



Isolated diastolic hypertension and target organ damage: Findings from the STANISLAS cohort

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Abstract

Background: Isolated diastolic hypertension (IDH) is defined as diastolic blood pressure (DBP) ≥ 80 mmHg and systolic blood pressure (SBP) < 130 mmHg according to 2017 ACC/AHA guidelines. The effective cardiovascular risk linked to IDH is debated.

Hypothesis: IDH might contribute marginally to hypertension-related target organ damage (TOD) development.

Methods: In this cross-sectional analysis 1605 subjects from the STANISLAS cohort, a large familial longitudinal study from Eastern France, were included. Participants were categorized according to average values at 24-h ABP recording as having normal BP (SBP < 130 /DBP < 80 mmHg); combined hypertension (SBP ≥ 130 /DBP ≥ 80 mmHg or on antihypertensive treatment); IDH (SBP < 130 /DBP > 80 mmHg); isolated systolic hypertension (ISH: SBP ≥ 130 /DBP < 80 mmHg). The association between hypertension status and TOD was assessed by multivariable-adjusted logistic models.

Results: Using normotension as reference, IDH was not significantly associated with NTproBNP levels (adjusted odds ratio [OR] 1.04 [95%CI 0.82;1.32], $p = .750$), microalbuminuria (OR 0.99 [0.69; 1.42], $p = .960$), diastolic dysfunction (OR 1.53 [0.88; 2.68], $p = .130$), left ventricular (LV) mass index (OR per 10 g/m² increase 1.07 [0.95; 1.21], $p = .250$), LV longitudinal strain (global: OR 1.07 [0.99; 1.14], $p = .054$; subendocardial: OR 1.06 [0.99; 1.13], $p = .087$), carotid intima media thickness (OR 1.27 [0.79; 2.06], $p = .320$), reduced ankle-brachial index (< 0.9 ; OR 1.59 [0.19; 13.55], $p = .670$) and pulse wave velocity (PWV; OR 1.07 [0.93; 1.23], $p = .360$). In contrast, combined hypertension and ISH were independently associated with LV mass index and PWV increase (all $p \leq .01$).

Conclusions: IDH was not significantly associated with TOD. Further studies are needed to clarify the clinical role of IDH. Registration: URL: <https://www.clinicaltrials.gov>; Unique identifier: NCT01391442.

KEYWORDS

ambulatory blood pressure, isolated diastolic hypertension, target organ damage

1 | INTRODUCTION

Hypertension is a major risk factor for cardiovascular morbidity and mortality.¹ Current guidelines classified hypertension into isolated diastolic (IDH), isolated systolic (ISH), and systolic and diastolic mixed (or combined) hypertension based on the elevation of systolic and/or diastolic blood pressure (DBP) values.^{2,3} IDH is a less prevalent hypertension definition,⁴ and is classified as elevated diastolic BP with a systolic BP within the normal range.^{2,3} Thresholds for hypertension diagnosis are defined as office systolic BP values (SBP) ≥ 140 mmHg and/or diastolic BP values (DBP) ≥ 90 mmHg according to the 2018 European Society of Cardiology (ESC)/European Society of Hypertension (ESH) Guidelines.³ In 2017 American College of Cardiology (ACC)/American Heart Association (AHA) guidelines lowered the cut-off from 140/90 to 130/80 mmHg.² This change had major clinical and socioeconomic implications due to the increased number of eligible patients for treatment in a cohort classically defined at low risk for cardiovascular events.⁵⁻⁷ A recent longitudinal analysis that included 8703 adults failed to show a significant association between IDH, as defined by the 2017 ACC/AHA guidelines, and increased risk for cardiovascular outcomes.⁸ In line, previous observations demonstrated that IDH is usually not associated with cardiovascular outcomes independently of baseline systolic BP.^{5,9,10} On the other side, some studies showed a slight but significant association between diastolic hypertension and cardiovascular risk,¹¹⁻¹⁴ although the larger part of these reports did not exclusively investigate IDH but diastolic hypertension in the setting of combined hypertension. Given this context, the effective cardiovascular risk linked to IDH is debatable. The *Suivi Temporaire Annuel Non-Invasif de la Santé des Lorrains Assurés Sociaux* (STANISLAS) cohort is a longitudinal transgenerational study from the Nancy region of France characterized by a familial structure and a long follow-up (up to 23 years). In this cohort, individuals underwent to an extensive cardiovascular evaluation and the hypertensive status was evaluated by ambulatory BP monitoring (ABPM), that provides a more accurate diagnosis and a better prediction of cardiovascular risk compared to office BP.^{2,3} Moreover, several papers have suggested that 24-h average blood pressure (BP) is superior to office BP in relation to hypertension target organ damage.¹⁵ The STANISLAS cohort therefore offers the unique opportunity of studying the early changes induced by cardiovascular risk factors in initially healthy subjects.¹⁶ The aim of the present study is to determine the association between IDH, identified by 24-h ambulatory BP monitoring, and markers of target organ damage in a populational cohort with detailed cardiovascular phenotyping and long follow-up.

