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Title: Climacteric: the journal of the International Menopause Society.

Title Abbrev: Climacteric

Citation: 2000 Sep;3(3):153-4

Article: How should we give progestogen?

Author: Sturdee D

NLM Unique ID: 9810959 Verify: PubMed

PubMed UI: 11910615

ISSN: 1369-7137 (Print)

Publisher: Parthenon Pub., New York:

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Received: Jul 02, 2006 (03:40 PM EST)

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Editorial

How should we give progestogen?

David W. Sturdee Editor-in-Chief

Side-effects from hormone replacement therapy (HRT) have always been one of the main reasons for patient dissatisfaction, with the progestogen component more often being the cause of this. Such problems are more common with sequential regimens and better long-term compliance is achieved with continuous combined therapy regimens^{1,2}. Additional progestogen is obligatory for women with a uterus to protect the endometrium from hyperplasia and carcinoma, but how should it best be given? We have the option of giving progestogen by the oral, transdermal, vaginal or rectal routes, and each has its particular merits. However, as the endometrium is the only tissue where the effects of progestogens are necessary, it is logical to deliver the hormone directly into the uterine cavity. Intrauterine contraceptive devices that release progestogen have been tried for many years and currently the Mirena® intrauterine system (IUS) is available in several countries. However, the additional benefit, a significant reduction in the menstrual blood loss, means that it is becoming used increasingly for controlling menorrhagia as well. Using an IUS for postmenopausal women as part of their HRT regimen is a natural progression. The development of intrauterine progestogen systems is comprehensively reviewed in this issue by Frits Riphagen3. It is likely that this route of administration will prove to be most effective and acceptable, especially for long-term continuous combined therapy.

In previous issues, in Editorials^{4,5}, original scientific articles^{6,7} and a review⁸, this Journal has highlighted concerns about the exaggerated and

unsupported claims for the benefits of various complementary medicines, and, in particular, for phytoestrogens. Another controlled study of phytoestrogens in this issue, by Kotsopoulos and colleagues9, reports that the effect of soy supplement on menopausal symptoms is no different to that of a placebo. A further complementary medicine for which much has been claimed, particularly in relation to osteoporosis prevention, is natural progesterone cream 10, but again there are no validated data. Using such a preparation alone with the intention of protecting the skeleton is unlikely to cause any harm, but some women are now using it as part of their HRT regimen, in the hope that it will avoid some of the progestogenic side-effects. But there are no data to show that this preparation will protect the endometrium from the proliferative effect of estrogen, and this is clearly demonstrated in the study by Barry Wren and colleagues11, also in this issue. Furthermore, the very low levels of progesterone in the plasma suggest that any significant clinical effect would be unlikely.

Unfortunately, our patients do not read these papers and anyway often seem more ready to accept the attractive anecdotal promotions in the media. We have a responsibility to inform our patients about the relative risks and benefits of hormone therapies, but some of the media will have a different agenda and motive in publishing unvalidated advertisements. What are we going to do about this?

We welcome correspondence on this and any other subjects relevant to the Journal.

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B. G. Wren

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