

Hao-Jie Zhu

List of Publications by Year in descending order

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87
papers

2,911
citations

147566

31
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182168

51
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89
all docs

89
docs citations

89
times ranked

3414
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 1 | Transcriptional Regulation of Carboxylesterase 1 in Human Liver: Role of the Nuclear Receptor Subfamily 1 Group H Member 3 and Its Splice Isoforms. <i>Drug Metabolism and Disposition</i> , 2022, 50, 43-48. | 1.7 | 0 |
| 2 | Plasma Carboxylesterase 1 Predicts Methylphenidate Exposure: A Proof-of-Concept Study Using Plasma Protein Biomarker for Hepatic Drug Metabolism. <i>Clinical Pharmacology and Therapeutics</i> , 2022, 111, 878-885. | 2.3 | 3 |
| 3 | Effects of Overexpression of Fibroblast Growth Factor 15/19 on Hepatic Drug Metabolizing Enzymes. <i>Drug Metabolism and Disposition</i> , 2022, 50, 468-477. | 1.7 | 2 |
| 4 | Contributions of Cathepsin A and Carboxylesterase 1 to the Hydrolysis of Tenofovir Alafenamide in the Human Liver, and the Effect of CES1 Genetic Variation on Tenofovir Alafenamide Hydrolysis. <i>Drug Metabolism and Disposition</i> , 2022, 50, 243-248. | 1.7 | 1 |
| 5 | Physiologically-based pharmacokinetic modeling to predict methylphenidate exposure affected by interplay among carboxylesterase 1 pharmacogenetics, drug-drug interactions, and sex. <i>Journal of Pharmaceutical Sciences</i> , 2022, , . | 1.6 | 0 |
| 6 | Genome-Wide Association Study for the Genetic Determinants of Thiopurine S-Methyltransferase Protein Expression in the Liver. <i>FASEB Journal</i> , 2022, 36, . | 0.2 | 0 |
| 7 | Impact of carboxylesterase 1 genetic polymorphism on trandolapril activation in human liver and the pharmacokinetics and pharmacodynamics in healthy volunteers. <i>Clinical and Translational Science</i> , 2021, 14, 1380-1389. | 1.5 | 4 |
| 8 | Tissue-Specific Proteomics Analysis of Anti-COVID-19 Nucleoside and Nucleotide Prodrug-Activating Enzymes Provides Insights into the Optimization of Prodrug Design and Pharmacotherapy Strategy. <i>ACS Pharmacology and Translational Science</i> , 2021, 4, 870-887. | 2.5 | 9 |
| 9 | Effect of CES1 genetic variation on enalapril steady-state pharmacokinetics and pharmacodynamics in healthy subjects. <i>British Journal of Clinical Pharmacology</i> , 2021, 87, 4691-4700. | 1.1 | 5 |
| 10 | Data-independent acquisition (DIA): An emerging proteomics technology for analysis of drug-metabolizing enzymes and transporters. <i>Drug Discovery Today: Technologies</i> , 2021, 39, 49-56. | 4.0 | 26 |
| 11 | Developing a SWATH capillary LC-MS/MS method for simultaneous therapeutic drug monitoring and untargeted metabolomics analysis of neonatal plasma. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2021, 1179, 122865. | 1.2 | 5 |
| 12 | Intranasal lipopolysaccharide administration prevents chronic stress-induced depression- and anxiety-like behaviors in mice. <i>Neuropharmacology</i> , 2021, 200, 108816. | 2.0 | 13 |
| 13 | Activation of Tenofovir Alafenamide and Sofosbuvir in the Human Lung and Its Implications in the Development of Nucleoside/Nucleotide Prodrugs for Treating SARS-CoV-2 Pulmonary Infection. <i>Pharmaceutics</i> , 2021, 13, 1656. | 2.0 | 7 |
| 14 | Comparative Proteomics Analysis of Human Liver Microsomes and S9 Fractions. <i>Drug Metabolism and Disposition</i> , 2020, 48, 31-40. | 1.7 | 30 |
| 15 | Biliary Excretion-Mediated Food Effects and Prediction. <i>AAPS Journal</i> , 2020, 22, 124. | 2.2 | 7 |
| 16 | FRACPREDA-2D-PRM: A Fraction Prediction Algorithm-Assisted 2D Liquid Chromatography-Based Parallel Reaction Monitoring-Mass Spectrometry Approach for Measuring Low-Abundance Proteins in Human Plasma. <i>Proteomics</i> , 2020, 20, 2000175. | 1.3 | 3 |
| 17 | Genome-wide pQTL analysis of protein expression regulatory networks in the human liver. <i>BMC Biology</i> , 2020, 18, 97. | 1.7 | 49 |
| 18 | Carboxylesterase 1 and Precision Pharmacotherapy: Pharmacogenetics and Nongenetic Regulators. <i>Drug Metabolism and Disposition</i> , 2020, 48, 230-244. | 1.7 | 62 |

