

Evidence based general practice: a retrospective study of interventions in one training practice

P Gill, A C Dowell, R D Neal, N Smith, P Heywood, A E Wilson

Abstract

Objectives—To estimate the proportion of interventions in general practice that are based on evidence from clinical trials and to assess the appropriateness of such an evaluation.

Design—Retrospective review of case notes.

Setting—One suburban training general practice.

Subjects—122 consecutive doctor-patient consultations over two days.

Main outcome measures—Proportions of interventions based on randomised controlled trials (from literature search with Medline, pharmaceutical databases, and standard textbooks), on convincing non-experimental evidence, and without substantial evidence.

Results—21 of the 122 consultations recorded were excluded due to insufficient data; 31 of the interventions were based on randomised controlled trial evidence and 51 based on convincing non-experimental evidence. Hence 82/101 (81%) of interventions were based on evidence meeting our criteria.

Conclusions—Most interventions within general practice are based on evidence from clinical trials, but the methods used in such trials may not be the most appropriate to apply to this setting.

Introduction

The recent enthusiasm for developing evidence based medical practice has been the subject of debate.¹

The establishment of the Cochrane Collaboration² and the publication of a new journal, *Evidence-Based Medicine*,³ highlight the fact that the research and scientific basis of medicine is currently subject to close scrutiny.

Last year the *Lancet* published a paper⁴ that challenged previously held beliefs that less than 20% of medical practice was based on scientific evidence.⁵ The authors assessed the evidence base for the treatment of 109 medical patients in hospital. Their findings, that up to 80% of acute hospital interventions had a scientific rationale, produced much comment⁶⁻⁸ and a challenge from the authors to repeat the study in other clinical settings. By applying the same methodology we investigated the degree to which general practice is evidence based.

Methods

Consecutive consultations over two days were reviewed by retrospective analysis of case notes from a suburban training general practice. For each consultation, two of the authors independently recorded the primary diagnosis and intervention before reaching consensus. The primary diagnosis was defined as the first problem recorded for the consultation and the primary intervention as "the treatment or manoeuvre that represented the practitioner's attempt to cure, alleviate, or care for the patient in respect of the primary diagnosis."⁴ The evidence for the interventions was then searched for in Medline (1966-95), standard textbooks, and pharmaceutical companies' databases.

We classified the interventions as did Ellis *et al*: (i) intervention based on evidence from randomised controlled trial; (ii) intervention based on convincing non-experimental evidence; (iii) intervention without substantial evidence, not meeting criterion (i) or (ii). To assess non-randomised controlled trial interventions we held a consensus meeting of our academic team of five general practitioners and a non-medical arbiter. Only interventions with unanimous consensus of the team were allocated to either groups (ii) or (iii).

Results

Of the 122 consultations recorded, 21 were excluded as the patients were referred (six patients) or sent for investigations (five patients) to hospitals; the remaining 10 patients had insufficient data, leaving a study sample of 101 diagnosis-intervention pairs.

Primary interventions were classified "evidence based" if they fulfilled the criteria for category (i) or (ii), with the result that 81% of patients (82/101) had received evidence based interventions (tables 1 and 2).⁹⁻³³ The remaining 19% (table 3) were judged to have received treatment that had no substantial evidence from our search.

Centre for Research in Primary Care, Leeds University, Leeds LS2 9LN

P Gill, research tutor

A C Dowell, director

R D Neal, research fellow

N Smith, research fellow

P Heywood, deputy director

A E Wilson, lecturer

Correspondence to: Dr Dowell.

