Shiori Hasegawa

List of Publications by Year
in descending order

22	370	1051969 10	889612 19
papers	citations	h-index	g-index

25	25	25	419
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Analysis of chemotherapy-induced peripheral neuropathy using the Japanese Adverse Drug Event Report database. Scientific Reports, 2021, 11, 11324.	1.6	6
2	Analysis of Drug-Induced Gastrointestinal Obstruction and Perforation Using the Japanese Adverse Drug Event Report Database. Frontiers in Pharmacology, 2021, 12, 692292.	1.6	6
3	Gentamicin-induced hearing loss: A retrospective study using the Food and Drug Administration Adverse Event Reporting System and a toxicological study using drugâ^'gene network analysis. Heliyon, 2021, 7, e07429.	1.4	2
4	Pharmacovigilance study of anti-infective-related acute kidney injury using the Japanese adverse drug event report database. BMC Pharmacology & Toxicology, 2021, 22, 47.	1.0	5
5	Analysis of immuneâ€related adverse events caused by immune checkpoint inhibitors using the Japanese Adverse Drug Event Report database. Pharmacoepidemiology and Drug Safety, 2020, 29, 1279-1294.	0.9	28
6	Assessment of Reyeâ€™s syndrome profile with data from the US Food and Drug Administration Adverse Event Reporting System and the Japanese Adverse Drug Event Report databases using the disproportionality analysis. SAGE Open Medicine, 2020, 8, 205031212097417.	0.7	2
7	Analysis of drug-induced interstitial lung disease using the Japanese Adverse Drug Event Report database. SAGE Open Medicine, 2020, 8, 205031212091826.	0.7	17
8	Evaluation of anti-infective-related Clostridium difficile-associated colitis using the Japanese Adverse Drug Event Report database. International Journal of Medical Sciences, 2020, 17, 921-930.	1.1	7
9	Analysis of drug-induced hearing loss by using a spontaneous reporting system database. PLoS ONE, 2019, 14, e0217951.	1.1	21
10	Adverse reaction profiles of hemorrhagic adverse reactions caused by direct oral anticoagulants analyzed using the Food and Drug Administration Adverse Event Reporting System (FAERS) database and the Japanese Adverse Drug Event Report (JADER) database. International Journal of Medical Sciences, 2019, 16, 1295-1303.	1.1	22
11	Adverse event profiles of ifosfamide-induced encephalopathy analyzed using the Food and Drug Administration Adverse Event Reporting System and the Japanese Adverse Drug Event Report databases. Cancer Chemotherapy and Pharmacology, 2019, 84, 1097-1105.	1.1	10
12	Adverse event profiles of solvent-based and nanoparticle albumin-bound paclitaxel formulations using the Food and Drug Administration Adverse Event Reporting System. SAGE Open Medicine, 2019, 7, 205031211983601.	0.7	6
13	Evaluation of pregabalin-induced adverse events related to falls using the FDA adverse event reporting system and Japanese Adverse Drug Event Report databases. Journal of Clinical Pharmacy and Therapeutics, 2019, 44, 285-291.	0.7	7
14	Contraceptives as possible risk factors for postpartum depression: A retrospective study of the food and drug administration adverse event reporting system, 2004â€"2015. Nursing Open, 2018, 5, 131-138.	1.1	11
15	Analysis of fall-related adverse events among older adults using the Japanese Adverse Drug Event Report (JADER) database. Journal of Pharmaceutical Health Care and Sciences, 2018, 4, 32.	0.4	13
16	Analysis of adverse events of renal impairment related to platinum-based compounds using the Japanese Adverse Drug Event Report database. SAGE Open Medicine, 2018, 6, 205031211877247.	0.7	8
17	Adverse events of smoking cessation treatments (nicotine replacement therapy and non-nicotine) Tj ETQq1 1 0.784314 rgBT /Overlock Reporting System, 2004â^'2016. SAGE Open Medicine, 2018, 6, 205031211877795.	0.7	17
18	Evaluation of Drug-Induced Photosensitivity Using the Japanese Adverse Drug Event Report (JADER) Database. Biological and Pharmaceutical Bulletin, 2017, 40, 2158-2165.	0.6	29

#	Article	IF	Citations
19	Thromboembolic adverse event study of combined estrogen-progestin preparations using Japanese Adverse Drug Event Report database. PLoS ONE, 2017, 12, e0182045.	1.1	10
20	Comparison of the adverse event profiles of conventional and liposomal formulations of doxorubicin using the FDA adverse event reporting system. PLoS ONE, 2017, 12, e0185654.	1.1	39
21	Drug-induced gingival hyperplasia: a retrospective study using spontaneous reporting system databases. Journal of Pharmaceutical Health Care and Sciences, 2017, 3, 19.	0.4	56
22	Time-to-Onset Analysis of Drug-Induced Long QT Syndrome Based on a Spontaneous Reporting System for Adverse Drug Events. PLoS ONE, 2016, 11, e0164309.	1.1	36