

BJP The British Journal of Psychiatry Psych

Economic evaluation of a randomised controlled trial for anorexia nervosa in adolescents

SARAH BYFORD, BARBARA BARRETT, CHRIS ROBERTS, ANDREW CLARK, VANESSA EDWARDS, NICOLA SMETHURST and SIMON G. GOWERS

BJP 2007, 191:436-440.

Access the most recent version at DOI: [10.1192/bjp.bp.107.036806](https://doi.org/10.1192/bjp.bp.107.036806)

Supplementary Material

Supplementary material can be found at:
<http://bjp.rcpsych.org/content/suppl/2007/11/01/191.5.436.DC1.html>

References

This article cites 0 articles, 0 of which you can access for free at:
<http://bjp.rcpsych.org/content/191/5/436#BIBL>

Reprints/permissions

To obtain reprints or permission to reproduce material from this paper, please write to permissions@rcpsych.ac.uk

You can respond to this article at

<http://bjp.rcpsych.org/cgi/eletter-submit/191/5/436>

Downloaded from

<http://bjp.rcpsych.org/> on February 17, 2013
Published by [The Royal College of Psychiatrists](#)

Economic evaluation of a randomised controlled trial for anorexia nervosa in adolescents[†]

SARAH BYFORD, BARBARA BARRETT, CHRIS ROBERTS, ANDREW CLARK, VANESSA EDWARDS, NICOLA SMETHURST and SIMON G. GOWERS

Background Young people with anorexia nervosa are often admitted to hospital for treatment. As well as being disruptive to school, family and social life, in-patient treatment is expensive, yet cost-effectiveness evidence is lacking.

Aims Cost-effectiveness analysis of three treatment strategies for adolescents with anorexia nervosa.

Method UK multicentre randomised, controlled trial comparing in-patient psychiatric treatment, specialist out-patient treatment and general out-patient treatment. Outcomes and costs assessed at baseline, 1 and 2 years.

Results There were 167 young people in the trial. There were no statistically significant differences in clinical outcome between the three groups at 2 years. The specialist out-patient group was less costly over the 2-year follow-up (mean total cost £26 738) than the in-patient (£34 531) and general out-patient treatment (£40 794) groups, but this result was not statistically significant. Exploration of the uncertainty associated with the costs and effects of the three treatments suggests that specialist out-patient treatment has the highest probability of being cost-effective.

Conclusions On the basis of cost-effectiveness, these results support the provision of specialist out-patient services for adolescents with anorexia nervosa.

Declaration of interest None. Funding detailed in Acknowledgements.

Anorexia nervosa is commonly associated with severe physical, psychological and social impairments, high levels of mortality (National Collaborating Centre for Mental Health, 2004) and a significant cost burden (Striegel-Moore *et al*, 2000; Simon *et al*, 2005). Young people with anorexia nervosa are often admitted to hospital for treatment. This is disruptive to school, family and social life, and in-patient treatment is an expensive option, yet evidence to support its cost-effectiveness is lacking (Romeo *et al*, 2005). A recent systematic review did not identify any economic evaluations of treatments for anorexia nervosa (National Collaborating Centre for Mental Health, 2004). One subsequent modelling study has been reported (Crow & Nyman, 2004). However, the authors acknowledge the limitations of modelling and the need for controlled trials. We report the results of an economic evaluation of psychiatric in-patient, specialist out-patient and general out-patient services for adolescents with anorexia nervosa carried out alongside a randomised controlled clinical evaluation – the Treatment Outcome for Child and adolescent Anorexia Nervosa (TOuCAN) trial (ISRCTN39345394).

METHOD

The aim of the TOuCAN trial was to explore the clinical and cost-effectiveness of in-patient, specialist out-patient and general out-patient services for adolescents with anorexia nervosa. The main economic hypotheses were that: (a) specialist out-patient services would be more cost-effective than general out-patient treatment in community child and adolescent mental health services (CAMHS); and (b) out-patient services would be more cost-effective than in-patient services.

