

Michel Lazdunski

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.

382
papers

35,214
citations

100
h-index

168
g-index

386
ext. papers

36,989
ext. citations

6.3
avg, IF

6.72
L-index

#	Paper	IF	Citations
382	Targeting the TREK-1 potassium channel via riluzole to eliminate the neuropathic and depressive-like effects of oxaliplatin. <i>Neuropharmacology</i> , 2018 , 140, 43-61	5.5	38
381	Acute and long-term cardioprotective effects of the Traditional Chinese Medicine MLC901 against myocardial ischemia-reperfusion injury in mice. <i>Scientific Reports</i> , 2017 , 7, 14701	4.9	13
380	MLC901 Favors Angiogenesis and Associated Recovery after Ischemic Stroke in Mice. <i>Cerebrovascular Diseases</i> , 2016 , 42, 139-54	3.2	21
379	Role of the TREK2 potassium channel in cold and warm thermosensation and in pain perception. <i>Pain</i> , 2014 , 155, 2534-2544	8	95
378	Activation of TREK-1 by morphine results in analgesia without adverse side effects. <i>Nature Communications</i> , 2013 , 4, 2941	17.4	69
377	Pharmacology of ASIC channels. <i>Environmental Sciences Europe</i> , 2013 , 2, 155-171	5	12
376	Black mamba venom peptides target acid-sensing ion channels to abolish pain. <i>Nature</i> , 2012 , 490, 552-5	50.4	283
375	Asic3 is a neuronal mechanosensor for pressure-induced vasodilation that protects against pressure ulcers. <i>Nature Medicine</i> , 2012 , 18, 1205-7	50.5	74
374	Oxaliplatin-induced cold hypersensitivity is due to remodelling of ion channel expression in nociceptors. <i>EMBO Molecular Medicine</i> , 2011 , 3, 266-78	12	256
373	Spinal cord plasticity and acid-sensing ion channels involvement in a rodent model of irritable bowel syndrome. <i>European Journal of Pain</i> , 2011 , 15, 335-43	3.7	31
372	Acid-sensing ion channels in postoperative pain. <i>Journal of Neuroscience</i> , 2011 , 31, 6059-66	6.6	127
371	Structural elements for the generation of sustained currents by the acid pain sensor ASIC3. <i>Journal of Biological Chemistry</i> , 2009 , 284, 31851-9	5.4	46
370	Acid-sensing ion channel 3 in retinal function and survival 2009 , 50, 2417-26		38
369	Extracellular acidification exerts opposite actions on TREK1 and TREK2 potassium channels via a single conserved histidine residue. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009 , 106, 14628-33	11.5	109
368	The mechano-activated K ⁺ channels TRAAK and TREK-1 control both warm and cold perception. <i>EMBO Journal</i> , 2009 , 28, 1308-18	13	270
367	Sea anemone toxins affecting potassium channels. <i>Progress in Molecular and Subcellular Biology</i> , 2009 , 46, 99-122	3	32
366	ASIC3, a sensor of acidic and primary inflammatory pain. <i>EMBO Journal</i> , 2008 , 27, 3047-55	13	298

365 Deletion of the acid-sensing ion channel ASIC3 prevents gastritis-induced acid hyperresponsiveness of the stomach-brainstem axis. *Pain*, **2008**, 134, 245-253 8 46

364 Acid sensing ion channels in dorsal spinal cord neurons. *Journal of Neuroscience*, **2008**, 28, 1498-508 6.6 89

363 Mtap2 is a constituent of the protein network that regulates twik-related K⁺ channel expression and trafficking. *Journal of Neuroscience*, **2008**, 28, 8545-52 6.6 48

362 A tarantula peptide against pain via ASIC1a channels and opioid mechanisms. *Nature Neuroscience*, **2007**, 10, 943-5 25.5 211

361 Altered acetylcholine, bradykinin and cutaneous pressure-induced vasodilation in mice lacking the TREK1 potassium channel: the endothelial link. *EMBO Reports*, **2007**, 8, 354-9 6.5 73

360 Up- and down-regulation of the mechano-gated K(2P) channel TREK-1 by PIP (2) and other membrane phospholipids. *Pflugers Archiv European Journal of Physiology*, **2007**, 455, 97-103 4.6 61

359 Regulation of the Mechano-Gated K2P Channel TREK-1 by Membrane Phospholipids. *Current Topics in Membranes*, **2007**, 59, 155-70 2.2 18

358 Polyunsaturated fatty acids are cerebral vasodilators via the TREK-1 potassium channel. *Circulation Research*, **2007**, 101, 176-84 15.7 107

357 Antipsychotics inhibit TREK but not TRAAK channels. *Biochemical and Biophysical Research Communications*, **2007**, 354, 284-9 3.4 48

356 Does sumoylation control K2P1/TWIK1 background K⁺ channels?. *Cell*, **2007**, 130, 563-9 56.2 66

355 Silencing acid-sensing ion channel 1a alters cone-mediated retinal function. *Journal of Neuroscience*, **2006**, 26, 5800-9 6.6 65

354 Species diversity and peptide toxins blocking selectivity of ether-a-go-go-related gene subfamily K⁺ channels in the central nervous system. *Molecular Pharmacology*, **2006**, 69, 1673-83 4.3 47

353 Regulation of sensory neuron-specific acid-sensing ion channel 3 by the adaptor protein Na⁺/H⁺ exchanger regulatory factor-1. *Journal of Biological Chemistry*, **2006**, 281, 1796-807 5.4 32

352 Membrane potential-regulated transcription of the resting K⁺ conductance TASK-3 via the calcineurin pathway. *Journal of Biological Chemistry*, **2006**, 281, 28910-8 5.4 26

351 Neurotoxicity and other pharmacological activities of the snake venom phospholipase A2 OS2: the N-terminal region is more important than enzymatic activity. *Biochemistry*, **2006**, 45, 5800-16 3.2 54

350 The background K(+) channel TASK-3 is regulated at both the transcriptional and post-transcriptional levels. *Biochemical and Biophysical Research Communications*, **2006**, 348, 1350-7 3.4 6

349 Four novel tarantula toxins as selective modulators of voltage-gated sodium channel subtypes. *Molecular Pharmacology*, **2006**, 69, 419-29 4.3 132

348 FMRFamide-gated sodium channel and ASIC channels: a new class of ionotropic receptors for FMRFamide and related peptides. *Peptides*, **2006**, 27, 1138-52 3.8 98

347	Deletion of the background potassium channel TREK-1 results in a depression-resistant phenotype. <i>Nature Neuroscience</i> , 2006 , 9, 1134-41	25.5	295
346	TREK-1, a K ⁺ channel involved in polymodal pain perception. <i>EMBO Journal</i> , 2006 , 25, 2368-76	13	323
345	AKAP150, a switch to convert mechano-, pH- and arachidonic acid-sensitive TREK K(+) channels into open leak channels. <i>EMBO Journal</i> , 2006 , 25, 5864-72	13	88
344	The receptor site of the spider toxin PcTx1 on the proton-gated cation channel ASIC1a. <i>Journal of Physiology</i> , 2006 , 570, 339-54	3.9	75
343	Lysophosphatidic acid-operated K ⁺ channels. <i>Journal of Biological Chemistry</i> , 2005 , 280, 4415-21	5.4	77
342	Solution structure of APETx2, a specific peptide inhibitor of ASIC3 proton-gated channels. <i>Protein Science</i> , 2005 , 14, 2003-10	6.3	54
341	A phospholipid sensor controls mechanogating of the K ⁺ channel TREK-1. <i>EMBO Journal</i> , 2005 , 24, 44-53	13	192
340	Cross-talk between the mechano-gated K2P channel TREK-1 and the actin cytoskeleton. <i>EMBO Reports</i> , 2005 , 6, 642-8	6.5	110
339	Solution structure of APETx1 from the sea anemone <i>Anthopleura elegantissima</i> : a new fold for an HERG toxin. <i>Proteins: Structure, Function and Bioinformatics</i> , 2005 , 59, 380-6	4.2	29
338	ASIC2b-dependent regulation of ASIC3, an essential acid-sensing ion channel subunit in sensory neurons via the partner protein PICK-1. <i>Journal of Biological Chemistry</i> , 2004 , 279, 19531-9	5.4	84
337	Acid-sensing ion channel 2 is important for retinal function and protects against light-induced retinal degeneration. <i>Journal of Neuroscience</i> , 2004 , 24, 1005-12	6.6	92
336	A new sea anemone peptide, APETx2, inhibits ASIC3, a major acid-sensitive channel in sensory neurons. <i>EMBO Journal</i> , 2004 , 23, 1516-25	13	296
335	ARF6-dependent interaction of the TWIK1 K ⁺ channel with EFA6, a GDP/GTP exchange factor for ARF6. <i>EMBO Reports</i> , 2004 , 5, 1171-5	6.5	61
334	Knockout of the ASIC2 channel in mice does not impair cutaneous mechanosensation, visceral mechanonociception and hearing. <i>Journal of Physiology</i> , 2004 , 558, 659-69	3.9	97
333	The 2P-domain K ⁺ channels: role in apoptosis and tumorigenesis. <i>Pflügers Archiv European Journal of Physiology</i> , 2004 , 448, 261-73	4.6	117
332	Solution structure of Phrixotoxin 1, a specific peptide inhibitor of Kv4 potassium channels from the venom of the theraphosid spider <i>Phrixotrichus auratus</i> . <i>Protein Science</i> , 2004 , 13, 1197-208	6.3	43
331	How nerve growth factor drives physiological and inflammatory expressions of acid-sensing ion channel 3 in sensory neurons. <i>Journal of Biological Chemistry</i> , 2003 , 278, 48907-13	5.4	117
330	Mechanisms underlying excitatory effects of group I metabotropic glutamate receptors via inhibition of 2P domain K ⁺ channels. <i>EMBO Journal</i> , 2003 , 22, 5403-11	13	161

