

Medical Sociology and Health Experiences Research Group

Publications

Francine Toye, Jennifer MacLellan, Sharon Dixon, Abigail McNiven, (2023). Understanding primary care perspectives on supporting women's health needs: a qualitative study, *British Journal of General Practice* 2023.

MacLean A, Wild C, Hunt K, et al (2023). Impact of Long Covid on the school experiences of children and young people: a qualitative study *BMJ Open* 2023.

Tanvi Rai, Bakita Kasadha, Shema Tariq, Sabrina Keating, Lisa Hinton, Angelina Namiba, Catherine Pope, (2023) Infant feeding as a transgressive practice in the context of HIV in the UK: A qualitative interview study, *Women's Studies International Forum*, Volume 101, 2023.

Cervantée E.K. Wild, Maria Inês Gandolfo Conceição, Miho Iwakuma, Sasha Lewis-Jackson, Rie Toyomoto, Alicia Regina Navarro Dias de Souza, Vinita Mahtani-Chugani, Rika Sakuma Sato, Tanvi Rai, (2023). Perceptions of government guidance and citizen responses during the COVID-19 pandemic: A cross-country analysis, *SSM - Qualitative Research in Health*, Volume 4, 2023.

Anna Dowrick, Rachel Grob, Akiko Sawada, Anne Thier, Christine Holmberg, Rika Sakuma Sato, (2023). Navigating responsible bio-political citizenship: Cross-country comparison of stigma in Covid-19 illness narratives in Germany, Japan, the UK and the USA, *SSM - Qualitative Research in Health*.

Anna Dowrick, Tanvi Rai, Lisa Hinton, Bella Eacott, Sylvan Baker, Maria Askew, Sue Ziebland, Louise Locock (2023) Health inequalities, ethnic minorities and COVID19: interactive theatre workshop drawing on a qualitative interview study. *The Lancet*, Volume 400, Supplement 1, 2022, Page S9.

Alice Maclean, Kate Hunt, Ashley Brown, Jane A. Evered, Anna Dowrick, Andrea Fokkens, Rachel Grob, Susan Law, Louise Locock, Michelle Marcinow, Lorraine Smith, Anna Urbanowicz, Nientke Verheij, Cervantee Wild, (2023). Negotiation of collective and individual candidacy for long Covid healthcare in the early phases of the Covid-19 pandemic: Validated, diverted and rejected candidacy, *SSM - Qualitative Research in Health*, Volume 3, 2023.

Suzanne Sayuri Ii, Hannah Jobling, Kelly Devenney & Sue Ziebland (2022) Connecting worlds: social work educators' perceptions on the role of lived experience in pedagogic practice, *Social Work Education*.

Anna Dowrick, Tanvi Rai, Lisa Hinton, Bella Eacott, Sylvan Baker, Maria Askew, Sue Ziebland, Louise Locock. (2022) Health inequalities, ethnic minorities and COVID19: interactive theatre workshop drawing on a qualitative interview study. *The Lancet*, November 2022.

Kasadha B, Tariq S, Nyatsanza F, Freeman-Romilly N, Namiba A, Rai T. (2021) Stakeholder engagement is essential to maximise the impact of research on infant feeding in the context of HIV. *Therapeutic Advances in Infectious Disease*. January 2021.

Mikulak M., Ryan S., Ma R., Martin S., Stewart J., Davidon S., Stepney M. (2021) Health professionals' identified barriers to trans healthcare: a qualitative interview study, *British Journal of General Practice*

Ghio, D., Greenwell, K., Muller, I., Roberts, A., McNiven, A. and Santer, M. (2021), Psychosocial needs of adolescents and young adults with eczema: A secondary analysis of qualitative data to inform a behaviour change intervention. *Br. J. Health Psychol.*, 26.

de Vere Hunt IJ, McNiven A, Roberts A, et al (2021) 'Not just a piece of skin in front of you'—a qualitative exploration of the experiences of adolescents with eczema and psoriasis with healthcare professionals *BMJ Open* 2021.

Greenwell K, Ghio D, Muller I, et al (2021) Taking charge of eczema self-management: a qualitative interview study with young people with eczema *BMJ Open* 2021.

Magdalena Mikulak (2021): For whom is ignorance bliss? ignorance, its functions and transformative potential in trans health, *Journal of Gender Studies*.

de Vere Hunt I., McNiven A., McPherson T. (2020) [A qualitative exploration of the experiences of adolescents with alopecia areata and their messages for healthcare professionals. *British Journal of Dermatology*.](#)

Ghio D., Muller I., Greenwell K., Roberts A., McNiven A., Langan SM., Santer M. (2020) ['It's like the bad guy in a movie who just doesn't die': a qualitative exploration of young people's adaptation to eczema and implications for self-care. *British Journal of Dermatology* 10.](#)

Ip A., Muller I., Geraghty AWA., McNiven A., Little P., Santer M. (2020) Young people's perceptions of acne and acne treatments: secondary analysis of qualitative interview data. *Br J Dermatol*, Vol 183.

I. de Vere Hunt, A. McNiven and T. McPherson. (2020) 'You're harnessed into that rollercoaster no matter what': a qualitative exploration of the psychological impact of alopecia areata in adolescence. *British Association of Dermatologists* 183 (Suppl. 1).

Ziebland, S., Grob, R. and Schlesinger, M., (2020). Polyphonic perspectives on health and care: Reflections from two decades of the DIPEX project. *Journal of Health Services Research & Policy*.

Madeley, A.M., Williams, V. and McNiven, A., 2019. An interpretative phenomenological study of midwives supporting home birth for women with complex needs. *British Journal of Midwifery*, 27(10), pp.625-632.

Ip, A., Muller, I., Geraghty, A.W.A., McNiven, A., Little, P. and Santer, M., 2019. Young people's perceptions of acne and acne treatments: secondary analysis of qualitative interview data. *British Journal of Dermatology*.

Ghio, D., Muller, I., Greenwell, K., Roberts, A., McNiven, A., Langan, S.M. and Santer, M., 2020. 'It's like the bad guy in a movie who just doesn't die': a qualitative exploration of young people's adaptation to eczema and implications for self-care. *British Journal of Dermatology*, 182(1), pp.112-118.

MacArtney, J.I., Andersen, R.S., Malmström, M., Rasmussen, B. and Ziebland, S. (2020), The convivial and the pastoral in patient–doctor relationships: a multi-country study of patient stories of care, choice and medical authority in cancer diagnostic processes. *Sociol Health Illn*, 42: 844-861. doi:10.1111/1467-9566.13067

Albury, C., Strain, W.D., Le Brocq, S., Logue, J., Lloyd, C., Tahrani, A. and Language Matters working group, 2020. The importance of language in engagement between health-care professionals and people living with obesity: a joint consensus statement. *The Lancet Diabetes & Endocrinology*, 8(5), pp.447-455.

Dougall, G., Franssen, M., Tucker, K.L., Yu, L.M., Hinton, L., Rivero-Arias, O., Abel, L., Allen, J., Band, R.J., Chisholm, A. and Crawford, C., 2020. Blood pressure monitoring in high-risk pregnancy to

improve the detection and monitoring of hypertension (the BUMP 1 and 2 trials): protocol for two linked randomised controlled trials. *BMJ open*, 10(1).

Locock L, Graham C, King J, et al. Understanding how front-line staff use patient experience data for service improvement: an exploratory case study evaluation. Southampton (UK): NIHR Journals Library; 2020 Mar. (Health Services and Delivery Research, No. 8.13.) Available from: <https://www.ncbi.nlm.nih.gov/books/NBK554766/>

Morgan, B., Yu, L-M., Solomon, T., Ziebland, S. 2020. Assessing health research grant applications: A retrospective comparative review of a one-stage versus a two-stage application assessment process. *PLOS ONE* 15(3)e0230118 <https://doi.org/10.1371/journal.pone.0230118>

Turk, A., Fleming, J., Powell, J., Atherton, H. 2020. Exploring UK doctors' attitudes towards online patient feedback: Thematic analysis of survey data. *Digital Health*. 6/ 2055207620908140 DOI: 10.1177/2055207620908148

Powell, J., Williams, V., Atherton, H., Bennett, K., Yang, Y., Davoudianfar, M., Hellsing, A., Martin, A., Mollison, J., Shanyinde, M., Yu, L.M., Griffiths, K.M. 2020. Effectiveness and Cost-Effectiveness of a Self-Guided Internet Intervention for Social Anxiety Symptoms in a General Population Sample: Randomized Controlled Trial. *J Med Internet Res* 2020;22(1):e16804 DOI: 10.2196/16804

Muinga, N., Magare, S., Monda, J., English, M., Fraser, H., Powell, J., Paton, C. 2020. Digital health Systems in Kenyan Public Hospitals: a mixed-methods survey. *BMC Med Inform Decis Mak* 20(2)2020. <https://doi.org/10.1186/s12911-019-1005-7>

MacArtney, J.I., Andersen, R.S., Malmström, M., Rasmussen, B. and Ziebland, S. (2020), The convivial and the pastoral in patient–doctor relationships: a multi-country study of patient stories of care, choice and medical authority in cancer diagnostic processes. *Sociol Health Illn*. doi:10.1111/1467-9566.13067

Locock, L., Montgomery, C., Parkin, S., Chisholm, A., Bostock, J., Dopson, S., ... Ziebland, S. (2020). How do frontline staff use patient experience data for service improvement? Findings from an ethnographic case study evaluation. *Journal of Health Services Research & Policy*. <https://doi.org/10.1177/1355819619888675>

Rolfe, U., Pope, C., Crouch, R. 2020 Paramedic performance when managing patients experiencing mental health issues – Exploring paramedics' Presentation of Self. *International Emergency Nursing* <https://doi.org/10.1016/j.ienj.2019.100828>

Smith, S., Alvand, A., Locock, L., Ryan, S., Smith, J., Bayliss, L., Wilson, H., Price, A. 2019. Partial or total knee replacement? Identifying patients' information needs on knee replacement surgery: a qualitative study to inform a decision aid. *Quality of Life Research*. DOI: 10.1007/s11136-019-02381-9

Rai, T., Bruton, J., Kall, M., Ma, R., Pufall, E., Day, S., Delpech, V., Ward, H. 2019. Experience of primary care for people with HIV: a mixed-method analysis. *BJGP Open*. [bjgpopen19X101665](https://doi.org/10.3399/bjgpopen19X101665). <https://doi.org/10.3399/bjgpopen19X101665>

Bridges, J., Pope, C., Braithwaite, J. 2019. Making health care responsive to the needs of older people. *Age and Ageing*. 48(6)785-788 10.1093/ageing/afz085

Tompson, A., Nicholson, B.D., Ziebland, S., Evans, J., Bankhead, C. 2019. Quality improvements of safety-netting guidelines for cancer in UK primary care: insights from a qualitative interview study of GPs. *British Journal of General Practice*. DOI: <https://doi.org/10.3399/bjgp19X706565>

Powell, J., Atherton, H., Williams, V., Mazanderani, F., Dudhwala, F., Woolgar, S., Boylan, A.M., Fleming, J., Kirkpatrick, S., Martin, A., van Velthoven, M., de Jongh, A., Findlay, D., Locock, L., Ziebland, S. 2019 Using online patient feedback to improve NHS services: the INQUIRE multimethod study. *Health Services and Delivery Research* 2019. 7(38) DOI: 10.3310/hsdr07380

Price, J., Wilcox, M., Kabudula, C.W., Herbst, K., Hinton, L., Kahn, K., Harnden, A. 2019. Care pathways during a child's final illness in rural South Africa: Findings from a social autopsy study. *Plos One* <https://doi.org/10.1371/journal.pone.0224284>

Powell, J. 2019. Trust me I'm a chatbot: why AI in healthcare won't pass the Turing test. *JMIR Publications* DOI: 10.2196/16222

Stevenson, F., Hall, L., Sequin, M., Atherton, H., Barnes, R., Leydon, G., Pope, C., Murray, E., Ziebland, S. 2019. General Practitioner's use of online resources during medical visits: managing the boundary between inside and outside the clinic. *Sociology of Health & Illness*. 41/S1/65-81 doi: 10.1111/1467-9566.12833

Evans, J., MacArtney, J.I., Bankhead, C., Albury, C., Jones, D., Ziebland, S., Nicholson, B.D. 2019. How do GPs and patients share the responsibility for cancer safety netting follow-up actions? A qualitative interview study of GPs and patients in Oxfordshire, UK. *BMJ Open* 9(9)e029316. DOI: 10.1136/bmjopen-2019-029316

Boylan, A.-M., Williams, V., & Powell, J. 2019. Online patient feedback: a scoping review and stakeholder consultation to guide health policy. *Journal of Health Services Research & Policy*. <https://doi.org/10.1177/1355819619870837>

Harvey, J., Powell, J. 2019. Harnessing mobile devices to support the delivery of community-based clinical care: a participatory evaluation. *BMC Medical Informatics and Decision Making*. 19 (1) 134 DOI: 10.1186/s12911-019-0869-x

Ferrando, P., Gould, D.W., Walmsley, E., Richards-Belle, A., Canter, R., Saunders, S., Harrison, D.A., Harvey, S., Heyland, D.K., Hinton, L., McColl, E., Richardson, A., Richardson, M., Wright, S.E., Rowan, K.M. 2019. Family satisfaction with critical care in the UK: a multicentre cohort study. *BMJ Open* 9(8)e028956 DOI: 10.1136/bmjopen-2019-028956

Methodological decisions influence the identification of potential core outcomes in pre-eclampsia related studies: a systematic review and sensitivity analysis informing guidelines for future core outcome set developers. 2019. Duffy, J.M.N., Hirsch, M., Ziebland, S., McManus, R., International Collaboration to Harmonise Outcomes in Pre-eclampsia (iHOPE). *BJOG: Int J Obstet Gy*

Moscrop, A., Ziebland, S., Roberts, N., Papanikitas, A. 2019 A systematic review of reasons for and against asking patients about their socioeconomic contexts. *International Journal for Equity in Health*. 18(1)112

Albury, C. Hall, A.Syed, A. Ziebland, S. Stokoe, E. Roberts, N. Webb, H. Aveyard, P. Communication practices for delivering health behaviour change conversations in primary care: A systematic review and thematic synthesis. Accepted by *BMC family practice*, July 2019

Rikke Sand Andersen, John McArtney, Birgit H. Rasmussen, Britt-Marie Bernhardson, Senada Hajdarevic, Marlene Malmström & Sue Ziebland (2019) Caring as sharing. Negotiating the moral boundaries of receiving care, *Critical Public Health*, DOI: 10.1080/09581596.2019.1623381

Nagraj, S., Hinton, L., Praveen, D., Kennedy, S., Norton, R., Hirst, J. 2019 Women's and healthcare providers' perceptions of long-term complications associated with hypertension and diabetes in pregnancy: a qualitative study.

Ryan, S. 2019. NHS Inquiries and Investigations; an Exemplar in Peculiarity and Assumption. The Political [Quarterly](https://doi.org/10.1111/1467-923X.12703). doi.org/10.1111/1467-923X.12703

Atherton, H., Fleming, J., Williams, V., Powell, J. 2019 Online patient feedback: a cross sectional survey of the attitudes and experiences of United Kingdom health care professionals. *Journal of Health Services Research & Policy*. DOI:10.1177/1355819619844540

Katherine Runswick-Cole & Sara Ryan (2019) Liminal still? Unmothering disabled children, *Disability & Society*, DOI: 10.1080/09687599.2019.1602509

Farre, A., Ryan, S., McNiven, A., et al. (2019). The impact of arthritis on the educational and early work experiences of young people: a qualitative secondary analysis. *International Journal of Adolescent Medicine and Health*, 0(0), pp. -. Retrieved 27 Mar. 2019, from doi:10.1515/ijamh-2018-0240

Powell, J. and Deetjen, U., 2019. Characterizing the Digital Health Citizen: Mixed-Methods Study Deriving a New Typology. *Journal of medical Internet research*, 21(3), p.e11279.

Dalton, L., Rapa, E., Ziebland, S., Rochat, T., Kelly, B., Hanington, L., Bland, R., Yousafzai, A., Stein, A., Betancourt, T. and Bluebond-Langner, M., 2019. Communication with children and adolescents about the diagnosis of a life-threatening condition in their parent. *The Lancet*, 393(10176), pp.1164-1176.

