Paal S Andersen

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/313843/paal-s-andersen-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.

175
papers7,139
citations46
h-index77
g-index182
ext. papers8,271
ext. citations4.9
avg, IF5.43
L-index

#	Paper	IF	Citations
175	Temporal and Spatial Variation of the Skin-Associated Bacteria from Healthy Participants and Atopic Dermatitis Patients <i>MSphere</i> , 2022 , 7, e0091721	5	1
174	Skin Microbiome in Patients with Hand Eczema and Healthy Controls: A Three-week Prospective Study. <i>Acta Dermato-Venereologica</i> , 2021 ,	2.2	1
173	Hand eczema and temporal variation of Staphylococcus aureus clonal complexes: A prospective observational study. <i>Journal of the American Academy of Dermatology</i> , 2021 ,	4.5	1
172	Changes in Skin and Nasal Microbiome and Staphylococcal Species Following Treatment of Atopic Dermatitis with Dupilumab. <i>Microorganisms</i> , 2021 , 9,	4.9	3
171	Staphylococcal Communities on Skin Are Associated with Atopic Dermatitis and Disease Severity. <i>Microorganisms</i> , 2021 , 9,	4.9	10
170	Large-Scale Staphylococcus aureus Foodborne Disease Poisoning Outbreak among Primary School Children. <i>Microbiology Research</i> , 2021 , 12, 43-52	1	5
169	Complete genome of a methicillin-resistant Staphylococcus vitulinus from Danish ground beef meat carrying a mecA2 resistance gene and a novel ccr allotype. <i>Journal of Global Antimicrobial Resistance</i> , 2020 , 23, 221-223	3.4	
168	Cross-sectional study identifies lower risk of Staphylococcus aureus nasal colonization in Danish blood donors with hidradenitis suppurativa symptoms. <i>British Journal of Dermatology</i> , 2020 , 183, 387-3	18 9	2
167	Genomic analyses of clonal complex 45 isolates does not distinguish nasal carriage from bacteraemia. <i>Microbial Genomics</i> , 2020 , 6,	4.4	4
166	High persister cell formation by clinical strains belonging to clonal complex 30. <i>Microbiology (United Kingdom)</i> , 2020 , 166, 654-658	2.9	2
165	eHealth: Disease activity measures are related to the faecal gut microbiota in adult patients with ulcerative colitis. <i>Scandinavian Journal of Gastroenterology</i> , 2020 , 55, 1291-1300	2.4	1
164	Colonization with Staphylococcus aureus in patients with hand eczema: Prevalence and association with severity, atopic dermatitis, subtype and nasal colonization. <i>Contact Dermatitis</i> , 2020 , 83, 442-449	2.7	5
163	Alteration of Bacterial Communities in Anterior Nares and Skin Sites of Patients Undergoing Arthroplasty Surgery: Analysis by 16S rRNA and Staphylococcal-Specific Gene Sequencing. <i>Microorganisms</i> , 2020 , 8,	4.9	7
162	Effect of Co-inhabiting Coagulase Negative Staphylococci on Quorum Sensing, Host Factor Binding, and Biofilm Formation. <i>Frontiers in Microbiology</i> , 2019 , 10, 2212	5.7	15
161	Prevalence of infective endocarditis in patients with positive blood cultures: a Danish nationwide study. <i>European Heart Journal</i> , 2019 , 40, 3237-3244	9.5	16
160	Temporal variation of Staphylococcus aureus clonal complexes in atopic dermatitis: a follow-up study. <i>British Journal of Dermatology</i> , 2019 , 180, 181-186	4	12
159	Cohort description: The Danish Blood Donor Carriage Study. <i>Clinical Epidemiology</i> , 2019 , 11, 885-900	5.9	5

(2017-2019)