329	Recombinant production and solution structure of PcTx1, the specific peptide inhibitor of ASIC1a proton-gated cation channels. <i>Protein Science</i> , 2003 , 12, 1332-43	6.3	92
328	Linolenic acid prevents neuronal cell death and paraplegia after transient spinal cord ischemia in rats. <i>Journal of Vascular Surgery</i> , 2003 , 38, 564-75	3.5	90
327	Novel mammalian group XII secreted phospholipase A2 lacking enzymatic activity. <i>Biochemistry</i> , 2003 , 42, 11494-503	3.2	93
326	APETx1, a new toxin from the sea anemone <i>Anthopleura elegantissima</i> , blocks voltage-gated human ether-a-go-go-related gene potassium channels. <i>Molecular Pharmacology</i> , 2003 , 64, 59-69	4.3	102
325	K ⁺ -dependent cerebellar granule neuron apoptosis. Role of task leak K ⁺ channels. <i>Journal of Biological Chemistry</i> , 2003 , 278, 32068-76	5.4	157
324	Proinflammatory mediators, stimulators of sensory neuron excitability via the expression of acid-sensing ion channels. <i>Journal of Neuroscience</i> , 2002 , 22, 10662-70	6.6	288
323	Secreted phospholipase A2 potentiates glutamate-induced calcium increase and cell death in primary neuronal cultures. <i>Journal of Neuroscience Research</i> , 2002 , 67, 634-45	4.4	49
322	A potent protective role of lysophospholipids against global cerebral ischemia and glutamate excitotoxicity in neuronal cultures. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2002 , 22, 821-34	7.3	77
321	ASIC-like, proton-activated currents in rat hippocampal neurons. <i>Journal of Physiology</i> , 2002 , 539, 485-94	5.9	174
320	An intracellular proton sensor commands lipid- and mechano-gating of the K ⁽⁺⁾ channel TREK-1. <i>EMBO Journal</i> , 2002 , 21, 2968-76	13	168
319	p11, an annexin II subunit, an auxiliary protein associated with the background K ⁺ channel, TASK-1. <i>EMBO Journal</i> , 2002 , 21, 4439-48	13	114
318	Interfacial kinetic and binding properties of the complete set of human and mouse groups I, II, V, X, and XII secreted phospholipases A2. <i>Journal of Biological Chemistry</i> , 2002 , 277, 48535-49	5.4	272
317	The multivalent PDZ domain-containing protein CIPP is a partner of acid-sensing ion channel 3 in sensory neurons. <i>Journal of Biological Chemistry</i> , 2002 , 277, 16655-61	5.4	61
316	Protein kinase C stimulates the acid-sensing ion channel ASIC2a via the PDZ domain-containing protein PICK1. <i>Journal of Biological Chemistry</i> , 2002 , 277, 50463-8	5.4	96
315	Novel tarantula toxins for subtypes of voltage-dependent potassium channels in the Kv2 and Kv4 subfamilies. <i>Molecular Pharmacology</i> , 2002 , 62, 48-57	4.3	153
314	Molecular basis of the voltage-dependent gating of TREK-1, a mechano-sensitive K ⁽⁺⁾ channel. <i>Biochemical and Biophysical Research Communications</i> , 2002 , 292, 339-46	3.4	77
313	Nonsteroid anti-inflammatory drugs inhibit both the activity and the inflammation-induced expression of acid-sensing ion channels in nociceptors. <i>Journal of Neuroscience</i> , 2001 , 21, 8026-33	6.6	420
312	ATP-sensitive potassium channels (K(ATP)) in retina: a key role for delayed ischemic tolerance. <i>Brain Research</i> , 2001 , 890, 118-29	3.7	67

311	Lipid and mechano-gated 2P domain K(+) channels. <i>Current Opinion in Cell Biology</i> , 2001 , 13, 422-8	9	250
310	Zn ²⁺ and H ⁺ are coactivators of acid-sensing ion channels. <i>Journal of Biological Chemistry</i> , 2001 , 276, 35361-7	5.4	158
309	A TREK-1-like potassium channel in atrial cells inhibited by beta-adrenergic stimulation and activated by volatile anesthetics. <i>Circulation Research</i> , 2001 , 89, 336-42	15.7	122
308	Genomic and functional characteristics of novel human pancreatic 2P domain K(+) channels. <i>Biochemical and Biophysical Research Communications</i> , 2001 , 282, 249-56	3.4	144
307	Activation of the nuclear factor-kappaB is a key event in brain tolerance. <i>Journal of Neuroscience</i> , 2001 , 21, 4668-77	6.6	244
306	On the Functional Diversity of Secreted Phospholipases A2: Cloning of Novel Mammalian Enzymes and HIV-1 Antiviral Properties. <i>Medical Science Symposia Series</i> , 2001 , 81-84		
305	Axonal transport of TREK and TRAAK potassium channels in rat sciatic nerves. <i>NeuroReport</i> , 2000 , 11, 927-30	1.7	23
304	The bee venom peptide tertiapin underlines the role of I(KACh) in acetylcholine-induced atrioventricular blocks. <i>British Journal of Pharmacology</i> , 2000 , 131, 569-77	8.6	63
303	Molecular and functional properties of two-pore-domain potassium channels. <i>American Journal of Physiology - Renal Physiology</i> , 2000 , 279, F793-801	4.3	450
302	Human TREK2, a 2P domain mechano-sensitive K ⁺ channel with multiple regulations by polyunsaturated fatty acids, lysophospholipids, and Gs, Gi, and Gq protein-coupled receptors. <i>Journal of Biological Chemistry</i> , 2000 , 275, 28398-405	5.4	246
301	Isolation of a tarantula toxin specific for a class of proton-gated Na ⁺ channels. <i>Journal of Biological Chemistry</i> , 2000 , 275, 25116-21	5.4	356
300	Cloning and recombinant expression of a structurally novel human secreted phospholipase A2. <i>Journal of Biological Chemistry</i> , 2000 , 275, 39823-6	5.4	130
299	Lysophospholipids open the two-pore domain mechano-gated K(+) channels TREK-1 and TRAAK. <i>Journal of Biological Chemistry</i> , 2000 , 275, 10128-33	5.4	279
298	TWIK-2, an inactivating 2P domain K ⁺ channel. <i>Journal of Biological Chemistry</i> , 2000 , 275, 28722-30	5.4	92
297	Ischemic spinal cord injury induced by aortic cross-clamping: prevention by riluzole. <i>European Journal of Cardio-thoracic Surgery</i> , 2000 , 18, 174-81	3	53
296	Cloning and recombinant expression of human group IIF-secreted phospholipase A(2). <i>Biochemical and Biophysical Research Communications</i> , 2000 , 279, 223-8	3.4	58
295	Cloning and expression of human TRAAK, a polyunsaturated fatty acids-activated and mechano-sensitive K(+) channel. <i>FEBS Letters</i> , 2000 , 471, 137-40	3.8	91
294	Molecular cloning, functional expression and chromosomal localization of an amiloride-sensitive Na(+) channel from human small intestine. <i>FEBS Letters</i> , 2000 , 471, 205-10	3.8	63

293	M-type KCNQ2-KCNQ3 potassium channels are modulated by the KCNE2 subunit. <i>FEBS Letters</i> , 2000 , 480, 137-41	3.8	62
292	Novel human secreted phospholipase A(2) with homology to the group III bee venom enzyme. <i>Journal of Biological Chemistry</i> , 2000 , 275, 7492-6	5.4	137
291	Prevention of ischemic spinal cord injury: comparative effects of magnesium sulfate and riluzole. <i>Journal of Vascular Surgery</i> , 2000 , 32, 179-89	3.5	54
290	Cloning of a new mouse two-P domain channel subunit and a human homologue with a unique pore structure. <i>Journal of Biological Chemistry</i> , 1999 , 274, 11751-60	5.4	100
289	Secreted phospholipases A(2), a new class of HIV inhibitors that block virus entry into host cells. <i>Journal of Clinical Investigation</i> , 1999 , 104, 611-8	15.9	104
288	On the diversity of secreted phospholipases A(2). Cloning, tissue distribution, and functional expression of two novel mouse group II enzymes. <i>Journal of Biological Chemistry</i> , 1999 , 274, 31195-202	5.4	147
287	Cloning and recombinant expression of a novel mouse-secreted phospholipase A2. <i>Journal of Biological Chemistry</i> , 1999 , 274, 19152-60	5.4	98
286	Mechano- or acid stimulation, two interactive modes of activation of the TREK-1 potassium channel. <i>Journal of Biological Chemistry</i> , 1999 , 274, 26691-6	5.4	322
285	Both group IB and group IIA secreted phospholipases A2 are natural ligands of the mouse 180-kDa M-type receptor. <i>Journal of Biological Chemistry</i> , 1999 , 274, 7043-51	5.4	114
284	TRAAK is a mammalian neuronal mechano-gated K ⁺ channel. <i>Journal of Biological Chemistry</i> , 1999 , 274, 1381-7	5.4	269
283	The pre-transmembrane 1 domain of acid-sensing ion channels participates in the ion pore. <i>Journal of Biological Chemistry</i> , 1999 , 274, 10129-32	5.4	70
282	Cloning and functional expression of a novel degenerin-like Na ⁺ channel gene in mammals. <i>Journal of Physiology</i> , 1999 , 519 Pt 2, 323-33	3.9	79
281	H ⁽⁺⁾ -gated cation channels. <i>Annals of the New York Academy of Sciences</i> , 1999 , 868, 67-76	6.5	181
280	Kv2.1/Kv9.3, an ATP-dependent delayed-rectifier K ⁺ channel in pulmonary artery myocytes. <i>Annals of the New York Academy of Sciences</i> , 1999 , 868, 438-41	6.5	15
279	Inhalational anesthetics activate two-pore-domain background K ⁺ channels. <i>Nature Neuroscience</i> , 1999 , 2, 422-6	25.5	545
278	Effects of phrixotoxins on the Kv4 family of potassium channels and implications for the role of Ito1 in cardiac electrogenesis. <i>British Journal of Pharmacology</i> , 1999 , 126, 251-63	8.6	126
277	Mutually protective actions of kainic acid epileptic preconditioning and sublethal global ischemia on hippocampal neuronal death: involvement of adenosine A1 receptors and K(ATP) channels. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 1999 , 19, 1296-308	7.3	114
276	Reply:. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 1999 , 118, 1157	1.5	2

