Henrik K Jensen

List of Publications by Year in Descending Order

138 papers	3,525 citations	29 h-index	56 g-index
150 ext. papers	4,299 ext. citations	4.5 avg, IF	4.8 L-index

#	Paper	IF	Citations
138	Recurrent atrial arrhythmia in a randomised controlled trial comparing contact force-guided and contact force-blinded ablation for typical atrial flutter.. *Journal of Interventional Cardiac Electrophysiology*, **2022**, 1	2.4	0
137	Aortic Dissections in the Population-Based Danish National Patient Registry from 1996-2016: A Validation Study.. *Clinical Epidemiology*, **2022**, 14, 51-58	5.9	0
136	Long-term outcomes in a randomized controlled trial of multimodality imaging-guided left ventricular lead placement in cardiac resynchronization therapy.. *Europace*, **2022**,	3.9	1
135	Electrocardiographic Findings, Arrhythmias, and Left Ventricular Involvement in Familial ST-Depression Syndrome.. *Circulation: Arrhythmia and Electrophysiology*, **2022**, 101161CIRCEP121010688	6.4	0
134	Cardiac computed tomography-verified right ventricular lead position and outcomes in cardiac resynchronization therapy.. *Journal of Interventional Cardiac Electrophysiology*, **2022**, 1	2.4	
133	Diagnostic Yield of Genetic Testing in Young Patients With Atrioventricular Block of Unknown Cause.. *Journal of the American Heart Association*, **2022**, e025643	6	
132	Clinical and Functional Characterization of Ryanodine Receptor 2 Variants Implicated in Calcium-Release Deficiency Syndrome. *JAMA Cardiology*, **2021**,	16.2	3
131	Polygenic Risk Score-Enhanced Risk Stratification of Coronary Artery Disease in Patients With Stable Chest Pain. *Circulation Genomic and Precision Medicine*, **2021**, 14, e003298	5.2	2
130	Discontinuation of oral anticoagulation and risk of stroke and death after ablation for typical atrial flutter: A nation-wide Danish cohort study. *International Journal of Cardiology*, **2021**, 333, 110-116	3.2	2
129	Complications of implantable cardioverter-defibrillator treatment in arrhythmogenic right ventricular cardiomyopathy. *Europace*, **2021**,	3.9	2
128	Reply to: TFC ECG in arrhythmogenic cardiomyopathy: Inadequate mixture of criteria?. *International Journal of Cardiology*, **2021**, 323, 203	3.2	
127	Repeat pulmonary vein isolation in patients with atrial fibrillation: low ablation index is associated with increased risk of recurrent arrhythmia. *Scandinavian Cardiovascular Journal*, **2021**, 55, 29-34	2	1
126	Severity of congenital long QT syndrome disease manifestation and risk of depression, anxiety, and mortality: a nationwide study. *Europace*, **2021**,	3.9	2
125	Cardiac ryanodine receptor calcium release deficiency syndrome. *Science Translational Medicine*, **2021**, 13,	17.5	18
124	Long-term outcomes in young patients with atrioventricular block of unknown aetiology. *European Heart Journal*, **2021**, 42, 2060-2068	9.5	6
123	Genotype-phenotype correlation in arrhythmogenic right ventricular cardiomyopathy-risk of arrhythmias and heart failure. *Journal of Medical Genetics*, **2021**,	5.8	1
122	Rapid Exclusion of COVID Infection With the Artificial Intelligence Electrocardiogram. *Mayo Clinic Proceedings*, **2021**, 96, 2081-2094	6.4	2

121	Human RyR2 (Ryanodine Receptor 2) Loss-of-Function Mutations: Clinical Phenotypes and In Vitro Characterization. *Circulation: Arrhythmia and Electrophysiology*, **2021**, 14, e010013	6.4	2
120	Genetic Variant Score and Arrhythmogenic Right Ventricular Cardiomyopathy Phenotype in Plakophilin-2 Mutation Carriers. *Cardiology*, **2021**, 146, 763-771	1.6	1