158	Antimicrobial Resistance and Virulence Gene Profiles of Methicillin-Resistant and -Susceptible From Food Products in Denmark. <i>Frontiers in Microbiology</i> , 2019 , 10, 2681	5.7	18
157	Association of Disease Severity With Skin Microbiome and Filaggrin Gene Mutations in Adult Atopic Dermatitis. <i>JAMA Dermatology</i> , 2018 , 154, 293-300	5.1	62
156	Transcriptomic profiling of interacting nasal staphylococci species reveals global changes in gene and non-coding RNA expression. <i>FEMS Microbiology Letters</i> , 2018 , 365,	2.9	7
155	Genomic analysis reveals different mechanisms of fusidic acid resistance in Staphylococcus aureus from Danish atopic dermatitis patients. <i>Journal of Antimicrobial Chemotherapy</i> , 2018 , 73, 856-861	5.1	14
154	Demographic fluctuation of community-acquired antibiotic-resistant Staphylococcus aureus lineages: potential role of flimsy antibiotic exposure. <i>ISME Journal</i> , 2018 , 12, 1879-1894	11.9	6
153	Confirmation of an IRAK3 polymorphism as a genetic marker predicting response to anti-TNF treatment in rheumatoid arthritis. <i>Pharmacogenomics Journal</i> , 2018 , 18, 81-86	3.5	21
152	Genetically determined high activity of IL-12 and IL-18 in ulcerative colitis and TLR5 in Crohns disease were associated with non-response to anti-TNF therapy. <i>Pharmacogenomics Journal</i> , 2018 , 18, 87-97	3.5	27
151	Associations between functional polymorphisms and response to biological treatment in Danish patients with psoriasis. <i>Pharmacogenomics Journal</i> , 2018 , 18, 494-500	3.5	27
150	Identification of a PVL-negative SCCmec-IVa sublineage of the methicillin-resistant Staphylococcus aureus CC80 lineage: understanding the clonal origin of CA-MRSA. <i>Clinical Microbiology and Infection</i> , 2018 , 24, 273-278	9.5	6
149	Dissemination and Characteristics of a Novel Plasmid-Encoded Carbapenem-Hydrolyzing Class D Lactamase, OXA-436, Found in Isolates from Four Patients at Six Different Hospitals in Denmark. <i>Antimicrobial Agents and Chemotherapy</i> , 2018 , 62,	5.9	18
148	Human Genetic Susceptibility to Native Valve Endocarditis in Patients With Bacteremia: Genome-Wide Association Study. <i>Frontiers in Microbiology</i> , 2018 , 9, 640	5.7	8
147	Genetic polymorphisms associated with psoriasis and development of psoriatic arthritis in patients with psoriasis. <i>PLoS ONE</i> , 2018 , 13, e0192010	3.7	23
146	The Skin Microbiome in Atopic Dermatitis Potential Treatment Target?. <i>Current Dermatology Reports</i> , 2018 , 7, 199-208	1.5	2
145	Human genetic variation in GLS2 is associated with development of complicated Staphylococcus aureus bacteremia. <i>PLoS Genetics</i> , 2018 , 14, e1007667	6	9
144	Staphylococcus aureus Bacteremia in Children Aged 5-18 Years-Risk Factors in the New Millennium. <i>Journal of Pediatrics</i> , 2018 , 203, 108-115.e3	3.6	7
143	Genetically determined high activities of the TNF-alpha, IL23/IL17, and NFkB pathways were associated with increased risk of ankylosing spondylitis. <i>BMC Medical Genetics</i> , 2018 , 19, 165	2.1	31
142	High Interlaboratory Reproducibility and Accuracy of Next-Generation-Sequencing-Based Bacterial Genotyping in a Ring Trial. <i>Journal of Clinical Microbiology</i> , 2017 , 55, 908-913	9.7	59
141	Staphylococcus aureus CC395 harbours a novel composite staphylococcal cassette chromosome mec element. <i>Journal of Antimicrobial Chemotherapy</i> , 2017 , 72, 1002-1005	5.1	12

