List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Targeting the OXE receptor with a selective antagonist inhibits allergenâ€induced pulmonary inflammation in nonâ€human primates. British Journal of Pharmacology, 2022, 179, 322-336.	5.4	6
2	Metabolism of anti-inflammatory OXE (oxoeicosanoid) receptor antagonists by nonhuman primates. European Journal of Pharmaceutical Sciences, 2022, 172, 106144.	4.0	1
3	Novel highly potent OXE receptor antagonists with prolonged plasma lifetimes that are converted to active metabolites in vivo in monkeys. British Journal of Pharmacology, 2020, 177, 388-401.	5.4	10
4	Inhibition of allergenâ€induced dermal eosinophilia by an oxoeicosanoid receptor antagonist in nonâ€human primates. British Journal of Pharmacology, 2020, 177, 360-371.	5.4	10
5	Targeting the OXE receptor as a potential novel therapy for asthma. Biochemical Pharmacology, 2020, 179, 113930.	4.4	14
6	Metabolism and pharmacokinetics of a potent N-acylindole antagonist of the OXE receptor for the eosinophil chemoattractant 5-oxo-6,8,11,14-eicosatetraenoic acid (5-oxo-ETE) in rats and monkeys. European Journal of Pharmaceutical Sciences, 2018, 115, 88-99.	4.0	6
7	Novel Highly Potent and Metabolically Resistant Oxoeicosanoid (OXE) Receptor Antagonists That Block the Actions of the Granulocyte Chemoattractant 5-Oxo-6,8,11,14-Eicosatetraenoic Acid (5-oxo-ETE). Journal of Medicinal Chemistry, 2018, 61, 5934-5948.	6.4	7
8	Structure-activity relationship study of β -oxidation resistant indole-based 5-oxo-6,8,11,14-eicosatetraenoic acid (5-oxo-ETE) receptor antagonists. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 4770-4776.	2.2	4
9	In vivo α-hydroxylation of a 2-alkylindole antagonist of the OXE receptor for the eosinophil chemoattractant 5-oxo-6,8,11,14-eicosatetraenoic acid in monkeys. Biochemical Pharmacology, 2017, 138, 107-118.	4.4	8
10	Design and synthesis of affinity chromatography ligands for the purification of 5-hydroxyeicosanoid dehydrogenase. Bioorganic and Medicinal Chemistry, 2017, 25, 116-125.	3.0	4
11	Pharmacokinetics and Metabolism of Selective Oxoeicosanoid (OXE) Receptor Antagonists and Their Effects on 5-Oxo-6,8,11,14-eicosatetraenoic Acid (5-Oxo-ETE)-Induced Granulocyte Activation in Monkeys. Journal of Medicinal Chemistry, 2016, 59, 10127-10146.	6.4	14
12	5-Oxo-ETE and Inflammation. , 2016, , 185-210.		1
13	Biosynthesis and actions of 5-oxoeicosatetraenoic acid (5-oxo-ETE) on feline granulocytes. Biochemical Pharmacology, 2015, 96, 247-255.	4.4	14
14	Stereoselective synthesis of two highly potent 5-oxo-ETE receptor antagonists. Tetrahedron Letters, 2015, 56, 6896-6899.	1.4	11
15	Biosynthesis, biological effects, and receptors of hydroxyeicosatetraenoic acids (HETEs) and oxoeicosatetraenoic acids (oxo-ETEs) derived from arachidonic acid. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2015, 1851, 340-355.	2.4	234
16	Inhibition of 5-Oxo-6,8,11,14-eicosatetraenoic Acid-Induced Activation of Neutrophils and Eosinophils by Novel Indole OXE Receptor Antagonists. Journal of Medicinal Chemistry, 2014, 57, 364-377.	6.4	27
17	Two Potent OXE-R Antagonists: Assignment of Stereochemistry. ACS Medicinal Chemistry Letters, 2014, 5, 815-819.	2.8	13
18	Base-dependent formation of cis and trans olefins and their application in the synthesis of 5-oxo-ETE receptor antagonists. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 3385-3388.	2.2	7

