Francesca Lugani

List of Publications by Year
in descending order

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44 papers	2,188 citations	430874 18 h-index	254184 43 g-index
46 all docs	46 docs citations	46 times ranked	3968 citing authors

#	Article	IF	Citations
1	Refractory Minimal Change Disease and Focal Segmental Glomerular Sclerosis Treated With Anakinra. Kidney International Reports, 2022, 7, 121-124.	0.8	6
2	Parathyroid hormone and phosphate homeostasis in patients with Bartter and Gitelman syndrome: an international cross-sectional study. Nephrology Dialysis Transplantation, 2022, 37, 2474-2486.	0.7	5
3	An update on COVID-19 in paediatric and young adults with nephrotic syndrome,Âreceiving chronic immunosuppression during the Omicron pandemic. Journal of Nephrology, 2022, 35, 1775-1776.	2.0	4
4	Case Report: Atypical Manifestations Associated With FOXP3 Mutations. The â€œFil Rougeâ€•of Treg Between IPEX Features and Other Clinical Entities?. Frontiers in Immunology, 2022, 13, 854749.	4.8	6
5	Proteomics and Extracellular Vesicles as Novel Biomarker Sources in Peritoneal Dialysis in Children. International Journal of Molecular Sciences, 2022, 23, 5655.	4.1	4
6	Renal involvement and StrÃ¸mme syndrome. CKJ: Clinical Kidney Journal, 2021, 14, 439-441.	2.9	3
7	A novel nonsense variation in the albumin gene (c.1309 A>T) causing analbuminaemia. British Journal of Biomedical Science, 2021, 78, 154-157.	1.3	2
8	Rituximab vs Low-Dose Mycophenolate Mofetil in Recurrence of Steroid-Dependent Nephrotic Syndrome in Children and Young Adults. JAMA Pediatrics, 2021, 175, 631.	6.2	21
9	Vaccines and Disease Relapses in Children with Nephrotic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2021, 16, 937-938.	4.5	12
10	Improving data quality in observational research studies: Report of the Cure Glomerulonephropathy (CureGN) network. Contemporary Clinical Trials Communications, 2021, 22, 100749.	1.1	7
11	Human or Chimeric Monoclonal Anti-CD20 Antibodies for Children with Nephrotic Syndrome: A Superiority Randomized Trial. Journal of the American Society of Nephrology: JASN, 2021, 32, 2652-2663.	6.1	30
12	Neutrophil Extracellular Traps-DNase Balance and Autoimmunity. Cells, 2021, 10, 2667.	4.1	23
13	Randomised controlled trial comparing rituximab to mycophenolate mofetil in children and young adults with steroid-dependent idiopathic nephrotic syndrome: study protocol. BMJ Open, 2021, 11, e052450.	1.9	5
14	Recurrent Hypoglycemia in a Case of Congenital Analbuminemia. Case Reports in Endocrinology, 2020, 2020, 1-6.	0.4	1
15	Serum indoxyl sulfate concentrations associate with progression of chronic kidney disease in children. PLoS ONE, 2020, 15, e0240446.	2.5	19
16	A novel insertion (c.1098dupT) in the albumin gene causes analbuminemia in a consanguineous family. European Journal of Medical Genetics, 2019, 62, 144-148.	1.3	4
17	Impaired Systolic and Diastolic Left Ventricular Function in Children with Chronic Kidney Disease - Results from the 4C Study. Scientific Reports, 2019, 9, 11462.	3.3	20
18	Diagnosis, Phenotype, and Molecular Genetics of Congenital Analbuminemia. Frontiers in Genetics, 2019, 10, 336.	2.3	22

