Carotid Atherosclerosis and Electrocardiographic Left Ventricular Hypertrophy in the General Population: The Namwon Study

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This study aimed to investigate the relationship between carotid atherosclerosis and left ventricular hypertrophy on electrocardiogram (ECG-LVH) on adults living in the community. A total of 9,266 adults who participated in the Namwon Study were included in this analysis. Carotid atherosclerosis, including intima-media thickness (IMT) and plaques, were assessed using high-resolution B-mode ultrasound. ECG-LVH was determined using the Sokolow-Lyon voltage (SokV) and Cornell voltage (CorV) criteria. The prevalence of ECG-LVH was 12.7% using the SokV criteria and 9.7% using the CorV criteria. After full adjustment, compared to the lowest quartile of common carotid artery IMT (CCA-IMT), the odds ratios and 95% confidence intervals for ECG-LVH of the carotid IMT quartiles 2, 3, and 4 increased linearly as follows: 1.54 (1.24-1.90), 1.62 (1.31-2.02), and 1.91 (1.54-2.38), respectively, for the SokV criteria (p<0.001); and 1.33 (1.05-1.68), 1.41 (1.11-1.78), and 1.48 (1.16-1.88), respectively, for the CorV criteria (p=0.003). Positive associations between the carotid bulb IMT (CB-IMT) quartiles and the ECG-LVH were also observed, although the magnitudes of association between CB-IMT and ECG-LVH were slightly lower than those of CCA-IMT. However, no significant association between carotid plaques and ECG-LVH as defined by the SokV or CorV criteria was found. The present study demonstrated that increased carotid IMT, but not carotid plaques, is significantly associated with LVH defined by various ECG criteria in a large population.

Key Words: Hypertrophy, Left Ventricular; Carotid Intima-Media Thickness; Plaque, Atherosclerotic; Adult

INTRODUCTION

Cardiovascular disease (CVD) accounts for nearly one-third of all deaths worldwide, and coronary artery disease (CAD) is the main contributor to CVD.12 Although the total CVD mortality rate has decreased over a 30-year period in South Korea, the CAD mortality rate has continuously increased until recently.3 Left ventricular hypertrophy (LVH) is a strong predictor of CVD morbidity and mortality, including CAD, stroke, and congestive heart failure.4-6 Epidemiological studies have shown that LVH on the electrocardiogram (ECG-LVH) is a potent risk factor for future CVD events and death.7,8 Moreover, randomized clinical trials have shown that regression of ECG-LVH during medical treatment is associated with a decrease in the prevalence and incidence of CVD.9-11

Evaluation of carotid arteries with B-mode ultrasound is a reliable, cost-effective, and widely used method for the assessment of carotid atherosclerosis. Epidemiological studies have determined that carotid atherosclerotic findings, such as carotid intima-media thickness (IMT) and plaque, are a surrogate of subclinical atherosclerosis and a predictor of CVD.12-14 Moreover, it has recently been reported that carotid ultrasonographic indices are potent predictors of death and cardiovascular events in patients with CAD, suggesting that carotid plaque might have
greater prognostic significance than IMT.\textsuperscript{15,16}

Although carotid atherosclerosis and LVH are principal risk factors for CVD morbidity and mortality, little is known about the relationship between carotid atherosclerosis and ECG-LVH in the general population.\textsuperscript{17}

Therefore, to date, the relationship between carotid atherosclerosis and ECG-LVH remains undetermined. To gain additional insights into the potential role of carotid atherosclerosis in the development of LVH, it is important to examine the effect of carotid atherosclerotic phenotypes on ECG-LVH. Thus, in this large cross-sectional population-based study, we evaluated the association between carotid atherosclerosis and LVH (defined by commonly used electrocardiographic voltage criteria) in Korean adults.

