

Munis Dundar

List of Publications by Year in descending order

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202
papers

2,490
citations

393982

19
h-index

243296

44
g-index

235
all docs

235
docs citations

235
times ranked

3415
citing authors

#	ARTICLE	IF	CITATIONS
1	Familial Mediterranean Fever (FMF) in Turkey. <i>Medicine (United States)</i> , 2005, 84, 1-11.	0.4	651
2	General Report & Recommendations in Predictive, Preventive and Personalised Medicine 2012: White Paper of the European Association for Predictive, Preventive and Personalised Medicine. <i>EPMA Journal</i> , 2012, 3, 14.	3.3	218
3	Loss of Dermatan-4-Sulfotransferase 1 Function Results in Adducted Thumb-Clubfoot Syndrome. <i>American Journal of Human Genetics</i> , 2009, 85, 873-882.	2.6	134
4	Loss of dermatan sulfate epimerase (DSE) function results in musculocontractural Ehlers-Danlos syndrome. <i>Human Molecular Genetics</i> , 2013, 22, 3761-3772.	1.4	78
5	Apolipoprotein E3/E3 Genotype Decreases the Risk of Pituitary Dysfunction after Traumatic Brain Injury due to Various Causes: Preliminary Data. <i>Journal of Neurotrauma</i> , 2008, 25, 1071-1077.	1.7	71
6	Circulating microRNAs in patients with non-alcoholic fatty liver disease. <i>World Journal of Hepatology</i> , 2014, 6, 613.	0.8	67
7	Genotype-phenotype correlation in children with familial Mediterranean fever in a Turkish population. <i>Pediatrics International</i> , 2008, 50, 208-212.	0.2	64
8	Common Familial Mediterranean Fever gene mutations in a Turkish cohort. <i>Molecular Biology Reports</i> , 2011, 38, 5065-5069.	1.0	56
9	Genetic background of supernumerary teeth. <i>European Journal of Dentistry</i> , 2015, 09, 153-158.	0.8	54
10	An autosomal recessive adducted thumb-club foot syndrome observed in Turkish cousins. <i>Clinical Genetics</i> , 2008, 51, 61-64.	1.0	42
11	Inherited diseases and syndromes leading to aortic aneurysms and dissections. <i>European Journal of Cardio-thoracic Surgery</i> , 2009, 35, 931-940.	0.6	42
12	Progress towards the "Golden Age" of biotechnology. <i>Current Opinion in Biotechnology</i> , 2013, 24, S6-S13.	3.3	32
13	A case with adducted thumb and club foot syndrome. <i>Clinical Dysmorphology</i> , 2001, 10, 291-293.	0.1	27
14	Diagnosis of intracranial calcification and hemorrhage in pediatric patients: Comparison of quantitative susceptibility mapping and phase images of susceptibility-weighted imaging. <i>Diagnostic and Interventional Imaging</i> , 2017, 98, 707-714.	1.8	27
15	5,10-Methylenetetrahydrofolate reductase C677T gene polymorphism in Behcet's patients with or without ocular involvement. <i>British Journal of Ophthalmology</i> , 2005, 89, 1634-1637.	2.1	26
16	The prevalence of non-classic adrenal hyperplasia among Turkish women with hyperandrogenism. <i>Gynecological Endocrinology</i> , 2010, 26, 139-143.	0.7	24
17	Scottish frequency of the common G985 mutation in the medium-chain acyl-CoA dehydrogenase (MCAD) gene and the role of MCAD deficiency in sudden infant death syndrome (SIDS). <i>Journal of Inherited Metabolic Disease</i> , 1993, 16, 991-993.	1.7	22
18	A novel acropectoral syndrome maps to chromosome 7q36. <i>Journal of Medical Genetics</i> , 2001, 38, 304-309.	1.5	21

