Leen M 't Hart

List of Publications by Year
in descending order

114 papers	5,487 citations	94433 37 h-index	95266 68 g-index
121 all docs	121 docs citations	121 times ranked	9327 citing authors

#	Article	IF	Citations
1	Response to Comment on Dawed et al. Genome-Wide Meta-analysis Identifies Genetic Variants Associated With Glycemic Response to Sulfonylureas. Diabetes Care 2021;44:2673â€“2682. Diabetes Care, 2022, 45, e82-e83.	8.6	0
2	Diabetes risk loci-associated pathways are shared across metabolic tissues. BMC Genomics, 2022, 23, 368.	2.8	0
3	Clinical profiles of postâ€load glucose subgroups and their association with glycaemic traits over time: An IMIâ€DIRECT study. Diabetic Medicine, 2021, 38, e14428.	2.3	2
4	Discovery of predictors of sudden cardiac arrest in diabetes: rationale and outline of the RESCUED (REcognition of Sudden Cardiac arrest vUlnErability in Diabetes) project. Open Heart, 2021, 8, e001554.	2.3	5
5	Performance of Risk Assessment Models for Prevalent or Undiagnosed Type 2 Diabetes Mellitus in a Multi-Ethnic Populationâ€”The Helius Study. Global Heart, 2021, 16, 13.	2.3	2
6	Replication and cross-validation of type 2 diabetes subtypes based on clinical variables: an IMI-RHAPSODY study. Diabetologia, 2021, 64, 1982-1989.	6.3	44
7	Long RNA Sequencing and Ribosome Profiling of Inflamed Î²-Cells Reveal an Extensive Translatome Landscape. Diabetes, 2021, 70, 2299-2312.	0.6	10
8	Profiles of Glucose Metabolism in Different Prediabetes Phenotypes, Classified by Fasting Glycemia, 2-Hour OGTT, Glycated Hemoglobin, and 1-Hour OGTT: An IMI DIRECT Study. Diabetes, 2021, 70, 2092-2106.	0.6	17
9	Distinct Molecular Signatures of Clinical Clusters in People With Type 2 Diabetes: An IMI-RHAPSODY Study. Diabetes, 2021, 70, 2683-2693.	0.6	26
10	Performance of prediction models for nephropathy in people with type 2 diabetes: systematic review and external validation study. BMJ, The, 2021, 374, n2134.	6.0	24
11	Processes Underlying Glycemic Deterioration in Type 2 Diabetes: An IMI DIRECT Study. Diabetes Care, 2021, 44, 511-518.	8.6	16
12	Plasma protein N-glycosylation is associated with cardiovascular disease, nephropathy, and retinopathy in type 2 diabetes. BMJ Open Diabetes Research and Care, 2021, 9, e002345.	2.8	14
13	Genome-Wide Meta-analysis Identifies Genetic Variants Associated With Glycemic Response to Sulfonylureas. Diabetes Care, 2021, 44, 2673-2682.	8.6	23
14	The power of genetic diversity in genome-wide association studies of lipids. Nature, 2021, 600, 675-679.	27.8	353
15	NACHO: an R package for quality control of NanoString nCounter data. Bioinformatics, 2020, 36, 970-971.	4.1	13
16	Heritability estimates for 361 blood metabolites across 40 genome-wide association studies. Nature Communications, 2020, 11, 39.	12.8	64
17	Genetics and Not Shared Environment Explains Familial Resemblance in Adult Metabolomics Data. Twin Research and Human Genetics, 2020, 23, 145-155.	0.6	6
18	CONQUER: an interactive toolbox to understand functional consequences of GWAS hits. NAR Genomics and Bioinformatics, 2020, 2, lqaa085.	3.2	3

