

Enhancement by isosorbide dinitrate of haemodynamic effects of dopamine in chronic congestive cardiac failure

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SUMMARY The haemodynamic effects of dopamine and isosorbide dinitrate alone and in combination were studied in 8 patients with chronic congestive cardiac failure. In 7 patients dopamine, $6\mu\text{g}/\text{kg}$ per min, increased mean cardiac index from 2.0 to 3.0 l/min per m^2 ($P < 0.0005$); mean stroke volume index from 20 to 27 ml/m^2 ($P < 0.0025$), and mean left ventricular stroke work index from 14 to 20 g/m^2 ($P < 0.0025$). Mean left ventricular filling pressure did not change significantly. Isosorbide, 5 to 10 mg sublingually, reduced mean left ventricular filling pressure from 29 to 24 mmHg ($P < 0.0005$) and mean arterial pressure from 80 to 73 mmHg ($P < 0.01$), with no significant change in mean cardiac index, stroke volume index, or left ventricular stroke work index. When dopamine was reinfused after isosorbide administration mean cardiac index increased to 3.2 l/min per m^2 , stroke volume index to 31 ml/m^2 ($P < 0.05$ vs dopamine alone), and left ventricular stroke work index to 23 g/m^2 ($P < 0.0125$ vs dopamine alone). Mean left ventricular filling pressure rose slightly to 25 mmHg but this was significantly less than for dopamine alone ($P < 0.0005$). In an eighth patient, whose left ventricular filling pressure fell to 12 mmHg after isosorbide, the reinfusion of dopamine was associated with severe bradycardia and hypotension, which responded to the intravenous administration of atropine.

Afterload reduction with isosorbide combined with inotropic stimulation with dopamine may produce greater improvement in left ventricular performance in patients with congestive cardiac failure than either drug alone. Such treatment should be used with caution possibly in patients whose left ventricular filling pressure falls into the lower range after isosorbide alone.

Isosorbide dinitrate has been shown to be effective in reducing left ventricular filling pressure in patients with chronic congestive cardiac failure (Franciosa *et al.*, 1974; Gray *et al.*, 1975; Williams *et al.*, 1977) though the improvement in other aspects of left ventricular function such as cardiac output appears to be modest.

This study was designed to examine the possibility that the combination of isosorbide and an inotropic agent may be a more effective form of therapy for patients requiring intensive treatment for severe congestive cardiac failure. Dopamine is a short acting inotropic drug which has been shown to be effective in the treatment of congestive cardiac failure (Beregovitch *et al.*, 1974) and was, therefore, selected for the study.

Patients and methods

Eight patients with severe congestive cardiac failure were studied. Informed consent was obtained in all cases. Five patients were studied during an intensive treatment programme for severe intractable congestive cardiac failure. The remaining 3 patients were studied at the time of haemodynamic assessment for long-term isosorbide therapy for less severe, though disabling congestive cardiac failure. The aetiology was ischaemic heart disease in all but one patient who developed severe congestive cardiac failure in the early postoperative period after mitral valve replacement for mitral stenosis. This patient's left ventricular function was found to be normal at cardiac catheterisation before the operation. There was no evidence on clinical and echocardiographic grounds for prosthetic valve dysfunction. Isotope ventriculography indicated a severe reduction in

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Table 1 Patient data

Case	Age	Sex	History	Catheterisation findings	Clinical course
1	51	M	Recent mitral valve replacement; developed severe CCF postoperatively	—	Discharged on ISDN and diuretics; died after 3 months
2	62	M	3 MIs; admitted in pulmonary oedema	—	Discharged on ISDN and diuretics
3	58	M	2 MIs; severe exertional dyspnoea	Diffuse impairment of LV function; 3 vessel CAD	Discharged on ISDN and diuretics
4	52	M	2 MIs; angina and severe exertional dyspnoea	Diffuse impairment of LV function; 2 vessel disease	Discharged on ISDN and diuretics
5	52	M	Massive AMI requiring IABP assist; severe dyspnoea since discharge	—	Discharged on ISDN and diuretics
6	67	M	1 MI, angina and severe exertional dyspnoea; developed severe CCF while awaiting surgery	Large dyskinetic area of LV; 2 vessel CAD	Underwent LV aneurysmectomy; died perioperatively
7	49	M	2 MIs admitted in severe CCF	Diffuse impairment of LV function; 3 vessel CAD	IABP assist required because of systemic hypotension; died after 2 weeks
8	62	M	1 MI; angina and exertional dyspnoea	Diffuse impairment of LV function	Discharged on ISDN and diuretics