BMJ 1996;312:819-21

Table 1—Interventions (n=31) substantiated by evidence from randomised controlled trials

Primary diagnosis	Primary treatment	No of patients	Reference No
Hypertension	Bendrofluazide	2	9
	Metoprolol	1	10
	Atenolol	1	11
	Captopril	1	12
	Nifedipine	1	13
	Hydralazine	1	14
Asthma	Salmeterol	1	15
	Beclomethasone	2	16
Bronchospasm	Salbutamol	1	17
Depression	Fluoxetine	1	18
Anxiety	Diazepam	1	19
Insomnia	Temazepam	1	20
Urinary tract infection	Trimethoprim	3	21
	Cephadrine	2	22
Chest infection	Amoxycillin	1	23
Conjunctivitis	Chloramphenicol	1	24
Athlete's foot	Clotrimazole	1	25
Fungal skin infection	Terbinafine	1	26
Thrush (cutaneous)	Miconazole	1	27
Painful shoulder	Triamcinolone	2	28
Low back pain	Exercise	1	29
Acne	Minocycline	1	30
Eczema	Hydrocortisone	1	31
Hay fever	Terfenidine	1	32
Allergic conjunctivitis	Sodium cromoglycate	1	33

Table 2—Interventions (n=51) substantiated by convincing non-experimental evidence

Primary diagnosis	Primary treatment	No of patients
Hypercholesterolaemia	Supervision of weight	1
Valvular heart disease	Warfarin	1
Asthma	Prednisolone	1
Depression	Counselling	2
Migraine	Migril	1
Anxiety	Reassurance	1
Worries about gastric cancer	Counselling	1
Cellulitis	Co-fluampicil	1
Soft tissue infection	Co-fluampicil	1
Infected eczema	Erythromycin	1
Boil in ear	Flucloxacillin	1
Soft tissue infection	Flucloxacillin	1
Throat infection	Penicillin V	2
Tonsillitis	Penicillin V	1
Chest infection	Penicillin V	1
	Erythromycin	1
Impetigo	Fucidic acid	1
Oral thrush	Nystatin	1
Chest infection resolving	Nil	2
Back pain	Dihydrocodeine	1
Low back pain	Ibuprofen	1
	Mefenamic acid	1
Resolving foot pain	Nil	1
Sprained toe	Nil	1
Skin tag	Cryotherapy	1
Allergic rash	Chlorpheniramine	1
Gastritis	Gaviscon	1
Oesophagitis	Gaviscon	1
Diarrhoea	Rehydration	1
Viral gastroenteritis	Fluids	1
Otitis externa	Gentamycin/hydrocortisone	1
	Aural toilet	1
Ear wax	Syringing	2
Herpes zoster	Codeine phosphate	1
Viral illness	Paracetamol	1
	Nil	1
Hypothyroidism	Thyroxine	2
Contraception	Oral contraceptive pill	4
Menopause	Hormone replacement therapy	1
Pregnant 20 weeks	Antenatal check	1
Abscess	Incision and drainage	1
Insect bites	Terfenadine	1
Infected bite	Incision and drainage	1

Table 3—Interventions (n=19) without substantial evidence

Primary diagnosis	Primary treatment	No of patients
Cerebrovascular accident	Dipyridole	1
Headache	Carbamazepine	1
Paronychia	Erythromycin	1
Laryngitis	Oxytetracycline	1
Fungal skin infection	E45 cream	1
Viral illness	Pseudoephedrine	1
Tennis elbow	Fenbufen	1
Varicose eczema	Providine iodine	1
	Unguentum Merck	1
Traumatic skin ulcers	Hydrocortisone	1
Warts	Cryotherapy	1
	Curettage	1
Heat rash	Cetirizine	1
	Chlorpheniramine	1
Chronic abdominal pain	Cisapride	1
Bladder spasm	Indoramin	1
Vaginal tear	Perineal massage	1
Cervical polyp	Nil	1
Ankle swelling	Bendrofluazide	1

Discussion

This pilot study has shown that the majority of interventions within general practice are based on evidence. This is comparable with the findings of the study set in an acute unit in Oxford.⁴

Our study has some limitations. As it is a retrospective study within one training practice, the results cannot be generalised. By limiting our search to databases such as Medline, which we acknowledge are not comprehensive,³⁴ we may have failed to find all the

evidence available. Neither did we make any attempt to assess the methodological quality of the trials identified. Nevertheless, we believe that our study raises some points that are worthy of debate.