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19	Metabolic Age Based on the BBMRI-NL ¹ H-NMR Metabolomics Repository as Biomarker of Age-related Disease. Circulation Genomic and Precision Medicine, 2020, 13, 541-547.	3.6	50
20	Predicting and elucidating the etiology of fatty liver disease: A machine learning modeling and validation study in the IMI DIRECT cohorts. PLoS Medicine, 2020, 17, e1003149.	8.4	47
21	Metformin and statin use associate with plasma protein N-glycosylation in people with type 2 diabetes. BMJ Open Diabetes Research and Care, 2020, 8, e001230.	2.8	8
22	Integration of epidemiologic, pharmacologic, genetic and gut microbiome data in a drugâ€“metabolite atlas. Nature Medicine, 2020, 26, 110-117.	30.7	54
23	The endothelial function biomarker soluble Eâ€selectin is associated with nonalcoholic fatty liver disease. Liver International, 2020, 40, 1079-1088.	3.9	17
24	Plasma Metabolomics Identifies Markers of Impaired Renal Function: A Meta-analysis of 3089 Persons with Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 2275-2287.	3.6	24
25	Post-load glucose subgroups and associated metabolic traits in individuals with type 2 diabetes: An IMI-DIRECT study. PLoS ONE, 2020, 15, e0242360.	2.5	7
26	Title is missing!. , 2020, 17, e1003149.		0
27	Title is missing!. , 2020, 17, e1003149.		0
28	Title is missing!. , 2020, 17, e1003149.		0
29	Title is missing!. , 2020, 17, e1003149.		0
30	Title is missing!. , 2020, 17, e1003149.		0
31	Variation in the Plasma Membrane Monoamine Transporter (PMAT) (Encoded by SLC29A4) and Organic Cation Transporter 1 (OCT1) (Encoded by SLC22A1) and Gastrointestinal Intolerance to Metformin in Type 2 Diabetes: An IMI DIRECT Study. Diabetes Care, 2019, 42, 1027-1033.	8.6	43
32	Visit-to-visit variability of glycemia and vascular complications: the Hoorn Diabetes Care System cohort. Cardiovascular Diabetology, 2019, 18, 170.	6.8	23
33	A Type 1 Diabetes Genetic Risk Score Can Identify Patients With GAD65 Autoantibodyâ€“Positive Type 2 Diabetes Who Rapidly Progress to Insulin Therapy. Diabetes Care, 2019, 42, 208-214.	8.6	35
34	Cohort Profile: The Hoorn Studies. International Journal of Epidemiology, 2018, 47, 396-396j.	1.9	24
35	Metabolite ratios as potential biomarkers for type 2 diabetes: a DIRECT study. Diabetologia, 2018, 61, 117-129.	6.3	32
36	HbA1c is associated with altered expression in blood of cell cycle- and immune response-related genes. Diabetologia, 2018, 61, 138-146.	6.3	10

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37	Blood Metabolomic Measures Associate With Present and Future Glycemic Control in Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 4569-4579.	3.6	25
38	A SNP panel for identification of DNA and RNA specimens. BMC Genomics, 2018, 19, 90.	2.8	47
39	Integrative network analysis highlights biological processes underlying GLP-1 stimulated insulin secretion: A DIRECT study. PLoS ONE, 2018, 13, e0189886.	2.5	9
40	Rare and low-frequency coding variants alter human adult height. Nature, 2017, 542, 186-190.	27.8	544
41	A Genome-Wide Association Study of IVGTT-Based Measures of First-Phase Insulin Secretion Refines the Underlying Physiology of Type 2 Diabetes Variants. Diabetes, 2017, 66, 2296-2309.	0.6	102
42	The Hoorn Diabetes Care System (DCS) cohort. A prospective cohort of persons with type 2 diabetes treated in primary care in the Netherlands. BMJ Open, 2017, 7, e015599.	1.9	58
43	BclI Glucocorticoid Receptor Polymorphism in Relation to Arterial Stiffening and Cardiac Structure and Function: The Hoorn and CODAM Studies. American Journal of Hypertension, 2017, 30, 286-294.	2.0	2
44	Variants in Pharmacokinetic Transporters and Glycemic Response to Metformin: A Metgen Metaâ€Analysis. Clinical Pharmacology and Therapeutics, 2017, 101, 763-772.	4.7	79
45	A Common Gene Variant in Glucokinase Regulatory Protein Interacts With Glucose Metabolism on Diabetic Dyslipidemia: the Combined CODAM and Hoorn Studies. Diabetes Care, 2016, 39, 1811-1817.	8.6	21
46	Variation in the glucose transporter gene SLC2A2 is associated with glycemic response to metformin. Nature Genetics, 2016, 48, 1055-1059.	21.4	165
47	BclI glucocorticoid receptor polymorphism in relation to cardiovascular variables: the Hoorn and CODAM studies. European Journal of Endocrinology, 2015, 173, 455-464.	3.7	15
48	Low mitochondrial DNA content associates with familial longevity: the Leiden Longevity Study. Age, 2014, 36, 9629.	3.0	28
49	SULF2 strongly prediposes to fasting and postprandial triglycerides in patients with obesity and type 2 diabetes mellitus. Obesity, 2014, 22, 1309-1316.	3.0	33
50	Sexâ€specific effects of naturally occurring variants in the dopamine receptor D2 locus on insulin secretion and Type 2 diabetes susceptibility. Diabetic Medicine, 2014, 31, 1001-1008.	2.3	12
51	Exome sequencing-driven discovery of coding polymorphisms associated with common metabolic phenotypes. Diabetologia, 2013, 56, 298-310.	6.3	119
52	The Role of Pharmacogenetics in Drug Disposition and Response of Oral Glucose-Lowering Drugs. Clinical Pharmacokinetics, 2013, 52, 833-854.	3.5	27
53	Preserved GLP-1 and exaggerated GIP secretion in type 2 diabetes and relationships with triglycerides and ALT. European Journal of Endocrinology, 2013, 169, 421-430.	3.7	52
54	Effects of induced hyperinsulinaemia with and without hyperglycaemia on measures of cardiac vagal control. Diabetologia, 2013, 56, 1436-1443.	6.3	25