Abbreviations: CCF, congestive cardiac failure; MI, myocardial infarction; LV, left ventricle; CAD, coronary artery disease (> 50% stenosis); IABP, intra-aortic balloon pump; ISDN, isosorbide dinitrate.

left ventricular ejection fraction with diffuse hypokinesia. He was, therefore, assumed to have sustained severe heart muscle damage in the intra-operative or early postoperative period.

Patient data are presented in Table 1.

All studies were carried out in the coronary care unit. No drugs had been given within 4 hours of the start of the study though current treatment was not modified. A Waters thermodilution catheter was advanced to the pulmonary artery from a subclavian vein using a Desilet Hoffman No. 8 cannula. A 20 gauge cannula was inserted into the radial or brachial artery transcutaneously. Lead II of the electrocardiogram was recorded. Pulmonary arterial and systemic arterial pressures were measured using Elcomatic transducers and displayed on a Sanborn 780-6A Viso Scope (Hewlett-Packard).

For convenience, changes in the pulmonary arterial end-diastolic pressure were taken to represent changes in left ventricular filling pressure (Bouchard *et al.*, 1971; Mantle *et al.*, 1976; Bussmann *et al.*, 1977).

Cardiac output was measured by the thermodilution technique (Ganz and Swan, 1972) using a Waters Cardiac Output Computer.

Control readings of heart rate, pulmonary arterial and systemic arterial pressures, and cardiac output were made. In addition, the frequency of ventricular ectopic beats when present was recorded during a 10-minute control period. Dopamine 6 µg/kg per min was then infused using a Harvard pump for 15 minutes at which time measurements were repeated. The infusion was then discontinued. After 15 minutes and further control measurements isosorbide 5 mg was administered sublingually. If no fall in left ventricular filling pressure was seen a further 5 mg was given. After 10 minutes measure-

ments were repeated. Dopamine was then reinfused at the same dose rate for 15 minutes when measurements were repeated.

Although the study was completed without incident in 7 patients, in one (case 8) a pronounced sinus bradycardia (40 beats/minute with junctional escape beats) developed after 5 minutes of the second dopamine infusion. There was hypotension (systolic arterial pressure 60 mmHg) with diaphoresis associated with a profound fall in pulmonary arterial pressure (pulmonary arterial end-diastolic pressure < 10 mmHg). The bradycardia was promptly reversed by discontinuing the dopamine infusion and the administration of atropine 0.6 mg intravenously. Systemic arterial and pulmonary arterial pressures gradually returned to control within 45 minutes. Until this episode haemodynamic changes had followed a similar pattern to that observed in the other patients. There were no further complications.

CALCULATIONS

Mean arterial pressure (\overline{AP}) mmHg

$$= D + \frac{S - D}{3} \quad (\text{Yang } et al., 1972)$$

where D = arterial end-diastolic pressure and S = peak systolic arterial pressure. Mean pulmonary arterial pressure (\overline{PA}) was calculated in the same way.

Cardiac index l/min per m² = cardiac output (CO)/body surface area.

Stroke volume index (SVI) ml/m² = stroke volume/body surface area.

Left ventricular stroke work index g m/m²
= (\overline{AP}) - PAEDP) × SVI × 0.0136

where PAEDP = pulmonary arterial end-diastolic pressure.

Total systemic vascular resistance, dynes s cm⁻⁵
 $= \overline{AP}/CO \times 80$

Pulmonary vascular resistance, dynes s cm⁻⁵
 $= (\overline{PA} - PAEDP)/CO \times 80$

Statistical analysis was by the paired Student's *t* test.

Results

HAEMODYNAMIC EFFECTS

(Data for the patient who developed severe bradycardia are not included but illustrated separately in Fig. 5).