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19	The Age Structure, Stringency Policy, Income, and Spread of Coronavirus Disease 2019: Evidence From 209 Countries. <i>Frontiers in Psychology</i> , 2020, 11, 632192.	1.1	21
20	The frequencies of Y chromosome microdeletions in infertile males. <i>Turkish Journal of Urology</i> , 2018, 44, 389-392.	1.3	20
21	Sacrococcygeal teratoma in a fetus with prenatally diagnosed partial trisomy 10q (10q24.3â†’qter) and partial monosomy 17p (p13.3â†’pter). <i>Prenatal Diagnosis</i> , 2007, 27, 365-368.	1.1	19
22	Nitric oxide synthase in diabetic rat testicular tissue and the effects of pentoxifylline therapy. <i>Systems Biology in Reproductive Medicine</i> , 2016, 62, 22-30.	1.0	19
23	Female-to-male transsexual with 47,XXX karyotype. <i>Biological Psychiatry</i> , 2000, 48, 1116-1117.	0.7	18
24	Genetic background, nutrition and obesity: a review. <i>European Review for Medical and Pharmacological Sciences</i> , 2019, 23, 1751-1761.	0.5	17
25	Increased vitamin D receptor gene expression and rs11568820 and rs4516035 promoter polymorphisms in autistic disorder. <i>Molecular Biology Reports</i> , 2018, 45, 541-546.	1.0	16
26	Bacteriophages presence in nature and their role in the natural selection of bacterial populations. <i>Acta Biomedica</i> , 2020, 91, e2020024.	0.2	16
27	Sexing Greater Flamingo Chicks from Feather Bulb DNA. <i>Waterbirds</i> , 2007, 30, 450-453.	0.2	15
28	Maternal uniparental isodisomy is responsible for serious molybdenum cofactor deficiency. <i>Developmental Medicine and Child Neurology</i> , 2010, 52, 868-872.	1.1	15
29	Loss of dermatanâ€”sulfotransferase 1 (D4ST1/ <i>CHST14</i>) function represents the first dermatan sulfate biosynthesis defect, â€œdermatan sulfateâ€”deficient adducted thumbâ€”clubfoot syndromeâ€”. <i>Human Mutation</i> , 2011, 32, 484-485.	1.1	15
30	Biotechnology worldwide and the â€”European Biotechnology Thematic Networkâ€” Association (EBTNA). <i>Current Opinion in Biotechnology</i> , 2011, 22, S7-S14.	3.3	15
31	A Glutamine Repeat Variant of the RUNX2 Gene Causes Cleidocranial Dysplasia. <i>Molecular Syndromology</i> , 2015, 6, 50-53.	0.3	15
32	A brief overview of global biotechnology. <i>Biotechnology and Biotechnological Equipment</i> , 2021, 35, S5-S14.	0.5	14
33	Natural compounds as inhibitors of SARS-CoV-2 endocytosis: A promising approach against COVID-19. <i>Acta Biomedica</i> , 2020, 91, e2020008.	0.2	14
34	A pilot study on the preventative potential of alpha-cyclodextrin and hydroxytyrosol against SARS-CoV-2 transmission. <i>Acta Biomedica</i> , 2020, 91, e2020022.	0.2	14
35	The Deletion Polymorphism of the Angiotensin-Converting Enzyme Gene Is Associated with Acute Aortic Dissection. <i>Tohoku Journal of Experimental Medicine</i> , 2009, 219, 33-37.	0.5	13
36	Is there relation between COL4A1/A2 mutations and antenatally detected fetal intraventricular hemorrhage?. <i>Child's Nervous System</i> , 2014, 30, 419-424.	0.6	12

