCV) spontaneous clearance in hemodialysis individuals: a case-control study. <i>BMC Infectious Diseases</i> , <b>2021</b> , 21, 875	4	
91	Paraoxonase 1 gene variants concerning cardiovascular mortality in conventional cigarette smokers and non-smokers treated with hemodialysis. <i>Scientific Reports</i> , <b>2021</b> , 11, 19467	4.9	0
90	Alpha-1 Antitrypsin Z Variant (AAT PI*Z) as a Risk Factor for Intrahepatic Cholestasis of Pregnancy. <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 720465	4.5	1
89	MiRNA-149 as a Candidate for Facial Clefting and Neural Crest Cell Migration. <i>Journal of Dental Research</i> , <b>2021</b> , 220345211038203	8.1	2
88	Association of rs699947 (-2578 C/A) and rs2010963 (-634 G/C) Single Nucleotide Polymorphisms of the Gene, VEGF-A and Leptin Serum Level, and Cardiovascular Risk in Patients with Excess Body Mass: A Case-Control Study. <i>Journal of Clinical Medicine</i> , <b>2020</b> , 9,	5.1	10
87	Paraoxonase 1 gene polymorphisms concerning non-insulin-dependent diabetes mellitus nephropathy in hemodialysis patients. <i>Journal of Diabetes and Its Complications</i> , <b>2020</b> , 34, 107687	3.2	4
86	Polymorphism rs368234815 of interferon- $\lambda$ gene and generation of antibodies to hepatitis B virus surface antigen in extracorporeal dialysis patients. <i>Expert Review of Vaccines</i> , <b>2020</b> , 19, 293-303	5.2	4
85	PAX7 nucleotide variants and the risk of non-syndromic orofacial clefts in the Polish population. <i>Oral Diseases</i> , <b>2019</b> , 25, 1608-1618	3.5	7
84	FP679IFNL4 rs368234815 POLYMORPHISM AND SPONTANEOUS CLEARANCE OF HEPATITIS C VIRUS IN HEMODIALYSIS PATIENTS. <i>Nephrology Dialysis Transplantation</i> , <b>2019</b> , 34,	4.3	1
83	Calcium-sensing receptor gene (CASR) polymorphisms and CASR transcript level concerning dyslipidemia in hemodialysis patients: a cross-sectional study. <i>BMC Nephrology</i> , <b>2019</b> , 20, 436	2.7	2
82	Association of CDKAL1 nucleotide variants with the risk of non-syndromic cleft lip with or without cleft palate. <i>Journal of Human Genetics</i> , <b>2018</b> , 63, 397-406	4.3	4
81	GREM2 nucleotide variants and the risk of tooth agenesis. <i>Oral Diseases</i> , <b>2018</b> , 24, 591-599	3.5	6
80	Common variants in DLG1 locus are associated with non-syndromic cleft lip with or without cleft palate. <i>Clinical Genetics</i> , <b>2018</b> , 93, 784-793	4	20
79	Correlations of indoleamine 2,3-dioxygenase, interferon- $\beta$ , and anti-HBs antibodies in hemodialysis patients. <i>Vaccine</i> , <b>2018</b> , 36, 4454-4461	4.1	2

78	The assessment of GWAS - identified polymorphisms associated with infertility risk in Polish women with endometriosis. <i>Ginekologia Polska</i> , <b>2018</b> , 89, 304-310	1	8
77	ENHO, RXRA, and LXRA polymorphisms and dyslipidaemia, related comorbidities and survival in haemodialysis patients. <i>BMC Medical Genetics</i> , <b>2018</b> , 19, 194	2.1	5
76	The Calcium-Sensing Receptor Gene Polymorphism rs1801725 and Calcium-Related Phenotypes in Hemodialysis Patients. <i>Kidney and Blood Pressure Research</i> , <b>2018</b> , 43, 719-734	3.1	3
75	Further Evidence of the Association of the Diacylglycerol Kinase Kappa (DGKK) Gene With Hypospadias. <i>Urology Journal</i> , <b>2018</b> , 15, 272-276	0.9	5
74	Involvement of 17 $\beta$ hydroxysteroid dehydrogenase type gene 1 937 A>G polymorphism in infertility in Polish Caucasian women with endometriosis. <i>Journal of Assisted Reproduction and Genetics</i> , <b>2017</b> , 34, 789-794	3.4	7
