Modulation of Metoprolol Pharmacokinetics and Hemodynamics by Diphenhydramine Coadministration during Exercise Testing in Healthy Premenopausal Women

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ABSTRACT

Premenopausal women may be most vulnerable to acute coronary syndromes at a point in their menstrual cycle when their plasma estrogen levels are the lowest during and immediately after menstruation. Metoprolol is a first-line drug in the management of patients with acute coronary syndrome; however, when metoprolol was marketed in 1982, women were largely excluded from clinical trials. Furthermore, the over-the-counter antihistamine diphenhydramine inhibited the metabolism of the CYP2D6 substrate metoprolol in healthy, young men with pharmacokinetic and pharmacodynamic consequences. The pharmacokinetics and pharmacodynamics of metoprolol and its interaction with diphenhydramine were investigated in a randomized, double-blind, crossover, placebo-controlled manner. Metoprolol is extensively metabolized in humans into three major metabolites: O-hydroxymetoprolol (around 10% of the administered dose), O-desmethylmetoprolol, and deaminated metoprolol (Borg et al., 1975; Lennard, 1985). O-Desmethylmetoprolol is further metabolized to form a carboxylic acid metabolite (metoprolol acid) with the latter accounting for approximately 65% of the administered dose. All these metabolites together account for around 85% of the administered dose (Godbillon and Duval, 1984). alpha-Hydroxymetoprolol and O-desmethylmetoprolol were found to have significant beta-blocking activity when tested in cats. However, their ED50 values were around 9 to 10 times (heart rate), 5 to 8 times (contractile force), and 2 to 7 times (vasodilatation) higher than those of metoprolol (Borg et al., 1975). The alpha-hydroxylation pathway is controlled predominantly by the cytochrome P450 isof orm CYP2D6. This cytochrome P450 isof orm is subject to a genetic polymorphism with around 6 to 10% of the white population, the so-called PMs, lacking this enzyme due to the inheritance of two mutant CYP2D6 null alleles. The other 90% of white persons have been classified as EMs, although more recently gene multiplications were

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ABBREVIATIONS: PM, poor metabolizer; EM, extensive metabolizer; LVOT, left ventricular outflow tract; HR, heart rate; BP, blood pressure; VTI, velocity time integral; SV, stroke volume; SVI, stroke volume index; CO, cardiac output; CI, cardiac index; RPP, rate-pressure product; BSA, body surface area; PK/PD, pharmacokinetics/pharmacodynamics; AUC0–∞, area under the concentration-time curve; HPLC, high-performance liquid chromatography; AUEC, area under the effect curve.
found to result in an ultrarapid metabolizer status (3–5% of white persons) (Johansson et al., 1993), and around 10 to 15% of white persons are now referred to as intermediate metabolizers due to the presence of certain rare CYP2D6 alleles that result in reduced CYP2D6 activity (Raimundo et al., 2004).

Metoprolol is administered as a racemic mixture of R(+)- and S(-)-metoprolol. S-Metoprolol possesses 500 times greater β₁-receptor affinity than R-metoprolol (Wahlund et al., 1990). Although S-metoprolol has 33-fold greater β₁-receptor blocking activity on rabbit heart compared with the R-enantiomer, the latter possesses 10-fold greater β₂-receptor blocking activity in the rabbit ciliary process (Nathanson, 1988).

Although a multitude of mechanistic and observational studies suggest a protective effect of estrogen substitution in postmenopausal women against heart disease, its use was recently discouraged when the popular equine estrogens were associated with an increase of venous thromboembolic events in a placebo-controlled study (Hully et al., 1998). Nevertheless, epidemiological data have clearly shown that women in their reproductive years have a low incidence of heart disease and that the cardiovascular risk increases after menopause when endogenous hormone levels are naturally low (Lerner and Kannel, 1986). Our group has recently reported that premenopausal women with at least one known risk factor of coronary artery disease were most likely to suffer acute myocardial infarctions or unstable angina attacks during or immediately after menstruation, suggesting that relatively low levels of circulating estrogen may contribute or act as a trigger for acute coronary events in this young, female population (Hamelin et al., 2003). Although the treatment of women with heart disease is usually based on extrapolations of data obtained in men, sex-specific differences in the activities of metabolic enzymes and the pharmacokinetics/pharmacodynamics (PK/PD) of many drugs exist (Labbé et al., 2000; Meibohm et al., 2002). Our group has recently reported that the activity of CYP2D6 as determined by the metabolic ratio of dextromethorphan/dextrorphan was significantly greater in 56 postmenopausal female EMs compared with 86 male EMs. The β₁-receptor antagonist metoprolol, a prominent CYP2D6 substrate, is a first-line treatment choice in the management of patients presenting with acute coronary syndromes. Because women were essentially excluded from clinical trials when metoprolol was brought to market in 1982, pharmacokinetic/pharmacodynamic data on this drug in women are scarce. We were interested in studying the pharmacokinetics and hemodynamics of metoprolol in healthy premenopausal women as a representative group for all women with fluctuating endogenous hormone levels during the reproductive years. The study was performed as close to the menstruation as possible to standardize for the potential effects of fluctuations in endogenous hormones on metoprolol’s disposition while approaching the period of increased risk for acute events in this population.

