

Lourdes Isaac

List of Publications by Year in descending order

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57
papers

1,729
citations

304368

22
h-index

301761

39
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58
all docs

58
docs citations

58
times ranked

1930
citing authors

#	ARTICLE	IF	CITATIONS
1	Systemic Lupus Erythematosus and Deficiencies of Early Components of the Complement Classical Pathway. <i>Frontiers in Immunology</i> , 2016, 7, 55.	2.2	208
2	Leptospiral Immunoglobulin-like Proteins Interact With Human Complement Regulators Factor H, FHL-1, FHR-1, and C4BP. <i>Journal of Infectious Diseases</i> , 2012, 205, 995-1004.	1.9	132
3	Immune Evasion of <i>Leptospira</i> Species by Acquisition of Human Complement Regulator C4BP. <i>Infection and Immunity</i> , 2009, 77, 1137-1143.	1.0	97
4	Functional Characterization of LcpA, a Surface-Exposed Protein of <i>Leptospira</i> spp. That Binds the Human Complement Regulator C4BP. <i>Infection and Immunity</i> , 2010, 78, 3207-3216.	1.0	90
5	Leptospirosis: Aspects of Innate Immunity, Immunopathogenesis and Immune Evasion From the Complement System. <i>Scandinavian Journal of Immunology</i> , 2011, 73, 408-419.	1.3	82
6	Immune Evasion by Pathogenic <i>Leptospira</i> Strains: The Secretion of Proteases that Directly Cleave Complement Proteins. <i>Journal of Infectious Diseases</i> , 2014, 209, 876-886.	1.9	82
7	Interaction of <i>Leptospira</i> Elongation Factor Tu with Plasminogen and Complement Factor H: A Metabolic Leptospiral Protein with Moonlighting Activities. <i>PLoS ONE</i> , 2013, 8, e81818.	1.1	72
8	Plasmin cleaves fibrinogen and the human complement proteins C3b and C5 in the presence of <i>Leptospira</i> interrogans proteins: A new role of LigA and LigB in invasion and complement immune evasion. <i>Immunobiology</i> , 2016, 221, 679-689.	0.8	72
9	Complement Evasion by Pathogenic <i>Leptospira</i> . <i>Frontiers in Immunology</i> , 2016, 7, 623.	2.2	63
10	Pathogenic <i>Leptospira</i> Species Acquire Factor H and Vitronectin via the Surface Protein LcpA. <i>Infection and Immunity</i> , 2015, 83, 888-897.	1.0	57
11	Is immunity in diabetic patients influencing the susceptibility to infections? Immunoglobulins, complement and phagocytic function in children and adolescents with type 1 diabetes mellitus. <i>Pediatric Diabetes</i> , 2005, 6, 206-212.	1.2	49
12	Genetic analysis of complement C1s deficiency associated with systemic lupus erythematosus highlights alternative splicing of normal C1s gene. <i>Molecular Immunology</i> , 2008, 45, 1693-1702.	1.0	44
13	The Serine Protease Pic From Enterohaggregative <i>Escherichia coli</i> Mediates Immune Evasion by the Direct Cleavage of Complement Proteins. <i>Journal of Infectious Diseases</i> , 2015, 212, 106-115.	1.9	41
14	Complement components, regulators and receptors are produced by human monocyte-derived dendritic cells. <i>Immunobiology</i> , 2007, 212, 151-157.	0.8	35
15	Pathogenic <i>Leptospira</i> Secreted Proteases Target the Membrane Attack Complex: A Potential Role for Thermolysin in Complement Inhibition. <i>Frontiers in Microbiology</i> , 2017, 8, 958.	1.5	35
16	Acquisition of negative complement regulators by the saprophyte <i>Leptospira biflexa</i> expressing LigA or LigB confers enhanced survival in human serum. <i>Immunology Letters</i> , 2016, 173, 61-68.	1.1	33
17	Fine Mapping of the Interaction between C4b-Binding Protein and Outer Membrane Proteins LigA and LigB of Pathogenic <i>Leptospira</i> interrogans. <i>PLoS Neglected Tropical Diseases</i> , 2015, 9, e0004192.	1.3	33
18	Plasma levels of complement proteins from the alternative pathway in patients with age-related macular degeneration are independent of Complement Factor H Tyr442His polymorphism. <i>Molecular Vision</i> , 2012, 18, 2288-99.	1.1	31

