- 1 British Thoracic and Tuberculosis Association. Short course chemotherapy in pulmonary tuberculosis. Lancet 1976;ii:1102-4
- Monie RDM, Hunter J.M., Rocchicoli K, White J, Campbell IA, Kilpatrick GS. Survey of pulmonary tuberculosis in south and west Wales (1976-8). *Br Med J* 1982;284:571-3.
   Medical Research Council Tuberculosis and Chest Diseases Unit. Treatment of pulmonary
- tuberculosis in patients notified in England and Wales in 1978-9: chemotherapy and hospital admission. Thorax 1985:40:113-20.
- Buechner HA. Short course chemotherapy for tuberculosis. Ann Intern Med 1981;94:277-8.
- 5 Buechner HA. Short course chemotherapy for tuberculosis-a story of flawed studies. Am Rev Respir Dis 1981;245:655
- 6 Buechner HA. When is pulmonary tuberculosis cured? Lancet 1982;i:1462
- 7 Anonymous. When is pulmonary tuberculosis cured? [Editorial]. Lancet 1982;i:1163-4.
   8 Citron KM, Angel JH, Somner AR. Short course chemotherapy for tuberculosis: a story of flawed
- studies. Am Rev. Respir Dis 1981;124:658.
  9 Darbyshire J, Fox W, Girling DJ, Nunn AJ, Mitchinson DA. Short course chemotherapy for
- tuberculosis: a story of flawed studies. Am Rev Respir Dis 1981;**124**:658-60. ox W. Compliance of patients and physicians: experience and lessons from tuberculosis—II. Br
- Med 7 1983.287.101-5 11 British Medical Research Council. Co-operative controlled trials of a standard regimen of
- streptomycin, PAS and isoniazid and three alternative regimens of chemotherapy in Britain. *Tubercle* 1973;54:99-129. 12 Gow JG. Management of extra-pulmonary tuberculosis. Br Med 7 1982;285:894-5
- Croîton JW. Treatment of tuberculosis. Br. Med J 1979;i:52.
   Parsons M. Tuberculous meningüis—a handbook for clinicians. Oxford: Oxford University Press,
- 1979 15 Gow JG, Barbos AS. Genitourinary tuberculosis. A study of 1117 cases over a period of 34 years.
- Br J Urol 1984;56:449-55. 16 Byram D, Hatton P, Williams SE, Pearson SB. The form and presentation of tuberculosis over a
- 10 year interval in Leeds. Br J Dis Chest 1985;79:152-60.
   17 Wardman AG, Williams SE, Curzon PGD, Page RL, Cooke NJ. Tuberculosis: who should rescribe? Br Med 7 1982;284:569-71
- 18 Laidlaw M. Renal tuberculosis. In: Chisholm GD, Williams DI, eds. Scientific foundation of workey. 2nd ed. London: William Heinemann Medical Books, 1982:222-7.
   Malkasian GD. Respiratory diseases and pregnancy. In: Iffy L, Kaminetzky HA, eds. *Principles*
- and practice of obstetrics and perinatology. Vol 2. New York: John Wiley and Sons, 1981:1311-4. 20 Barnes J. Female genital tuberculosis. In: Lecture notes on gynaecology. 5th ed. Oxford: Blackwell
- Scientific Publications, 1983:67-70. 21 Griffiths DLL. The treatment of spinal tuberculosis. In: McKibbin B, ed. Recent advances in
- Orintins DLL. The treatment of spinal tuberculosis. In: McNibin B, ed. Recent advances in orthopaedics. Vol 3. Edinburgh: Churchill Livingstone, 1979:1-17.
   Sutherland AM. Gynaecological tuberculosis. Br J Hosp Med 1979;22:569-76.
   Wong SH, Lau WY, Ho KK, Fan ST, Yiu TF, Chan SL. The management of urinary tuberculosis—a logical approach. Br J Urol 1984;56:349-53.
- 24 British Thoracic Society. A controlled trial of 6 months' chemotherapy in pulmonary tuberculosis Final report: results during the 36 months after the end of chemotherapy and beyond. BrJ Dis Chest 1984:78-330-6