73	Polymorphisms of T helper cell cytokine-associated genes and survival of hemodialysis patients - a prospective study. <i>BMC Nephrology</i> , <b>2017</b> , 18, 165	2.7	2
72	Circulating Interferon-3, Responsiveness to HBV Vaccination, and HBV/HCV Infections in Haemodialysis Patients. <i>BioMed Research International</i> , <b>2017</b> , 2017, 3713025	3	4
71	Monocyte Chemotactic Protein-1 (Cytokine, Receptors, and Gene Polymorphisms) in Hepatitis. <i>Biomarkers in Disease</i> , <b>2017</b> , 927-955		1
70	Involvement of adropin and adropin-associated genes in metabolic abnormalities of hemodialysis patients. <i>Life Sciences</i> , <b>2016</b> , 160, 41-46	6.8	5
69	Polymorphic variants in the vitamin D pathway genes and the risk of ovarian cancer among non-carriers of mutations. <i>Oncology Letters</i> , <b>2016</b> , 11, 1181-1188	2.6	15
68	Antibodies to HBV surface antigen in relation to interferon- $\beta$ in hemodialysis patients. <i>Vaccine</i> , <b>2016</b> , 34, 4866-4874	4.1	5
67	Associations of the calcium-sensing receptor gene CASR rs7652589 SNP with nephrolithiasis and secondary hyperparathyroidism in haemodialysis patients. <i>Scientific Reports</i> , <b>2016</b> , 6, 35188	4.9	9
66	Nucleotide Variants of the BH4 Biosynthesis Pathway Gene GCH1 and the Risk of Orofacial Clefts. <i>Molecular Neurobiology</i> , <b>2016</b> , 53, 769-776	6.2	1
65	Polymorphisms of Vitamin D Signaling Pathway Genes and Calcium-Sensing Receptor Gene in respect to Survival of Hemodialysis Patients: A Prospective Observational Study. <i>International Journal of Endocrinology</i> , <b>2016</b> , 2016, 2383216	2.7	5
64	Association between DNMT3L polymorphic variants and the risk of endometriosis-associated infertility. <i>Molecular Medicine Reports</i> , <b>2016</b> , 13, 1040-6	2.9	
63	De novo EDA mutations: Variable expression in two Egyptian families. <i>Archives of Oral Biology</i> , <b>2016</b> , 68, 21-8	2.8	6
62	Replication study for the association of seven genome- GWAS-identified Loci with susceptibility to ovarian cancer in the Polish population. <i>Pathology and Oncology Research</i> , <b>2015</b> , 21, 307-13	2.6	4
61	Involvement of vascular endothelial growth factor -460 C/T, +405 G/C and +936 C/T polymorphisms in the development of endometriosis. <i>Biomedical Reports</i> , <b>2015</b> , 3, 220-224	1.8	9

60	Association investigation of rs3757247 and rs4880 polymorphisms with the type 1 diabetes and diabetes long-term complications risk in the Polish population. <i>Biomedical Reports</i> , <b>2015</b> , 3, 327-332	1.8	4
59	WNT10A coding variants and maxillary lateral incisor agenesis with associated dental anomalies. <i>European Journal of Oral Sciences</i> , <b>2015</b> , 123, 1-8	2.3	15
58	Monocyte Chemotactic Protein-1 (Cytokine, Receptors, and Gene Polymorphisms) in Hepatitis. <i>Exposure and Health</i> , <b>2015</b> , 1-29	8.8	
57	Polymorphic variants in the dopamine receptor D2 in women with endometriosis-related infertility. <i>Molecular Medicine Reports</i> , <b>2015</b> , 12, 3055-60	2.9	1
56	Polymorphic variants in vitamin D signaling pathway genes and the risk of endometriosis-associated infertility. <i>Molecular Medicine Reports</i> , <b>2015</b> , 12, 7109-15	2.9	18
