

Eric Sijbrands

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

506
papers

55,054
citations

97
h-index

227
g-index

553
ext. papers

64,787
ext. citations

9.7
avg, IF

6.97
L-index

#	Paper	IF	Citations
506	Interplay of Dinner Timing and MTNR1B Type 2 Diabetes Risk Variant on Glucose Tolerance and Insulin Secretion: A Randomized Crossover Trial.. <i>Diabetes Care</i> , 2022 ,	14.6	3
505	Response to Comment on Dawed et al. Genome-Wide Meta-analysis Identifies Genetic Variants Associated With Glycemic Response to Sulfonylureas. <i>Diabetes Care</i> 2021;44:2673-2682.. <i>Diabetes Care</i> , 2022 , 45, e82-e83	14.6	
504	Ancestral diversity improves discovery and fine-mapping of genetic loci for anthropometric traits-The Hispanic/Latino Anthropometry Consortium.. <i>Human Genetics and Genomics Advances</i> , 2022 , 3, 100099	0.8	0
503	Polygenic scores, diet quality, and type 2 diabetes risk: An observational study among 35,759 adults from 3 US cohorts.. <i>PLoS Medicine</i> , 2022 , 19, e1003972	11.6	2
502	Multi-ancestry genetic study of type 2 diabetes highlights the power of diverse populations for discovery and translation.. <i>Nature Genetics</i> , 2022 ,	36.3	7
501	Extending precision medicine tools to populations at high risk of type 2 diabetes.. <i>PLoS Medicine</i> , 2022 , 19, e1003989	11.6	0
500	Genome-wide association analyses highlight etiological differences underlying newly defined subtypes of diabetes. <i>Nature Genetics</i> , 2021 , 53, 1534-1542	36.3	7
499	Obesity Partially Mediates the Diabetogenic Effect of Lowering LDL Cholesterol. <i>Diabetes Care</i> , 2021 ,	14.6	4
498	The Clinical Genome Resource (ClinGen) Familial Hypercholesterolemia Variant Curation Expert Panel consensus guidelines for LDLR variant classification.. <i>Genetics in Medicine</i> , 2021 ,	8.1	6
497	Recessive Genome-wide Meta-analysis Illuminates Genetic Architecture of Type 2 Diabetes. <i>Diabetes</i> , 2021 ,	0.9	0
496	Cross-Laboratory Standardization of Preclinical Lipidomics Using Differential Mobility Spectrometry and Multiple Reaction Monitoring. <i>Analytical Chemistry</i> , 2021 ,	7.8	8
495	Genome-Wide Meta-analysis Identifies Genetic Variants Associated With Glycemic Response to Sulfonylureas. <i>Diabetes Care</i> , 2021 , 44, 2673-2682	14.6	5
494	Effects of Sex, Age, and Apolipoprotein E Genotype on Brain Ceramides and Sphingosine-1-Phosphate in Alzheimer® Disease and Control Mice. <i>Frontiers in Aging Neuroscience</i> , 2021 , 13, 765252	5.3	1
493	The comparative effect of exposure to various risk factors on the risk of hyperuricaemia: diet has a weak causal effect. <i>Arthritis Research and Therapy</i> , 2021 , 23, 75	5.7	5
492	Cardiometabolic risk factors for COVID-19 susceptibility and severity: A Mendelian randomization analysis. <i>PLoS Medicine</i> , 2021 , 18, e1003553	11.6	37
491	The impact of non-additive genetic associations on age-related complex diseases. <i>Nature Communications</i> , 2021 , 12, 2436	17.4	10
490	Genome-wide gene-diet interaction analysis in the UK Biobank identifies novel effects on hemoglobin A1c. <i>Human Molecular Genetics</i> , 2021 , 30, 1773-1783	5.6	3

