Desheng Liang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6918051/publications.pdf

Version: 2024-02-01

129	2,257	304743	289244
papers	citations	22 h-index	40 g-index

134	134	134	3484
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Amyotrophic lateral sclerosis (ALS) linked mutation in Ubiquilin 2 affects stress granule assembly via TIAâ€1. CNS Neuroscience and Therapeutics, 2022, 28, 105-115.	3.9	13
2	Ectopic Expression of FVIII in HPCs and MSCs Derived from hiPSCs with Site-Specific Integration of ITGA2B Promoter-Driven BDDF8 Gene in Hemophilia A. International Journal of Molecular Sciences, 2022, 23, 623.	4.1	4
3	Site-Specific Integration of TRAIL in iPSC-Derived Mesenchymal Stem Cells for Targeted Cancer Therapy. Stem Cells Translational Medicine, 2022, 11, 297-309.	3.3	16
4	Genome sequencing demonstrates high diagnostic yield in children with undiagnosed global developmental delay/intellectual disability: A prospective study. Human Mutation, 2022, 43, 568-581.	2.5	12
5	Comprehensive Analysis of Congenital Adrenal Hyperplasia Using Long-Read Sequencing. Clinical Chemistry, 2022, 68, 927-939.	3.2	23
6	Analysis of the relationship between phenotypes and genotypes in 60 Chinese patients with propionic acidemia: a fourteen-year experience at a tertiary hospital. Orphanet Journal of Rare Diseases, 2022, 17, 135.	2.7	9
7	Identification of four novel mutations in BTK from six Chinese families with X-linked agammaglobulinemia. Clinica Chimica Acta, 2022, 531, 48-55.	1.1	3
8	Cas14a1-Mediated Nucleic Acid Diagnostics for Spinal Muscular Atrophy. Biosensors, 2022, 12, 268.	4.7	3
9	Targeted B-domain deletion restores F8 function in human endothelial cells and mice. Signal Transduction and Targeted Therapy, 2022, 7, .	17.1	4
10	Noninvasive prenatal diagnosis for pregnancies at risk for Î²â€thalassaemia: a retrospective study. BJOG: an International Journal of Obstetrics and Gynaecology, 2021, 128, 448-457.	2.3	12
11	NovelGZF1pathogenic variants identified in two Chinese patients with Larsen syndrome. Clinical Genetics, 2021, 99, 281-285.	2.0	0
12	IL-24 armored CAR19-T cells show enhanced antitumor activity and persistence. Signal Transduction and Targeted Therapy, 2021, 6, 14.	17.1	4
13	Detection of Spinal Muscular Atrophy Using a Duplexed Real-Time PCR Approach With Locked Nucleic Acid-Modified Primers. Annals of Laboratory Medicine, 2021, 41, 101-107.	2.5	5
14	Loss of PIGK function causes severe infantile encephalopathy and extensive neuronal apoptosis. Human Genetics, 2021, 140, 791-803.	3.8	6
15	Behavioral and Gene Expression Analysis of Stxbp6-Knockout Mice. Brain Sciences, 2021, 11, 436.	2.3	3
16	Targeted addition of mini-dystrophin into rDNA locus of Duchenne muscular dystrophy patient-derived iPSCs. Biochemical and Biophysical Research Communications, 2021, 545, 40-45.	2.1	5
17	Cas12a and Lateral Flow Strip-Based Test for Rapid and Ultrasensitive Detection of Spinal Muscular Atrophy. Biosensors, 2021, 11, 154.	4.7	9
18	Molecular diagnosis for 55 fetuses with skeletal dysplasias by wholeâ€exome sequencing: A retrospective cohort study. Clinical Genetics, 2021, 100, 219-226.	2.0	13

#	Article	IF	Citations
19	CRISPR/Cas12a-Based Ultrasensitive and Rapid Detection of JAK2 V617F Somatic Mutation in Myeloproliferative Neoplasms. Biosensors, 2021, 11, 247.	4.7	10
20	CRISPR/Cas9-Mediated in vivo Genetic Correction in a Mouse Model of Hemophilia A. Frontiers in Cell and Developmental Biology, 2021, 9, 672564.	3.7	10
21	Simultaneous Identification of Both MFSD8 and RDH12 Pathogenic Variants in a Chinese Family Affected With Retinitis Pigmentosa. Frontiers in Genetics, 2021, 12, 715100.	2.3	1
