

Talia Gutman

List of Publications by Year in descending order

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Version: 2024-02-01

37
papers

1,068
citations

471509

17
h-index

434195

31
g-index

39
all docs

39
docs citations

39
times ranked

1244
citing authors

#	ARTICLE	IF	CITATIONS
1	Reporting guideline for priority setting of health research (REPRISE). BMC Medical Research Methodology, 2019, 19, 243.	3.1	112
2	Establishing a Core Outcome Set for Peritoneal Dialysis: Report of the SONG-PD (Standardized) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 70 Diseases, 2020, 75, 404-412.	1.9	92
3	Identifying Outcomes Important to Patients with Glomerular Disease and Their Caregivers. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 673-684.	4.5	66
4	Establishing core outcome domains in pediatric kidney disease: report of the Standardized Outcomes in Nephrologyâ€™Children and Adolescents (SONG-KIDS) consensus workshops. Kidney International, 2020, 98, 553-565.	5.2	58
5	Patient and Caregiver Priorities for Outcomes in CKD: A Multinational Nominal Group Technique Study. American Journal of Kidney Diseases, 2020, 76, 679-689.	1.9	56
6	Communication during childhood cancer: Systematic review of patient perspectives. Cancer, 2020, 126, 701-716.	4.1	50
7	Patient and Caregiver Perspectives on Terms Used to Describe Kidney Health. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 937-948.	4.5	47
8	Child and Parental Perspectives on Communication and Decision Making in Pediatric CKD: A Focus Group Study. American Journal of Kidney Diseases, 2018, 72, 547-559.	1.9	46
9	Identifying Important Outcomes for Young People With CKD and Their Caregivers: A Nominal Group Technique Study. American Journal of Kidney Diseases, 2019, 74, 82-94.	1.9	42
10	Experiences and Perspectives of Dietary Management Among Patients on Hemodialysis: An Interview Study. , 2018, 28, 411-421.		36
11	Patientsâ€™ and caregiversâ€™ perspectives on access to kidney replacement therapy in rural communities: systematic review of qualitative studies. BMJ Open, 2020, 10, e037529.	1.9	24
12	Core Outcome Domains for Trials in Autosomal Dominant Polycystic Kidney Disease: An International Delphi Survey. American Journal of Kidney Diseases, 2020, 76, 361-373.	1.9	23
13	KHAâ€™CARI Guideline recommendations for renal biopsy. Nephrology, 2019, 24, 1205-1213.	1.6	22
14	Establishing a Core Outcome Set for Autosomal Dominant Polycystic Kidney Disease: Report of the Standardized Outcomes in Nephrologyâ€™Polycystic Kidney Disease (SONG-PKD) Consensus Workshop. American Journal of Kidney Diseases, 2021, 77, 255-263.	1.9	21
15	Standardised Outcomes in Nephrologyâ€™Polycystic Kidney Disease (SONG-PKD): study protocol for establishing a core outcome set in polycystic kidney disease. Trials, 2017, 18, 560.	1.6	20
16	Identifying patientâ€™important outcomes in polycystic kidney disease: An international nominal group technique study. Nephrology, 2019, 24, 1214-1224.	1.6	20
17	Standardized Outcomes in Nephrologyâ€™Glomerular Disease (SONG-GD): establishing a core outcome set for trials in patients with glomerular disease. Kidney International, 2019, 95, 1280-1283.	5.2	20
18	Recruitment and retention in clinical trials in chronic kidney disease: report from national workshops with patients, caregivers and health professionals. Nephrology Dialysis Transplantation, 2020, 35, 755-764.	0.7	19

#	ARTICLE	IF	CITATIONS
19	Developing Consensus-Based Outcome Domains for Trials in Children and Adolescents With CKD: An International Delphi Survey. <i>American Journal of Kidney Diseases</i> , 2020, 76, 533-545.	1.9	19
20	Range and Variability of Outcomes Reported in Randomized Trials Conducted in Patients With Polycystic Kidney Disease: A Systematic Review. <i>American Journal of Kidney Diseases</i> , 2020, 76, 213-223.	1.9	16
21	Principles and strategies for involving patients in research in chronic kidney disease: report from national workshops. <i>Nephrology Dialysis Transplantation</i> , 2020, 35, 1585-1594.	0.7	14
22	Patient needs and priorities for patient navigator programmes in chronic kidney disease: a workshop report. <i>BMJ Open</i> , 2020, 10, e040617.	1.9	14
23	Living kidney donor and recipient perspectives on their relationship: longitudinal semi-structured interviews. <i>BMJ Open</i> , 2019, 9, e026629.	1.9	13
24	Patient-led identification and prioritization of exercise interventions for fatigue on dialysis: a workshop report. <i>CKJ: Clinical Kidney Journal</i> , 2021, 14, 831-839.	2.9	11
25	Patients' Perspectives on Access to Dialysis and Kidney Transplantation in Rural Communities in Australia. <i>Kidney International Reports</i> , 2022, 7, 591-600.	0.8	11
26	Patient and Caregiver Experiences and Attitudes about Their Involvement in Research in Chronic Kidney Disease. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2022, 17, 215-227.	4.5	10
27	Clinicians' perspectives on equity of access to dialysis and kidney transplantation for rural people in Australia: a semistructured interview study. <i>BMJ Open</i> , 2022, 12, e052315.	1.9	10
28	Clinicians' and researchers' perspectives on establishing and implementing core outcomes in haemodialysis: semistructured interview study. <i>BMJ Open</i> , 2018, 8, e021198.	1.9	9
29	Identifying and integrating patient and caregiver perspectives in clinical practice guidelines for percutaneous renal biopsy. <i>Nephrology</i> , 2019, 24, 395-404.	1.6	9
30	"A sword of Damocles": patient and caregiver beliefs, attitudes and perspectives on presymptomatic testing for autosomal dominant polycystic kidney disease: a focus group study. <i>BMJ Open</i> , 2020, 10, e038005.	1.9	5
31	Controversy and Debate Series on Core Outcome Sets. Paper 5: Large-scale, mixed-methods, knowledge exchange to establish core outcomes – the SONG approach. <i>Journal of Clinical Epidemiology</i> , 2020, 125, 225-228.	5.0	5
32	"Can I go to Glasgow?" Learnings from patient involvement at the 17th Congress of the International Society for Peritoneal Dialysis (ISPD). <i>Peritoneal Dialysis International</i> , 2020, 40, 12-25.	2.3	5
33	A Core Outcome Set for Trials in Glomerular Disease. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2022, 17, 53-64.	4.5	4
34	Perspectives of Clinicians on Shared Decision Making in Pediatric CKD: A Qualitative Study. <i>American Journal of Kidney Diseases</i> , 2022, 80, 241-250.	1.9	3
35	Patient and caregiver perspectives on blood pressure in children with chronic kidney disease. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 1330-1339.	0.7	2
36	A focus group study of self-management in patients with glomerular disease.. <i>Kidney International Reports</i> , 2021, 7, 56-67.	0.8	2

#	ARTICLE	IF	CITATIONS
37	Child and caregiver perspectives on access to psychosocial and educational support in pediatric chronic kidney disease: a focus group study. <i>Pediatric Nephrology</i> , 2023, 38, 249-260.	1.7	2