

ORIGINAL RESEARCH

U-Shaped Association Between Monocyte-Lymphocyte Ratio and Risk of Cardiac Conduction Block

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Purpose: Inflammation plays a critical role in the development of cardiac conduction block (CCB), which is associated with an increased risk of morbidity and mortality. The monocyte-lymphocyte ratio (MLR) acts as a novel inflammatory marker; however, its association with CCB has not yet been studied. This study aimed to investigate the association between MLR and CCB risk.

Patients and Methods: In total, 82,472 CCB-free participants were identified from the Kailuan study. MLR was calculated using the monocyte count/lymphocyte count. The participants were stratified based on quartiles of MLR levels. Incident CCB and its subtypes were ascertained from electrocardiograms at biennial follow-up visits. The Cox proportional hazards model and restricted cubic spline analysis were used to investigate the association between MLR with CCB and its subtypes.

Results: During a median follow-up of 10.4 years, 3222 incident CCB cases were observed. A U-shaped association was observed between MLR and CCB risk ($P_{\text{nonlinearity}} < 0.05$). After multivariate adjustment, individuals in the highest MLR quartile had a hazard ratio (HR) of 1.212 (95% CI: 1.097-1.340; Q4 vs Q2), while those in the lowest MLR quartile had an HR of 1.106 (95% CI: 1.000-1.224; Q1 vs Q2). Sensitivity and subgroup analyses yielded consistent results. The U-shaped association persisted for atrioventricular block (AVB) in subtype analyses.

Conclusion: MLR was significantly associated with an increased risk of new-onset CCB. Assessing MLR may have clinical relevance for predicting CCB risk, providing valuable insights for preventive strategies and patient management.

Pre-Registered Clinical Trial Number: The pre-registered clinical trial number is ChiCTR-TNC-11001489.

Keywords: cardiac arrhythmia, cardiac conduction block, inflammation, monocyte-lymphocyte ratio, risk factors

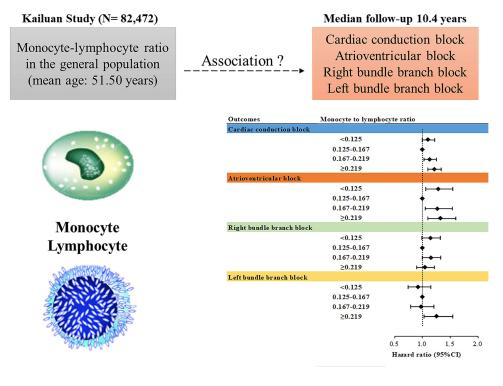
Introduction

Cardiac conduction block (CCB) is characterised by a disease manifested as the delay or interruption in the transmission of an impulse within the conduction system, including the sinoatrial, atrioventricular block (AVB), and bundle branch block (BBB). Regardless of the type of conduction block, disease progression in severe cases significantly increases the risk of cardiovascular morbidity and mortality. Although pacemaker implantation is an effective treatment for end-stage conduction block diseases, it carries the risk of infection and thrombosis and is associated with long-term adverse cardiac remodelling. Therefore, it is particularly important to identify high-risk individuals with conduction block and implement early measures to avoid illness onset.

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Graphical Abstract



The monocyte-lymphocyte ratio (MLR) is a novel inflammatory biomarker that can effectively reflect the balance between inherent and acquired immune responses.^{8,9} Previous epidemiological evidence suggested that MLR was an independent predictor of all-cause mortality and cardiovascular mortality in the general population.¹⁰ Studies have shown that inflammation plays an important role in the initiation of CCB.^{11,12} However, data on the association of inflammatory biomarkers with CCB are relatively limited. A longitudinal cohort study found that lymphocyte count (LC) showed a U-shaped association with the risk of incident arrhythmias.¹³ Compared with LC or monocyte count (MC) alone, MLR provides a more stable and accurate evaluation of immune status.^{9,14} Therefore, we conducted a prospective cohort study to investigate the relationship between MLR and new-onset CCB and its subtypes.

