

Andrew N Shelling

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/4933279/andrew-n-shelling-publications-by-year.pdf>

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

119
papers

5,830
citations

37
h-index

74
g-index

122
ext. papers

6,402
ext. citations

4.9
avg, IF

5.29
L-index

#	Paper	IF	Citations
119	Nanotechnology-Enabled COVID-19 mRNA Vaccines. <i>Encyclopedia</i> , 2021 , 1, 773-780		6
118	Dual or multiple drug loaded nanoparticles to target breast cancer stem cells.. <i>RSC Advances</i> , 2020 , 10, 19089-19105	3.7	21
117	Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. <i>American Journal of Human Genetics</i> , 2020 , 107, 837-848	11	12
116	A Predictor of Early Disease Recurrence in Patients With Breast Cancer Using a Cell-free RNA and Protein Liquid Biopsy. <i>Clinical Breast Cancer</i> , 2020 , 20, 108-116	3	5
115	Relationships of maternal body mass index and plasma biomarkers with childhood body mass index and adiposity at 6 years: The Children of SCOPE study. <i>Pediatric Obesity</i> , 2019 , 14, e12537	4.6	11
114	Human Sensory LTP Predicts Memory Performance and Is Modulated by the ValMet Polymorphism. <i>Frontiers in Human Neuroscience</i> , 2019 , 13, 22	3.3	11
113	Gene-by-Activity Interactions on Obesity Traits of 6-Year-Old New Zealand European Children: A Children of SCOPE Study. <i>Pediatric Exercise Science</i> , 2018 , 30, 69-80	2	4
112	Re: "Widespread prevalence of a CREBRF variant among Māori and Pacific children is associated with weight and height in early childhood". <i>International Journal of Obesity</i> , 2018 , 42, 1389-1391	5.5	5
111	Discordant association of the CREBRF rs373863828 A allele with increased BMI and protection from type 2 diabetes in Māori and Pacific (Polynesian) people living in Aotearoa/New Zealand. <i>Diabetologia</i> , 2018 , 61, 1603-1613	10.3	35
110	The futility of fertility research? Barriers to embryo research in New Zealand. <i>New Zealand Medical Journal</i> , 2018 , 131, 63-70	0.8	
109	Genomic medicine must reduce, not compound, health inequities: the case for hauora-enhancing genomic resources for New Zealand. <i>New Zealand Medical Journal</i> , 2018 , 131, 81-89	0.8	11
108	Analysis of association of gene variants with obesity traits in New Zealand European children at 6 years of age. <i>Molecular BioSystems</i> , 2017 , 13, 1524-1533		12
107	Development of biodegradable PLGA nanoparticles surface engineered with hyaluronic acid for targeted delivery of paclitaxel to triple negative breast cancer cells. <i>Materials Science and Engineering C</i> , 2017 , 76, 593-600	8.3	64
106	MicroRNA profiling of ovarian granulosa cell tumours reveals novel diagnostic and prognostic markers. <i>Clinical Epigenetics</i> , 2017 , 9, 72	7.7	15
105	Gene-by-environment interactions of the CLOCK, PEMT, and GHRELIN loci with average sleep duration in relation to obesity traits using a cohort of 643 New Zealand European children. <i>Sleep Medicine</i> , 2017 , 37, 19-26	4.6	7
104	Characterizing nuclear and mitochondrial DNA in spent embryo culture media: genetic contamination identified. <i>Fertility and Sterility</i> , 2017 , 107, 220-228.e5	4.8	68
103	Multimodal Assessment of Estrogen Receptor mRNA Profiles to Quantify Estrogen Pathway Activity in Breast Tumors. <i>Clinical Breast Cancer</i> , 2017 , 17, 139-153	3	2

