

Stefania Cheli

List of Publications by Year in descending order

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Version: 2024-02-01

34
papers

397
citations

932766
10
h-index

794141
19
g-index

34
all docs

34
docs citations

34
times ranked

752
citing authors

#	ARTICLE	IF	CITATIONS
1	Fast clearance of anti-TNF \pm agents unrelated to antidrug antibodies: a case report. <i>European Journal of Clinical Pharmacology</i> , 2022, 78, 891-893.	0.8	3
2	Therapeutic drug monitoring and pharmacogenetics of antipsychotics and antidepressants in real life settings: A 5-year single centre experience. <i>World Journal of Biological Psychiatry</i> , 2021, 22, 34-45.	1.3	9
3	Single nucleotide polymorphisms to predict taxanes toxicities and effectiveness in cancer patients. <i>Pharmacogenomics Journal</i> , 2021, 21, 491-497.	0.9	2
4	ABCC4 single-nucleotide polymorphisms as markers of tenofovir disoproxil fumarate-induced kidney impairment. <i>Pharmacogenomics Journal</i> , 2021, 21, 586-593.	0.9	8
5	In linezolid underexposure, pharmacogenetics matters: The role of CYP3A5. <i>Biomedicine and Pharmacotherapy</i> , 2021, 139, 111631.	2.5	8
6	Case of Suboptimal Linezolid Exposure: Is There a Role for Pharmacogenetics?. <i>Therapeutic Drug Monitoring</i> , 2020, 42, 347-348.	1.0	2
7	Neonatal Outcomes in Maternal Depression in Relation to Intrauterine Drug Exposure. <i>Frontiers in Pediatrics</i> , 2019, 7, 309.	0.9	16
8	Is a pharmacogenomic panel useful to estimate the risk of oxaliplatin-related neurotoxicity in colorectal cancer patients?. <i>Pharmacogenomics Journal</i> , 2019, 19, 465-472.	0.9	16
9	ABCB1 c.3435C>T polymorphism is associated with platinum toxicity: a preliminary study. <i>Cancer Chemotherapy and Pharmacology</i> , 2019, 83, 803-808.	1.1	8
10	Pharmacogenetic variants in bipolar disorder with elevated treatment resistance and intolerance: Towards a personalized pattern of care. <i>Bipolar Disorders</i> , 2019, 21, 288-291.	1.1	4
11	Different effects of glucocorticoids on darunavir plasma concentrations. <i>European Journal of Clinical Pharmacology</i> , 2019, 75, 733-735.	0.8	3
12	Clinical and genetic factors associated with increased risk of severe liver toxicity in a monocentric cohort of HIV positive patients receiving nevirapine-based antiretroviral therapy. <i>BMC Infectious Diseases</i> , 2018, 18, 556.	1.3	14
13	Clinical and genetic determinants of nevirapine plasma trough concentration. <i>SAGE Open Medicine</i> , 2018, 6, 205031211878086.	0.7	11
14	Variant alleles in factor V, prothrombin, plasminogen activator inhibitor-1, methylenetetrahydrofolate reductase and risk of thromboembolism in metastatic colorectal cancer patients treated with first-line chemotherapy plus bevacizumab. <i>Pharmacogenomics Journal</i> , 2017, 17, 331-336.	0.9	10
15	Pharmacokinetics and Pharmacogenetics of Selective Serotonin Reuptake Inhibitors During Pregnancy: An Observational Study. <i>Therapeutic Drug Monitoring</i> , 2017, 39, 197-201.	1.0	17
16	Pharmacogenetics-based optimisation of atazanavir treatment: potential role of new genetic predictors. <i>Drug Metabolism and Personalized Therapy</i> , 2017, 32, 115-117.	0.3	0
17	HLA-B*57. <i>Pharmacogenetics and Genomics</i> , 2017, 27, 81-82.	0.7	0
18	Fluoropyrimidine-Associated Toxicity in Two Gastrointestinal Cancer Patients: Potential Role of Common DPYD Polymorphisms. <i>Chemotherapy</i> , 2017, 62, 323-326.	0.8	1

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19	The combination of pharmacogenetic and pharmacokinetic analyses to optimize clomipramine dosing in major depression: a case report. <i>Journal of Clinical Pharmacy and Therapeutics</i> , 2017, 42, 119-121.	0.7	0
20	Renal function in <scp>HIV</scp>/<scp>HBV</scp> coâ€infecte and <scp>HBV</scp> monoâ€infecte patients on a longâ€term treatment with tenofovir in real life setting. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2017, 44, 191-196.	0.9	8
21	Genetic Factors Associated with Platinum Toxicity: A Preliminary Study. <i>Annals of Oncology</i> , 2017, 28, vi95.	0.6	0
22	Pharmacogenetic approach to losartan in Marfan patients: a starting point to improve dosing regimen?. <i>Drug Metabolism and Personalized Therapy</i> , 2016, 31, 157-163.	0.3	5
23	The role of Toll-like receptor 4 polymorphisms in vaccine immune response. <i>Pharmacogenomics Journal</i> , 2016, 16, 96-101.	0.9	12
24	LightSNiP assay is a good strategy for pharmacogenetics test. <i>Frontiers in Pharmacology</i> , 2015, 6, 114.	1.6	5
25	SLC29A1 polymorphism and prediction of anaemia severity in patients with chronic hepatitis C receiving triple therapy with telaprevir. <i>Journal of Antimicrobial Chemotherapy</i> , 2015, 70, 1155-1160.	1.3	9
26	Undetected Toxicity Risk in Pharmacogenetic Testing for Dihydropyrimidine Dehydrogenase. <i>International Journal of Molecular Sciences</i> , 2015, 16, 8884-8895.	1.8	13
27	DPD and UGT1A1 deficiency in colorectal cancer patients receiving triplet chemotherapy with fluoropyrimidines, oxaliplatin and irinotecan. <i>British Journal of Clinical Pharmacology</i> , 2015, 80, 581-588.	1.1	52
28	<i>ITPA</i> and <i>SLC29A1</i> Genotyping for the Prediction of Ribavirin Dose Reduction in Anti-HCV Triple Therapy with Protease Inhibitors. <i>Pharmacology</i> , 2015, 96, 163-166.	0.9	2
29	The first steps towards the era of personalised vaccinology: predicting adverse reactions. <i>Pharmacogenomics Journal</i> , 2015, 15, 284-287.	0.9	14
30	Predictive testing for DPD deficiency in a patient with familial history of fluoropyrimidine-associated toxicity. <i>Personalized Medicine</i> , 2014, 11, 259-262.	0.8	4
31	Levofloxacin-induced seizures in a patient without predisposing risk factors: the impact of pharmacogenetics. <i>European Journal of Clinical Pharmacology</i> , 2013, 69, 1611-1613.	0.8	17
32	ACE inhibitors and ribavirin-associated cough: a common undefined predisposing factor?. <i>European Journal of Clinical Pharmacology</i> , 2013, 69, 743-745.	0.8	4
33	Expression Profiling of FSHD-1 and FSHD-2 Cells during Myogenic Differentiation Evidences Common and Distinctive Gene Dysregulation Patterns. <i>PLoS ONE</i> , 2011, 6, e20966.	1.1	39
34	Remodeling of the chromatin structure of the facioscapulohumeral muscular dystrophy (FSHD) locus and upregulation of FSHD-related gene 1 (FRG1) expression during human myogenic differentiation. <i>BMC Biology</i> , 2009, 7, 41.	1.7	81