55	Association between polymorphisms at the GREM1 locus and the risk of nonsyndromic cleft lip with or without cleft palate in the Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2015</b> , 103, 847-56		9
54	Association of Retinoid X Receptor Alpha Gene Polymorphism with Clinical Course of Chronic Glomerulonephritis. <i>Medical Science Monitor</i> , <b>2015</b> , 21, 3671-81	3.2	3
53	Clinical aspects of vitamin D-binding protein gene polymorphisms in hemodialysis patients. <i>Polish Archives of Internal Medicine</i> , <b>2015</b> , 125, 8-17	1.9	4
52	Effect of interferon $\beta$ gene polymorphisms, rs8099917 and rs12979860, on response to hepatitis B virus vaccination and hepatitis B or C virus infections among hemodialysis patients. <i>Polish Archives of Internal Medicine</i> , <b>2015</b> , 125, 894-902	1.9	7
51	EVC gene polymorphisms and risks of isolated hypospadias - a preliminary study. <i>Central European Journal of Urology</i> , <b>2015</b> , 68, 257-62	0.9	
50	T helper cell-related cytokine gene polymorphisms and vitamin D pathway gene polymorphisms as predictors of survival probability in patients on renal replacement therapy. <i>Polish Archives of Internal Medicine</i> , <b>2015</b> , 125, 511-20	1.9	3
49	An analysis of polymorphisms within the Wnt signaling pathway in relation to ovarian cancer risk in a Polish population. <i>Molecular Diagnosis and Therapy</i> , <b>2014</b> , 18, 85-91	4.5	33
48	Genetic variants in BRIP1 (BACH1) contribute to risk of nonsyndromic cleft lip with or without cleft palate. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2014</b> , 100, 670-8		6
47	Nucleotide variants of the cancer predisposing gene CDH1 and the risk of non-syndromic cleft lip with or without cleft palate. <i>Familial Cancer</i> , <b>2014</b> , 13, 415-21	3	18
46	A novel WNT10A mutation causes non-syndromic hypodontia in an Egyptian family. <i>Archives of Oral Biology</i> , <b>2014</b> , 59, 722-8	2.8	18
45	Single nucleotide polymorphisms of vitamin D binding protein, vitamin D receptor and retinoid X receptor alpha genes and response to hepatitis B vaccination in renal replacement therapy patients. <i>Expert Review of Vaccines</i> , <b>2014</b> , 13, 1395-403	5.2	15
44	Association of aldosterone synthase (CYP11B2) gene -344T/C polymorphism with the risk of primary chronic glomerulonephritis in the Polish population. <i>JRAAS - Journal of the Renin-Angiotensin-Aldosterone System</i> , <b>2014</b> , 15, 553-8	3	7
43	T-cell cytokine gene polymorphisms and vitamin D pathway gene polymorphisms in end-stage renal disease due to type 2 diabetes mellitus nephropathy: comparisons with health status and other main causes of end-stage renal disease. <i>Journal of Diabetes Research</i> , <b>2014</b> , 2014, 120317	3.9	9

42	Association of common variants in PAH and LAT1 with non-syndromic cleft lip with or without cleft palate (NSCL/P) in the Polish population. <i>Archives of Oral Biology</i> , <b>2014</b> , 59, 363-9	2.8	1
41	Searching for new genes and loci involved in cleft lip and palate in the Polish population □ genome-wide association study. <i>Journal of Medical Science</i> , <b>2014</b> , 83, 265-268	1.6	3
40	Vitamin D receptor gene BsmI, FokI, ApaI and TaqI polymorphisms and the risk of systemic lupus erythematosus. <i>Molecular Biology Reports</i> , <b>2013</b> , 40, 803-10	2.8	44
39	Nucleotide variants of genes encoding components of the Wnt signalling pathway and the risk of non-syndromic tooth agenesis. <i>Clinical Genetics</i> , <b>2013</b> , 84, 429-40	4	42
