Munis Dundar

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7645382/publications.pdf

Version: 2024-02-01

202	2,490	394421	243625
papers	citations	19 h-index	44 g-index

235	235	235	3415
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Familial Mediterranean Fever (FMF) in Turkey. Medicine (United States), 2005, 84, 1-11.	1.0	651
2	General Report & Recommendations in Predictive, Preventive and Personalised Medicine 2012: White Paper of the European Association for Predictive, Preventive and Personalised Medicine. EPMA Journal, 2012, 3, 14.	6.1	218
3	Loss of Dermatan-4-Sulfotransferase 1 Function Results in Adducted Thumb-Clubfoot Syndrome. American Journal of Human Genetics, 2009, 85, 873-882.	6.2	134
4	Loss of dermatan sulfate epimerase (DSE) function results in musculocontractural Ehlersâ€“Danlos syndrome. Human Molecular Genetics, 2013, 22, 3761-3772.	2.9	78
5	Apolipoprotein E3/E3 Genotype Decreases the Risk of Pituitary Dysfunction after Traumatic Brain Injury due to Various Causes: Preliminary Data. Journal of Neurotrauma, 2008, 25, 1071-1077.	3.4	71
6	Circulating microRNAs in patients with non-alcoholic fatty liver disease. World Journal of Hepatology, 2014, 6, 613.	2.0	67
7	Genotypeâ€“phenotype correlation in children with familial Mediterranean fever in a Turkish population. Pediatrics International, 2008, 50, 208-212.	0.5	64
8	Common Familial Mediterranean Fever gene mutations in a Turkish cohort. Molecular Biology Reports, 2011, 38, 5065-5069.	2.3	56
9	Genetic background of supernumerary teeth. European Journal of Dentistry, 2015, 09, 153-158.	1.7	54
10	An autosomal recessive adducted thumb-club foot syndrome observed in Turkish cousins. Clinical Genetics, 2008, 51, 61-64.	2.0	42
11	Inherited diseases and syndromes leading to aortic aneurysms and dissections. European Journal of Cardio-thoracic Surgery, 2009, 35, 931-940.	1.4	42
12	Progress towards the â€˜Golden Ageâ€™ of biotechnology. Current Opinion in Biotechnology, 2013, 24, S6-S13.	6.6	32
13	A case with adducted thumb and club foot syndrome. Clinical Dysmorphology, 2001, 10, 291-293.	0.3	27
14	Diagnosis of intracranial calcification and hemorrhage in pediatric patients: Comparison of quantitative susceptibility mapping and phase images of susceptibility-weighted imaging. Diagnostic and Interventional Imaging, 2017, 98, 707-714.	3.2	27
15	5,10-Methylenetetrahydrofolate reductase C677T gene polymorphism in Behcet's patients with or without ocular involvement. British Journal of Ophthalmology, 2005, 89, 1634-1637.	3.9	26
16	The prevalence of non-classic adrenal hyperplasia among Turkish women with hyperandrogenism. Gynecological Endocrinology, 2010, 26, 139-143.	1.7	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). Journal of Inherited Metabolic Disease, 1993, 16, 991-993.	3.6	22
18	A novel acropectoral syndrome maps to chromosome 7q36. Journal of Medical Genetics, 2001, 38, 304-309.	3.2	21