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37	Ameliorative effects of pentoxifylline on NOS induced by diabetes in rat kidney. <i>Renal Failure</i> , 2016, 38, 605-613.	0.8	12
38	Genetic tests for low- and middle-income countries: a literature review. <i>Genetics and Molecular Research</i> , 2017, 16, .	0.3	12
39	The effects of streptozotocin-induced diabetes on ghrelin expression in rat testis: biochemical and immunohistochemical study. <i>Folia Histochemica Et Cytobiologica</i> , 2015, 53, 26-34.	0.6	12
40	Detection of p16 promotor hypermethylation in "Maras powder" and tobacco users. <i>Cancer Epidemiology</i> , 2009, 33, 47-50.	0.8	11
41	Idiopathic hirsutism: local and peripheral expression of aromatase (CYP19A) and 5 α -reductase genes (SRD5A1 and SRD5A2). <i>Fertility and Sterility</i> , 2011, 96, 479-482.	0.5	11
42	Expression of Biologically Active Human Interferon Gamma in the Milk of Transgenic Mice Under the Control of the Murine Whey Acidic Protein Gene Promoter. <i>Biochemical Genetics</i> , 2011, 49, 251-257.	0.8	11
43	The role of TNF- α and PAI-1 gene polymorphisms in familial Mediterranean fever. <i>Modern Rheumatology</i> , 2013, 23, 140-145.	0.9	11
44	The association of endothelin-1 levels with renal survival in polycystic kidney disease patients. <i>Journal of Nephrology</i> , 2019, 32, 83-91.	0.9	11
45	TIE1 as a Candidate Gene for Lymphatic Malformations with or without Lymphedema. <i>International Journal of Molecular Sciences</i> , 2020, 21, 6780.	1.8	11
46	Prenatal Diagnosis of a Fetus with Partial Trisomy 7p. <i>Fetal Diagnosis and Therapy</i> , 2007, 22, 229-232.	0.6	10
47	The Effect of Maras Powder on DNA Methylation and Micronucleus Formation in Human Buccal Tissue. <i>Journal of Toxicology and Environmental Health - Part A: Current Issues</i> , 2008, 71, 396-404.	1.1	10
48	The Frequency of CYP 21 Gene Mutations in Turkish Women with Hyperandrogenism. <i>Experimental and Clinical Endocrinology and Diabetes</i> , 2009, 117, 205-208.	0.6	10
49	Lack of association between the Glu298Asp polymorphism of endothelial nitric oxide synthase and slow coronary flow in the Turkish population. <i>Canadian Journal of Cardiology</i> , 2009, 25, e69-e72.	0.8	10
50	Unbalanced 3;22 translocation with 22q11 and 3p deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 2791-2795.	0.7	10
51	A Novel COL4A3 Mutation Causes Autosomal-Recessive Alport Syndrome in a Large Turkish Family. <i>Genetic Testing and Molecular Biomarkers</i> , 2013, 17, 260-264.	0.3	10
52	Neurological Manifestations in Familial Mediterranean Fever: Results of 22 Children from a Reference Center in Kayseri, an Urban Area in Central Anatolia, Turkey. <i>Neuropediatrics</i> , 2017, 48, 079-085.	0.3	10
53	Comprehensive genotyping of Turkish women with hirsutism. <i>Journal of Endocrinological Investigation</i> , 2019, 42, 1077-1087.	1.8	10
54	Current state and prospects of biotechnology in Central and Eastern European countries. Part I: Visegrad countries (CZ, H, PL, SK). <i>Critical Reviews in Biotechnology</i> , 2019, 39, 114-136.	5.1	10

