Bad behaviour does not equal research fraud

I was not impressed by Dr Geggie's article offering a survey of the attitudes of newly appointed consultants towards research fraud (Journal of Medical Ethics 2001;27:344-6). Indeed, by mixing up categories of misconduct from what is at most "bad behaviour" to the very serious, he is not entirely beyond reproach himself. I remind readers that Dr Geggie suggested that 55.7% of the respondents had observed (from the title) "research fraud".

If the term "research fraud" is to have any meaning, it must be reserved for conduct that consciously and deliberately attempts to impose a fraud on others. The US National Academy of Sciences report, On Being a Scientist: Responsible Conduct in Research distinguishes clearly between "misallocation of credit, honest errors, and errors caused through negligence" and "deception, making up data, changing or misreporting data or results, and plagiarism". The former are "ethical transgressions...that generally remain internal to the scientific community... dealt with locally through peer review, administrative action, and the system of appointments and evaluations". The latter "strike at the heart of the values on which science is based". The White House's Office of Science and Technology Policy reached similar conclusions, restricting research misconduct to "fabrication, falsification and plagiarism".

I agree with these assessments.

In Dr Geggie's paper, deception would include deliberate falsification of data (category 3 of Dr Geggie's table 1), cheating (4A) and deliberately plagiarising without attribution (4D). We then move into fairyland, because Dr Geggie next asks "have you ever been omitted from a paper for which any of the authors have not made a sufficient contribution to warrant credit for the work" or "has your name ever been omitted from a paper for which you had a substantial contribution?..." and, surprise, surprise, over half the consultants we reported in the questionnaire fits this definition. The same dictionary includes in its definition of "fraud" the phrases "use of false representations to gain unjust advantage" and "person or thing not fulfilling expectation or description". The behaviour described in questions 1, 2, and 3—that is inappropriate authorship, omission of data for publication, fits this definition. In the paper (Journal of Medical Ethics 2001;27:344-6). For the purposes of my article I stated that consultants who had answered "yes" to questions 1, 2, or 3 of table 1 had reported "observed misconduct" and I stand by this conclusion. The dictionary on my bookshelf (Concise Oxford Dictionary (6th ed)) defines "misconduct" as "improper conduct, esp. adulter; bad management" and I think that what the consultants were reporting in the questionnaire fits this definition. The same dictionary includes in its definition of "fraud" the phrases "use of false representations to gain unjust advantage" and "person or thing not fulfilling expectation or description". The behaviour described in questions 1, 2, and 3—that is inappropriate authorship, omission of data for publication, fits this definition.

In the article I tried to use the term "fraud" as a more general term than "misconduct" and I apologise if this caused confusion. I think, however, that the majority of readers (and certainly the majority of my colleagues) would have had something approaching the above definitions in mind when they read the article (although several colleagues have told me to get a more up-to-date dictionary)! Whilst I acknowledge that Professor Williamson makes a valid point, I do think there is a danger of any rational discussion becoming bogged down in semantics and I think it is clear from my article what the initial data was and how I have derived my variables.

Professor Williamson dismisses the medical consultants (not cleaners!) who answered yes to questions 1 and 2 as having "more often delusions of grandeur" on their part. That is a point of view—but I suspect the majority of respondents to my questionnaire do not share it. Indeed I think his attitude risks sweeping a real problem under the carpet and, whilst I am aware of attempts to standardise authorship, I think this is a problem which requires further investigation.

I find the overall tone of this letter and phrases such as "into fairyland" and "give me a break" inflammatory and this, unfortunately, distracts from some valid points.

Professor Williamson's main criticism appears to be that he has learnt nothing new from my paper. The original idea for this study came after a lively discussion in a bar about the prevalence of research fraud (a subject of which I knew very little). After this discussion I did a literature search and could find only a few references to research fraud and misconduct. I decided to perform a survey in an attempt to answer the questions I had. I wish that Professor Williamson had been present at the original discussion—he could have saved me a lot of trouble! On a more serious note, I am sure that the professor is aware that in the "hierarchy of evidence" questionnaire surveys rank higher than anecdotal evidence and personal experience (although only just!). The value of my article is that it adds evidence to that it adds evidence to a subject in which there is a marked paucity of hard facts.