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55	The CTRB1/2 Locus Affects Diabetes Susceptibility and Treatment via the Incretin Pathway. Diabetes, 2013, 62, 3275-3281.	0.6	96
56	BclI Glucocorticoid Receptor Polymorphism Is Associated With Greater Body Fatness: The Hoorn and CODAM Studies. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E595-E599.	3.6	31
57	A gene variant near ATM is significantly associated with metformin treatment response in type 2 diabetes: a replication and meta-analysis of five cohorts. Diabetologia, 2012, 55, 1971-1977.	6.3	107
58	Glucocorticoid receptor gene polymorphisms are associated with reduced firstâ€phase glucoseâ€stimulated insulin secretion and disposition index in women, but not in men. Diabetic Medicine, 2012, 29, e211-6.	2.3	15
59	Common Variants in the Type 2 Diabetes KCNQ1 Gene Are Associated with Impairments in Insulin Secretion During Hyperglycaemic Glucose Clamp. PLoS ONE, 2012, 7, e32148.	2.5	37
60	The heritability of beta cell function parameters in a mixed meal test design. Diabetologia, 2011, 54, 1043-1051.	6.3	11
61	PS5 - 25. Pharmacogenetics in type 2 diabetes: Genetic variation influences metformin treatment response. Nederlands Tijdschrift Voor Diabetologie, 2011, 9, 107-108.	0.0	0
62	Combined Gene and Protein Expression of Hormone-Sensitive Lipase and Adipose Triglyceride Lipase, Mitochondrial Content, and Adipocyte Size in Subcutaneous and Visceral Adipose Tissue of Morbidly Obese Men. Obesity Facts, 2011, 4, 407-416.	3.4	29
63	The association between the â˜374T/A polymorphism of the receptor for advanced glycation endproducts gene and blood pressure and arterial stiffness is modified by glucose metabolism status: the Hoorn and CoDAM studies. Journal of Hypertension, 2010, 28, 285-293.	0.5	21
64	Genetic association analysis of LARS2 with type 2 diabetes. Diabetologia, 2010, 53, 103-110.	6.3	10
65	Sex Steroids Affect Triglyceride Handling, Glucose-Dependent Insulinotropic Polypeptide, and Insulin Sensitivity. Diabetes Care, 2010, 33, 1831-1833.	8.6	31
66	Gene Variants in the Novel Type 2 Diabetes Loci CDC123/CAMK1D, THADA, ADAMTS9, BCL11A, and MTNR1B Affect Different Aspects of Pancreatic Î²-Cell Function. Diabetes, 2010, 59, 293-301.	0.6	125
67	Combined Risk Allele Score of Eight Type 2 Diabetes Genes Is Associated With Reduced First-Phase Glucose-Stimulated Insulin Secretion During Hyperglycemic Clamps. Diabetes, 2010, 59, 287-292.	0.6	51
68	The Association of Mitochondrial Content with Prevalent and Incident Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1909-1915.	3.6	21
69	The Impact of Genetic Variation in theG6PC2Gene on Insulin Secretion Depends on Glycemia. Journal of Clinical Endocrinology and Metabolism, 2010, 95, E479-E484.	3.6	22
70	Association of Polymorphism in the Receptor for Advanced Glycation End Products (RAGE) Gene with Circulating RAGE Levels. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 5174-5180.	3.6	86
71	Combined effects of single-nucleotide polymorphisms in GCK, GCKR, G6PC2 and MTNR1B on fasting plasma glucose and type 2 diabetes risk. Diabetologia, 2009, 52, 1866-1870.	6.3	71
72	Genetic influences on the insulin response of the beta cell to different secretagogues. Diabetologia, 2009, 52, 2570-2577.	6.3	31

