Serum Homocysteine Levels in Patients with Retinal Vein Occlusion in a Spanish Population

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Abstract

Purpose: Our aim was a) to compare serum Hcy levels in patients with RVO and population-based controls, and b) to evaluate whether hyperhomocysteinemia is a risk factor for RVO.

Patients and methods: Prospective case-control study of all patients diagnosed with RVO at a tertiary-care hospital, and age- and sex-matched controls taking part in a population-based prospective cohort in the same geographic area. Differences in serum Hcy between both groups were analyzed by a general linear model adjusted for age, body mass index (BMI), glomerular filtration rate (GFR), hypertension, dyslipidemia, diabetes, tobacco use and alcohol intake. Besides, we tested the relationship between hyperhomocysteinemia (> 15 µmol/L) and RVO, by a stepwise logistic regression analysis.

Results: RVO patients (n = 85) had a higher prevalence of hypertension (p = 0.002), diabetes (p = 0.008), and dyslipidemia (p = 0.04) than controls (n = 82). Adjusted median Hcy levels were higher in RVO patients (p < 0.0001). Adjusted OR for hyperhomocysteinemia were 4.4 (95% CI, 2.0-9.3; p < 0.0001) in the overall sample, and 2.6 (95% CI, 1.04-6.6; p = 0.04) and 6.1 (95% CI, 1.7-21.6; p = 0.005) for men and women, respectively.

Conclusion: Patients with RVO have higher serum Hcy levels than age- and sex-matched population-based controls. Hyperhomocysteinemia seems to be a risk factor for RVO, independent of age, BMI, GFR and classical vascular risk factors.

Keywords
Retinal vein occlusion, Homocysteine, Hyperhomocysteinemia, Risk factor

Introduction

Retinal vein occlusion (RVO) is one of the most common causes of vision loss in the elderly and has been associated with an increase in cardiovascular mortality [1]. Classical vascular risk factors (VRF) represent the main etiopathogenic factors for RVO. In fact, it is usually thought that the disorder may be considered as a manifestation of systemic atherosclerosis [2].

On the other hand, several studies have suggested that a hypercoagulable state could be involved in some patients with RVO, although this has not been conclusively established [3,4]. In line with this, we have recently shown that acquired thrombophilia (antiphospholipid syndrome and hyperhomocysteinemia) should be considered in the clinical assessment of patients with RVO, whereas genetic thrombophilia must be only ruled out in patients aged < 50 years [5-7].
Homocysteine (Hcy) is a sulfhydryl-containing amino acid formed as an intermediate product of methionine and cysteine metabolism [8]. The first report suggesting a proatherogenic effect of Hcy was published by McCully in 1969 [9]. Since then, increased Hcy levels (above 15 µmol/L) have been related to increased risk of stroke, myocardial infarction and venous thrombosis [10-12].

Several mechanisms have been proposed to explain the possible contribution of serum Hcy to development of RVO, and include the activation of factor V, the increased oxidation of low-density lipoprotein (LDL), the inhibition of plasminogen activator binding, and the activation of protein C [13].

There are some studies analyzing the relationship between serum Hcy levels and RVO [14-19]. Nevertheless, most of them are retrospective in nature, have substantial heterogeneity, adjustment for potential confounder is not well defined, and well-characterized population-based controls are scarce. For that reasons, meta-analysis on the association between serum Hcy and RVO, have reported a high level of heterogeneity among studies, included case-series studies, and only indirectly compared cases and controls. Therefore, the results have been questioned [15,20,21]. In the most recent systematic review and meta-analysis of case-controls studies, Li, et al. [15] founded some evidence that plasma Hcy was related to a moderate increase in RVO. However, the same limitations reported for previous meta-analysis have been found: substantial heterogeneity among included studies, publication and selection bias, lack of adjustment for important confounding factors, unclear definition or selection of controls, and absence of population-based data.

To get deeper insight and shed some light in the association between Hcy and RVO, we have carried out a prospective case-control study with a well-defined cohort of patients diagnosed with RVO and age- and sex-matched population-based controls.

**Patients and Methods**

We conducted a prospective case-control study, from July 2012 to January 2015, at a tertiary-care center that serves as a reference hospital for a population of 350.000 inhabitants in Northern Spain. Based on clinical, fundoscopic and angiographic criteria, all patients diagnosed with RVO at the Division of Ophthalmology, were referred and assessed at our Internal Medicine outpatient clinic department.

A randomly selected age- and sex-matched control group of 82 subjects taking part in a population-based prospective cohort study in the same geographic area [22] were also screened for plasma Hcy levels between April 2013 and January 2015. Those with renal failure (defined as a glomerular filtration rate < 60 ml/min/1.72 m² or a previous diagnosis of chronic kidney disease) or previous arterial or venous RVO were excluded from the present study. Serum vitamin B₁₂ or folate deficiency was ruled out at baseline, and none of the cases and controls received vitamin B₁₂ or folic acid supplementation. All the subjects gave written informed consent. The study protocol was approved by the ethical committee and conducted in accordance with the Declaration of Helsinki.

