

Ahmed A Othman, Fcp

List of Publications by Year in descending order

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Version: 2024-02-01

61
papers

2,856
citations

293460

24
h-index

198040

52
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62
all docs

62
docs citations

62
times ranked

2177
citing authors

#	ARTICLE	IF	CITATIONS
1	Evaluation of the potential drug interactions mediated through P-gp, OCT2, and MATE1/2K with filgotinib in healthy subjects. <i>Clinical and Translational Science</i> , 2022, 15, 361-370.	1.5	6
2	Assessment of the Effect of Filgotinib on the Pharmacokinetics of Atorvastatin, Pravastatin, and Rosuvastatin in Healthy Adult Participants. <i>Clinical Pharmacology in Drug Development</i> , 2022, 11, 235-245.	0.8	8
3	Evaluation of the potential for pharmacokinetic interaction between tirabrutinib and levonorgestrel/ethinyl estradiol in healthy female volunteers. <i>Clinical and Translational Science</i> , 2022, , .	1.5	1
4	Utility of Modeling and Simulation Approach to Support the Clinical Relevance of Dissolution Specifications: a Case Study from Upadacitinib Development. <i>AAPS Journal</i> , 2022, 24, 39.	2.2	2
5	Oral Glucose Tolerance Test: An Informative Endpoint or an Added Burden in Metformin <sc>Drug-Drug Interaction Studies?. <i>Clinical Pharmacology and Therapeutics</i> , 2022, 112, 453-455.	2.3	4
6	Characterization of the Effect of Upadacitinib on the Pharmacokinetics of Bupropion, a Sensitive Cytochrome P450 2B6 Probe Substrate. <i>Clinical Pharmacology in Drug Development</i> , 2021, 10, 299-306.	0.8	4
7	Exposure-Response Analyses for Upadacitinib Efficacy in Subjects With Atopic Dermatitis- Analyses of Phase 2b Study to Support Selection of Phase 3 Doses. <i>Journal of Clinical Pharmacology</i> , 2021, 61, 628-635.	1.0	8
8	Therapeutic Protein Drug Interaction Potential in Subjects With Psoriasis: An Assessment Based on Population Pharmacokinetic Analyses of Sensitive Cytochrome P450 Probe Substrates. <i>Journal of Clinical Pharmacology</i> , 2021, 61, 307-318.	1.0	3
9	Effect of Upadacitinib on the Pharmacokinetics of Rosuvastatin or Atorvastatin in Healthy Subjects. <i>Clinical Pharmacology in Drug Development</i> , 2021, 10, 1335-1344.	0.8	2
10	Exposure-Response Analyses of the Effects of Venetoclax, a Selective BCL-2 Inhibitor, on B-Lymphocyte and Total Lymphocyte Counts in Women with Systemic Lupus Erythematosus. <i>Clinical Pharmacokinetics</i> , 2020, 59, 335-347.	1.6	5
11	Effects of Upadacitinib Coadministration on the Pharmacokinetics of Sensitive Cytochrome P450 Probe Substrates: A Study With the Modified Cooperstown 5+1 Cocktail. <i>Journal of Clinical Pharmacology</i> , 2020, 60, 86-95.	1.0	13
12	Exposure-Response Relationships for Efficacy and Safety of Risankizumab in Phase II and III Trials in Psoriasis Patients. <i>Clinical Pharmacology and Therapeutics</i> , 2020, 107, 378-387.	2.3	13
13	Preferential Inhibition of JAK1 Relative to JAK3 by Upadacitinib: Exposure-Response Analyses of Ex Vivo Data From 2 Phase 1 Clinical Trials and Comparison to Tofacitinib. <i>Journal of Clinical Pharmacology</i> , 2020, 60, 188-197.	1.0	30
14	Pharmacokinetics of Upadacitinib in Healthy Subjects and Subjects With Rheumatoid Arthritis, Crohn's Disease, Ulcerative Colitis, or Atopic Dermatitis: Population Analyses of Phase 1 and 2 Clinical Trials. <i>Journal of Clinical Pharmacology</i> , 2020, 60, 528-539.	1.0	26
15	Exposure-Response Relationships for the Efficacy and Safety of Risankizumab in Japanese Subjects with Psoriasis. <i>Clinical Pharmacokinetics</i> , 2020, 59, 575-589.	1.6	15
16	Meta-Analyses of Clinical Efficacy of Risankizumab and Adalimumab in Chronic Plaque Psoriasis: Supporting Evidence of Risankizumab Superiority. <i>Clinical Pharmacology and Therapeutics</i> , 2020, 107, 435-442.	2.3	14
17	Exposure-Response Analyses for Upadacitinib Efficacy and Safety in the Crohn's Disease CELEST Study and Bridging to the Extended-Release Formulation. <i>Clinical Pharmacology and Therapeutics</i> , 2020, 107, 639-649.	2.3	10
18	Exposure-Response Analyses of Upadacitinib Efficacy and Safety in Phase II and III Studies to Support Benefit-Risk Assessment in Rheumatoid Arthritis. <i>Clinical Pharmacology and Therapeutics</i> , 2020, 107, 994-1003.	2.3	21