Trial design

The design of this multicentre randomised controlled trial is described in detail in the

accompanying paper (Gowers *et al*, 2007, this issue). In brief, adolescents aged 12–18 years with a DSM-IV diagnosis of anorexia nervosa were recruited to the study from community CAMHS in the north-west of England between March 2000 and December 2003. No exclusions were made on the grounds of clinical severity, but the responsible clinician reserved the right to refer for acute medical management according to urgent need. Participants with severe intellectual difficulties or severe, chronic comorbid physical conditions affecting digestion and/or metabolism were excluded. Randomisation was carried out by an independent statistical centre. Stochastic minimisation was used to control for gender, age (above and below 16 years) and body mass index (BMI) (above and below 15.5).

Interventions

In-patient psychiatric treatment was for 6 weeks in the first instance, extended as clinically indicated and determined by the treating service. The in-patient services used a multidisciplinary psychiatric approach with the aim of normalising eating, restoring healthy weight and facilitating psychological (cognitive) change. Specialist out-patient treatment, described in detail elsewhere (Gowers & Smyth, 2004; Gowers, 2006), was manualised and comprised individual cognitive-behavioural therapy (CBT), parental counselling with the participant, dietary therapy and multi-modal feedback. Treatment as usual in community CAMHS generally involved a multidisciplinary, individual- and family-based approach, with variable dietetic and paediatric liaison.

Clinical outcomes

Research assessors masked to treatment allocation carried out assessments at baseline, 1 and 2 years after trial entry. The *a priori* primary outcome measure for the clinical and economic evaluation was the Morgan-Russell Average Outcome Scale (MRAOS; Morgan & Hayward, 1988), adjusted for adolescents. Full details of outcome measures, their reliability and validity, are reported in the accompanying paper (Gowers *et al*, 2007, this issue).

Cost

The economic evaluation took a broad service-providing perspective, including that of the health, social services, education,

[†]See pp. 427–435, this issue.

voluntary and private sectors. Information on resource use was collected in interview at the 1- and 2-year follow-up assessments using the Child and Adolescent Service Use Schedule (CA-SUS), developed by the authors in previous research with young people and adapted for the purpose of the current study (Byford *et al*, 1999; Harrington *et al*, 2000; Barrett *et al*, 2006). Data on hospital contacts were collected from clinical records to avoid patients revealing their treatment group to the research assessors.

All unit costs were for the financial year 2003–2004. Costs in the second year were discounted at a rate of 3.5%, as recommended by the National Institute for Health and Clinical Excellence (National Institute for Clinical Excellence, 2004). This was varied from 0 to 6% in sensitivity analysis.

All National Health Service (NHS) hospital contacts, including the trial interventions, were costed using NHS reference costs (Department of Health, 2004). The unit costs of private sector in-patient stays were collected through direct personal communication with each facility. Unit costs of community health and social services were taken from national publications (Curtis & Netten, 2004). The costs of schooling came from a number of sources including various Ofsted reports (the inspectorate and regulatory body for schools in England, see <http://www.ofsted.gov.uk>) and published documents (Berridge *et al*, 2002; Independent Schools Council, 2004). Medications were costed using the British National Formulary (British Medical Association & Royal Pharmaceutical Society of Great Britain, 2004). Where necessary, unit costs were inflated to 2003–2004 costs using the Hospital and Community Health Services inflation indices (Curtis & Netten, 2004).

Statistical methods

All economic analyses were carried out on an intention-to-treat basis using a statistical analysis plan drawn up prior to the analysis of the data. The primary analysis was of total costs over 2 years for the sample of young people with complete economic data.

Although costs were not normally distributed, analyses compared mean costs in the three groups using analysis of covariance with covariates for pre-specified baseline characteristics: site (Liverpool and Manchester), gender, age at baseline,

baseline BMI and baseline MRAOS score. The robustness of the parametric tests was confirmed using bootstrapping (Efron & Tibshirani, 1993), as recommended by Barber & Thompson (1998). The impact of drop-out was assessed by comparing the baseline characteristics of participants who had missing data with those who had full economic data.