Methods

Study Design and Population

The Kailuan study is an ongoing prospective cohort study that based on a community in Tangshan. The details of this study have been described elsewhere. ^{15,16} In current study, we first enrolled 101,510 individuals (81,110 men and 20,400 women, aged 18–98 years) who participated baseline examination (2006–2007). After excluding participants with missing data pertaining to the MC, LC, or electrocardiogram (ECG) (n=8274), with a history of CCB, myocardial infarction, atrial fibrillation, or heart failure (n=3974), and who did not attend any of the follow-up examinations (n=6790), a total of 82,472 participants (65,164 men and 17,308 women) were included in the current analysis (Supplementary Figure 1). These participants complete face-to-face questionnaire surveys, clinical examinations, laboratory tests and electrocardiograph (ECG) measurement during follow-up period.

This study complied with the Helsinki Declaration and was approved by the ethics committee of Kailuan General Hospital. All participants provided written informed consent prior to inclusion in the study.

Assessment of Incident CCB

CCB diagnoses were based on standard 12-lead ECGs at biennial follow-up visit. According to the relevant US AHA guidelines, CCB was defined as the presence of any type of heart block, including atrioventricular block (AVB), complete right bundle branch block (CRBBB), incomplete right bundle branch block (iRBBB), complete left bundle branch block (CLBBB), incomplete left bundle branch block (iLBBB), left anterior fascicular block (LAFB), left posterior fascicular block (LPFB), and nonspecific intraventricular conduction block (NS-IVCB). The right bundle branch block (RBBB) included the CRBBB and iRBBB, and the left bundle branch block (LBBB) included the CLBBB, iLBBB, LAFB, and LPFB. Detailed definitions of the endpoints are listed in Supplementary Table 1. The final diagnosis was confirmed by two cardiologists.

All participants were followed up through assessments updated every two years from the baseline examination until the onset of CCB, the date of all-cause mortality, or the last follow-up, whichever came first. The participants were monitored until December 31, 2019, for all outcomes.

Assessment of MLR

During each examination, the participants underwent venous blood collection after fasting for at least 8 hours. Routine blood tests, including MC and LC measurements, were performed using a full blood count analyzer (Sysmex XT-1800i, Sysmex Corporation). The MLR was calculated as MC/LC.

Data Collection and Definitions

Demographic data and lifestyle information, including alcohol intake, smoking status, physical activity, snoring, and medical history, were obtained through face-to-face interviews using a standardised questionnaire, as described in further detail previously. Anthropometric parameters and blood pressure were measured during biennial physical examinations. The detailed measurement procedures have been described previously. Body mass index (BMI) was determined by dividing the body weight (kg) by the square of the height (m²). During the physical examination, blood samples were collected from the antecubital vein following a fasting period of at least 8 hours overnight.

Biochemical parameters, including triglyceride (TG), low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), total cholesterol (TC), fasting blood glucose (FBG), and serum creatinine (SCr) levels, were measured using an autoanalyzer (Hitachi 747; Hitachi, Tokyo, Japan). Levels of high-sensitivity C-reactive protein (hs-CRP) were assessed using an autoanalyzer (Cias Latex CRP-H, Kanto Chemical, Tokyo, Japan). Hypertension was defined as systolic blood pressure \geq 140 mmHg, diastolic blood pressure \geq 90 mmHg, utilisation of antihypertensive medication, or a self-reported history of hypertension. Diabetes was defined as an FBG level \geq 7.0 mmol/L, a history of diabetes, or utilisation of antidiabetic medication. Hyperlipidaemia was defined as TG level \geq 2.3 mmol/L, TC level \geq 6.2 mmol/L, LDL-c level \geq 4.1 mmol/L, HDL-c level <1.0 mmol/L, having a history of hyperlipidaemia, or usage of any lipid-lowering medication. The estimated glomerular filtration rate (eGFR) was defined as eGFR calculated using the CKD-EPI method.