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19	Clinical Pharmacokinetics of Upadacitinib: Review of Data Relevant to the Rheumatoid Arthritis Indication. <i>Clinical Pharmacokinetics</i> , 2020, 59, 531-544.	1.6	38
20	Clinical Pharmacokinetics and Pharmacodynamics of Risankizumab in Psoriasis Patients. <i>Clinical Pharmacokinetics</i> , 2020, 59, 311-326.	1.6	20
21	Upadacitinib in adults with moderate to severe atopic dermatitis: 16-week results from a randomized, placebo-controlled trial. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 877-884.	1.5	242
22	Efficacy of Upadacitinib in a Randomized Trial of Patients With Active Ulcerative Colitis. <i>Gastroenterology</i> , 2020, 158, 2139-2149.e14.	0.6	171
23	Efficacy and Safety of Upadacitinib in a Randomized Trial of Patients With Crohn's Disease. <i>Gastroenterology</i> , 2020, 158, 2123-2138.e8.	0.6	189
24	Efficacy and safety of upadacitinib in Japanese patients with rheumatoid arthritis (SELECT-SUNRISE): a placebo-controlled phase IIb/III study. <i>Rheumatology</i> , 2020, 59, 3303-3313.	0.9	41
25	Population Pharmacokinetics of the Interleukin-23 Inhibitor Risankizumab in Subjects with Psoriasis and Crohn's Disease: Analyses of Phase I and II Trials. <i>Clinical Pharmacokinetics</i> , 2019, 58, 375-387.	1.6	25
26	Upadacitinib Versus Placebo or Adalimumab in Patients With Rheumatoid Arthritis and an Inadequate Response to Methotrexate: Results of a Phase III, Double-Blind, Randomized Controlled Trial. <i>Arthritis and Rheumatology</i> , 2019, 71, 1788-1800.	2.9	284
27	Pharmacokinetics of Risankizumab in Asian Healthy Subjects and Patients With Moderate to Severe Plaque Psoriasis, Generalized Pustular Psoriasis, and Erythrodermic Psoriasis. <i>Journal of Clinical Pharmacology</i> , 2019, 59, 1656-1668.	1.0	24
28	Development of In Vitro-In Vivo Correlation for Upadacitinib Extended-Release Tablet Formulation. <i>AAPS Journal</i> , 2019, 21, 108.	2.2	15
29	Exposure-Response Analyses of Upadacitinib Efficacy in Phase II Trials in Rheumatoid Arthritis and Basis for Phase III Dose Selection. <i>Clinical Pharmacology and Therapeutics</i> , 2019, 106, 1319-1327.	2.3	13
30	Upadacitinib as monotherapy in patients with active rheumatoid arthritis and inadequate response to methotrexate (SELECT-MONOTHERAPY): a randomised, placebo-controlled, double-blind phase 3 study. <i>Lancet</i> , The, 2019, 393, 2303-2311.	6.3	237
31	Population Pharmacokinetics of Risankizumab in Healthy Volunteers and Subjects with Moderate to Severe Plaque Psoriasis: Integrated Analyses of Phase I-III Clinical Trials. <i>Clinical Pharmacokinetics</i> , 2019, 58, 1309-1321.	1.6	25
32	Population Pharmacokinetics of Upadacitinib Using the Immediate-Release and Extended-Release Formulations in Healthy Subjects and Subjects with Rheumatoid Arthritis: Analyses of Phase I-III Clinical Trials. <i>Clinical Pharmacokinetics</i> , 2019, 58, 1045-1058.	1.6	42
33	Models of Variability and Circadian Rhythm in Heart Rate, Blood Pressure, and QT Interval for Healthy Subjects Who Received Placebo in Phase I Trials. <i>Clinical and Translational Science</i> , 2019, 12, 470-480.	1.5	8
34	Characterization of the Effect of Hepatic Impairment on Upadacitinib Pharmacokinetics. <i>Journal of Clinical Pharmacology</i> , 2019, 59, 1188-1194.	1.0	16
35	Pharmacokinetics, Safety, and Tolerability of the Dual Inhibitor of Tumor Necrosis Factor- α and Interleukin 17A, ABBV-257, in Healthy Volunteers and Patients With Rheumatoid Arthritis. <i>Clinical Pharmacology in Drug Development</i> , 2019, 8, 492-502.	0.8	1
36	Lack of Effect of 12-Week Treatment with Risankizumab on the Pharmacokinetics of Cytochrome P450 Probe Substrates in Patients with Moderate to Severe Chronic Plaque Psoriasis. <i>Clinical Pharmacokinetics</i> , 2019, 58, 805-814.	1.6	25