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19	Quantification of Lipid Mediator Metabolites in Human Urine from Asthma Patients by Electrospray Ionization Mass Spectrometry: Controlling Matrix Effects. Analytical Chemistry, 2013, 85, 7866-7874.	6.5	44
20	Quantification of <i>in vivo</i> oxidative damage in <i>Caenorhabditis elegans</i> during aging by endogenous F3â€isoprostane measurement. Aging Cell, 2013, 12, 214-223.	6.7	39
21	The eosinophil chemoattractant 5-oxo-ETE and the OXE receptor. Progress in Lipid Research, 2013, 52, 651-665.	11.6	71
22	5-Oxo-ETE Receptor Antagonists. Journal of Medicinal Chemistry, 2013, 56, 3725-3732.	6.4	22
23	5-Oxo-ETE is a major oxidative stress-induced arachidonate metabolite in B lymphocytes. Free Radical Biology and Medicine, 2011, 50, 1297-1304.	2.9	15
24	5-Oxo-15-HETE: Total synthesis and bioactivity. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 1857-1860.	2.2	5
25	C20-trifluoro-5-oxo-ETE: A metabolically stable 5-oxo-ETE derivative. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 1987-1990.	2.2	4
26	Enhanced formation of 5-oxo-6,8,11,14-eicosatetraenoic acid by cancer cells in response to oxidative stress, docosahexaenoic acid and neutrophil-derived 5-hydroxy-6,8,11,14-eicosatetraenoic acid. Carcinogenesis, 2011, 32, 822-828.	2.8	19
27	Novel Eicosapentaenoic Acid-derived F3-isoprostanes as Biomarkers of Lipid Peroxidation. Journal of Biological Chemistry, 2009, 284, 23636-23643.	3.4	44
28	5-Oxo-ETE and the OXE receptor. Prostaglandins and Other Lipid Mediators, 2009, 89, 98-104.	1.9	94
29	Oxidative stress-induced changes in pyridine nucleotides and chemoattractant 5-lipoxygenase products in aging neutrophils. Free Radical Biology and Medicine, 2009, 47, 62-71.	2.9	32
30	A new approach to the synthesis of polyunsaturated deuterated isoprostanes: Total synthesis of d4-5-epi-8,12-iso-iPF3α-VI and d4-8,12-iso-iPF3α-VI. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 6755-6758.	2.2	3
31	Substrate Selectivity of 5-Hydroxyeicosanoid Dehydrogenase and Its Inhibition by 5-Hydroxy-Δ ⁶ -Long-Chain Fatty Acids. Journal of Pharmacology and Experimental Therapeutics, 2009, 329, 335-341.	2.5	15
32	Eicosapentaenoic-acid-derived isoprostanes: Synthesis and discovery of two major isoprostanes. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 5523-5527.	2.2	12
33	Human Neutrophils Convert the Sebum-derived Polyunsaturated Fatty Acid Sebaleic Acid to a Potent Granulocyte Chemoattractant. Journal of Biological Chemistry, 2008, 283, 11234-11243.	3.4	28
34	Structural Requirements for Activation of the 5-Oxo-6 <i>E</i> ,8 <i>Z</i> , 11 <i>Z</i> ,14 <i>Z</i> -eicosatetraenoic Acid (5-Oxo-ETE) Receptor: Identification of a Mead Acid Metabolite with Potent Agonist Activity. Journal of Pharmacology and Experimental Therapeutics, 2008, 325, 698-707.	2.5	59
35	Enantio- and Stereospecific Syntheses of 15(R)-Me-PGD2, A Potent and Selective DP2â `Receptor Agonist. Journal of Organic Chemistry, 2008, 73, 7213-7218.	3.2	13
36	Neurofurans, Novel Indices of Oxidant Stress Derived from Docosahexaenoic Acid. Journal of Biological Chemistry, 2008, 283, 6-16.	3.4	73

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37	Agonist and Antagonist Effects of 15R-Prostaglandin (PG) D2 and 11-Methylene-PGD2 on Human Eosinophils and Basophils. Journal of Pharmacology and Experimental Therapeutics, 2007, 320, 173-179.	2.5	19
38	Regulation of 5-hydroxyeicosanoid dehydrogenase activity in monocytic cells. Biochemical Journal, 2007, 403, 157-165.	3.7	28
39	Reductive deprotection of silyl groups with Wilkinson's catalyst/catechol borane. Tetrahedron Letters, 2007, 48, 5289-5292.	1.4	11
