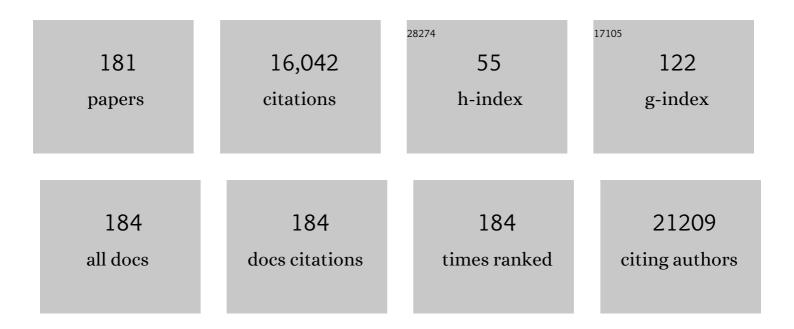
Jan H M Schellens

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Dihydropyrimidine Dehydrogenase Phenotyping Using Pretreatment Uracil: A Note of Caution Based on a Large Prospective Clinical Study. Clinical Pharmacology and Therapeutics, 2022, 112, 62-68.	4.7	32
2	Bioanalytical LC–MS/MS validation of therapeutic drug monitoring assays in oncology. Biomedical Chromatography, 2020, 34, e4623.	1.7	13
3	Pembrolizumab After Two or More Lines of Previous Therapy in Patients With Recurrent or Metastatic SCLC: Results From the KEYNOTE-028 and KEYNOTE-158 Studies. Journal of Thoracic Oncology, 2020, 15, 618-627.	1.1	254
4	Circulating epithelial tumor cell analysis in CSF in patients with leptomeningeal metastases. Neurology, 2020, 94, e521-e528.	1.1	40
5	Evaluating the role of ENOSF1 and TYMS variants as predictors in fluoropyrimidine-related toxicities: An IPD meta-analysis. Pharmacological Research, 2020, 152, 104594.	7.1	17
6	No relation between docetaxel administration route and highâ€grade diarrhea incidence. Pharmacology Research and Perspectives, 2020, 8, e00633.	2.4	9
7	Dabrafenib plus trametinib in patients with BRAFV600E-mutated biliary tract cancer (ROAR): a phase 2, open-label, single-arm, multicentre basket trial. Lancet Oncology, The, 2020, 21, 1234-1243.	10.7	297
8	Phase 1 study of the pan-HER inhibitor dacomitinib plus the MEK1/2 inhibitor PD-0325901 in patients with KRAS-mutation-positive colorectal, non-small-cell lung and pancreatic cancer. British Journal of Cancer, 2020, 122, 1166-1174.	6.4	30
9	Phase I pharmacological study of continuous chronomodulated capecitabine treatment. Pharmaceutical Research, 2020, 37, 89.	3.5	12
10	Quantification of the pharmacokinetic-toxicodynamic relationship of oral docetaxel co-administered with ritonavir. Investigational New Drugs, 2020, 38, 1526-1532.	2.6	1
11	Phase I study of lapatinib plus trametinib in patients with KRAS-mutant colorectal, non-small cell lung, and pancreatic cancer. Cancer Chemotherapy and Pharmacology, 2020, 85, 917-930.	2.3	29
12	Population Pharmacokinetics of MCLA-128, a HER2/HER3 Bispecific Monoclonal Antibody, in Patients with Solid Tumors. Clinical Pharmacokinetics, 2020, 59, 875-884.	3.5	13
13	Pharmacokinetics of Capecitabine and Four Metabolites in a Heterogeneous Population of Cancer Patients: A Comprehensive Analysis. CPT: Pharmacometrics and Systems Pharmacology, 2019, 8, 940-950.	2.5	19
14	Comparison of toxicity and effectiveness between fixed-dose and body surface area-based dose capecitabine. Therapeutic Advances in Medical Oncology, 2019, 11, 175883591983896.	3.2	10
15	DPYD genotype-guided dose individualisation of fluoropyrimidine therapy: who and how? – Authors' reply. Lancet Oncology, The, 2019, 20, e67.	10.7	2
16	Neutropenia and docetaxel exposure in metastatic castrationâ€resistant prostate cancer patients: A metaâ€analysis and evaluation of a clinical cohort. Cancer Medicine, 2019, 8, 1406-1415.	2.8	13
17	Development and validation of LC-MS/MS methods for the quantification of the novel anticancer agent guadecitabine and its active metabolite βâ€decitabine in human plasma, whole blood and urine. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2019, 1109, 132-141.	2.3	10
18	A phase I study of the HDM2 antagonist SAR405838 combined with the MEK inhibitor pimasertib in patients with advanced solid tumours. British Journal of Cancer, 2019, 120, 286-293.	6.4	39

