

Olaf van Tellingen

List of Publications by Year in descending order

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Version: 2024-02-01

181
papers

13,035
citations

26630

56
h-index

24258

110
g-index

184
all docs

184
docs citations

184
times ranked

14984
citing authors

#	ARTICLE	IF	CITATIONS
1	Disruption of the mouse <i>mdr1a</i> P-glycoprotein gene leads to a deficiency in the blood-brain barrier and to increased sensitivity to drugs. <i>Cell</i> , 1994, 77, 491-502.	28.9	2,163
2	Overcoming the blood-brain tumor barrier for effective glioblastoma treatment. <i>Drug Resistance Updates</i> , 2015, 19, 1-12.	14.4	706
3	Potent and specific inhibition of the breast cancer resistance protein multidrug transporter in vitro and in mouse intestine by a novel analogue of fumitremogin C. <i>Molecular Cancer Therapeutics</i> , 2002, 1, 417-25.	4.1	371
4	The Effect of Bcrp1 (Abcg2) on the In vivo Pharmacokinetics and Brain Penetration of Imatinib Mesylate (Gleevec): Implications for the Use of Breast Cancer Resistance Protein and P-Glycoprotein Inhibitors to Enable the Brain Penetration of Imatinib in Patients. <i>Cancer Research</i> , 2005, 65, 2577-2582.	0.9	338
5	Absence or pharmacological blocking of placental P-glycoprotein profoundly increases fetal drug exposure. <i>Journal of Clinical Investigation</i> , 1999, 104, 1441-1447.	8.2	314
6	Drug-induced histone eviction from open chromatin contributes to the chemotherapeutic effects of doxorubicin. <i>Nature Communications</i> , 2013, 4, 1908.	12.8	310
7	Selective induction of chemotherapy resistance of mammary tumors in a conditional mouse model for hereditary breast cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 12117-12122.	7.1	279
8	Bmi1 Controls Tumor Development in an Ink4a/Arf-Independent Manner in a Mouse Model for Glioma. <i>Cancer Cell</i> , 2007, 12, 328-341.	16.8	264
9	P-Glycoprotein and Breast Cancer Resistance Protein: Two Dominant Transporters Working Together in Limiting the Brain Penetration of Topotecan. <i>Clinical Cancer Research</i> , 2007, 13, 6440-6449.	7.0	252
10	Mechanism of the Pharmacokinetic Interaction between Methotrexate and Benzimidazoles. <i>Cancer Research</i> , 2004, 64, 5804-5811.	0.9	222
11	Knockout of cytochrome P450 3A yields new mouse models for understanding xenobiotic metabolism. <i>Journal of Clinical Investigation</i> , 2007, 117, 3583-3592.	8.2	210
12	Long-lasting suppression of hippocampal cell proliferation and impaired cognitive performance by methotrexate in the rat. <i>Behavioural Brain Research</i> , 2008, 186, 168-175.	2.2	209
13	Pharmacokinetics, Brain Delivery, and Efficacy in Brain Tumor-Bearing Mice of Glutathione Pegylated Liposomal Doxorubicin (2B3-101). <i>PLoS ONE</i> , 2014, 9, e82331.	2.5	207
14	Tumour-specific proline vulnerability uncovered by differential ribosome codon reading. <i>Nature</i> , 2016, 530, 490-494.	27.8	202
15	MRP2 (ABCC2) transports taxanes and confers paclitaxel resistance and both processes are stimulated by probenecid. <i>International Journal of Cancer</i> , 2005, 116, 824-829.	5.1	189
16	Effect of the ATP-binding cassette drug transporters ABCB1, ABCG2, and ABCC2 on erlotinib hydrochloride (Tarceva) disposition in vitro and in vivo pharmacokinetic studies employing <i>Bcrp1</i> ^{-/-} / <i>Mdr1a/1b</i> ^{-/-} (triple-knockout) and wild-type mice. <i>Molecular Cancer Therapeutics</i> , 2008, 7, 2280-2287.	4.1	183
17	Increased penetration of paclitaxel into the brain by inhibition of P-Glycoprotein. <i>Clinical Cancer Research</i> , 2003, 9, 2849-55.	7.0	183
18	Boronic acid-based inhibitor of autotaxin reveals rapid turnover of LPA in the circulation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 7257-7262.	7.1	182

