

# Elaine M Leslie

## List of Publications by Year in descending order

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50  
papers

4,538  
citations

172457

29  
h-index

223800

46  
g-index

50  
all docs

50  
docs citations

50  
times ranked

4551  
citing authors

#	ARTICLE	IF	CITATIONS
1	Multidrug resistance proteins: role of P-glycoprotein, MRP1, MRP2, and BCRP (ABCG2) in tissue defense. <i>Toxicology and Applied Pharmacology</i> , 2005, 204, 216-237.	2.8	1,222
2	Toxicological relevance of the multidrug resistance protein 1, MRP1 (ABCC1) and related transporters. <i>Toxicology</i> , 2001, 167, 3-23.	4.2	364
3	Modulation of Multidrug Resistance Protein 1 (MRP1/ABCC1) Transport and ATPase Activities by Interaction with Dietary Flavonoids. <i>Molecular Pharmacology</i> , 2001, 59, 1171-1180.	2.3	228
4	Arsenic Transport by the Human Multidrug Resistance Protein 1 (MRP1/ABCC1). <i>Journal of Biological Chemistry</i> , 2004, 279, 32700-32708.	3.4	227
5	CFTR directly mediates nucleotide-regulated glutathione flux. <i>EMBO Journal</i> , 2003, 22, 1981-1989.	7.8	193
6	Membrane Topology of the Multidrug Resistance Protein (MRP). <i>Journal of Biological Chemistry</i> , 1997, 272, 23623-23630.	3.4	189
7	Transport of the $^3\text{H}$ -O-Glucuronide Conjugate of the Tobacco-specific Carcinogen 4-(Methylnitrosamino)-1-(3-pyridyl)-1-butanol (NNAL) by the Multidrug Resistance Protein 1 (MRP1). <i>Journal of Biological Chemistry</i> , 2001, 276, 27846-27854.	3.4	147
8	Methods To Evaluate Biliary Excretion of Drugs in Humans: An Updated Review. <i>Molecular Pharmaceutics</i> , 2006, 3, 198-211.	4.6	136
9	Comparative Toxicity of Arsenic Metabolites in Human Bladder Cancer EJ-1 Cells. <i>Chemical Research in Toxicology</i> , 2011, 24, 1586-1596.	3.3	129
10	Bioflavonoid Stimulation of Glutathione Transport by the 190-kDa Multidrug Resistance Protein 1 (MRP1). <i>Drug Metabolism and Disposition</i> , 2003, 31, 11-15.	3.3	125
11	Differential Inhibition of Rat and Human Na <sup>+</sup> -Dependent Taurocholate Cotransporting Polypeptide (NTCP/SLC10A1) by Bosentan: A Mechanism for Species Differences in Hepatotoxicity. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007, 321, 1170-1178.	2.5	119
12	Arsenic-glutathione conjugate transport by the human multidrug resistance proteins (MRPs/ABCCs). <i>Journal of Inorganic Biochemistry</i> , 2012, 108, 141-149.	3.5	119
13	In Vitro-In Vivo Correlation of Hepatobiliary Drug Clearance in Humans. <i>Clinical Pharmacology and Therapeutics</i> , 2007, 81, 406-413.	4.7	117
14	A naturally occurring mutation in MRP1 results in a selective decrease in organic anion transport and in increased doxorubicin resistance. <i>Pharmacogenetics and Genomics</i> , 2002, 12, 321-330.	5.7	112
15	ATPase activity of purified and reconstituted multidrug resistance protein MRP1 from drug-selected H69AR cells. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1999, 1461, 69-82.	2.6	89
16	Use of Tc-99m Mebrofenin as a Clinical Probe to Assess Altered Hepatobiliary Transport: Integration of In Vitro, Pharmacokinetic Modeling, and Simulation Studies. <i>Pharmaceutical Research</i> , 2008, 25, 1851-1860.	3.5	86
17	Use of Sandwich-Cultured Hepatocytes To Evaluate Impaired Bile Acid Transport as a Mechanism of Drug-Induced Hepatotoxicity. <i>Molecular Pharmaceutics</i> , 2007, 4, 911-918.	4.6	80
18	Functional and Structural Consequences of Cysteine Substitutions in the NH <sub>2</sub> Proximal Region of the Human Multidrug Resistance Protein 1 (MRP1/ABCC1). <i>Biochemistry</i> , 2003, 42, 5214-5224.	2.5	73