140	Spread of avian pathogenic Escherichia coli ST117 O78:H4 in Nordic broiler production. <i>BMC Genomics</i> , 2017 , 18, 13	4.5	54
139	Methicillin-resistant and -susceptible Staphylococcus aureus from retail meat in Denmark. <i>International Journal of Food Microbiology</i> , 2017 , 249, 72-76	5.8	47
138	A universal primer-independent next-generation sequencing approach for investigations of norovirus outbreaks and novel variants. <i>Scientific Reports</i> , 2017 , 7, 813	4.9	14
137	Staphylococcus aureus colonization in atopic eczema and its association with filaggrin gene mutations. <i>British Journal of Dermatology</i> , 2017 , 177, 1394-1400	4	63
136	Whole-genome comparison of urinary pathogenic Escherichia coli and faecal isolates of UTI patients and healthy controls. <i>International Journal of Medical Microbiology</i> , 2017 , 307, 497-507	3.7	32
135	Genomic characterization, phylogenetic analysis, and identification of virulence factors in Aerococcus sanguinicola and Aerococcus urinae strains isolated from infection episodes. <i>Microbial Pathogenesis</i> , 2017 , 112, 327-340	3.8	9
134	Nasal and pharyngeal carriage of methicillin-resistant Staphylococcus sciuri among hospitalised patients and healthcare workers in a Serbian university hospital. <i>PLoS ONE</i> , 2017 , 12, e0185181	3.7	6
133	Genome analysis of Clostridium perfringens isolates from healthy and necrotic enteritis infected chickens and turkeys. <i>BMC Research Notes</i> , 2017 , 10, 270	2.3	13
132	A broad range quorum sensing inhibitor working through sRNA inhibition. <i>Scientific Reports</i> , 2017 , 7, 9857	4.9	45
131	The associations between socioeconomic status and risk of Staphylococcus aureus bacteremia and subsequent endocarditis - a Danish nationwide cohort study. <i>BMC Infectious Diseases</i> , 2017 , 17, 589	4	16
130	Commercial Biocides Induce Transfer of Prophage 113 from Human Strains of to Livestock CC398. <i>Frontiers in Microbiology</i> , 2017 , 8, 2418	5.7	15
129	Genomic relatedness of Staphylococcus pettenkoferi isolates of different origins. <i>Journal of Medical Microbiology</i> , 2017 , 66, 601-608	3.2	8
128	Whole-genome sequencing of bloodstream Staphylococcus aureus isolates does not distinguish bacteraemia from endocarditis. <i>Microbial Genomics</i> , 2017 , 3,	4.4	14
127	Use of WGS data for investigation of a long-term NDM-1-producing Citrobacter freundii outbreak and secondary in vivo spread of blaNDM-1 to Escherichia coli, Klebsiella pneumoniae and Klebsiella oxytoca. <i>Journal of Antimicrobial Chemotherapy</i> , 2016 , 71, 3117-3124	5.1	37
126	Vancomycin gene selection in the microbiome of urban Rattus norvegicus from hospital environment. <i>Evolution, Medicine and Public Health</i> , 2016 , 2016, 219-26	3	7
125	Methicillin-Resistant Staphylococcus aureus CC398 in Humans and Pigs in Norway: A "One Health" Perspective on Introduction and Transmission. <i>Clinical Infectious Diseases</i> , 2016 , 63, 1431-1438	11.6	66
124	Adaptation of Escherichia coli traversing from the faecal environment to the urinary tract. <i>International Journal of Medical Microbiology</i> , 2016 , 306, 595-603	3.7	11
123	Draft Genome Sequences of Two Avian Pathogenic Escherichia coli Strains of Clinical Importance, E44 and E51. <i>Genome Announcements</i> , 2016 , 4,		4

122	Cross-Talk between and Other Staphylococcal Species via the Quorum Sensing System. <i>Frontiers in Microbiology</i> , 2016 , 7, 1733	5.7	50
121	Familial Clustering of Staphylococcus aureus Bacteremia in First-Degree Relatives: A Danish Nationwide Cohort Study. <i>Annals of Internal Medicine</i> , 2016 , 165, 390-8	8	9
120	Molecular characterisation of the clonal emergence of high-level ciprofloxacin-monoresistant Haemophilus influenzae in the Region of Southern Denmark. <i>Journal of Global Antimicrobial Resistance</i> , 2016 , 5, 67-70	3.4	7
119	Copy number variation of scavenger-receptor cysteine-rich domains within DMBT1 and Crohn@ disease. <i>European Journal of Human Genetics</i> , 2016 , 24, 1294-300	5.3	6
118	Description and characterization of a penicillin-resistant Streptococcus dysgalactiae subsp. equisimilis clone isolated from blood in three epidemiologically linked patients. <i>Journal of Antimicrobial Chemotherapy</i> , 2016 , 71, 3376-3380	5.1	18
117	Evidence for Human Adaptation and Foodborne Transmission of Livestock-Associated Methicillin-Resistant Staphylococcus aureus. <i>Clinical Infectious Diseases</i> , 2016 , 63, 1349-1352	11.6	73
116	In vivo expression of antimicrobial peptides in atopic dermatitis. <i>Experimental Dermatology</i> , 2016 , 25, 3-9	4	22
115	Investigation of a possible outbreak of carbapenem-resistant Acinetobacter baumannii in Odense, Denmark using PFGE, MLST and whole-genome-based SNPs. <i>Journal of Antimicrobial Chemotherapy</i> , 2015 , 70, 1965-8	5.1	42
114	Mapping the Evolution of Hypervirulent Klebsiella pneumoniae. <i>MBio</i> , 2015 , 6, e00630	7.8	194
113	Echocardiographic evaluation of pre-diagnostic development in young relatives genetically predisposed to hypertrophic cardiomyopathy. <i>International Journal of Cardiovascular Imaging</i> , 2015 , 31, 1511-8	2.5	4
112	Surface Glycopolymers Are Crucial for In Vitro Anti-Wall Teichoic Acid IgG-Mediated Complement Activation and Opsonophagocytosis of Staphylococcus aureus. <i>Infection and Immunity</i> , 2015 , 83, 4247-5	5 3.7	25
111	Non-toxigenic tox gene-bearing Corynebacterium ulcerans in a traumatic ulcer from a human case and his asymptomatic dog. <i>Microbes and Infection</i> , 2015 , 17, 717-9	9.3	9
110	Use of whole-genome sequencing for detection of the spread of VIM-4-producing Escherichia coli between two patients in Denmark. <i>International Journal of Antimicrobial Agents</i> , 2015 , 45, 327-9	14.3	2
109	Characterisation of an IMP-7-producing ST357 Pseudomonas aeruginosa isolate detected in Denmark using whole genome sequencing. <i>International Journal of Antimicrobial Agents</i> , 2015 , 45, 200-7	1 ^{14.3}	11
108	Meticillin-resistant Staphylococcus aureus CC398 is an increasing cause of disease in people with no livestock contact in Denmark, 1999 to 2011. <i>Eurosurveillance</i> , 2015 , 20,	19.8	106
107	Staphylococcus aureus and the ecology of the nasal microbiome. <i>Science Advances</i> , 2015 , 1, e1400216	14.3	126
106	Investigation of a possible outbreak of NDM-5-producing ST16 Klebsiella pneumoniae among patients in Denmark with no history of recent travel using whole-genome sequencing. <i>Journal of Global Antimicrobial Resistance</i> , 2015 , 3, 219-221	3.4	23
105	Using Whole Genome Analysis to Examine Recombination across Diverse Sequence Types of Staphylococcus aureus. <i>PLoS ONE</i> , 2015 , 10, e0130955	3.7	27

