

Erik J Björck

List of Publications by Year in descending order

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Version: 2024-02-01

39
papers

1,403
citations

361045

20
h-index

344852

36
g-index

40
all docs

40
docs citations

40
times ranked

2229
citing authors

#	ARTICLE	IF	CITATIONS
1	Recurrent Sites for New Centromere Seeding. <i>Genome Research</i> , 2004, 14, 1696-1703.	2.4	135
2	Integration of whole genome sequencing into a healthcare setting: high diagnostic rates across multiple clinical entities in 3219 rare disease patients. <i>Genome Medicine</i> , 2021, 13, 40.	3.6	116
3	Stickler syndrome caused by COL2A1 mutations: genotypeâ€“phenotype correlation in a series of 100 patients. <i>European Journal of Human Genetics</i> , 2010, 18, 872-880.	1.4	114
4	Ovarian cancer linked to lynch syndrome typically presents as early-onset, non-serous epithelial tumors. <i>Gynecologic Oncology</i> , 2011, 121, 462-465.	0.6	110
5	Periodontal Ehlers-Danlos Syndrome Is Caused by Mutations in C1R and C1S , which Encode Subcomponents C1r and C1s of Complement. <i>American Journal of Human Genetics</i> , 2016, 99, 1005-1014.	2.6	100
6	Mutated VH genes and preferential VH3-21 use define new subsets of mantle cell lymphoma. <i>Blood</i> , 2003, 101, 4047-4054.	0.6	99
7	The Genetic Basis of Pachyonychia Congenita. <i>Journal of Investigative Dermatology Symposium Proceedings</i> , 2005, 10, 21-30.	0.8	98
8	Genetic variation in ICF syndrome: Evidence for genetic heterogeneity. <i>Human Mutation</i> , 2000, 16, 509-517.	1.1	92
9	High expression of cyclin B1 predicts a favorable outcome in patients with follicular lymphoma. <i>Blood</i> , 2005, 105, 2908-2915.	0.6	54
10	Evidence for a common Spinocerebellar ataxia type 7 (SCA7) founder mutation in Scandinavia. <i>European Journal of Human Genetics</i> , 2000, 8, 918-922.	1.4	49
11	Hereditary Spastic Paraplegia 3A Associated With Axonal Neuropathy. <i>Archives of Neurology</i> , 2007, 64, 706.	4.9	42
12	Prevalence of colonic neoplasia and advanced lesions in the normal population: a prospective population-based colonoscopy study. <i>Scandinavian Journal of Gastroenterology</i> , 2012, 47, 184-190.	0.6	36
13	Mantle cell lymphomas acquire increased expression of CCL4, CCL5 and 4-1BB-L implicated in cell survival. <i>International Journal of Cancer</i> , 2006, 118, 2092-2097.	2.3	35
14	Mutation screening of fumarate hydratase by multiplex ligation-dependent probe amplification: detection of exonic deletion in a patient with leiomyomatosis and renal cell cancer. <i>Cancer Genetics and Cytogenetics</i> , 2008, 183, 83-88.	1.0	30
15	No evidence for a genetic modifier for renal cell cancer risk in HLRCC syndrome. <i>Familial Cancer</i> , 2010, 9, 245-251.	0.9	26
16	To belong or not to belong: nursing studentsâ€™ interactions with clinical learning environments â€“ an observational study. <i>BMC Medical Education</i> , 2016, 16, 197.	1.0	26
17	C1R Mutations Trigger Constitutive Complement 1 Activation in Periodontal Ehlers-Danlos Syndrome. <i>Frontiers in Immunology</i> , 2019, 10, 2537.	2.2	26
18	Dystrophia Helsinglandica: a new type of hereditary corneal recurrent erosions with late subepithelial fibrosis. <i>Acta Ophthalmologica</i> , 2009, 87, 659-665.	0.6	25