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73	Neuropeptide Y polymorphism significantly magnifies diabetes and cardiovascular disease risk in obesity: the Hoorn Study. European Journal of Clinical Nutrition, 2009, 63, 150-152.	2.9	33
74	Genetic association analysis of 13 nuclear-encoded mitochondrial candidate genes with type II diabetes mellitus: the DAMAGE study. European Journal of Human Genetics, 2009, 17, 1056-1062.	2.8	14
75	Variants of CDKAL1 and IGF2BP2 affect first-phase insulin secretion during hyperglycaemic clamps. Diabetologia, 2008, 51, 1659-1663.	6.3	116
76	Lessons that can be learned from patients with diabetogenic mutations in mitochondrial DNA: implications for common type 2 diabetes. Current Opinion in Clinical Nutrition and Metabolic Care, 2007, 10, 693-697.	2.5	19
77	Impaired glucagon-like peptide-1-induced insulin secretion in carriers of transcription factor 7-like 2 (TCF7L2) gene polymorphisms. Diabetologia, 2007, 50, 2443-2450.	6.3	284
78	New Insights in the Molecular Pathogenesis of the Maternally Inherited Diabetes and Deafness Syndrome. Endocrinology and Metabolism Clinics of North America, 2006, 35, 385-396.	3.2	29
79	Mitochondrial diabetes and its lessons for common Type 2 diabetes. Biochemical Society Transactions, 2006, 34, 819-823.	3.4	36
80	Inflammation and apoptosis genes and the risk of restenosis after percutaneous coronary intervention. Pharmacogenetics and Genomics, 2006, 16, 747-754.	1.5	14
81	Progressive Sensorineural Hearing Impairment in Maternally Inherited Diabetes Mellitus and Deafness (MIDD). Otology and Neurotology, 2006, 27, 802-808.	1.3	5
82	Novel Mitochondrial DNA Length Variants and Genetic Instability in a Family with Diabetes and Deafness. Experimental and Clinical Endocrinology and Diabetes, 2006, 114, 168-174.	1.2	8
83	The HADHSC Gene Encoding Short-Chain L-3-Hydroxyacyl-CoA Dehydrogenase (SCHAD) and Type 2 Diabetes Susceptibility: The DAMAGE Study. Diabetes, 2006, 55, 3193-3196.	0.6	7
84	Tumor necrosis factorâ€Î± plays an important role in restenosis development. FASEB Journal, 2005, 19, 1998-2004.	0.5	73
85	Evidence that the Mitochondrial Leucyl tRNA Synthetase (LARS2) Gene Represents a Novel Type 2 Diabetes Susceptibility Gene. Diabetes, 2005, 54, 1892-1895.	0.6	48
86	Molecular mechanisms of mitochondrial diabetes (MIDD). Annals of Medicine, 2005, 37, 213-221.	3.8	121
87	The Gly482Ser Variant in the Peroxisome Proliferator-Activated Receptor Î³ Coactivator-1 is not Associated with Diabetes-Related Traits in Non-Diabetic German and Dutch Populations. Experimental and Clinical Endocrinology and Diabetes, 2004, 112, 253-257.	1.2	32
88	Mitochondrial Diabetes. Diabetes, 2004, 53, S103-S109.	0.6	376
89	Genetic Factors and Insulin Secretion. Diabetes, 2004, 53, S26-S30.	0.6	68
90	Lack of association between gene variants in the ALMS1 gene and Type 2 diabetes mellitus. Diabetologia, 2003, 46, 1023-1024.	6.3	14