MATERIALS AND METHODS

1. Study population

A total of 10,667 subjects aged 45-74 years were enrolled in the baseline survey of the Namwon Study between 2004 and 2007. The Namwon Study, whose cohort profile was published previously,\textsuperscript{18} is an ongoing population-based prospective study designed to examine the prevalence, incidence, and risk factors for chronic disease in rural areas of Korea. Of the total subjects, 444 patients with missing or poor carotid ultrasonography images, and 791 subjects with missing information on ECG, were excluded. Finally, a total of 9,286 participants were included in the analyses after further excluding 166 subjects whose records lacked information regarding anthropometric or laboratory measures. This study was conducted in accordance with the Declaration of Helsinki guidelines. All subjects were well-informed of the study contents and provided informed consent. The study protocol was independently approved by the Institutional Review Board of Chonnam National University Hospital (I-2007-07-062).

2. Carotid ultrasonography

A high-resolution B-mode ultrasound system (SONOACE 9900; Medison, Seoul, Korea) equipped with a 7.5-MHz linear array transducer was used to measure carotid IMT and to determine the presence of carotid plaque. IMT was defined as the distance from the intima-lumen interface to the media-adventitia interface on the far wall in a region free of plaque. Carotid plaque was defined as focal structures that encroached into the lumen by an area at least 100% thicker than the surrounding IMT value of the common carotid artery (CCA), carotid bulb (CB), or internal carotid artery (ICA). IMT of the CCA (CCA-IMT) was measured between the origin of the CB and a point 10 mm proximal to the CCA. IMT of the CB (CB-IMT) was measured in the CB region. In the present study, the CCA-IMT and CB-IMT values were determined as the average of the maximum IMT values of the right and left CCA and CB, respectively. Two sonographers performed carotid ultrasonographs. The correlation coefficients for inter-sonographer and intra-sonographer variability of the carotid IMT were 0.86 and 0.90, respectively. The kappa coefficients of the carotid plaques were 0.76 for inter-sonographer and 0.85 for intra-sonographer agreement. One trained reader analyzed still images obtained from carotid ultrasound. The correlation coefficient for intra-reader variability of carotid IMT was 0.94. The kappa coefficient of carotid plaques was 0.91 for intra-reader agreement.

3. Electrocardiographic measurement

Trained research technicians performed a 12-lead ECG examination using a 12-channel ECG device (PageWriter-200; Hewlett-Packard, MA, USA). Patients were able to freely breathe and did not talk during the procedure, and ECG recordings were performed in a comfortable supine position at a speed of 25 mm/sec and a voltage of 0.1 mV/mm. The heart rate (30 sec), pulse rate interval, QRS duration, QT interval, and corrected QT interval were obtained from ECG records. Research staff trained by cardiologists measured the R-wave amplitudes in leads aVL, V5, and V6, and the S-wave amplitudes in leads V1 and V3. LVH was determined as the voltage sum of SV 1+RV 5 or V 6 ≥35 mm using the Sokolow-Lyon voltage (SokV) criteria\textsuperscript{19} and as RaVL+SV 3 ≥28 mm for males and ≥20 mm for females using the Cornell voltage (CorV) criteria.\textsuperscript{20}

4. Measurements and assessments of covariates

The participants’ heights were measured to the nearest 0.1 cm; their weights were measured to the nearest 0.1 kg in light clothing without shoes. Body mass index (BMI) was calculated as weight divided by height squared (kg/m\textsuperscript{2}). After a 5-min rest in the sitting position, blood pressure was measured to the nearest 2 mmHg on the right upper arm using a standard mercury sphygmomanometer (Baumanometer; WA Baum Co., Inc., Copiague, NY, USA). Three consecutive measurements of systolic blood pressure (SBP) and diastolic blood pressure (DBP) were performed at 1-min intervals, and the average values were used in the analysis. Venous blood samples and urinary specimens were collected after a 12-h overnight fast. Total cholesterol (TC), high-density lipoprotein (HDL) cholesterol, triglycerides, and fasting blood glucose (FBG) were analyzed using an automatic analyzer with enzymatic methods (Model 7600 Chemical Analyzer; Hitachi Ltd., Tokyo, Japan). High-sensitivity C-reactive protein (hsCRP) was measured by latex-enhanced nephelometry using a high-sensitivity assay analyzer (Behring Nephelometer II; Dade-Behring Diagnostics, Marburg, Germany).