The Lancet Editorial, 2019. Let's talk with children about life-threatening diseases. 393,1072

Crocker, J., A. Hughes-Morley, S. Petit-Zeman and S. Rees (2015). "Assessing the impact of patient and public involvement on recruitment and retention in clinical trials: a systematic review." *Trials* 16(Suppl 2): O91-O91.

Crocker, J., Rees, S., Locock, L., Petit-Zeman, S., Chant, A., Treweek, S., Cook, J., Farrar, N., Woolfall, K. and Bulbulia, R., 2015. Developing a patient and public involvement intervention to enhance recruitment and retention in surgical trials (PIRRIST): study protocol. *Trials*, 16(S2), p.P89.

Crocker, J.C., Rees, S., Locock, L., Petit-Zeman, S., Chant, A., Treweek, S., Cook, J.A., Farrar, N., Woolfall, K., Bostock, J. and Bowman, L., 2016. Developing a patient and public involvement intervention to enhance recruitment and retention in UK surgical trials (PIRRIST). *International Journal of Surgery*, (36), pp.S140-S141. <http://dx.doi.org/10.1016%2Fj.ijsu.2016.11.045>

Crocker, J., J. Bostock and R. Bulbulia (2016). PPI: calling all surgical researchers... *Bulletin of the Royal College of Surgeons* 98(1): 30-33.

Crocker, J., N. Farrar, S. Treweek, S. Petit-Zeman, A. Chant, J. Bostock, K. Woolfall, Locock, S. Rees and R. Bulbulia (2018). "#15 Applying the IDEAL framework to a methodological complex intervention (PIRRIST) (poster presentation)." *International Journal of Surgery* 59: S4.

Crocker, J., Rees, S., Locock, L., Petit-Zeman, S., Chant, A., Treweek, S., Cook, J., Farrar, N., Woolfall, K., Bostock, J. and Harmston, R., 2018. # 3 PIRRIST: A patient and public involvement (PPI) intervention to enhance recruitment and retention in surgical trials (oral presentation). *International Journal of Surgery*, 59, pp.S1-S2.

Mazanderani, F., Hughes, N., Hardy, C., Sillence, E. and Powell, J., 2019. Health information work and the enactment of care in couples and families affected by Multiple Sclerosis. *Sociology of health & illness*. 41(2)pp 395-410 doi: 10.1111/1467-9566.12842

Albury, C., Stokoe, E., Ziebland, S., Webb, H. and Aveyard, P., 2018. GP-delivered brief weight loss interventions: a cohort study of patient responses and subsequent actions, using conversation analysis in UK primary care. *Br J Gen Pract*, 68(674), pp.e646-e653.

Crocker, J.C., Pratt-Boyden, K., Hislop, J., Rees, S., Locock, L., Olszowski, S., Chant, A., Treweek, S., Cook, J.A., Woolfall, K., Farrar, N., Bostock, J., Bulbulia, R. 2019. Patient and public involvement (PPI) in UK surgical trials: a survey and focus groups with stakeholders to identify practices, views, and experiences. *Trials*. 20(1)119. <https://doi.org/10.1186/s13063-019-3183-0>

Ziebland, S.B., Stevenson, F., Pope, C., Greenhalgh, T., Murray, E., Atherton, H.C., Barnes, R., Leydon, G. and Seguin, M., 2018. General Practitioners use of online resources during medical visits: Managing the boundary between inside and outside the clinic. *Sociology of Health and Illness*.

Duffy JMN, Thompson T, Hinton L, Salinas M, McManus RJ, Ziebland S, The International Collaboration to Harmonise Outcomes in Pre-eclampsia (iHOPE) Qualitative Research Group. 2019 What outcomes should researchers select, collect, and report in pre-eclampsia research? A qualitative study exploring the views of women with lived experience of pre-eclampsia. *BJOG An International Journal of Obstetrics & Gynaecology*. <https://doi.org/10.1111/1471-0528.15616>

Locock, L., Kirkpatrick, S., Brading, L., Sturme, G., Cornwell, J., Churchill, N., Glenn, R. 2019. Involving service users in the qualitative analysis of patient narratives to support healthcare quality improvement.

Cook, J.A., Merritt, N., Rees, J.L., Crocker, J.C., Hopewell, S., Dritsaki, M., Beard, D.J., Rangan, A., Cooper, C., Kottam, L., Farrar-Hockley, D., Thomas, M., Earle, R., Carr, A.J. 2018. Patch-augmented rotator cuff surgery (PARCS) study-protocol for a feasibility study.

Greaves, F., Joshi, I., Campbell, M., Roberts, S., Patel, N., Powell, J. 2018. What is an appropriate level of evidence for a digital health intervention? *The Lancet*. [https://doi.org/10.1016/S0140-6736\(18\)33129-5](https://doi.org/10.1016/S0140-6736(18)33129-5)

Moore, H., Solomons, W., Hinton, L. 2018. Life-threatening complications in childbirth: a discursive analysis of fathers' accounts. *Journal of reproductive and infant psychology* pp 1-11. <https://doi.org/10.1080/02646838.2018.1546382>

Crocker, J.C., Ricci-Cabello, I., Servet, M., Parker, A., Hirst, J.A., Chant, A., Petit-Zeman, S., Evans, D., Rees, S. 2018. Impact of patient and public involvement on enrolment and retention in clinical trials: systematic review and meta-analysis.

Hajdarevic, S., Rasmussen, B. H., Hasle, T. L. O., & Ziebland, S. (2018). Qualitative cross-country comparison of whether, when and how people diagnosed with lung cancer talk about cigarette smoking in narrative interviews. *BMJ open*, 8(11), e023934.

Stepney, M., Aveyard, P., Begh, R. 2018. GP and nurse perceptions of e-cigarettes in England: a qualitative interview study. *Br J Gen Pract*. bjpgp18X699821. DOI <https://doi.org/10.3399/bjpgp18X699821>

Duffy, J.M.N., Ziebland, S., Von Dadelszen, P., McManus, R.J. 2018. Tackling poorly selected, collected, and reported outcomes in obstetrics and gynecology research. *American Journal of Obstetrics and Gynecology* doi: <https://doi.org/10.1016/j.ajog.2018.09.023>.

Dixon S. 2015. The FGM enhanced dataset: how are we going to discuss this with our patients? (letter) *Br J Gen Pract*. 65 (641) 629-629 DOI: <https://doi.org/10.3399/bjpgp15X687781>

Darbyshire, J.L., Hinton, L. 2018 Using patient narratives to design an intervention to reduce noise in the Intensive Care Unit. *The Journal of Health Design*. 3(2)

<https://doi.org/10.21853/JHD.2018.51>

Dixon S, Agha K, Ali F, El-Hindi L, Kelly B, Locock L, Otoo-Oyortey N, Penny S, Plugge E, Hinton L. (2018) Female genital mutilation (FGM) in the UK- where are we, where do we go next? How to co-design research with communities? *BMC Research Involvement and Engagement*.

4(1)29 doi: 10.1186/s40900-018-0103-5

Ryan S, Julian G. (2016) #107days: Personal reflection on an exercise (transformative, subversive or accidental?) in effective action. In M.Kent and K.Ellis [Eds] *Disability and Social Media*. pp25-40. Routledge; Abingdon.

Julian G, Ryan S. 2018. #JusticeforLB. in S. Carr and P. Beresford (Eds) *Social Policy First Hand*. pp319-323, Policy Press; Bristol.

Ryan, S., 2005. 'People don't do odd, do they?' mothers making sense of the reactions of others towards their learning disabled children in public places. *Children's Geographies*, 3(3), pp.291-305.

Ryan S. (2005) 'Busy behaviour' in the 'Land of the Golden M': going out with learning disabled children in public places. *Journal of Applied Research in Intellectual Disabilities*, 18(1), 65-74. doi.org/10.1111/j.1468-3148.2004.00225.x

Ryan, S. (2006) It takes two to tango... but what if one can't dance and the other doesn't want to?: A response to van de Ven et al. *Disability and Society*, 21 (1), pp91-92

doi.org/10.1080/096875905003757

Ryan S. (2008) 'I used to worry about what other people thought but now I just think... well I don't care': Shifting accounts of learning difficulties in public places. *Health & Place*, 14(4), 730-739.

doi.org/10.1016/j.healthplace.2007.11.004

Ryan S, Runswick-Cole K. (2008) Repositioning mothers: Mothers, disabled children and disability studies. *Disability & Society*, 23(3), 199-210. doi.org/10.1080/09687590801953937

Pyer, M., Horton, J., Tucker, F., Ryan, S, Kraftl, P., (2010) Children, young people and 'disability': challenging children's geographies? *Children's Geographies*. 8 (1), 1-8. [Co-editor of special journal issue] doi.org/10.1080/14733280903500059

Searle A, Neville P, Ryan S, Waylen A. The Role of the Clinical Nurse Specialist From the Perspective of Parents of Children Born With Cleft Lip and/or Palate in the United Kingdom: A Qualitative Study. *Clinical Nurse Specialist*. 2018 May 1;32(3):121-8. doi: 10.1097/NUR.0000000000000371

Dasgupta, R., Speakman, M., Ray, A., Powell, J., Modi, S., Carolan-Rees, G., & Hacking, N. (2018). MP62-09 Prostate artery embolisation versus TURP; a multicentric prospective comparison: the uk-rope study. *The Journal of Urology*, 199(4), e835. <https://doi.org/10.1016/j.juro.2018.02.2010>

Albury, C., Stokoe, E., Ziebland, S., Webb, H., Aveyard, P. 2018. GP delivered brief weight loss interventions: In-consultation responses, and subsequent patient action. *British Journal of General Practice*.

Seguin, M., Hall, L., Atherton, H., Barnes, R., Leydon, G., Murray, E., Pope, C., Ziebland, S., Stevenson, F.A.. 2018. Protocol paper for the 'Harnessing resources from the internet to maximise outcomes from GP consultations (HaRI) study: a mixed qualitative methods study. *BMJ Open* 2018;8:e024188. doi: 10.1136/bmjopen-2018-024188

Ziebland, S., Rasmussen, B., MacArtney, J., Hajdarevic, S., Sand Andersen, R. 2018. How wide is the Goldilocks Zone in your health system? *Journal of Health Services Research & Policy*. 1355819618790985 DOI 10.1177/1355819618790985

Robinson H., Williams V., Curtis F., Bridle C., Jones AW. 2018. Facilitators and barriers to physical activity following pulmonary rehabilitation in COPD: a systematic review of qualitative studies. *Primary Care Respiratory Medicine*. 28(1)19 10.1038/s41533-018-0085-7

Brant, H.D., Atherton, H., Bikker, A., Porqueddu, T., Salisbury, C., McKinstry, B., Campbell, J., Gibson, A., Ziebland, S. 2018. Receptionists' role in new approaches to consultations in primary care: a focused ethnographic study. *Br J Gen Pract* bjgp18X697505. DOI: <https://doi.org/10.3399/bjgp18X697505>

Malmströma, M., Rasmussena, B.H., Bernhardsonc, B.M., Hajdarevicd, S., Erikssonc, L.E., Anderseng, R.S., MacArtney, J.I. 2018. It is important that the process goes quickly, isn't it?" A qualitative multicountry study of colorectal or lung cancer patients' narratives of the timeliness of diagnosis and quality of care. *European Journal of Oncology Nursing*. 34 82-88 DOI: 10.1016/j.ejon.2018.04.002

Hinton, L., Locock, L., Long, A-M., & Knight, M. 2018. What can make things better for parents when babies need abdominal surgery in their first year of life? : A qualitative interview study in the UK *BMJ Open* , vol 8 , e020921 . DOI: 10.1136/bmjopen-2017-020921

Stepney, M, Kirkpatrick, S, Locock, L, Prinjha, S, Ryan, S. 2018. 'A licence to drive? Neurological illness, loss and disruption.' *Sociology of Health and Illness*. <https://doi.org/10.1111/1467-9566.12754>

Evans, J., Ziebland, S., MacArtney, J.I., Bankhead, C.R., Rose, P.W., and Nicholson, B.D. 2018. GPs' understanding and practice of safety netting for potential cancer presentations: a qualitative study in primary care. *Br J Gen Pract* 8 May 2018 DOI: <https://doi.org/10.3399/bjgp18X696233>

Muinga N., Magare S., Monda J., Kamau O., Houston S., Fraser H., Powell J., English M., Paton C. 2018. Implementing an open source electronic health record system in kenyan health care facilities: Case *JMIR Med Inform* 2018; 6(2):e22

Powell J, Speakman M, Longford N, DasGupta R, Modi S, Dyer J, Harris M, Carolan-Rees G, Hacking N. 2018. The UK ROPE Study: efficacy and safety of prostate artery embolisation for benign prostatic hyperplasia. An observational study and propensity matched comparison with transurethral resection of the prostate. *BJUI*. <https://doi.org/10.1111/bju.14249>

van Velthoven MH., Atherton H., Powell J. 2018. A cross sectional survey of the UK public to understand use of online ratings and reviews of health services. *Patient Education and Counseling*. 101(9)1690-1696 <https://doi.org/10.1016/j.pec.2018.04.001>

Eassey, D., Reddel, H.K., Foster, J.M., Kirkpatrick, S., Locock, L., Ryan, K., & Smith, L. 2018 "...I've said I wish I was dead, you'd be better off without me": A systematic review of people's experiences of living with severe asthma. *Journal of Asthma*. DOI: 10.1080/02770903.2018.1452034

Stevens, R.J., Evans, J., Oke, J., Smart, B., Hobbs, F.D.R., Holloway, E., Horwood, J., Judd, M., Locock, L., McLellan, J., Perera, R. 2018. Kidney Age, not Kidney Disease. *CMAJ* April 03, 2018 190 (13) E389-E393; DOI: <https://doi.org/10.1503/cmaj.170674>

Tucker, K. L., Bowen, L., Crawford, C., Mallon, P., Hinton, L., Lee, M.M., Oke, J., Taylor, K.S., Heneghan, C., Bankhead, C., Mackillop, L., James, T., Oakeshott, P., Chappell, L.C., McManus, R.J. 2018. The feasibility and acceptability of self-testing for proteinuria during pregnancy: A mixed methods approach. *Pregnancy Hypertension-an International Journal of Womens Cardiovascular Health*. Vol 12, 161-168. <https://doi.org/10.1016/j.preghy.2017.11.009>

Atherton, H., Salisbury, C., Ziebland, S., Campbell, J., McKinstry, B., Brant, H., Bikker, A., Gibson, A. 2018. The potential of alternatives to face-to-face consultation in general practice, and the impact on different patient groups: a mixed-methods case study. *NIHR Health Services and Delivery Research*, volume 6, number 20

Knowles S., Hays R., Senra H., Bower P., Locock L., Protheroe J., Sanders C., Daker-White G. 2018. Empowering people to help speak up about safety in primary care: Using codesign to involve patients and professionals in developing new interventions for patients with multimorbidity. *Health Expectations*. 21(2)539-548 <https://doi.org/10.1111/hex.12648>

McNiven, A. 2018. 'Disease, illness, affliction? Don't know': Ambivalence and ambiguity in the narratives of young people about having acne. *Health*. <https://doi.org/10.1177%2F1363459318762035>

Van Velthoven M, Powell J, Powell G. Problematic smartphone use: digital approaches to an emerging public health problem. *Digital Health* 2018. 5 Mar 2018. <https://doi.org/10.1177/2055207618759167>

Boylan, A.M.R., Locock, L., Machin, L. 2018. From waste product to blood, brains and narratives: Developing a pluralist sociology of contributions to health research. *Sociology of Health & Illness*. <https://doi.org/10.1111/1467-9566.12715>

Kirkpatrick, S., Locock, L., Farre, A., Ryan, S., Salisbury, H., McDonagh, J.E. 2018. Untimely illness: When diagnosis does not match age-related expectations. *Health Expectations* 2018 (1-11) DOI: 10.1111/hex.12669

Atherton, H.C., Brant, H., Ziebland, S., Bikker, A., Campbell, J., Gibson, A., McKinstry, B., Porqueddu, T. and Salisbury, C. 2018. Alternatives to the face-to-face consultation in general practice : focused ethnographic case study. *British Journal of General Practice* 68 (669): e293-e300 DOI: <https://doi.org/10.3399/bjgp18X694853>

Ryan, S. 2018 Honouring a life and narrative work: John's story. *Arts and Humanities in Higher Education*, 17(1)58-68 doi: 10.1177/1474022217729178

Hinton, L., Dumelow, C., Rowe, R., Hollowell, J. (2018) Birthplace choices: what are the information needs of women when choosing where to give birth in England? A qualitative study using online and face to face focus groups. *BMC Pregnancy and Childbirth*. 2018(18)12 DOI 10.1186/s12884-017-1601-4

Tucker, K. L., Taylor, K.S., Crawford, C., Hodgkinson, J.A., Bankhead, C., Carver, T., Ewers, E., Glogowska, M., Greenfield, S.M., Ingram, L., Hinton, L., Khan, K.S., Locock, L., Mackillop, L., McCourt, C., Pirie, A.M., Stevens, R., McManus, R.J. 2017. Blood pressure self-monitoring in pregnancy: examining feasibility in a prospective cohort study. *BMC Pregnancy and Childbirth*. 17:442 <https://doi.org/10.1186/s12884-017-1605-0>

Hinton, L., Tucker, K. L., Greenfield, S.M., Hodgkinson, J.A., Mackillop, L., McCourt, C., Carver, T., Crawford, C., Glogowska, M., Locock, L., Selwood, M., Taylor, K.S., McManus, R.J. 2017. Blood

pressure self-monitoring in pregnancy (BuMP) feasibility study; a qualitative analysis of women's experiences of self-monitoring. *BMC Pregnancy and Childbirth*. 17:427.