275	Riluzole prevents ischemic spinal cord injury caused by aortic crossclamping. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 1999 , 117, 881-9	1.5	65
274	Secretory phospholipase A2 potentiates glutamate-induced rat striatal neuronal cell death in vivo. <i>Neuroscience Letters</i> , 1999 , 274, 167-70	3.3	33
273	MIT(1), a black mamba toxin with a new and highly potent activity on intestinal contraction. <i>FEBS Letters</i> , 1999 , 461, 183-8	3.8	72
272	Chapter 12 Potassium Channels with Two P Domains. <i>Current Topics in Membranes</i> , 1999 , 46, 199-222	2.2	11
271	Chapter 5 Isk: A Novel Type of Potassium Channel Regulatory Subunit. <i>Current Topics in Membranes</i> , 1999 , 46, 67-84	2.2	2
270	IKs, a slow and intriguing cardiac K ⁺ channel and its associated long QT diseases. <i>Trends in Cardiovascular Medicine</i> , 1998 , 8, 207-14	6.9	17
269	H ⁽⁺⁾ -gated cation channels: neuronal acid sensors in the Na ^C /DEG family of ion channels. <i>Current Opinion in Neurobiology</i> , 1998 , 8, 418-24	7.6	458
268	Identification, functional expression and chromosomal localisation of a sustained human proton-gated cation channel. <i>FEBS Letters</i> , 1998 , 433, 257-60	3.8	101
267	The KCNQ2 potassium channel: splice variants, functional and developmental expression. Brain localization and comparison with KCNQ3. <i>FEBS Letters</i> , 1998 , 438, 171-6	3.8	111
266	Sea anemone peptides with a specific blocking activity against the fast inactivating potassium channel Kv3.4. <i>Journal of Biological Chemistry</i> , 1998 , 273, 6744-9	5.4	149
265	A new member of the amiloride-sensitive sodium channel family in <i>Drosophila melanogaster</i> peripheral nervous system. <i>Biochemical and Biophysical Research Communications</i> , 1998 , 246, 210-6	3.4	45
264	Mapping of human potassium channel genes TREK-1 (KCNK2) and TASK (KCNK3) to chromosomes 1q41 and 2p23. <i>Genomics</i> , 1998 , 51, 478-9	4.3	21
263	A structural homologue of colipase in black mamba venom revealed by NMR floating disulphide bridge analysis. <i>Journal of Molecular Biology</i> , 1998 , 283, 205-19	6.5	62
262	The Phe-Met-Arg-Phe-amide-activated sodium channel is a tetramer. <i>Journal of Biological Chemistry</i> , 1998 , 273, 8317-22	5.4	93
261	Cloning and expression of a novel pH-sensitive two pore domain K ⁺ channel from human kidney. <i>Journal of Biological Chemistry</i> , 1998 , 273, 30863-9	5.4	292
260	dGNaC1, a gonad-specific amiloride-sensitive Na ⁺ channel. <i>Journal of Biological Chemistry</i> , 1998 , 273, 9424-9	5.4	36
259	Mutations causing neurodegeneration in <i>Caenorhabditis elegans</i> drastically alter the pH sensitivity and inactivation of the mammalian H ⁺ -gated Na ⁺ channel MDEG1. <i>Journal of Biological Chemistry</i> , 1998 , 273, 15418-22	5.4	85
258	Genetic analysis of the beta subunit of the epithelial Na ⁺ channel in essential hypertension. <i>Hypertension</i> , 1998 , 32, 129-37	8.5	115

257	Involvement of Isk-associated K ⁺ channel in heart rate control of repolarization in a murine engineered model of Jervell and Lange-Nielsen syndrome. <i>Circulation Research</i> , 1998 , 83, 95-102	15.7	120
256	Disruption of mitochondrial respiration inhibits volume-regulated anion channels and provokes neuronal cell swelling. <i>Journal of Neuroscience</i> , 1998 , 18, 3117-23	6.6	66
255	Modes of regulation of shab K ⁺ channel activity by the Kv8.1 subunit. <i>Journal of Biological Chemistry</i> , 1997 , 272, 8774-80	5.4	72
254	Cloning, chromosomal mapping, and expression of a novel human secretory phospholipase A2. <i>Journal of Biological Chemistry</i> , 1997 , 272, 15745-52	5.4	215
253	New modulatory alpha subunits for mammalian Shab K ⁺ channels. <i>Journal of Biological Chemistry</i> , 1997 , 272, 24371-9	5.4	164
252	Localization of structural elements of bee venom phospholipase A2 involved in N-type receptor binding and neurotoxicity. <i>Journal of Biological Chemistry</i> , 1997 , 272, 7173-81	5.4	82
251	A modulatory subunit of acid sensing ion channels in brain and dorsal root ganglion cells. <i>Journal of Biological Chemistry</i> , 1997 , 272, 29778-83	5.4	411
250	Molecular mechanism and functional significance of the MinK control of the KvLQT1 channel activity. <i>Journal of Biological Chemistry</i> , 1997 , 272, 16713-6	5.4	103
249	Genotype-phenotype analysis of a newly discovered family with Liddle's syndrome. <i>Journal of Hypertension</i> , 1997 , 15, 1091-100	1.9	71
248	The acid-sensitive ionic channel subunit ASIC and the mammalian degenerin MDEG form a heteromultimeric H ⁺ -gated Na ⁺ channel with novel properties. <i>Journal of Biological Chemistry</i> , 1997 , 272, 28819-22	5.4	182
247	Molecular cloning of a non-inactivating proton-gated Na ⁺ channel specific for sensory neurons. <i>Journal of Biological Chemistry</i> , 1997 , 272, 20975-8	5.4	440
246	The structure, function and distribution of the mouse TWIK-1 K ⁺ channel. <i>FEBS Letters</i> , 1997 , 402, 28-32	3.8	99
245	A proton-gated cation channel involved in acid-sensing. <i>Nature</i> , 1997 , 386, 173-7	50.4	1111
244	The amiloride-sensitive Na ⁺ channel: from primary structure to function. <i>Comparative Biochemistry and Physiology A, Comparative Physiology</i> , 1997 , 118, 193-200		23
243	Comparative expression of the inward rectifier K ⁺ channel GIRK2 in the cerebellum of normal and weaver mutant mice. <i>Brain Research</i> , 1997 , 753, 8-17	3.7	35
242	The potassium channel opener (-)-cromakalim prevents glutamate-induced cell death in hippocampal neurons. <i>Journal of Neurochemistry</i> , 1997 , 69, 1570-9	6	66
241	Endocytic properties of the M-type 180-kDa receptor for secretory phospholipases A2. <i>Journal of Biological Chemistry</i> , 1996 , 271, 250-7	5.4	99
240	Dominant negative chimeras provide evidence for homo and heteromultimeric assembly of inward rectifier K ⁺ channel proteins via their N-terminal end. <i>FEBS Letters</i> , 1996 , 378, 64-8	3.8	36