2 | METHODS

2.1 | Study population

This cross-sectional study is derived from the STANISLAS cohort, a single-center familial longitudinal cohort comprised of 1006 families (4295 subjects) from the Nancy region recruited in 1993-1995 at the

Center for Preventive Medicine. The cohort was established with the primary objective of investigating gene-gene and gene-environment interactions in the field of cardiovascular diseases. The families were deemed healthy, free of declared acute and/or chronic illness, in order to assess the effect of genetics on the variability of intermediate phenotypes on the transition toward disease. From 2011 to 2019, 1705 survivors of the original cohort underwent their fourth examination (STANISLAS-V4) at our department, as previously described.¹⁷ The research protocol was approved by the local Ethics Committee (Comité de Protection des Personnes Est III-Nancy-France) and all study participants gave a written informed consent to participate. The informed written consent was approved previously by the local ethics committee (ClinicalTrials.gov identifier NCT01391442).¹⁶

For the present study, 1605 adult patients (i.e., ≥ 18 years old and with ambulatory BP measurements) attending STANISLAS-V4 were included in the analyses (Figure 1). All participants were scheduled to attend the Centre d'Investigation Clinique Plurithématique Pierre Drouin at Nancy University Hospital at 8 AM after a 12- to 14-h fast. All subjects underwent blood and urine sampling for laboratory analysis. Medical history, medications, anthropometric parameters were also recorded.

2.2 | Blood pressure measurements

Office BP was measured three times in all participants, at 1-min intervals, using an electronic sphygmomanometer after the participant had rested for at least 10 min and calculated as the mean of the three measurements. As previously described,¹⁶ all participants underwent a 24-h recording of ABPM using the Spacelabs 90207 ambulatory monitor (Spacelabs Medical, WA). The monitoring cuff was placed around the participant's nondominant arm. The BP system was programmed to acquire measurements every 15 min from 6 AM to 10 PM and every 30 min from 10 PM to 6 AM. Self-reported sleep-wake times were used to divide ABP monitoring data into daytime and nocturnal periods. BP indices were calculated from 24-h, daytime, and nighttime averaged measurements according to current guidelines.¹⁸ In addition, participants had to complete a diary describing their main daily activities (e.g., eating, sleeping) and were asked to avoid excessive exercise during the 24-h recording. Central reading of the recordings was performed by a trained technician blinded to the participants' clinical features. Data were considered for further analysis if they met the following criteria: the recording lasted ≥ 24 h, $\geq 70\%$ of the expected number of readings were available, no missing data for >2 consecutive hourly intervals, and ≥ 2 valid measurements were obtained per hour. Mean 24 h diastolic and systolic BP were computed as the sum of non-missing available BP values on the period of 24 h divided by the number of non-missing readings recorded during the same period. Similarly, mean daytime and night-time diastolic and systolic BP were calculated as the sum of non-missing BP values recorded during day time or night-time divided on the number of non-missing readings recorded during the same period. Participants were categorized based on the average values at 24-h ABP recording as having normal BP (SBP < 130 /DBP < 80 mmHg; $n = 957$); combined

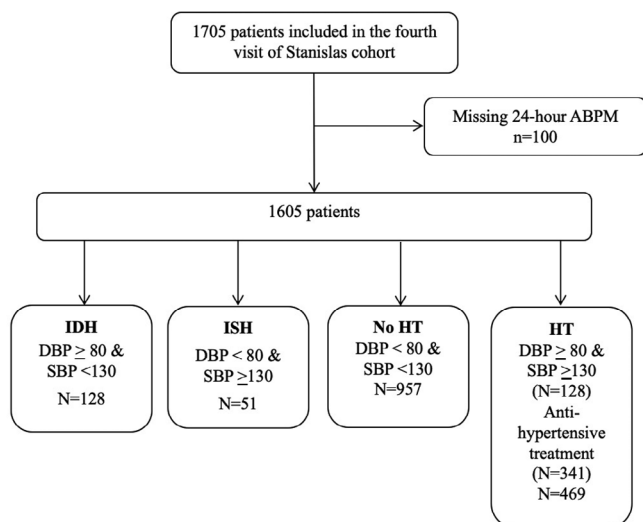


FIGURE 1 Study flowchart. ABPM, ambulatory blood pressure monitoring; DBP, diastolic blood pressure; HT, systolic-diastolic hypertension; IDH, isolated diastolic hypertension; ISH, isolated systolic hypertension; SBP, systolic blood pressure

hypertension (SDH; SBP \geq 130 /DBP \geq 80 mmHg or on antihypertensive treatment; $n = 469$); isolated diastolic hypertension (SBP <130/DBP \geq 80 mmHg; $n = 128$); isolated systolic hypertension (SBP \geq 130/DBP <80 mmHg; $n = 51$). The patient was considered a dipper if SBP and DBP reduced more than 10% between daytime and nighttime, otherwise he was considered a nondipper.¹⁸

2.3 | Assessment of markers of target organ damage

Echocardiographic exams were performed by an experienced sonographer, in the left lateral decubitus position with a commercially available standard ultrasound scanner (Vivid 9, General Electric Medical Systems, Horten, Norway) using a 2.5 MHz phased-array transducer (M5S), as previously described.¹⁶ The echo/Doppler examination was performed according to EAE/ASE recommendations¹⁹ and longitudinal left ventricular (LV) deformation parameters (strain) were obtained using speckle tracking echocardiography. LV diastolic dysfunction (DD) was assessed according to the 2009 ASE/EACVI recommendations.²⁰ We used this algorithm instead of the most recent 2016 ASE/EACVI scoring system because the new criteria commonly results in a marked reduction of DD, as also recently demonstrated by our group.²¹ Image acquisitions were shown to be highly reproducible.¹⁷

Carotid intima media thickness (cIMT) measurements were performed by high-resolution echo-tracking (Wall Track System; Pie Medical, Maastricht, The Netherlands) on the right common carotid artery at 1–2 cm below the carotid bifurcation and the retained value was obtained as the mean of four measurements.²² Both reproducibility and agreement (intra/interoperator/devices) were excellent.¹⁶

Carotid to femoral pulse wave velocity (PWV) measurement was performed with Complior® (Alam Medical, Vincennes, France) and

Sphygmocor® CVMS (AtCor Medical, Sydney, NSW, Australia) devices according to the European Network for Noninvasive Investigation of Large Arteries recommendations.²³ The protocol was previously discussed in detail.¹⁶

The ankle-brachial index (ABI) was calculated as the systolic pressure at the ankle, divided by the systolic pressure at the arm. Systolic BP was measured with a continuous Doppler machine using a BP cuff. An ABI between 0.9 and 1.2 is considered normal, while an index <0.9 suggests the presence of arterial disease.²⁴ The measurements were performed according to a standardized protocol²⁵ on the side (arm and ankle) with the highest SBP.