Cost-effectiveness was assessed through the calculation of incremental cost-effectiveness ratios (ICER) – the additional costs of one intervention compared with another divided by the additional effects of one intervention compared with another (Van Hout *et al*, 1994), in this case using the MRAOS measure of effectiveness. When more than two strategies are compared, ICERs are calculated using rules of dominance and extended dominance (Johannesson & Weinstein, 1993). Strategies are ranked by cost, from the least expensive to the most expensive, and if a strategy is more expensive and less effective than the previous strategy, it is said to be dominated and is excluded from the calculation of ICERs. This process compares strategies in terms of observed differences in costs and effects, regardless of the statistical significance of the difference.

Uncertainty around the cost and effectiveness estimates was represented by cost-effectiveness acceptability curves (Van Hout *et al*, 1994; Fenwick *et al*, 2001). Repeat re-sampling from the costs and effectiveness data (bootstrapping) was used to generate a distribution of mean costs and effects for the three treatments. These distributions were used to calculate the probability that each of the treatments is the optimal choice, subject to a range of possible maximum values (ceiling ratio, λ) that a decision-maker might be willing to pay for a unit improvement in MRAOS score. Cost-effectiveness acceptability curves are presented by plotting these probabilities for a range of possible values of the ceiling ratio. These curves incorporate the uncertainty that exists around the estimates of mean costs and effects as a result of sampling variation and uncertainty regarding the maximum cost-effectiveness ratio that a decision-maker would consider acceptable (Fenwick & Byford, 2005).

Missing data were explored in three sensitivity analyses using the following data: (a) hospital cost data collected from clinical records and available for a larger sample of young people than full economic data from the CA-SUS; (b) hospital cost

data collected from records plus missing non-hospital cost data imputed using the last value carried forward approach for participants with missing year-2 data; and (c) hospital cost data collected from records plus mean imputation by randomised group of missing non-hospital cost data. The results of all sensitivity analyses are reported in the data supplement to the online version of this paper.

RESULTS

Participants

There were 167 young people entering the trial and they were randomised to in-patient care ($n=57$), specialist out-patient services ($n=55$) or treatment as usual by general CAMHS ($n=55$). Full economic data for the 2-year follow-up period were available for 135 young people (81%), 47 in the in-patient group, 45 in the specialist out-patient group and 43 in the general out-patient group. A comparison of baseline characteristics (site, age, gender, BMI and MRAOS score) revealed no significant differences between those included in the economic evaluation and those who were missing, and there was no difference overall in missing data between the three treatment groups. Length of follow-up varied somewhat (range 99–118 weeks), however there was no significant difference in length of follow-up between the three treatment groups (mean 105 weeks in the in-patient and general out-patient groups and 106 in the specialist out-patient group).

Outcomes

There were no significant differences between the three groups by intention to treat at either 1- or 2-year follow-up on the MRAOS (in-patient 2-year global score 8.3, specialist out-patient 8.4, general out-patient 8.3; $P=0.838$). Full clinical outcome data are reported by Gowers *et al* (2007, this issue).

Resource use

Table 1 details the mean number of contacts young people had with all services over the 2-year follow-up period. Resource use differed little between the groups except for in-patient and out-patient contacts. The general out-patient group spent more time in hospital and had a greater number of out-patient attendances on average than the specialist out-patient or in-patient groups. The specialist out-patient group spent the least amount of time in hospital.

Exploration of hospital contacts over time reveals that a larger proportion of days were spent in hospital in the first year (in-patient group 62 days, specialist out-patient 35, general out-patient 65) than the second year (in-patient group 12 days, specialist out-patient 20, general out-patient 24). The in-patient treatment of those allocated to out-patient treatment generally occurred after assigned treatment had ended. Details of adherence to treatment are given in the accompanying paper (Gowers *et al*, 2007, this issue).

Hospital contacts reported in Table 1 include all specialties. However, the vast majority of contacts were psychiatric (71% of in-patient admissions and 90% of in-patient days) or paediatric (20% of admissions and 10% of in-patient days). Other specialties (9% of admissions and 0.2% of in-patient days) included gastroenterology, general medicine, haematology, intensive care unit, obstetrics, orthopaedics, plastic surgery and urology.

Time in education was similar across the three groups, however on average participants spent a significant proportion of the 2-year follow-up period out of education (approximately 10 out of the 24 months of follow-up).

