Leif C Groop

List of Publications by Year in descending order

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248 papers 73,522 citations

86 h-index 946 246 g-index

266 all docs 266 docs citations

times ranked

266

88687 citing authors

#	Article	IF	Citations
1	Subgroups of patients with young-onset type 2 diabetes in India reveal insulin deficiency as a major driver. Diabetologia, 2022, 65, 65-78.	2.9	34
2	Validation of the classification for type 2 diabetes into five subgroups: a report from the ORIGIN trial. Diabetologia, 2022, 65, 206-215.	2.9	31
3	The role of circulating galectin-1 in type 2 diabetes and chronic kidney disease: evidence from cross-sectional, longitudinal and Mendelian randomisation analyses. Diabetologia, 2022, 65, 128-139.	2.9	7
4	Two New Mutations in the <i>CEL</i> Gene Causing Diabetes and Hereditary Pancreatitis: How to Correctly Identify MODY8 Cases. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1455-e1466.	1.8	12
5	Mapping the Cord Blood Transcriptome of Pregnancies Affected by Early Maternal Anemia to Identify Signatures of Fetal Programming. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 1303-1316.	1.8	8
6	Rare coding variants in 35 genes associate with circulating lipid levels—A multi-ancestry analysis of 170,000 exomes. American Journal of Human Genetics, 2022, 109, 81-96.	2.6	24
7	Data-driven subgroups of type 2 diabetes, metabolic response, and renal risk profile after bariatric surgery: a retrospective cohort study. Lancet Diabetes and Endocrinology, the, 2022, 10, 167-176.	5.5	32
8	Multi-ancestry genome-wide association study of gestational diabetes mellitus highlights genetic links with type 2 diabetes. Human Molecular Genetics, 2022, 31, 3377-3391.	1.4	47
9	Obesity-associated Blunted Subcutaneous Adipose Tissue Blood Flow After Meal Improves After Bariatric Surgery. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 1930-1938.	1.8	2
10	A multigenerational study on phenotypic consequences of the most common causal variant of HNF1A-MODY. Diabetologia, 2022, 65, 632-643.	2.9	7
11	Lipid-Associated Variants near ANGPTL3 and LPL Show Parent-of-Origin Specific Effects on Blood Lipid Levels and Obesity. Genes, 2022, 13, 91.	1.0	0
12	Multi-ancestry genetic study of type 2 diabetes highlights the power of diverse populations for discovery and translation. Nature Genetics, 2022, 54, 560-572.	9.4	250
13	Novel Subgroups of Type 2 Diabetes Display Different Epigenetic Patterns That Associate With Future Diabetic Complications. Diabetes Care, 2022, 45, 1621-1630.	4.3	15
14	Genome-wide meta-analysis and omics integration identifies novel genes associated with diabetic kidney disease. Diabetologia, 2022, 65, 1495-1509.	2.9	16
15	Adultâ€onset diabetes in Middle Eastern immigrants to Sweden: Novel subgroups and diabetic complicationsâ€"The All New Diabetes in Scania cohort diabetic complications and ethnicity. Diabetes/Metabolism Research and Reviews, 2021, 37, e3419.	1.7	21
16	Glucose-Dependent Insulinotropic Peptide in the High-Normal Range Is Associated With Increased Carotid Intima-Media Thickness. Diabetes Care, 2021, 44, 224-230.	4.3	20
17	Genetic analysis of obstructive sleep apnoea discovers a strong association with cardiometabolic health. European Respiratory Journal, 2021, 57, 2003091.	3.1	85
18	Sex-dimorphic genetic effects and novel loci for fasting glucose and insulin variability. Nature Communications, 2021, 12, 24.	5.8	87

