Direct and Indirect endocrine-mediated suppression of human endometrial CD8+ T cell cytotoxicity

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Regulation of endometrial (EM) CD8+ T cells is essential for successful reproduction and protection against pathogens. Suppression of CD8+ T cells is necessary for a tolerogenic environment that promotes implantation and pregnancy. However, the mechanisms regulating this process remain unclear. Sex hormones are known to control immune responses directly on immune cells and indirectly through the tissue environment. When the actions of estradiol (E2), progesterone (P) and TGFβ on EM CD8+ T cells were evaluated, cytotoxic activity, perforin and granzymes were directly suppressed by E2 and TGFβ but not P. Moreover, incubation of polarized EM epithelial cells with P, but not E2, increased TGFβ secretion. These findings suggest that E2 acts directly on CD8+ T cell to suppress cytotoxic activity while P acts indirectly through induction of TGFβ production. Understanding the mechanisms involved in regulating endometrial CD8+ T cells is essential for optimizing reproductive success and developing protective strategies against genital infections and gynecological cancers.

The mucosal immune system in the human female reproductive tract (FRT) is precisely regulated to optimize conditions for successful fertilization, implantation, and pregnancy while providing protection against sexually transmitted infections (STIs)1. Unlike other sites in the FRT, the endometrium must protect itself from sexually transmitted pathogens while allowing for the presence of allogeneic spermatozoa and supporting an immunologically distinct fetus. Key to this unique balance within the FRT are the sex hormones, estradiol (E2) and progesterone (P), which are secreted by the ovary and whose levels fluctuate during the menstrual cycle and thereby regulate multiple aspects of innate and adaptive immunity2–5. During the proliferative phase, E2 is present at low concentrations. However, E2 levels surge at ovulation, decline rapidly, before increasing again during secretory phase, when fertilization and implantation occur. In contrast, P levels remain low throughout the proliferative and ovulatory phases, before rising and peaking in mid-secretory phase. E2 and P act directly on FRT immune cells, epithelial cells and fibroblasts to regulate their function, and modulate the production of cytokines, chemokines and growth factors, which in turn regulate the innate and adaptive immune systems. These direct and indirect effects of sex hormones are essential for successful reproduction, and influence susceptibility to STIs in ways that are site-specific in the upper (endometrium, endocervix) and lower (ectocervix, vagina) FRT.

T cells are central to adaptive immune protection in the FRT and represent a major constituent of leukocytes (30–60%) in the endometrium6–9. To measure the cytotoxic potential of CD8+ T cells, White et al. evaluated the ability of FRT mixed cell suspensions to kill targets using a redirected lysis assay10. In addition to demonstrating cytotoxic activity throughout the FRT, these studies demonstrated that endometrial CD3+CD8+ T cell cytotoxic activity is markedly influenced by the stage of the menstrual cycle. Cytotoxic activity was elevated during the proliferative phase of the cycle and suppressed during the secretory phase. Interestingly, cytotoxic activity in the endometrium from postmenopausal women was significantly higher than activity at any point of the cycle in premenopausal women. Cycle dependent suppression occurred without changes in total CD8+ T cell numbers. More recently, using a time-lapse imaging system to measure direct killing by purified CD8+ T cells from the endometrium, we confirmed changes in cytotoxic activity during the menstrual cycle and increased activity following menopause11. Maximal suppression of CD8+ T cell cytotoxic activity in the endometrium coincided with the time when fertilization and implantation is likely to occur, and therefore may represent a mechanism to avoid recognition and rejection of sperm and fetal cells by maternal endometrial CD8+ T cells. 

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Results

Estradiol and TGFβ suppress cytotoxic activity in CD8+T cells from the endometrium. We have previously demonstrated that cytotoxic activity of EM CD8+T cells is suppressed during the secretory phase of the menstrual cycle and enhanced after menopause. In the present study, we investigated whether sex hormones are directly involved in the suppression of CD8+T cell cytotoxic activity. T cells were pretreated for 48 h with E2 or P at concentrations known to have physiological effects in vivo or with TGFβ for 2 h prior to measuring cell killing. To measure cytotoxic activity, control and treated CD8+T cells were co-cultured with CFSE-labelled allogeneic target cells, and the number of dead cells quantified by time-lapse imaging as described before. Based on our earlier studies for optimal killing, we selected an Effector:Target ratio of 1:1 and measured average cell killing during the first 6 h in culture. As seen in Fig. 1a–c, in representative time-course studies, relative to target cells alone, we detected a significant increase in dead target cells in the presence of allogeneic EM CD8+T cells. Moreover, when CD8+T cells were pretreated with E2 (Fig. 1a) or TGFβ (Fig. 1c), but not P (Fig. 1b), cell killing was markedly reduced. Figure 1d–f indicates the effect of hormones and TGFβ treatment on cytotoxicity measured both as total number of dead cells and as fold change in dead target cell numbers, the latter being done to compare experiments with different background mortalities of target cells alone. Consistent with our representative findings, relative to controls, E2 (Fig. 1d) and TGFβ (Figs. 1f), but not P (Fig. 1e), significantly reduced cell killing. Figure 1f confirms our earlier results that CD8+T cell cytotoxic activity is significantly reduced following TGFβ treatment of EM CD8+T cells.

