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Features of randomised trials designed by the NPEU Perinatal Trials Service during Adrian Grant’s directorship

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Abstract

Adrian Grant pioneered methodological innovations in the randomised trials organised by the Perinatal Trials Service established at the national Perinatal Epidemiology Unit in Oxford, UK. This Commentary discusses these innovations, and shows the wide range of trials designed under his directorship.

Background

In his article recording Adrian Grant’s pioneering use of evidence synthesis in perinatal medicine between 1980 and 1992, Iain Chalmers [1] quoted from a 1984 letter published in The Lancet which Iain and I had co-authored with Adrian [2]. In that letter we alluded to some of the additional principles - beyond the need for systematic review of existing evidence - which became methodological features of the wide range of randomised trials organised by the Perinatal Trials Service (PTS) established at the national Perinatal Epidemiology Unit (NPEU). The trials are listed in Appendix 1.

Methodological features

Appropriate size

Our 1984 letter concluded with a warning that, to avoid the dangers of false inferences from non-randomised comparisons and small randomised trials, many perinatal controlled trials require sample sizes larger than any single unit can generate within a reasonable length of time. Although one centre was sufficient to obtain sufficient sample sizes to address some questions, for other questions, multicentre (often international) trials were needed.

Secure, random allocation

The NPEU PTS used a variety of contextually appropriate methods for secure random allocation - sequentially numbered sealed opaque envelopes, sequentially numbered drug vials, and central random allocation when there was sufficient time to make a call and where reasonable telecommunications existed.

Appropriate design

Most of the trials used two-armed, individually randomised designs. Where appropriate, more complex designs were employed, including factorial trials and a cluster randomised trial.

Involving the views of care-givers and patients and their families

The PTS recognised that our work needed to address questions considered important by caregivers and families, so they were involved in deciding which questions to address, trial design and conduct, and dissemination of results. Taking account of families meant that many PTS trials investigated long term outcomes, such as pain, dyspareunia and incontinence for women, and disability for children.
Facilitating infrastructure
A programme of randomised trials to support these methodological underpinnings needed an infrastructure and the PTS was established in 1982 “to provide a service to busy clinicians who wish to mount large simple-in-design randomized trials...[aiming] to identify moderate, but clinically useful, effects of promising treatments for the most important problems in perinatal care” [3].

The PTS had a flexible five-person core staff and others employed to work on specific trials. This continuity of staff enabled us to build standard operating systems. International trials needed particularly careful coordination, and the provision of trials materials in a number of languages. The eclampsia trial [4], for example was preceded by a pilot study in Argentina, with materials in Spanish.

Adrian Grant’s legacy for perinatal trials
Over the years that Adrian Grant designed perinatal trials in the NPEU, the above methodological innovations and others are listed in the Table 1 below.

Adrian continued to support trials in Aberdeen after his move to direct the Health Services Research Unit there (https://www.abdn.ac.uk/hsru) in 1994, and then for the National Institute for Health Research (http://www.nihr.ac.uk).

Conclusions
Many people are grateful for Adrian’s methodological rigor, his innovative approaches, and his generosity of support, mentoring and teaching. The lives of many babies and their families have been improved by Adrian’s pioneering work in perinatal trials, and the PTS that Adrian created has gone on to become a highly successful Clinical Trials Unit (https://www.npeu.ox.ac.uk/ctu).

Table 1 Adrian Grant’s methodological innovations
- Identification and prioritisation of important questions
- Systematic reviews
- Alliance of patients, carers, and clinicians
- Efficient trial conduct
- Secure randomisation
- Appropriate design, outcomes, and size
- Support for participants
- Newsletters
- Integral economic evaluation
- Embedded methodological research
- Long term follow up
- Feedback of trial results

Appendix

Randomised trials designed by the NPEU Perinatal Trials Service during Adrian Grant’s directorship

Table 2 Antenatal interventions
- Chorion villus sampling vs amniocentesis [5]
- Cervical cerclage [6, 7]
- Breast shells and Hoffman’s exercises [8]
- Formal fetal movement counting [9]
- Placental grading by ultrasonography [10]
- Anti-convulsants for eclampsia [4]
- Low dose aspirin [11, 12]
- Fish-oil supplementation [13]
- ‘Know your Midwife’ [14]
- Social support [15]