N-terminal pro-B-type natriuretic peptide (NT-proBNP) was measured using a multiplex assay (CVDII panel, Olink Proteomics AB®, Uppsala, Sweden). The final assay readout was expressed as normalized protein expression values, which is an arbitrary unit on a log₂ scale in which a higher value corresponds to higher protein expression.

2.4 | Statistical methods

Proportions were compared using χ^2 test and were expressed as number (proportion as percentage). Continuous variables were expressed as mean \pm SD or median (interquartile range [IQR: Q1; Q3]) and compared using one-way ANOVA and Kruskal–Wallis depending on the normality of the distribution. We focused on the association between noninvasive markers of target organ damage and IDH, with a cutoff of DBP \geq 80 mmHg and SBP <130 mmHg at 24-h ABP recording. Descriptive statistics were computed on all patients' characteristics in the sample overall and according to hypertension status. Multinomial logistic regression models were used to assess the associations between the dependent variable hypertension (HT) categories (No HT [reference], IDH, ISH, and SDH) and independent variables (echocardiographic, vascular and biological markers of target organ damage). The log linearity assumptions of the relationship between HT status and the continuous variables of target organ damage were assessed using restricted cubic splines according to the Harrell's rule.²⁶ We use the Wald test for assessing the linearity assumptions. The variables were then categorized, as required, to meet the model assumptions. Each model was adjusted for gender, age, waist circumference, smoking status, total cholesterol, glycemia, lipid lowering agents, self-declared hypertension status, glomerular filtration rate, and hemoglobin. The two-tailed significance level was set at $p < .05$. All analyses were performed using SAS version 9.4.6 (SAS Institute Inc., Cary, NC) and R version 3.6.1 (2019-07-05).

3 | RESULTS

3.1 | Participants' baseline characteristics

The characteristics of the population are given in Table 1. The study examined 1605 subjects, 128 with IDH, 51 with ISH, 469 with SDH

TABLE 1 Population characteristics in the overall cohort and according to hypertension status