22	Prenatal Diagnosis in a Fetus With X-Linked Recessive Chondrodysplasia Punctata: Identification and Functional Study of a Novel Missense Mutation in ARSE. Frontiers in Genetics, 2021, 12, 722694.	2.3	2
23	Increase in diagnostic yield achieved for 174 whole-exome sequencing cases reanalyzed 1â€“2Âyears after initial analysis. Clinica Chimica Acta, 2021, 523, 163-168.	1.1	8
24	Novel variants in OSGEP leading to Galloway-Mowat syndrome by altering its subcellular localization. Clinica Chimica Acta, 2021, 523, 297-303.	1.1	3
25	An Episomal CRISPR/Cas12a System for Mediating Efficient Gene Editing. Life, 2021, 11, 1262.	2.4	4
26	Noninvasive Prenatal Testing of Methylmalonic Acidemia cblC Type Using the cSMART Assay for MMACHC Gene Mutations. Frontiers in Genetics, 2021, 12, 750719.	2.3	6
27	28 novel mutations identified from 33 Chinese patients with cilia-related kidney disorders. Clinica Chimica Acta, 2020, 501, 207-215.	1.1	4
28	A study on a cohort of 301 Chinese patients with isolated methylmalonic acidemia. Journal of Inherited Metabolic Disease, 2020, 43, 409-423.	3.6	32
29	Development and validation of a haplotypeâ€free technique for nonâ€invasive prenatal diagnosis of spinal muscular atrophy. Journal of Clinical Laboratory Analysis, 2020, 34, e23046.	2.1	7
30	The rare Alus element-mediated chimerism of multiple de novo complex rearrangement sequences in GAN result in giant axonal neuropathy. Clinica Chimica Acta, 2020, 502, 91-98.	1.1	4
31	Variable phenotypes and outcomes associated with the MMACHC c.609G>A homologous mutation: long term follow-up in a large cohort of cases. Orphanet Journal of Rare Diseases, 2020, 15, 200.	2.7	13
32	REDBot: Natural language process methods for clinical copy number variation reporting in prenatal and products of conception diagnosis. Molecular Genetics & Genomic Medicine, 2020, 8, e1488.	1.2	3
33	Analysis of 70 patients with hydrocephalus due to cobalamin C deficiency. Neurology, 2020, 95, e3129-e3137.	1.1	11
34	A novel GJB1 mutation associated with Xâ€linked Charcotâ€“Marieâ€“Tooth disease in a large Chinese family pedigree. Molecular Genetics & Genomic Medicine, 2020, 8, e1127.	1.2	4
35	Mesenchymal stem cells derived from iPSCs expressing interleukin-24 inhibit the growth of melanoma in the tumor-bearing mouse model. Cancer Cell International, 2020, 20, 33.	4.1	14
36	Six novel Mutation analysis of the androgen receptor gene in 17 Chinese patients with androgen insensitivity syndrome. Clinica Chimica Acta, 2020, 506, 180-186.	1.1	6

#	Article	IF	Citations
37	Long-read DNA sequencing fully characterized chromothripsis in a patient with Langerâ€“Giedion syndrome and Cornelia de Lange syndrome-4. Journal of Human Genetics, 2020, 65, 667-674.	2.3	24
38	Gene Therapy for Hemophilia A: Where We Stand. Current Gene Therapy, 2020, 20, 142-151.	2.0	7
39	Biochemical and genetic characteristics of 40 neonates with carnitine deficiency. Journal of Central South University (Medical Sciences), 2020, 45, 1164-1171.	0.1	0
40	Ectopic expression of factor VIII in MSCs and hepatocytes derived from rDNA targeted hESCs. Clinica Chimica Acta, 2019, 495, 656-663.	1.1	2
41	Noninvasive fetal genotyping in pregnancies at risk for PKU using a comprehensive quantitative cSMART assay for PAH gene mutations: a clinical feasibility study. BJOG: an International Journal of Obstetrics and Gynaecology, 2019, 126, 1466-1474.	2.3	11
42	ssODN-Mediated In-Frame Deletion with CRISPR/Cas9 Restores FVIII Function in Hemophilia A-Patient-Derived iPSCs and ECs. Molecular Therapy - Nucleic Acids, 2019, 17, 198-209.	5.1	23
43	Clinical utility of noninvasive prenatal screening for expanded chromosome disease syndromes. Genetics in Medicine, 2019, 21, 1998-2006.	2.4	158