Statistical Analysis

Baseline characteristics are summarised as mean ± standard deviation (SD), median (25th percentile, 75th percentile), or frequency (percentages), as appropriate. Variances between groups were evaluated using one-way analysis of variance or nonparametric tests for continuous variables and chi-square tests for categorical variables. CCB probabilities were estimated using the Kaplan-Meier method and compared using the Log rank test. Cox proportional hazard regression models were used to assess the association between MLR levels and the risk of CCB and its subtypes by calculating the hazard ratio (HR) and corresponding CI (95%). The MLR quartile with the lowest incidence of CCB was selected as the reference group. Model 1 was adjusted for age (continuous variable, years) and sex (male or female). Model 2 was adjusted for age, sex, current smoking (yes or no), current drinking (yes or no), physical activity (active or inactive), and BMI (<24 kg/m², 24–28 kg/m², or ≥28 kg/m²). Model 3 was adjusted for all covariates in model 2 and additionally adjusted for hypertension, diabetes, hyperlipidaemia, eGFR [continuous, mL/(min·1.73 m²)], hs-CRP (continuous, mg/

L), and snoring (ever or never). Model 4 was adjusted for all covariates in model 3 and additionally adjusted for the use of antihypertensive, hypoglycaemic, and lipid-lowering medications (yes or no) at baseline. To fully assess potential nonlinear relationships, the association between the MLR and the risk of CCB and its subtypes was assessed on a continuous scale using restricted cubic splines, where the optimal df was determined by minimising the model's Akaike information criterion.²⁶

Further analyses were conducted to assess the robustness of the association between the MLR and the risk of CCB. First, stratified analyses based on baseline age (<60 and ≥60 years) and sex were used to examine the consistency of the effect. Second, to evaluate the robustness of our results, sensitivity analyses were conducted by (1) excluding new-onset cardiac conduction block cases within the first 2 years of follow-up; (2) excluding new-onset myocardial infarction, atrial fibrillation, and heart failure during follow-up; (3) using time-dependent Cox regression models in which the level of MLR and covariates were updated at each follow-up and the most recent measurements were used to estimate risk; and (4) performing competing risk regression using the Fine-Gray model to address the potentially confounding issue of death.

Statistical analyses were performed using SAS software (version 9.4; SAS Institute, Cary, NC, USA). All tests were two-sided, and P < 0.05 was considered statistically significant.

Results

The cohort consisted of 82,472 participants with a mean age of 51.50 years, and 79.01% were male. The baseline characteristics of the participants based on MLR quartiles are summarised in Table 1. Individuals with higher MLR levels were generally older; less likely to be current smokers, drinkers, or snorers; and exhibited higher body mass index and hs-CRP levels but lower eGFR levels.

During the median follow-up of 10.4 years, 3222 new-onset CCB events occurred, corresponding to an incidence of 3.76 per 1000 person-years. For the CCB subtypes, there were 949, 1506, and 700 incident cases of AVB, RBBB, and