Characteristics	Overall (N = 1605)	IDH (N = 128) SBP <130/DBP >80 mmHg	ISH (N = 51) SBP ≥130/DBP <80 mmHg	SDH (N = 469) SBP ≥130/DBP ≥80 mmHg	No hypertension (N = 957) SBP <130/DBP <80 mmHg	p-value
Age, years	56 [35; 60]	55 [36; 59]	58 [33; 63]	59 [56; 63]	41 [33; 59]	<.0001 ^a
Male, n (%)	774 (48.2)	75 (58.6)	35 (68.6)	282 (60.1)	382 (39.9)	<.0001 ^a
BMI, Kg/m ²	25.2 [22.5; 28.6]	25.0 [22.3; 27.8]	24.9 [23.8; 28.3]	27.8 [24.8; 31.8]	24.2 [21.9; 27.3]	<.0001 ^a
Waist circumference (cm)	90.0 [80.0; 99.0]	91.0 [81.0; 98.0]	91.0 [85.0; 99.0]	97.0 [89.0; 106.0]	85.0 [77.0; 94.0]	<.0001 ^a
Diabetes, n (%)	67 (4.2)	2 (1.6)	0 (0.0)	48 (10.3)	17 (1.8)	<.0001 ^a
Hypertension duration, years	NA	2.5 [2.0; 9.0]	5.0 [1.0; 10.0]	8.0 [4.0; 13.0]	NA	.13
PVD, n (%)	8 (0.5)	0 (0.0)	1 (2.0)	4 (0.9)	3 (0.3)	.19
Smoker status, n (%)						
Current	330 (20.7)	24 (18.8)	9 (18.0)	73 (15.7)	224 (23.5)	<.0001 ^a
Former	517 (32.4)	45 (35.2)	12 (24.0)	202 (43.4)	258 (27.0)	
Never	751 (47.0)	59 (46.1)	29 (58.0)	190 (40.9)	473 (49.5)	
Heart rate, bpm	63 (58; 69)	64 (59; 70)	61 (54; 70)	64 (58; 70)	63 (58; 69)	.15
Hemoglobin, g/L	14.71 ± 1.20	15.04 ± 1.11	14.91 ± 1.12	14.86 ± 1.22	14.59 ± 1.19	<.0001 ^a
Total cholesterol, g/L	2.1 ± 0.4	2.1 ± 0.3	2.0 ± 0.4	2.1 ± 0.4	2.1 ± 0.4	.3
LDL, g/L	1.34 ± 0.34	1.34 ± 0.31	1.29 ± 0.35	1.34 ± 0.36	1.34 ± 0.33	.75
HDL, g/L	0.57 [0.48; 0.66]	0.56 [0.47; 0.67]	0.54 [0.45; 0.61]	0.54 [0.45; 0.62]	0.58 [0.49; 0.68]	<.0001 ^a
Triglycerides, g/L	0.92 [0.68; 1.27]	0.93 [0.65; 1.39]	0.87 [0.66; 1.14]	1.12 [0.81; 1.49]	0.85 [0.64; 1.14]	<.0001 ^a
Glycemia, g/L	0.88 [0.83; 0.95]	0.88 [0.81; 0.93]	0.89 [0.85; 0.95]	0.94 [0.87; 1.03]	0.87 [0.81; 0.92]	<.0001 ^a
eGFR CKD-EPI, ml/min/1.73m ²	96.1 [86.7; 106.6]	95.7 [87.7; 105.7]	95.0 [87.0; 107.7]	91.4 [79.7; 99.2]	98.2 [89.4; 110.4]	<.0001 ^a
C-reactive protein, mg/L	1.5 [0.7; 3.2]	1.5 [0.9; 3.3]	1.1 [0.7; 3.0]	2.0 [1.0; 3.7]	1.3 [0.7; 2.8]	<.0001 ^a
Microalbuminuria, mg/L	6.0 [3.6; 9.9]	6.4 [3.4; 11.2]	6.1 [3.4; 11.9]	6.6 [3.8; 11.6]	5.7 [3.5; 9.1]	.009 ^a
NT-ProBNP (Olink)	3.55 [2.87; 4.28]	3.40 [2.81; 4.27]	3.66 [2.63; 4.36]	3.68 [3.04; 4.55]	3.49 [2.81; 4.17]	<.0001 ^a
Lipid lowering therapy, n (%)	257 (16.0)	9 (7.0)	8 (15.7)	172 (36.7)	68 (7.1)	<.0001 ^a
Diabetes treatment, n (%)	59 (3.7)	2 (1.6)	0 (0.0)	46 (9.8)	11 (1.1)	<.0001 ^a
Antihypertensive drugs, n (%)	341 (21.2)	0	0	341 (72.7)	0	NA
^b Non-dipper status, n (%)	669 (48.6)	60 (57.1)	25 (61.0)	174 (44.7)	410 (48.7)	.048
LV-GLS, %	-20.8 [-22.5; -18.8]	-19.9 [-22.0; -18.4]	-19.9 [-21.6; -18.1]	-20.3 [-22.2; -17.9]	-21.1 [-22.7; -19.2]	<.0001 ^a
Left ventricular mass index, g/m ²	73.9 [62.8; 87.4]	73.5 [64.4; 87.1]	83.6 [71.6; 94.9]	82.4 [71.9; 96.5]	69.1 [59.4; 81.6]	<.0001 ^a
Left ventricular hypertrophy, n (%)	233 (14.7)	18 (14.4)	10 (20.4)	109 (23.7)	96 (10.1)	<.0001 ^a
Left atrial volume index, ml/m ²	21.9 [17.7; 26.5]	21.3 [17.5; 25.7]	22.5 [19.0; 26.6]	23.7 [18.6; 28.9]	21.3 [17.3; 25.6]	<.0001 ^a
E/e' ratio	6.2 [5.2; 7.4]	6.2 [5.3; 7.5]	6.6 [5.9; 8.4]	7.1 [6.0; 8.4]	5.8 [5.0; 6.8]	<.0001 ^a
E/A ratio	1.10 [0.87; 1.41]	1.07 [0.84; 1.28]	1.19 [0.93; 1.48]	0.91 [0.78; 1.13]	1.20 [0.96; 1.53]	<.0001 ^a

(Continues)

TABLE 1 (Continued)

Characteristics	Overall (N = 1605)	IDH (N = 128) SBP <130/DBP >80 mmHg	ISH (N = 51) SBP ≥130/DBP <80 mmHg	SDH (N = 469) SBP ≥130/DBP ≥80 mmHg	No hypertension (N = 957) SBP <130/DBP <80 mmHg	p-value
Diastolic dysfunction, n (%)	293 (18.7)	23 (18.5)	10 (20.0)	157 (34.2)	103 (11.0)	<.0001 ^a
LVEDV index, ml/m ²	47.8 [40.0; 56.5]	48.9 [39.1; 61.4]	47.4 [39.2; 61.7]	47.1 [38.9; 57.2]	47.8 [40.4; 55.7]	.35
LVESV index, ml/m ²	16.3 [12.9; 20.5]	17.4 [12.9; 22.1]	16.4 [13.3; 22.5]	15.8 [11.9; 20.2]	16.5 [13.4; 20.4]	.023 ^a
Left ventricular ejection fraction, %	65.4 [61.2; 69.4]	65.4 [60.2; 69.8]	66.4 [60.1; 70.8]	66.4 [62.1; 70.8]	64.9 [61.0; 68.7]	.0009 ^a
Pulse wave velocity, m/s	8.2 [7.3; 9.3]	8.4 [7.6; 9.1]	9.2 [7.8; 10.3]	9.1 [8.0; 10.3]	7.8 [7.1; 8.8]	<.0001 ^a
Carotid intima media thickness, μm	616.0 [526.0; 717.2]	634.3 [537.0; 722.0]	628.0 [544.0; 766.0]	692.0 [603.2; 791.0]	579.0 [500.0; 674.0]	<.0001 ^a
Ankle-brachial index < 0.9, n (%)	16 (1.0)	1 (0.8)	1 (2.0)	5 (1.1)	9 (1.0)	.90

Note: Data are presented as mean ± SD or median and interquartile range [Q1; Q3].