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55	The role of androgen receptor CAG repeat polymorphism in androgen excess disorder and idiopathic hirsutism. <i>Journal of Endocrinological Investigation</i> , 2020, 43, 1271-1281.	1.8	10
56	Cytogenetic results of patients with infertility in middle anatolia, Turkey: do heterochromatin polymorphisms affect fertility?. <i>Journal of Reproduction and Infertility</i> , 2010, 11, 179-81.	1.0	10
57	Adrenal axis functions in patients with familial Mediterranean fever. <i>Clinical Rheumatology</i> , 2006, 25, 458-461.	1.0	9
58	How the I1307K adenomatous polyposis coli gene variant contributes in the assessment of risk of colorectal cancer, but not stomach cancer, in a Turkish population. <i>Cancer Genetics and Cytogenetics</i> , 2007, 177, 95-97.	1.0	9
59	Frank-ter Haar syndrome with unusual clinical features. <i>European Journal of Medical Genetics</i> , 2009, 52, 247-249.	0.7	9
60	Atypical presentation and a novel mutation in ALMS1: implications for clinical and molecular diagnostic strategies for Alstr�m syndrome. <i>Clinical Genetics</i> , 2013, 83, 96-98.	1.0	9
61	Is idiopathic hirsutism (IH) really idiopathic? mRNA expressions of skin steroidogenic enzymes in women with IH. <i>European Journal of Endocrinology</i> , 2015, 173, 447-454.	1.9	9
62	The molecular basis and genotype�phenotype correlations of congenital adrenal hyperplasia (CAH) in Anatolian population. <i>Molecular Biology Reports</i> , 2019, 46, 3677-3690.	1.0	9
63	The role of TNF-� and PAI-1 gene polymorphisms in familial Mediterranean fever. <i>Modern Rheumatology</i> , 2013, 23, 140-145.	0.9	9
64	A molecular analysis of familial Mediterranean fever disease in a cohort of Turkish patients. <i>Annals of Saudi Medicine</i> , 2012, 32, 343-348.	0.5	9
65	ICR1 Epimutations in 11p15 are Restricted to Patients with Silver-Russell Syndrome Features. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2008, 21, 59-62.	0.4	8
66	Healthcare in overview of Turkey. <i>EPMA Journal</i> , 2010, 1, 587-594.	3.3	8
67	COVID-19 vaccine candidates and vaccine development platforms available worldwide. <i>Journal of Pharmaceutical Analysis</i> , 2021, 11, 675-682.	2.4	8
68	Pilot study for the evaluation of safety profile of a potential inhibitor of SARS-CoV-2 endocytosis. <i>Acta Biomedica</i> , 2020, 91, e2020009.	0.2	8
69	A case of ambiguous genitalia presenting with a 45,X/46,Xr(Y)(p11.2;q11.23)/47,X,idi(Y)(p11.2),idi(Y)(p11.2) karyotype. <i>Annales De G�n�tique</i> , 2001, 44, 5-8.	0.4	7
70	Etiopathogenesis of Sheehan�s Syndrome: Roles of Coagulation Factors and TNF-Alpha. <i>International Journal of Endocrinology</i> , 2014, 2014, 1-6.	0.6	7
71	Genotoxic Effects of some Antituberculosis Drugs and Mixtures in Rats. <i>Drug Research</i> , 2015, 65, 219-222.	0.7	7
72	Research of genetic bases of hereditary non-syndromic hearing loss. <i>Turk Pediatri Arsivi</i> , 2017, 52, 122-132.	0.9	7

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73	Comparing expression levels of PERIOD genes PER1, PER2 and PER3 in chronic insomnia patients and medical staff working in the night shift. <i>Sleep Medicine</i> , 2020, 73, 101-105.	0.8	7
74	Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium. <i>Functional and Integrative Genomics</i> , 2022, 22, 291-315.	1.4	7
75	Current State of Biotechnology in Turkey. <i>Current Opinion in Biotechnology</i> , 2011, 22, S3-S6.	3.3	6
76	Comparison between American and European legislation in the therapeutical and alimentary bacteriophage usage. <i>Acta Biomedica</i> , 2020, 91, e2020023.	0.2	6
77	A c.1244G A (p.Arg415Gln) mutation in SH3BP2 gene causes cherubism in a Turkish family: Report of a family with review of the literature. <i>Medicina Oral, Patologia Oral Y Cirugia Bucal</i> , 2014, 19, e340-e344.	0.7	5
78	The Association of Brain-Derived Neurotrophic Factor Gene Polymorphism with Obstructive Sleep Apnea Syndrome and Obesity. <i>Lung</i> , 2016, 194, 839-846.	1.4	5
79	Pharmacologically active fractions of <i>Sideritis</i> spp. and their use in inherited eye diseases. <i>The EuroBiotech Journal</i> , 2017, 1, 6-10.	0.5	5
80	Current state and prospects of biotechnology in Central and Eastern European countries. Part II: new and preaccession EU countries(CRO, RO, B&H, SRB). <i>Critical Reviews in Biotechnology</i> , 2019, 39, 137-155.	5.1	5
81	Possible Role of the <i>RORC</i> Gene in Primary and Secondary Lymphedema: Review of the Literature and Genetic Study of Two Rare Causative Variants. <i>Lymphatic Research and Biology</i> , 2021, 19, 129-133.	0.5	5
82	Patient with Weismann-Netter and Stuhl (Toxopachyosteosis) Syndrome with Communicant Hydrocephalus and Arachnoid Cyst. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2000, 13, 211-5.	0.4	4
83	A family with two different chromosomal translocations. <i>Annales De G�n�tologie</i> , 2002, 45, 185-187.	0.4	4
84	Polycystic kidney disease, biliary dysgenesis in a patient with Larsen's syndrome. <i>Clinical Genetics</i> , 1997, 51, 408-411.	1.0	4
85	Prediction, prevention and personalisation of medication for the prenatal period: genetic prenatal tests for both rare and common diseases. <i>EPMA Journal</i> , 2011, 2, 181-195.	3.3	4
86	Segregation Analysis of Rare NRP1 and NRP2 Variants in Families with Lymphedema. <i>Genes</i> , 2020, 11, 1361.	1.0	4
87	Two rare <i>PROX1</i> variants in patients with lymphedema. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1424.	0.6	4
88	<i>NOTCH1</i> : Review of its role in lymphatic development and study of seven families with rare pathogenic variants. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1529.	0.6	4
89	Current and Future Therapeutic Strategies for Limb Girdle Muscular Dystrophy Type R1: Clinical and Experimental Approaches. <i>Pathophysiology</i> , 2021, 28, 238-249.	1.0	4
90	The Effects of Long-Term Diabetes on Ghrelin Expression in Rat Stomachs. <i>Advances in Clinical and Experimental Medicine</i> , 2015, 24, 401-407.	0.6	4