## Surgery and the pill

After major surgery the relative risk of overt, clinical, deep venous thrombosis in women taking oestrogen containing combined oral contraceptives is about two, as compared with non-users.<sup>12</sup> This estimate is approximate for many reasons: the risk appeared higher in early studies,<sup>3</sup> whereas the increasing use of formulations containing less than 50 µg oestrogen implies that the real risk of all forms of venous thromboembolism may have come down.4

Metabolic studies show numerous changes in factors associated with blood coagulation and fibrinolysis, whose overall effect, even with our current pills, would be expected to predispose to deep vein thrombosis.<sup>12</sup> Anaesthesia and the combined oral contraceptives have additive effects on at least one important factor (both reduce the activity of antithrombin III).<sup>5</sup> Even if the relative risk with 30-35 µg oestrogen pills is now less than two the attributable risk if women continued to take them during and after major surgery would be high, because venous thromboembolism after major surgery is one of the commonest serious complications of the combined oral contraceptive.<sup>12</sup> The risk is greatest after major orthopaedic, abdominal, and cancer surgery or any surgery that entails giving a hypotensive anaesthetic.

On p 516 there is another risk to which attention is drawn-namely, unwanted pregnancy when the oral contraceptive pill is discontinued before surgery. This is a risk that needs to be clearly explained to the woman and her partner, and the couple should be helped to organise alternative means of contraception. (Many couples who have become used to the convenience of the combined oral contraceptive for several years have no notion of how easy it is to conceive unless barrier methods are used with obsessional care.) Steps must also be taken to ensure that an early pregnancy is not present before proceeding with elective anaesthesia and surgery.

To reduce this risk of pregnancy further three questions need to be answered: For which procedures is it unnecessary to discontinue the combined pill? Do the restrictions apply to the progestogen only pill, or other oestrogen free preparations? and How long before and after surgery should oral contraceptives be avoided?

The risk of deep venous thrombosis after minor surgery, such as most dental procedures and laparoscopy, with a short duration of anaesthesia and full mobilisation the same day, is vanishingly small. Hence any extra risk associated with the combined oral contraceptive would be more than outweighed by the risk of pregnancy. The only exception would be minor procedures to the legs themselves-notably varicose vein surgery and injection sclerotherapy. Metabolic studies have generally failed to show any important effects of progestogens in promoting intravascular coagulation<sup>6</sup>; thus the progestogen only pill need not be avoided over the time of elective surgery, however major. Indeed, the injectable progestogens such as Depo-Provera might well be offered as perioperative "cover" to women on the waiting list for a major procedure.

Finally, the time of avoidance of the combined oral contraceptive should be reduced to the minimum commensurate with safety. Published epidemiological studies suggest that the excess risk of deep venous thrombosis is unrelated to duration of use and reverts to normal within less than one month.<sup>7</sup> Some changes in the coagulation and fibrinolytic systems have been detected about six weeks after discontinuing oestrogens, but others—such as the decreased activity of antithrombin III-have reverted to normal within four weeks.8 Avoiding combined oral contraceptives for six weeks preoperatively may be the counsel of perfection, but four weeks is probably more realistic. Postoperatively most agree that the combined oral contraceptive may be restarted two weeks after the patient is fully mobile.910

Taking all factors into consideration, I believe that the following-which is essentially that which will appear in the forthcoming issue number 10 of the British National Formulary, para 7.3.1-represents good policy: oestrogen containing contraceptives should be discontinued (and adequate alternative contraceptive arrangements made) four weeks before major elective surgery; they should normally be started again at the first menses occurring at least two weeks after the procedure. When discontinuation is not possiblefor example, after trauma or if, by oversight, a patient admitted for an elective procedure is still taking an oestrogen containing oral contraceptive-prophylactic low dose subcutaneous heparin should be considered.5 These recommendations do not apply to minor surgery with short duration of anaesthesia and early mobilisation-for example, laparoscopic sterilisation or tooth extraction-or to women taking oestrogen free contraceptives.