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19	A cost analysis of upfront DPYD genotype–guided dose individualisation in fluoropyrimidine-based anticancer therapy. European Journal of Cancer, 2019, 107, 60-67.	2.8	65
20	Impact of Older Age on the Exposure of Paclitaxel: a Population Pharmacokinetic Study. Pharmaceutical Research, 2019, 36, 33.	3.5	6
21	Enzyme linked immunosorbent assay for the quantification of nivolumab and pembrolizumab in human serum and cerebrospinal fluid. Journal of Pharmaceutical and Biomedical Analysis, 2019, 164, 128-134.	2.8	47
22	Development, validation, and clinical application of a high-performance liquid chromatography-tandem mass spectrometry assay for the quantification of total intracellular β-decitabine nucleotides and genomic DNA incorporated β-decitabine and 5-methyl-2′-deoxycytidine. Journal of Pharmaceutical and Biomedical Analysis, 2019, 164, 16-26.	2.8	9
23	Effectiveness and safety of reducedâ€dose fluoropyrimidine therapy in patients carrying the <i>DPYD</i> *2A variant: A matched pair analysis. International Journal of Cancer, 2019, 144, 2347-2354.	5.1	40
24	Efficacy and safety of dabrafenib (D) and trametinib (T) in patients (pts) with <i>BRAF</i> V600E–mutated biliary tract cancer (BTC): A cohort of the ROAR basket trial Journal of Clinical Oncology, 2019, 37, 187-187.	1.6	66
25	Solubility and bioavailability improvement of pazopanib hydrochloride. International Journal of Pharmaceutics, 2018, 544, 181-190.	5.2	21
26	The impact of liver resection on the dihydrouracil:uracil plasma ratio in patients with colorectal liver metastases. European Journal of Clinical Pharmacology, 2018, 74, 737-744.	1.9	8
27	Intracellular pharmacokinetics of gemcitabine, its deaminated metabolite 2′,2′â€difluorodeoxyuridine and their nucleotides. British Journal of Clinical Pharmacology, 2018, 84, 1279-1289.	2.4	26
28	Combined BRAF, EGFR, and MEK Inhibition in Patients with <i>BRAF</i> V600E-Mutant Colorectal Cancer. Cancer Discovery, 2018, 8, 428-443.	9.4	448
29	Bioanalytical assay for the quantification of the ALK inhibitor lorlatinib in mouse plasma using liquid chromatography-tandem mass spectrometry. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2018, 1083, 204-208.	2.3	16
30	PARP Inhibitors in the Treatment of Triple-Negative Breast Cancer. Clinical Pharmacokinetics, 2018, 57, 427-437.	3.5	87
31	Thermal stability study of crystalline and novel spray-dried amorphous nilotinib hydrochloride. Journal of Pharmaceutical and Biomedical Analysis, 2018, 148, 182-188.	2.8	7
32	Capecitabineâ€based treatment of a patient with a novel <i>DPYD</i> genotype and complete dihydropyrimidine dehydrogenase deficiency. International Journal of Cancer, 2018, 142, 424-430.	5.1	15
33	A drug–drug interaction study to assess the effect of the CYP1A2 inhibitor fluvoxamine on the pharmacokinetics of dovitinib (TKI258) in patients with advanced solid tumors. Cancer Chemotherapy and Pharmacology, 2018, 81, 73-80.	2.3	2
34	Clinical Pharmacogenetics Implementation Consortium (CPIC) Guideline for Dihydropyrimidine Dehydrogenase Genotype and Fluoropyrimidine Dosing: 2017 Update. Clinical Pharmacology and Therapeutics, 2018, 103, 210-216.	4.7	407
35	Improved pharmacodynamic (PD) assessment of low dose PARP inhibitor PD activity for radiotherapy and chemotherapy combination trials. Radiotherapy and Oncology, 2018, 126, 443-449.	0.6	17
36	Clinical Pharmacokinetics of Systemically Administered Antileishmanial Drugs. Clinical Pharmacokinetics, 2018, 57, 151-176.	3.5	55