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19	Modulation of the blood-brain barrier in oncology: therapeutic opportunities for the treatment of brain tumours?. <i>Cancer Treatment Reviews</i> , 2004, 30, 415-423.	7.7	174
20	Concerns about anti-angiogenic treatment in patients with glioblastoma multiforme. <i>BMC Cancer</i> , 2009, 9, 444.	2.6	166
21	Carcinogen and Anticancer Drug Transport by Mrp2 in Vivo: Studies Using <i>Mrp2</i> (<i>Abcc2</i>) Knockout Mice. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2006, 318, 319-327.	2.5	146
22	Restricted brain penetration of the tyrosine kinase inhibitor erlotinib due to the drug transporters P-gp and BCRP. <i>Investigational New Drugs</i> , 2012, 30, 443-449.	2.6	135
23	The effect of P-gp (<i>Mdr1a/1b</i>), BCRP (<i>Bcrp1</i>) and P-gp/BCRP inhibitors on the in vivo absorption, distribution, metabolism and excretion of imatinib. <i>Investigational New Drugs</i> , 2009, 27, 31-40.	2.6	132
24	Preclinical pharmacokinetics of paclitaxel and docetaxel. <i>Anti-Cancer Drugs</i> , 1998, 9, 1-17.	1.4	125
25	Modulation of oral bioavailability of anticancer drugs: from mouse to man. <i>European Journal of Pharmaceutical Sciences</i> , 2000, 12, 103-110.	4.0	125
26	Blood-brain barrier and chemotherapeutic treatment of brain tumors. <i>Expert Review of Neurotherapeutics</i> , 2006, 6, 1199-1209.	2.8	124
27	Validity of bioluminescence measurements for noninvasive in vivo imaging of tumor load in small animals. <i>BioTechniques</i> , 2007, 43, S7-S13, S30.	1.8	121
28	The Functional Role of P-Glycoprotein in the Blood-brain Barrier. <i>Journal of Pharmaceutical Sciences</i> , 1997, 86, 881-884.	3.3	119
29	Chemotherapy-related cognitive dysfunction: current animal studies and future directions. <i>Brain Imaging and Behavior</i> , 2013, 7, 453-459.	2.1	118
30	Low systemic exposure of oral docetaxel in mice resulting from extensive first-pass metabolism is boosted by ritonavir. <i>Cancer Research</i> , 2002, 62, 6158-64.	0.9	116
31	Tissue distribution, metabolism and excretion of paclitaxel in mice. <i>Anti-Cancer Drugs</i> , 1996, 7, 78-86.	1.4	106
32	ABCB1, ABCG2, and PTEN Determine the Response of Glioblastoma to Temozolomide and ABT-888 Therapy. <i>Clinical Cancer Research</i> , 2014, 20, 2703-2713.	7.0	105
33	A Phase I Study of the P-Glycoprotein Antagonist Tariquidar in Combination with Vinorelbine. <i>Clinical Cancer Research</i> , 2009, 15, 3574-3582.	7.0	101
34	Effects of the Selective MPS1 Inhibitor MPS1-IN-3 on Glioblastoma Sensitivity to Antimitotic Drugs. <i>Journal of the National Cancer Institute</i> , 2013, 105, 1322-1331.	6.3	94
35	Uncoupling DNA damage from chromatin damage to detoxify doxorubicin. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 15182-15192.	7.1	93
36	Mouse breast cancer resistance protein (<i>Bcrp1/Abcg2</i>) mediates etoposide resistance and transport, but etoposide oral availability is limited primarily by P-glycoprotein. <i>Cancer Research</i> , 2003, 63, 1339-44.	0.9	89