PRIMARY DIAGNOSES AND PRIMARY INTERVENTIONS

General practice is characterised by patients presenting with multiple and ill defined problems; a specific diagnosis may not be reached within a single consultation. We accepted the primary definition of the clinical problem by the general practitioner at face value. In using the primary diagnosis as denominator we have not only reduced the complexity of general practice but also lost some of its reality.⁶ We recognise that it may be difficult to allocate a specific diagnosis to a symptom, such as "painful tongue," in the same way as, in a hospital setting, "non-cardiac chest pain" may have various causes.

General practice consultations may be triggered by a variety of circumstances (certification or external pressure, for example), whereas a more critical, often clinical, event usually precipitates a hospital admission. Clinical problems have many facets, hence diagnoses and interventions are often multiple, particularly when physical, psychological, and social elements are considered. The diagnosis-treatment pair "depression/counselling" can tell only a fraction of the story of a complex interaction. There are patients in whom the disease is neither clear nor relevant to the patient's problem,³⁵ and the presence of any disease is not always proved by investigations. Also, there is pressure to record a medical diagnosis to justify treatment.³⁶ On the other hand, secondary diagnoses and social problems may not be recorded.³⁷ Nevertheless, we agree with Bridges-Webb *et al* that doctors' diagnoses remain relevant, if not absolutely valid, since doctors are likely to base their recommendations on the labels they report.³⁸ This difficulty in separating out primary events is not unique to general practice, and in all specialties assigning an appropriate diagnostic label or labels this must be considered as an integral part of evidence based practice.

ASSESSMENT OF RANDOMISED CONTROLLED TRIALS

The study raises several questions concerning the appropriateness and quality of randomised control trial evidence. Firstly, randomised controlled trials may not necessarily indicate the most cost effective current treatment for general practice. For example, use of a third generation cephalosporin may be substantiated by a randomised controlled trial for a urinary tract infection, but in an uncomplicated case trimethoprim may be just as effective—and certainly cheaper.

Further questions that arose during our literature search concern the issue of how endpoints for randomised controlled trials can be measured. There is clear evidence that angiotensin converting enzyme inhibitors and calcium channel blockers reduce blood pressure, but none that they reduce cardiovascular morbidity and mortality. Should this be regarded as "good" evidence? How should we deal with randomised controlled trials that are apparently outdated by evidence that another treatment is available (for example, oral ketoconazole rather than topical clotrimazole for candida albicans), or evidence from other sources suggesting later that a treatment can be harmful (temazepam capsules, for example)? Do randomised controlled trials have to be compared against placebo, or is it acceptable to compare with the currently accepted "standard" treatment, even though this may not have previously been tested against placebo in a randomised controlled trial?

Evidence based practice has to accept the possibility that evidence from randomised controlled trials has not necessarily the value of a "gold standard" but has more

Key messages

- Standard definitions of diagnosis and interventions in general practice are unclear
- 81% of general practice can be described as evidence based using this method of assessment
- Evidence derived from different methodologies may be important for the assessment of the evidence base of general practice

the value of a coffee future—likely to be altered by tomorrow's experience. Furthermore, some interventions were originally assessed within secondary care but their main use is in the community.

THE SEARCH FOR APPROPRIATE PARADIGMS

Despite the healthy debate resulting from the Oxford study,⁶⁻⁸ both patients and policy makers might want doctors to try to base as many of their interventions as possible on evidence from clinical trials. There may be a temptation to produce a league table of specialties and settings—for example, inpatient medicine might be found to be “better than” general practice by 1%.

We could question whether it is feasible or even desirable to pursue the goal of 100% evidence based practice. Much of the work within general practice, as well as in other settings, consists of medicine that combines science with art, sociology, mythology, and pastoral care. These aspects of care must be incorporated into an appropriate paradigm of evidence based practice rather than that determined solely by clinical trials.

Linked to this is the search for appropriateness of the methods used to provide the evidence. We believe that for general practice, and possibly in other settings too, the most important evidence may be found in developing alternative methodologies which complement conclusions from randomised controlled trials.