Abbreviations: BMI, body mass index; CV, cardiovascular; DBP, diastolic blood pressure; eGFR, estimated glomerular filtration rate; GLS, global longitudinal strain; IDH, isolated diastolic hypertension; LV, left ventricular; LVEDV, left ventricular end-diastolic volume; LVESV, left ventricular end-systolic volume; NA, not applicable; PVD, peripheral vascular disease; SBP, systolic blood pressure; SDH, systolic-diastolic mixed hypertension.

^aStatistically significant difference between groups.

^b1377 patients available for analysis.

and 957 without hypertension. Overall, the mean age was 56 (IQR 35; 60) years, 48.2% were male, the mean BMI was 25.2 (IQR 22.5; 28.6) kg/m² and the mean estimated glomerular filtration rate was 96.1 [86.7; 106.6] ml/min/1.73 m². A history of diabetes, hypercholesterolemia and current smoking habit was identified in the 4.2%, 16%, and 20.7% of the sample, respectively. Participants with IDH were more likely to be younger, women and current smoker than those who had other types of hypertension, but not of normotensives. Non-dipper BP pattern was slightly more prevalent in IDH and ISH compared with SDH.

Subjects with IDH showed higher prevalence of left ventricular hypertrophy and diastolic dysfunction compared to normotensives, but lower than other groups of hypertensives. Among other markers of target organ damage, microalbuminuria level and cIMT were higher in participants with IDH compared to those without hypertension and ISH, but lower than SDH. On the contrary, estimated glomerular filtration rate and LV-GLS showed lower values in IDH compared to normotensives, and intermediate between ISH and SDH. Finally, participants with IDH showed the lower value of NTproBNP and similar prevalence of pathologically reduced ABI compared to other BP categories.

3.2 | Associations between hypertension classes and markers of target organ damage

In models adjusted for gender, age, waist circumference, smoking status, total cholesterol, glycemia, lipid lowering agents, self-declared hypertension status, glomerular filtration rate and hemoglobin, the majority of associations in the “crude model” between IDH and markers of target organ damage became no longer significant (Table 2). In particular, with normotension as reference, IDH was not significantly associated with NTproBNP levels ($p = .750$), microalbuminuria ($p = .960$), diastolic dysfunction ($p = .130$), LV mass index ($p = .250$), LV longitudinal strain (global, $p = .054$; subendocardial, $p = .087$), cIMT ($p = .320$), pathologically reduced ABI (<0.9 ; $p = .670$) and PWV ($p = .360$). In contrast, mixed hypertension and ISH appeared strongly and directly associated with LV mass index and PWV increase, meanwhile only SDH was associated with NTproBNP concentrations (all $p \leq .010$). Using spline-based analyses, we did not find any evidence of a nonlinear association between hypertension classes (compared to normal BP) and markers of target organ damage, except for the association between SDH and NTproBNP (Figure 2). Sensitivity analysis based on office BP measurements showed similar findings (Table S1).

4 | DISCUSSION

The key finding of our study was that an isolated increase in diastolic BP, without a concurrent rise in systolic values at 24-h ABP recording, was not significantly associated with markers of target organ damage. Concurrently, we showed that combined hypertension was associated

TABLE 2 Crude and adjusted association between markers of target organ damage and hypertension categories