119	An Exploratory Analysis of Proprotein Convertase Subtilisin/Kexin Type 9 Inhibition and Aortic Stenosis in the FOURIER Trial. *JAMA Cardiology*, **2020**, 5, 709-713	16.2	25
118	Absence of ECG Task Force Criteria does not rule out structural changes in genotype positive ARVC patients. *International Journal of Cardiology*, **2020**, 317, 152-158	3.2	2
117	The ABO Locus is Associated with Increased Fibrin Network Formation in Patients with Stable Coronary Artery Disease. *Thrombosis and Haemostasis*, **2020**, 120, 1248-1256	7	1
116	Genetic Risk of Coronary Artery Disease, Features of Atherosclerosis, and Coronary Plaque Burden. *Journal of the American Heart Association*, **2020**, 9, e014795	6	9
115	A randomized trial of contact force in atrial flutter ablation. *Europace*, **2020**, 22, 947-955	3.9	5
114	Targeted molecular genetic testing in young sudden cardiac death victims from Western Denmark. *International Journal of Legal Medicine*, **2020**, 134, 111-121	3.1	7
113	Incidence, Predictors, and Success of Ventricular Tachycardia Catheter Ablation in Arrhythmogenic Right Ventricular Cardiomyopathy (from the Nordic ARVC Registry). *American Journal of Cardiology*, **2020**, 125, 803-811	3	3
112	Pregnancies, ventricular arrhythmias, and substrate progression in women with arrhythmogenic right ventricular cardiomyopathy in the Nordic ARVC Registry. *Europace*, **2020**, 22, 1873-1879	3.9	3
111	Atrial fibrillation as a clinical characteristic of arrhythmogenic right ventricular cardiomyopathy: Experience from the Nordic ARVC Registry. *International Journal of Cardiology*, **2020**, 298, 39-43	3.2	5
110	Primary Prevention of Sudden Cardiac Death With Implantable Cardioverter-Defibrillator Therapy in Patients With Arrhythmogenic Right Ventricular Cardiomyopathy. *American Journal of Cardiology* , **2019**, 123, 1156-1162	3	5
109	Outcome after catheter ablation for left atrial flutter. *Scandinavian Cardiovascular Journal*, **2019**, 53, 133-140	2	5
108	Long-term proarrhythmic pharmacotherapy among patients with congenital long QT syndrome and risk of arrhythmia and mortality. *European Heart Journal*, **2019**, 40, 3110-3117	9.5	16
107	Appropriate use of genetics in a young patient with atrioventricular block and family history of sudden cardiac death. *HeartRhythm Case Reports*, **2019**, 5, 169-172	1	3
106	The ABO locus is associated with increased platelet aggregation in patients with stable coronary artery disease. *International Journal of Cardiology*, **2019**, 286, 152-158	3.2	4
105	Aetiologies and temporal trends of atrioventricular block in young patients: a 20-year nationwide study. *Europace*, **2019**, 21, 1710-1716	3.9	8
104	Electrically vs. imaging-guided left ventricular lead placement in cardiac resynchronization therapy: a randomized controlled trial. *Europace*, **2019**, 21, 1369-1377	3.9	15

103	Radiofrequency ablation lesions in low-, intermediate-, and normal-voltage myocardium: an in vivo study in a porcine heart model. *Europace*, **2019**, 21, 1919-1927	3.9	11
102	Lipoprotein(a), PCSK9 Inhibition, and Cardiovascular Risk. *Circulation*, **2019**, 139, 1483-1492	16.7	288
101	Left ventricular regional remodeling and lead position during cardiac resynchronization therapy. *Heart Rhythm*, **2018**, 15, 1542-1549	6.7	1
100	Implantable cardioverter-defibrillator therapy and device-related complications in young patients with inherited cardiomyopathies or channelopathies: a 17-year cohort study. *Europace*, **2018**, 20, 1849-1855	3.9	3
99	Voltage and pace-capture mapping of linear ablation lesions overestimates chronic ablation gap size. *Europace*, **2018**, 20, 2028-2035	3.9	2