40	Airway epithelial cells synthesize the lipid mediator 5-oxo-ETE in response to oxidative stress. Free Radical Biology and Medicine, 2007, 42, 654-664.	2.9	43
41	Total Synthesis of 8,12-iso-iPF3α-VI, an EPA-Derived Isoprostane: Stereoselective Introduction of the Fifth Asymmetric Center. Journal of Organic Chemistry, 2006, 71, 1370-1379.	3.2	33
42	Metabolism of 5-hydroxy-6,8,11,14-eicosatetraenoic acid by human endothelial cells. Biochemical and Biophysical Research Communications, 2006, 350, 151-156.	2.1	18
43	Oxidized derivatives of ω-3 fatty acids: identification of IPF3α-VI in human urine. Journal of Lipid Research, 2006, 47, 2515-2524.	4.2	32
44	An efficient preparation of stereospecific \hat{l}^2 -hydroxy nitriles. Tetrahedron Letters, 2005, 46, 161-164.	1.4	4
45	A new synthetic approach for 4(S)-hydroxycyclopent-2-enone: a precursor to prostanoid synthesis. Tetrahedron Letters, 2005, 46, 6325-6328.	1.4	9
46	iPF2α-III-17,18,19,20-d4: Total synthesis and metabolism. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 1613-1617.	2.2	8
47	Synthesis of 15R-PGD2: a potential DP2 receptor agonist. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 1873-1876.	2.2	14
48	Chronic melatonin therapy fails to alter amyloid burden or oxidative damage in old Tg2576 mice: implications for clinical trials. Brain Research, 2005, 1037, 209-213.	2.2	100
49	An Efficient Preparation of Stereospecific ?-Hydroxy Nitriles ChemInform, 2005, 36, no.	0.0	0
50	Effects of Prostaglandin D ₂ and 5-Lipoxygenase Products on the Expression of CD203c and CD11b by Basophils. Journal of Pharmacology and Experimental Therapeutics, 2005, 312, 627-634.	2.5	51
51	Effects of Prostaglandin D ₂ , 15-Deoxy-Δ ^{12,14} -prostaglandin J ₂ , and Selective DP ₁ and DP ₂ Receptor Agonists on Pulmonary Infiltration of Eosinophils in Brown Norway Rats. Journal of Pharmacology and Experimental Therapeutics, 2005, 313, 64-69	2.5	67
52	Biochemistry, biology and chemistry of the 5-lipoxygenase product 5-oxo-ETE. Progress in Lipid Research, 2005, 44, 154-183.	11.6	124
53	5-Oxo-ETE regulates tone of guinea pig airway smooth muscle via activation of Ca2+pools and Rho-kinase pathway. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2004, 287, L631-L640.	2.9	14
54	5-Oxo-6,8,11,14-eicosatetraenoic Acid Stimulates the Release of the Eosinophil Survival Factor Granulocyte/Macrophage Colony-stimulating Factor from Monocytes. Journal of Biological Chemistry, 2004, 279, 28159-28164.	3.4	27

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55	Oxidative Stress Stimulates the Synthesis of the Eosinophil Chemoattractant 5-Oxo-6,8,11,14-eicosatetraenoic Acid by Inflammatory Cells. Journal of Biological Chemistry, 2004, 279, 40376-40384.	3.4	44
56	Vitamin E reduces amyloidosis and improves cognitive function in Tg2576 mice following repetitive concussive brain injury. Journal of Neurochemistry, 2004, 90, 1541-1541.	3.9	1
57	Vitamin E reduces amyloidosis and improves cognitive function in Tg2576 mice following repetitive concussive brain injury. Journal of Neurochemistry, 2004, 90, 758-764.	3.9	147
58	Silyl Group Deprotection by Pd/C/H2. A Facile and Selective Method ChemInform, 2004, 35, no.	0.0	0
59	Silyl group deprotection by Pd/C/H2. A facile and selective method. Tetrahedron Letters, 2004, 45, 1973-1976.	1.4	19
60	Total synthesis of isoprostanes: discovery and quantitation in biological systems. Chemistry and Physics of Lipids, 2004, 128, 35-56.	3.2	41