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91	Future Biotechnology. The EuroBiotech Journal, 2019, 3, 53-56.	0.5	4
92	Genetic testing for ocular coloboma. The EuroBiotech Journal, 2017, 1, 29-31.	0.5	4
93	Genetic testing for autonomic dysfunction or dysautonomias. Acta Biomedica, 2020, 91, e2020002.	0.2	4
94	Ethics committees for clinical experimentation at international level with a focus on Italy. Acta Biomedica, 2020, 91, e2020016.	0.2	4
95	A novel missense mutation outside the <sc>DNAJ</sc> domain of <sc><i>DNAJC21</i></sc> is associated with <sc>Shwachmanâ€™Diamond</sc> syndrome. British Journal of Haematology, 2022, 197, .	1.2	4
96	A Turner patient with a 45,X,t(1;2) (q41;p11.2) karyotype. Annales De GÃ©nÃ©tique, 2002, 45, 181-183.	0.4	3
97	Isolated congenital anonychia cases with coincident chromosomal fragility. Annales De GÃ©nÃ©tique, 2004, 47, 381-386.	0.4	3
98	Genetic expressions of thrombophilic factors in patients with Sheehanâ€™s syndrome. Gynecological Endocrinology, 2016, 32, 908-911.	0.7	3
99	Prenatal diagnosis of upper extremity malformations with ultrasonography: Diagnostic features and perinatal outcome. Journal of Clinical Ultrasound, 2017, 45, 267-276.	0.4	3
100	Genetic testing for central areolar choroidal dystrophy. The EuroBiotech Journal, 2017, 1, 23-25.	0.5	3
101	The effects of O ⁶ -methyl guanine DNA-methyl transferase promotor methylation and CpG1, CpG2, CpG3 and CpG4 methylation on treatment response and their prognostic significance in patients with glioblastoma. Balkan Journal of Medical Genetics, 2020, 23, 33-41.	0.5	3
102	BRCA Variations Risk Assessment in Breast Cancers Using Different Artificial Intelligence Models. Genes, 2021, 12, 1774.	1.0	3
103	A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia. Clinica Chimica Acta, 2022, 529, 61-66.	0.5	3
104	Can the classical euchromatic variants of 9q12/qh+ cause recurrent abortions?. Genetic Counseling, 2008, 19, 281-6.	0.1	3
105	<i>CDH5</i>, a Possible New Candidate Gene for Genetic Testing of Lymphedema. Lymphatic Research and Biology, 2021, , .	0.5	3
106	Neonatal Diabetes, Congenital Hypothyroidism, and Congenital Glaucoma Coexistence: A Case of GLIS3 Mutation. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2023, 15, 426-430.	0.4	3
107	Germline landscape of BRCA by 7-site collaborations as a BRCA consortium in Turkey. Breast, 2022, 65, 15-22.	0.9	3
108	Congenital alacrima in a patient with G (Opitz Frias) syndrome. Human Genetics, 1996, 97, 540-542.	1.8	2