Our group has further demonstrated that the classic over-the-counter prototype antihistamine diphenhydramine [2-(diphenylmethoxy)-N,N-dimethylethylamine] inhibits the oral, nonrenal, and partial metabolic clearances of racemic metoprolol to α-hydroxymetoprolol, thus increasing metoprolol area under the concentration-time curve (AUC)₀⁻∞ and prolonging the negative chronotropic and inotropic effects of the drug in EM but not PM men (Hamelin et al., 2000).

However, the exact nature of the interaction (solely pharmacokinetic versus pharmacokinetic/pharmacodynamic) as well as the potential differential effects of diphenhydramine on the disposition of metoprolol’s enantiomers are still unknown. Thus, the goals of the present study were to determine 1) the racemic and stereoselective disposition of metoprolol; 2) the resultant hemodynamic effects of metoprolol; and 3) the interaction between the CYP2D6 inhibitor diphenhydramine and metoprolol in healthy, premenopausal women immediately after menstruation and before ovulation.

Materials and Methods
This study was approved by the Laval Hospital Ethics Committee, and all volunteers provided written informed consent before participating in this study.

Volunteers. Twenty young, healthy, nonsmoking white women (age range 18–40 years, mean age 26.8 ± 8 years, weight range 49–100 kg, and mean weight 60 ± 12 kg) not consuming oral contraceptives and having regular menstrual cycles were recruited from the Quebec city area. The participants were recruited according to their CYP2D6 activity and were either EMs (n = 16) or PMs (n = 4) as determined by phenotyping and genotyping (described below). The general health status of the participants was determined based on a general questionnaire and a physical examination, including electrocardiogram and routine laboratory tests to determine renal and hepatic function. Volunteers followed their menstruation and ovulation schedule for 1 to 3 months before participation in the study. A Conceive LH (Quidel Corp., San Diego, CA) kit that accurately predicts ovulation was provided during that period, and an ovulation test was also done on the study day mornings to ensure that the subjects were not ovulating that day. During screening, all volunteers also underwent two-dimensional echocardiography using SONOS 5500 echocardiograph (Philips Medical Systems, Bothell, WA) to rule out stenosis and to determine the left ventricular outflow tract (LVOT) area.

Phenotyping and Genotyping for CYP2D6 Activity. The CYP2D6 phenotype was determined by using dextromethorphan (3-methoxy-17-methylmorphinan monohydrate) as the probe drug as described previously (Labbé et al., 2000). Individuals with a dextromethorphan/dextrorphan metabolic ratio of >0.3 were considered poor metabolizers. A 10-ml blood sample was obtained for genotyping. DNA was extracted from peripheral blood lymphocytes, and CYP2D6*1A, CYP2D6*3, CYP2D6*4, CYP2D6*5, CYP2D6*6, CYP2D6*7, and CYP2D6*8 alleles were determined using a classic multiplex-polymerase chain reaction (Stuven et al., 1996).

Study Design. This study was conducted in a randomized, double-blind, crossover, placebo-controlled manner and was carried out after menstruation but before ovulation over a 2-month period (Fig. 1). During each study arm, diphenhydramine (50 mg) or placebo (lactose) was administered to the women three times daily for 5 days. A single oral dose of 100 mg of metoprolol tartrate was administered on the morning of the third study day. The T₀½ value of diphenhydramine ranges between 4 and 9.2 h (Spector et al., 1980; Bliden et al., 1986). Hence, at the time of metoprolol administration, diphenhydramine plasma concentrations were at steady state. Diphenhydramine or placebo administration was continued beyond metoprolol administration, until the fifth study day, i.e., the end of the study arm. Randomization tables were prepared by the biostatistician of the research center, whereas a hospital pharmacist dedicated to research protocols controlled the randomization schedule and dispensed crushed placebo tablets and diphenhydramine capsules hidden inside identical-looking colored hard gelatin capsules.

Hemodynamic Assessment. The individual workload necessary to obtain an exercise heart rate of 140 beats/min on a stationary
upright bicycle was determined for each individual before metoprolol administration (time 0). This workload was applied at final 4 min of each 8-min exercise test at 0.75, 1.5, 2.25, 3, 4, 8, and 12 h after metoprolol administration. Heart rate (HR) and blood pressure (BP) were obtained using an automated blood pressure monitor (Quinton, Bothel, Washington), and continuous-wave Doppler recordings of flow velocity were obtained from the suprasternal notch using a nonimaging transducer connected to a SONOS 5500 echocardiograph during rest and exercise at various time points. The nonimaging transducer was angulated to record the signal with maximal flow velocity in the ascending aorta. The Doppler velocity signals were analyzed to obtain the following parameters: aortic velocity time integral (VTI), acceleration time, and ejection time. The values of stroke volume (SV), stroke volume index (SVI), cardiac output (CO), and cardiac index (CI) and rate-pressure product (RPP) were calculated as indicated below: 1) SV = VTI × LVOT area, 2) SVI = SV/body surface area (BSA), 3) CO = SV × HR, 4) CI = CO/BSA, and 5) RPP = HR × systolic BP.