44	Translocation breakpoint disrupting the host SNHG14 gene but not coding genes or snoRNAs in typical Prader-Willi syndrome. Journal of Human Genetics, 2019, 64, 647-652.	2.3	12
45	Identification of pathogenic mutations in 6 Chinese families with multiple exostoses by whole-exome sequencing and multiplex ligation-dependent probe amplification. Medicine (United States), 2019, 98, e15692.	1.0	2
46	Gene Therapy for Hemophilia and Duchenne Muscular Dystrophy in China. Human Gene Therapy, 2018, 29, 146-150.	2.7	5
47	Truncating mutations of HIBCH tend to cause severe phenotypes in cases with HIBCH deficiency: a case report and brief literature review. Journal of Human Genetics, 2018, 63, 851-855.	2.3	11
48	Identification of a novel MIP frameshift mutation associated with congenital cataract in a Chinese family by whole-exome sequencing and functional analysis. Eye, 2018, 32, 1359-1364.	2.1	6
49	Seamless Genetic Conversion of SMN2 to SMN1 via CRISPR/Cpf1 and Single-Stranded Oligodeoxynucleotides in Spinal Muscular Atrophy Patient-Specific Induced Pluripotent Stem Cells. Human Gene Therapy, 2018, 29, 1252-1263.	2.7	50
50	Restoration of SMN expression in mesenchymal stem cells derived from gene-targeted patient-specific iPSCs. Journal of Molecular Histology, 2018, 49, 27-37.	2.2	6
51	Diagnosis of Joubert Syndrome 10 in a Fetus with Suspected Dandy-Walker Variant by WES: A Novel Splicing Mutation in OFD1. BioMed Research International, 2018, 2018, 1-7.	1.9	9
52	Paired CRISPR/Cas9 Nickases Mediate Efficient Site-Specific Integration of F9 into rDNA Locus of Mouse ESCs. International Journal of Molecular Sciences, 2018, 19, 3035.	4.1	19
53	Deficiency in GnRH receptor trafficking due to a novel homozygous mutation causes idiopathic hypogonadotropic hypogonadism in three prepubertal siblings. Gene, 2018, 669, 42-46.	2.2	7
54	XRCC2 mutation causes meiotic arrest, azoospermia and infertility. Journal of Medical Genetics, 2018, 55, 628-636.	3.2	54

#	Article	IF	Citations
55	Generation of reporter hESCs by targeting EGFP at the CD144 locus to facilitate the endothelial differentiation. Development Growth and Differentiation, 2018, 60, 205-215.	1.5	2
56	Three Novel Mutations in <i>FBN1</i> and <i>TGFBR2</i> in Patients with the Syndromic Form of Thoracic Aortic Aneurysms and Dissections. International Heart Journal, 2018, 59, 1059-1068.	1.0	9
57	De Novo Mutations of CCNK Cause a Syndromic Neurodevelopmental Disorder with Distinctive Facial Dysmorphism. American Journal of Human Genetics, 2018, 103, 448-455.	6.2	17
58	Novel GATAD2B loss-of-function mutations cause intellectual disability in two unrelated cases. Journal of Human Genetics, 2017, 62, 513-516.	2.3	12
59	Characterization of chromosomal abnormalities in pregnancy losses reveals critical genes and loci for human early development. Human Mutation, 2017, 38, 669-677.	2.5	28
60	Novel variants in PAX6 gene caused congenital aniridia in two Chinese families. Eye, 2017, 31, 956-961.	2.1	14
61	Clinical outcomes for couples containing a reciprocal chromosome translocation carrier without preimplantation genetic diagnosis. International Journal of Gynecology and Obstetrics, 2017, 136, 304-308.	2.3	7
62	A lesson from a reported pathogenic variant in Peutz-Jeghers syndrome: a case report. Familial Cancer, 2017, 16, 417-422.	1.9	3
63	Functional properties of peripheral <scp>CD</scp>8⁺ÂT cells in patients with repeated implantation failure. American Journal of Reproductive Immunology, 2017, 78, e12704.	1.2	6
64	A clear bias in parental origin of de novo pathogenic CNVs related to intellectual disability, developmental delay and multiple congenital anomalies. Scientific Reports, 2017, 7, 44446.	3.3	19