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37	Dabrafenib and Trametinib Treatment in Patients With Locally Advanced or Metastatic <i>BRAF</i> V600–Mutant Anaplastic Thyroid Cancer. Journal of Clinical Oncology, 2018, 36, 7-13.	1.6	630
38	Diagnostic and Therapeutic Strategies for Fluoropyrimidine Treatment of Patients Carrying Multiple DPYD Variants. Genes, 2018, 9, 585.	2.4	10
39	Standard fluoropyrimidine dosages in chemoradiation therapy result in an increased risk of severe toxicity in DPYD variant allele carriers. European Journal of Cancer, 2018, 104, 210-218.	2.8	14
40	Development and validation of a quantitative method for thymidine phosphorylase activity in peripheral blood mononuclear cells. Nucleosides, Nucleotides and Nucleic Acids, 2018, 37, 436-454.	1.1	1
41	DPYD genotype-guided dose individualisation of fluoropyrimidine therapy in patients with cancer: a prospective safety analysis. Lancet Oncology, The, 2018, 19, 1459-1467.	10.7	238
42	Quantitative bioanalytical assay for the tropomyosin receptor kinase inhibitor larotrectinib in mouse plasma and tissue homogenates using liquid chromatography-tandem mass spectrometry. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2018, 1102-1103, 167-172.	2.3	10
43	Simultaneous population pharmacokinetic modelling of plasma and intracellular PBMC miltefosine concentrations in New World cutaneous leishmaniasis and exploration of exposure–response relationships. Journal of Antimicrobial Chemotherapy, 2018, 73, 2104-2111.	3.0	11
44	Foodâ€effect study on uracil and dihydrouracil plasma levels as marker for dihydropyrimidine dehydrogenase activity in human volunteers. British Journal of Clinical Pharmacology, 2018, 84, 2761-2769.	2.4	26
45	Therapeutic drug monitoring of small molecule kinase inhibitors in oncology in a realâ€world cohort study: does age matter?. British Journal of Clinical Pharmacology, 2018, 84, 2770-2778.	2.4	14
46	Review of Chromatographic Bioanalytical Assays for the Quantitative Determination of Marine-Derived Drugs for Cancer Treatment. Marine Drugs, 2018, 16, 246.	4.6	16
47	Macrophage Activation Marker Neopterin: A Candidate Biomarker for Treatment Response and Relapse in Visceral Leishmaniasis. Frontiers in Cellular and Infection Microbiology, 2018, 8, 181.	3.9	15
48	Bioanalytical liquid chromatography-tandem mass spectrometric assay for the quantification of the ALK inhibitors alectinib, brigatinib and lorlatinib in plasma and mouse tissue homogenates. Journal of Pharmaceutical and Biomedical Analysis, 2018, 161, 136-143.	2.8	22
49	89Zr-labeled CEA-targeted IL-2 variant immunocytokine in patients with solid tumors: CEA-mediated tumor accumulation and role of IL-2 receptor-binding. Oncotarget, 2018, 9, 24737-24749.	1.8	24
50	Evaluation of BGJ398, a Fibroblast Growth Factor Receptor 1-3 Kinase Inhibitor, in Patients With Advanced Solid Tumors Harboring Genetic Alterations in Fibroblast Growth Factor Receptors: Results of a Global Phase I, Dose-Escalation and Dose-Expansion Study. Journal of Clinical Oncology, 2017, 35, 157-165.	1.6	345
51	How Much Longer Will We Put Up With \$100,000 Cancer Drugs?. Cell, 2017, 168, 579-583.	28.9	74
52	Pretreatment serum uracil concentration as a predictor of severe and fatal fluoropyrimidine-associated toxicity. British Journal of Cancer, 2017, 116, 1415-1424.	6.4	94
53	Molecular Pathways: Targeting the Protein Kinase Wee1 in Cancer. Clinical Cancer Research, 2017, 23, 4540-4544.	7.0	106