Pharmacokinetic Analysis. Serial blood samples were obtained from the subjects at 0, 0.25, 0.5, 0.75, 1, 1.25, 1.5, 1.75, 2, 2.25, 2.5, 3, 3.5, 4, 6, 8, 10, 12, 24, 28, 32, 36, and 48 h after metoprolol administration. Samples were obtained through an indwelling catheter up to 12 h and by venous puncture thereafter. Blood samples were immediately spun down, and plasma was harvested and frozen at −20°C until analysis. Urine collection was done from 0 to 12. 12 to 24, and 24 to 48 h after metoprolol dosing. Urine volume and pH were determined, and aliquots were frozen at −20°C until analysis. Subjects reported for the study after overnight fasting and were provided a light snack at 2 h and lunch at 4 h postmetoprolol. Subjects were instructed to abstain from alcohol, grapefruit juice, cruciferous vegetables, char broiled food, any form of medication (including any over-the-counter medications, drugs belonging to an alternate system of medicine, herbal supplements, vitamins, and minerals) from at least 2 days before starting diphenhydramine/placebo administration until the end of the study arm.

HPLC Analysis of Metoprolol and α-Hydroxymetoprolol in Biological Samples. Racemic metoprolol [1-isopropylamino]-3-(4-(2-methoxyethyl)phenoxy)-2-propanol and its α-hydroxy metabolite [1-(3-hydroxy-2-methoxyethyl)phenoxy]-3-isopropylamino)-2-propanol were determined in plasma and urine using a previously reported HPLC method from our laboratory (Hamelin et al., 2000). The method was found to be highly reproducible with inter- and intraday coefficients of variation below 5%.

R- and S-metoprolol were determined with modifications of a previously published method (Lanchote et al., 2000). The analysis was carried out using a Shimadzu HPLC system with fluorescence detection (λex = 229 and λem = 298). The resolution was achieved on a Chiralcel OD-H analytical column (250 × 4.6 mm; Daicel Chemical Industries Limited, Exton, PA) using a C18 guard column (Waters, Milford, MA). The mobile phase consisted of n-hexane/isopropanol/diethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time. After addition of 400 ng of alprenolol [internal standard, 1-((methyldiethylamine/trifluoroacetic acid (92:8:0.15:0.025), which was recirculated in a closed loop system to analyze up to 45 samples at a time.
For compartmental pharmacokinetic analysis, nonlinear regression with iteratively reweighted least square estimation was used to fit one- \((n = 18)\) or two- \((n = 2)\)-compartment models to S-metoprolol concentration-time data according to the following criteria: value of objective function, Akaike and Schwartz criteria, correlation matrix, distribution of residuals, and visual fit. The fitting procedure was repeated by changing the start values for pharmacokinetic parameters to ensure that the nonlinear regression algorithm converged at the global rather than a local minimum. The results from compartmental analysis were used as input variables for S-metoprolol PK/PD analysis (see below).

**Metoprolol Pharmacodynamics.** The total area under the response-time curve (AUEC) in EMs and PMs on metoprolol with or without diphenhydramine coadministration was calculated for three pharmacodynamic response markers during exercise, i.e., heart rate, blood pressure and rate-pressure product, by a calculation of the cumulative reduction from the baseline values over the 12-h study duration. Baseline values were the maximum values of the three response markers. The AUEC was estimated by the linear trapezoidal rule (Microsoft Excel; Microsoft, Mississauga, ON, Canada). The maximum effect compared with baseline, i.e., \(E_{\text{max}}\), was read directly from the response-time data.

**S-Metoprolol PK/PD Modeling.** Integrated PK/PD modeling was used to relate S-metoprolol plasma concentrations to three pharmacodynamic response markers, i.e., heart rate, blood pressure, and rate-pressure product. Pharmacodynamic parameters were modeled as changes relative to baseline values (highest values for any individual in this case). The results of compartmental pharmacokinetic modeling (absorption rate constant, elimination rate constant, apparent volume of distribution, and lag time) of S-metoprolol along with S-metoprolol plasma concentrations were modeled relative to effect-time profiles for each individual. Based on objective function, residual distribution, visual goodness-of-fit, and physiological reality of the parameter estimates, a direct-link, direct-response pharmacodynamic model using a sigmoid \(E_{\text{max}}\) relationship, with the effect compartment in the central compartment, was used (Holford and Sheiner, 1991; Meibohm and Derendorf, 1997): \(E = (E_{\text{max}} \times C^{n})/(EC_{50}^{n} + C^{n})\), where \(n\) is the shape factor, \(E_{\text{max}}\) is the maximum hemodynamic effect, \(EC_{50}\) is the plasma concentration needed to achieve half of \(E_{\text{max}}\) and \(E\) and \(C\) refer to the observed response parameter and plasma concentration values. The values of \(E_{\text{max}}\) and \(EC_{50}\) were determined by the modeling procedure. All calculations were performed with the Scientist software (MicroMath Inc., Salt Lake City, UT).

