## Bart van de Sluis

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8636042/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Clearance of p16Ink4a-positive senescent cells delays ageing-associated disorders. Nature, 2011, 479, 232-236.	13.7	2,806
2	Low-density lipoproteins cause atherosclerotic cardiovascular disease. 1. Evidence from genetic, epidemiologic, and clinical studies. A consensus statement from the European Atherosclerosis Society Consensus Panel. European Heart Journal, 2017, 38, 2459-2472.	1.0	2,292
3	Low-density lipoproteins cause atherosclerotic cardiovascular disease: pathophysiological, genetic, and therapeutic insights: a consensus statement from the European Atherosclerosis Society Consensus Panel. European Heart Journal, 2020, 41, 2313-2330.	1.0	776
4	Chronic inflammation induces telomere dysfunction and accelerates ageing in mice. Nature Communications, 2014, 5, 4172.	5.8	596
5	Identification of a new copper metabolism gene by positional cloning in a purebred dog population. Human Molecular Genetics, 2002, 11, 165-173.	1.4	334
6	Increased expression of BubR1 protects against aneuploidy and cancer and extends healthy lifespan. Nature Cell Biology, 2013, 15, 96-102.	4.6	229
7	A Proinflammatory Gut Microbiota Increases Systemic Inflammation and Accelerates Atherosclerosis. Circulation Research, 2019, 124, 94-100.	2.0	226
8	A novel role for XIAP in copper homeostasis through regulation of MURR1. EMBO Journal, 2004, 23, 244-254.	3.5	201
9	COMMD1 is linked to the WASH complex and regulates endosomal trafficking of the copper transporter ATP7A. Molecular Biology of the Cell, 2015, 26, 91-103.	0.9	200
10	CCC- and WASH-mediated endosomal sorting of LDLR is required for normal clearance of circulating LDL. Nature Communications, 2016, 7, 10961.	5.8	165
11	The NADPH oxidase Nox4 has anti-atherosclerotic functions. European Heart Journal, 2015, 36, 3447-3456.	1.0	150
12	Two-Step Senescence-Focused Cancer Therapies. Trends in Cell Biology, 2018, 28, 723-737.	3.6	145
13	The S/T-Rich Motif in the DNAJB6 Chaperone Delays Polyglutamine Aggregation and the Onset of Disease in a Mouse Model. Molecular Cell, 2016, 62, 272-283.	4.5	140
14	Distinct Wilson's Disease Mutations in ATP7B Are Associated With Enhanced Binding to COMMD1 and Reduced Stability of ATP7B. Gastroenterology, 2007, 133, 1316-1326.	0.6	133
15	The ubiquitously expressed MURR1 protein is absent in canine copper toxicosis. Journal of Hepatology, 2003, 39, 703-709.	1.8	131
16	COMMD1 disrupts HIF-1α/β dimerization and inhibits human tumor cell invasion. Journal of Clinical Investigation, 2010, 120, 2119-2130.	3.9	109
17	Increased Activity of Hypoxia-Inducible Factor 1 Is Associated with Early Embryonic Lethality in Commd1 Null Mice. Molecular and Cellular Biology, 2007, 27, 4142-4156.	1.1	107
18	Tuning NF-κB activity: A touch of COMMD proteins. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2013, 1832, 2315-2321.	1.8	97

