RENAL RESPONSE TO HYDRALAZINE IN CHF/Pierpont et al.

Effect of Hydralazine on Renal Failure in Patients with Congestive Heart Failure

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SUMMARY Hydralazine is known to improve cardiac function in patients with congestive heart failure (CHF), but its effects on renal function in CHF are less clear. Sodium retention is known to occur with long-term use of hydralazine to treat hypertension; if this occurs in patients with CHF it could be deleterious. Therefore, in a metabolic unit we studied renal effects of hydralazine in patients with stable class III or IV CHF. In a single-blind study, the patients were given placebo twice daily for 3 days (period P-1), 100 mg of oral hydralazine twice daily for 3 days (period P-H), and placebo for 3 more days (period P-2). The average 24-hour creatinine clearance was 69.7 ± 7.7 ml/min (mean ± SEM) in P-1, increased to 76.3 ± 9.0 ml/min with hydralazine (p < 0.01) and fell again when hydralazine was stopped (P-2) to 68.5 ± 7.8 ml/min (p < 0.02). Though the slight improvement in sodium excretion was not statistically significant (60.2 ± 12.1 mEq in P-1, 64.5 ± 12.4 mEq in P-H, 52.3 ± 7.7 mEq in P-2), serum osmolality decreased from 288 ± 1.8 mosM in P-1 to 283 ± 1.9 mosM in P-H (p < 0.02) and rose to 286 ± 1.9 mosM in P-2 (NS). During the three periods, serum sodium, chloride, potassium, carbon dioxide, blood urea nitrogen, creatinine and glucose were unchanged, as were weight and urine volume. Systolic blood pressure was 109.6 ± 3.6 mm Hg in P-1, 110.1 ± 3.9 mm Hg in P-H (NS), and 114.2 ± 5.0 mm Hg in P-2 (p < 0.05). Diastolic blood pressure, heart rate and respirations were unchanged. Thus, we found no evidence of sodium or water retention during hydralazine administration in patients with CHF, and renal function was actually improved, as evidenced by the increased creatinine clearance.

HYDRAZLINE improves left ventricular performance in patients with congestive heart failure (CHF) and may be effective for long-term oral therapy of patients with chronic CHF. However, hydralazine can lead to sodium and fluid retention when used to treat hypertension, and sodium retention has been reported during administration of hydralazine in a patient with CHF. Such a response to hydralazine in patients with CHF might limit the value of this drug for long-term therapy or necessitate larger doses of diuretics. The present study assesses the effects of hydralazine on renal function and sodium balance in patients with CHF.

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Methods

Nine male patients with New York Heart Association functional class III or IV heart failure (three class IV and six class III) who were on conventional therapy were studied. The patients were ages 47–64 years (average 59 years). Three of the patients had primary myocardial disease and six had ischemic heart disease. Exclusion criteria included: insulin-dependent diabetes mellitus, severe pulmonary disease (Po2 < 50 mm Hg or PCO2 > 50 mm Hg), renal insufficiency (BUN > 40 mg/dl or creatinine > 2.0 mg/dl), abnormal serum electrolytes, angina requiring nitrate therapy, myocardial infarction within the past 3 months, significant primary heart valve disease, hypertension requiring natriuretic antihypertensive agents or heart failure due to restrictive heart disease.

After giving written informed consent, the patients were admitted to a special diagnostic and treatment unit (SDTU) of the Minneapolis Veterans Administration Medical Center for evaluation. All patients were clinically stable with a steady body weight when admitted to the SDTU. All patients were
receiving a constant dose of diuretics, and eight of nine
were taking digitalis at the time of admission. These
medications, as well as any other medication the
patient was taking regularly, were continued at a con-
stant dose throughout the study period. One patient
had been receiving isosorbide dinitrate, which was
stopped more than 24 hours before entry into the
study.

On admission to the SDTU a dietary history was
taken by a nutritionist and a diet constructed to match
the usual sodium intake for each patient. Sodium and
fluid intake were kept constant for the duration of the
study and patients were allowed their normal level of
activity. Supine blood pressure, heart rate, respira-
tions and temperature were monitored four times
daily, and the patients were weighed twice daily.

The study protocol consisted of three periods of 3
days each. The first period consisted of 3 days of
placebo administration (two tablets twice daily). This
was followed by 3 days of active hydralazine therapy,
100 mg (two tablets twice daily) and another 3-day
placebo period. The investigators, but not the patients,
were in the study when hydralazine was being given. Each morn-
ing blood was drawn for determination of urea
nour, creatinine, glucose, sodium, chloride, total
CO₂, potassium and osmolality. In the last five
patients studied, blood samples for plasma renin ac-
tivity (PRA) were drawn on the third day of each
period in the early morning with the patient supine.
Urine was collected at 8-hour intervals throughout the
study and analyzed for creatinine, potassium, sodium
and osmolality. Creatinine clearances were calculated
for each of the three daily 8-hour urine collections
using the serum creatinine value obtained at the end of
the day.

