Reduction in the Size of a Uterine Leiomyoma following Discontinuation of an Estrogen-Progestin Contraceptive

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Abstract
The effects of estrogen-progestin oral contraceptives on the volume of uterine leiomyomata is not well characterized. In this case report, a 45-year-old woman with a symptomatic uterine leiomyoma was observed to have a 47% reduction in myoma volume after discontinuation of an oral contraceptive. The volume of uterine leiomyomata may be influenced by oral contraceptives.

Introduction
Uterine leiomyomata are the most common pelvic tumors of women. The etiology of uterine leiomyomata is not completely characterized, but it appears that these tumors arise after a somatic mutation in a progenitor cell [1]. Uterine leiomyomata are monoclonal [2], and many are cytogenetically abnormal [3]. Genes in the high mobility group family, including HMGIC [4], are mutated in some leiomyomata.

Estrogen and progestins influence the volume of uterine leiomyomata. Administration of the gonadotropin-releasing hormone (GnRH) agonist leuprolide acetate suppresses circulating estradiol and progesterone and is associated with an approximately 45% reduction in mean uterine volume in women with leiomyomata [5]. In women with leiomyomata being treated with a GnRH agonist, the simultaneous administration of estrogen or progestin can block the decrease in leiomyomata and uterine volume that would be expected with GnRH agonist treatment [6, 7]. The effects of estrogen-progestin oral contraceptives on the uterine leiomyomata volume are not well studied.

Case Report
A 45-year-old woman presented with right lower quadrant pain. The patient received regular gynecologic care and had no chronic medical problems. She had taken a combination estrogen-progestin contraceptive containing 0.35 µg of ethinyl estradiol and 0.35 mg of norgestrel for more than 5 years. She had regular menses, of normal flow and duration. She did not smoke cigarettes. The pain developed over a few days and was crampy. She had no uterine bleeding. Examination of the abdomen demonstrated no masses and no peritoneal signs. Pelvic examination revealed a normal vulva and vagina. The cervix was normal. The uterus was enlarged to a size corresponding to 12-14 gestational weeks, with a mass at the right fundal...
Discussion

The size of uterine leiomyomata is influenced by estradiol and progesterone. In a hypoestrogenic state (menopause or treatment with a GnRH agonist), the administration of estrogens and progestins can cause an increase in uterine volume in women with leiomyomata. Sener et al. [8] randomized menopausal women with small asymptomatic uterine leiomyomata to receive 1 of 2 hormone replacement regimens. The women who were randomized to receive transdermal estradiol (50 µg) and medroxyprogesterone acetate (5 mg daily) had a 38% increase in the diameter of their largest myoma. Two of the 20 women on this dose of hormone replacement had exceptionally large increases (300 and 400%) in the diameter of the leiomyoma. This suggests that a minority of menopausal women are especially sensitive to the effects of exogenous estrogens and progestins on leiomyoma volume. A second group of women were randomized to receive conjugated equine estrogen (0.625 mg daily) and medroxyprogesterone acetate (2.5 mg daily). This hormone replacement regimen caused no change in the diameter of the largest leiomyoma. One explanation for this finding is that in a hypoestrogenic state, there is a relationship between the dose of exogenous estrogens and/or progestins and the magnitude of the change in leiomyoma diameter.

Carr et al. [7] randomized women with uterine leiomyomata to receive either leuprolide acetate depot (3.75 mg i.m. every 4 weeks) plus a placebo pill or leuprolide acetate depot plus medroxyprogesterone acetate (20 mg daily). The women who received the leuprolide acetate plus a placebo pill had a 26% reduction in uterine volume after 12 weeks of treatment. The women who received the leuprolide acetate plus medroxyprogesterone acetate had no change in the uterine volume after 12 weeks of treatment. This suggests that the progestin, medroxyprogesterone acetate, blocks the effect of a hypoestrogenic state on uterine volume in women with leiomyomata. The effects of estrogens and progestins on leiomyoma volume in premenopausal women is not well studied. A small number of case reports suggests that in an occasional woman with leiomyomata, treatment with estrogen-progestin oral contraceptives can be associated with an increase in uterine volume [9, 10]. However, given the small number of such cases which have
been reported, and the high prevalence of leiomyomata and oral contraceptive use, it is probable that only a minority of women with leiomyomata are likely to experience a change in uterine volume with oral contraceptive use. Randomized studies will be needed to definitively address this issue.

The case presented in this report is an example of a large change in uterine leiomyoma volume associated with the discontinuation of an oral contraceptive. In women on oral contraceptives, with clinically significant uterine leiomyomata, clinicians might consider the discontinuation of the oral contraceptive as low-risk intervention that might be associated with a decrease in uterine volume.

References


Uterine Myomata and Contraceptives

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