65	First report on an X-linked hypohidrotic ectodermal dysplasia family with X chromosome inversion: Breakpoint mapping reveals the pathogenic mechanism and preimplantation genetics diagnosis achieves an unaffected birth. Clinica Chimica Acta, 2017, 475, 78-84.	1.1	3
66	WDR73 missense mutation causes infantile onset intellectual disability and cerebellar hypoplasia in a consanguineous family. Clinica Chimica Acta, 2017, 464, 24-29.	1.1	10
67	Notable Carrier Risks for Individuals Having Two Copies of SMN1 in Spinal Muscular Atrophy Families with 2â€copy Alleles: Estimation Based on Chinese Metaâ€analysis Data. Journal of Genetic Counseling, 2017, 26, 72-78.	1.6	5
68	Targeted exome sequencing identifies novel compound heterozygous mutations in P3H1 in a fetus with osteogenesis imperfecta type VIII. Clinica Chimica Acta, 2017, 464, 170-175.	1.1	4
69	Enhanced tumor growth inhibition by mesenchymal stem cells derived from iPSCs with targeted integration of interleukin24 into rDNA loci. Oncotarget, 2017, 8, 40791-40803.	1.8	20
70	De novo exonic deletion of KDM6A in a Chinese girl with Kabuki syndrome: A case report and brief literature review. American Journal of Medical Genetics, Part A, 2016, 170, 1613-1621.	1.2	19
71	Morphometric analysis and developmental comparison of embryos from carriers with balanced chromosomal rearrangements in preimplantation genetic diagnosis cycles. Reproduction, Fertility and Development, 2016, 28, 1953.	0.4	0
72	In situ genetic correction of F8 intron 22 inversion in hemophilia A patient-specific iPSCs. Scientific Reports, 2016, 6, 18865.	3.3	43

#	Article	IF	Citations
73	A New Next-Generation Sequencing-Based Assay for Concurrent Preimplantation Genetic Diagnosis of Charcot-Marie-Tooth Disease Type 1A and Aneuploidy Screening. Journal of Genetics and Genomics, 2016, 43, 155-159.	3.9	10
74	Novel SIL1 nonstop mutation in a Chinese consanguineous family with Marinesco-SjÃ¶gren syndrome and Dandy-Walker syndrome. Clinica Chimica Acta, 2016, 458, 1-4.	1.1	11
75	A novel MSX1 intronic mutation associated with autosomal dominant non-syndromic oligodontia in a large Chinese family pedigree. Clinica Chimica Acta, 2016, 461, 135-140.	1.1	4
76	Three novel mutations of STK11 gene in Chinese patients with Peutzâ€“Jeghers syndrome. BMC Medical Genetics, 2016, 17, 77.	2.1	8
77	Molecular cytogenetic analysis of early spontaneous abortions conceived from varying assisted reproductive technology procedures. Molecular Cytogenetics, 2016, 9, 79.	0.9	23
78	A novel de novo POGZ mutation in a patient with intellectual disability. Journal of Human Genetics, 2016, 61, 357-359.	2.3	31
79	Clinical and molecular investigation in Chinese patients with glutaric aciduria type I. Clinica Chimica Acta, 2016, 453, 75-79.	1.1	11
80	Chromosomal analysis of blastocysts from balanced chromosomal rearrangement carriers. Reproduction, 2016, 151, 455-464.	2.6	7
81	Targeting of the human F8 at the multicopy rDNA locus in Hemophilia a patient-derived iPSCs using TALENickases. Biochemical and Biophysical Research Communications, 2016, 472, 144-149.	2.1	28
82	A novel FOXL2 mutation in a Chinese family with blepharophimosis, ptosis, epicanthus inversus syndrome. Human Genome Variation, 2015, 2, 15008.	0.7	1
83	De Novo ring chromosome 11 and non-reciprocal translocation of 11p15.3-pter to 21qter in a patient with congenital heart disease. Molecular Cytogenetics, 2015, 8, 88.	0.9	4
84	Targeted next-generation sequencing identifies novel compound heterozygous mutations of DYNC2H1 in a fetus with short rib-polydactyly syndrome, type III. Clinica Chimica Acta, 2015, 447, 47-51.	1.1	20
