The Combined Effect of Ear Lobe Crease and Conventional Risk Factor in the Diagnosis of Angiographically Diagnosed Coronary Artery Disease and the Short-Term Prognosis in Patients Who Underwent Coronary Stents

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Abstract: The role of diagonal ear lobe crease (DELC) in coronary artery disease (CAD) diagnosis and prognosis remains controversial. In this study, we aimed to assess the combined effect of DELC with other conventional risk factors in the diagnosis and prognosis of CAD in Chinese patients who underwent angiography and coronary stent implantation.

The study consisted of 956 consecutive patients who underwent angiography. The DELC was identified as no DELC, unilateral, and bilateral DELC. The conventional risk factors for CAD were recorded.

Our data showed that the overall presence of DELC is associated with CAD risk. Stratification analyses revealed that the diagnostic value of DELC was mostly significant in those with >4 risk factors. Also in patients with >4 risk factors, the presence of bilateral DELC remains to be associated with higher hs-CRP level, higher severity of CAD, and higher possibility of developing major adverse cardiac events after successful percutaneous coronary intervention (PCI).

Our study confirmed the relation of DELC with CAD in Chinese patients; more importantly, our data suggest the combination of DELC and CAD risk factors will help to predict the incidence of CAD and may predict the prognosis after successfully PCI.

Abbreviations: CAD = coronary artery disease, CVD = cardiovascular disease, DELC = diagonal ear lobe crease, DM = diabetes mellitus, HDL-C = high-low-density lipoprotein cholesterol, LDL-C = low-density lipoprotein cholesterol, MACE = Major adverse cardiac events, OR = Odd ration, PCI = percutaneous coronary intervention.
bypass cardiac surgery, and development of congestive heart failure (MACEs) were considered as end points, including death, nonfatal myocardial infarction, or repeat revascularization. The outcome was evaluated at the end of 12th month. Major adverse cardiovascular events (MACEs) were recorded. The number of risk factor was recorded for each patient.

Traditional risk factors for CAD, including arterial hypertension (blood pressure \( \geq 140/90 \text{mmHg} \) or antihypertensive therapy), serum total cholesterol, serum high- low-density lipoprotein cholesterol (HDL-C), low-density lipoprotein cholesterol (LDL-C), triglycerides, creatinine, current or prior smokers, diabetes mellitus (DM), and a family history of CVD were recorded. The number of risk factor was recorded for each patient.

PCI Procedure and Follow-Up

PCI was performed according to the American Heart Association/American College of Cardiology guidelines.\(^{15}\) Antegrade perfusion was graded by Thrombolysis In Myocardial Infarction\(^ {16}\) criteria. All patients received drug-eluting stent. After the procedures, all patients were on dual antiplatelet therapy with aspirin and clopidogrel for one year. Other treatment including statins, angiotensin-converting enzyme inhibitor, and B-blocker were given to patients. The clinical outcome was evaluated at the end of 12th month. Major adverse cardiac events (MACEs) were considered as end points, including cardiac death, acute myocardial infarction (AMI), new PCI, bypass cardiac surgery, and development of congestive heart failure.\(^ {17}\)

Statistical Analysis

Data on quantitative characteristics are expressed as means ± SD. Data on qualitative characteristics are expressed as percent values or absolute numbers, as indicated. Differences in demographic characteristics and vascular risk factors between patients and controls were compared by using Student t test or analysis of variance for continuous variables and the \( \chi^2 \) test for all categorical variables. The DELC frequencies in subjects with and without CAD were compared by \( \chi^2 \) analysis or Fisher exact test. Multivariate logistic regression analysis was used to determine the influence of DELC on CAD risk, controlling potential confounding conventional risk factors. A forward stepwise (Likelihood Ratio) procedure was used for multivariable analysis. Data were analyzed with the SAS9.3 (SAS Inc, Cary, NC). The results were considered statistically significant at \( P < 0.01 \) using a 2-tailed test.

