Relation between phase of the menstrual cycle and asthma presentations in the emergency department

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**Background:** The phase of the menstrual cycle is thought to influence the course of asthma in women. One recent study reported a large increase in exacerbations during the perimenstrual phase, while another found a preovulatory increase. A study was undertaken to determine the relation between phase of the menstrual cycle and acute asthma exacerbation in patients presenting to the emergency department (ED).

**Methods:** All women aged 18–54 years presenting with a diagnosis of acute asthma exacerbation were considered for enrolment in the study. Women who were pregnant, postmenopausal, following hysterectomy, or with a >28 day menstrual cycle or incomplete reproductive history were excluded. The 792 eligible women were classified by menstrual phase based on both date of symptom onset and date of ED visit.

**Results:** When classified by date of symptom onset, 28% were preovulatory (days 5 to 11), 25% were periovulatory (days 12 to 18), 21% were postovulatory (days 19 to 25), and 27% were perimenstrual (days 26 to 4; p = 0.03). When classified by date of ED visit, 28% were preovulatory, 22% were periovulatory, 22% were postovulatory, and 27% were perimenstrual (p = 0.004). Using either approach, there were no significant differences in demographic factors or in asthma severity of women in the various menstrual phase groups.

**Conclusion:** Acute asthma exacerbations do not markedly increase during the perimenstrual phase. The results support the suggestion that both preovulatory and perimenstrual phases are actual triggers of asthma exacerbation in some women, or that these two phases serve as “co-factors” that worsen other recognised triggers of acute asthma.

**Methods**

The study was conducted at the Brooklyn Hospital Center emergency department between January 1997 and December 1999. Patients were enrolled and interviewed 24 hours per day, 7 days per week throughout the study period by trained research assistants. All acute asthma patients were screened for inclusion. Patients were asked the date of their last menstrual period during the triage process in the emergency department, whether they were pregnant, their medications for inclusion. Patients were asked the date of their last menstrual period during the triage process in the emergency department, whether they were pregnant, their medications, or if they were undergoing a surgical or surgical or surgical. No other questions were asked about their menstrual cycle. Inclusion criteria were female sex, age 18–54, physician diagnosis of acute asthma, and last menstrual period within 28 days of the ED visit. Since the cycle was idealised to 28 days, patients were sorted into their respective menstrual phase based on the range from their last menstrual cycle. Patients were excluded if they were pregnant, on oral contraceptives, had a surgical or surgical. No other questions were asked about their menstrual cycle. Inclusion criteria were female sex, age 18–54, physician diagnosis of acute asthma, and last menstrual period within 28 days of the ED visit. Since the cycle was idealised to 28 days, patients were sorted into their respective menstrual phase based on the range from their last menstrual period. Patients were excluded if they were pregnant, on oral contraceptives, had a surgical or surgical. No other questions were asked about their menstrual cycle. Inclusion criteria were female sex, age 18–54, physician diagnosis of acute asthma, and last menstrual period within 28 days of the ED visit. Since the cycle was idealised to 28 days, patients were sorted into their respective menstrual phase based on the range from their last menstrual period.
physiological menopause, or the visit was not their first ED visit for an asthma exacerbation during the study period.

Data collected about each study subject included demographic factors, past medical history (including asthma triggers and menstrual history), current medications, compliance, and details of the current asthma episode. Measures of asthma severity included history of intubation, oral steroids, or ED visit within the past year, initial peak expiratory flow (PEF), final PEF, and proportion admitted to the hospital.

As in two earlier studies, the menstrual cycle was assumed to have a duration of 28 days and women were divided into the following four categories with day 1 considered to be the onset of menstruation: preovulatory (days 5 to 11), periovulatory (days 12 to 18), postovulatory (days 19 to 25), and perimenstrual (days 26 to 4). Unique to this study, patients were classified into a phase based on two points in time: both the day of symptom onset and the day of the ED visit. Patients were excluded from that specific analysis if the date of symptom onset was missing (9%), or the date of symptom onset produced a cycle that was greater than 28 days or a cycle that was unable to be classified (91%). This resulted in a difference in sample size depending on method of classification (n = 560 for date of symptom onset, n = 792 for date of ED visit).

