

Rozemarijn Snoek

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/905572/publications.pdf>

Version: 2024-02-01

10
papers

235
citations

1307594

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h-index

1474206

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g-index

10
all docs

10
docs citations

10
times ranked

447
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetics-first approach improves diagnostics of ESKD patients <50 years old. Nephrology Dialysis Transplantation, 2022, 37, 349-357.	0.7	27
2	FC 012PRIMARY KIDNEY DISEASE IMPACTS OUTCOME IN CKD PREGNANCIES: COMPLICATIONS IN COL4A3-5 RELATED DISEASE (ALPORT SYNDROME) VS OTHER CKD PREGNANCIES. Nephrology Dialysis Transplantation, 2021, 36, .	0.7	0
3	Preimplantation Genetic Testing for Monogenic Kidney Disease. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 1279-1286.	4.5	27
4	Pregnancy in Advanced Kidney Disease: Clinical Practice Considerations on a Challenging Combination. Nephron, 2020, 144, 185-189.	1.8	10
5	Defects in t6A tRNA modification due to GON7 and YRDC mutations lead to Galloway-Mowat syndrome. Nature Communications, 2019, 10, 3967.	12.8	66
6	Importance of Genetic Diagnostics in Adult-Onset Focal Segmental Glomerulosclerosis. Nephron, 2019, 142, 351-358.	1.8	10
7	NPHP1 (Nephrocystin-1) Gene Deletions Cause Adult-Onset ESRD. Journal of the American Society of Nephrology: JASN, 2018, 29, 1772-1779.	6.1	74
8	Accuracy of diagnosis and counseling of fetal brain anomalies prior to 24 weeks of gestational age. Journal of Maternal-Fetal and Neonatal Medicine, 2018, 31, 2188-2194.	1.5	10
9	Assessing Nephron Hyperplasia in Fetal Congenital Solitary Functioning Kidneys by Measuring Renal Papilla Number. American Journal of Kidney Diseases, 2018, 72, 465-467.	1.9	5
10	Importance of reliable variant calling and clear phenotyping when reporting on gene panel testing in renal disease. Kidney International, 2017, 92, 1325-1327.	5.2	6