EFFECTS OF DOPAMINE

Mean cardiac index increased from 2.0 to 3.0 l/min/m² ($P < 0.0005$), stroke volume index from 20 to 27 ml/m² ($P < 0.0025$), and left ventricular stroke work index from 14 to 20 g m/m² ($P < 0.0025$). Mean heart rate increased from 105 to 114 beats/minute ($P < 0.05$), while mean pulmonary vascular resistance fell from 197 to 158 dynes s cm⁻⁵ ($P < 0.025$) and mean systemic vascular resistance from 1840 to 1328 dynes s cm⁻⁵ ($P < 0.01$). Mean

pulmonary arterial pressure, pulmonary arterial end-diastolic pressure, and mean arterial pressure did not change significantly.

EFFECTS OF ISOSORBIDE

Mean pulmonary arterial end-diastolic pressure fell from 29 to 24 mmHg ($P < 0.0005$), mean arterial pressure from 80 to 73 mmHg ($P < 0.01$), and mean pulmonary arterial pressure from 37 to 32 mmHg ($P < 0.01$). There was no significant change in mean heart rate, cardiac index, stroke volume index, pulmonary vascular resistance, and left ventricular stroke work index. Mean systemic

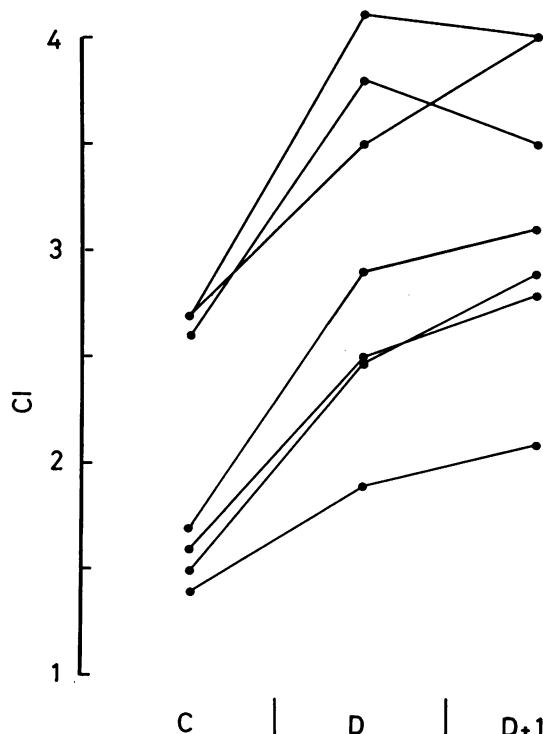


Fig. 1 Effects of dopamine and dopamine plus isosorbide on cardiac index (CI) l/min per m². C = control D = dopamine. I = isosorbide.

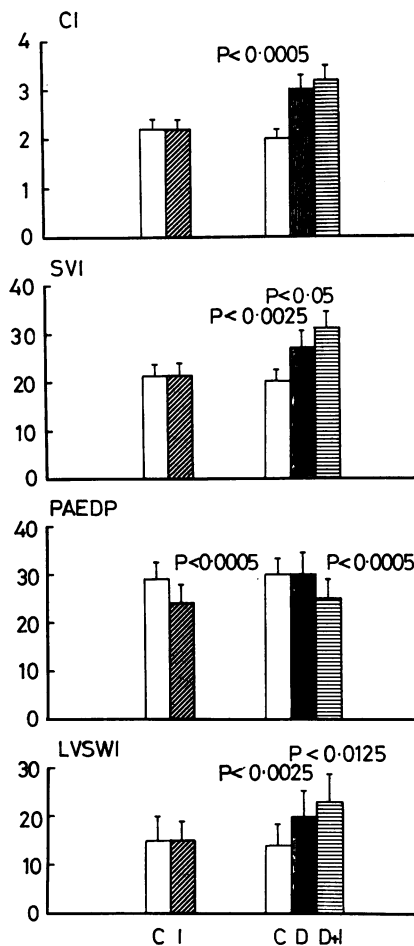


Fig. 2 Haemodynamic effects of dopamine (D) and isosorbide (I) alone and in combination. CI, cardiac index, l/min per m²; SVI, stroke volume index, ml/m²; PAEDP, pulmonary arterial end-diastolic pressure, mmHg; LVSWI, left ventricular stroke work index, g m/m²; C, control. Values indicated are mean values. Bars represent standard error of mean.