102	Gene expression profiling of breast tumours from New Zealand patients. <i>New Zealand Medical Journal</i> , 2017 , 130, 40-56	0.8	3
101	Nuclear and mitochondrial DNA in blastocoele fluid and embryo culture medium: evidence and potential clinical use. <i>Human Reproduction</i> , 2016 , 31, 1653-61	5.7	33
100	Oocyte mitochondrial deletions and heteroplasmy in a bovine model of ageing and ovarian stimulation. <i>Molecular Human Reproduction</i> , 2016 , 22, 261-71	4.4	15
99	A novel EGR-1 dependent mechanism for YB-1 modulation of paclitaxel response in a triple negative breast cancer cell line. <i>International Journal of Cancer</i> , 2016 , 139, 1157-70	7.5	27
98	Stage I granulosa cell tumours: A management conundrum? Results of long-term follow up. <i>Gynecologic Oncology</i> , 2015 , 138, 285-91	4.9	32
97	Maternal age and ovarian stimulation independently affect oocyte mtDNA copy number and cumulus cell gene expression in bovine clones. <i>Human Reproduction</i> , 2015 , 30, 1410-20	5.7	32
96	Nanoparticle therapeutics: Technologies and methods for overcoming cancer. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2015 , 97, 140-51	5.7	58
95	New Zealand University students knowledge of fertility decline in women via natural pregnancy and assisted reproductive technologies. <i>Human Fertility</i> , 2015 , 18, 208-14	1.9	15
94	Nanoparticulate carriers: an emerging tool for breast cancer therapy. <i>Journal of Drug Targeting</i> , 2015 , 23, 97-108	5.4	10
93	NOS1AP Polymorphisms Modify QTc Interval Duration But Not Cardiac Arrest Risk in Hypertrophic Cardiomyopathy. <i>Journal of Cardiovascular Electrophysiology</i> , 2015 , 26, 1346-51	2.7	3
92	The genetics of premature ovarian failure: current perspectives. <i>International Journal of Women's Health</i> , 2015 , 7, 799-810	2.8	56
91	Assessing embryo quality by combining non-invasive markers: early time-lapse parameters reflect gene expression in associated cumulus cells. <i>Human Reproduction</i> , 2015 , 30, 1850-60	5.7	28
90	The role of SPRASA in female fertility. <i>Reproductive Sciences</i> , 2015 , 22, 452-61	3	8
89	The role of FOXL2 in the pathogenesis of adult ovarian granulosa cell tumours. <i>Gynecologic Oncology</i> , 2014 , 133, 382-7	4.9	26
88	Expression of a Mutant <i>kcnj2</i> Gene Transcript in Zebrafish 2014 , 2014, 1-14		6
87	SPACA3 gene variants in a New Zealand cohort of infertile and fertile couples. <i>Human Fertility</i> , 2014 , 17, 106-13	1.9	2
86	Single nucleotide polymorphisms in arrhythmia genes modify the risk of cardiac events and sudden death in long QT syndrome. <i>Heart Rhythm</i> , 2014 , 11, 76-82	6.7	46
85	FSH receptor gene variants are rarely associated with premature ovarian failure. <i>Reproductive BioMedicine Online</i> , 2013 , 26, 396-9	4	18

84	Adult granulosa cell tumours (GCT): clinicopathological outcomes including FOXL2 mutational status and expression. <i>Gynecologic Oncology</i> , 2013 , 131, 325-9	4.9	20
83	Comparative study of microRNA regulation on FOXL2 between adult-type and juvenile-type granulosa cell tumours in vitro. <i>Gynecologic Oncology</i> , 2013 , 129, 209-15	4.9	10
82	Community detection of long QT syndrome with a clinical registry: an alternative to ECG screening programs?. <i>Heart Rhythm</i> , 2013 , 10, 233-8	6.7	51
81	Comparison of responses of human melanoma cell lines to MEK and BRAF inhibitors. <i>Frontiers in Genetics</i> , 2013 , 4, 66	4.5	37
80	Expression of a Mutant <i>kcnj2</i> Gene Transcript in Zebrafish 2013 , 2013, 324839		5
79	Mutations in inhibin and activin genes associated with human disease. <i>Molecular and Cellular Endocrinology</i> , 2012 , 359, 113-20	4.4	11
78	Comparison of growth factor signalling pathway utilisation in cultured normal melanocytes and melanoma cell lines. <i>BMC Cancer</i> , 2012 , 12, 141	4.8	17
77	Nanocarrier systems for delivery of siRNA to ovarian cancer tissues. <i>Expert Opinion on Drug Delivery</i> , 2012 , 9, 743-54	8	6
76	In vivo testing of microRNA-mediated gene knockdown in zebrafish. <i>Journal of Biomedicine and Biotechnology</i> , 2012 , 2012, 350352		8
75	YB-1, the E2F pathway, and regulation of tumor cell growth. <i>Journal of the National Cancer Institute</i> , 2012 , 104, 133-46	9.7	79
74	Ethnic differences in disease presentation of uterine cancer in New Zealand women. <i>Journal of Family Planning and Reproductive Health Care</i> , 2012 , 38, 239-45		8
73	The transcriptional targets of mutant FOXL2 in granulosa cell tumours. <i>PLoS ONE</i> , 2012 , 7, e46270	3.7	56
72	Centrosomal dysregulation in human metastatic melanoma cell lines. <i>Cancer Genetics</i> , 2011 , 204, 477-85	2.3	11
71	Symptoms and signs associated with syncope in young people with primary cardiac arrhythmias. <i>Heart Lung and Circulation</i> , 2011 , 20, 593-8	1.8	25
70	Prospective, population-based long QT molecular autopsy study of postmortem negative sudden death in 1 to 40 year olds. <i>Heart Rhythm</i> , 2011 , 8, 412-9	6.7	121
69	Elevated serum gastrin levels in Jervell and Lange-Nielsen syndrome: a marker of severe KCNQ1 dysfunction?. <i>Heart Rhythm</i> , 2011 , 8, 551-4	6.7	22
68	Zebrafish as a model for long QT syndrome: the evidence and the means of manipulating zebrafish gene expression. <i>Acta Physiologica</i> , 2010 , 199, 257-76	5.6	35
67	Predictive and prognostic molecular markers for cancer medicine. <i>Therapeutic Advances in Medical Oncology</i> , 2010 , 2, 125-48	5.4	138