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37	Characterization of the Effect of Renal Impairment on Upadacitinib Pharmacokinetics. <i>Journal of Clinical Pharmacology</i> , 2019, 59, 856-862.	1.0	29
38	Exposure-response analyses demonstrate no evidence of interleukin 17A contribution to efficacy of ABT-122 in rheumatoid or psoriatic arthritis. <i>Rheumatology</i> , 2019, 58, 352-360.	0.9	17
39	The JAK1 Inhibitor Upadacitinib Has No Effect on the Pharmacokinetics of Levonorgestrel and Ethinylestradiol: A Study in Healthy Female Subjects. <i>Journal of Clinical Pharmacology</i> , 2019, 59, 510-516.	1.0	14
40	Pharmacokinetics of Upadacitinib With the Clinical Regimens of the Extended-Release Formulation Utilized in Rheumatoid Arthritis Phase 3 Trials. <i>Clinical Pharmacology in Drug Development</i> , 2019, 8, 208-216.	0.8	42
41	Pharmacokinetics of the B-Cell Lymphoma 2 (Bcl-2) Inhibitor Venetoclax in Female Subjects with Systemic Lupus Erythematosus. <i>Clinical Pharmacokinetics</i> , 2018, 57, 1185-1198.	1.6	7
42	Metabolism and Disposition of a Novel Selective $\alpha 7$ Neuronal Acetylcholine Receptor Agonist ABT-126 in Humans: Characterization of the Major Roles for Flavin-Containing Monooxygenases and UDP-Glucuronosyl Transferase 1A4 and 2B10 in Catalysis. <i>Drug Metabolism and Disposition</i> , 2018, 46, 429-439.	1.7	5
43	Population Pharmacokinetics of the TNF- α and IL-17A Dual-Variable Domain Antibody ABT-122 in Healthy Volunteers and Subjects With Psoriatic or Rheumatoid Arthritis: Analysis of Phase 1 and 2 Clinical Trials. <i>Journal of Clinical Pharmacology</i> , 2018, 58, 803-813.	1.0	11
44	Pharmacokinetics of ABT-122, a TNF- α and IL-17A-Targeted Dual-Variable Domain Immunoglobulin, in Healthy Subjects and Patients with Rheumatoid Arthritis: Results from Three Phase I Trials. <i>Clinical Pharmacokinetics</i> , 2018, 57, 613-623.	1.6	17
45	Population Pharmacokinetics of Upadacitinib in Healthy Subjects and Subjects with Rheumatoid Arthritis: Analyses of Phase I and II Clinical Trials. <i>Clinical Pharmacokinetics</i> , 2018, 57, 977-988.	1.6	64
46	Use of Early Clinical Trial Data to Support Thorough QT Study Waiver for Upadacitinib and Utility of Food Effect to Demonstrate ECG Assay Sensitivity. <i>Clinical Pharmacology and Therapeutics</i> , 2018, 103, 836-842.	2.3	13
47	Dual inhibition of tumour necrosis factor and interleukin-17A with ABT-122: open-label long-term extension studies in rheumatoid arthritis or psoriatic arthritis. <i>Rheumatology</i> , 2018, 57, 1972-1981.	0.9	30
48	Safety and efficacy of upadacitinib in patients with rheumatoid arthritis and inadequate response to conventional synthetic disease-modifying anti-rheumatic drugs (SELECT-NEXT): a randomised, double-blind, placebo-controlled phase 3 trial. <i>Lancet</i> , The, 2018, 391, 2503-2512.	6.3	280
49	Assessment of effect of CYP3A inhibition, CYP induction, OATP1B inhibition, and high-fat meal on pharmacokinetics of the JAK1 inhibitor upadacitinib. <i>British Journal of Clinical Pharmacology</i> , 2017, 83, 2242-2248.	1.1	49
50	Levodopa-Carbidopa Intestinal Gel Pharmacokinetics: Lower Variability than Oral Levodopa-Carbidopa. <i>Journal of Parkinson's Disease</i> , 2017, 7, 275-278.	1.5	29
51	Use of a Novel Artificial Intelligence Platform on Mobile Devices to Assess Dosing Compliance in a Phase 2 Clinical Trial in Subjects With Schizophrenia. <i>JMIR MHealth and UHealth</i> , 2017, 5, e18.	1.8	87
52	Therapeutic protein-drug interaction assessment for daclizumab high-yield process in patients with multiple sclerosis using a cocktail approach. <i>British Journal of Clinical Pharmacology</i> , 2016, 82, 160-167.	1.1	25
53	Population Pharmacokinetics of Daclizumab High-Yield Process in Healthy Volunteers and Subjects with Multiple Sclerosis: Analysis of Phase III Clinical Trials. <i>Clinical Pharmacokinetics</i> , 2016, 55, 943-955.	1.6	18
54	A Phase IIb Study of ABT-494, a Selective JAK1 Inhibitor, in Patients With Rheumatoid Arthritis and an Inadequate Response to Anti-Tumor Necrosis Factor Therapy. <i>Arthritis and Rheumatology</i> , 2016, 68, 2867-2877.	2.9	149