#	Article	IF	Citations
19	The Age Structure, Stringency Policy, Income, and Spread of Coronavirus Disease 2019: Evidence From 209 Countries. Frontiers in Psychology, 2020, 11, 632192.	2.1	21
20	The frequencies of Y chromosome microdeletions in infertile males. Turkish Journal of Urology, 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). Prenatal Diagnosis, 2007, 27, 365-368.	2.3	19
22	Nitric oxide synthase in diabetic rat testicular tissue and the effects of pentoxifylline therapy. Systems Biology in Reproductive Medicine, 2016, 62, 22-30.	2.1	19
23	Female-to-male transsexual with 47,XXX karyotype. Biological Psychiatry, 2000, 48, 1116-1117.	1.3	18
24	Genetic background, nutrition and obesity: a review. European Review for Medical and Pharmacological Sciences, 2019, 23, 1751-1761.	0.7	17
25	Increased vitamin D receptor gene expression and rs11568820 and rs4516035 promoter polymorphisms in autistic disorder. Molecular Biology Reports, 2018, 45, 541-546.	2.3	16
26	Bacteriophages presence in nature and their role in the natural selection of bacterial populations. Acta Biomedica, 2020, 91, e2020024.	0.3	16
27	Sexing Greater Flamingo Chicks from Feather Bulb DNA. Waterbirds, 2007, 30, 450-453.	0.3	15
28	Maternal uniparental isodisomy is responsible for serious molybdenum cofactor deficiency. Developmental Medicine and Child Neurology, 2010, 52, 868-872.	2.1	15
29	Loss of dermatanâ€4â€sulfotransferase 1 (D4ST1/ CHST14) function represents the first dermatan sulfate biosynthesis defect, â€œdermatan sulfateâ€deficient adducted thumbâ€“clubfoot syndromeâ€• Human Mutation, 2011, 32, 484-485.	2.5	15
30	Biotechnology worldwide and the â€˜European Biotechnology Thematic Networkâ€™ Association (EBTNA). Current Opinion in Biotechnology, 2011, 22, S7-S14.	6.6	15
31	A Glutamine Repeat Variant of the RUNX2 Gene Causes Cleidocranial Dysplasia. Molecular Syndromology, 2015, 6, 50-53.	0.8	15
32	A brief overview of global biotechnology. Biotechnology and Biotechnological Equipment, 2021, 35, S5-S14.	1.3	14
33	Natural compounds as inhibitors of SARS-CoV-2 endocytosis: A promising approach against COVID-19. Acta Biomedica, 2020, 91, e2020008.	0.3	14
34	A pilot study on the preventative potential of alpha-cyclodextrin and hydroxytyrosol against SARS-CoV-2 transmission. Acta Biomedica, 2020, 91, e2020022.	0.3	14
35	The Deletion Polymorphism of the Angiotensin-Converting Enzyme Gene Is Associated with Acute Aortic Dissection. Tohoku Journal of Experimental Medicine, 2009, 219, 33-37.	1.2	13
36	Is there relation between COL4A1/A2 mutations and antenatally detected fetal intraventricular hemorrhage?. Child's Nervous System, 2014, 30, 419-424.	1.1	12

#	Article	IF	Citations
37	Ameliorative effects of pentoxifylline on NOS induced by diabetes in rat kidney. Renal Failure, 2016, 38, 605-613.	2.1	12
38	Genetic tests for low- and middle-income countries: a literature review. Genetics and Molecular Research, 2017, 16, .	0.2	12
39	The effects of streptozotocin-induced diabetes on ghrelin expression in rat testis: biochemical and immunohistochemical study. Folia Histochemica Et Cytobiologica, 2015, 53, 26-34.	1.5	12
40	Detection of p16 promotor hypermethylation in â€œMaras powderâ€•and tobacco users. Cancer Epidemiology, 2009, 33, 47-50.	1.9	11
41	Idiopathic hirsutism: local and peripheral expression of aromatase (CYP19A) and 5Î±-reductase genes (SRD5A1 and SRD5A2). Fertility and Sterility, 2011, 96, 479-482.	1.0	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. Biochemical Genetics, 2011, 49, 251-257.	1.7	11
43	The role of TNF-Î± and PAI-1 gene polymorphisms in familial Mediterranean fever. Modern Rheumatology, 2013, 23, 140-145.	1.8	11
44	The association of endothelin-1 levels with renal survival in polycystic kidney disease patients. Journal of Nephrology, 2019, 32, 83-91.	2.0	11
45	TIE1 as a Candidate Gene for Lymphatic Malformations with or without Lymphedema. International Journal of Molecular Sciences, 2020, 21, 6780.	4.1	11
46	Prenatal Diagnosis of a Fetus with Partial Trisomy 7p. Fetal Diagnosis and Therapy, 2007, 22, 229-232.	1.4	10
47	The Effect of Maras Powder on DNA Methylation and Micronucleus Formation in Human Buccal Tissue. Journal of Toxicology and Environmental Health - Part A: Current Issues, 2008, 71, 396-404.	2.3	10
48	The Frequency of CYP 21 Gene Mutations in Turkish Women with Hyperandrogenism. Experimental and Clinical Endocrinology and Diabetes, 2009, 117, 205-208.	1.2	10
49	Lack of association between the Glu298Asp polymorphism of endothelial nitric oxide synthase and slow coronary flow in the Turkish population. Canadian Journal of Cardiology, 2009, 25, e69-e72.	1.7	10
50	Unbalanced 3;22 translocation with 22q11 and 3p deletion syndrome. American Journal of Medical Genetics, Part A, 2010, 152A, 2791-2795.	1.2	10
51	A NovelCOL4A3Mutation Causes Autosomal-Recessive Alport Syndrome in a Large Turkish Family. Genetic Testing and Molecular Biomarkers, 2013, 17, 260-264.	0.7	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. Neuropediatrics, 2017, 48, 079-085.	0.6	10
53	Comprehensive genotyping of Turkish women with hirsutism. Journal of Endocrinological Investigation, 2019, 42, 1077-1087.	3.3	10
54	Current state and prospects of biotechnology in Central and Eastern European countries. Part I: Visegrad countries (CZ, H, PL, SK). Critical Reviews in Biotechnology, 2019, 39, 114-136.	9.0	10