Information concerning cigarette smoking, alcohol consumption, and medication (hypertension, diabetes, and dyslipidemia) was obtained with a standardized questionnaire by trained research staff. Smoking status was classified as never, former, or current smoker. Alcohol consumption was calculated using the usual quantity of alcohol consumed and drinking frequency, and grouped into the following categories: none, 0-1.9 drinks/day, 2.0-3.9 drinks/day, and ≥4.0 drinks/day.
5. Statistical analysis

Data was presented as means ± standard deviation for continuous variables and frequencies (percentages) for categorical variables. Characteristics were compared between subjects with and without LVH using the chi-squared test and the independent t-test. Carotid IMT values were classified into quartiles for CCA-IMT (quartile 1: median value 0.594, interquartile range 0.549-0.609, range [min-max] 0.387-0.641 mm; quartile 2: median value 0.688, interquartile range 0.670-0.698, range [min-max], 0.642-0.732 mm; quartile 3: median value 0.775, interquartile range 0.733-0.779, range [min-max] 0.733-0.822 mm; quartile 4: median value 0.876, interquartile range 0.859-0.959, range [min-max] 0.823-1.839 mm) and for CB-IMT (quartile 1, median value 0.688, interquartile range 0.640-0.707, range [min-max] 0.420-0.733 mm; quartile 2: median value 0.785, interquartile range 0.777-0.823, range [min-max] 0.734-0.830 mm, quartile 3: median value 0.875, interquartile range 0.869-0.914, range [min-max] 0.831-0.932 mm; quartile 4: median value 1.01, interquartile range 0.966-1.10, range [min-max] 0.933-2.070 mm). The differences and linear trends in the proportions of ECG-LVH by carotid IMT quartile, and the presence of carotid plaque, were compared using the independent t-test. Furthermore, independent associations between carotid atherosclerosis and ECG-LVH were analyzed sequentially using multiple logistic regression. Model 1 was adjusted for age and gender and model 2 was further adjusted for smoking status, alcohol consumption, BMI, heart rate, and TC were lower. The proportion of patients taking anti-hypertensive and anti-diabetic medications was lower than in the control group.

2. LVH prevalence by carotid IMT and plaque

The prevalence of LVH defined by the SokV criteria increased with increasing CCA-IMT quartiles (7.0%, 11.6%, 13.9%, and 18.0%, respectively) (p < 0.001) and that of LVH defined by the CorV criteria showed a similar trend according to increasing quartiles (6.0%, 9.2%, 10.8%, and 12.4%, respectively) (p < 0.001). The prevalence of LVH defined by the SokV criteria increased by 8.1%, 12.0%, 14.4%, and 16.5% (p-trend < 0.001) for the first, second, third, and fourth CB-IMT quartiles, respectively, and that defined by the CorV criteria increased by 8.2%, 9.3%, 9.8%, and 11.3% (p < 0.001), respectively. According to the SokV criteria, the prevalence of LVH was 15.7% in subjects with carotid plaque and 11.3% in those without carotid plaque (p < 0.001). However, according to the CorV criteria, no significant difference was observed between subjects with carotid plaque and those without (p=0.658; Table 3).

3. Association between carotid IMT, plaque, and LVH

The associations between carotid atherosclerosis and ECG-LVH using logistic regression analysis are presented in Table 4. Using both the SokV and CorV criteria, the age- and gender-adjusted ORs for LVH increased as the CCA-IMT quartiles increased (p < 0.001). After multivariate adjustment, compared to the lowest CCA-IMT quartile, the ORs and 95% CIs for ECG-LVH carotid IMT quartiles increased linearly and were 1.54 (1.24-1.90) for using the CorV criteria. Significant differences in smoking status and alcohol consumption between the control and LVH groups were observed using both the SokV and CorV criteria.

Using the SokV criteria, SBP and DBP, HDL cholesterol, and hsCRP were significantly higher in subjects with LVH than in the control group, while BMI, waist circumference, heart rate, and TC were lower. The proportion of patients taking anti-diabetic and anti-dyslipidemic medications was lower than in the control group. Using the CorV criteria, BMI, SBP and DBP, FBG, TC, and hsCRP were significantly higher in subjects with LVH compared with control subjects. The proportion of patients taking anti-hypertensive and anti-diabetic medications was higher in subjects with LVH defined by the CorV criteria than in the control group.