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19	Maternal isodisomy of chromosome 9 with no impact on the phenotype in a woman with two isochromosomes: i(9p) and i(9q)., 1999, 87, 49-52.		23
20	Distinction in gene expression profiles demonstrated in parathyroid adenomas by high-density oligoarray technology. <i>European Journal of Endocrinology</i> , 2005, 152, 459-470.	1.9	22
21	A new corneal disease with recurrent erosive episodes and autosomalâ€dominant inheritance. <i>Acta Ophthalmologica</i> , 2008, 86, 758-763.	0.6	22
22	Early prenatal diagnosis of the ICF syndrome. <i>Prenatal Diagnosis</i> , 2000, 20, 828-831.	1.1	20
23	Contradictions in clinical teachersâ€™ engagement in educational development: an activity theory analysis. <i>Advances in Health Sciences Education</i> , 2019, 24, 125-140.	1.7	20
24	Molecular characterization of parathyroid tumors from two patients with hereditary colorectal cancer syndromes. <i>Familial Cancer</i> , 2012, 11, 355-362.	0.9	11
25	A retrospective study of extracolonic, non-endometrial cancer in Swedish Lynch syndrome families. <i>Hereditary Cancer in Clinical Practice</i> , 2018, 16, 16.	0.6	11
26	Molecular Cytogenetic Approach to the Diagnosis of Splenic Lymphoma: A Case Report of Blastoid Mantle Cell Lymphoma. <i>Leukemia and Lymphoma</i> , 2003, 44, 1229-1234.	0.6	9
27	Identifying keys to success in clinical learning: a study of two interprofessional learning environments. <i>Journal of Interprofessional Care</i> , 2015, 29, 156-158.	0.8	9
28	Navigating without a map: how medical students interact with clinical learning environments. <i>Studies in Higher Education</i> , 2019, 44, 275-286.	2.9	5
29	Collaborative knotworking â€“ transforming clinical teaching practice through faculty development. <i>BMC Medical Education</i> , 2020, 20, 497.	1.0	5
30	Predicting Outcome in Colonoscopic High-risk Surveillance. <i>Anticancer Research</i> , 2015, 35, 4813-9.	0.5	5
31	A retrospective two centre study of Birt-Hogg-DubÃ© syndrome reveals a pathogenic founder mutation in FLCN in the Swedish population. <i>PLoS ONE</i> , 2022, 17, e0264056.	1.1	5
32	Shifting to team-based faculty development: a programme designed to facilitate change in medical education. <i>Higher Education Research and Development</i> , 2022, 41, 269-283.	1.9	4
33	Room for improvement: One third of Lynch syndrome patients presenting for genetic testing in a highly specialised centre in Stockholm already have cancer. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 18.	0.6	4
34	Genetic variation in ICF syndrome: Evidence for genetic heterogeneity. <i>Human Mutation</i> , 2000, 16, 509.	1.1	4
35	Participation in a clinical learning environment. <i>Clinical Teacher</i> , 2015, 12, 284-285.	0.4	3
36	Congenital dyserythropoietic anemia type 1: A case with novel compound heterozygous mutations in the <i>C15orf41</i> gene. <i>American Journal of Hematology</i> , 2018, 93, E213.	2.0	3

#	ARTICLE	IF	CITATIONS
37	Merged Testing for Colorectal Cancer Syndromes and Re-evaluation of Genetic Variants Improve Diagnostic Yield: results from a nationwide prospective cohort. <i>Genes Chromosomes and Cancer</i> , 2022, , .	1.5	2
38	Discordant structural chromosomal aberrations in chorionic villi and amniotic fluid leading to a formation of an isochromosome 21: a case report. <i>Molecular Cytogenetics</i> , 2021, 14, 30.	0.4	1
39	Schäpflin-Schulz-Passarge syndrome with multiple angiomas on the tongue: a new feature?. <i>International Journal of Dermatology</i> , 2021, 60, 641-642.	0.5	1