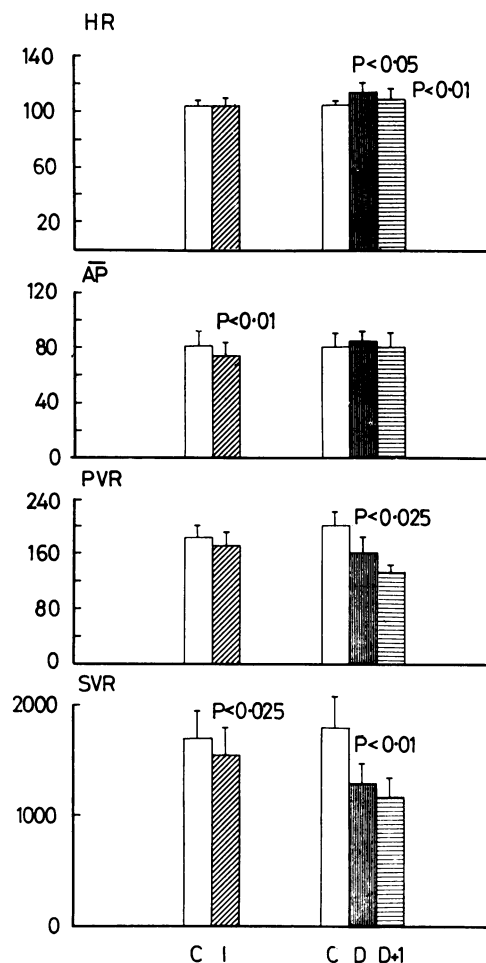


Fig. 3 Haemodynamic effects of dopamine (D) and isosorbide (I) alone and in combination. HR, heart rate, beats/min; AP, mean systemic arterial pressure, mmHg; PVR, pulmonary vascular resistance, dynes $s\ cm^{-5}$; SVR, systemic vascular resistance, dynes $s\ cm^{-5}$; C, control. Values indicated are mean values. Bars indicate standard error of mean.

vascular resistance fell from 1703 to 1554 dynes $s\ cm^{-5}$ ($P < 0.025$).

COMBINED EFFECTS OF DOPAMINE AND ISOSORBIDE

There was a further increase in cardiac index in 5 patients and a slight fall in 2 compared with dopamine alone (Fig. 1). Compared with dopamine alone stroke volume index increased from 27 to 31 ml/m² ($P < 0.05$) and left ventricular stroke work index from 20 to 23 g m/m² ($P < 0.0125$). Compared with dopamine alone mean pulmonary arterial end-diastolic pressure fell from 30 to 25 mm

Hg ($P < 0.0005$), and mean pulmonary arterial pressure from 40 to 34 mmHg ($P < 0.0025$). There was a further fall in pulmonary and systemic vascular resistance in all but one patient compared with dopamine alone, with no significant change in the mean. There was a small but significant fall in mean heart rate (mean 5 beats/minute, $P < 0.01$ dopamine alone). Mean values for mean arterial pressure did not change significantly.

Changes in mean values are illustrated in Fig. 2 and 3; individual data are shown in Tables 2 to 4.

EFFECTS ON FREQUENCY OF VENTRICULAR PREMATURE BEATS

Neither drug alone or in combination produced ventricular premature beats or significantly increased their frequency when present except in one patient (case 1) in whom they occurred frequently (35/minute) during the control period. The frequency increased to 55/minute during dopamine. During combined drug administration the frequency was 44/minute. There were no serious arrhythmias such as ventricular tachycardia. No patient complained of angina pectoris during the study.

Discussion

The results of this study indicate that inotropic stimulation of the failing heart with dopamine is effective in improving certain aspects of left ventricular function, i.e. cardiac output and stroke volume. However, another important aspect of left ventricular performance, left ventricular filling pressure, may remain severely impaired. In the patient with severe congestive cardiac failure this assumes great importance since the consequences of a high left ventricular filling pressure are those of pulmonary venous congestion and consequent dyspnoea. Further, in the acutely ill patient with severe congestive cardiac failure, pulmonary arterial oxygenation may be compromised leading to systemic arterial hypoxaemia which may further impair cardiac performance, particularly in the presence of ischaemic heart disease.

