Genistein reduces angiogenesis and apoptosis in women with endometrial hyperplasia

Abstract: Endometrial hyperplasia without cytological atypia is commonly treated with progestins, but other treatments may be available with equivalent efficacy and reduced side effects. Here, we evaluate the effect of genistein aglycone on angiogenesis and apoptosis-related markers in women with endometrial hyperplasia. Premenopausals (n=38) with nonatypical endometrial hyperplasia were administered either genistein aglycone (54 mg/day, n=19) or norethisterone acetate (10 mg/day, n=19) on days 16–25 of the menstrual cycle and evaluated for 6 months. Biopsies were taken during hysteroscopy at baseline and 6 months, and symptoms including excessive uterine bleeding were assessed at baseline and 3 and 6 months following recruitment. The expression of angiogenesis (Vegf), epithelial (Egf and Tgf), and apoptosis-related (Bax, Bel-2, and Casp-9) molecules, were assessed in uterine biopsies at baseline and after 6 months of therapy. Follicle-stimulating hormone, luteinizing hormone, estradiol, SHBG, and progesterone levels were also measured. After 6 months, 42% of genistein aglycone-administered patients had a significant improvement of symptoms compared to 47% of norethisterone acetate subjects. No significant differences were noted in hormone levels for any treatment. Gene expression revealed a significant reduction in Vegf, Egf, and Tgf (p<0.05 versus baseline), and an increase in proapoptotic molecules (Bax and Casp-9), with a concomitant decrease in Bel-2 values (p<0.05) in both groups. These results suggest that genistein aglycone might be useful for the management of endometrial hyperplasia without atypia in women who cannot or do not wish to be treated with progestin.

Keywords: genistein, endometrial hyperplasia, Vegf, Bel-2, Bax, Casp-9

Introduction

Endometrial hyperplasia is defined as a glandular proliferation that results in an increase in the ratio of glands to stroma, and is most common in the premenopausal period. In about 12% of women during this time, chronically high levels of estrogen, not offset by adequate progesterone production (anovulation) may be responsible for endometrial tissue hyperplasias and the often-correlated symptoms of menometrorrhagia. Under these circumstances, the endometrium initially loses the capacity for secretory maturation, and subsequently, with the persistence of the estrogenic stimulus, can assume a morphology very similar to endometrial cancer.

Chronic exaggerated estrogenic stimulation may induce synthesis of its estrogen receptors (ERs) and also of progesterone receptors, which are thus present at high levels in the hyperplastic endometrium. Moreover, estradiol (E2) induces the synthesis of different mitogenic factors, including VEGF and its receptor, EGF, which are promoters of endometrial proliferation. In addition, a significant correlation...
between steroid-receptor expression and TGFβ expression has also been demonstrated. TGFβ acts as a cell-growth inhibitor, and ERα inhibits TGFβ signaling via a non-genomic mechanism.

In a recent study, we tested the therapeutic action of the isoflavone genistein on endometrial hyperplasia. Genistein in fact has a weak effect in stimulating the growth of the endometrium when administered in menopause, but in an environment rich in estrogen, it acts as an antiestrogen. Genistein binds the ER, either as an agonist or antagonist, acting as a selective ER modulator, with full agonistic activity on the α-receptor and partial agonistic on the β-receptor, but with a greater affinity on the latter. In that study, we showed that genistein and estrogens compete for ER binding, with a resultant significant decline in receptor expression. In addition, genistein administration has antiestrogenic activity, decreasing the expression of α-receptors, mainly present in the uterus, which results in regression of hyperplasia comparable to that seen with treatment with progestins.

In this study, we evaluated the gene expression of the known growth factors (Vegf, EGF, and Tgfβ) and of some of the key molecules that regulate apoptosis (Bax, Casp-9, and Bcl-2) which might be involved in the development of hyperplasia in premenopausal women. The aim of the study was to understand if the positive effects of genistein on endometrial hyperplasia are at least in part mediated through apoptotic and growth-related pathways.

Subjects and methods
Between January and November 2010, 38 consecutive premenopausal women with abnormal uterine bleeding and diagnosed with nonatypical endometrial hyperplasia were enrolled in this observational study. All procedures were approved by the Ethics Committee of Messina University Hospital (approved December 2009, 47/09), and the protocol followed the principles of the Declaration of Helsinki. All study participants gave their written informed consent. All subjects were counseled by an expert gynecologist, and based on their own choice, received either genistein 54 mg daily (n=19; Genivis; Mastelli, Italy) or norethisterone acetate 10 mg/day on cycle days 16–25 (n=19). Subjects were treated for 6 months, or until the regression of hyperplasia (Figure 1).

