Sarah M Greene

List of Publications by Year
in descending order

52 papers	3,188 citations	172457 29 h-index	206112 48 g-index
52 all docs	52 docs citations	52 times ranked	4335 citing authors

#	Article	IF	Citations
1	Recommendations for strengthening the role of embedded researchers to accelerate implementation in health systems: Findings from a state-of-the-art (SOTA) conference workgroup. Healthcare, 2021, 8, 100455.	1.3	16
2	Rethinking ethical oversight in the era of the learning health system. Healthcare, 2020, 8, 100462.	1.3	7
3	"You Really Do Have to Know the Local Context" IRB Administrators and Researchers on the Implications of the NIH Single IRB Mandate for Multisite Genomic Studies. Journal of Empirical Research on Human Research Ethics, 2019, 14, 286-295.	1.3	6
4	We need to talk: Provider conversations with peers and patients about a medical error. Journal of Patient Safety and Risk Management, 2019, 24, 140-146.	0.6	0
5	Collaborating on Data, Science, and Infrastructure: The 20-Year Journey of the Cancer Research Network. EGEMS (Washington, DC), 2019, 7, 7.	2.0	6
6	Celebrating a Quarter-Century of Public Domain Research: 25th Annual Conference of the Health Care Systems Research Network. Journal of Patient-centered Research and Reviews, 2019, 6, 218-223.	0.9	3
7	We've Only Just Begun — Insights from a 25-Year Journey to Accelerate Health Care Transformation Through Delivery System Research. EGEMS (Washington, DC), 2019, 7, 19.	2.0	4
8	From Principles to Practice: Real-World Patient and Stakeholder Engagement in Breast Cancer Research. , 2018, 22, 17-232.		7
9	Primary care physicians' willingness to disclose oncology errors involving multiple providers to patients. BMJ Quality and Safety, 2016, 25, 787-795.	3.7	18
10	Oversight on the borderline: Quality improvement and pragmatic research. Clinical Trials, 2015, 12, 457-466.	1.6	57
11	Providers' Perceptions of Communication Breakdowns in Cancer Care. Journal of General Internal Medicine, 2014, 29, 1122-1130.	2.6	58
12	Cancer research network: using integrated healthcare delivery systems as platforms for cancer survivorship research. Journal of Cancer Survivorship, 2013, 7, 55-62.	2.9	11
13	Patients' and family members' views on patient-centered communication during cancer care. Psycho-Oncology, 2013, 22, 2487-2495.	2.3	106
14	Potential Role of Community-Based Healthcare System Data in Research on Survivors of Adolescent and Young Adult Cancer. Journal of Adolescent and Young Adult Oncology, 2013, 2, 53-58.	1.3	2
15	More than words: Patients' views on apology and disclosure when things go wrong in cancer care. Patient Education and Counseling, 2013, 90, 341-346.	2.2	75
16	Toward Patient-Centered Cancer Care: Patient Perceptions of Problematic Events, Impact, and Response. Journal of Clinical Oncology, 2012, 30, 1784-1790.	1.6	128
17	Implementing the Learning Health System: From Concept to Action. Annals of Internal Medicine, 2012, 157, 207.	3.9	276
18	The Cancer Message Literacy Tests: Psychometric analyses and validity studies. Patient Education and Counseling, 2012, 89, 69-75.	2.2	20

