### FROM THE CEU/FACULTY EXAMINATIONS

The CEU recommends that women should be informed about the lack of data on the efficacy of POEC when using liver enzyme-inducers and be offered an IUD as an alternative. Neither regimens of POEC (two tablets at first presentation followed by one tablet 12 h later or the three tablets single regimen) are within the product licence for Levonelle and Levonelle-2.<sup>3</sup> Clinicians may consider using the regimen that is most acceptable to individual women using liver enzyme-inducers but acknowledge the lack of an evidence base for this.

#### Disclaimer

The advice given in this Members' Enquiry Response has been prepared by the FFPRHC Clinical Effectiveness Unit team. It is based on a structured search and review of published evidence available at the time of preparation. The advice given here should be considered as guidance only. Adherence to it will not ensure a successful outcome in every case and it may not include all acceptable methods of care aimed at the same results. This response has been prepared as a service to FFPRHC members, but is not an official Faculty Guidance product; Faculty Guidance is produced by a different and lengthier process. It is not intended to be construed or to

serve as a standard of medical care. Such standards are determined on the basis of all clinical data available for an individual case and are subject to change as scientific knowledge advances. Members are welcome to reproduce this Response by photocopying or other means, in order to share the information with colleagues.

### References

- 1 Faculty of Family Planning and Reproductive Health Care Clinical Effectiveness Unit. FFPRHC Guidance. Emergency contraception (April 2003). *J Fam Plann Reprod Health Care* 2003; **29**(2): 9–16.
- 2 British National Formulary, Vol. 48, September 2004. London, UK: British Medical Association and the Royal Pharmaceutical Society of Great Britain, 2004. http://www.bnf.org.
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# MEMBERSHIP OF THE FACULTY OF FAMILY PLANNING AND REPRODUCTIVE HEALTH CARE

The MFFP Examination consists of:

Part 1 (A or B) New format Multiple Choice Question paper (MCQ)

**Part 1A Examination:** For those who have not passed the Part 1 MRCOG nor received exemption from Part 1 MRCOG. This 2-hour paper consists of 60 MCQs based on basic, applied and clinical science.

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Part 1 (A and B) examinations will be held on Friday 21 October 2005 (applications must be received by 1 July 2005).

## Part 2 Examination (Dissertation or Case Reports)

☐ Part 2 - Dissertation or Case Reports

Submission of one Dissertation (10 000 words) or two Case Reports (2500  $\pm$  500 words each).

Approval of the Dissertation or Case Reports titles by the Dissertation/Case Reports Convenor **must be obtained before** the candidate starts work on the Dissertation or Case Reports **and before** the candidate applies to sit the Part 2 (CRQ, MEQ, OSCE) component. Guidance notes and proposal form, plus exemption form/information, are available on request (see below).

## Part 2 Examination (CRQ, MEQ, OSCE)

☐ Part 2 – CRQ, MEQ, OSCE

Critical Reading Question examination paper (CRQ)

Modified Essay Question examination paper (MEQ)

Objective Structured Clinical Examination (OSCE)

Applications for the **Part 2** held in **June 2006** must be received by **1 December 2005**.

Please consult the revised Examination regulations for changes to entry requirements.

The qualification is subject to re-certification every 5 years.

Revised regulations (June 2004), application forms and dissertation documents are available on application to: Miss Denise Newell, Examination Secretary, Faculty of Family Planning and Reproductive Health Care of the Royal College of Obstetricians and Gynaecologists, 27 Sussex Place, Regent's Park, London NW1 4RG, UK. Tel: +44 (0) 20 7724 5629. Fax: +44 (0) 20 7723 5333. E-mail: denise@ffprhc.org.uk. Website: www.ffprhc.org.uk