Table I Baseline Characteristics of the Study Population

	Total (n=82472)	Q1 (n=19913)	Q2 (n=20599)	Q3 (n=21499)	Q4 (n=20461)
MLR score		<0.125	0.125-0.167	0.167-0.219	≥0.219
Age, year	51.50±12.68	50.72±12.07	50.29±12.35	51.18±12.76	53.80±13.17
Male, N (%)	65164 (79.01)	15283 (76.75)	16380 (79.52)	17359 (80.74)	16142 (78.89)
Current smoker, N (%)	28404 (34.44)	7195 (36.13)	7224 (35.07)	7489 (34.83)	6496 (31.75)
Current drinker, N (%)	31115 (37.73)	7897 (39.66)	8071 (39.18)	8078 (37.57)	7069 (34.55)
Physical activity, N (%)	12511 (15.17)	3324 (16.69)	3016 (14.64)	3057 (14.22)	3114 (15.22)
Snorer, N (%)	32128 (38.96)	8607 (43.22)	8101 (39.33)	8023 (37.32)	7397 (36.15)
Body mass index, N (%)					
Normal/underweight (<24 kg/m²)	32889 (39.88)	8068 (40.52)	8300 (40.29)	8598 (39.99)	7923 (38.72)
Overweight (24–28 kg/m²)	34342 (41.64)	8330 (41.83)	8591 (41.71)	8866 (41.24)	8555 (41.81)
Obese (≥28 kg/m²)	15241 (18.48)	3515 (17.65)	3708 (18.00)	4035 (18.77)	3983 (19.47)
Monocytes, (10 ⁹ /L)	0.42±0.28	0.23±0.11	0.34±0.18	0.43±0.21	0.67±0.35
Lymphocytes, (10 ⁹ /L)	2.35±1.33	2.56±1.86	2.37±1.23	2.28±1.11	2.20±0.94
hs-CRP, (mg/L)	0.87 (0.30-2.30)	0.81 (0.30-2.10)	0.82 (0.30-2.17)	0.86 (0.30-2.30)	0.93 (0.33–2.68)
eGFR, [mL/(min·1.73m²)]	82.30±22.88	84.56±22.23	84.37±22.58	82.23+23.53	78.10±22.54
Hypertension, N (%)	29735 (36.05)	7201 (36.16)	7269 (35.29)	7820 (36.37)	7445 (36.39)
Diabetes, N (%)	7667 (9.30)	1879 (9.44)	1819 (8.83)	1916 (8.91)	2053 (10.03)
Hyperlipemia, N (%)	35993 (43.64)	8009 (40.22)	8552 (41.52)	9502 (44.20)	9930 (48.53)
Use of antihypertensive agents, N (%)	9297 (11.27)	2339 (11.75)	2182 (10.59)	2274 (10.58)	2502 (12.23)
Use of hypoglycemic medications, N (%)	2006 (2.43)	490 (2.46)	463 (2.25)	495 (2.30)	558 (2.73)
Use of lipid lowering medications, N (%)	793 (0.96)	225 (1.13)	175 (0.85)	167 (0.78)	226 (1.10)

Abbreviations: MLR, monocyte to lymphocyte ratio; hs-CRP, high-sensitivity C-reactive protein; eGFR, indicates estimated glomerular filtration rate.

LBBB, respectively. The cumulative incidences of CCB and its subtypes according to the MLR quartile groups are depicted in Supplementary Figure 2 (Log rank test, P<0.05). The cubic spline models (Figure 1) showed a U-shaped association between MLR and the risk of CCB and AVB (P_{non-linearity} < 0.05) and a nearly linear association between MLR levels and LBBB risk (P_{non-linearity} = 0.90).

The second MLR quartile had the lowest CCB incidence. Both low and high MLR were associated with an increased risk of CCB. The multivariable adjusted hazard ratio (HR) for the Q1 group was 1.106 (95% CI, 1.000-1.224), and for Q3 and Q4, it was 1.134 (95% CI, 1.026-1.253) and 1.212 (95% CI, 1.097-1.340), respectively (Figure 2; Supplementary Table 2). For the CCB subtypes, a similar trend was observed for AVB but not for RBBB and LBBB (Figure 2; Supplementary Table 2). The lowest incidence of the LBBB subtype was observed in the lowest MLR quartile. In the multivariate-adjusted models, elevated MLR levels were significantly associated with an increased risk of LBBB (Supplementary Table 3).

The association between MC or LC alone and incident CCB was further tested in age- and sex-adjusted and multivariate-adjusted models. Neither the MC nor LC levels were associated with the risk of CCB (Supplementary Table 4).

Subgroup analyses of MLR levels and CCB risk are shown in Table 2. There were significant interactions among MLR, age, and sex (all $P_{\text{interaction}} < 0.05$). The association between MLR and new-onset CCB was more prominent in male participants and younger individuals, and these trends were also observed in the AVB subtype.

