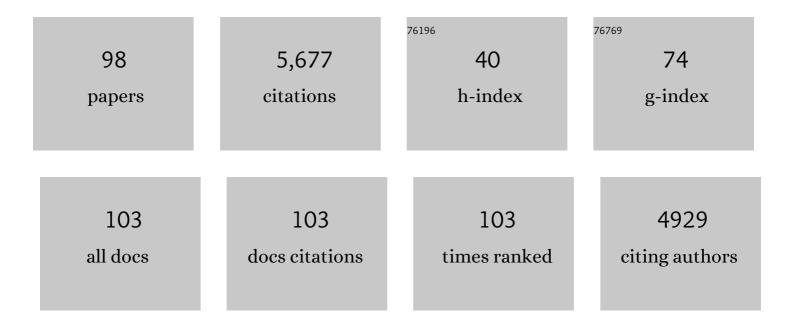
Michael J Devito

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Developmental Exposure to Brominated Diphenyl Ethers Results in Thyroid Hormone Disruption. Toxicological Sciences, 2002, 66, 105-116.	1.4	448
2	Effects of Short-Term in Vivo Exposure to Polybrominated Diphenyl Ethers on Thyroid Hormones and Hepatic Enzyme Activities in Weanling Rats. Toxicological Sciences, 2001, 61, 76-82.	1.4	410
3	Incorporating New Technologies Into Toxicity Testing and Risk Assessment: Moving From 21st Century Vision to a Data-Driven Framework. Toxicological Sciences, 2013, 136, 4-18.	1.4	230
4	Polybrominated Dibenzo-p-Dioxins, Dibenzofurans, and Biphenyls: Inclusion in the Toxicity Equivalency Factor Concept for Dioxin-Like Compounds. Toxicological Sciences, 2013, 133, 197-208.	1.4	197
5	Short-term in vivo exposure to the water contaminant triclosan: Evidence for disruption of thyroxine. Environmental Toxicology and Pharmacology, 2007, 24, 194-197.	2.0	193
6	Thyroid-Hormone–Disrupting Chemicals: Evidence for Dose-Dependent Additivity or Synergism. Environmental Health Perspectives, 2005, 113, 1549-1554.	2.8	179
7	Possible mechanisms of thyroid hormone disruption in mice by BDE 47, a major polybrominated diphenyl ether congener. Toxicology and Applied Pharmacology, 2008, 226, 244-250.	1.3	179
8	In Vitro Metabolism of Pyrethroid Pesticides by Rat and Human Hepatic Microsomes and Cytochrome P450 Isoforms. Drug Metabolism and Disposition, 2009, 37, 221-228.	1.7	161
9	Comparisons of estimated human body burdens of dioxinlike chemicals and TCDD body burdens in experimentally exposed animals Environmental Health Perspectives, 1995, 103, 820-831.	2.8	160
10	Use of toxic equivalency factors for risk assessment for dioxins and related compounds. Toxicology, 1995, 105, 391-401.	2.0	138
11	Identification of Rat and Human Cytochrome P450 Isoforms and a Rat Serum Esterase That Metabolize the Pyrethroid Insecticides Deltamethrin and Esfenvalerate. Drug Metabolism and Disposition, 2007, 35, 1664-1671.	1.7	122
12	Short-term Exposure to Triclosan Decreases Thyroxine In Vivo via Upregulation of Hepatic Catabolism in Young Long-Evans Rats. Toxicological Sciences, 2010, 113, 367-379.	1.4	121
13	An Intuitive Approach for Predicting Potential Human Health Risk with the Tox21 10k Library. Environmental Science & Technology, 2017, 51, 10786-10796.	4.6	120
14	Development of a Refined Database of Mammalian Relative Potency Estimates for Dioxin-like Compounds. Toxicological Sciences, 2006, 89, 4-30.	1.4	115
15	Antiestrogenic action of 2,3,7,8-tetrachlorodibenzo-p-dioxin: Tissue-specific regulation of estrogen receptor in CD1 mice. Toxicology and Applied Pharmacology, 1992, 113, 284-292.	1.3	110
16	Endocrine disrupting chemical emissions from combustion sources: diesel particulate emissions and domestic waste open burn emissions. Atmospheric Environment, 2005, 39, 801-811.	1.9	106
17	Developmental triclosan exposure decreases maternal, fetal, and early neonatal thyroxine: A dynamic and kinetic evaluation of a putative mode-of-action. Toxicology, 2012, 300, 31-45.	2.0	104
18	Species Differences in the in Vitro Metabolism of Deltamethrin and Esfenvalerate: Differential Oxidative and Hydrolytic Metabolism by Humans and Rats. Drug Metabolism and Disposition, 2006, 34, 1764-1771.	1.7	92

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19	Comparative Responsiveness of Hypothyroxinemia and Hepatic Enzyme Induction in Long-Evans Rats Versus C57BL/6J Mice Exposed to TCDD-like and Phenobarbital-like Polychlorinated Biphenyl Congeners. Toxicological Sciences, 2002, 68, 372-380.	1.4	87