We thank staff at The Street Lane Practice for their help.
Funding: None.
Conflict of interest: None.

- 1 Smith R. The scientific basis of health services. *BMJ* 1995;311:961-2.
- 2 Cochrane's legacy. *Lancet* 1992;340:1131-2.
- 3 Davidoff F, Haynes B, Sackett D, Smith R. Evidence based medicine. *BMJ* 1995;310:1085-6.
- 4 Ellis J, Mulligan I, Rowe J, Sackett DL. Inpatient general medicine is evidence based. *Lancet* 1995;346:407-10.
- 5 Office of Technology Assessment of the Congress of the United States. *Assessing the efficacy and safety of medical technologies*. Washington, DC: US Government Printing Office, 1978.
- 6 Bradley F, Field J. Evidence-based medicine. *Lancet* 1995;346:838.
- 7 Aveyard P. Evidence-based medicine. *Lancet* 1995;346:838.
- 8 Evidence-based medicine in its place [editorial]. *Lancet* 1995;346:785.

- 9 Medical Research Council Working Party. MRC trial of treatment of mild hypertension: principal results. *BMJ* 1985;291:97-104.
- 10 Morley CA, Cavalcanti C, Perrins EJ, Sutton R. A comparison of once daily atenolol and metoprolol SA in mild to moderate hypertension. *Br J Clin Pharmacol* 1983;15:715-7.
- 11 Hansson L, Aberg H, Karlberg BE, Westerlund A. Controlled study of atenolol in treatment of hypertension. *BMJ* 1975;ii:367-70.
- 12 Drayer JJ, Weber MA. Monotherapy of essential hypertension with a converting-enzyme inhibitor. *Hypertension* 1983;5:108-13.
- 13 Hallin L, Andren L, Hansson L. Controlled trial of nifedipine and bendroflumethiazide in hypertension. *Journal of Cardiovasc Pharmacol* 1983;5:1083-5.
- 14 Silas JH, Ramsay LE, Freestone S. Hydralazine once daily in hypertension. *BMJ* 1982;284:1602-4.
- 15 Fitzpatrick MF, Mackay T, Driver H, Douglas NJ. Salmeterol in nocturnal asthma: a double blind, placebo controlled trial of a long acting inhaled β 2 agonist. *BMJ* 1990;301:1365-8.
- 16 Agnew RA, Walker DJ, Phillips LA. A prospective study of respiratory infection in asthmatic patients treated with beclomethasone dipropionate and sodium cromoglycate. *Clin Allergy* 1977;7:183-8.
- 17 Hasham F, Kennedy JD, Jones RS. Actions of salbutamol, disodium cromoglycate, and placebo administered as aerosols in acute asthma. *Arch Dis Child* 1981;56:722-5.
- 18 Come SJ, Hall JR. A double-blind comparative study of fluoxetine and dothiepin in the treatment of depression in general practice. *Int Clin Psychopharmacol* 1989;4:245-54.
- 19 Rickels K, Csanalosi I, Greisman P, Mirman MJ, Morris RJ, Weise CC, et al. Ketazolam and diazepam in anxiety: a controlled study. *J Clin Pharmacol* 1980;20:581-9.
- 20 Wheatley D. Insomnia in general practice: the role of temazepam and a comparison with zopiclone. *Acta Psychiatr Scand* 1986;332(suppl):142-8.
- 21 Lacey RW, Lord VL, Gunasekera HK, Leiberman PJ, Luxton DE. Comparison of trimethoprim alone with trimethoprim sulphamethoxazole in the treatment of respiratory and urinary infections with particular reference to selection of trimethoprim resistance. *Lancet* 1980;ii:1270-3.
- 22 Low RA, Clarke TK. Cephradine in urinary tract infections: a double-blind comparison with ampicillin. *Curr Med Res Opin* 1975;3:211-7.
- 23 Richards JG. Doxycycline and amoxicillin in respiratory infections: a comparative assessment in general practice. *Curr Med Res Opin* 1980;6:393-7.