54	A Phase lb Dose-Escalation Study of Encorafenib and Cetuximab with or without Alpelisib in Metastatic <i>BRAF</i> -Mutant Colorectal Cancer. Cancer Discovery, 2017, 7, 610-619.	9.4	194

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55	A phase I study of SAR405838, a novel human double minute 2 (HDM2) antagonist, in patients with solid tumours. European Journal of Cancer, 2017, 76, 144-151.	2.8	92
56	A Phase I Dose-Escalation Study of the Safety and Pharmacokinetics of Pictilisib in Combination with Erlotinib in Patients with Advanced Solid Tumors. Oncologist, 2017, 22, 1491-1499.	3.7	23
57	A dose-escalation study of bi-daily once weekly oral docetaxel either as ModraDoc001 or ModraDoc006 combined with ritonavir. European Journal of Cancer, 2017, 86, 217-225.	2.8	23
58	An LC–MS/MS method for quantification of the active abiraterone metabolite Δ(4)-abiraterone (D4A) in human plasma. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2017, 1068-1069, 119-124.	2.3	11
59	Clinical trial simulations in paediatric oncology: AÂfeasibility study from the Innovative Therapies for Children with Cancer Consortium. European Journal of Cancer, 2017, 85, 78-85.	2.8	5
60	Dihydrofolate Reductase/Thymidylate Synthase Fine-Tunes the Folate Status and Controls Redox Homeostasis in Plants. Plant Cell, 2017, 29, 2831-2853.	6.6	64
61	Liquid chromatography-tandem mass spectrometric assay for the quantitative determination of the tyrosine kinase inhibitor quizartinib in mouse plasma using salting-out liquid-liquid extraction. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2017, 1061-1062. 300-305.	2.3	7
62	Improving the solubility of nilotinib through novel spray-dried solid dispersions. International Journal of Pharmaceutics, 2017, 529, 294-302.	5.2	28
63	<i>BRAF</i> Mutations as Predictive Biomarker for Response to Anti-EGFR Monoclonal Antibodies. Oncologist, 2017, 22, 864-872.	3.7	56
64	Potential Benefit of Low-Dose Candesartan in Trastuzumab-Induced Cardiotoxic Effects—Reply. JAMA Oncology, 2017, 3, 279.	7.1	0
65	Treatment Algorithm for Homozygous or Compound Heterozygous DPYD Variant Allele Carriers With Low-Dose Capecitabine. JCO Precision Oncology, 2017, 1, 1-10.	3.0	8
66	Treatment of Peritoneal Dissemination in Stomach Cancer Patients With Cytoreductive Surgery and Hyperthermic Intraperitoneal Chemotherapy (HIPEC): Rationale and Design of the PERISCOPE Study. JMIR Research Protocols, 2017, 6, e136.	1.0	17
67	Pharmacokinetics of Selected Anticancer Drugs in Elderly Cancer Patients: Focus on Breast Cancer. Cancers, 2016, 8, 6.	3.7	28
68	Pronounced betweenâ€subject and circadian variability in thymidylate synthase and dihydropyrimidine dehydrogenase enzyme activity in human volunteers. British Journal of Clinical Pharmacology, 2016, 82, 706-716.	2.4	44
69	Bevacizumab combined with docetaxel, oxaliplatin, and capecitabine, followed by maintenance with capecitabine and bevacizumab, as firstâ€line treatment of patients with advanced HER2â€negative gastric cancer: A multicenter phase 2 study. Cancer, 2016, 122, 1434-1443.	4.1	31
70	Rs895819 in <scp><i>MIR27A</i></scp> improves the predictive value of <scp><i>DPYD</i></scp> variants to identify patients at risk of severe fluoropyrimidineâ€associated toxicity. International Journal of Cancer, 2016, 138, 2752-2761.	5.1	28
71	Reply to T. Magnes et al. Journal of Clinical Oncology, 2016, 34, 2434-2435.	1.6	3