EM CD8+T cell expression of perforin and granzyme A is inhibited by E2. To identify the underlying mechanisms involved in suppression of cytotoxic T cell activity, we evaluated the expression of perforin, granzyme A (GZA) and granzyme B (GZB) following preincubation of CD8+T cells with E2 or P. EM tissues were digested to obtain mixed cell suspensions and stained for intracellular cytokotoxic molecules prior to analysis by flow cytometry as previously described. Figure 2a contains representative histogram overlays of CD8+T cells for perforin, GZA and GZB indicating the downward shift in expression of perforin and GZA in response to E2 but not P. Analysis of five separate experiments with different subjects (Fig. 2b), demonstrated that E2 significantly reduced the intracellular content (MFI) of perforin and GZA in CD8+T cells. In contrast, E2 had no significant effect on GZB. Hormone treatment also affected the percentage of cells expressing cytotoxic molecules, as seen in Fig. 2c, with a significant decrease in the percentage of cells positive for perforin and GZA cells following incubation with E2. Overall, these studies demonstrate that E2 decreases the expression of perforin and GZA in EM CD8+T cells, which correlates with reduced cell cytotoxicity. Further, it indicates that these changes are hormone specific given that P had no effect on either perforin, GZA, and GZB intracellular expression or the percentage of positive cells.

EM CD103+ and CD103- CD8+T cell expression of perforin and granzyme A is inhibited by estradiol. In previous studies, we demonstrated that human EM CD8+T cells consist of tissue resident cells and non-resident CD8+T cells that are CD103+ and CD103-, respectively. When cytotoxicity was compared between CD103+ and CD103–CD8+T cells, direct cytotoxic activity and expression of cytotoxic molecules was lower in CD103+ compared to CD103–T cells. Therefore, in this study we explored whether the observed effects of E2 in reducing cytotoxic molecule expression were due to effects on both CD103+ and CD103–T...
cell populations. Mixed cell suspensions from EM tissues were incubated with E2 for 48 h prior to measuring granular content by flow cytometry. Figure 3a shows a representative example of the content of perforin, GZA and GZB before and after treatment with estradiol. As shown in Fig. 3b, using matched samples from multiple patients, E2 significantly suppressed the content of perforin and GZA (MFI) in both tissue resident (CD103+) and non-resident (CD103−) CD8+T cells. In contrast, E2 had no effect on GZB. Interestingly, as seen in Fig. 3c, E2 reduced the percent positive CD103+ and CD103−CD8+T cells containing perforin but had no effect on the percentage of cells that were GZA or GZB positive. These findings suggest that E2 regulates both CD103+ and CD103−CD8+T cells and that suppression of the levels of cytotoxic molecules is unique in that each of the cytotoxic responses to E2 are separate and distinct. This is consistent with our previous results that menopausal status equally affects resident and non-resident T cells11, suggesting that tissue residency does not modify sensitivity to hormonal control.

Perforin, granzyme A and B Content in EM CD8+T Cells is suppressed by TGFβ. Next, we investigated the underlying mechanisms of TGFβ-mediated cytotoxicity suppression. CD8+T cells in mixed cell suspensions were incubated with TGFβ prior to perforin, GZA, and GZB analysis by flow cytometry at 48 h. As seen in the histograms (Fig. 4a) and analyses of intracellular content (Fig. 4b), TGFβ significantly reduced the concentration of all three intracellular cytotoxic molecules. In our patients we observe a wide range of perforin expression from 0.5 to 32.6% as shown in this study and in our previous publication11. The representative plots presented in Figs. 2 and 4 are from two different patients. Figure 2 is from a patient with low perforin expression, while Fig. 4 is from a patient with high perforin expression. These profiles are examples that reflect the range in expression profiles between patients. We also found that TGFβ reduced the percentage of total CD8+T cells that were perforin positive, but had no effect on the percentage of cells expressing GZA and GZB (Fig. 4c). We next investigated the effect of TGFβ on cytotoxic molecules in CD103+ and CD103−CD8+T cell compartments (Fig. 5a). As seen in Fig. 5b, TGFβ significantly reduced the expression of perforin, GZA and GZB within both CD103+ and CD103−CD8+T cell populations. Interestingly, TGFβ reduced the number of perforin positive cells but had no effect on cells positive for GZA or GZB (Fig. 5c).