85	Rare intracranial cholesterol deposition and a homozygous mutation of LDLR in a familial hypercholesterolemia patient. Gene, 2015, 569, 313-317.	2.2	5
86	Pregnancy-related complications and adverse pregnancy outcomes in multiple pregnancies resulting from assisted reproductive technology: a meta-analysis of cohort studies. Fertility and Sterility, 2015, 103, 1492-1508.e7.	1.0	109
87	Novel de novo nonsense mutation of the PHEX gene (p.Lys50Ter) in a Chinese patient with hypophosphatemic rickets. Gene, 2015, 565, 150-154.	2.2	4
88	Assisted reproductive technology and risk of congenital malformations: a meta-analysis based on cohort studies. Archives of Gynecology and Obstetrics, 2015, 292, 777-798.	1.7	75
89	Partial trisomy 2q33.3-q37.3 in a patient with an inverted duplicated neocentric marker chromosome. Molecular Cytogenetics, 2015, 8, 10.	0.9	4
90	DMD mutation spectrum analysis in 613 Chinese patients with dystrophinopathy. Journal of Human Genetics, 2015, 60, 435-442.	2.3	47

#	Article	IF	Citations
91	Pachygyria, seizures, hypotonia, and growth retardation in a patient with an atypical 1.33Mb inherited microduplication at 22q11.23. Gene, 2015, 569, 46-50.	2.2	9
92	Two novel NIPBL gene mutations in Chinese patients with Cornelia de Lange syndrome. Gene, 2015, 555, 476-480.	2.2	5
93	Noninvasive Prenatal Testing for Wilson Disease by Use of Circulating Single-Molecule Amplification and Resequencing Technology (cSMART). Clinical Chemistry, 2015, 61, 172-181.	3.2	85
94	Brief report adult patient presenting an interstitial (9) (q21.32q31.1) direct duplication resulting from the malsegregation of a paternal balanced insertional translocation. Birth Defects Research Part A: Clinical and Molecular Teratology, 2014, 100, 294-299.	1.6	4
95	Identification of a novel SHOX mutation in a Chinese family with isolated Madelung deformity. Journal of Genetics, 2014, 93, 809-812.	0.7	0
96	TALE nickase mediates high efficient targeted transgene integration at the human multi-copy ribosomal DNA locus. Biochemical and Biophysical Research Communications, 2014, 446, 261-266.	2.1	45
97	A novel deletion mutation in RS1 gene caused X-linked juvenile retinoschisis in a Chinese family. Eye, 2014, 28, 1364-1369.	2.1	5
98	Common genetic variants on 1p13.2 associate with risk of autism. Molecular Psychiatry, 2014, 19, 1212-1219.	7.9	85
99	Copy Number Variation Sequencing for Comprehensive Diagnosis of Chromosome Disease Syndromes. Journal of Molecular Diagnostics, 2014, 16, 519-526.	2.8	106
100	Phenotypic expansion of the interstitial 16p13.3 duplication: A case report and review of the literature. Gene, 2013, 531, 502-505.	2.2	10
101	Nonâ€invasive prenatal testing of fetal whole chromosome aneuploidy by massively parallel sequencing. Prenatal Diagnosis, 2013, 33, 409-415.	2.3	134
102	Nonviral Gene Targeting at rDNA Locus of Human Mesenchymal Stem Cells. BioMed Research International, 2013, 2013, 1-10.	1.9	10
103	An Xp21.3p11.4 duplication observed in a boy with intellectual deficiency and speech delay and his asymptomatic mother. Birth Defects Research Part A: Clinical and Molecular Teratology, 2013, 97, 467-470.	1.6	4
104	MiR-223 Regulates Human Embryonic Stem Cell Differentiation by Targeting the IGF-1R/Akt Signaling Pathway. PLoS ONE, 2013, 8, e78769.	2.5	22
105	Mutation Analysis in Chinese Patients with Cornelia de Lange Syndrome. Genetic Testing and Molecular Biomarkers, 2012, 16, 1130-1134.	0.7	4
106	Novel compound heterozygous mutation of MLYCD in a Chinese patient with malonic aciduria. Molecular Genetics and Metabolism, 2012, 105, 79-83.	1.1	13