I am sorry that Professor Williamson was unimpressed with my paper. One of the problems with the very nature of the subject of research fraud/misconduct mitigates against good quality research. I am sure that readers would welcome any double-blinded randomised controlled trials (RCTs) on the subject and, if the professor wishes to carry out some research in this area I am sure that the scientific community would welcome it.

Dr Geggie

The Murdoch Childrens Research Institute, 10th Floor, Royal Children's Hospital, Flemington Road, Parkville Vic 3052, Australia; williamsb@cryptic.rch.unimelb.edu.au

Author's reply

Professor Williamson makes a valid point about the term "research fraud" and I agree that the term covers a number of different categories of unethical behaviour. I also pointed out that "Research fraud can take many forms", that "inclusion of data or result...and plagiarism". The former are "ethical transgressions...that generally remain internal to the scientific community... dealt...", and, surprise, surprise, over half the consultants we reported in the questionnaire fits this definition. The same dictionary includes in its definition of "fraud" the phrases "use of false representations to gain unjust advantage" and "person or thing not fulfilling expectation or description". The behaviour described in questions 1, 2, and 3—that is inappropriate authorship, omission of data for publication, fits this definition. In the article I tried to use the term "fraud" as a more general term than "misconduct" and I apologise if this caused confusion. I think, however, that the majority of readers (and certainly the majority of my colleagues) would have had something approaching the above definitions in mind when they read the article (although several colleagues have told me to get a more up-to-date dictionary)! Whilst I acknowledge that Professor Williamson makes a valid point, I do think there is a danger of any rational discussion becoming bogged down in semantics and I think it is clear from my article what the initial data was and how I have derived my variables.

Professor Williamson dismisses the medical consultants (not cleaners!) who answered yes to questions 1 and 2 as having "more often delusions of grandeur" on their part. That is a point of view—but I suspect the majority of respondents to my questionnaire do not share it. Indeed I think his attitude risks sweeping a real problem under the carpet and, whilst I am aware of attempts to standardise authorship, I think this is a problem which requires further investigation.

I find the overall tone of this letter and phrases such as "into fairyland" and "give me a break" inflammatory and this, unfortunately, distracts from some valid points.

Professor Williamson's main criticism appears to be that he has learnt nothing new from my paper. The original idea for this study came after a lively discussion in a bar about the prevalence of research fraud (a subject of which I knew very little). After this discussion I did a literature search and could find only a few references to research fraud and misconduct. I decided to perform a survey in an attempt to answer the questions I had. I wish that Professor Williamson had been present at the original discussion—he could have saved me a lot of trouble! On a more serious note, I am sure that the professor is aware that in the "hierarchy of evidence" questionnaire surveys rank higher than anecdotal evidence and personal experience (although only just!). The value of my article is that it adds evidence to that it adds evidence to a subject in which there is a marked paucity of hard facts.

I am sorry that Professor Williamson was unimpressed with my paper. One of the problems with the very nature of the subject of research fraud/misconduct mitigates against good quality research. I am sure that readers would welcome any double-blinded randomised controlled trials (RCTs) on the subject and, if the professor wishes to carry out some research in this area I am sure that the scientific community would welcome it.

D Geggie

Accident and Emergency Department, Arrowe Park Hospital, Arrowe Park Road, Upton, Wirral, Merseyside CH49 5PE, UK; DAVID@DGEGGIE.fsireserve.co.uk

Ethical implications of consent and confidentiality

Recently a prospective, observational clinical study was carried out in the department of ophthalmology, at a district general hospital. The main purpose of the study was to evaluate the medicolegal and ethical implication of consent and confidentiality in ophthalmic practice, in accordance with the guidelines provided by medical law.

One hundred patients, who had been referred by optometrists to ophthalmologists, were included in the study. The general ophthalmic services (GOS) 18 form, a referral form used by optometrists for referring patients to ophthalmologists, allows optometrists to share a patient's medical information with ophthalmologists, was used as a study tool to evaluate the percentage of patients giving signatory consent.

Data was also collected regarding patients' awareness about the medicolegal implications of consent and their views about their medical information being shared among different health care professionals in ophthalmic practice.