104	Genetic Variations in Pattern Recognition Receptor Loci Are Associated with Anti-TNF Response in Patients with Rheumatoid Arthritis. <i>PLoS ONE</i> , 2015 , 10, e0139781	3.7	26
103	Polymorphisms in the Toll-Like Receptor and the IL-23/IL-17 Pathways Were Associated with Susceptibility to Inflammatory Bowel Disease in a Danish Cohort. <i>PLoS ONE</i> , 2015 , 10, e0145302	3.7	43
102	Effectiveness of anti-tumour necrosis factor-litherapy in Danish patients with inflammatory bowel diseases. <i>Danish Medical Journal</i> , 2015 , 62,	3.8	4
101	Role of urinary cathelicidin LL-37 and human Elefensin 1 in uncomplicated Escherichia coli urinary tract infections. <i>Infection and Immunity</i> , 2014 , 82, 1572-8	3.7	54
100	Genetic susceptibility and genotype-phenotype association in 588 Danish children with inflammatory bowel disease. <i>Journal of Crohnks and Colitis</i> , 2014 , 8, 678-85	1.5	19
99	Selection of unique Escherichia coli clones by random amplified polymorphic DNA (RAPD): Evaluation by whole genome sequencing. <i>Journal of Microbiological Methods</i> , 2014 , 103, 101-3	2.8	15
98	Polymorphisms in the inflammatory pathway genes TLR2, TLR4, TLR9, LY96, NFKBIA, NFKB1, TNFA, TNFRSF1A, IL6R, IL10, IL23R, PTPN22, and PPARG are associated with susceptibility of inflammatory bowel disease in a Danish cohort. <i>PLoS ONE</i> , 2014 , 9, e98815	3.7	75
97	Anti-TNF treatment response in rheumatoid arthritis patients is associated with genetic variation in the NLRP3-inflammasome. <i>PLoS ONE</i> , 2014 , 9, e100361	3.7	43
96	Associations between functional polymorphisms in the NFB signaling pathway and response to anti-TNF treatment in Danish patients with inflammatory bowel disease. <i>Pharmacogenomics Journal</i> , 2014 , 14, 526-34	3.5	98
95	Origin and evolution of European community-acquired methicillin-resistant Staphylococcus aureus. <i>MBio</i> , 2014 , 5, e01044-14	7.8	75
94	Risk factors for Staphylococcus aureus nasal colonization in Danish middle-aged and elderly twins. <i>European Journal of Clinical Microbiology and Infectious Diseases</i> , 2013 , 32, 1321-6	5.3	23
93	High-quality and -quantity DNA extraction from frozen archival blood clots for genotyping of single-nucleotide polymorphisms. <i>Genetic Testing and Molecular Biomarkers</i> , 2013 , 17, 501-3	1.6	16
92	Genome Sequence of Staphylococcus aureus Strain CA-347, a USA600 Methicillin-Resistant Isolate. <i>Genome Announcements</i> , 2013 , 1,		20
91	The epidemic of extended-spectrum-Elactamase-producing Escherichia coli ST131 is driven by a single highly pathogenic subclone, H30-Rx. <i>MBio</i> , 2013 , 4, e00377-13	7.8	288
90	Complete Genome Sequence of the Epidemic and Highly Virulent CTX-M-15-Producing H30-Rx Subclone of Escherichia coli ST131. <i>Genome Announcements</i> , 2013 , 1,		38
89	Penetrance of hypertrophic cardiomyopathy in children and adolescents: a 12-year follow-up study of clinical screening and predictive genetic testing. <i>Circulation</i> , 2013 , 127, 48-54	16.7	90
88	Human Edefensin-2 as a marker for disease severity and skin barrier properties in atopic dermatitis. <i>British Journal of Dermatology</i> , 2013 , 169, 587-93	4	34
87	Rapid differentiation between livestock-associated and livestock-independent Staphylococcus aureus CC398 clades. <i>PLoS ONE</i> , 2013 , 8, e79645	3.7	71