489	Sequencing cell-free fetal DNA in pregnant women with GCK-MODY: a proof-of-concept study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 ,	5.6	2
488	Lipoprotein(a) is robustly associated with aortic valve calcium. <i>Heart</i> , 2021 , 107, 1422-1428	5.1	5
487	Large-Scale Analysis of Apolipoprotein CIII Glycosylation by Ultrahigh Resolution Mass Spectrometry. <i>Frontiers in Chemistry</i> , 2021 , 9, 678883	5	0
486	The trans-ancestral genomic architecture of glycemic traits. <i>Nature Genetics</i> , 2021 , 53, 840-860	36.3	44
485	Determinants of penetrance and variable expressivity in monogenic metabolic conditions across 77,184 exomes. <i>Nature Communications</i> , 2021 , 12, 3505	17.4	5
484	Genetic Loci and Physiologic Pathways Involved in Gestational Diabetes Mellitus Implicated Through Clustering. <i>Diabetes</i> , 2021 , 70, 268-281	0.9	5
483	A Polygenic Score for Type 2 Diabetes Risk Is Associated With Both the Acute and Sustained Response to Sulfonylureas. <i>Diabetes</i> , 2021 , 70, 293-300	0.9	5
482	Sex-dimorphic genetic effects and novel loci for fasting glucose and insulin variability. <i>Nature Communications</i> , 2021 , 12, 24	17.4	30
481	Interaction of diabetes genetic risk and successful lifestyle modification in the Diabetes Prevention Programme. <i>Diabetes, Obesity and Metabolism</i> , 2021 , 23, 1030-1040	6.7	3
480	Sharing ICU Patient Data Responsibly Under the Society of Critical Care Medicine/European Society of Intensive Care Medicine Joint Data Science Collaboration: The Amsterdam University Medical Centers Database (AmsterdamUMCdb) Example. <i>Critical Care Medicine</i> , 2021 , 49, e563-e577	1.4	18
479	Genetic susceptibility, obesity and lifetime risk of type 2 diabetes: The ARIC study and Rotterdam Study. <i>Diabetic Medicine</i> , 2021 , 38, e14639	3.5	2
478	Breakfast partly restores the anti-inflammatory function of high-density lipoproteins from patients with type 2 diabetes mellitus. <i>Atherosclerosis Plus</i> , 2021 , 44, 43-43		
477	Genetic analysis of dietary intake identifies new loci and functional links with metabolic traits. <i>Nature Human Behaviour</i> , 2021 ,	12.8	5
476	Genome-wide Association Study of Lipid Traits in Youth With Type 2 Diabetes. <i>Journal of the Endocrine Society</i> , 2021 , 5, bvab139	0.4	0
475	Monogenic Diabetes in Youth With Presumed Type 2 Diabetes: Results From the Progress in Diabetes Genetics in Youth (ProDiGY) Collaboration. <i>Diabetes Care</i> , 2021 ,	14.6	6
474	The effect of monomeric and oligomeric FLAVAnols in patients with type 2 diabetes and microalbuminuria (FLAVA-trial): A double-blind randomized controlled trial. <i>Clinical Nutrition</i> , 2021 , 40, 5587-5594	5.9	0
473	HDL associates with insulin resistance and beta-cell dysfunction in South Asian families at risk of type 2 diabetes. <i>Journal of Diabetes and Its Complications</i> , 2021 , 35, 107993	3.2	0
472	A novel integrated QSP model of in vivo human glucose regulation to support the development of a glucagon/GLP-1 dual agonist. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2021 ,	4.5	1