61	F2-isoprostanes as indices of lipid peroxidation in inflammatory diseases. Chemistry and Physics of Lipids, 2004, 128, 165-171.	3.2	116
62	Increased F2 isoprostane plasma levels in patients with congestive heart failure are correlated with antioxidant status and disease severity. Journal of Cardiac Failure, 2004, 10, 334-338.	1.7	86
63	5-Oxo-6,8,11,14-eicosatetraenoic acid induces the infiltration of granulocytes into human skin. Journal of Allergy and Clinical Immunology, 2003, 112, 768-774.	2.9	52
64	Vitamin E Reduces Progression of Atherosclerosis in Low-Density Lipoprotein Receptor-Deficient Mice With Established Vascular Lesions. Circulation, 2003, 107, 521-523.	1.6	75
65	15 <i>R</i> -Methyl-Prostaglandin D ₂ Is a Potent and Selective CRTH2/DP ₂ Receptor Agonist in Human Eosinophils. Journal of Pharmacology and Experimental Therapeutics, 2003, 304, 349-355.	2.5	78
66	15-Deoxy-Δ12,1412,14-prostaglandins D2 and J2 Are Potent Activators of Human Eosinophils. Journal of Immunology, 2002, 168, 3563-3569.	0.8	108
67	Increase of Brain Oxidative Stress in Mild Cognitive Impairment. Archives of Neurology, 2002, 59, 972.	4.5	574
68	Local and systemic increase in lipid peroxidation after moderate experimental traumatic brain injury. Journal of Neurochemistry, 2002, 80, 894-898.	3.9	63
69	Brains of Aged Apolipoprotein E-Deficient Mice Have Increased Levels of F2-Isoprostanes, In Vivo Markers of Lipid Peroxidation. Journal of Neurochemistry, 2002, 73, 736-741.	3.9	51
70	The first total synthesis of iPF4α-VI and its deuterated analog. Tetrahedron Letters, 2002, 43, 2801-2805.	1.4	20
71	An efficient approach to the synthesis of LTB4 and ï‰-substituted LTB4 metabolites. Tetrahedron Letters, 2002, 43, 6063-6066.	1.4	5
72	lsoprostane Activation of the Nuclear Hormone Receptor Ppar. Advances in Experimental Medicine and Biology, 2002, 507, 351-355.	1.6	14

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73	Eotaxin and RANTES enhance 5-oxo-6,8,11,14-eicosatetraenoic acid–induced eosinophil chemotaxis. Journal of Allergy and Clinical Immunology, 2001, 107, 272-278.	2.9	37
74	Prostaglandin F2α Receptor-Dependent Regulation of Prostaglandin Transport. Molecular Pharmacology, 2001, 59, 1506-1513.	2.3	14
75	Prostaglandin D2 is a potent chemoattractant for human eosinophils that acts via a novel DP receptor. Blood, 2001, 98, 1942-1948.	1.4	308
76	iPF2α-III metabolism. First total synthesis of 2,3-dinor iPF2α-III, a primary β-oxidation metabolite. Tetrahedron Letters, 2001, 42, 8277-8280.	1.4	6
77	A photoaffinity probe for 5-hydroxyeicosanoid dehydrogenase suitable for radioiodination. Tetrahedron Letters, 2001, 42, 4445-4448.	1.4	13
78	Quantitative Analysis of 5-Oxo-6,8,11,14-eicosatetraenoic Acid by Electrospray Mass Spectrometry Using a Deuterium-Labeled Internal Standard. Analytical Biochemistry, 2001, 295, 262-266.	2.4	10
79	No evidence for lipid peroxidation in severe preeclampsia. American Journal of Obstetrics and Gynecology, 2001, 185, 572-578.	1.3	80
80	Lipid Peroxidation and Platelet Activation in Murine Atherosclerosis. Circulation, 2001, 104, 1940-1945.	1.6	34
81	Absence of 12/15-Lipoxygenase Expression Decreases Lipid Peroxidation and Atherogenesis in Apolipoprotein E–Deficient Mice. Circulation, 2001, 103, 2277-2282.	1.6	225
82	Down's syndrome is associated with increased 8,12― <i>iso</i> â€iPF _{2î±} â€VI levels: Evidence for enhanced lipid peroxidation in vivo. Annals of Neurology, 2000, 48, 795-798.	5.3	62