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37	Multidrug Resistance Protein 2 Is an Important Determinant of Paclitaxel Pharmacokinetics. <i>Clinical Cancer Research</i> , 2006, 12, 6125-6132.	7.0	88
38	Absence of Both Cytochrome <i>P</i> 450 3A and P-glycoprotein Dramatically Increases Docetaxel Oral Bioavailability and Risk of Intestinal Toxicity. <i>Cancer Research</i> , 2009, 69, 8996-9002.	0.9	88
39	Functionally Overlapping Roles of Abcg2 (Bcrp1) and Abcc2 (Mrp2) in the Elimination of Methotrexate and Its Main Toxic Metabolite 7-Hydroxymethotrexate <i>In vivo</i> . <i>Clinical Cancer Research</i> , 2009, 15, 3084-3093.	7.0	87
40	An Experimenters' Guide to Glioblastoma Invasion Pathways. <i>Trends in Molecular Medicine</i> , 2018, 24, 763-780.	6.7	86
41	The influence of the P-glycoprotein inhibitor zosuquidar trihydrochloride (LY335979) on the brain penetration of paclitaxel in mice. <i>Cancer Chemotherapy and Pharmacology</i> , 2004, 53, 173-178.	2.3	85
42	Improved Brain Penetration and Antitumor Efficacy of Temozolomide by Inhibition of ABCB1 and ABCG2. <i>Neoplasia</i> , 2018, 20, 710-720.	5.3	84
43	Phase I and Pharmacokinetic Study of Oral Paclitaxel. <i>Journal of Clinical Oncology</i> , 2000, 18, 2468-2475.	1.6	83
44	P-glycoprotein (P-gp/Abcb1), Abcc2, and Abcc3 Determine the Pharmacokinetics of Etoposide. <i>Clinical Cancer Research</i> , 2010, 16, 130-140.	7.0	79
45	Crizotinib Inhibits Metabolic Inactivation of Gemcitabine in c-Met-driven Pancreatic Carcinoma. <i>Cancer Research</i> , 2013, 73, 6745-6756.	0.9	79
46	Sensitivity and Acquired Resistance of BRCA1;p53-Deficient Mouse Mammary Tumors to the Topoisomerase I Inhibitor Topotecan. <i>Cancer Research</i> , 2010, 70, 1700-1710.	0.9	76
47	Abcc2 (Mrp2), Abcc3 (Mrp3), and Abcg2 (Bcrp1) are the main determinants for rapid elimination of methotrexate and its toxic metabolite 7-hydroxymethotrexate <i>in vivo</i> . <i>Molecular Cancer Therapeutics</i> , 2009, 8, 3350-3359.	4.1	74
48	The importance of drug-transporting P-glycoproteins in toxicology. <i>Toxicology Letters</i> , 2001, 120, 31-41.	0.8	72
49	Efficacy of novel P-glycoprotein inhibitors to increase the oral uptake of paclitaxel in mice. <i>Investigational New Drugs</i> , 2004, 22, 219-229.	2.6	71
50	Prolonged Ezh2 Depletion in Glioblastoma Causes a Robust Switch in Cell Fate Resulting in Tumor Progression. <i>Cell Reports</i> , 2015, 10, 383-397.	6.4	70
51	The co-solvent Cremophor EL limits absorption of orally administered paclitaxel in cancer patients. <i>British Journal of Cancer</i> , 2001, 85, 1472-1477.	6.4	68
52	P-glycoprotein and breast cancer resistance protein restrict the brain penetration of the CDK4/6 inhibitor palbociclib. <i>Investigational New Drugs</i> , 2015, 33, 1012-1019.	2.6	68
53	Glycosylated extracellular vesicles released by glioblastoma cells are decorated by CCL18 allowing for cellular uptake via chemokine receptor CCR8. <i>Journal of Extracellular Vesicles</i> , 2018, 7, 1446660.	12.2	64
54	Determination of paclitaxel and metabolites in mouse plasma, tissues, urine and faeces by semi-automated reversed-phase high-performance liquid chromatography. <i>Biomedical Applications</i> , 1995, 664, 383-391.	1.7	63