**Statistical analysis**
All analyses were performed using SPSS (Chicago, Illinois). Density of symptom onset (or ED visits) was plotted as a function of the day of menstrual cycle using Splus 4.0 (MathSoft Inc, Seattle, WA). The $\chi^2$ goodness of fit test was used to assess the association between phase of the menstrual cycle and asthma exacerbation. We chose a priori that women would be distributed equally across the four phases of the menstrual cycle and the analysis was done to compare the actual proportions in the four phases to 0.25. All tests are two tailed with p values <0.05 considered statistically significant.

**RESULTS**
There were 4987 patient visits to the ED for an asthma exacerbation during the study period of which 3208 (64%) were women. A total of 2161 women were excluded because their menstrual period was longer than 28 days (n = 522), surgical or physiological menopause (n = 420), multiple visits (>2 ED visits, n = 729), oral contraceptives (n = 275), and no menstrual date listed (n = 215). Of the remaining 1047 women, another 255 were excluded because the present visit was the second visit to the ED for asthma during that menstrual cycle leaving 792 women with acute asthma. When the data were analysed including all ED visits (that is, more than one ED visit by a single patient), the results did not materially differ from results obtained with the dataset based on only the first ED visit. Moreover, there were no significant differences in mean age, race, median family income (based on home ZIP code), or percentage actively smoking between the study group and those excluded for the reasons cited above.

Figure 1 shows the relative proportion of ED visits according to phase of menstrual cycle when classified by either symptom onset or day of ED visit. The two methods of classification yielded similar results. Asthma exacerbations began more often during the preovulatory (28%) and perimenstrual phases (27%) of the cycle than in the periovulatory or postovulatory phases (p = 0.03). Likewise, ED visits occurred more commonly among women in the preovulatory (28%) and perimenstrual (27%) phases (p = 0.004). Figure 2 shows the density of ED asthma visits plotted as a function of the day of menstrual cycle.

Using either approach, there were no significant differences in demographic factors or in asthma severity of women according to menstrual phases. To facilitate comparison with prior studies, table 1 shows data according to date of ED visit for acute asthma in relation to blood hormone levels in idealised menstrual cycle (B).

**DISCUSSION**
Our results show that ED visits for acute asthma are slightly more common during the preovulatory and perimenstrual phases of the menstrual cycle. This pattern was present whether the menstrual phase was estimated from the day of symptom onset or the day of the ED visit. These findings differ from Skobeloff et al who found that almost 50% of 182 ED visits for acute asthma occurred during the perimenstrual phase. Our results are more consistent with the recent study by Zimmerman et al on behalf of the Multicenter Airway Research Collaboration (MARC) investigators (www.emnetusa.org). They found that only 21% of 288 ED visits for acute asthma exacerbation occurred in the perimenstrual phase, whereas 33% occurred in the preovulatory phase. In the present study involving 792 ED visits for asthma, we noted...
more modest increases in both the preovulatory (28%) and perimenstrual (27%) phases.

It is unclear whether the observed associations represent a hormonal effect on a certain subset of the female asthmatic population or the effect of confounding factors such as differential smoking, infection, or exposure to allergens. In the MARC study only 13% (95% CI 9 to 18) of women reported reproductive factors as personal asthma triggers.28

Exactly how the normal menstrual cycle (fig 1) interacts with female asthma is unknown, but it seems clear that the initial ED study27 overstated the importance of perimenstrual asthma and the benefits of oestrogen. In some premenopausal women administration of oestradiol has decreased symptoms, cyclic variability in PEF, and airway reactivity.25 However, in postmenopausal women, oestrogen replacement therapy can cause worsening of some asthma symptoms.26 In addition, in non-asthmatic women, oestrogen replacement therapy is associated with an increased chance of developing asthma.27 Nevertheless, comparisons of endogenous hormonal fluctuations with exogenous hormonal administration require cautious interpretation; uncritical mixing of these concepts may contribute to apparent contradictions in the literature.