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55	Efficacy and Safety of ABT-494, a Selective JAK1 Inhibitor, in a Phase IIb Study in Patients With Rheumatoid Arthritis and an Inadequate Response to Methotrexate. <i>Arthritis and Rheumatology</i> , 2016, 68, 2857-2866.	2.9	172
56	Pharmacokinetics, Safety and Tolerability of ABT-494, a Novel Selective JAK 1 Inhibitor, in Healthy Volunteers and Subjects with Rheumatoid Arthritis. <i>Clinical Pharmacokinetics</i> , 2016, 55, 1547-1558.	1.6	92
57	Blockade of the High-Affinity Interleukin-2 Receptors with Daclizumab High-Yield Process: Pharmacokinetic/Pharmacodynamic Analysis of Single- and Multiple-Dose Phase I Trials. <i>Clinical Pharmacokinetics</i> , 2016, 55, 121-130.	1.6	9
58	Jejunal Infusion of Levodopa/Carbidopa Intestinal Gel Versus Oral Administration of Levodopa/Carbidopa Tablets in Japanese Subjects with Advanced Parkinson's Disease: Pharmacokinetics and Pilot Efficacy and Safety. <i>Clinical Pharmacokinetics</i> , 2015, 54, 975-984.	1.6	18
59	Population pharmacokinetics of levodopa in subjects with advanced Parkinson's disease: levodopa/carbidopa intestinal gel infusion vs oral tablets. <i>British Journal of Clinical Pharmacology</i> , 2014, 78, 94-105.	1.1	37
60	Population Pharmacokinetics of Daclizumab High-Yield Process in Healthy Volunteers: Integrated Analysis of Intravenous and Subcutaneous, Single- and Multiple-Dose Administration. <i>Clinical Pharmacokinetics</i> , 2014, 53, 907-918.	1.6	28
61	The H3 antagonist ABT-288 is tolerated at significantly higher exposures in subjects with schizophrenia than in healthy volunteers. <i>British Journal of Clinical Pharmacology</i> , 2014, 77, 965-974.	1.1	13