Lack of Association between 
ABO, PPAP2B, ADAMST7, PIK3CG, and EDNRA and Carotid Intima-Media 
Thickness, Carotid Plaques, and Cardiovascular Disease in 
Patients with Rheumatoid Arthritis

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Introduction. Rheumatoid arthritis (RA) is a polygenic disease associated with accelerated atherosclerosis and increased cardiovascular (CV) mortality. Recent studies have identified the ABO rs579459, PPAP2B rs17114036, and ADAMTS7 rs3825807 polymorphisms as genetic variants associated with coronary artery disease and the PIK3CG rs17398575 and EDNRA rs1878406 polymorphisms as the most significant signals related to the presence of carotid plaque in nonrheumatic Caucasian individuals. Accordingly, we evaluated the potential relationship between these 5 polymorphisms and subclinical atherosclerosis (assessed by carotid intima-media thickness (cIMT) and presence/absence of carotid plaques) and CV disease in RA. Material and Methods. 2140 Spanish RA patients were genotyped for the 5 polymorphisms by TaqMan assays. Subclinical atherosclerosis was evaluated in 620 of these patients by carotid ultrasonography technology. Results. No statistically significant differences were found when each polymorphism was assessed according to cIMT values and presence/absence of carotid plaques in RA, after adjusting the results for potential confounders. Moreover, no significant differences were obtained when RA patients were stratified according to the presence/absence of CV disease after adjusting for potential confounders. Conclusion. Our results do not confirm association between ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 and subclinical atherosclerosis and CV disease in RA.
1. Introduction

Rheumatoid arthritis (RA) is an autoimmune disease characterized by inflammation of synovial tissues in the joints and pannus formation and erosion [1]. This pathology affects up to 1% of the Caucasian population [2] and is associated with increased risk of cardiovascular (CV) disease and CV death [3]. This is the result of a process of accelerated atherosclerosis [4]. Although the etiology of RA is still unknown, traditional CV risk factors, chronic inflammation [5], and genetic background [6–9] have been implicated in the augmented CV mortality detected in this disorder. Subclinical atherosclerosis has been observed in patients with RA [10], even in those without traditional CV risk factors [10]. In this regard, both the assessment of carotid intima-media thickness (cIMT) and the presence of carotid plaques have been proposed to be useful markers as predictors of CV events in low and intermediate risk groups of nonrheumatic individuals [11] and in RA patients [12, 13].

Recently, a large-scale study of coronary artery disease (CAD) performed in nonrheumatic Caucasian individuals has identified 13 novel loci harboring one or more polymorphisms that were associated with this pathology. Moreover, this study confirmed the association of 10 out of 12 previously reported CAD loci [14]. With respect to this, the polymorphisms ABO (histoblood group ABO system transferase) rs579459, PPAP2B (phosphatidic acid phosphatase type 2B) rs17114036, and ADAMTS7 (metalloproteinase with thrombospondin type 1 motif, 7) rs3825807 have been identified as significant genetic variants associated with CAD in nonrheumatic Caucasian individuals [14]. In addition, a recent meta-analysis of carotid intima-media thickness (cIMT) and presence/absence of carotid plaque performed in nonrheumatic Caucasian individuals has identified five new loci for common cIMT and plaque [15]. In this regard, the polymorphisms PIK3CG (phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit gamma isoform) rs17398575 and EDNRA (endothelin receptor type A) rs1878406 have been revealed as the most significant signals associated with presence of carotid plaques in nonrheumatic Caucasian individuals [15].

Taking all these considerations together, we aimed to determine, for the first time, the potential association between the ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms and subclinical atherosclerosis (assessed by the evaluation of cIMT values and presence/absence of carotid plaques) and CV disease in a large and well-characterized cohort of RA patients.

2. Patients and Methods

2.1. Patients and Study Protocol. For this purpose, a set of 2140 Spanish patients with RA were included in the present study. Blood samples were obtained from patients recruited from Hospital Lucus Augusti (Lugo), Hospital Marqués de Valdecilla (Santander), Hospital de Bellvitge (Barcelona), and Hospital Clínico San Carlos, Hospital La Paz, Hospital La Princesa, Hospital Gregorio Marañon and Hospital 12 de Octubre (Madrid). A subject’s written consent was obtained in all the cases. The Ethics Committees of the corresponding hospitals approved the purpose of the work. All the patients fulfilled the 1987 American College of Rheumatology (ACR) and the 2010 classification criteria for RA [16, 17]. In all the cases, patients were assessed for ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms. In addition, cIMT and presence/absence of carotid plaques were determined by carotid ultrasonography (US) in 620 of these patients.

