Beth Anne Tarini

List of Publications by Year
in descending order

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80 papers	2,650 citations	249298 26 h-index	214428 50 g-index
80 all docs	80 docs citations	80 times ranked	3772 citing authors

#	Article	IF	Citations
1	Healthcare team communication training in the United States: A scoping review. Health Communication, 2023, 38, 1821-1846.	1.8	3
2	Pediatrician Adherence to Guidelines for Diagnosis and Management of High Blood Pressure. Journal of Pediatrics, 2022, 242, 12-17.e1.	0.9	18
3	Genomics and Newborn Screening: Perspectives of Public Health Programs. International Journal of Neonatal Screening, 2022, 8, 11.	1.2	5
4	493: FOOD INSECURITY IN THE PEDIATRIC INTENSIVE CARE UNIT. Critical Care Medicine, 2022, 50, 238-238.	0.4	0
5	Parentsâ€™ Experiences and Needs Regarding Infant Sickle Cell Trait Results. Pediatrics, 2022, 149, .	1.0	2
6	The Value of Cognitive Pretesting: Improving Validity and Revealing Blind Spots through the Development of a Newborn Screening Parent Experiences Survey. International Journal of Neonatal Screening, 2021, 7, 41.	1.2	4
7	The Effect of BabySeq on Pediatric and Genomic Researchâ€”More Than Baby Steps. JAMA Pediatrics, 2021, 175, 1107.	3.3	3
8	Clusters of adverse childhood experiences and unmet need for care coordination. Child Abuse and Neglect, 2021, 122, 105334.	1.3	6
9	Associations between adverse childhood experiences and need and unmet need for care coordination. International Journal of Care Coordination, 2021, 24, 125-132.	0.3	3
10	Systems Integration: The Next Frontier in Newborn-Screening Timeliness. Journal of Public Health Management and Practice, 2020, 26, E8-E15.	0.7	1
11	Vulnerable Child Syndrome and Newborn Screening Carrier ResultsÂforÂCystic Fibrosis or Sickle Cell. Journal of Pediatrics, 2020, 224, 44-50.e1.	0.9	13
12	Comparison of Video, App, and Standard Consent Processes on Decision-Making for Biospecimen Research: A Randomized Controlled Trial. Journal of Empirical Research on Human Research Ethics, 2020, 15, 252-260.	0.6	11
13	How a baby with classic galactosemia was nearly missed: When the test succeeds but system fails. American Journal of Medical Genetics, Part A, 2020, 182, 1750-1753.	0.7	1
14	Primary Care Providersâ€™ Preferences and Concerns Regarding Specific Visual Displays for Returning Hemoglobin A1c Test Results to Patients. Medical Decision Making, 2019, 39, 796-804.	1.2	3
15	Misclassification of VLCAD carriers due to variable confirmatory testing after a positive NBS result. Journal of Community Genetics, 2019, 10, 447-451.	0.5	3
16	Newborn Screening Collection and Delivery Processes in Michigan Birthing Hospitals: Strategies to Improve Timeliness. Maternal and Child Health Journal, 2018, 22, 1436-1443.	0.7	3
17	Family History Collection Practices: National Survey of Pediatric Primary Care Providers. Clinical Pediatrics, 2018, 57, 537-546.	0.4	3
18	An Assessment of Public Preferences for Newborn Screening Using Bestâ€“Worst Scaling. Journal of Pediatrics, 2018, 201, 62-68.e1.	0.9	11

