

A Population Pharmacokinetic and Pharmacodynamic Clinical Trial in Cancer Patients

Clinical Pharmacokinetics

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Extrapolation of pharmacokinetic interaction data of proton pump inhibitors obtained in healthy subjects for oral targeted therapies in cancer patients. <i>International Journal of Pharmacokinetics</i> , 2018, 3, 93-97.	0.5	0
2	Utilization of data below the analytical limit of quantitation in pharmacokinetic analysis and modeling: promoting interdisciplinary debate. <i>Bioanalysis</i> , 2018, 10, 1229-1248.	1.5	17
3	Abemaciclib Is Active in Preclinical Models of Ewing Sarcoma via Multipronged Regulation of Cell Cycle, DNA Methylation, and Interferon Pathway Signaling. <i>Clinical Cancer Research</i> , 2018, 24, 6028-6039.	7.0	41
4	The role of abemaciclib in treatment of advanced breast cancer. <i>Therapeutic Advances in Medical Oncology</i> , 2018, 10, 175883591877692.	3.2	14
5	Abemaciclib: The Newest CDK4/6 Inhibitor for the Treatment of Breast Cancer. <i>Annals of Pharmacotherapy</i> , 2019, 53, 178-185.	1.9	27
6	Pharmacokinetics and Metabolite Profiling of Trepibutone in Rats Using Ultra-High Performance Liquid Chromatography Combined With Hybrid Quadrupole-Orbitrap and Triple Quadrupole Mass Spectrometers. <i>Frontiers in Pharmacology</i> , 2019, 10, 1266.	3.5	5
7	Drug-drug interactions in breast cancer patients treated with CDK4/6 inhibitors. <i>Cancer Treatment Reviews</i> , 2019, 74, 21-28.	7.7	31
8	Management of targeted therapies in cancer patients with chronic kidney disease, or on haemodialysis: An Associazione Italiana di Oncologia Medica (AIOM)/Societa' Italiana di Nefrologia (SIN) multidisciplinary consensus position paper. <i>Critical Reviews in Oncology/Hematology</i> , 2019, 140, 39-51.	4.4	11
9	Management of toxicities associated with targeted therapies for HR-positive metastatic breast cancer: a multidisciplinary approach is the key to success. <i>Breast Cancer Research and Treatment</i> , 2019, 176, 483-494.	2.5	28
10	CNS penetration of the CDK4/6 inhibitor ribociclib in non-tumor bearing mice and mice bearing pediatric brain tumors. <i>Cancer Chemotherapy and Pharmacology</i> , 2019, 84, 447-452.	2.3	19
11	CDK4/6 Inhibitors Expand the Therapeutic Options in Breast Cancer: Palbociclib, Ribociclib and Abemaciclib. <i>BioDrugs</i> , 2019, 33, 125-135.	4.6	75
12	Involvement of Phosphatase and Tensin Homolog in Cyclin-Dependent Kinase 4/6 Inhibitor-Induced Blockade of Glioblastoma. <i>Frontiers in Pharmacology</i> , 2019, 10, 1316.	3.5	3
13	Drug Exposure to Establish Pharmacokinetic-Response Relationships in Oncology. <i>Clinical Pharmacokinetics</i> , 2020, 59, 123-135.	3.5	5
14	Pharmacokinetic/Pharmacodynamic Modeling of the Anti-Cancer Effect of Dexamethasone in Pancreatic Cancer Xenografts and Anticipation of Human Efficacious Doses. <i>Journal of Pharmaceutical Sciences</i> , 2020, 109, 1169-1177.	3.3	5
15	Clinical Pharmacokinetics and Pharmacodynamics of the Cyclin-Dependent Kinase 4 and 6 Inhibitors Palbociclib, Ribociclib, and Abemaciclib. <i>Clinical Pharmacokinetics</i> , 2020, 59, 1501-1520.	3.5	43
16	Development and Application of a Mechanistic Population Modeling Approach to Describe Abemaciclib Pharmacokinetics. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2020, 9, 523-533.	2.5	7
17	A Phase II Study of Abemaciclib in Patients with Brain Metastases Secondary to Hormone Receptor-Positive Breast Cancer. <i>Clinical Cancer Research</i> , 2020, 26, 5310-5319.	7.0	102
18	CDK4/6 Inhibitors in Breast Cancer Treatment: Potential Interactions with Drug, Gene, and Pathophysiological Conditions. <i>International Journal of Molecular Sciences</i> , 2020, 21, 6350.	4.1	34