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55	Determination of doxorubicin and metabolites in murine specimens by high-performance liquid chromatography. <i>Biomedical Applications</i> , 1998, 712, 129-143.	1.7	62
56	Multiple low dose therapy as an effective strategy to treat EGFR inhibitor-resistant NSCLC tumours. <i>Nature Communications</i> , 2020, 11, 3157.	12.8	59
57	Entrapment by Cremophor EL decreases the absorption of paclitaxel from the gut. <i>Cancer Chemotherapy and Pharmacology</i> , 2002, 49, 119-125.	2.3	57
58	<sc>ABC</sc>1 and <sc>ABC</sc>2 restrict the brain penetration of a panel of novel <sc>EZH</sc>2 inhibitors. <i>International Journal of Cancer</i> , 2015, 137, 2007-2018.	5.1	57
59	Characterisation of tumour vasculature in mouse brain by USPIO contrast-enhanced MRI. <i>British Journal of Cancer</i> , 2008, 98, 1784-1789.	6.4	56
60	Impact of Abcc2 (Mrp2) and Abcc3 (Mrp3) on the <i>in vivo</i> Elimination of Methotrexate and its Main Toxic Metabolite 7-hydroxymethotrexate. <i>Clinical Cancer Research</i> , 2008, 14, 8152-8160.	7.0	56
61	High-grade glioma mouse models and their applicability for preclinical testing. <i>Cancer Treatment Reviews</i> , 2009, 35, 714-723.	7.7	56
62	PI3K-mTOR Pathway Inhibition Exhibits Efficacy Against High-grade Glioma in Clinically Relevant Mouse Models. <i>Clinical Cancer Research</i> , 2017, 23, 1286-1298.	7.0	56
63	Therapy-resistant tumor microvascular endothelial cells contribute to treatment failure in glioblastoma multiforme. <i>Oncogene</i> , 2013, 32, 1539-1548.	5.9	55
64	Identification of a Druggable Pathway Controlling Glioblastoma Invasiveness. <i>Cell Reports</i> , 2017, 20, 48-60.	6.4	55
65	The impact of P-glycoprotein and breast cancer resistance protein on the brain pharmacokinetics and pharmacodynamics of a panel of MEK inhibitors. <i>International Journal of Cancer</i> , 2018, 142, 381-391.	5.1	55
66	Tumor microvasculature supports proliferation and expansion of glioma-propagating cells. <i>International Journal of Cancer</i> , 2009, 125, 1222-1230.	5.1	53
67	Cognitive impact of cytotoxic agents in mice. <i>Psychopharmacology</i> , 2015, 232, 17-37.	3.1	53
68	Isolation, purification, and biological activity of mono- and dihydroxylated paclitaxel metabolites from human feces. <i>Cancer Chemotherapy and Pharmacology</i> , 1995, 36, 299-304.	2.3	52
69	Rapid and Robust Transgenic High-Grade Glioma Mouse Models for Therapy Intervention Studies. <i>Clinical Cancer Research</i> , 2010, 16, 3431-3441.	7.0	52
70	Buparlisib is a brain penetrable pan-PI3K inhibitor. <i>Scientific Reports</i> , 2018, 8, 10784.	3.3	52
71	Magnetic resonance imaging-based detection of glial brain tumors in mice after antiangiogenic treatment. <i>International Journal of Cancer</i> , 2008, 122, 1981-1986.	5.1	51
72	High Impact of Oatp1a/1b Transporters on In Vivo Disposition of the Hydrophobic Anticancer Drug Paclitaxel. <i>Clinical Cancer Research</i> , 2011, 17, 294-301.	7.0	49