### Table 1. Agreement of LVH using Kappa statistics

| Sokolow-Lyon criteria | LVH | Normal | Total |
|-----------------------|-----|--------|-------|
| LVH                   | 264 | 915    | 1,179 |
| Normal                | 632 | 7,455  | 8,087 |
| Total                 | 896 | 8,370  | 9,266 |

LVH: left ventricular hypertrophy.
**Table 2.** Baseline characteristics according to the presence of LVH

|                        | Sokolow-Lyon criteria | Cornell criteria |
|------------------------|------------------------|------------------|
|                        | Normal (n=8,087)       | LVH (n=1,179)    | Normal (n=8,370) | LVH (n=896) |
| Age, years             | 61.2±7.9              | 63.7±7.2         | 61.3±7.9         | 63.8±7.1    |
| Female, %              | 5,082 (62.8)          | 539 (45.7%)      | 4,851 (58.0)     | 770 (85.9)  |
| Body mass index, kg/m² | 24.5±3.1              | 23.5±2.8         | 24.3±3.0         | 24.7±3.1    |
| Waist circumference, cm| 86.5±8.4              | 83.4±7.9         | 86.1±8.3         | 86.2±9.1    |
| Heart rate, beats/50 sec| 35.6±5.5              | 34.7±5.3         | 35.5±5.4         | 35.9±6.0    |
| Systolic blood pressure, mmHg | 124.1±17.4          | 134.2±20.5       | 124.3±17.4       | 135.5±21.0  |
| Diastolic blood pressure, mmHg | 79.8±10.0           | 82.5±10.8        | 79.7±9.9         | 84.3±11.3   |
| Fasting blood glucose, mg/dl | 104.7±24.6          | 104.6±23.0       | 104.4±24.2       | 107.7±25.9  |
| Total cholesterol, mg/dl | 188.9±37.0           | 186.5±35.7       | 187.6±36.6       | 198.2±37.8  |
| HDL-cholesterol, mg/dl | 47.5±11.9            | 48.2±12.6        | 47.6±11.9        | 47.4±12.5   |
| hs-CRP, mg/dl*         | 0.071 (0.069-0.073)   | 0.078 (0.072-0.084) | 0.071 (0.069-0.073) | 0.080 (0.073-0.087) |
| Anti-hypertensive medications | 1,714 (21.2)        | 314 (26.6)       | 1,726 (20.6)     | 302 (33.7)  |
| Anti-diabetic medications | 627 (7.8)           | 67 (5.7)         | 602 (7.2)        | 92 (10.3)   |
| Anti-dyslipidemic medications | 424 (5.2)        | 44 (3.7)         | 430 (5.1)        | 38 (4.2)    |
| Smoking status         | 5,438 (67.2)         | 643 (54.5)       | 5,323 (63.6)     | 758 (84.6)  |
| Alcohol consumption, drinks/day | 4,453 (55.1)    | 540 (45.8)       | 4,439 (53.0)     | 554 (61.8)  |
| Never                  | 1,476 (18.3)         | 301 (25.5)       | 1,718 (20.5)     | 59 (6.6)    |
| Former                 | 1,173 (14.5)         | 235 (19.9)       | 1,329 (15.9)     | 79 (8.8)    |
| Current                | 417 (5.2)            | 102 (8.7)        | 494 (5.9)        | 25 (2.8)    |
| ≥ 4.0                  | 504 (6.2)            | 125 (10.6)       | 606 (7.2)        | 23 (2.6)    |

Values are expressed as means ± standard deviation or geometric means (95% confidence intervals)* for continuous variables, and frequency (percentage) for categorical variables. LVH, left ventricular hypertrophy; HDL, high-density lipoprotein; hs-CRP, high sensitivity C-reactive protein.

