# Public Involvement Journal Club – articles discussed

Abelson, J., Li, K., Wilson, G., Shields, K., Schneider, C., and Boesveld, S. (2015) Supporting quality public and patient engagement in health system organizations: development and usability testing of the Public and Patient Engagement Evaluation Tool. *Health Expectations* 19(4) pp. 817-827 doi:10.1111/hex.12378. Available from: [*http://onlinelibrary.wiley.com/doi/10.1111/hex.12378/abstract*](http://onlinelibrary.wiley.com/doi/10.1111/hex.12378/abstract)

Barber, R. et. al (2011) Can the impact of public involvement in research be evaluated? A mixed methods study. *Health Expectations* doi: 10.1111/j.1369-7625.2010.00660.x. <http://onlinelibrary.wiley.com/doi/10.1111/j.1369-7625.2010.00660.x/pdf>

Barnes M, Newman J, Knops A & Sullivan H. (2003) Constituting ‘the public’ in public participation. *Public Administration* 81 (2) pp. 379-399. <http://onlinelibrary.wiley.com/doi/10.1111/1467-9299.00352/abstract>

Becker, S, Sempik, J & Bryman, A. (2010) Advocates, agnostics and adversaries: researchers’ perceptions of service user involvement in social policy research. *Social Policy & Society* 9(3) pp. 355-366. <http://journals.cambridge.org/action/displayFulltext?type=1&fid=7788187&jid=SPS&volumeId=9&issueId=03&aid=7788185>.

Beresford, P. (2013) From ‘other’ to involved: user involvement in research: an emerging paradigm. *Nordic Social Work Research* 3(2) pp. 139-148 DOI: 10.1080/2156857X.2013.835138

Blacksher, E. (2013) Participatory and Deliberative Practices in Health: Meanings, Distinctions, and Implications for Health Equity. *Journal of Public Deliberation* 9(1) article 6. Available from: <http://www.publicdeliberation.net/jpd/vol9/iss1/art6/>

Braun, K. and Schultz, S. (2010) **“**… a certain amount of engineering involved'': Constructing the public in participatory governance arrangements**.** *Public Understanding of Science* 2010 19: 403 Originally published online 9 October 2009 DOI: 10.1177/0963662509347814. Link to the article: [http://pus.sagepub.com/content/19/4/403.full.pdf+html](http://pus.sagepub.com/content/19/4/403.full.pdf%2Bhtml)

Brown, Rob (2016) unpublished paper presented at a conference ‘Systemic cooperation and the trustworthy state’. This paper summarises some key aspects of Rob’s doctoral work.

Cornwall, Andrea (2008) Unpacking ‘Participation’: models, meanings and practices. *Community Development Journal*, 43(3), pp. 269-283 <http://cdj.oxfordjournals.org/content/43/3/269>

Crocker, J., Boylan, A., Boystock, J. and Locock, L. (2016) Is it worth it? Patient and public views on the impact of their involvement in health research and its assessment: a UK- based qualitative interview study. *Health Expectations* doi: 10.1111/hex.12479

Dawson, S., Campbell, S., Giles, S., Morris, R., and Cheraghi-Sohi, S. (2018) Black and minority ethnic group involvement in health and social care research: A systematic review. *Health Expectations* 21: 3-22. DOI: 10.1111/hex.12597

Edelman, N. and Barron, D. (2015) Evaluation of public involvement in research: times for a major re-think? *Journal of Health Services Research & Policy*. Published online 27 October 2015 doi:10.1177/1355819615612510 [http://hsr.sagepub.com/content/early/2015/10/27/1355819615612510.full.pdf+html](http://hsr.sagepub.com/content/early/2015/10/27/1355819615612510.full.pdf%2Bhtml)

Fotaki, M. (2014) Can consumer choice replace trust in the National Health Service in England? Towards developing an affective psychosocial conception of trust in health care. *Sociology of Health & Illness* 36(8) pp. 1276–1294. <http://onlinelibrary.wiley.com/doi/10.1111/1467-9566.12170/pdf>

