

Carmen GÃ³mez-Traseira

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

181
citations

1477746

6
h-index

1281420

11
g-index

11
all docs

11
docs citations

11
times ranked

304
citing authors

#	ARTICLE	IF	CITATIONS
1	Anakinra hypersensitivity reaction in a paediatric patient with autoinflammatory syndrome. <i>Rheumatology</i> , 2021, 61, e17-e18.	0.9	1
2	Drug Provocation Tests for Assessing Antibiotic Hypersensitivity. <i>Pediatric Infectious Disease Journal</i> , 2020, 39, 835-839.	1.1	3
3	Fixed food eruption caused by peanut confirmed by open oral food challenge and in vitro cellular testing. <i>Contact Dermatitis</i> , 2020, 83, 227-229.	0.8	3
4	Psychometric Field Study of Hereditary Angioedema Quality of Life Questionnaire for Adults: HAE-QoL. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2016, 4, 464-473.e4.	2.0	48
5	Trimethoprimâ€“sulfamethoxazole (cotrimoxazole) desensitization in an <scp>HIV</scp>â€“infected 5â€“yrâ€“old girl. <i>Pediatric Allergy and Immunology</i> , 2015, 26, 287-289.	1.1	10
6	Selective hypersensitivity reactions to acetaminophen: A 13-case series. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2014, 2, 343-345.	2.0	16
7	Hereditary angioedema caused by the p.Thr309Lys mutation in the F12 gene: AÂ“multifactorial disease. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 132, 986-989.e5.	1.5	19
8	Hypersensitivity to nabumetone: cross reactivity with naproxen. <i>Annals of Allergy, Asthma and Immunology</i> , 2013, 111, 74-75.	0.5	3
9	Paracetamol-Induced Fixed Drug Eruption at an Unusual Site. <i>Recent Patents on Inflammation and Allergy Drug Discovery</i> , 2013, 7, 268-270.	3.9	6
10	Development of a disease-specific quality of life questionnaire for adult patients with hereditary angioedema due to C1 inhibitor deficiency (HAE-QoL): Spanish multi-centre research project. <i>Health and Quality of Life Outcomes</i> , 2012, 10, 82.	1.0	52
11	Usefulness of abdominal ultrasonography in the follow-up of patients with hereditary C1-inhibitor deficiency. <i>Annals of Allergy, Asthma and Immunology</i> , 2009, 102, 483-486.	0.5	20