#	Article	IF	Citations
55	The role of androgen receptor CAG repeat polymorphism in androgen excess disorder and idiopathic hirsutism. Journal of Endocrinological Investigation, 2020, 43, 1271-1281.	3.3	10
56	Cytogenetic results of patients with infertility in middle anatolia, Turkey: do heterochromatin polymorphisms affect fertility?. Journal of Reproduction and Infertility, 2010, 11, 179-81.	1.0	10
57	Adrenal axis functions in patients with familial Mediterranean fever. Clinical Rheumatology, 2006, 25, 458-461.	2.2	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. Cancer Genetics and Cytogenetics, 2007, 177, 95-97.	1.0	9
59	Frank-ter Haar syndrome with unusual clinical features. European Journal of Medical Genetics, 2009, 52, 247-249.	1.3	9
60	Atypical presentation and a novel mutation in ALMS1: implications for clinical and molecular diagnostic strategies for AlstrÃ¶m syndrome. Clinical Genetics, 2013, 83, 96-98.	2.0	9
61	Is idiopathic hirsutism (IH) really idiopathic? mRNA expressions of skin steroidogenic enzymes in women with IH. European Journal of Endocrinology, 2015, 173, 447-454.	3.7	9
62	The molecular basis and genotypeâ€“phenotype correlations of congenital adrenal hyperplasia (CAH) in Anatolian population. Molecular Biology Reports, 2019, 46, 3677-3690.	2.3	9
63	The role of TNF-Î± and PAI-1 gene polymorphisms in familial Mediterranean fever. Modern Rheumatology, 2013, 23, 140-145.	1.8	9
64	A molecular analysis of familial Mediterranean fever disease in a cohort of Turkish patients. Annals of Saudi Medicine, 2012, 32, 343-348.	1.1	9
65	ICR1 Epimutations in 11p15 are Restricted to Patients with Silver-Russell Syndrome Features. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 59-62.	0.9	8
66	Healthcare in overview of Turkey. EPMA Journal, 2010, 1, 587-594.	6.1	8
67	COVID-19 vaccine candidates and vaccine development platforms available worldwide. Journal of Pharmaceutical Analysis, 2021, 11, 675-682.	5.3	8
68	Pilot study for the evaluation of safety profile of a potential inhibitor of SARS-CoV-2 endocytosis. Acta Biomedica, 2020, 91, e2020009.	0.3	8
69	A case of ambiguous genitalia presenting with a 45,X/46,Xr(Y)(p11.2;q11.23)/47,X,idic(Y)(p11.2),idic(Y)(p11.2) karyotype. Annales De GÃ©nÃ©tique, 2001, 44, 5-8.	0.4	7
70	Etiopathogenesis of Sheehanâ€™s Syndrome: Roles of Coagulation Factors and TNF-Alpha. International Journal of Endocrinology, 2014, 2014, 1-6.	1.5	7
71	Genotoxic Effects of some Antituberculosis Drugs and Mixtures in Rats. Drug Research, 2015, 65, 219-222.	1.7	7
72	Research of genetic bases of hereditary non-syndromic hearing loss. Turk Pediatri Arsivi, 2017, 52, 122-132.	0.9	7

