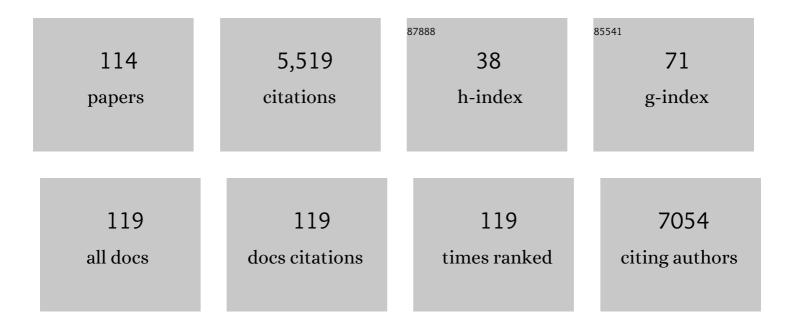
Hans M G Princen

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genetic and Pharmacologic Inactivation of ANGPTL3 and Cardiovascular Disease. New England Journal of Medicine, 2017, 377, 211-221.	27.0	633
2	Mouse Models for Atherosclerosis and Pharmaceutical Modifiers. Arteriosclerosis, Thrombosis, and Vascular Biology, 2007, 27, 1706-1721.	2.4	470
3	Fibrates down-regulate IL-1–stimulated C-reactive protein gene expression in hepatocytes by reducing nuclear p50-NFκB–C/EBP-β complex formation. Blood, 2003, 101, 545-551.	1.4	211
4	The AT04A vaccine against proprotein convertase subtilisin/kexin type 9 reduces total cholesterol, vascular inflammation, and atherosclerosis in APOE*3Leiden.CETP mice. European Heart Journal, 2017, 38, 2499-2507.	2.2	176
5	Evidence for anti-inflammatory activity of statins and PPARα activators in human C-reactive protein transgenic mice in vivo and in cultured human hepatocytes in vitro. Blood, 2004, 103, 4188-4194.	1.4	166
6	No Effect of Consumption of Green and Black Tea on Plasma Lipid and Antioxidant Levels and on LDL Oxidation in Smokers. Arteriosclerosis, Thrombosis, and Vascular Biology, 1998, 18, 833-841.	2.4	165
7	Alirocumab inhibits atherosclerosis, improves the plaque morphology, and enhances the effects of a statin. Journal of Lipid Research, 2014, 55, 2103-2112.	4.2	165
8	Niacin Increases HDL by Reducing Hepatic Expression and Plasma Levels of Cholesteryl Ester Transfer Protein in <i>APOE*3Leiden.CETP</i> Mice. Arteriosclerosis, Thrombosis, and Vascular Biology, 2008, 28, 2016-2022.	2.4	161
9	Rosuvastatin Reduces Atherosclerosis Development Beyond and Independent of Its Plasma Cholesterol–Lowering Effect in APOE*3-Leiden Transgenic Mice. Circulation, 2003, 108, 1368-1374.	1.6	157
10	Metformin Lowers Plasma Triglycerides by Promoting VLDL-Triglyceride Clearance by Brown Adipose Tissue in Mice. Diabetes, 2014, 63, 880-891.	0.6	129
11	Perfluoroalkyl Sulfonates Cause Alkyl Chain Length–Dependent Hepatic Steatosis and Hypolipidemia Mainly by Impairing Lipoprotein Production in APOE*3-Leiden CETP Mice. Toxicological Sciences, 2011, 123, 290-303.	3.1	118
12	Acyl-CoA:Cholesterol Acyltransferase Inhibitor Avasimibe Reduces Atherosclerosis in Addition to Its Cholesterol-Lowering Effect in ApoE*3-Leiden Mice. Circulation, 2001, 103, 1778-1786.	1.6	115
13	Serum carotenoids and vitamins in relation to markers of endothelial. European Journal of Epidemiology, 2004, 19, 915-921.	5.7	112
14	The Cholesterol-Raising Factor from Coffee Beans, Cafestol, as an Agonist Ligand for the Farnesoid and Pregnane X Receptors. Molecular Endocrinology, 2007, 21, 1603-1616.	3.7	107
15	Torcetrapib Does Not Reduce Atherosclerosis Beyond Atorvastatin and Induces More Proinflammatory Lesions Than Atorvastatin. Circulation, 2008, 117, 2515-2522.	1.6	89
16	Insulin suppresses bile acid synthesis in cultured rat hepatocytes by down-regulation of cholesterol 7α-hydroxylase and sterol 27-hydroxylase gene transcription. Hepatology, 1995, 21, 501-510.	7.3	88
17	Fenofibrate increases HDL-cholesterol by reducing cholesteryl ester transfer protein expression. Journal of Lipid Research, 2007, 48, 1763-1771.	4.2	86