107	Three patients with Wolfâ€Hirschhorn syndrome carrying a satellited chromosome 4p. Birth Defects Research Part A: Clinical and Molecular Teratology, 2012, 94, 549-552.	1.6	1
108	Targeting of the Human Coagulation Factor IX Gene at rDNA Locus of Human Embryonic Stem Cells. PLoS ONE, 2012, 7, e37071.	2.5	21

#	Article	IF	Citations
109	MicroRNA let-7c inhibits Bcl-xl expression and regulates ox-LDL-induced endothelial apoptosis. BMB Reports, 2012, 45, 464-469.	2.4	48
110	Mutation analysis of theSRY, NR5A1, andDHHgenes in six Chinese 46,XY women. Journal of Maternal-Fetal and Neonatal Medicine, 2011, 24, 863-866.	1.5	3
111	MicroRNAs expression in ox-LDL treated HUVECs: MiR-365 modulates apoptosis and Bcl-2 expression. Biochemical and Biophysical Research Communications, 2011, 410, 127-133.	2.1	77
112	A patient with apparently reciprocal translocation and cryptic 10p deletion. American Journal of Medical Genetics, Part A, 2011, 155, 1753-1755.	1.2	0
113	A family with partial duplication/deletion 4p due to a balanced t (4; 15) (p16.2; p11.2) translocation. American Journal of Medical Genetics, Part A, 2011, 155, 656-659.	1.2	4
114	A novel single-base deletion mutation of the RUNX2 gene in a Chinese family with cleidocranial dysplasia. Genetics and Molecular Research, 2011, 10, 3539-3544.	0.2	10
115	A novel GPR143 splicing mutation in a Chinese family with X-linked congenital nystagmus. Molecular Vision, 2011, 17, 715-22.	1.1	12
116	Parental origin of apparently balanced de novo complex chromosomal rearrangements investigated by microdissection, whole genome amplification, and microsatelliteâ€mediated haplotype analysis. Clinical Genetics, 2010, 78, 548-553.	2.0	11
117	A non-viral vector for potential DMD gene therapy study by targeting a minidystrophin-GFP fusion gene into the hrDNA locus. Acta Biochimica Et Biophysica Sinica, 2009, 41, 1053-1060.	2.0	10
118	Non-viral ex vivo transduction of human hepatocyte cells to express factor VIII using a human ribosomal DNA-targeting vector. Journal of Thrombosis and Haemostasis, 2007, 5, 347-351.	3.8	26
119	Molecular analysis of hearing loss associated with enlarged vestibular aqueduct in the mainland Chinese: a unique SLC26A4 mutation spectrum. Journal of Human Genetics, 2007, 52, 492-497.	2.3	39
120	A syndactyly type IV locus maps to 7q36. Journal of Human Genetics, 2007, 52, 561-564.	2.3	35
121	Functional analysis of three genetic disorder related PITX2 mutants. Science Bulletin, 2006, 51, 164-169.	1.7	1
122	Investigation of hrDNA targeting vector-mediated tumor-specific suicide gene therapy for hepatocellular carcinoma. Science Bulletin, 2006, 51, 2342-2350.	1.7	1
123	A father and son with mental retardation, a characteristic face, inv(12), and insertion trisomy 12p12.3-p11.2. American Journal of Medical Genetics, Part A, 2006, 140A, 238-244.	1.2	8
124	Confirmation and refinement of a genetic locus of congenital motor nystagmus in Xq26.3-q27.1 in a Chinese family. Human Genetics, 2005, 116, 128-131.	3.8	13
125	Expression of reconstructive hFVIII in the hrDNA by using hrDNA targeting vector. Science Bulletin, 2005, 50, 2187-2192.	1.7	4
126	Silica nanoparticle is a possible safe carrier for gene therapy. Science Bulletin, 2005, 50, 2323-2327.	1.7	10

#	Article	IF	Citations
127	The Combination of Suicide Gene Therapy and Radiation Enhances the Killing of Nasopharyngeal Carcinoma Xenographs. Journal of Radiation Research, 2004, 45, 281-289.	1.6	8
128	Mutation in PITX2 is associated with ring dermoid of the cornea. Journal of Medical Genetics, 2004, 41, e129-e129.	3.2	39
129	A novel fusion suicide gene yeast CDglyTK plays a role in radio-gene therapy of nasopharyngeal carcinoma. Cancer Gene Therapy, 2004, 11, 790-796.	4.6	16