Baseline laboratory measurements included follicle-stimulating hormone (FSH), luteinizing hormone (LH), E2, progesterone, and SHBG. During hysteroscopy, a biopsy sample was taken for pathology assessment and a small piece separately stored at −20°C for determination of messenger RNA (mRNA) expression of Vegf, Egf, Tgfβ, Bax, Casp-9, and Bcl-2.

E2, FSH, LH, SHBG, and progesterone were routinely evaluated as previously described. Symptomatic and pathological descriptions followed previously published criteria.

After 3 and 6 months, all women were asked for symptoms and changes in uterine bleeding, and an ultrasound evaluation of endometrial thickness was performed. After 6 months, all baseline data points were again measured.

For the gene-expression study, the extraction of total mRNA was performed from endometrial biopsies under sterile conditions using Trizol (Invitrogen, Italy) following the manufacturer’s protocol and evaluated by quantitative real-time polymerase chain reaction (PCR). For each sample, 5 mg of mRNA was reverse-transcribed into complementary deoxyribonucleic acid (cDNA) and 3 mL of cDNA was amplified in duplicate using the TaqMan Universal PCR Master Mix containing primer and TaqMan validated probes designed to specifically target human Vegf, Egf, Tgfβ, Bax, Casp-9, and Bcl-2 (all reagents from Applied Biosystems, USA). For the reaction, we used an SDS 7300 RealTime PCR instrument (Applied Biosystems). The result was expressed as number of copies of the target gene compared to the housekeeping gene (Actb), and the 2^{-ΔΔCt} mean values of both groups were compared with those of an arbitrary calibrator.

Statistical analysis was performed with SPSS 11 (SPSS, USA). To compare data between groups, Student’s t-test
was used for parametric data, and the Mann–Whitney U-test for nonparametric data. The significance of difference was assessed for parametric data by a two-way repeated-measures analysis of variance, followed by post hoc analyses where indicated. Dichotomous variables were analyzed with the $\chi^2$ test and the Fisher’s exact test, when appropriate. A $P$-value $<0.05$ was considered statistically significant.

### Results

The two study groups (Figure 1) showed similar characteristics regarding age (genistein, 47.2±3.1 years; norethisterone acetate, 47.2±3.4 years), body mass index (genistein, 24±1.1; norethisterone acetate, 23.5±1.5), parity (genistein, 1.3±1.2; norethisterone acetate, 1.2±1.1), and endometrial thickness evaluated at days 8–10 of the menstrual cycle (genistein, 7±3 mm; norethisterone acetate, 7±2.6 mm). After 3 months, ten genistein subjects had significant improvement in bleeding, seven had moderate improvement, and two subjects had persistent symptoms.

In the norethisterone acetate-treated group, nine subjects had significant improvement in bleeding, five had moderate improvement, and four subjects had persistent symptoms. During this study period, one subject was withdrawn because she started antihypertensive therapy.

At the second follow-up, the abnormal bleeding in eleven genistein-administered subjects was completely absent (further confirmed by histology in nine women), reduced in four, and persistent in three. In the norethisterone acetate-treated group, clinical improvement was reported in eleven subjects and a mild improvement in two, while four subjects showed persistent clinical symptoms. Histological examination of biopsies confirmed regression of hyperplasia in the eleven women from both groups, and its persistence in the remaining subjects. Two women with persistent bleeding, one in each group, decided to undergo to surgical ablation.

A nonparametric test showed a significant decrease ($P<0.05$) of symptoms in both genistein and norethisterone acetate groups compared to baseline after 6 months.

After 6 months of therapy, levels of $E_2$, FSH, LH, and SHBG were assayed. No significant differences were found for responders or nonresponders (data not shown). It is noteworthy that $E_2$ levels remained elevated in all nonresponders, while a tendency for decreased $E_2$ was observed for genistein-responsive subjects (baseline, 79.94±49.47 pg/mL; 6 months, 64.27±23.69 pg/mL; not significant). In the norethisterone acetate group, 14 subjects reported water retention, and none in the genistein group.