	Blood pressure categories Reference: no hypertension	Univariate		Multivariable ^c	
		Odds ratio [95% CI]	p-value	Odds ratio [95% CI]	p-value
		1.00		1.00	
Left ventricular mass index, per increment of 10 g/m ²	Systolic-diastolic hypertension	1.49 [1.40; 1.59]	<.0001 ^a	1.19 [1.09; 1.32]	.0003 ^a
	Isolated diastolic hypertension	1.19 [1.07; 1.32]	.001 ^a	1.07 [0.95; 1.21]	.250
	Isolated systolic hypertension	1.41 [1.22; 1.63]	<.0001 ^a	1.25 [1.05; 1.47]	.010 ^a
Left ventricular hypertrophy, n (%)	Systolic-diastolic hypertension	2.76 [2.05; 3.74]	<.0001 ^a	1.52 [0.98; 2.35]	.064
	Isolated diastolic hypertension	1.49 [0.87; 2.57]	.150	1.22 [0.69; 2.18]	.490
	Isolated systolic hypertension	2.28 [1.10; 4.70]	.026 ^a	2.03 [0.4; 4.39]	.073
Left atrial volume index, per increment of 10 ml/m ²	Systolic-diastolic hypertension	1.67 [1.43; 1.95]	<.0001 ^a	1.39 [1.11; 1.77]	.005 ^a
	Isolated diastolic hypertension	0.98 [0.74; 1.30]	.910	0.86 [0.64; 1.17]	.340
	Isolated systolic hypertension	1.45 [0.99; 2.11]	.054	1.22 [0.81; 1.84]	.340
LV-GLS, %	Systolic-diastolic hypertension	1.11 [1.07; 1.16]	<.0001 ^a	1.04 [0.98; 1.10]	.170
	Isolated diastolic hypertension	1.09 [1.03; 1.16]	.006 ^a	1.07 [0.99; 1.14]	.054
	Isolated systolic hypertension	1.09 [0.99; 1.19]	.057	1.04 [0.94; 1.15]	.450
Subendocardial LV-GLS, %	Systolic-diastolic hypertension	1.09 [1.06; 1.14]	<.0001 ^a	1.03 [0.97; 1.09]	.270
	Isolated diastolic hypertension	1.08 [0.99; 1.17]	.012 ^a	1.06 [0.99; 1.13]	.087
	Isolated systolic hypertension	1.08 [0.99; 1.17]	.075	1.03 [0.94; 1.13]	.510
E/e' ratio	Systolic-diastolic hypertension	1.56 [1.45; 1.68]	<.0001 ^a	1.17 [1.05; 1.29]	.003 ^a
	Isolated diastolic hypertension	1.19 [1.06; 1.34]	.003 ^a	1.15 [1.01; 1.30]	.034 ^a
	Isolated systolic hypertension	1.42 [1.23; 1.65]	<.0001 ^a	1.36 [1.15; 1.60]	.0002 ^a
E/A ratio	Systolic-diastolic hypertension	0.13 [0.09; 0.18]	<.0001 ^a	0.85 [0.48; 1.50]	.570
	Isolated diastolic hypertension	0.47 [0.29; 0.77]	.003 ^a	0.69 [0.36; 1.31]	.250
	Isolated systolic hypertension	0.87 [0.45; 1.69]	.69	1.65 [0.79; 3.46]	.180
Diastolic dysfunction, n (%)	Systolic-diastolic hypertension	4.19 [3.17; 5.56]	<.0001 ^a	1.53 [0.99; 2.35]	.055
	Isolated diastolic hypertension	1.84 [1.12; 3.02]	.016 ^a	1.53 [0.88; 2.68]	.130
	Isolated systolic hypertension	2.02 [0.98; 4.16]	.057	1.60 [0.71; 3.62]	.260
LVEDV index, per increment of 10 ml/m ²	Systolic-diastolic hypertension	0.99 [0.91; 1.08]	.850	0.99 [0.87; 1.14]	.980
	Isolated diastolic hypertension	1.13 [0.98; 1.29]	.096	1.13 [0.96; 1.32]	.140
	Isolated systolic hypertension	1.11 [0.90; 1.37]	.320	0.99 [0.79; 1.25]	.970
LVESV index, per increment of 10 ml/m ²	Systolic-diastolic hypertension	0.90 [0.75; 1.08]	.260	1.14 [0.88; 1.47]	.320
	Isolated diastolic hypertension	1.23 [0.95; 1.59]	.120	1.23 [0.92; 1.65]	.170
	Isolated systolic hypertension	1.13 [0.75; 1.70]	.560	0.98 [0.62; 1.55]	.930
^b LV ejection fraction, %	Systolic-diastolic hypertension	1.55 [1.23; 1.94]	.0002 ^a	0.99 [0.70; 1.39]	.950
	Isolated diastolic hypertension	1.22 [0.83; 1.77]	.310	1.14 [0.76; 1.69]	.530
	Isolated systolic hypertension	1.39 [0.78; 2.44]	.260	1.30 [0.71; 2.37]	.390
^b Carotid intima media thickness, μm	Systolic-diastolic hypertension	3.94 [3.07; 5.05]	<.0001 ^a	1.05 [0.71; 1.57]	.800
	Isolated diastolic hypertension	1.61 [1.11; 2.34]	.012 ^a	1.27 [0.79; 2.06]	.320
	Isolated systolic hypertension	1.55 [0.88; 2.73]	.130	0.85 [0.42; 1.76]	.670
Pulse wave velocity, m/s	Systolic-diastolic hypertension	1.57 [1.46; 1.69]	<.0001 ^a	1.17 [1.05; 1.31]	.006 ^a
	Isolated diastolic hypertension	1.21 [1.08; 1.37]	.002 ^a	1.07 [0.93; 1.23]	.360
	Isolated systolic hypertension	1.49 [1.29; 1.74]	<.0001 ^a	1.30 [1.08; 1.56]	.005 ^a
Ankle-brachial index < 0.9	Systolic-diastolic hypertension	1.14 [0.38; 3.42]	.81	2.03 [0.36; 11.40]	.42
	Isolated diastolic hypertension	0.85 [0.11; 6.73]	.87	1.59 [0.19; 13.55]	.67
	Isolated systolic hypertension	2.12 [0.26; 17.09]	.48	4.49 [0.49; 41.44]	.19
Microalbuminuria, per increment of 100 mg/L	Systolic-diastolic hypertension	1.25 [0.97; 1.61]	.089	0.94 [0.74; 1.21]	.630
	Isolated diastolic hypertension	0.99 [0.54; 1.85]	1.00	0.99 [0.69; 1.42]	.960
	Isolated systolic hypertension	0.97 [0.34; 2.75]	.950	0.87 [0.36; 2.08]	.750

(Continues)

TABLE 2 (Continued)

	Blood pressure categories Reference: no hypertension	Univariate		Multivariable ^c	
		Odds ratio [95% CI]	p-value	Odds ratio [95% CI]	p-value
C-reactive protein, per increment of 10 mg/L	Systolic-diastolic hypertension	1.54 [1.23; 1.92]	.0002 ^a	1.25 [0.92; 1.71]	.160
	Isolated diastolic hypertension	1.28 [0.89; 1.83]	.170	1.32 [0.93; 1.87]	.120
	Isolated systolic hypertension	0.91 [0.39; 2.08]	.820	0.91 [0.41; 2.02]	.810
NT-ProBNP, (Olink)	Systolic-diastolic hypertension	1.40 [1.246; 1.58]	<.0001 ^a	1.29 [1.07; 1.56]	.009 ^a
	Isolated diastolic hypertension	0.99 [0.81; 1.13]	.950	1.04 [0.82; 1.32]	.750
	Isolated systolic hypertension	1.03 [0.76; 1.39]	.830	1.04 [0.74; 1.47]	.820

Abbreviations: GLS, global longitudinal strain; LV, left ventricular; LVEDV, left ventricular end-diastolic volume; LVESV, left ventricular end-systolic volume.