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19	A double edged-sword - The Complement System during SARS-CoV-2 infection. <i>Life Sciences</i> , 2021, 272, 119245.	2.0	27
20	Impaired dendritic cell differentiation and maturation in the absence of C3. <i>Molecular Immunology</i> , 2008, 45, 1952-1962.	1.0	26
21	On the Three-Finger Protein Domain Fold and CD59-Like Proteins in <i>Schistosoma mansoni</i> . <i>PLoS Neglected Tropical Diseases</i> , 2013, 7, e2482.	1.3	26
22	Screening for C3 Deficiency in Newborns Using Microarrays. <i>PLoS ONE</i> , 2009, 4, e5321.	1.1	22
23	Strategies used by <i>Leptospira</i> spirochetes to evade the host complement system. <i>FEBS Letters</i> , 2020, 594, 2633-2644.	1.3	22
24	C57BL/6 and A/J Mice Have Different Inflammatory Response and Liver Lipid Profile in Experimental Alcoholic Liver Disease. <i>Mediators of Inflammation</i> , 2015, 2015, 1-11.	1.4	20
25	Deficiency of the Human Complement Regulatory Protein Factor H Associated with Low Levels of Component C9. <i>Scandinavian Journal of Immunology</i> , 2008, 68, 445-455.	1.3	19
26	Native conformations of human complement components C3 and C4 show different dependencies on thioester formation. <i>Biochemical Journal</i> , 1998, 329, 705-712.	1.7	17
27	Skipping of exon 30 in C5 gene results in complete human C5 deficiency and demonstrates the importance of C5d and CUB domains for stability. <i>Molecular Immunology</i> , 2009, 46, 2116-2123.	1.0	17
28	<i>Pasteurella pneumotropica</i> Evades the Human Complement System by Acquisition of the Complement Regulators Factor H and C4BP. <i>PLoS ONE</i> , 2014, 9, e111194.	1.1	17
29	Complement C5 controls liver lipid profile, promotes liver homeostasis and inflammation in C57BL/6 genetic background. <i>Immunobiology</i> , 2016, 221, 822-832.	0.8	17
30	<i>Trypanosoma cruzi</i> : Killing and enhanced uptake by resident peritoneal macrophages treated with alpha-2-macroglobulin. <i>Parasitology Research</i> , 1990, 76, 545-552.	0.6	16
31	Homozygous hereditary C3 deficiency due to a premature stop codon. <i>Journal of Clinical Immunology</i> , 2002, 22, 321-330.	2.0	16
32	The complement component C5 promotes liver steatosis and inflammation in murine non-alcoholic liver disease model. <i>Immunology Letters</i> , 2016, 177, 53-61.	1.1	16
33	<i>Leptospira interrogans</i> Secreted Proteases Degrade Extracellular Matrix and Plasma Proteins From the Host. <i>Frontiers in Cellular and Infection Microbiology</i> , 2018, 8, 92.	1.8	16
34	Nonsense-codon-mediated decay in human hereditary complement C3 deficiency. <i>Immunogenetics</i> , 2004, 55, 667-673.	1.2	14
35	Basal physiological parameters of two congenic mice strains: C5 deficient C57BL/6 and C5 sufficient A/J. <i>Immunology Letters</i> , 2014, 159, 47-54.	1.1	14
36	Association of complement factor H Y402H polymorphism and age-related macular degeneration in Brazilian patients. <i>Acta Ophthalmologica</i> , 2010, 88, e165-9.	0.6	13