18	Systemic PFOS and PFOA exposure and disturbed lipid homeostasis in humans: what do we know and what not?. Critical Reviews in Toxicology, 2021, 51, 141-164.	3.9	78

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19	Effect of Low Dose Atorvastatin Versus Diet-Induced Cholesterol Lowering on Atherosclerotic Lesion Progression and Inflammation in Apolipoprotein E*3–Leiden Transgenic Mice. Arteriosclerosis, Thrombosis, and Vascular Biology, 2005, 25, 161-167.	2.4	77
20	Atorvastatin increases HDL cholesterol by reducing CETP expression in cholesterol-fed APOE*3-Leiden.CETP mice. Atherosclerosis, 2008, 197, 57-63.	0.8	76
21	Osteoarthritis development is induced by increased dietary cholesterol and can be inhibited by atorvastatin in APOE*3Leiden.CETP mice—a translational model for atherosclerosis. Annals of the Rheumatic Diseases, 2014, 73, 921-927.	0.9	67
22	Anacetrapib reduces progression of atherosclerosis, mainly by reducing non-HDL-cholesterol, improves lesion stability and adds to the beneficial effects of atorvastatin. European Heart Journal, 2015, 36, 39-50.	2.2	65
23	PCSK9 inhibition fails to alter hepatic LDLR, circulating cholesterol, and atherosclerosis in the absence of ApoE. Journal of Lipid Research, 2014, 55, 2370-2379.	4.2	59
24	Cafestol, the Cholesterol-Raising Factor in Boiled Coffee, Suppresses Bile Acid Synthesis by Downregulation of Cholesterol 7α-Hydroxylase and Sterol 27-Hydroxylase in Rat Hepatocytes. Arteriosclerosis, Thrombosis, and Vascular Biology, 1997, 17, 3064-3070.	2.4	57
25	Antioxidants and Coronary Heart Disease. Annals of Medicine, 1994, 26, 429-434.	3.8	56
26	Increased Fecal Bile Acid Excretion in Transgenic Mice With Elevated Expression of Human Phospholipid Transfer Protein. Arteriosclerosis, Thrombosis, and Vascular Biology, 2003, 23, 892-897.	2.4	56
27	Dietary Plant Stanol Esters Reduce VLDL Cholesterol Secretion and Bile Saturation in Apolipoprotein E*3-Leiden Transgenic Mice. Arteriosclerosis, Thrombosis, and Vascular Biology, 2001, 21, 1046-1052.	2.4	49
28	Vitamin E inhibits lipid peroxidation-induced adhesion molecule expression in endothelial cells and decreases soluble cell adhesion molecules in healthy subjects. Cardiovascular Research, 2003, 57, 563-571.	3.8	49
29	Bexarotene Induces Dyslipidemia by Increased Very Low-Density Lipoprotein Production and Cholesteryl Ester Transfer Protein-Mediated Reduction of High-Density Lipoprotein. Endocrinology, 2009, 150, 2368-2375.	2.8	49
30	Resveratrol protects against atherosclerosis, but does not add to the antiatherogenic effect of atorvastatin, in APOE*3-Leiden.CETP mice. Journal of Nutritional Biochemistry, 2013, 24, 1423-1430.	4.2	49
31	Alirocumab, evinacumab, and atorvastatin triple therapy regresses plaque lesions and improves lesion composition in mice. Journal of Lipid Research, 2020, 61, 365-375.	4.2	48
32	Acyl-coenzyme A: Cholesterol acyltransferase inhibitor, avasimibe, stimulates bile acid synthesis and cholesterol 7?-hydroxylase in cultured rat hepatocytes andin vivo in the rat. Hepatology, 1999, 30, 491-500.	7.3	47