#	Article	IF	Citations
73	Comparing expression levels of PERIOD genes PER1, PER2 and PER3 in chronic insomnia patients and medical staff working in the night shift. Sleep Medicine, 2020, 73, 101-105.	1.6	7
74	Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium. Functional and Integrative Genomics, 2022, 22, 291-315.	3.5	7
75	Current State of Biotechnology in Turkey. Current Opinion in Biotechnology, 2011, 22, S3-S6.	6.6	6
76	Comparison between American and European legislation in the therapeutical and alimentary bacteriophage usage. Acta Biomedica, 2020, 91, e2020023.	0.3	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. Medicina Oral, Patologia Oral Y Cirugia Bucal, 2014, 19, e340-e344.	1.7	5
78	The Association of Brain-Derived Neurotrophic Factor Gene Polymorphism with Obstructive Sleep Apnea Syndrome and Obesity. Lung, 2016, 194, 839-846.	3.3	5
79	Pharmacologically active fractions of Sideritis spp. and their use in inherited eye diseases. The EuroBiotech Journal, 2017, 1, 6-10.	1.0	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). Critical Reviews in Biotechnology, 2019, 39, 137-155.	9.0	5
81	Possible Role of the RORC Gene in Primary and Secondary Lymphedema: Review of the Literature and Genetic Study of Two Rare Causative Variants. Lymphatic Research and Biology, 2021, 19, 129-133.	1.1	5
82	Patient with Weismann-Netter and Stuhl (Toxopachyosteosis) Syndrome with Communicant Hydrocephalus and Arachnoid Cyst. Journal of Pediatric Endocrinology and Metabolism, 2000, 13, 211-5.	0.9	4
83	A family with two different chromosomal translocations. Annales De GÃ©nÃ©tique, 2002, 45, 185-187.	0.4	4
84	Polycystic kidney disease, biliary dysgenesis in a patient with Larsen's syndrome. Clinical Genetics, 1997, 51, 408-411.	2.0	4
85	Prediction, prevention and personalisation of medication for the prenatal period: genetic prenatal tests for both rare and common diseases. EPMA Journal, 2011, 2, 181-195.	6.1	4
86	Segregation Analysis of Rare NRP1 and NRP2 Variants in Families with Lymphedema. Genes, 2020, 11, 1361.	2.4	4
87	Two rare PROX1 variants in patients with lymphedema. Molecular Genetics & Genomic Medicine, 2020, 8, e1424.	1.2	4
88	NOTCH1: Review of its role in lymphatic development and study of seven families with rare pathogenic variants. Molecular Genetics & Genomic Medicine, 2021, 9, e1529.	1.2	4
89	Current and Future Therapeutic Strategies for Limb Girdle Muscular Dystrophy Type R1: Clinical and Experimental Approaches. Pathophysiology, 2021, 28, 238-249.	2.2	4
90	The Effects of Long-Term Diabetes on Ghrelin Expression in Rat Stomachs. Advances in Clinical and Experimental Medicine, 2015, 24, 401-407.	1.4	4

#	Article	IF	Citations
91	Future Biotechnology. The EuroBiotech Journal, 2019, 3, 53-56.	1.0	4
92	Genetic testing for ocular coloboma. The EuroBiotech Journal, 2017, 1, 29-31.	1.0	4
93	Genetic testing for autonomic dysfunction or dysautonomias. Acta Biomedica, 2020, 91, e2020002.	0.3	4
94	Ethics committees for clinical experimentation at international level with a focus on Italy. Acta Biomedica, 2020, 91, e2020016.	0.3	4
95	A novel missense mutation outside the <scp>DNAJ</scp> domain of <scp>DNAJC21</scp> is associated with <scp>Shwachmanâ€“Diamond</scp> syndrome. British Journal of Haematology, 2022, 197, .	2.5	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.	1.7	3
99	Prenatal diagnosis of upper extremity malformations with ultrasonography: Diagnostic features and perinatal outcome. Journal of Clinical Ultrasound, 2017, 45, 267-276.	0.8	3
100	Genetic testing for central areolar choroidal dystrophy. The EuroBiotech Journal, 2017, 1, 23-25.	1.0	3
101	The effects of O⁶-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.	2.4	3
103	A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia. Clinica Chimica Acta, 2022, 529, 61-66.	1.1	3
104	Can the classical euchromatic variants of 9q12/qh+ cause recurrent abortions?. Genetic Counseling, 2008, 19, 281-6.	0.1	3
105	CDH5, a Possible New Candidate Gene for Genetic Testing of Lymphedema. Lymphatic Research and Biology, 2021, , .	1.1	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.	0.9	3
107	Germline landscape of BRCAs by 7-site collaborations as a BRCA consortium in Turkey. Breast, 2022, 65, 15-22.	2.2	3
108	Congenital alacrima in a patient with G (Opitz Frias) syndrome. Human Genetics, 1996, 97, 540-542.	3.8	2