72	Phase II Study of WEE1 Inhibitor AZD1775 Plus Carboplatin in Patients With <i>TP53</i> -Mutated Ovarian Cancer Refractory or Resistant to First-Line Therapy Within 3 Months. Journal of Clinical Oncology, 2016, 34, 4354-4361.	1.6	241

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73	Liquid chromatography–tandem mass spectrometric assay for ponatinib and N-desmethyl ponatinib in mouse plasma. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2016, 1023-1024, 24-29.	2.3	11
74	Development and validation of a rapid and sensitive UPLC–MS/MS method for determination of uracil and dihydrouracil in human plasma. Journal of Pharmaceutical and Biomedical Analysis, 2016, 126, 75-82.	2.8	39
75	Phase I Study Evaluating WEE1 Inhibitor AZD1775 As Monotherapy and in Combination With Gemcitabine, Cisplatin, or Carboplatin in Patients With Advanced Solid Tumors. Journal of Clinical Oncology, 2016, 34, 4371-4380.	1.6	203
76	Improving safety of fluoropyrimidine chemotherapy by individualizing treatment based on dihydropyrimidine dehydrogenase activity – Ready for clinical practice?. Cancer Treatment Reviews, 2016, 50, 23-34.	7.7	76
77	Antidrug Antibody Formation in Oncology: Clinical Relevance and Challenges. Oncologist, 2016, 21, 1260-1268.	3.7	87
78	Liquid chromatographyâ;¿tandem mass spectrometric assay for therapeutic drug monitoring of the B-Raf inhibitor encorafenib, the EGFR inhibitors afatinib, erlotinib and gefitinib and the Oâ;¿ desmethyl metabolites of erlotinib and gefitinib in human plasma. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2016, 1033-1034, 390-398.	2.3	30
79	Pharmaceutical development of an oral tablet formulation containing a spray dried amorphous solid dispersion of docetaxel or paclitaxel. International Journal of Pharmaceutics, 2016, 511, 765-773.	5.2	40
80	Liquid chromatography–tandem mass spectrometric assay for the T790M mutant EGFR inhibitor osimertinib (AZD9291) in human plasma. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2016, 1031, 80-85.	2.3	38
81	Increased risk of severe fluoropyrimidine-associated toxicity in patients carrying a G to C substitution in the first 28-bp tandem repeat of the thymidylate synthase 2R allele. International Journal of Cancer, 2016, 138, 245-253.	5.1	23
82	Pharmacokinetics and excretion of 14C-omacetaxine in patients with advanced solid tumors. Investigational New Drugs, 2016, 34, 565-574.	2.6	3
83	Patients homozygous for DPYD c.1129-5923C>G/haplotype B3 have partial DPD deficiency and require a dose reduction when treated with fluoropyrimidines. Cancer Chemotherapy and Pharmacology, 2016, 78, 875-880.	2.3	17
84	A dose escalating phase I study of GLPG0187, a broad spectrum integrin receptor antagonist, in adult patients with progressive high-grade glioma and other advanced solid malignancies. Investigational New Drugs, 2016, 34, 184-192.	2.6	46
85	Angiotensin Il–Receptor Inhibition With Candesartan to Prevent Trastuzumab-Related Cardiotoxic Effects in Patients With Early Breast Cancer. JAMA Oncology, 2016, 2, 1030.	7.1	160
86	Recent developments in the chromatographic bioanalysis of approved kinase inhibitor drugs in oncology. Journal of Pharmaceutical and Biomedical Analysis, 2016, 130, 244-263.	2.8	26
87	Renal function, body surface area, and age are associated with risk of early-onset fluoropyrimidine-associated toxicity in patients treated with capecitabine-based anticancer regimens in daily clinical care. European Journal of Cancer, 2016, 54, 120-130.	2.8	40