Table 3 Intrapartum interventions
- Dublin intrapartum fetal heart rate monitoring [16–19]
- Vacuum extraction vs forceps (Portsmouth operative delivery) [20–22]
- Vacuum extraction: different cups [23]
- Fetal scalp electrode [24]
- Perineal management (Berkshire) [25, 26]
- Perineal suture (Southmead) [27]
- Catgut for the repair of perineal trauma [28, 29]
- Ipswich perineal repair [30–33]

Table 4 Postnatal interventions
- Pelvic floor exercises [34]
- Salt and Savlon bath concentrate [35]
- Ultrasound and pulsed electromagnetic energy treatment for perineal trauma [36, 37]

Table 5 Neonatal interventions
- Neonatal ventriculomegaly [38, 39]
- Dexamethasone [40, 41]
- Prophylactic ethamsylate [42, 43]
- Surfactant [44]
- Extracorporeal membrane oxygenation [45–53]
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DE is the sole contributor. The author read and approved the final manuscript.

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The authors declare that they have no competing interests.

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References
1. Chalmers I, Grant AM. Rationale for and work of the perinatal trials service. Early Hum Dev. 1980;5:1–5.
2. Chalmers I, Elbourne D, Grant A. Phenobarbitone and periventricular haemorrhage. Lancet. 1984;2:345–49.
3. Grant AM. Rational for and work of the perinatal trials service. Early Hum Dev. 1982;6(2):79–85.
4. Grant AM. Some answers to questions raised about the Dublin trial. Birth. 1989;16(3):255–6.
5. Garcia J, Anderson J, Vacca A, Elbourne DR, Grant AM, Chalmers I. Views of women and their medical and midwifery attendants about instrumental delivery using vacuum extraction and forceps. J Psychosom Obstet Gynaecol. 1985;1:9–17.
6. Carmody F, Grant AM, Mutch LM, Vacca A, Chalmers I. Follow up of babies delivered in a randomized controlled comparison of vacuum extraction and forceps delivery. Acta Obstet Gynecol Scand. 1986;65(7):763–6.
7. Rush RW, Isaacs S, McPherson K, Jones L, Chalmers I, Grant AM. A randomized controlled trial of glycerol-impregnated chromic catgut with untreated chromic catgut for the repair of perineal trauma. Br J Obstet Gynaecol. 1989;96(3):426–30.
8. Grant AM. Rationale for and work of the perinatal trials service. Early Hum Dev. 1981;5(7):497–8.
9. Rush RW, Isaacs S, McPherson K, Jones L, Chalmers I, Grant AM. A randomized controlled trial of cervical cerclage in women at high risk of spontaneous preterm delivery. Br J Obstet Gynaecol. 1984;91(8):724–30.
10. Alexander JM, Grant AM, Campbell MJ. Randomized controlled trial of breast shells and Hoffman’s exercises for inverted and non-protractile nipples. Br Med J. 1992;304:1030–2.
11. CLASP (Collaborative Low-dose Aspirin Study in Pregnancy) Collaborative Group. CLASP: a randomised trial of low-dose aspirin for the prevention and treatment of pre-eclampsia among 9364 pregnant women. Lancet. 1994;343:619–24.
12. Rotchell YE, Cruickshank JK, Phillips Gay M, Griffiths J, Stewart A, Farrell B, Ayers S, Hennis A, Grant AM, Duley L, Collins RE. Barbados low dose aspirin scheme - a randomised controlled trial to compare three types of fetal scalp electrode. Br J Obstet Gynaecol. 1992;99(9):806–4.
13. Olsen SF, Sorensen JD, Secher NJ, Hedegaard M, Henriksen TB, Hansen HS, Ayers S, Hennis A, Grant AM, Duley L, Collins RE. CLASP: a randomised trial of low-dose aspirin for the prevention of pre-eclampsia and its complications. Br J Obstet Gynaecol. 1998;105:286–92.
14. Olsen SF, Sorensen JD, Secher NJ, Hedegaard M, Henriksen TB, Hansen HS, Ayers S, Hennis A, Grant AM. Randomized controlled trial of the effect of fish-oil supplement on pregnancy duration. The lancet. 1992;339(8808):1003–7.
