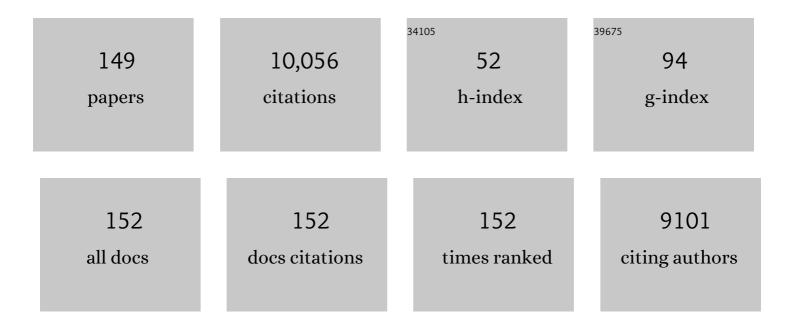
Diana S Rose

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Global pattern of experienced and anticipated discrimination against people with schizophrenia: a cross-sectional survey. Lancet, The, 2009, 373, 408-415.	13.7	888
2	Evidence for effective interventions to reduce mental-health-related stigma and discrimination. Lancet, The, 2016, 387, 1123-1132.	13.7	776
3	Stigma: ignorance, prejudice or discrimination?. British Journal of Psychiatry, 2007, 190, 192-193.	2.8	545
4	Patients' perspectives on electroconvulsive therapy: systematic review. BMJ: British Medical Journal, 2003, 326, 1363-0.	2.3	378
5	The WPA- Lancet Psychiatry Commission on the Future of Psychiatry. Lancet Psychiatry,the, 2017, 4, 775-818.	7.4	305
6	Discrimination in health care against people with mental illness. International Review of Psychiatry, 2007, 19, 113-122.	2.8	302
7	The importance of relationships in mental health care: A qualitative study of service users' experiences of psychiatric hospital admission in the UK. BMC Health Services Research, 2008, 8, 92.	2.2	272
8	Development and Psychometric Properties of the Mental Health Knowledge Schedule. Canadian Journal of Psychiatry, 2010, 55, 440-448.	1.9	242
9	Patients' views and readmissions 1 year after involuntary hospitalisation. British Journal of Psychiatry, 2009, 194, 49-54.	2.8	186
10	Barriers and facilitators of disclosures of domestic violence by mental health service users: qualitative study. British Journal of Psychiatry, 2011, 198, 189-194.	2.8	173
11	Clinical outcomes of Joint Crisis Plans to reduce compulsory treatment for people with psychosis: a randomised controlled trial. Lancet, The, 2013, 381, 1634-1641.	13.7	148
12	Development and psychometric evaluation of the Discrimination and Stigma Scale (DISC). Psychiatry Research, 2013, 208, 33-40.	3.3	145
13	Coercion and Treatment Satisfaction Among Involuntary Patients. Psychiatric Services, 2010, 61, 286-292.	2.0	144
14	Lost in the shadows: reflections on the dark side of co-production. Health Research Policy and Systems, 2020, 18, 43.	2.8	144
15	Cancer diagnosis in people with severe mental illness: practical and ethical issues. Lancet Oncology, The, 2010, 11, 797-804.	10.7	139
16	Psychiatric patients' views on why their involuntary hospitalisation was right or wrong: a qualitative study. Social Psychiatry and Psychiatric Epidemiology, 2012, 47, 1169-1179.	3.1	137
17	Selecting outcome measures in mental health: the views of service users. Journal of Mental Health, 2011, 20, 336-346.	1.9	136
18	The importance of content and face validity in instrument development: lessons learnt from service users when developing the Recovering Quality of Life measure (ReQoL). Quality of Life Research, 2018, 27, 1893-1902.	3.1	128

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19	Experiences of discrimination among people Using mental health services in England 2008-2011. British Journal of Psychiatry, 2013, 202, s58-s63.	2.8	127
20	250 labels used to stigmatise people with mental illness. BMC Health Services Research, 2007, 7, 97.	2.2	124
21	Hearing the voices of service user researchers in collaborative qualitative data analysis: the case for multiple coding. Health Expectations, 2013, 16, e89-99.	2.6	121
22	Influence of Time to Change's social marketing interventions on stigma in England 2009-2011. British Journal of Psychiatry, 2013, 202, s77-s88.	2.8	119
23	Service user/survivor-led research in mental health: epistemological possibilities. Disability and Society, 2017, 32, 773-789.	2.2	119
24	Developing the evidence base of patient and public involvement in health and social care research: the case for measuring impact. International Journal of Consumer Studies, 2011, 35, 628-632.	11.6	118
25	Can't surf, won't surf: The digital divide in mental health. Journal of Mental Health, 2012, 21, 395-403.	1.9	114
26	VOICE: Developing a new measure of service users' perceptions of inpatient care, using a participatory methodology. Journal of Mental Health, 2012, 21, 57-71.	1.9	112