Costs

Table 2 details the total mean costs per participant over the 2-year follow-up period. There were no statistically significant differences between the three groups. In terms of observed differences, the specialist out-patient group was consistently cheaper than the other two groups and the general out-patient group was the most expensive of the three. The bootstrapped results differed little and are thus not reported here. Hospital costs constitute the greatest proportion of total costs (93% in each group), with few community health and social services being used.

Cost-effectiveness analysis

Using the rules of dominance described in the Method section, specialist out-patient treatment (bootstrapped mean cost per participant £26 797; bootstrapped mean effect 8.35) dominates the in-patient group (£34 371; 8.26) and the general out-patient group (£40 520; 8.26) since it is both cheaper and more effective. Figure 1 illustrates the uncertainty associated with the costs and effects of the three treatments at 2 years and demonstrates that if decision-makers were willing to pay nothing for a unit

Table 1 Use of resources during the 2-year follow-up period

Service	In-patient (n=47)	Specialist out-patient (n=45)	General out-patient (n=43)
	Mean (s.d.)		
Secondary health services			
In-patient nights	73 (124)	55 (114)	89 (159)
Out-patient appointments	23 (20)	26 (22)	31 (24)
Day patient contacts	4 (12)	1 (7)	1 (5)
Accident and emergency contacts	0 (1)	1 (2)	0 (1)
Community health and social services contacts			
General practitioner	7 (9)	7 (9)	6 (8)
Practice nurse	2 (5)	3 (10)	1 (4)
Dietician	0 (3)	0 (0)	0 (0)
District nurse	0 (0)	0 (2)	0 (0)
Health visitor	0 (1)	0 (0)	0 (0)
Community paediatrician	0 (1)	0 (0)	0 (3)
Community psychiatric nurse	0 (1)	1 (7)	1 (5)
Clinical psychologist	1 (8)	0 (1)	0 (0)
Counsellor	0 (1)	0 (2)	0 (1)
Family therapist	1 (8)	0 (0)	0 (2)
Dentist	0 (0)	0 (1)	0 (0)
School doctor	0 (2)	0 (0)	0 (0)
School nurse	1 (6)	1 (2)	1 (2)
Social worker	1 (2)	1 (4)	0 (1)
Eating disorders association	0 (1)	0 (1)	0 (0)
Family therapy	0 (1)	0 (1)	0 (0)
Foster care, days	0 (0)	0 (0)	2 (13)
Education			
State day school, months	11 (9)	10 (9)	12 (10)
Independent day school, months	1 (4)	2 (6)	1 (5)
Independent boarding school, months	0 (1)	0 (0)	0 (2)
Hospital school, months	1 (3)	1 (2)	2 (4)
Home tuition, months	0 (1)	1 (3)	1 (4)
School counsellor, contacts	1 (7)	1 (7)	0 (0)
Education welfare officer, contacts	0 (0)	0 (1)	0 (1)

increase in MRAOS score, there is a 78% chance of specialist out-patient services being the most cost-effective strategy, 16% for in-patient services and only 6% for general CAMHS. The probability of specialist out-patient services being the most cost-effective strategy decreases with increasing levels of willingness to pay for gains in effectiveness, levelling out at around 47%, but remains higher than the other two strategies over the full range of willingness to pay values shown, and beyond.

Figure 1 suggests that the probability of our first hypothesis being true is high, i.e. that specialist out-patient services are more cost-effective than general out-patient

services. Figure 2 depicts the cost-effectiveness acceptability curve for the second hypothesis, i.e. in-patient *v.* out-patient services (specialist combined with general), and shows that there is a greater probability of out-patient services being more cost-effective than in-patient services for the full range of values of willingness to pay.

DISCUSSION

This paper reports the results of the first economic evaluation of alternative strategies for the treatment of anorexia nervosa using primary data collected from a randomised controlled trial.