Information on the main demographic data, clinical characteristics, CV risk factors, and CV events of patients enrolled in the study is shown in Table 1. Three hundred and seventy-seven (17.6%) of these 2140 patients had experienced CV events. Definitions of CV events and traditional CV risk factors were established as previously described [4, 13].

2.2. Genotyping. DNA from patients was obtained from peripheral blood using standard methods.

Table 1: Demographic and clinical characteristics of the Spanish patients with RA included in the study.

| Clinical feature | % (n/N) |
|------------------|---------|
| Patients         | 2140    |
| Main characteristics |       |
| Age at the time of disease onset (years, mean ± SD) | 52.4 ± 14.9 |
| Follow-up (years, mean ± SD) | 12.2 ± 8.8 |
| Percentage of women | 75.1 |
| Rheumatoid factor positive* | 67.9 (1388/2044) |
| Anti-CCP antibodies positive | 59.2 (1058/1786) |
| Shared epitope positive | 62.4 (736/1180) |
| Erosions | 54.7 (940/1718) |
| Extra-articular manifestations** | 31.5 (540/1715) |
| Cardiovascular risk factors |       |
| Hypertension | 38.7 (817/2108) |
| Diabetes mellitus | 12.2 (258/2108) |
| Dyslipidemia | 37.3 (786/2108) |
| Obesity | 16.8 (355/2108) |
| Smoking habit | 23.8 (502/2108) |
| Patients with cardiovascular events | 17.6 (377/2140) |
| Ischemic heart disease | 8.3 (179/2140) |
| Heart failure | 5.3 (113/2140) |
| Cerebrovascular accident | 5.3 (114/2140) |
| Peripheral arteriopathy | 2.3 (50/2140) |

RA: rheumatoid arthritis; SD: standard deviation; Anti-CCP antibodies: anticyclic citrullinated peptide antibodies.

* At least two determinations were required for analysis of this result.

** Extra-articular manifestations of the disease (if RA patients experienced at least one of the following manifestations: nodular disease, Felty’s syndrome, pulmonary fibrosis, rheumatoid vasculitis, or secondary Sjogren’s syndrome) [4].
The ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms were genotyped with TaqMan predesigned single-nucleotide polymorphism genotyping assays in a 7900HT Real-Time polymerase chain reaction (PCR) system, according to the conditions recommended by the manufacturer (Applied Biosystems, Foster City, CA, USA). Negative controls and duplicate samples were included to check the accuracy of genotyping.

2.3. Carotid US Examination. Measurement of the cIMT and presence/absence of carotid plaques were performed in 620 patients from Lugo and Santander by carotid US. Patients from Santander were assessed using a commercially available scanner, MyLab 70, Esaote (Genoa, Italy), equipped with 7–12 MHz linear transducer and the automated software-guided technique radiofrequency—Quality Intima Media Thickness in real-time (QIMT, Esaote, Maastricht, Holland)—was used [18]. Patients from Lugo were assessed using high-resolution B-mode ultrasound, Hewlett Packard SONOS 5500, with a 10-MHz linear transducer as previously reported [19]. cIMT was measured at the far wall of the right and left common carotid arteries, 10 mm from the carotid bifurcation, over the proximal 15 mm long segment. cIMT was determined as the average of three measurements in each common carotid artery. The final cIMT was the largest average cIMT (left or right). The plaque criteria in the accessible extracranial carotid tree (common carotid artery, bulb, and internal carotid artery) were cIMT >1.5 mm, protrusion at least 50% greater than the surrounding cIMT, or arterial lumen encroaching >0.5 mm, according to Mannheim consensus criteria [20]. The carotid plaques were counted in each territory and defined as no plaque, unilateral plaque, or bilateral plaques [18, 20–22]. Agreement between the two US methods in patients with RA was previously reported [21]. Two experts with high experience and close collaboration in the assessment of subclinical atherosclerosis in RA from Santander (AC) and Lugo (CGJ) performed the studies.