* p-values were obtained using analysis of variance for continuous variables and the chi-square test for categorical variables.
TABLE 3. Prevalence of LVH assessed with carotid IMT and plaque

|                | Sokolow-Lyon criteria | Cornell criteria |
|----------------|-----------------------|-----------------|
|                | Normal               | LVH             | p-difference (p-trend) | Normal | LVH             | p-difference (p-trend) |
| CCA-IMT*       |                       |                 |                         |        |                 |                         |
| Quartile 1 (n=2,216) | 2,061 (93.0)         | 155 (7.0)       | <0.001 (<0.001)        | 2,082 (94.0) | 134 (6.0)       | <0.001 (<0.001)        |
| Quartile 2 (n=2,360) | 2,086 (88.4)         | 274 (11.6)      |                         | 2,144 (90.8) | 216 (9.2)       |                         |
| Quartile 3 (n=2,289) | 1,970 (86.1)         | 319 (13.9)      |                         | 2,041 (89.2) | 248 (10.8)      |                         |
| Quartile 4 (n=2,401) | 1,970 (82.0)         | 431 (18.0)      |                         | 2,103 (87.6) | 298 (12.4)      |                         |
| CB-IMT†,‡      |                       |                 |                         |        |                 |                         |
| Quartile 1 (n=2,389) | 2,196 (91.9)         | 193 (8.1)       | <0.001 (<0.001)        | 2,192 (91.8) | 197 (8.2)       |                         |
| Quartile 2 (n=2,216) | 1,950 (88.0)         | 266 (12.0)      |                         | 2,009 (90.7) | 207 (9.3)       |                         |
| Quartile 3 (n=2,337) | 2,001 (85.6)         | 336 (14.4)      |                         | 2,107 (90.2) | 230 (9.8)       |                         |
| Quartile 4 (n=2,313) | 1,931 (83.5)         | 382 (16.5)      |                         | 2,052 (88.7) | 261 (11.3)      |                         |
| Carotid plaque  |                       |                 |                         |        |                 |                         |
| Absent (n=6,307) | 5,592 (88.7)         | 715 (11.3)      | <0.001                  | 5,703 (90.4) | 604 (9.6)       |                         |
| Present (n=2,959) | 2,495 (84.3)         | 464 (15.7)      |                         | 2,667 (90.1) | 292 (9.9)       |                         |

Values are expressed as number (percentile).
LVH, left ventricular hypertrophy; CCA, common carotid artery; IMT, intima-media thickness; CB, carotid bulb.

*Quartile 1: median value 0.594, interquartile range 0.549-0.609, range (min-max) 0.387-0.641 mm; quartile 2: median value 0.688, interquartile range 0.670-0.698, range (min-max) 0.642-0.732 mm, quartile 3: median value 0.775, interquartile range 0.733-0.779, range (min-max) 0.733-0.822 mm; quartile 4: median value 0.876, interquartile range 0.859-0.959, range (min-max) 0.823-1.839 mm.
†Quartile 1, median value 0.688, interquartile range 0.640-0.707, range (min-max) 0.420-0.733 mm; quartile 2: median value 0.785, interquartile range 0.777-0.823, range (min-max) 0.734-0.830 mm, quartile 3: median value 0.875, interquartile range 0.869-0.914, range (min-max) 0.831-0.932 mm; quartile 4: median value 1.01, interquartile range 0.966-1.10, range (min-max) 0.933-2.070 mm.
‡Eleven subjects with missing CB-IMT information were excluded.

TABLE 4. Associations between carotid atherosclerosis and LVH using multiple logistic regression