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109	Genetic testing for familial exudative vitreoretinopathy. The EuroBiotech Journal, 2017, 1, 51-53.	0.5	2
110	Genetic testing for pattern dystrophies. The EuroBiotech Journal, 2017, 1, 86-88.	0.5	2
111	Clinical Characteristics of Cases with Spinal Muscular Atrophy. Guncel Pediatri, 2016, 14, 18-22.	0.1	2
112	Genetic testing for gyrate atrophy of the choroid and retina. The EuroBiotech Journal, 2017, 1, 54-56.	0.5	2
113	Genetic testing for Ebstein anomaly. The EuroBiotech Journal, 2018, 2, 55-57.	0.5	2
114	Anadoluâ€™daki Tâ±bbÄ±n DoÄŸuÄŸu, DÄ¼nyadaki Ä°lk Tâ±p Okulu Olarak: Gevher Nesibe Tâ±p Medresesi ve DarÄ¼ÄŸifasâ±. Bilimname: DÄ¼ÄŸÄ¼nce Platformu, 0, , 79-103.	0.3	2
115	Evaluation of the Results of Cases Prenatally Diagnosed as VSD. Erciyes Tip Dergisi, 2012, 34, 111-115.	0.1	1
116	A new syndrome of microtia with unilateral renal agenesis and short stature. American Journal of Medical Genetics, Part A, 2012, 158A, 1837-1840.	0.7	1
117	A case of SRY positive 46, XX male with speaking disorder. Journal of Biotechnology, 2015, 208, S85.	1.9	1
118	Expression of Ghrelin and GHSR-1a in Long Term Diabetic Rat's Kidney. Brazilian Archives of Biology and Technology, 2016, 59, .	0.5	1
119	Developments in biotechnology. Journal of Biotechnology, 2017, 256, S7.	1.9	1
120	Genetic testing for Usher syndrome. The EuroBiotech Journal, 2017, 1, 108-110.	0.5	1
121	Genetic testing for aortic valve stenosis. The EuroBiotech Journal, 2018, 2, 61-63.	0.5	1
122	Mutations in the ARAP3 Gene in Three Families with Primary Lymphedema Negative for Mutations in Known Lymphedema-Associated Genes. International Journal of Genomics, 2020, 2020, 1-9.	0.8	1
123	Are new genome variants detected in SARS-CoV-2 expected considering population dynamics in viruses?. The EuroBiotech Journal, 2021, 5, 1-3.	0.5	1
124	COVID-19 vaccines: Where do we stand?. The EuroBiotech Journal, 2021, 5, 4-7.	0.5	1
125	Genetic testing for Bietti crystalline dystrophy. The EuroBiotech Journal, 2017, 1, 20-22.	0.5	1
126	Genetic testing for Stargardt macular dystrophy. The EuroBiotech Journal, 2017, 1, 105-107.	0.5	1

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127	Genetic testing for tetralogy of Fallot. The EuroBiotech Journal, 2018, 2, 71-73.	0.5	1
128	Quality assurance of genetic laboratories and the EBTNA practice certification, a simple standardization assurance system for a laboratory network. The EuroBiotech Journal, 2018, 2, 215-222.	0.5	1
129	Holt-Oram syndrome in two generations with translocation t(9;15)(p12;q11.2). Annals of Saudi Medicine, 2008, 28, 209.	0.5	1
130	Holt-Oram syndrome in two generations with translocation t(9;15)(p12;q11.2). Annals of Saudi Medicine, 2008, 28, 209-212.	0.5	1
131	Genetic Disorders of Pituitary Development in Patients with Sheehan'S Syndrome. Acta Endocrinologica, 2016, 12, 413-417.	0.1	1
132	Advances in biotechnology: Genomics and genome editing. The EuroBiotech Journal, 2017, 1, 2-9.	0.5	1
133	Genetic testing for color vision deficiency. The EuroBiotech Journal, 2017, 1, 32-34.	0.5	1
134	Genetic testing for non syndromic retinitis pigmentosa. The EuroBiotech Journal, 2017, 1, 92-95.	0.5	1
135	Genetic testing for Refsum disease. The EuroBiotech Journal, 2017, 1, 89-91.	0.5	1
136	Propranolol decreases DRD3 and SLC1A2 gene expression in patients with essential tremor. Universa Medicina, 2020, 39, 105-112.	0.1	1
137	A very rare cause of arthrogryposis multiplex congenita: a novel mutation in <i>TOR1A</i> . Journal of Pediatric Endocrinology and Metabolism, 2022, 35, 845-850.	0.4	1
138	Diagnosing Alström syndrome in a patient followed up with syndromic obesity for years. Intractable and Rare Diseases Research, 2022, 11, 84-86.	0.3	1
139	A case with Waardenburg syndrome presenting with two separate translocations ??? one reciprocal and one complex. Clinical Dysmorphology, 2001, 10, 65-66.	0.1	0
140	The increasing importance of Medical Genetics in Turkey. Current Opinion in Biotechnology, 2011, 22, S42-S43.	3.3	0
141	Prenatally detected de novo 46, XX, t(21;21)(p12;p12) at chorionic villus sampling. Current Opinion in Biotechnology, 2011, 22, S107.	3.3	0
142	Analysing the role of MDM2 SNP309 in patients with glioblastoma multiforme. Current Opinion in Biotechnology, 2013, 24, S98.	3.3	0
143	Innovations in biotechnology. Journal of Biotechnology, 2015, 208, S5.	1.9	0
144	A novel nonsense mutation in GALNS gene in family with MPS4A diagnosed child. Journal of Biotechnology, 2016, 231, S108.	1.9	0