2.4. Statistical Analysis. Statistical power for CV events was calculated using CaTS—Power Calculator for Two Stage Association Studies (http://www.sph.umich.edu/csg/abecasis/CaTS/). All genotype data were checked for deviation from Hardy-Weinberg equilibrium (HWE) using http://ihg.gsdf.de/cgi-bin/hw/hwa1.pl.

cIMT values are displayed as mean and standard deviation (SD). The association between genotypes and alleles of each polymorphism and cIMT values was tested using unpaired t-test to compare between 2 groups and one-way analysis of variance (ANOVA) to compare more than two groups. Comparisons of means were adjusted for sex, age at the time of US study, follow-up time, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) as potential confounders. Similarly, no statistically significant differences were detected when each polymorphism was evaluated according to the presence/absence of carotid plaques in RA, after adjusting the results for sex, age at the time of US study, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) as potential confounders. Similarly, no statistically significant differences were detected when each polymorphism was evaluated according to the presence/absence of carotid plaques in RA, after adjusting the results for sex, age at the time of US study, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) as potential confounders.

Differences in the genotypic and allelic frequencies of each polymorphism according to the presence/absence of carotid plaques were calculated by χ² or Fisher tests when necessary (expected values below 5). Strengths of associations were estimated using odds ratios (OR) and 95% confidence intervals (CI). Results were adjusted for sex, age at the time of US study, and traditional CV risk factors by logistic regression.

The relationship between genotypes and alleles of each polymorphism and CV events that occurred in the follow-up was tested using Cox regression adjusted for sex, age at RA diagnosis, and traditional CV risk factors. For that purpose, we used the most frequent genotype and allele as reference; the end of follow-up was the first date among the end of the study, date of death, or date of CV event. Follow-up time was estimated as the difference between the RA diagnosis date and the end of follow-up. Patients without CV events in the follow-up time or dying by any non-CV cause were considered as censored. Results were expressed as hazard ratios (HR) with 95% confidence interval (CI).

Statistical significance was defined as P < 0.05. All analyses were performed with STATA statistical software 12/SE (Stata Corp., College Station, TX, USA).

3. Results

This study had >80% of power to detect genotypic OR >1.4 for ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406.

The ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 genotype distributions were in Hardy-Weinberg equilibrium. The genotyping success was greater than 97% in all the cases.

Table 2 describes the potential association between the ABO rs579459, PPAP2B rs17114036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms and both subclinical atherosclerosis and CV disease in RA patients. As shown in Table 2, no statistically significant differences were found when each polymorphism was assessed according to the evaluation of the cIMT in RA patients, after adjusting the results for sex, age at the time of US study, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) as potential confounders. Similarly, no statistically significant differences were detected when each polymorphism was evaluated according to the presence/absence of carotid plaques in RA, after adjusting the results for potential confounder factors (Table 2). Moreover, when RA patients were stratified according to the presence or absence of CV disease, no significant differences were obtained after adjusting the results for sex, age at RA diagnosis, and traditional CV risk factors (Table 2). It was also the case when RA patients were stratified according to the presence/absence of ischemic heart disease (data not shown).

4. Discussion

CV disease is the most common cause of premature mortality in patients with RA being a consequence of accelerated atherosclerosis [23]. Since a genetic background [6–9] has been involved in the accelerated atherosclerosis in RA, several


Table 2: Association between AB0, PPAP2B, ADAMTS7, PIK3CG, and EDNRA genotypes and alleles and cIMT, presence/absence of carotid plaques, and CV events in RA patients.