It is important to plan ahead: not only must this matter be discussed in the outpatient department but also women must be given at least a month's warning of their admission or (preferably) a definite date from the clinic. Other risk factors must receive due weight, notably obesity and a family history of venous thrombosis. The latter implies that the patient should be investigated preoperatively for any factors that might either promote thrombosis or impair fibrinolysis. It is also helpful to instruct women to restart the combined oral

contraceptive on the first day of their postoperative menses since this provides immediate contraception.<sup>11</sup>

**JOHN GUILLEBAUD** 

Medical Director Margaret Pyke Centre for Study and Training in Family Planning, London WIV STW

- 1 Stadel BV. Oral contraceptives and cardiovascular disease. N Engl J Med 1981;305:612-8
- Stadel BV. Oral contraceptives and cardiovascular disease. N Engl J Med 1981;305:672-7
- 3 Vessey MP. Female hormones and vascular disease: epidemiologic overview. British Journ Family Planning 1980;6(suppl):1-12. 4 Bottiger LE, Westerholm B. Oral contraceptives and thromboembolic disease: Swedish
- experience. Acta Med Scand 1979:190:455-63 5 Sagar S, Stamatakis JD, Thomas DP, Kakkar VV. Oral contraceptives, anti-thrombin III
- Statiatakis JD, Holnas DF, Katkar VV. Olar Collideepives, and thromon fir activity, and post-operative deep-vein thrombosis. Lancet 1976;i:509-111.
  Fotherby K. Low doses of gestagens as fertility regulating agents. In: Diczfalusy E, ed. Advances in fertility regulation. Copenhagen: Scriptor, 1977:283-312.
  Vessey MP, Doll R, Fairbairn AS, et al. Post-operative thromboembolism and the use of oral contraceptives. Br Med J 1970;iii:123-6.
- von Kaulla E, Droegemueller W, Aoki N, von Kaulla KN. Anti-thrombin III depression and
- thrombin generation acceleration in women taking oral contraceptives. Am J Obstet Gynecol 1971;109:868-73.

9 Anonymous. Any questions? Br Med J 1981;282:1857.
10 Department of Health and Social Security. Handbook of contraceptive practice. London: Department of Health and Social Security, 1984:26, para 2.3.11.

11 Loudon NB. Starting on the pill. Br Med J 1977;ii:521-2.

## Setting standards in general practice

The Royal College of General Practitioners launched its "quality initiative" in 1983,1 and the council of the college has now proposed radical measures that will transform the initiative from a voluntary exercise undertaken by keen college members into a strategy for the whole of general practice (29 June, p 1981).<sup>2</sup> But the consultation document, Towards Quality in General Practice, makes two challengeable assumptions: firstly, that "quality of care" can be described and assessed; and, secondly, that quality of care is synonymous with quality of doctor.<sup>3</sup>

Quality of care has been measured in terms of structures (personnel, buildings, equipment), processes (the consultation, prescribing, screening, what we do), and outcome (changes in individual and community health). Measures of outcome are difficult to obtain, but until valid indicators are developed judgments about quality of care can be made only with reservation. Computerisation, both of family practitioner committees and of practices, is one way of obtaining data for producing measures of outcome. The college says in the report that the computerisation of general practice is essential, but it must emphasise the benefit from such investment and seek earmarked funding in the form of grants to individual practices.

The second assumption-that improving the quality of general practitioners by asking "What sort of doctor?" will result in raised standards of care-leads to the controversial proposals for the membership examination being an entry standard for new principals in National Health Service general practice, "accreditation" of new principals by continuous assessment after two years of "higher training," and "reaccreditation" at intervals, also by continuous peer assessment, linked to remuneration by a new "performance sensitive" contract. These are radical proposals, yet the continuing public criticism of the stereotyped general practitioner ("You feel you're wasting your time talking to him"; "Before you've finished saying 'I've got a sore throat' you've got a prescription and you're out"<sup>4</sup>) and the concern of college examiners at the standard of some candidates mean that there is clearly a need to examine the issues of quality and standards.