Conversely isosorbide has been shown to reduce left ventricular filling pressure with little effect on cardiac output. Indeed, falls in cardiac output have been reported (Williams *et al.*, 1977) possibly as a result of a fall in left ventricular filling pressure below optimal levels (Crexells *et al.*, 1973). Thus, though pulmonary venous congestion may be effectively relieved, an associated fall in cardiac output may compromise renal perfusion leading to further sodium and water retention in patients whose cardiac output is already reduced to a critical level.

Table 2 Haemodynamic effects of dopamine

Case No.	HR		AP		PA		PAEDP		CI		SVI		LVSWI		PVR		SVR
	C	D	C	D	C	D	C	D	C	D	C	D	C	D	C	D	C
1	118	136	71	72	25	23	18	16	1.6	2.5	14	19	10	14	207	133	2103
2	110	125	80	87	53	59	44	48	1.4	1.9	13	15	6	8	248	231	2207
3	102	112	—	—	32	28	24	20	1.7	2.9	17	26	—	—	200	118	—
4	95	85	73	80	38	39	30	26	1.5	2.5	16	29	9	21	256	254	2336
5	110	130	70	88	40	48	32	36	2.7	3.5	24	27	12	19	133	152	1167
6	100	100	125	112	43	41	32	32	2.6	3.8	27	39	34	42	200	112	2273
7	97	110	62	67	41	44	32	34	2.7	4.1	28	37	11	17	138	103	954
Mean	105	114	80	84	39	40	30	30	2.0	3.0	20	27	14	20	197	158	1840
± SEM	3.4	7.3	10.2	7.1	3.6	4.9	3.3	4.4	0.2	0.3	2.6	3.5	4.5	5.2	20	25	274
P	<0.05		NS		NS		NS		<0.0005		<0.0025		<0.0025		<0.025		

Abbreviations: HR, heart rate; AP, mean arterial pressure; PA, mean pulmonary arterial pressure; PAEDP, pulmonary arterial end-diastolic pressure; CI, cardiac index; SVI, stroke volume index; LVSWI, left ventricular stroke work index; PVR, pulmonary vascular resistance; SVR, systemic vascular resistance; SEM, standard error of mean; C, control; D, dopamine.

Table 3 Haemodynamic effects of isosorbide

Case No.	HR		AP		PA		PAEDP		CI		SVI		LVSWI		PVR		SVR
	C	I	C	I	C	I	C	I	C	I	C	I	C	I	C	I	C
1	119	120	71	64	23	17	16	10	1.8	1.6	15	13	11	9	187	207	1893
2	110	117	87	83	54	50	44	40	1.4	1.5	13	13	8	8	276	258	2400
3	108	108	—	—	29	25	22	18	1.7	1.8	16	16	—	—	181	170	—
4	90	86	63	63	38	28	30	20	1.8	2.0	20	22	9	13	213	194	1680
5	115	110	70	62	40	35	32	28	2.7	3.2	23	29	12	13	133	98	1167
6	98	98	125	110	37	33	28	24	2.8	2.7	28	28	37	33	156	160	2174
7	95	95	62	55	41	33	32	25	2.9	2.5	30	26	12	11	131	136	902
Mean	105	105	80	73	37	32	29	24	2.2	2.2	21	21	15	15	182	175	1703
± SEM	4.4	5.0	10.7	9.1	4.0	4.2	3.6	3.8	0.2	0.2	2.7	2.8	4.9	4.1	21	21	259
P	NS		<0.01		<0.01		<0.0005		NS		NS		NS		NS		

Abbreviations: HR, heart rate; AP, mean arterial pressure; PA, mean pulmonary arterial pressure; PAEDP, pulmonary arterial end-diastolic pressure; CI, cardiac index; SVI, stroke volume index; LVSWI, left ventricular stroke work index; PVR, pulmonary vascular resistance; SVR, systemic vascular resistance; SEM, standard error of mean; C, control; I, isosorbide.