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|----|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 19 | Liquid Chromatography-Tandem Mass Spectrometry (LC-MS/MS)-Based Proteomics of Drug-Metabolizing Enzymes and Transporters. <i>Molecules</i> , 2020, 25, 2718. | 1.7 | 21 |
| 20 | Chemoproteomic Identification of Serine Hydrolase RBBP9 as a Valacyclovir-Activating Enzyme. <i>Molecular Pharmaceutics</i> , 2020, 17, 1706-1714. | 2.3 | 9 |
| 21 | Pharmacokinetics of gemcitabine and its amino acid ester prodrug following intravenous and oral administrations in mice. <i>Biochemical Pharmacology</i> , 2020, 180, 114127. | 2.0 | 13 |
| 22 | Antidepressive properties of microglial stimulation in a mouse model of depression induced by chronic unpredictable stress. <i>Progress in Neuro-Psychopharmacology and Biological Psychiatry</i> , 2020, 101, 109931. | 2.5 | 28 |
| 23 | Acetaminophen-Induced Liver Injury Alters Expression and Activities of Cytochrome P450 Enzymes in an Age-Dependent Manner in Mouse Liver. <i>Drug Metabolism and Disposition</i> , 2020, 48, 326-336. | 1.7 | 25 |
| 24 | Absolute Quantitation of Drug-Metabolizing Cytochrome P450 Enzymes and Accessory Proteins in Dog Liver Microsomes Using Label-Free Standard-Free Analysis Reveals Interbreed Variability. <i>Drug Metabolism and Disposition</i> , 2019, 47, 1314-1324. | 1.7 | 24 |
| 25 | Label-free absolute protein quantification with data-independent acquisition. <i>Journal of Proteomics</i> , 2019, 200, 51-59. | 1.2 | 60 |
| 26 | Potential Regulation of UGT2B10 and UGT2B7 by miR-485-5p in Human Liver. <i>Molecular Pharmacology</i> , 2019, 96, 674-682. | 1.0 | 6 |
| 27 | Response to the Comments on "Determining Allele-Specific Protein Expression (ASPE) Using a Novel Quantitative Concatamer Proteomics Method", <i>Journal of Proteome Research</i> , 2019, 18, 1458-1459. | 1.8 | 0 |
| 28 | Functional Study of Carboxylesterase 1 Protein Isoforms. <i>Proteomics</i> , 2019, 19, e1800288. | 1.3 | 13 |
| 29 | Crataegus Special Extract WS 1442 Effects on eNOS and microRNA 155. <i>Planta Medica</i> , 2018, 84, 1094-1100. | 0.7 | 4 |
| 30 | Comparison of protein expression between human livers and the hepatic cell lines HepG2, Hep3B, and Huh7 using SWATH and MRM-HR proteomics: Focusing on drug-metabolizing enzymes. <i>Drug Metabolism and Pharmacokinetics</i> , 2018, 33, 133-140. | 1.1 | 42 |
| 31 | A sensitive liquid chromatography-tandem mass spectrometry method for the quantification of valacyclovir and its metabolite acyclovir in mouse and human plasma. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2018, 1092, 447-452. | 1.2 | 9 |
| 32 | Determining Allele-Specific Protein Expression (ASPE) Using a Novel Quantitative Concatamer Based Proteomics Method. <i>Journal of Proteome Research</i> , 2018, 17, 3606-3612. | 1.8 | 20 |
| 33 | Effect of biphenyl hydrolase-like (BPHL) gene disruption on the intestinal stability, permeability and absorption of valacyclovir in wildtype and Bphl knockout mice. <i>Biochemical Pharmacology</i> , 2018, 156, 147-156. | 2.0 | 4 |
| 34 | Consequences of Phenytoin Exposure on Hepatic Cytochrome P450 Expression during Postnatal Liver Maturation in Mice. <i>Drug Metabolism and Disposition</i> , 2018, 46, 1241-1250. | 1.7 | 7 |
| 35 | Short- and Long-term Effects of Phenytoin Exposure on the Liver Proteome of Neonatal and Adult Mice Using SWATH-MS Technology. <i>FASEB Journal</i> , 2018, 32, 563.2. | 0.2 | 0 |
| 36 | Influence of peptide transporter 2 (PEPT2) on the distribution of cefadroxil in mouse brain: A microdialysis study. <i>Biochemical Pharmacology</i> , 2017, 131, 89-97. | 2.0 | 21 |