88	Exposure and Tumor Fn14 Expression as Determinants of Pharmacodynamics of the Anti-TWEAK Monoclonal Antibody RG7212 in Patients with Fn14-Positive Solid Tumors. Clinical Cancer Research, 2016, 22, 858-867.	7.0	28
89	Development of a Tumour Growth Inhibition Model to Elucidate the Effects of Ritonavir on Intratumoural Metabolism and Anti-tumour Effect of Docetaxel in a Mouse Model for Hereditary Breast Cancer. AAPS Journal, 2016, 18, 362-371.	4.4	4
90	EpCAM-based flow cytometry in cerebrospinal fluid greatly improves diagnostic accuracy of leptomeningeal metastases from epithelial tumors. Neuro-Oncology, 2016, 18, 855-862.	1.2	57

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91	Crizotinib-induced fatal fulminant liver failure. Lung Cancer, 2016, 93, 17-19.	2.0	22
92	Liquid chromatography–tandem mass spectrometric assay for the tyrosine kinase inhibitor afatinib in mouse plasma using salting-out liquid–liquid extraction. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2016, 1012-1013, 118-123.	2.3	18
93	Prospective DPYD genotyping to reduce the risk of fluoropyrimidine-induced severe toxicity: Ready for prime time. European Journal of Cancer, 2016, 54, 40-48.	2.8	110
94	Upfront Genotyping of <i>DPYD</i> * <i>2A</i> to Individualize Fluoropyrimidine Therapy: A Safety and Cost Analysis. Journal of Clinical Oncology, 2016, 34, 227-234.	1.6	279
95	Liquid chromatography–tandem mass spectrometric assay for the simultaneous determination of the irreversible BTK inhibitor ibrutinib and its dihydrodiol-metabolite in plasma and its application in mouse pharmacokinetic studies. Journal of Pharmaceutical and Biomedical Analysis, 2016, 118, 123-131.	2.8	39
96	Pharmacodynamics and pharmacokinetics of oral topotecan in patients with advanced solid tumours and impaired renal function. British Journal of Clinical Pharmacology, 2015, 80, 253-266.	2.4	10
97	Tailored Tamoxifen Treatment for Breast Cancer Patients: A Perspective. Clinical Breast Cancer, 2015, 15, 241-244.	2.4	18
98	Variability in bioavailability of small molecular tyrosine kinase inhibitors. Cancer Treatment Reviews, 2015, 41, 412-422.	7.7	103
99	Treatment Individualization in Colorectal Cancer. Current Colorectal Cancer Reports, 2015, 11, 335-344.	0.5	17
100	The use of combinations of monoclonal antibodies in clinical oncology. Cancer Treatment Reviews, 2015, 41, 859-867.	7.7	79
101	A Phase I Monotherapy Study of RG7212, a First-in-Class Monoclonal Antibody Targeting TWEAK Signaling in Patients with Advanced Cancers. Clinical Cancer Research, 2015, 21, 258-266.	7.0	32
102	Liquid chromatography-tandem mass spectrometric assay for the PI3K/mTOR inhibitor GSK2126458 in mouse plasma and tumor homogenate. Journal of Pharmaceutical and Biomedical Analysis, 2015, 107, 403-408.	2.8	4
103	Improved pharmacodynamic assay for dihydropyrimidine dehydrogenase activity in peripheral blood mononuclear cells. Bioanalysis, 2015, 7, 519-529.	1.5	15
104	Long-term safety and anti-tumour activity of olaparib monotherapy after combination with carboplatin and paclitaxel in patients with advanced breast, ovarian or fallopian tube cancer. British Journal of Cancer, 2015, 113, 396-402.	6.4	42
105	Development of an LC–MS/MS assay for the quantitative determination of the intracellular 5-fluorouracil nucleotides responsible for the anticancer effect of 5-fluorouracil. Journal of Pharmaceutical and Biomedical Analysis, 2015, 110, 58-66.	2.8	30
106	Incorporation of concentration data below the limit of quantification in population pharmacokinetic analyses. Pharmacology Research and Perspectives, 2015, 3, e00131.	2.4	127