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37	Complement System in Alcohol-Associated Liver Disease. <i>Immunology Letters</i> , 2021, 236, 37-50.	1.1	13
38	Chemical Chaperones Curcumin and 4-Phenylbutyric Acid Improve Secretion of Mutant Factor H R127H by Fibroblasts from a Factor H-Deficient Patient. <i>Journal of Immunology</i> , 2012, 189, 3242-3248.	0.4	12
39	Dysregulation of adaptive immune responses in complement C3-deficient patients. <i>European Journal of Immunology</i> , 2015, 45, 915-921.	1.6	12
40	Studies of the binding of ficolin-2 and ficolin-3 from the complement lectin pathway to <i>Leptospira biflexa</i> , <i>Pasteurella pneumotropica</i> and Diarrheagenic <i>Escherichia coli</i> . <i>Immunobiology</i> , 2015, 220, 1177-1185.	0.8	12
41	<i>Trypanosoma cruzi</i> : Plasma levels of alpha-2-macroglobulin during experimental murine infections with reticulotropic and myotropic strains. <i>Zeitschrift für Parasitenkunde (Berlin, Germany)</i> , 1990, 76, 726-728.	0.8	11
42	Complement Immune Evasion by Spirochetes. <i>Current Topics in Microbiology and Immunology</i> , 2017, 415, 215-238.	0.7	10
43	Culture-attenuated pathogenic <i>Leptospira</i> lose the ability to survive to complement-mediated-killing due to lower expression of factor H binding proteins. <i>Microbes and Infection</i> , 2019, 21, 377-385.	1.0	10
44	Skipping of exon 27 in C3 gene compromises TED domain and results in complete human C3 deficiency. <i>Immunobiology</i> , 2016, 221, 641-649.	0.8	9
45	<i>Leptospira interrogans</i> thermolysin refolded at high pressure and alkaline pH displays proteolytic activity against complement C3. <i>Biotechnology Reports (Amsterdam, Netherlands)</i> , 2018, 19, e00266.	2.1	7
46	<i>Leptospira</i> and Leptospirosis. , 2015, , 1973-1990.		6
47	Role of Murine Complement Component C5 in Acute in Vivo Infection by Pathogenic <i>Leptospira interrogans</i> . <i>Frontiers in Cellular and Infection Microbiology</i> , 2018, 8, 63.	1.8	6
48	Interaction of human complement factor H variants Tyr402 and His402 with <i>Leptospira</i> spp.. <i>Frontiers in Immunology</i> , 2011, 2, 44.	2.2	5
49	Cytokine Profile in Early Infection by <i>Leptospira interrogans</i> in A/J Mice. <i>Journal of Immunology Research</i> , 2019, 2019, 1-13.	0.9	5
50	A new model of outbred genetically selected mice which present a strong acute inflammatory response in the absence of complement component C5. <i>Inflammation Research</i> , 2009, 58, 204-209.	1.6	4
51	Binding of human complement C1 sesterase inhibitor to <i>Leptospira</i> spp.. <i>Immunobiology</i> , 2018, 223, 183-190.	0.8	4
52	Contribution of Complement System pathways to the killing of <i>Leptospira</i> spp.. <i>Microbes and Infection</i> , 2020, 22, 550-557.	1.0	3
53	Lytic, agglutinating, and opsonizing effect of alpha-2-macroglobulin on sheep red blood cells. <i>Inflammation</i> , 1990, 14, 259-266.	1.7	2
54	Phagocytic activity mediated via Fc gamma R, Fc gamma 4R, and CR3 and H2O2 release during ontogeny of mouse macrophages. <i>Developmental and Comparative Immunology</i> , 1994, 18, 443-454.	1.0	2

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55	The Role of Properdin in Killing of Non-Pathogenic <i>Leptospira biflexa</i> . <i>Frontiers in Immunology</i> , 2020, 11, 572562.	2.2	2
56	Simple Method To Distinguish between Primary and Secondary C3 Deficiencies. <i>Vaccine Journal</i> , 2003, 10, 216-220.	3.2	1
57	Complement Resistance Assays. <i>Methods in Molecular Biology</i> , 2020, 2134, 187-198.	0.4	0