

Sacubitril/valsartan improves hypertension, nighttime blood pressure and heart failure: a case-report

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Objective

Sacubitril/valsartan, a neprilysin inhibitor/ angiotensin receptor blocker (ARB) combination, has been shown to effectively reduce ambulatory 24-h blood pressure in patients with hypertension. Improvements in heart failure have also been reported. This case report highlights the validity of sacubitril/valsartan for the treatment of hypertension, nocturnal hypertension and heart failure.

Clinical course up to our clinic

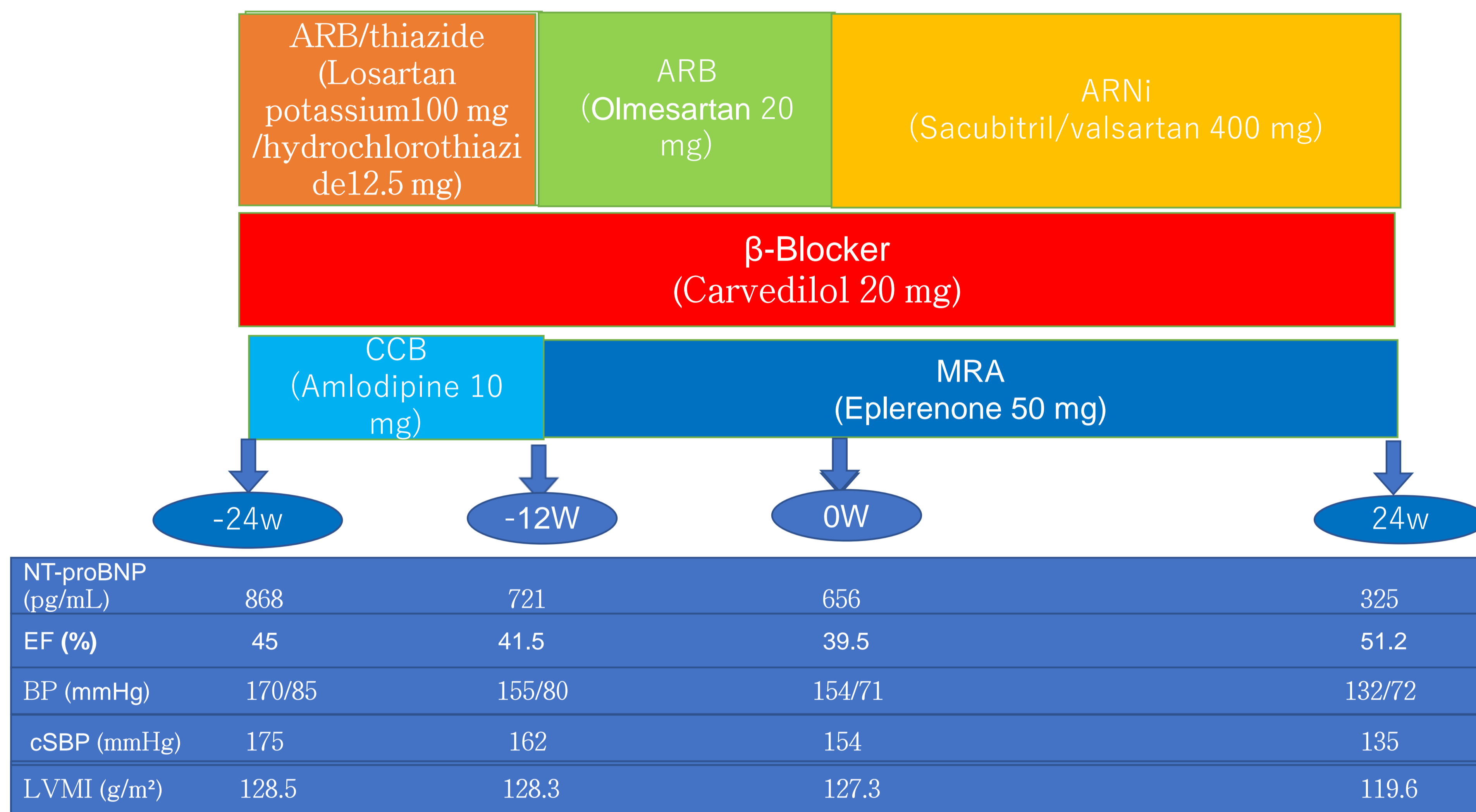
Patient: 75-year-old hypertensive male with heart failure
Chief complaint: Shortness of breath when walking on flat ground
Past history: Diagnosed with severe essential hypertension without DM, dyslipidemia, coronary artery disease, stroke, or valvular heart disease 15 years earlier.
He was referred to us from another clinic 1 year earlier.