83	Increased 8,12-iso-iPF2?-VI in Alzheimer's disease: Correlation of a noninvasive index of lipid peroxidation with disease severity. Annals of Neurology, 2000, 48, 809-812.	5.3	341
84	The design and synthesis of a 5-HETE affinity chromatography ligand for 5-hydroxyeicosanoid dehydrogenase. Tetrahedron Letters, 2000, 41, 5807-5811.	1.4	8
85	The synthesis of a 5-HETE photoaffinity ligand. Tetrahedron Letters, 2000, 41, 6313-6317.	1.4	7
86	Intramolecular sulfur-assisted NaBH4 reduction of esters synthesis of 5-oxo-ETE and 5-oxo-12-HETE. Tetrahedron Letters, 2000, 41, 5653-5657.	1.4	16
87	Prothrombinase Acceleration by Oxidatively Damaged Phospholipids. Journal of Biological Chemistry, 2000, 275, 22925-22930.	3.4	26
88	Specific Analysis in Plasma and Urine of 2,3-Dinor-5,6-dihydro-isoprostane F2α-III, a Metabolite of Isoprostane F2α-III and an Oxidation Product of γ-Linolenic Acid. Journal of Biological Chemistry, 2000, 275, 2499-2504.	3.4	39
89	Increased 8,12â€isoâ€iPF2â€VI in Alzheimer's disease: Correlation of a noninvasive index of lipid peroxidation with disease severity. Annals of Neurology, 2000, 48, 809-812.	5.3	4
90	Alcohol-induced generation of lipid peroxidation products in humans. Journal of Clinical Investigation, 1999, 104, 805-813.	8.2	216

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91	Ongoing Prothrombotic State in Patients With Antiphospholipid Antibodies: A Role for Increased Lipid Peroxidation. Blood, 1999, 93, 3401-3407.	1.4	69
92	Biological Inactivation of 5-oxo-6,8,11,14-Eicosatetraenoic Acid by Human Platelets. Blood, 1999, 93, 1086-1096.	1.4	36
93	Leukotrienes, Lipoxins, and Hydroxyeicosatetraenoic Acids. , 1999, 120, 213-247.		14
94	Isoprostanes: Formation, Analysis and Use As Indices of Lipid Peroxidation in Vivo. Journal of Biological Chemistry, 1999, 274, 24441-24444.	3.4	339
95	A new method for the preparation of olefins from vicinal diols. Tetrahedron Letters, 1999, 40, 4019-4022.	1.4	18
96	Synthesis of iPF2α-V: a new route. Tetrahedron Letters, 1999, 40, 6167-6170.	1.4	10
97	A convenient strategy for the synthesis of β,γ-unsaturated aldehydes and acids. A construction of skipped dienes. Tetrahedron Letters, 1999, 40, 7179-7183.	1.4	14
98	Vitamin E suppresses isoprostane generation in vivo and reduces atherosclerosis in ApoE-deficient mice. Nature Medicine, 1998, 4, 1189-1192.	30.7	496
99	Total Synthesis of 17,17,18,18-d4-iPF2α-VI and Quantification of iPF2α-VI in Human Urine by Gas Chromatography/Mass Spectrometry. Analytical Biochemistry, 1998, 262, 45-56.	2.4	19
100	Syntheses and identification of the most abundant urinary type VI isoprostanes. Tetrahedron Letters, 1998, 39, 7039-7042.	1.4	16
101	Dielsâ^'Alder Approach to Isoprostanes. Total Synthesis of iPF2α-V. Journal of the American Chemical Society, 1998, 120, 11953-11961.	13.7	43
102	Total Synthesis of a Potent Proinflammatory 5-Oxo-ETE and Its 6,7-Dihydro Biotransformation Product. Journal of Organic Chemistry, 1998, 63, 337-342.	3.2	62
103	The Total Synthesis of 5-Oxo-12(S)-hydroxy-6(E),8(Z),10(E),14(Z)-eicosatetraenoic Acid and Its 8,9-trans-Isomer and Their Identification in Human Platelets. Journal of Organic Chemistry, 1998, 63, 8976-8982.	3.2	15
104	The Total Synthesis of Tritiated and Deuterated 5-Oxo-ETE, a Novel Inflammatory Mediator. Journal of Organic Chemistry, 1998, 63, 4098-4102.	3.2	19
105	The Isoprostanes, A New Class of Natural Products: Synthesis and Biosynthesis. Synthesis, 1998, 1998, 569-580.	2.3	42
106	Calcium/Calmodulin-dependent Conversion of 5-Oxoeicosanoids to 6,7-Dihydro Metabolites by a Cytosolic Olefin Reductase in Human Neutrophils. Journal of Biological Chemistry, 1998, 273, 20951-20959.	3.4	13
