Thoughts on patriarchy

I felt I could not let the diatribe¹ posing as a scientific article by Professor Malcolm Potts and his wife, Dr Martha Campbell, go without comment. When one reads an article that contains gross inaccuracy, even in part, the credibility of the rest and of the journal running it is thrown into question.

The couple has an understandable modern problem with St Augustine but that is no excuse for their absurdly inaccurate summary of over a hundred theological works,² that he "saw God as an arbitrary judge who could be placated by bribes". Nor does it justify ridiculing him from a 21st century perspective with no attempt to understand his historical or personal context, or indeed any of their other (highly selected) examples. To then link patron saints in a *non sequitur* to bribery and the etymology of 'grace' (wrongly)³ to the same is as ridiculous as it is irrelevant. (I note they are only able to quote Potts' own authority for this.)

To limit the development of Christian sexual ethics only to the context of sexual exploitation tolerated in the Roman Empire is profoundly misleading. They show no awareness of the contributing influences of Judaism, Hellenism, Stoicism or Gnosticism.⁴

Just how blinkered the authors' historical viewpoint is can be seen from the fact that the discussion is limited to one theologian among many, to only the Western tradition, ignoring Eastern and Celtic perspectives completely, and subsequently narrowing within that to Roman Catholicism.

Islam fares worse, seeming only to be represented by the Taliban in the authors' minds. If the two faiths both have "liberal and conservative interpretations", why is discussion limited to the negative ones?

The most one can learn from this article is something of the prejudices of its authors. This is not the first time Malcolm Potts has been accused of a lack of objectivity.⁵

Space precludes a more detailed response but I wonder how this got through the Journal's peer review process. It would have been sensible to mention Martha Campbell's membership of the Editorial Advisory Board as a competing interest. In a journal aspiring to international scientific respectability, authors should be obliged to give balanced and informed discussion. If polemic is to be part of the offering, it should be flagged as such.

I am certainly no friend of patriarchy, but a scientific article is not the place to abandon balance in favour of rhetoric.

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Competing interest

The author is in training as a Reader in the Church of England.

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Reply

We appreciate Dr Shepherd's broader view of other strains of Catholicism.¹ However, our article was not a book, but a glance at those streams of patriarch that are doing most damage to women's health. We did not try to be

comprehensive and our discussion focuses not on the liberal wings of Christianity and Islam but rather on the conservative, because these represent seven-eighths of their history, and the shadow of conservatism seems to last longest in the area of human sexuality.

Dr Shepherd is also correct that striving to understand the historical context in which key actors such as Augustine of Hippo lived is central to historical scholarship. But it is precisely that historical context which we find so brutally patriarchal. We took part of our analysis from careful historians of Christianity, such as Susan Haskins,² Marina Warner,³ Elaine Pagels⁴ and Uta Ranke-Heinemann.⁵ In addition, some of the insights into Augustine came from a patristic historian and theologian, the late Francis Xavier Murphy, who taught moral theology at the Academia Alfonsiana in Rome and who was a *peritus* (expert adviser) to the Second Vatican Council.⁶ He was a personal friend and mentor of one of us (MP).

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Implant removal by modifying access

I read with great interest the review articles by Diana Mansour and colleagues on 'Methods of accurate localisation of the non-palpable subdermal contraceptive implant' and 'Removal of non-palpable etonogestrel implants' in the January and April 2008 issues of the Journal, respectively.^{1,2} I totally agree with the author's suggestion to modify access with 2–3 ml of local anaesthetic applied under the palpable portion. This helped me greatly to modify the procedure, depending upon the patient's clinical circumstances and requirements as outlined in the following two cases.

Case 1: A 24-year-old mother of two children was fitted with Implanon® 18 months previously. The patient requested removal of the Implanon for her next conception. On examination the distal end of the implant was non-palpable and the proximal end was palpable with difficulty. An ultrasound scan was done as suggested in the article. The ultrasound report noted that: "the Implanon is located in the left arm, the lower edge is marked and is at a depth of 1.6 cm. The upper edge is palpable". After explaining the possible difficulties, I obtained fully informed, written consent from the patient for removal of her implant. I made a 4 mm incision at the proximal, barely palpable, end of the Implanon and dissected with small curved mosquito forceps and finally managed to get the Implanon out without any surgical complications. Here I realised the importance of having an assistant and the use of small skin retractors, which were mentioned in the article but which were not available in our clinic. Consequently we ordered

two small skin retractors with immediate effect for future use.