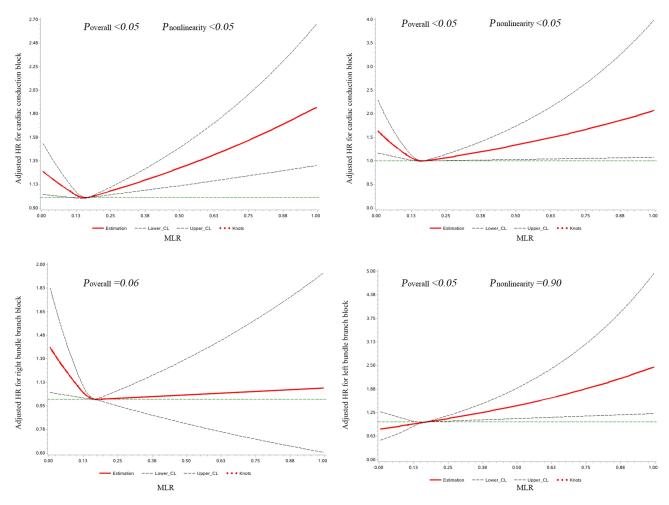


Figure I Associations of MLR with the risk of cardiac conduction block and its subtypes using restricted cubic spline regression models (model 4). Point estimates (solid line) and 95% confidence intervals (dashed lines) were obtained by restricted cubic spline models.

Abbreviations: HR, hazard ratio; CI, confidence interval; MLR, monocyte to lymphocyte ratio.

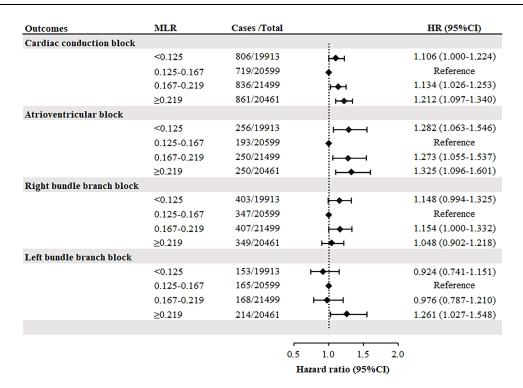


Figure 2 Forest plot for multivariate adjusted hazard ratio for association of monocyte to lymphocyte ratio categories and cardiac conduction block and its subtypes (model 4).

Abbreviations: HR, hazard ratio; Cl, confidence interval; MLR, monocyte to lymphocyte ratio.

A series of sensitivity analyses were performed to examine the robustness of the association between MLR levels and CCB risk, such as 1) excluding new-onset cardiac conduction block within the first two years of follow-up (n=81218), 2) excluding new-onset myocardial infarction, atrial fibrillation, and heart failure during follow-up (n=80059), and 3) using time-varying MLR as exposures. No significant changes were observed in sensitivity analysis. Meanwhile, the competing risk analysis using the Fine-Grey model showed a trend similar to that of the original analysis (Supplementary Table 5).

Discussion

Our study, based on a large prospective cohort, revealed a U-shaped relationship between MLR and overall CCB risk. Regarding the specific subtypes of CCB, an increase in MLR was significantly associated with an elevated risk of AVB and LBBB, but not RBBB. Conversely, a decrease in the MLR was significantly associated with an increased risk of developing AVB.

MLR is a novel inflammatory biomarker that is less affected by various physiological and pathological conditions and can effectively reflect the balance between inherent and acquired immune responses. Compared with MC or LC alone, MLR provides a more stable and accurate evaluation of the body's immune status. ^{9,14} In a study with a mean follow-up of 11 years, Hua et al found that MLR was a strong independent predictor of all-cause mortality and cardiovascular mortality in the general population. ¹⁰ In a retrospective analysis by Liu et al in China, a high MLR was predictive of the risk of ischemic stroke. ²⁷ Another study conducted in the Netherlands revealed a strong relationship between MLR levels and heart failure markers and predicted heart failure hospitalisations during the follow-up period in patients with coronary artery disease. ²⁸ Our study revealed a significant association between MLR and CCB risk, whereas no significant association was observed between MC or LC alone and CCB. This suggests that the immune status reflected by MLR is closely associated with CCB occurrence.