#	ARTICLE	IF	CITATIONS
19	Can Integrating Health Literacy into the Patient-centered Medical Home Help us Weather the Perfect Storm?. Journal of General Internal Medicine, 2012, 27, 588-594.	2.6	12
20	Health literacy and cancer prevention: Two new instruments to assess comprehension. Patient Education and Counseling, 2012, 88, 54-60.	2.2	37
21	Improving Modern Cancer Care Through Information Technology. American Journal of Preventive Medicine, 2011, 40, S198-S207.	3.0	117
22	If we only knew what we know: principles for knowledge sharing across people, practices, and platforms. Translational Behavioral Medicine, 2011, 1, 15-25.	2.4	15
23	The quality of cancer patient experience: perspectives of patients, family members, providers and experts. BMJ Quality and Safety, 2010, 19, 484-489.	3.7	64
24	Media Messages About Cancer: What Do People Understand?. Journal of Health Communication, 2010, 15, 126-145.	2.4	26
25	Bioinformatics. American Journal of Preventive Medicine, 2010, 38, 646-651.	3.0	16
26	The process is the product: a new model for multisite IRB review of data-only studies. IRB: Ethics & Human Research, 2010, 32, 1-6.	0.8	8
27	Looking at Research Consent Forms through a Participant-Centered Lens: The PRISM Readability Toolkit. American Journal of Health Promotion, 2009, 23, 371-375.	1.7	15
28	Diffusion of aromatase inhibitors for breast cancer therapy between 1996 and 2003 in the Cancer Research Network. Breast Cancer Research and Treatment, 2008, 107, 397-403.	2.5	35
29	Understanding highâ€quality cancer care. Cancer, 2008, 112, 934-942.	4.1	109
30	Survivorship research based in integrated healthcare delivery systems. Cancer, 2008, 112, 2617-2626.	4.1	17
31	Positive, Negative, and Disparate-Womenâ€™s Differing Long-Term Psychosocial Experiences of Bilateral or Contralateral Prophylactic Mastectomy. Breast Journal, 2008, 14, 25-32.	1.0	87
32	Web-Based Smoking-Cessation Programs. American Journal of Preventive Medicine, 2008, 34, 373-381.	3.0	257
33	What Women Wish They Knew Before Prophylactic Mastectomy. Cancer Nursing, 2007, 30, 285-291.	1.5	57
34	Quality of Life After Bilateral Prophylactic Mastectomy. Annals of Surgical Oncology, 2007, 14, 686-694.	1.5	62
35	Impact of IRB Requirements on a Multicenter Survey of Prophylactic Mastectomy Outcomes. Annals of Epidemiology, 2006, 16, 275-278.	1.9	47
36	A review finds that multicenter studies face substantial challenges but strategies exist to achieve Institutional Review Board approval. Journal of Clinical Epidemiology, 2006, 59, 784-790.	5.0	94

#	Article	IF	Citations
37	How health care systems can begin to address the challenge of limited literacy. Journal of General Internal Medicine, 2006, 21, 884-887.	2.6	189
38	Contentment With Quality of Life Among Breast Cancer Survivors With and Without Contralateral Prophylactic Mastectomy. Journal of Clinical Oncology, 2006, 24, 1350-1356.	1.6	92
39	Interest in an Online Smoking Cessation Program and Effective Recruitment Strategies: Results From Project Quit. Journal of Medical Internet Research, 2006, 8, e14.	4.3	47
40	Building a Research Consortium of Large Health Systems: The Cancer Research Network. Journal of the National Cancer Institute Monographs, 2005, 2005, 3-11.	2.1	172
41	Race and Ethnicity: Comparing Medical Records to Self-Reports. Journal of the National Cancer Institute Monographs, 2005, 2005, 72-74.	2.1	48
42	RESPONSE: Re: Efficacy of Breast Cancer Screening in the Community According to Risk Level. Journal of the National Cancer Institute, 2005, 97, 1704-1705.	6.3	1
43	Efficacy of Breast Cancer Screening in the Community According to Risk Level. Journal of the National Cancer Institute, 2005, 97, 1035-1043.	6.3	60
44	Finding Common Ground: Patient-Centeredness and Evidence-Based Chronic Illness Care. Journal of Alternative and Complementary Medicine, 2005, 11, s-7-s-15.	2.1	326
45	Women's Decision-Making Roles Regarding Contralateral Prophylactic Mastectomy. Journal of the National Cancer Institute Monographs, 2005, 2005, 55-60.	2.1	56
46	Building a Virtual Cancer Research Organization. Journal of the National Cancer Institute Monographs, 2005, 2005, 12-25.	2.1	143
47	Measuring and Improving Performance in Multicenter Research Consortia. Journal of the National Cancer Institute Monographs, 2005, 2005, 26-32.	2.1	30
48	The Coordinated Clinical Studies Network: A Multidisciplinary Alliance to Facilitate Research and Improve Care.. , 2005, 9, 33-5.		2
49	Organizational barriers to physician participation in cancer clinical trials. American Journal of Managed Care, 2005, 11, 413-21.	1.1	66
50	The Role of Research in Integrated Health Care Systems: The HMO Research Network. , 2004, 8, 10-7.		4
51	A computerized system to facilitate medical record abstraction in cancer research (United States). Cancer Causes and Control, 2003, 14, 469-476.	1.8	8
52	Training, Quality Assurance, and Assessment of Medical Record Abstraction in a Multisite Study. American Journal of Epidemiology, 2003, 157, 546-551.	3.4	69