Overend et al. Trials (2016) 17:217
DOI 10.1186/s13063-016-1361-x

Trials

ERRATUM Open Access

Erratum to: CASPER plus (CollAborative care in screen-positive ElderRs with major depressive disorder): study protocol for a randomised controlled trial

Karen Overend[1], Helen Lewis[1], Della Bailey[1], Kate Bosanquet[1], Carolyn Chew-Graham[4], David Ekers[3], Samantha Gascoyne[1], Deborah Hems[1], John Holmes[2], Ada Keding[1], Dean McMillan[1], Shaista Meer[2], Jodi Meredith[1], Natasha Mitchell[1], Sarah Nutbrown[1], Steve Parrott[1], David Richards[5], Gemma Traviss[2], Dominic Trépel[1], Rebecca Woodhouse[1] and Simon Gilbody[1*]

Erratum

After publication of this work [1], we noted that we inadvertently failed to include the complete list of all coauthors. The full list of authors has now been added, and the Authors' contributions and Competing interests section modified accordingly. We are publishing this erratum to update the author list, which is as follows: Karen Overend, Helen Lewis, Della Bailey, Kate Bosanquet, Carolyn Chew-Graham, David Ekers, Samantha Gascoyne, Deborah Hems, John Holmes, Ada Keding, Dean McMillan, Shaista Meer, Jodi Meredith, Natasha Mitchell, Sarah Nutbrown, Steve Parrott, David Richards, Gemma Traviss, Dominic Trépel, Rebecca Woodhouse and Simon Gilbody.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

All authors contributed to the design and development of the study protocol and were members of the project management team. KO collected data, led the qualitative study, contributed to analysis, assisted with trial co-ordination and drafted the manuscript. HL contributed to the study design and development, managed and co-ordinated the overall study and helped draft the manuscript. DB delivered the trial intervention, trained case managers, collected data, contributed to analysis and to drafting the manuscript. KB was local coordinator for the central York site, collected data, contributed to analysis and to drafting the manuscript. CCG designed the process evaluation, supervised qualitative data collection and initial analysis, and contributed to drafting the manuscript. DE was PI for the Durham site and contributed to the study design and development. SGas collected data and contributed to analysis. DH delivered the trial intervention, trained case managers, collected data, contributed to analysis and to drafting the manuscript. JH was PI for the Leeds site and contributed to the study design and development. AK designed statistical analysis for the study and contributed to the drafting of the manuscript. DM contributed to the study design, co-ordination and development. NM contributed to the study design and to drafting the manuscript. SM was local study co-ordinator for the Leeds site. SN collected data, assisted with trial co-ordination, contributed to analysis and to drafting the manuscript. SP designed the economic analysis of health outcomes. DR contributed to the study design. GT was local study co-ordinator and contributed to drafting the manuscript. DT developed the economic analysis of health outcomes. RW collected data and contributed to analysis. SG conceived of the study, directed its design and coordination and contributed to drafting the manuscript. JM delivered the trial intervention, collected data and contributed to analysis. All authors read and approved the final manuscript.

* Correspondence: simon.gilbody@york.ac.uk
[1]Department of Health Sciences, University of York, Seebohm Rowntree, Building Heslington, York YO10 5DD, UK

Author details

[1]Department of Health Sciences, University of York, Seebohm Rowntree, Building Heslington, York YO10 5DD, UK. [2]Leeds Institute of Health Sciences, University of Leeds, Charles Thackrah, Building, 101 Clarendon Road, Leeds LS2 9LJ, UK. [3]Centre for Mental Health Research, University of Durham, Durham TS17 6BH, UK. [4]Research Institute, Primary Care and Health Sciences, Keele University, Keele ST5 5BG, UK. [5]Washington Singer Laboratories, School of Psychology, University of Exeter, Perry Road, Exeter EX4 4QG, UK.

Received: 22 April 2016 Accepted: 22 April 2016
Published online: 27 April 2016

Reference

1. Overend K, Lewis H, Bailey D, Bosanquet K, Chew-Graham C, Ekers D, Gascoyne S, Hems D, Holmes J, Keding A, McMillan D, Meer S, Mitchell N, Nutbrown S, Parrott S, Richards D, Traviss G, Trépel D, Woodhouse R and Gilbody S. CASPER plus (CollAborative care in Screen-Positive ElderRs with major depressive disorder): study protocol for a randomised controlled trial. Trials. 2014;15:451.