- 24 Trimethoprim-Polymixin B Sulphate Ophthalmic Ointment Study Group. Trimethoprim-polymixin B sulphate ophthalmic ointment versus chloramphenicol ointment in the treatment of bacterial conjunctivitis—a review of four clinical studies. *J Antimicrob Chemother* 1989;23:261-6.
- 25 Smith EB, Graham JL, Ulrich JA. Topical clotrimazole in tinea pedis. *South Med J* 1977;70:47-8.
- 26 Evans EGV, Shah JM, Joshipura RC. One week treatment of tinea corporis and tinea cruris with terbinafine (Lamisal) 1% cream: a placebo controlled study. *Journal of Dermatological Treatment* 1992;3:181-4.
- 27 Cullin SI. Cutaneous candidiasis: treatment with miconazole nitrate. *Cutis* 1977;19:126-9.
- 28 Petri M, Dobrow R, Neiman R, Whiting-O'Keefe Q, Seaman WE. Randomised double-blind, placebo-controlled study of the treatment of the painful shoulder. *Arthritis Rheum* 1987;30:1040-5.
- 29 Frost H, Klaber Moffett JA, Moser JS, Fairbank JC. Randomised controlled trial for evaluation of fitness programme for patients with chronic low back pain. *BMJ* 1995; 310:151-4.
- 30 Cullen SI, Cohan RH. Minocycline therapy in acne vulgaris. *Cutis* 1976;17:1208-10.
- 31 Brogden RN, Pinder RM, Sawyer PR, Speight TM, Avery GS. Hydrocortisone 17-butyrate: a new topical corticosteroid preliminary report. *Drugs* 1976;12:249-57.
- 32 Ciprandi G, Iudice A, Tosca MA, Ruffoni S, Buscaglia S, Canonica GW. Comparative effects of terfenadine and loratadine in the treatment of hay fever. *J Investigat Allergol Clin Immunol* 1991;1:368-72.
- 33 Ciprandi G, Cerqueti PM, Sacca S, Cilli P, Canonica GW. Levocabastine versus cromolyn sodium in the treatment of pollen-induced conjunctivitis. *Ann Allergy* 1990;65:156-8.
- 34 Dickersin K, Scherer R Lefevre C. Identifying relevant studies for systematic reviews. *BMJ* 1994;309:1286-91.
- 35 Marsland DW, Wood M, Mayo F. *Content of family practice*. New York: Appleton-Century-Crofts, 1980.
- 36 Howie JGR. Diagnosis: the Achilles heel. *J R Coll Gen Pract* 1972;22:310.
- 37 Gelbach S. Comparing methods of data collection in an academic ambulatory practice. *J Med Educ* 1979;54:730-2.
- 38 Bridges-Webb C, Britt H, Miles DA, Neary S, Charles J, Traynor V. Morbidity and treatment in general practice in Australia 1990-1991. *Med J Aust* 1992;157:S1-56.

(Accepted 19 February 1996)

Survey of general practitioners' views of consultants' non-urgent referral of outpatients to other consultants

Royal Sussex County Hospital, Brighton BN2 5BE
S Bridger, medical registrar
S R Cairns, consultant gastroenterologist

S Bridger, S R Cairns

There are an estimated nine million new outpatient referrals each year.¹ Between 60% and 80% of these referrals originate from general practitioners.² Additional referrals arise predominantly from hospital departments. A study in Leicester found a cross

referral rate of at least 17% from one clinic to another.³ In the years preceding fundholding the practice of referral between consultants seems to have been accepted.⁴ However, fundholding general practitioners have become increasingly aware of the cost and logistic implications of this practice and have begun to express their concerns.⁵

We investigated local general practitioners' preferences about the practice of non-urgent referral of outpatients between consultants.

Patients, methods, and results

We sent a questionnaire to all 165 general practitioners in the Brighton area. These questionnaires consisted of 10 specimen case histories, and on each of

Correspondence to:
Dr Cairns.

BMJ 1996;312:821-2