**Clinical variables**

Data were collected using a prespecified standardized questionnaire, in a computerizing database. We assessed the following clinical variables: age, sex, weight, height, body mass index (BMI), current tobacco use, alcohol intake (> 20 g per day), high blood pressure (equal or greater than 140/90 mm Hg or being on antihypertensive agents), dyslipidemia (serum total cholesterol or triglyceride levels greater than 230 mg/dl and 150 mg/dl respectively or being on lipid-lowering drugs), diabetes mellitus (according to the ADA criteria) [23], past or present history of thromboembolic disease in another location different from retinal, history of ischemic heart disease, stroke or peripheral arterial disease, type of RVO (central or branch), and prescribed treatments.

**Laboratory parameters and Hcy measurement**

Routine biochemical parameters were measured by standard automated methods in an ADVIA 2400 Chemistry System autoanalyzer (Siemens). Blood samples were obtained from an antecubital vein in the morning, after a requested 12-hour overnight fast.

We estimated glomerular filtration rate (eGFR) using the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equations [24].

We measured total serum Hcy by chemiluminescent immunoassay (Immulite 2000 Xpi; Siemens*). Serum sample releases Hcy from its binding proteins. It is converted to S-adenosyl-homocysteine (SAH) in the presence of S-adenosyl-L-Hcy hydrodase and dithiothreitol (DTT). After 30 minute incubation, the treated sample is transferred to a second reaction tube containing a SAH-coated polystyrene bead and an alkaline phosphatase-labeled antibody specific for SAH. During 30 minute incubation, the converted SAH from the sample pretreatment competes with immobilized SAH for binding alkaline phosphatase-labeled anti-SAH antibody conjugate. Unbound enzyme conjugate is removed by a centrifugal wash. The bound label is then quantified using the dioxetane substrate to produce light. Light is emitted when the chemiluminescent substrate reacts with the al-
adjusted for age, BMI, hypertension, dyslipidemia, diabetes mellitus, and tobacco use and alcohol intake. Besides, further adjustment for past history of ischemic heart disease, stroke, and peripheral arterial disease and thromboembolic disorders was performed.

Finally, we have developed a stepwise logistic regression analysis to test the relationship between hyperhomocysteinemia, (defined as a serum Hcy level above 15 µmol/L) and RVO. Odds ratios (ORs) (95% CI) were calculated and regression models were adjusted for the same covariates above mentioned. All analyses were conducted using SPSS 20.0 (Chicago, IL, USA). A p value < 0.05 was considered statistically significant in all the calculations.

## Results

Eighty-five patients (45 men and 40 women; mean age, 69 ± 12 years) and 82 controls (43 men and 39 women; mean age, 67 ± 9 years), were included in the study. Fifty-five patients (64.7%) had central RVO and 30 (35.3%) had branch RVO.

### Statistical analysis

Baseline characteristics of the population were calculated for the total sample and for men and women separately. Serum Hcy was not normally distributed and was log-transformed before analysis. Results were expressed as mean ± SD, median (interquartile range-IQR-) or percentages, as appropriate. Student’s t test or Mann-Whitney U-test were used to determine the differences between groups for continuous variables, and \( \chi^2 \)-test for categorical variables. We constructed a general linear model to test the differences in serum Hcy levels between cases and controls. Multivariable models were adjusted for age, BMI, hypertension, dyslipidemia, diabetes mellitus, and tobacco use and alcohol intake. Besides, further adjustment for past history of ischemic heart disease, stroke, and peripheral arterial disease and thromboembolic disorders was performed.

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### Table 1: Baseline characteristics of the study sample.