#	Article	IF	Citations
109	Genetic testing for familial exudative vitreoretinopathy. The EuroBiotech Journal, 2017, 1, 51-53.	1.0	2
110	Genetic testing for pattern dystrophies. The EuroBiotech Journal, 2017, 1, 86-88.	1.0	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.	1.0	2
113	Genetic testing for Ebstein anomaly. The EuroBiotech Journal, 2018, 2, 55-57.	1.0	2
114	Anadoluâ€™daki TÄ±bbÄ±n DoÄŸuÅŸu, DÃ¼nyadaki Ä°lk TÄ±p Okulu Olarak: Gevher Nesibe TÄ±p Medresesi ve DarÃ¼ÅŸÅŸifasÄ±. Bilimname: DÃ¼ÅŸÃ¼nce Platformu, 0, , 79-103.	0.4	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.	1.2	1
117	A case of SRY positive 46, XX male with speaking disorder. Journal of Biotechnology, 2015, 208, S85.	3.8	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.	3.8	1
120	Genetic testing for Usher syndrome. The EuroBiotech Journal, 2017, 1, 108-110.	1.0	1
121	Genetic testing for aortic valve stenosis. The EuroBiotech Journal, 2018, 2, 61-63.	1.0	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.	1.6	1
123	Are new genome variants detected in SARS-CoV-2 expected considering population dynamics in viruses?. The EuroBiotech Journal, 2021, 5, 1-3.	1.0	1
124	COVID-19 vaccines: Where do we stand?. The EuroBiotech Journal, 2021, 5, 4-7.	1.0	1
125	Genetic testing for Bietti crystalline dystrophy. The EuroBiotech Journal, 2017, 1, 20-22.	1.0	1
126	Genetic testing for Stargardt macular dystrophy. The EuroBiotech Journal, 2017, 1, 105-107.	1.0	1

#	Article	IF	Citations
127	Genetic testing for tetralogy of Fallot. The EuroBiotech Journal, 2018, 2, 71-73.	1.0	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.	1.0	1
129	Holt-Oram syndrome in two generations with translocation t(9;15)(p12;q11.2). Annals of Saudi Medicine, 2008, 28, 209.	1.1	1
130	Holt-Oram syndrome in two generations with translocation t(9;15)(p12;q11.2). Annals of Saudi Medicine, 2008, 28, 209-212.	1.1	1
131	Genetic Disorders of Pituitary Development in Patients with Sheehan'S Syndrome. Acta Endocrinologica, 2016, 12, 413-417.	0.3	1
132	Advances in biotechnology: Genomics and genome editing. The EuroBiotech Journal, 2017, 1, 2-9.	1.0	1
133	Genetic testing for color vision deficiency. The EuroBiotech Journal, 2017, 1, 32-34.	1.0	1
134	Genetic testing for non syndromic retinitis pigmentosa. The EuroBiotech Journal, 2017, 1, 92-95.	1.0	1
135	Genetic testing for Refsum disease. The EuroBiotech Journal, 2017, 1, 89-91.	1.0	1
136	Propranolol decreases DRD3 and SLC1A2 gene expression in patients with essential tremor. Universa Medicina, 2020, 39, 105-112.	0.2	1
137	A very rare cause of arthrogryposis multiplex congenita: a novel mutation in TOR1A. Journal of Pediatric Endocrinology and Metabolism, 2022, 35, 845-850.	0.9	1
138	Diagnosing AlstrÃ¶m syndrome in a patient followed up with syndromic obesity for years. Intractable and Rare Diseases Research, 2022, 11, 84-86.	0.9	1
139	A case with Waardenburg syndrome presenting with two separate translocations ??? one reciprocal and one complex. Clinical Dysmorphology, 2001, 10, 65-66.	0.3	0
140	The increasing importance of Medical Genetics in Turkey. Current Opinion in Biotechnology, 2011, 22, S42-S43.	6.6	0
141	Prenatally detected de novo 46, XX, t(21;21)(p12;p12) at chorionic villus sampling. Current Opinion in Biotechnology, 2011, 22, S107.	6.6	0
142	Analysing the role of MDM2 SNP309 in patients with glioblastoma multiforme. Current Opinion in Biotechnology, 2013, 24, S98.	6.6	0
143	Innovations in biotechnology. Journal of Biotechnology, 2015, 208, S5.	3.8	0
144	A novel nonsense mutation in GALNS gene in family with MPS4A diagnosed child. Journal of Biotechnology, 2016, 231, S108.	3.8	0