|                | LVH defined using Sokolow-Lyon criteria | LVH defined using Cornell criteria |
|----------------|-----------------------------------------|----------------------------------|
|                | Age- and gender-adjusted | Multivariate adjusted* | Age- and gender-adjusted | Multivariate adjusted* |
| CCA-IMT*       |                         |                                |                         |                         |
| Quartile 1     | 1.00 (ref.)              | 1.00 (ref.)                    | 1.00 (ref.)              | 1.00 (ref.)              |
| Quartile 2     | 1.53 (1.24-1.89)         | 1.54 (1.24-1.90)               | 1.45 (1.15-1.82)         | 1.33 (1.05-1.68)         |
| Quartile 3     | 1.72 (1.40-2.12)         | 1.62 (1.31-2.02)               | 1.68 (1.33-2.11)         | 1.41 (1.11-1.78)         |
| Quartile 4     | 2.12 (1.72-2.61)         | 1.91 (1.54-2.38)               | 2.02 (1.60-2.54)         | 1.48 (1.16-1.88)         |
| p-trend        | <0.001                   | <0.001                         | <0.001                   | 0.003                    |
| CB-IMT†        |                         |                                |                         |                         |
| Quartile 1     | 1.00 (ref.)              | 1.00 (ref.)                    | 1.00 (ref.)              | 1.00 (ref.)              |
| Quartile 2     | 1.33 (1.09-1.63)         | 1.25 (1.02-1.53)               | 1.10 (0.89-1.36)         | 1.06 (0.85-1.31)         |
| Quartile 3     | 1.45 (1.19-1.76)         | 1.32 (1.08-1.61)               | 1.22 (0.99-1.51)         | 1.14 (0.92-1.41)         |
| Quartile 4     | 1.57 (1.29-1.91)         | 1.41 (1.15-1.72)               | 1.45 (1.17-1.78)         | 1.29 (1.04-1.60)         |
| p-trend        | <0.001                   | 0.002                          | <0.001                   | 0.016                    |
| Carotid plaque |                         |                                |                         |                         |
| Absent         | 1.00 (ref.)              | 1.00 (ref.)                    | 1.00 (ref.)              | 1.00 (ref.)              |
| Present        | 1.10 (0.96-1.25)         | 1.01 (0.87-1.16)               | 1.06 (0.91-1.24)         | 0.92 (0.78-1.08)         |

Values are expressed as odds ratios (95% confidence intervals).
LVH, left ventricular hypertrophy; CCA, common carotid artery; IMT, intima-media thickness; CB, carotid bulb.

*Adjusted for age, gender, smoking status, alcohol consumption, body mass index, heart rate, systolic blood pressure, fasting blood glucose, total cholesterol, high-density lipoprotein cholesterol, high-sensitivity C-reactive protein (log), anti-hypertensive medication, anti-diabetic medication, and anti-dyslipidemic medication.

the second quartile, 1.62 (1.31-2.02) for the third quartile, and 1.91 (1.54-2.38) for the fourth quartile using the SokLV criteria (p<0.001) and 1.33 (1.05-1.68) for the second quarteR, 1.41 (1.11-1.78) for the third quartile, and 1.48 (1.16-1.88) for the fourth quartile using the Cornell criteria (p=0.003). In addition, the age- and gender-adjusted ORs
for LVH increased as the CB-IMT quartiles increased (p < 0.001) using both the SokV and CorV criteria. After full adjustment, compared to the lowest CB-IMT quartile, the ORs and 95% CI for LVH increased for both the SokV (1.25 (1.02-1.53), 1.32 (1.08-1.61), and 1.41 (1.15-1.72) for the second, third, and fourth quartiles, respectively; p < 0.001) and the CorV (1.06 (0.85-1.31), 1.14 (0.92-1.41), and 1.29 (1.04-1.60) for the second, third, and fourth quartiles, respectively; p=0.017) criteria. However, no significant association between carotid plaque and LVH (defined by the SokV or CorV criteria) was found in a logistic regression analysis.

**DISCUSSION**

We explored the cross-sectional associations between carotid atherosclerosis and ECG-LVH in a community-based Korean population aged 45-74 years. Carotid IMT, including CCA-IMT and CB-IMT, was independently and linearly associated with ECG-LVH, while this association was more strongly present in ECG-LVH defined by the SokV criteria than by the CorV criteria. However, no significant association between carotid plaque and ECG-LVH was determined.

Although echocardiography is the preferred method used to diagnose LVH in clinical practice, ECG is commonly used as the first-line instrument for detecting LVH due to its convenience, good reproducibility, cost-effectiveness, and availability in large cohort settings. However, ECG-LVH may be a better predictor of cardiovascular complications than LVH detected by echocardiography (Echo-LVH). Both ECG and echocardiography should be performed as they predict total and cardiovascular mortality independently of each other; ECG-LVH provides independent prognostic information on cardiovascular risk even after adjusting for left ventricular mass. Recent studies have shown that both ECG-LVH and Echo-LVH are independent risk factors for various CVDs. ECG-LVH and Echo-LVH can be used alternately for the prediction of stroke or congestive heart failure because, both have been shown to be similarly associated with increased risk. Therefore, ECG-LVH, independent of Echo-LVH, is regarded as an electrophysiological marker for future CVD.