33	The BCR-ABL1 Inhibitors Imatinib and Ponatinib Decrease Plasma Cholesterol and Atherosclerosis, and Nilotinib and Ponatinib Activate Coagulation in a Translational Mouse Model. Frontiers in Cardiovascular Medicine, 2018, 5, 55.	2.4	47
34	Differential Effects of Amlodipine and Atorvastatin Treatment and Their Combination on Atherosclerosis in ApoE*3-Leiden Transgenic Mice. Journal of Cardiovascular Pharmacology, 2003, 42, 63-70.	1.9	44
35	Innovative pharmaceutical interventions in cardiovascular disease: Focusing on the contribution of non-HDL-C/LDL-C-lowering versus HDL-C-raisingA systematic review and meta-analysis of relevant preclinical studies and clinical trials. European Journal of Pharmacology, 2015, 763, 48-63.	3.5	44
36	Cafestol Increases Serum Cholesterol Levels in Apolipoprotein E*3-Leiden Transgenic Mice by Suppression of Bile Acid Synthesis. Arteriosclerosis, Thrombosis, and Vascular Biology, 2000, 20, 1551-1556.	2.4	42

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37	<scp>APOE</scp> * <scp>3Leiden</scp> . <scp>CETP</scp> transgenic mice as model for pharmaceutical treatment of the metabolic syndrome. Diabetes, Obesity and Metabolism, 2014, 16, 537-544.	4.4	42
38	Increased lipogenesis and resistance of lipoproteins to oxidative modification in two patients with glycogen storage disease type 1a. Journal of Pediatrics, 2002, 140, 256-260.	1.8	40
39	Regulation of low density lipoprotein receptor activity in primary cultures of human hepatocytes by serum lipoproteins. Hepatology, 1986, 6, 1356-1360.	7.3	39
40	LDL Oxidation and Extent of Coronary Atherosclerosis. Arteriosclerosis, Thrombosis, and Vascular Biology, 1998, 18, 193-199.	2.4	39
41	Inhibition of macrophage proliferation dominates plaque regression in response to cholesterol lowering. Basic Research in Cardiology, 2020, 115, 78.	5.9	37
42	CYP7A1 A-278C Polymorphism Affects the Response of Plasma Lipids after Dietary Cholesterol or Cafestol Interventions in Humans. Journal of Nutrition, 2004, 134, 2200-2204.	2.9	36
43	Niacin Reduces Atherosclerosis Development in APOE*3Leiden.CETP Mice Mainly by Reducing NonHDL-Cholesterol. PLoS ONE, 2013, 8, e66467.	2.5	36
44	Hepatic low-density lipoprotein receptor–related protein deficiency in mice increases atherosclerosis independent of plasma cholesterol. Blood, 2004, 103, 3777-3782.	1.4	35
45	Negative effects of rofecoxib treatment on cardiac function after ischemia-reperfusion injury in APOE*3Leiden mice are prevented by combined treatment with thromboxane prostanoid-receptor antagonist S18886 (terutroban)*. Critical Care Medicine, 2008, 36, 2576-2582.	0.9	35
46	Raman spectroscopic investigation of atorvastatin, amlodipine, and both on atherosclerotic plaque development in APOE*3 Leiden transgenic mice. Atherosclerosis, 2002, 164, 65-71.	0.8	34
47	The dual PPARα/γ agonist tesaglitazar blocks progression of preâ€existing atherosclerosis in <i>APOE*3Leiden.CETP</i> transgenic mice. British Journal of Pharmacology, 2009, 156, 1067-1075.	5.4	34
48	Design of a Targeted Peptide Nucleic Acid Prodrug To Inhibit Hepatic Human Microsomal Triglyceride Transfer Protein Expression in Hepatocytesâ€. Bioconjugate Chemistry, 2002, 13, 295-302.	3.6	33