#	Article	IF	Citations
145	A case of XYY male patient with micropenis. Journal of Biotechnology, 2016, 231, S109.	3.8	0
146	Perspectives of biotechnology. Journal of Biotechnology, 2016, 231, S4.	3.8	0
147	Editorial. Journal of Biotechnology, 2016, 231, S1-S3.	3.8	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.	1.5	0
149	Prenatal diagnosis of a foetus with partial monosomy 4p and partial trisomy 13q. Journal of Biotechnology, 2017, 256, S76.	3.8	0
150	The effect of CYP2C19 * 2 polymorphism on clopidogrel resistance in COPD patients. Journal of Biotechnology, 2017, 256, S80.	3.8	0
151	Frequency of chromosome variants in families with recurrent pregnancy loss and statistical analysis of infertility. Journal of Biotechnology, 2017, 256, S76.	3.8	0
152	Genetic testing for congenital stationary night blindness. The EuroBiotech Journal, 2017, 1, 38-40.	1.0	0
153	Genetic testing in translational ophthalmology. The EuroBiotech Journal, 2017, 1, 1-5.	1.0	0
154	Genetic testing for Norrie disease. The EuroBiotech Journal, 2017, 1, 77-79.	1.0	0
155	Editorial. Journal of Biotechnology, 2018, 280, S1-S2.	3.8	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.	0.9	0
157	Detection of mutations in CML patients resistant to tyrosine kinase inhibitor: imatinib mesylate therapy. Medical Oncology, 2021, 38, 120.	2.5	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.	1.1	0
161	Genetic testing for infantile nystagmus. The EuroBiotech Journal, 2017, 1, 57-59.	1.0	0
162	Genetic testing for retinitis punctata albescens/fundus albipunctatus. The EuroBiotech Journal, 2017, 1, 96-98.	1.0	0

#	Article	IF	Citations
163	Genetic testing for X-linked juvenile retinoschisis. The EuroBiotech Journal, 2017, 1, 111-113.	1.0	0
164	Genetic testing for Sorsbyâ€™s fundus dystrophy. The EuroBiotech Journal, 2017, 1, 102-104.	1.0	0
165	Genetic testing for Bardet-Biedl syndrome. The EuroBiotech Journal, 2017, 1, 14-16.	1.0	0
166	Genetic testing for inherited eye misalignment. The EuroBiotech Journal, 2017, 1, 60-62.	1.0	0
167	Genetic testing for corneal dystrophies and other corneal Mendelian diseases. The EuroBiotech Journal, 2017, 1, 41-44.	1.0	0
168	Genetic testing for optic atrophy. The EuroBiotech Journal, 2017, 1, 83-85.	1.0	0
169	Genetic testing for Mendelian myopia. The EuroBiotech Journal, 2017, 1, 74-76.	1.0	0
170	Genetic testing for achromatopsia. The EuroBiotech Journal, 2017, 1, 11-13.	1.0	0
171	Genetic testing for Mendelian cataract. The EuroBiotech Journal, 2017, 1, 66-69.	1.0	0
172	Genetic testing for Leber congenital amaurosis. The EuroBiotech Journal, 2017, 1, 63-65.	1.0	0
173	Genetic testing for Mendelian glaucoma. The EuroBiotech Journal, 2017, 1, 70-73.	1.0	0
174	Genetic testing for Best vitelliform macular dystrophy. The EuroBiotech Journal, 2017, 1, 17-19.	1.0	0
175	Genetic testing for enhanced S-cone syndrome. The EuroBiotech Journal, 2017, 1, 48-50.	1.0	0
176	Genetic testing for Senior-Loken syndrome. The EuroBiotech Journal, 2017, 1, 99-101.	1.0	0
177	Genetic testing for cone rod dystrophies. The EuroBiotech Journal, 2017, 1, 35-37.	1.0	0
178	Genetic testing for choroideremia. The EuroBiotech Journal, 2017, 1, 26-28.	1.0	0
179	Genetic testing for ocular albinism and oculocutaneous albinism. The EuroBiotech Journal, 2017, 1, 80-82.	1.0	0
180	Genetic testing for Doyne honeycomb retinal dystrophy. The EuroBiotech Journal, 2017, 1, 45-47.	1.0	0