239	Inner ear defects induced by null mutation of the isk gene. <i>Neuron</i> , 1996 , 17, 1251-64	13.9	349
238	A pH-sensitive yeast outward rectifier K ⁺ channel with two pore domains and novel gating properties. <i>Journal of Biological Chemistry</i> , 1996 , 271, 4183-7	5.4	85
237	Effects on behaviour and EEG of single chain phospholipases A2 from snake and bee venoms injected into rat brain: search for a functional antagonism. <i>Basic and Clinical Pharmacology and Toxicology</i> , 1996 , 78, 341-7		16
236	K(V)LQT1 and Isk (minK) proteins associate to form the I(Ks) cardiac potassium current. <i>Nature</i> , 1996 , 384, 78-80	50.4	1389
235	A new K ⁺ channel beta subunit to specifically enhance Kv2.2 (CDRK) expression. <i>Journal of Biological Chemistry</i> , 1996 , 271, 26341-8	5.4	85
234	The mammalian degenerin MDEG, an amiloride-sensitive cation channel activated by mutations causing neurodegeneration in <i>Caenorhabditis elegans</i> . <i>Journal of Biological Chemistry</i> , 1996 , 271, 10433-8	5.4	265
233	Structure and regulation of the amiloride-sensitive epithelial sodium channel. <i>Ion Channels</i> , 1996 , 4, 115-67		22
232	Localization and regulation by steroids of the alpha, beta and gamma subunits of the amiloride-sensitive Na ⁺ channel in colon, lung and kidney. <i>Pflugers Archiv European Journal of Physiology</i> , 1995 , 430, 299-307	4.6	188
231	Biophysical, pharmacological and developmental properties of ATP-sensitive K ⁺ channels in cultured myotomal muscle cells from <i>Xenopus</i> embryos. <i>Pflugers Archiv European Journal of Physiology</i> , 1995 , 429, 607-16	4.6	
230	Cloning of the amiloride-sensitive FMRFamide peptide-gated sodium channel. <i>Nature</i> , 1995 , 378, 730-3	50.4	355
229	Molecular properties of neuronal G-protein-activated inwardly rectifying K ⁺ channels. <i>Journal of Biological Chemistry</i> , 1995 , 270, 28660-7	5.4	214
228	Structural elements of secretory phospholipases A2 involved in the binding to M-type receptors. <i>Journal of Biological Chemistry</i> , 1995 , 270, 5534-40	5.4	101
227	Functional degenerin-containing chimeras identify residues essential for amiloride-sensitive Na ⁺ channel function. <i>Journal of Biological Chemistry</i> , 1995 , 270, 11735-7	5.4	102
226	Identification of the binding domain for secretory phospholipases A2 on their M-type 180-kDa membrane receptor. <i>Journal of Biological Chemistry</i> , 1995 , 270, 28869-73	5.4	56
225	The human 180-kDa receptor for secretory phospholipases A2. Molecular cloning, identification of a secreted soluble form, expression, and chromosomal localization. <i>Journal of Biological Chemistry</i> , 1995 , 270, 8963-70	5.4	152
224	Molecular cloning and functional expression of a novel amiloride-sensitive Na ⁺ channel. <i>Journal of Biological Chemistry</i> , 1995 , 270, 27411-4	5.4	233
223	Multifunctional activity of the extracellular domain of the M-type (180 kDa) membrane receptor for secretory phospholipases A2. <i>Biochemistry</i> , 1995 , 34, 13146-51	3.2	81
222	Coexistence of two classes of glibenclamide-inhibitable ATP-regulated K ⁺ channels in avian skeletal muscle. <i>Pflugers Archiv European Journal of Physiology</i> , 1995 , 431, 117-24	4.6	4

221	Kalicludines and kaliseptine. Two different classes of sea anemone toxins for voltage sensitive K ⁺ channels. <i>Journal of Biological Chemistry</i> , 1995 , 270, 25121-6	5.4	159
220	Expression cloning in K ⁺ transport defective yeast and distribution of HBP1, a new putative HMG transcriptional regulator. <i>Nucleic Acids Research</i> , 1994 , 22, 3685-8	20.1	42
219	Identification of the Ca ²⁺ current activated by vasoconstrictors in vascular smooth muscle cells. <i>Pflugers Archiv European Journal of Physiology</i> , 1994 , 429, 1-6	4.6	18
218	CGRP-induced activation of KATP channels in follicular <i>Xenopus</i> oocytes. <i>Pflugers Archiv European Journal of Physiology</i> , 1994 , 428, 604-9	4.6	16
217	Glutamate-induced overexpression of NMDA receptor messenger RNAs and protein triggered by activation of AMPA/kainate receptors in rat hippocampus following forebrain ischemia. <i>Brain Research</i> , 1994 , 659, 67-74	3.7	57
216	Expression of group II phospholipase A2 in rat brain after severe forebrain ischemia and in endotoxic shock. <i>Brain Research</i> , 1994 , 651, 353-6	3.7	96
215	Cloning provides evidence for a family of inward rectifier and G-protein coupled K ⁺ channels in the brain. <i>FEBS Letters</i> , 1994 , 353, 37-42	3.8	252
214	Molecular cloning of a murine N-type calcium channel alpha 1 subunit. Evidence for isoforms, brain distribution, and chromosomal localization. <i>FEBS Letters</i> , 1994 , 338, 1-5	3.8	37
213	Effects of two chemically related new Ca ²⁺ channel antagonists, SR33557 (fantofarone) and SR33805, on the L-type cardiac channel. <i>European Journal of Pharmacology</i> , 1994 , 263, 101-5	5.3	9
212	Pharmacological properties of ATP-sensitive K ⁺ channels in mammalian skeletal muscle cells. <i>European Journal of Pharmacology</i> , 1993 , 236, 419-26	5.3	38
211	Localization of a potassium channel gene (KCNE1) to 21q22.1-q22.2 by in situ hybridization and somatic cell hybridization. <i>Genomics</i> , 1993 , 15, 243-5	4.3	25
210	Memory processing and apamin induce immediate early gene expression in mouse brain. <i>Molecular Brain Research</i> , 1993 , 18, 17-22		60
209	Endothelin and vasopressin activate low conductance chloride channels in aortic smooth muscle cells. <i>Pflugers Archiv European Journal of Physiology</i> , 1993 , 425, 156-63	4.6	58
208	Single-channel properties and regulation of pinacidil/glibenclamide-sensitive K ⁺ channels in follicular cells from <i>Xenopus</i> oocyte. <i>Pflugers Archiv European Journal of Physiology</i> , 1993 , 424, 113-21	4.6	33
207	The protein IsK is a dual activator of K ⁺ and Cl ⁻ channels. <i>Nature</i> , 1993 , 365, 850-2	50.4	131
206	Molecular cloning and functional expression of different molecular forms of rat amiloride-binding proteins. <i>FEBS Journal</i> , 1993 , 216, 679-87		43
205	Expression cloning of an epithelial amiloride-sensitive Na ⁺ channel. A new channel type with homologies to <i>Caenorhabditis elegans</i> degenerins. <i>FEBS Letters</i> , 1993 , 318, 95-9	3.8	295
204	Characterization of the Sulfonylurea-Sensitive ATP-Modulated Potassium Channel 1993 , 61-75		3

203	Purification, affinity labeling, and reconstitution of voltage-sensitive potassium channels. <i>Methods in Enzymology</i> , 1992 , 207, 556-64	1.7	4
202	Receptor-mediated regulation of Isk, a very slowly activating, voltage-dependent K ⁺ channel in <i>Xenopus</i> oocytes. <i>Biochemical and Biophysical Research Communications</i> , 1992 , 184, 1135-41	3.4	19
201	Scyllatoxin, a blocker of Ca(2+)-activated K ⁺ channels: structure-function relationships and brain localization of the binding sites. <i>Biochemistry</i> , 1992 , 31, 648-54	3.2	58
200	ATP/ADP binding sites are present in the sulfonylurea binding protein associated with brain ATP-sensitive K ⁺ channels. <i>Biochemistry</i> , 1992 , 31, 6328-32	3.2	40
199	Effects of the level of mRNA expression on biophysical properties, sensitivity to neurotoxins, and regulation of the brain delayed-rectifier K ⁺ channels Kv1.2. <i>Biochemistry</i> , 1992 , 31, 12463-8	3.2	65
198	Effectors of ATP-sensitive K ⁺ channels inhibit the regulatory effects of somatostatin and GH-releasing factor on growth hormone secretion. <i>Biochemical and Biophysical Research Communications</i> , 1992 , 187, 1007-14	3.4	17
197	Characterization and partial purification from pheochromocytoma cells of an endogenous equivalent of scyllatoxin, a scorpion toxin which blocks small conductance Ca(2+)-activated K ⁺ channels. <i>Brain Research</i> , 1992 , 599, 230-6	3.7	10
196	Behavioral effects of modulators of ATP-sensitive K ⁺ channels in the rat dorsal pallidum. <i>European Journal of Pharmacology</i> , 1992 , 217, 71-7	5.3	6
195	ISK, a slowly activating voltage-sensitive K ⁺ channel. Characterization of multiple cDNAs and gene organization in the mouse. <i>FEBS Letters</i> , 1992 , 301, 168-72	3.8	27
194	Developmental expression of voltage-sensitive K ⁺ channels in mouse skeletal muscle and C2C12 cells. <i>FEBS Letters</i> , 1992 , 310, 162-6	3.8	46
193	A small-conductance charybdotoxin-sensitive, apamin-resistant Ca(2+)-activated K ⁺ channel in aortic smooth muscle cells (A7r5 line and primary culture). <i>Pflugers Archiv European Journal of Physiology</i> , 1992 , 420, 417-23	4.6	24
192	Nucleotide diphosphates activate the ATP-sensitive potassium channel in mouse skeletal muscle. <i>Pflugers Archiv European Journal of Physiology</i> , 1992 , 422, 185-92	4.6	42
191	TMB-8 (8-(N,N-diethylamino) octyl-3,4,5-trimethoxybenzoate) inhibits the ATP-sensitive K ⁺ channel. <i>European Journal of Pharmacology</i> , 1992 , 226, 175-7		10
190	Altered chloride ion channel kinetics associated with the delta F508 cystic fibrosis mutation. <i>Nature</i> , 1991 , 354, 526-8	50.4	600
189	Properties of receptors for neurotoxic phospholipases A2 in different tissues. <i>Neurochemical Research</i> , 1991 , 16, 651-8	4.6	44
188	Indolizinsulphonates. A class of blockers with dual but discriminative effects on L-type Ca ²⁺ channel activity and excitation-contraction coupling in skeletal muscle. <i>Pflugers Archiv European Journal of Physiology</i> , 1991 , 419, 651-6	4.6	21
187	A new non-voltage-dependent, epithelial-like Na ⁺ channel in vascular smooth muscle cells. <i>Pflugers Archiv European Journal of Physiology</i> , 1991 , 419, 401-8	4.6	38
186	A voltage, calcium, and ATP sensitive non selective cation channel in human colonic tumor cells. <i>Biochemical and Biophysical Research Communications</i> , 1991 , 176, 1196-203	3.4	24

