

# C Roland Wolf

## List of Publications by Year in descending order

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39  
papers

1,534  
citations

393982

19  
h-index

329751

37  
g-index

43  
all docs

43  
docs citations

43  
times ranked

2817  
citing authors

#	ARTICLE	IF	CITATIONS
1	Through a glass, darkly? HepaRG and HepG2 cells as models of human phase I drug metabolism. <i>Drug Metabolism Reviews</i> , 2022, 54, 46-62.	1.5	12
2	Nrf2 activation does not affect adenoma development in a mouse model of colorectal cancer. <i>Communications Biology</i> , 2021, 4, 1081.	2.0	1
3	Quantifying ERK activity in response to inhibition of the BRAFV600E-MEK-ERK cascade using mathematical modelling. <i>British Journal of Cancer</i> , 2021, 125, 1552-1560.	2.9	6
4	Glutathione-S-transferase P promotes glycolysis in asthma in association with oxidation of pyruvate kinase M2. <i>Redox Biology</i> , 2021, 47, 102160.	3.9	23
5	Application of hepatic cytochrome b/P450 reductase null (HBRN) mice to study the role of cytochrome b in the cytochrome P450-mediated bioactivation of the anticancer drug ellipticine. <i>Toxicology and Applied Pharmacology</i> , 2019, 366, 64-74.	1.3	2
6	Constitutive Androstane Receptor 1 is Constitutively Bound to Chromatin and 'Primed' for Transactivation in Hepatocytes. <i>Molecular Pharmacology</i> , 2019, 95, 97-105.	1.0	12
7	Discovery of common and rare genetic risk variants for colorectal cancer. <i>Nature Genetics</i> , 2019, 51, 76-87.	9.4	377
8	Drug-induced chromatin accessibility changes associate with sensitivity to liver tumor promotion. <i>Life Science Alliance</i> , 2019, 2, e201900461.	1.3	6
9	Identification of Novel Pathways of Osimertinib Disposition and Potential Implications for the Outcome of Lung Cancer Therapy. <i>Clinical Cancer Research</i> , 2018, 24, 2138-2147.	3.2	21
10	Cytochrome b 5 impacts on cytochrome P450-mediated metabolism of benzo[a]pyrene and its DNA adduct formation: studies in hepatic cytochrome b 5 /P450 reductase null (HBRN) mice. <i>Archives of Toxicology</i> , 2018, 92, 1625-1638.	1.9	26
11	Measuring <i>in vivo</i> responses to endogenous and exogenous oxidative stress using a novel haem oxygenase 1 reporter mouse. <i>Journal of Physiology</i> , 2018, 596, 105-127.	1.3	22
12	Interaction between polymorphisms in aspirin metabolic pathways, regular aspirin use and colorectal cancer risk: A case-control study in unselected white European populations. <i>PLoS ONE</i> , 2018, 13, e0192223.	1.1	5
13	Xenobiotic CAR Activators Induce Dlk1-Dio3 Locus Noncoding RNA Expression in Mouse Liver. <i>Toxicological Sciences</i> , 2017, 158, 367-378.	1.4	7
14	Novel Pathways of Ponatinib Disposition Catalyzed By CYP1A1 Involving Generation of Potentially Toxic Metabolites. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2017, 363, 12-19.	1.3	29
15	Progress in identifying epigenetic mechanisms of xenobiotic-induced non-genotoxic carcinogenesis. <i>Current Opinion in Toxicology</i> , 2017, 3, 62-70.	2.6	7
16	Application of Mice Humanized for CYP2D6 to the Study of Tamoxifen Metabolism and Drug-Drug Interaction with Antidepressants. <i>Drug Metabolism and Disposition</i> , 2017, 45, 17-22.	1.7	14
17	Loss of Tet1-Associated 5-Hydroxymethylcytosine Is Concomitant with Aberrant Promoter Hypermethylation in Liver Cancer. <i>Cancer Research</i> , 2016, 76, 3097-3108.	0.4	71
18	Cyp2c70 is responsible for the species difference in bile acid metabolism between mice and humans. <i>Journal of Lipid Research</i> , 2016, 57, 2130-2137.	2.0	221