20	From the Cover: Three-Dimensional (3D) HepaRG Spheroid Model With Physiologically Relevant Xenobiotic Metabolism Competence and Hepatocyte Functionality for Liver Toxicity Screening. Toxicological Sciences, 2017, 159, 124-136.	1.4	85
21	Physiologically Based Pharmacokinetic Modeling of Deltamethrin: Development of a Rat and Human Diffusion-Limited Model. Toxicological Sciences, 2010, 115, 330-343.	1.4	79
22	Dose–Response Relationships for Disposition and Hepatic Sequestration of Polyhalogenated Dibenzo-p-dioxins, Dibenzofurans, and Biphenyls Following Subchronic Treatment in Mice. Toxicological Sciences, 1998, 46, 223-234.	1.4	78
23	Dose-Response Relationships in Mice Following Subchronic Exposure to 2,3,7,8-Tetrachlorodibenzo-p-dioxin: CYP1A1, CYP1A2, Estrogen Receptor, and Protein Tyrosine Phosphorylation. Toxicology and Applied Pharmacology, 1994, 124, 82-90.	1.3	76
24	A Chemical Category-Based Prioritization Approach for Selecting 75 Per- and Polyfluoroalkyl Substances (PFAS) for Tiered Toxicity and Toxicokinetic Testing. Environmental Health Perspectives, 2019, 127, 14501.	2.8	75
25	Dose–Response Relationships for Polyhalogenated Dioxins and Dibenzofurans Following Subchronic Treatment in Mice. Toxicology and Applied Pharmacology, 1997, 147, 267-280.	1.3	74
26	The Next Generation of Risk Assessment Multi-Year Study—Highlights of Findings, Applications to Risk Assessment, and Future Directions. Environmental Health Perspectives, 2016, 124, 1671-1682.	2.8	74
27	Comparison of the Use of a Physiologically Based Pharmacokinetic Model and a Classical Pharmacokinetic Model for Dioxin Exposure Assessments. Environmental Health Perspectives, 2005, 113, 1666-1668.	2.8	67
28	Developmental triclosan exposure decreases maternal and neonatal thyroxine in rats. Environmental Toxicology and Chemistry, 2010, 29, 2840-2844.	2.2	67
29	Use of a Physiologically Based Pharmacokinetic Model for Rats to Study the Influence of Body Fat Mass and Induction of CYP1A2 on the Pharmacokinetics of TCDD. Environmental Health Perspectives, 2006, 114, 1394-1400.	2.8	64
30	Comprehensive Analyses and Prioritization of Tox21 10K Chemicals Affecting Mitochondrial Function by in-Depth Mechanistic Studies. Environmental Health Perspectives, 2018, 126, 077010.	2.8	60
31	A Pharmacokinetic Model of cis- and trans-Permethrin Disposition in Rats and Humans With Aggregate Exposure Application. Toxicological Sciences, 2012, 130, 33-47.	1.4	58
32	Correlation of tissue concentrations of the pyrethroid bifenthrin with neurotoxicity in the rat. Toxicology, 2011, 290, 1-6.	2.0	56
33	The Power of Resolution: Contextualized Understanding of Biological Responses to Liver Injury Chemicals Using High-throughput Transcriptomics and Benchmark Concentration Modeling. Toxicological Sciences, 2019, 169, 553-566.	1.4	54
34	Toxicology of Dioxins and Related Chemicals. , 1994, , 139-162.		52
35	Evidence for Dose-Additive Effects of Pyrethroids on Motor Activity in Rats. Environmental Health Perspectives, 2009, 117, 1563-1570.	2.8	51
36	Environmentally Relevant Mixtures in Cumulative Assessments: An Acute Study of Toxicokinetics and Effects on Motor Activity in Rats Exposed to a Mixture of Pyrethroids. Toxicological Sciences, 2012, 130, 309-318.	1.4	49

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37	Physiologically Based Pharmacokinetic Model for Developmental Exposures to TCDD in the Rat. Toxicological Sciences, 2004, 80, 115-133.	1.4	47
38	Relative Potencies of Polychlorinated Dibenzo-p-dioxins, Dibenzofurans, and Biphenyls Derived from Hepatic Porphyrin Accumulation in Mice. Toxicology and Applied Pharmacology, 1996, 138, 98-109.	1.3	46