185	Activation by cromakalim of pre- and post-synaptic ATP-sensitive K ⁺ channels in substantia nigra. <i>Biochemical and Biophysical Research Communications</i> , 1991 , 174, 909-14	3.4	65
184	Two different types of channels are targets for potassium channel openers in <i>Xenopus oocytes</i> . <i>FEBS Letters</i> , 1991 , 287, 75-9	3.8	19
183	Identification of different receptor types for toxic phospholipases A2 in rabbit skeletal muscle. <i>FEBS Letters</i> , 1991 , 293, 29-33	3.8	35
182	Riluzole prevents hyperexcitability produced by the mast cell degranulating peptide and dendrotoxin I in the rat. <i>European Journal of Pharmacology</i> , 1991 , 193, 223-9	5.3	26
181	Anticonvulsant and sleep-waking influences of riluzole in a rat model of absence epilepsy. <i>European Journal of Pharmacology</i> , 1991 , 199, 371-3	5.3	28
180	Conditional immortalization of normal and dysgenic mouse muscle cells by the SV40 large T antigen under the vimentin promoter control. <i>Developmental Biology</i> , 1991 , 148, 517-28	3.1	24
179	MCD peptide and dendrotoxin I activate c-fos and c-jun expression by acting on two different types of K ⁺ channels. A discrimination using the K ⁺ channel opener lemakalim. <i>Brain Research</i> , 1991 , 554, 22-9	3.7	11
178	Increase of sodium channels in demyelinated lesions of multiple sclerosis. <i>Brain Research</i> , 1991 , 556, 311-6	3.7	100
177	Specific hippocampal lesions indicate the presence of sulfonylurea binding sites associated to ATP-sensitive K ⁺ channels both post-synaptically and on mossy fibers. <i>Brain Research</i> , 1991 , 540, 340-4	3.7	41
176	K ⁺ efflux pathways and neurotransmitter release associated to hippocampal ischemia: effects of glucose and of K ⁺ channel blockers. <i>Brain Research</i> , 1991 , 539, 155-8	3.7	24
175	Effect of apamin, a toxin that inhibits Ca(2+)-dependent K ⁺ channels, on learning and memory processes. <i>Brain Research</i> , 1991 , 551, 322-6	3.7	104
174	Solution structure of leiurotoxin I by 1H magnetic resonance 1991 , 563-564		
173	Localization of the gene for amiloride binding protein on chromosome 7 and RFLP analysis in cystic fibrosis families. <i>Human Genetics</i> , 1990 , 85, 587-9	6.3	12
172	[3H]phenamil binding protein of the renal epithelium Na ⁺ channel. Purification, affinity labeling, and functional reconstitution. <i>Biochemistry</i> , 1990 , 29, 1039-45	3.2	26
171	Purification and pharmacological characterization of peptide toxins from the black mamba (<i>Dendroaspis polylepis</i>) venom. <i>Toxicon</i> , 1990 , 28, 847-56	2.8	61
170	Mast cell degranulating peptide induces the expression of the c-fos proto-oncogene in hippocampus. <i>European Journal of Pharmacology</i> , 1990 , 180, 179-81	5.3	7
169	Sulfonylurea binding sites associated with ATP-regulated K ⁺ channels in the central nervous system: autoradiographic analysis of their distribution and ontogenesis, and of their localization in mutant mice cerebellum. <i>Brain Research</i> , 1990 , 519, 29-43	3.7	137
168	Brain ischemia alters the density of binding sites for glibenclamide, a specific blocker of ATP-sensitive K ⁺ channels. <i>Brain Research</i> , 1990 , 526, 147-52	3.7	36

167	Saxitoxin-sensitive Na ⁺ channels: presynaptic localization in cerebellum and hippocampus of neurological mutant mice. <i>Brain Research</i> , 1990 , 533, 196-202	3.7	16
166	Solution conformation of leiurotoxin I (scyllatoxin) by 1H nuclear magnetic resonance. Resonance assignment and secondary structure. <i>FEBS Letters</i> , 1990 , 260, 249-53	3.8	45
165	Induction of normal ultrastructure by CGRP treatment in dysgenic myotubes. <i>FEBS Letters</i> , 1990 , 263, 147-52	3.8	17
164	Regional expression of a MCD-peptide and dendrotoxin I-sensitive voltage-dependent potassium channel in rat brain. <i>FEBS Letters</i> , 1990 , 263, 163-5	3.8	18
163	Ca ²⁺ channel blockers inhibit secretory C1-channels in intestinal epithelial cells. <i>Biochemical and Biophysical Research Communications</i> , 1990 , 171, 1022-8	3.4	13
162	Regulation of the ATP-sensitive potassium channel. <i>Ion Channels</i> , 1990 , 2, 205-22		21
161	Phenotypic and functional reversion of muscular dysgenesis by heterotypic fibroblast-myotube fusion in vitro. <i>Advances in Experimental Medicine and Biology</i> , 1990 , 280, 139-46	3.6	2
160	Pyrethroid receptor in the insect sodium channel: alteration of its properties in pyrethroid-resistant flies. <i>Biochemistry</i> , 1989 , 28, 1673-1677	3.2	85
159	Identification and properties of a novel type of Na ⁺ -permeable amiloride-sensitive channel in thyroid cells. <i>FEBS Journal</i> , 1989 , 183, 499-505		21
158	Dendrotoxin-binding brain membrane protein displays a K ⁺ channel activity that is stimulated by both cAMP-dependent and endogenous phosphorylations. <i>Biochemistry</i> , 1989 , 28, 6455-60	3.2	89
157	Charybdotoxin is a new member of the K ⁺ channel toxin family that includes dendrotoxin I and mast cell degranulating peptide. <i>Biochemistry</i> , 1989 , 28, 9708-14	3.2	51
156	Ca ²⁺ channel blockers prevent seizures induced by a class of K ⁺ channel inhibitors. <i>European Journal of Pharmacology</i> , 1989 , 160, 173-7	5.3	25
155	K ⁺ channel openers decrease seizures in genetically epileptic rats. <i>European Journal of Pharmacology</i> , 1989 , 167, 181-3	5.3	54
154	Polypeptide constitution of receptors for apamin, a neurotoxin which blocks a class of Ca ²⁺ -activated K ⁺ channels. <i>FEBS Letters</i> , 1989 , 248, 150-154	3.8	15
153	The inotropic effect of endothelin-1 on rat atria involves hydrolysis of phosphatidylinositol. <i>FEBS Letters</i> , 1989 , 249, 143-6	3.8	92
152	Charybdotoxin blocks dendrotoxin-sensitive voltage-activated K ⁺ channels. <i>FEBS Letters</i> , 1989 , 250, 519-22	3.8	52
151	Analogies and differences in the mode of action and properties of binding sites (localization and mutual interactions) of two K ⁺ channel toxins, MCD peptide and dendrotoxin I. <i>Brain Research</i> , 1989 , 495, 45-57	3.7	47
150	Subtypes of K ⁺ channels differentiated by the effect of K ⁺ channel openers upon K ⁺ channel blocker-induced seizures. <i>Brain Research</i> , 1989 , 495, 189-92	3.7	42

149	Antidiabetic sulfonylureas: localization of binding sites in the brain and effects on the hyperpolarization induced by anoxia in hippocampal slices. <i>Brain Research</i> , 1989 , 486, 159-64	3-7	216
148	K ⁺ channels openers prevent epilepsy induced by the bee venom peptide MCD. <i>European Journal of Pharmacology</i> , 1989 , 159, 329-30	5-3	52
147	Monoclonal Antibodies That Coimmunoprecipitate the 1,4-Dihydropyridine and Phenylalkylamine Receptors and Reveal the Ca ²⁺ Channel Structure 1989 , 47-56		
146	The secondary structure of the toxin ATX Ia from <i>Anemonia sulcata</i> in aqueous solution determined on the basis of complete sequence-specific ¹ H-NMR assignments. <i>FEBS Journal</i> , 1988 , 171, 177-92		24
145	The Na ⁺ /K ⁺ /Cl ⁻ cotransport in C6 glioma cells. Properties and role in volume regulation. <i>FEBS Journal</i> , 1988 , 171, 425-33		36
144	The regulation of the intracellular pH in cells from vertebrates. <i>FEBS Journal</i> , 1988 , 174, 3-14		263
143	The regulation of the cytoplasmic free Ca ²⁺ concentration in aortic smooth muscle cells (A7r5 line) after stimulation by vasopressin and bombesin. <i>FEBS Journal</i> , 1988 , 176, 47-52		21
142	Calcium channels: molecular pharmacology, structure and regulation. <i>Journal of Membrane Biology</i> , 1988 , 104, 81-105	2-3	333
141	Calcium channel inhibitors that bind to plant cell membranes block calcium entry into protoplasts. <i>Biochemistry</i> , 1988 , 27, 764-768	3-2	118
140	The receptor site for the bee venom mast cell degranulating peptide. Affinity labeling and evidence for a common molecular target for mast cell degranulating peptide and dendrotoxin I, a snake toxin active on K ⁺ channels. <i>Biochemistry</i> , 1988 , 27, 1827-32	3-2	65
139	Transmembrane ionic transport systems and hypertension. <i>American Journal of Medicine</i> , 1988 , 84, 3-9	2-4	5
138	Properties of structure and interaction of the receptor for omega-conotoxin, a polypeptide active on Ca ²⁺ channels. <i>Biochemical and Biophysical Research Communications</i> , 1988 , 150, 1051-62	3-4	80
137	Transmembrane ionic transport systems and hypertension. <i>American Journal of Medicine</i> , 1988 , 84, 3-9	2-4	124
136	Interaction of insecticides of the pyrethroid family with specific binding sites on the voltage-dependent sodium channel from mammalian brain. <i>Brain Research</i> , 1988 , 459, 44-53	3-7	80
135	Cerebral glucose utilization after administration of apamin, a toxin active on Ca ²⁺ -dependent K ⁺ channels. <i>Brain Research</i> , 1988 , 451, 274-84	3-7	265
134	Distribution of voltage-dependent Na ⁺ channels identified by high-affinity receptors for tetrodotoxin and saxitoxin in rat and human brains: quantitative autoradiographic analysis. <i>Brain Research</i> , 1988 , 448, 128-39	3-7	21
133	High affinity receptors for the bee venom MCD peptide. Quantitative autoradiographic localization at different stages of brain development and relationship with MCD neurotoxicity. <i>Brain Research</i> , 1988 , 446, 106-12	3-7	28
132	Somatostatin activates glibenclamide-sensitive and ATP-regulated K ⁺ channels in insulinoma cells via a G-protein. <i>FEBS Letters</i> , 1988 , 242, 94-6	3-8	47