<https://doi.org/10.1186/s12884-017-1592-1>

J.M.N. Duffy, M. Hirsch, L. Pealing, M. Showell, K.S. Khan, S. Ziebland, R.J. McManus on behalf of the International Collaboration to Harmonise Outcomes in Pre-eclampsia (iHOPE) (UK, New Zealand). 2017. Inadequate safety reporting in pre-eclampsia trials: a systematic evaluation. *BJOG*, Volume 125, Issue 7, June 2018, pages 795-803. <https://doi.org/10.1111/1471-0528.14969>

Chapple, A & Ziebland, S. Challenging Explanations for the lack of Senior Women in Science? Reflections from Successful Women Scientists at an elite British University. *International Journal of Gender, Science and Technology*. [S.l.], v. 9, n. 3, p. 298-315, Jan. 2018. ISSN 2040-0748

MacArtney, J., Malmström, M., Overgaard Nielsen, T., Evans, J., Bernhardson, B.M., Hajdarevic, S., Chapple, A., Eriksson, L.E., Locock, L., Rasmussen, B., Vedsted, P., Tishelman, C., Sand Andersen, R., Ziebland, S. 2017. Patients' initial steps to cancer diagnosis in Denmark, England and Sweden: what can a qualitative, cross-country comparison of narrative interviews tell us about potentially modifiable factors? *BMJ Open* 2017;7:e018210. doi: 10.1136/bmjopen-2017-018210

Chapple, A., Ziebland, S. 2018. Methodological and Practical Issues in Cross-National Qualitative Research: Lessons From the Literature and a Comparative Study of the Experiences of People Receiving a Diagnosis of Cancer. *Qualitative Health Research*. 28(5)789-799. Article first published online: November 2, 2017; Issue published: April 1, 2018,

<https://doi.org/10.1177/1049732317736284>

Ridge D, Broom A. Kokanović, R. Ziebland, S. Hill, N. 2017 Depression at work, authenticity in question: Experiencing, concealing and revealing. *Health* 2017; 1-18 DOI 10.1177/1363459317739437

Keltie, K., Elneil, S., Monga, A., Patrick, H., Powell, J., Campbell, B., Sims, AJ. 2017. Complications following vaginal mesh procedures for stress urinary incontinence: An 8 year study of 92,246 women. *Nature Scientific Reports*. doi:10.1038/s41598-017-11821-w

"Williams, V., Ryan, S. 2017. "I just know": exploring self-knowledge in chronic obstructive pulmonary disease. *Palgrave Communications*. Vol 3. 17089 DOI 10.1057/palcomms.2017.89

<https://www.nature.com/articles/palcomms201789#supplementary-information>"

Conducting a team-based multi-sited focused ethnography in primary care A.P. Bikker, H. Atherton, H. Brant, T. Porqueddu, J.L. Campbell, A. Gibson, B. McKinstry, C. Salisbury and S. Ziebland *BMC Medical Research Methodology* (2017) 17:139 DOI 10.1186/s12874-017-0422-5

Dudhwala F., Boylan A-M., Williams V., Powell J. 2017. VIEWPOINT: What counts as online patient feedback, and for whom? *Digital Health*. Vol 3. 1-3. <https://doi.org/10.1177%2F2055207617728186>

Duffy, J.M.N., Hirsch, M., Gale, C., Pealing, L., Kawsar, A., Showell, M., Williamson, P.R., Khan, K.S., Ziebland, S., McManus, R.J. 2017 A systematic review of primary outcomes and outcome measure reporting in randomized trials evaluating treatments for pre-eclampsia.

Pickles K, Eassey D, Reddel HK, Locock L, Kirkpatrick S, Smith L. "This illness diminishes me. What it does is like theft": A qualitative synthesis of people's experiences of living with asthma. *Health Expect*. 2017;00:1-18. <https://doi.org/10.1111/hex.12605>

Salisbury H., Dixon S., Papanikitas AN. 2017. Everyday clinical dilemmas. *InnovAiT: Education and inspiration for general practice*. 10(8)442-447. <https://doi.org/10.1177%2F1755738017710963>

Howick, J., Rees, S. 2017. Overthrowing barriers to empathy in healthcare: empathy in the age of the Internet . Journal of the Royal Society of Medicine, 0(0)1-6 DOI: 10.1177/0141076817714443

Kearney A., Williamson P., Young B., Bagley H., Gamble C., Denegri S., Muir D., Simon NA., Thomas S., Elliot JT., Bulbeck H., Crocker JC., Planner C., Vale C., Clarke M., Sprosen T., Woolfall K. 2017. Priorities for methodological research on patient and public involvement in clinical trials: A modified Delphi process. Health Expectations. 20(16)1401-1410 <https://doi.org/10.1111/hex.12583>

Albasri, A., O'Sullivan, J.O., Roberts, N.R., Prinjha, S.P., McManus, R.M., Sheppard, J.S. 2017. A comparison of blood pressure in community pharmacies with ambulatory, home and general practitioner office readings: systematic review and meta-analysis. Journal of Hypertension. DOI: 10.1097/HJH.0000000000001443

Parkin, S. (2017) Synthesising Qualitative Research Methods (Observation and Participation) to Provide Deeper Understandings of Substance Use: A Commentary. J Addict Res Ther 8:327. doi:10.4172/2155-6105.1000327

Farmer A, Williams V, Velardo C, Shah SA, Yu LM, Rutter H, Jones L, Williams N, Heneghan C, Price J, Hardinge M, Tarassenko L. 2017. Self-Management Support Using a Digital Health System Compared With Usual Care for Chronic Obstructive Pulmonary Disease: Randomized Controlled Trial. J Med Internet Res 2017;19(5):e144. DOI: 10.2196/jmir.7116 PMID: 28468749

Swan J, Gkeredakis E, Manning R, Nicolini D, Sharp D, Powell J. 2017. Improving the capabilities of NHS organisations to use evidence: a qualitative study of redesign projects in clinical commissioning groups. Health Services and Delivery Research 2018;5(18). DOI: 10.3310/hsdr05180

Malouf, R., McLeish, J., Ryan, S., Gray, R., Redshaw, M. 2017. We both just wanted to be normal parents': a qualitative study of the experience of maternity care for women with learning disability. BMJ Open, 7(3) <http://dx.doi.org/10.1136/bmjopen-2016-015526>

Duffy, J.M.N., Hirsch, M., Kawsar, A., Gale, C., Pealing, L., Plana, M.N., Showell, M., Williamson, P.R., Khan, K.S., Ziebland, S., McManus, R.J. 2017. Outcome reporting across randomised controlled trials evaluating therapeutic interventions for pre-eclampsia: a systematic review. BJOG DOI 10.1111/1471-0528.14702

Duffy, J.M.N., Rolph, R., Gale, C., Hirsch, M., Khan, K.S., Ziebland, S., McManus, R.J. Core Outcome Sets in Women's and Newborn Health: A Systematic Review.

Velthoven MVV, Powell J. Do health apps need endorsement? Challenges for giving advice about which health apps are safe and effective to use. Digital Health, 2017. 10 April 2017. doi: 10.1177/2055207617701342

Powell J, Atherton H, Williams V, Martin A, Bennett K, Bennett, A, Mollison J, Yu L-M, Yang Y, Locock L, Davoudianfar M, Griffiths KM. Effectiveness and cost-effectiveness of a fully self-guided internet-based intervention for sub-clinical social anxiety symptoms: study protocol for a randomised controlled trial. Digital Health, 2017. 10 April 2017. doi:10.1177/2055207617702272

Knight, F., Ridge, D., McShane, R., Ryan, S., Griffith, L. 2017. Care, Control, and the Electroconvulsive Therapy Ritual: Making Sense of Polarized Patient Narratives. Qualitative Health Research.

Coxon K., Chisholm A., Malouf R., Rowe R., Hollowell J. 2017. What influences birth place preferences, choices and decision-making amongst healthy women with straightforward pregnancies in the UK? A qualitative evidence synthesis using a 'best fit' framework approach. BMC Pregnancy and Childbirth. 17:103. <https://doi.org/10.1186/s12884-017-1279-7>

Duffy JMN., Hirsch M., Gale C., Pealing L., Kawsar A., Showell M., Williamson P., Khan K., Ziebland S., McManus RJ. 2017. Primary outcome and outcome measure reporting in pre-eclampsia trials. A systematic review of 79 randomised trials reporting data from 31 615 participants. *BJOG* 124. 122-123

Duffy JMN., Hirsch M., Kawsar A., Pealing L., Showell M., Williamson P., Khan K., Ziebland S., McManus RJ. 2017. Completeness of safety reporting in 79 randomised trials, 31 615 participants, evaluating therapeutic interventions for pre-eclampsia: A systematic review. *BJOG*. 124, 37-37

Duffy, J. M. N., Hirsch, M., Kawsar, A., Gale, C., Pealing, L., Plana, M., Showell, M., Williamson, P., Khan, K., Ziebland, S., McManus, R.J. 2017. Completeness of outcome reporting across 79 randomised trials, 31 615 participants, evaluating therapeutic interventions. for pre-eclampsia: a systematic review. *BJOG* 124, 123-123.

Duffy JMN., van't Hooft J., Gale C., Brown M., Grobman W., Fitzpatrick R., Karumanchi SA., Lucas N., Magee L., Mol B., Stark M., Thangaratnam S., Wilson M., von Dadelszen P., Williamson P., Khan K., Ziebland S., McManus RJ. 2017. Developing a pre-eclampsia core outcome set. Round 1 results: 283 healthcare professionals, 41 researchers and 112 patients from 55 countries participated. *BJOG*. 124, 151-152.

Ovseiko, P.V., Chapple, A., Edmunds, L.D. & Ziebland, S. 2017. Advancing gender equality through the Athena SWAN Charter for Women in Science: an exploratory study of women's and men's perceptions. *Health Research Policy and Systems*. 2017(15)12 DOI: 10.1186/s12961-017-0177-9

Ryan, S., Hislop, J., Ziebland, S. 2017 Do we all agree what "good health care" looks like? Views from those who are "seldom heard" in health research, policy and service improvement. *Health Expectations*. DOI: 10.1111/hex.12528

Holt, T.A., Dalton, A., Marshall, T., Fay, M., Qureshi, N., Kirkpatrick, S., Hislop, J., Lasserson, D., Kearley, K., Mollison, J., Yu, L.M., Hobbs, F.D.R., and Fitzmaurice, D. 2017. Automated Software System to Promote Anticoagulation and Reduce Stroke Risk. *Stroke*. 2017(48)787-790
<https://doi.org/10.1161/STROKEAHA.116.015468>

Hartmann-Boyce, J., Boylan, A.M., Jebb, S.A., Fletcher, B., Aveyard, P. 2017. Cognitive and behavioural strategies for self-directed weight loss: systematic review of qualitative studies. *Obesity Reviews*. doi: 10.1111/obr.12500

Stepney, M. 2017 Psychoanalysis/Psychoanalytic Geography, in, *The International Encyclopedia of Geography: People, the Earth, Environment and Technology*. Book Chapter.

Newhouse, N., Martin, A., Jawad, S., Yu, L.M., Davoudianfar, M., Locock, L., Ziebland, S., Powell, J. 2016. Randomised feasibility study of a novel experience-based internet intervention to support self-management in chronic asthma. *BMJ Open*. 6(12) <http://dx.doi.org/10.1136/bmjopen-2016-013401>

Stewart A, Hughes ND, Simkin S, Locock L, Ferrey AE, Kapur N, Gunnell D, Hawton K. (2016) Navigating an unfamiliar world: how parents of young people who self-harm experience support and treatment. *Child and Adolescent Mental Health*. doi:10.1111/camh.12205

Ziebland, S., Powell, J., Jenkinson, Perera, R., Martin, A., Locock, L., Kelly, L., Newhouse, N., Farmer, A. 2016 Examining the role of patients experiences as a resource for choice and decision-making in health care: a creative, interdisciplinary mixed-method study in digital health. 2016. Programme Grants for Applied Research. 4 (17) DOI 10.3310/pgfar04170

Ann H. Kelly, Hayley MacGregor and Catherine M. Montgomery. The publics of Public Health in Africa. *Critical Public Health*

Helen Atherton and Sue Ziebland. (2016) What do we need to consider when planning, implementing and researching the use of alternatives to face-to-face consultations in primary healthcare? *Digital Health* Vol 2: 1-13 DOI 10.1177/2055207616675559 dhj.sagepub.com.

Locock, L., Boylan, A-M., Snow, R., Staniszewska. 2016. The power of symbolic capital in patient and public involvement in health research. *Health Expectations*

Catherine M. Montgomery (2016): From Standardization to Adaptation: Clinical Trials and the Moral Economy of Anticipation, *Science as Culture*, DOI:10.1080/09505431.2016.1255721

Powell J, Newhouse N, Martin A, Jawad S, Yu LM, Davoudianfar M, Locock L, Ziebland S. A novel experience-based internet intervention for smoking cessation: feasibility randomised controlled trial. *BMC Public Health* 16(1):1156

Jones, C.H., Glogowska, M., Locock, L., Lasserson, D.S. 2016 Embedding new technologies in practice - a normalization process theory study of point of care testing. *Bio Med Central Health Serv Res* DOI: 10.1186/s12913-016-1834-3

Kidd J, Ziebland S. Narratives of experience of mental health and illness on healthtalk.org. *BJPsych Bull* 40(5):273-276 Oct 2016

Powell, J., Newhouse, N., Boylan, A.M., Williams, V. 2016. Digital health citizens and the future of the NHS. *Digital Health* DOI: 10.1177/2055207616672033

Beecham, E., Oostendorp, L., Crocker, J., Kelly, P., Dinsdale, A., Hemsley, J., Russell, J., Jones, L. and Bluebond-Langner, M. 2016. Keeping all options open: Parents' approaches to advance care planning. *Health Expect.* doi:10.1111/hex.12500

Snow, R., (2016) Do patients think you are learning the right things? *Oxford Medical Gazette* <http://www.omsg-online.com/do-patients-think-youre-learning-the-right-things/>

Deetjen, U., Powell, J.A. 2017. Internet use and health: Connecting secondary data through spatial microsimulation. *Digital Health*.