All data were analyzed using the paired t test and
differences were considered statistically significant for
p values less than 0.05. The mean of all values ob-
tained during each 3-day period were compared, and
analysis of the data was repeated by averaging the
values from the two placebo periods and comparing
them with the values from the hydralazine period.
Combining the placebo periods would tend to correct
for trends during the course of hospitalization that
might occur independent of hydralazine therapy. In
order to examine the data for possible changes within
each drug period, an additional analysis of the more
pertinent variables was performed using an analysis of
variance.

Results

Table 1 shows the mean values during each period
for mean arterial pressure (calculated as the diastolic
pressure plus one-third of the pulse pressure), heart
rate, body weight, serum creatinine, serum sodium,
serum chloride, serum potassium and urine volume.
None of these parameters changes significantly during
hydralazine therapy when compared with each
placebo period or with the average of the two placebo
periods.

Table 1. Clinical Data from the Three Study Periods

<table>
<thead>
<tr>
<th>Study period</th>
<th>P₁</th>
<th>H</th>
<th>P₃</th>
</tr>
</thead>
<tbody>
<tr>
<td>BP (mm Hg)</td>
<td>85 ± 2.1</td>
<td>84 ± 2.0</td>
<td>87 ± 3.4</td>
</tr>
<tr>
<td>HR (beats/min)</td>
<td>83 ± 3.0</td>
<td>84 ± 3.3</td>
<td>84 ± 2.7</td>
</tr>
<tr>
<td>Weight (lbs)</td>
<td>178 ± 9.8</td>
<td>170 ± 9.8</td>
<td>177 ± 10.1</td>
</tr>
<tr>
<td>Creat (mg/ml)</td>
<td>1.4 ± 0.15</td>
<td>1.4 ± 0.18</td>
<td>1.4 ± 0.16</td>
</tr>
<tr>
<td>BUN (mg/dl)</td>
<td>25 ± 3.9</td>
<td>24 ± 4.7</td>
<td>25 ± 4.0</td>
</tr>
<tr>
<td>Na⁺ (mEq/l)</td>
<td>141 ± 1.0</td>
<td>139 ± 1.5</td>
<td>140 ± 1.6</td>
</tr>
<tr>
<td>Cl⁻ (mEq/l)</td>
<td>100 ± 1.4</td>
<td>99 ± 1.9</td>
<td>99 ± 1.9</td>
</tr>
<tr>
<td>K⁺ (mEq/l)</td>
<td>4.3 ± 0.1</td>
<td>4.3 ± 0.1</td>
<td>4.2 ± 0.1</td>
</tr>
<tr>
<td>Urine vol (l/24 hr)</td>
<td>2.1 ± 0.26</td>
<td>2.2 ± 0.27</td>
<td>2.1 ± 0.27</td>
</tr>
</tbody>
</table>

Values are average ± SEM. None of the changes were
statistically significant when compared from one period to
another.

Abbreviations: BP = mean blood pressure; HR = heart
rate; Creat = creatinine; BUN = blood urea nitrogen;
Na⁺ = serum sodium; Cl⁻ = serum chloride; K⁺ = serum
potassium; P₁ = first placebo period; P₃ = second placebo
period; H = hydralazine.

were no significant changes in respiratory rate, serum
creatinine, glucose, serum total CO₂, creatinine excretion, sodium
balance or combined sodium plus potassium excretion.
Potassium excretion did not change from the first
placebo period (102 ± 10.5 mEq/24 hours) to the
drug period (103 ± 11.8 mEq/24 hours), but the
decrease to 98 ± 11.1 mEq/24 hours in the second
placebo period was statistically significant (p < 0.05)
when compared with the active drug period. Because
of this decrease during the second placebo period,
potassium excretion was statistically higher (p < 0.05)
during hydralazine therapy compared with the
average of the two placebo periods.

The effect of hydralazine on creatinine clearance,
sodium excretion and serum osmolality are shown in
figure 1. The average of all values during each period
is shown for every patient, as well as the mean ± SEM
for the whole group. Creatinine clearance increased
from an average of 69.7 ± 7.7 ml/min during the
first placebo period to 76.3 ± 9.0 ml/min during
hydralazine therapy (p < 0.01). In no patient did
creatinine clearance fall during administration of
hydralazine. The creatinine clearance fell again during
the second placebo period to 68.5 ± 7.8 ml/min,
which was significantly lower than during hydralazine
(p < 0.02). This value was not significantly different
from the first placebo period. When compared with
the average of the two placebo periods, the increase
in creatinine clearance during hydralazine was sig-
nificant (p < 0.01).