Case 2: A 38-year-old Zimbabwean woman was referred to me by a colleague for removal of a Norplant® device inserted 6 years ago in Zimbabwe. I explained to the patient that I am not trained in Norplant insertion or removal, although I am trained and experienced in Implanon insertion and removal. Consequently I suggested referral to a doctor trained in Norplant insertion/removal. The patient declined referral and insisted that I undertake the removal procedure. On examination, four of the rods were palpable and close together and a few millimetres above the insertion scar. Although the rods on either side were about 1.5 cm away, they were superficial and easily palpable. I explained to the patient that the middle four rods could be removed by means of the 'pop-out' technique with one incision and the remaining two rods could be removed by the same technique but with two separate incisions, similar to Implanon removal. The patient happily gave written consent to this suggested treatment plan and accepted the risks and complications (if any) of the procedure such as multiple scars, infection, bruises, nerve and vessel injury and their consequences.

Even though the Mansour *et al.* article mentioned a vertical incision for removal I went through the original horizontal scar and carefully dissected tissues vertically and removed four rods by directing each of them to the incision. Rather than attempting the heroic removal of the remaining two rods (located laterally and above) through the same incision, I felt it was much safer and easier to employ two small incisions (<2 mm each) and use the 'pop-out' technique, similar to Implanon removal. This avoided a T-incision with the old scar and I hope the patient will end up with only the single, old incision scar. A patient review visit after 1 week confirmed a nicely healing wound. I discharged the patient from the clinic since she had not decided on her future contraception, except the use of condoms. The patient was extremely happy to have avoided referral to an unknown clinic for removal of her Norplant.

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Norplant removal forceps

I was interested to read the article by Diana Mansour *et al.*¹ reviewing removal techniques for contraceptive implants.

In common, I am sure, with many other readers, I am continuing to see patients requiring Norplant® removal who have had this implant fitted outside the UK.

Our primary care trust has opted for a single-use instruments policy, and we have had to get rid of our desktop sterilisers, so we are no longer able to reuse the modified vasectomy forceps ("Norplant removers") with a 2.2 mm diameter, referred to in the article.

The best that any of the single-use manufacturers I have contacted can come up with is disposable vasectomy forceps with an internal diameter of approximately 4 mm, which allow the Norplant rods to slip during "U" removals (Figure 1, right).

I am left with the unsatisfactory situation of using the wrong instruments, or referring to the

acute Trust for removal, where they still re-sterilise instruments.

Do the authors know of a source for the correct single-use instruments, or can a manufacturer be encouraged to develop the appropriate items?

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Reference

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Figure 1 Disposable vasectomy forceps used for Norplant® removal

Reply

We thank Anne Bennett for her comments¹ related to our article describing removal techniques for contraceptive implants.²

Many primary care organisations are bringing in 'single-use instruments policies' as a result of new NHS guidance on decontamination aimed at improving the quality of surgical instrument reprocessing across the health care sectors in England. Finding manufacturers who are prepared to supply small numbers of specialist disposable surgical instruments is difficult, however we have good news.

Disposable modified vasectomy forceps can now be obtained for about £2 from the supplier mentioned below. This company is also making disposable 'deep' implant removal kits, which include mosquito forceps and small Langenbecks skin retractors.

Contact: Unisurge International Ltd, Unit N, Dales Manor Business Park, East Way, Sawston, Cambridge CB22 4TJ, UK. Tel: 01223 839911. E-mail: info@unisurge.com.

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Underuse of the IUD

I read the article on the underuse of the intrauterine device (IUD)¹ in the July issue with great interest and noted that none of the references are prior to 1983. That is 25 years ago, however the real blow to the IUD's popularity was in the late-1970s when the Dalkon Shield[®] was pilloried and vilified as a source of pelvic infection and sterility, on what I believe was false scientific evidence. At that time there was a UK IUD network, organised by Professor R Snowden of Exeter University to which 20 major family planning clinics belonged.

Every device fitted and every subsequent patient visit was recorded and sent to Exeter. In 1977, over 40 000 fittings had been recorded, of which 7282 were Dalkon Shields. There were

only two cases of pelvic infection and both were cases known to me in Glasgow: one in a prostitute with gonorrhoea and the other in a woman who had tried to abort herself with a knitting needle. As a direct result of organised adverse publicity, the manufacturers of the Dalkon Shield went out of business and the reputation of all IUDs worldwide suffered a slump from which it has never recovered. A similar campaign against Depo-Provera[®] was mounted from the USA in the early 1980s and injectable methods never achieved the popularity they deserve for similar commercial reasons. Women are ill-served by 'market forces'.