**Statistical Analysis.** The analysis of Kolmogorov-Smirnov and Levine were used to test the normality and variance homogeneity of the data. Because all data were normally distributed and variances were equal, data were expressed as mean ± standard deviation in all statistical analysis. Pharmacokinetic parameters of racemic and enantiomeric metoprolol in the whole population \((n = 20)\) and in EMs \((n = 16)\) and PMs \((n = 4)\) were compared using a cross-nested design with two experimental factors (metabolizer and medication, i.e., administration of metoprolol with or without diphenhydramine) and medication randomly assigned to subjects as the nested factor. A mixed model analysis was also performed with an interaction term between the fixed factors (metabolizer and medication). Pharmacokinetic parameters of metoprolol enantiomers were analyzed using the same statistical approach with a fourth factor added to the model to compare the \(R\)-enantiomer with the \(S\)-enantiomer.

For comparison of pharmacodynamic parameters in the whole population and in EMs and PMs, a cross-nested design was used to analyze changes of mean cardiac index, blood pressure, and rate-pressure product data. This design was performed with two fixed factors (metabolizer and medication), one random factor (subject within groups), and a repeated factor (time) nested into the random factor subject. Different statistical models were tested, and the final analysis was done with heterogeneity between metabolizers (covariance structures not similar). The multivariate normality was verified using Mardia tests (Mardia, 1974). The results were considered significant with \(P \leq 0.05\). All analyses were conducted using the statistical package SAS (SAS Institute, Cary, NC).

**Results**

**Clinical Study**

Sixteen \((n = 16)\) EMs and four PMs were recruited for the study. All subjects completed the study and reported no significant adverse effects other than somnolence in some subjects (six EMs and all four PMs) during the diphenhydramine coadministration arm of the study. Results of phenotyping using dextromethorphan were in line with those of genotyping. Seven of 16 EMs were found to be homozygous for the wild-type allele, whereas nine subjects were heterozygous with one wild-type and one mutant allele (CYP2D6*3 in one subject, CYP2D6*4 in seven subjects, and CYP2D6*5 in one subject). Two of the four PMs were homozygous for CYP2D6*4, and two were heterozygous (CYP2D6*4/ CYP2D6*5).

**Pharmacokinetics**

**Influence of CYP2D6 Phenotype on Racemic Metoprolol Pharmacokinetics in Women.** The mean pharmacokinetic profile of racemic metoprolol in 16 EM and four PM women is presented in Fig. 2, and calculated noncompartmental pharmacokinetic parameters are summarized in Table 1. The plasma concentration-time profiles of metoprolol were significantly different in EMs compared with PMs \((P < 0.05)\). \(C_{\text{max}}\) and AUC\(_{0-\infty}\) values were 2- and 4-fold higher, respectively, in PMs on placebo compared with EMs on placebo. This was the result of an approximately 7-fold lower CL/F, an 8-fold lower CL\(_{NR}\), and a 300-fold lower partial metabolic CL\(_{\text{MET-OH-MET}}\) in PMs compared with EMs \((P < 0.05)\). The \(t_{1/2}\) value was 2.5-fold longer in PMs compared with EMs \((P < 0.0001)\). In contrast, \(T_{\text{max}}\) and CL\(_{R}\) values were similar among the phenotypes \((P > 0.05); \text{Table 1})\.

**Fig. 2.** Pharmacokinetic profile of racemic metoprolol in 16 extensive and four poor metabolizer women after the administration of a single oral dose of 100 mg of metoprolol tartrate with or without concomitant administration of diphenhydramine or placebo to steady state. Results presented as mean ± S.D. MET, metoprolol; PCB, placebo; DPH, diphenhydramine.
Coadministration of diphenhydramine resulted in a 30% increase in CL/F, CLNR, and CLR of the two enantiomers in the presence or absence of diphenhydramine during exercise. After the placebo and diphenhydramine phase. The changes were the result of a 2.6- and 2.2-fold decrease in CL/F and CLNR for the R- and S-metoprolol enantiomers, respectively (P < 0.05). Inhibition of metoprolol hepatic elimination brought R- and S-metoprolol pharmacokinetic parameters closer to those of PMs, but inhibition was not complete (Table 3; Fig. 3). Diphenhydramine did not significantly affect exposure to either metoprolol enantiomer in PMs (Table 3; P > 0.05).