Progesterone increases the basolateral secretion of TGFβ by polarized EM epithelial cells. In addition to their directs effects, sex hormones are known to indirectly regulate immune cells by acting on adjacent cells in the tissue environment. TGFβ is known to be synthesized and secreted by EM epithelial cells and stromal cells from the uterus18,19. Our studies in a rodent model demonstrated that polarized EM epithelial cells preferentially secrete biologically active TGFβ into the basolateral compartment18. Therefore, we analyzed TGFβ production by polarized human EM epithelial cells. As seen in Fig. 6a, TGFβ was secreted constitutively by polarized epithelial cells with preferential release basolaterally, in the direction of underlying tissues. Since

Figure 1. Effect of E2, P and TGFβ on endometrial CD8+T cells cytotoxicity. Purified endometrial CD8+T cells (effector cells, E) were pre-treated with E2 (5 × 10−8 M), P (1 × 10−7 M) for 48 h or TGFβ (10 ng/ml) for 2 h before co-culture with CFSE-stained allogeneic blood CD4+T cells (target cells, T) using a E:T ratio of 1:1. Cytotoxicity was measured over time by using quantitative time-lapse microscopy. Representative time-course of the kinetics of cytotoxicity over a period of 6 h in the absence (control) or presence of E2 (a), P (b) and TGFβ (c). Target alone are allogeneic blood CD4+T cells. Comparison of E2 (d, n = 7), P (e, n = 5) and TGFβ (f, n = 16) treatment in the mean number of dead target cells (left graph) and fold change in the mean number of dead target cells after E2, P and TGFβ treatment compared to control (right graph). Each dot represents a different patient. Min to Max (d–f, right graph) are shown. *P<0.05; ** P<0.01; Wilcoxon matched-pairs signed rank test.
we have previously demonstrated that human polarized epithelial cells are hormonally responsive to E2, we hypothesized that E2 might increase TGFβ secretion by epithelial cells as a secondary mechanism for regulating CD8+T cell cytotoxic activity. To test this, polarized EM epithelial cells were grown to confluence and incubated

**Figure 2.** Effect of E2 and P on intracellular cytotoxic molecules in endometrial CD8+T cells. Mixed cell suspensions from endometrial tissues were pre-treated with E2 (5 × 10^{-8} M), P (1 × 10^{-7} M) for 48 h and stained for the intracellular cytotoxic molecules perforin, granzyme A and granzyme B for analysis by flow cytometry. (a) Representative histograms of intracellular perforin (left), granzyme A (middle) and granzyme B (right) expression in endometrial CD8+T cells treated with E2 and P. Gate boundaries (horizontal line) were set by fluorescence minus one (FMO) to indicate the gating of positive cells. (b) Median fluorescence intensity (MFI) of perforin (left), granzyme A (middle), granzyme B (right) and (c) percentage of perforin+ (left), granzyme A+ (middle), granzyme B+ (right) in endometrial CD8+T cells following incubation with E2 and P. Each dot represents a different patient (n = 5). *P < 0.05; Friedman test with Dunn’s multiple comparisons test.
Figure 3. Effect of E2 on intracellular cytotoxic molecules in endometrial CD103+ and CD103−CD8+T cells. Mixed cell suspensions from endometrium were pre-treated with E2 (5 × 10^{-8} M) for 48 h and stained for the intracellular cytotoxic molecules perforin, granzyme A and granzyme B for analysis by flow cytometry. (a) Representative dot plots of intracellular perforin (left), granzyme A (middle) and granzyme B (right) expression in endometrial CD103+ and CD103−CD8+T cells treated with E2. Top row shows control group (without E2) and bottom row shows E2 treated group. (b) Median fluorescence intensity (MFI) of perforin (left), granzyme A (middle), granzyme B (right) and (c) percentage of perforin+ (left), granzyme A+ (middle), granzyme B+ (right) in endometrial CD103+ and CD103−CD8+T cells following incubation with E2. Each dot represents a different patient (n = 5). *P < 0.05; Friedman test with Dunn’s multiple comparisons test.
with E2 or P for 48 h prior to collection of basolateral media. Unexpectedly, as shown in a representative experiment (Fig. 6b), we found that P increased the basolateral secretion of TGFβ relative to controls and that E2 had no effect on TGFβ production. To more fully analyze this response, we normalized the data from 6 experiments to express TGFβ production as the fold change relative to control cells, to compare experiments with different