471	Anxiety and depression in diabetes care: longitudinal associations with health-related quality of life. <i>Scientific Reports</i> , 2020 , 10, 8307	4.9	7
470	Trajectories of BMI Before Diagnosis of Type 2 Diabetes: The Rotterdam Study. <i>Obesity</i> , 2020 , 28, 1149-1156	8.1	7
469	Association of the IgG -glycome with the course of kidney function in type 2 diabetes. <i>BMJ Open Diabetes Research and Care</i> , 2020 , 8,	4.5	11
468	Smoking-by-genotype interaction in type 2 diabetes risk and fasting glucose. <i>PLoS ONE</i> , 2020 , 15, e0230815	3.7	4
467	Genetics of diabetes mellitus and diabetes complications. <i>Nature Reviews Nephrology</i> , 2020 , 16, 377-390	14.9	172
466	Health economic evaluation of screening and treating children with familial hypercholesterolemia early in life: Many happy returns on investment?. <i>Atherosclerosis</i> , 2020 , 304, 1-8	3.1	11
465	Precision medicine in diabetes: a Consensus Report from the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD). <i>Diabetologia</i> , 2020 , 63, 1671-1693	10.3	33
464	Precision Medicine in Diabetes: A Consensus Report From the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD). <i>Diabetes Care</i> , 2020 , 43, 1617-1635	14.6	75
463	Lipoprotein(a) plasma levels are not associated with incident microvascular complications in type 2 diabetes mellitus. <i>Diabetologia</i> , 2020 , 63, 1248-1257	10.3	6
462	Comprehensive genomic analysis of dietary habits in UK Biobank identifies hundreds of genetic associations. <i>Nature Communications</i> , 2020 , 11, 1467	17.4	25
461	Metformin and statin use associate with plasma protein -glycosylation in people with type 2 diabetes. <i>BMJ Open Diabetes Research and Care</i> , 2020 , 8,	4.5	2
460	Mendelian Randomization Study of Obesity and Cerebrovascular Disease. <i>Annals of Neurology</i> , 2020 , 87, 516-524	9.4	26
459	Sex difference in the incidence of microvascular complications in patients with type 2 diabetes mellitus: a prospective cohort study. <i>Acta Diabetologica</i> , 2020 , 57, 725-732	3.9	3
458	Reducing the Clinical and Public Health Burden of Familial Hypercholesterolemia: A Global Call to Action. <i>JAMA Cardiology</i> , 2020 , 5, 217-229	16.2	85
457	Melatonin Effects on Glucose Metabolism: Time To Unlock the Controversy. <i>Trends in Endocrinology and Metabolism</i> , 2020 , 31, 192-204	8.8	46
456	Interpreting the Benefit of Simvastatin-Ezetimibe in Patients 75 Years or Older. <i>JAMA Cardiology</i> , 2020 , 5, 234	16.2	2
455	The Need for Precision Medicine to be Applied to Diabetes. <i>Journal of Diabetes Science and Technology</i> , 2020 , 14, 1122-1128	4.1	7
454	Sex differences in cardiometabolic risk factors, pharmacological treatment and risk factor control in type 2 diabetes: findings from the Dutch Diabetes Pearl cohort. <i>BMJ Open Diabetes Research and Care</i> , 2020 , 8,	4.5	6

453	Lifetime risk to progress from pre-diabetes to type 2 diabetes among women and men: comparison between American Diabetes Association and World Health Organization diagnostic criteria. <i>BMJ Open Diabetes Research and Care</i> , 2020 , 8,	4.5	4
452	Analysis of Glucocorticoid-Related Genes Reveal as a New Candidate Gene for Type 2 Diabetes. <i>Journal of the Endocrine Society</i> , 2020 , 4, bvaa121	0.4	2
451	High prevalence of impaired awareness of hypoglycemia and severe hypoglycemia among people with insulin-treated type 2 diabetes: The Dutch Diabetes Pearl Cohort. <i>BMJ Open Diabetes Research and Care</i> , 2020 , 8,	4.5	17
450	Smoking-by-genotype interaction in type 2 diabetes risk and fasting glucose 2020 , 15, e0230815		
449	Smoking-by-genotype interaction in type 2 diabetes risk and fasting glucose 2020 , 15, e0230815		
448	Smoking-by-genotype interaction in type 2 diabetes risk and fasting glucose 2020 , 15, e0230815		
447	Smoking-by-genotype interaction in type 2 diabetes risk and fasting glucose 2020 , 15, e0230815		
446	A Polygenic Lipodystrophy Genetic Risk Score Characterizes Risk Independent of BMI in the Diabetes Prevention Program. <i>Journal of the Endocrine Society</i> , 2019 , 3, 1663-1677	0.4	6
445	Genetic Risk Scores for Diabetes Diagnosis and Precision Medicine. <i>Endocrine Reviews</i> , 2019 , 40, 1500-1520	2.2	94
444	Polyunsaturated Fatty Acid Desaturation Is a Mechanism for Glycolytic NAD Recycling. <i>Cell Metabolism</i> , 2019 , 29, 856-870.e7	24.6	42
443	Mendelian Randomization Analysis of Hemoglobin A as a Risk Factor for Coronary Artery Disease. <i>Diabetes Care</i> , 2019 , 42, 1202-1208	14.6	17
442	Clinical aspects of transgenerational epigenetics 2019 , 465-483		
441	Predictors and patterns of eating behaviors across childhood: Results from The Generation R study. <i>Appetite</i> , 2019 , 141, 104295	4.5	13
440	Novel metabolic indices and incident type 2 diabetes among women and men: the Rotterdam Study. <i>Diabetologia</i> , 2019 , 62, 1581-1590	10.3	26
439	An integrative cross-omics analysis of DNA methylation sites of glucose and insulin homeostasis. <i>Nature Communications</i> , 2019 , 10, 2581	17.4	31
438	Exome sequencing of 20,791 cases of type 2 diabetes and 24,440 controls. <i>Nature</i> , 2019 , 570, 71-76	50.4	129
437	Mortality Risk Associated With Truncating Founder Mutations in Titin. <i>Circulation Genomic and Precision Medicine</i> , 2019 , 12, e002436	5.2	3
436	Dietary Sargassum fusiforme improves memory and reduces amyloid plaque load in an Alzheimer® disease mouse model. <i>Scientific Reports</i> , 2019 , 9, 4908	4.9	32