Table 2 Total cost (£) per young person over the 2-year follow-up period

Sector	In-patient (n=47)			Specialist out-patient (n=45)			General out-patient (n=43)			ANOVA ¹ P
	Mean (s.d.)	Minimum	Maximum	Mean (s.d.)	Minimum	Maximum	Mean (s.d.)	Minimum	Maximum	
Secondary health care	32 015 (51 541)	69	280 580	24 724 (46 231)	103	244 043	37 746 (62 046)	68	269 236	0.456
Primary health care	380 (640)	4	2773	385 (873)	0	4325	245 (361)	0	2078	0.503
Education	2098 (2115)	0	8783	1595 (1456)	0	6393	2654 (2228)	0	12 093	0.088
Other community services ²	37 (110)	0	513	35 (104)	0	445	150 (806)	0	5244	0.504
Total 2-year cost	34 531 (52 439)	86	282 508	26 738 (46 809)	462	244 174	40 794 (63 652)	1483	274 838	0.426
Total cost per week	325 (487)	1	2588	253 (442)	4	2288	386 (600)	14	2603	0.423

1. Adjusted for site, gender, age at baseline, baseline body mass index and baseline Morgan–Russell Average Outcome Scale score.
 2. Includes community social, voluntary and private sector services.

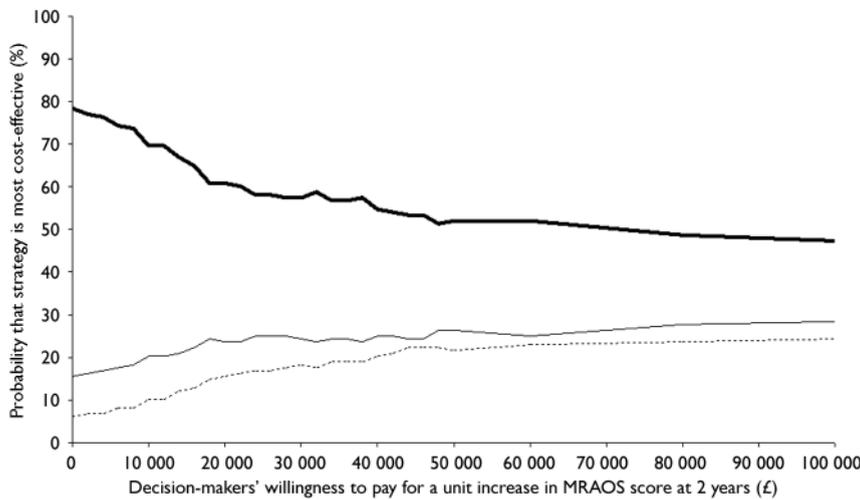


Fig. 1 Cost-effectiveness acceptability curve for Morgan–Russell Average Outcome Scale (MRAOS) score. —, Specialist out-patient; —, in-patient; - - -, general out-patient.

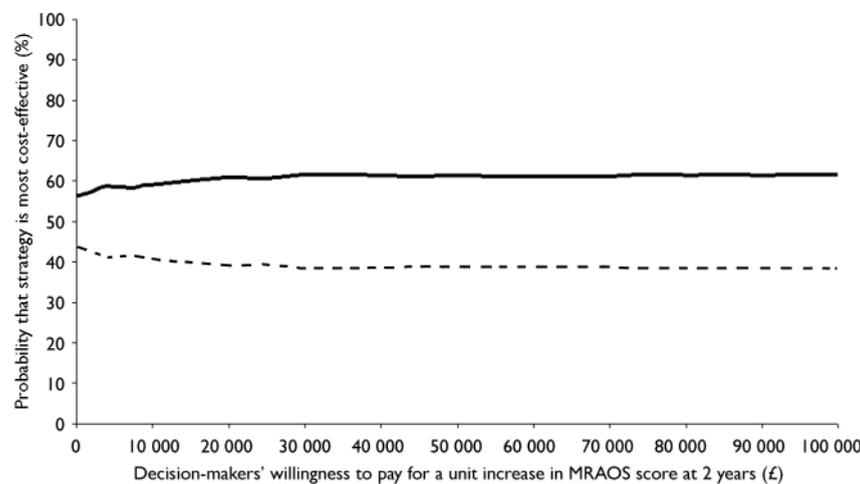


Fig. 2 Cost-effectiveness acceptability curve for Morgan–Russell Average Outcome Scale (MRAOS) score. —, Out-patient; - - -, in-patient.

Clinical outcomes, resource use and cost

There were no statistically significant differences in clinical outcomes between the three groups, with results demonstrating similar improvements in all groups over time. The specialist out-patient group was observed to be the cheapest of the three groups and the general out-patient group the most expensive. However, these differences were not statistically significant. These findings were robust to changes in the discount rate and in analyses of missing data.