Table 4 Haemodynamic effects of dopamine vs dopamine and isosorbide

Case No.	HR		AP		PA		PAEDP		CI		SVI		LVSWI		PVR		SVR
	D	D+I	D	D+I	D	D+I	D	D+I	D	D+I	D	D+I	D	D+I	D	D+I	D
1	136	130	72	65	23	16	16	10	2.5	2.8	19	22	14	16	133	102	1371
2	125	122	87	87	59	50	48	40	1.9	2.1	15	18	8	11	231	186	1832
3	112	112	—	—	28	27	20	18	2.9	3.1	26	28	—	—	118	124	—
4	85	76	80	63	39	30	26	20	2.5	2.9	29	38	21	22	254	167	1561
5	130	130	88	88	48	46	36	34	3.5	4.0	27	31	19	23	152	133	1117
6	100	94	112	120	41	33	32	24	3.8	3.5	39	37	42	48	112	124	1400
7	110	100	67	58	44	36	34	26	4.1	4.0	37	40	17	17	103	106	687
Mean	114	109	84	80	40	34	30	25	3.0	3.2	27	31	20	23	158	135	1328
± SEM	7.3	8.2	7.1	10.4	4.9	4.7	4.4	4.1	0.3	0.3	3.6	3.4	5.2	5.8	25	13	176
P	<0.01		NS		<0.0025		<0.0005		NS		<0.05		<0.0125		NS		

Abbreviations: HR, heart rate; AP, mean arterial pressure; PA, mean pulmonary arterial pressure; PAEDP, pulmonary arterial end-diastolic pressure; CI, cardiac index; SVI, stroke volume index; LVSWI, left ventricular stroke work index; PVR, pulmonary vascular resistance; SVR, systemic vascular resistance; SEM, standard error of mean; D, dopamine; I, isosorbide.

The principal actions of isosorbide on the peripheral vasculature appear to be those of arteriolar vasodilatation and venodilatation (Mason and Braunwald, 1965). The former reduces left ventricular afterload which Cohn (1973) argues is primarily responsible for the haemodynamic improvement

reported in patients with congestive cardiac failure treated with certain vasodilators; the fall in the left ventricular filling pressure observed with vasodilator therapy may be secondary to the reduction in afterload or because of dilatation of venous capacitance vessels.

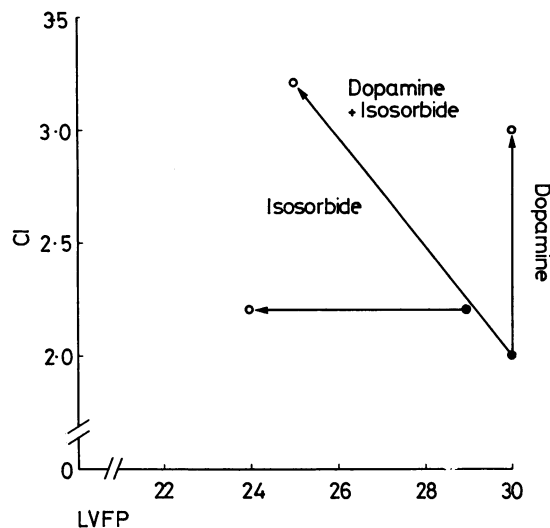


Fig. 4 Relation between left ventricular filling pressure (LVFP) mmHg and cardiac index (CI) l/min per m². The arrows show the direction of change from control (closed circles) to the intervention indicated (open circles).

Dopamine is an inotropic drug which has a vasoconstrictor effect on certain peripheral vascular beds (Mark *et al.*, 1970; Higgins *et al.*, 1973). In congestive cardiac failure these actions may have the undesirable effects of increasing left ventricular afterload and venous return; the former through arteriolar vasoconstriction and the latter through constriction of venous capacitance vessels. These effects may explain the failure of dopamine to reduce left ventricular filling pressure in congestive cardiac failure (Loeb *et al.*, 1971).

The hypothesis that combined therapy with an inotropic drug and a vasodilator will have a synergistic effect on the failing heart has been supported by the observation of this study. Thus the principal cardiac therapeutic effect of dopamine (increase in cardiac output and stroke volume) has been combined with that of isosorbide (reduction in left ventricular filling pressure) to achieve a more comprehensive improvement in left ventricular performance (Fig. 4). Afterload reduction by isosorbide, therefore, augments the effects of dopamine on stroke volume and cardiac output which in turn maintains arterial pressure. These effects, possibly combined with the venodilating effect of isosorbide, lead to a reduction in left ventricular filling pressure.

