Effect of nifedipine on serum theophylline concentrations and asthma control

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ABSTRACT The effect of adding slow release nifedipine to oral theophylline has been studied in eight patients with stable but symptomatic asthma, a double blind placebo controlled crossover protocol being used. No change in asthma control occurred during the nifedipine treatment period as assessed by serial peak flow measurements and symptom scores. Serum theophylline concentrations were significantly lower after nifedipine than after placebo (6.8 v 9.7 μg/ml) and in three patients were well below the therapeutic range (<4 μg/ml).

Although calcium antagonists reduce bronchoconstriction induced by exercise, histamine, and antigen, resting airway calibre is not altered to any great extent. They may, however, have a beneficial effect when combined with other treatment for asthma. Nifedipine potentiates the bronchodilator effect of inhaled salbutamol and intravenous terbutaline in acute studies in asthmatic subjects and it enhances the airways smooth muscle relaxation induced by theophylline in isolated guinea pig trachea. We have studied asthma control in a group of patients with stable but symptomatic asthma when nifedipine was added for two weeks to their usual asthma treatment, which included theophylline.

Several drugs are known to influence theophylline pharmacokinetics and these are important clinically in view of the narrow therapeutic range of theophylline. Nifedipine is widely used to treat angina and hypertension in asthmatic patients, in whom β adrenoceptor antagonists are contraindicated. We have therefore evaluated the effect of nifedipine on serum theophylline concentrations.

Methods

We recruited 10 patients (six male, four female), with a mean age of 56 (range 49–70) years and with stable but symptomatic asthma. All gave informed consent to participate in the study, which had been approved by East Birmingham Hospital Research and Ethical Subcommittee. All subjects had shown at least 15% reversibility in FEV1 in response to inhaled salbutamol and the FEV1/FVC ratio was less than 60% in all cases. None of the patients had had exacerbations of their asthma for at least one month before entering the study. In addition to slow release theophylline twice daily (8 am and 8 pm), all patients were using inhaled β2 agonists and inhaled corticosteroids. One patient was using a sodium cromoglycate inhaler and one was taking salbutamol spandets. All of these treatments were continued unchanged throughout the study. No patient was having oral corticosteroids or any other form of medication and none was known to have any medical condition apart from asthma. One patient smoked and his consumption of cigarettes remained constant throughout the study. No specific dietary restrictions were placed on the patients, although they were asked to keep their consumption of food, including tea and coffee as constant as possible, especially on the days they came to the clinic.

Before entry into the study the dose of slow release theophylline was adjusted to ensure serum concentrations of over 8 μg/ml. Serum concentrations of theophylline were measured six hours after the last dose of slow release theophylline (that is, at 2 pm) at each of the four clinic attendances, as described below. This is likely to represent a near peak concentration. An enzyme mediated immunoassay (Emit, Syva Company) was used. Assay batches contained quality control samples and the between batch coefficient of variation of the assay was 6.2%. The addition of an excess of nifedipine (100 ng/ml) to samples of serum...
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containing theophylline did not influence serum theophylline measurements.

Once satisfactory serum theophylline concentrations had been achieved, patients were asked to record symptoms and the best achievable peak expiratory flow (PEF) each day at 8 am, 12 noon, 6 pm, and 10 pm, before they used their bronchodilator inhaler. Symptoms included perceived severity of asthma, degree of dyspnoea, cough, and disturbance of sleep due to asthma, each being assessed on a 0–10 scale (that is, no symptoms = 0, very severe symptoms = 10). No change was made to their usual treatment during the first week, which served as a “practice” period. At the end of the week any problems with recordings were resolved and blood was taken for measurement of serum theophylline concentrations. These were found to be stable and within the therapeutic range in all 10 patients. Subjects were started on slow release nifedipine 20 mg twice daily or an identical placebo tablet twice daily in addition to their usual treatment in a randomised double blind fashion. Measurements of PEF and symptom recordings were continued for a further two weeks, after which further blood was taken for serum theophylline assay. Subjects who had received nifedipine were changed to placebo and vice versa. Recordings were continued for another two weeks, at the end of which a final serum theophylline measurement was made.

Compliance with treatment was checked by tablet counts at the end of each of the two treatment periods.

Statistical comparisons were made by means of Student’s t test.

Results

Two men (one from each limb of the study) had to withdraw because of troublesome headache, which developed within 24 hours of starting nifedipine. The remaining eight subjects tolerated the treatment without side effects.

There was no significant difference in mean PEF measurements or combined symptom scores between the placebo and the nifedipine period (table). The combined symptom score represents the sum of the means of the two week periods of recordings of the four assessments of symptoms.

The mean (SEM) serum theophylline concentration was similar after two weeks’ placebo to the concentration immediately before entry to the randomised part of the study (9.7 (0.5) µg/ml after placebo compared with 9.6 (0.9) µg/ml after one week’s practice). After nifedipine the mean theophylline concentration was significantly lower (6.8 (1.0) µg/ml) than after either the placebo or the practice period (p < 0.02 for both comparisons; figure). In three patients serum theophylline concentrations after nifedipine were 4 µg/ml or less, representing falls of 64%, 56%, and 50% from the levels seen after placebo.

Mean peak expiratory flow and combined symptom scores (with standard errors in parentheses) during placebo and nifedipine treatment periods in eight asthmatic subjects

<table>
<thead>
<tr>
<th></th>
<th>Combined symptom score</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>8 am</strong></td>
<td><strong>12 noon</strong></td>
</tr>
<tr>
<td>Placebo</td>
<td>318 (14)</td>
</tr>
<tr>
<td>Nifedipine</td>
<td>326 (17)</td>
</tr>
</tbody>
</table>

Serum theophylline concentrations in eight asthmatic subjects after two weeks’ treatment with nifedipine and placebo (means with 1 standard error).
Discussion

In this double blind placebo controlled crossover study of nifedipine treatment for two weeks there was no change in subjective or objective measurements of asthma control. The absence of any deterioration despite a fall in serum theophylline concentration while patients were taking nifedipine may reflect the stability of asthma in our patients. An alternative explanation is that the nifedipine was producing some benefit to compensate for the lower serum concentrations of theophylline. In patients with brittle asthma, who may be particularly dependent on theophylline treatment, the fall in serum theophylline concentration is potentially hazardous. Optimum serum theophylline concentrations are from 8 to 20 μg/ml. In a recent study of long term treatment with theophylline in combination with β2 agonists the effect of theophylline on PEF and FEV₁ in theophylline responsive patients increased with increasing plasma concentrations. Concentrations of 4 μg/ml or below, as found in three of our patients after nifedipine, would not be expected to be of any therapeutic value.

The fall in theophylline concentrations during nifedipine administration may be explained by the results of a study in normal volunteers by Jackson and colleagues. They found that serum theophylline concentrations after an infusion of lysine theophylline were significantly lower in the presence of nifedipine during steady state chronic oral dosing owing to an increase in the mean volume of distribution of theophylline. Alternatively, nifedipine might alter theophylline absorption, although this would not account for the results with intravenous theophylline, or nifedipine might increase theophylline clearance by increasing liver blood flow. The latter is unlikely since, although nifedipine does increase liver blood flow, theophylline is subject to low hepatic extraction with a small first pass loss. Its clearance is more dependent on the activity of liver enzymes than on changes in liver blood flow.

We suggest that, when nifedipine is used to treat angina or hypertension in patients with brittle asthma who are taking oral theophylline, serum theophylline concentrations should be monitored during the first few weeks so that appropriate dose adjustments can be made.

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References

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