Observed differences in total mean cost per participant were almost entirely due to differences in the length of time spent in hospital. Secondary healthcare costs accounted for over 90% of all costs and, of this, almost 90% was due to in-patient stays. The majority of in-patient stays took place in the first year. Although not randomised to psychiatric in-patient services, the general out-patient service group spent almost as much time in hospital as the in-patient group, suggesting that general CAMHS were less successful at maintaining these young people in the community than specialist out-patient services. With the exception of CAMHS, participants used very few community health and social services. Months in education was similar across the groups on average, but highlighted the significant proportion of time participants spent out of education, presumably as a result of their illness.

At almost £17 000 per year on average, the annual service costs of caring for this group of young people were high. Although much higher than the cost of conditions generally treated in the community, for

example conduct disorder with annual service cost estimates varying between £1300 and £3200 (Harrington *et al*, 2000; Romeo *et al*, 2006), this figure is similar to the cost of a cohort of young people treated in child and adolescent psychiatric in-patient wards, estimated to be £24 000 per admission (Green *et al*, 2007). Although slightly higher than the costs reported in this paper, the mean length of stay was longer at 116 days.

Cost-effectiveness

Specialist out-patient services were found to be the dominant treatment option in terms of incremental cost-effectiveness (more effective and less costly). Exploration of the associated uncertainty supported this finding. In terms of our hypotheses, the data suggest that specialist out-patient services have a higher probability of being cost-effective than general out-patient services and that out-patient services (specialist combined with general) have a higher probability of being cost-effective than psychiatric in-patient services.

Limitations

Despite substantial differences in observed cost data, these differences did not reach statistical significance. This may be due to inadequate sample sizes for the economic evaluation. Sample size calculations were based on the primary outcome measure, the MRAOS. Calculations on the basis of cost or cost-effectiveness were not feasible at the design stage because of the lack of any relevant published cost data. Although acknowledging this limitation, the use of a decision-making approach to the economic evaluation provides probabilistic evidence of the cost-effectiveness of the alternative treatment strategies, given the data currently available. Although larger trials may be considered in future research, this must be balanced against the cost of additional research in a disease area where low prevalence rates necessitate multicentre evaluation. Analysis of patients excluded because of missing economic data did not suggest any bias; patients included in the economic evaluation did not differ significantly from those excluded and there was no evidence to suggest any bias in missing data between the three treatment groups.

SARAH BYFORD, MSc, BARBARA BARRETT, MSc, King's College London, Centre for the Economics of Mental Health, Institute of Psychiatry, London; CHRIS ROBERTS, PhD, Biostatistics Group, Division of Epidemiology and Health Sciences, University of Manchester; ANDREW CLARK, FRCPsych, Bolton Salford and Trafford Mental Health NHS Trust, Manchester; VANESSA EDWARDS, BA, Peninsular College of Medicine and Dentistry, Exeter; NICOLA SMETHURST, BA, SIMON G. GOWERS, FRCPsych, University of Liverpool, Section of Adolescent Psychiatry, Academic Unit, Chester, UK

Correspondence: Sarah Byford, Box No 24, Centre for the Economics of Mental Health, Institute of Psychiatry, De Crespigny Park, London SE5 8AF, UK. Email: s.byford@iop.kcl.ac.uk

(First received 9 February 2007, final revision 10 May 2007, accepted 29 June 2007)

ACKNOWLEDGEMENTS

This project is funded by the UK National Health Service Health Technology Assessment programme, Grant No. 97/42/02 and supported by the North West MHEN. This paper is dedicated to Professor Richard Harrington.