435	The emerging concept of "individualized cholesterol-lowering therapy": A change in paradigm. <i>Pharmacology & Therapeutics</i> , 2019 , 199, 111-116	13.9	16
434	Genome-wide meta-analysis of macronutrient intake of 91,114 European ancestry participants from the cohorts for heart and aging research in genomic epidemiology consortium. <i>Molecular Psychiatry</i> , 2019 , 24, 1920-1932	15.1	30
433	Dietary antioxidant capacity and risk of type 2 diabetes mellitus, prediabetes and insulin resistance: the Rotterdam Study. <i>European Journal of Epidemiology</i> , 2019 , 34, 853-861	12.1	32
432	Statin treatment increases lipoprotein(a) levels in subjects with low molecular weight apolipoprotein(a) phenotype. <i>Atherosclerosis</i> , 2019 , 289, 201-205	3.1	24
431	Quality of dietary fat and genetic risk of type 2 diabetes: individual participant data meta-analysis. <i>BMJ, The</i> , 2019 , 366, l4292	5.9	23
430	Genome-wide Association Study of Change in Fasting Glucose over time in 13,807 non-diabetic European Ancestry Individuals. <i>Scientific Reports</i> , 2019 , 9, 9439	4.9	3
429	Gain-of-Function Claims for Type-2-Diabetes-Associated Coding Variants in SLC16A11 Are Not Supported by the Experimental Data. <i>Cell Reports</i> , 2019 , 29, 778-780	10.6	4
428	Genome-Wide Association Study of Diabetic Kidney Disease Highlights Biology Involved in Glomerular Basement Membrane Collagen. <i>Journal of the American Society of Nephrology: JASN</i> , 2019 , 30, 2000-2016	12.7	66
427	Metabolite Profiles of Incident Diabetes and Heterogeneity of Treatment Effect in the Diabetes Prevention Program. <i>Diabetes</i> , 2019 , 68, 2337-2349	0.9	13
426	The effect of guideline revisions on vascular complications of type 2 diabetes. <i>Therapeutic Advances in Endocrinology and Metabolism</i> , 2019 , 10, 2042018819875408	4.5	3
425	The SLC16A11 risk haplotype is associated with decreased insulin action, higher transaminases and large-size adipocytes. <i>European Journal of Endocrinology</i> , 2019 , 180, 99-107	6.5	10
424	Protein-coding variants implicate novel genes related to lipid homeostasis contributing to body-fat distribution. <i>Nature Genetics</i> , 2019 , 51, 452-469	36.3	44
423	Genetic Ancestry Markers and Difference in A1c Between African American and White in the Diabetes Prevention Program. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 328-336	5.6	9
422	Translocon Declogger Ste24 Protects against IAPP Oligomer-Induced Proteotoxicity. <i>Cell</i> , 2018 , 173, 62-73.e9	56.2	34
421	Transcription factor 7-like 2 gene links increased in vivo insulin synthesis to type 2 diabetes. <i>EBioMedicine</i> , 2018 , 30, 295-302	8.8	8
420	Refining the accuracy of validated target identification through coding variant fine-mapping in type 2 diabetes. <i>Nature Genetics</i> , 2018 , 50, 559-571	36.3	221
419	Transethnic Evaluation Identifies Low-Frequency Loci Associated With 25-Hydroxyvitamin D Concentrations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018 , 103, 1380-1392	5.6	18
418	Metabolomics insights into early type 2 diabetes pathogenesis and detection in individuals with normal fasting glucose. <i>Diabetologia</i> , 2018 , 61, 1315-1324	10.3	66