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19	Olaparib, Monotherapy or with Ionizing Radiation, Exacerbates DNA Damage in Normal Tissues: Insights from a New p21 Reporter Mouse. <i>Molecular Cancer Research</i> , 2016, 14, 1195-1203.	1.5	24
20	Aldo-keto reductases are biomarkers of NRF2 activity and are co-ordinately overexpressed in non-small cell lung cancer. <i>British Journal of Cancer</i> , 2016, 115, 1530-1539.	2.9	31
21	The Role of Protein-Protein and Protein-Membrane Interactions on P450 Function. <i>Drug Metabolism and Disposition</i> , 2016, 44, 576-590.	1.7	39
22	An Enhanced In Vivo Stable Isotope Labeling by Amino Acids in Cell Culture (SILAC) Model for Quantification of Drug Metabolism Enzymes*. <i>Molecular and Cellular Proteomics</i> , 2015, 14, 750-760.	2.5	7
23	Cytochrome b5 Is a Major Determinant of Human Cytochrome P450 CYP2D6 and CYP3A4 Activity In Vivo. <i>Molecular Pharmacology</i> , 2015, 87, 733-739.	1.0	30
24	The Hepatic Reductase Null (HRN <sup>Δ,ϕ</sup> ) and Reductase Conditional Null (RCN) mouse models as suitable tools to study metabolism, toxicity and carcinogenicity of environmental pollutants. <i>Toxicology Research</i> , 2015, 4, 548-562.	0.9	13
25	Activation Status of the Pregnane X Receptor Influences Vemurafenib Availability in Humanized Mouse Models. <i>Cancer Research</i> , 2015, 75, 4573-4581.	0.4	23
26	Altered Protein S-Glutathionylation Identifies a Potential Mechanism of Resistance to Acetaminophen-Induced Hepatotoxicity. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2015, 355, 137-144.	1.3	25
27	Defining Human Pathways of Drug Metabolism In Vivo through the Development of a Multiple Humanized Mouse Model. <i>Drug Metabolism and Disposition</i> , 2015, 43, 1679-1690.	1.7	22
28	HDAC Inhibitors Increase NRF2-Signaling in Tumour Cells and Blunt the Efficacy of Co-Administered Cytotoxic Agents. <i>PLoS ONE</i> , 2014, 9, e114055.	1.1	21
29	Phenobarbital Induces Cell Cycle Transcriptional Responses in Mouse Liver Humanized for Constitutive Androstane and Pregnane X Receptors. <i>Toxicological Sciences</i> , 2014, 139, 501-511.	1.4	60
30	Phenobarbital-Mediated Tumor Promotion in Transgenic Mice with Humanized CAR and PXR. <i>Toxicological Sciences</i> , 2014, 140, 259-270.	1.4	50
31	Epigenetic profiles as defined signatures of xenobiotic exposure. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , 2014, 764-765, 3-9.	0.9	53
32	In Vivo Regulation of Human Glutathione Transferase GSTP by Chemopreventive Agents. <i>Cancer Research</i> , 2014, 74, 4378-4387.	0.4	35
33	Cytochrome b5 and epoxide hydrolase contribute to benzo[a]pyrene-DNA adduct formation catalyzed by cytochrome P450 1A1 under low NADPH:P450 oxidoreductase conditions. <i>Toxicology</i> , 2014, 318, 1-12.	2.0	41
34	The Gerhard Zbinden memorial lecture. <i>Toxicology Letters</i> , 2002, 127, 3-17.	0.4	9
35	Role of the Conserved Phenylalanine 181 of NADPH-Cytochrome P450 Oxidoreductase in FMN Binding and Catalytic Activity. <i>Biochemistry</i> , 2001, 40, 13439-13447.	1.2	14
36	Determinants of specificity for aflatoxin B1-8,9-epoxide in Alpha-class glutathione S-transferases. <i>Biochemical Journal</i> , 1999, 339, 95-101.	1.7	11

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37	Crystal structure of the FMN-binding domain of human cytochrome P450 reductase at 1.93 Å resolution. <i>Protein Science</i> , 1999, 8, 298-306.	3.1	78
38	<sup>1</sup> H, <sup>15</sup> N and <sup>13</sup> C NMR resonance assignment, secondary structure and global fold of the FMN-binding domain of human cytochrome P450 reductase. <i>Journal of Biomolecular NMR</i> , 1997, 10, 63-75.	1.6	30
39	Increased levels of alpha-class and pi-class glutathione S-transferases in cell lines resistant to 1-chloro-2,4-dinitrobenzene. <i>FEBS Journal</i> , 1993, 217, 671-676.	0.2	19