131	Molecular properties of structure and regulation of the calcium channel. <i>Annals of the New York Academy of Sciences</i> , 1988 , 522, 134-49	6.5	7
130	Targets for calcium channel blockers in mammalian skeletal muscle and their respective functions in excitation-contraction coupling. <i>Biochemical and Biophysical Research Communications</i> , 1988 , 156, 1324-32	3.4	28
129	Molecular mechanism of action of the vasoconstrictor peptide endothelin. <i>Biochemical and Biophysical Research Communications</i> , 1988 , 157, 977-85	3.4	306
128	Existence of different populations of the dendrotoxin I binding protein associated with neuronal K ⁺ channels. <i>Biochemical and Biophysical Research Communications</i> , 1988 , 153, 231-40	3.4	44
127	Amiloride and its analogs as tools to inhibit Na ⁺ transport via the Na ⁺ channel, the Na ⁺ /H ⁺ antiport and the Na ⁺ /Ca ²⁺ exchanger. <i>Biochimie</i> , 1988 , 70, 1285-90	4.6	84
126	Affinity labeling of the digitalis-binding site. <i>Methods in Enzymology</i> , 1988 , 156, 323-33	1.7	7
125	Receptors for diphenylbutylpiperidine neuroleptics in brain, cardiac, and smooth muscle membranes. Relationship with receptors for 1,4-dihydropyridines and phenylalkylamines and with Ca ²⁺ channel blockade. <i>European Journal of Pharmacology</i> , 1987 , 141, 261-8	5.3	36
124	The brain response to the bee venom peptide MCD. Activation and desensitization of a hippocampal target. <i>Brain Research</i> , 1987 , 418, 235-44	3.7	46
123	Autoradiographic analysis in rat brain of the postnatal ontogeny of voltage-dependent Na ⁺ channels, Ca ²⁺ -dependent K ⁺ channels and slow Ca ²⁺ channels identified as receptors for tetrodotoxin, apamin and (-)-desmethoxyverapamil. <i>Brain Research</i> , 1987 , 417, 21-32	3.7	55
122	Identification in mammalian brain of an endogenous substance with Na ⁺ channel blocking activities similar to those of tetrodotoxin. <i>Brain Research</i> , 1987 , 417, 327-34	3.7	7
121	Ciguatoxin and brevetoxins share a common receptor site on the neuronal voltage-dependent Na ⁺ channel. <i>FEBS Letters</i> , 1987 , 219, 355-9	3.8	258
120	Two potent central convulsant peptides, a bee venom toxin, the MCD peptide, and a snake venom toxin, dendrotoxin I, known to block K ⁺ channels, have interacting receptor sites. <i>Biochemical and Biophysical Research Communications</i> , 1987 , 143, 383-9	3.4	64
119	The antidiabetic sulfonylurea glibenclamide is a potent blocker of the ATP-modulated K ⁺ channel in insulin secreting cells. <i>Biochemical and Biophysical Research Communications</i> , 1987 , 146, 21-5	3.4	112
118	Photoaffinity labelling and phosphorylation of a 165 kilodalton peptide associated with dihydropyridine and phenylalkylamine-sensitive calcium channels. <i>Biochemical and Biophysical Research Communications</i> , 1987 , 147, 1137-45	3.4	85
117	Identification and affinity labeling of very high affinity binding sites for the phenylalkylamine series of Ca ⁺ channel blockers in the Drosophila nervous system. <i>Biochemistry</i> , 1987 , 26, 6311-5	3.2	36
116	Monoclonal antibodies that coimmunoprecipitate the 1,4-dihydropyridine and phenylalkylamine receptors and reveal the Ca ²⁺ channel structure. <i>Biochemistry</i> , 1987 , 26, 5-9	3.2	68
115	Molecular properties of amiloride action and of its Na ⁺ transporting targets. <i>Kidney International</i> , 1987 , 32, 785-93	9.9	91
114	Restoration of dysgenic muscle contraction and calcium channel function by co-culture with normal spinal cord neurons. <i>Nature</i> , 1987 , 330, 563-6	50.4	59

113	The calcium channel antagonists receptor from rabbit skeletal muscle. Reconstitution after purification and subunit characterization. <i>FEBS Journal</i> , 1987 , 164, 525-31		38
112	[3H]nitrendipine receptors as markers of a class of putative voltage-sensitive Ca ²⁺ channels in normal human skeletal muscle and in muscle from Duchenne muscular dystrophy patients. <i>Muscle and Nerve</i> , 1986 , 9, 148-51	3.4	16
111	Expression of apamin receptor in muscles of patients with myotonic muscular dystrophy. <i>Nature</i> , 1986 , 319, 678-80	50.4	113
110	Interaction of guanidinium and guanidinium derivatives with the Na ⁺ /H ⁺ exchange system. <i>FEBS Journal</i> , 1986 , 154, 241-5		38
109	The Na ⁺ /H ⁺ exchange system in glial cell lines. Properties and activation by an hyperosmotic shock. <i>FEBS Journal</i> , 1986 , 160, 211-9		59
108	Purification, subunit structure and pharmacological effects on cardiac and smooth muscle cells of a polypeptide toxin isolated from the marine snail <i>Conus tessulatus</i> . <i>FEBS Journal</i> , 1986 , 161, 787-92		14
107	Biochemical characterization of the Na ⁺ /K ⁺ /Cl ⁻ co-transport in chick cardiac cells. <i>Biochemical and Biophysical Research Communications</i> , 1986 , 134, 326-31	3.4	60
106	[3H]phenamil, a radiolabelled diuretic for the analysis of the amiloride-sensitive Na ⁺ channels in kidney membranes. <i>Biochemical and Biophysical Research Communications</i> , 1986 , 135, 25-32	3.4	32
105	The activity of the Na ⁺ /H ⁺ antiporter in cultured cardiac cells is dependent on the culture conditions used. <i>FEBS Letters</i> , 1986 , 196, 163-6	3.8	25
104	A polypeptide toxin from the coral <i>Goniopora</i> . Purification and action on Ca ²⁺ channels. <i>FEBS Letters</i> , 1986 , 202, 331-6	3.8	15
103	Tetrodotoxin-sensitive and tetrodotoxin-resistant Na ⁺ channels differ in their sensitivity to Cd ²⁺ and Zn ²⁺ . <i>European Journal of Pharmacology</i> , 1986 , 122, 245-50	5.3	73
102	The interaction of polypeptide neurotoxins with tetrodotoxin-resistant Na ⁺ channels in mammalian cardiac cells. Correlation with inotropic and arrhythmic effects. <i>European Journal of Pharmacology</i> , 1986 , 120, 161-70	5.3	33
101	The voltage-dependent sodium channel is co-localized with the acetylcholine receptor at the vertebrate neuromuscular junction. <i>Biochemical and Biophysical Research Communications</i> , 1986 , 139, 196-201	3.4	10
100	Antibodies reveal the cytolocalization and subunit structure of the 1,4-dihydropyridine component of the neuronal Ca ²⁺ channel. <i>Biochemical and Biophysical Research Communications</i> , 1986 , 139, 996-1002	3.4	27
99	[3H]-tetrodotoxin binding in neuronal and non-neuronal spinal cord cultures. <i>Biochemical and Biophysical Research Communications</i> , 1986 , 138, 1250-6	3.4	2
98	Polypeptide toxins as tools to study voltage-sensitive Na ⁺ channels. <i>Annals of the New York Academy of Sciences</i> , 1986 , 479, 204-20	6.5	52
97	Axonal transport of Na ⁺ ,K ⁺ -ATPase identified as a ouabain binding site in rat sciatic nerve. <i>Neuroscience Letters</i> , 1986 , 64, 177-83	3.3	29
96	Immunochemical analysis of subunit structures of 1,4-dihydropyridine receptors associated with voltage-dependent Ca ²⁺ channels in skeletal, cardiac, and smooth muscles. <i>Biochemistry</i> , 1986 , 25, 3492-5	3.2	96