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73	Intratumoural administration of cisplatin in slow-release devices: II. pharmacokinetics and intratumoural distribution. <i>Cancer Chemotherapy and Pharmacology</i> , 1991, 27, 347-353.	2.3	48
74	Impact of Abcc2 [Multidrug Resistance-Associated Protein (Mrp) 2], Abcc3 (Mrp3), and Abcg2 (Breast) Tj ETQq0 0 0 rgBT /Overlock 10 T 7-Hydroxymethotrexate. <i>Drug Metabolism and Disposition</i> , 2011, 39, 1338-1344.	3.3	48
75	Abcc4 Together with Abcb1 and Abcg2 Form a Robust Cooperative Drug Efflux System That Restricts the Brain Entry of Camptothecin Analogues. <i>Clinical Cancer Research</i> , 2013, 19, 2084-2095.	7.0	48
76	Tryptophan depletion results in tryptophan-to-phenylalanine substitutants. <i>Nature</i> , 2022, 603, 721-727.	27.8	47
77	Combined Therapy of AXL and HDAC Inhibition Reverses Mesenchymal Transition in Diffuse Intrinsic Pontine Glioma. <i>Clinical Cancer Research</i> , 2020, 26, 3319-3332.	7.0	44
78	The oral route for the administration of cytotoxic drugs: strategies to increase the efficiency and consistency of drug delivery. <i>Investigational New Drugs</i> , 2000, 18, 231-241.	2.6	43
79	Determination of imatinib mesylate and its main metabolite (CGP74588) in human plasma and murine specimens by ion-pairing reversed-phase high-performance liquid chromatography. <i>Biomedical Chromatography</i> , 2007, 21, 747-754.	1.7	43
80	<i>In vivo</i> disposition of doxorubicin is affected by mouse Oatp1a/1b and human OATP1A/1B transporters. <i>International Journal of Cancer</i> , 2014, 135, 1700-1710.	5.1	43
81	<i>SLC1A3</i> contributes to <i>Asparaginase</i> resistance in solid tumors. <i>EMBO Journal</i> , 2019, 38, e102147.	7.8	41
82	A phase I and pharmacokinetic study of bi-daily dosing of oral paclitaxel in combination with cyclosporin A. <i>Cancer Chemotherapy and Pharmacology</i> , 2001, 47, 347-354.	2.3	39
83	P-glycoprotein and Mrp1 collectively protect the bone marrow from vincristine-induced toxicity in vivo. <i>British Journal of Cancer</i> , 2003, 89, 1776-1782.	6.4	39
84	Cannulation of the jugular vein in mice: a method for serial withdrawal of blood samples. <i>Laboratory Animals</i> , 2003, 37, 181-187.	1.0	37
85	Neurobiological changes by cytotoxic agents in mice. <i>Behavioural Brain Research</i> , 2016, 299, 19-26.	2.2	36
86	The G2 checkpoint is a node-based molecular switch. <i>FEBS Open Bio</i> , 2017, 7, 439-455.	2.3	36
87	Phase I trial and pharmacological study of a 3-hour paclitaxel infusion in children with refractory solid tumours: a SFOP study. <i>British Journal of Cancer</i> , 2001, 84, 604-610.	6.4	35
88	Determination of polyoxyethyleneglycerol triricinoleate 35 (Cremophor EL) in plasma by pre-column derivatization and reversed-phase high-performance liquid chromatography. <i>Biomedical Applications</i> , 1996, 681, 355-362.	1.7	34
89	OATP1A/1B Transporters Affect Irinotecan and SN-38 Pharmacokinetics and Carboxylesterase Expression in Knockout and Humanized Transgenic Mice. <i>Molecular Cancer Therapeutics</i> , 2014, 13, 492-503.	4.1	33
90	EZH2 Is Overexpressed in <i>BRCA1</i> -like Breast Tumors and Predictive for Sensitivity to High-Dose Platinum-Based Chemotherapy. <i>Clinical Cancer Research</i> , 2019, 25, 4351-4362.	7.0	33

