

# Fernando Corvillo

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/678596/publications.pdf>

Version: 2024-02-01

14  
papers

242  
citations

1163117

8  
h-index

1058476

14  
g-index

16  
all docs

16  
docs citations

16  
times ranked

342  
citing authors

#	ARTICLE	IF	CITATIONS
1	Complement Genetic Variants and FH Desialylation in <i>S. pneumoniae</i> -Haemolytic Uraemic Syndrome. <i>Frontiers in Immunology</i> , 2021, 12, 641656.	4.8	14
2	Complement Factor D (adipsin) Levels Are Elevated in Acquired Partial Lipodystrophy (Barraquer-Simons syndrome). <i>International Journal of Molecular Sciences</i> , 2021, 22, 6608.	4.1	7
3	Immunological features of patients affected by Barraquer-Simons syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 9.	2.7	11
4	The FXII c.-4T>C Polymorphism as a Disease Modifier in Patients With Hereditary Angioedema Due to the FXII p.Thr328Lys Variant. <i>Frontiers in Genetics</i> , 2020, 11, 1033.	2.3	9
5	Evidence of ongoing complement activation on adipose tissue from an 11-year-old girl with Barraquer-Simons syndrome. <i>Journal of Dermatology</i> , 2020, 47, 1439-1444.	1.2	6
6	An overview of lipodystrophy and the role of the complement system. <i>Molecular Immunology</i> , 2019, 112, 223-232.	2.2	21
7	Nephritic Factors: An Overview of Classification, Diagnostic Tools and Clinical Associations. <i>Frontiers in Immunology</i> , 2019, 10, 886.	4.8	52
8	Complement as a diagnostic tool in immunopathology. <i>Seminars in Cell and Developmental Biology</i> , 2019, 85, 86-97.	5.0	33
9	Lipodistrofia parcial adquirida y glomerulopatía C3: la desregulación del sistema del complemento como mecanismo común. <i>Nefrología</i> , 2018, 38, 258-266.	0.4	8
10	Autoantibodies Against Perilipin 1 as a Cause of Acquired Generalized Lipodystrophy. <i>Frontiers in Immunology</i> , 2018, 9, 2142.	4.8	23
11	Three cases of C3 glomerulopathy with anti-factor H autoantibodies. <i>Molecular Immunology</i> , 2018, 102, 209.	2.2	0
12	Acquired partial lipodystrophy and C3 glomerulopathy: Dysregulation of the complement system as a common mechanism. <i>Nefrología</i> , 2018, 38, 258-266.	0.4	6
13	Serum properdin consumption as a biomarker of C5 convertase dysregulation in C3 glomerulopathy. <i>Clinical and Experimental Immunology</i> , 2016, 184, 118-125.	2.6	24
14	Testing the Activity of Complement Convertases in Serum/Plasma for Diagnosis of C4NeF-Mediated C3 Glomerulonephritis. <i>Journal of Clinical Immunology</i> , 2016, 36, 517-527.	3.8	26