66	Inhibin and premature ovarian failure. <i>Human Reproduction Update</i> , 2010 , 16, 39-50	15.8	78
65	Array comparative genomic hybridization for the detection of submicroscopic copy number variations of the X chromosome in women with premature ovarian failure. <i>Human Reproduction</i> , 2010 , 25, 3159-60; author reply 3160-1	5.7	16
64	Premature ovarian failure. <i>Reproduction</i> , 2010 , 140, 633-41	3.8	183
63	Identification and expression analysis of <i>kcnh2</i> genes in the zebrafish. <i>Biochemical and Biophysical Research Communications</i> , 2010 , 396, 817-24	3.4	24
62	Mutational analysis of inhibin alpha gene revealed three novel variations in Indian women with premature ovarian failure. <i>Fertility and Sterility</i> , 2010 , 94, 90-8	4.8	18
61	Posthumous diagnosis of long QT syndrome from neonatal screening cards. <i>Heart Rhythm</i> , 2010 , 7, 481-6	6.7	49
60	FOXL2 copy number changes in the molecular pathogenesis of BPES: unique cohort of 17 deletions. <i>Human Mutation</i> , 2010 , 31, E1332-47	4.7	16
59	Characterization of single-nucleotide polymorphisms relevant to inflammatory bowel disease in commonly used gastrointestinal cell lines. <i>Inflammatory Bowel Diseases</i> , 2010 , 16, 282-95	4.5	11
58	Progress in the study of genetic disease: bringing new light to complex problems. <i>Postgraduate Medical Journal</i> , 2009 , 85, 505-7	2	8
57	Biophysical properties of 9 KCNQ1 mutations associated with long-QT syndrome. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2009 , 2, 417-26	6.4	33
56	Misdiagnosis of long QT syndrome as epilepsy at first presentation. <i>Annals of Emergency Medicine</i> , 2009 , 54, 26-32	2.1	80
55	Activin is a potent growth suppressor of epithelial ovarian cancer cells. <i>Cancer Letters</i> , 2009 , 285, 157-65	9.9	22
54	Investigating the association between inhibin alpha gene promoter polymorphisms and premature ovarian failure. <i>Fertility and Sterility</i> , 2009 , 91, 62-6	4.8	30
53	Interactions among genes influencing bacterial recognition increase IBD risk in a population-based New Zealand cohort. <i>Human Immunology</i> , 2009 , 70, 440-6	2.3	23
52	Absence of 566C>T mutation in exon 7 of the FSHR gene in Indian women with premature ovarian failure. <i>International Journal of Gynecology and Obstetrics</i> , 2009 , 105, 265-6	4	9
51	Sex of bovine embryos may be related to mothers' preovulatory follicular testosterone. <i>Biology of Reproduction</i> , 2008 , 78, 812-5	3.9	68
50	Single nucleotide polymorphisms in human Paneth cell defensin A5 may confer susceptibility to inflammatory bowel disease in a New Zealand Caucasian population. <i>Digestive and Liver Disease</i> , 2008 , 40, 723-30	3.3	18
49	Identification of large gene deletions and duplications in KCNQ1 and KCNH2 in patients with long QT syndrome. <i>Heart Rhythm</i> , 2008 , 5, 1275-81	6.7	70