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91	Preclinical Mouse Models To Study Human OATP1B1- and OATP1B3-Mediated Drugâ€“Drug Interactions <i>in Vivo</i> . <i>Molecular Pharmaceutics</i> , 2015, 12, 4259-4269.	4.6	32
92	ATP-binding cassette transporters restrict drug delivery and efficacy against brain tumors even when blood-brain barrier integrity is lost. <i>Cell Reports Medicine</i> , 2021, 2, 100184.	6.5	32
93	A simple and sensitive assay for the quantitative analysis of paclitaxel in human and mouse plasma and brain tumor tissue using coupled liquid chromatography and tandem mass spectrometry. <i>Journal of Mass Spectrometry</i> , 2004, 39, 1506-1512.	1.6	31
94	MELK Inhibition in Diffuse Intrinsic Pontine Glioma. <i>Clinical Cancer Research</i> , 2018, 24, 5645-5657.	7.0	30
95	The effect of different doses of cyclosporin A on the systemic exposure of orally administered paclitaxel. <i>Anti-Cancer Drugs</i> , 2001, 12, 351-358.	1.4	29
96	Extensive Metabolism and Hepatic Accumulation of Gemcitabine After Multiple Oral and Intravenous Administration in Mice. <i>Drug Metabolism and Disposition</i> , 2008, 36, 1606-1615.	3.3	29
97	Mps1 inhibitors synergise with low doses of taxanes in promoting tumour cell death by enhancement of errors in cell division. <i>British Journal of Cancer</i> , 2018, 118, 1586-1595.	6.4	29
98	Differential effects of anticoagulants on tumor development of mouse cancer cell lines B16, K1735 and CT26 in lung. <i>Clinical and Experimental Metastasis</i> , 2009, 26, 171-178.	3.3	28
99	In vitro transport of gimatecan (7-t-butoxyiminomethylcamptothecin) by breast cancer resistance protein, P-glycoprotein, and multidrug resistance protein 2. <i>Molecular Cancer Therapeutics</i> , 2007, 6, 3307-3313.	4.1	27
100	Metabolism and excretion of paclitaxel after oral administration in combination with cyclosporin A and after i.v. administration. <i>Anti-Cancer Drugs</i> , 2000, 11, 813-820.	1.4	26
101	Dual mTORC1 and mTORC2 inhibitor Palomid 529 penetrates the Bloodâ€“Brain Barrier without restriction by ABCB1 and ABCG2. <i>International Journal of Cancer</i> , 2013, 133, 1222-1233.	5.1	26
102	EZN-2208 (PEG-SN38) Overcomes ABCG2-Mediated Topotecan Resistance in BRCA1-Deficient Mouse Mammary Tumors. <i>PLoS ONE</i> , 2012, 7, e45248.	2.5	24
103	Effect of the drug transporters ABCG2, Abcg2, ABCB1 and ABCC2 on the disposition, brain accumulation and myelotoxicity of the aurora kinase B inhibitor barasertib and its more active form barasertib-hydroxy-QPA. <i>Investigational New Drugs</i> , 2013, 31, 1125-1135.	2.6	24
104	Determination of oxaliplatin in human plasma and plasma ultrafiltrate by graphite-furnace atomic-absorption spectrometry. <i>Analytical and Bioanalytical Chemistry</i> , 2005, 382, 1484-1490.	3.7	23
105	Analytical methods for the determination of vinca alkaloids in biological specimens: A survey of the literature. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 1991, 9, 1077-1082.	2.8	22
106	Plasma pharmacokinetics, tissue disposition, excretion and metabolism of vinorelbine in mice as determined by high performance liquid chromatography. <i>Investigational New Drugs</i> , 1993, 11, 141-150.	2.6	21
107	Targeting core (mutated) pathways of high-grade gliomas: challenges of intrinsic resistance and drug efflux. <i>CNS Oncology</i> , 2013, 2, 271-288.	3.0	21
108	MEK/MELK inhibition and bloodâ€“brain barrier deficiencies in atypical teratoid/rhabdoid tumors. <i>Neuro-Oncology</i> , 2020, 22, 58-69.	1.2	21

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109	Determination of topotecan in human and mouse plasma and in mouse tissue homogenates by reversed-phase high-performance liquid chromatography. <i>Biomedical Chromatography</i> , 2007, 21, 1191-1200.	1.7	20
110	Paclitaxel in self-micro emulsifying formulations: oral bioavailability study in mice. <i>Investigational New Drugs</i> , 2011, 29, 768-776.	2.6	20
111	ABCB1 Attenuates the Brain Penetration of the PARP Inhibitor AZD2461. <i>Molecular Pharmaceutics</i> , 2018, 15, 5236-5243.	4.6	20
112	Tissue disposition, excretion and metabolism of vinblastine in mice as determined by high-performance liquid chromatography. <i>Cancer Chemotherapy and Pharmacology</i> , 1993, 32, 286-292.	2.3	19
113	Sildenafil is not a useful modulator of ABCB1 and ABCG2 mediated drug resistance in vivo. <i>European Journal of Cancer</i> , 2013, 49, 2059-2064.	2.8	19
114	Molecular Imaging of ABCB1 and ABCG2 Inhibition at the Human Blood-Brain Barrier Using Elacridar and ¹¹ C-Erlotinib PET. <i>Journal of Nuclear Medicine</i> , 2018, 59, 973-979.	5.0	19
115	Monitoring Carboplatin Concentrations in Saliva. <i>Therapeutic Drug Monitoring</i> , 1995, 17, 465-470.	2.0	18
116	Topoisomerase I/II inhibitor irinotecan administered as a 24 h infusion. <i>Anti-Cancer Drugs</i> , 1999, 10, 17-24.	1.4	18
117	Trabectedin (ET-743, Yondelis [®]) is a substrate for P-glycoprotein, but only high expression of P-glycoprotein confers the multidrug resistance phenotype. <i>Investigational New Drugs</i> , 2007, 25, 1-7.	2.6	18
118	Expression and Cellular Distribution of P-Glycoprotein and Breast Cancer Resistance Protein in Amyotrophic Lateral Sclerosis Patients. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 266-276.	1.7	17
119	Validated method for the determination of the novel organo-ruthenium anticancer drug NAMI-A in human biological fluids by Zeeman atomic absorption spectrometry. <i>Fresenius' Journal of Analytical Chemistry</i> , 2001, 369, 442-445.	1.5	15
120	A serum and platelet-rich plasma serotonin assay using liquid chromatography tandem mass spectrometry for monitoring of neuroendocrine tumor patients. <i>Clinica Chimica Acta</i> , 2017, 469, 130-135.	1.1	15
121	Metabolism of paclitaxel in mice. <i>Anti-Cancer Drugs</i> , 2003, 14, 203-209.	1.4	14
122	Metabolism of docetaxel in mice. <i>Cancer Chemotherapy and Pharmacology</i> , 2005, 56, 299-306.	2.3	14
123	Strategies to target drugs to gliomas and CNS metastases of solid tumors. <i>Journal of Neurology</i> , 2016, 263, 428-440.	3.6	14
124	Development and validation of a method to determine the unbound paclitaxel fraction in human plasma. <i>Analytical Biochemistry</i> , 2004, 324, 11-15.	2.4	12
125	Evaluation of Human Plasma Protein Binding of Trabectedin (Yondelis [®] ; ET-743). <i>Current Clinical Pharmacology</i> , 2009, 4, 38-42.	0.6	12
126	Disposition and toxicity of trabectedin (ET-743) in wild-type and mdr1 gene (P-gp) knock-out mice. <i>Investigational New Drugs</i> , 2010, 28, 145-155.	2.6	12