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19	Relationship between insulin sensitivity and gene expression in human skeletal muscle. BMC Endocrine Disorders, 2021, 21, 32.	0.9	6
20	Accuracy of 1-Hour Plasma Glucose During the Oral Glucose Tolerance Test in Diagnosis of Type 2 Diabetes in Adults: A Meta-analysis. Diabetes Care, 2021, 44, 1062-1069.	4.3	25
21	Combined lifestyle factors and the risk of LADA and type 2 diabetes – Results from a Swedish population-based case-control study. Diabetes Research and Clinical Practice, 2021, 174, 108760.	1.1	8
22	Genetic factors affect the susceptibility to bacterial infections in diabetes. Scientific Reports, 2021, 11, 9464.	1.6	2
23	HLA class I genes modulate disease risk and age at onset together with DR-DQ in Chinese patients with insulin-requiring type 1 diabetes. Diabetologia, 2021, 64, 2026-2036.	2.9	8
24	Determinants of penetrance and variable expressivity in monogenic metabolic conditions across 77,184 exomes. Nature Communications, 2021, 12, 3505.	5.8	49
25	Replication and cross-validation of type 2 diabetes subtypes based on clinical variables: an IMI-RHAPSODY study. Diabetologia, 2021, 64, 1982-1989.	2.9	44
26	Novel Reclassification of Adult Diabetes Is Useful to Distinguish Stages of \hat{l}^2 -Cell Function Linked to the Risk of Vascular Complications: The DOLCE Study From Northern Ukraine. Frontiers in Genetics, 2021, 12, 637945.	1.1	15
27	Distinct Molecular Signatures of Clinical Clusters in People With Type 2 Diabetes: An IMI-RHAPSODY Study. Diabetes, 2021, 70, 2683-2693.	0.3	26
28	Reduced expression of OXPHOS and DNA damage genes is linked to protection from microvascular complications in long-term type 1 diabetes: the PROLONG study. Scientific Reports, 2021, 11, 20735.	1.6	7
29	TIGER: The gene expression regulatory variation landscape of human pancreatic islets. Cell Reports, 2021, 37, 109807.	2.9	45
30	Urinary extracellular vesicles: Assessment of preâ€analytical variables and development of a quality control with focus on transcriptomic biomarker research. Journal of Extracellular Vesicles, 2021, 10, e12158.	5.5	26
31	Genome-Wide Association Study of Peripheral Artery Disease. Circulation Genomic and Precision Medicine, 2021, 14, e002862.	1.6	24
32	Genome-wide association analyses highlight etiological differences underlying newly defined subtypes of diabetes. Nature Genetics, 2021, 53, 1534-1542.	9.4	81
33	Elevated circulating follistatin associates with an increased risk of type 2 diabetes. Nature Communications, 2021, 12, 6486.	5.8	31
34	Genotypes of HLA, TCF7L2, and FTO as potential modifiers of the association between sweetened beverage consumption and risk of LADA and type 2 diabetes. European Journal of Nutrition, 2020, 59, 127-135.	1.8	6
35	Glucocorticoid induces human beta cell dysfunction by involving riborepressor GAS5 LincRNA. Molecular Metabolism, 2020, 32, 160-167.	3.0	37
36	Genetic Discrimination Between LADA and Childhood-Onset Type 1 Diabetes Within the MHC. Diabetes Care, 2020, 43, 418-425.	4.3	23

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37	Liver nucleotide biosynthesis is linked to protection from vascular complications in individuals with long-term type 1 diabetes. Scientific Reports, 2020, 10, 11561.	1.6	8
38	Low-cost exercise interventions improve long-term cardiometabolic health independently of a family history of type 2 diabetes: a randomized parallel group trial. BMJ Open Diabetes Research and Care, 2020, 8, e001377.	1.2	3
39	Discovery of rare variants associated with blood pressure regulation through meta-analysis of 1.3 million individuals. Nature Genetics, 2020, 52, $1314-1332$.	9.4	91
40	Genome-wide association analysis of type 2 diabetes in the EPIC-InterAct study. Scientific Data, 2020, 7, 393.	2.4	19
41	Physical Activity, Genetic Susceptibility, and the Risk of Latent Autoimmune Diabetes in Adults and Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e4112-e4123.	1.8	11
42	Subtypes of Type 2 Diabetes Determined From Clinical Parameters. Diabetes, 2020, 69, 2086-2093.	0.3	103
43	Epigenetic markers associated with metformin response and intolerance in drug-na $ ilde{A}^-$ ve patients with type 2 diabetes. Science Translational Medicine, 2020, 12, .	5.8	34
44	Genetic Predisposition to Coronary Artery Disease in Type 2 Diabetes Mellitus. Circulation Genomic and Precision Medicine, 2020, 13, e002769.	1.6	5
45	The mutational constraint spectrum quantified from variation in 141,456 humans. Nature, 2020, 581, 434-443.	13.7	6,140
46	Transcript expression-aware annotation improves rare variant interpretation. Nature, 2020, 581, 452-458.	13.7	142
47	Metabolic Effects of Gastric Bypass Surgery: Is It All About Calories?. Diabetes, 2020, 69, 2027-2035.	0.3	24
48	Absence of Islet Autoantibodies and Modestly Raised Glucose Values at Diabetes Diagnosis Should Lead to Testing for MODY: Lessons From a 5-Year Pediatric Swedish National Cohort Study. Diabetes Care, 2020, 43, 82-89.	4.3	68
49	Metabolic and Genetic Determinants of Glucose Shape After Oral Challenge in Obese Youths: A Longitudinal Study. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 534-542.	1.8	8
50	Glucose-dependent insulinotropic peptide and risk of cardiovascular events and mortality: a prospective study. Diabetologia, 2020, 63, 1043-1054.	2.9	18
51	Genetic variant effects on gene expression in human pancreatic islets and their implications for T2D. Nature Communications, 2020, 11, 4912.	5.8	89
52	MuscleAtlasExplorer: a web service for studying gene expression in human skeletal muscle. Database: the Journal of Biological Databases and Curation, 2020, 2020, .	1.4	2
53	Hydroxysteroid $17\cdot\hat{l}^2$ dehydrogenase 13 variant increases phospholipids and protects against fibrosis in nonalcoholic fatty liver disease. JCI Insight, 2020, 5, .	2.3	62
54	Heterogeneity of diabetes $\hat{a}\in$ An Indian perspective. Diabetes and Metabolic Syndrome: Clinical Research and Reviews, 2019, 13, 3065-3067.	1.8	4