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19	Monoclonal Antibodies That Inhibit the Transport Function of the 190-kDa Multidrug Resistance Protein, MRP. <i>Journal of Biological Chemistry</i> , 1999, 274, 15420-15426.	3.4	71
20	Cellular arsenic transport pathways in mammals. <i>Journal of Environmental Sciences</i> , 2016, 49, 38-58.	6.1	71
21	Identification of Proline Residues in the Core Cytoplasmic and Transmembrane Regions of Multidrug Resistance Protein 1 (MRP1/ABCC1) Important for Transport Function, Substrate Specificity, and Nucleotide Interactions. <i>Journal of Biological Chemistry</i> , 2004, 279, 12325-12336.	3.4	63
22	Selenium-dependent and -independent transport of arsenic by the human multidrug resistance protein 2 (MRP2/ABCC2): implications for the mutual detoxification of arsenic and selenium. <i>Carcinogenesis</i> , 2010, 31, 1450-1455.	2.8	60
23	Structural Requirements for Functional Interaction of Glutathione Tripeptide Analogs with the Human Multidrug Resistance Protein 1 (MRP1). <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2003, 304, 643-653.	2.5	50
24	Acquired Cadmium Resistance in Metallothionein-I/II(â€“/â€“) Knockout Cells: Role of the T-Type Calcium Channel Cacn1g1 in Cadmium Uptake. <i>Molecular Pharmacology</i> , 2006, 69, 629-639.	2.3	47
25	Nitric oxide prodrugs and metallochemotherapeutics: JS-K and CB-3-100 enhance arsenic and cisplatin cytotoxicity by increasing cellular accumulation. <i>Molecular Cancer Therapeutics</i> , 2004, 3, 709-14.	4.1	47
26	A Novel Pathway for Arsenic Elimination: Human Multidrug Resistance Protein 4 (MRP4/ABCC4) Mediates Cellular Export of Dimethylarsinic Acid (DMA <sup>V</sup> ) and the Diglutathione Conjugate of Monomethylarsonous Acid (MMA <sup>III</sup> ). <i>Molecular Pharmacology</i> , 2014, 86, 168-179.	2.3	45
27	Monomethylarsenic Diglutathione Transport by the Human Multidrug Resistance Protein 1 (MRP1/ABCC1). <i>Drug Metabolism and Disposition</i> , 2011, 39, 2298-2304.	3.3	38
28	Character and Temporal Evolution of Apoptosis in Acetaminophen-Induced Acute Liver Failure*. <i>Critical Care Medicine</i> , 2013, 41, 2543-2550.	0.9	37
29	Polymorphic variants of MRP4/ABCC4 differentially modulate the transport of methylated arsenic metabolites and physiological organic anions. <i>Biochemical Pharmacology</i> , 2016, 120, 72-82.	4.4	32
30	Modulation of trabectedin (ET-743) hepatobiliary disposition by multidrug resistance-associated proteins (Mrps) may prevent hepatotoxicity. <i>Toxicology and Applied Pharmacology</i> , 2008, 228, 17-23.	2.8	27
31	Arsenic Triglutathione [As(GS) <sub>3</sub> ] Transport by Multidrug Resistance Protein 1 (MRP1/ABCC1) Is Selectively Modified by Phosphorylation of Tyr920/Ser921 and Glycosylation of Asn19/Asn23. <i>Molecular Pharmacology</i> , 2016, 90, 127-139.	2.3	21
32	Characterization of Arsenic Hepatobiliary Transport Using Sandwich-Cultured Human Hepatocytes. <i>Toxicological Sciences</i> , 2015, 145, 307-320.	3.1	20
33	Metabolism of a Phenylarsenical in Human Hepatic Cells and Identification of a New Arsenic Metabolite. <i>Environmental Science &amp; Technology</i> , 2018, 52, 1386-1392.	10.0	17
34	Detection of Ophthalmic Acid in Serum from Acetaminophen-Induced Acute Liver Failure Patients Is More Frequent in Non-Survivors. <i>PLoS ONE</i> , 2015, 10, e0139299.	2.5	16
35	Multidrug Resistance Protein 4 (MRP4/ABCC4) Protects Cells from the Toxic Effects of Halobenzoquinones. <i>Chemical Research in Toxicology</i> , 2017, 30, 1815-1822.	3.3	16
36	Biotransformation and transport of the tobacco-specific carcinogen 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) in bile duct-cannulated wild-type and MRP2/ABCC2-deficient (TR) Wistar rats. <i>Carcinogenesis</i> , 2007, 28, 2650-2656.	2.8	15