Deetjen, U., Powell, J. 2016. Connecting secondary data through spatial microsimulation. *Digital Health* 2 2055207616666588; DOI: 10.1177/2055207616666588

Snow, R., Crocker, J., Talbot, K., Moore, J., Salisbury, H. (2016). Does hearing the patient perspective improve consultation skills? *Medical Teacher* [1-7]

Lim, M.S.M., Bowden-Jones, H., Salinas, M., Price, J., Goodwin, G.M., Geddes, J. & Rogers, R.D. 2016. The experience of gambling problems in British professional footballers: a preliminary qualitative study. *Addiction Research and Theory*. <http://dx.doi.org/10.1080/16066359.2016.1212338>

Snow, R., (2016) What to call junior doctors <http://blogs.bmj.com/bmj/2016/08/12/rosamund-snow-what-to-call-junior-doctors-a-patients-perspective/>

Montgomery, C.M., 2016. Adaptive trials for tuberculosis: early reflections on theory and practice. *The International Journal of Tuberculosis and Lung Disease* Volume 20, Number 8, 1 August 2016, pp. 1091-1098(8) DOI: <http://dx.doi.org/10.5588/ijtld.16.0166>

Broom, A., Kirby, E., Kenny, K.E., MacArtney, J. I., & Good, P. 2016. Moral ambivalence and the valorisation of informal care. *The Sociological Review*. DOI: 10.1111/1467-954X.12400

- Devi, R., Singh, S., Powell, J., Fulton, E., Rees, K. 2016. Are there benefits of using web-based interventions in the secondary prevention of coronary heart disease? *Heart* 0:1–2. doi:10.1136/heartjnl-2015-309138
- Montgomery, C.M., Pool, R. 2016. From 'trial community' to 'experimental publics': how clinical research shapes public participation. *Critical Public Health* DOI:10.1080/09581596.2016.1212161
- Snow, R., (2016) What makes a real patient? <http://blogs.bmj.com/bmj/2016/07/19/rosamund-snow-what-makes-a-real-patient/>
- Williams, V., Kinnear, D., Victor, C. 2016 'The little things count': Delivering dignified care. *Nursing Times* 112: 27/28, 28-29.
- Parkin, S. 2016. Observant participation with people who inject drugs in street-based settings: reflections on a method used during applied ethnographic research *Addiction Research & Theory* DOI: 10.1080/16066359.2016.1196675
- Ferrey, A.E., Hughes, N.D., Simkin, S., Locock, L., Stewart, A., Kapur, N., Gunnell, D., Hawton, K. 2016. Changes in parenting strategies after a young person's self-harm: a qualitative study. *Child and Adolescent Psychiatry and Mental Health* DOI: 10.1186/s13034-016-0110-y
- Knight, M., Acosta, C., Brocklehurst, P., Cheshire, A., Fitzpatrick, K., Hinton, L., Jokinen, M., Kemp, B., Kurinczuk, J.J., Lewis, G., Lindquist, A., Locock, L., Nair, M., Patel, N., Quigley, M., Ridge, D., Sellars, S., Shah, A. 2016. Beyond maternal death: improving the quality of maternal care through national studies of 'near-miss' maternal morbidity. *NIHR Journals Library, PGfAR, Vol 4 Issue 9* DOI: 10.3310/pgfar04090
- Crocker, J.C., Boylan, A.M., Bostock, J., 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
- McQuaid, F., Pask, S., Locock, L., Davis, E., Stevens, Z., Plumb, J., and Snape, M.D. 2016. Attitudes towards antenatal vaccination, Group B streptococcus and participation in clinical trials: Insights from focus groups and interviews of parents and healthcare professionals. *Vaccine* 34(34):4056-61. pii: S0264-410X(16)30436-4. doi: 10.1016/j.vaccine.2016.06.024
- Ashdown, H., Räisänen, U., Wang, K., Ziebland, S., Harnden, A. 2016. Prescribing antibiotics to 'at-risk' children with influenza-like illness in primary care: qualitative study. *BMJ Open* 2016; 6:e011497 doi: 10.1136/bmjopen-2016-011497
- Searle, A., Ryan, S., Waylen, A. 2016. Health Professional communication in the diagnosis and care of infants born with cleft lip and palate in the U.K. *Journal of Neonatal Nursing*. doi:10.1016/j.jnn.2016.04.005
- Richardson, J., Liddle, J., Mallen, C., Roddy, E., Hider, S., Prinjha, S., Ziebland, S. 2016. A joint effort over a period of time: factors affecting use of urate-lowering therapy for long-term treatment of gout. *BMC Musculoskeletal Disorders* 17:249 DOI: 10.1186/s12891-016-1117-5
- Prinjha, S., Chapple, A., Feneley, R., Mangnall, J. 2016 Exploring the information needs of people living with a long-term indwelling urinary catheter: a qualitative study. *Journal of Advanced Nursing* 72(6):1335-46. doi: 10.1111/jan.12923
- Brant, H., Atherton, H., Ziebland, S., McKinstry, B., Campbell, J.L., Salisbury, C. 2016. Using alternatives to face-to-face consultations: a survey of prevalence and attitudes in general practice. *British Journal of General Practice Online First* DOI: 10.3399/bjgp16X685597

Campbell, B., Tabiri-Essuman, J., Gallo, H., Verdiel, V., Mandava, L., Azhar, MA., Powell, J. 2016. Public consultation changes guidance on the use of health-care interventions. An observational study. *Health Expectations* DOI: 10.1111/hex.12476

Snow, R., Gilbert, D. 2016 Why some patients are keeping their heads down. *British Medical Journal*. 353. doi: <http://dx.doi.org/10.1136/bmj.i2458>

Duffy, J.M.N., van 't Hooft, J., Gale, C., Brown, M., Grobman, W., Fitzpatrick, R., Ananth Karumanchi, S., Lucas, N., Magee, L., Mol, B., Stark, M., Thangaratnam, S., Wilson, M., von Dadelszen, P., Williamson, P., Khan, K.S., Ziebland, S., McManus RJ. 2016. A protocol for developing, disseminating, and implementing a core outcome set for pre-eclampsia. *Pregnancy Hypertension* DOI: <http://dx.doi.org/10.1016/j.preghy.2016.04.008>

Darbyshire, J., Grieg, P., Vollam, S., Young, D., Hinton, L. 2016 "I can remember sort of vivid people...but to me they were plasticine." Delusions on the intensive care unit: what do patients think is going on? *PLOS One* <http://dx.doi.org/10.1371/journal.pone.0153775>

Boaz, A., Robert, G., Locock, M.L., Sturmey, G., Gager, M., Vougioukalou, S., Ziebland, S., Fielden, J. 2016 What patients do and their impact on implementation process and outcomes during (and after) participatory quality improvement projects in English acute hospitals: Reflections from an ethnographic study. *Journal of Health Organization and Management*. 30 (2), [258-278](https://doi.org/10.1108/JHOM-02-2015-0027).
doi.org/10.1108/JHOM-02-2015-0027

Hinton, L. 2016. The intensive care unit was so noisy I couldn't sleep. *BMJ* 353 doi: <http://dx.doi.org/10.1136/bmj.i2150>

Shah, A., Kemp, B., Sellers, S., Hinton, L., Brocklehurst, P., Kurinczuk, J.J., Knight, M., O'Connor, M. 2016 Towards optimising local reviews of severe incidents in maternity care: messages from a comparison of local and external reviews. *BMJ Qual Saf*. pii: bmjqs-2015-004960. doi: 10.1136/bmjqs-2015-004960. [Epub ahead of print]

Oyebode, O., Patrick, H., Walker, A., Campbell, B., Powell, J. 2016. The Ghost in the machine? The value of expert advice in the production of evidence-based guidance: A mixed-methods study of the NICE Interventional Procedures Programme. *International Journal of Technology Assessment in Health Care*

Kelly, A., 2016 Growing up in care. *BMJ* 2016; 352 doi: <http://dx.doi.org/10.1136/bmj.i1085>

Locock, N., Nettleton, S., Kirkpatrick, S., Ryan, S., Ziebland, S., Locock, M.L. 2016. 'I knew before I was told': Breaches, cues and clues in the diagnostic assemblage. *Soc Sci Med*. 154:85-92. doi: 10.1016/j.socscimed.2016.02.037. [Epub ahead of print]

Watson, J., Salisbury, C., Atherton, H., Campbell, J., McKinstry, B., Ziebland, S. 2016. Proliferation of private online healthcare companies. *BMJ*. 23;352:i1076. doi: 10.1136/bmj.i1076.

Deetjen U., Powell JA. 2016. Informational and emotional elements in online support groups: a Bayesian approach to large-scale content analysis. *J Am Med Inform Assoc*. 2016 Feb 15. pii: ocv190. doi: 10.1093/jamia/ocv190. [Epub ahead of print]

Greenhalgh T., Annandale E., Ashcroft R., Barlow J., Black N., Bleakley A., Boaden R., Braithwaite J., Britten N., Carnevale F., Checkland K., Cheek J., Clark A., Cohn S., Coulehan J., Crabtree B., Cummins S., Davidoff F., Davies H., Dingwall R., Dixon-Woods M., Elwyn G., Engebretsen E., Ferlie E., Fulop N., Gabbay J., Gagnon MP., Galasinski D., Garside R., Gilson L., Griffiths P., Hawe P., Helderma JK., Hodges B., Hunter D., Kearney M., Kitzinger C., Kitzinger J., Kuper A., Kushner S.,

May AL., Legare F., Lingard L., Locock L., Ziebland S. 2016. An open letter to The BMJ editors on qualitative research. *BMJ*. ;352:i563. doi: 10.1136/bmj.i563.

Luchtenberg M., Maeckelberghe E., Locock L., Powell L., Verhagen AA. 2016 A Response to the Open Peer Commentaries on "Young People's Experiences of Participation in Clinical Trials: Reasons for Taking Part". *Am J Bioeth*. 16(2):W10-2. doi: 10.1080/15265161.2015.1125968

Snow, R., (2016) A year of patients' thoughts. <http://blogs.bmj.com/bmj/2016/01/18/rosamund-snow-a-year-of-patients-thoughts/>

Ferrey, A.E., Hughes, N.D., Simkin, S., Locock, L., Stewart, A., Kapur, N., Gunnell, D., Hawton, K. 2016. The impact of self-harm by young people on parents and families: a qualitative study. *BMJ Open* 6:e009631 doi:10.1136/bmjopen-2015-009631

Williams, V., Kinnear, D., Victor, C. 2016. 'It's the little things that count': Healthcare professionals' views on delivering dignified care: A qualitative study. *Journal of Advanced Nursing*. 72(4), 782–790 DOI: 10.1111/jan.12878

Snow, R., (2016) I never asked to be ICE'd. *British Medical Journal*, 354

Greig, P., Snow, R. (in review). The culture of risk: are train drivers safer than doctors? Analysis, *British Medical Journal*

Snow, R. (in press). Using one's own experience and body for research alongside others' narratives: challenges, advantages, and learning points. *Sage Research Methods Cases*. London: Sage.

Snow, R. (in press). The Patient Voice. Chapter in: Papanikitas, A., Spicer, J. (eds.) *Handbook of Primary Care Ethics*, FL: CRC Press

Fletcher, B.R., Hinton, L., Bray, E.P., Hayen, A., Hobbs, F.D.R., Mant, J., Potter, J.F., McManus, R.J. 2016. Self-monitoring blood pressure in patients with hypertension: an internet-based survey of UK GP. *British Journal of General Practice* DOI: 10.3399/bjgp16X687037

Newhouse, N. 2016. Bump2bump: Online peer support in first-time pregnancy. ORA, (2016) CHI EA '16 Proceedings of the 2016 CHI Conference Extended Abstracts on Human Factors in Computing Systems

McNiven, A., 2016. Pregnancy Loss, Skin-based Body Modifications and Agency.

Simmonds, R., Evans, J., Feder, G., Blakeman, T., Lasserson, D., Murray, E., Bennert, K., Locock, L., Horwood, J. 2016. Understanding tensions and identifying clinician agreement on improvements to early-stage chronic kidney disease monitoring in primary care: a qualitative study. *BMJ Open* 6:e010337. doi:10.1136/bmjopen-2015-010337.

Snow, R. 2016. What patients wish you knew. (In press) Opening chapter in: Firth, J., Cox, T., & Conlon, C. (eds.) *Oxford Textbook of Medicine*. OUP

Stepney, M., Stepney, P. 2016 Young women, alcohol and loss in the night-time economy. *Handbook of Traumatic Loss: A Guide to Theory and Practice*.

Chapple, A., Prinjha, S., Feneley, R., and Ziebland, S., 2015. Drawing on Accounts of Long-Term Urinary Catheter Use: Design for the "Seemingly Mundane" Sage Pub © The Author(s) 2015. DOI: 10.1177/1049732315570135

Anderson, C., Kirkpatrick, S., Ridge, D., Kokanovic, R., Tanner, C., 2015. Starting antidepressant use: a qualitative synthesis of UK and Australian data. *BMJ Open* 2015;5:12 e008636 doi:10.1136/bmjopen-2015-008636

Richardson, J., Liddle, J., Mallen, C., Roddy, E., Prinjha, S., Ziebland, S., Hider, S., 2015. "Why me? I don't fit the mould ... I am a freak of nature": a qualitative study of women's experience of gout. *BMC Women's Health*. 15:122 DOI: 10.1186/s12905-015-0277-z

Kinnear, D., Victor, C., Williams, V., 2015. What facilitates the delivery of dignified care to older people? A survey of health care professionals. *BMC Res Notes*. 28;8(1):826. doi: 10.1186/s13104-015-1801-9.

Montgomery, C.M. 2015. 'HIV has a Woman's Face': Vaginal Microbicides and a Case of Ambiguous Failure. *Anthropology and Medicine*. 22(3):250-62. doi: 10.1080/13648470.2015.1077200.

Devi, R., Singh, S.J., Powell, J., Fulton, E.A., Igbinedion, E., Rees, K. 2015. Internet-based interventions for the secondary prevention of coronary heart disease. *Cochrane Database Syst Rev*. 22;(12):CD009386. doi: 10.1002/14651858.CD009386.pub2.

Anderson, C., Kirkpatrick, S., 2015. Narrative interviewing. *International Journal of Clinical Pharmact*. June 2016. 38, Issue 3, pp 631–634 DOI: 10.1007/s11096-015-0222-0

Luchtenberg M., Maeckelberghe E., Locock L., Powell L., Verhagen A.A. 2015. Young People's Experiences of Participation in Clinical Trials: Reasons for Taking Part. *Am J Bioeth*. 15(11):3-13. doi: 10.1080/15265161.2015.1088974.

Harvey, J., Dopson, S., McManus, R., Powell, J. 2015. Factors influencing the adoption of self-management solutions: an interpretive synthesis of the literature on stakeholder experiences. *Implementation Science*. 10.159 doi: 10.1186/s13012-015-0350-x.

Powell, J., Boylan, A.M., Greaves, F. 2015. Harnessing patient feedback data: A challenge for policy and service improvement. *Digital Health* 0(0)1-3 DOI: 10.1177/20552076156

Cockbain, B.C., Thompson, S., Salisbury, H., Mitter, P., and Martos, L., 2015. A collaborative strategy to improve geriatric medical education. *Age and Ageing*. 44(6):1036-9 doi: 10.1093/ageing/afv100

Parkin, S. 2015. Salutogenesis: Contextualising place and space in the policies and politics of recovery from drug dependence (UK). *Int J Drug Policy*. 33:21-6. doi: 10.1016/j.drugpo.2015.10.002

Salisbury, H., and Dixon, S., 2015. Collecting the data on FGM will do more harm than good. *PULSE*.