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19	Abemaciclib Does Not Have a Clinically Meaningful Effect on Pharmacokinetics of CYP1A2, CYP2C9, CYP2D6, and CYP3A4 Substrates in Patients with Cancer. <i>Drug Metabolism and Disposition</i> , 2020, 48, 796-803.	3.3	8
20	Predicting Clinical Effects of CYP3A4 Modulators on Abemaciclib and Active Metabolites Exposure Using Physiologically Based Pharmacokinetic Modeling. <i>Journal of Clinical Pharmacology</i> , 2020, 60, 915-930.	2.0	34
21	Physiologically Based Pharmacokinetic Modeling of Central Nervous System Pharmacokinetics of CDK4/6 Inhibitors to Guide Selection of Drug and Dosing Regimen for Brain Cancer Treatment. <i>Clinical Pharmacology and Therapeutics</i> , 2021, 109, 494-506.	4.7	16
22	Therapeutic drug monitoring of oral targeted antineoplastic drugs. <i>European Journal of Clinical Pharmacology</i> , 2021, 77, 441-464.	1.9	110
23	Serum concentration of the CKD4/6 inhibitor abemaciclib, but not of creatinine, strongly predicts hematological adverse events in patients with breast cancer: a preliminary report. <i>Investigational New Drugs</i> , 2021, 39, 272-277.	2.6	12
24	Severe hypoglycaemia under abemaciclib administration in a patient with breast cancer: A case report. <i>Molecular and Clinical Oncology</i> , 2021, 14, 61.	1.0	1
25	Simple and Rapid Method for Determination of Abemaciclib in Human Serum using Supported Liquid Extraction Pretreatment and LC-MS/MS Analysis. <i>Indonesian Journal of Pharmaceutics</i> , 2020, 2, .	0.6	0
26	In vitro to Clinical Translation of Combinatorial Effects of Doxorubicin and Abemaciclib in Rb-Positive Triple Negative Breast Cancer: A Systems-Based Pharmacokinetic/Pharmacodynamic Modeling Approach. <i>Breast Cancer: Targets and Therapy</i> , 2021, Volume 13, 87-105.	1.8	5
27	A phase 1b study of the Notch inhibitor crenigacestat (LY3039478) in combination with other anticancer target agents (taladegib, LY3023414, or abemaciclib) in patients with advanced or metastatic solid tumors. <i>Investigational New Drugs</i> , 2021, 39, 1089-1098.	2.6	19
28	Japanese subpopulation analysis of MONARCH 2: phase 3 study of abemaciclib plus fulvestrant for treatment of hormone receptor-positive, human epidermal growth factor receptor 2-negative breast cancer that progressed on endocrine therapy. <i>Breast Cancer</i> , 2021, 28, 1038-1050.	2.9	10
29	Quantification of abemaciclib and metabolites: evolution of bioanalytical methods supporting a novel oncolytic agent. <i>Bioanalysis</i> , 2021, 13, 711-724.	1.5	8
30	CDK4/6 Inhibitors in Melanoma: A Comprehensive Review. <i>Cells</i> , 2021, 10, 1334.	4.1	31
31	Myocardial dysfunction caused by abemaciclib: a case report. <i>International Cancer Conference Journal</i> , 2021, 10, 324-328.	0.5	3
32	Inhibiting CDK4/6 in Breast Cancer with Palbociclib, Ribociclib, and Abemaciclib: Similarities and Differences. <i>Drugs</i> , 2021, 81, 317-331.	10.9	173
33	Abemaciclib is synergistic with doxorubicin in osteosarcoma pre-clinical models via inhibition of CDK4/6â€“Cyclin Dâ€“Rb pathway. <i>Cancer Chemotherapy and Pharmacology</i> , 2022, 89, 31-40.	2.3	7
35	Therapeutic Drug Monitoring of Protein Kinase Inhibitors in Breast Cancer Patients. <i>Prague Medical Report</i> , 2021, 122, 243-256.	0.8	5
36	An UHPLC-MS/MS method for quantification of the CDK4/6 inhibitor abemaciclib in human serum. <i>Journal of Mass Spectrometry and Advances in the Clinical Lab</i> , 2022, 24, 15-21.	2.4	3
37	Therapeutic potential of CDK4/6 inhibitors in renal cell carcinoma. <i>Nature Reviews Urology</i> , 2022, 19, 305-320.	3.8	9