| Genotype | cIMT mm (n = 620) | Presence/absence of carotid plaques (n = 620) | Presence/absence of CV events (n = 2140) |
|----------|-----------------|---------------------------------|---------------------------------|
|          | Mean ± SD | P* | P** | OR [95% CI]** | P*** | HR [95% CI]*** |
| AB0 rs579459 |
| TT | 0.73 ± 0.17 | 0.91 | — | 1 (Ref.) | — | 1 (Ref.) |
| TC | 0.74 ± 0.17 | 0.23 | 0.66 [0.33–1.30] | 0.79 | 1.05 [0.72–1.53] |
| CC | 0.71 ± 0.17 | 0.63 | 0.71 [0.18–2.86] | 0.22 | 0.59 [0.26–1.37] |
| T  | 0.74 ± 0.17 | 0.86 | — | 1 (Ref.) | — | 1 (Ref.) |
| C   | 0.73 ± 0.17 | 0.29 | 0.76 [0.45–1.26] | 0.47 | 0.89 [0.67–1.20] |
| PPAP2B rs17114036 |
| AA | 0.74 ± 0.17 | 0.64 | — | 1 (Ref.) | — | 1 (Ref.) |
| AG | 0.73 ± 0.19 | 0.55 | 1.14 [0.71–1.86] | 0.99 | 0.99 [0.58–1.70] |
| GG | 0.74 ± 0.23 | 0.10 | 0.72 [0.03–1.54] | ¥ | ¥ |
| A  | 0.74 ± 0.17 | 0.99 | — | 1 (Ref.) | — | 1 (Ref.) |
| G  | 0.74 ± 0.19 | 0.10 | 0.49 [0.21–1.15] | 0.48 | 0.83 [0.50–1.39] |
| ADAMTS7 rs3825807 |
| AA | 0.76 ± 0.17 | 0.46 | — | 1 (Ref.) | — | 1 (Ref.) |
| AG | 0.73 ± 0.17 | 0.44 | 0.86 [0.58–1.28] | 0.10 | 0.59 [0.39–1.20] |
| GG | 0.72 ± 0.17 | 0.72 | 0.92 [0.57–1.49] | 0.73 | 0.92 [0.56–1.49] |
| A  | 0.74 ± 0.17 | 0.34 | — | 1 (Ref.) | — | 1 (Ref.) |
| G  | 0.73 ± 0.17 | 0.53 | 0.86 [0.55–1.36] | 0.43 | 0.89 [0.69–1.17] |
| PIK3CG rs17398575 |
| GG | 0.73 ± 0.16 | 0.1 | — | 1 (Ref.) | — | 1 (Ref.) |
| GA | 0.74 ± 0.18 | 0.34 | 1.38 [0.72–2.64] | 0.99 | 1.00 [0.68–1.47] |
| AA | 0.75 ± 0.19 | 0.86 | 0.86 [0.17–4.26] | 0.95 | 1.02 [0.47–2.23] |
| G  | 0.73 ± 0.17 | 0.09 | — | 1 (Ref.) | — | 1 (Ref.) |
| A  | 0.74 ± 0.18 | 0.56 | 1.17 [0.70–1.95] | 0.96 | 1.00 [0.75–1.36] |
| EDNRA rs1878406 |
| CC | 0.73 ± 0.16 | 0.61 | — | 1 (Ref.) | — | 1 (Ref.) |
| CT | 0.74 ± 0.19 | 0.28 | 1.54 [0.70–3.38] | 0.44 | 0.83 [0.53–1.32] |
| TT | 0.73 ± 0.14 | 0.80 | 0.76 [0.10–6.04] | 0.85 | 1.10 [0.40–2.90] |
| C  | 0.74 ± 0.16 | 0.32 | — | 1 (Ref.) | — | 1 (Ref.) |
| T  | 0.74 ± 0.18 | 0.47 | 1.28 [0.66–2.47] | 0.62 | 0.91 [0.62–1.33] |

cIMT: carotid intima-media thickness; CV: cardiovascular; RA: rheumatoid arthritis; SD: standard deviation; OR: odds ratio; CI: confidence interval; HR: hazard ratios.
* Adjusted for sex, age at the time of ultrasonography study, follow-up time, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) using analysis of covariance (ANCOVA).
** Adjusted for sex, age at the time of ultrasonography study, follow-up time, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) by logistic regression.
*** Adjusted for sex, age at RA diagnosis, follow-up time, and traditional CV risk factors (hypertension, diabetes mellitus, dyslipidemia, obesity, and smoking habit) using Cox regression.
¥ Model does not converge.

Studies have been focused on the search of genetic markers that may improve the identification of RA patients at risk of experiencing CV events. Previous studies have described the AB0 rs579459, PPAP2B rs17114036, and ADAMTS7 rs3825807 polymorphisms as genetic variants associated with coronary artery disease [14] and the PIK3CG rs17398575 and EDNRA rs1878406 polymorphisms as the most significant signals related to the presence of carotid plaque [15] in nonrheumatic Caucasian individuals. Taking into account these considerations, we assessed for the first time the potential association between these 5 polymorphisms and subclinical atherosclerosis and CV events in RA patients.

Nevertheless, in contrast to the results obtained in nonrheumatic Caucasian individuals [14], the relationship between the AB0 rs579459, PPAP2B rs17114036, ADAMTS7...
rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms and CV events was not statistically significant. Moreover, unlike nonrheumatic individuals [15], we did not disclose association between the ABO rs579459, PPAP2B rs1714036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms and cIMT values or presence/absence of carotid plaques.

Although atherosclerosis and RA are chronic inflammatory diseases that share similar pathophysiological mechanisms [24], additional factors so far unknown may explain the differences observed in terms of genetic predisposition. Further studies to better characterize the genetic susceptibility for accelerated atherosclerosis in RA are under way.