#	Article	IF	CITATIONS
19	The COMMD Family Regulates Plasma LDL Levels and Attenuates Atherosclerosis Through Stabilizing the CCC Complex in Endosomal LDLR Trafficking. Circulation Research, 2018, 122, 1648-1660.	2.0	94
20	A human-like bile acid pool induced by deletion of hepatic Cyp2c70 modulates effects of FXR activation in mice. Journal of Lipid Research, 2020, 61, 291-305.	2.0	93
21	Mechanism of leukemogenesis by the inv(16) chimeric gene CBFB/PEBP2B-MHY11. Oncogene, 2004, 23, 4297-4307.	2.6	92
22	Characterization of COMMD protein–protein interactions in NF-κB signalling. Biochemical Journal, 2006, 398, 63-71.	1.7	85
23	A liverâ€specific long noncoding RNA with a role in cell viability is elevated in human nonalcoholic steatohepatitis. Hepatology, 2017, 66, 794-808.	3.6	80
24	News on the molecular regulation and function of hepatic low-density lipoprotein receptor and LDLR-related protein 1. Current Opinion in Lipidology, 2017, 28, 241-247.	1.2	76
25	Gene expression profiling of liver cells after copper overload in vivo and in vitro reveals new copper-regulated genes. Journal of Biological Inorganic Chemistry, 2007, 12, 495-507.	1.1	72
26	Endosomal sorting of Notch receptors through COMMD9-dependent pathways modulates Notch signaling. Journal of Cell Biology, 2015, 211, 605-617.	2.3	62
27	The canine copper toxicosis gene MURR1 does not cause non-Wilsonian hepatic copper toxicosis. Journal of Hepatology, 2003, 38, 164-168.	1.8	60
28	The Menkes and Wilson disease genes counteract in copper toxicosis in Labrador retrievers: a new canine model for copper-metabolism disorders. DMM Disease Models and Mechanisms, 2016, 9, 25-38.	1.2	60
29	Liver-Specific Commd1 Knockout Mice Are Susceptible to Hepatic Copper Accumulation. PLoS ONE, 2011, 6, e29183.	1.1	57
30	Evidence for non-random distribution of Fcγ receptor genotype combinations. Immunogenetics, 2003, 55, 240-246.	1.2	56
31	Endonucleases: new tools to edit the mouse genome. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2014, 1842, 1942-1950.	1.8	56
32	Reduced Life- and Healthspan in Mice Carrying a Mono-Allelic BubR1 MVA Mutation. PLoS Genetics, 2012, 8, e1003138.	1.5	52
33	The copper-transporting capacity of ATP7A mutants associated with Menkes disease is ameliorated by COMMD1 as a result of improved protein expression. Cellular and Molecular Life Sciences, 2012, 69, 149-163.	2.4	52
34	Ccne1 Overexpression Causes Chromosome Instability in Liver Cells and Liver Tumor Development in Mice. Gastroenterology, 2019, 157, 210-226.e12.	0.6	50
35	Relevance of animal models for understanding mammalian copper homeostasis. American Journal of Clinical Nutrition, 2008, 88, 840S-845S.	2.2	48
36	COMMD1 Promotes pVHL and O2-Independent Proteolysis of HIF-1α via HSP90/70. PLoS ONE, 2009, 4, e7332.	1.1	47

#	Article	IF	CITATIONS
37	Nuclearâ€Cytosolic Transport of COMMD1 Regulates NFâ€̂PB and HIFâ€1 Activity. Traffic, 2009, 10, 514-527.	1.3	47
38	Impaired Hepatic Vitamin A Metabolism in NAFLD Mice Leading to Vitamin A Accumulation in Hepatocytes. Cellular and Molecular Gastroenterology and Hepatology, 2021, 11, 309-325.e3.	2.3	46
39	The Copper Metabolism MURR1 Domain Protein 1 (COMMD1) Modulates the Aggregation of Misfolded Protein Species in a Client-Specific Manner. PLoS ONE, 2014, 9, e92408.	1.1	45
40	The life cycle of the low-density lipoprotein receptor. Current Opinion in Lipidology, 2015, 26, 82-87.	1.2	43
41	Identification of p38 MAPK and JNK as new targets for correction of Wilson diseaseâ€causing ATP7B mutants. Hepatology, 2016, 63, 1842-1859.	3.6	42
42	Cu,Zn Superoxide Dismutase Maturation and Activity Are Regulated by COMMD1. Journal of Biological Chemistry, 2010, 285, 28991-29000.	1.6	39
43	Tumor necrosis factor receptor 1 gain-of-function mutation aggravates nonalcoholic fatty liver disease but does not cause insulin resistance in a murine model*. Hepatology, 2013, 57, 566-576.	3.6	36
44	Accumulation of 5-oxoproline in myocardial dysfunction and the protective effects of OPLAH. Science Translational Medicine, 2017, 9, .	5.8	36
45	Chemokine-Like Receptor 1 Deficiency Does Not Affect the Development of Insulin Resistance and Nonalcoholic Fatty Liver Disease in Mice. PLoS ONE, 2014, 9, e96345.	1.1	36
46	Elevated mutant dynorphin A causes Purkinje cell loss and motor dysfunction in spinocerebellar ataxia type 23. Brain, 2015, 138, 2537-2552.	3.7	34
47	Copper Metabolism Domain-Containing 1 Represses Genes That Promote Inflammation and Protects Mice From Colitis and Colitis-Associated Cancer. Gastroenterology, 2014, 147, 184-195.e3.	0.6	33
48	Taking One Step Back in Familial Hypercholesterolemia. Arteriosclerosis, Thrombosis, and Vascular Biology, 2020, 40, 973-985.	1.1	33
49	Cholangiopathy and Biliary Fibrosis in Cyp2c70-Deficient Mice Are Fully Reversed by Ursodeoxycholic Acid. Cellular and Molecular Gastroenterology and Hepatology, 2021, 11, 1045-1069.	2.3	31
50	Haploid genetic screens identify SPRING/C12ORF49 as a determinant of SREBP signaling and cholesterol metabolism. Nature Communications, 2020, 11, 1128.	5.8	30
51	The hepatocyte IKK:NF-κB axis promotes liver steatosis by stimulating de novo lipogenesis and cholesterol synthesis. Molecular Metabolism, 2021, 54, 101349.	3.0	28
52	Refined genetic and comparative physical mapping of the canine copper toxicosis locus. Mammalian Genome, 2000, 11, 455-460.	1.0	26
53	Deregulated Renal Calcium and Phosphate Transport during Experimental Kidney Failure. PLoS ONE, 2015, 10, e0142510.	1.1	26
54	Functional understanding of the versatile protein copper metabolism MURR1 domain 1 (COMMD1) in copper homeostasis. Annals of the New York Academy of Sciences, 2014, 1314, 6-14.	1.8	25