The setting of professional standards is an accepted

function of an academic body, and provided its methods are both valid and widely acceptable to doctors and patients any attempt to raise standards of patient care should be welcomed. Although the membership examination has been carefully developed and validated<sup>5-7</sup> it is not without its critics.<sup>8</sup> A better method of checking on the standard of vocational training might be the forms of continuous assessment that the college is developing,<sup>3</sup> combined perhaps with a shorter "part 1" examination.' The membership examination might

patients and colleagues will increasingly expect. It is surprising that a document that looks towards 1990 makes no mention of the Declaration of Alma-Ata, with its patient centred goal of "health for all by 2000."10 Perhaps that highlights the doctor centred approach of this document -"What sort of doctor?" not "What sort of health?"" Primary health care demands participation among doctors, nurses, social workers, lay community workers, and patients. Patients are mentioned in the college document but only as a potential influence on the service who should receive more information.

then be left as a voluntary indicator of achievement, which

The concluding call for additional government resources is at best inopportune-it sits uncomfortably next to the statement: "Personal satisfaction is derived from doing the job of general practice well-it is fostered by the nature of the doctor patient relationship, and is encouraged by the intellectual stimulus and satisfaction of comparing one's own clinical experiences with those of colleagues."

Even so, this document deserves careful reading by all general practitioners and will doubtless generate wide debate beyond the college. While some of the proposals may be seen as too extreme, the underlying principles of objective inquiry into structures, processes, and outcome followed by standard setting by consensus and local peer assessment merit serious consideration.

**P D CAMPION** 

Senior Lecturer in General Practice University of Liverpool, Liverpool L69 3BX

- 1 Irvine D. Quality of care in general practice: our outstanding problem. J R Coll Gen Pract 1983:33:521-3
- 2 Royal College of General Practitioners. Towards quality in general practice: a council consultation document. London: Royal College of General Practitioners, 1985. Royal College of General Practitioners. "What sort of doctor?" London: Royal College of General
- Practitioners, 1985. (Report from General Practice, 23.)
- Cartwright A. Patients and their doctors. London: Routledge and Kegan Paul, 1967. Walker JH, Stanley IM, Venables TL, Gambrill EC, Hodgkin GKH. The MRCGP examination and its methods. J R Coll Gen Pract 1983;33:662-75. 6 Walker JH, Stanley IM, Venables TL, Gambrill EC, Hodgkin GKH. The MRCGP examination
- water JH, stanley IM, venables TL, Gambrill EC, Hodgkin GKH. The WRVEOF examination and its methods. J R Coll Gen Pract 1983;33:732-7.
  Walker JH, Stanley IM, Venables TL, Gambrill EC, Hodgkin GKH. The MRCGP examination and its methods. J R Coll Gen Pract 1983;33:804-10.
  Marinker M. The MRCGP revisited. J R Coll Gen Pract 1984;34:529-32.
- Stanley IM, Belton A, Freeman P, et al. A method of assessment during vocational training: report of a pilot study. J R Coll Gen Pract 1985;35:9-14.

Horder J. Alma-Ata Declaration. Br Med J 1983;286:191-4.
 Townsend P, Davidson N. Inequalities in health. London: Penguin, 1982.

## Molecular genetics of acute intermittent porphyria

Acute intermittent porphyria is the most important of the porphyria diseases and is clinically distinguishable from the others by the dominance of gastrointestinal and neuropsychiatric symptoms and the absence of skin photosensitivity. The most prominent symptoms are abdominal pain, vomiting, and constipation, with less commonly paralysis or paresis and psychological abnormalities of various types. The main signs of the disease are tachycardia -a useful index of activity-and hypertension, which is present in over half of patients. All of these clinical mani-