The results of our study show that only 15% of GOs 18 forms contained written consent by the patients for information to be shared by their optometrist. The remaining 85% were referred without obtaining an express written consent.

These results were further supported by the information generated from a questionnaire filled in by patients attending the eye clinics. The questionnaire includes the following three simple questions: responses are as follows:...
Are you aware about the fact that you should be consented by your optometrist on the referral form? Yes/No (46% / 54%) Did your optometrist explain to you about the consent statement mentioned on the GOS 18 form? Yes/No (40% / 60%) Would you like the ophthalmologist to make available your medical information to your optometrist/ophthalmic medical practitioner? Yes/No (85% / 15%)

Conclusion
Only a few GOS 18 forms contained patients’ written consent for information to be sent back to the referring optometrist. Fifteen per cent of the patients surveyed said they did not wish information to be shared with their optometrists. Therefore we should be careful about sending back information to optometrists where signatory consent has not been given.

Optometrists need to be aware of this potential issue. In the light of the increasingly close relationship between optometrists and optometrists (especially where they share care for glaucoma and postoperative cataract patients) it is important for the optometrist that consent is given if feedback is required.

Discussion
A good doctor-patient relationship can be defined by the three Cs: (i) Confidentiality, (ii) Consent, and (iii) Competence. If any of these three components are missing the doctor-patient relationship could be damaged and the flow of communication in both directions inhibited.

A promise on the part of the doctor to maintain patient confidentiality is central if patients are to be allowed to speak freely. If information is shared without the patient’s consent then the faith of the patient in the treating optometrist could be damaged and the doctor-patient relationship could be undermined. Consent in its most basic form must be evidenced in the patient’s written or verbal consent to have their information shared with their optometrists. Therefore we hold patients to an expectation that they will not wish their medical records to be sent to their optometrists. Such an expectation should be respected.

Consent is an integral part of the GOS 18 referral form but our study shows that it is taken for granted and is not handled in accordance with guidelines set out in medical law.

Y Khan
SHO Ophthalmology, Ashford Hospital, London Road, Middlesex, TW15 3AA, UK; yasii1399@hotmail.com

R J Stirling
Darlington Memorial Hospital, Hollyhurst Road, Darlington, DL3 6HX, UK

Tuskegee’s Truths: Rethinking the Tuskegee Syphills Study

No one interested in the ethics of biomedical research will have failed to hear about the Tuskegee syphilis study, or, to give it its full title, the US Public Health Service’s Tuskegee Study of Untreated Syphilis in the Negro Male. This study, conducted from 1932 to 1972, on black (African American) males in Tuskegee, Alabama, has, with complete justification, become the paradigm of moral depravity in the field of biomedical research. Virtually every rule of good, ethical research was broken during this “research” over a period of 40 years, down to denying participants even the knowledge, let alone the option, of a remedy when it became available.

In recent years, the Tuskegee syphilis study has received renewed public attention, for two reasons. First, in 1996, 24 years after the cessation of the study, President Clinton provided a formal federal apology, saying to the survivors that “[w]e can look at you in the eye and finally say on behalf of the American people, what the United States government did was shameful, and I am sorry”. With this apology, Clinton not only accepted moral responsibility—something not easily done by governments in the affairs of state, domestic or foreign—but also contributed to addressing African Americans’ distrust of health care and biomedical research, a distrust fuelled by the legacy of Tuskegee.

Second, echoes of Tuskegee have been heard in the ongoing debate about the ethics of biomedical research financed or conducted in the developing world by government agencies and companies from the developed world, in particular in regard to HIV/AIDS. Such research raises ethical questions, relating to key issues such as exploitation and justice, informed consent, and duties of beneficence. David J Rothman at the MacLean Center for Clinical Medical Ethics at the University of Chicago. (Her analysis is particularly clear here, and in other “reproductive ethics” chapters, especially in the distinction between genetic and stem cell research, which is recently approved by the UK parliament, affecting women differently from men?) The obvious reason is that large numbers of enucleated eggs will be required, and that enucleated eggs come from women, taken in a painful and difficult procedure. But as Mahowald markedly notes, so far as the further step of human cloning goes: “Interestingly, while some bioethicists expressed concerns about the impact of human cloning on cloned individuals, none, to my knowledge, indicated that there were gender differences to worry about as well” (pages 281–2).