4 ¹⁷	Precision medicine in diabetes: an opportunity for clinical translation. <i>Annals of the New York Academy of Sciences</i> , 2018 , 1411, 140-152	6.5	22
4 ¹⁶	Re-analysis of public genetic data reveals a rare X-chromosomal variant associated with type 2 diabetes. <i>Nature Communications</i> , 2018 , 9, 321	17.4	50
4 ¹⁵	Cardiovascular risk in patients with familial hypercholesterolemia using optimal lipid-lowering therapy. <i>Journal of Clinical Lipidology</i> , 2018 , 12, 409-416	4.9	23
4 ¹⁴	Evaluating the contribution of rare variants to type 2 diabetes and related traits using pedigrees. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, 379-384	11.5	21
4 ¹³	Genetic Variation Augments Incretin Resistance and Influences Response to a Sulfonylurea and Metformin: The Study to Understand the Genetics of the Acute Response to Metformin and Glipizide in Humans (SUGAR-MGH). <i>Diabetes Care</i> , 2018 , 41, 554-561	14.6	27
4 ¹²	Group cognitive behavioural therapy and weight regain after diet in type 2 diabetes: results from the randomised controlled POWER trial. <i>Diabetologia</i> , 2018 , 61, 790-799	10.3	11
4 ¹¹	Six-Year Diabetes Incidence After Genetic Risk Testing and Counseling: A Randomized Clinical Trial. <i>Diabetes Care</i> , 2018 , 41, e25-e26	14.6	5
4 ¹⁰	Genetic Evidence That Carbohydrate-Stimulated Insulin Secretion Leads to Obesity. <i>Clinical Chemistry</i> , 2018 , 64, 192-200	5.5	47
4 ⁰⁹	A Genome-Wide Association Study of Diabetic Kidney Disease in Subjects With Type 2 Diabetes. <i>Diabetes</i> , 2018 , 67, 1414-1427	0.9	71
4 ⁰⁸	High Diabetes Distress Among Ethnic Minorities Is Not Explained by Metabolic, Cardiovascular, or Lifestyle Factors: Findings From the Dutch Diabetes Pearl Cohort. <i>Diabetes Care</i> , 2018 , 41, 1854-1861	14.6	11
4 ⁰⁷	Clinical Genetic Testing for Familial Hypercholesterolemia: JACC Scientific Expert Panel. <i>Journal of the American College of Cardiology</i> , 2018 , 72, 662-680	15.1	215
4 ⁰⁶	Use of monomeric and oligomeric flavanols in the dietary management of patients with type 2 diabetes mellitus and microalbuminuria (FLAVA trial): study protocol for a randomized controlled trial. <i>Trials</i> , 2018 , 19, 379	2.8	3
4 ⁰⁵	Reversal of Aging-Induced Increases in Aortic Stiffness by Targeting Cytoskeletal Protein-Protein Interfaces. <i>Journal of the American Heart Association</i> , 2018 , 7,	6	14
4 ⁰⁴	Plasma protein N-glycan signatures of type 2 diabetes. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2018 , 1862, 2613-2622	4	28
4 ⁰³	Genetic inactivation of ANGPTL4 improves glucose homeostasis and is associated with reduced risk of diabetes. <i>Nature Communications</i> , 2018 , 9, 2252	17.4	71
4 ⁰²	Polyunsaturated Fatty Acid Desaturase-Mediated NAD ⁺ Recycling Permits Ongoing Glycolysis and Cell Proliferation. <i>FASEB Journal</i> , 2018 , 32, 672.4	0.9	
4 ⁰¹	Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. <i>Nature Genetics</i> , 2018 , 50, 26-41	36.3	186
4 ⁰⁰	Genetics and biobanks converge to resolve a vexing knowledge gap in diabetes. <i>Lancet Diabetes and Endocrinology</i> , 2018 , 6, 87-89	18.1	

