

# Annamaria Fra

## List of Publications by Year in descending order

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37  
papers

2,787  
citations

218381

26  
h-index

344852

36  
g-index

38  
all docs

38  
docs citations

38  
times ranked

3053  
citing authors

#	ARTICLE	IF	CITATIONS
1	De novo formation of caveolae in lymphocytes by expression of VIP21-caveolin.. Proceedings of the National Academy of Sciences of the United States of America, 1995, 92, 8655-8659.	3.3	555
2	Caveolin-1 and -2 in the Exocytic Pathway of MDCK Cells. Journal of Cell Biology, 1998, 140, 795-806.	2.3	283
3	Developmental regulation of IgM secretion: The role of the carboxy-terminal cysteine. Cell, 1990, 60, 781-790.	13.5	248
4	Cutting Edge: Scavenging of Inflammatory CC Chemokines by the Promiscuous Putatively Silent Chemokine Receptor D6. Journal of Immunology, 2003, 170, 2279-2282.	0.4	181
5	A photo-reactive derivative of ganglioside GM1 specifically cross-links VIP21-caveolin on the cell surface. FEBS Letters, 1995, 375, 11-14.	1.3	169
6	Progressively impaired proteasomal capacity during terminal plasma cell differentiation. EMBO Journal, 2006, 25, 1104-1113.	3.5	139
7	Differential Recognition and Scavenging of Native and Truncated Macrophage-Derived Chemokine (Macrophage-Derived Chemokine/CC Chemokine Ligand 22) by the D6 Decoy Receptor. Journal of Immunology, 2004, 172, 4972-4976.	0.4	132
8	Quality control of ER synthesized proteins: an exposed thiol group as a three-way switch mediating assembly, retention and degradation.. EMBO Journal, 1993, 12, 4755-4761.	3.5	124
9	Degradation of unassembled soluble Ig subunits by cytosolic proteasomes: evidence that retrotranslocation and degradation are coupled events. FASEB Journal, 2000, 14, 769-778.	0.2	96
10	Cysteines as Redox Molecular Switches and Targets of Disease. Frontiers in Molecular Neuroscience, 2017, 10, 167.	1.4	95
11	Identification and characterisation of eight novel SERPINA1 Null mutations. Orphanet Journal of Rare Diseases, 2014, 9, 172.	1.2	60
12	Generation and characterization of a mouse lymphatic endothelial cell line. Cell and Tissue Research, 2006, 325, 91-100.	1.5	56
13	CHOP-independent apoptosis and pathway-selective induction of the UPR in developing plasma cells. Molecular Immunology, 2010, 47, 1356-1365.	1.0	56
14	Three New Alpha1-Antitrypsin Deficiency Variants Help to Define a C-Terminal Region Regulating Conformational Change and Polymerization. PLoS ONE, 2012, 7, e38405.	1.1	43
15	The efficiency of cysteine-mediated intracellular retention determines the differential fate of secretory IgA and IgM in B and plasma cells. European Journal of Immunology, 1994, 24, 2477-2482.	1.6	42
16	Polymers of Z Î± <sub>1</sub> -antitrypsin are secreted in cell models of disease. European Respiratory Journal, 2016, 47, 1005-1009.	3.1	41
17	Exposed Thiols Confer Localization in the Endoplasmic Reticulum by Retention Rather than Retrieval. Journal of Biological Chemistry, 1996, 271, 26138-26142.	1.6	40
18	Mechanisms of interleukin-6 protection against ischemiaâ€“reperfusion injury in rat liver. Cytokine, 2006, 34, 131-142.	1.4	37

#	ARTICLE	IF	CITATIONS
19	Real-world clinical applicability of pathogenicity predictors assessed on <i>SERPINA1</i> mutations in alpha-1-antitrypsin deficiency. <i>Human Mutation</i> , 2018, 39, 1203-1213.	1.1	36
20	The Endoplasmic Reticulum as a Site of Protein Degradation. <i>Sub-Cellular Biochemistry</i> , 1993, 21, 143-168.	1.0	36
21	The Decrease of Mineralcorticoid Receptor Drives Angiogenic Pathways in Colorectal Cancer. <i>PLoS ONE</i> , 2013, 8, e59410.	1.1	30
22	Genomic organization and transcriptional analysis of the human genes coding for caveolin-1 and caveolin-2. <i>Gene</i> , 2000, 243, 75-83.	1.0	29
23	Different molecular behavior of CD40 mutants causing hyper-IgM syndrome. <i>Blood</i> , 2010, 116, 5867-5874.	0.6	29
24	Aberrant disulphide bonding contributes to the ER retention of alpha1-antitrypsin deficiency variants. <i>Human Molecular Genetics</i> , 2016, 25, 642-650.	1.4	28
25	Molecular characterization of the new defective P <sub>brescia</sub> alpha1-antitrypsin allele. <i>Human Mutation</i> , 2009, 30, E771-E781.	1.1	27
26	Human Caveolin-1 and Caveolin-2 Are Closely Linked Genes Colocalized with WI-5336 in a Region of 7q31 Frequently Deleted in Tumors. <i>Genomics</i> , 1999, 56, 355-356.	1.3	26
27	Heteropolymerization of $\alpha$ -1-antitrypsin mutants in cell models mimicking heterozygosity. <i>Human Molecular Genetics</i> , 2018, 27, 1785-1793.	1.4	24
28	The pathological Trento variant of alpha $\alpha$ -1-antitrypsin (E75V) shows nonclassical behaviour during polymerization. <i>FEBS Journal</i> , 2017, 284, 2110-2126.	2.2	23
29	Intermittent C1-Inhibitor Deficiency Associated with Recessive Inheritance: Functional and Structural Insight. <i>Scientific Reports</i> , 2018, 8, 977.	1.6	22
30	Neuroserpin: structure, function, physiology and pathology. <i>Cellular and Molecular Life Sciences</i> , 2021, 78, 6409-6430.	2.4	16
31	Intrahepatic heteropolymerization of M and Z alpha-1-antitrypsin. <i>JCI Insight</i> , 2020, 5, .	2.3	16
32	Phenotypic behavior of C2C12 myoblasts upon expression of the dystrophy-related caveolin $\beta$ P104L and TFT mutants. <i>FEBS Letters</i> , 2007, 581, 5099-5104.	1.3	13
33	Characterisation of a type II functionally-deficient variant of alpha-1-antitrypsin discovered in the general population. <i>PLoS ONE</i> , 2019, 14, e0206955.	1.1	13
34	Cellular Models for the Serpinopathies. <i>Methods in Molecular Biology</i> , 2018, 1826, 109-121.	0.4	9
35	The molecular species responsible for $\alpha$ -1-antitrypsin deficiency are suppressed by a small molecule chaperone. <i>FEBS Journal</i> , 2021, 288, 2222-2237.	2.2	8
36	The Importance of N186 in the Alpha-1-Antitrypsin Shutter Region Is Revealed by the Novel Bologna Deficiency Variant. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5668.	1.8	5

#	ARTICLE	IF	CITATIONS
37	ALPHA1-ANTITRYPSIN DEFICIENCY: A 25-YEAR EXPERIENCE. Chest, 2020, 157, A177.	0.4	0