5. Conclusion

Our results do not confirm association of ABO rs579459, PPAP2B rs1714036, ADAMTS7 rs3825807, PIK3CG rs17398575, and EDNRA rs1878406 polymorphisms with subclinical atherosclerosis and CV disease in RA patients.

Conflict of Interests

The authors declare that they have no conflict of interests.

Authors’ Contribution

Raquel López-Mejías, Fernanda Genre, and Mercedes García-Bermúdez carried out genotyping, participated in the design of the study and data analysis, and helped to draft the paper. Begoña Ubilla, José A. Miranda-Filloy, Trinitario Pina, Carlos González-Juanatey, Luis Rodríguez-Rodríguez, Alejandro Balsa, Dora Pascual-Salcedo, Francisco J. López-Longo, Patricia Carreira, and Ricardo Blanco participated in the acquisition and interpretation of data and helped to draft the paper. Santos Castañeda and Benjamin Fernández-Gutiérrez have been involved in the acquisition and interpretation of data and in revising it critically for important intellectual content. Javier Llorca carried out the analysis and interpretation of the data. Carlos González-Juanatey and Alfonso Corrales performed the carotid US examination and they have been involved in the acquisition and interpretation of data and coordination and helped to draft the paper. José A. Miranda-Filloy and Miguel A. González-Gay have made substantial contributions to conception and design of the study, acquisition of data, and coordination and have helped to draft the paper and have given final approval of the version to be published. Raquel López-Mejías, Fernanda Genre, and Mercedes García-Bermúdez had equal contribution. Dr. Javier Martín and Dr. Miguel A. González-Gay shared senior authorship in this study.

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References

[1] L. Klareskog, A. I. Catrina, and S. Paget, “Rheumatoid arthritis,” The Lancet, vol. 373, no. 9664, pp. 659–672, 2009.
[2] S. E. Gabriel and K. Michaud, “Epidemiological studies in incidence, prevalence, mortality, and comorbidity of the rheumatic diseases,” Arthritis Research and Therapy, vol. 11, no. 3, article 229, 2009.
[3] C. P. Chung, A. Oeser, P. Raggi et al., “Increased coronary-artery atherothrombosis in rheumatoid arthritis: relationship to disease duration and cardiovascular risk factors,” Arthritis and Rheumatism, vol. 52, no. 10, pp. 3045–3053, 2005.
[4] M. A. Gonzalez-Gay, C. Gonzalez-Juanatey, M. J. Lopez-Diaz et al., “HLA-DRB1 and persistent chronic inflammation contribute to cardiovascular events and cardiovascular mortality in patients with rheumatoid arthritis,” Arthritis Care and Research, vol. 57, no. 1, pp. 125–132, 2007.
[5] P. H. Dessein, G. R. Norton, A. J. Woodiwiss, B. I. Joffe, and F. Wolfe, “Influence of nonclassical cardiovascular-risk factors on the accuracy of predicting subclinical atherosclerosis in rheumatoid arthritis,” Journal of Rheumatology, vol. 34, no. 5, pp. 943–951, 2007.
[6] R. Lopez-Mejias, M. Garcia-Bermudex, C. Gonzalez-Juanatey et al., “NFKBL94ATTG INS/Del polymorphism (rs2836249) is associated with cardiovascular disease in patients with rheumatoid arthritis,” Atherosclerosis, vol. 224, pp. 426–429, 2012.
[7] L. Rodriguez-Rodriguez, R. Lopez-Mejias, M. Garcia-Bermudex, C. Gonzalez-Juanatey, M. A. Gonzalez-Gay, and J. Martin, “Genetic markers of cardiovascular disease in rheumatoid arthritis,” Mediators of Inflammation, vol. 2012, Article ID 574817, 14 pages, 2012.
[8] R. Lopez-Mejias, F. Genre, M. Garcia-Bermudex et al., “The ZC3H12C rs11556924 polymorphism is associated with increased carotid intima-media thickness in patients with rheumatoid arthritis,” Arthritis Research and Therapy, vol. 15, article R152, 2013.
[9] R. Lopez-Mejias, F. Genre, M. Garcia-Bermudex et al., “The 1q23.3 genomic region—rs964184— is associated with cardiovascular disease in patients with rheumatoid arthritis,” Tissue Antigens, vol. 82, pp. 344–347, 2013.
[10] M. Gonzalez-Juanatey, J. Llorca, A. Testa, J. Revuelta, C. Garcia-Porrua, and M. A. Gonzalez-Gay, “Increased prevalence of severe subclinical atherosclerotic findings in long-term treated rheumatoid arthritis patients without clinically evident...
atherosclerotic disease,” *Medicine*, vol. 82, no. 6, pp. 407–413, 2003.