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|----|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 37 | Ethanol Interactions With Dexmethylphenidate and dl-Methylphenidate Spheroidal Oral Drug Absorption Systems in Healthy Volunteers. <i>Journal of Clinical Psychopharmacology</i> , 2017, 37, 419-428. | 0.7 | 16 |
| 38 | A Comprehensive Functional Assessment of Carboxylesterase 1 Nonsynonymous Polymorphisms. <i>Drug Metabolism and Disposition</i> , 2017, 45, 1149-1155. | 1.7 | 24 |
| 39 | Institutional profile of pharmacogenetics within University of Michigan College of Pharmacy. <i>Pharmacogenomics</i> , 2017, 18, . | 0.6 | 2 |
| 40 | Targeted absolute quantitative proteomics with SILAC internal standards and unlabeled full-length protein calibrators (TAQSI). <i>Rapid Communications in Mass Spectrometry</i> , 2016, 30, 553-561. | 0.7 | 24 |
| 41 | Regulatory effects of genomic translocations at the human carboxylesterase-1 (CES1) gene locus. <i>Pharmacogenetics and Genomics</i> , 2016, 26, 197-207. | 0.7 | 18 |
| 42 | Dabigatran etexilate activation is affected by the CES1 genetic polymorphism G143E (rs71647871) and gender. <i>Biochemical Pharmacology</i> , 2016, 119, 76-84. | 2.0 | 72 |
| 43 | Association of Oseltamivir Activation with Gender and Carboxylesterase 1 Genetic Polymorphisms. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2016, 119, 555-561. | 1.2 | 33 |
| 44 | Sacubitril Is Selectively Activated by Carboxylesterase 1 (CES1) in the Liver and the Activation Is Affected by CES1 Genetic Variation. <i>Drug Metabolism and Disposition</i> , 2016, 44, 554-559. | 1.7 | 54 |
| 45 | CES1P1 variant $\delta^{816A}>C$ is not associated with hepatic carboxylesterase 1 expression and activity or antihypertensive effect of trandolapril. <i>European Journal of Clinical Pharmacology</i> , 2016, 72, 681-687. | 0.8 | 11 |
| 46 | Carboxylesterase 1-Mediated Drug-Drug Interactions between Clopidogrel and Simvastatin. <i>Biological and Pharmaceutical Bulletin</i> , 2015, 38, 292-297. | 0.6 | 22 |
| 47 | An ex vivo approach to botanical drug interactions: A proof of concept study. <i>Journal of Ethnopharmacology</i> , 2015, 163, 149-156. | 2.0 | 7 |
| 48 | The influence of the CYP2C19*10 allele on clopidogrel activation and CYP2C19*2 genotyping. <i>Pharmacogenetics and Genomics</i> , 2014, 24, 381-386. | 0.7 | 13 |
| 49 | Clopidogrel Bioactivation and Risk of Bleeding in Patients Cotreated With Angiotensin-Converting Enzyme Inhibitors After Myocardial Infarction: A Proof-of-Concept Study. <i>Clinical Pharmacology and Therapeutics</i> , 2014, 96, 713-722. | 2.3 | 23 |
| 50 | The Effects of Milk Thistle (<i>Silybum marianum</i>) on Human Cytochrome P450 Activity. <i>Drug Metabolism and Disposition</i> , 2014, 42, 1611-1616. | 1.7 | 46 |
| 51 | Carboxylesterase 1 (CES1) genetic polymorphisms and oseltamivir activation. <i>European Journal of Clinical Pharmacology</i> , 2013, 69, 733-734. | 0.8 | 13 |
| 52 | Isopropylphenidate: An Ester Homolog of Methylphenidate with Sustained and Selective Dopaminergic Activity and Reduced Drug Interaction Liability. <i>Journal of Child and Adolescent Psychopharmacology</i> , 2013, 23, 648-654. | 0.7 | 9 |
| 53 | Carboxylesterase 1 as a Determinant of Clopidogrel Metabolism and Activation. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2013, 344, 665-672. | 1.3 | 160 |
| 54 | An Assessment of Pharmacokinetics and Antioxidant Activity of Free Silymarin Flavonolignans in Healthy Volunteers: A Dose Escalation Study. <i>Drug Metabolism and Disposition</i> , 2013, 41, 1679-1685. | 1.7 | 71 |