MacArtney, J., 2015. Cancer self-health programmes: an ethos for negotiating multiplicities of healthcare. *Health Sociology Review*, DOI: 10.1080/14461242.2015.1082436

Pilcher, K.; Martin, W. and Williams, V. 2015 Issues of collaboration, representation, meaning and emotions: Utilising participant-led visual diaries to capture the everyday lives of people in mid to later life. *International Journal of Social Research Methodology*. 19(6)pp677-692
<http://dx.doi.org/10.1080/13645579.2015.1086199>

Liddle, J., Roddy, E., Mallen, C.D., Hider, S.L., Prinjha, S., Ziebland, S., Richardson, J.C., 2015. Mapping patients' experiences from initial symptoms to gout diagnosis: a qualitative exploration. *BMJ Open* 2015;5:e008323. doi:10.1136/bmjopen-2015-008323

- Parkin, S.G. 2015 Colliding intervention in the spatial management of street-based injecting and drug-related litter within settings of public convenience (UK). *Space and Polity, Special Issue (Drugs, Law, People, Place and the State)*. 20(1):1-20 DOI: 10.1080/13562576.2015.1077548
- Hinton, L., Locock, L., Knight, M., 2015. Support for mothers and their families after life-threatening illness in pregnancy and childbirth: a qualitative study in primary [care. BJGP.org](http://care.bjgp.org). DOI: 10.3399/bjgp15X686461
- Greenhalgh, T., Snow, R., Ryan, S., Rees, S., and Salisbury, H., 2015. Six 'biases' against patients and carers in evidence-based medicine. *BMC Medicine* September, 13:200 DOI:10.1186/s12916-015-0437-x
- Richards, T., Snow, R., & Schroter, S. (2015) Logging the BMJ's 'patient journey'. *British Medical Journal* ;351, h4396
- Snow, R., Crocker, J.C., Crowe., 2015. Missed opportunities for impact in patient and carer involvement: a mixed methods case study of research priority setting. *Research Involvement and Engagement*. 1 (7) DOI 10.1186/s40900-015-0007-6
- Fixsen, A., Ridge, D., Kirkpatrick, S., Foot, D., 2015. Interaction Rituals and Jumbled Emotions Among "Relative Strangers": Simulated Patient Work on a Trainee Complementary Therapy Practitioner Program. *SageOpen*. DOI: 10.1177/2158244015595090
- Heaton, J., Räisänen, U., and Salinas, M., 2015. 'Rule your condition, don't let it rule you': young adults' sense of mastery in their accounts of growing up with a chronic illness. *Sociology of Health & Illness*. doi: 10.1111/1467-9566.12298
- Rodas-Perez, C., Clarke, A., Powell, J., Thorogood, M. 2015. Challenges for providing genetic counselling in Colombian genetic clinics - the viewpoint of the physicians providing genetic consultations. *Journal of Community Genetics*. 6(3):301-11. DOI: 10.1007/s12687-015-0237-1
- Ferrey, A.E., Hawton, K., Simkin, S., Hughes, N., Stewart, A., Locock, L. 2015. "As a parent, there is no rulebook": A new resource for parents and carers of young people who self-harm. *The Lancet Psychiatry*. Volume 2, No.7, p577-579. DOI: [http://dx.doi.org/10.1016/S2215-0366\(15\)00182-0](http://dx.doi.org/10.1016/S2215-0366(15)00182-0)
- Kelly, L., Ziebland, S., Jenkinson, C., 2015. Measuring the effects of online health information: Scale validation for the e-Health Impact Questionnaire. *Patient Educ Couns*. pii: S0738-3991(15)00281-5. doi: 10.1016/j.pec.2015.06.008.
- Hardinge, M., Rutter, H., Velardo, C., Shah, S., Williams, V., Tarassenko, L. and Farmer, A. 2015. Using a mobile health application to support self- management in chronic obstructive pulmonary disease: a six-month cohort study. *BMC Medical Informatics and Decision Making*. DOI: 10.1186/s12911-015-0171-5
- Balatsoukas, P., Kennedy, C.M., Buchan, I., Powell, J., Ainsworth, J. 2015. The Role of Social Network Technologies in Online Health Promotion: a Narrative Review of Theoretical and Empirical Factors Influencing Intervention Effectiveness. *Journal of Medical Internet Research* 17(6):e141. doi: 10.2196/jmir.3662.
- Boaz, A., Locock, L., Ward, V., 2015. Whose evidence is it anyway? *Evidence & Policy: A Journal of Research, Debate and Practice*. 11(2), pp. 145-148(4)
- McNiven, A. 2015. 'Geographies of dying and death' in relation to pregnancy losses: ultrasonography experiences. *Social & Cultural Geography* 16(2)233-246

Anderson, C., Kirkpatrick, S., 2015. The missing piece in antidepressant treatment. *Chemist and Druggist*.

Chapple, A., Prinjha, S., & Feneley, R., 2014. Comparing Transurethral and Suprapubic Catheterization for Long-term Bladder Drainage. *Journal of Wound, Ostomy and Continence Nursing* 42(2)170-5 DOI: <http://doi.org/x57>

Newhouse, N., Lupiáñez-Villanueva, F., Codagnone, C., Atherton, H., 2015. Patient Use of Email for Health Care Communication Purposes Across 14 European Countries: An Analysis of Users According to Demographic and Health-Related Factors. *J Med Internet Res.* 17 (3): e58. (2015) DOI: 10.2196/jmir.3700

Coulter, A., Entwistle, V. A., Eccles, A., Ryan, S., Shepperd, S. and Perera, R., 2013. Personalised care planning for adults with chronic or long-term health conditions. *The Cochrane Library*, Issue 5. Art. No.: CD010523 <http://dx.doi.org/10.1002/14651858.CD010523>

Crocker, J.C., Beecham, E., Kelly, P., Dinsdale, A.P., Hemlsey, J., Jones, L., Bluebond-Langner, M., 2015. Inviting parents to take part in paediatric palliative care research: a mixed-methods examination of selection bias. *Palliative Medicine* 29(3):231-240.

Lowson, K., Jenks, M., Filby, A., Carr, L., Campbell, B., Powell, J. 2015. Examining the implementation of NICE Guidance: cross-sectional survey of the use of NICE interventional procedures guidance by NHS Trusts. *Implementation Science.* 10:93 DOI: 10.1186/s13012-015-0283-4

Chapple, A., Ziebland, S., Hawton, K., 2015. Taboo and the different death? Perceptions of those bereaved by suicide or other traumatic death. *Sociology of Health and Illness.* 37. 4 pp 610–625. DOI: 10.1111/1467-9566.12224 [46, 52]

Robert, G., Cornwell, J., Locock, L., Purushotham, A., Sturmey, G., Gager, M., 2015. Patients and staff as codesigners of healthcare services. *BMJ.* 350, g7714 doi:<http://dx.doi.org/10.1136/bmj.g7714>

Panesar, S., Diaper, M., Forsyth, D., Carson-Stevens, A., Ahluwalia, S., Rees, S., Marshall, M., Vincent, C., 2015. Don't let primary care patients slip through the nets. *HSJ* [online]

Ziebland, S., Lavie-Ajayi, M., and Lucius-Hoene, G., 2015. The role of the Internet for people with chronic pain: examples from the DIPEX International Project. *British Journal of Pain.* Vol. 9(1) 62–64. DOI:10.1177/2049463714555438

Locock, L., Boylan, A-M., 2015. Biosamples as gifts? How participants in biobanking projects talk about donation. *Health Expectations.* doi: 10.1111/hex.12376

Snow, R. (2015). Do you know what your patient is thinking? *British Medical Journal*, 350, h375.

Mike Bury: Biographical disruption and long-term and other health conditions Locock L., Ziebland S. Chapter in book. Palgrave Macmillan UK *The Palgrave Handbook of Social Theory in Health, Illness and Medicine*, Part IV, pp 582-598 ISBN 978-1-349-47022-8

Ridge, D., Kokanovic, R., Broom, A., Kirkpatrick, S., Anderson, C., Tanner, C., 2015. “My dirty little habit”: Patient constructions of antidepressant use and the ‘crisis’ of legitimacy. *Social Science & medicine.* doi:10.1016/j.socscimed.2015.10.012

Hughes, N.D., Locock, L., Simkin, S., Stewart, A., Ferrey, A.E., Gunnell, D., Kapur, N., Hawton, K. 2015. Making Sense of an Unknown Terrain: How Parents Understand Self-Harm in Young People. *Qual Health Res.* DOI: 10.1177/1049732315603032

Gonçalves-Bradley, D.C., Boylan, A.M., Koshiaris, C., Vazquez Montes, M., Ford, G.A., Lasserson, D.S., 2015. GPs' adherence to guidelines for structured assessments of stroke survivors in the community and care homes. *Fam Pract.* [Epub ahead of print] doi:10.1093/fampra/cmz074

Fletcher, B.R., Hinton, L., Hartmann-Boyce, J., Roberts, N.W., Bobrovitz, N., McManus, R.J., 2015. Self-monitoring blood pressure in hypertension, patient and provider perspectives; a systematic review and thematic synthesis. *Patient Education and Counselling*
<http://dx.doi.org/10.1016/j.pec.2015.08.026>

Hinton, L., Locock, L., Knight, M., 2015. Maternal critical care: what can we learn from patient experience? A qualitative study. *BMJ Open.* 2015;5:e006676 doi:10.1136/bmjopen-2014-006676

Locock, L., Ziebland, S., and Bury, M., 2015. 'Biographical disruption' and its application to long-term conditions and other aspects of health. In F Collyer (ed). *Handbook of Social Theory for Health and Medicine*. Basingstoke: Palgrave Macmillan

Crocker, J. C., Beecham, E., Kelly, P., Dinsdale, A. P., Hemsley, J., Jones, L., & Bluebond-Langner, M., 2015. Inviting parents to take part in paediatric palliative care research: A mixed-methods examination of selection bias. *Palliative Medicine*, 29(3), 231–240. doi:10.1177/0269216314560803

Ziebland, S., Ryan, S., 2015. On interviewing people with pets: reflections from qualitative research on people with long-term conditions. *Sociology of Health and Illness*. (Wiley Online Library) DOI: 10.1111/1467-9566.12176

Lezaun, J. & Montgomery, C.M. 2015 The Pharmaceutical Commons: Sharing & Exclusion in Global Health Drug Development. *Science, Technology & Human Values*. 40(1)3-29 DOI: 10.1177/0162243914542349

Ziebland, S., Chapple, A., & Evans, J., 2014. Barriers to shared decisions in the most serious of cancers: a qualitative study of patients with pancreatic cancer treated in the UK. *Health Expectations*. DOI: 10.1111/hex.12319

Kinnear, D., Williams, V., Victor, C. 2014. The meaning of dignified care: an exploration of health and social care professionals' perspectives. *BMC Research Notes*. 7.854 DOI: 10.1186/1756-0500-7854

Julie Evans, Helen Salisbury, sue Ziebland, Alison Chapple, Pippa Corrie. Intermittent gastrointestinal non-specific symptoms may also precede a pancreatic cancer diagnosis [letter/rapid response] *BMJ*. 20/11/2014 (<http://www.bmj.com/content/349/bmj.g6385/rapid-responses>)

Hodgkinson, J. A., Tucker, K. L., Crawford, C., Greenfield, Sheila M., Heneghan, C., Hinton, L., Khan, K., Locock, L., Mackillop, L., McCourt, C., Selwood, M., McManus, R.J., 2014. Is self monitoring of blood pressure in pregnancy safe and effective? *BMJ* 349: g6616.
<http://dx.doi.org/10.1136/bmj.g6616>

Ziebland, S., Locock, L., Fitzpatrick, R., Stokes, T., Robert, G., O'Flynn, N., Bennert, K., Ryan, S., Thomas, V., and Martin, A., 2014. Informing the development of NICE (National Institute for Health & Care Excellence) quality standards through secondary analysis of qualitative narrative interviews on patients experiences. *Health Serv Deliv Res* 2014; 2 (45) Doi: 10.3310/hsdr02450

Hinton, L., Locock, L., Knight, M., 2014. Experiences of quality care of women with near-miss maternal morbidities in the UK. *BJOG* 121 (Suppl. 4): 20-23 <http://dx.doi.org/10.1111/1471-0528.12800>

Devi, R., Powell, J., Singh, S. 2014. A Web-Based Program Improves Physical Activity Outcomes in a Primary Care Angina Population: Randomized Controlled Trial. *Journal of Medical Internet Research*. 16(9)e186

Delamothe, T., Snow, R., & Godlee, F. (2014) Why the Assisted Dying Bill should become law in England and Wales. *BMJ* 349: g4349.

Craig, J.A., Mahon, J., Yellowlees, A., Barata, T., Glanville, J., Arber, M., Mandava, L., Powell, J., Figueiredo, F. 2014. Epithelium off Photochemical Corneal Collagen Cross-Linkage Using Riboflavin and Ultraviolet A for Keratoconus and Keratectasia: A Systematic Review and Meta-Analysis. *The Ocular Surface*. 12(3)202-14 DOI:10.1016/j.jtos.2014.05.002

Ziebland, S. and Hunt, K., 2014. Using secondary analysis of qualitative data of patient experiences of health care to inform health services research and policy. *Journal of Health Services Research & Policy*, 19(3), pp.177–182. <http://dx.doi.org/10.1177/1355819614524187>

Taylor-Phillips, S., Clarke, A., Grove, A., Swan, J., Parsons, H., Gkeredakis, E., Mills, P., Powell, J., Nicolini, D., Roginski, C., Scarborough, H. 2014. Coproduction in commissioning decisions : is there an association with decision satisfaction for commissioners working in the NHS? A cross-sectional survey 2010/2011. *BMJ Open*. 4(6)4:e004810 doi:10.1136/bmjopen-2014-004810

Snow, R., Sandall, J., Humphrey, C., (2014). The use of clinical targets in diabetes patient education: analysis of the expectations and impact of a structured self-management programme in Type 1 diabetes, *Diabetic Medicine* 31:6

Locock, L., Robert, G., Boaz, A., Vougioukalou, S., Shuldham, C., Fielden, J., Ziebland, S., Gager, M., Tollyfield, R. and Pearcey, J., 2014. Using a national archive of patient experience narratives to promote local patient-centered quality improvement: an ethnographic process evaluation of ‘accelerated’ experience-based co-design. *Journal of health services research & policy*: 19: 200-7 <http://dx.doi.org/10.1177/1355819614531565>

O'Neill, B., Ziebland, S. and Valderas Francisco Lupiáñez-Villanueva, J., 2014. User-generated online health content: A survey of UK Internet users. *Journal of medical Internet research*, 16(4), e118. <http://dx.doi.org/10.2196/jmir.3187>

O'Neill, B., Ziebland, S., Valderas, J., Lupiáñez-Villanueva, F. 2014. User-generated online health content: a survey of Internet users in the United Kingdom. *Journal of medical Internet research*. 16(4):e118. doi: 10.2196/jmir.3187.

Hinton, L., Locock, L., and Knight, M., 2014. Partner Experiences of “Near-Miss” Events in Pregnancy and Childbirth in the UK: A Qualitative Study. *PloS one*, 9(4), e91735. <http://dx.doi.org/10.1371/journal.pone.0091735>

Poote, A.E., French, D.P., Dale, J., Powell, J. 2014. Automated self-assessment in a primary care student health centre setting: A cross-sectional study. *Journal of Telemedicine and Telecare*. 20(3)123-7 doi: 10.1177/1357633X14529246.

Prinjha S. & Chapple A. (2014) Penile sheaths and types of catheter. *Nursing and Residential Care*. Volume 16, No. 4

Soar, S., Ryan, S. and Salisbury, H., 2014. Using patients’ experiences in e-learning design. *The Clinical Teacher*, 11(2), pp.80–83. <http://dx.doi.org/10.1111/tct.12166>

Coulter, A., Locock, L., Ziebland, Z. and Calabrese, J., 2014. Collecting data on patient experience is not enough: they must be used to improve care. *BMJ*, 348:g2225

<http://dx.doi.org/10.1136/bmj.g2225>

Locock, L., Robert, G., Boaz, A., Vougioukalou, S., Shuldham, C., Fielden, J., Ziebland, S., Gager, M., Tollyfield, R. and Pearcey, J., 2014. Testing accelerated experience-based co-design: a qualitative study of using a national archive of patient experience narrative interviews to promote rapid patient-centred service improvement. *Health Serv Deliv Res.* 2014;2(4) DOI: 10.3310/hsdr02040

Evans, J., Chapple, A., Salisbury, H., Corrie, P. and Ziebland, S., 2014. "It can't be very important because it comes and goes"—patients' accounts of intermittent symptoms preceding a pancreatic cancer diagnosis: a qualitative study. *BMJ Open*, 4:e004215.

<http://dx.doi.org/10.1136/bmjopen-2013-004215>

Kuluski, K., Dow, C., Locock, L., Lyons, R. F. and Lasserson, D., 2014. Life interrupted and life regained? Coping with stroke at a young age. *International Journal of Qualitative Studies on Health and Well-being*, 9.

Laidlaw, A., Salisbury, H., Doherty, E. M. and Wiskin, C., 2014. National survey of clinical communication assessment in medical education in the United Kingdom (UK). *BMC Medical Education*, 14:10. <http://dx.doi.org/10.1186/1472-6920-14-10>.

Keltie, K., Cole, H., Arber, M., Patrick, H., Powell, J., Campbell, B., Sims, A., 2014. Identifying complications of interventional procedures from UK routine healthcare databases: a systematic search for methods using clinical codes. *BMC Med Res Methodol.* 14(1):126. doi: 10.1186/1471-2288-14-126

Locock, L., Robert, G., Boaz, A., Shuldham, C., Fielden, J., Ziebland, S., 2014. Testing accelerated experience-based co-design: using a national archive of patient experience narrative interviews to promote rapid patient-centred service improvement. In Keating, M.A., Montgomery, K., and McDermott, A, (eds) *Patient-centred healthcare: Achieving co-ordination, communication and innovation*. Palgrave Macmillan

Snow, R. (2014). Real patient participation in simulations. In: Nestel, D. & Bearman, M. (eds.) *Simulated Patient Methodology: Theory, Evidence and Practice*,. Oxford: Wiley Blackwell.