Gibson, A, Britten, N & Lynch, J. (2012) Theoretical directions for an emancipator concept of patient and public involvement. *Health* 16(5) pp. 531-547. <http://hea.sagepub.com/content/16/5/531.short>

Gradinger, F., Britten, N., Wyatt, K., Froggatt, K., Gibson, A., Jacoby, A., Lobban, F., Mayes, D., Snape, D., Rawcliffe, T., and Popay, J. (2013) Values associated with public involvement in health and social care research: a narrative review. To be published in *Health Expectations* doi: 10.1111/hex.12158. <http://onlinelibrary.wiley.com/doi/10.1111/hex.12158/pdf>

Green, G. (2016) Power to the people: To what extent has public involvement in applied health research achieved this? *Research Involvement and Engagement* (2016) 2:28 DOI 10.1186/s40900-016-0042-y

Hamilton, C., Hoens, A., Backman, C., McKinnon, A., McQuitty, S., English, K., and Li, L. (2017) ‘An empirically based conceptual framework for fostering meaningful patient engagement in research’ *Health Expectations*. First published: 6 October 2017 DOI: 10.1111/hex.12635 Available from: <http://onlinelibrary.wiley.com/doi/10.1111/hex.12635/full>

Johannesen, J. (2018) “The trouble with PPI (Public and Patient Involvement)”Keynote presentation, Cochrane Colloquium, Edinburgh, Tuesday September 18, 2018

Knaapen, L. and Lehoux, P. (2016) Three Conceptual Models of Patient and Public Involvement in Standard-setting: From Abstract Principles to Complex Practice. *Science as Culture*, 25:2, pp. 239-263. <http://www.tandfonline.com/doi/full/10.1080/09505431.2015.1125875>

Li, K., Abelson, J., Giacomini, M., and Contandriopoulos, D. (2015) Conceptualizing the use of public involvement in health policy decision-making. *Social Science & Medicine* 138 pp 14-21. <http://www.sciencedirect.com/science/article/pii/S0277953615003056>

Locock, L., Boylan, A.-M., Snow, R. and Staniszewska, S. (2016), The power of symbolic capital in patient and public involvement in health research. Health Expect. doi:10.1111/hex.12519 Available from: <http://onlinelibrary.wiley.com/doi/10.1111/hex.12519/abstract?campaign=wolearlyview>

Maguire, K. and Britten, N. (2018) ‘You’re there because you are unprofessional’: patient and public involvement as liminal knowledge spaces. *Sociology of Health and Illness* 40(3) pp 463-477. doi: 10.1111/1467-9566.12655 Available from: <https://onlinelibrary.wiley.com/doi/epdf/10.1111/1467-9566.12655>

Martin, G. (2008) ‘Ordinary people only’ : knowledge, representativeness and the publics of public participation in healthcare, *Sociology of Health and Illness* 30:1 pp 35-54 <http://onlinelibrary.wiley.com/doi/10.1111/j.1467-9566.2007.01027.x/abstract>

McCoy, M., Jongsma, K., Friesen, P., Dunn, M., Plunkett Neuhaus, C., Rand, L., and Sheehan, M. (2018) National Standards for Public Involvement in Research: missing the forest for the trees. *J Med Ethics*; 0:1-4. Doi: 10.1136/medethics-2018-105088.

Morenike Oluwatoyin Folayan & Bridget Haire (2016) Communitarian societies and public engagement in public health. *Critical Public Health*, DOI: 10.1080/09581596.2016.1252035. To link to this article: <http://dx.doi.org/10.1080/09581596.2016.1252035>

Osborne, S., Radnor, Z., and Strokosch, K. (2016, forthcoming) Co-production and the co-creation of value in public services: a suitable case for treatment? *Public Management Review*. Available from: <http://www.tandfonline.com/doi/full/10.1080/14719037.2015.1111927>

Prior, L. (2003) Belief, knowledge and expertise: the emergence of the lay expert in medical sociology. *Sociology of Health & Illness* Vol. 25 Silver Anniversary Issue, pp. 41–57. Available from: <https://onlinelibrary.wiley.com/doi/pdf/10.1111/1467-9566.00339>