Findings at first visit to our clinic

Symptoms: Shortness of breath on exertion, palpitations.
Vital signs: BP 175/85 mmHg, HR 95 bpm, SpO₂ (on room air) 95%
Physical examination: edema of both lower extremities
Heart failure data: NT-pro BNP 868 pg/mL, EF 40.5%,
Arterial stiffness data: cSBP 175 mmHg, AI 91%
Cardiac hypertrophy data: LVMI 128.5 g/m²
Medication prescribed by previous doctor: amlodipine 10 mg, candesartan 20 mg.

Table 1. Transition of treatment

Period 1 (12 weeks)	Period 2 (12 weeks)	Period 3 (24 weeks)
ARB/thiazide (Losartan potassium 100 mg /hydrochlorothiazide 12.5 mg)	ARB (Olmesartan 20 mg)	ARNi (Sacubitril/valsartan 400 mg)
β-Blocker (Carvedilol 20 mg)	β-Blocker (Carvedilol 20 mg)	β-Blocker (Carvedilol 20 mg)
Ca-channel blocker (CCB) (Amlodipine 10 mg)	Mineralocorticoid receptor antagonist (MRA) (Eplerenone 50 mg)	MRA (Eplerenone 50 mg)



Treatment progress at our clinic

After period 1 of treatment: blood pressure: 155/80 mmHg, heart rate: 65 bpm, NT-pro BNP: 721 pg/mL, EF: 41.5%, cSBP: 162 mmHg, LVMI: 128.3 g/m², no improvement in symptoms.

Hence, treatment changed from period 1 to period 2.

After period 2 of treatment (pre period 3): Inadequate response.

Hence, treatment changed from period 2 to period 3.

Since period 3 of treatment (pre period 4) showed an inadequate response, treatment was changed from period 3 to period 4.

Table 2. Comparison of changes in blood pressure before and after treatment period 3

	Pre period 3	Post period 3
Brachial blood pressure (mmHg)	154/71	132/72
Central systolic blood pressure (mmHg)	154	135
Heart rate (bpm)	68	65
Daytime blood pressure (mmHg)	135/68	134/63
Nocturnal blood pressure (mmHg)	136/65	121/67

Table 3. Comparison of transthoracic echocardiography results before and after treatment period 3

	Pre period 3	Post period 3
EF (%) (Simpson)	39.5	51.2
LVMI (g/m ²)	127.3	119.6
TR (m/s)	2.91	2.65
LA diameter (mm)	40.1	37.8
Septal E/e'	15.1	14.5
Septal e'	3.1	3.3

Table 4. Comparison of laboratory data before and after treatment period 3

	Pre period 3	Post period 3
NT-pro BNP (pg/mL)	656	325
HbA1c (%)	5.6	5.7
LDL-cholesterol (mg/dL)	118	108
HDL-cholesterol (mg/dL)	54	55
TG (mg/dL)	108	98
eGFR (%)	54.8	53.8
Na (mEq/L)	142	141
K (mEq/L)	4.2	4.1
Cl (mEq/L)	108	106
AST (IU/L)	21	23
ALT (IU/L)	25	26
Hb (g/dL)	12.9	13.1

Conclusion

- In our patient, changing from an ARB to sacubitril/valsartan resulted in an improvement in nighttime and central blood pressure, left ventricular hypertrophy and diastolic function, and recovery of EF, with a change from heart failure with reduced ejection fraction (HFrEF) to heart failure with preserved ejection fraction (HFpEF). His NT-proBNP levels and heart failure symptoms also improved.
- Our experience suggests that sacubitril/valsartan might be useful in patients with heart failure who have nocturnal hypertension.

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COI Disclosure

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The authors have no financial conflicts of interest to disclose concerning the presentation.