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|----|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 55 | Limitations of In Vitro Assessments of the Drug Interaction Potential of Botanical Supplements. <i>Planta Medica</i> , 2012, 78, 1421-1427. | 0.7 | 31 |
| 56 | A discriminative analytical method for detection of CES1A1 and CES1A2/CES1A3 genetic variants. <i>Pharmacogenetics and Genomics</i> , 2012, 22, 215-218. | 0.7 | 12 |
| 57 | A sensitive LC-MS/MS assay for the simultaneous analysis of the major active components of silymarin in human plasma. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2012, 902, 1-9. | 1.2 | 41 |
| 58 | Evaluation of organic cation transporter 3 (SLC22A3) inhibition as a potential mechanism of antidepressant action. <i>Pharmacological Research</i> , 2012, 65, 491-496. | 3.1 | 51 |
| 59 | A liquid chromatography/tandem mass spectrometry assay for the analysis of atomoxetine in human plasma and <i>in vitro</i> cellular samples. <i>Biomedical Chromatography</i> , 2012, 26, 1364-1370. | 0.8 | 8 |
| 60 | Prediction and In Vitro Evaluation of Selected Protease Inhibitor Antiviral Drugs as Inhibitors of Carboxylesterase 1: A Potential Source of Drug-Drug Interactions. <i>Pharmaceutical Research</i> , 2012, 29, 972-982. | 1.7 | 27 |
| 61 | An <i>in vitro</i> evaluation of guanfacine as a substrate for P-glycoprotein. <i>Neuropsychiatric Disease and Treatment</i> , 2011, 7, 501. | 1.0 | 3 |
| 62 | Enantiospecific determination of dl-methylphenidate and dl-ethylphenidate in plasma by liquid chromatography-tandem mass spectrometry: Application to human ethanol interactions. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2011, 879, 783-788. | 1.2 | 34 |
| 63 | Identification of selected therapeutic agents as inhibitors of carboxylesterase 1: Potential sources of metabolic drug interactions. <i>Toxicology</i> , 2010, 270, 59-65. | 2.0 | 46 |
| 64 | Interaction of organic cation transporter 3 (SLC22A3) and amphetamine. <i>Journal of Neurochemistry</i> , 2010, 114, 142-149. | 2.1 | 34 |
| 65 | Activation of the Antiviral Prodrug Oseltamivir Is Impaired by Two Newly Identified Carboxylesterase 1 Variants. <i>Drug Metabolism and Disposition</i> , 2009, 37, 264-267. | 1.7 | 75 |
| 66 | Berberine promotes glucagon-like peptide-1 (7 ³⁶) amide secretion in streptozotocin-induced diabetic rats. <i>Journal of Endocrinology</i> , 2009, 200, 159-165. | 1.2 | 101 |
| 67 | Age- and Sex-Related Expression and Activity of Carboxylesterase 1 and 2 in Mouse and Human Liver. <i>Drug Metabolism and Disposition</i> , 2009, 37, 1819-1825. | 1.7 | 83 |
| 68 | Role of carboxylesterase 1 and impact of natural genetic variants on the hydrolysis of trandolapril. <i>Biochemical Pharmacology</i> , 2009, 77, 1266-1272. | 2.0 | 64 |
| 69 | Aripiprazole brain concentration is altered in P-glycoprotein deficient mice. <i>Schizophrenia Research</i> , 2009, 110, 90-94. | 1.1 | 35 |
| 70 | Enantiospecific gas chromatographic-mass spectrometric analysis of urinary methylphenidate: Implications for phenotyping. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2008, 862, 140-149. | 1.2 | 19 |
| 71 | Two CES1 Gene Mutations Lead to Dysfunctional Carboxylesterase 1 Activity in Man: Clinical Significance and Molecular Basis. <i>American Journal of Human Genetics</i> , 2008, 82, 1241-1248. | 2.6 | 202 |
| 72 | Antipsychotic Drugs Inhibit the Function of Breast Cancer Resistance Protein. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2008, 103, 336-341. | 1.2 | 51 |