95	The Na ⁺ /H ⁺ antiport of eukaryotic cells: relationship between the kinetic properties of the system and its physiological function. <i>Biochimie</i> , 1986 , 68, 1279-85	4.6	9
94	Quantitative autoradiographic mapping in rat brain of the receptor of apamin, a polypeptide toxin specific for one class of Ca ²⁺ -dependent K ⁺ channels. <i>Brain Research</i> , 1986 , 382, 239-49	3.7	78
93	The role of the Na ⁺ /H ⁺ exchange system in the regulation of the internal pH in cultured cardiac cells. <i>FEBS Journal</i> , 1985 , 149, 1-4		69
92	Abnormal transverse tubule system and abnormal amount of receptors for Ca ²⁺ channel inhibitors of the dihydropyridine family in skeletal muscle from mice with embryonic muscular dysgenesis. <i>Developmental Biology</i> , 1985 , 112, 458-66	3.1	71
91	Axonal transport of the voltage-dependent Na ⁺ channel protein identified by its tetrodotoxin binding site in rat sciatic nerves. <i>Brain Research</i> , 1985 , 345, 153-8	3.7	53
90	Purification, sequence, and pharmacological properties of sea anemone toxins from <i>Radianthus paumotensis</i> . A new class of sea anemone toxins acting on the sodium channel. <i>Biochemistry</i> , 1985 , 24, 3554-61	3.2	68
89	The voltage-dependent Na ⁺ channel of insect nervous system identified by receptor sites for tetrodotoxin, and scorpion and sea anemone toxins. <i>Biochemical and Biophysical Research Communications</i> , 1985 , 131, 1226-33	3.4	43
88	Characterization of the Ca ²⁺ coordination site regulating binding of Ca ²⁺ channel inhibitors d-cis-diltiazem, (+/-)bepridil and (-)desmethoxyverapamil to their receptor site in skeletal muscle transverse tubule membranes. <i>Biochemical and Biophysical Research Communications</i> , 1985 , 132, 49-55	3.4	39
87	Differentiation of receptor sites for [3H]nitrendipine in chick hearts and physiological relation to the slow Ca ²⁺ channel and to excitation-contraction coupling. <i>FEBS Journal</i> , 1984 , 139, 673-81		62
86	Characterization, solubilization, affinity labeling and purification of the cardiac Na ⁺ channel using Tityus toxin gamma. <i>FEBS Journal</i> , 1984 , 141, 651-60		90
85	Solubilization of the nitrendipine receptor from skeletal muscle transverse tubule membranes. Interactions with specific inhibitors of the voltage-dependent Ca ²⁺ channel. <i>FEBS Journal</i> , 1984 , 142, 449-55		58
84	Properties of receptors for the Ca ²⁺ -channel blocker verapamil in transverse-tubule membranes of skeletal muscle. Stereospecificity, effect of Ca ²⁺ and other inorganic cations, evidence for two categories of sites and effect of nucleoside triphosphates. <i>FEBS Journal</i> , 1984 , 144, 211-5		71
83	Tityus gamma toxin, a high affinity effector of the Na ⁺ channel in muscle, with a selectivity for channels in the surface membrane. <i>Pflugers Archiv European Journal of Physiology</i> , 1984 , 400, 22-7	4.6	57
82	The effect of Tityus serrulatus scorpion toxin gamma on Na channels in neuroblastoma cells. <i>Pflugers Archiv European Journal of Physiology</i> , 1984 , 401, 297-303	4.6	55
81	Different functional states of tetrodotoxin sensitive and tetrodotoxin resistant Na ⁺ channels occur during the in vitro development of rat skeletal muscle. <i>Pflugers Archiv European Journal of Physiology</i> , 1984 , 402, 121-8	4.6	61
80	Activation of the voltage-dependent Ca ²⁺ channel in rat heart cells by dihydropyridine derivatives. <i>Biochemical and Biophysical Research Communications</i> , 1984 , 125, 405-12	3.4	42
79	Molecular properties of the apamin-binding component of the Ca ²⁺ -dependent K ⁺ channel. Radiation-inactivation, affinity labelling and solubilization. <i>FEBS Journal</i> , 1984 , 142, 1-6		37
78	Autoradiographic localization of tetrodotoxin-sensitive Na ⁺ channels in rat brain. <i>Neuroscience Letters</i> , 1984 , 52, 31-5	3.3	13

77	Comparative changes of levels of nitrendipine Ca ²⁺ channels, of tetrodotoxin-sensitive Na ⁺ channels and of ouabain-sensitive (Na ⁺ + K ⁺)-ATPase following denervation of rat and chick skeletal muscle. <i>FEBS Letters</i> , 1984 , 172, 114-8	3.8	30
76	Intracellular pH measurements using the fluorescence of 9-aminoacridine. <i>FEBS Letters</i> , 1984 , 172, 275-278	3.8	9
75	Autoradiographic localization of apamin-sensitive Ca ²⁺ -dependent K ⁺ channels in rat brain. <i>European Journal of Pharmacology</i> , 1984 , 100, 135-6	5.3	17
74	A search for an Ouabain-like substance from the electric organ of <i>Electrophorus electricus</i> which led to arachidonic acid and related fatty acids. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1984 , 769, 245-52	3.8	67
73	[³ H] verapamil binding sites in skeletal muscle transverse tubule membranes. <i>Biochemical and Biophysical Research Communications</i> , 1984 , 118, 239-45	3.4	63
72	Purification of the dihydropyridine receptor of the voltage-dependent Ca ²⁺ channel from skeletal muscle transverse tubules using (+) [³ H]PN 200-110. <i>Biochemical and Biophysical Research Communications</i> , 1984 , 122, 1357-66	3.4	128
71	The coexistence in rat muscle cells of two distinct classes of Ca ²⁺ -dependent K ⁺ channels with different pharmacological properties and different physiological functions. <i>Biochemical and Biophysical Research Communications</i> , 1984 , 118, 669-74	3.4	172
70	Structure-function relationships in the inhibition of synaptosomal dopamine uptake by phencyclidine and analogues: potential correlation with binding site identified with [³ H]phencyclidine. <i>Biochemical Pharmacology</i> , 1984 , 33, 700-2	6	29
69	Identification of an amiloride sensitive Na ⁺ /H ⁺ exchange system in brain synaptosomes. <i>Brain Research</i> , 1984 , 301, 371-4	3.7	29
68	A micro-radioimmunoassay for apamin. <i>Toxicon</i> , 1984 , 22, 985-8	2.8	10
67	Classification of Na channel receptors specific for various scorpion toxins. <i>Pflugers Archiv European Journal of Physiology</i> , 1983 , 397, 164-5	4.6	50
66	Specific photoaffinity labeling of the digitalis binding site of the sodium and potassium ion activated adenosinetriphosphatase induced by energy transfer. <i>Biochemistry</i> , 1983 , 22, 4685-90	3.2	38
65	[³ H]TCP: a new tool with high affinity for the PCP receptor in rat brain. <i>Brain Research</i> , 1983 , 280, 194-7	3.7	141
64	Ethylisopropyl-amiloride: a new and highly potent derivative of amiloride for the inhibition of the Na ⁺ /H ⁺ exchange system in various cell types. <i>Biochemical and Biophysical Research Communications</i> , 1983 , 116, 86-90	3.4	155
63	Ontogenic appearance of Na ⁺ channels characterized as high affinity binding sites for tetrodotoxin during development of the rat nervous and skeletal muscle systems. <i>Biochemical and Biophysical Research Communications</i> , 1983 , 110, 894-901	3.4	44
62	Determination of the molecular size of the nitrendipine-sensitive Ca ²⁺ channel by radiation inactivation. <i>Biochemical and Biophysical Research Communications</i> , 1983 , 111, 878-83	3.4	64
61	Affinity labelling of the tetrodotoxin-binding component of the Na ⁺ channel. <i>Biochemical and Biophysical Research Communications</i> , 1983 , 114, 126-30	3.4	15
60	Ontogenic appearance of Ca ²⁺ channels characterized as binding sites for nitrendipine during development of nervous, skeletal and cardiac muscle systems in the rat. <i>FEBS Letters</i> , 1983 , 164, 75-9	3.8	70

