STUDIES ON DIGITALIS. IX. EFFECTS OF OUABAIN ON THE NONFAILING HUMAN HEART

By DEAN T. MASON and EUGENE BRAUNWALD

(From the Cardiology Branch, National Heart Institute, Bethesda, Md.)

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In spite of the widespread use of digitalis in clinical medicine and the extensive experimental work that has been carried out with this drug, its action on the nonfailing human heart still requires clarification. A variety of experiments on open-chest, presumably normal, anesthetized dogs has clearly shown that digitalis glycosides have a positive inotropic action (1-4). The applicability of these observations to the normal heart in intact human subjects is limited because the possibility cannot be excluded that depression of myocardial contractility by the anesthetic agents and the surgical exposure of the heart modified the response to the digitalis glycoside in the animal experiments. Hemodynamic studies in subjects with normal hearts and in patients with heart disease without congestive heart failure have suggested that digitalis does not improve myocardial contractility, and may even have a depressing effect, since its administration results either in no change in the cardiac output, or in a decline (5-9). It is now clear that measurement of the cardiac output alone does not necessarily provide an assessment of ventricular contractility (10). The unavailability, in the past, of a reliable method for studying contractility in the hearts of intact human subjects has been responsible for the lack of information concerning the crucial question of how digitalis affects the normal, or the diseased but nonfailing human heart.

Recently, Sarnoff and Mitchell have defined myocardial contractility as follows: "When from any given end-diastolic pressure or fiber length, the ventricle produces more external stroke work and more external stroke power, an increase in ventricular contractility is said to have taken place. Implicit in this definition is an increased rate of development of tension when contractility increases" (11). Rushmer has also found that an increased ventricular contractility is reflected by an increase in the peak rate of development of intraventricular pressure (12). The observations of these and other investigators (1, 4, 13-19) are all compatible with the view that the rate at which ventricular pressure rises during isometric contraction is determined by the contractile state of the myocardium and by the conditions under which the ventricle contracts, i.e., the ventricular end-diastolic pressure, arterial pressure, and heart rate.

In 1927, Wiggers and Stimson demonstrated that digitalis resulted in a more rapid rise of left ventricular pressure in the dog, and they concluded that, like epinephrine, the glycosides improve myocardial contraction (1, 16). Measurement of the instantaneous rate of rise of intraventricular pressure in man has been fraught with considerable difficulty because of the serious artifacts in the pressure tracings obtained at cardiac catheterization by the standard catheter-manometer systems. The recent development of a cardiac catheter with a high-fidelity micromanometer mounted at its tip (20), however, has largely solved this problem, since it is now possible with this instrument to record intraventricular pressure pulses faithfully and without significant distortion. In a previous communication from this laboratory, the use of the continuously computed rate of change of intraventricular pressure in assessing the functional status of the heart in man was presented (21). The objective of the present investigation was to apply this technic in the determination of the effects of a cardiac glycoside, ouabain, on the normal and on the diseased but nonfailing human ventricle.

SUBJECTS AND METHODS

The effects of ouabain were determined in a total of ten patients who ranged in age between 14 and 45 years. The rate of change (dp/dt) in the right ventricular pressure pulse was studied in six patients after detailed clinical examination, as well as right and left heart catheterization, had been carried out. Four of these subjects, who were referred because of precordial systolic mur-
murs, were considered to have normal cardiovascular systems. One patient, W.A., was studied 1 year following successful closure of an atrial septal defect, and the clinical and hemodynamic studies did not reveal any abnormalities. The sixth patient, R.B., had a small coronary arteriovenous fistula emptying into the pulmonary artery. The pressures in the right side of the heart were within normal limits. There was no cardiac enlargement or diminished cardiac reserve, and the shunt into the pulmonary artery was so small that it could not be detected by the inhaled Krypton test (22) or by selective angiocardiography. The effects of ouabain on the dp/dt of the left ventricular pressure pulse was determined in four patients who had uncomplicated ostium secundum atrial septal defects; the presence of the interatrial communication permitted passage of the catheter into the left ventricle. None of these four patients had ever experienced congestive heart failure. Although they did not have normal hearts, their left ventricular end-diastolic pressures were within normal limits (23), and their atrial septal defects were not considered to have placed an abnormal hemodynamic burden on their left ventricles.

All studies were performed with the patient in the basal, postabsorptive state. After placement of the catheter into the ventricle, 15 minutes were permitted to elapse for the patient to reach a stable state. Three control measurements of intraventricular pressures and the dp/dt of the right (or left) ventricular pressure pulse were then made at 5-minute intervals. Ouabain, 0.30 mg to 0.60 mg, representing doses ranging between 0.0076 mg per kg and 0.0126 mg per kg (average = 0.0096 mg per kg), was then infused through the cardiac catheter over a 10-minute period. Measurements of ventricular pressure and dp/dt of the right (or left) ventricular pressure pulse were carried out at 5-minute intervals for the next 60 minutes.

