Charlotte Esser

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Aryl Hydrocarbon Receptor Activation by Benzo[a]pyrene Prevents Development of Septic Shock and Fatal Outcome in a Mouse Model of Systemic Salmonella enterica Infection. Cells, 2022, 11, 737.	4.1	4
2	Trajectory Shifts in Interdisciplinary Research of the Aryl Hydrocarbon Receptor—A Personal Perspective on Thymus and Skin. International Journal of Molecular Sciences, 2021, 22, 1844.	4.1	3
3	Functional screening identifies aryl hydrocarbon receptor as suppressor of lung cancer metastasis. Oncogenesis, 2020, 9, 102.	4.9	24
4	The impact of COVIDâ€19 lockâ€downs for European (female) immunologists – our views as members of the EFIS gender and diversity task force. European Journal of Immunology, 2020, 50, 1855-1857.	2.9	5
5	COVID-19 research: toxicological input urgently needed!. Archives of Toxicology, 2020, 94, 2547-2548.	4.2	2
6	Beyond sequencing: fast and easy microbiome profiling by flow cytometry. Archives of Toxicology, 2019, 93, 2703-2704.	4.2	3
7	Proximal <i>Lck</i> Promoter–Driven <i>Cre</i> Function Is Limited in Neonatal and Ineffective in Adult γδT Cell Development. Journal of Immunology, 2019, 203, 569-579.	0.8	19
8	AHR and the issue of immunotoxicity. Current Opinion in Toxicology, 2018, 10, 91-97.	5.0	4
9	Benzo(a)pyrene attenuates the pattern-recognition-receptor induced proinflammatory phenotype of murine macrophages by inducing IL-10 expression in an aryl hydrocarbon receptor-dependent manner. Toxicology, 2018, 409, 80-90.	4.2	14
10	The AHR represses nucleotide excision repair and apoptosis and contributes to UV-induced skin carcinogenesis. Cell Death and Differentiation, 2018, 25, 1823-1836.	11.2	56
11	Aryl hydrocarbon receptor activation by benzo(a)pyrene inhibits proliferation of myeloid precursor cells and alters the differentiation state as well as the functional phenotype of murine bone marrow-derived macrophages. Toxicology Letters, 2018, 296, 106-113.	0.8	16
12	AHR in the skin: From the mediator of chloracne to a therapeutic panacea?. Current Opinion in Toxicology, 2017, 2, 79-86.	5.0	5
13	Indole-3-carbinol, a plant nutrient and AhR-Ligand precursor, supports oral tolerance against OVA and improves peanut allergy symptoms in mice. PLoS ONE, 2017, 12, e0180321.	2.5	29
14	Aryl Hydrocarbon Receptor in Keratinocytes Is Essential for Murine SkinÂBarrier Integrity. Journal of Investigative Dermatology, 2016, 136, 2260-2269.	0.7	97
15	The aryl hydrocarbon receptor promotes aging phenotypes across species. Scientific Reports, 2016, 6, 19618.	3.3	67
16	Balancing intestinal and systemic inflammation through cell type-specific expression of the aryl hydrocarbon receptor repressor. Scientific Reports, 2016, 6, 26091.	3.3	54
17	The Aryl Hydrocarbon Receptor in Immunity: Tools and Potential. Methods in Molecular Biology, 2016, 1371, 239-257.	0.9	36
18	The Aryl Hydrocarbon Receptor in Barrier Organ Physiology, Immunology, and Toxicology. Pharmacological Reviews, 2015, 67, 259-279.	16.0	393