These practical chapters show Mahowald’s clinical knowledge to good advantage (although a philosopher, she is professor in the college, the Department of Obstetrics and Gynecology, the Committee on Genetics and the MacLean Center for Clinical Medical Ethics at the University of Chicago.) Her analysis is particularly clear here, and in other “reproductive ethics” chapters, especially in the distinction she draws between genetic, gynecological, and lactational motherhood. She rightly draws our attention to a fourth form of motherhood which can also now occur: the provision of enucleated eggs, into which another set of genes is inserted. Which of these is “real” motherhood?

W A Landman

Reference

Genes, Women, Equality

Far too often it is still assumed that if feminist bioethics has any role to play, its contributions are purely in reproductive ethics. For instance, one of the beliefs behind Mahowald’s: Women, Equality should dispel that delusion once and for all, with a second illusion: that the new genetics is gender-neutral. Mahowald is not a bioethical Luddite: she is not concerned to attack the new genetics, but to make good the failure of bioethicists and scientists to explore the differential impact of the new genetics on women. Specific implications may be seen in reproductive ethics, but different fields form the bulk of the book, with chapters on genetic counselling; genetics research; allocation of genetic services; culture and sex selection; misattributed parenthood and cystic fibrosis; sickle cell disease and carrier testing; breast cancer susceptibility testing; preimplantation genetic diagnosis and abortion; genomic alternation; genetically linked alcoholism, employment and insurance testing, and human cloning.

This last chapter is a prime example of the need for Mahowald’s sort of analysis: how often is it recognised that even therapeutic cloning and stem cell research, so recently approved by the UK parliament, affect women differently from men? The obvious reason is that large numbers of enucleated eggs will be required, and that enucleated eggs come from women, taken in a painful and difficult procedure. But as Mahowald mildly notes, so far as the further step of human cloning goes: “Interestingly, while some bioethicists expressed concerns about the impact of human cloning on cloned individuals, none, to my knowledge, indicated that there were gender differences to worry about as well” (pages 281–2).
There are also a series of “mid-level theory” chapters, such as that on Disabilities, feminism and caregiving, which is informed by the split in feminist thought between disability rights feminists such as Adrienne Asch, who distrust genetic screening and correlated abortion on the grounds that they pitting against the disabled, and other feminists such as Christine Overall,1 or Mahowald herself, who distinguish between the legitimacy of eugenics with disabilities and advocacy for disabled people (and their carers, usually women). The analysis in both the “specific issues” and the “mid-level” chapters will be of enormous use to both practitioners and academics.

Mahowald also attempts to provide a normative foundation for the two less theoretical sorts of chapters, particularly in her chapter 4, Gender justice in genetics. Here she explores what she terms a feminist standpoint approach or egalitarian feminist model, which directs our attention towards power imbalances. Where inequalities result from rectifiable social power imbalances rather than unalterable and value-neutral differences, the standpoint of the less powerful group should be privileged over that of the more powerful, in this model. “Some differences entail inequalities; others are merely associated with them.” Inequality that is provided by mere association, such as the continued association of women with caregiving, are more easily rectifiable; biological differences, such as the fact that it is women who give birth, are harder to remedy, but we should try to minimise their impact. Certainly we should not allow necessary biological inequality to become an excuse for avoidable social inequality, but that is what some aspects of the new genetics risk doing.

The feminist standpoint model is frequently contrasted with a conservative libertarian model, as the theoretical overview which has so far occupied us in the new genetics. While I agree with this part of Mahowald’s analysis, I am less convinced that the liberal feminist model is always wrong, even though I do not count myself a liberal feminist. There is a certain risk of demonisation of the liberal feminist view, which Mahowald does not always avoid; it is not the same as libertarianism. On the whole, however, this is a vital book for anyone interested in the new genetics—even for those who don’t actually think they are also interested in feminism.