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55	Fostering improved human islet research: a European perspective. Diabetologia, 2019, 62, 1514-1516.	2.9	13
56	Risk of diabetes-associated diseases in subgroups of patients with recent-onset diabetes: a 5-year follow-up study. Lancet Diabetes and Endocrinology, the, 2019, 7, 684-694.	5.5	364
57	Human pancreatic islet three-dimensional chromatin architecture provides insights into the genetics of type 2 diabetes. Nature Genetics, 2019, 51, 1137-1148.	9.4	208
58	Genome-Wide Association Study of Diabetic Kidney Disease Highlights Biology Involved in Glomerular Basement Membrane Collagen. Journal of the American Society of Nephrology: JASN, 2019, 30, 2000-2016.	3.0	135
59	Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.	5.8	84
60	Loss of ZnT8 function protects against diabetes by enhanced insulin secretion. Nature Genetics, 2019, 51, 1596-1606.	9.4	96
61	Clusters provide a better holistic view of type 2 diabetes than simple clinical features. Lancet Diabetes and Endocrinology,the, 2019, 7, 668-669.	5.5	24
62	FOETAL for NCD—FOetal Exposure and Epidemiological Transitions: the role of Anaemia in early Life for Non-Communicable Diseases in later life: a prospective preconception study in rural Tanzania. BMJ Open, 2019, 9, e024861.	0.8	15
63	Genome editing of human pancreatic beta cell models: problems, possibilities and outlook. Diabetologia, 2019, 62, 1329-1336.	2.9	20
64	Interaction Between Overweight and Genotypes of HLA, TCF7L2, and FTO in Relation to the Risk of Latent Autoimmune Diabetes in Adults and Type 2 Diabetes. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4815-4826.	1.8	22
65	Exome sequencing of 20,791Âcases of type 2 diabetes and 24,440Âcontrols. Nature, 2019, 570, 71-76.	13.7	248
66	Roadmap for a precision-medicine initiative in the Nordic region. Nature Genetics, 2019, 51, 924-930.	9.4	22
67	1-Hour Post-OGTT Glucose Improves the Early Prediction of Type 2 Diabetes by Clinical and Metabolic Markers. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 1131-1140.	1.8	53
68	The associations of daylight and melatonin receptor 1B gene rs10830963 variant with glycemic traits: the prospective PPP-Botnia study. Annals of Medicine, 2019, 51, 58-67.	1.5	7
69	The functional impact of G protein-coupled receptor 142 (Gpr142) on pancreatic \hat{l}^2 -cell in rodent. Pflugers Archiv European Journal of Physiology, 2019, 471, 633-645.	1.3	24
70	Multiethnic Genome-Wide Association Study of Diabetic Retinopathy Using Liability Threshold Modeling of Duration of Diabetes and Glycemic Control. Diabetes, 2019, 68, 441-456.	0.3	54
71	Preserving Insulin Secretion in Diabetes by Inhibiting VDAC1 Overexpression and Surface Translocation in \hat{l}^2 Cells. Cell Metabolism, 2019, 29, 64-77.e6.	7.2	100
72	Novel subgroups of adult-onset diabetes and their association with outcomes: a data-driven cluster analysis of six variables. Lancet Diabetes and Endocrinology, the, 2018, 6, 361-369.	5.5	1,430