### R-Metoprolol/S-Metoprolol Ratios
For EMs in the placebo phase, CL/F and CLNR of R-metoprolol were 1.7 times faster than that of S-metoprolol (P < 0.05), but there was no significant difference in the renal clearance of either enantiomer (CLR R/S-metoprolol 1.05; P > 0.05). As a consequence, R/S-metoprolol AUC0–inf was 0.67 (P < 0.05). Interestingly, diphenhydramine coadministration reduced the differences in disposition between the two enantiomers. Diphenhydramine decreased CL/F and CLNR approximately 2.6-fold for the R- compared with approximately 2.2-fold for the S-enantiomer, resulting in an R/S-metoprolol ratio of the clearances of approximately 1.4 (P > 0.05 for difference in CL/F and CLNR between enantiomers). In contrast, after diphenhydramine, CLR values of the two enantiomers were similar to each other (CLR R/S-metoprolol 1.02; P > 0.05) and to those values observed during the placebo phase. After the changes in clearances caused by diphenhydramine administration, AUC0–inf R-metoprolol increased 2.1-fold compared with 1.8-fold for S-metoprolol, resulting in an R/S-metoprolol ratio of AUC0–inf of 0.76 (P > 0.05 for differences in AUC0–inf between enantiomers). In PMs, the AUC0–inf R/S-metoprolol ratio was found to be 0.98 without and 1.00 with diphenhydramine coadministration, showing no stereoselectivity. Correspondingly, there was no significant difference (P > 0.05) in the CL/F, CLNR, and CLR of the two enantiomers in the placebo and diphenhydramine phase. The R/S-metoprolol ratios for AUC0–inf CL/F, CLNR, and CLR were significantly different between EMs and PMs (P < 0.0001).

### Pharmacodynamics
Changes in four hemodynamic response markers, i.e., heart rate, rate-pressure product, stroke volume index, and cardiac index, in response to the administration of metoprolol in the presence or absence of diphenhydramine during exercise are shown in Fig. 4. Metoprolol administration also affected these parameters while obtained at rest. However,
changes at rest were small and hence are not reported herein.

**Exercise Heart Rate.** Metoprolol administration resulted in a significant change in mean exercise heart rate over 12 h for PMs and EMs ($P < 0.001$; Fig. 4A). PMs and EMs on placebo followed a significantly different heart rate profile over time ($P < 0.05$ from 1.5 to 12 h). The mean heart rate was reduced by 31 and 24% compared with baseline values at 1.5 h post metoprolol in PMs and EMs ($P < 0.05$). Compared with baseline values, the heart rate was still reduced by 26% (placebo) and 28% (diphenhydramine) in PMs but only by 8% in EMs (placebo) at 12 h post metoprolol (PMs, $P < 0.05$; EMs, $P > 0.05$). However, EMs on diphenhydramine had heart rate reduced by 17% compared with baseline values at 12-h postmetoprolol ($P < 0.05$). Diphenhydramine coadministration had no significant ($P > 0.05$) influence on the heart rate profile of PMs. Starting at 4 h, the heart rate profile of EMs on placebo evolved distinctly from the heart rate profile of EMs on diphenhydramine and was significantly different ($P < 0.05$) at 4, 6 ($P = 0.06$), 8, and 12 h.

**Exercise Rate-Pressure Product.** In all 20 volunteers, rate-pressure product was significantly affected ($P = 0.0001$; Fig. 4B) by metoprolol administration regardless of placebo or diphenhydramine cotreatment. The rate-pressure product profile of PMs was significantly different from that of EMs ($P = 0.0007$ from 1.5 to 12 h). Initially, mean rate-pressure product had maximal decreases of 46 and 35% compared with baseline values at 1.5 h post metoprolol in PMs and EMs on placebo, respectively (PMs versus EMs; $P = 0.0004$). Compared with baseline values, the rate-pressure product was still reduced by about 27% (placebo and diphenhydramine phase) in PMs but only by around 8% in EMs (placebo) at 12 h post metoprolol (PMs, $P = 0.0005$; EMs, $P = 0.02$ compared with baseline). However, the rate-pressure product of EMs on diphenhydramine was reduced by 14% compared with baseline values at 12 h post metoprolol ($P = 0.004$). Diphenhydramine coadministration significantly ($P < 0.05$) altered the rate pressure profile of EMs, shifting it toward the profile of PMs (Fig. 4B). However, diphenhydramine coadministration had no significant effect ($P > 0.05$) on the rate-pressure product profile of PMs.

**Exercise Stroke Volume Index.** Stroke volume index values changed significantly over time in all 20 volunteers ($P = 0.0001$; Fig. 4C). There was no significant difference between the profile of PMs and EMs ($P > 0.05$). However, diphenhydramine coadministration significantly affected the

