TECHNIQUE

Twenty four hour continuous non-invasive finger blood pressure monitoring: a novel approach to the evaluation of treatment in patients with autonomic failure

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Abstract
Occasional sphygmomanometric readings are not an effective way of evaluating the effect of treatment in patients with hypoadrenergic orthostatic hypotension. A novel non-invasive portable device (Portapres) was used to monitor 24 hour continuous finger blood pressure before and during chronic volume expansion in a 66 year old woman with severe orthostatic hypotension. In both conditions pressures while she was standing were lowest in the morning. Her tolerance to standing and walking increased during the day and, as a consequence of a higher upright mean blood pressure, was improved after treatment. Mean blood pressure during sleep was increased after treatment.

Continuous 24 hour non-invasive finger blood pressure monitoring is a promising technique for the evaluation of the effect of treatment in patients with autonomic failure. It provides information about situations in daily life that cannot be obtained by laboratory tests or conventional sphygmomanometric measurements.

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Ordinary daily activities greatly influence blood pressure in patients with orthostatic hypotension caused by autonomic failure. Therefore, evaluation of the effect of treatment is difficult because features such as supine hypertension during night sleep or the effect of mild exercise on blood pressure cannot be detected by conventional blood pressure measurements. We report on the results obtained by a novel approach based on continuous, non-invasive, 24 hour finger blood pressure monitoring in a patient with severe hypoadrenergic orthostatic hypotension, before and during treatment with fludrocortisone.

Case report
A 66 year old woman with a 3 year history of symptomatic orthostatic hypotension was referred to our hospital. On admission sphygmomanometric systolic and diastolic blood pressure in the morning was 168/94 mm Hg supine and 86/55 mm Hg after 1–2 min standing. Physical examination and routine blood chemistry were normal. Supine noradrenaline was low (69 ng/l), with an abnormally small increase upon standing (122 ng/l). A diagnosis of pure autonomic failure was established.

Blood pressure was measured over the 24 hours by the Portapres device, a portable recorder to measure beat-to-beat finger blood pressure non-invasively in ambulatory conditions. With a trained operator, it is as easy to obtain good ambulatory recordings with the Portapres as with common non-invasive intermittent monitors (for example, Spacelabs). In addition, Portapres does not cause any discomfort to the patient, does not appreciably disturb sleeping or interfere with daily activities, and can be repeated as needed in the same subject. In a recent paper it was shown that the blood pressure values provided continuously throughout the 24 hours by the Portapres are close to those simultaneously obtained by beat-to-beat ambulatory intra-arterial recordings.

In our patient a 24 hour recording was performed before and after 7 days of treatment with fludrocortisone 0·1 mg once a day, combined with 12g head-up tilt sleep, and a 150 mmol sodium diet. During both recordings, started at 1200, the patient engaged in the same activities and performed a 15 min walk in the afternoon (1600) and the next morning (1000). Effects of standing were assessed before she went to bed (2230) and immediately after waking up (0630). Each blood pressure recording was stored on a magnetic tape cassette and further analysed by a computer to obtain beat-to-beat systolic, diastolic, mean blood pressure, and pulse pressure values. Mean 24 h diastolic blood pressure was similar before (63 mm Hg) and during treatment (64 mm Hg). Conversely, 24 h systolic
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Blood pressure and pulse pressure increased from 119 to 131 mm Hg and from 56 to 67 mm Hg. Blood pressure in both instances did not show the drop usually seen during nocturnal sleep (fig 1). Blood pressures in the afternoon and in the evening were lower before than during treatment, but were similar in the early morning. However, the severity of postural symptoms before treatment forced the patient to spend most of the morning in bed and thereby masked low upright pressures. During treatment symptoms improved markedly and morning activities were less limited. The large fall in blood pressure induced by toilet visits was reduced during treatment. Lying supine awake in the early afternoon (siesta) caused a substantial blood pressure increase particularly before treatment.

Orthostatic tolerance improved over the day. Before treatment mean finger arterial pressure after 1–2 min standing was 33 mm Hg in the morning and 56 mm Hg in the evening, the drops in blood pressure from values while supine control being 64% and 42%, respectively. During treatment upright mean blood pressure measurements in the morning (50 mm Hg) and in the evening (61 mm Hg) were higher and the drop in blood pressure was 47% and 35% from supine control. Before treatment the patient could stand without symptoms in the morning for only 1 minute, but during treatment she could stand for nearly 3 min.

Before and during treatment she could not walk for long in the morning without becoming dizzy (fig 2). The tolerance to walking in the morning amounted to 8 min only, before treatment, compared with 15 minutes during treatment. Both before and during treatment blood pressure increased in the afternoon.

Figure 1 24 h systolic and diastolic blood pressure profiles in our patient before (upper panel) and during treatment (lower panel) with fludrocortisone, head-up tilt sleep, and salt loading. Data are shown as 5 min averages. Black bars indicate, from left hand to right hand side, siesta (1400–1530), walking (w) in the afternoon, night sleep (2300–0630) and walking in the morning (w). Arrows indicate the times of meals and visits to the toilet during the night and early morning.

Figure 2 Walking in the morning (upper panels) and in the afternoon (lower panels), before (left panels) and during treatment (right panels). Walking periods are indicated by the black bars. Zero refers to the beginning of walking. Arrows mark the moments of dizziness and presyncopal symptoms that compelled the patient to sit down on a chair (recovery). Note the increase in sitting and walking blood pressure in the afternoon compared with the morning, both before and during treatment.
and the patient could walk without major symptoms. The main effect of treatment was the increase of the level of morning and afternoon systolic blood pressure and pulse pressure during walking and sitting.

**Discussion**

In our patient continuous non-invasive blood pressure monitoring during the 24 hours by the Portapres device showed the real effect of treatment on orthostatic hypotension and the benefit on postural symptoms and patient’s quality of life. It extended to everyday life the observations made by means of standardised laboratory tests—that is, that a relative small increase in mean blood pressure during treatment is sufficient to improve perfusion of the brain and to delay the occurrence of symptoms during various activities over the day.1 7 The clinical information yielded by the Portapres recording showed how treatment affected blood pressure and symptoms during everyday life at home. In the past such information has come only from beat-to-beat intra-arterial recordings,8 which are too invasive for routine use. Non-invasive blood pressure measurements—such as, conventional sphygmomanometric measurements or intermittent blood pressure samples taken by automatic portable recorder—however, cannot be used to assess blood pressure variations throughout the day.9 Our data show that these problems can be overcome by the Portapres device, which will be helpful to the clinical management of individuals with impaired autonomic control of circulation, especially when new treatments are evaluated.10

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