

Richard M Nixon

List of Publications by Year in descending order

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34
papers

1,848
citations

279798

23
h-index

377865

34
g-index

34
all docs

34
docs citations

34
times ranked

2994
citing authors

#	ARTICLE	IF	CITATIONS
1	Relative benefit-risk comparing diclofenac to other traditional non-steroidal anti-inflammatory drugs and cyclooxygenase-2 inhibitors in patients with osteoarthritis or rheumatoid arthritis: a network meta-analysis. <i>Arthritis Research and Therapy</i> , 2015, 17, 66.	3.5	175
2	Non-parametric methods for cost-effectiveness analysis: the central limit theorem and the bootstrap compared. <i>Health Economics (United Kingdom)</i> , 2010, 19, 316-333.	1.7	148
3	The Effect of Disease, Functional Status, and Relapses on the Utility of People with Multiple Sclerosis in the UK. <i>Value in Health</i> , 2007, 10, 54-60.	0.3	138
4	Using mixed treatment comparisons and meta-regression to perform indirect comparisons to estimate the efficacy of biologic treatments in rheumatoid arthritis. <i>Statistics in Medicine</i> , 2007, 26, 1237-1254.	1.6	125
5	Developing Appropriate Methods for Cost-Effectiveness Analysis of Cluster Randomized Trials. <i>Medical Decision Making</i> , 2012, 32, 350-361.	2.4	119
6	Methods for incorporating covariate adjustment, subgroup analysis and between-centre differences into cost-effectiveness evaluations. <i>Health Economics (United Kingdom)</i> , 2005, 14, 1217-1229.	1.7	109
7	Parametric modelling of cost data: some simulation evidence. <i>Health Economics (United Kingdom)</i> , 2005, 14, 421-428.	1.7	84
8	Parametric modelling of cost data in medical studies. <i>Statistics in Medicine</i> , 2004, 23, 1311-1331.	1.6	83
9	No Evidence of Disease Activity: Indirect Comparisons of Oral Therapies for the Treatment of Relapsing-Remitting Multiple Sclerosis. <i>Advances in Therapy</i> , 2014, 31, 1134-1154.	2.9	83
10	Biologic drugs for rheumatoid arthritis in the medicare program: A cost-effectiveness analysis. <i>Arthritis and Rheumatism</i> , 2008, 58, 939-946.	6.7	82
11	Using multilevel models for assessing the variability of multinational resource use and cost data. <i>Health Economics (United Kingdom)</i> , 2005, 14, 185-196.	1.7	69
12	How Sensitive Are Cost-Effectiveness Analyses to Choice of Parametric Distributions?. <i>Medical Decision Making</i> , 2005, 25, 416-423.	2.4	65
13	Valsartan vs. other angiotensin II receptor blockers in the treatment of hypertension: a meta-analytical approach. <i>International Journal of Clinical Practice</i> , 2009, 63, 766-775.	1.7	61
14	Randomised controlled trial of educational package on management of menorrhagia in primary care: the Anglia menorrhagia education study. <i>BMJ: British Medical Journal</i> , 1999, 318, 1246-1250.	2.3	47
15	Cost-Effectiveness of Disease-Modifying Therapies in the Management of Multiple Sclerosis for the Medicare Population. <i>Value in Health</i> , 2009, 12, 657-665.	0.3	47
16	Statistical Methods for Cost-Effectiveness Analyses That Use Data from Cluster Randomized Trials. <i>Medical Decision Making</i> , 2012, 32, 209-220.	2.4	46
17	METHODS FOR COVARIATE ADJUSTMENT IN COST-EFFECTIVENESS ANALYSIS THAT USE CLUSTER RANDOMISED TRIALS. <i>Health Economics (United Kingdom)</i> , 2012, 21, 1101-1118.	1.7	44
18	Bayesian Hierarchical Models for Cost-Effectiveness Analyses that Use Data from Cluster Randomized Trials. <i>Medical Decision Making</i> , 2010, 30, 163-175.	2.4	42

#	ARTICLE	IF	CITATIONS
19	Addressing the issues that arise in analysing multicentre cost data, with application to a multinational study. <i>Journal of Health Economics</i> , 2006, 25, 1015-1028.	2.7	41
20	A case study using the ProACT ^{URL} and BRAT frameworks for structured benefit risk assessment. <i>Biometrical Journal</i> , 2016, 58, 8-27.	1.0	34
21	Multilevel models for estimating incremental net benefits in multinational studies. <i>Health Economics (United Kingdom)</i> , 2007, 16, 815-826.	1.7	29
22	The Distribution of the Cost of Multiple Sclerosis in the UK: How Do Costs Vary by Illness Severity?. <i>Value in Health</i> , 2007, 10, 386-389.	0.3	29
23	Network meta-analysis combining individual patient and aggregate data from a mixture of study designs with an application to pulmonary arterial hypertension. <i>BMC Medical Research Methodology</i> , 2015, 15, 34.	3.1	29
24	Multilevel models for cost-effectiveness analyses that use cluster randomised trial data: An approach to model choice. <i>Statistical Methods in Medical Research</i> , 2016, 25, 2036-2052.	1.5	24
25	A comparison of clinical assessment with ultrasound in the management of secondary postpartum haemorrhage. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2002, 104, 113-115.	1.1	17
26	The Rheumatoid Arthritis Drug Development Model: a case study in Bayesian clinical trial simulation. <i>Pharmaceutical Statistics</i> , 2009, 8, 371-389.	1.3	16
27	High correlation of VAS pain scores after 2 and 6 weeks of treatment with VAS pain scores at 12 weeks in randomised controlled trials in rheumatoid arthritis and osteoarthritis: meta-analysis and implications. <i>Arthritis Research and Therapy</i> , 2016, 18, 73.	3.5	11
28	Evidence That Longer Androgen Receptor Polyglutamine Repeats Are a Causal Factor for Genital Abnormalities. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 3207-3210.	3.6	11
29	Efficacy and safety of diclofenac in osteoarthritis: Results of a network meta-analysis of unpublished legacy studies. <i>Scandinavian Journal of Pain</i> , 2017, 16, 74-88.	1.3	10
30	Management of menorrhagia: an audit of practices in the Anglia menorrhagia education study. <i>BMJ: British Medical Journal</i> , 2001, 322, 523-524.	2.3	8
31	Bayesian evaluation of breast cancer screening using data from two studies. <i>Statistics in Medicine</i> , 2003, 22, 1661-1674.	1.6	8
32	Using short-term evidence to predict six-month outcomes in clinical trials of signs and symptoms in rheumatoid arthritis. <i>Pharmaceutical Statistics</i> , 2009, 8, 150-162.	1.3	8
33	Imputation of a true endpoint from a surrogate: application to a cluster randomized controlled trial with partial information on the true endpoint. <i>BMC Medical Research Methodology</i> , 2003, 3, 17.	3.1	3
34	Review and comparison of methodologies for indirect comparison of clinical trial results: an illustration with ranibizumab and aflibercept. <i>Expert Review of Pharmacoeconomics and Outcomes Research</i> , 2016, 16, 793-801.	1.4	3