399	Challenges and Opportunities for Cancer Predisposition Cascade Screening for Hereditary Breast and Ovarian Cancer and Lynch Syndrome in Switzerland: Findings from an International Workshop. <i>Public Health Genomics</i> , 2018 , 21, 121-132	1.9	6
398	Fine-mapping type 2 diabetes loci to single-variant resolution using high-density imputation and islet-specific epigenome maps. <i>Nature Genetics</i> , 2018 , 50, 1505-1513	36.3	675
397	Genetic Determinants of Glycemic Traits and the Risk of Gestational Diabetes Mellitus. <i>Diabetes</i> , 2018 , 67, 2703-2709	0.9	17
396	Type 2 diabetes genetic loci informed by multi-trait associations point to disease mechanisms and subtypes: A soft clustering analysis. <i>PLoS Medicine</i> , 2018 , 15, e1002654	11.6	180
395	A Global Overview of Precision Medicine in Type 2 Diabetes. <i>Diabetes</i> , 2018 , 67, 1911-1922	0.9	52
394	Eating behavior and body composition across childhood: a prospective cohort study. <i>International Journal of Behavioral Nutrition and Physical Activity</i> , 2018 , 15, 96	8.4	28
393	ClinVar database of global familial hypercholesterolemia-associated DNA variants. <i>Human Mutation</i> , 2018 , 39, 1631-1640	4.7	55
392	Short-term vascular hemodynamic responses to isometric exercise in young adults and in the elderly. <i>Clinical Interventions in Aging</i> , 2018 , 13, 509-514	4	7
391	Pleiotropic Effect of Human ApoE4 on Cerebral Ceramide and Saturated Fatty Acid Levels. <i>Journal of Alzheimer's Disease</i> , 2017 , 60, 769-781	4.3	3
390	Rare and low-frequency coding variants alter human adult height. <i>Nature</i> , 2017 , 542, 186-190	50.4	412
389	Proprotein convertase subtilisin/kexin 9 inhibition in patients with familial hypercholesterolemia: Initial clinical experience. <i>Journal of Clinical Lipidology</i> , 2017 , 11, 674-681	4.9	22
388	Pharmacogenetics in type 2 diabetes: precision medicine or discovery tool?. <i>Diabetologia</i> , 2017 , 60, 800-807	4.3	41
387	Low-density lipoprotein receptor-negative compound heterozygous familial hypercholesterolemia: Two lifetime journeys of lipid-lowering therapy. <i>Journal of Clinical Lipidology</i> , 2017 , 11, 301-305	4.9	3
386	Novel protein biomarkers associated with coronary artery disease in statin-treated patients with familial hypercholesterolemia. <i>Journal of Clinical Lipidology</i> , 2017 , 11, 682-693	4.9	18
385	Genetic Variation at the Sulfonylurea Receptor, Type 2 Diabetes, and Coronary Heart Disease. <i>Diabetes</i> , 2017 , 66, 2310-2315	0.9	17
384	Effect of diet-induced weight loss on lipoprotein(a) levels in obese individuals with and without type 2 diabetes. <i>Diabetologia</i> , 2017 , 60, 989-997	10.3	17
383	Oxidized LDL, Gamma-Glutamyltransferase and Adverse Outcomes in Older Adults. <i>Journal of the American Geriatrics Society</i> , 2017 , 65, e77-e82	5.6	1
382	Variation in Maturity-Onset Diabetes of the Young Genes Influence Response to Interventions for Diabetes Prevention. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017 , 102, 2678-2689	5.6	12