Figure 4. Effect of TGFβ on intracellular cytotoxic molecules in endometrial CD8+T cells. Mixed cell suspensions from endometrial tissues were pre-treated with TGFβ (10 ng/ml) for 48 h and stained for the intracellular cytotoxic molecules perforin, granzyme A and granzyme B for analysis by flow cytometry. (a) Representative histograms of intracellular perforin (left), granzyme A (middle) and granzyme B (right) expression in endometrial CD8+T cells treated with TGFβ. Gate boundaries (horizontal line) were set by fluorescence minus one (FMO) to indicate the gating of positive cells. (b) Median fluorescence intensity (MFI) of perforin (left), granzyme A (middle), granzyme B (right) and (c) percentage of perforin+ (left), granzyme A+ (middle), granzyme B+ (right) in endometrial CD8+T cells following incubation with TGFβ. Each dot represents a different patient (n = 7). *P < 0.05, Wilcoxon matched-pairs signed rank test.
Figure 5. Effect of TGFβ on intracellular cytotoxic molecules in endometrial CD103+ and CD103−CD8+ T cells. Mixed cell suspensions from endometrial were pre-treated with TGFβ (10 ng/ml) for 48 h and stained for the intracellular cytotoxic molecules perforin, granzyme A and granzyme B for analysis by flow cytometry. (a) Representative dot plots of intracellular perforin (left), granzyme A (middle) and granzyme B (right) expression in endometrial CD103+ and CD103−CD8+ T cells treated with TGFβ. Top row shows control group (without TGFβ) and bottom row shows TGFβ treated group. (b) Median fluorescence intensity (MFI) of perforin (left), granzyme A (middle), granzyme B (right) and (c) percentage of perforin+ (left), granzyme A+ (middle), granzyme B+ (right) in endometrium CD103+ and CD103−CD8+ T cells following incubation with TGFβ. Each dot represents a different patient (n = 7). *P < 0.05; Friedman test with Dunn’s multiple comparisons test.
secretion backgrounds by control cells. As shown in Fig. 6c, following normalization, P but not E2 consistently stimulated the secretion of TGFβ by EM epithelial cells. Overall, these studies suggest that during the secretory phase menstrual cycle, under the influence of progesterone, EM epithelial cells secrete increased amounts of TGFβ basolaterally into the tissue compartment to suppress CD8+T cell cytolytic activity.

Discussion
Our study demonstrates that E2 and TGFβ directly suppress EM CD8+T cell cytotoxic activity by downregulating intracellular perforin, GZA, or GZB expression. Further, we demonstrate that whereas E2 acts directly on CD8+T cells to suppress killing, P has no direct effect. However, P upregulates TGFβ secretion by EM epithelial cells, which could represent an indirect mechanism of control of cytotoxic activity in vivo. Overall, as depicted in Fig. 7, our findings demonstrate direct and indirect hormone-dependent mechanisms in the human endometrium able to suppress CD8+T cell cytolytic capacity in premenopausal women during the secretory phase of the menstrual cycle, without which successful implantation and pregnancy would be unlikely to occur.

CD8+T cells are key effector cells for immune protection, but their function in the endometrium is tightly regulated to avoid rejection of the semi-allogeneic fetus. Our previous studies demonstrate that EM CD8+T cell cytotoxic activity is suppressed in premenopausal women compared to postmenopausal women, particularly during the secretory phase of the menstrual cycle, suggesting that sex hormones are key regulators of CD8+T cell cytotoxicity. However, the direct effect of sex hormones in suppression of cytotoxic function of EM CD8+T cells had never been addressed before. Therefore, we investigated whether sex hormones directly contribute to the suppression of CD8+T cell cytotoxic activity seen in premenopausal women.

We found that E2 directly suppresses cytotoxic activity of EM CD8+T cells, and that the reduction in killing capacity is due to decreased expression of the cytotoxic molecules perforin, GZA, or GZB in CD8+T cells. In contrast to our studies which showed no effect of E2 on GZB expression, Navarro et al. found that E2 increased the expression of GZB in blood. Differences observed are most likely due to their use of buffy-coat blood suspension containing CD8+T cells that were retrovirally transduced and activated, whereas we used purified EM CD8+T cells without any in vitro activation stimulus to measure the effects of E2. Furthermore, to test GZB production, they used a preparation of total T cells, containing both CD4+ and CD8+T cells, while we used purified CD8+T cells. Whether activation status alters EM CD8+T cell responsiveness to E2 remains to be determined.

In contrast to E2, we found that P had no direct effect on EM CD8+T cell cytotoxic activity or cytotoxic granule content. This lack of effect is probably due to the previously reported absence of nuclear P receptors in CD8+T cells and contrasts with findings showing P changes in cytokine production and reduction in T-cell proliferation following P treatment of peripheral blood CD8+T cells, which the authors attributed to P membrane