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38	Pharmacokinetics and therapeutic drug monitoring of anticancer protein/kinase inhibitors. <i>Therapie</i> , 2022, 77, 157-170.	1.0	2
39	Abemaciclib in Patients with End-Stage Renal Disease and Advanced Estrogen Receptor-Positive, Human Epidermal Growth Factor Receptor 2-Negative Breast Cancer: A Report of 2 Cases. <i>Case Reports in Oncology</i> , 2022, 15, 305-311.	0.7	2
44	Identification of Novel Mutant (R132H) Isocitrate Dehydrogenase 1 Inhibitors for Glioma Therapy. <i>Journal of Computational Biophysics and Chemistry</i> , 2022, 21, 647-661.	1.7	1
45	Effects of ABCB1 and ABCG2 polymorphisms on the pharmacokinetics of abemaciclib. <i>European Journal of Clinical Pharmacology</i> , 2022, 78, 1239-1247.	1.9	6
46	Development and Validation of a Novel LC-MS/MS Method for the Simultaneous Determination of Abemaciclib, Palbociclib, Ribociclib, Anastrozole, Letrozole, and Fulvestrant in Plasma Samples: A Prerequisite for Personalized Breast Cancer Treatment. <i>Pharmaceuticals</i> , 2022, 15, 614.	3.8	14
47	A New Lc-MS/Ms Method for the Simultaneous Quantification of Abemaciclib, its Main Active Metabolites M2 and M20, and Letrozole for Therapeutic Drug Monitoring. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0
48	The Emerging Role of Cyclin-Dependent Kinase Inhibitors in Treating Diet-Induced Obesity: New Opportunities for Breast and Ovarian Cancers?. <i>Cancers</i> , 2022, 14, 2709.	3.7	2
49	Continuous treatment with abemaciclib leads to sustained and efficient inhibition of breast cancer cell proliferation. <i>Oncotarget</i> , 2022, 13, 864-875.	1.8	6
50	Targeting CDK4 and 6 in Cancer Therapy: Emerging Preclinical Insights Related to Abemaciclib. <i>Oncologist</i> , 0, , .	3.7	11
51	A new LC-MS/MS method for the simultaneous quantification of abemaciclib, its main active metabolites M2 and M20, and letrozole for therapeutic drug monitoring. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2022, 1207, 123403.	2.3	4
52	The Cyclin-Dependent Kinase 4/6 Inhibitor Abemaciclib Is Tolerated Better than Palbociclib by Advanced Breast Cancer Patients with High Serum Albumin Levels. <i>Biological and Pharmaceutical Bulletin</i> , 2022, 45, 1476-1481.	1.4	2
53	Imidazole and Biphenyl Derivatives as Anti-cancer Agents for Glioma Therapeutics: Computational Drug Repurposing Strategy. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2023, 23, 1085-1101.	1.7	2
54	Effects of ABCB1 and ABCG2 Polymorphisms on the Pharmacokinetics of Abemaciclib Metabolites (M2, M20, M18). <i>Anticancer Research</i> , 2023, 43, 1283-1289.	1.1	3
55	Abemaciclib. <i>Journal of Clinical Pharmacy and Therapeutics</i> , 2022, 47, 1-10.	0.4	0
56	Abemaciclib does not increase the corrected QT interval in healthy participants. <i>Clinical and Translational Science</i> , 2023, 16, 1617-1627.	3.1	0