49	PXR agonism decreases plasma HDL levels in ApoEâŽ3-Leiden.CETP mice. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2009, 1791, 191-197.	2.4	33
50	Genetic variation in the rate-limiting enzyme in cholesterol catabolism (cholesterol 7α-hydroxylase) influences the progression of atherosclerosis and risk of new clinical events. Clinical Science, 2005, 108, 539-545.	4.3	32
51	Low dose of the liver X receptor agonist, AZ876, reduces atherosclerosis in APOE*3Leiden mice without affecting liver or plasma triglyceride levels. British Journal of Pharmacology, 2011, 162, 1553-1563.	5.4	32
52	Distribution of perfluorooctanesulfonate and perfluorooctanoate into human plasma lipoprotein fractions. Toxicology Letters, 2012, 210, 360-365.	0.8	32
53	Salsalate attenuates diet induced nonâ€alcoholic steatohepatitis in mice by decreasing lipogenic and inflammatory processes. British Journal of Pharmacology, 2015, 172, 5293-5305.	5.4	29
54	Anti-PCSK9 antibodies inhibit pro-atherogenic mechanisms in APOE*3Leiden.CETP mice. Scientific Reports, 2019, 9, 11079.	3.3	29

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55	Fenofibrate Increases Very Low Density Lipoprotein Triglyceride Production Despite Reducing Plasma Triglyceride Levels in APOE*3-Leiden.CETP Mice. Journal of Biological Chemistry, 2010, 285, 25168-25175.	3.4	28
56	Olmesartan and pravastatin additively reduce development of atherosclerosis in APOE*3Leiden transgenic mice. Journal of Hypertension, 2007, 25, 2454-2462.	0.5	27
57	Aliskiren inhibits atherosclerosis development and improves plaque stability in APOE*3Leiden.CETP transgenic mice with or without treatment with atorvastatin. Journal of Hypertension, 2012, 30, 107-116.	0.5	27
58	Anacetrapib reduces (V)LDL cholesterol by inhibition of CETP activity and reduction of plasma PCSK9. Journal of Lipid Research, 2015, 56, 2085-2093.	4.2	27
59	Effects of amlodipine, atorvastatin and combination of both on advanced atherosclerotic plaque in APOE*3-Leiden transgenic mice. Journal of Molecular and Cellular Cardiology, 2003, 35, 109-118.	1.9	26
60	Cholesterol 7α-Hydroxylase Deficiency in Mice on an APOE*3-Leiden Background Impairs Very-Low-Density Lipoprotein Production. Arteriosclerosis, Thrombosis, and Vascular Biology, 2004, 24, 768-774.	2.4	24
61	In Vivo Magnetic Resonance Imagingâ€Based Detection of Heterogeneous Endothelial Response in Thoracic and Abdominal Aorta to Shortâ€Term Highâ€Fat Diet Ascribed to Differences in Perivascular Adipose Tissue in Mice. Journal of the American Heart Association, 2020, 9, e016929.	3.7	24
62	Bile acids exert negative feedback control on bile acid synthesis in cultured pig hepatocytes by suppression of cholesterol 71±-hydroxylase activity. Hepatology, 1990, 12, 1209-1215.	7.3	23
63	Atorvastatin accelerates clearance of lipoprotein remnants generated by activated brown fat to further reduce hypercholesterolemia and atherosclerosis. Atherosclerosis, 2017, 267, 116-126.	0.8	23
