understanding of biological process seems beyond reason.

References

- (1) Kottow M H. Classical medicine v alternative medical practices. *Journal of medical ethics* 1992; 18: 18-22.
- (2) Hume D. A treatise of human nature. London: Penguin, 1984: 124 and 139.
- (3) Coulter H L. Divided legacy: a history of the schism in medical thought. Washington DC: Wehawken Book Co, 1977, vol I: 337-418.
- (4) Bulletin general de therapeutique, VIII. 1835: 158-191.
- (5) Armstrong D. The problem of the whole person in holistic medicine. *Holistic medicine* 1986; 1: 27–36.
- (6) Jewson N K. Disappearance of the sick man from medical cosmologies, 1770-1870. Sociology 1976; 10: 225-244.

ADRIAN RENTON MFPHM St Mary's Hospital Medical School, Norfolk Place, London W2 1PG

'Unprincipled QALYs'

SIR

There's just one point of theoretical interest in Mr Cubbon's response to my Unprincipled QALYs (1,2). He complains that since his approach is utilitarian and since utilitarian approaches are as old as the hills he 'cannot believe that the essence of the approach for which I was arguing can be satisfactorily attacked with ephemeral slogans such as "ageism", "sexism" and "Thatcherism".

I think Cubbon must have meant to complain that the slogans were 'contemporary' rather 'ephemeral'; but of course his approach was not attacked with slogans but with arguments to the effect that it could be fairly characterised by those slogans, that they represented its ideology and that the ideology was flawed. In so far as Thatcherism is an ideology, some of the Greeks (ancient) could have been Thatcherite, just as there could have been fascists among them. There were certainly sexists among them.

References

(1) Harris J. Unprincipled QALYs: a

- response to Cubbon. Journal of medical ethics 1991; 17: 185-188.
- (2) Cubbon J. The principle of QALY maximisation as the basis for allocating health care resources. *Journal of medical ethics* 1991; 17: 181-184.

JOHN HARRIS

The Centre for Social Ethics and Policy, University of Manchester, Oxford Road, Manchester M13 9PL

Response to Saunders and Singh

SH

I was pleased to read the comments about my article Enforced death: enforced life (Journal of Medical Ethics, 1991; 17: 144) by Dame Cicely Saunders and Surinder Singh and I should like to say something about their interesting remarks.

was, as Saunders correctly surmised, unaware that both the Association for Palliative Medicine and the European Association for Palliative Care '...consider that the direct and intentional killing of patients is unacceptable'. However, I would have been surprised to hear otherwise and being aware of this fact does not incline me to change my views in the slightest, since it is people with such views that I am primarily interested in influencing. I would have been interested to know whether these organisations consider that the indirect and intentional killing of patients is acceptable; do they consider that it is OK to kill one's patients provided that one can say with hand on heart that all one was really aiming at was relieving their pain?

As far as intentional killing goes I can see no moral distinction between direct killing and indirect killing or 'allowing to die' as it is sometimes sanitisingly called. To my mind, anyone who considers that even in the kinds of circumstances I described, the direct killing of those who request to be killed is wrong, and yet would condone indirect killing in such circumstances, has to explain why one is acceptable while the other is not. And they have to do so in a way that does not refer to the spurious moral distinction between killing and letting die.

I note with interest that Saunders seems to agree with me both that there are cases where palliative care is unavailable and that there are cases where it is unavailing (ie ineffective) and hence that there will be people who will have life *enforced* on them, unless they are killed at their request. Naturally, as Saunders rightly points out, in such circumstances patients will have to rely on the compassion and understanding of their doctors and other carers. It's just that I think understanding and compassion in such circumstances, would lead palliative carers to kill their patients; to fail to do so would, in my opinion, be indicative of a failure of real understanding and a lack of genuine compassion.

Turning now to Dr Singh's remark, let me first of all deny that it is disingenuous of me to have said that doctors who refuse, in circumstances such as I have described, to kill their patients, 'guard their own quality of life at the patient's expense', since I believe it to be true. A doctor who fails to kill a terminally ill patient who is suffering irremediable pain or distress, who would prefer to be dead and has rationally asked to be killed, cannot claim to be doing so out of compassion for the patient. She cannot refuse help to such a patient and claim in good faith that she has done so because she understands that patient's distress and suffering. My guess is that it is a naive adherence to the idea that direct killing is always wrong (though indirect killings may not be) that persuades doctors who wish to remain 'good guys', to refrain from killing even in in extremis.

It could be claimed that it is because they believe that to enter into the practice of killing certain patients in order to help them could result in their becoming the kinds of people who might kill other patients against their will, that doctors do not want to add killing to their battery of clinical skills. But then to do this is to place the absolute against killing above the absolute against causing suffering (in this case by the omission to kill) and this seems to me to be questionable.

I agree with Singh that it is legitimate that requests for death from patients on account of poor quality of life, should 'be treated with the utmost caution and fully explored'. I also agree with him that the implementation of acts of euthanasia must 'be treated with great circumspection and caution'. Indeed, given that I only believe that doctors should kill patients who. devastatingly awful circumstances, autonomously and rationally request to be killed, I hope that it was obvious in my article that my attitude to euthanasia is probably as cautious as it is possible to be, without being absolutist. I do not, as

he seems to imagine I might, support 'early recourse' to euthanasia. Nor do I, as Saunders seemed to think, support the legalisation of euthanasia; I believe that to legalise euthanasia would be to set foot on a slippery slope towards a situation where unrequested euthanasia would become more likely. I do. however, believe in what Singh refers to as a central tenet of medical ethics, Primum non nocere or 'first do no harm'. It's just that I believe that to force a person to live a life that he would rather not live, a life that for him has negative rather than positive value, and is destined to end soon anyway, is to harm him most awfully, and that to relieve him of the burden of that life would be to help in the compassionate and understanding way that I believe should be central to the enterprise of medicine.

Wolf Wolfensberger (1) has recently written that he believes we are living in a deathmaking society. Wolfensberger is wary of euthanasia and along with countless others I share his alarm at the possibility that utilitarian ethics is leading us into bad places. I do not wish to live in a world where old people and sick people and disabled people are killed because they are an encumbrance on others. But the euthanasia to which I have offered support is not about deathmaking; on the contrary, it could be argued, it is about respecting and even venerating life.

Like hospice care aimed at allowing people to live their lives fully to the end, requested euthanasia of the limited kind that I believe is decent and morally right, would allow the individual whose life is in question, to die gracefully. By

contrast it could be argued, forcing a person who wants to die to live on in pain and distress, when one could by killing, help him, is to disregard the things that make life valuable. In such a case killing the person could be seen as life-enhancing while forcing life on him by refusing death, could be seen as death-making.

References

(1) Wolfensberger, W. The genocide of handicapped and afflicted people. Syracuse, NY: Author, 1987.

> GAVIN J FAIRBAIRN Senior Lecturer in Education. North East Wales Institute, 23 Range Road, Whalley Range, Manchester M16 8FS

News and notes

Manuscripts sought

The editor of the Journal of Information Ethics (published by McFarland and Co) would be gratified to hear from potential contributors. JIE is a new, semi-annual periodical that deals with ethics in all areas of information or knowledge production and dissemination.

This includes, but is not limited to, library and information science, education for these professions, technology, government publication and legislation,

display, graphic computer security, database management, peer review, privacy, cyberspace, and information liability approached from sociological, philosophical, theoretical, and applied perspectives.

Please query before producing a manuscript. Contact Robert Hauptman, Editor, The Journal of Information Ethics, LRS, St Cloud State University, St Cloud, MN 56301, USA. Telephone: (612) 255-4822.