RESULTS

Based on the results of angiography, 446 subjects were assigned to the CAD group, and 510 cases were assigned to non-CAD group. Demographic characteristics of CAD groups and non-CAD groups are presented in Table 1. The CAD group had older age, higher systolic blood pressure level, higher BMI, LDL, CRP level, had higher percentage of male, DM, smoker, and CVD family history than non-CAD group (All \( P < 0.05 \)).

Table 2 shows the comparison of overall DELC presence between CAD and non-CAD groups. Our data showed there is a significant difference in the overall DELC appearance between the 2 groups. The CAD group had markedly increased frequencies of bilateral DELC compared with non-CAD group (39% vs 27%, \( P < 0.001 \)). To determine whether DELC is associated with the CAD risk, we performed the logistic regression analyses. Our results showed that the presence of bilateral DELC represented a significantly higher risk for CAD with adjustment with age, sex, BMI, blood pressure, LDL, smoking status, DM, and family history of CVD (adjusted odds ratio [OR] = 1.917, compared with no DELC subgroup, adjusted \( P < 0.001 \)). However, the unilateral presence of DELC was not associated with CAD risk (\( P = 0.183 \)).

To analyze the interaction between DELC and conventional risk factors for CAD, we next stratified the enrolled subjects by the numbers of risk factors each patient carried. Table 3 shows the frequencies of DELC in each stratification subgroups. In those with \( <3 \) risk factors, the DELC frequencies were similar between CAD and non-CAD groups (\( P = 0.303 \)). In patients with 3 to 4 risk factors, a significant difference in DELC frequencies was noted (\( P = 0.039 \)). In this subgroup,
those with bilateral DELC had 1.882 times higher chance to develop CAD (95% confidence interval [CI]: 1.089–3.253, \(P = 0.023\)). A significant difference in DELC frequencies was noted more evident in those with >4 risk factors \((P < 0.001)\). The presence of bilateral DELC in this subgroup represented 4.047 times higher chance to have CAD (95% CI: 2.267–7.226, \(P < 0.001\)). Unilateral DELC was not associated with the CAD risk in all subgroups.

**TABLE 2.** The Comparison of Overall DELC Presence Between CAD and Non-CAD Groups

|                  | CAD+  | Non-CAD  | Adjusted OR | 95% CI  | Adjusted \(P\) |
|------------------|-------|----------|-------------|---------|----------------|
| No DELC          | 107   | 0.24     | 165         | 0.32    |                |
| Unilateral DELC  | 165   | 37%      | 205         | 40%     | 1.241          | 0.903          | 1.706          | 0.183          |
| Bilateral DELC   | 174   | 39%      | 140         | 27%     | 1.917          | 1.378          | 2.665          | <0.001         |

\(CAD = \) coronary artery disease, \(CI = \) confidence interval, \(DELC = \) diagonal ear lobe crease, \(OR = \) odds ratio.

**FIGURE 1.** The association of diagonal ear lobe crease (DELC) with the coronary artery disease (CAD) severity. (A) The association of DELC with the CAD severity determined by the number of diseased vessels. (B) The association of DELC with the CAD severity determined by Gensini scores. Those patients with bilateral DELC had significantly higher severity of CAD compared with those without DELC and with unilateral DELC.
We next analyzed the association of DELC with the severity of CAD by the number of diseased vessels and by the Gensini scores. We found that the overall DELC appearance did not affect the number of disease vessels (Figure 1A) and Gensini scores (Figure 1B). There was no significant difference in the disease severity among patients with <3 or 3 to 4 risk factors. Only among the patients with >4 risk factors, there is a significant increase in the diseased vessel numbers (Figure 1A-D) and Gensini scores (Figure 1B-D) among patients with bilateral DELC, compared with those without DELC and those with unilateral DELC. The bilateral DELC carriers had significantly higher diseased number than those without or with only unilateral DELC (2.3 ± 0.2 vs 1.7 ± 0.1, P = 0.013; vs 1.8 ± 0.1, P = 0.021 Figure 1D). Likewise, patients with bilateral DELC had markedly higher Gensini scores (78.3 ± 10.6) than those without DELC (47.3 ± 11.2, P = 0.006) and those with unilateral DELC (58.1 ± 8.6, P = 0.010).