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73	Refining the accuracy of validated target identification through coding variant fine-mapping in type 2 diabetes. Nature Genetics, 2018, 50, 559-571.	9.4	356
74	N1-methylnicotinamide is a signalling molecule produced in skeletal muscle coordinating energy metabolism. Scientific Reports, 2018, 8, 3016.	1.6	42
75	HAPT2D: high accuracy of prediction of T2D with a model combining basic and advanced data depending on availability. European Journal of Endocrinology, 2018, 178, 331-341.	1.9	12
76	Genome-wide meta-analysis identifies novel determinants of circulating serum progranulin. Human Molecular Genetics, 2018, 27, 546-558.	1.4	15
77	Family history of diabetes and its relationship with insulin secretion and insulin sensitivity in Iraqi immigrants and native Swedes: a population-based cohort study. Acta Diabetologica, 2018, 55, 233-242.	1.2	13
78	Haplotype Sharing Provides Insights into Fine-Scale Population History and Disease in Finland. American Journal of Human Genetics, 2018, 102, 760-775.	2.6	57
79	A Genome-Wide Association Study of Diabetic Kidney Disease in Subjects With Type 2 Diabetes. Diabetes, 2018, 67, 1414-1427.	0.3	136
80	Overweight, obesity and the risk of LADA: results from a Swedish case–control study and the Norwegian HUNT Study. Diabetologia, 2018, 61, 1333-1343.	2.9	63
81	Role of osteopontin and its regulation in pancreatic islet. Biochemical and Biophysical Research Communications, 2018, 495, 1426-1431.	1.0	8
82	A variant within the FTO confers susceptibility to diabetic nephropathy in Japanese patients with type 2 diabetes. PLoS ONE, 2018, 13, e0208654.	1.1	30
83	Fine-mapping type 2 diabetes loci to single-variant resolution using high-density imputation and islet-specific epigenome maps. Nature Genetics, 2018, 50, 1505-1513.	9.4	1,331
84	Turning Vice into Virtue: Using Batch-Effects to Detect Errors in Large Genomic Data Sets. Genome Biology and Evolution, 2018, 10, 2697-2708.	1.1	7
85	Discovering human diabetes-risk gene function with genetics and physiological assays. Nature Communications, 2018, 9, 3855.	5.8	47
86	First Genome-Wide Association Study of Latent Autoimmune Diabetes in Adults Reveals Novel Insights Linking Immune and Metabolic Diabetes. Diabetes Care, 2018, 41, 2396-2403.	4.3	99
87	A genomeâ€wide association study suggests new evidence for an association of the <scp>NADPH</scp> Oxidase 4 (<i><scp>NOX</scp>4</i>) gene with severe diabetic retinopathy in type 2 diabetes. Acta Ophthalmologica, 2018, 96, e811-e819.	0.6	52
88	Predictors of responses to clinicâ€based childhood obesity care. Pediatric Diabetes, 2018, 19, 1351-1356.	1.2	12
89	Obstructive sleep apnoea and the risk for coronary heart disease and type 2 diabetes: a longitudinal population-based study in Finland. BMJ Open, 2018, 8, e022752.	0.8	54
90	Novel diabetes subgroups – Authors' reply. Lancet Diabetes and Endocrinology,the, 2018, 6, 440-441.	5.5	4