D Dickenson

Reference

On Dying Well: An Anglican Contribution to the Debate on Euthanasia

Board for Social Responsibility of the Church of England, Church House Publishing, 2000, £4.95, 94 pages, 0 7151 6387 9

For any reader interested in euthanasia, On Dying Well gives an accessible yet detailed account of the Church of England’s view on the subject. First published in 1975, this short report is the product of the Church’s Board for Social Responsibility, which brought together theologians, philosophers, lawyers, and medical professionals to form a working party with the remit of examining euthanasia. The second edition of On Dying Well leaves most of the original working party report findings unaltered, but adds a new introduction by the then Bishop of St Albans, the chairman of the British Medical Association’s ethics committee from 1989 to 1997. Other changes to the first edition are a redrafted chapter on the legal questions surrounding euthanasia, in light of new cases pertinent to the debate and additional reflections on the report’s medical content. Also included is an updated bibliography, listing publications produced after 1975 through to 1993 Joint Submission to the House of Lords Select Committee on Medical Ethics made by the House of Bishops of the Church of England and the Roman Catholic Bishops’ Conference of England and Wales.

On Dying Well is a report of broad-ranging scope which, not surprisingly, robustly rejects the legalisation of voluntary euthanasia. This rejection operates on two interlocking but both principled, theologised grounds: one is the medicalisation of the notion of autonomy which ignores an “ultimate accountability”: the point at which the doctor-patient relationship ends. Professor Horder rejects what he terms an “unbridled” notion of autonomy which ignores an “ultimate accountability” for her hospice work in the care of the dying. She is awarded the $1m Hilton Humanitarian Prize for her hospice work in the care of the dying. The philosophical debate surrounding euthanasia has reached an impasse. The “solution” for now, appears to lie in allowing death with at least some degree of self regulation but without appeals to changes in the law.

L Campbell

Bio Engagement: Making Christian Difference through Bioethics Today


This book is concerned with advocating a pro-life stance rather than with detailed discussions of the medical ethics of biotechnology. The essays are written from the particular Christian perspective of conservative evangelicalism and the writers are committed to the verbal inspiration of scripture. It is a book which will be of interest to a certain section of the Christian church. One constraint with this approach is the sense of the discussion taking place “in house”, calling Christians of like mind to defend a pro-life stance on healthcare. Occasionally writers with contrasting points of view are mentioned, as there is sustained discussion of their arguments. The pervasive advocacy of the pro-life view only occasionally yields to a recognition of the complexity of the issues and the general, unacademic, level of the discussions does not provide any detailed argument and support for any reader interested in the subject.
for the ethical presupposition of the book, which is simply assumed.

This is a work that takes its cue from the idea of “engagement”; the wide ranging discussions of biology and biotechnology currently taking place are not covered, rather the focus is on abortion, and to a lesser extent, euthanasia. Several essays show how Christians holding pro-life views can best make their position known, how to defend their views when moral perplexities are presented in clinical practice or training. Case studies are used to support the overall programme of the book and are not subject to the analysis which might have been expected.

More technical, ethical discussion occurs in the section on Law and policy, where there is an awareness of the complexity of the issues relating to the use of unfertilised eggs which might later be fertilised. And in Casey’s essay about how the law will shape our life and death decisions: the case of the human embryo, there is detailed discussion of various important court cases in the United States. But here again the argument seems to presuppose too much to make his case convincing no matter how interesting the hints towards ethical solutions might be. In keeping with the theme “making a difference”, Casey concludes his paper with a proposal for legislation. There is an interesting criticism in this section of the liberal outlook, namely that liberalism ends in its response to those who object to liberalism.

Overall then there is a paucity of technical discussion of medical ethics in this book. Even the practical discussions are couched in the manner of advocacy rather than analysis. The particular Christian perspective does not seem likely to make the book much useful from the point of view of developing discussion even among Christian ethicists. Christian medical practitioners who share the outlook of the authors will be encouraged to advance their views in the present climate of opinion, where decisive battles are won by political lobbying. But even some who are sympathetic to the stance of this book, including this reviewer, will demand more sustained argument and sophisticated presentation of their views.

N Fennemore

Textbook of Research Ethics: Theory and Practice

Research ethics is high on the agenda of medical ethics and regulatory topics worldwide at the moment. In this field it is thus important to have good textbooks, to educate health care professionals, researchers, ethicists, students, and policy makers about the nature of research ethics, its history and social contexts. Professor Louise’s book is thus published in an increasingly competitive marketplace.