64	Results, meta-analysis and a first evaluation of UNOxR, the urinary nitrate-to-nitrite molar ratio, as a measure of nitrite reabsorption in experimental and clinical settings. Amino Acids, 2018, 50, 799-821.	2.7	23
65	Variable cartilage degradation in mice with diet-induced metabolic dysfunction: food for thought. Osteoarthritis and Cartilage, 2018, 26, 95-107.	1.3	23
66	Rosuvastatin Reduces Plasma Lipids by Inhibiting VLDL Production and Enhancing Hepatobiliary Lipid Excretion in ApoE*3-Leiden Mice. Journal of Cardiovascular Pharmacology, 2005, 45, 53-60.	1.9	21
67	Anti-Atherosclerotic Effect of Amlodipine, Alone and in Combination With Atorvastatin, in APOE*3-Leiden/hCRP Transgenic Mice. Journal of Cardiovascular Pharmacology, 2006, 47, 89-95.	1.9	21
68	Niacin reduces plasma CETP levels by diminishing liver macrophage content in CETP transgenic mice. Biochemical Pharmacology, 2012, 84, 821-829.	4.4	21
69	Dose Effects of Ammonium Perfluorooctanoate on Lipoprotein Metabolism in APOE*3-Leiden.CETP Mice. Toxicological Sciences, 2019, 168, 519-534.	3.1	20
70	Structural Aspects of Bile Acids Involved in the Regulation of Cholesterol 7alpha-Hydroxylase and Sterol 27-Hydroxylase. FEBS Journal, 1995, 228, 596-604.	0.2	19
71	Genetic Analysis of Indicators of Cholesterol Synthesis and Absorption: Lathosterol and Phytosterols in Dutch Twins and Their Parents. Twin Research and Human Genetics, 2003, 6, 307-314.	1.0	17
72	Plasma annexin A5 level relates inversely to the severity of coronary stenosis. Biochemical and Biophysical Research Communications, 2007, 356, 674-680.	2.1	17

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73	Anacetrapib, but not evacetrapib, impairs endothelial function in CETP-transgenic mice in spite of marked HDL-C increase. Atherosclerosis, 2017, 257, 186-194.	0.8	17
74	Colestilan decreases weight gain by enhanced NEFA incorporation in biliary lipids and fecal lipid excretion. Journal of Lipid Research, 2013, 54, 1255-1264.	4.2	15
75	Inflammatory cytokine oncostatin M induces endothelial activation in macro- and microvascular endothelial cells and in APOE*3Leiden.CETP mice. PLoS ONE, 2018, 13, e0204911.	2.5	15
76	Icosabutate Exerts Beneficial Effects Upon Insulin Sensitivity, Hepatic Inflammation, Lipotoxicity, and Fibrosis in Mice. Hepatology Communications, 2020, 4, 193-207.	4.3	15
77	Normal Oxidative Stress and Enhanced Lipoprotein Resistance to In Vitro Oxidation in Hypertriglyceridemia. Arteriosclerosis, Thrombosis, and Vascular Biology, 2000, 20, 2434-2440.	2.4	14
78	Both Transient and Continuous Corticosterone Excess Inhibit Atherosclerotic Plaque Formation in APOE*3-Leiden.CETP Mice. PLoS ONE, 2013, 8, e63882.	2.5	14
79	Absence of an atheroprotective effect of the garlic powder printanor in APOE*3-Leiden transgenic mice. Atherosclerosis, 2004, 177, 291-297.	0.8	13
80	Preferential campesterol incorporation into various tissues in apolipoprotein E*3-Leiden mice consuming plant sterols or stanols. Metabolism: Clinical and Experimental, 2008, 57, 1241-1247.	3.4	13
81	The impact of metabolic syndrome and CRP on vascular phenotype in type 2 diabetes mellitus. European Journal of Internal Medicine, 2008, 19, 115-121.	2.2	13