15. Flint C, Poulengeris P, Grant AM. The ‘know your Midwife’ scheme - a randomised trial of continuity of care by a team of midwives. Midwifery. 1995;11:5–6.
16. MacDonald D, Grant AM, Sheridan-Pereira M, Boylan P, Chalmers I. The Dublin randomized controlled trial of intrapartum fetal heart rate monitoring. Am J Obstet Gynecol. 1985;152(5):524–9.
17. Garcia J, Cory M, MacDonald D, Elbourne DR, Grant AM. Mothers’ views of continuous electronic fetal heart monitoring and intermittent auscultation in a randomized controlled trial. Birth. 1985;12(2):79–86.
18. Grant AM. Some answers to questions raised about the Dublin trial. Birth. 1986;13(4):255–6.
19. Grant AM, O’Brien N, Joy M-T, Hennessy E, MacDonald D. Cerebral palsy among children born during the Dublin randomized trial of intrapartum monitoring. Lancet 1989;86(749):1233–1236.
20. Vacca A, Grant AM, Wyatt G, Chalmers I. Portsmouth operative delivery trial: a comparison of vacuum extraction and forceps delivery. Br J Obstet Gynaecol. 1983;90(12):1107–12.
21. Garcia J, Anderson J, Vacca A, Elbourne DR, Grant AM, Chalmers I. Views of women and their medical and midwifery attendants about instrumental delivery using vacuum extraction and forceps. J Psychosom Obstet Gynaecol. 1985;4:1–9.
22. Carmody F, Grant AM, Mutch LM, Vacca A, Chalmers I. Follow up of babies delivered in a randomized controlled comparison of vacuum extraction and forceps delivery. Acta Obstet Gynecol Scand. 1986;65(7):763–6.
23. Carmody F, Grant AM, Somchitwong M. Vacuum extraction: a randomized controlled comparison of the next generation cup with the original bird cup. J Perinat Med. 1986;14:95–100.
24. Needs L, Grant AM, Sleep J, Ayers S, Henson G. A randomized controlled trial to compare three types of fetal scalp electrode. Br J Obstet Gynaecol. 1992;99(3):302–6.
25. Sleep J, Grant AM, Garcia J, Elbourne DR, Spencer JAD, Chalmers I. West Berkshire perineal management trial. Br Med J. 1984;289:587–90.
26. Sleep J, Grant AM. West Berkshire perineal management trial: three year follow up. Br Med J. 1987;295:749–51.
27. Mahomed K, Grant AM, Ashurst H, James D. The Southmead perinatal suture study. A randomized comparison of suture materials and suturing techniques for repair of perineal trauma. Br J Obstet Gynaecol 1989;86(11):1272–1280.
28. Spencer JAD, Grant AM, Elbourne DR, Garcia J, Sleep J. A randomized comparison of glycerol-impregnated chromic catgut with untreated chromic catgut for the repair of perineal trauma. Br J Obstet Gynaecol. 1989;96:426–30.
29. Grant AM, Sleep J, Ashurst H, Spencer JAD. Dyspareunia associated with the use of glycerol-impregnated catgut to repair perineal trauma - report of a three-year follow-up study. Br J Obstet Gynaecol. 1989;96:741–3.
30. Mackirdt C, Gordon B, Fern E, Ayers S, Truesdale A, Grant AM. The Ipswich childbirth study: 2. A randomised comparison of polyglycolin 910 with chromic catgut for postpartum perineal repair. Br J Obstet Gynaecol. 1998;105(4):441–5.
31. Gordon B, Mackirdt C, Fern E, Truesdale A, Ayers S, Grant AM. The Ipswich childbirth study: 1. A randomised evaluation of two stage postpartum perineal repair leaving the skin unsutured. Br J Obstet Gynaecol. 1998;105(4):435–40.
32. Grant AM, Gordon B, Mackirdt C, Fern E, Truesdale A, Ayers S, Grant AM. The Ipswich childbirth study: one year follow-up of alternative methods used in perineal repair. Br J Obstet Gynaecol. 2001;108:34–40.
33. Petrou S, Gordon B, Mackirdt C, Fern E, Ayers S, Grant AM, Truesdale A, McCandlish R. How cost-effective is it to leave perineal skin unsutured? British J Midwifery. 2001;9(4):209–14.