39	Dioxins: model chemicals for assessing receptor-mediated toxicity. Toxicology, 1995, 102, 115-123.	2.0	45
40	Evaluation of 5-day In Vivo Rat Liver and Kidney With High-throughput Transcriptomics for Estimating Benchmark Doses of Apical Outcomes. Toxicological Sciences, 2020, 176, 343-354.	1.4	45
41	Exposure assessment to dioxins from the use of tampons and diapers Environmental Health Perspectives, 2002, 110, 23-28.	2.8	42
42	A Pharmacodynamic Analysis of TCDD-Induced Cytochrome P450 Gene Expression in Multiple Tissues: Dose- and Time-Dependent Effects. Toxicology and Applied Pharmacology, 1998, 151, 294-310.	1.3	41
43	Subchronic Exposure of [3H]-2,3,7,8-Tetrachlorodibenzo-p-dioxin (TCDD) in Female B6C3F1 Mice: Relationship of Steady-State Levels to Disposition and Metabolism. Toxicological Sciences, 2001, 61, 241-255.	1.4	39
44	Inhibition of Human and Rat CYP1A2 by TCDD and Dioxin-like Chemicals. Toxicological Sciences, 2005, 84, 225-231.	1.4	38
45	Comparative Ability of Various PCBs, PCDFs, and TCDD to Induce Cytochrome P450 1A1 and 1A2 Activity Following 4 Weeks of Treatment. Toxicological Sciences, 1993, 20, 125-130.	1.4	37
46	EGF and TGF-alpha Expression Influence the Developmental Toxicity of TCDD: Dose Response and AhR Phenotype in EGF, TGF-alpha, and EGF + TGF-alpha Knockout Mice. Toxicological Sciences, 2003, 71, 84-95.	1.4	37
47	Dose–Response Relationships for Induction of CYP1A1 and CYP1A2 Enzyme Activity in Liver, Lung, and Skin in Female Mice Following Subchronic Exposure to Polychlorinated Biphenyls. Toxicology and Applied Pharmacology, 2000, 167, 157-172.	1.3	35
48	2,3,7,8-Tetrachlorodibenzo-p-dioxin in Pregnant Long Evans Rats: Disposition to Maternal and Embryo/Fetal Tissues. Toxicological Sciences, 1998, 45, 129-136.	1.4	33
49	Repeated dose toxicity and relative potency of 1,2,3,4,6,7-hexachloronaphthalene (PCN 66) 1,2,3,5,6,7-hexachloronaphthalene (PCN 67) compared to 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD) for induction of CYP1A1, CYP1A2 and thymic atrophy in female Harlan Sprague–Dawley rats. Toxicology, 2012. 301. 85-93.	2.0	32
50	Opposite Effects of 2,2′,4,4′,5,5′-Hexachlorobiphenyl and 2,3,7,8-Tetrachlorodibenzo-p-dioxin on the Antibody Response to Sheep Erythrocytes in Mice,. Fundamental and Applied Toxicology, 1997, 37, 141-149.	1.9	31
51	Evaluation and Optimization of Pharmacokinetic Models for <i>in Vitro</i> to <i>in Vivo</i> Extrapolation of Estrogenic Activity for Environmental Chemicals. Environmental Health Perspectives, 2018, 126, 97001.	2.8	31
52	Toxicokinetics of perfluorohexanoic acid (PFHxA), perfluorooctanoic acid (PFOA) and perfluorodecanoic acid (PFDA) in male and female Hsd:Sprague dawley SD rats following intravenous or gavage administration. Xenobiotica, 2020, 50, 722-732.	0.5	31
53	Sensitivity of the SRBC PFC assay versus ELISA for detection of immunosuppression by TCDD and TCDD-like congeners. Toxicology, 2000, 156, 1-11.	2.0	30
54	Induction of Oxidative Stress in Brain Tissues of Mice after Subchronic Exposure to 2,3,7,8-Tetrachlorodibenzo-p-dioxin. Toxicological Sciences, 1998, 42, 23-27.	1.4	28

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55	Coordinated Changes in Xenobiotic Metabolizing Enzyme Gene Expression in Aging Male Rats. Toxicological Sciences, 2008, 106, 263-283.	1.4	28
56	Predictive Modeling of a Mixture of Thyroid Hormone Disrupting Chemicals That Affect Production and Clearance of Thyroxine. International Journal of Toxicology, 2009, 28, 368-381.	0.6	28
57	Induction of Cytochrome P450 Isoenzymes after Toxicokinetic Interactions between 2,3,7,8-Tetrachlorodibenzo-p-dioxin and 2,2′,4,4′,5,5′-Hexachlorobiphenyl in the Liver of the Mouse. Fundamental and Applied Toxicology, 1995, 25, 264-270.	1.9	27