That the enhancement of the effects of an inotropic drug by a vasodilator may be achieved with an attenuation or reduction in myocardial oxygen consumption is an attractive hypothesis. A reduc-

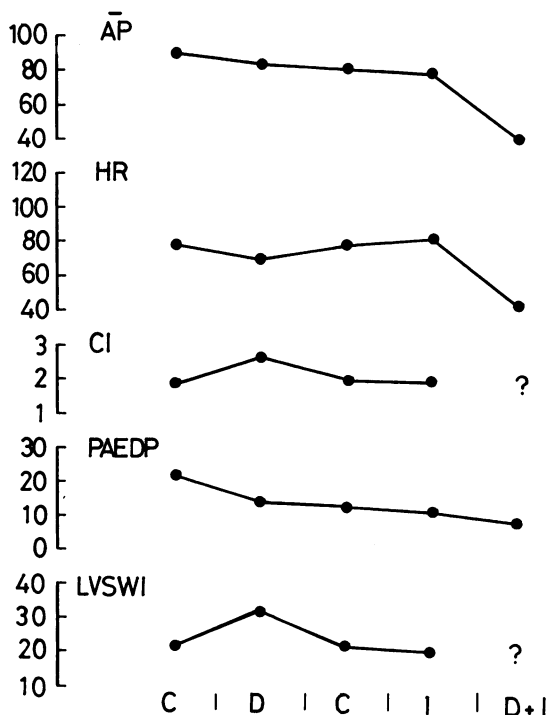


Fig. 5 Haemodynamic changes in the patient (case 8) who developed severe bradycardia and hypotension. \bar{AP} , mean systemic arterial pressure, mmHg; HR, heart rate, beats/minute; CI, cardiac index, l/min per m²; PAEDP, pulmonary arterial end-diastolic pressure, mmHg; LVSWI, left ventricular stroke work index, g m/m²; C, control; D, dopamine; I, isosorbide.

tion in left ventricular wall tension achieved by the vasodilator consequent upon reductions in preload and afterload would reduce myocardial oxygen consumption though an increase in myocardial contractility would have the opposite effect (Graham *et al.*, 1968). Further investigation is required to determine the net effect of changes in such variables which influence myocardial oxygen consumption. The attenuation of inotropically induced increases in myocardial oxygen consumption would clearly be an advantage in patients with ischaemic heart disease, particularly those with recent myocardial infarction.

Although the patients presented in this study were those with chronic congestive cardiac failure it is possible that the use of combined dopamine and isosorbide therapy may have a place in the management of patients with congestive cardiac failure or cardiogenic shock after acute myocardial infarction. In addition to the advantage of such therapy with regard to myocardial oxygen consumption discussed above, there is the possibility that it may be

applicable in hypotensive states where isosorbide therapy alone may be contraindicated, since in this study the reinfusion of dopamine after isosorbide administration counteracted the hypotensive effect of the latter.

The reason for the occurrence of the severe bradycardia in one patient is obscure. It occurred in association with a pronounced fall in left ventricular filling pressure which may have caused the observed precipitous fall in cardiac output and arterial pressure. However, such a chain of events would be expected to produce a tachycardia through the baroreceptor reflex. A similar syndrome has been reported in association with intravenous nitroglycerin administration (Come and Pitt, 1976) in patients with acute myocardial infarction. These authors suggested several possible mechanisms including vagal stimulation through pressure or length receptors in the right atrium or left ventricle. Since the bradycardia in our patient responded quickly to atropine, vagal discharge appears to be the final common pathway involved in its genesis. It is possible that a lesser degree of vagal discharge occurred in all patients when dopamine was reinfused, which may explain the observed small fall in heart rate compared with the first infusion period.

Thus, though combined therapy with dopamine and isosorbide offers a promising new approach in the treatment of severe congestive cardiac failure it should be used with caution, perhaps more so in the patient whose left ventricular filling pressure falls in the lower range after isosorbide administration.

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