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19	Low levels of urinary epidermal growth factorÂpredict chronic kidney disease progressionÂin children. Kidney International, 2019, 96, 214-221.	5.2	43
20	Isolated nocturnal and isolated daytime hypertension associate with altered cardiovascular morphology and function in children with chronic kidney disease. Journal of Hypertension, 2019, 37, 2247-2255.	0.5	45
21	Congenital analbuminemia in a patient affected by hypercholesterolemia: A case report. World Journal of Clinical Cases, 2019, 7, 466-472.	0.8	7
22	Clinical Characteristics and Treatment Patterns of Children and Adults With IgA Nephropathy or IgA Vasculitis: Findings From the CureGN Study. Kidney International Reports, 2018, 3, 1373-1384.	0.8	39
23	A novel splicing mutation in the ALB gene causing analbuminaemia in a Portuguese woman. Pathology, 2018, 50, 679-682.	0.6	4
24	Whole-Exome Sequencing in Adults With Chronic Kidney Disease. Annals of Internal Medicine, 2018, 168, 100.	3.9	154
25	Metabolic acidosis is common and associates with disease progression in children with chronic kidney disease. Kidney International, 2017, 92, 1507-1514.	5.2	66
26	Glucokinase mutations in pediatric patients with impaired fasting glucose. Acta Diabetologica, 2017, 54, 913-923.	2.5	11
27	Urine Proteome Biomarkers in Kidney Diseases. I. Limits, Perspectives, and First Focus on Normal Urine. Biomarker Insights, 2016, 11, BMI.S26229.	2.5	22
28	A novel splicing mutation in the albumin gene (c.270+1G>T) causes analbuminaemia in a German infant. Annals of Clinical Biochemistry, 2016, 53, 615-619.	1.6	6
29	A nucleotide deletion and frame-shift cause analbuminemia in a Turkish family. Biochemia Medica, 2016, 26, 264-271.	2.7	5
30	Novel INF2 mutations in an Italian cohort of patients with focal segmental glomerulosclerosis, renal failure and Charcot-Marie-Tooth neuropathy. Nephrology Dialysis Transplantation, 2014, 29, iv80-iv86.	0.7	28
31	Discovery of new risk loci for IgA nephropathy implicates genes involved in immunity against intestinal pathogens. Nature Genetics, 2014, 46, 1187-1196.	21.4	505
32	Osteocalcin regulates murine and human fertility through a pancreas-bone-testis axis. Journal of Clinical Investigation, 2014, 124, 5522-5522.	8.2	0
33	Hyperglycaemia and Î²-cell antibodies: Is it always pre-type 1 diabetes?. Diabetes Research and Clinical Practice, 2013, 100, e20-e22.	2.8	4
34	A novel mutation in the albumin gene (c.1A>C) resulting in analbuminemia. European Journal of Clinical Investigation, 2013, 43, 72-78.	3.4	18
35	Mutations in DSTYK and Dominant Urinary Tract Malformations. New England Journal of Medicine, 2013, 369, 621-629.	27.0	119
36	A Retrotransposon Insertion in the 5â€² Regulatory Domain of Ptf1a Results in Ectopic Gene Expression and Multiple Congenital Defects in Danforth's Short Tail Mouse. PLoS Genetics, 2013, 9, e1003206.	3.5	20

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37	Osteocalcin regulates murine and human fertility through a pancreas-bone-testis axis. Journal of Clinical Investigation, 2013, 123, 2421-2433.	8.2	233
38	Comment on: Clinical application of best practice guidelines for genetic diagnosis of MODY2. Diabetes Research and Clinical Practice, 2012, 95, e29-e30.	2.8	2
39	Wolfram Syndrome: New Mutations, Different Phenotype. PLoS ONE, 2012, 7, e29150.	2.5	55
40	Long-term home parenteral nutrition in children with chronic intestinal failure: A 15-year experience at a single Italian centre. Digestive and Liver Disease, 2011, 43, 28-33.	0.9	57
41	Genome-wide association study identifies susceptibility loci for IgA nephropathy. Nature Genetics, 2011, 43, 321-327.	21.4	528
42	Neonatal Diabetes Caused by Pancreatic Agenesia. Diabetes Care, 2010, 33, e112-e112.	8.6	0
43	Strong Association Between Time Watching Television and Blood Glucose Control in Children and Adolescents With Type 1 Diabetes. Diabetes Care, 2007, 30, e137-e137.	8.6	2
44	To the Editor: Alcaligenes as a pathogen in airways chronic infection in cystic fibrosis. Pediatric Pulmonology, 2003, 35, 412-413.	2.0	8