Snow, R., Sandall, J., Humphrey, C., (2014). Understanding the impact of patient education on the lives of people with diabetes, inside and outside the health system. *Diabetic Medicine special supplement* 31:1 98-99

Braden, O., Gonçalves, D., Ricci-Cabello, I., Ziebland, S., & Valderas, J., 2014. An Overview of Self-Administered Health Literacy Instruments. *PloS one*, 9(12), e109110. DOI: 10.1371/journal.pone.0109110

Crocker, J., Beecham, E., Kelly, P., Dinsdale, A., Hemsley, J., Jones, L., Bluebond-Langner, M. 2014. Recruiting parents to pediatric palliative Care Research: impact of Low invitation rate on sample bias.

Kelly, P., Crocker, J., Bluebond-Langner, M., Rajapaske, D., 2014. Unleashing the Potential of Routine Clinical Records as a Data Source: Use of Consensus Methods for Instrument Development in Investigating Breakthrough Pain in Children with Life-Limiting Conditions. *Journal of Palliative Care* 30, 195 – 196

Williams, V., Hardinge, M., Ryan, S., Farmer, A., 2014. Patients' experience of identifying and managing exacerbations in COPD: a qualitative study. *NPJ Primary Care Respiratory*

Medicine 24:14062 <http://dx.doi.org/10.1038/npjpcrm.2014.62>

Scarborough, H., D'Andreta, D., Evans, S., Marabelli, M., Newell, S., Powell, J., Swan, J., 2014. Networked innovation in the health sector: comparative qualitative study of the role of Collaborations for Leadership in Applied Health Research and Care in translating research into practice. *Health Services and Delivery Research*. 2(13). DOI: 10.3310/hsdr02130.

Nicolini, D., Powell, J., Korica, M., 2014. Keeping knowledgeable: how NHS chief executive officers mobilise knowledge and information in their daily work. *Health Services and Delivery Research*. 2(26) <http://dx.doi.org/10.3310/hsdr02260>

Devi, R., Carpenter, C., Powell, J., Singh, S., 2014. Exploring the experience of using a web-based cardiac rehabilitation programme in a primary care angina population: a qualitative study. *International Journal of Therapy and Rehabilitation*. 21(9):434-440.

Witty, K., Branney, P., Bullen, K., White, A., Evans, J., and Eardley, I., 2014. Engaging men with penile cancer in qualitative research: reflections from an interview-based study. *Nurse researcher*, 21(3), pp.13-19. <http://dx.doi.org/10.7748/nr2014.01.21.3.13.e1218>

Chapple, A., Prinjha, S. and Salisbury, H., 2014. How users of an indwelling urinary catheter talk about sex and sexuality: a qualitative study. *The British Journal of General Practice*. 64(623), e364-e371 <http://dx.doi.org/10.3399/bjgp14X680149>

Prinjha S. & Chapple A (2014) Living with a urinary catheter *International Continence News* Vol. 10 (1) page 25

Parkin, S.G. 2014. *An Applied Visual Sociology: Picturing Harm Reduction*. (Book) Ashgate Publications Ltd

Prinjha S & Chapple A. (2014) Patients' experiences of living with an indwelling urinary catheter *British Journal of Neuroscience Nursing* 10 (2): 62 .

Farmer, A., Toms, C., Hardinge, M., Williams, V., Rutter, H. and Tarassenko, L. 2014. Self-management support using an internet-linked tablet computer (the EDGE platform) based intervention in chronic obstructive pulmonary disease: protocol for the EDGE-COPD randomized controlled trial.

Chapple, A., Prinjha, S. and Mangnall, J., 2013. Changing a urethral or suprapubic catheter: The patient's perspective. *British Journal of Community Nursing*, 18(12), pp.591-6

Mazanderani, F., O'Neill, B. and Powell, J., 2013. "People power" or "pester power"? YouTube as a forum for the generation of evidence and patient advocacy. *Patient Education and Counseling* 93(3), pp.420–25. <http://dx.doi.org/10.1016/j.pec.2013.06.006>

Kelly, L., Jenkinson, C. and Ziebland, S., 2013. Measuring the effects of online health information for patients: Item generation for an e-health impact questionnaire. *Patient Education and Counseling*, 93(3), pp.433-38. <http://dx.doi.org/10.1016/j.pec.2013.03.012>

France, E. F., Hunt, K., Dow, C., and Wyke, S., 2013. Do Men's and Women's Accounts of Surviving a Stroke Conform to Frank's Narrative Genres? *Qualitative Health Research* 23(12) 1649-1659. <http://dx.doi.org/10.1177/1049732313509895>

Holt, T. A., Fitzmaurice, D. A., Marshall, T., Fay, M., Qureshi, N., Dalton, A. R. H., Hobbs, F. D. R., Lasserson, D. S., Kearley, K., Hislop, J. and Jin, J., 2013. AUtomed Risk Assessment for Stroke in Atrial Fibrillation (AURAS-AF) - an automated software system to promote anticoagulation and reduce stroke risk: study protocol for a cluster randomised controlled trial. *Trials*, 14(385).

<http://dx.doi.org/10.1186/1745-6215-14-385>

Hughes, N., Locock, L. and Ziebland, S., 2013. Personal identity and the role of 'carer' among relatives and friends of people with multiple sclerosis. *Social Science & Medicine*, 96, pp.78-85.

<http://dx.doi.org/10.1016/j.socscimed.2013.07.023>

Todkill, T., Powell, J. Participant experiences of an internet-based intervention and randomised control trial: interview study. *BMC Public Health* 2013;13:1017 doi:10.1186/1471-2458-13-1017

Raftery, J., Powell, J., 2013. Health Technology Assessment in the UK. *The Lancet*, 382(9900), pp.1278-85. [http://dx.doi.org/10.1016/S0140-6736\(13\)61724-9](http://dx.doi.org/10.1016/S0140-6736(13)61724-9)

Snow, R., Sandall, J., Humphrey, C., (2013). What happens when patients know more than their doctors? *BMJ Open* 3:11

Mavrodaris A, Powell J, Thorogood M. 2013. Prevalences of dementia and cognitive impairment among older people in sub-Saharan Africa: a systematic review. *Bulletin of the World Health Organization*. 2013;91(10):773-783. doi:10.2471/BLT.13.118422.

Witty, K., Branney, P., Evans, J., Bullen, K., White, A. and Eardley, I., 2013. The impact of surgical treatment for Penile Cancer – Patients' Perspectives. *European Journal of Oncology Nursing*, 17(5), pp.661–7. <http://dx.doi.org/10.1016/j.ejon.2013.06.004>

Kwon, J. H., Powell, J. and Chalmers, A., 2013. How Level of Realism Influences Anxiety in Virtual Reality Environments. *International Journal of Human Computer Studies*. 71(10), pp.978–987. <http://dx.doi.org/10.1016/j.ijhcs.2013.07.003>

Almond, S. C., Salisbury, H. and Ziebland, S., 2012. Women's experience of coronary heart disease: why is it different? *British Journal of Cardiac Nursing*, 7(4), pp.165-70.

Petit-Zeman, S. and Locock, L., 2013. Health care: Bring on the evidence. *Nature*, 501, pp.160-1. <http://dx.doi.org/1038/501160a>

Salisbury, H., A'Court, C., Jones, C., Fleming, S., Gonçalves, D. and Thompson, M., 2013. The RTI Clinical Iceberg. *The British Journal of General Practice*, 63(614), pp.461-462(2).

<http://dx.doi.org/10.3399/bjgp13X671524>

Hinton, L. and Miller, T., 2013. Mapping men's anticipations and experiences in the reproductive realm: (in)fertility journeys. *Reproductive BioMedicine Online*, 27(3), pp.244–52.

<http://dx.doi.org/10.1016/j.rbmo.2013.06.008>

Chapple, A., Ziebland, S., Simkin, S. and Hawton, K., 2013. How people bereaved by suicide perceive newspaper reporting: qualitative study. *The British Journal of Psychiatry*, 203(3), pp.228-32.

<http://dx.doi.org/10.1192/bjp.bp.112.114116>

Kirkpatrick, S., Locock, L., Giles, M. F. and Lasserson, D. S., 2013. Non-focal neurological symptoms associated with classical presentations of transient ischaemic attack: qualitative analysis of

interviews with patients. PLoS One, 8(6), e66351. PMID: 23776662
<http://dx.doi.org/10.1371/journal.pone.0066351>

Liew, S. M., Blacklock, C., Hislop, J., Glasziou, P. and Mant, D., 2013. Cardiovascular risk scores: qualitative study of how primary care practitioners understand and use them. The British Journal of General Practice, 63(611), e401-e407. <http://dx.doi.org/10.3399/bjgp13X668195>

Barber, V., Linsell, L., Locock, L., Juszczak, E., Powell, L., Shakeshaft, C., Colman, J., Lean, K. and Brocklehurst, P., 2013. Electronic fetal monitoring during labour and anxiety levels in women taking part in a RCT. British Journal of Midwifery, 21(6), pp.394-403

Clarke, A., Taylor-Phillips, S., Swan, J., Gkeredakis, E., Mills, P., Powell, J., Nicolini, D., Roginski, C., Scarbrough, H. and Grove, A., 2013. Evidence-based commissioning in the English NHS: who uses which sources of evidence? A survey 2010/2011. BMJ Open, 3(5), e002714 [article]
<http://dx.doi.org/10.1136/bmjopen-2013-002714>

Branney, P., Witty, K., Evans, J., Bullen, K., 2013. Helping men cope with penile cancer. BMJ (Clinical research ed.) 346:f2101 2013. [response to Arya, M., Kalsi, J., Kelly, J., Muneer, A., Malignant and premalignant lesions of the penis. BMJ2013;346:f1149. (6 March.)]

Mazanderani, F., Locock, L. and Powell, J., 2013. Biographical value: towards a conceptualisation of the commodification of illness narratives in contemporary healthcare. Sociology of Health & Illness, 35(6), pp.891–905 <http://dx.doi.org/10.1111/1467-9566.12001>

Prinjha S & Chapple A (2013) Living with an indwelling urinary catheter. Nursing Times Vol 109 No 44, 12-14

Ryan, S., 2013. Capturing diagnostic journeys of life on the autism spectrum, in J. Davidson and M. Orsini [Eds] Worlds of Autism; Across the Spectrum of Neurological Difference. University of Minnesota Press. (Book Chapter)

Ziebland, S., Coulter, A., Calabrese, J. and Locock, L., 2013. (Eds) Understanding and Using Health Experiences: improving patient care. Oxford: Oxford University Press

Ryan, S., 2013. Engagement and inclusivity in researching patients' experiences, Ziebland, S., Coulter, A., Calabrese, J. and Locock, L., (Eds) Understanding and Using Health Experiences: improving patient care, Oxford University Press

France, E. F., Wyke, S., Ziebland, S., Entwistle, V. A. and Hunt, K., 2013. What parents say about disclosing the end of their pregnancy due to fetal abnormality. Midwifery, 29(1), pp.24-32.
<http://dx.doi.org/10.1016/j.midw.2011.10.006> [25]

Powell, J., Hamborg, T., Stallard, N., Burls, A., McSorley, J., Bennett, K., Griffiths, K. M. and Christensen, H., 2013. Effectiveness of a Web-Based Cognitive-Behavioral Tool to Improve Mental Well-Being in the General Population: Randomized Controlled Trial. Journal of Medical Internet Research, 15(1), e2. <http://dx.doi.org/10.2196/jmir.2240>

Hall, S., Gray, N., Browne, S., Ziebland, S. and Campbell, N., 2012. A qualitative exploration of the role of primary care in supporting colorectal cancer patients. Supportive Care in Cancer, 20(12), pp.3071-8. <http://dx.doi.org/10.1007/s00520-012-1434-7>

O'Neill, B., Mazanderani, F. and Powell, J., 2012. People power or pester power? Multiple sclerosis, YouTube, and the liberation procedure. *The Lancet*, 380(S3), pp.S20.

[http://dx.doi.org/10.1016/S0140-6736\(13\)60376-1](http://dx.doi.org/10.1016/S0140-6736(13)60376-1)

Rowe, R., Kurinczuk, J., Locock, L. and Fitzpatrick, R., 2012. Women's experience of transfer from midwifery unit to hospital obstetric unit during labour. *BioMed Central Pregnancy and Childbirth*, 12(129). <http://dx.doi.org/10.1186/1471-2393-12-129>

Foy, R., Locock, L., Purdy, S., O'Donnell, C., Gray, N., Doran, T. and Davies, H., 2013. Research shapes policy: but the dynamics are subtle. *Public Money and Management*, 33(1), pp.9-14.

Kitson, A. L., Dow, C., Calabrese, J. D., Locock, L. and Athlin, A. M., 2013. Stroke survivors' experiences of the fundamentals of care: A qualitative analysis. *International Journal of Nursing Studies*, 50(3), pp.392-403. <http://dx.doi.org/10.1016/j.ijnurstu.2012.09.017>

Gill, P. J., Hislop, J., Mant, D. and Harnden, A., 2012. General practitioners' views on quality markers for children in UK primary care: a qualitative study. *BMC Family Practice*, 13(92).

<http://dx.doi.org/10.1186/1471-2296-13-92>

Ziebland, S. and Kokanovic, R., 2012. Emotions and chronic illness. *Chronic Illness*, 8(3), pp.159-162. DOI: 10.1177/1742395312451281

Dow, C. M., Roche, P. A. and Ziebland, S., 2012. Talk of frustration in the narratives of people with chronic pain. *Chronic Illness*, 8(3), pp.176-191. <http://dx.doi.org/10.1177/1742395312443692>

Ryan, S. and Raisanen, U., 2012. "The brain is such a delicate thing": an exploration of fear and seizures among young people with epilepsy. *Chronic Illness*, 8(3), pp.214-24.

<http://dx.doi.org/10.1177/1742395312449666>

Hinton, L., Kurinczuk, J. J. and Ziebland, S., 2012. Reassured or fobbed off? Perspectives on infertility consultations in primary care (a qualitative study). *The British Journal of General Practice* 62(599), pp.e438-e445(8). <http://dx.doi.org/10.3399/bjgp12X649133>

Ziebland, S. and Wyke, S., 2012. Health and illness in a connected world: how might sharing experiences on the internet affect people's health? *Milbank Quarterly*, 90(2), pp.219-249.

<http://dx.doi.org/10.1111/j.1468-0009.2012.00662.x>

Chapple, A., Ziebland, S., Hawton, K. 2012. A Proper, Fitting Explanation? Suicide Bereavement and Perceptions of the Coroner's Verdict. *Crisis*, 33(4), pp.230-38. <http://dx.doi.org/10.1027/0227-5910/a000139>

Ryan, S. and Salisbury, H., 2012. 'You know what boys are like': pre-diagnosis experiences of parents of children with autism spectrum conditions. *The British Journal of General Practice*, 62(598), pp.e378-e383(6) <http://dx.doi.org/10.3399/bjgp12X641500>

Ziebland, S., 2012. Why listening to health care users really matters. *Journal of Health Services Research & Policy*, 17(2), pp.68-9 <http://dx.doi.org/10.1258/jhsrp.2012.012025>

Locock, L., Mazanderani, F. and Powell, J., 2012. Metaphoric language and the articulation of emotions by people affected by Motor Neurone Disease. *Chronic Illness*, 8(3), pp.201-213.