Figure 6. Constitutive production and regulation of TGFβ secretion in endometrial epithelial cells.
Endometrial epithelial cells were isolated and cultured until confluence and high transepithelial resistance was reached. Polarized endometrial epithelial cells were treated with E2 (5 × 10⁻⁸ M) or P (1 × 10⁻⁷ M) for 48 h. Apical and basolateral conditioned media (CM) were collected and assayed total TGFβ by ELISA. (a) Level of total TGFβ secreted following 48 h accumulation of constitutive apical and basolateral (n = 14). (b) Representative experiment showing the basolateral secretion of TGFβ by polarized endometrial epithelial cells following treatment with E2 or P for 48 h. (c) Graph represents the fold change in total TGFβ in basolateral CM following E2 or P treatment compared with untreated controls (n = 6). The mean and SEM (a and b) and Min to Max (c) are shown. *P < 0.05; **P < 0.01; Wilcoxon matched-pairs signed rank test (a) and Friedman test with Dunn's multiple comparisons test (b and c).
These differences suggest a finite specificity of P effects that are unique to the class of P receptor present or to functional differences between peripheral blood CD8+T cells and EM CD8+T cells. Confirming our previous results, we found that EM cytotoxic function was rapidly suppressed by TGFβ after short-time treatment. In the present study, we demonstrate that the suppressive effects of TFGβ are due to a reduction in the content of intracellular cytotoxic molecules (perforin, GZA and GZB). Given that in our experimental setting, cells were pre-treated with TGFβ for 2 h prior to cytotoxicity analysis and 48 h prior to intracellular cytotoxic molecule assessment, the underlying mechanism for rapid regulation of cytotoxic capacity remains unclear, and further investigation is warranted. Others have demonstrated that perforin and granzymes work synergistically to induce apoptosis in target cells recognized by CD8+T cells. In addition to reductions in GZA and GZB, our findings show a reduction in both the levels of intracellular content of perforin as well as the percent positive cells. This is consistent with reports by others showing that cytotoxic function is suppressed by TGFβ through the reduction of perforin positive CD8+T cell numbers in the gastrointestinal tract, and may indicate a general mechanism by which TGFβ controls cytotoxic activity of CD8+T cells in tissues. To the best of our knowledge, our findings are the first to demonstrate that E2 and TGFβ in culture reduce both the number of percent perforin positive CD8+T cells as well as the intracellular content of cytotoxic molecules of these cells. We and others have demonstrated that TGFβ is produced by epithelial cells and fibroblasts from the FRT and under hormonal control in a rodent model. Interestingly, in the human, TGFβ mRNA expression in the endometrium is increased during the secretory phase of the menstrual cycle, when EM CD8+T cells cytotoxic activity is maximally suppressed. Our current study extends these findings by demonstrating that P acts directly on human EM epithelial cells to stimulate the secretion of TGFβ. Interestingly, we found that E2 had no
effect on epithelial cell TGFβ production, in contrast to our previous findings with the rodent model. Consistent with our results, early studies in non-human primates also found that P increased mRNA of TGFβ2 in the endometrium. We have previously demonstrated broad effects of E2 treatment on epithelial cells, including stimulation of intracellular genes, such as Cytosolic 5'-nucleotidase and P receptor, and the secretion of SLPI along with other antimicrobial peptides with broad antimiicrobial, antiviral and anti-fungal activity. Our findings in the present study highlight the selectivity of hormonal effects on epithelial cells. Whether after its secretion in response to P, other hormonally dependent steps further process TGFβ to increase biological activity remains to be determined. Taken together, these findings suggest E2 and P precisely regulate adaptive immunity in the endometrium, by acting indirectly on epithelial cells to increase the production of TGFβ.

The menstrual cycle is divided into a proliferative and a secretory phase, during which fertilization and implantation occurs. Others have shown that shortly after ovulation, during the secretory phase of the menstrual cycle, levels of P rise sharply and are sustained for 10–12 days. Accompanying this increase in P is a secondary mid-secretory rise in E2 which is essential for implantation. In contrast, the proliferative phase of the cycle is dominated by E2. These hormonal changes control the cyclical remodeling of the endometrium in premenopausal women to optimize conditions for implantation. Based on our findings in the present study, we propose that in premenopausal women, E2 directly suppresses cytotoxic activity by EM CD8+T cells and reduces the intracellular content of perforin and GZβ. With ovulation and unique to the endometrium, P induces the production of TGFβ by epithelial cells, which in turn acts on EM CD8+T cells to further suppress cytotoxic activity, and reduce the expression of perforin, GZA and GZB, achieving maximum suppression during the secretory phase of the menstrual cycle (Fig. 7).

In our previous studies, we demonstrated that cytotoxic activity is significantly increased in postmenopausal women. Following menopause, sex hormones levels (E2 and P) drop and remain low and constant. Our results in this study suggest that the reduction in sex hormones removes the direct and indirect suppressive mechanisms from EM CD8+T cells, resulting in an increase in cytotoxic activity and intracellular content of cytotoxic molecules, consistent with our previous findings.

We and others have shown that CD8+ T cells in the endometrium consist of both CD103+ and CD103− T cells, likely representing tissue resident and non-resident memory T cells respectively. We demonstrated that cytotoxic activity and intracellular content of cytotoxic molecules was significantly higher in CD103− compared to CD103+ EM CD8+ T cells. Here we investigated whether the regulatory effects of E2 and TGFβ were preferential to either subset. We found that E2 and TGFβ affected CD103+ and CD103− CD8+ T cells equally and reduced the content of cytotoxic molecules in both subsets. These findings suggest that tissue-resident and non-resident CD8+ T cells are equally susceptible to regulation by sex hormones, and that transition into tissue-residency status does not modify sensitivity to regulation by sex hormones and the tissue environment. Our results here are also in agreement with our previous observation that postmenopausal women tended to have decreased expression of GZβ and increased expression of GZA compared to premenopausal women in both CD103− and CD103+ CD8+ T cells. This would suggest that removal of sex hormone suppression after menopause induces changes in intracellular content of cytotoxic molecules in both cellular subsets.