381	Low-density lipoprotein receptor mutational analysis in diagnosis of familial hypercholesterolemia. <i>Current Opinion in Lipidology</i> , 2017 , 28, 120-129	4.4	26
380	Greater preclinical atherosclerosis in treated monogenic familial hypercholesterolemia vs. polygenic hypercholesterolemia. <i>Atherosclerosis</i> , 2017 , 263, 405-411	3.1	47
379	An Expanded Genome-Wide Association Study of Type 2 Diabetes in Europeans. <i>Diabetes</i> , 2017 , 66, 2888-2902	3.9	414
378	Mining the Genome for Therapeutic Targets. <i>Diabetes</i> , 2017 , 66, 1770-1778	0.9	11
377	Thyroid dysfunction in patients with Down syndrome: Results from a multi-institutional registry study. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 1539-1545	2.5	19
376	A Low-Frequency Inactivating Variant Enriched in the Finnish Population Is Associated With Fasting Insulin Levels and Type 2 Diabetes Risk. <i>Diabetes</i> , 2017 , 66, 2019-2032	0.9	29
375	The anti-inflammatory function of high-density lipoprotein in type II diabetes: A systematic review. <i>Journal of Clinical Lipidology</i> , 2017 , 11, 712-724.e5	4.9	19
374	Genetically Driven Hyperglycemia Increases Risk of Coronary Artery Disease Separately From Type 2 Diabetes. <i>Diabetes Care</i> , 2017 , 40, 687-693	14.6	34
373	Carotid artery plaques and intima medial thickness in familial hypercholesterolemia patients on long-term statin therapy: A case control study. <i>Atherosclerosis</i> , 2017 , 256, 62-66	3.1	14
372	Serum Levels of Apolipoproteins and Incident Type 2 Diabetes: A Prospective Cohort Study. <i>Diabetes Care</i> , 2017 , 40, 346-351	14.6	31
371	Impact of common genetic determinants of Hemoglobin A1c on type 2 diabetes risk and diagnosis in ancestrally diverse populations: A transethnic genome-wide meta-analysis. <i>PLoS Medicine</i> , 2017 , 14, e1002383	11.6	223
370	A Loss-of-Function Splice Acceptor Variant in <i>IS</i> Protective for Type 2 Diabetes. <i>Diabetes</i> , 2017 , 66, 2903-2914	3.9	32
369	A Mendelian Randomization Study of Metabolite Profiles, Fasting Glucose, and Type 2 Diabetes. <i>Diabetes</i> , 2017 , 66, 2915-2926	0.9	27
368	Individual and partner's level of occupation and the association with HbA _{1c} levels in people with Type 2 diabetes mellitus: the Dutch Diabetes Pearl cohort. <i>Diabetic Medicine</i> , 2017 , 34, 1623-1628	3.5	3
367	The pharmacogenetics of metformin. <i>Diabetologia</i> , 2017 , 60, 1648-1655	10.3	46
366	Metabolomics based markers predict type 2 diabetes in a 14-year follow-up study. <i>Metabolomics</i> , 2017 , 13, 104	4.7	50
365	Testing the direction of effects between child body composition and restrictive feeding practices: results from a population-based cohort. <i>American Journal of Clinical Nutrition</i> , 2017 , 106, 783-790	7	66
364	Soluble LR11 associates with aortic root calcification in asymptomatic treated male patients with familial hypercholesterolemia. <i>Atherosclerosis</i> , 2017 , 265, 299-304	3.1	4