[11] V. Nambi, L. Chambless, A. R. Folsom et al., “Carotid intima-media thickness and presence or absence of plaque improves prediction of coronary heart disease risk. The ARIC (Atherosclerosis Risk In Communities) Study,” *Journal of the American College of Cardiology*, vol. 55, no. 15, pp. 1600–1607, 2010.

[12] M. R. Evans, A. Escalante, D. F. Battafarano, G. L. Freeman, D. H. O’Leary, and I. Del Rincón, “Carotid atherosclerosis predicts incident acute coronary syndromes in rheumatoid arthritis,” *Arthritis and Rheumatism*, vol. 63, no. 5, pp. 1211–1220, 2011.

[13] C. Gonzalez-Juanatey, J. Llorca, J. Martin, and M. A. Gonzalez-Gay, “Carotid intima-media thickness predicts the development of cardiovascular events in patients with rheumatoid arthritis,” *Seminars in Arthritis and Rheumatism*, vol. 38, no. 5, pp. 366–371, 2009.

[14] H. Schunkert, I. R. König, S. Kathiresan et al., “Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease,” *Nature Genetics*, vol. 43, pp. 333–338, 2011.

[15] J. C. Bis, M. Kavousi, N. Franceschini et al., “Meta-analysis of genome-wide association studies from the CHARGE consortium identifies common variants associated with carotid intima media thickness and plaque,” *Nature Genetics*, vol. 43, pp. 940–947, 2011.

[16] F. C. Arnott, S. M. Edworthy, D. A. Bloch et al., “The American Rheumatism Association 1987 revised criteria for the classification of rheumatoid arthritis,” *Arthritis and Rheumatism*, vol. 31, no. 3, pp. 315–324, 1988.

[17] D. Aletaha, T. Neogi, and A. J. Silman, “2010 Rheumatoid arthritis classification criteria: an American College of Rheumatology/European League Against Rheumatism collaborative initiative,” *Annals of the Rheumatic Diseases*, vol. 69, pp. 1580–1588, 2010.

[18] A. Corrales, C. Gonzalez-Juanatey, M. E. Peiro, R. Blanco, J. Llorca, and M. A. Gonzalez-Gay, “Carotid ultrasound is useful for the cardiovascular risk stratification of patients with rheumatoid arthritis: results of a population-based study,” *Annals of the Rheumatic Diseases*, no. 4, pp. 722–727, 2014.

[19] C. Gonzalez-Juanatey, J. Llorca, C. García-Porrua, J. Martin, and M. A. Gonzalez-Gay, “Effect of anti-tumor necrosis factor α therapy on the progression of subclinical atherosclerosis in severe rheumatoid arthritis,” *Arthritis Care and Research*, vol. 55, no. 1, pp. 150–153, 2006.

[20] P.-J. Touboul, M. G. Hennerici, S. Meairs et al., “Mannheim carotid intima-media thickness consensus (2004–2006): an update on behalf of the advisory board of the 3rd and 4th Watching the Risk Symposium 13th and 15th European Stroke Conferences, Mannheim, Germany, 2004, and Brussels, Belgium, 2006,” *Cerebrovascular Diseases*, vol. 23, no. 1, pp. 75–80, 2007.

[21] E. Naredo, I. Möller, M. Gutiérrez et al., “Multi-examiner reliability of automated radio frequency-based ultrasound measurements of common carotid intima-media thickness in rheumatoid arthritis,” *Rheumatology (Oxford)*, vol. 50, no. 10, pp. 1860–1864, 2011.

[22] A. Corrales, J. A. Parra, C. González-Juanatey et al., “Cardiovascular risk stratification in rheumatic diseases: carotid ultrasound is more sensitive than Coronary Artery Calcification Score to detect subclinical atherosclerosis in patients with rheumatoid arthritis,” *Annals of the Rheumatic Diseases*, vol. 72, pp. 1764–1770, 2013.

[23] M. A. Gonzalez-Gay, C. Gonzalez-Juanatey, and J. Martin, “Rheumatoid arthritis: a disease associated with accelerated atherogenesis,” *Seminars in Arthritis and Rheumatism*, vol. 35, no. 1, pp. 8–17, 2005.

[24] V. Pasceri and E. T. H. Yeh, “A tale of two diseases: atherosclerosis and rheumatoid arthritis,” *Circulation*, vol. 100, no. 21, pp. 2124–2126, 1999.