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145	A case of XYY male patient with micropenis. Journal of Biotechnology, 2016, 231, S109.	1.9	0
146	Perspectives of biotechnology. Journal of Biotechnology, 2016, 231, S4.	1.9	0
147	Editorial. Journal of Biotechnology, 2016, 231, S1-S3.	1.9	0
148	The effect of parental 5,10-methylenetetrahydrofolate reductase 677C/T and 1298A/C gene polymorphisms on response to single-dose methotrexate in tubal ectopic pregnancy. Journal of Maternal-Fetal and Neonatal Medicine, 2017, 30, 1232-1237.	0.7	0
149	Prenatal diagnosis of a foetus with partial monosomy 4p and partial trisomy 13q. Journal of Biotechnology, 2017, 256, S76.	1.9	0
150	The effect of CYP2C19 * 2 polymorphism on clopidogrel resistance in COPD patients. Journal of Biotechnology, 2017, 256, S80.	1.9	0
151	Frequency of chromosome variants in families with recurrent pregnancy loss and statistical analysis of infertility. Journal of Biotechnology, 2017, 256, S76.	1.9	0
152	Genetic testing for congenital stationary night blindness. The EuroBiotech Journal, 2017, 1, 38-40.	0.5	0
153	Genetic testing in translational ophthalmology. The EuroBiotech Journal, 2017, 1, 1-5.	0.5	0
154	Genetic testing for Norrie disease. The EuroBiotech Journal, 2017, 1, 77-79.	0.5	0
155	Editorial. Journal of Biotechnology, 2018, 280, S1-S2.	1.9	0
156	The Story of a Ship Journey, Malaria, and the HBB Gene IVS-II-745 Mutation: Circassian Immigration to Cyprus. Global Medical Genetics, 2021, 08, 069-071.	0.4	0
157	Detection of mutations in CML patients resistant to tyrosine kinase inhibitor: imatinib mesylate therapy. Medical Oncology, 2021, 38, 120.	1.2	0
158	Megarbane syndrome. Indian Journal of Human Genetics, 2008, 14, 27.	0.7	0
159	Overview of the Healthcare System in Turkey. Advances in Predictive, Preventive and Personalised Medicine, 2012, , 167-187.	0.6	0
160	Autozygosity in a Turkish family with scoliosis, blindness, and arachnodactyly syndrome. Annals of Saudi Medicine, 2015, 35, 462-467.	0.5	0
161	Genetic testing for infantile nystagmus. The EuroBiotech Journal, 2017, 1, 57-59.	0.5	0
162	Genetic testing for retinitis punctata albescens/fundus albipunctatus. The EuroBiotech Journal, 2017, 1, 96-98.	0.5	0

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163	Genetic testing for X-linked juvenile retinoschisis. The EuroBiotech Journal, 2017, 1, 111-113.	0.5	0
164	Genetic testing for Sorsby's fundus dystrophy. The EuroBiotech Journal, 2017, 1, 102-104.	0.5	0
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