48	Gender-stratified analysis of DLG5 R30Q in 4707 patients with Crohn disease and 4973 controls from 12 Caucasian cohorts. <i>Journal of Medical Genetics</i> , 2008 , 45, 36-42	5.8	37
47	Long QT and Brugada syndrome gene mutations in New Zealand. <i>Heart Rhythm</i> , 2007 , 4, 1306-14	6.7	35
46	Association of DLG5 variants with inflammatory bowel disease in the New Zealand Caucasian population and meta-analysis of the DLG5 R30Q variant. <i>Inflammatory Bowel Diseases</i> , 2007 , 13, 1069-76	4.5	18
45	Genome-wide association study identifies novel breast cancer susceptibility loci. <i>Nature</i> , 2007 , 447, 1087-93	5.4	1957
44	Genetic variation in human disease and a new role for copy number variants. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2007 , 622, 33-41	3.3	28
43	Genes, diet and inflammatory bowel disease. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2007 , 622, 70-83	3.3	64
42	Nutrigenomics and gut health: meeting report from an international conference in Auckland, New Zealand, April 30, May 1-3, 2006. <i>Genes and Nutrition</i> , 2007 , 2, 157-60	4.3	2
41	Has toll-like receptor 4 been prematurely dismissed as an inflammatory bowel disease gene? Association study combined with meta-analysis shows strong evidence for association. <i>American Journal of Gastroenterology</i> , 2007 , 102, 2504-12	0.7	102
40	Brugada syndrome masquerading as febrile seizures. <i>Pediatrics</i> , 2007 , 119, e1206-11	7.4	47
39	Functional analysis of the human inhibin alpha subunit variant A257T and its potential role in premature ovarian failure. <i>Human Reproduction</i> , 2007 , 22, 3241-8	5.7	40
38	Triallelic single nucleotide polymorphisms and genotyping error in genetic epidemiology studies: MDR1 (ABCB1) G2677/T/A as an example. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2007 , 16, 1185-92	4.92	45
37	Mutational analysis of betaglycan/TGF-betaRIII in premature ovarian failure. <i>Fertility and Sterility</i> , 2007 , 87, 210-2	4.8	15
36	Modeling inflammatory bowel disease: the zebrafish as a way forward. <i>Expert Review of Molecular Diagnostics</i> , 2007 , 7, 177-93	3.8	10
35	Expression of TGF-beta1, TGF-beta2, TGF-beta3 and the receptors TGF-betaRI and TGF-betaRII in placentomes of artificially inseminated and nuclear transfer derived bovine pregnancies. <i>Placenta</i> , 2006 , 27, 307-16	3.4	13
34	An investigation into FOXE1 polyalanine tract length in premature ovarian failure. <i>Molecular Human Reproduction</i> , 2006 , 12, 145-9	4.4	14
33	Mutational analysis of BMP15 and GDF9 as candidate genes for premature ovarian failure. <i>Fertility and Sterility</i> , 2006 , 86, 1009-12	4.8	65
32	Mutational screening of FOXO3A and FOXO1A in women with premature ovarian failure. <i>Fertility and Sterility</i> , 2006 , 86, 1518-21	4.8	89
31	The genetic basis of premature ovarian failure. <i>Australian and New Zealand Journal of Obstetrics and Gynaecology</i> , 2006 , 46, 242-4	1.7	78