The book has many merits. It consists of five chapters and two appendices. The first chapter summarises several of the iconic scandals in medical ethics, notably the Tuskegee syphilis study and the Nazi doctors’ atrocities. The historical scholarship of this chapter is solid, if not original, and it is useful to have it gathered here in a convenient place. There is a serious debate to be had about the pedagogy and philosophy of starting research ethics from a consideration of its worst breaches; does this blind students to the myriad of more minor but more pressing problems that arise in “routine” research? Does it encourage the thought that ethics is about hard cases only? Does it take seriously the fact that research ethics is, historically speaking, an evolving discipline? Nevertheless, this is the traditional place to start, and Louise does it very well.

The second chapter summarises some of the main theoretical positions in bioethics. This is the weakest chapter in the book, several times stating a thumbnail sketch of a position (for example, lesbian ethics) with a list of pros and cons before moving on. A survey is always vulnerable to this fault, of course. More seriously, in my view, these views are set out, but few of them are seriously illustrated in the remainder of the book. A more ambitious book would have given a more thorough sketch of what, for instance, Gilligan’s ethics of care would say about HIV vaccine trials in the Third World. The promise of non-utilitarian, non-principalist approaches in research ethics has often been asserted, but rarely has a serious attempt been made to apply them in a systematic way.

Chapters three and four analyse the ethical issues that arise in study design and implementation. These chapters are very competently done, but focus in the main on consent and confidentiality, with interesting material (and this is unusual) on conflicts of interest and the rights and duties of the controller of research data relating to publication and use. Chapter five is a valuable account of the regulatory and legal frameworks governing research in the United States. This chapter is of little direct applicability for non-US readers, but is most informative for both students of comparative bioethics and for researchers who wish their studies to have US sites or to be acceptable to US sponsors and regulators.

In an increasingly globalised research context, this is a valuable asset. On the other hand, there is relatively little in the book about international research.

The book closes with a pair of appendices, one setting out in some detail the various kinds of research study design and the other setting out the principles of US law. Both of these are very useful and helpful, and should be of great assistance to readers who are unfamiliar with either—a situation common enough, since research ethics interests both the researcher (who may know no law or ethics) and the ethicist (whose grasp of research design may be weak).

The book is not comprehensive: gaps include health services and nursing research, and human rights in medical research. A useful feature is the inclusion of exercises for the reader, although these are very taxing at times, and no model solutions are provided.

In summary this is a valuable textbook, which aptly summarises much of current scholarship in research ethics. It would be suitable for undergraduate or postgraduate courses in medical ethics and in research methods. I find the price of the book rather high, considering the existence of more affordable alternatives. That said, the teacher of research ethics will find this a very useful addition to his or her library.

R E Ashcroft

NOTICES

6th World Congress of Bioethics

The 6th World Congress of Bioethics will be held in Brasilia, from 30 October to 3 November 2002. The congress is sponsored by the International Association of Bioethics, the Brazilian Society of Bioethics, the Center for Studies and Research in Bioethics (University of Brazil), the Brazilian Federal Council of Medicine and the Latin American and Caribbean Federation of Bioethics Institutions.

The theme of the congress is bioethics: power and injustice.

For further information please see: www.bioethicscongress.org.br

News from the Centre for the Study of Global Ethics, the University of Birmingham

The new weekend short course programme includes Global Bioethics (June) and Development Ethics (September).

The new MSc in Global Ethics begins in October 2002, with modules including: Global Ethics; Globalisation and Governance; Conflict Humanitarian Aid; Non-Governmental Organisations in a Changing International Context; Research Methods, and practical placements in non-governmental organisations, as well as the above course topics.

For further information please contact: Helen Harris on +44 (0)121 693 4687.

www.jmedethics.com
Bad behaviour does not equal research fraud

Bob Williamson

*J Med Ethics* 2002 28: 207
doi: 10.1136/jme.28.3.207

Updated information and services can be found at:
http://jme.bmj.com/content/28/3/207.1

These include:

**Email alerting service**

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/