98	Outcome of clinical management in relatives of sudden cardiac death victims. *International Journal of Cardiology*, **2018**, 262, 45-50	3.2	8
97	Multifocal atrial and ventricular premature contractions with an increased risk of dilated cardiomyopathy caused by a Na1.5 gain-of-function mutation (G213D). *International Journal of Cardiology*, **2018**, 257, 160-167	3.2	14
96	Heart transplantation in arrhythmogenic right ventricular cardiomyopathy - Experience from the Nordic ARVC Registry. *International Journal of Cardiology*, **2018**, 250, 201-206	3.2	14
95	Cardiac magnetic resonance characteristics in young survivors of aborted sudden cardiac death. *European Journal of Radiology*, **2018**, 105, 141-147	4.7	1
94	Recurrence after pulmonary vein isolation is associated with low contact force. *Scandinavian Cardiovascular Journal*, **2018**, 52, 28-33	2	2
93	The QTc interval and risk of cardiac events in bulimia nervosa: A long-term follow-up study. *International Journal of Eating Disorders*, **2018**, 51, 1331-1338	6.3	7
92	Coronary Plaque Burden and Adverse Plaque Characteristics Are Increased in Healthy Relatives of Patients With Early‐Onset Coronary Artery Disease. *JACC: Cardiovascular Imaging*, **2017**, 10, 1128-1135	8.4	9
91	A 45-SNP genetic risk score is increased in early-onset coronary artery disease but independent of familial disease clustering. *Atherosclerosis*, **2017**, 257, 172-178	3.1	8
90	Nationwide experience of catecholaminergic polymorphic ventricular tachycardia caused by RyR2 mutations. *Heart*, **2017**, 103, 901-909	5.1	13
89	Phenotypic characterization of late onset cardiac amyloidosis caused by the transthyretin mutation TTRA45S, p.(Ala65Ser). *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2017**, 24, 70-71	2.7	1
88	A genetic risk score predicts cardiovascular events in patients with stable coronary artery disease. *International Journal of Cardiology*, **2017**, 241, 411-416	3.2	12
87	Is the knowledge of contact force beneficial in pulmonary vein antrum isolation?. *Scandinavian Cardiovascular Journal*, **2017**, 51, 129-137	2	7
86	The SH2B3 and KCNK5 loci may be implicated in regulation of platelet count, volume, and maturity. *Thrombosis Research*, **2017**, 158, 86-92	8.2	3

85	Loss-of-activity-mutation in the cardiac chloride-bicarbonate exchanger AE3 causes short QT syndrome. *Nature Communications*, **2017**, 8, 1696	17.4	46
84	Long-Term Cardiovascular Risk in Heterozygous Familial Hypercholesterolemia Relatives Identified by Cascade Screening. *Journal of the American Heart Association*, **2017**, 6,	6	9
83	Coronary artery disease-associated genetic variants and biomarkers of inflammation. *PLoS ONE*, **2017**, 12, e0180365	3.7	14
82	Transcatheter left atrial appendage occlusion in patients with atrial fibrillation and a high bleeding risk using aspirin alone for post-implant antithrombotic therapy. *EuroIntervention*, **2017**, 12, 2075-2082	3.1	56
81	Adding the implantable cardioverter-defibrillator to cardiac resynchronization therapy is associated with improved long-term survival in ischaemic, but not in non-ischaemic cardiomyopathy. *Europace*, **2016**, 18, 413-9	3.9	13
80	Failure of ICD therapy in lethal arrhythmogenic right ventricular cardiomyopathy type 5 caused by the p.Ser358Leu mutation. *HeartRhythm Case Reports*, **2016**, 2, 217-222	1	2
79	Variable primary phenotypic manifestations in a rare familial form of Wolff-Parkinson-White syndrome and hypertrophic cardiomyopathy. *Europace*, **2016**, 18, 397	3.9	1
78	Cardiovascular risk factor control is insufficient in young patients with coronary artery disease. *Vascular Health and Risk Management*, **2016**, 12, 219-27	4.4	5