Sodium excretion increased from 60.2 ± 12.1 mEq/
24 hours during placebo to 64.5 ± 12.4 mEq/24
hours during hydralazine and decreased again to 52.3
± 7.7 mEq/24 hours after hydralazine was stopped.
These changes were not statistically significant when
the hydralazine response was compared with each
placebo period separately or with the pooled placebo
periods.
Serum osmolality decreased during hydralazine in seven of the eight patients in whom it was measured. The mean serum osmolality dropped from 288 ± 2.6 mosM to 283 ± 2.7 mosM during hydralazine (p < 0.02), and returned toward baseline during the second placebo period (286 ± 2.7 mosM). The change in serum osmolality during hydralazine compared with the average of the two placebo periods was significant (p < 0.02).

Although neither sodium excretion nor mean arterial blood pressure changed significantly during administration of hydralazine in these nine patients, it is of interest to examine the relationship of these two variables in individual patients. Figure 2 shows the changes in blood pressure and sodium excretion from the combined placebo periods to hydralazine administration for each patient. Sodium excretion was lower during hydralazine in only three of nine patients. A decrease in mean arterial blood pressure occurred during hydralazine in all three of these patients. The largest decrease in sodium excretion occurred in the patients who had the largest decrease in blood pressure. In the three patients whose mean arterial blood pressure increased, sodium excretion also increased. In no patient did the blood pressure increase and sodium excretion decrease, and three patients had an increase in sodium excretion despite a decrease in blood pressure.

Creatinine clearance and serum osmolality were further analyzed using the analysis of variance for single-factor experiments with repeated measures on the same elements. This analysis also showed the increase in creatinine clearance during hydralazine to be significant (p < 0.01), as well as the decrease in serum osmolality (p < 0.005), with no differences between the first and second placebo periods. Examination of the data for changes within each period revealed no significant differences from day 0 day in any period for either creatinine clearance or serum osmolality. We conclude that the patients were indeed stable during the initial placebo periods, and that they returned to control levels and stayed unchanged during the second placebo period, with no evidence of prolonged effect after cessation of hydralazine. Also, because there was no significant change from day to day during hydralazine therapy, there is no evidence of tolerance or cumulative effect over the 3-day hydralazine period.

PRA increased during hydralazine administration in all five patients in whom it was measured. The average PRA increased from 4.2 ± 1.18 ng/ml/hr during the first placebo period to 8.5 ± 2.05 ng/ml/hr during hydralazine (p < 0.02), and decreased slightly to 7.6 ± 3.13 ng/ml/hr during the second placebo period.

Discussion

Most studies of the renal effects of hydralazine were performed in hypertensive patients in the early 1950s. These studies were stimulated in part by Reubi's find-
ing of increased renal plasma flow with no change in glomerular filtration rate when six patients with hypertension and four normal subjects were given 10 mg of hydralazine subcutaneously. Similar results in patients with hypertension were subsequently reported by other investigators.9,14

In other studies in hypertensive patients, the renal hemodynamic responses to hydralazine were more variable.15-17 Falch et al.16 found no change in effective renal plasma flow in 11 patients with hypertension, but the dose of hydralazine was only 25 mg given orally, twice a day. In six patients with essential hypertension, Gjorup and Hilden16 observed a variable response to intravenous hydralazine. They suggested that if the decrease in blood pressure was not marked, renal blood flow increased, whereas it decreased if blood pressure dropped markedly. Our data suggest that this might also be true in patients with CHF, as sodium retention during hydralazine, if it occurred, appeared to be associated with a fall in blood pressure. Schroeder18 found no change in urea clearance or excretion of phenyl red in 50 hypertensives treated with 50-120 mg of hydralazine for 10-30 days.

Despite the variability of some of these studies, the evidence suggests an increase in renal blood flow with little change in glomerular filtration rate as the usual response of hypertensive patients given hydralazine. A similar response is likely in normal subjects if the data obtained by Moyer et al.19 in nonhypertensive dogs are applicable to humans.

Because the pathophysiology of CHF differs significantly from that of hypertension, we cannot readily assume that the renal responses to hydralazine would be the same in the two groups. In hypertension, the renal homeostatic mechanisms are altered toward increased salt excretion.20 In contrast, in CHF, the kidneys tend to retain salt and water, at least partly because of renal vasoconstriction.21 Thus, it might be expected that a vasodilator could improve salt excretion in CHF if renal perfusion is enhanced without a decrease in blood pressure.