#	Article	IF	Citations
181	Evaluation of aortic intima-media thickness in newborns with Down syndrome. Advances in Clinical and Experimental Medicine, 2017, 26, 1253-1256.	1.4	0
182	Prof. Mariapia Viola-Magni â€“ An Appreciation. The EuroBiotech Journal, 2018, 2, 1-1.	1.0	0
183	Genetic testing for ventricular septal defect. The EuroBiotech Journal, 2018, 2, 51-54.	1.0	0
184	Genetic testing for Marfan syndrome. The EuroBiotech Journal, 2018, 2, 35-37.	1.0	0
185	Genetic testing for vascular Ehlers-Danlos syndrome and other variants with fragility of the middle arteries. The EuroBiotech Journal, 2018, 2, 42-44.	1.0	0
186	Genetic testing for Marfan-like disorders. The EuroBiotech Journal, 2018, 2, 38-41.	1.0	0
187	Genetic testing for atrial septal defect. The EuroBiotech Journal, 2018, 2, 45-47.	1.0	0
188	THE ASSOCIATION OF BRAIN-DERIVED NEUROTROPHIC FACTOR GENE POLYMORPHISM WITH OBSTRUCTIVE SLEEP APNEA SYNDROME and OBESITY. , 2018, , .		0
189	Genetic testing for cerebral cavernous malformations. The EuroBiotech Journal, 2018, 2, 83-85.	1.0	0
190	Genetic testing for atrioventricular septal defect. The EuroBiotech Journal, 2018, 2, 48-50.	1.0	0
191	Genetic testing for bicuspid aortic valve. The EuroBiotech Journal, 2018, 2, 67-70.	1.0	0
192	Genetic testing for coarctation of aorta. The EuroBiotech Journal, 2018, 2, 64-66.	1.0	0
193	Genetic testing for pulmonary stenosis. The EuroBiotech Journal, 2018, 2, 58-60.	1.0	0
194	Genetic testing for hereditary hemorrhagic telangiectasia. The EuroBiotech Journal, 2018, 2, 32-34.	1.0	0
195	A Potential Method to Help Predict Genetic Diseases and Arrange Healthcare: Copy Number Variations Analysis. Erciyes Medical Journal, 2019, , .	0.0	0
196	Reflections on Emerging Technologies in Nanomedicine. Erciyes Medical Journal, 2020, , .	0.0	0
197	Enhancer of zeste homolog 2 (EZH2) gene inhibition via 3-Deazaneplanocin A (DZNep) in human liver cells and it is relation with fibrosis. Turkish Journal of Biochemistry, 2020, 45, 737-745.	0.5	0
198	Bacteriophages in food supplements obtained from natural sources. Acta Biomedica, 2020, 91, e2020025.	0.3	0

#	Article	IF	Citations
199	Propranolol significantly reduced DNA polymerase Î² expression in patients with essential tremor. Universa Medicina, 2021, 40, 207-215.	0.2	0
200	Evaluation of Utilizing the Distinct Genes as Predictive Biomarkers in Late-Onset Alzheimer's Disease. Global Medical Genetics, 0, , .	0.9	0
201	Congenital alacrima in a patient with G (Opitz Frias) syndrome. Human Genetics, 1996, 97, 540-542.	3.8	0
202	INVESTIGATION OF CDKL5 GENE MUTATIONS IN AUTISTIC PATIENTS ACCOMPANIED WITH INTRACTABLE SEIZURES, AUTISTIC DISORDER AND SEIZURE IN INFANCY AND EARLY CHILDHOOD. Cumhuriyet Medical Journal, 0, , .	0.1	0