107	Increased Formation of Distinct F ₂ Isoprostanes in Hypercholesterolemia. Circulation, 1998, 98, 2822-2828.	1.6	266
108	ldentification of Two Major F2 Isoprostanes, 8,12-Iso- and 5-epi-8,12-Iso-isoprostane F2α-VI, in Human Urine. Journal of Biological Chemistry, 1998, 273, 29295-29301.	3.4	78

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109	Increased F ₂ â€isoprostanes in Alzheimer's disease: evidence for enhanced lipid peroxidation <i>in vivo</i> . FASEB Journal, 1998, 12, 1777-1783.	0.5	396
110	Functional Characterization of the Ocular Prostaglandin F2α (PGF2α) Receptor. Journal of Biological Chemistry, 1997, 272, 27147-27154.	3.4	97
111	Synthesis of 10,11-Dihydro-12-oxo-LTB4, a Key Biochemical Intermediate. Journal of Organic Chemistry, 1997, 62, 325-330.	3.2	18
112	First total synthesis of isoprostane IPF2α-III. Tetrahedron Letters, 1997, 38, 3339-3342.	1.4	23
113	High-Pressure Liquid Chromatography of Oxo-Eicosanoids Derived from Arachidonic Acid. Analytical Biochemistry, 1997, 247, 17-24.	2.4	16
114	Increased Formation of the Isoprostanes IPF _{2α} -I and 8-Epi-Prostaglandin F _{2α} in Acute Coronary Angioplasty. Circulation, 1997, 96, 3314-3320.	1.6	185
115	Molecular Cloning, Expression and Characterization of Mouse Leukotriene C4 Synthase. FEBS Journal, 1996, 238, 606-612.	0.2	34
116	Total synthesis of 12-epi-PGF2α. Tetrahedron Letters, 1996, 37, 779-782.	1.4	31
117	Deblocking of dithioacetals and oxathioacetals using periodic acid under mild nonaqueous conditions. Tetrahedron Letters, 1996, 37, 4331-4334.	1.4	43
118	Total synthesis of a novel isoprostane IPF2α-I and its identification in biological fluids. Tetrahedron Letters, 1996, 37, 4849-4852.	1.4	54
119	Effects of oxo and dihydro metabolites of 12-hydroxy-5,8,10,14-eicosatetraenoic acid on chemotaxis and cytosolic calcium levels in human neutrophils. Journal of Leukocyte Biology, 1995, 57, 257-263.	3.3	15
120	Total synthesis of proinflammatory dihydro-12-KETE metabolites. Tetrahedron Letters, 1995, 36, 513-516.	1.4	5
121	Regioncontrolled formation of iodohy dnns and expoxides from Vic-diols. Tetrahedron Letters, 1995, 36, 7367-7370.	1.4	23
122	12-Oxo-LTB4, a Key Pivotal Intermediate in LTB4 Metabolism. Journal of Organic Chemistry, 1995, 60, 1806-1813.	3.2	27
123	An Efficient Synthesis of 4(S)-Hydroxycyclopent-2-enone. Journal of Organic Chemistry, 1995, 60, 7548-7551.	3.2	42
124	Synthesis of 12-KETE and its 8,9-trans-isomer. Tetrahedron Letters, 1994, 35, 4051-4054.	1.4	16
125	Total synthesis of 11-R,12-R-dihydroxyeicosatrienoic acid, a metabolite of the cytochrome P-450 epoxygenase pathway. Tetrahedron Letters, 1994, 35, 6239-6242.	1.4	6
126	Total Synthesis of 8-epi-PGF2.alpha A Novel Strategy for the Synthesis of Isoprostanes. Journal of the American Chemical Society, 1994, 116, 10829-10830.	13.7	58

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127	A free radical route to syn lactones and other prostanoid intermediates in isoprostaglandin synthesis Tetrahedron Letters, 1993, 34, 8245-8248.	1.4	42
128	Calcium mobilization and right-angle light scatter responses to 12-oxo-derivatives of arachidonic acid in neutrophils: evidence for the involvement of the leukotriene B4 receptor. Biochimica Et Biophysica Acta - Molecular Cell Research, 1991, 1133, 102-106.	4.1	11
129	[1] Nomenclature. Methods in Enzymology, 1990, 187, 1-9.	1.0	21