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**TABLE 3**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>EM (n = 16)</th>
<th>PM (n = 4)</th>
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<tbody>
<tr>
<td></td>
<td>R-MET-PCB</td>
<td>S-MET-PCB</td>
</tr>
<tr>
<td></td>
<td>R-MET-PCB</td>
<td>S-MET-PCB</td>
</tr>
<tr>
<td>C$_{\text{max}}$ (μmol/l)</td>
<td>0.38 ± 0.29$^a$</td>
<td>0.51 ± 0.23$^b$</td>
</tr>
<tr>
<td>t$_{\text{max}}$ (h)</td>
<td>1.69 ± 0.63</td>
<td>1.72 ± 0.68</td>
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<tr>
<td>AUC$_{\text{0–48h}}$ (μmol · h/l)</td>
<td>1.63 ± 0.92$^a$</td>
<td>2.44 ± 1.25$^b$</td>
</tr>
<tr>
<td>CL$_{\text{R}}$ (l/h)</td>
<td>147 ± 124$^a$</td>
<td>88.7 ± 68.4$^b$</td>
</tr>
<tr>
<td>CL$_{\text{Qc}}$ (l/h)</td>
<td>4.10 ± 2.18</td>
<td>3.86 ± 2.01</td>
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$^a P < 0.05$ EM R-MET/PCB versus PM R-MET/PCB.  
$^b P < 0.05$ EM S-MET/PCB versus PM S-MET/PCB.  
$^c P < 0.05$ EM R-MET/PCB versus EM R-MET/DPH.  
$^d P < 0.05$ EM S-MET/PCB versus EM S-MET/DPH.
The profile of EMs on diphenhydramine dropped lower than the profile of EMs on placebo and followed closely the profile of PMs (regardless of the treatment) at most times. Concomitant diphenhydramine did not significantly alter the stroke volume index profile of PMs.

**Exercise Cardiac Index.** Cardiac index changed significantly in all 20 subjects over the 12-h study period ($P = 0.0001$; Fig. 4D). In the placebo phase, PMs followed a significantly different cardiac index profile over time compared with the EMs ($P = 0.009$). Initially, mean cardiac index had a maximal decrease of around 27% and 17% compared with baseline in PMs and EMs on placebo, respectively (PMs versus EMs; $P < 0.05$). Coadministration of diphenhydramine did not significantly affect the maximum effects on cardiac index in PMs and EMs (27 and 19% decrease for PMs and EMs, respectively; $P > 0.05$ for the treatment effect). However, diphenhydramine coadministration significantly influenced the cardiac index profile of EMs ($P < 0.05$ at 3, 4, 8, and 12 h compared with EMs on placebo) with the profile becoming similar to that of PMs receiving either cotreatment (PMs versus EMs on diphenhydramine; $P > 0.05$ from 3 to 12 h). Diphenhydramine coadministration had no significant effect on this response marker in PMs ($P > 0.05$).

**S-Metoprolol PK/PD Modeling**

The results of the pharmacokinetic/pharmacodynamic modeling of S-metoprolol data with various pharmacodynamic response markers are summarized in Table 4. PMs had a significantly greater AUEC_{(heart rate)} and AUEC_{(rate-pressure product)} compared with EMs on placebo (irrespective of the cotreatment; $P \leq 0.006$). Concomitant administration of diphenhydramine resulted in a significant increase in AUECs for heart rate and rate-pressure product values in EMs ($P = 0.01$ compared with placebo coadministration) but not in PMs. In contrast, AUEC for systolic blood pressure was neither influenced by the phenotype nor the cotreatment ($P > 0.05$). $E_{\text{max}}$ for heart rate, blood pressure, and rate-pressure product values were not significantly different between EMs and PMs on either cotreatment. $E_{50}$ for heart rate ($P = 0.009$), $E_{50}$ for systolic blood pressure ($P = 0.03$), and $E_{50}$ for rate-pressure product ($P = 0.01$) were significantly higher in PMs compared with EMs. Although diphenhydramine coadministration had no significant effect on $E_{\text{max}}$ and $E_{50}$ values, it significantly increased the AUECs for heart rate and rate-pressure product in EMs ($P < 0.05$ at 3, 4, 8, and 12 h).
TABLE 4
Results of pharmacokinetic/pharmacodynamic modeling of S-metoprolol pharmacokinetic data with various pharmacodynamic response markers in 20 healthy, young women (16 extensive and four poor metabolizers) in the absence or presence of diphenhydramine dosed to steady state. Results presented as mean ± S.D.