Data on renal vascular effects of hydralazine in patients with heart failure are scarce. Two studies22,23 have reported on the renal responses of patients with mitral stenosis given hydralazine, but these patients are not comparable to our study group. Judson et al.23 included five patients with “cardiac insufficiency” in their series, but four of the five were also hypertensive. Our study therefore represents one of the first attempts to analyze the effects of hydralazine systematically in nonhypertensive patients with CHF due to left ventricular dysfunction. In an uncontrolled study, Packer et al.24 noted a brisk diuresis in some of their refractory heart failure patients placed on maintenance hydralazine therapy, but no further analysis of renal function is described. Preliminary reports of improved renal blood flow with hydralazine administration in heart failure patients have also been presented by LeJemtel et al.25 Mantle et al.26 and Cogan et al.27 In the patients of Cogan et al.,27 glomerular filtration increased insignificantly.

Because we wanted to keep the study noninvasive and were reluctant to water-load class III or IV heart failure patients, we did not measure para-aminohippurate or inulin clearance in our patients. Nonetheless, creatinine clearance increased, consistent with a rise in glomerular filtration rate. Because hydralazine is known to increase cardiac output in patients with CHF,1,2 it is likely that the improved creatinine clearance was accompanied by an increase in renal blood flow. We can thus infer that in contrast to patients with hypertension, glomerular filtration rate and renal blood flow are increased by hydralazine in patients with CHF. This response could in part be attributed to the fact that renal blood flow and glomerular filtration rate are functionally reduced in CHF, probably because of renal vasoconstriction, which can then be relaxed as the cardiac output increases.

Although we did not measure PRA in all patients, PRA did increase during the period of hydralazine administration in all five patients in whom it was measured. An increase in PRA also occurs when hydralazine is given to hypertensives, and has been thought to be related to the decrease in blood pressure. In this subgroup of five patients, renin increased even though blood pressure did not fall significantly and the control PRA tended to be elevated. In these five patients, no relationship was apparent between the change in mean blood pressure and change in PRA. Curtis et al.28 and Gavras et al.29 noted a wide range of values for resting PRA in heart failure patients receiving diuretics, with many patients having quite high levels. They postulated a role of renin in maintaining the abnormally high peripheral resistance observed in heart failure and noted hemodynamic improvement with inhibition of the renin-angiotensin system. These observations suggest that activation of the renin-angiotensin system may tend to counteract the vasodilator effect of hydralazine in CHF, and could also contribute to sustained renal vasoconstriction. The combination of a converting enzyme inhibitor with hydralazine might therefore be appropriate for vasodilator therapy in heart failure. The rapidity of onset of changes in PRA and the duration of effect have not been studied, and our data are far too few to help define the time course of changes in PRA.

In this study, neither weight loss nor diuresis occurred during the 3-day hydralazine period, despite the effect on creatinine clearance. Our patients were considered stable and reasonably well compensated before entry into the study. It is entirely possible that hydralazine administration to patients with CHF and acute volume overload would initiate or enhance diuresis. However, because of the instability of patients with volume overload during hospitalization, the present study was limited to patients who were relatively free of edema.

The decrease in serum osmolality that occurred with hydralazine is probably related to changes in serum sodium concentration. Increased sodium excretion or water retention or both could alter the serum sodium concentration and the osmolality. As we noted no increase in body weight or decrease in urine output,
the slight (but statistically insignificant) increase in sodium excretion may be the major reason for the decrease in osmolality.

The absence of a significant decrease in blood pressure or increase in heart rate is consistent with previous observations with hydralazine in heart failure.\(^1\)\(^2\) The dose of hydralazine chosen has been confirmed effective in increasing cardiac output, which allows systemic vascular resistance to fall without changing blood pressure significantly. Although we did not measure central hemodynamics, it is reasonable to presume that similar responses were attained in the patients in this study.

In summary, our data show improved renal function during hydralazine administration in patients with CHF, as manifested by increased creatinine clearance. Although an occasional patient may tend to retain sodium in response to hydralazine, this apparently is confined to patients whose blood pressure falls. Although our data suggest that sodium retention is not likely to be an important factor in the majority of patients, the patients in our study received hydralazine only for a period of 3 days. Extrapolation of these findings to chronic therapy must be made with caution, and the efficacy of long-term hydralazine therapy in CHF remains to be established by long-term, controlled clinical trials.

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References

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