Objective evaluations of airway function during the menstrual cycle in women with stable asthma have yielded inconsistent results. Some studies have noted worsening symptoms and decreases in PEF in the premenstrual and menstrual period,17 18 20 while others found no changes in symptoms or spirometric parameters.29 Other studies found a discordance between symptoms and spirometric measurements during the menstrual cycle.20 21 In asthmatic women responsiveness to methacholine25 and adenosine monophosphate11 increases in the luteal phase (days 24 to 28) compared with the follicular phase (days 12 to 16) of the menstrual cycle.

Few studies have distinguished between women who report perimenstrual worsening of asthma and those who do not. In a prospective study Shames et al24 found increased use of bronchodilators and decreased morning PEF during the perimenstrual phase in women with reported menstrual asthma compared with women without menstrual asthma. However, daily symptom scores, evening PEF, differences in forced expiratory volume in 1 second between the luteal and follicular phases, and results of methacholine challenge were not significantly different for the two groups. Agarwal and Shah found significant cyclical differences in PEF in asthmatic women who perceived a menstrual linked deterioration which was not noted in women without such deterioration.29 In a small study Chandler et al29 noted that, although only a small percentage of patients noted a prior premenstrual exacerbation, all had a fall in PEF or exacerbation in symptoms which improved with oestradiol.

This amelioration was not mediated, however, by $\beta_2$ receptors since there was no change in $\beta_2$ receptor density during the menstrual cycle in those with premenstrual asthma. Even if one considers that airway function worsens in the perimenstrual period, our data suggest that it does not result in more asthma exacerbations requiring acute medical care.

This study has some potential limitations. Although the sample size is relatively large, all the subjects were from one site in an urban location which may limit the generalisability of the findings. Two hundred and fifteen women were excluded because their menstrual data were missing. Only women with physiological reproductive cycle were included, so women with hysterectomies but intact ovaries were excluded as well as those on birth control pills. In addition, inaccurate reporting of the subjects’ last menstrual period could have resulted in misclassification of the phase of the menstrual cycle. Only laboratory analyses could provide more accurate classification and allow the inclusion of women with menstrual cycles longer than 28 days. Finally, we note the possibility that perimenstrual and preovulatory presentations with acute asthma may have been increased by the patients’ concomitant ingestion of aspirin or other analgesics that worsen asthma. We were unable to test this hypothesis, which merits further study.

In conclusion, our data do not support claims that asthma exacerbations markedly increase during the perimenstrual phase. It is possible that both preovulatory and perimenstrual phases may have some adverse impact. For specific subsets of women, these phases could be actual triggers of an asthma exacerbation or they might serve as ‘co-factors’ that worsen other recognised triggers of acute asthma. These menstrual phase findings, together with a growing body of evidence for sex differences in asthma,14 support efforts to explore hormonal influences on asthma in women.

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REFERENCES
A clinic service offering isoniazid preventive therapy to HIV infected adults reduces the incidence of tuberculosis

This study followed 1655 HIV infected male employees of a South African gold mining company (median age 37 years) for a median duration of 22.1 months. Isoniazid preventive therapy, 300 mg daily, was offered at clinic enrolment to 679 subjects who attended with no evidence of active tuberculosis (TB). A total of 254 TB episodes met the study criteria. Before enrolment at the clinic the incidence of TB was 11.9 per 100 person-years, which reduced to 9.0 per 100 person-years following intervention. The adjusted incidence rate ratio for the effect of clinic enrolment on the incidence of TB was 0.68 (95% CI 0.48 to 0.96; p = 0.03). The overall reduction in the incidence of TB was 38%, and 42% in those with no prior history of TB. Failure to attain results as high as in previous studies may have been due to the fact that all the subjects who attended the clinic were assessed, irrespective of whether they received isoniazid.

Despite the reduction, the incidence rate is still high at 9 per 100 person-years. As the authors point out, long term prevention of TB needs to be considered as well as the emergence of isoniazid resistant strains. The follow up period in the study was short, and evidence suggests that the protective strategies of TB preventive therapy, particularly those without rifampicin, wane over time. Further studies to assess secondary prevention of TB—although not currently recommended—would be interesting, particularly in view of the prolonged survival now offered by antiretroviral therapy.

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