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19	Survey of family history taking and genetic testing in pediatric practice. Journal of Community Genetics, 2017, 8, 109-115.	0.5	19
20	An Assessment of a Shortened Consent Form for the Storage and Research Use of Residual Newborn Screening Blood Spots. Journal of Empirical Research on Human Research Ethics, 2017, 12, 335-342.	0.6	6
21	Adolescent bariatric surgery: a systematic review of recommendation documents. Surgery for Obesity and Related Diseases, 2017, 13, 1768-1779.	1.0	20
22	Graphics help patients distinguish between urgent and non-urgent deviations in laboratory test results. Journal of the American Medical Informatics Association: JAMIA, 2017, 24, 520-528.	2.2	72
23	Biobank participant support of newborn screening for disorders with variable treatment and intervention options. Journal of Community Genetics, 2016, 7, 291-302.	0.5	2
24	Effect of â€œPink Eyeâ€•Label on Parentsâ€™ Intent to Use Antibiotics and Perceived Contagiousness. Clinical Pediatrics, 2016, 55, 543-548.	0.4	13
25	A Quality Improvement Collaborative to Improve Pediatric Primary Care Genetic Services. Pediatrics, 2016, 137, e20143874.	1.0	9
26	Primary Care Providersâ€™ Experiences Notifying Parents of Cystic Fibrosis Newborn Screening Results. Clinical Pediatrics, 2015, 54, 67-75.	0.4	16
27	Parent and Public Interest in Whole-Genome Sequencing. Public Health Genomics, 2015, 18, 151-159.	0.6	27
28	Primary Care Providers' Initial Evaluation of Children with Global Developmental Delay: A Clinical Vignette Study. Journal of Pediatrics, 2015, 167, 1404-1408.e1.	0.9	13
29	Framing optional genetic testing in the context of mandatory newborn screening tests. BMC Medical Informatics and Decision Making, 2015, 15, 50.	1.5	7
30	Understanding Outcomes in Adolescent Bariatric Surgery. Pediatrics, 2015, 136, e312-e314.	1.0	14
31	Primary-care providersâ€™ perceived barriers to integration of genetics services: a systematic review of the literature. Genetics in Medicine, 2015, 17, 169-176.	1.1	201
32	Comprehensive Evaluation of the Child With Intellectual Disability or Global Developmental Delays. Pediatrics, 2014, 134, e903-e918.	1.0	412
33	Genetic services and attitudes in primary care pediatrics. American Journal of Medical Genetics, Part A, 2014, 164, 449-455.	0.7	32
34	The Value of Time in Assessing the Effectiveness of Newborn Screening for Congenital Adrenal Hyperplasia. JAMA Pediatrics, 2014, 168, 515.	3.3	3
35	Effects of Undergoing Multiplex Genetic Susceptibility Testing on Parent Attitudes towards Testing Their Children. Annals of Behavioral Medicine, 2014, 47, 388-394.	1.7	5
36	Emerging Issues in Public Health Genomics. Annual Review of Genomics and Human Genetics, 2014, 15, 461-480.	2.5	39

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37	Parental Permission for Pilot Newborn Screening Research: Guidelines From the NBSTRN. Pediatrics, 2014, 133, e410-e417.	1.0	20
38	Screening Practices for Identifying Type 2 Diabetes in Adolescents. Journal of Adolescent Health, 2014, 54, 139-143.	1.2	28
39	Parentsâ€™ interest in whole-genome sequencing of newborns. Genetics in Medicine, 2014, 16, 78-84.	1.1	73
40	Anticipating the arrival of low-penetrance genetic testing to primary care medicine. Journal of Community Genetics, 2013, 4, 285-288.	0.5	2
41	Growing Up in the Genomic Era: Implications of Whole-Genome Sequencing for Children, Families, and Pediatric Practice. Annual Review of Genomics and Human Genetics, 2013, 14, 535-555.	2.5	54
42	The Perils of SNP Microarray Testing: Uncovering Unexpected Consanguinity. Pediatric Neurology, 2013, 49, 50-53.	1.0	7
43	Lessons that newborn screening in the USA can teach us about biobanking and large-scale genetic studies. Personalized Medicine, 2013, 10, 81-87.	0.8	6
44	Health Supervision for Children With Marfan Syndrome. Pediatrics, 2013, 132, e1059-e1072.	1.0	49
45	Ethical and Policy Issues in Genetic Testing and Screening of Children. Pediatrics, 2013, 131, 620-622.	1.0	326
46	Effects of Hypothetical Type 2 Diabetes Genetic Testing on Parentsâ€™ Efforts to Prevent Diabetes in Children. Clinical Pediatrics, 2013, 52, 821-828.	0.4	1
47	Influence of â€œGERDâ€•Label on Parentsâ€™ Decision to Medicate Infants. Pediatrics, 2013, 131, 839-845.	1.0	72
48	Blindness in Walnut Grove: How Did Mary Ingalls Lose Her Sight?. Pediatrics, 2013, 131, 404-406.	1.0	1
49	Personalized medicine in primary care: the need for relevance. Personalized Medicine, 2013, 10, 515-517.	0.8	0
50	Family History in Primary Care Pediatrics. Pediatrics, 2013, 132, S203-S210.	1.0	19
51	Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 477-88.	0.4	10
52	Decision analysis, economic evaluation, and newborn screening: challenges and opportunities. Genetics in Medicine, 2012, 14, 703-712.	1.1	34
53	Communicating With Parents About Newborn Screening. JAMA Pediatrics, 2012, 166, 95.	3.6	4
54	Decision Analysis, Economic Evaluation, and Newborn Screening. Obstetrical and Gynecological Survey, 2012, 67, 758-760.	0.2	1