<table>
<thead>
<tr>
<th>Experiment</th>
<th>EM-PCB</th>
<th>EM-DPH</th>
<th>PM-PCB</th>
<th>PM-DPH</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exercise heart rate</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Observed $E_{\text{max}}$ ± S.D. (beats/min)</td>
<td>37 ± 5</td>
<td>42 ± 8</td>
<td>46 ± 11</td>
<td>45 ± 8</td>
</tr>
<tr>
<td>$E_{\text{max}}$ ± S.D. (μmol/l)</td>
<td>0.19 ± 0.09</td>
<td>0.14 ± 0.08</td>
<td>0.25 ± 0.08</td>
<td>0.17 ± 0.08</td>
</tr>
<tr>
<td>$AUEC$ ± S.D. (beats·h/min)</td>
<td>275 ± 63</td>
<td>350 ± 88</td>
<td>423 ± 85</td>
<td>408 ± 78</td>
</tr>
<tr>
<td>Exercise systolic blood pressure</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Observed $E_{\text{max}}$ ± S.D. (mm Hg)</td>
<td>31 ± 10</td>
<td>36 ± 13</td>
<td>43 ± 6</td>
<td>37 ± 12</td>
</tr>
<tr>
<td>$E_{\text{max}}$ ± S.D. (μmol/l)</td>
<td>0.13 ± 0.09</td>
<td>0.21 ± 0.15</td>
<td>0.24 ± 0.07</td>
<td>0.31 ± 0.07</td>
</tr>
<tr>
<td>$AUEC$ ± S.D. (mm Hg·h)</td>
<td>215 ± 95</td>
<td>245 ± 73</td>
<td>278 ± 80</td>
<td>231 ± 141</td>
</tr>
<tr>
<td>Exercise rate-pressure product</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Observed $E_{\text{max}}$ ± S.D. (beats·mm Hg/min)</td>
<td>7688 ± 1971</td>
<td>8504 ± 2011</td>
<td>9626 ± 1758</td>
<td>9905 ± 2876</td>
</tr>
<tr>
<td>$E_{\text{max}}$ ± S.D. (μmol/l)</td>
<td>0.09 ± 0.06</td>
<td>0.15 ± 0.10</td>
<td>0.20 ± 0.08</td>
<td>0.22 ± 0.04</td>
</tr>
<tr>
<td>$AUEC$ ± S.D. (beats·mm Hg·h/min)</td>
<td>57,952 ± 16,193</td>
<td>71,838 ± 17,949</td>
<td>85,435 ± 15,382</td>
<td>84,359 ± 23,239</td>
</tr>
</tbody>
</table>

Discussion

This is the first study to report an extensive assessment of pharmacokinetics and hemodynamics of metoprolol in young, healthy premenopausal women (EMs and PMs) with and without the administration of a moderate inhibitor of CYP2D6, namely, the over-the-counter antihistamine diphenhydramine in a randomized, double-blind, and placebo-controlled manner. The study was conducted under controlled conditions at a point in the menstrual cycle when circulating estrogens in the body are in the low range, i.e., after menstruation and before ovulation. Conducting the study at this point in the menstrual cycle is very pertinent, considering the increased cardiovascular vulnerability of premenopausal women during and right after the menstruation (Hamelin et al., 2003). Furthermore, conducting the study at a precise time period during the menstrual cycle is important for minimizing any potential interindividual variability in drug metabolism caused by cyclic hormonal changes (Walle et al., 1996; Benton et al., 2000). The study is also highly relevant considering the prescription-free availability of many classic antihistamines that carry the risk of interacting with a coadministered CYP2D6 substrate such as metoprolol in this age group and in older women.

The pharmacokinetic profiles of racemetric metoprolol as well as of $R$- and $S$-metoprolol observed in PMs and EMs in our study are similar to those reported previously (Lennard et al., 1983). In our study, the average AUC$_{(0-\infty)}$, S/R-metoprolol ratio was 1.0 in PMs and 1.5 in EMs, which is similar to literature values (Lennard et al., 1983). Of interest, the total clearance of $R$-metoprolol was significantly greater than that of $S$-metoprolol in EMs on placebo, and coadministration of diphenhydramine decreased the clearance and eliminated the difference between the enantiomers. This suggests that diphenhydramine has a greater inhibitory effect on the metabolism of $R$-metoprolol. Since others have shown that O-demethylation was significantly stereoselective for $R$-metoprolol (Murthy et al., 1990; Kim et al., 1993; Mautz et al., 1995), one may speculate that diphenhydramine inhibits O-demethylation of metoprolol to a greater extent.

Our group has previously demonstrated that, in vitro, diphenhydramine can competitively inhibit metoprolol metabolism with a Ki value of 2 μM (Hamelin et al., 2000). This in vitro inhibition persisted in vivo in healthy, young EM men, resulting in significantly decreased metoprolol clearance and thus more pronounced and significantly prolonged negative chronotropic and inotropic effects of metoprolol. In the present study, we demonstrated that diphenhydramine administration to steady-state modulated the pharmacokinetics and hemodynamics of metoprolol in healthy young premenopausal EM women to a similar extent as previously found in men. Diphenhydramine coadministration shifted the heart rate profile of EMs toward that of PMs receiving either cotreatment, thus demonstrating a prolongation of the negative chronotropic effect of metoprolol in EMs. Similarly, the stroke volume index profile of EMs during diphenhydramine coadministration was lower than their profile during placebo coadministration and followed the profile of PMs, even though the heart rate profiles of EMs on concomitant diphenhydramine and that of PMs was lower than that of EMs on concomitant placebo. This indicates a greater negative inotropic effect in PMs (on either cotreatment) and in EMs on diphenhydramine compared with EMs on placebo.