82	Inhibition and induction of bile acid synthesis by ketoconazole effects on bile formation in the rat. Lipids, 1989, 24, 759-764.	1.7	12
83	Dual targeting of hepatic fibrosis and atherogenesis by icosabutate, an engineered eicosapentaenoic acid derivative. Liver International, 2020, 40, 2860-2876.	3.9	12
84	No effects of atorvastatin (10mg/d or 80mg/d) on nitric oxide, prostacyclin, thromboxane and oxidative stress in type 2 diabetes mellitus patients of the DALI study. Pharmacological Research, 2015, 94, 1-8.	7.1	11
85	Comment on "Hypercholesterolemia with consumption of PFOA-laced Western diets is dependent on strain and sex of mice―by Rebholz S.L. et al. Toxicol. Rep. 2016 (3) 46–54. Toxicology Reports, 2016, 3, 306-309.	3.3	11
86	Oncostatin M reduces atherosclerosis development in APOE*3Leiden.CETP mice and is associated with increased survival probability in humans. PLoS ONE, 2019, 14, e0221477.	2.5	10
87	Beneficial effects of elafibranor on NASH in E3L.CETP mice and differences between mice and men. Scientific Reports, 2021, 11, 5050.	3.3	10
88	Free cytoplasmic messenger ribonucleoprotein complexes from rabbit reticulocytes. Molecular Biology Reports, 1979, 5, 59-64.	2.3	9
89	Cholesterol 7α-Hydroxylase Deficiency in Mice on an APOE*3-Leiden Background Increases Hepatic ABCA1 mRNA Expression and HDL-Cholesterol. Arteriosclerosis, Thrombosis, and Vascular Biology, 2006, 26, 2724-2730.	2.4	8
90	The APOE <mml:math <br="" xmlns:mml="http://www.w3.org/1998/Math/MathML">id="M1"><mml:msup><mml:mrow /><mml:mrow><mml:mo>â^—</mml:mo></mml:mrow></mml:mrow </mml:msup></mml:math> 3-Leiden Heterozygous Glucokinase Knockout Mouse as Novel Translational Disease Model for Type 2 Diabetes, Dyslipidemia, and Diabetic Atherosclerosis. Journal of Diabetes Research, 2019, 2019, 1-13.	2.3	8

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91	CETP does not affect triglyceride production or clearance in APOE*3-Leiden mice. Journal of Lipid Research, 2010, 51, 97-102.	4.2	7
92	Effects of mineral oil administration on the pharmacokinetics, metabolism and pharmacodynamics of atorvastatin and pravastatin in mice and dogs. European Journal of Pharmaceutical Sciences, 2021, 161, 105776.	4.0	7
93	Well-Characterized Garlic-Derived Materials Are Not Hypolipidemic in APOE*3-Leiden Transgenic Mice. Journal of Nutrition, 2004, 134, 1500-1503.	2.9	6
94	Novel high-intensive cholesterol-lowering therapies do not ameliorate knee OA development in humanized dyslipidemic mice. Osteoarthritis and Cartilage, 2021, 29, 1314-1323.	1.3	6
95	Osteoarthritis development is induced by increased dietary cholesterol in APOE*3Leiden.CETP mice, a translational model for atherosclerosis, and can be inhibited by atorvastatin. Osteoarthritis and Cartilage, 2013, 21, S65-S66.	1.3	5
96	HOE 402 lowers serum cholesterol levels by reducing VLDL-lipid production, and not by induction of the LDL receptor, and reduces atherosclerosis in wild-type and LDL receptor-deficient mice. Biochemical Pharmacology, 2002, 63, 1755-1761.	4.4	4
97	α-Tocopherol levels in plasma in new-onset, insulin-dependent diabetes mellitus. European Journal of Internal Medicine, 2004, 15, 371-374.	2.2	4
