

# Sevgi Bilgic Eltan

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7449618/publications.pdf>

Version: 2024-02-01

13  
papers

140  
citations

1478505

6  
h-index

1281871

11  
g-index

13  
all docs

13  
docs citations

13  
times ranked

330  
citing authors

#	ARTICLE	IF	CITATIONS
1	Evolution and long-term outcomes of combined immunodeficiency due to CARMIL2 deficiency. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022, 77, 1004-1019.	5.7	19
2	Adverse COVID-19 outcomes in immune deficiencies: Inequality exists between subclasses. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022, 77, 282-295.	5.7	29
3	Expanding the Clinical and Immunological Phenotypes and Natural History of MALT1 Deficiency. <i>Journal of Clinical Immunology</i> , 2022, 42, 634-652.	3.8	12
4	A boy with a novel homozygous <i>ZAP70</i> mutation presenting with a dermatological phenotype and postnatal decrease in CD8 <sup>+</sup> T cells. <i>Pediatric Allergy and Immunology</i> , 2022, 33, e13756.	2.6	3
5	Clinical and Laboratory Factors Affecting the Prognosis of Severe Combined Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2022, 42, 1036-1050.	3.8	6
6	Comparing the levels of CTLA-4-dependent biological defects in patients with LRBA deficiency and CTLA-4 insufficiency. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022, 77, 3108-3123.	5.7	7
7	A Patient with Novel ICOS Mutation Presented with Progressive Loss of B Cells. <i>Journal of Clinical Immunology</i> , 2021, 41, 251-255.	3.8	5
8	A set of clinical and laboratory markers differentiates hyper-IgE syndrome from severe atopic dermatitis. <i>Clinical Immunology</i> , 2021, 223, 108645.	3.2	7
9	Lymphopenia with Low T and NK Cells in a Patient with USB1 Mutation, Rare Findings in Clericuzio-Type Poikiloderma with Neutropenia. <i>Journal of Clinical Immunology</i> , 2021, 41, 1106-1111.	3.8	0
10	Stepwise Reversal of Immune Dysregulation Due to STAT1 Gain-of-Function Mutation Following Ruxolitinib Bridge Therapy and Transplantation. <i>Journal of Clinical Immunology</i> , 2021, 41, 769-779.	3.8	26
11	Diagnostic Modalities Based on Flow Cytometry for Chronic Granulomatous Disease: A Multicenter Study in a Well-Defined Cohort. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020, 8, 3525-3534.e1.	3.8	7
12	Clinical features and accompanying findings of Pseudo-Bartter Syndrome in cystic fibrosis. <i>Pediatric Pulmonology</i> , 2020, 55, 2011-2016.	2.0	12
13	Management of Systemic Hypersensitivity Reactions to Gonadotropin-Releasing Hormone Analogues during Treatment of Central Precocious Puberty. <i>Hormone Research in Paediatrics</i> , 2020, 93, 66-72.	1.8	7