59	New Ouabain Derivatives to Covalently Label the Digitalis Binding Site. <i>Current Topics in Membranes and Transport</i> , 1983 , 271-276		4
58	Apamin, a neurotoxin specific for one class of Ca ²⁺ -dependent K ⁺ channels. <i>Cell Calcium</i> , 1983 , 4, 421-8	4	86
57	Specific binding and pharmacological interactions of apamin, the neurotoxin from bee venom, with guinea pig colon. <i>Life Sciences</i> , 1982 , 31, 437-43	6.8	44
56	Specific binding of toxin II from <i>Centruroides suffusus suffusus</i> to the sodium channel in electroplaque membranes. <i>Biochemistry</i> , 1982 , 21, 5628-34	3.2	16
55	Properties of Na ⁺ channels in fibroblasts. <i>Biochemical and Biophysical Research Communications</i> , 1982 , 107, 202-8	3.4	11
54	The amino acid sequence of toxin V from <i>Anemonia sulcata</i> . <i>Biochemical and Biophysical Research Communications</i> , 1982 , 107, 272-8	3.4	35
53	Biochemical evidence for pharmacological similarities between alpha-adrenoreceptors and voltage-dependent Na ⁺ and Ca ⁺⁺ channels. <i>Biochemical and Biophysical Research Communications</i> , 1982 , 106, 967-73	3.4	28
52	Biochemical properties of the brain phencyclidine receptor. <i>European Journal of Pharmacology</i> , 1982 , 81, 531-42	5.3	77
51	Identification of a protein component of the Ca ²⁺ -dependent K ⁺ channel by affinity labelling with apamin. <i>Biochemical and Biophysical Research Communications</i> , 1982 , 107, 1577-82	3.4	41
50	Na ⁺ channels with binding sites of high and low affinity for tetrodotoxin in different excitable and non-excitable cells. <i>FEBS Journal</i> , 1982 , 124, 199-203		55
49	Lipid-soluble toxins thought to be specific for Na ⁺ channels block Ca ²⁺ channels in neuronal cells. <i>Nature</i> , 1982 , 297, 79-8	50.4	74
48	A cardiac tetrodotoxin binding component: biochemical identification, characterization, and properties. <i>Biochemistry</i> , 1981 , 20, 1279-85	3.2	49
47	Freeze-fracture study of cardiotoxin action on axonal membrane and axonal membrane lipid vesicles. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1981 , 643, 101-14	3.8	23
46	Purification and pharmacological properties of eight sea anemone toxins from <i>Anemonia sulcata</i> , <i>Anthopleura xanthogrammica</i> , <i>Stoichactis giganteus</i> , and <i>Actinodendron plumosum</i> . <i>Biochemistry</i> , 1981 , 20, 5245-52	3.2	130
45	The specificity of the sodium channel for monovalent cations. <i>FEBS Journal</i> , 1981 , 119, 437-42		33
44	¹ H nuclear-magnetic-resonance studies of the conformation of cardiotoxin VII2 from <i>Naja mossambica mossambica</i> . <i>FEBS Journal</i> , 1981 , 120, 467-75		27
43	Properties of the interaction of the sodium channel with permeant monovalent cations. <i>FEBS Journal</i> , 1980 , 106, 71-83		14
42	Toxin-induced K ⁺ efflux through the Na ⁺ channel of neuroblastoma cells. <i>FEBS Journal</i> , 1980 , 111, 265-73		10

41	Synthesis of new, highly radioactive tetrodotoxin derivatives and their binding properties to the sodium channel. <i>FEBS Journal</i> , 1980 , 104, 617-25		83
40	Binding of phencyclidine to rat brain membranes: technical aspect. <i>European Journal of Pharmacology</i> , 1980 , 68, 73-7	5.3	25
39	An in vitro system to study the action potential sodium channel. <i>FEBS Letters</i> , 1980 , 121, 265-8	3.8	9
38	Photoaffinity labelling of a small protein component of a purified (Na ⁺ -K ⁺)ATPase. <i>FEBS Letters</i> , 1979 , 98, 373-6	3.8	33
37	Interaction of molecules of the phencyclidine series with cardiac cells. Association with the muscarinic receptor. <i>FEBS Letters</i> , 1979 , 103, 133-7	3.8	11
36	Photoaffinity labeling of the digitalis receptor in the (sodium + potassium)-activated adenosinetriphosphatase. <i>Biochemistry</i> , 1979 , 18, 135-40	3.2	61
35	A search for the apamin receptor in the central nervous system. <i>Toxicon</i> , 1979 , 17, 176-9	2.8	7
34	Specificity and interactions at the cationic sites of the axonal (Na ⁺ , K ⁺)-activated adenosinetriphosphatase. <i>FEBS Journal</i> , 1978 , 85, 561-70		24
33	The 1H nuclear-magnetic-resonance spectra of Neurotoxin I and cardiotoxin Vii4 from <i>Naja mossambica mossambica</i> . <i>FEBS Journal</i> , 1978 , 92, 361-71		44
32	Isolation and partial characterization of rat CNS axolemma enriched fractions. <i>Brain Research</i> , 1978 , 147, 339-52	3.7	72
31	Properties of association of cardiotoxin with lipid vesicles and natural membranes. A fluorescence study. <i>FEBS Letters</i> , 1978 , 85, 103-8	3.8	59
30	1H n.m.r. studies of a neurotoxin and a cardiotoxin from <i>Naja mossambica mossambica</i> : amide proton resonances. <i>Biochemical and Biophysical Research Communications</i> , 1977 , 76, 1071-8	3.4	30
29	Molecular mechanism of the cardiotoxic action of a polypeptide neurotoxin from sea anemone on cultured embryonic cardiac cells. <i>Biochemistry</i> , 1977 , 16, 3850-5	3.2	31
28	Mechanistic analysis of the (Na ⁺ ,K ⁺)ATPase using new pseudosubstrates. <i>Biochemistry</i> , 1977 , 16, 2957-65.2		27
27	Effects of neurotoxins (veratridine, sea anemone toxin, tetrodotoxin) on transmitter accumulation and release by nerve terminals in vitro. <i>Biochemistry</i> , 1977 , 16, 1838-44	3.2	82
26	Non-equivalence of the sites of yeast phenylalanyl-tRNA synthetase during catalysis. <i>FEBS Journal</i> , 1977 , 73, 7-15		31
25	Molecular mechanism of cardiotoxin action on axonal membranes. <i>Biochemistry</i> , 1976 , 15, 3171-5	3.2	63
24	Pre-existence of the active site in zymogens, the interaction of trypsinogen with the basic pancreatic trypsin inhibitor (Kunitz). <i>FEBS Letters</i> , 1976 , 63, 240-4	3.8	47

23	(Na ⁺ , K ⁺)-activated adenosinetriphosphatase of axonal membranes, cooperativity and control. Steady-state analysis. <i>FEBS Journal</i> , 1976 , 65, 293-306		86
22	Constitution and properties of axonal membranes of crustacean nerves. <i>Biochemistry</i> , 1975 , 14, 5500-113,2		100
21	Structure-function relationship in the binding of snake neurotoxins to the torpedo membrane receptor. <i>Biochemistry</i> , 1975 , 14, 2081-91	3.2	106
20	Structure-function relationships and site of action of apamin, a neurotoxic polypeptide of bee venom with an action on the central nervous system. <i>Biochemistry</i> , 1975 , 14, 2521-5	3.2	97
19	The non-equivalence of the active sites and the mechanism of a mutationally altered E. coli alkaline phosphatase. <i>Biochemical and Biophysical Research Communications</i> , 1975 , 63, 529-34	3.4	7
18	Negative cooperativity and half of the sites reactivity. Alkaline phosphatases of Escherichia coli with Zn ²⁺ , Co ²⁺ , Cd ²⁺ , Mn ²⁺ , and Cu ²⁺ in the active sites. <i>Biochemistry</i> , 1974 , 13, 3754-62	3.2	49
17	The association of anhydrotrypsin with the pancreatic trypsin inhibitors. <i>Biochemistry</i> , 1974 , 13, 4205-113,2		57
16	Intestinal alkaline phosphatase. Physical properties and quaternary structure. <i>Biochemistry</i> , 1974 , 13, 1783-8	3.2	166
15	Intestinal alkaline phosphatase. Catalytic properties and half of the sites reactivity. <i>Biochemistry</i> , 1974 , 13, 1788-95	3.2	120
14	Guanidination of lysine-15 in the active site of the basic pancreatic trypsin inhibitor. Implications for complex formation with trypsin and chymotrypsin. <i>FEBS Journal</i> , 1974 , 42, 505-10		13
13	The interaction between alpha-chymotrypsin and pancreatic trypsin inhibitor (Kunitz inhibitor). Kinetic and thermodynamic properties. <i>FEBS Journal</i> , 1973 , 38, 365-72		66
12	Trypsin--pancreatic secretory inhibitor (Kazal inhibitor) interaction. Kinetic and thermodynamic properties. <i>Biochemistry</i> , 1973 , 12, 2841-6	3.2	55
11	Structure-function relationships of neurotoxins isolated from Naja haje venom. Physicochemical properties and identification of the active site. <i>Biochemistry</i> , 1972 , 11, 1681-91	3.2	75
10	Trypsin-pancreatic trypsin inhibitor association. Dynamics of the interaction and role of disulfide bridges. <i>Biochemistry</i> , 1972 , 11, 2967-77	3.2	241
9	Zymogen-enzyme transformations. On the mechanism of activation of prophospholipase A. <i>FEBS Journal</i> , 1972 , 30, 37-47		48
8	Flip-Flop Mechanisms and Half-Site Enzymes. <i>Current Topics in Cellular Regulation</i> , 1972 , 6, 267-310		66
7	Flip-flop mechanisms in enzymology. A model: the alkaline phosphatase of Escherichia coli. <i>FEBS Journal</i> , 1971 , 20, 124-39		148
6	The conformational properties of the basic pancreatic trypsin-inhibitor. <i>FEBS Journal</i> , 1971 , 23, 401-11		77

5	Structure-function relationships of the acyl-carrier protein of Escherichia coli. <i>FEBS Journal</i> , 1971 , 23, 412-20	28
4	The conformation of small proteins. The state-diagram of a neurotoxin of Androctonus Australis Hector. <i>FEBS Journal</i> , 1970 , 14, 549-55	12
3	The Cu ² plus-alkaline phosphatase of Escherichia coli. <i>FEBS Journal</i> , 1970 , 17, 239-45	25
2	INHIBITION DE LA PHOSPHATASE ALCALINE INTESTINALE. <i>Canadian Journal of Biochemistry and Physiology</i> , 1962 , 40, 1619-1639	1
1	PHOSPHATASE ALCALINE INTESTINALE: CINÉTIQUE DE L'HYDROLYSE DU PHOSPHATE DE p-NITROPHÉNYLE. <i>Canadian Journal of Chemistry</i> , 1961 , 39, 1298-1308	0.9 7