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91	Controllability in an islet specific regulatory network identifies the transcriptional factor NFATC4, which regulates Type 2 Diabetes associated genes. Npj Systems Biology and Applications, 2018, 4, 25.	1.4	25
92	Liver blood dynamics after bariatric surgery: the effects of mixed-meal test and incretin infusions. Endocrine Connections, 2018, 7, 888-896.	0.8	12
93	Activation of imidazoline receptor I 2 , and improved pancreatic \hat{I}^2 -cell function in human islets. Journal of Diabetes and Its Complications, 2018, 32, 813-818.	1.2	3
94	Melatonin receptor 1B gene rs10830963 polymorphism, depressive symptoms and glycaemic traits. Annals of Medicine, 2018, 50, 704-712.	1.5	6
95	Genetic inactivation of ANGPTL4 improves glucose homeostasis and is associated with reduced risk of diabetes. Nature Communications, 2018, 9, 2252.	5.8	99
96	Silencing of the FTO gene inhibits insulin secretion: An in vitro study using GRINCH cells. Molecular and Cellular Endocrinology, 2018, 472, 10-17.	1.6	23
97	DOLORisk: study protocol for a multi-centre observational study to understand the risk factors and determinants of neuropathic pain. Wellcome Open Research, 2018, 3, 63.	0.9	26
98	Bariatric Surgery Enhances Splanchnic Vascular Responses in Patients With Type 2 Diabetes. Diabetes, 2017, 66, 880-885.	0.3	13
99	Impaired hepatic lipid synthesis from polyunsaturated fatty acids in TM6SF2 E167K variant carriers with NAFLD. Journal of Hepatology, 2017, 67, 128-136.	1.8	97
100	MECHANISMS IN ENDOCRINOLOGY: Epigenetic modifications and gestational diabetes: a systematic review of published literature. European Journal of Endocrinology, 2017, 176, R247-R267.	1.9	42
101	Aortic diameter at age 65 in men with newly diagnosed type 2 diabetes. Scandinavian Cardiovascular Journal, 2017, 51, 202-206.	0.4	13
102	Functional Investigations of <i>HNF1A</i> Identify Rare Variants as Risk Factors for Type 2 Diabetes in the General Population. Diabetes, 2017, 66, 335-346.	0.3	54
103	Early metabolic markers identify potential targets for the prevention of type 2 diabetes. Diabetologia, 2017, 60, 1740-1750.	2.9	96
104	Continuous and simultaneous determination of venous blood metabolites. Talanta, 2017, 171, 270-274.	2.9	5
105	An Expanded Genome-Wide Association Study of Type 2 Diabetes in Europeans. Diabetes, 2017, 66, 2888-2902.	0.3	615
106	The rs7903146 Variant in the <i>TCF7L2</i> Gene Increases the Risk of Prediabetes/Type 2 Diabetes in Obese Adolescents by Impairing \hat{l}^2 -Cell Function and Hepatic Insulin Sensitivity. Diabetes Care, 2017, 40, 1082-1089.	4.3	50
107	Automated pathway and reaction prediction facilitates in silico identification of unknown metabolites in human cohort studies. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2017, 1071, 58-67.	1.2	16
108	A Low-Frequency Inactivating <i>AKT2</i> Variant Enriched in the Finnish Population Is Associated With Fasting Insulin Levels and Type 2 Diabetes Risk. Diabetes, 2017, 66, 2019-2032.	0.3	47

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109	Nonsuppressed Glucagon After Glucose Challenge as a Potential Predictor for Glucose Tolerance. Diabetes, 2017, 66, 1373-1379.	0.3	25
110	Differentiation of Diabetes by Pathophysiology, Natural History, and Prognosis. Diabetes, 2017, 66, 241-255.	0.3	454
111	Exome-wide association study of plasma lipids in >300,000 individuals. Nature Genetics, 2017, 49, 1758-1766.	9.4	470
112	Effects of meal and incretins in the regulation of splanchnic blood flow. Endocrine Connections, 2017, 6, 179-187.	0.8	21
113	Heterozygous RFX6 protein truncating variants are associated with MODY with reduced penetrance. Nature Communications, 2017, 8, 888.	5.8	95
114	Glucose-Induced Changes in Gene Expression in Human Pancreatic Islets: Causes or Consequences of Chronic Hyperglycemia. Diabetes, 2017, 66, 3013-3028.	0.3	61
115	The Genetic Landscape of Renal Complications in Type 1 Diabetes. Journal of the American Society of Nephrology: JASN, 2017, 28, 557-574.	3.0	101
116	Sequence data and association statistics from 12,940 type 2 diabetes cases and controls. Scientific Data, 2017, 4, 170179.	2.4	31
117	Genetic determinants of circulating GIP and GLP-1 concentrations. JCI Insight, 2017, 2, .	2.3	46
118	The impact of Roux-en-Y gastric bypass surgery on normal metabolism in a porcine model. PLoS ONE, 2017, 12, e0173137.	1.1	10
119	Glucose-Dependent Insulinotropic Polypeptide Stimulates Osteopontin Expression in the Vasculature via Endothelin-1 and CREB. Diabetes, 2016, 65, 239-254.	0.3	41
120	A Variant of GJD2, Encoding for Connexin 36, Alters the Function of Insulin Producing \hat{l}^2 -Cells. PLoS ONE, 2016, 11, e0150880.	1.1	19
121	The genetic architecture of type 2 diabetes. Nature, 2016, 536, 41-47.	13.7	952
122	Increased Melatonin Signaling Is a Risk Factor for Type 2 Diabetes. Cell Metabolism, 2016, 23, 1067-1077.	7.2	194
123	α-Hydroxybutyric Acid Is a Selective Metabolite Biomarker of Impaired Glucose Tolerance. Diabetes Care, 2016, 39, 988-995.	4.3	93
124	Analysis of 589,306 genomes identifies individuals resilient to severe Mendelian childhood diseases. Nature Biotechnology, 2016, 34, 531-538.	9.4	273
125	Smoking and the Risk of LADA: Results From a Swedish Population-Based Case-Control Study. Diabetes Care, 2016, 39, 794-800.	4.3	26
126	Excess maternal transmission of variants in the THADA gene to offspring with type 2 diabetes. Diabetologia, 2016, 59, 1702-1713.	2.9	19