^aStatistically significant.

^bLog linearity assumption not respected. Variable categorized according to the median value (≥median).

^cMultivariate adjusted for: gender, age, waist circumference, smoking status, total cholesterol, glycemia, lipid lowering agents, hypertension status, glomerular filtration rate, and hemoglobin.

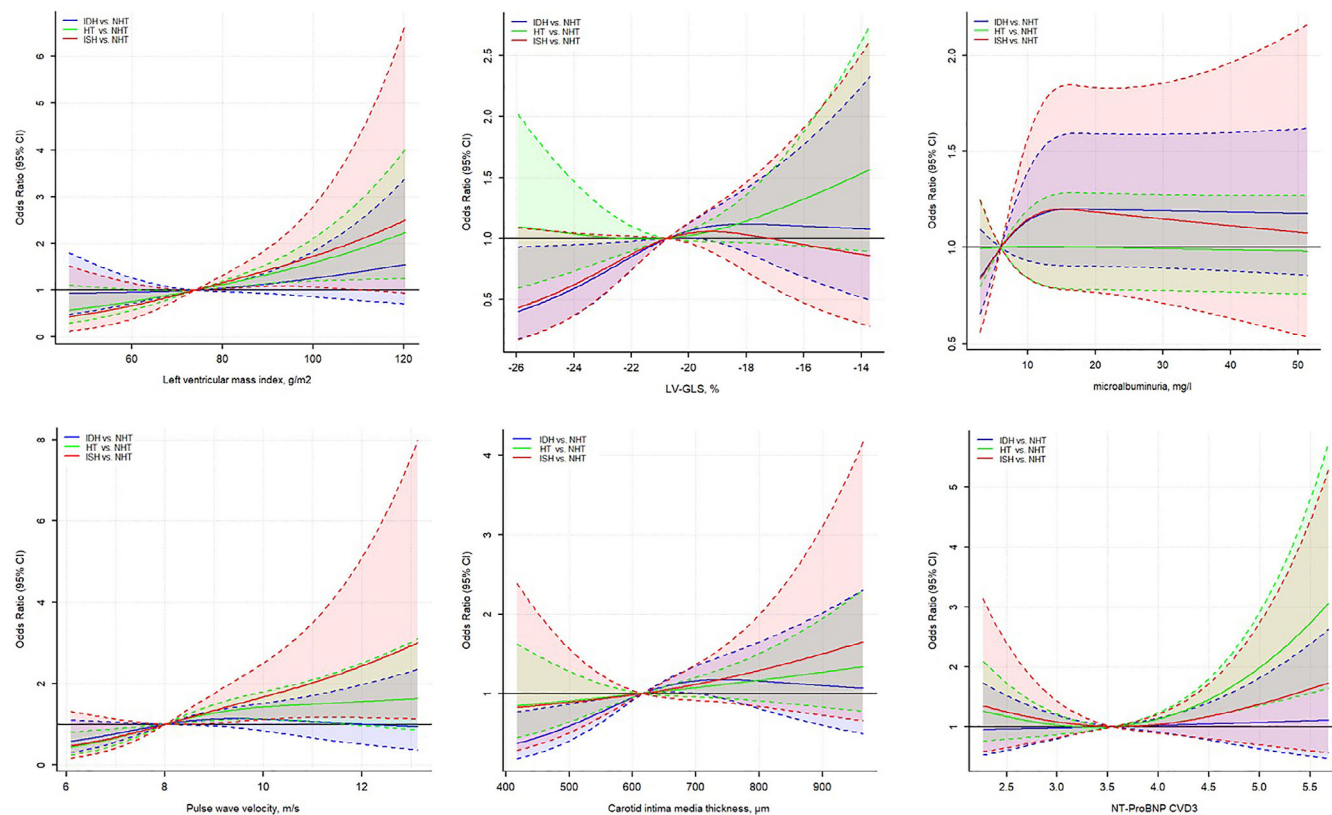


FIGURE 2 “Spline” graphical representation of the association between hypertension categories and markers of target organ damage. Solid line: hazard ratio; dashed lines: 95% confidence interval. Blue line: isolated diastolic hypertension (IDH); green line: systolic-diastolic hypertension (HT); red line: isolated systolic hypertension (ISH). GLS, global longitudinal strain

with a higher NTproBNP concentration, meanwhile both SDH and ISH were correlated with LV mass index and PWV increase.

While a raise in systolic BP was previously reported to be associated with an increased risk of cardiovascular disease and adverse outcomes,²⁷ diastolic hypertension seems to generally confer a low

cardiovascular risk.⁵⁻⁷ In a recent longitudinal analysis that included 8703 US adults from National Health and Nutrition Examination Survey (NHANES) III (1988–1994), NHANES 1999–2014 and the Give Us a Clue to Cancer and Heart Disease (CLUE) II cohort, there was no significant association between IDH, as defined by the 2017

ACC/AHA guidelines (diastolic BP \geq 80 mmHg), and incident atherosclerotic cardiovascular disease, heart failure, or chronic kidney disease.⁸ Another study conducted by ABP monitoring on 8,341 untreated subjects showed that IDH (diastolic BP \geq 80 mmHg) did not increase the risk of total mortality, cardiovascular mortality or stroke, while ISH (systolic BP \geq 130 mmHg) and mixed hypertension were associated with increased cardiovascular risk.²⁸ It should be noted that few other studies showed an association between diastolic hypertension and cardiovascular risk,^{11–14} together with a beneficial effect of antihypertensive treatment on cardiovascular mortality and morbidity. Although these evidences might appear in contradiction with the abovementioned observations, the larger part of these reports did not exclusively investigate IDH but diastolic hypertension in the setting of SDH, without differentiating the effect of solely diastolic or systolic BP lowering.