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37	Accumulation and Transport of Roxarsone, Arsenobetaine, and Inorganic Arsenic Using the Human Immortalized Caco-2 Cell Line. <i>Journal of Agricultural and Food Chemistry</i> , 2016, 64, 8902-8908.	5.2	14
38	The novel p.Ser263Phe mutation in the human high-affinity choline transporter 1 (CHT1/ <i>SLC5A7</i> ) causes a lethal form of fetal akinesia syndrome. <i>Human Mutation</i> , 2019, 40, 1676-1683.	2.5	14
39	Multidrug Resistance Protein 1 (MRP1/ABCC1)-Mediated Cellular Protection and Transport of Methylated Arsenic Metabolites Differs between Human Cell Lines. <i>Drug Metabolism and Disposition</i> , 2018, 46, 1096-1105.	3.3	12
40	Redox metabolism of ingested arsenic: Integrated activities of microbiome and host on toxicological outcomes. <i>Current Opinion in Toxicology</i> , 2019, 13, 90-98.	5.0	11
41	Absolute quantitation of acetaminophen-modified human serum albumin in acute liver failure patients by liquid chromatography/tandem mass spectrometry. <i>Rapid Communications in Mass Spectrometry</i> , 2018, 32, 1573-1582.	1.5	10
42	Glutathione Transferase P1 Interacts Strongly with the Inner Leaflet of the Plasma Membrane. <i>Drug Metabolism and Disposition</i> , 2011, 39, 1122-1126.	3.3	8
43	Human red blood cell uptake and sequestration of arsenite and selenite: Evidence of seleno-bis(S-glutathionyl) arsinium ion formation in human cells. <i>Biochemical Pharmacology</i> , 2020, 180, 114141.	4.4	7
44	Studies of selenium and arsenic mutual protection in human HepG2 cells. <i>Chemico-Biological Interactions</i> , 2020, 327, 109162.	4.0	7
45	Transporters and Toxicity: Insights From the International Transporter Consortium Workshop 4. <i>Clinical Pharmacology and Therapeutics</i> , 2022, 112, 527-539.	4.7	4
46	Biliary excretion of arsenic by human HepaRG cells is stimulated by selenide and mediated by the multidrug resistance protein 2 (MRP2/ABCC2). <i>Biochemical Pharmacology</i> , 2021, 193, 114799.	4.4	2
47	Efflux transporters in anti-cancer drug resistance: Molecular and functional identification and characterization of multidrug resistance proteins (MRPs/ABCCs). , 2020, , 31-65.		1
48	Determining site occupancy of acetaminophen covalent binding to target proteins in vitro. <i>Analytical Science Advances</i> , 2021, 2, 263-271.	2.8	0
49	Selenide stimulates the biliary excretion of arsenic in human HepaRG cells. <i>FASEB Journal</i> , 2021, 35, .	0.5	0
50	Regulation of Arsenic Triglutathione [As(GS) <sub>3</sub> ] Transport by the Human Multidrug Resistance Protein 1 (MRP1/ABCC1) Through Post-translational Modification. <i>FASEB Journal</i> , 2011, 25, 1b502.	0.5	0