(2010-2013)

86	Genome analysis of Staphylococcus aureus ST291, a double locus variant of ST398, reveals a distinct genetic lineage. <i>PLoS ONE</i> , 2013 , 8, e63008	3.7	15
85	Rapid detection, differentiation and typing of methicillin-resistant Staphylococcus aureus harbouring either mecA or the new mecA homologue mecA(LGA251). <i>Clinical Microbiology and Infection</i> , 2012 , 18, 395-400	9.5	242
84	Genetic variability in beta-defensins is not associated with susceptibility to Staphylococcus aureus bacteremia. <i>PLoS ONE</i> , 2012 , 7, e32315	3.7	8
83	Screening of congenital heart disease patients using multiplex ligation-dependent probe amplification: early diagnosis of syndromic patients. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 720-5	2.5	24
82	Influence of host genetics and environment on nasal carriage of staphylococcus aureus in danish middle-aged and elderly twins. <i>Journal of Infectious Diseases</i> , 2012 , 206, 1178-84	7	32
81	Cardiac myosin binding protein-C mutations in families with hypertrophic cardiomyopathy: disease expression in relation to age, gender, and long term outcome. <i>Circulation: Cardiovascular Genetics</i> , 2012 , 5, 156-66		94
80	Staphylococcus aureus CC398: host adaptation and emergence of methicillin resistance in livestock. <i>MBio</i> , 2012 , 3,	7.8	504
79	Effects of a 17q21 chromosome gene variant, tobacco smoke and furred pets on infant wheeze. <i>Genes and Immunity</i> , 2012 , 13, 94-7	4.4	22
78	A novel Myosin essential light chain mutation causes hypertrophic cardiomyopathy with late onset and low expressivity. <i>Biochemistry Research International</i> , 2012 , 2012, 685108	2.4	22
77	Genome sequence of Staphylococcus aureus strain 11819-97, an ST80-IV European community-acquired methicillin-resistant isolate. <i>Journal of Bacteriology</i> , 2012 , 194, 1625-6	3.5	28
76	Human Edefensin 3 (DEFB103) and its influence on Staphylococcus aureus nasal carriage. <i>International Journal of Infectious Diseases</i> , 2011 , 15, e388-94	10.5	14
75	Genome wide peripheral blood leukocyte DNA methylation microarrays failed to identify associations with Inflammatory Bowel DiseasesP-176 <i>Inflammatory Bowel Diseases</i> , 2011 , 17, S65	4.5	
74	Alpha-defensin DEFA1A3 gene copy number elevation in Danish Crohn@ disease patients. <i>Digestive Diseases and Sciences</i> , 2011 , 56, 3517-24	4	19
73	The KCNE genes in hypertrophic cardiomyopathy: a candidate gene study. <i>Journal of Negative Results in BioMedicine</i> , 2011 , 10, 12		8
72	Prevalence of infective endocarditis in patients with Staphylococcus aureus bacteraemia: the value of screening with echocardiography. <i>European Journal of Echocardiography</i> , 2011 , 12, 414-20		108
71	Determination of beta-defensin genomic copy number in different populations: a comparison of three methods. <i>PLoS ONE</i> , 2011 , 6, e16768	3.7	37
70	CACNA1C (rs1006737) is associated with schizophrenia. <i>Molecular Psychiatry</i> , 2010 , 15, 119-21	15.1	147
69	Cloning and occurrence of czrC, a gene conferring cadmium and zinc resistance in methicillin-resistant Staphylococcus aureus CC398 isolates. <i>Antimicrobial Agents and Chemotherapy</i> , 2010 , 54, 3605-8	5.9	102