98	No effects of PCSK9-inhibitor treatment on spatial learning, locomotor activity, and novel object recognition in mice. Behavioural Brain Research, 2021, 396, 112875.	2.2	3
99	Common Variants Associated With OSMR Expression Contribute to Carotid Plaque Vulnerability, but Not to Cardiovascular Disease in Humans. Frontiers in Cardiovascular Medicine, 2021, 8, 658915.	2.4	3
100	Effects of Inhibition or Deletion of PCSK9 (Proprotein Convertase Subtilisin/Kexin Type 9) on Intracerebral Hemorrhage Volumes in Mice. Stroke, 2020, 51, e297-e298.	2.0	2
101	Chronic Oral Administration of Mineral Oil Compared With Corn Oil: Effects on Gut Permeability and Plasma Inflammatory and Lipid Biomarkers. Frontiers in Pharmacology, 2021, 12, 681455.	3.5	2
102	Alirocumab, monoclonal antibody to PCSK9, dose-dependently decreases atherosclerosis, improves plaque stability and shows additive effects with atorvastatin in apoe*3leiden.cetp mice. Atherosclerosis, 2014, 235, e19.	0.8	1
103	The APOE*3Leiden.GK +/- mouse as novel translational model for dyslipidemia, type 2 diabetes and macrovascular complications. Atherosclerosis, 2016, 252, e226-e227.	0.8	1
104	Inflammatory Cytokine Oncostatin M Induces Endothelial Activation in vitro and in APOE*3Leiden.CETP Mice. Atherosclerosis Supplements, 2018, 32, 19.	1.2	1
105	Triple Treatment With Alirocumab And Evinacumab On Top Of Atorvastatin Regresses Lesion Size And Improves Plaque Phenotype In Apoe*3leiden.Cetp Mice. Atherosclerosis, 2019, 287, e12.	0.8	1
106	FRI0528â€HIGH INTENSIVE THERAPEUTIC LOWERING OF SYSTEMIC CHOLESTEROL DOES NOT AMELIORATE O DEVELOPMENT IN KNEE JOINTS OF HUMANIZED DYSLIPIDEMIC MICE. , 2019, , .	A	1
107	DUAL PPAR-ALPHA/GAMMA AGONIST TESAGLITAZAR BLOCKS PROGRESSION OF PRE-EXISTING ATHEROSCLEROSIS IN APOE*3LEIDEN.CETP TRANSGENIC MICE. Atherosclerosis Supplements, 2008, 9, 209.	1.2	0
108	P28 RESVERATROL PROTECTS AGAINST ATHEROSCLEROSIS DEVELOPMENT IN APOE*3-LEIDEN.CETP MICE. Atherosclerosis Supplements, 2010, 11, 22.	1.2	0

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109	P70 THE EFFECT OF ALISKIREN ON ATHEROSCLEROSIS DEVELOPMENT IN APOE*3LEIDEN.CETP TRANSGENIC MICE WITH AND WITHOUT TREATMENT WITH ATORVASTATIN. Atherosclerosis Supplements, 2010, 11, 31.	1.2	0
110	P328 APOE*3LEIDEN.CETP TRANSGENIC MICE AS MODEL FOR THE METABOLIC SYNDROME. Atherosclerosis Supplements, 2010, 11, 86.	1.2	0
111	172 NIACIN, ATORVASTATIN AND FENOFIBRATE DECREASE PLASMA CETP BY REDUCTION OF THE HEPATIC MACROPHAGE CONTENT IN APOE*3-LEIDEN.CETP MICE. Atherosclerosis Supplements, 2011, 12, 38.	1.2	0
112	Affitope-based anti-PCSK9 (proprotein convertase subtilisin/kexin type 9) vaccine (ATO4A) reduces atherosclerosis in APOE*3Leiden.CETP mice. Atherosclerosis, 2016, 252, e253-e254.	0.8	0
113	Cardiovascular safety of BCR-ABL1 tyrosine kinase inhibitors: imatinib and ponatinib decrease plasma cholesterol and atherosclerosis in APOE3*Leiden.CETP Mice. Atherosclerosis, 2017, 263, e29-e30.	0.8	0
114	The Structurally Engineered Fatty Acid Icosabutate Improves Lipid Metabolism And Reduces Severity Of Atherogenesis In Mice. Atherosclerosis, 2019, 287, e55.	0.8	0