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127	DNA methylation of loci within <i>ABCG1 </i> and <i>PHOSPHO1 </i> in blood DNA is associated with future type 2 diabetes risk. Epigenetics, 2016, 11, 482-488.	1.3	152
128	Single-Cell Sequencing of Human Pancreatic Isletsâ€"New Kids on the Block. Cell Metabolism, 2016, 24, 523-524.	7.2	7
129	A reference panel of 64,976 haplotypes for genotype imputation. Nature Genetics, 2016, 48, 1279-1283.	9.4	2,421
130	A principal component meta-analysis on multiple anthropometric traits identifies novel loci for body shape. Nature Communications, 2016, 7, 13357.	5.8	74
131	Adhesion G Protein-Coupled Receptor G1 (ADGRG1/GPR56) and Pancreatic \hat{l}^2 -Cell Function. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 4637-4645.	1.8	53
132	Trans-ancestry meta-analyses identify rare and common variants associated with blood pressure and hypertension. Nature Genetics, 2016, 48, 1151-1161.	9.4	261
133	Genome-Wide Association Study of the Modified Stumvoll Insulin Sensitivity Index Identifies <i>BCL2</i> and <i>FAM19A2</i> as Novel Insulin Sensitivity Loci. Diabetes, 2016, 65, 3200-3211.	0.3	67
134	Impact of the TCF7L2 genotype on risk of hypoglycaemia and glucagon secretion during hypoglycaemia. Endocrine Connections, 2016, 5, 53-60.	0.8	2
135	Blood-based biomarkers of age-associated epigenetic changes in human islets associate with insulin secretion and diabetes. Nature Communications, 2016, 7, 11089.	5.8	201
136	Histone acetylation of glucose-induced thioredoxin-interacting protein gene expression in pancreatic islets. International Journal of Biochemistry and Cell Biology, 2016, 81, 82-91.	1.2	20
137	CART is overexpressed in human type 2 diabetic islets and inhibits glucagon secretion and increases insulin secretion. Diabetologia, 2016, 59, 1928-1937.	2.9	24
138	Atrial Natriuretic Peptide in the High Normal Range Is Associated With Lower Prevalence of Insulin Resistance. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 1372-1380.	1.8	17
139	Epigenetic regulation of glucose-stimulated osteopontin (OPN) expression in diabetic kidney. Biochemical and Biophysical Research Communications, 2016, 469, 108-113.	1.0	33
140	Epigenetic regulation of the thioredoxin-interacting protein (TXNIP) gene by hyperglycemia in kidney. Kidney International, 2016, 89, 342-353.	2.6	70
141	Harmonising and linking biomedical and clinical data across disparate data archives to enable integrative cross-biobank research. European Journal of Human Genetics, 2016, 24, 521-528.	1.4	27
142	Influence of Familial Renal Glycosuria Due to Mutations in the SLC5A2 Gene on Changes in Glucose Tolerance over Time. PLoS ONE, 2016, 11, e0146114.	1.1	22
143	A Genome-Wide mQTL Analysis in Human Adipose Tissue Identifies Genetic Variants Associated with DNA Methylation, Gene Expression and Metabolic Traits. PLoS ONE, 2016, 11, e0157776.	1.1	88
144	Type 2 diabetes mellitus. Nature Reviews Disease Primers, 2015, 1, 15019.	18.1	1,308