58	Environmentally relevant mixing ratios in cumulative assessments: A study of the kinetics of pyrethroids and their ester cleavage metabolites in blood and brain; and the effect of a pyrethroid mixture on the motor activity of rats. Toxicology, 2014, 320, 15-24.	2.0	25
59	RELATIONSHIP BETWEEN CYP1A ENZYME ACTIVITIES AND PROTEIN LEVELS IN RATS TREATED WITH 2,3,7,8-TETRACHLORODIBENZO-p-DIOXIN. Journal of Toxicology and Environmental Health - Part A: Current Issues, 1996, 47, 379-394.	1.1	24
60	<i>In vitro</i> metabolism of thyroxine by rat and human hepatocytes. Xenobiotica, 2014, 44, 391-403.	0.5	24
61	Human and animal evidence of potential transgenerational inheritance of health effects: An evidence map and state-of-the-science evaluation. Environment International, 2018, 115, 48-69.	4.8	22
62	Environmentally relevant pyrethroid mixtures: A study on the correlation of blood and brain concentrations of a mixture of pyrethroid insecticides to motor activity in the rat. Toxicology, 2016, 359-360, 19-28.	2.0	18
63	Arsenite malignantly transforms human prostate epithelial cells in vitro by gene amplification of mutated KRAS. PLoS ONE, 2019, 14, e0215504.	1.1	16
64	Relative potency based on hepatic enzyme induction predicts immunosuppressive effects of a mixture of PCDDS/PCDFS and PCBS. Toxicology and Applied Pharmacology, 2008, 227, 477-484.	1.3	15
65	Evaluating Sufficient Similarity of Botanical Dietary Supplements: Combining Chemical and In Vitro Biological Data. Toxicological Sciences, 2019, 172, 316-329.	1.4	15
66	Using Tox21 High-Throughput Screening Assays for the Evaluation of Botanical and Dietary Supplements. Applied in Vitro Toxicology, 2019, 5, 10-25.	0.6	15
67	Relative Potency for Altered Humoral Immunity Induced by Polybrominated and Polychlorinated Dioxins/Furans in Female B6C3F1/N Mice. Toxicological Sciences, 2014, 139, 488-500.	1.4	14
68	Follow that botanical: Challenges and recommendations for assessing absorption, distribution, metabolism and excretion of botanical dietary supplements. Food and Chemical Toxicology, 2018, 121, 194-202.	1.8	14
69	Methods for evaluating variability in human health dose–response characterization. Human and Ecological Risk Assessment (HERA), 2020, 26, 1755-1778.	1.7	13
70	Lack of antiandrogenic effects in adult male rats following acute exposure to 2,2-bis(4-chlorophenyl)-1,1-dichloroethylene (p,p′-DDE). Toxicology, 2002, 174, 69-78.	2.0	12
71	Benchmark Concentrations for Untargeted Metabolomics Versus Transcriptomics for Liver Injury Compounds in <i>In Vitro</i> Liver Models. Toxicological Sciences, 2021, 181, 175-186.	1.4	11
72	Exploration of xenobiotic metabolism within cell lines used for Tox21 chemical screening. Toxicology in Vitro, 2021, 73, 105109.	1.1	10

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73	The impact of exposure to a mixture of eighteen polyhalogenated aromatic hydrocarbons on thyroid function: Estimation of an interaction threshold. Journal of Agricultural, Biological, and Environmental Statistics, 2007, 12, 96-111.	0.7	9
74	Development of a Quantitative Model of Pregnane X Receptor (PXR) Mediated Xenobiotic Metabolizing Enzyme Induction. Bulletin of Mathematical Biology, 2010, 72, 1799-1819.	0.9	9
75	Tissue time course and bioavailability of the pyrethroid insecticide bifenthrin in the Long-Evans rat. Xenobiotica, 2016, 46, 430-438.	0.5	9
76	In Vivo Acute Exposure to Polychlorinated Biphenyls: Effects on Free and Total Thyroxine in Rats. International Journal of Toxicology, 2009, 28, 382-391.	0.6	8
