

Henrik Nissen

List of Publications by Year in descending order

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27
papers

2,771
citations

706676

14
h-index

721071

23
g-index

27
all docs

27
docs citations

27
times ranked

3177
citing authors

#	ARTICLE	IF	CITATIONS
1	Safety and Efficacy of Myval Implantation in Patients with Severe Bicuspid Aortic Valve Stenosis—A Multicenter Real-World Experience. <i>Journal of Clinical Medicine</i> , 2022, 11, 443.	1.0	14
2	Eight-year outcomes for patients with aortic valve stenosis at low surgical risk randomized to transcatheter vs. surgical aortic valve replacement. <i>European Heart Journal</i> , 2021, 42, 2912-2919.	1.0	159
3	Early and late risk of ischemic stroke after TAVR as compared to a nationwide background population. <i>Clinical Research in Cardiology</i> , 2020, 109, 791-801.	1.5	13
4	Five-Year Clinical and Echocardiographic Outcomes From the NOTION Randomized Clinical Trial in Patients at Lower Surgical Risk. <i>Circulation</i> , 2019, 139, 2714-2723.	1.6	229
5	Durability of Transcatheter and Surgical Bioprosthetic Aortic Valves in Patients at Lower Surgical Risk. <i>Journal of the American College of Cardiology</i> , 2019, 73, 546-553.	1.2	252
6	Short- and Long-Term Mortality and Stroke Risk After Transcatheter Aortic Valve Implantation. <i>American Journal of Cardiology</i> , 2018, 121, 78-85.	0.7	15
7	Mortality prediction after transcatheter treatment of failed bioprosthetic aortic valves utilizing various international scoring systems: Insights from the Valve-in-Valve International Data (VIVID). <i>Catheterization and Cardiovascular Interventions</i> , 2018, 92, 1163-1170.	0.7	8
8	Effect of permanent pacemaker on mortality after transcatheter aortic valve replacement. <i>Scandinavian Cardiovascular Journal</i> , 2017, 51, 40-46.	0.4	15
9	Novel ELN mutation in a family with supra-aortic stenosis and intracranial aneurysm. <i>European Journal of Medical Genetics</i> , 2017, 60, 110-113.	0.7	21
10	Two-Year Outcomes in Patients With Severe Aortic Valve Stenosis Randomized to Transcatheter Versus Surgical Aortic Valve Replacement. <i>Circulation: Cardiovascular Interventions</i> , 2016, 9, .	1.4	155
11	No clinical effect of prosthesis-patient mismatch after transcatheter versus surgical aortic valve replacement in intermediate- and low-risk patients with severe aortic valve stenosis at mid-term follow-up: an analysis from the NOTION trial. <i>European Journal of Cardio-thoracic Surgery</i> , 2016, 50, 721-728.	0.6	47
12	Transcatheter Versus Surgical Aortic Valve Replacement in Patients With Severe Aortic Valve Stenosis. <i>Journal of the American College of Cardiology</i> , 2015, 65, 2184-2194.	1.2	779
13	Transcatheter Aortic Valve Implantation in Failed Bioprosthetic Surgical Valves. <i>JAMA - Journal of the American Medical Association</i> , 2014, 312, 162.	3.8	762
14	Cancer risk among patients with congenital heart defects: a nationwide follow-up study. <i>Cardiology in the Young</i> , 2014, 24, 40-46.	0.4	23
15	The Nordic Aortic Valve Intervention (NOTION) trial comparing transcatheter versus surgical valve implantation: study protocol for a randomised controlled trial. <i>Trials</i> , 2013, 14, 11.	0.7	41
16	Familial hypercholesterolaemia in Finland: common, rare and mild mutations of the LDL receptor and their clinical consequences. <i>Annals of Medicine</i> , 2001, 33, 410-421.	1.5	45
17	Evaluation of a clinically applicable mutation screening technique for genetic diagnosis of familial hypercholesterolemia and familial defective apolipoprotein B. <i>Clinical Genetics</i> , 1998, 53, 433-439.	1.0	8
18	Mutation screening of the LDLR gene and ApoB gene in patients with a phenotype of familial hypercholesterolemia and normal values in a functional LDL receptor/apolipoprotein B assay. <i>Clinical Genetics</i> , 1998, 54, 79-82.	1.0	9

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19	Effects of amlodipine and isosorbide dinitrate on exercise-induced and ambulatory ischemia in patients with chronic stable angina pectoris. <i>Cardiovascular Drugs and Therapy</i> , 1997, 11, 629-635.	1.3	6
20	Common founder mutation in the LDL receptor gene causing familial hypercholesterolaemia in the Icelandic population. , 1997, 10, 36-44.		35
21	Diagnostic strategy, genetic diagnosis and identification of new mutations in intermittent porphyria by denaturing gradient gel electrophoresis. , 1997, 9, 122-130.		29
22	Molecular genetics of familial hypercholesterolemia in Israel. <i>Human Genetics</i> , 1996, 98, 581-586.	1.8	36
23	Unusual xanthomas in a young patient with heterozygous familial hypercholesterolemia and type III hyperlipoproteinemia. , 1996, 65, 149-154.		9
24	Clinically applicable mutation screening in familial hypercholesterolemia. , 1996, 8, 168-177.		37
25	Clinically applicable mutation screening in familial hypercholesterolemia. <i>Human Mutation</i> , 1996, 8, 168-177.	1.1	4
26	A frequent HhaI polymorphism in intron 9 of the low density lipoprotein receptor gene detected by the denaturing gradient gel electrophoresis technique. <i>Clinical Genetics</i> , 1995, 48, 221-222.	1.0	4
27	Assessment of the NMR visibility of intraerythrocytic sodium by flame photometric and ion-competitive studies. <i>Magnetic Resonance in Medicine</i> , 1989, 10, 388-398.	1.9	16