130	Synthesis of two analogues of arachidonic acid and their reactions with 12-Lipoxygenase. Tetrahedron, 1990, 46, 6301-6310.	1.9	29
131	Inversion of configurations of contiguous carbinol centres: application to the synthesis of both enantiomers of natural products from the same enantiomerically pure starting material. Carbohydrate Research, 1990, 202, 93-104.	2.3	6
132	Fractional conversion of thromboxane A2 and B2 to urinary 2,3-dinor-thromboxane B2 and 11-dehydrothromboxane B2 in the cynomolgus monkey. Biochimica Et Biophysica Acta - General Subjects, 1989, 992, 71-77.	2.4	15
133	Photoaffinity labelling of the human platelet thromboxane A2/prostaglandin H2 receptor. Biochimica Et Biophysica Acta - Molecular Cell Research, 1989, 1012, 184-190.	4.1	27
134	[4 + 2] Cycloaddition reaction of dibenzyl azodicarboxylate and glycals. Journal of the American Chemical Society, 1989, 111, 2995-3000.	13.7	78
135	Sensitivity of immunoaffinity-purified porcine 5-lipoxygenase to inhibitors and activating lipid hydroperoxides. Biochemical Pharmacology, 1989, 38, 2313-2321.	4.4	47
136	Total synthesis of leukotriene E4 metabolites and precursors to radiolabeled forms of those metabolites. Journal of Organic Chemistry, 1989, 54, 3635-3640.	3.2	32
137	The lipoxins. International Journal of Biochemistry & Cell Biology, 1988, 20, 753-758.	0.5	18
138	Inversion of configurations of contiguous carbinol centers. Journal of Organic Chemistry, 1988, 53, 4421-4422.	3.2	15
139	Biochemical Studies on Mammalian Lipoxygenases,b. Annals of the New York Academy of Sciences, 1988, 524, 12-26.	3.8	21
140	Preparation of complex aminoglycosides: a new strategy. Journal of the American Chemical Society, 1988, 110, 5229-5231.	13.7	41
141	NAD(P)H-dependent reduction of 12-ketoeicosatetraenoic acid to 12(R)-and 12(S)-hydroxyeicosatetraenoic acid by rat liver microsomes. Biochemical and Biophysical Research Communications, 1988, 156, 1083-1089.	2.1	23
142	Lipoxin Syntheses by Arachidonate 12- and 5-Lipoxygenases Purified from Porcine Leukocytes. , 1988, 229, 15-26.		7
143	Lipoxygenase Metabolites. , 1988, , 171-206.		0

144 The Lipoxins: Synthesis and Biosynthesis. , 1988, 229, 79-92.

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145	Pharmacology of L-655,240 (3-[1-(4-chlorobenzyl)-5-fluoro-3-methyl-indol-2-yl]2,2-dimethylpropanoic) Tj ETQq1 1 Pharmacology, 1987, 135, 193-201.	0.784314 3.5	rgBT /Over 37
146	Metabolism and excretion of peptide leukotrienes in the anesthetized rat. Lipids and Lipid Metabolism, 1987, 921, 486-493.	2.6	22
147	Lipoxin synthesis by arachidonate 12-lipoxygenase purified from porcine leukocytes. Biochemical and Biophysical Research Communications, 1987, 149, 1063-1069.	2.1	35
148	Lipoxin synthesis by arachidonate 5-lipoxygenase purified from porcine leukocytes. Biochemical and Biophysical Research Communications, 1987, 144, 996-1002.	2.1	47
149	[4 + 2] Cycloaddition of azodicarboxylate and glycals: a novel and simple method for the preparation of 2-amino-2-deoxy carbohydrates. Journal of the American Chemical Society, 1987, 109, 285-286.	13.7	58
150	Synthesis of 5S-hydroxy-14, 15 LTA4 a biogenic precursor to the lipoxins. Tetrahedron Letters, 1987, 28, 3449-3452.	1.4	20
151	Formation of lipoxin B by the pure reticulocyte lipoxygenase via sequential oxygenation of the substrate. FEBS Journal, 1987, 169, 593-601.	0.2	60
152	The total synthesis of 12-HETE (12-hydroxyeicosatetraenoic acid) and 12,20-diHETE. Journal of Organic Chemistry, 1986, 51, 789-793.	3.2	94
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