107	Clinical relevance of DPYD variants c.1679T>C, c.1236G>A/HapB3, and c.1601G>A as predictors of severe fluoropyrimidine-associated toxicity: a systematic review and meta-analysis of individual patient data. Lancet Oncology, The, 2015, 16, 1639-1650.	10.7	277
108	Annexin A1 expression in a pooled breast cancer series: association with tumor subtypes and prognosis. BMC Medicine, 2015, 13, 156.	5.5	51

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109	Observations on Three Endpoint Properties and Their Relationship to Regulatory Outcomes of European Oncology Marketing Applications. Oncologist, 2015, 20, 683-691.	3.7	7
110	Translating <i>DPYD</i> genotype into DPD phenotype: using the <i>DPYD</i> gene activity score. Pharmacogenomics, 2015, 16, 1275-1284.	1.3	81
111	Systematic Review of Biomarkers To Monitor Therapeutic Response in Leishmaniasis. Antimicrobial Agents and Chemotherapy, 2015, 59, 1-14.	3.2	62
112	Phase I/II study with ruthenium compound NAMI-A and gemcitabine in patients with non-small cell lung cancer after first line therapy. Investigational New Drugs, 2015, 33, 201-214.	2.6	327
113	Updated efficacy of the MEK inhibitor trametinib (T), BRAF inhibitor dabrafenib (D), and anti-EGFR antibody panitumumab (P) in patients (pts) with BRAF V600E mutated (BRAFm) metastatic colorectal cancer (mCRC) Journal of Clinical Oncology, 2015, 33, 103-103.	1.6	43
114	Semiphysiological versus Empirical Modelling of the Population Pharmacokinetics of Free and Total Cefazolin during Pregnancy. BioMed Research International, 2014, 2014, 1-9.	1.9	17
115	NT-23 * PHASE 1/2A STUDY OF GLUTATHIONE PEGYLATED LIPOSOMAL DOXORUBICIN (2B3-101) IN BREAST CANCER PATIENTS WITH BRAIN METASTASES (BCBM) OR RECURRENT HIGH GRADE GLIOMAS (HGG). Neuro-Oncology, 2014, 16, v163-v163.	1.2	10
116	Quantitative determination of azacitidine triphosphate in peripheral blood mononuclear cells using liquid chromatography coupled with high-resolution mass spectrometry. Journal of Pharmaceutical and Biomedical Analysis, 2014, 90, 7-14.	2.8	15
117	Effects of low-fat and high-fat meals on steady-state pharmacokinetics of lapatinib in patients with advanced solid tumours. Investigational New Drugs, 2014, 32, 481-488.	2.6	39
118	Practical Guidelines for Therapeutic Drug Monitoring of Anticancer Tyrosine Kinase Inhibitors: Focus on the Pharmacokinetic Targets. Clinical Pharmacokinetics, 2014, 53, 305-325.	3.5	190
119	Predictive Value of CYP3A and ABCB1 Phenotyping Probes for the Pharmacokinetics of Sunitinib: the ClearSun Study. Clinical Pharmacokinetics, 2014, 53, 261-269.	3.5	23
120	Liquid chromatography-tandem mass spectrometric assay for the light sensitive survivin suppressant sepantronium bromide (YM155) in mouse plasma. Journal of Pharmaceutical and Biomedical Analysis, 2014, 92, 144-148.	2.8	2
121	Liquid chromatography–tandem mass spectrometric assay for the PARP inhibitor rucaparib in plasma. Journal of Pharmaceutical and Biomedical Analysis, 2014, 88, 626-629.	2.8	14
122	Phase II and Pharmacological Study of Oral Docetaxel Plus Cyclosporin A in Anthracycline Pre-Treated Metastatic Breast Cancer. Current Clinical Pharmacology, 2014, 9, 139-147.	0.6	9
123	Pharmacodynamic assay of thymidylate synthase activity in peripheral blood mononuclear cells. Analytical and Bioanalytical Chemistry, 2013, 405, 2495-2503.	3.7	7