68	HLA-A alleles and infectious mononucleosis suggest a critical role for cytotoxic T-cell response in EBV-related Hodgkin lymphoma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 6400-5	11.5	88
67	Detecting 22q11.2 deletions by use of multiplex ligation-dependent probe amplification on DNA from neonatal dried blood spot samples. <i>Journal of Molecular Diagnostics</i> , 2010 , 12, 147-51	5.1	29
66	Fabry disease mimicking hypertrophic cardiomyopathy: genetic screening needed for establishing the diagnosis in women. <i>European Journal of Heart Failure</i> , 2010 , 12, 535-40	12.3	63
65	Associations between COX-2 polymorphisms, blood cholesterol and risk of acute coronary syndrome. <i>Atherosclerosis</i> , 2010 , 209, 155-62	3.1	23
64	Infliximab dependency is related to decreased surgical rates in adult Crohn@ disease patients. European Journal of Gastroenterology and Hepatology, 2010 , 22, 1196-203	2.2	12
63	Echocardiographic strain imaging to assess early and late consequences of sarcomere mutations in hypertrophic cardiomyopathy. <i>Circulation: Cardiovascular Genetics</i> , 2009 , 2, 314-21		111
62	Diagnostic yield, interpretation, and clinical utility of mutation screening of sarcomere encoding genes in Danish hypertrophic cardiomyopathy patients and relatives. <i>Human Mutation</i> , 2009 , 30, 363-70	, 4.7	94
61	Sharing data between LSDBs and central repositories. <i>Human Mutation</i> , 2009 , 30, 493-5	4.7	17
60	PPARgamma Pro12Ala polymorphism and risk of acute coronary syndrome in a prospective study of Danes. <i>BMC Medical Genetics</i> , 2009 , 10, 52	2.1	22
59	Infliximab dependency in children with Crohn@ disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2009 , 29, 792-9	6.1	19
58	The role of sarcomere gene mutations in patients with idiopathic dilated cardiomyopathy. <i>European Journal of Human Genetics</i> , 2009 , 17, 1241-9	5.3	71
57	Polymorphisms in inflammation genes, tobacco smoke and furred pets and wheeze in children. <i>Pediatric Allergy and Immunology</i> , 2009 , 20, 614-23	4.2	10
56	Micro-exons of the cardiac myosin binding protein C gene: flanking introns contain a disproportionately large number of hypertrophic cardiomyopathy mutations. <i>European Journal of Human Genetics</i> , 2008 , 16, 1062-9	5.3	16
55	A 3.2 Mb deletion on 18q12 in a patient with childhood autism and high-grade myopia. <i>European Journal of Human Genetics</i> , 2008 , 16, 312-9	5.3	14
54	Genetic and environmental factors as predictors of disease severity and extent at time of diagnosis in an inception cohort of inflammatory bowel disease, Copenhagen County and City 2003-2005. Journal of Crohnks and Colitis, 2008, 2, 162-9	1.5	15
53	Multiplex ligation-dependent probe amplification technique for copy number analysis on small amounts of DNA material. <i>Analytical Chemistry</i> , 2008 , 80, 9363-8	7.8	18
52	Idiopathic restrictive cardiomyopathy in children is caused by mutations in cardiac sarcomere protein genes. <i>Heart</i> , 2008 , 94, 1478-84	5.1	148
51	Single-strand conformation polymorphism analysis using capillary array electrophoresis for large-scale mutation detection. <i>Nature Protocols</i> , 2007 , 2, 1458-66	18.8	20