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19	Chemical warfare in the First World War: reflections 100Âyears later. Archives of Toxicology, 2014, 88, 1909-1911.	4.2	23
20	Filling the gaps: need for research on cell-specific xenobiotic metabolism in the skin. Archives of Toxicology, 2013, 87, 1873-1875.	4.2	6
21	Functions of the aryl hydrocarbon receptor in the skin. Seminars in Immunopathology, 2013, 35, 677-691.	6.1	149
22	Natural Aryl Hydrocarbon Receptor Ligands Control Organogenesis of Intestinal Lymphoid Follicles. Science, 2011, 334, 1561-1565.	12.6	706
23	Aryl Hydrocarbon Receptor Is Critical for Homeostasis of Invariant Î ³ δT Cells in the Murine Epidermis. Journal of Immunology, 2011, 187, 3104-3110.	0.8	134
24	2,3,7,8-Tetrachlorodibenzo-p-Dioxin Impairs Stable Establishment of Oral Tolerance in Mice. Toxicological Sciences, 2010, 118, 98-107.	3.1	46
25	Langerhans Cell Maturation and Contact Hypersensitivity Are Impaired in Aryl Hydrocarbon Receptor-Null Mice. Journal of Immunology, 2009, 182, 6709-6717.	0.8	126
26	The immune phenotype of AhR null mouse mutants: Not a simple mirror of xenobiotic receptor over-activation. Biochemical Pharmacology, 2009, 77, 597-607.	4.4	65
27	Small Chemicals, Bioactivation, and the Immune System – A Fragile Balance of iâ€Tox and Benefits?. Chemistry and Biodiversity, 2009, 6, 2138-2143.	2.1	14
28	The aryl hydrocarbon receptor in immunity. Trends in Immunology, 2009, 30, 447-454.	6.8	460
29	Promoter analysis of TCDD-inducible genes in a thymic epithelial cell line indicates the potential for cell-specific transcription factor crosstalk in the AhR response. Toxicology and Applied Pharmacology, 2008, 232, 268-279.	2.8	39
30	A toolbox of novel murine house-keeping genes identified by meta-analysis of large scale gene expression profiles. Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms, 2008, 1779, 830-837.	1.9	34
31	Data sieving analysis as a novel method to asses immunotoxic exposure to dioxins retrospectively. International Immunopharmacology, 2006, 6, 1374-1375.	3.8	1
32	Transcriptional signatures of immune cells in aryl hydrocarbon receptor (AHR)-proficient and AHR-deficient mice. Biological Chemistry, 2006, 387, 1219-26.	2.5	27
33	Role of the aryl hydrocarbon receptor in thymocyte emigrationin vivo. European Journal of Immunology, 2005, 35, 2738-2747.	2.9	37
34	Detection of a novel population of fetal thymocytes characterized by preferential emigration and a TCRÎ ³ Î′+ T cell fate after dioxin exposure. International Immunopharmacology, 2005, 5, 1659-1674.	3.8	10
35	Effects of a single dose of 2,3,7,8-tetrachlorodibenzo-p-dioxin, given at post-puberty, in senescent mice. Toxicology Letters, 2005, 157, 89-98.	0.8	9
36	Signaling via the AHR leads to enhanced usage of CD44v10 by murine fetal thymic emigrants: possible role for CD44 in emigration. International Immunopharmacology, 2004, 4, 805-818.	3.8	12

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37	Identification of dioxin-responsive elements (DREs) in the 5′ regions of putative dioxin-inducible genes. Chemico-Biological Interactions, 1996, 100, 97-112.	4.0	86
38	Evidence for the promotion of positive selection of thymocytes by Ah receptor agonist 2,3,7,8-tetrachlorodibenzo-p-dioxin. European Journal of Pharmacology - Environmental Toxicology and Pharmacology Section, 1995, 293, 413-427.	0.8	25
39	Evidence for the promotion of positive selection of thymocytes by Ah receptor agonist 2,3,7,8-tetrachlorodibenzodioxin. European Journal of Pharmacology, 1995, 293, 413-427.	3.5	3
40	Ontogenic development of murine fetal thymocytes is accelerated by 3,3′,4,4′-tetrachlorobiphenyl. International Journal of Immunopharmacology, 1993, 15, 841-852.	1.1	23