

Hemiarthroplasty and total hip arthroplasty for treating primary intracapsular fracture of the hip: a systematic review and cost-effectiveness analysis

C Carroll,^{1*} M Stevenson,¹ A Scope,¹ P Evans¹
and S Buckley²

¹School of Health and Related Research (SchARR), University of Sheffield, Sheffield, UK

²Consultant Orthopaedic Surgeon and Honorary Senior Lecturer, University of Sheffield, Sheffield, UK

*Corresponding author



Executive summary

Health Technology Assessment 2011; Vol. 15: No. 36
DOI: 10.3310/hta15360

Health Technology Assessment
NIHR HTA programme
www.hta.ac.uk



Executive summary

Background

Hip fracture is a common problem in people aged ≥ 60 years. The annual rate of hip fracture in women in the UK has been reported to be exponentially distributed and to be 20 per 10,000, 38 per 10,000 and 73 per 10,000 at 65, 70 and 75 years of age, respectively. Only 5% of fractures occur in men and women under the age of 60 years. Owing to increasingly ageing populations, the absolute number of hip fractures is expected to rise. Half of all hip fractures are displaced intracapsular fractures, i.e. unstable fractures in which the blood supply to the femoral head may be impaired, affecting the rate of fracture healing. The treatment for displaced intracapsular fractures is currently determined by the mobility and functional demands of the patient. There is no consensus regarding the optimal treatment for individuals who are cognitively intact and have high pre-fracture mobility or function: the two options are hemiarthroplasty (HA) or total hip arthroplasty (THA).

The principal outcomes associated with hip arthroplasty are dislocation, revision rates and quality of life. THA is particularly associated with higher rates of dislocation, whereas HA is particularly associated with pain, infection, loosening of the joint and acetabular erosion. Postoperative complications such as loosening and acetabular erosion can necessitate revision surgery. Revision rates may therefore be higher for HA than for THA.

Objectives

The purpose of this report is to assess the clinical effectiveness and cost-effectiveness evidence of THA compared with HA in patients with displaced intracapsular fracture who are cognitively intact and have high pre-fracture mobility or function.

Methods

A systematic review of the evidence for the clinical effectiveness and cost-effectiveness of THA compared with HA was performed. The primary outcomes of interest were dislocation, revision and reoperation rates. An information specialist made a systematic search of 11 databases of published and unpublished literature from their inception to December 2010. There was no restriction by language, date or study design. Two reviewers screened all titles and abstracts of the citations retrieved by the search to identify both clinical effectiveness and cost-effectiveness studies that satisfied the inclusion criteria, and extracted relevant data from all included studies. The references of all included studies were also checked for further relevant citations. Additionally, exploratory modelling was conducted using the differential costs and quality of life associated with THA compared with HA that were reported in a direct head-to-head randomised controlled trial (RCT) with 2-year follow-up.

Results

A single literature search was conducted for both clinical effectiveness and cost-effectiveness reviews and identified 532 unique citations. Fourteen citations satisfied the inclusion criteria

for the clinical effectiveness review. This represented eight separate trials with 972 participants. Meta-analysis of the six trials found a near significant increased risk of dislocation within 1 year for THA compared with HA [relative risk (RR) 3.98, 95% confidence interval (CI) 0.98 to 16.12, $p=0.05$], but meta-analysis of seven trials found a statistically significant increased risk of dislocation for patients treated with THA (RR 2.40, 95% CI 1.41 to 2.76, $p=0.01$) for all follow-up periods up to 13 years. Meta-analysis of five trials found a statistically non-significant 59% reduced risk of revision within 1 year for THA compared with HA (RR 0.41, 95% CI 0.16 to 1.03, $p=0.06$), but meta-analysis of seven trials found a statistically significant 69% reduced risk of revision for patients treated with THA compared with HA (RR 0.31, 95% CI 0.17 to 0.59, $p=0.0003$) for all follow-up periods up to 13 years.

Meta-analyses of the five and seven trials, respectively, found a statistically non-significant increased risk of any surgery (reduction of dislocations, revisions and all other surgical interventions) both within 1 year and for all follow-up periods for THA compared with HA ($p=0.46$ and 0.75 , respectively). Meta-analyses of five and seven trials, respectively, found a statistically non-significant 9% reduced risk of mortality within 1 year, and a non-significant 4% increased risk of mortality for all follow-up periods, for THA compared with HA ($p=0.60$ and 0.81 , respectively).

Independent subgroup analyses also indicate that study quality, the surgical approach taken (lateral or posterior), the use of cement and the use of unipolar or bipolar prostheses in HA are not statistically significant confounding variables affecting any of these outcomes, when comparing the data on THA and HA reported for the RCTs identified for this review.