50	Type 1 diabetes risk analysis on dried blood spot samples from population-based newborns: design and feasibility of an unselected case-control study. <i>Paediatric and Perinatal Epidemiology</i> , 2007 , 21, 507-	- 17 7	30
49	Rapid tumour-like growth of giant filiform polyposis in a patient without a history of chronic bowel inflammation. <i>Apmis</i> , 2007 , 115, 1306-10	3.4	10
48	Prospective study of interaction between alcohol, NSAID use and polymorphisms in genes involved in the inflammatory response in relation to risk of colorectal cancer. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2007 , 624, 88-100	3.3	61
47	Whole genome amplification on DNA from filter paper blood spot samples: an evaluation of selected systems. <i>Genetic Testing and Molecular Biomarkers</i> , 2007 , 11, 65-71		30
46	Mutations in the Kv1.5 channel gene KCNA5 in cardiac arrest patients. <i>Biochemical and Biophysical Research Communications</i> , 2007 , 354, 776-82	3.4	21
45	Chronic inflammation: importance of NOD2 and NALP3 in interleukin-1beta generation. <i>Clinical and Experimental Immunology</i> , 2007 , 147, 227-35	6.2	607
44	Optimization of capillary array electrophoresis single-strand conformation polymorphism analysis for routine molecular diagnostics. <i>Electrophoresis</i> , 2006 , 27, 3816-22	3.6	19
43	Screening of 99 Danish patients with congenital heart disease for GATA4 mutations. <i>Genetic Testing and Molecular Biomarkers</i> , 2006 , 10, 277-80		16
42	NOD2/CARD15 gene polymorphisms in Crohn@ disease: a genotype-phenotype analysis in Danish and Portuguese patients and controls. <i>Digestion</i> , 2005 , 72, 156-63	3.6	33
41	One third of Danish hypertrophic cardiomyopathy patients with MYH7 mutations have mutations [corrected] in MYH7 rod region. <i>European Journal of Human Genetics</i> , 2005 , 13, 161-5	5.3	37
40	Mutations in the HERG K+-ion channel: a novel link between long QT syndrome and sudden infant death syndrome. <i>American Journal of Cardiology</i> , 2005 , 95, 433-4	3	58
39	High-throughput single-strand conformation polymorphism analysis on a microfabricated capillary array electrophoresis device. <i>Electrophoresis</i> , 2005 , 26, 1834-42	3.6	40
38	High-Throughput Mutation Screening 2005 , 71-100		
37	Potassium Must Be Considered in Congenital Long QT Syndrome. <i>Cardiology</i> , 2005 , 5, 54-58		3
36	Apo E in multiple sclerosis and optic neuritis: the apo E-epsilon4 allele is associated with progression of multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2005 , 11, 511-5	5	29
35	Disease concordance, zygosity, and NOD2/CARD15 status: follow-up of a population-based cohort of Danish twins with inflammatory bowel disease. <i>American Journal of Gastroenterology</i> , 2005 , 100, 248	6 ⁻⁹ 72	81
34	Genetic and environmental factors in monozygotic twins with Crohn@ disease and their first-degree relatives: a case report. <i>Digestion</i> , 2005 , 71, 262-5	3.6	5
33	Clinical and genetic characteristics of alpha cardiac actin gene mutations in hypertrophic cardiomyopathy. <i>Journal of Medical Genetics</i> , 2004 , 41, e10	5.8	39

32	Genetic and phenotypic characterization of mutations in myosin-binding protein C (MYBPC3) in 81 families with familial hypertrophic cardiomyopathy: total or partial haploinsufficiency. <i>European Journal of Human Genetics</i> , 2004 , 12, 673-7	5.3	57
31	Does KCNE5 play a role in long QT syndrome?. Clinica Chimica Acta, 2004, 345, 49-53	6.2	10
30	Outcome of clinical versus genetic family screening in hypertrophic cardiomyopathy with focus on cardiac beta-myosin gene mutations. <i>Cardiovascular Research</i> , 2003 , 57, 347-57	9.9	29
29	High-throughput single strand conformation polymorphism mutation detection by automated capillary array electrophoresis: validation of the method. <i>Human Mutation</i> , 2003 , 21, 116-22	4.7	48
28	Capillary electrophoresis-based single strand DNA conformation analysis in high-throughput mutation screening. <i>Human Mutation</i> , 2003 , 21, 455-65	4.7	79
27	Single-strand conformation polymorphism analysis using capillary electrophoresis. <i>Current Protocols in Human Genetics</i> , 2003 , Chapter 7, Unit 7.12	3.2	1
26	Automated mutation screening using dideoxy fingerprinting and capillary array electrophoresis. <i>Human Mutation</i> , 2001 , 18, 451-7	4.7	20
25	The Val606Met mutation in the cardiac beta-myosin heavy chain gene in patients with familial hypertrophic cardiomyopathy is associated with a high risk of sudden death at young age. <i>American Journal of Cardiology</i> , 2001 , 87, 1315-7	3	24
24	A response calculus for immobilized T cell receptor ligands. <i>Journal of Biological Chemistry</i> , 2001 , 276, 49125-32	5.4	25
23	Role of the T cell receptor ligand affinity in T cell activation by bacterial superantigens. <i>Journal of Biological Chemistry</i> , 2001 , 276, 33452-7	5.4	54
22	Myosin light chain mutations in familial hypertrophic cardiomyopathy: phenotypic presentation and frequency in Danish and South African populations. <i>Journal of Medical Genetics</i> , 2001 , 38, E43	5.8	60
21	Recent developments in high-throughput mutation screening. <i>Pharmacogenomics</i> , 2001 , 2, 387-99	2.6	37
20	Predictors of coronary in-stent restenosis: importance of angiotensin-converting enzyme gene polymorphism and treatment with angiotensin-converting enzyme inhibitors. <i>Journal of the American College of Cardiology</i> , 2001 , 38, 1434-9	15.1	37
19	Screening for Mutations and Polymorphisms in the Genes KCNH2 and KCNE2 Encoding the Cardiac HERG/MiRP1 Ion Channel: Implications for Acquired and Congenital Long Q-T Syndrome. <i>Clinical Chemistry</i> , 2001 , 47, 1390-1395	5.5	68
18	Development and application of linkage analysis in genetic diagnosis of familial hypertrophic cardiomyopathy. <i>Journal of Medical Genetics</i> , 2001 , 38, 193-8	5.8	6
17	Long QT syndrome with a high mortality rate caused by a novel G572R missense mutation in KCNH2. <i>Clinical Genetics</i> , 2000 , 57, 125-30	4	12
16	A novel missense mutation, Leu390Val, in the cardiac beta-myosin heavy chain associated with pronounced septal hypertrophy in two families with hypertrophic cardiomyopathy. <i>Scandinavian Cardiovascular Journal</i> , 2000 , 34, 558-63	2	8
15	High throughput mutation screening by automated capillary electrophoresis. <i>Combinatorial Chemistry and High Throughput Screening</i> , 2000 , 3, 393-409	1.3	21