38	DNMT1, DNMT3A and DNMT3B gene variants in relation to ovarian cancer risk in the Polish population. <i>Molecular Biology Reports</i> , <b>2013</b> , 40, 4893-9	2.8	44
37	Vitamin D receptor gene BsmI and FokI polymorphisms in relation to ovarian cancer risk in the Polish population. <i>Genetic Testing and Molecular Biomarkers</i> , <b>2013</b> , 17, 183-7	1.6	23
36	Association of the interleukin-12 polymorphic variants with the development of antibodies to surface antigen of hepatitis B virus in hemodialysis patients in response to vaccination or infection. <i>Molecular Biology Reports</i> , <b>2013</b> , 40, 6899-911	2.8	17
35	IL4R and IL13 polymorphic variants and development of antibodies to surface antigen of hepatitis B virus in hemodialysis patients in response to HBV vaccination or infection. <i>Vaccine</i> , <b>2013</b> , 31, 1766-70	4.1	10
34	Polymorphic variants of DNMT3A and the risk of endometriosis. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , <b>2013</b> , 166, 81-5	2.4	5
33	Novel PAX9 mutation associated with syndromic tooth agenesis. <i>European Journal of Oral Sciences</i> , <b>2013</b> , 121, 403-11	2.3	14
32	A mutation in mouse Pak1ip1 causes orofacial clefting while human PAK1IP1 maps to 6p24 translocation breaking points associated with orofacial clefting. <i>PLoS ONE</i> , <b>2013</b> , 8, e69333	3.7	7
31	Genotype and haplotype analysis of WNT genes in non-syndromic cleft lip with or without cleft palate. <i>European Journal of Oral Sciences</i> , <b>2012</b> , 120, 1-8	2.3	34
30	Polymorphic variants at 10q25.3 and 17q22 loci and the risk of non-syndromic cleft lip and palate in the Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2012</b> , 94, 42-6		21
29	Polymorphic variants of genes involved in homocysteine metabolism in celiac disease. <i>Molecular Biology Reports</i> , <b>2012</b> , 39, 3123-30	2.8	19
28	Association of DVL2 and AXIN2 gene polymorphisms with cleft lip with or without cleft palate in a Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2012</b> , 94, 943-50		17
27	Antibodies to hepatitis B virus surface antigen and interleukin 12 and interleukin 18 gene polymorphisms in hemodialysis patients. <i>BMC Nephrology</i> , <b>2012</b> , 13, 75	2.7	18
26	Contribution of IL12A and IL12B polymorphisms to the risk of cervical cancer. <i>Pathology and Oncology Research</i> , <b>2012</b> , 18, 997-1002	2.6	17
25	Folate and choline metabolism gene variants in relation to ovarian cancer risk in the Polish population. <i>Molecular Biology Reports</i> , <b>2012</b> , 39, 5553-60	2.8	26

24	No association of monocyte chemoattractant protein-1 -2518 A/G polymorphism with the risk of primary glomerulonephritis in the Polish population. <i>Molecular Biology Reports</i> , <b>2012</b> , 39, 5933-41	2.8	5
23	Novel MSX1 mutation in a family with autosomal-dominant hypodontia of second premolars and third molars. <i>Archives of Oral Biology</i> , <b>2012</b> , 57, 790-5	2.8	32
22	Polymorphic variants of folate and choline metabolism genes and the risk of endometriosis-associated infertility. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , <b>2011</b> , 157, 67-72	2.4	7
21	Polymorphisms in CHDH gene and the risk of tooth agenesis. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2011</b> , 91, 169-76		4
20	Polymorphisms of stress-related genes and the risk of nonsyndromic cleft lip with or without cleft palate. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2011</b> , 91, 948-55		8