Given that cytotoxic activity following menses returns during the follicular phase of the menstrual cycle, suggests either that the absence of P and TGFβ reverses intracellular granular content and cytolytic activity or that a new population of CD8+ T cells enters the endometrium. That cytolytic potential can be restored was suggested from studies in which in vitro neutralization of TGFβ partially restored perforin expression in gut CD8+ T cells. Moreover, using an inhibitor that blocks TGFβ signaling in EM CD8+ T cells, we rescued cytotoxic activity in some samples. Whether cytolytic potential is restored as hormone levels change with the menstrual cycle or the result of recruitment of CD8+ T cells to the endometrium with the start of a new menstrual cycle remains to be determined. Further elucidation of EM CD8+T lymphocyte (CTL) activity and its implications for successful reproduction and protection against sexually transmitted diseases, including HIV, will provide essential information for the development of vaccines and immunotherapies targeted to the FRT.

Regarding study limitations, in addition to the mechanisms addressed in our study, alternative mechanisms of regulation of cytotoxic activity by sex hormone remain to be explored. For example, while we detected significantly lower levels of cytotoxic molecules following E2 treatment, whether the observed effects were due to direct actions of E2 on the production of cytotoxic molecules or secondary to decreased activation status of CD8+ T cells remains to be determined. Further, in addition to the perforin/granzyme pathway, the FAS/FASL pathway also mediates cell death. In preliminary studies we have observed no modification of sFASL production by EM CD8+ T cells after incubation with hormones under resting conditions. Further research is needed to address whether the FAS/FASL pathway is significantly modified by E2 after T cell stimulation, and if so, the potential contribution of the FAS/FASL pathway to cytotoxic function of EM CD8+T cells.

Another important aspect not addressed in this study is the extent to which sex hormones modify degranulation of T cells. In previous studies, we demonstrated a significant increase in degranulation capacity of CD8+ T cells from the FRT after menopause, suggesting a potential implication of sex hormones in regulation of degranulation capacity. Future studies are needed to investigate the extent to which E2 and P directly modify degranulation of CD8+ T cells. Furthermore, in addition to cytotoxic molecules, FRT CD8+ T cells can secrete array of cytokines and chemokines upon stimulation, including IFN-γ and TNFα. In preliminary studies, we observed no differences in secretion of IFN-γ and TNFα in CD8+ T cells treated with E2 or P under resting conditions; however, whether cytokine production after stimulation is modified by sex hormones remains to be addressed in future studies.

Although our study is focused on CD8+ T cells, a full understanding of the regulation of cytotoxic activity in the FRT during the menstrual cycle will require the functional evaluation of CD4+ T cells and NK cells. We have previously demonstrated decreased proportions of CD4+ T cells in the EM from postmenopausal women. However, whether E2 and P modify the presence and function of EM CD4+ T cells throughout the menstrual cycle remains to be explored.
cycle is unknown. Endometrial NK cells, which have overlapping cytotoxic functions with CD8+ T cells, increase in numbers from approximately 3% of leukocytes in proliferative phase to 30% in secretory phase tissues where they surround the spiral arteries, due to increased chemotaxis and proliferation. Whether sex hormones directly influence cytotoxic function of endometrial NK cells is unclear. Previous studies using peripheral blood NK cells have shown contradictory evidence of a menstrual cycle effect on cytotoxicity. However, others have demonstrated that endometrial NK cells in the early proliferative phase have lower cytotoxicity compared to other stages of the menstrual cycle. Together these findings suggest that sex hormones modulate cytotoxic activity of multiple cell types as part of a larger program to optimize conditions in the FRT for successful reproduction.

In conclusion, our study demonstrates two unique hormone-dependent regulatory mechanisms that regulate CD8+ T cell cytotoxic function in the uterine endometrium, one which involves the direct effect of E2 and the other, the indirect effect of P-mediated epithelial cell production of TGFβ. In response to E2 and TGFβ, CD103+ and CD103− CD8+ T cell cytolytic potential is suppressed. These results are central to our understanding of hormonal regulation of cytotoxic activity during the menstrual cycle, during pregnancy and following menopause, and to the development of therapeutics for both the prevention and control sexually transmitted infections and gynecological cancers.