The associations of DELC presence with blood pressure, serum HDL-C, LDL-C level, and CRP at diagnosis were also studied. We found that the mean levels of these variables were similar among patients without DELC, unilateral, and bilateral DELC, even after stratification by the number of risk factors (data not shown). As for the serum CRP levels, the mean CRP level was higher in those with bilateral DELC compared with those without DELC, but did not reach significant difference (P = 0.052, Figure 2A). After stratification with the number of risk factors, we observed that patients with bilateral DELC from subgroup with >4 risk factors had a significantly higher mean CRP level (2.45 [2.08–2.51] mg/L) than those without (1.65 [1.45–2.05] mg/L, P = 0.014) or only with unilateral DELC (1.92 [1.64–2.2] mg/L, P = 0.016) (Figure 2D). In patient with <3 and with 3 to 4 risks, the CRP levels were not noted significantly different (both P > 0.05, Figure 2B and C).

Of all CAD patients, there were 276 patients who underwent successful PCI and were follow-up for 12 month. Of all patients, 17 had MACE during follow-up period (6.15%). The overall DELC frequencies between patients with MACE and those without MACE were not significantly different (P = 0.155, Table 4). Logistic regression analyses with conventional risk factors did not reveal any association of DELC, both unilateral and bilateral, with MACE incidence. However, when we stratified these patients according to the number of risk factors each one carried, we found that in only patients with >4 risk factors, the presence of bilateral DELC represented a significantly increased risk for MACE incidence (OR = 5.568, P = 0.028, Table 5).
To exclude that the worse prognosis after PCI in the presence of DELC is due to a worse atherosclerotic coronary artery disease, we stratified all patients based on their Gensini scores (<30, 31–59, and >60). Similarly, we found that the DELC presence was not associated with the MACE in patients with Gensini scores <30 and between 31 and 59; however, in those with Gensini scores >60, the bilateral DELC had dramatically increased chance to develop MACE (OR = 6.002, P = 0.019, Table 6).

**DISCUSSION**

In this study, we emphasized the interaction between DELC presence and the conventional risk factors in the diagnosis of angiographically determined CAD and the clinical outcome in patients who underwent successful stent procedures. Our data showed that the overall presence of DELC is associated with CAD incidence. Stratification analyses revealed that the diagnostic value of DELC was mostly

**TABLE 3. The Interaction Between DELC and the Number of Conventional Risk Factors for CAD**

| DELC Status | CAD N | % | Non-CAD N | % | OR | 95% CI | P  |
|-------------|-------|---|-----------|---|----|--------|----|
| No DELC     | 23    | 21%| 40        | 24%| 1  |        |    |
| Unilateral DELC | 53  | 48%| 64        | 39%| 1.44 | 0.768  | 2.701 | 0.255 |
| Bilateral DELC | 34  | 31%| 61        | 37%| 0.969| 0.5    | 1.88  | 0.927 |
| Patients with <3 risk factors |
| No DELC     | 45    | 31%| 69        | 38%| 1  |        |    |
| Unilateral DELC | 45  | 31%| 67        | 37%| 1.03 | 0.605  | 1.754 | 0.914 |
| Bilateral DELC | 54  | 38%| 44        | 24%| 1.882| 1.089  | 3.253 | 0.023 |
| Patients with 3–4 risk factors |
| No DELC     | 34    | 18%| 56        | 34%| 1  |        |    |
| Unilateral DELC | 67  | 36%| 74        | 45%| 1.491| 0.87   | 2.557 | 0.146 |
| Bilateral DELC | 86  | 46%| 35        | 21%| 4.047| 2.267  | 7.226 | <0.001 |
| Patients >4 risk factors |

CAD = coronary artery disease, CI = confidence interval, DELC = diagonal ear lobe crease, OR = odds ratio.