Subtle damage to certain organs can be detected in hypertensive patients early in the disease and can occur years before overt clinical events occur.²⁹ Evaluation of markers of target organ damage constitute an easy to assess, early and reliable surrogate of hypertension-related clinical events.²⁴ In fact, it has been shown that target organ damage have an independent prognostic significance and strongly increases cardiovascular risk, irrespectively of the involved structure (heart, kidney, brain, or vessels).³ Wei et al.³⁰ explored the association of target organ damage with 24-h systolic and diastolic BP levels and ambulatory hypertension subtypes in a large cohort of untreated Chinese patients. They found that 24-h SBP and mixed hypertension were the major determinants of target organ damage and its severity irrespectively of age and target organ, whereas 24-h diastolic BP and IDH only related to the urinary albumin/creatinine ratio below middle age. Another study by Lin et al.³¹ investigated the impact of ambulatory ISH, SDH and IDH (cut-off for systolic BP \geq 140 mmHg and diastolic BP \geq 90 mmHg) on target organ damage in a population of 171 subjects, showing that IDH had only a marginal effect in determining target organ damage when compared to ISH and mixed hypertension. In a large metanalysis that involved 2485 patients with and without hypertension, SBP was more closely associated with target organ damage (LV mass index, cIMT, PWV, and urinary protein excretion) compared to diastolic BP.³² In a study conducted in 972 hypertensive patients at high cardiovascular risk, systolic BP was showed to be linearly associated with ABI, while diastolic BP was not.³³ In line, Powell et al.³⁴ showed a strong association between ISH and SDH and incident symptomatic peripheral artery disease in middle-aged and older American women, whereas IDH was not associated with an increased risk. Finally, in a previous analysis of STANISLAS cohort, Ferreira et al.²² showed a linear positive association between systolic BP increase and cIMT thickness, meanwhile the association with diastolic BP was weaker and not significant.

We also demonstrated that, with normotension as reference, combined hypertension was associated with a higher NTproBNP concentration, meanwhile both SDH and ISH, but not IDH, were correlated with LV mass index and PWV increase. Prior observations suggested that LV mass is more closely related to systolic BP,

whereas LV wall thickness correlates better with diastolic BP.^{32,35,36} Our finding is in line with previous evidences and corroborates the hypothesis that wall stress is mainly related to systolic BP and is a key determinant of LV hypertrophy development.³⁷ Concurrently, LV wall stress was demonstrated to be a main determinant of natriuretic peptides increase,³⁸ likely explaining the described correlation between mixed hypertension and NTproBNP. Although natriuretic peptides are not classically regarded as a marker of target organ damage, recent evidences showed that plasma NT-proBNP is a strong prognostic marker in hypertensive patients.³⁹ Systolic BP and arterial stiffness have been demonstrated to be closely related and may behave reciprocally as cause or effect, interacting in a vicious cycle. In this regard, PWV is considered a reliable measure to quantify arterial stiffness.⁴⁰ Our results confirm previous observations showing the association between systolic BP, but not diastolic BP, and PWV.⁴¹

In our analysis we found a trend toward the association between GLS change and IDH. GLS has been demonstrated as a sensitive tool to recognize early subclinical systolic dysfunction in newly diagnosed hypertensive patients without LVH,⁴² even when ejection fraction and other strain components are normal.⁴³ Interestingly, a recent echocardiographic study conducted on STANISLAS cohort identified a significant association between layer-specific strain variables and self-reported dyspnea, suggesting that refined strain components (i.e., subendocardial strain) could help identify early stages of increased LV filling pressure.⁴⁴

Altogether, our results appear to be consistent with previous observations showing a lack of association between IDH and target organ damage. These findings might also provide an explanation for the inconclusive association between IDH and adverse cardiovascular outcomes frequently reported in the scientific literature.

4.1 | Limitations

The main limitation of our study is its observational design; therefore, it is not possible to establish a causal link to the results obtained. In addition, given our sample size, we could not adjust our analysis for every possible cardiovascular risk variable. The cross-sectional design allowed us to assess only intermediate signs of target organ damage; therefore, our findings cannot be extrapolated to the incidence of hard cardiovascular or renal end-points. We had no assessment of microvascular damage besides microalbuminuria and therefore cannot rule out the impact of IDH on this target. The reclassification of our population according to ESC/ESH BP thresholds³ identified a very small number of IDH patients (N = 11), preventing any reliable sensitivity analysis in this setting (Figure S1). Different duration of hypertension among subgroups might underestimated the effect of BP on TOD in patients with a shorter follow-up, although it was not statistically significant. Finally, the results of this analysis cannot be extended to general hypertensive population, as they refer to an initially healthy population with a low cardiovascular risk at the time of evaluation and constituted by a mix of already diagnosed and newly-diagnosed hypertensives during STANISLAS research visit.

4.2 | Conclusions

In our study IDH was not significantly associated with target organ damage. This finding might suggest that the clinical significance of IDH in the absence of elevated systolic BP is questionable. Further studies are needed to clarify the causative role of IDH in the development of target organ damage and cardiovascular outcomes.

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CONFLICT OF INTEREST

The authors declare no conflicts of interest.

DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author upon reasonable request. All data will be made available upon request.

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SUPPORTING INFORMATION

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