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127	Phase I study of Carzelesin (U-80,244) given (4-weekly) by intravenous bolus schedule. <i>British Journal of Cancer</i> , 1999, 79, 1454-1461.	6.4	11
128	Effect of the drug transporters ABCB1, ABCC2, and ABCG2 on the disposition and brain accumulation of the taxane analog BMS-275,183. <i>Investigational New Drugs</i> , 2014, 32, 1083-1095.	2.6	11
129	Plasma membrane targeting by short chain sphingolipids inserted in liposomes improves anti-tumor activity of mitoxantrone in an orthotopic breast carcinoma xenograft model. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2015, 94, 207-219.	4.3	11
130	The vascular compartment hampers accurate determination of teniposide penetration into brain tumor tissue. <i>Cancer Chemotherapy and Pharmacology</i> , 1997, 40, 330-334.	2.3	10
131	A population analysis of the pharmacokinetics of Cremophor EL using nonlinear mixed-effect modelling. <i>Cancer Chemotherapy and Pharmacology</i> , 2002, 50, 16-24.	2.3	10
132	The effect of P-glycoprotein and cytochrome P450 3a on the oral bioavailability of vinorelbine in mice. <i>Cancer Chemotherapy and Pharmacology</i> , 2006, 57, 819-825.	2.3	10
133	Abstract 5687: Development of glutathione pegylated liposomal doxorubicin (2B3-101) for the treatment of brain cancer. , 2012, , .		10
134	Cremophor EL pharmacokinetics in a phase I study of paclitaxel (Taxol®) and carboplatin in non-small cell lung cancer patients. <i>Anti-Cancer Drugs</i> , 2000, 11, 687-694.	1.4	9
135	Determination of cyclosporin A in human and mouse plasma by reversed-phase high-performance liquid chromatography. <i>Biomedical Applications</i> , 2001, 763, 201-206.	1.7	9
136	Clinical pharmacokinetics of an amorphous solid dispersion tablet of elacridar. <i>Drug Delivery and Translational Research</i> , 2017, 7, 125-131.	5.8	9
137	ATP-binding cassette transporters limit the brain penetration of Wee1 inhibitors. <i>Investigational New Drugs</i> , 2018, 36, 380-387.	2.6	8
138	Retrospective analysis of serum testosterone levels by LC-MS/MS in chemically castrated prostate cancer patients: Biological variation and analytical performance specifications. <i>Clinica Chimica Acta</i> , 2021, 521, 70-75.	1.1	8
139	Phase I and pharmacologic study of weekly doxorubicin and 1 h infusional paclitaxel in patients with advanced breast cancer. <i>Anti-Cancer Drugs</i> , 1998, 9, 665-763.	1.4	7
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