Previous evidence suggests that inflammatory responses and immune dysfunction play important roles in CCB. ^{13,29} Our study demonstrated that elevated MLR was significantly associated with an increased risk of AVB and LBBB. This phenomenon can be explained using several mechanisms. First, increased chemotaxis of monocytes in the circulatory

Table 2 The Associations Between the Level of MLR and the Risk of CCB and Its Subtypes Stratified by Age and Sex

	P _{interaction}	HR (95% CI)				
		QI	Q2	Q3	Q4	
Cardiac conduction block, N (%)						
Age<60	<0.05	1.087 (0.967–1.220)	Ref.	1.165 (1.037–1.307)	1.265 (1.124–1.424)	
Age≥60		1.179 (0.963–1.444)	Ref.	1.035 (0.849–1.262)	1.080 (0.895-1.304)	
Male	<0.05	1.156 (1.032–1.295)	Ref.	1.125 (1.005–1.259)	1.224 (1.093–1.371)	
Female		0.954 (0.768–1.184)	Ref.	1.164 (0.939–1.443)	1.152 (0.929–1.427)	
Atrioventricular block, N (%)						
Age<60	<0.05	1.164 (0.933–1.451)	Ref.	1.245 (0.998–1.553)	1.429 (1.142–1.788)	
Age≥60		1.692 (1.183–2.420)	Ref.	1.341 (0.935–1.923)	1.165 (0.816–1.664)	
Male	<0.05	1.368 (1.107–1.691)	Ref.	1.305 (1.055–1.614)	1.372 (1.107–1.701)	
Female		1.022 (0.686–1.521)	Ref.	1.161 (0.776–1.738)	1.129 (0.753–1.694)	
Right bundle branch block, N (%)						
Age<60	0.41	1.141 (0.969–1.343)	Ref.	1.211 (1.029–1.425)	1.084 (0.909–1.291)	
Age≥60		1.173 (0.864–1.593)	Ref.	0.964 (0.711–1.307)	0.938 (0.700–1.257)	
Male	0.27	1.167 (0.994–1.371)	Ref.	1.109 (0.945–1.303)	1.064 (0.901–1.257)	
Female		1.084 (0.787-1.492)	Ref.	1.341 (0.976–1.842)	0.961 (0.682–1.355)	
Left bundle branch block, N (%)						
Age<60	0.34	0.991 (0.767–1.281)	Ref.	1.078 (0.837–1.389)	1.383 (1.078–1.774)	
Age≥60		0.760 (0.493-1.172)	Ref.	0.741 (0.493–1.114)	0.975 (0.679–1.401)	
Male	0.31	0.999 (0.779–1.281)	Ref.	0.956 (0.748-1.222)	1.193 (0.940–1.512)	
Female		0.721 (0.448–1.159)	Ref.	1.047 (0.670–1.638)	1.488 (0.986–2.244)	

Notes: Model adjusted for age, sex, smoking, alcohol consumption, body mass index, physical activity, diabetes, hypertension, hyperlipidemia, eGFR, hs-CRP, snoring, antihypertensive, lipid-lowering and glucose-lowering drugs.Q1: MLR<0.125, Q2: MLR 0.125–0.167, Q3: MLR 0.167–0.219, Q4: MLR≥0.219.