In the present study, the heart rate and rate-pressure product AUEC values increased significantly from EMs on placebo to EMs on diphenhydramine to PMs on either cotreatment (Table 4). This increase corresponds to a significant increase in the plasma AUC$_{(0-\infty)}$ of the drug from EMs on placebo to EMs on diphenhydramine to PMs on either cotreatment (Tables 1 and 3). No significant increase in systolic blood pressure AUEC values, similar to the increases in heart rate and rate-pressure product, were observed. However, this is not entirely unexpected. Although a linear relationship between log of plasma levels and exercise heart rate variations has been described in the literature, no such association exists for antihypertensive activity of metoprolol (Bengtsson et al., 1975). It is also possible that the potential differences cannot be identified because blood pressure is a more complex regulated parameter than heart rate where numerous competing and compensatory mechanisms are interacting simultaneously. No significant differences in the
$E_{\text{max}}$, heart rate, blood pressure, and rate-pressure product were observed between EMs and PMs (regardless of the co-treatment), despite the fact that PMs (regardless of the co-treatment) and EMs on diphenhydramine had a higher $AUC(0-\infty)$ and $C_{\text{max}}$ of metoprolol and a higher AUEC. This could be explained by considering that these subjects, despite their phenotype and co-treatment, are reaching the observed $E_{\text{max}}$ for the tested response markers at a certain time point with the administered dose. Hence, any increase in the concentration of S-metoprolol in the plasma (whether due to concomitant diphenhydramine as for EMs or due to phenotype) has no significant additional effect on this parameter. The $EC_{50}$ heart rate, systolic blood pressure, and rate-pressure product values were significantly higher in PMs (placebo) than EMs (placebo), indicating that EMs had a higher sensitivity toward S-metoprolol than PMs. Diphenhydramine coadministration was found to increase the $EC_{50}$ heart rate and rate pressure parameters in EMs but not in PMs. One may speculate that diphenhydramine- or metoprolol-related metabolites or a combination of both contribute to these observations regarding the $EC_{50}$. Diphenhydramine or one of its metabolites might change the concentration-effect relationship by counteracting the effects of metoprolol on heart rate. On the other hand, the amounts of $\alpha$-hydroxymetoprolol and/or $O$-demethylmetoprolol and their contribution to the hemodynamic effects may play a role. Even though their in vitro activity is 2 to 10 times less than that of metoprolol, both $\alpha$-hydroxymetoprolol and metoprolol acid were reported to reach higher concentrations than S-metoprolol in plasma after a single dose (Mistry et al., 2002). In fact, the $AUC(0-\infty)$ of racemic $\alpha$-hydroxymetoprolol was 21% higher than that of S-metoprolol in our study. Hence, the concentration and activity would be high enough to actually contribute to the observed heart rate response to S-metoprolol. Since PMs had less $\alpha$-hydroxymetoprolol (AUC of 0.06 $\mu$mol · h/l for PMs on placebo compared with AUC of 3.04 $\mu$mol · h/l for EMs on placebo), they require higher S-metoprolol concentrations to reach the same effect, i.e., $EC_{50}$ was higher. Similarly, diphenhydramine coadministration decreased the formation of $\alpha$-hydroxymetoprolol by 15.5% and possibly that of other metoprolol metabolites, thereby requiring more S-metoprolol, i.e., higher $EC_{50}$ values for the same heart rate response.

Interestingly, although the EM women in this study had a 300-fold higher CL$_{\text{MET-s-OH-MET}}$ (racemic metoprolol), a 6-fold higher CL$_{\text{NR}}$ (S-metoprolol), and a 3.6-fold lower $AUC(0-\infty)$ (S-metoprolol) compared with their PM counterparts during the placebo phase, PMs had merely between 1.3- and 1.5-fold higher AUEC and between 1.5- and 2.5-fold higher $EC_{50}$. These differences in pharmacokinetics and pharmacodynamics could possibly be explained by significant synergistic cardiovascular effects of the metabolites in EMs. In contrast, a mere 3-fold decrease in CL$_{\text{MET-s-OH-MET}}$ (racemic metoprolol), a 2.3-fold decrease in CL$_{\text{NR}}$ (S-metoprolol), and a 1.8-fold increase in AUC$(0-\infty)$ (S-metoprolol) in EMs, caused by the coadministration of diphenhydramine instead of placebo, resulted in an around 1.3-fold increase in AUEC and about 1.5-fold increase in $EC_{50}$. This implies that the small changes in metoprolol disposition in EMs produced by diphenhydramine coadministration resulted in nearly as much pharmacodynamic effects as seen in PMs. This pharmacodynamic modulation could also be related to the cardiovascular effects of diphenhydramine (Zareba et al., 1997; Khalifa et al., 1999).

A potential weakness of this study is that the pharmacodynamic parameters were measured only for a period of 12 h postmetoprolol, which might have caused an underestimation of pharmacodynamic parameters in some volunteers. However, given the length of the protocol, it would have been beyond the physical capacity of the volunteers to extend the study longer.

The current study reports an extensive assessment of metoprolol pharmacokinetics (racemic and enantiomeric) and hemodynamics in young healthy premenopausal EM and PM women at the time of the menstrual cycle when they may be most predisposed to acute coronary syndromes (i.e., after menstruation and before ovulation). Significant differences in the PK/PD relationships for EM and PM women were observed and need to be taken into consideration in clinical practice. Furthermore, caution is warranted when the over-the-counter antihistamine diphenhydramine is part of a chronic therapeutic regimen, especially because relatively small, although significant effects on metoprolol’s disposition result in relatively large pharmacodynamic effects.

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References


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