Regarding the expression of the genes studied in the endometrial tissue, we found that $Vegf$ was significantly reduced after 6 months of treatment in the genistein-treated group ($P<0.01$) and in the norethisterone acetate-treated group ($P<0.01$) compared to basal expression levels (Figure 2A). The reduction observed in the genistein group was significantly more pronounced than that observed in the norethisterone acetate-treated group ($P<0.05$). Expression of $Egf$ was also significantly reduced in the two study groups after 6 months of treatment ($P<0.005$, Figure 2B). Moreover, in the two groups, even gene expression of $Tgfb$ decreased significantly after 6 months ($P<0.005$, Figure 2C). Angiogenic, growth-related, and apoptotic molecule gene expression showed a significant change after 6 months. In particular, a significantly enhanced expression of the proapoptotic protein was observed in the genistein group.
molecules Casp-9 and Bax (Figure 3A and C, respectively) was noted after 6 months of treatment in both treatment groups \( (P<0.05 \text{ for both}) \). Finally, a concomitant decrease in the expression of Bcl-2 was observed after 6 months in both study groups \( (P<0.05, \text{Figure 3B}) \).

**Discussion**

The data presented here demonstrate a modulating action of genistein on the apoptotic and growth-related pathways in the hyperplastic endometrium. Clinical efficacy was essentially identical in the genistein- and progesterone-treated groups,
while the genistein group demonstrated a more favorable adverse-event profile.

In the presence of a hyperestrogenic environment in premenopausal women, with endometrial hyperplasia genistein showed a significant reduction in hyperplasia and related symptoms,\(^8\) compatible with the antiestrogenic effects of genistein that have previously been shown in vitro.\(^1^4\) It is known that more than 80% of hyperplasia cases without atypia and more than 50% of cases of complex hyperplasia with atypia spontaneously regress.\(^1^5,1^6\) For this reason, we further studied some of the possible molecular pathways involved in the mechanism of action of genistein aglycone in promoting the reduction of endometrial hyperplasia in subjects without signs of atypia. Growth factors involved in the establishment of endometrial hyperplasia act through kinase-associated receptors, and stimulate tissue and vascular growth. In this study, we demonstrated the inhibitory effect of genistein on the expression of Vegf, Egrf, and Tgfb, all of which are regulated by estrogens in the endometrium.\(^1^7,1^8\) Therefore, genistein acted in two different ways – reducing the cascade of kinase activation that follows growth-factor binding, and inhibiting the production of the mRNA for these growth factors – acting through ER\(\beta\), which is a known inhibitor of proliferation at the transcriptional level. We hypothesize that elevated estrogen levels promote enhanced ER activation, which is responsible for the activation of proliferative genes and the downregulation of antiproliferative genes. Therefore, pro/antiapoptotic homeostasis is altered in favor of cell survival and proliferation. Since deregulation of proliferation and apoptosis is known to contribute to neoplastic transformation and growth, maintaining the homeostasis of BAX, Bcl-2, and caspase 9 expression is of importance to avoid the onset of atypia in the hyperplastic endometrium. Genistein, by interfering with estrogen binding and receptor expression, is indirectly able to induce apoptosis and stop endometrial proliferation, as suggested by the observed increase in Casp-9 and Bax gene expression and Bcl-2 reduction.

The strengths of this paper are the well-defined population sample, the high adherence to the study protocol, and the demonstration of genistein’s molecular mechanism of action in this well-defined condition. A potential limitation of this report is the small sample of subjects observed and the lack of follow-up information after the end of treatment. However, our data are supported by solid demonstration of genistein’s mechanism of action, somewhat similar to progestin, but with a stronger safety profile.\(^1^9–2^1\)

The effects of genistein were studied on the same pathways that are known to be affected by progesterone-receptor activation, which have already been observed in different subsets by other authors, and our data are in agreement with their results.\(^1^5\)

**Conclusion**

The present study provides support for the possible use of genistein as an alternative therapy for those women that prefer to use a natural approach for the treatment of endometrial hyperplasia or those who have experienced undesirable side effects with progesterone therapy.

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**Author contributions**

All authors contributed toward data analysis, drafting and revising the paper and agree to be accountable for all aspects of the work.

**Disclosure**

The authors report no conflicts of interest in this work.

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