Abbreviations: MLR, monocyte to lymphocyte ratio; hs-CRP, high-sensitivity C-reactive protein; eGFR, estimated glomerular filtration rate; Ref, reference; HR, hazard ratio and CI, confidence interval.

system, along with enhanced infiltration of macrophages in the heart, plays an important role in the development of cardiac fibrosis, ^{30–33} which serves as a crucial pathological basis for conduction system abnormalities. ^{29,34} Second, agerelated degenerative changes in the conduction system contribute to the pathogenesis of the conduction block, ³⁵ and ageing is often accompanied by adaptive immune system suppression and chronic low-grade inflammation, reflected as a decreased level of T lymphocytes and an increase in circulatory monocyte level. ³⁶ Additionally, a recent Mendelian randomization analysis provided evidence of a causal relationship between elevated MC and decreased LC and the deposition of abnormal cardiac proteins, which are known to be significantly associated with the occurrence of conduction blocks. ^{37,38} Overall, the precise mechanisms underlying the association between high MLR and increased risk of CCB are still not fully understood.

Additionally, our study revealed a significant association between a decreased MLR and the risk of new-onset AVB. Similar results were observed in a previous study focusing on overall bradyarrhythmia events. The specific mechanisms underlying the association between decreased MLR and increased risk of AVB remain elusive. Several factors might have contributed to this observation. Monocytes and macrophages play important roles in inflammation initiation and remission. When their levels are too low, it may impede the inflammatory remission process, leading to a chronic low inflammatory state that can contribute to cardiac conduction system remodelling. Additionally, certain subsets of lymphocytes, such as Th17 cells, exhibit pro-inflammatory properties. The increased number of these lymphocyte subsets can directly participate in or indirectly modulate cardiac remodelling by releasing various inflammatory mediators. Together, decreased MC or increased LC may play an etiological role in the conduction block, but the specific mechanisms require further investigation.

The results of the subgroup analyses in males were consistent with our main findings regarding the association between MLR with CCB and AVB. However, no statistically significant association was observed among females. Similar findings were previously reported by Yang et al, who found that the effect of MLR on bradyarrhythmia was

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more pronounced in males than in females.¹³ This difference may be related to the effect of oestrogen, as it can reduce the risk of cardiac fibrosis⁴¹ and enhance the antioxidant capacity of the myocardial tissue,⁴² which may counteract the risk associated with MLR. Stratification analysis by age showed a significant association between higher MLR and the risk of CCB and its subtypes in participants aged < 60 years. This may be partly because ageing itself being a risk factor for developing conduction block,³⁶ which could diminish the effect of MLR in older individuals. This suggests that more attention should be paid to higher MLR in young and middle-aged individuals to prevent the occurrence of CCB.

This study had several major strengths, including a large sample size, long-term follow-up, and the use of multiple sensitivity analyses to verify the stability of the results. Outcome event data were collected through follow-ups rather than relying solely on hospitalisation reports. This approach allows for the identification of subclinical CCB, thereby avoiding underestimation of the true incidence. To the best of our knowledge, this is the first cohort study to examine the association between MLR and CCB. Despite these strengths, it is important to acknowledge the limitations of this study. First, the observational study design made it impossible to establish cause-effect relationships among the investigated variables. Further basic and Mendelian randomization studies are necessary to confirm these findings. Moreover, this was a single-centre study; despite adjusting for possible confounders, residual confounding cannot be ruled out. Finally, the study population was limited to occupational northern Chinese individuals, which may limit the generalizability of the results to other populations or regions.

Conclusion

Our study, based on a large prospective cohort, found a U-shaped association between the MLR and CCB risk. A similar association persisted for AVB; however, a linear association was observed for LBBB. Further studies are required to confirm our findings and explore the underlying mechanisms.

Data Sharing Statement

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Ethical Approval Statement

The ethics committee of Kailuan General Hospital (approval number: 2006-05). [2006] Yilunzi No. 5.

Acknowledgments

We thank all the survey teams of the Kailuan Study Group for their contribution and the study participants who contributed their information.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Funding

There is no funding to report.

Disclosure

The authors declared no conflict of interest.

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