

Marijan Saraga

List of Publications by Year in descending order

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papers

994
citations

623734

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501196

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all docs

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docs citations

29
times ranked

1652
citing authors

#	ARTICLE	IF	CITATIONS
1	Copy-Number Disorders Are a Common Cause of Congenital Kidney Malformations. <i>American Journal of Human Genetics</i> , 2012, 91, 987-997.	6.2	201
2	The copy number variation landscape of congenital anomalies of the kidney and urinary tract. <i>Nature Genetics</i> , 2019, 51, 117-127.	21.4	144
3	Genetic Drivers of Kidney Defects in the DiGeorge Syndrome. <i>New England Journal of Medicine</i> , 2017, 376, 742-754.	27.0	120
4	Mutations in <i>DSTYK</i> and Dominant Urinary Tract Malformations. <i>New England Journal of Medicine</i> , 2013, 369, 621-629.	27.0	119
5	Exome-wide Association Study Identifies <i>GREB1L</i> Mutations in Congenital Kidney Malformations. <i>American Journal of Human Genetics</i> , 2017, 101, 789-802.	6.2	63
6	Alcohol Intoxication in Pediatric Age: Ten-year Retrospective Study. <i>Croatian Medical Journal</i> , 2009, 50, 151-156.	0.7	35
7	Role of mitotic, pro-apoptotic and anti-apoptotic factors in human kidney development. <i>Pediatric Nephrology</i> , 2006, 21, 627-636.	1.7	34
8	Ciliogenesis in normal human kidney development and post-natal life. <i>Pediatric Nephrology</i> , 2012, 27, 55-63.	1.7	33
9	The Role of Direct Radionuclide Cystography in Evaluation of Vesicoureteral Reflux. <i>Scandinavian Journal of Urology and Nephrology</i> , 1996, 30, 367-371.	1.4	31
10	Expression of intermediate filaments, EGF and TGF- β in early human kidney development. <i>Journal of Molecular Histology</i> , 2008, 39, 227-235.	2.2	24
11	Involvement of FGF and BMP family proteins and VEGF in early human kidney development. <i>Histology and Histopathology</i> , 2008, 23, 853-62.	0.7	21
12	Copy Number Variant Analysis and Genome-wide Association Study Identify Loci with Large Effect for Vesicoureteral Reflux. <i>Journal of the American Society of Nephrology: JASN</i> , 2021, 32, 805-820.	6.1	17
13	Expression and localization of DAB1 and Reelin during normal human kidney development. <i>Croatian Medical Journal</i> , 2019, 60, 521-531.	0.7	16
14	Immunohistochemical and electronmicroscopic features of mesenchymal-to-epithelial transition in human developing, postnatal and nephrotic podocytes. <i>Histochemistry and Cell Biology</i> , 2017, 147, 481-495.	1.7	15
15	Immunohistochemical expression pattern of RIP5, FGFR1, FGFR2 and HIP2 in the normal human kidney development. <i>Acta Histochemica</i> , 2019, 121, 531-538.	1.8	14
16	FHR-5 Serum Levels and CFHR5 Genetic Variations in Patients With Immune Complex-Mediated Membranoproliferative Glomerulonephritis and C3-Glomerulopathy. <i>Frontiers in Immunology</i> , 2021, 12, 720183.	4.8	12
17	Expression of intermediate filaments and desmosomal proteins during differentiation of the human spinal cord. <i>Acta Histochemica</i> , 2002, 104, 157-166.	1.8	10
18	Glomeruli from patients with nephrin mutations show increased number of ciliated and poorly differentiated podocytes. <i>Acta Histochemica</i> , 2018, 120, 748-756.	1.8	10

#	ARTICLE	IF	CITATIONS
19	Connexin Signaling in the Juxtaglomerular Apparatus (JGA) of Developing, Postnatal Healthy and Nephrotic Human Kidneys. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8349.	4.1	10
20	Epidemiology of 10-year paediatric renal biopsies in the region of southern Croatia. <i>BMC Nephrology</i> , 2020, 21, 65.	1.8	10
21	Validation of distinct pathogenic patterns in a cohort of membranoproliferative glomerulonephritis patients by cluster analysis. <i>CKJ: Clinical Kidney Journal</i> , 2020, 13, 225-234.	2.9	9
22	Epidemiology of renal disease in children in the region of southern Croatia: a 10-year review of regional renal biopsy databases. <i>Medical Science Monitor</i> , 2007, 13, CR172-6.	1.1	9
23	Mechanism of cystogenesis in nephrotic kidneys: a histopathological study. <i>BMC Nephrology</i> , 2014, 15, 3.	1.8	7
24	Changing Pattern of Acute Alcohol Intoxications in Children. <i>Medical Science Monitor</i> , 2018, 24, 5123-5131.	1.1	7
25	Spatio-temporal patterning of different connexins in developing and postnatal human kidneys and in nephrotic syndrome of the Finnish type (CNF). <i>Scientific Reports</i> , 2020, 10, 8756.	3.3	7
26	Intrarenal Reflux in the Light of Contrast-Enhanced Voiding Urosonography. <i>Frontiers in Pediatrics</i> , 2021, 9, 642077.	1.9	7
27	Types of Parenchymal Changes Diagnosed on DMSA Scans of Kidneys Affected by Different Grades of Vesicoureteral Reflux. <i>Medical Science Monitor</i> , 2021, 27, e929617.	1.1	4
28	Differences in Immunohistochemical and Ultrastructural Features between Podocytes and Parietal Epithelial Cells (PECs) Are Observed in Developing, Healthy Postnatal, and Pathologically Changed Human Kidneys. <i>International Journal of Molecular Sciences</i> , 2022, 23, 7501.	4.1	4
29	The Spectrum of Parenchymal Changes in Kidneys Affected by Intrarenal Reflux, Diagnosed by Contrast-Enhanced Voiding Urosonography and DMSA Scan. <i>Frontiers in Pediatrics</i> , 0, 10, .	1.9	1