**Methods**

**Study subjects.** Studies were performed with Dartmouth College Institutional Review Board approval. All investigations were conducted according to the principles expressed in the Declaration of Helsinki and carried out with the approval to use discarded tissues from hysterectomies from the Committee for the Protection of Human Subjects (CPHS), Dartmouth-Hitchcock Medical Center, and with written informed consent obtained from the patients before surgery. Indications for surgery were benign conditions such as fibroids and prolapse (age from 31 to 82); tissue samples selected were distant from the sites of pathology and without pathological lesions as determined by a pathologist. Women were not on oral contraceptives before hysterectomy. Menopausal status was determined by a pathologist based on the histological evaluation of sections of the endometrium (endometrial dating). Post-menopausal status was characterized by an atrophic endometrium. Information regarding genital infections was not available. Blood Leuko Paks from women were obtained from our IRB-approved collection facility at Dartmouth–Hitchcock Medical Center. Blood donors were anonymous, no information regarding age or hormonal status was available and only female donors were used in this study.

**Tissue processing.** EM tissues were transferred to the laboratory immediately after surgery and processed as previously described. Average tissue weight obtained was 2.6 ± 2.0 g. Tissues were rinsed with 1× HBSS (Hanks balanced salt solution) supplemented with phenol red, 100 U/ml penicillin, 100 µg/ml streptomycin (all Thermo Scientific HyClone, Logan, UT), and 0.35 mg/ml NaCO3 (Fisher Scientific, Pittsburgh, PA). Tissues were then minced under sterile conditions into 1–2 mm fragments and digested using an enzyme mixture containing 0.05% collagenase type IV (Sigma-Aldrich, St. Louis, MO) and 0.01% DNAse (Worthington Biochemical, Lakewood, NJ) for 1 h at 37°C. Type IV collagenase was selected based on preliminary studies to ensure non-cleavage of surface markers. After digestion, cells were dispersed through a 250-µm nylon mesh screen (Small Parts, Miami Lakes, FL), washed, and resuspended in complete medium consisting of DMEM/F12 medium without phenol red (Invitrogen Life Technologies, Carlsbad, CA) supplemented with 20 mM HEPES, 2 mM L-glutamine (Invitrogen Life Technologies, Carlsbad, CA), and 10% heat-inactivated defined Fetal Bovine Serum (FBS) (HyClone, Logan, UT, USA). Epithelial cell sheets were recovered from 20-µm mesh filters by rinsing and backwashing the filter with complete medium, centrifuged at 500 × g for 5 min and analyzed for cell number and viability. To establish a cell culture system of polarized human EM epithelial cells with both apical and basolateral compartments, EM epithelial cells were cultured in Matrigel matrix (BD Biosciences) coated Falcon cell culture inserts in 24-well companion culture plates (Fisher Scientific). Apical and basolateral compartments contained 300 and 500 µl of complete medium, respectively, which was changed every 2 days. Tight junction formation of epithelial cell monolayers from EM was assessed by periodically measuring transepithelial resistance (TER) using an EVOM electrode and Voltohmmeter (World Precision Instruments, Sarasota, FL), as described previously.

**Isolation and culture of EM epithelial cells.** Epithelial cell sheets were recovered from 20-µm mesh filters by rinsing and backwashing the filter with complete medium, centrifuged at 500 × g for 5 min and analyzed for cell number and viability. To establish a cell culture system of polarized human EM epithelial cells with both apical and basolateral compartments, EM epithelial cells were cultured in Matrigel matrix (BD Biosciences) coated Falcon cell culture inserts in 24-well companion culture plates (Fisher Scientific). Apical and basolateral compartments contained 300 and 500 µl of complete medium, respectively, which was changed every 2 days. Tight junction formation of epithelial cell monolayers from EM was assessed by periodically measuring transepithelial resistance (TER) using an EVOM electrode and Voltohmmeter (World Precision Instruments, Sarasota, FL), as described previously.

**Preparation of EM CD8+ T cells.** Following removal of dead cells from the filter pass through suspension of stromal cells, CD8+ T cells were isolated using negative magnetic bead selection with the CD8+ T cell isolation kit (Miltenyi Biotec, Auburn, CA) following instructions with minor modifications. This negative selection protocol delivers untouched CD3+CD8+ T cells. Additionally, anti-fibroblast microbeads (Miltenyi Biotec) were added in combination with the microbeads supplied with the kit to ensure depletion of stromal fibroblasts present in the mixed cell suspension as described before. After two rounds of negative selection, purity of the CD8+ T cell population was higher than 90%, with approximately 2% contamination with non-immune cells, 2% CD3- cells and 1–2% contamination with CD4+ T cells. The average number of total CD8+ T cells recovered per gram of tissue was 2 × 10⁶. Following isolation, purified CD8+ T cells were resuspended in immune cell media consisting of X-VIVO 15 Media (Lonza, Walkersville, MD) supplemented with 10% charcoal stripped human AB serum.
Preparation of blood CD4+T cells. Blood CD4+T cells were purified using negative magnetic beads selection with the CD4+T cell isolation kit (Miltenyi Biotech) following isolation of PBMC by standard Ficoll density gradient centrifugation\(^\text{16,50}\). Freshly isolated blood CD4+T cells were stained with CFSE (Cell Division Tracker Kit; BioLegend) as recommended by the manufacturer. The cells were then resuspended in immune cell media as allogeneic target cells prior to cytotoxicity assays.