Five studies reported Harris Hip Score (HHS). Two studies reported a statistically significant ($p<0.05$) difference after 1 or 2 years in favour of THA, and the three other studies reported the average HHS for study survivors at all follow-up points to be higher (i.e. better) for individuals receiving THA than for those receiving HA. The three remaining studies also reported hip scores using different scales: two studies reported statistically significant differences in favour of THA compared with HA, one after 2 years and one after 3 years, and the third reported that individuals receiving THA reported less pain and better ambulation than those receiving HA. The only statistically significant differences between groups for peri- and postoperative adverse events or complications reported by any study were higher numbers of patients receiving blood transfusion for THA than for HA in one study and higher percentages of patients experiencing acetabular erosion or loosening for HA than for THA in two studies.

Three papers were found that reported the cost-effectiveness of THA compared with HA, although they performed only a cost-utility analysis. An additional paper reported the usage of resources and patient utility recorded in an RCT. The conclusion from the cost-utility analyses was that THA was more cost-effective than HA with an expected 1.53 quality-adjusted life-years (QALYs) being provided at a cost of US\$3000. The cost per QALY ratio of US\$1960 would be viewed as extremely cost-effective using standard UK cost-effectiveness thresholds. A further estimate of the cost-effectiveness of THA compared with HA was also calculated by the authors of this report using data from a published trial which had a follow-up period of 2 years. Even when the utility benefits are constrained to this 2-year horizon, the cost per QALY is $<£25,000$. When the time horizon is extrapolated to more realistic values, the cost per QALY decreases, reaching a value $<£10,000$ with a horizon of only 5 years. This value would be seen as cost-effective under current cost-effectiveness thresholds. Furthermore, longer-term consequences, such as the likely reduced rates of revision associated with THA compared with HA, have not been incorporated in the model. Therefore, the results presented are likely to be unfavourable to THA and the cost-effectiveness of THA is likely to be better than reported.

Discussion

This review conducted a comprehensive and sensitive search for relevant evidence and identified eight RCTs, as well as three ongoing studies. The evidence from the eight relevant RCTs identified indicates that the risk of dislocation is significantly increased for those patients treated with THA than for those with HA, and that the risk of revision is significantly reduced for those treated with THA compared with HA. Patients treated with THA are also more likely to report better function and mobility and less pain than those treated with HA. There are no significant differences in terms of other effectiveness or safety outcomes.

Exploratory modelling was undertaken that showed that THA is likely to be cost-effective compared with HA even when the limitations of the data and methodology are considered. The exploratory model did not consider future revisions or dislocations or differential mortality rates; however, these omissions are expected to strengthen the conclusion that THA is more cost-effective than HA.

Conclusions

Meta-analysis of eight RCTs indicates that THA is more effective than HA in terms of rates of revision, and also more effective in terms of function, pain and mobility, but less effective than HA in terms of rates of dislocation. THA appears to be more cost-effective than HA. It is likely that THA will be associated with increased costs in the initial 2-year period, but the longer-term costs, due to potentially lower revision rates associated with THA, have not been estimated. The capacity and experience of surgeons to perform THA have not been explored and these would need to be addressed at local level were THA to become recommended for active, elderly patients in whom THA is not contraindicated.

Funding

The National Institute for Health Research Health Technology Assessment programme.

Publication

Carroll C, Stevenson M, Scope A, Evans P, Buckley S. Hemiarthroplasty and total hip arthroplasty for treating primary intracapsular fracture of the hip: a systematic review and cost-effectiveness analysis. *Health Technol Assess* 2011;**15**(36).



How to obtain copies of this and other HTA programme reports

An electronic version of this title, in Adobe Acrobat format, is available for downloading free of charge for personal use from the HTA website (www.hta.ac.uk). A fully searchable DVD is also available (see below).

Printed copies of HTA journal series issues cost £20 each (post and packing free in the UK) to both public **and** private sector purchasers from our despatch agents.

Non-UK purchasers will have to pay a small fee for post and packing. For European countries the cost is £2 per issue and for the rest of the world £3 per issue.

How to order:

- fax (with **credit card details**)
- post (with **credit card details** or **cheque**)
- phone during office hours (**credit card** only).

Additionally the HTA website allows you to either print out your order or download a blank order form.

Contact details are as follows:

Synergie UK (HTA Department)
Digital House, The Loddon Centre
Wade Road
Basingstoke
Hants RG24 8QW

Email: orders@hta.ac.uk
Tel: 0845 812 4000 – ask for 'HTA Payment Services'
(out-of-hours answer-phone service)
Fax: 0845 812 4001 – put 'HTA Order' on the fax header

Payment methods

Paying by cheque

If you pay by cheque, the cheque must be in **pounds sterling**, made payable to *University of Southampton* and drawn on a bank with a UK address.