LIST OF PUBLICATIONS

14	Adult-onset familial hypertrophic cardiomyopathy caused by a novel mutation, R694C, in the MYH7 gene. <i>Clinical Genetics</i> , 1999 , 56, 244-6	4	5
13	Recessive Romano-Ward syndrome associated with compound heterozygosity for two mutations in the KVLQT1 gene. <i>European Journal of Human Genetics</i> , 1999 , 7, 724-8	5.3	61
12	High-throughput single-strand conformation polymorphism analysis by automated capillary electrophoresis: robust multiplex analysis and pattern-based identification of allelic variants. <i>Human Mutation</i> , 1999 , 13, 318-27	4.7	58
11	A single strand conformation polymorphism/heteroduplex (SSCP/HD) method for detection of mutations in 15 exons of the KVLQT1 gene, associated with long QT syndrome. <i>Clinica Chimica Acta</i> , 1999 , 280, 113-25	6.2	32
10	Familial hypertrophic cardiomyopathy associated with a novel missense mutation affecting the ATP-binding region of the cardiac beta-myosin heavy chain. <i>Journal of Molecular and Cellular Cardiology</i> , 1999 , 31, 745-50	5.8	15
9	Novel donor splice site mutation in the KVLQT1 gene is associated with long QT syndrome. <i>Journal of Cardiovascular Electrophysiology</i> , 1998 , 9, 620-4	2.7	26
8	Mutation detection by cleavase in combination with capillary electrophoresis analysis: Application to mutations causing hypertrophic cardiomyopathy and long-QT syndrome*. <i>Molecular Diagnosis and Therapy</i> , 1998 , 3, 105-111		6
7	Sequence analysis and identification of the pyrKDbF operon from Lactococcus lactis including a novel gene, pyrK, involved in pyrimidine biosynthesis. <i>Journal of Bacteriology</i> , 1996 , 178, 5005-12	3.5	48
6	The B form of dihydroorotate dehydrogenase from Lactococcus lactis consists of two different subunits, encoded by the pyrDb and pyrK genes, and contains FMN, FAD, and [FeS] redox centers. <i>Journal of Biological Chemistry</i> , 1996 , 271, 29359-65	5.4	46
5	Two genes encoding uracil phosphoribosyltransferase are present in Bacillus subtilis. <i>Journal of Bacteriology</i> , 1995 , 177, 271-4	3.5	36
4	Uracil uptake in Escherichia coli K-12: isolation of uraA mutants and cloning of the gene. <i>Journal of Bacteriology</i> , 1995 , 177, 2008-13	3.5	45
3	Nucleotide metabolism in Lactococcus lactis: salvage pathways of exogenous pyrimidines. <i>Journal of Bacteriology</i> , 1994 , 176, 1514-6	3.5	26
2	Two different dihydroorotate dehydrogenases in Lactococcus lactis. <i>Journal of Bacteriology</i> , 1994 , 176, 3975-82	3.5	49
1	Characterization of the upp gene encoding uracil phosphoribosyltransferase of Escherichia coli K12. <i>FEBS Journal</i> , 1992 , 204, 51-6		54