363	IgG glycan patterns are associated with type 2 diabetes in independent European populations. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2017 , 1861, 2240-2249	4	64
362	Introduction of the DiaGene study: clinical characteristics, pathophysiology and determinants of vascular complications of type 2 diabetes. <i>Diabetology and Metabolic Syndrome</i> , 2017 , 9, 47	5.6	11
361	Type 2 Diabetes Variants Disrupt Function of SLC16A11 through Two Distinct Mechanisms. <i>Cell</i> , 2017 , 170, 199-212.e20	56.2	94
360	ADAMTS13 activity as a novel risk factor for incident type 2 diabetes mellitus: a population-based cohort study. <i>Diabetologia</i> , 2017 , 60, 280-286	10.3	11
359	The Genetic Landscape of Renal Complications in Type 1 Diabetes. <i>Journal of the American Society of Nephrology: JASN</i> , 2017 , 28, 557-574	12.7	69
358	Sequence data and association statistics from 12,940 type 2 diabetes cases and controls. <i>Scientific Data</i> , 2017 , 4, 170179	8.2	22
357	The Genetic Basis of Type 2 Diabetes in Hispanics and Latin Americans: Challenges and Opportunities. <i>Frontiers in Public Health</i> , 2017 , 5, 329	6	18
356	Identification of a novel proinsulin-associated SNP and demonstration that proinsulin is unlikely to be a causal factor in subclinical vascular remodelling using Mendelian randomisation. <i>Atherosclerosis</i> , 2017 , 266, 196-204	3.1	2
355	A stable isotope method for in vivo assessment of human insulin synthesis and secretion. <i>Acta Diabetologica</i> , 2016 , 53, 935-944	3.9	4
354	Diet-induced weight loss and markers of endothelial dysfunction and inflammation in treated patients with type 2 diabetes. <i>Clinical Nutrition ESPEN</i> , 2016 , 15, 101-106	1.3	11
353	Analysis of protein-coding genetic variation in 60,706 humans. <i>Nature</i> , 2016 , 536, 285-91	50.4	6940
352	Detecting celiac disease in patients with Down syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2016 , 170, 3098-3105	2.5	7
351	Peripheral Blood Transcriptomic Signatures of Fasting Glucose and Insulin Concentrations. <i>Diabetes</i> , 2016 , 65, 3794-3804	0.9	18
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29	Patent foramen ovale and hypercoagulability as combined risk factors for stroke. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2003 , 12, 114-8	2.8	17
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27	Low-density lipoprotein receptor gene mutations and cardiovascular risk in a large genetic cascade screening population. <i>Circulation</i> , 2002 , 106, 3031-6	16.7	86
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25	Review of first 5 years of screening for familial hypercholesterolaemia in the Netherlands. <i>Lancet, The</i> , 2001 , 357, 165-8	40	343
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12	Alcohol consumption and mortality among women. <i>New England Journal of Medicine</i> , 1995 , 332, 1245-50	59.2	586
11	Effect of insulin resistance, apoE2 allele, and smoking on combined hyperlipidemia. <i>Arteriosclerosis and Thrombosis: A Journal of Vascular Biology</i> , 1994 , 14, 1576-80		21
10	Aetiological differences between novel subtypes of diabetes derived from genetic associations		2
9	Refining the accuracy of validated target identification through coding variant fine-mapping in type 2 diabetes		4
8	Rare Non-coding Variation Identified by Large Scale Whole Genome Sequencing Reveals Unexplained Heritability of Type 2 Diabetes		2
7	Rare coding variants in 35 genes associate with circulating lipid levels in a multi-ancestry analysis of 170,000 exomes		2
6	Fine-mapping of an expanded set of type 2 diabetes loci to single-variant resolution using high-density imputation and islet-specific epigenome maps		18
5	Clustering of Type 2 Diabetes Genetic Loci by Multi-Trait Associations Identifies Disease Mechanisms and Subtypes		5
4	Protein-Coding Variants Implicate Novel Genes Related to Lipid Homeostasis Contributing to Body Fat Distribution		1

3	Genetic discovery and translational decision support from exome sequencing of 20,791 type 2 diabetes cases and 24,440 controls from five ancestries	2
2	Comprehensive genomic analysis of dietary habits in UK Biobank identifies hundreds of genetic loci and establishes causal relationships between educational attainment and healthy eating	4
1	The Clinical Genome Resource (ClinGen) Familial Hypercholesterolemia Variant Curation Expert Panel consensus guidelines for LDLR variant classification	1