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91	A Case of a De Novo A3243G Mutation in Mitochondrial DNA in a Patient with Diabetes and Deafness. Archives of Physiology and Biochemistry, 2002, 110, 186-188.	2.1	18
92	Variations in Insulin Secretion in Carriers of Gene Variants in IRS-1 and -2. Diabetes, 2002, 51, 884-887.	0.6	26
93	Variations in Insulin Secretion in Carriers of the E23K Variant in the KIR6.2 Subunit of the ATP-Sensitive K+ Channel in the Â-Cell. Diabetes, 2002, 51, 3135-3138.	0.6	45
94	HLA-DQ polymorphism and degree of heteroplasmy of the A3243G mitochondrial DNA mutation in maternally inherited diabetes and deafness. Diabetic Medicine, 2000, 17, 841-847.	2.3	23
95	Characterization of a novel mitochondrial DNA deletion in a patient with a variant of the Pearson marrowâ€“pancreas syndrome. European Journal of Human Genetics, 2000, 8, 195-203.	2.8	40
96	Focal segmental glomerulosclerosis associated with mitochondrial cytopathy. Kidney International, 2000, 58, 1851-1858.	5.2	90
97	Reduced second phase insulin secretion in carriers of a sulphonylurea receptor gene variant associating with Type II diabetes mellitus. Diabetologia, 2000, 43, 515-519.	6.3	35
98	Maternally inherited diabetes and deafness (MIDD): unusual occult exocrine pancreatic manifestation in an affected German family. Experimental and Clinical Endocrinology and Diabetes, 2000, 108, 81-85.	1.2	16
99	Prevalence of Variants in Candidate Genes for Type 2 Diabetes Mellitus in The Netherlands: The Rotterdam Study and the Hoorn Study1. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 1002-1006.	3.6	44
100	Altered beta-cell characteristics in impaired glucose tolerant carriers of a GAA trinucleotide repeat polymorphism in the frataxin gene. Diabetes, 1999, 48, 924-926.	0.6	15
101	Search for mitochondrial DNA mutations in migraine subgroups. Cephalalgia, 1999, 19, 20-22.	3.9	4
102	Search for Mitochondrial Dna Mutations in Migraine Subgroups. Cephalalgia, 1999, 19, 20-22.	3.9	71
103	Variants in the sulphonylurea receptor gene: association of the exon 16-3t variant with Type II diabetes mellitus in Dutch Caucasians. Diabetologia, 1999, 42, 617-620.	6.3	79
104	Prevalence of Variants in Candidate Genes for Type 2 Diabetes Mellitus in The Netherlands: The Rotterdam Study and the Hoorn Study. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 1002-1006.	3.6	45
105	Association of a leucine(7)-to-proline(7) polymorphism in the signal peptide of neuropeptide Y with high serum cholesterol and LDL cholesterol levels. Nature Medicine, 1998, 4, 1434-1437.	30.7	214
106	Maternally Inherited Diabetes and Deafness: A Diabetic Subtype Associated with a Mutation in Mitochondrial DNA. Hormone and Metabolic Research, 1997, 29, 50-55.	1.5	29
107	Detection of Mitochondrial DNA Deletions in Human Skin Fibroblasts of Patients with Pearson's Syndrome by Two-color Fluorescence In Situ Hybridization. Journal of Histochemistry and Cytochemistry, 1997, 45, 55-61.	2.5	38
108	An Insulin Receptor Mutant (Asp707â†' Ala), Involved in Leprechaunism, Is Processed and Transported to the Cell Surface but Unable to Bind Insulin. Journal of Biological Chemistry, 1996, 271, 18719-18724.	3.4	21

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109	Heteroplasmy levels of a mitochondrial gene mutation associated with diabetes mellitus decrease in leucocyte DNA upon aging. Human Mutation, 1996, 7, 193-197.	2.5	87
110	The molecular basis and clinical characteristics of Maternally Inherited Diabetes and Deafness (MIDD), a recently recognized diabetic subtype. Experimental and Clinical Endocrinology and Diabetes, 1996, 104, 205-211.	1.2	22
111	Heteroplasmy levels of a mitochondrial gene mutation associated with diabetes mellitus decrease in leucocyte DNA upon aging. Human Mutation, 1996, 7, 193-197.	2.5	3
112	Absence of the Gly40-Ser Mutation in the Glucagon Receptor Among Diabetic Patients in the Netherlands. Diabetes Care, 1995, 18, 1400-1401.	8.6	6
113	Prevalence of maternally inherited diabetes and deafness in diabetic populations in the Netherlands. Diabetologia, 1994, 37, 1169-1170.	6.3	52
114	Genome-Wide Meta-Analysis Identifies the Organic Anion-Transporting Polypeptide Gene <i>SLCO1B1</i> and Statins as Modifiers of Glycemic Response to Sulfonylureas. SSRN Electronic Journal, 0, , .	0.4	0