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145	The Influence of Age and Sex on Genetic Associations with Adult Body Size and Shape: A Large-Scale Genome-Wide Interaction Study. PLoS Genetics, 2015, 11, e1005378.	1.5	331
146	Discovery and Fine-Mapping of Glycaemic and Obesity-Related Trait Loci Using High-Density Imputation. PLoS Genetics, 2015, 11, e1005230.	1.5	77
147	Country of birth modifies the association of fatty liver index with insulin action in Middle Eastern immigrants to Sweden. Diabetes Research and Clinical Practice, 2015, 110, 66-74.	1.1	10
148	New genetic loci link adipose and insulin biology to body fat distribution. Nature, 2015, 518, 187-196.	13.7	1,328
149	Genetic studies of body mass index yield new insights for obesity biology. Nature, 2015, 518, 197-206.	13.7	3,823
150	Modelling of OGTT curve identifies 1 h plasma glucose level as a strong predictor of incident type 2 diabetes: results from two prospective cohorts. Diabetologia, 2015, 58, 87-97.	2.9	106
151	Age- and Sex-Specific Causal Effects of Adiposity on Cardiovascular Risk Factors. Diabetes, 2015, 64, 1841-1852.	0.3	63
152	Genetics and neonatal diabetes: towards precision medicine. Lancet, The, 2015, 386, 934-935.	6.3	11
153	Biomarkers of rapid chronic kidney disease progression in type 2 diabetes. Kidney International, 2015, 88, 888-896.	2.6	124
154	Prevalence and risk factors of gestational diabetes in Punjab, North India: results from a population screening program. European Journal of Endocrinology, 2015, 173, 257-267.	1.9	75
155	Identification and Functional Characterization of G6PC2 Coding Variants Influencing Glycemic Traits Define an Effector Transcript at the G6PC2-ABCB11 Locus. PLoS Genetics, 2015, 11, e1004876.	1.5	95
156	Genetics of Type 2 Diabetesâ€"Pitfalls and Possibilities. Genes, 2015, 6, 87-123.	1.0	337
157	The impact of low-frequency and rare variants on lipid levels. Nature Genetics, 2015, 47, 589-597.	9.4	310
158	The genetics of diabetic complications. Nature Reviews Nephrology, 2015, 11, 277-287.	4.1	124
159	Impact of age, BMI and HbA1c levels on the genome-wide DNA methylation and mRNA expression patterns in human adipose tissue and identification of epigenetic biomarkers in blood. Human Molecular Genetics, 2015, 24, 3792-813.	1.4	223
160	A novel atlas of gene expression in human skeletal muscle reveals molecular changes associated with aging. Skeletal Muscle, 2015, 5, 35.	1.9	78
161	Genetic fine mapping and genomic annotation defines causal mechanisms at type 2 diabetes susceptibility loci. Nature Genetics, 2015, 47, 1415-1425.	9.4	365
162	Identification of novel genes for glucose metabolism based upon expression pattern in human islets and effect on insulin secretion and glycemia. Human Molecular Genetics, 2015, 24, 1945-1955.	1.4	89

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163	Genetics of Type 2 Diabetes: It Matters From Which Parent We Inherit the Risk. Review of Diabetic Studies, 2015, 12, 233-242.	0.5	28
164	Gene-Lifestyle Interaction and Type 2 Diabetes: The EPIC InterAct Case-Cohort Study. PLoS Medicine, 2014, 11, e1001647.	3.9	180
165	Pharmacogenetic meta-analysis of genome-wide association studies of LDL cholesterol response to statins. Nature Communications, 2014, 5, 5068.	5.8	216
166	A Central Role for GRB10 in Regulation of Islet Function in Man. PLoS Genetics, 2014, 10, e1004235.	1.5	164
167	Distribution and Medical Impact of Loss-of-Function Variants in the Finnish Founder Population. PLoS Genetics, 2014, 10, e1004494.	1.5	351
168	Chromosome X-Wide Association Study Identifies Loci for Fasting Insulin and Height and Evidence for Incomplete Dosage Compensation. PLoS Genetics, 2014, 10, e1004127.	1.5	61
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