Rycroft-Malone, J., Seers K., Titchen, A., Harvey, G., Kitson, A. and McCormack, B. (2004) What counts as evidence in evidence-based practice? *Journal of Advanced Nursing* 47(1), pp.81–90 <http://onlinelibrary.wiley.com/doi/10.1111/j.1365-2648.2004.03068.x/pdf>

Rowe, G. and Frewer, L.J. (2005) A Typology of Public Engagement Mechanisms. *Science, Technology, & Human Values* [online]. 30 (2), pp. 251-290. [Accessed 2 June 2015]. [http://sth.sagepub.com/content/30/2/251.full.pdf+html](http://sth.sagepub.com/content/30/2/251.full.pdf%2Bhtml)

Shippee N et al (2013) Patient and service user engagement in research: a systematic review and synthesized framework. *Health Expectations* doi:10.1111/hex.12090. <http://onlinelibrary.wiley.com/doi/10.1111/hex.12090/abstract>

Staley, K. (2017) ‘Changing what researchers “think and do”: Is this how involvement impacts on research?’ *Research for All*, 1 (1), 158–67. DOI 10.18546/RFA.01.1.13.

Staley K. et al. (2012) ‘The missing links’: understanding how context and mechanism influence the impact of public involvement in research. *Health Expectations* doi: 10.1111/hex.12017. <http://onlinelibrary.wiley.com/doi/10.1111/hex.12017/full>

Staley, K. (2015) ‘Is it worth doing?’ Measuring the impact of patient and public involvement in research. *Research Involvement and Engagement* 2015, 1:6 doi:10.1186/s40900-015-0008-5. Available from: <http://www.researchinvolvement.com/content/1/1/6>

S. Staniszewska, J. Brett, I. Simera, K. Seers, C. Mockford, S. Goodlad, D. G. Altman, D. Moher, R. Barber, S. Denegri, A. Entwistle, P. Littlejohns, C. Morris, R. Suleman, V. Thomas and C. Tysall (2017) GRIPP2 reporting checklists: tools to improve reporting of patient and public involvement in research. *Research Involvement and Engagement* (2017) 3:13 DOI 10.1186/s40900-017-0062-2

Thompson, J., Bissell, P., Cooper, C., Armitage, C. and Barber, R. (2012) Credibility and the ‘professionalized’ lay expert: Reflections on the dilemmas and opportunities of public involvement in health research. *Health* 16(6) pp 602-618. [http://hea.sagepub.com/content/16/6/602.full.pdf+html](http://hea.sagepub.com/content/16/6/602.full.pdf%2Bhtml)

Tritter, J. (2009) Revolution or evolution: the challenges of conceptualizing patient and public involvement in a consumerist world. *Health Expectations* 12, pp. 275-87. <http://onlinelibrary.wiley.com/doi/10.1111/j.1369-7625.2009.00564.x/abstract;jsessionid=3FEFB89BB28F752114A790C18ADE1C4F.d03t02>

Ward et al. (2009) Critical perspectives on ‘consumer involvement’ in health research: epistemological dissonance and the know-do gap. *Journal of Sociology* 46(1) pp. 63-82. <http://jos.sagepub.com/content/46/1/63>

Wilson, P., Mathie, E., Keenan, J., McNeilly, E., Goodman, C., Howe, A., Poland, F., Staniszewska, S., Kendall, S., Munday, D., Cowe, M., and Peckham S. ‘ReseArch with Patient and Public invOlvement: a RealisT evaluation – the RAPPORT study’ Scientific summary. *Health Services and Delivery Research* 2015; Vol. 3: No. 38. doi: 10.3310/hsdr03380 NIHR Journals Library [www.journalslibrary.nihr.ac.uk](http://www.journalslibrary.nihr.ac.uk/) Available from: <http://www.journalslibrary.nihr.ac.uk/__data/assets/pdf_file/0013/152302/ScientificSummary-hsdr03380.pdf>

Rosie Davies

12 September 2018