19	Folate and choline metabolism gene variants and development of uterine cervical carcinoma. <i>Clinical Biochemistry</i> , <b>2011</b> , 44, 596-600	3.5	33
18	Polymorphisms located in the region containing BHMT and BHMT2 genes as maternal protective factors for orofacial clefts. <i>European Journal of Oral Sciences</i> , <b>2010</b> , 118, 325-32	2.3	24
17	Associations of folate and choline metabolism gene polymorphisms with orofacial clefts. <i>Journal of Medical Genetics</i> , <b>2010</b> , 47, 809-15	5.8	60
16	Polymorphic variants of genes related to arginine metabolism and the risk of orofacial clefts. <i>Archives of Oral Biology</i> , <b>2010</b> , 55, 861-6	2.8	3
15	Association between genetic variants of reported candidate genes or regions and risk of cleft lip with or without cleft palate in the polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2010</b> , 88, 538-45		47
14	Lack of association of polymorphic variants of genes encoding zinc transporters with the risk of orofacial cleft-affected pregnancies. <i>Folia Histochemica Et Cytobiologica</i> , <b>2010</b> , 48, 618-23	1.4	2
13	Identification of microdeletions in candidate genes for cleft lip and/or palate. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , <b>2009</b> , 85, 42-51		44
12	Polymorphic variants of genes encoding main antioxidant enzymes and the risk of CL/P-affected pregnancies. <i>Clinical Biochemistry</i> , <b>2007</b> , 40, 416-9	3.5	5
11	A novel c.581C>T transition localized in a highly conserved homeobox sequence of MSX1: is it responsible for oligodontia?. <i>Journal of Applied Genetics</i> , <b>2006</b> , 47, 159-64	2.5	35
10	Natural selection and molecular evolution in primate PAX9 gene, a major determinant of tooth development. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2006</b> , 103, 5676-81	11.5	42
9	Association of 677C>T polymorphism of methylenetetrahydrofolate reductase (MTHFR) gene with bipolar disorder and schizophrenia. <i>Neuroscience Letters</i> , <b>2006</b> , 400, 267-71	3.3	50
8	Maternal MTR genotype contributes to the risk of non-syndromic cleft lip and palate in the Polish population. <i>Clinical Genetics</i> , <b>2006</b> , 69, 512-7	4	69
7	A novel mutation in PAX9 causes familial form of molar oligodontia. <i>European Journal of Human Genetics</i> , <b>2006</b> , 14, 173-9	5.3	42

6	Axis inhibition protein 2 (AXIN2) polymorphisms may be a risk factor for selective tooth agenesis. <i>Journal of Human Genetics</i> , <b>2006</b> , 51, 262-266	4.3	80
5	Gene symbol: IRF6. Disease: Van der Woude syndrome. <i>Human Genetics</i> , <b>2005</b> , 116, 534	6.3	3
4	Expression of modified Cry1Ac gene of <i>Bacillus thuringiensis</i> in transgenic tobacco plants. <i>Molecular Biotechnology</i> , <b>2004</b> , 26, 17-26	3	8
3	Novel mutation in the paired box sequence of PAX9 gene in a sporadic form of oligodontia. <i>European Journal of Oral Sciences</i> , <b>2003</b> , 111, 272-6	2.3	59
2	Molecular basis of non-syndromic tooth agenesis: mutations of MSX1 and PAX9 reflect their role in patterning human dentition. <i>European Journal of Oral Sciences</i> , <b>2003</b> , 111, 365-70	2.3	76
1	The novel polymorphic variants within the paired box of the PAX9 gene are associated with selective tooth agenesis. <i>Folia Histochemica Et Cytobiologica</i> , <b>2001</b> , 39, 111-2	1.4	6