The methods of recording intraventricular pressure and dp/dt have been presented in detail previously (21). Briefly, a Telco intracardiac micromanometer was employed. This manometer has been shown to maintain a uniform response to frequencies as high as 200 cycles per second (21). The dp/dt of ventricular pressure was continuously determined with an R-C differentiating circuit having a time constant of 9.4 x 10^{-4} seconds, which provided differentiation of linear amplitude without phase distortion to 50 cycles per second (24).

RESULTS

The results are presented in Table I. The means of the three values of the peak dp/dt de-

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<table>
<thead>
<tr>
<th>Patient</th>
<th>Age</th>
<th>Weight</th>
<th>Diagnosis</th>
<th>Ouabain dose</th>
<th>Condition</th>
<th>Vent. peak dp/dt</th>
<th>Heart rate</th>
<th>Vent. pres. s/d</th>
<th>Intensification of pre-systolic dp/dt</th>
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<td></td>
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<td>235</td>
<td>111</td>
<td>23/2</td>
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<td></td>
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<td>O</td>
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<td>65.9</td>
<td>Normal</td>
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<td>191</td>
<td>83</td>
<td>17/3</td>
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<td>O</td>
<td>210</td>
<td>73</td>
<td>18/3</td>
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<td>C</td>
<td>140</td>
<td>67</td>
<td>24/6</td>
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<td></td>
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<td>176</td>
<td>67</td>
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<td></td>
<td></td>
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<td>O</td>
<td>224</td>
<td>88</td>
<td>18/2</td>
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<tr>
<td>W.A.</td>
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<td>POASD</td>
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<td>R.B.</td>
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<td>Cor. A-V fist.</td>
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<td></td>
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<td>C</td>
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<td>64</td>
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<td>-</td>
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<td></td>
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<td></td>
<td></td>
<td>O</td>
<td>1,000</td>
<td>58</td>
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<td>C</td>
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<td>102</td>
<td>100/3</td>
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<td>R.M.</td>
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<td>53.9</td>
<td>ASD</td>
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<td>C</td>
<td>485</td>
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<td></td>
<td></td>
<td>C</td>
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<td>81</td>
<td>94/5</td>
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<td></td>
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<td></td>
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<td>O</td>
<td>1,058</td>
<td>78</td>
<td>109/4</td>
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</table>

* Abbreviations: dp/dt = rate of change; vent. pres. = ventricular pressure; s/d = systolic/diastolic; C = control observation before ouabain; O = observation after ouabain; + indicates intensification of pre-systolic dp/dt; − indicates no intensification of pre-systolic dp/dt; POASD = postoperative atrial septal defect; Cor. A-V Fist. = small coronary arteriovenous fistula; ASD = atrial septal defect.
There was little variation among these three values recorded in any given subject before the infusion of ouabain; the average deviation of the single measurements from the patient's mean value ranged from 0 to 6.6% and averaged 2.0% of the mean value in the ten patients studied.

In the six patients in whom the effects of ouabain on the right ventricular peak dp/dt were determined, the average values for this variable during the control period ranged between 140 mm Hg per second and 291 mm Hg per second. An increase in the peak dp/dt occurred in every patient, and the effect was usually appreciable by 10 minutes after the onset of injection. The maximal increases were noted between 25 and 60 minutes (average = 42 minutes) following the onset of the injection of the drug. The maximal value for the peak dp/dt exceeded the average of the control values by between 19 mm Hg per second and 177 mm Hg per second, an increase that averaged 63 mm Hg per second in the six subjects. Expressed as a percentage, the maximal value for the peak dp/dt exceeded the average of the control values by between 9.9 and 75.3%, an increase that averaged 31.5% of the control values in the six subjects. The changes in the peak dp/dt following the administration of ouabain to one of these patients, J.B., are illustrated in Figure 1.

In the four patients in whom the effects of ouabain on the left ventricular peak dp/dt were studied, the average of the peak values for this variable during the control period ranged between 677 mm Hg per second and 1,005 mm Hg per second. An increase in the peak dp/dt occurred in every patient and became maximal between 20 and 50 minutes (average = 33 minutes) following the onset of the injection of the drug. The maximal dp/dt exceeded the average of the control values by between 180 mm Hg per second and 490 mm Hg per second (average = 277 mm Hg per second). Expressed as a percentage, the maximal peak dp/dt exceeded the average of the control values by between 26.3 and 48.8% (average = 35.5%). The recording of dp/dt before and after ouabain in one of these patients, R.N., is illustrated in Figure 2.