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|----|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 73 | Interactions of attention-deficit/hyperactivity disorder therapeutic agents with the efflux transporter P-glycoprotein. <i>European Journal of Pharmacology</i> , 2008, 578, 148-158. | 1.7 | 30 |
| 74 | Sertraline and Its Metabolite Desmethylsertraline, but not Bupropion or Its Three Major Metabolites, Have High Affinity for P-Glycoprotein. <i>Biological and Pharmaceutical Bulletin</i> , 2008, 31, 231-234. | 0.6 | 57 |
| 75 | Risperidone and Paliperidone Inhibit P-Glycoprotein Activity In Vitro. <i>Neuropsychopharmacology</i> , 2007, 32, 757-764. | 2.8 | 84 |
| 76 | Methylphenidate and its ethanol transesterification metabolite ethylphenidate: brain disposition, monoamine transporters and motor activity. <i>Behavioural Pharmacology</i> , 2007, 18, 39-51. | 0.8 | 42 |
| 77 | Long-Term Consequences of Methamphetamine Exposure in Young Adults Are Exacerbated in Glial Cell Line-Derived Neurotrophic Factor Heterozygous Mice. <i>Journal of Neuroscience</i> , 2007, 27, 8816-8825. | 1.7 | 66 |
| 78 | Sensitive quantification of atomoxetine in human plasma by HPLC with fluorescence detection using 4-(4,5-diphenyl-1H-imidazole-2-yl) benzoyl chloride derivatization. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2007, 846, 351-354. | 1.2 | 29 |
| 79 | A novel HPLC fluorescence method for the quantification of methylphenidate in human plasma. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2007, 858, 91-95. | 1.2 | 32 |
| 80 | Population pharmacokinetic analysis of drug-drug interactions among risperidone, bupropion, and sertraline in CF1 mice. <i>Psychopharmacology</i> , 2006, 183, 490-499. | 1.5 | 43 |
| 81 | Evaluation of antipsychotic drugs as inhibitors of multidrug resistance transporter P-glycoprotein. <i>Psychopharmacology</i> , 2006, 187, 415-423. | 1.5 | 110 |
| 82 | Characterization of P-glycoprotein Inhibition by Major Cannabinoids from Marijuana. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2006, 317, 850-857. | 1.3 | 157 |
| 83 | THE ROLE OF THE POLYMORPHIC EFFLUX TRANSPORTER P-GLYCOPROTEIN ON THE BRAIN ACCUMULATION OF d-METHYLPHENIDATE AND d-AMPHETAMINE. <i>Drug Metabolism and Disposition</i> , 2006, 34, 1116-1121. | 1.7 | 19 |
| 84 | Pharmacokinetics of Olanzapine After Single-Dose Oral Administration of Standard Tablet Versus Normal and Sublingual Administration of an Orally Disintegrating Tablet in Normal Volunteers. <i>Journal of Clinical Pharmacology</i> , 2006, 46, 164-171. | 1.0 | 53 |
| 85 | Reversal of P-Glycoprotein Mediated Multidrug Resistance in K562 Cell Line by a Novel Synthetic Calmodulin Inhibitor, E6. <i>Biological and Pharmaceutical Bulletin</i> , 2005, 28, 1974-1978. | 0.6 | 20 |
| 86 | Glutamate up-regulates P-glycoprotein expression in rat brain microvessel endothelial cells by an NMDA receptor-mediated mechanism. <i>Life Sciences</i> , 2004, 75, 1313-1322. | 2.0 | 91 |
| 87 | Effect of E6, a novel calmodulin inhibitor, on activity of P-glycoprotein in purified primary cultured rat brain microvessel endothelial cells. <i>Acta Pharmacologica Sinica</i> , 2003, 24, 1143-9. | 2.8 | 11 |