The advantages of frameless stereotactic biopsy over frame-based biopsy

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Record Status
This is a critical abstract of an economic evaluation that meets the criteria for inclusion on NHS EED. Each abstract contains a brief summary of the methods, the results and conclusions followed by a detailed critical assessment on the reliability of the study and the conclusions drawn.

Health technology
The use of frameless stereotactic biopsy was investigated. The technique consisted of a neuronavigation system, based on an infrared LED-based system (Easy Guide Neuro, Philips Medical Systems), and a novel stereotactic guide that was adaptable to a variety of instruments. The developed guide was freely adjustable to reach the entire cranium and would lock in place rigidly, allowing fine correction of the trajectory setting. The guide arm was fixed to a Mayfield clamp and the joints of the arm were locked simultaneously without producing tensional movement. This technique was compared with frame-based stereotactic biopsy. The stereotactic frames employed were the Codman-Roberts-Wells (Radioing) and the Resell (Elektra Instruments) systems. All biopsies were performed under general anaesthetic, induced prior to application of the base ring.

Type of intervention
Diagnosis.

Economic study type
Cost-effectiveness analysis.

Study population
The study population comprised samples of biopsies performed in patients who had undergone stereotactic biopsies.

Setting
The setting was secondary care. The economic study was carried out in London, UK.

Dates to which data relate
The effectiveness analysis was carried out between September 1996 and April 1999. The dates to which the resource use data related were not stated. The price year was not stated.

Source of effectiveness data
The effectiveness data were derived from a single study.

Link between effectiveness and cost data
The costing was undertaken prospectively on the same patient sample as that used in the effectiveness study.

Study sample
No study sample size seems to have been determined in the planning phase of the study. In addition, no power calculations were performed retrospectively. A total of 155 stereotactic biopsies were performed, of which 79 were undertaken with a stereotactic frame and 76 with a frameless biopsy. The selection of the technique for each case was
determined by the availability of the image guidance system (IGS) and the IGS research fellow. The frame-based group comprised 49 male and 30 female patients, and the mean age was 52.8 (+/- 15.9) years. The imaging modality employed to target the biopsy was computed tomography (CT) in 89% and magnetic resonance imaging (MRI) in 11% of the frame-based procedures. The frameless group comprised 42 male and 34 female patients, and the mean age was 54.9 (+/- 14.7) years. The imaging modality employed to target the biopsy was CT in 32% and MRI in 68% of the frameless procedures.

**Study design**
The study was a prospective cohort study that was carried out at two departments of neurosurgery (Royal Free Hospital and National Hospital for Neurology and Neurosurgery, London). The duration of follow-up was not stated, but it would appear that the patients were followed up until they were discharged from hospital. The authors did not record any loss to follow-up.

**Analysis of effectiveness**
All of the patients included in the study were accounted for in the analysis. The primary health outcomes used in the analysis were the radiological characteristics of the lesions, pathological diagnosis, the duration of the operations, and complications. The maximum dimension of each lesion was measured, the depth of the lesion below the nearest skin surface ascertained and the lesion classified according to the lobe(s) involved, side, eloquence and brain structure(s) affected. The duration of the anaesthetic was obtained from the anaesthetic records and the duration of surgery from the operating room computer log. Intra-operative complications were obtained from the ward record and pathology service records. There were no significant differences between the two groups in terms of their demographics, lesion site, size or pathologies.

**Effectiveness results**
Radiological characteristics of the lesions.

The distribution, dimensions and locations of the lesions were similar for the frame-based and frameless series. However, the depth of the lesions below the nearest skin surface was found to be significantly greater in the frame-based series (47.2 +/- 20.6 mm) than in the frameless series of cases (35.2 +/- 18.6 mm), (p<0.0001). No difference between the two groups was found in the frequency of lesions within eloquent regions.

Pathological diagnosis.

There were no significant differences in the proportions of pathologies between the frame-based and frameless series. The final pathological diagnosis was classified as high-grade gliomas (60% for the frame-based series versus 68% for the frameless series), low-grade gliomas (16% versus 8%), other tumours (13% versus 16%), inflammatory conditions (6% versus 7%) and non-diagnosed (5% versus 1%). The correct diagnosis was reached on smear examination in 130 (87%) of the 150 diagnostic specimens. These rates were similar for both groups.

Duration of operations.