<http://dx.doi.org/10.1177/1742395312443390>

- Chapple, A., Evans, J. and Ziebland, S., 2012. An Alarming Prognosis: How People Affected by Pancreatic Cancer Use (and Avoid) Internet Information, *Policy & Internet*, 4(2), pp.1-20. <http://dx.doi.org/10.1515/1944-2866.1162>
- Khan, N., Rose, P. W. and Evans, J., 2012. Defining cancer survivorship: a transparent approach is needed. *Journal of Cancer Survivorship*, 6(1), pp.33-36. <http://dx.doi.org/10.1007/s11764-011-0194-6>
- NF, Khan., Harrison, S., Rose, P. W., Ward, A. and Evans, J., 2012. Interpretation and acceptance of the term 'cancer survivor': a United Kingdom-based qualitative study. *European Journal of Cancer Care*, 21(2), pp.177-86. <http://dx.doi.org/10.1111/j.1365-2354.2011.01277.x>
- Mazanderani, F., Powell, J. and Locock, L., 2012. Being differently the same: an exploration of the mediation of identity tensions in the sharing of illness experience. *Social Science and Medicine*. 74(4), pp.546-53. <http://dx.doi.org/10.1016/j.socscimed.2011.10.036>
- France, E. F., Locock, L., Hunt, K., Ziebland, S., Field, K. and Wyke, S., 2012. Imagined futures: how experiential knowledge of disability affects parents decision making about fetal abnormality. *Health Expectations*, 15(2), pp.139-156. <http://dx.doi.org/10.1111/j.1369-7625.2011.00672>
- Venn, S. and Arber, S., 2012. Understanding older peoples' decisions about the use of sleeping medication: issues of control and autonomy. *Sociology of Health and Illness*, available online, DOI: 10.1111/j.1467-9566.2012.01468.x
- Evans, J., Ziebland, S. and Pettitt, A. R., 2012. Incurable, invisible and inconclusive: watchful waiting for chronic lymphocytic leukaemia and implications for doctor–patient communication. *European Journal of Cancer Care*, 21(1), pp.67-77. doi: 10.1111/j.1365-2354.2011.01278.
- Entwistle, V. A., France, E. F., Wyke, S., Jepson, R., Hunt, K., Ziebland, S. and Thompson, A., 2011. How information about other people's personal experiences can help with healthcare decision-making: A qualitative study. *Patient Education and Counseling*, 85(3), e291-e98, <http://dx.doi.org/10.1016/j.pec.2011.05.014>
- Browne, S., Dowie, A. L., Mitchell, L., Wyke, S., Ziebland, S., Campbell, N., and Macleod, U. 2011. Patients' needs following colorectal cancer diagnosis: where does primary care fit in?. *The British Journal of General Practice*, 61(592), e692-e699. <http://dx.doi.org/10.3399/bjgp11X606582>
- Khan, N. F., Evans, J. and Rose, P. W., 2011. A qualitative study of unmet needs and interactions with primary care among cancer survivors. *British Journal of Cancer*, 105, S46–S51. <http://dx.doi.org/10.1038/bjc.2011.422>
- Ridge, D. and Ziebland, S., 2012. Understanding depression through a 'coming out' framework. *Sociology of health & illness*, 34(5), pp.730-45. <http://dx.doi.org/10.1111/j.1467-9566.2011.01409.x>
- Locock, L. and Smith, L., 2011. Personal experiences of taking part in clinical trials – A qualitative study. *Patient Education and Counseling*, 84(3), pp.303-09. <http://dx.doi.org/10.1016/j.pec.2011.06.002> [49]
- Chapple, A., Evans, J., McPherson, A. and Payne, S., 2011. Patients with pancreatic cancer and relatives talk about preferred place of death and what influenced their preferences: a qualitative

study. *BMJ Supportive & Palliative Care*, 1(3), pp.291-295.

<http://dx.doi.org/10.1136/bmjspcare-2011-000091>

Ziebland, S., Evans, J., and Chapple, A., 2011. Qualitative studies of the benefits and adverse effects of the availability of cancer information and support on the internet. In Whitten, P., Kreps, G., & Eastin, M.S. (Eds.) *E-Health: The Advent of Online Cancer Information Systems*, Hampton Press: USA. June 2011.

Ryan, K., Todres, L. and Alexander, J., 2011. Calling, Permission, and Fulfillment: The Interembodied Experience of Breastfeeding. *Qualitative Health Research*, 21(6), pp.731-742

<http://dx.doi.org/10.1177/1049732310392591>

France, E. F., Wyke, S., Ziebland, S., Entwistle, V. A. and Hunt, A., 2011. How personal experiences feature in women's accounts of use of information for decisions about antenatal diagnostic testing for foetal abnormality. *Social Science & Medicine*, 72(5), pp.755-62.

<http://dx.doi.org/10.1016/j.socscimed.2010.11.031>

Chapple, A. and Ziebland, S., 2011. How the Internet is changing the experience of bereavement by suicide: A qualitative study in the UK. *Health*, 15(2), pp.173-87.

<http://dx.doi.org/10.1177/1363459309360792>

Ziebland, S., Evans, J. and Toynbee, P., 2011. Exceptionally good? Positive experiences of NHS care and treatment surprises lymphoma patients: a qualitative interview study. *Health Expectations*, 14(1), pp.21-8.<http://dx.doi.org/10.1111/j.1369-7625.2010.00609>

Locock, L. and Smith, L., 2011. Personal benefit, or benefitting others? Deciding whether to take part in clinical trials. *Clinical Trials*, 8(1), pp.85-93. [49]

<http://dx.doi.org/10.1177/1740774510392257>

Chapple, A., Swift, C. and Ziebland, S., 2011. The role of spirituality and religion for those bereaved due to a traumatic death. *Mortality*, 16(1), pp.1-19.

<http://dx.doi.org/10.1080/13576275.2011.535998>

Prinjha, S., Evans, J. and Ziebland, S., 2011. Women's distress about unexpected DCIS uncertainties and information provision (response to Pryke et al.). *Journal of Medical Screening*, 18(2), pp.105-6

<http://dx.doi.org/10.1258/jms.2011.011077>

Prinjha, S., Evans, J., Ziebland, S. and McPherson, A., 2011. A mastectomy for something that wasn't even truly invasive cancer': Women's understandings of having a mastectomy for screen-detected DCIS, a qualitative study. *Journal of Medical Screening*, 18(1), pp.34-40.

<http://dx.doi.org/10.1258/jms.2011.010143>

Hinton, L., Kurinczuk, J. and Ziebland, S., 2010. Infertility; isolation and the internet: a qualitative interview study. *Patient Education and Counseling*, 81(3), pp.436-41.

<http://dx.doi.org/10.1016/j.pec.2010.09.023>

Locock, L. and Brown, J. B., 2010. All in the same boat'? Patient and carer attitudes to peer support and social comparison in Motor Neurone Disease (MND). *Social Science & Medicine*, 71(8), pp.1498 – 1505 <http://dx.doi.org/10.1016/j.socscimed.2010.06.043>

Ryan, S., 2010. 'Meltdowns', surveillance and managing emotions; going out with children with autism. *Health & Place*, 16(5), pp.868 - 875 <http://dx.doi.org/10.1016/j.healthplace.2010.04.012>

Seale, C., Charteris-Black, J., Macfarlane, A. and McPherson, A., 2010. Interviews and Internet Forums: A Comparison of Two Sources of Qualitative Data. *Qualitative Health Research*, 20(5), pp.595-606. <http://dx.doi.org/10.1177/1049732309354094>

Chapple, A. and Ziebland, S., 2010. Viewing the body after bereavement due to a traumatic death: a qualitative study in the UK. *BMJ*, 340, c2032-c2032 <http://dx.doi.org/10.1136/bmj.c2032>

Ryan, K. M., Bissell, P. and Alexander, J., 2010. Moral work in women's narratives of breastfeeding. *Social Science & Medicine*, 70 (6), pp.591-598. <http://dx.doi.org/10.1016/j.socscimed.2009.11.023>

Locock, L., Field, K., McPherson, A. and Boyd, P., 2010. Women's accounts of the physical sensation of chorionic villus sampling (CVS) and amniocentesis: expectations and experience. *Midwifery*, 26(1), pp.64-75. <http://dx.doi.org/10.1016/j.midw.2008.02.002>

Locock, L., Ziebland, S. and Dumelow, C., 2009. Biographical disruption, abruption and repair in the context of Motor Neurone Disease. *Sociology of Health & Illness*, 31(7), pp.1043-1058. <http://dx.doi.org/10.1111/j.1467-9566.2009.01176.x>

Lowe, P., Powell, J., Thorogood, M. and Locock, L., 2009. 'Making it All Normal': The Role of the Internet in Problematic Pregnancy. *Qualitative Health Research*, 19(10), 1476 – 1484 <http://dx.doi.org/10.1177/1049732309348368>

Hunt, K., France, E., Wyke, S., Field, K. and Ziebland, S., 2009. My brain couldn't move from planning a birth to planning a funeral': a qualitative study of parents' experiences of decisions after ending a pregnancy for fetal abnormality. *International Journal of Nursing Studies*, 46(8), pp.1111-21. <http://dx.doi.org/10.1016/j.ijnurstu.2008.12.004>

Shariff, F., Carter, J., Dow, C., Polley, M., Salinas, M. and Ridge, D., 2009. Mind and Body Management Strategies for Chronic Pain and Rheumatoid Arthritis. *Qualitative Health Research*, 19(8), pp.1037-49. <http://dx.doi.org/10.1177/1049732309341189>

Hilton, S., Emslie, C., Hunt, K., Chapple, A. and Ziebland S., 2009. Disclosing a Cancer Diagnosis to Friends and Family: a gendered analysis of young men's and women's experiences. *Qualitative Health Research*, 19(6), pp.744-54. <http://dx.doi.org/10.1177/1049732309334737>

Prinjha, S., Field, K. and Rowan, K., 2009. What patients think about ICU follow-up services: a qualitative study. *Critical Care*, 13(2), R46 <http://dx.doi.org/10.1186/cc7769>

Emslie, C., Browne, S., McLeod, U., Rozmovits, L., Mitchell, E. and Ziebland S., 2009. 'Getting through' not 'going under'. A qualitative study of gender and spousal support after diagnosis with colorectal cancer. *Social Science & Medicine*, 68(6), pp.1169-75. <http://dx.doi.org/10.1016/j.socscimed.2009.01.004>

Forrest, G., Plumb, C., Ziebland, S. and Stein, A., 2009. Breast cancer in young families: a qualitative interview study of fathers and their role and communication with their children following the diagnosis of maternal breast cancer. *Psycho-Oncology*, 18(1), pp.96–103. <http://dx.doi.org/10.1002/pon.1387>

- Newman, M., Ziebland, S. and Barker, K., 2009. Patients views of a multimedia resource featuring experiences of rheumatoid arthritis: pilot evaluation of www.healthtalkonline.org. Health Informatics Journal. 15(2), pp.147-59. <http://dx.doi.org/10.1177/1460458209102974>
- Ryan, S. and Raisanen, U., 2009. 'It's like you are just a spectator in this thing': experiencing social life the 'aspie' way. Emotion, Space and Society, 1(2), pp.135-143
<http://dx.doi.org/10.1016/j.emospa.2009.02.001>
- Smith, S. M., Campbell, N. C., MacLeod, U., Lee, A. J., Raja, A., Wyke, S., Ziebland, S. B., Duff, E. M., Ritchie, L. D. and Nicolson, M. C., 2009. Factors contributing to the time taken to consult with symptoms of lung cancer: a cross-sectional study. Thorax, 64(6), pp.523-31.
<http://dx.doi.org/10.1136/thx.2008.096560>
- Ryan, S., Runswick Cole, K., 2009. From Advocate to Activist? Mapping Experiences of Mothers of Children On the Autism Spectrum. Journal of Applied Research in Intellectual Disabilities, 22(1), pp.43-53. <http://dx.doi.org/10.1111/j.1468-3148.2008.00438.x>
- Chapple, A., Ziebland, S., Hewitson, P. and McPherson, A., 2008. What affects the uptake of screening for bowel cancer using the faecal occult blood test (FOBT): a qualitative study. Social Science & Medicine, 66(12), pp.2425-2435. <http://dx.doi.org/10.1016/j.socscimed.2008.02.009>
- Locock, L., Alexander, J. and Rozmovits, L., 2008. Women's responses to nausea and vomiting in pregnancy. Midwifery, 24(2), pp.143-152. <http://dx.doi.org/10.1016/j.midw.2006.12.001>
- Hilton, S., Hunt, K., Emslie, C., Salinas, M. and Ziebland, S., 2008. Have men been overlooked? A comparison of young men and women's experiences of chemotherapy-induced alopecia. Psycho-Oncology, 17(6), pp.577-83. <http://dx.doi.org/10.1002/pon.1272>
- Ziebland, S. and Herxheimer, A., 2008. How patients' experiences contribute to decision making: illustrations from DIPEX (personal experiences of health and illness). Journal of Nursing Management, 16(4), pp.433-39. <http://dx.doi.org/10.1111/j.1365-2834.2008.00863.x>
- Seale, C., Charteris-Black, J., Dumelow, C., Locock, L. and Ziebland, S.,
Locock, L. and Kai, J.,
- Field, K., Prinjha, S. and Rowan, K., 2008. One patient amongst many: a qualitative analysis of Intensive Care Unit patients' experiences of transferring to the general ward. Critical Care, 12(1), R21 <http://dx.doi.org/10.1186/cc6795>
- Mildorf, J. 2008, Thought Presentation and Constructed Dialogue in Oral Stories: Limits and Possibilities of a Cross-Disciplinary Narratology. Partial Answers: Journal of Literature and the History of Ideas, 6(2), pp.279-300. <http://dx.doi.org/10.1353/pan.0.0025>
- Chapple, A., Ziebland, S., Hewitson, P. and McPherson, A., 2008. Why Men in the United Kingdom Still Want the Prostate Specific Antigen Test. Qualitative Health Research, 18(1), pp.56-64.
<http://dx.doi.org/10.1177/1049732307309000>
- Chapple, A., Ziebland, S., Brewster, S. and McPherson, A., 2007. Patients' perceptions of transrectal prostate biopsy: a qualitative study. European Journal of Cancer Care, 16(3), pp.215-21.
<http://dx.doi.org/10.1111/j.1365-2354.2006.00766.x>

Evans, J., Ziebland, S. and McPherson, A., 2007. Minimizing delays in ovarian cancer diagnosis: a reworking of Andersen's model of 'total patient delay'. *Family Practice*, 24(1), pp.48-55.
<http://dx.doi.org/10.1093/fampra/cml063>

Ziebland, S., Featherstone, K., Snowdon, C., Barker, K., Frost, H. and Fairbank, J., 2007. Does it matter if clinicians recruiting for a trial don't understand what the trial is really about? Qualitative study of surgeons' experiences of participation in a pragmatic multi-centre RCT. *Trials*, 8(4), pp.647-60.
<http://dx.doi.org/10.1186/1745-6215-8-4>

Ridge, D., Williams, I., Anderson, A. and Elford, J., 2007. Like a prayer: The role of spirituality and religion for people living with HIV in the UK. *Sociology of Health and Illness*, 30(3), pp.413-18. <http://dx.doi.org/10.1111/j.1467-9566.2007.01062.x>

Emslie, C., Ridge, D., Ziebland, S. and Hunt, K., 2007. Exploring men's and women's experiences of depression and engagement with health professionals: more similarities than differences? A qualitative interview study. *BMC Family Practice*, 8(1), p.43. <http://dx.doi.org/10.1186/1471-2296-8-43>

Ridge, D., Ziebland, S., Williams, I., Anderson, A. and Elford, J., 2007. Positive prevention: contemporary issues facing HIV positive people negotiating sex in the UK. *Social Science and Medicine*, 65(4), pp.755-70. <http://dx.doi.org/10.1016/j.socscimed.2007.03.049>

Ryan, K. and Herxheimer, A., 2007. Personal Experiences of health and illness on the web: a resource for patients, carers and health professionals. *Journal of the Malta College of Pharmacy Practice*, 13, pp.27-30.

Herxheimer, A., 2007. Online review - Learning from personal experiences of health and illness: www.dipex.org. Australian Health Consumer.