77	The effects of 2,2′,4,4′,5,5′-hexachlorobiphenyl cotreatment on the disposition of 2,3,7,8-tetrachlorodibenzo-p-dioxin in mice. Toxicology Letters, 1995, 80, 131-137.	0.4	7
78	Development of a Quantitative Model Incorporating Key Events in a Hepatotoxic Mode of Action to Predict Tumor Incidence. Toxicological Sciences, 2010, 115, 253-266.	1.4	7
79	A PBPK model describing the pharmacokinetics of Î ³ -HBCD exposure in mice. Toxicology and Applied Pharmacology, 2021, 428, 115678.	1.3	7
80	KRAS-retroviral fusion transcripts and gene amplification in arsenic-transformed, human prostate CAsE-PE cancer cells. Toxicology and Applied Pharmacology, 2020, 397, 115017.	1.3	6
81	Toxicology of Dioxins and Dioxinlike Compounds. , 2005, , 137-157.		5
82	Genomic Profiling Reveals Unique Molecular Alterations in Hepatoblastomas and Adjacent Hepatocellular Carcinomas in B6C3F1 Mice. Toxicologic Pathology, 2015, 43, 1114-1126.	0.9	5
83	The Influence of Obesity on the Pharmacokinetics of Dioxin in Mice: An Assessment Using Classical and PBPK Modeling. Toxicological Sciences, 2018, 164, 218-228.	1.4	5
84	The Importance of Pharmacokinetics in Determining the Relative Potency of 2,3,7,8-Tetrachlorodibenzo-p-dioxin and 2,3,7,8-Tetrachlorodibenzofuran. Toxicological Sciences, 1995, 24, 145-148.	1.4	4
85	Mutational analysis of pentabrominated diphenyl-induced hepatocellular tumors in rats and mice, tissue levels of PBDE congeners in rats and mice, and AhR genotyping of Wistar Han rats. Data in Brief, 2018, 21, 2125-2128.	0.5	4
86	Elevated Arsenic and Lead Concentrations in Natural Healing Clay Applied Topically as a Treatment for Ulcerative Dermatitis in Mice. Journal of the American Association for Laboratory Animal Science, 2020, 59, 212-220.	0.6	3
87	Pargyline and naltrexone fail to antagonize the gustatory avoidance response induced by 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine. Drug and Alcohol Dependence, 1986, 18, 293-299.	1.6	2
88	Employing a Mechanistic Model for the Mapk Pathway to Examine the Impact of Cellular all or None Behavior on Overall Tissue Response. Dose-Response, 2010, 8, dose-response.0.	0.7	2
89	Dose-response assessment of the dermal toxicity of Virginia cedarwood oil in F344/N rats and B6C3F1/N mice. Food and Chemical Toxicology, 2016, 98, 159-168.	1.8	2
90	F344/NTac Rats Chronically Exposed to Bromodichloroacetic Acid Develop Mammary Adenocarcinomas With Mixed Luminal/Basal Phenotype and <i>Tgfl²</i> Dysregulation. Veterinary Pathology, 2016, 53, 170-181.	0.8	2

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91	Ascorbic Acid Reduces and Diethyldithiocarbamate Potentiates Methamphetamine-induced Dopamine and Serotonin Depletions. Annals of the New York Academy of Sciences, 1987, 498, 527-529.	1.8	0
92	Subcellular Localization of TCDD Differs between the Liver, Lungs, and Kidneys after Acute and Subchronic Exposure: Species/Dose Comparisons and Possible Mechanism. Toxicological Sciences, 1996, 34, 265-275.	1.4	0
93	Interactive Effects between 2,3,7,8-Tetrachlorodibenzo-p-dioxin and 2,2′,4,4′,5,5′-Hexachlorobiphenyl in Female B6C3F1 Mice: Tissue Distribution and Tissue-Specific Enzyme Induction. Toxicological Sciences, 1996, 34, 118-131.	1.4	0
94	Dose-Response Modeling for 2,3,7,8-Tetrachlorodibenzo-p-Dioxin. , 2005, , 247-298.		0
95	Using a Chemical Mixture of Pyrethroid Pesticides to Determine Rodent Tissue Clearance Rates. Epidemiology, 2011, 22, S249-S250.	1.2	0
96	Extrapolating Dose in Vitro to Dose in Vivo of a Neurotoxic Pyrethroid Pesticide Using Empirical Approaches and a PBPK Model. ACS Symposium Series, 2012, , 229-241.	0.5	0
97	Foreword *. , 2013, , xxiii-xxiv.		0
98	An alternative to TURA. P2 Pollution Prevention Review, 1998, 8, 95-105.	0.0	0