77	Multimodality imaging-guided left ventricular lead placement in cardiac resynchronization therapy: a randomized controlled trial. *European Journal of Heart Failure*, **2016**, 18, 1365-1374	12.3	70
76	The TMEM43 Newfoundland mutation p.S358L causing ARVC-5 was imported from Europe and increases the stiffness of the cell nucleus. *European Heart Journal*, **2015**, 36, 872-81	9.5	40
75	Radiofrequency ablation of accessory pathways in patients with the Wolff-Parkinson-White syndrome: the long-term mortality and risk of atrial fibrillation. *Europace*, **2015**, 17, 117-22	3.9	25
74	Management of patients with Arrhythmogenic Right Ventricular Cardiomyopathy in the Nordic countries. *Scandinavian Cardiovascular Journal*, **2015**, 49, 299-307	2	14
73	Risk of arrhythmia induced by psychotropic medications: a proposal for clinical management. *European Heart Journal*, **2014**, 35, 1306-15	9.5	73
72	His or para-His pacing preserves left ventricular function in atrioventricular block: a double-blind, randomized, crossover study. *Europace*, **2014**, 16, 1189-96	3.9	101
71	The diagnostic performance of imaging methods in ARVC using the 2010 Task Force criteria. *European Heart Journal Cardiovascular Imaging*, **2014**, 15, 1219-25	4.1	56
70	Mutations in Danish patients with long QT syndrome and the identification of a large founder family with p.F29L in KCNH2. *BMC Medical Genetics*, **2014**, 15, 31	2.1	11
69	Truncating plakophilin-2 mutations in arrhythmogenic cardiomyopathy are associated with protein haploinsufficiency in both myocardium and epidermis. *Circulation: Cardiovascular Genetics*, **2014**, 7, 230-40		26
68	Cardiac magnetic resonance and electroanatomical mapping of acute and chronic atrial ablation injury: a histological validation study. *European Heart Journal*, **2014**, 35, 1486-95	9.5	89

67	Ablation for idiopathic ventricular arrhythmia - with and without arrhythmia-induced cardiomyopathy. *Scandinavian Cardiovascular Journal*, **2014**, 48, 130-7	2	2
66	Catheter ablation for ventricular tachycardia in ischaemic heart disease; acute success and long-term outcome. *Scandinavian Cardiovascular Journal*, **2014**, 48, 27-34	2	2
65	Mutated desmoglein-2 proteins are incorporated into desmosomes and exhibit dominant-negative effects in arrhythmogenic right ventricular cardiomyopathy. *Human Mutation*, **2013**, 34, 697-705	4.7	27
64	Protein expression studies of desmoplakin mutations in cardiomyopathy patients reveal different molecular disease mechanisms. *Clinical Genetics*, **2013**, 84, 20-30	4	24
63	Postmortem genetic testing of the ryanodine receptor 2 (RYR2) gene in a cohort of sudden unexplained death cases. *International Journal of Legal Medicine*, **2013**, 127, 139-44	3.1	22
62	Cascade screening in families with inherited cardiac diseases driven by cardiologists: feasibility and nationwide outcome in long QT syndrome. *Cardiology*, **2013**, 126, 131-7	1.6	16
61	Long QT interval in Turner syndrome--a high prevalence of LQTS gene mutations. *PLoS ONE*, **2013**, 8, e69614	3.7	23
60	Sudden cardiac death in young adults: environmental risk factors and genetic aspects of premature atherosclerosis. *Journal of Forensic Sciences*, **2012**, 57, 658-62	1.8	11
59	The prevalence of mutations in KCNQ1, KCNH2, and SCN5A in an unselected national cohort of young sudden unexplained death cases. *Journal of Cardiovascular Electrophysiology*, **2012**, 23, 1092-8	2.7	57
58	Amiodarone significantly decreases atrial fibrillation in patients undergoing surgery for lung cancer. *Annals of Thoracic Surgery*, **2012**, 94, 339-44; discussion 345-6	2.7	50