1

#	Article	IF	CITATIONS
55	Overexpression of A kinase interacting protein 1 attenuates myocardial ischaemia/reperfusion injury but does not influence heart failure development. Cardiovascular Research, 2016, 111, 217-226.	1.8	24
56	The hepatic WASH complex is required for efficient plasma LDL and HDL cholesterol clearance. JCI Insight, 2019, 4, .	2.3	24
57	Novel aspects of PCSK9 and lipoprotein receptors in renal disease-related dyslipidemia. Cellular Signalling, 2019, 55, 53-64.	1.7	23
58	T cell cholesterol efflux suppresses apoptosis and senescence and increases atherosclerosis in middle aged mice. Nature Communications, 2022, 13, .	5.8	21
59	Cholesterol-induced hepatic inflammation does not contribute to the development of insulin resistance in male LDL receptor knockout mice. Atherosclerosis, 2014, 232, 390-396.	0.4	20
60	A common variant in <i>CCDC93</i> protects against myocardial infarction and cardiovascular mortality by regulating endosomal trafficking of low-density lipoprotein receptor. European Heart Journal, 2020, 41, 1040-1053.	1.0	20
61	COMMD1: A Novel Protein Involved in the Proteolysis of Proteins. Cell Cycle, 2007, 6, 2091-2098.	1.3	16
62	Regulation of murine copper homeostasis by members of the COMMD protein family. DMM Disease Models and Mechanisms, 2021, 14, .	1.2	16
63	Function of the endolysosomal network in cholesterol homeostasis and metabolic-associated fatty liver disease (MAFLD). Molecular Metabolism, 2021, 50, 101146.	3.0	16
64	A cell-type-specific role for murine Commd1 in liver inflammation. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2014, 1842, 2257-2265.	1.8	15
65	The Direct and Indirect Roles of NF-κB in Cancer: Lessons from Oncogenic Fusion Proteins and Knock-in Mice. Biomedicines, 2018, 6, 36.	1.4	15
66	Partial Deletion of Tie2 Affects Microvascular Endothelial Responses to Critical Illness in A Vascular Bed and Organ-Specific Way. Shock, 2019, 51, 757-769.	1.0	14
67	Long Non-Coding RNAs Involved in Progression of Non-Alcoholic Fatty Liver Disease to Steatohepatitis. Cells, 2021, 10, 1883.	1.8	14
68	A20 deficiency in myeloid cells protects mice from diet-induced obesity and insulin resistance due to increased fatty acid metabolism. Cell Reports, 2021, 36, 109748.	2.9	14
69	Nuclear COMMD1 Is Associated with Cisplatin Sensitivity in Ovarian Cancer. PLoS ONE, 2016, 11, e0165385.	1.1	13
70	Naturally Occurring Variants in LRP1 (Low-Density Lipoprotein Receptor–Related Protein 1) Affect HDL (High-Density Lipoprotein) Metabolism Through ABCA1 (ATP-Binding Cassette A1) and SR-B1 (Scavenger) Tj E 1440-1453	ГQq0 <u>0</u> 0 rg	BT /Qverlock
71	Cardiac Function and Architecture Are Maintained in a Model of Cardiorestricted Overexpression of the Prorenin-Renin Receptor. PLoS ONE, 2014, 9, e89929.	1.1	12
72	The polarity protein Scrib limits atherosclerosis development in mice. Cardiovascular Research, 2019, 115, 1963-1974	1.8	11