124	Correction of peripheral blood mononuclear cell cytosolic protein for hemoglobin contamination. Analytical and Bioanalytical Chemistry, 2013, 405, 2391-2395.	3.7	9
125	Phase I study of lonafarnib (SCH66336) in combination with trastuzumab plus paclitaxel in Her2/neu overexpressing breast cancer: EORTC study 16023. Cancer Chemotherapy and Pharmacology, 2013, 71, 53-62.	2.3	21
126	Taxanes: Old drugs, new oral formulations. European Journal of Pharmacology, 2013, 717, 40-46.	3.5	53

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127	Pharmacokinetics of eribulin mesylate in patients with solid tumours receiving repeated oral rifampicin. British Journal of Clinical Pharmacology, 2013, 75, 507-521.	2.4	19
128	Oral Anticancer Drugs: Mechanisms of Low Bioavailability and Strategies for Improvement. Clinical Pharmacokinetics, 2013, 52, 399-414.	3.5	118
129	Concise Drug Review: Azacitidine and Decitabine. Oncologist, 2013, 18, 619-624.	3.7	221
130	Population pharmacokinetic–pharmacodynamic analysis for eribulin mesilateâ€associated neutropenia. British Journal of Clinical Pharmacology, 2013, 76, 412-424.	2.4	31
131	Lapatinib for Advanced or Metastatic Breast Cancer. Oncologist, 2012, 17, 536-542.	3.7	67
132	Concise Drug Review: Pazopanib and Axitinib. Oncologist, 2012, 17, 1081-1089.	3.7	74
133	Letter to the Editor Regarding "A Prospective, Controlled Study of the Botanical Compound Mixture LCS101 for Chemotherapyâ€Induced Hematological Complications in Breast Cancerâ€Iby Yaalâ€Hahoshen et al. (The Oncologist 2011;16:1197–1202). Oncologist, 2012, 17, 740-741.	3.7	2
134	Mass Balance Study of [14C]Eribulin in Patients with Advanced Solid Tumors. Drug Metabolism and Disposition, 2012, 40, 313-321.	3.3	27
135	Evaluation of a Pharmacology-Driven Dosing Algorithm of 3-Weekly Paclitaxel Using Therapeutic Drug Monitoring. Clinical Pharmacokinetics, 2012, 51, 607-617.	3.5	45
136	Disposition and metabolism of 14C-dovitinib (TKI258), an inhibitor of FGFR and VEGFR, after oral administration in patients with advanced solid tumors. Cancer Chemotherapy and Pharmacology, 2012, 70, 653-663.	2.3	16
137	Validation of a multiparameter flow cytometry method for the determination of phosphorylated extracellularâ€signalâ€regulated kinase and DNA in circulating tumor cells. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2012, 81A, 664-671.	1.5	24
138	Bioanalytical aspects of clinical mass balance studies in oncology. Bioanalysis, 2011, 3, 2637-2655.	1.5	6
139	Relationship between Single Nucleotide Polymorphisms and Haplotypes in <i>DPYD</i> and Toxicity and Efficacy of Capecitabine in Advanced Colorectal Cancer. Clinical Cancer Research, 2011, 17, 3455-3468.	7.0	168
140	Circulating tumor cells as pharmacodynamic biomarker in early clinical oncological trials. Cancer Treatment Reviews, 2011, 37, 579-589.	7.7	41
141	Characterization of the in vitro activity of AZD3409, a novel prenyl transferase inhibitor. Cancer Chemotherapy and Pharmacology, 2011, 67, 137-145.	2.3	5
142	Mass spectrometry in the quantitative analysis of therapeutic intracellular nucleotide analogs. Mass Spectrometry Reviews, 2011, 30, 321-343.	5.4	28
143	Trastuzumab. Oncologist, 2011, 16, 800-810.	3.7	84
144	Part 3: Pharmacogenetic Variability in Phase II Anticancer Drug Metabolism. Oncologist, 2011, 16, 992-1005.	3.7	25

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145	Part 1: Background, Methodology, and Clinical Adoption of Pharmacogenetics. Oncologist, 2011, 16, 811-819.	3.7	32
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