27	Power, Privilege and Knowledge: the Untenable Promise of Co-production in Mental "Healthâ€ Frontiers in Sociology, 2019, 4, 57.	2.0	108
28	DEVELOPMENT AND VALIDATION OF â€ [~] SURE': A PATIENT REPORTED OUTCOME MEASURE (PROM) FOR RECOVERY FROM DRUG AND ALCOHOL DEPENDENCE. Drug and Alcohol Dependence, 2016, 165, 159-167.	3.2	94
29	Close to the bench as well as at the bedside: involving service users in all phases of translational research. Health Expectations, 2012, 15, 389-400.	2.6	90
30	The mainstreaming of recovery. Journal of Mental Health, 2014, 23, 217-218.	1.9	89
31	"You're all going to hate the word â€~recovery' by the end of this― Service users' views of measu addiction recovery. Drugs: Education, Prevention and Policy, 2015, 22, 26-34.	iring 1.3	88
32	Discrimination against people with mental illness: what can psychiatrists do?. Advances in Psychiatric Treatment, 2010, 16, 53-59.	0.5	87
33	A model for developing outcome measures from the perspectives of mental health service users. International Review of Psychiatry, 2011, 23, 41-46.	2.8	87
34	England's Time to Change Antistigma Campaign: One-Year Outcomes of Service User-Rated Experiences of Discrimination. Psychiatric Services, 2012, 63, 451-457.	2.0	84
35	Information, consent and perceived coercion: patients' perspectives on electroconvulsive therapy. British Journal of Psychiatry, 2005, 186, 54-59.	2.8	83
36	What Do Clients Think of Cognitive Remediation Therapy?: A Consumer-Led Investigation of Satisfaction and Side Effects. American Journal of Psychiatric Rehabilitation, 2008, 11, 181-204.	0.7	83

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37	Service user involvement: impact and participation: a survey of service user and staff perspectives. BMC Health Services Research, 2014, 14, 491.	2.2	83
38	Exploring stigmatisation among people diagnosed with either bipolar disorder or borderline personality disorder: A critical realist analysis. Social Science and Medicine, 2014, 123, 7-17.	3.8	78
39	Patient and public involvement in health research: Ethical imperative and/or radical challenge?. Journal of Health Psychology, 2014, 19, 149-158.	2.3	77
40	Participatory research: real or imagined. Social Psychiatry and Psychiatric Epidemiology, 2018, 53, 765-771.	3.1	74
41	Achieving Continuity of Care: Facilitators and Barriers in Community Mental Health Teams. Implementation Science, 2011, 6, 23.	6.9	73
42	Illness careers and continuity of care in mental health services: A qualitative study of service users and carers. Social Science and Medicine, 2009, 69, 632-639.	3.8	72
43	Newspaper coverage of mental illness in England 2008-2011. British Journal of Psychiatry, 2013, 202, s64-s69.	2.8	71
44	Barriers to shared decision making in mental health care: qualitative study of the Joint Crisis Plan for psychosis. Health Expectations, 2016, 19, 448-458.	2.6	71
45	What are the research priorities of mental health service users?. Journal of Mental Health, 2002, 11, 1-3.	1.9	69
46	Newspaper coverage of mental illness in the UK, 1992-2008. BMC Public Health, 2011, 11, 796.	2.9	69
47	Collaborative research between users and professionals: peaks and pitfalls. Psychiatric Bulletin, 2003, 27, 404-406.	0.3	68
48	The psychiatric ward as a therapeutic space: systematic review. British Journal of Psychiatry, 2014, 205, 171-176.	2.8	68
49	The Feasibility and Acceptability to Service Users of CIRCuiTS, a Computerized Cognitive Remediation Therapy Programme for Schizophrenia. Behavioural and Cognitive Psychotherapy, 2016, 44, 288-305.	1.2	68
50	Family Caregivers' Experiences of Involuntary Psychiatric Hospital Admissions of Their Relatives – a Qualitative Study. PLoS ONE, 2011, 6, e25425.	2.5	64
51	How should we measure addiction recovery? Analysis of service provider perspectives using online Delphi groups. Drugs: Education, Prevention and Policy, 2014, 21, 310-323.	1.3	63
52	Joint crisis plans for people with borderline personality disorder: feasibility and outcomes in a randomised controlled trial. British Journal of Psychiatry, 2013, 202, 357-364.	2.8	55