For the frameless cases, the mean duration of the operation and anaesthetic was 54.2 (+/- 23.6) minutes. For the frame-based procedures, the mean duration of anaesthesia was 127.4 (+/- 31.9) minutes, whilst the mean duration of the operation was 56.3 (+/- 17.5) minutes. Thus, the mean duration of the anaesthetic was significantly longer in the frame-based series than in the frameless series, (p<0.0001). However, no significant differences were found in the duration of surgery between the two groups.

Complications.

The overall complication rate was significantly lower for the frameless series (14%) than for the frame-based series (22%), (p=0.0183). The rates of surgical complications were 6.6% (frameless) and 8.8% (frame-based), respectively. There was one fatality in each series. There were significantly more haemorrhagic complications (5 versus 0) and significantly more permanent neurological deficit (3 versus 1) in the frame-based series than in the frameless series.
(p=0.0008). There were also more infective complications in the frame-based series (10 versus 1).

**Clinical conclusions**
It can be concluded that frameless stereotactic biopsy resulted in a reduced anaesthetic time and fewer complications when compared with traditional frame-based stereotactic biopsy.

**Measure of benefits used in the economic analysis**
No summary health benefit measure was used in the economic analysis. Thus, the study has been classified as a cost-consequences analysis.

**Direct costs**
The resource use and the costs were reported separately. The direct costs included in the analysis were those of the hospital. These were operative procedures, ITU stay, ward stay, CT scans, MRI scans and capital costs. The cost of inpatient stay, day, surgery and imaging studies were obtained from the hospital finance department. The capital costs per case were derived for each technique by determination of the purchase costs, annual service charges, equipment lifetime and number of cases per annum. Discounting was not relevant, as the costs were incurred during a short time, and was not conducted. The study reported the average costs. The price year was not reported.

**Statistical analysis of costs**
The resource use and costs were treated stochastically. The statistical tests of significance employed were unpaired two-tailed t-tests (significance established at p<0.05).

**Indirect Costs**
The indirect costs were not included in the analysis.

**Currency**
UK pounds sterling ().
time, lower cost and fewer complications when compared with traditional frame-based stereotactic biopsy.

**CRD COMMENTARY - Selection of comparators**
The comparator, frame-based stereotactic biopsy, was justified on the grounds that it was the current 'gold' standard technique for the retrieval of histological specimens from targets within the brain. You should decide if this represents a widely used health technology in your own setting.

**Validity of estimate of measure of effectiveness**
The study used a prospective cohort study, which was appropriate for the study question. However, the internal validity of the study would have been enhanced had a randomised controlled trial been used, given that this is the 'gold' standard study design when comparing different health technologies, as it minimises the potential of both bias and confounding. The study sample was representative of the study population. Both patient groups were shown to be comparable at analysis in terms of the demographics, lesion site, size, and pathologies. The analysis of effectiveness seems to have been handled credibly. However, the authors pointed out that there remained the possibility that the high rate of adverse events in the frame-based series was related to the relative inexperience of the operators (as frameless biopsy was the preferred technique in the authors' setting), whereas frameless biopsy may have been more closely supervised. The authors employed appropriate statistical tests of significance to test for any differences between the two groups.

**Validity of estimate of measure of benefit**
The authors did not derive a summary measure of health benefit. The analysis was therefore categorised as a cost-consequences study.

**Validity of estimate of costs**
The perspective adopted in the economic analysis was not explicitly stated, but it would appear to have been that of the hospital. All the relevant categories of cost for this perspective, and all the relevant costs in each category, were included in the analysis. The costs and the quantities were reported separately, which will enhance the generalisability to other settings. Resource use was obtained from the authors' setting and statistical tests of significance were performed. No further analysis of the quantities was conducted. No statistical analysis of the prices was performed. Thus, any uncertainty surrounding these variables could not be taken into consideration. The authors did not perform any discounting since all the costs were incurred during a short time. The price year was not reported, making the results of any reflationary exercises questionable.

**Other issues**
The authors made appropriate comparisons of their findings with those from other studies that had found similar results. The issue of generalisability to other settings was addressed, and was further enhanced by the authors reporting the costs and the quantities separately. The authors do not appear to have presented their results selectively and their conclusions reflected the scope of the analysis. The authors reported no further limitations of their study.

**Implications of the study**
Based on their results, the authors appear to recommend the use of frameless stereotactic biopsy over frame-based techniques. The authors pointed out that the term "frameless stereotaxy" should only be applied to methods of image guidance where the instruments are directed with stereotactic precision to a preselected target by a navigation system, and not to hand-held methods.

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Other publications of related interest


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MeSH
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