34. Sleep J, Grant AM. Pelvic floor exercises in post-natal care. Midwifery. 1998;14:318–6.
35. Sleep J, Grant AM. Effects of salt and savon bath concentrate postpartum. Nurs Times. 1988;84:55–7.
36. Everett T, McIntosh J, Grant AM. Ultrasound therapy for persistent postnatal perineal pain and dyspareunia: a randomised placebo-controlled trial. Physiotherapy. 1992;78:263–7.
37. Grant AM, Sleep J, McIntosh J, Ashurst H. Ultrasound and pulsed electromagnetic energy treatment for perineal trauma. A randomized placebo-controlled trial. Br J Obstet Gynaecol. 1998;105(4):435–40.
38. Ventriculomegaly Trial Group. Randomised trial of early tapping in neonatal posthaemorrhagic ventricular dilatation. Arch dis child. 1990;65(1):3–10.
39. Ventriculomegaly Trial Group. Randomised trial of early tapping in neonatal posthaemorrhagic ventricular dilatation: results at 30 months. Arch dis child. 1994;70(2):129–36.
40. Collaborative Dexamethasone Trial Group. Dexamethasone therapy in neonatal chronic lung disease: an international placebo controlled trial. Pediatrics. 1991;88:421–7.
41. Jones RAK, Wincott E, Elbourne DR, Grant AM. Controlled trial of dexamethasone in neonatal chronic lung disease: a 3-year follow-up. Pediatrics. 1995;96:897–906.
42. The EC Ethamsylate Trial Group. The EC randomised controlled trial of prophylactic ethamsylate for very preterm neonates: early mortality and morbidity. Arch Dis Child. 1994;70:F201–5.
43. Elbourne DR, Ayers S, Dellagrammaticas H, Johnson A, Leloup M, Lenoir-Piat S. EC Ethamsylate trial group. Randomised controlled trial of prophylactic ethamsylate: follow up at 2 years of age. Arch Dis Child. 2001;84:F183–7.
44. OSRIS Collaborative Group. Early versus delayed neonatal administration of a synthetic surfactant - the judgment of OSRIS. Lancet. 1992;340:1363–9.
45. UK Collaborative ECMO Trial Group. UK collaborative randomised trial of neonatal extracorporeal membrane oxygenation. Lancet. 1996;348:75–82.
46. Howard S, Mugford M, Normand C, Elbourne DR, Grant AM, Field DJ, Johnson A. A cost effectiveness analysis of neonatal ECMO using existing evidence. Int J Technol Assess Health Care. 1996;12:80–92.
47. Snowdon C, Garcia J, Elbourne DR. Making sense of randomization: responses of parents of critically ill babies to random allocation of treatment in a clinical trial. Soc Sci Med. 1997;45:1337–55.
48. Snowdon C, Garcia J, Elbourne DR. Reactions of participants to the results of a randomised controlled trial: exploratory study. Br Med J. 1998;317:21–6.
49. Snowdon C, Elbourne DR, Garcia J. Zelen randomization: attitudes of parents participating in a neonatal clinical trial. Control Clin Trials. 1999;20:149–71.
50. Bennett CC, Johnson A, Field DJ, Elbourne DR. For the UK collaborative ECMO trial group. UK collaborative randomised trial of neonatal extracorporeal membrane oxygenation: follow-up to age 4 years. Lancet. 2001;357:1094–6.
51. Petrou S, Edwards L. Cost-effectiveness analysis of neonatal extracorporeal membrane oxygenation based on four year results from the UK collaborative ECMO trial. Arch Dis Child. 2004;89:F263–F8.
52. Petrou S, Bischof M, Bennett CC, Elbourne DR, Field DJ, McNally H. Cost-effectiveness of neonatal extracorporeal membrane oxygenation based on 7 year results from the United Kingdom collaborative ECMO trial. Pediatrics. 2006;117:1640–9.
53. McNally H, Bennett CC, Elbourne DR, Field DJ. For the UK collaborative ECMO trial group. United Kingdom collaborative randomized trial of neonatal extracorporeal membrane oxygenation: follow-up to age 7 years. Pediatrics. 2006;117:845–54.