53	â€~Our community is the worst': The influence of cultural beliefs on stigma, relationships with family and help-seeking in three ethnic communities in London. International Journal of Social Psychiatry, 2013, 59, 535-544.	3.1	54
54	Service user perspectives on the impact of a mental illness diagnosis. Epidemiologia E Psichiatria Sociale, 2010, 19, 140-147.	0.9	51

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55	What service users with psychotic disorders want in a mental health crisis or relapse: thematic analysis of joint crisis plans. Social Psychiatry and Psychiatric Epidemiology, 2014, 49, 1609-1617.	3.1	49
56	What are mental health service users' priorities for research in the UK?. Journal of Mental Health, 2008, 17, 520-530.	1.9	48
57	Service user perspectives on coercion and restraint in mental health. BJPsych International, 2017, 14, 59-61.	1.4	48
58	Service users' experiences of residential alternatives to standard acute wards: qualitative study of similarities and differences. British Journal of Psychiatry, 2010, 197, s26-s31.	2.8	45
59	Partnership, co-ordination of care and the place of user involvement. Journal of Mental Health, 2003, 12, 59-70.	1.9	44
60	The Response of Mental Health Services to Domestic Violence. Journal of the American Psychiatric Nurses Association, 2012, 18, 326-336.	1.0	43
61	Collaborative development of an electronic Personal Health Record for people with severe and enduring mental health problems. BMC Psychiatry, 2014, 14, 305.	2.6	43
62	Improving Therapeutic Relationships. Qualitative Health Research, 2015, 25, 1637-1647.	2.1	40
63	Understanding service user-defined continuity of care and its relationship to health and social measures: a cross-sectional study. BMC Health Services Research, 2012, 12, 145.	2.2	39
64	Patients' perceptions of depression and coronary heart disease: a qualitative UPBEAT-UK study. BMC Family Practice, 2013, 14, 38.	2.9	39
65	â€~Widening cross-disciplinary research for mental health': what is missing from the Research Councils UK mental health agenda?. Disability and Society, 2018, 33, 476-481.	2.2	37
66	Discrimination against people with a mental health diagnosis: qualitative analysis of reported experiences. Journal of Mental Health, 2014, 23, 88-93.	1.9	36
67	Consumers' views of electroconvulsive therapy: A qualitative analysis. Journal of Mental Health, 2004, 13, 285-293.	1.9	34
68	The mental health strategy for Europe: Why service user leadership in research is indispensable. Journal of Mental Health, 2012, 21, 219-226.	1.9	33
69	Continuity of care for people with psychotic illness: Its relationship to clinical and social functioning. International Journal of Social Psychiatry, 2013, 59, 5-17.	3.1	31
70	Randomised Controlled Trial of Joint Crisis Plans to Reduce Compulsory Treatment for People with Psychosis: Economic Outcomes. PLoS ONE, 2013, 8, e74210.	2.5	31
71	Can the therapeutic relationship predict 18 month outcomes for individuals with psychosis?. Psychiatry Research, 2014, 220, 585-591.	3.3	31
72	Service user and carer priorities in a Biomedical Research Centre for mental health. Journal of Mental Health, 2016, 25, 185-188.	1.9	31

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73	Eliciting users' views of ECT in two mental health trusts with a user-designed questionnaire. Journal of Mental Health, 2004, 13, 403-413.	1.9	30
74	Rapid progress or lengthy process? electronic personal health records in mental health. BMC Psychiatry, 2011, 11, 117.	2.6	30
75	Electronic personal health records for people with severe mental illness; a feasibility study. BMC Psychiatry, 2015, 15, 192.	2.6	29
76	'Having a diagnosis is a qualification for the job'. BMJ: British Medical Journal, 2003, 326, 1331-1331.	2.3	28
77	Madness strikes back. Journal of Community and Applied Social Psychology, 2008, 18, 638-644.	2.4	28
78	The great ambivalence: factors likely to affect service user and public acceptability of the pharmacogenomics of antidepressant medication. Sociology of Health and Illness, 2008, 30, 944-958.	2.1	28
79	The role of fear in mental health service users' experiences: a qualitative exploration. Social Psychiatry and Psychiatric Epidemiology, 2015, 50, 1079-1087.	3.1	28