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55	Ethical Issues in Neonatal and Pediatric Clinical Trials. Pediatric Clinics of North America, 2012, 59, 1205-1220.	0.9	56
56	Ethical Issues with Newborn Screening in the Genomics Era. Annual Review of Genomics and Human Genetics, 2012, 13, 381-393.	2.5	87
57	A Policy Impact Analysis of the Mandatory <scp>NCAA</scp> Sickle Cell Trait Screening Program. Health Services Research, 2012, 47, 446-461.	1.0	28
58	Storage and use of residual newborn screening blood spots: A public policy emergency. Genetics in Medicine, 2011, 13, 619-620.	1.1	24
59	Diagnosis of Diabetes using Hemoglobin A1c: Should Recommendations in Adults Be Extrapolated to Adolescents?. Journal of Pediatrics, 2011, 158, 947-952.e3.	0.9	82
60	Genetic Testing and Youth Sports. JAMA - Journal of the American Medical Association, 2011, 305, 1033.	3.8	12
61	Commentary: Children and Predictive Genomic Testing: Disease Prevention, Research Protection, and Our Future. Journal of Pediatric Psychology, 2011, 36, 1113-1121.	1.1	19
62	False-Positive Newborn Screening Result and Future Health Care Use in a State Medicaid Cohort. Pediatrics, 2011, 128, 715-722.	1.0	31
63	Response to the Commentary: Children and Predictive Genomic Testing. Journal of Pediatric Psychology, 2011, 36, 1128-1129.	1.1	0
64	Genetic Screening. Epidemiologic Reviews, 2011, 33, 148-164.	1.3	63
65	Parents' Interest in Predictive Genetic Testing for Their Children When a Disease Has No Treatment. Pediatrics, 2009, 124, e432-e438.	1.0	45
66	Neonatal Sepsis: Looking Beyond the Blood Culture. JAMA Pediatrics, 2009, 163, 12.	3.6	6
67	Afraid in the hospital: Parental concern for errors during a child's hospitalization. Journal of Hospital Medicine, 2009, 4, 521-527.	0.7	11
68	Waiving informed consent in newborn screening research: Balancing social value and respect. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2008, 148C, 23-30.	0.7	28
69	Delayed introduction of solids does not decrease the incidence of asthma or allergic rhinitis. Journal of Pediatrics, 2008, 153, 440.	0.9	1
70	A False-positive Newborn Screening Result: Goat's Milk Acidopathy: In Reply. Pediatrics, 2008, 122, 211-211.	1.0	2
71	Parents' Concern About Their Own and Their Children's Genetic Disease Risk. JAMA Pediatrics, 2008, 162, 1079.	3.6	31
72	The Current Revolution in Newborn Screening. JAMA Pediatrics, 2007, 161, 767.	3.6	50

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73	False-positive Results in Expanded Newborn Screening: In Reply. Pediatrics, 2007, 119, 415-415.	1.0	2
74	Institutional variation in ordering complete blood counts for children hospitalized with bronchiolitis. Journal of Hospital Medicine, 2007, 2, 69-73.	0.7	9
75	Toward Family-Centered Inpatient Medical Care: The Role of Parents as Participants in Medical Decisions. Journal of Pediatrics, 2007, 151, 690-695.e1.	0.9	31
76	State Newborn Screening in the Tandem Mass Spectrometry Era: More Tests, More False-Positive Results. Pediatrics, 2006, 118, 448-456.	1.0	122
77	Systematic Review of the Relationship Between Early Introduction of Solid Foods to Infants and the Development of Allergic Disease. JAMA Pediatrics, 2006, 160, 502.	3.6	116
78	Use of Serum Electrolyte Panels in Gastroenteritis. Pediatrics, 2005, 115, 1109-1109.	1.0	2
79	Does Presentation Format at the Pediatric Academic Societies' Annual Meeting Predict Subsequent Publication?. Pediatrics, 2003, 112, 1238-1241.	1.0	66
80	Is Duct Tape Occlusion Therapy as Effective as Cryotherapy for the Treatment of the Common Wart?. JAMA Pediatrics, 2002, 156, 975.	3.6	16