## Hilda Bastian

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1698404/publications.pdf

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35 papers

2,368 citations

623734 14 h-index 28 g-index

42 all docs 42 docs citations

times ranked

42

3885 citing authors

#	Article	IF	CITATIONS
1	What the systematic review of HPV vaccine clinical study reports does, and does not, reveal: commentary on JÃ,rgensen et al Systematic Reviews, 2020, 9, 41.	5.3	O
2	Improving consultations with children and adolescents. Drug and Therapeutics Bulletin, 2020, 58, 66-66.	0.3	0
3	Biomedical Journals and Preprint Services: Friends or Foes?. Clinical Chemistry, 2017, 63, 453-458.	3.2	15
4	Concern noted: a descriptive study of editorial expressions of concern in PubMed and PubMed Central. Research Integrity and Peer Review, 2017, 2, .	5.2	18
5	Nondisclosure of Financial Interest in Clinical Practice Guideline Development: An Intractable Problem?. PLoS Medicine, 2016, 13, e1002030.	8.4	9
6	A Stronger Post-Publication Culture Is Needed for Better Science. PLoS Medicine, 2014, 11, e1001772.	8.4	29
7	Presenting Unwelcome Research Findings. The Virtual Mentor: VM, 2014, 16, 98-102.	0.3	O
8	Evidence-based health information from the users' perspective – a qualitative analysis. BMC Health Services Research, 2013, 13, 405.	2.2	23
9	PRISMA for Abstracts: Reporting Systematic Reviews in Journal and Conference Abstracts. PLoS Medicine, 2013, 10, e1001419.	8.4	495
10	Core Competencies for Shared Decision Making Training Programs: Insights From an International, Interdisciplinary Working Group. Journal of Continuing Education in the Health Professions, 2013, 33, 267-273.	1.3	89
11	Integrating Values and Consumer Involvement in Guidelines with the Patient at the Center. Proceedings of the American Thoracic Society, 2012, 9, 262-268.	3.5	33
12	Choosing health technology assessment and systematic review topics: The development of priority-setting criteria for patients' and consumers' interests. International Journal of Technology Assessment in Health Care, 2011, 27, 348-356.	0.5	12
13	Patienteninformationen – vom Paternalismus zum Empowerment. Public Health Forum, 2011, 19, 7-8.	0.2	3
14	Comment on: Olsson et al. High Levels of Education Are Associated With an Increased Risk of Latent Autoimmune Diabetes in Adults: Results From the Nord-TrÃ,ndelag Health Study. Diabetes Care 2011;34:102–107. Diabetes Care, 2011, 34, e113-e113.	8.6	1
15	Integrating patients' views into health technology assessment: Analytic hierarchy process (AHP) as a method to elicit patient preferences. International Journal of Technology Assessment in Health Care, 2011, 27, 369-375.	0.5	142
16	Effects of treatment in women with gestational diabetes mellitus: systematic review and meta-analysis. BMJ: British Medical Journal, 2010, 340, c1395-c1395.	2.3	260
17	Seventy-Five Trials and Eleven Systematic Reviews a Day: How Will We Ever Keep Up?. PLoS Medicine, 2010, 7, e1000326.	8.4	812
18	Gesundheitsinformation.de und Informationsbed $\tilde{A}^{1}\!\!/\!$	0.2	0

#	Article	IF	Citations
19	Taking healthcare interventions from trial to practice. BMJ: British Medical Journal, 2010, 341, c3852-c3852.	2.3	168
20	FÃ⅓r BÃ⅓rger und Patienten – Die evidenzbasierten Gesundheitsinformationen des IQWiG. , 2009, , 183-197.		1
21	Health literacy and patient information: Developing the methodology for a national evidence-based health website. Patient Education and Counseling, 2008, 73, 551-556.	2.2	29
22	'They would say that, wouldn't they?' A reader's guide toauthor and sponsor biases in clinical research. Journal of the Royal Society of Medicine, 2006, 99, 611-614.	2.0	10
23	Consumer and researcher collaboration in trials: filling the gaps. Clinical Trials, 2005, 2, 3-4.	1.6	11
24	Variation in perceptions of risk between doctors and patients: risks look different when they are close to home. Australian Prescriber, 2003, 26, 20-21.	1.0	0
25	Allies or enemies? Evidenceâ€based medicine and consumer choice. Medical Journal of Australia, 2000, 172, 5-6.	1.7	9
26	A consumer trip into the world of the DALY calculations: An Alice-in-Wonderland experience. Reproductive Health Matters, 2000, 8, 113-116.	1.2	18
27	Perinatal Death Associated With Planned Home Birth in Australia. Obstetrical and Gynecological Survey, 1999, 54, 168-169.	0.4	1
28	Speaking Up for Ourselves: <i>The Evolution of Consumer Advocacy in Health Care</i> International Journal of Technology Assessment in Health Care, 1998, 14, 3-23.	0.5	71
29	Competing or Complementary?: Ethical Considerations and the Quality of Randomized Trials. International Journal of Technology Assessment in Health Care, 1996, 12, 247-263.	0.5	14
30	Raising the Standard: Practice Guidelines and Consumer Participation. International Journal for Quality in Health Care, 1996, 8, 485-490.	1.8	49
31	Personal Beliefs and Alternative Childbirth Choices: A Survey of 552 Women Who Planned to Give Birth at Home. Birth, 1993, 20, 186-192.	2.2	12
32	Confined, Managed, and Delivered. Obstetrical and Gynecological Survey, 1992, 47, 626-628.	0.4	0
33	Maternity alliance position on relationships between consumers and professional organisations. Women and Birth, 1992, 5, 6.	0.1	O
34	Confined, managed and delivered: the language of obstetrics. BJOG: an International Journal of Obstetrics and Gynaecology, 1992, 99, 92-93.	2.3	17
35	Obstetrics and Litigation. Obstetrical and Gynecological Survey, 1991, 46, 545-547.	0.4	0