80	Defining continuity of care from the perspectives of mental health service users and professionals: an exploratory, comparative study. Health Expectations, 2016, 19, 973-987.	2.6	28
81	CRIMSON [CRisis plan IMpact: Subjective and Objective coercion and eNgagement] Protocol: A randomised controlled trial of joint crisis plans to reduce compulsory treatment of people with psychosis. Trials, 2010, 11, 102.	1.6	27
82	Inpatient care 50Âyears after the process of deinstitutionalisation. Social Psychiatry and Psychiatric Epidemiology, 2014, 49, 665-671.	3.1	26
83	The contemporary state of service-user-led research. Lancet Psychiatry,the, 2015, 2, 959-960.	7.4	25
84	Views of the Therapeutic Environment (VOTE): Stakeholder involvement in measuring staff perceptions of acute in-patient care. International Journal of Nursing Studies, 2012, 49, 1403-1410.	5.6	24
85	User-generated quality standards for youth mental health in primary care: a participatory research design using mixed methods. BMJ Quality and Safety, 2014, 23, 857-866.	3.7	24
86	A randomised controlled trial of positive memory training for the treatment of depression within schizophrenia. BMC Psychiatry, 2015, 15, 85.	2.6	24
87	Up-Beat UK: A programme of research into the relationship between coronary heart disease and depression in primary care patients. BMC Family Practice, 2011, 12, 38.	2.9	23
88	The effect of disclosure of mental illness by interviewers on reports of discrimination experienced by service users: A randomized study. International Review of Psychiatry, 2011, 23, 47-54.	2.8	23
89	Advance Statements for Borderline Personality Disorder: A Qualitative Study of Future Crisis Treatment Preferences. Psychiatric Services, 2014, 65, 802-807.	2.0	23
90	Emerging consensus on measuring addiction recovery: Findings from a multi-stakeholder consultation exercise. Drugs: Education, Prevention and Policy, 2016, 23, 31-40.	1.3	23

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91	Service user led organisations in mental health today. Journal of Mental Health, 2016, 25, 254-259.	1.9	23
92	Exploring the potential use of patient and public involvement to strengthen Indonesian mental health care for people with psychosis: A qualitative exploration of the views of service users and carers. Health Expectations, 2020, 23, 377-387.	2.6	23
93	Public involvement in health outcomes research: lessons learnt from the development of the recovering quality of life (ReQoL) measures. Health and Quality of Life Outcomes, 2019, 17, 60.	2.4	22
94	Perceptions of User Involvement: a User-Led Study. International Journal of Social Psychiatry, 2010, 56, 389-401.	3.1	20
95	Psychiatrists' Perspectives on Antipsychotic Dose and the Role of Plasma Concentration Therapeutic Drug Monitoring. Therapeutic Drug Monitoring, 2014, 36, 486-493.	2.0	19
96	Costs of the police service and mental healthcare pathways experienced by individuals with enduring mental health needs. British Journal of Psychiatry, 2017, 210, 157-164.	2.8	19
97	Mental health in Europe. BMJ: British Medical Journal, 2005, 330, 613-614.	2.3	18
98	Service user produced knowledge. Journal of Mental Health, 2008, 17, 447-451.	1.9	18
99	Design in mind: eliciting service user and frontline staff perspectives on psychiatric ward design through participatory methods. Journal of Mental Health, 2016, 25, 114-121.	1.9	18
100	The effectiveness of joint crisis plans for people with borderline personality disorder: protocol for an exploratory randomised controlled trial. Trials, 2010, 11, 18.	1.6	16
101	Seeking help for obsessive compulsive disorder (<scp>OCD</scp>): a qualitative study of the enablers and barriers conducted by a researcher with personal experience of <scp>OCD</scp> . Psychology and Psychotherapy: Theory, Research and Practice, 2017, 90, 193-211.	2.5	16
102	The CIRCuiTS study (Implementation of cognitive remediation in early intervention services): protocol for a randomised controlled trial. Trials, 2018, 19, 183.	1.6	16