115, 1963-1974.

#	Article	IF	CITATIONS
73	NF-κB p65 serine 467 phosphorylation sensitizes mice to weight gain and TNFα-or diet-induced inflammation. Biochimica Et Biophysica Acta - Molecular Cell Research, 2017, 1864, 1785-1798.	1.9	9
74	Posttranscriptional Regulation of the Human LDL Receptor by the U2-Spliceosome. Circulation Research, 2022, 130, 80-95.	2.0	9
75	Cholesterol-Induced Hepatic Inflammation Does Not Underlie the Predisposition to Insulin Resistance in Dyslipidemic Female LDL Receptor Knockout Mice. Journal of Diabetes Research, 2015, 2015, 1-12.	1.0	7
76	Senescent cells in the development of cardiometabolic disease. Current Opinion in Lipidology, 2019, 30, 177-185.	1.2	7
77	Modeling Phenotypic Heterogeneity of Glycogen Storage Disease Type 1a Liver Disease in Mice by Somatic CRISPR/CRISPRâ€associated protein 9–Mediated Gene Editing. Hepatology, 2021, 74, 2491-2507.	3.6	7
78	Mice with a deficiency in Peroxisomal Membrane Protein 4 (PXMP4) display mild changes in hepatic lipid metabolism. Scientific Reports, 2022, 12, 2512.	1.6	7
79	Proteinuria converts hepatic heparan sulfate to an effective proprotein convertase subtilisin kexin type 9 enzyme binding partner. Kidney International, 2021, 99, 1369-1381.	2.6	6
80	Recycling the LDL receptor to combat atherosclerosis. Aging, 2018, 10, 3638-3640.	1.4	6
81	Gut microbiota dysbiosis augments atherosclerosis in LDLR-/- mice. Atherosclerosis, 2017, 263, e97.	0.4	3
82	Hypercholesterolemia in Progressive Renal Failure Is Associated with Changes in Hepatic Heparan Sulfate - PCSK9 Interaction. Journal of the American Society of Nephrology: JASN, 2021, 32, 1371-1388.	3.0	3
83	Apolipoprotein F is reduced in humans with steatosis and controls plasma triglycerideâ€rich lipoprotein metabolism. Hepatology, 2023, 77, 1287-1302.	3.6	3
84	News on the genetics of lipoprotein metabolism and cardiovascular disease. Current Opinion in Lipidology, 2013, 24, 185-186.	1.2	2
85	Transgene Design. Methods in Molecular Biology, 2011, 693, 89-101.	0.4	1
86	PS4 - 19. Expression of lipid genes, but not adipokine genes, in visceral adipose tissue is related to liver damage in obese individuals. Nederlands Tijdschrift Voor Diabetologie, 2011, 9, 103-103.	0.0	0
87	PS3 - 13. Enhanced TNF Signaling in Kupffer Cells is Sufficient to Induce NASH. Nederlands Tijdschrift Voor Diabetologie, 2012, 10, 108-108.	0.0	0
88	COMMD1 is an Immune Regulatory Gene That Plays a Role In IBD Pathogenesis. Inflammatory Bowel Diseases, 2012, 18, S104.	0.9	0
89	In-vivo genome editing using CRISPR-Cas9 to study lipid metabolism. Current Opinion in Lipidology, 2016, 27, 92-93.	1.2	0

90 COMMD1 in Copper Homeostasis. , 2019, , 57-63.

#	Article	IF	CITATIONS
91	Prevention of Triglyceridemia by (Non-)Anticoagulant Heparin(oids) Does Not Preclude Transplant Vasculopathy and Glomerulosclerosis. Frontiers in Cell and Developmental Biology, 2022, 10, 798088.	1.8	0