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Nurse IUD fitting

We are writing to respond to the Personal View article by Cooling and Dunster entitled 'Nurse intrauterine device training' that was published in the July issue of the Journal.¹ We wholeheartedly applaud the positive experience expressed within this article, and its conclusion that "... nurses who are recognised trainers and experienced in IUD insertion to supervise the training of doctors" as this underpins the work the FSRH Associate Nurses Working Group has been undertaking recently. However, we felt the need to respond to the not-so-positive comments about the Royal College of Nursing (RCN) guidance being "absurd in requiring the learner to observe the trainer doing five insertions in patients: one is usually sufficient ...". The RCN has been instrumental in the development of guidance in order to enhance and advance nursing practice, and has specifically demonstrated its support in enabling nurses working at a higher or specialist level within the area of contraception and sexual health to undertake the removal and insertion of intrauterine techniques (IUTs) and implants. Without this guidance, nurses who were performing these procedures, or who wished to do so, were opening themselves up to litigation should an error occur. The rationale behind the guidance stating five insertions is in the fact that some nurses (i.e. gynaecology nurses) may not have ever had the opportunity to observe an IUT fitting. It was always understood that those of us who had greater exposure to procedures would not necessarily need this level of observation.

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Pharmacy-based sexual health services and clinical governance

I read with interest the Editorial by Beth Taylor in the July 2008 issue of the Journal.¹

I agree that the time is right to develop vision to embrace additional providers of the sexual health service. The new pharmacist's contract is a welcome development in this direction.

As Beth Taylor highlighted, there is certainly a need for educational and training support from specialist services in order to avoid isolation. One way of developing a robust professional link would be the development of a linked Clinical Governance Plan with local specialist services. This will not only assure a safety net but also help in continued development of such services. This

would need to be thought out and planned carefully at the primary care trust (PCT) level. Moreover, treatment and patient group directions (PGDs), especially for STIs, would have to be carefully developed in the light of the recent alert on high-level azithromycin resistance in *Neisseria gonorrhoea*. Other issues such as partner notification would also need to be resolved through training input and professional pathway with local specialist services.

Another concern would be the prescription cost for medications, which already are free on the NHS. It would be an opportunity to work with the local PCTs to explore whether they would be willing to support some/all of the costs associated with a pharmacy site for Level 1/2 Sexual Health Service delivery. There are no data on demand for a Sexual Health Service with prescription charge. The uptake of the newly launched online chargeable repeat contraception service would give an indication of clients' willingness to pay for such services.

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Reference

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Treatment of perimenopausal menorrhagia with Implanon[®]

I write concerning the successful treatment of perimenopausal menorrhagia with Implanon[®] in a 53-year-old woman. The patient (date of birth early 1953) was seen at the surgery in October 2004 with symptoms of flushing and regular periods. In early 2006 she developed menorrhagia, which was investigated with normal hysteroscopy and intrauterine system (IUS) insertion in early August 2006. The IUS was expelled after 2 months *in situ* and after ongoing symptoms of polymenorrhagia. After some discussion with the patient regarding treatment options, she decided to trial Implanon insertion, aware that it was not a clinically recognised treatment option for menorrhagia. The insertion was carried out in early October 2006. On review in August 2008 the patient noted light bleeds in March and April 2007, and a 2-day light bleed in May 2008. She stated that she "would be willing to recommend Implanon to anybody".

I plan to write to the manufacturer concerned, namely Organon, concerning this important clinical effect of Implanon in the treatment of perimenopausal dysfunctional uterine bleeding. I would be interested to hear if other practitioners have anecdotal evidence of Implanon being used in this way.

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LETTERS TO THE EDITOR

Letters to the Editor are welcome and generally should not exceed 600 words or cite more than five references. For comments on material published in the most recent issue of the Journal, correspondence should be received within 4 weeks of dispatch of that Journal to be in time for inclusion in the next issue. When submitting letters correspondents should include their job title, a maximum of two qualifications and their address(es). A statement on competing interests should also be submitted for all letters. Letters may be submitted to the Editor or the Journal Editorial Office (details on page 205).