103	An Evaluation of New Services for Personality-Disordered Offenders: Staff and Service User Perspectives. International Journal of Social Psychiatry, 2010, 56, 186-195.	3.1	15
104	Holding blame at bay? â€~Gene talk' in family members' accounts of schizophrenia aetiology. BioSocieties 2012, 7, 273-293.	' 1.3	15
105	Exploring the potential of civic engagement to strengthen mental health systems in Indonesia (IGNITE): a study protocol. International Journal of Mental Health Systems, 2018, 12, 49.	2.7	15
106	A Comparison of Participant Information Elicited by Service User and Non-Service User Researchers. Psychiatric Services, 2011, 62, 210-213.	2.0	14
107	A pilot randomised controlled trial of personalised care for depressed patients with symptomatic coronary heart disease in South London general practices: the UPBEAT-UK RCT protocol and recruitment. BMC Psychiatry, 2012, 12, 58.	2.6	14
108	Taking part in a pharmacogenetic clinical trial: assessment of trial participants understanding of information disclosed during the informed consent process. BMC Medical Ethics, 2013, 14, 34.	2.4	14

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109	Protocol for a quasi-experimental study of the effectiveness and cost-effectiveness of mother and baby units compared with general psychiatric inpatient wards and crisis resolution team services (The) Tj ETQq1 1	0.084314	l∎gBT /Over
110	Service user views and service user research in the <i>Journal of Mental Health</i> . Journal of Mental Health, 2011, 20, 423-428.	1.9	13
111	The UPBEAT depression and coronary heart disease programme: using the UK medical research council framework to design a nurse-led complex intervention for use in primary care. BMC Family Practice, 2012, 13, 119.	2.9	13
112	"But what if nobody's going to sit down and have a real conversation with you?―Service user/survivor perspectives on human rights. Journal of Public Mental Health, 2013, 12, 184-192.	1.1	13
113	Staff and service users' views on a â€~Consent for Contact' research register within psychosis services: a qualitative study. BMC Psychiatry, 2014, 14, 377.	2.6	13
114	Improving recruitment to healthcare research studies: clinician judgements explored for opting mental health service users out of the time to change viewpoint survey. Journal of Mental Health, 2019, 28, 42-48.	1.9	13
115	Clinical and economic outcomes from the UK pilot psychiatric services for personality-disordered offenders. International Review of Psychiatry, 2011, 23, 61-69.	2.8	12
116	Working alliance and its relationship to outcomes in a randomized controlled trial (RCT) of antipsychotic medication. BMC Psychiatry, 2013, 13, 28.	2.6	12
117	Expectations of new treatment in rheumatoid arthritis: developing a patientâ€generated questionnaire. Health Expectations, 2015, 18, 995-1008.	2.6	12
118	Identifying uncertainties about the effects of treatments for schizophrenia. Journal of Mental Health, 2006, 15, 263-268.	1.9	11
119	Recovery concept in a Norwegian setting to be examined by the assertive community treatment model and mixed methods. International Journal of Mental Health Nursing, 2018, 27, 147-157.	3.8	11
120	How can the service user voice be best heard at psychiatric meetings?. British Journal of Psychiatry, 2013, 203, 88-89.	2.8	9
121	On personal epiphanies and collective knowledge in survivor research and action. Social Theory and Health, 2020, 18, 110-122.	1.8	9
122	How do managers and leaders in the National Health Service and social care respond to service user involvement in mental health services in both its traditional and emergent forms? The ENSUE study. Health Services and Delivery Research, 2014, 2, 1-94.	1.4	9
123	Perinatal mental health services in pregnancy and the year after birth: the ESMI research programme including RCT. Programme Grants for Applied Research, 2022, 10, 1-142.	1.0	9
124	Health services research: Is there anything to learn from mental health?. Journal of Health Services Research and Policy, 2005, 10, 1-2.	1.7	8
125	Predicting psychiatric inpatient costs. Social Psychiatry and Psychiatric Epidemiology, 2016, 51, 303-308.	3.1	8