30	Ovarian reserve tests for predicting fertility outcomes for assisted reproductive technology: the International Systematic Collaboration of Ovarian Reserve Evaluation protocol for a systematic review of ovarian reserve test accuracy. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , 2006 , 113, 1472-80	3.7	50
29	Ontogeny of AMPA and NMDA receptor gene expression in the developing sheep white matter and cerebral cortex. <i>Molecular Brain Research</i> , 2005 , 139, 242-50		11
28	Blepharophimosis and bilateral Duane syndrome associated with a FOXL2 mutation. <i>Clinical Genetics</i> , 2005 , 68, 520-3	4	13
27	INHA promoter polymorphisms are associated with premature ovarian failure. <i>Molecular Human Reproduction</i> , 2005 , 11, 779-84	4.4	46
26	The case for strategic international alliances to harness nutritional genomics for public and personal health. <i>British Journal of Nutrition</i> , 2005 , 94, 623-32	3.6	112
25	Altered placental lactogen and leptin expression in placentomes from bovine nuclear transfer pregnancies. <i>Biology of Reproduction</i> , 2004 , 71, 1862-9	3.9	22
24	A novel 30 bp deletion in the FOXL2 gene in a phenotypically normal woman with primary amenorrhoea: case report. <i>Human Reproduction</i> , 2004 , 19, 2767-70	5.7	42
23	Epigenetic regulation of inhibin alpha-subunit gene in prostate cancer cell lines. <i>Journal of Molecular Endocrinology</i> , 2004 , 32, 55-67	4.5	25
22	SPRASA, a novel sperm protein involved in immune-mediated infertility. <i>Human Reproduction</i> , 2004 , 19, 243-9	5.7	18
21	Isolation of genes differentially expressed in dominant and subordinate bovine follicles. <i>Endocrinology</i> , 2003 , 144, 3904-13	4.8	35
20	Activity, synthesis, storage, and messenger RNA of cyclooxygenase in intrauterine tissues of guinea pigs near term and during labor. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2003 , 68, 291-8	2.8	2
19	Physical and transcript map of the region between D6S264 and D6S149 on chromosome 6q27, the minimal region of allele loss in sporadic epithelial ovarian cancer. <i>Oncogene</i> , 2002 , 21, 387-99	9.2	23
18	Identification of novel mutations in FOXL2 associated with premature ovarian failure. <i>Molecular Human Reproduction</i> , 2002 , 8, 729-33	4.4	119
17	Molecular genetics of ovarian cancer. <i>Molecular Biotechnology</i> , 2001 , 19, 13-28	3	2
16	Analysis of the TGF beta functional pathway in epithelial ovarian carcinoma. <i>British Journal of Cancer</i> , 2001 , 85, 687-91	8.7	24
15	Molecular genetics of ovarian cancer : a technical overview. <i>Methods in Molecular Medicine</i> , 2001 , 39, 273-90		
14	Hierarchical mutation screening protocol for the BRCA1 gene. <i>Human Mutation</i> , 2000 , 16, 422-30	4.7	2
13	X chromosome defects and premature ovarian failure. <i>Australian and New Zealand Journal of Medicine</i> , 2000 , 30, 5-7		16

12	Evaluation of the possible protective role of adeno-associated virus type 2 infection in HPV-associated premalignant disease of the cervix. <i>Gynecologic Oncology</i> , 2000 , 78, 342-5	4.9	12
11	Evidence for susceptibility genes to familial Wilms tumour in addition to WT1, FWT1 and FWT2. <i>British Journal of Cancer</i> , 2000 , 83, 177-83	8.7	49
10	Inhibin: a candidate gene for premature ovarian failure. <i>Human Reproduction</i> , 2000 , 15, 2644-9	5.7	151
9	Role of p53 in drug resistance in ovarian cancer. <i>Lancet, The</i> , 1997 , 349, 744-5	4.0	38
8	Analysis of loss of heterozygosity and KRAS2 mutations in ovarian neoplasms: clinicopathological correlations. <i>Genes Chromosomes and Cancer</i> , 1997 , 18, 75-83	5	36
7	An integrated genetic map of Chromosome 6. <i>Mammalian Genome</i> , 1996 , 7, 157-9	3.2	7
6	Allele loss on chromosome arm 6q and fine mapping of the region at 6q27 in epithelial ovarian cancer. <i>Genes Chromosomes and Cancer</i> , 1996 , 15, 223-33	5	66
5	Localization of an epithelial-specific receptor kinase (EDDR1) to chromosome 6q16. <i>Genomics</i> , 1995 , 25, 584-7	4.3	19
4	The genetic analysis of ovarian cancer. <i>British Journal of Cancer</i> , 1995 , 72, 521-7	8.7	70
3	AF6 gene on chromosome band 6q27 maps distal to the minimal region of deletion in epithelial ovarian cancer. <i>Genes Chromosomes and Cancer</i> , 1995 , 14, 220-2	5	9
2	Fluorescence in situ hybridization analysis using cosmid probes to define chromosome 6q abnormalities in ovarian carcinoma cell lines. <i>Cancer Genetics and Cytogenetics</i> , 1994 , 77, 99-105		14
1	pH-Sensitive Nanoparticles Developed and Optimized Using Factorial Design for Oral Delivery of Gliclazide. <i>Journal of Pharmaceutical Innovation</i> , 1	1.8	1