In five of the ten patients, a small presystolic elevation of the ventricular dp/dt either became more prominent following ouabain (Figure 3) or appeared for the first time (Figure 4). These deflections, which followed the P wave of the electrocardiogram and occurred during atrial contraction, were followed by a return of the dp/dt to 0 before its major elevation during ventricular contraction. These deflections were too small to be measured with accuracy.

**DISCUSSION**

During the past several years, increasing evidence has been obtained which suggests that digitalis may exert a positive inotropic effect on the normal and the diseased but nonfailing human heart. Weissler and Grode (25) and Weissler and Warren (26) have reported that cardiac glycosides shorten the duration of mechanical systole, as determined from the phonocardiogram. In patients with heart disease but without heart failure undergoing cardiac operations, it has been
shown that cardiac glycosides augment the force of contraction, as measured with a myocardial strain-gauge arch (27). Since the strain-gauge arch was sewn to a segment of myocardium that was stretched by approximately 50% of its resting length, the reactions of the myocardial segment

**Fig. 2. Recordings of left ventricular (LV) pressure and rate of change (dp/dt) before and 30 minutes after ouabain administration.**

**Fig. 3. Recording of first derivative (dp/dt) of right ventricular (RV) pressure pulse during control period (top) and 25 minutes after ouabain administration (bottom). Letter a indicates the presystolic augmentation of the dp/dt during atrial systole, which became more prominent after ouabain.**
might not be representative of the reaction of the remainder of the ventricle. Furthermore, since the studies were carried out in patients with heart disease, under the conditions of general anesthesia with the chest and pericardium opened during total cardiopulmonary bypass and with the heart empty, the direct applicability of these observations to the unanesthetized intact patient was, of necessity, limited. The demonstration that digoxin diminished the total postexercise oxygen debt of patients with heart disease but without heart failure suggested the possibility that digitalis may be helpful in the clinical management of some of these patients (28). This study, however, also did not resolve the question of whether or not digitalis stimulates the normal or near normal heart of intact, unanesthetized individuals. Accordingly, the present investigation was undertaken.

In every patient, an increase in the rate of intraventricular pressure development (peak dp/dt) was noted, and this finding is interpreted to indicate that the glycoside exhibited a positive inotropic effect. It is now clearly appreciated that hemodynamic influences, besides the functional state of the myocardium, can affect the peak dp/dt (4). These influences are the heart rate, the ventricular end-diastolic pressure and volume, and the arterial diastolic pressure. Measurement of the peak dp/dt, however, is particularly suited for assessing the inotropic actions of digitalis in nonfailing hearts, since this drug does not appreciably alter these variables. Indeed, the slight slowing of the heart rate that occurred in five of the ten subjects would, by itself, tend to diminish the peak dp/dt. Ventricular end-diastolic pressures remained unchanged after ouabain. Ventricular end-diastolic volume, however, was not measured, and a change in this variable resulting from an alteration of ventricular compliance induced by ouabain cannot be completely excluded. The diastolic pressures in the pulmonary artery and the aorta were not measured in this investigation. However, the effects of acute digitalization on these pressures in patients without heart failure have been reported by a number of investigators (7, 29–32), and no consistent changes have been observed. Although systolic systemic arterial pressure usually rises, the diastolic pressure shows little alteration, presumably because of the concomitant slowing of heart rate. For the reasons outlined above, it is felt that the large and consistent elevations of the peak dp/dt that were observed in every patient can be attributed principally to a direct stimulating effect of ouabain on ventricular contraction. The development or intensification of a small, positive ventricular dp/dt during atrial systole in some patients suggests the possibility that this drug also stimulates the force of atrial contraction, resulting in a more rapid rise of intraventricular pressure during atrial systole.

**SUMMARY**

In four patients without heart disease and in two patients with minimal cardiac abnormalities and normal right ventricular function, 0.30 to 0.60 mg ouabain elevated the right ventricular peak rate of change (dp/dt) by 9.9 to 75.3% (average = 31.5%) of control values. In four patients
with uncomplicated atrial septal defects, in whom the left ventricular hemodynamic burden and left ventricular function were normal, 0.60 mg ouabain elevated the left ventricular peak dp/dt by 26.3 to 48.8% (average = 35.5%) of control values. These observations in intact, unanesthetized subjects indicate that ouabain is capable of stimulating the contractility of the nonfailing and of the normal human heart.

REFERENCES


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