57	Sodium current and potassium transient outward current genes in Brugada syndrome: screening and bioinformatics. *Canadian Journal of Cardiology*, **2012**, 28, 196-200	3.8	19
56	Molecular autopsy in young sudden cardiac death victims with suspected cardiomyopathy. *Forensic Science International*, **2012**, 219, 33-8	2.6	24
55	Mutations in the potassium channel subunit KCNE1 are associated with early-onset familial atrial fibrillation. *BMC Medical Genetics*, **2012**, 13, 24	2.1	67
54	Low disease prevalence and inappropriate implantable cardioverter defibrillator shock rate in Brugada syndrome: a nationwide study. *Europace*, **2012**, 14, 1025-9	3.9	23
53	Catheter ablation of arrhythmias in ebstein's anomaly: a multicenter study. *Journal of Cardiovascular Electrophysiology*, **2011**, 22, 1391-6	2.7	65
52	His and para-His pacing in AV block: feasibility and electrocardiographic findings. *Journal of Interventional Cardiac Electrophysiology*, **2011**, 31, 255-62	2.4	43
51	Adverse effect of right ventricular pacing prevented by biventricular pacing during long-term follow-up: a randomized comparison. *European Journal of Echocardiography*, **2011**, 12, 767-72		24
50	Prophylactic cavotricuspid isthmus block during atrial fibrillation ablation in patients without atrial flutter: a randomised controlled trial. *Heart*, **2009**, 95, 994-9	5.1	17

49	Identifying drug-induced repolarization abnormalities from distinct ECG patterns in congenital long QT syndrome: a study of sotalol effects on T-wave morphology. *Drug Safety*, **2009**, 32, 599-611	5.1	43
48	T(peak)T(end) interval in long QT syndrome. *Journal of Electrocardiology*, **2008**, 41, 603-8	1.4	48
47	Biventricular pacing preserves left ventricular performance in patients with high-grade atrio-ventricular block: a randomized comparison with DDD(R) pacing in 50 consecutive patients. *Europace*, **2008**, 10, 314-20	3.9	50
46	DDD(R)-pacing, but not AAI(R)-pacing induces left ventricular desynchronization in patients with sick sinus syndrome: tissue-Doppler and 3D echocardiographic evaluation in a randomized controlled comparison. *Europace*, **2008**, 10, 127-33	3.9	27
45	Superior transseptal approach to mitral valve is associated with a higher need for pacemaker implantation than the left atrial approach. *Annals of Thoracic Surgery*, **2007**, 83, 77-82	2.7	31
44	Prevention of atrial flutter with cryoablation may be proarrhythmogenic. *Annals of Thoracic Surgery* , **2007**, 83, 1717-23	2.7	10
43	A robust method for quantification of IKr-related T-wave morphology abnormalities **2007**,		3
42	Atrial incision affects the incidence of atrial tachycardia after mitral valve surgery. *Annals of Thoracic Surgery*, **2006**, 81, 509-13	2.7	25
41	The superior transseptal surgical approach to mitral valve creates slow conduction. *PACE - Pacing and Clinical Electrophysiology*, **2006**, 29, 719-26	1.6	17
40	The relationship of molecular genetic to clinical diagnosis of familial hypercholesterolemia in a Danish population. *Atherosclerosis*, **2005**, 180, 155-60	3.1	134
39	Ablation of atrial tachycardia after surgery for congenital and acquired heart disease using an electroanatomic mapping system: Which circuits to expect in which substrate?. *Heart Rhythm*, **2005**, 2, 64-72	6.7	87
38	Left ventricular lead performance in cardiac resynchronization therapy: impact of lead localization and complications. *PACE - Pacing and Clinical Electrophysiology*, **2005**, 28, 483-8	1.6	69