**TABLE 4. The Overall DELC Frequencies Between Patients With and Without MACE**

|         | With MACE | %  | Without MACE | %  | OR   | 95% CI | P   |
|---------|-----------|----|--------------|----|------|--------|-----|
| No DELC | 4         | 0.21| 98           | 0.38| 1    |        |     |
| Unilateral DELC | 5  | 26%| 94           | 36%| 1.303| 0.34   | 5.001| 0.699 |
| Bilateral DELC | 8  | 42%| 67           | 26%| 2.925| 0.847  | 10.107| 0.078 |

CAD = coronary artery disease, CI = confidence interval, DELC = diagonal ear lobe crease, OR = odds ratio.

**TABLE 5. The DELC Frequencies Between Patients With and Without MACE Stratified by the Number of Risk Factors**

| DELC Status | With MACE | %  | Without MACE | %  | Adjusted OR | 95% CI | Adjusted P |
|-------------|-----------|----|--------------|----|-------------|--------|------------|
| No DELC     | 0         | 0% | 25           | 35%| —           |        | —         |
| Unilateral DELC | 0  | 0%| 27           | 38%| —           |        | —         |
| Bilateral DELC | 0  | 0%| 19           | 27%| —           |        | —         |
| Patients with 3–4 risk factors |
| No DELC     | 2         | 40%| 38           | 39%| 1           |        | 0.149     | 8.374     | 0.914     |
| Unilateral DELC | 2  | 40%| 34           | 35%| 1.118       | 0.731  | 0.063     | 8.484     | 0.801     |
| Bilateral DELC | 1  | 20%| 26           | 27%| 0.731       | 0.594  | 0.058     | 8.484     | 0.801     |
| Patients with >4 risk factors |
| No DELC     | 2         | 14%| 35           | 90%| 1           |        | 0.252     | 10.132    | 0.623     |
| Unilateral DELC | 3  | 21%| 33           | 85%| 1.591       | 1.059  | 0.058     | 9.273     | 0.028     |
| Bilateral DELC | 7  | 48%| 22           | 57%| 5.568       | 4.047  | 0.058     | 29.273    | 0.028     |

CI = confidence interval, DELC = diagonal ear lobe crease, OR = odds ratio.
significant in those with >4 risk factors. DELC presence alone did not predict the MACE incidence after PCI; however, in patients with >4 risk factors, the presence of bilateral DELC remains to be associated with higher chance of developing MACE after successful PCI. This result confirmed a positive association between DELC and CAD risk, in addition, suggesting that the combination of DELC with risk factor each individual carriers will increase its diagnostic value in predicting the CAD and its outcome after PCI.

The association between DELC and ELC has been reported in several independent studies. DELC (both unilateral or bilateral) was significantly more common among American patients with an AMI than among age-matched control subjects.\(^\text{19}\) In Indian population, bilateral DELC was found to be significantly associated with angiographically documented CAD.\(^\text{19}\) In Irish patients, the association between an DELC and CAD was found to be significant, suggesting that it should be used as a physical sign predictive of the presence of coronary heart disease.\(^\text{20}\) DELC was reported more prevalent in male patients than female patients and the incidence of bilateral DELC remains to be associated with higher chance of developing MACE after successful PCI.

Another novelty of this study is that we investigated the role of DELC in the clinical outcome of patients who underwent successful coronary artery drug-eluting stents implantation. An autopsy study showed that ELC was one of the strongest independent risk factor for sudden cardiac death.\(^\text{24}\) Another follow-up study showed that a diagonal ELC is associated with increased all-cause and cardiac morbidity and mortality. Patients with ELC may be at higher risk for coronary events, even if currently without diagnostic evidence of CAD.\(^\text{23}\) Currently, there is no studying reporting the diagnostic role of DELC in patients underwent PCI. In this study, we found that in patients with >4 risk factors, the presence of bilateral DELC remains to be associated with higher chance of developing MACE after successful PCI.

### CONCLUSION

Taken together, our study confirmed the relation of DELC with CAD in Chinese patients; moreover, our data show that the value of DELC is more reliable in those with multiple risk factors, suggesting the importance of combination of DELC and CAD risk factors.

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