Hormone preparation. 17β-estradiol (E\(_2\), Calbiochem, Gibbstown, NJ) or progesterone (P, Calbiochem) was dissolved in 100% ethanol for an initial concentration of 1 × 10\(^{-3}\) M, evaporated to dryness and suspended in epithelial cells complete media containing 10% charcoal dextran-stripped FBS or immune cell complete media to a concentration of 1 × 10\(^{-5}\) M. Further dilutions were made to achieve final working concentration, and cells were treated with 5 × 10\(^{-8}\) M E\(_2\) and/or 1 × 10\(^{-7}\) M P. Both are standard hormone treatment concentrations used by our laboratory and each is within the physiological range of hormone concentration\(^\text{20}\). As a control, an equivalent amount of ethanol without dissolved hormone was initially evaporated.

Sex hormone treatment and TGFβ analysis of EM epithelial cells. Epithelial cells in culture were switched to complete media containing 10% of charcoal dextran-stripped FBS prior to hormone treatment. After 24 h, the media was replaced, and cells were treated with hormone for 48 h. Hormone (E\(_2\) or P) or ethanol control media was added to both the apical and basolateral compartments. After 48 h incubation, apical and basolateral conditional media (CM) were collected and stored immediately at −80ºC prior to TGFβ ELISA (R&D systems, Minneapolis, MN) according to the manufacturer's recommendations.

Cytotoxicity assay. Purified and preincubation of CD8+T cells with E\(_2\), P or TGFβ were co-cultured with CFSE-stained blood CD4+T cells in immune cell media, at an Effector:Target ratio of 1:1, in 96 well plates. Cytotox red (IncuCyte Cytotox Red, Essen Bioscience) was added to the media to stain dead cells. Plates were imaged every 15 min using the IncuCyte Zoom system (Essen Bioscience), and dead target cells were automatically quantified over time as double green (CFSE) and red (Cytotox) stained cells\(^\text{11}\).

Flow cytometry. Mixed cell suspensions from EM tissues were treated with TGFβ, E\(_2\) or P for 48 h prior to cell staining for flow cytometry analysis. For experiments in which intracellular cytotoxic molecules were analyzed, mixed EM CD8+T cells were examined by flow cytometry to identify CD103+ and CD103− subsets as described previously\(^\text{11}\). Suspensions were stained for surface markers with combinations of the following antibodies: CD45-AF700, CD3-APC-Cy7, CD103-BV711 (BioLegend), CD4-PE-Cy5.5 (eBioscience, San Diego, CA), CD8-BUV395 (BD Bioscience). To detect of perforin, granzyme A (GZA) and granzyme B (GZB), following surface staining, cells were fixed and permeabilized with Cytofix/cytoperm kit (BD) according to instructions. Intracellular staining of perforin, granzyme A and B was done using combinations of the following antibodies: anti-human Perforin-PE/Dazzle, Granzyme A-AF647 (BioLegend) and Granzyme B-BV421 (BD Bioscience). Analysis was performed on BioRad ZE5 flow cytometers (BioRad) using Everest software, and data analyzed with FlowJo software version v10 (Tree Star, Inc. Ashland, OR, www.flowjo.com). Expression of surface and intracellular markers was measured by the percentage of positive cells and measured median fluorescence intensity (MFI).

Statistics. Data analysis was performed using the GraphPad Prism version 8.3.0 for Windows (GraphPad Software, San Diego, CA, www.graphpad.com). A two-sided \(P\) value < 0.05 was considered statistically significant. Comparison of two groups was performed with the non-parametric Wilcoxon matched-pairs signed rank test. Comparison of three or more groups was performed applying the non-parametric Friedman test with Dunn’s multiple comparisons test.

Image generation. Figures 1 and 6 were created using GraphPad Prism version 8.3.0 for Windows (GraphPad Software, www.graphpad.com). Figures 2, 3, 4 and 5 were created using FlowJo software version v10 (www.flowjo.com) and GraphPad Prism version 8.3.0 for Windows (GraphPad Software, www.graphpad.com). Figure 7 was created using BioRender.com.

Ethics statement. The studies were performed with Dartmouth College Institutional Review Board approval. Approval to use discarded tissues from hysterectomies was obtained from the Committee for the Protection of Human Subjects (CPHS).

Data availability
The datasets generated for this study are available on request to the corresponding author.

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Author contributions
Z.S., C.W. and M.R.-G. designed the research and wrote the manuscript. M.R.-G., Z.S., and M.P. conducted experiments and acquired data and analyzed data. All authors approved the manuscript.

Competing interests
The authors declare no competing interests.

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