126	"This is not a Life Anyone would wantâ€â€"A Qualitative Study of Norwegian ACT Service users' Experience with Mental Health Treatment. Issues in Mental Health Nursing, 2018, 39, 519-526.	1.2	8

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127	Implementing civic engagement within mental health services in South East Asia: a systematic review and realist synthesis of current evidence. International Journal of Mental Health Systems, 2020, 14, 17.	2.7	8
128	Consenting for contact? Linking electronic health records to a research register within psychosis services, a mixed method study. BMC Health Services Research, 2015, 15, 199.	2.2	7
129	Improving the experience of care for adults using NHS mental health services: summary of NICE guidance. BMJ: British Medical Journal, 2012, 344, e1089-e1089.	2.3	6
130	Service users' and carers' views on research towards stratified medicine in psychiatry: a qualitative study. BMC Research Notes, 2015, 8, 489.	1.4	6
131	Critical qualitative research on â€~madness': knowledge making and activism among those designated â€~mad'. Wellcome Open Research, 2021, 6, 98.	1.8	6
132	A hidden activism and its changing contemporary forms: Mental health service users / survivors mobilising. Journal of Social and Political Psychology, 2018, 6, 728-744.	1.1	6
133	Stigma and schizophrenia – Authors' reply. Lancet, The, 2009, 373, 1336-1337.	13.7	5
134	Continuity of care for carers of people with severe mental illness: Results of a longitudinal study. International Journal of Social Psychiatry, 2013, 59, 663-670.	3.1	5
135	Linking a research register to clinical records in older adults' mental health services: a mixed-methods study. Alzheimer's Research and Therapy, 2015, 7, 15.	6.2	5
136	Patient involvement in improving the evidence base on mental health inpatient care: the PERCEIVE programme. Programme Grants for Applied Research, 2018, 6, 1-182.	1.0	5
137	Service user governors in mental health foundation trusts: accountability or business as usual?. Health Expectations, 2015, 18, 2892-2902.	2.6	4
138	Experience, madness theory, and politics. Philosophy, Psychiatry and Psychology, 2016, 23, 207-210.	0.4	4
139	An interpretative phenomenological analysis of the experience of receiving a diagnosis of bi-polar disorder. Journal of Mental Health, 2020, 29, 358-363.	1.9	4
140	The management of individuals with enduring moderate to severe mental health needs: a participatory evaluation of client journeys and the interface of mental health services with the criminal justice system in Cornwall. Health Services and Delivery Research, 2015, 3, 1-232.	1.4	4
141	Padrão global de discriminação experimentada e antecipada contra pessoas com esquizofrenia: estudo transversal. Revista Latinoamericana De Psicopatologia Fundamental, 2009, 12, 141-160.	0.0	4
142	UPBEAT-UK: a programme of research into the relationship between coronary heart disease and depression in primary care patients. Programme Grants for Applied Research, 2016, 4, 1-172.	1.0	4
143	Olanzapine and risperidone plasma concentration therapeutic drug monitoring: A feasibility study. Journal of Psychopharmacology, 2015, 29, 933-942.	4.0	3
144	Comment on the evaluation of the Time to Change anti-stigma campaign. The Psychiatrist, 2010, 34, 541-542.	0.3	2

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145	Changing nurses' views of the therapeutic environment: randomised controlled trial. BJPsych Open, 2019, 5, e17.	0.7	2
146	Strengthening Self-Determination of Persons with Mental Illness. , 2015, , 879-895.		2
147	Abnormal: Lectures at the Collège de France 1974–1975. By Michel Foucault (trans. G. Burchell). London: Verso. 2003. 88 pp. £20 (hb). ISBN 1 85984 539 8. British Journal of Psychiatry, 2004, 185, 443-443.	2.8	1
148	Psychometric properties of a new treatment expectation scale in rheumatoid arthritis: an application of item response theory. BMC Musculoskeletal Disorders, 2015, 16, 239.	1.9	1
149	Psychiatric Drugs: Key Issues and Service User Perspectives Jim Read, Macmillan Palgrave, 2009, £14.99 pb, 208 pp. ISBN 9780230549401. The Psychiatrist, 2010, 34, 215-215.	0.3	0