37	Radiofrequency ablation of atrial fibrillation: effectiveness and safety in 102 consecutive patients. *Heart*, **2005**, 91, 1611-2	5.1	1
36	The platelet polymorphism PlA2 is a genetic risk factor for myocardial infarction. *Journal of Internal Medicine*, **2004**, 255, 637-44	10.8	40
35	No genetic linkage or molecular evidence for involvement of the PCSK9, ARH or CYP7A1 genes in the Familial Hypercholesterolemia phenotype in a sample of Danish families without pathogenic mutations in the LDL receptor and apoB genes. *Atherosclerosis*, **2004**, 177, 415-22	3.1	29
34	Regional myocardial perfusion during chronic biventricular pacing and after acute change of the pacing mode in patients with congestive heart failure and bundle branch block treated with an atrioventricular sequential biventricular pacemaker. *European Journal of Heart Failure*, **2003**, 5, 179-86	12.3	23
33	LDL receptor mutation genotype and vascular disease phenotype in heterozygous familial hypercholesterolaemia. *Clinical Genetics*, **2002**, 61, 408-15	4	8
32	Tissue Doppler imaging predicts improved systolic performance and reversed left ventricular remodeling during long-term cardiac resynchronization therapy. *Journal of the American College of Cardiology*, **2002**, 40, 723-30	15.1	499

31	LDL-receptor gene mutations and the hypocholesterolemic response to statin therapy. *Clinical Genetics*, **2001**, 59, 397-405	4	12
30	Three dimensional echocardiography documents haemodynamic improvement by biventricular pacing in patients with severe heart failure. *British Heart Journal*, **2001**, 85, 514-20		45
29	Incomplete thromboxane inhibition with 100 mg of intravenous acetylsalicylic acid in patients with acute ST elevation myocardial infarction: a placebo-controlled pilot trial. *Thrombosis Research*, **2001** , 104, 175-80	8.2	6
28	Impact of acute biventricular pacing on left ventricular performance and volumes in patients with severe heart failure. A tissue doppler and three-dimensional echocardiographic study. *Cardiology*, **2001**, 95, 173-82	1.6	67
27	Flow cytometric assessment of effects of fluvastatin on low-density lipoprotein receptor activity in stimulated T-lymphocytes from patients with heterozygous familial hypercholesterolemia. *Journal of Clinical Pharmacology*, **2000**, 40, 421-9	2.9	4
26	Association of coronary heart disease with age-adjusted aortocoronary calcification in patients with familial hypercholesterolaemia. *Journal of Internal Medicine*, **2000**, 247, 479-84	10.8	12
25	Functional characterization of two low density lipoprotein receptor gene mutations by fluorescence flow cytometric assessment of receptor activity in stimulated human T-lymphocytes. *Clinical Genetics*, **2000**, 57, 110-5	4	5
24	Flow Cytometric Assessment of LDL Ligand Function for Detection of Heterozygous Familial Defective Apolipoprotein B-100. *Clinical Chemistry*, **2000**, 46, 224-233	5.5	3
23	LDL receptor mutations and ApoB mutations are not risk factors for ischemic cerebrovascular disease of the young, but lipids and lipoproteins are. *European Journal of Neurology*, **1999**, 6, 691-6	6	10
22	Linking genotype to aorto-coronary atherosclerosis: a model using familial hypercholesterolemia and aorto-coronary calcification. *Annals of Human Genetics*, **1999**, 63, 511-520	2.2	3
21	Flow cytometric assessment of LDL receptor activity in peripheral blood mononuclear cells compared to gene mutation detection in diagnosis of heterozygous familial hypercholesterolemia. *Cytometry*, **1999**, 36, 52-9		10
20	P1A1/A2 polymorphism of platelet glycoprotein IIIa and risk of acute coronary syndromes in heterozygous familial hypercholesterolemia. *Atherosclerosis*, **1999**, 143, 99-104	3.1	12