Chapple, A., Salinas, M., Ziebland, S., McPherson, A. and Macfarlane, A., 2007. Fertility Issues: The Perceptions and Experiences of Young Men Recently Diagnosed and Treated for Cancer. *Journal of Adolescent Health*, 40(1), pp.69-75 <http://dx.doi.org/10.1016/j.jadohealth.2006.07.010>

Chapple, A., Ziebland, S., Herxheimer, A. and McPherson, A., 2006. What people close to death say about euthanasia and assisted suicide: a qualitative study. *Journal of Medical Ethics*, 32(12), pp.706-10. <http://dx.doi.org/10.1136/jme.2006.015883>

Field, K., Ziebland, S., McPherson, A. and Lehman, R., 2006. Can I come off the tablets now? A qualitative analysis of heart failure patients' understanding of their medication. *Family Practice*, 23(6), pp.624-30. <http://dx.doi.org/10.1093/fampra/cml036>

Chapple, A., Ziebland, S. and McPherson, A., 2006. The specialist palliative care nurse: a qualitative study of the patients' perspective. *International Journal of Nursing Studies*, 43(8), pp.1011-22 <http://dx.doi.org/10.1016/j.ijnurstu.2005.11.007>

Ridge, D. T. and Ziebland, S., 2007. "The old me could never have done that': How people give meaning to recovery following depression'. *Qualitative Health Research*, 16(8), pp.1038-53. <http://dx.doi.org/10.1177/1049732306292132>

Chapple, A., 2006. Interviewing people with terminal illness: Practical and ethical issues. *Nurse Researcher*, 14(1), pp.50-58 <http://dx.doi.org/10.7748/nr2006.10.14.1.50.c6009>

- Prinjha, S., Evans, J. and McPherson, A., 2006. Women's information needs about ductal carcinoma in situ before mammographic screening and after diagnosis: a qualitative study. *Journal of Medical Screening*. 13(3), pp.110-114. <http://dx.doi.org/10.1258/096914106778440581>
- Ziebland, S., Evans, J., McPherson, A., 2006. The choice is yours? How women with ovarian cancer make sense of treatment choices. *Patient Education and Counseling*, 62(3), pp.361-67. <http://dx.doi.org/10.1016/j.pec.2006.06.014>
- Emslie, C., Ridge, D., Ziebland, S. and Hunt, K., 2006. Men's accounts of depression: reconstructing or resisting hegemonic masculinity? *Social Science & Medicine*, 62(9), pp.2246- 57 <http://dx.doi.org/10.1016/j.socscimed.2005.10.017>
- Ziebland, S. and McPherson, A., 2006. Making sense of qualitative data analysis: an introduction with illustrations from DIPEX (personal experiences of health and illness). *Medical Education*, 40(5), pp.405-14. <http://dx.doi.org/10.1111/j.1365-2929.2006.02467.x>
- Seale, C., Charteris-Black, J. and Ziebland, S., 2006. Gender, cancer experience and internet use: A comparative keyword analysis of interviews and online cancer support groups. *Social Science & Medicine*, 62(10), pp.2577-90 <http://dx.doi.org/10.1016/j.socscimed.2005.11.016>
- Forrest, G., Plumb, C., Ziebland, S. and Stein, A., 2006. Breast cancer in the family—children's perceptions of their mother's cancer and its initial treatment: qualitative study. *BMJ*, 332(7548), pp.998-1003 <http://dx.doi.org/10.1136/bmj.38793.567801.AE>
- Rose, P., Ziebland, S., Harnden, A., Mayon-White, R. and Mant, D., 2006. Why do General practitioners prescribe antibiotics for acute infective conjunctivitis in children? Qualitative interviews with GPs and a questionnaire survey of parents and teachers. *Family Practice*, 23(2), pp.226-32. <http://dx.doi.org/10.1093/fampra/cmi101>
- Locock, L. and Alexander, J., 2006. "'Just a bystander?'" Men's place in the process of fetal screening and diagnosis'. *Social Science and Medicine*, 62(6), pp.1349-59 [16] <http://dx.doi.org/10.1016/j.socscimed.2005.08.011>
- Herxheimer, A., 2006. Communicating patient-reported outcomes to the public: numbers don't work, patients' own words would. *PRO Newsletter*, 36 (Spring issue), pp.29-30.
- Alexander, J. and Ziebland, S., 2006. The web – bringing support and health information into the home: the communicative power of qualitative research. (Editorial). *International Journal of Nursing Studies*, 43(4), pp.389-391.
- Locock, L., Crawford, J. and Crawford, J., 2005. The Parents' Journey: continuing a pregnancy after a diagnosis of Patau's syndrome. *BMJ*, 331, pp.1186-9. <http://dx.doi.org/10.1136/bmj.331.7526.1186>
- Prinjha, S., Chapple, A., Herxheimer, A. and McPherson, A., 2005. Many people with epilepsy want to know more: a qualitative study. *Family Practice*, 22(4), pp.435-441 [8] <http://dx.doi.org/10.1093/fampra/cmi024>
- Fairhurst, K., Wyke, S., Seaman, P., Ziebland, S. and Glasier, A., 2005. "Not that sort of practice": The views and behaviour of primary care practitioners in a study of advance provision of emergency contraception. *Family Practice*, 22(3), pp.280-6. <http://dx.doi.org/10.1093/fampra/cmh728>

Ziebland, S., Wyke, S., Seaman, P., Fairhurst, K., Walker, J. and Glasier, A., 2005. What happened when Scottish women were given advanced supplies of emergency contraception? A survey and qualitative study of women's views and experiences. *Social Science and Medicine*, 60(8), pp.1767-79. <http://dx.doi.org/10.1016/j.socscimed.2004.08.022>

Botow, P., Fowler, J. and Ziebland, S., 2005. Using personal stories. In O'Connor AM, Llewelyn-Thomas H, Stacey D (Eds) IPDAS collaboration n background document 2005 http://ipdas.ohri.ca/IPDAS_background.pdf

Cheshire, J. and Ziebland, S., 2005. Narrative as a resource in accounts of the experience of illness. In: Jennifer Coates, Joanna Thornborrow (Eds), *The Sociolinguistics of Narrative* Amsterdam: John Benjamins, Ch2: 17 – 40.

McPherson, A., Prinjha, S. and Evans, J., 2005. A novel intervention aimed at increasing the emotional adjustment of breast cancer patients. In: Leonard R, Polychronis A, Miles A, eds. *The Effective Management of Breast Cancer*, Second edition. UK Key Advances in Clinical Practice Series. Aesculapius Medical Press, London, 283–293

Ziebland, S., 2004. The importance of being expert: the quest for cancer information on the Internet. *Social Science & Medicine*, 59(9), pp.1783-93. <http://dx.doi.org/10.1016/j.socscimed.2004.02.019>

Chapple, A. and Ziebland, S., 2004. The role of humor for men with testicular cancer. *Qualitative Health Research*. 14(8), pp.1123-39. <http://dx.doi.org/10.1177/1049732304267455>

Chapple, A. and McPherson, A., 2004. The decision to have a prosthesis: a qualitative study of men with testicular cancer. *Psycho-Oncology*, 13(9), pp.654-664. <http://dx.doi.org/10.1002/pon.787>

Chapple, A., Ziebland, S., McPherson, A. and Summerton, N., 2004. Lung cancer patients' perceptions of access to financial benefits: a qualitative study. *The British Journal General Practice*, 54(505), pp.589-94.

Rozmovits, L., Rose, P. and Ziebland, S., 2004. In the absence of evidence who chooses? A qualitative study of patients' needs after treatment for colorectal cancer. *Journal of Health Services Research & Policy*, 9(3), pp.159-64. <http://dx.doi.org/10.1258/1355819041403231>

Fairhurst, K., Ziebland, S., Wyke, D., Seaman, P. and Glasier, A., 2004. Emergency Contraception: why can't you give it away? Qualitative findings from an evaluation of advance provision of emergency contraception. *Contraception*, 70(1), pp.25-9. <http://dx.doi.org/10.1016/j.contraception.2004.02.012>

Chapple, A., Ziebland, S. and McPherson, A., 2004. Stigma, shame and blame experienced by patients with lung cancer: qualitative study. *BMJ*, 328(7454), 1470. DOI: 10.1136/bmj.38111.639734.7C

Baldwin, C., Hughes, J., Hope, T., Jacoby, R. and Ziebland, S., 2004. Ethics and dementia: the experience of family carers. *Progress in Neurology and Psychiatry*, 8(5), pp.24-8.

Glasier, A., Fairhurst, K., Wyke, S., Ziebland, S., Seaman, P., Walker, J. and Lakha, F., 2004. Advanced provision of emergency contraception does not reduce abortion rates. *Contraception*, 69(5), pp.361-66. <http://dx.doi.org/10.1016/j.contraception.2004.01.002>

Rozmovits, L. and Ziebland, S., 2004. What do patients with prostate or breast cancer want from an internet site? A qualitative study of information needs. *Patient Education and Counselling*, 53(1), pp.57-64 (corresponding author)

Ziebland, S., Chapple, A., Dumelow, C., Evans, J., Prinjha, S. and Rozmovits, L., 2004. How the Internet affects patients' experience of cancer. A qualitative study. *BMJ*, 328, 564. [1,3, 4,5,7] <http://dx.doi.org/10.1136/bmj.328.7439.564>

Rozmovits, L. and Ziebland, S., 2004. Expressions of loss of adulthood in the narratives of colorectal cancer patients. *Qualitative Health Research*, 14(2), pp.187-203 (corresponding author).

John, J. and Ziebland, S., 2004. Reported barriers to eating more fruit and vegetables before and after participation in a randomized controlled trial: a qualitative study. *Health Education Research*, 19(2), pp.165-74. <http://dx.doi.org/10.1093/her/cyg016>

Sillence, E., Briggs, P. and Herxheimer, A., 2004. Personal experiences matter: what patients think about hypertension information online. *He@lth Information on the Internet*, 42(1), pp.3-5 <http://www.hioti.org>

Herxheimer, A. and Ziebland, S., 2004. The DIPEX project: Collecting personal experiences of illness and health care: Hurwitz, B., Greenhalgh, T. and Skultans, V., eds. *Narrative Research in Health and Illness*. Blackwell, Ch6, pp.115 – 131.

Chapple, A., Ziebland, S. and McPherson, A., 2004. Qualitative study of men's perceptions of why treatment delays occur in the UK for those with testicular cancer. *The British Journal of General Practice*, 54(498), pp.25-32.

Herxheimer, A. and Ziebland, S., 2003. Illustrating Cochrane reviews with narrative clips describing patients' experiences of the interventions. *Cochrane Collaboration Methods Groups Newsletter*. 7, pp.5-6. (June)

John, J. H., Yudkin, P. L., Neil, H. A. W. and Ziebland, S., 2003. Does Stage of Change predict outcome in a primary-care intervention to encourage an increase in fruit and vegetable consumption? *Health Education Research*, 18(4), pp.429-38 (Corresponding author). <http://dx.doi.org/10.1093/her/cyf035>

Baldwin, C., Hughes, J., Hope, T., Jacoby, R. and Ziebland, S., 2003. Ethics and dementia: Mapping the literature by bibliometric analysis. *International Journal of Geriatric Psychiatry*, 18 (1), pp.41-54. <http://dx.doi.org/10.1002/gps.770>

Herxheimer, A., McPherson, A., Miller, R., Chapple, A., Shepperd, S. and Ziebland, S., [In Chinese *Journal of Evidence Based Medicine*](Chinese translation of updated version of ref 1)

Herxheimer, A., 2003. Clinical Trials: Finding and documenting the outcomes that matter most to patients. A proposal for a third dimension in trial design. *CERES News*, 33, pp.1-3.

Herxheimer, A., McPherson, A., Miller, R., Chapple, A., Shepperd, S., Ziebland, S. and Sanz, E., 2003. DIPEX (base de datos de experiencias de pacientes sobre su enfermedad): una propuesta multimedia para compartir experiencias e información sobre enfermedades entre pacientes y profesionales sanitarios. *Atención Primaria* 3:31 (6): 386-8. (Spanish translation of updated version of ref 1)

Herxheimer, A. and Ziebland, S., 2003. DIPEX: Fresh insights for medical practice. Editorial. Journal of the Royal Society of Medicine. 96(5), pp.209-10.

Moore, J., Ziebland, S. and Kennedy, S., 2002. 'People sometimes react funny if they're not told enough': Women's views about communicating the risks of diagnostic laparoscopy. Health Expectations, 5(4), pp.302-9. <http://dx.doi.org/10.1046/j.1369-6513.2002.00192.x>

Ziebland, S., Robertson, J., Jay, J. and Neil, A., 2002. Body image and weight change in middle age: a qualitative study. International Journal of Obesity, 26(8), pp.1083-91 (corresponding author). <http://dx.doi.org/10.1038/sj.ijo.0802049>

John, J., Ziebland, S., Yudkin, P., Roe, L. S. and Neil, H. A. W., 2002. Effects of fruit and vegetable consumption on plasma antioxidant concentrations and blood pressure: a randomised controlled trial. The Lancet, 359(9322), pp.1969-74. [http://dx.doi.org/10.1016/S0140-6736\(02\)98858-6](http://dx.doi.org/10.1016/S0140-6736(02)98858-6)

Edmunds, L. D. and Ziebland, S., 2002. Development and validation of the Day in the Life Questionnaire (DILQ) as a measure of fruit and vegetable consumption for 7-9 year olds. Health Education Research, 17(2), pp.211-20. <http://dx.doi.org/10.1093/her/17.2.211>

Hughes, J., Hope, T., Savulescu, J. and Ziebland, S., 2002. Carers Ethics and Dementia: a survey and review of the literature. International Journal of Geriatric Psychiatry, 17(1), pp.35-40. <http://dx.doi.org/10.1002/gps.515>

McPherson, A. and Herxheimer, A., 2002. Database of Individual Patients' Experiences – an update. The British Journal of General Practice, 52(954).

Chapple, A. and Ziebland, S., 2002. Prostate Cancer: Embodied experience and perceptions of masculinity. Sociology of Health and Illness, 24(6), pp.820-41.

Chapple, A., Ziebland, S., Shepperd, S., Miller, R., McPherson, A. and Herxheimer, A., 2002. Why men with prostate cancer want wider access to prostate specific antigen testing: qualitative study. BMJ, 325(7367), p.737.

Chapple, A., Ziebland, S., McPherson, A., Shepperd, S., Herxheimer, A. and Miller, R., 2002. Is 'watchful waiting' a real choice for men with prostate cancer? A qualitative study. The British Journal of Urology International, 90(3), pp.257-64. <http://dx.doi.org/10.1046/j.1464-410X.2002.02846.x>

Ziebland, S. and Fuller, A., 2001. Smoking cessation in pregnancy: what's a man to do? Health Education Journal, 60(3), pp.232-40 (corresponding author). <http://dx.doi.org/10.1177/001789690106000305>

Herxheimer, A., McPherson, A., Miller, R., Chapple, A., Shepperd, S., Kochen, M. M. and Ziebland, S., 2001. [DIPEX: A new approach in the UK - patients and health professionals are sharing experiences and information by using multimedia. The Informed Prescriber (Tokyo), 16 (9), pp.86-90. (Japanese translation of updated version of ref 5)

Herxheimer, A., McPherson, A., Miller, R., Shepperd, S. and Ziebland, S.

Herxheimer, A., McPherson, A., Miller, R., Chapple, A., Shepperd, S., Kochen, M. M. and Ziebland, S., 2001. A Database of Patients' Experiences (DIPEX): Patienten berichten von ihren

Krankheitserfahrungen. Über den Aufbau einer Datenbank im Internet. Z Allg Med, 77, pp.323-7.
(German translation of updated and abbreviated version of ref 1)

McPherson, A. and Herxheimer, A., 2001. DIPEX – A Database of Individual Patients' Experiences. The British Journal of General Practice, 51: 770.

Pope, C., Ziebland, S. and Mays, N., 2000. Qualitative research in health care: analysing qualitative data. BMJ, 320(7227), pp.114-6. <http://dx.doi.org/10.1136/bmj.320.7227.114>

Ziebland, S., 2000. DIPEX: a new resource linking narratives with evidence based information. Medical Sociology News, 26(3), pp.29-31.

Yaphe, J., Rigge, M., Herxheimer, A., McPherson, A., Miller, R., Shepperd, S. and Ziebland, S., 2000. The use of patients' stories by self-help groups: A survey of voluntary organizations in the UK on the register of the College of Health. Health Expectations, 3(3), pp.176-81 (corresponding author). <http://dx.doi.org/10.1046/j.1369-6513.2000.00092>

Herxheimer, A., McPherson, A., Miller, R., Shepperd, S., Yaphe, J. and Ziebland, S., 2000. Database of Patients' Experiences (DIPEX): a multi-media approach to sharing experiences and information. The Lancet, 355(9214), pp.1540-43 (corresponding author). [http://dx.doi.org/10.1016/S0140-6736\(00\)02174-7](http://dx.doi.org/10.1016/S0140-6736(00)02174-7)