Paying by credit card

You can order using your credit card by phone, fax or post.

Subscriptions

NHS libraries can subscribe free of charge. Public libraries can subscribe at a reduced cost of £100 for each volume (normally comprising 40–50 titles). The commercial subscription rate is £400 per volume (addresses within the UK) and £600 per volume (addresses outside the UK). Please see our website for details. Subscriptions can be purchased only for the current or forthcoming volume.

How do I get a copy of HTA on DVD?

Please use the form on the HTA website (www.hta.ac.uk/htacd/index.shtml). *HTA on DVD* is currently free of charge worldwide.

The website also provides information about the HTA programme and lists the membership of the various committees.

NIHR Health Technology Assessment programme

The Health Technology Assessment (HTA) programme, part of the National Institute for Health Research (NIHR), was set up in 1993. It produces high-quality research information on the effectiveness, costs and broader impact of health technologies for those who use, manage and provide care in the NHS. 'Health technologies' are broadly defined as all interventions used to promote health, prevent and treat disease, and improve rehabilitation and long-term care.

The research findings from the HTA programme directly influence decision-making bodies such as the National Institute for Health and Clinical Excellence (NICE) and the National Screening Committee (NSC). HTA findings also help to improve the quality of clinical practice in the NHS indirectly in that they form a key component of the 'National Knowledge Service'.

The HTA programme is needs led in that it fills gaps in the evidence needed by the NHS. There are three routes to the start of projects.

First is the commissioned route. Suggestions for research are actively sought from people working in the NHS, from the public and consumer groups and from professional bodies such as royal colleges and NHS trusts. These suggestions are carefully prioritised by panels of independent experts (including NHS service users). The HTA programme then commissions the research by competitive tender.

Second, the HTA programme provides grants for clinical trials for researchers who identify research questions. These are assessed for importance to patients and the NHS, and scientific rigour.

Third, through its Technology Assessment Report (TAR) call-off contract, the HTA programme commissions bespoke reports, principally for NICE, but also for other policy-makers. TARs bring together evidence on the value of specific technologies.

Some HTA research projects, including TARs, may take only months, others need several years. They can cost from as little as £40,000 to over £1 million, and may involve synthesising existing evidence, undertaking a trial, or other research collecting new data to answer a research problem.

The final reports from HTA projects are peer reviewed by a number of independent expert referees before publication in the widely read journal series *Health Technology Assessment*.

Criteria for inclusion in the HTA journal series

Reports are published in the HTA journal series if (1) they have resulted from work for the HTA programme, and (2) they are of a sufficiently high scientific quality as assessed by the referees and editors.

Reviews in *Health Technology Assessment* are termed 'systematic' when the account of the search, appraisal and synthesis methods (to minimise biases and random errors) would, in theory, permit the replication of the review by others.

The research reported in this issue of the journal was commissioned by the HTA programme as project number 09/108/02. The contractual start date was in September 2010. The draft report began editorial review in March 2011 and was accepted for publication in May 2011. As the funder, by devising a commissioning brief, the HTA programme specified the research question and study design. The authors have been wholly responsible for all data collection, analysis and interpretation, and for writing up their work. The HTA editors and publisher have tried to ensure the accuracy of the authors' report and would like to thank the referees for their constructive comments on the draft document. However, they do not accept liability for damages or losses arising from material published in this report.

The views expressed in this publication are those of the authors and not necessarily those of the HTA programme or the Department of Health.

Editor-in-Chief: Professor Tom Walley CBE
Series Editors: Dr Martin Ashton-Key, Professor Aileen Clarke, Dr Tom Marshall, Professor John Powell, Dr Rob Riemsma and Professor Ken Stein
Associate Editor: Dr Peter Davidson
Editorial Contact: edit@southampton.ac.uk

ISSN 1366-5278 (Print)

ISSN 2046-4924 (Online)

ISSN 2046-4932 (DVD)

© Queen's Printer and Controller of HMSO 2011. This work was produced by Carroll *et al.* under the terms of a commissioning contract issued by the Secretary of State for Health.

This journal is a member of and subscribes to the principles of the Committee on Publication Ethics (COPE) (<http://www.publicationethics.org/>).

This journal may be freely reproduced for the purposes of private research and study and may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NETSCC, Health Technology Assessment, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

Published by Prepress Projects Ltd, Perth, Scotland (www.prepress-projects.co.uk), on behalf of NETSCC, HTA.

Printed on acid-free paper in the UK by the Charlesworth Group.