19	Spectrum of LDL receptor gene mutations in Denmark: implications for molecular diagnostic strategy in heterozygous familial hypercholesterolemia. *Atherosclerosis*, **1999**, 146, 337-44	3.1	35
18	Intronic mutations at splice junctions in the low-density lipoprotein receptor gene. *Molecular and Cellular Probes*, **1999**, 13, 257-60	3.3	2
17	Identification of recurrent and novel mutations in the LDL receptor gene in Spanish patients with familial hypercholesterolemia. *Human Mutation*, **1998**, 11, 413-413	4.7	25
16	Flow cytometry with a monoclonal antibody to the low density lipoprotein receptor compared with gene mutation detection in diagnosis of heterozygous familial hypercholesterolemia. *Clinical Chemistry*, **1998**, 44, 966-972	5.5	11
15	Identification of recurrent and novel mutations in the LDL receptor gene in Spanish patients with familial hypercholesterolemia **1998**, 11, 413		1
14	A common W556S mutation in the LDL receptor gene of Danish patients with familial hypercholesterolemia encodes a transport-defective protein. *Atherosclerosis*, **1997**, 131, 67-72	3.1	28

13	A sequence variation: 713-8delC in the transforming growth factor-beta 1 gene has higher prevalence in osteoporotic women than in normal women and is associated with very low bone mass in osteoporotic women and increased bone turnover in both osteoporotic and normal women. *Bone*, **1997**, 20, 289-94	4.7	194
12	Phenotypic variation in patients heterozygous for familial defective apolipoprotein B (FDB) in three European countries. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **1997**, 17, 741-7	9.4	40
11	Two mutations in the same low-density lipoprotein receptor allele act in synergy to reduce receptor function in heterozygous familial hypercholesterolemia. *Human Mutation*, **1997**, 9, 437-44	4.7	22
10	PatientsSattitudes toward detection of heterozygous familial hypercholesterolemia. *Archives of Internal Medicine*, **1997**, 157, 553-560		9
9	Two mutations in the same low-density lipoprotein receptor allele act in synergy to reduce receptor function in heterozygous familial hypercholesterolemia **1997**, 9, 437		1
8	Identification of a novel mutation in exon 13 of the LDL receptor gene causing familial hypercholesterolemia in two Spanish families. *Biochimica Et Biophysica Acta - Molecular Basis of Disease*, **1996**, 1316, 1-4	6.9	7
7	The Trp23-Stop and Trp66-Gly mutations in the LDL receptor gene: common causes of familial hypercholesterolemia in Denmark. *Atherosclerosis*, **1996**, 120, 57-65	3.1	25
6	Allele-specific measurement of low-density lipoprotein receptor transcript levels. *Human Mutation*, **1996**, 8, 126-33	4.7	10
5	A frequent HhaI polymorphism in intron 9 of the low density lipoprotein receptor gene detected by the denaturing gradient gel electrophoresis technique. *Clinical Genetics*, **1995**, 48, 221-2	4	3
4	Characterization of a disease-causing Glu119-Lys mutation in the low-density lipoprotein receptor gene in two Danish families with heterozygous familial hypercholesterolemia. *Human Mutation*, **1994**, 4, 102-13	4.7	24
3	Incidence of the apolipoprotein B-3500 mutation in Denmark. *Clinica Chimica Acta*, **1994**, 230, 101-4	6.2	18
2	Comparison of the effects of xamoterol, atenolol and propranolol on breathlessness, fatigue and plasma electrolytes during exercise in healthy volunteers. *European Journal of Clinical Pharmacology*, **1991**, 41, 51-5	2.8	3
1	Pharmacokinetics and dynamic effects of diltiazem in the isolated guinea-pig heart. *Basic and Clinical Pharmacology and Toxicology*, **1988**, 62, 166-71		3