REFERENCES

- Barber, J. A. & Thompson, S. G. (1998)** Analysis and interpretation of cost data in randomised control trials: review of published studies. *BMJ*, **317**, 1195–2000.
- Barrett, B., Byford, S., Chitsabesan, P., et al (2006)** Mental health provision for young offenders: service use and cost. *British Journal of Psychiatry*, **188**, 541–546.
- Berridge, D., Beecham, J., Brodie, I., et al (2002)** *Costs and Consequences of Services for Troubled Adolescents: an Exploratory, Analytic Study* (Department of Health Report). University of Luton.
- British Medical Association & Royal Pharmaceutical Society of Great Britain (2004)** *British National Formulary 48*. BMJ Books & Pharmaceutical Press.
- Byford, S., Harrington, R., Torgerson, D., et al (1999)** Cost-effectiveness analysis of a home-based social work intervention for children and adolescents who have deliberately poisoned themselves. Results of a randomised controlled trial. *British Journal of Psychiatry*, **174**, 56–62.
- Crow, S. J. & Nyman, J. A. (2004)** The cost-effectiveness of anorexia nervosa treatment. *International Journal of Eating Disorders*, **35**, 155–160.
- Curtis, L. & Netten, A. (2004)** *Unit Costs of Health and Social Care*. Personal Social Services Research Unit.
- Department of Health (2004)** *NHS Reference Costs*. Department of Health.
- Efron, B. & Tibshirani, R. J. (1993)** *An Introduction to the Bootstrap*. Chapman & Hall.
- Fenwick, E. & Byford, S. (2005)** A guide to cost-effectiveness acceptability curves. *British Journal of Psychiatry*, **187**, 106–108.
- Fenwick, E., Claxton, K. & Sculpher, M. (2001)** Representing uncertainty: the role of cost-effectiveness acceptability curves. *Health Economics*, **10**, 779–787.
- Gowers, S. G. (2006)** Evidence based research in CBT with adolescent eating disorders. *Child and Adolescent Mental Health*, **11**, 9–12.
- Gowers, S. & Smyth, B. (2004)** The impact of a Motivational Assessment Interview on initial response to treatment in adolescent anorexia nervosa. *European Eating Disorders Review*, **12**, 87–93.
- Gowers S. G., Clark A., Roberts, C., et al (2007)** Clinical effectiveness of treatments for anorexia nervosa in adolescents. Randomised controlled trial. *British Journal of Psychiatry*, **191**, 427–435.
- Green, J., Jacobs, B., Beecham, J., et al (2007)** Inpatient treatment in child and adolescent psychiatry – an exploratory prospective study of health gain and costs. *Journal of Child Psychology and Psychiatry*, in press.
- Harrington, R., Peters, S., Green, J., et al (2000)** Randomised comparison of the effectiveness and costs of community and hospital based mental health services for children with behavioural disorders. *BMJ*, **321**, 1047–1050.
- Independent Schools Council (2004)** *Annual Census 2004*. Independent Schools Council.
- Johannesson, M. & Weinstein, M. C. (1993)** On the decision rules of cost-effectiveness analysis. *Journal of Health Economics*, **12**, 459–467.
- Morgan, H. G. & Hayward, A. E. (1988)** Clinical assessment of anorexia nervosa. The Morgan–Russell outcome assessment schedule. *British Journal of Psychiatry*, **152**, 367–372.
- National Collaborating Centre for Mental Health (2004)** *Core Interventions in the Treatment and Management of Anorexia Nervosa and Related Eating Disorders*. British Psychological Society & Gaskell.
- National Institute for Clinical Excellence (2004)** *Guide to the Methods of Technology Appraisals*. NICE.
- Romeo, R., Byford, S. & Knapp, M. (2005)** Economic evaluations of child and adolescent mental health interventions: a systematic review. *Journal of Child Psychology and Psychiatry*, **46**, 919–930.
- Romeo, R., Knapp, M. & Scott, S. (2006)** Economic cost of severe antisocial behaviour in children – and who pays it. *British Journal of Psychiatry*, **188**, 547–553.
- Simon, J., Schmidt, U. & Pilling, S. (2005)** The health service use and cost of eating disorders. *Psychological Medicine*, **35**, 1543–1551.
- Striegel-Moore, R. H., Leslie, D., Petrelli, S. A., et al (2000)** One-year use and cost of inpatient and outpatient services among female and male patients with an eating disorder: evidence from a national database of health insurance claims. *International Journal of Eating Disorders*, **27**, 381–389.
- Van Hout, B. A., Al, M. J., Gordon, G. S., et al (1994)** Costs, effects and c/e-ratios alongside a clinical trial. *Health Economics*, **3**, 309–319.