The present study showed that among carotid atherosclerotic indices, IMT, but not plaque, is significantly associated with ECG-LVH. It is not clear why this is so. One possible explanation is that increased IMT and atherosclerotic plaque formation reflect different stages, and distinct pathological aspects, of atherosclerosis, although they have been used as a surrogate of CVD. Carotid IMT thickening, which is associated with hypertrophy of the intima and media as a compensatory non-atherosclerotic alteration, represents an early stage of atherosclerosis. Plaque formation, which is more prevalent in CB or ICA than CCA, is a pathology of the intima and reflects a more advanced phase of atherosclerotic disease, caused by oxidation, inflammation, endothelial dysfunction, and smooth muscle cell proliferation. Epidemiological studies revealed that carotid IMT thickening is more closely associated with hypertension and ischemic stroke, while carotid plaque is more strongly associated with hyperlipidemia and CAD.

Variability in subjects and devices, as well as differences in IMT measurements, plaque definition, and methods of determining LVH (ECG or Echo) between previous studies and our study should be considered. Epidemiological studies have shown that carotid atherosclerosis was significantly associated with Echo-LVH. The study by Cuspidi et al. in untreated and uncomplicated essential hypertension found a significant association between carotid thickening and Echo-LVH defined as an increase in left ventricular mass. Cardiac hypertrophy defined as Echo-LVH was significantly associated with the presence of carotid plaque. Studies have shown that carotid atherosclerosis was significantly associated with ECG-LVH in 349 asymptomatic subjects. However, the abovementioned studies included only small sample sizes and hypertensive subjects, used plaque as a marker and not IMT, or used Echo-LVH. Our results revealed that carotid IMT thickening was positively associated with ECG-LVH in a large population, while carotid plaque was not.

In our study investigating LVH determined by both the SokV and CorV criteria, the strength of the association between carotid IMT and ECG-LVH was greater when using the CCA region for the IMT measurement compared with the CB region. A possible explanation for this is as follows. When plaque was present in the IMT measurement region of CCA and CB in our study, we measured IMT in a region free of plaque. Plaque was more prevalent in the CB and the origin of the ICA than in the CCA. Although ultrasound identifies later stages of atherosclerosis such as plaque, which may be identified either in the absence of, or in conjunction with, increased IMT, it cannot clearly identify the intermediate stages between increasing IMT and plaque formation. Thus, particularly in the CB region, IMT and plaque may not have been definitively separated in the present study. Compared with the other regions used for IMT measurements, the segment that is most useful for assessment, and which produces the most reproducible results for IMT measurement, is the CCA. Although there are no superior carotid artery segments clearly associated with CVD, CCA-IMT is associated with a higher relative risk for stroke, whereas ICA-IMT is associated with a higher relative risk for CHD.

The present study had certain limitations. First, our cross-sectional study design did not allow us to infer a causal relationship between carotid atherosclerosis and ECG-LVH. In addition, this study used the presence of carotid plaque as an outcome variable. However, more precise analyses using Doppler flow quantification to assess the degree of plaque stenosis are needed because carotid Doppler measurements, in addition to IMT and plaque evaluation, are independently associated with future...
Third, because this study measured only ECG-LVH instead of Echo-LVH, there may be a distortion in the results of this study that suggested a relationship between carotid atherosclerosis and LVH. Despite these limitations, our work is valuable as it provides insight into the potential role of carotid atherosclerosis in the development of LVH, as determined by ECG in large populations.

In conclusion, our study demonstrated that high carotid IMT, but not carotid plaque, was significantly associated with LVH as assessed by ECG in the general Korean population. These associations were more powerful in the CCA than in CB. Moreover, carotid IMT measured at both the CCA and CB was more strongly associated with ECG-LVH defined using the SokV criteria than using the CorV criteria. Further longitudinal, population-based studies are warranted to explore the relationship between carotid atherosclerosis and ECG-LVH. In addition, further research designs are needed to evaluate the clinical impact of carotid atherosclerosis on ECG-LVH.

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CONFLICT OF INTEREST STATEMENT

None declared.

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