| Variable                                | Cases N = 85 | Controls N = 82 | p    |
|-----------------------------------------|--------------|-----------------|------|
| Sex (men), %                            | 52.9         | 52.4            | 0.99 |
| Age (yrs), mean ± SD                    | 69.0 ± 12.3  | 67.2 ± 9.4      | 0.28 |
| Weight (Kg), mean ± SD                  | 75.2 ± 13.3  | 78.4 ± 14.9     | 0.15 |
| Height (cm), mean ± SD                  | 162.0 ± 8.9  | 162.1 ± 9.9     | 0.97 |
| BMI (Kg/m²), mean ± SD                  | 28.6 ± 4.6   | 29.8 ± 4.9      | 0.11 |
| Current smoker, %                       | 9.4          | 20.7            | 0.07 |
| Alcohol, %                              | 24.7         | 18.3            | 0.41 |
| Hypertension, %                         | 75.3         | 52.4            | 0.002|
| Dyslipidemia, %                         | 62.4         | 46.3            | 0.044|
| Diabetes, %                             | 25.9         | 9.8             | 0.008|
| HDL cholesterol (mg/dl), mean ± SD      | 56.6 ± 14.5  | 58.2 ± 12.8     | 0.26 |
| GFR, ml/min/1.73 m², median [IQR]       | 82.4 [69.6-91.7] | 88.2 [75.3-94.6] | 0.04 |
| Stroke, %                               | 4.7          | 6.2             | 0.74 |
| Ischemic heart disease, %               | 4.7          | 4.9             | 0.99 |
| Peripheral arterial disease, %          | 4.7          | 3.7             | 0.99 |
| DVT/PE, %                               | 0.0          | 3.7             | 0.11 |
| ASA, %                                  | 14.3         | 19.0            | 0.92 |
| Oral anticoagulants, %                  | 7.1          | 8.6             | 0.95 |
| Homocysteine (µmol/L), median [IQR]     | 15.1 [11.4-19.6] | 11.2 [9.0-14.6] | < 0.0001|
| Homocysteine > 15 µmol/L, %             | 50.6         | 23.2            | < 0.0001|

SD: Standard deviation; BMI: Body mass index; HDL: High density lipoprotein; GFR: Glomerular filtration rate (CKD-EPI equations); DVT: Deep venous thrombosis; PE: Pulmonary embolism; ASA: Acetylsalicylic acid; IQR: Interquartile range.
with RVO was the highest in the last quartile whereas it was the lowest in the group of controls. These figures are diametrically opposed in the first serum Hcy quartile.

Median serum Hcy levels as well as hyperhomocysteinemia defined as a cutoff value of 15 µmol/L were higher in cases than in controls (p < 0.0001, respectively). After adjusting for age, BMI, glomerular filtration rate, hypertension, dyslipidemia, diabetes mellitus, tobacco use and alcohol intake, differences in serum Hcy levels remained significant (15.6 vs. 14.2 µmol/L; p = 0.008).

(Table 2) shows linear general model adjusted for these covariates and stratified by sex. Associations between serum Hcy and RVO was significant in women (p = 0.02) and borderline significant in the group of men (p = 0.06). Further adjustment for past history of ischemic heart disease, stroke, peripheral arterial disease and thromboembolic disorders, did not virtually change these results (data not shown).

Unadjusted OR for hyperhomocysteinemia was 3.4 (95% CI, 1.7-6.6; p < 0.0001) in the overall sample, and 2.6 (95% CI, 1.1-6.3; p = 0.03) and 4.9 (95% CI, 1.7-14.5; p = 0.003) for men and women respectively. The correspondent figures after adjustment for confounders were 4.4 (95% CI, 2.0-9.3; p < 0.0001); 2.6 (95% CI, 1.04-6.6; p = 0.04), and 6.1 (95% CI, 1.7-21.6; p = 0.005).

**Discussion**

In the last years, Hcy has been recognized as an atherosclerotic and thrombotic risk factor [25,26]. However, its role in the pathological changes associated with atherosclerosis as well as the intrinsic mechanisms triggered by Hcy accumulation is unclear to date.

Hyperhomocysteinemia would act as an atherosclerotic risk factor in several ways. Firstly, Hcy metabolites can combine with LDL-cholesterol to produce aggregates, which may be taken by macrophages in the arterial intima, and contribute to the development of the atherosclerotic plaque [27]. Nevertheless, the effects of hyperhomocysteinemia on plaque rupture or instability are poorly understood although obstruction of vasa vasorum by aggregates of microorganisms with homocysteinylated low-density lipoproteins has been proposed to cause arterial wall ischemia and a microabscess of the intima, leading to vulnerable atherosclerotic plaque [28]. Secondly, Hcy may activate inflammatory responses that lead to the recruitment of monocytes to the arterial wall [27]. Thirdly, high Hcy levels may dysregulate lipid metabolism in vascular cells through activation of the sterol regulatory element-binding protein family of transcription factors [29]. Finally, hyperhomocysteinemia also increases smooth muscle cell proliferation, and its oxidation would form free radicals, inducing endothelial damage [30].

Although RVO pathogenesis remains still unclear, the disease is considered a manifestation of the atherosclerosis process [31]. In this sense, classical vascular risk factors, such as hypertension, diabetes, dyslipidemia and smoking have been reported as the main predisposing factors for RVO [32]. In this sense, it has been suggested that hyperhomocysteinemia may lead to an enhancement of the adverse effects of these classical risk factors [33].

The role of hyperhomocysteinemia as a risk factor for RVO is an ongoing matter of debate. Several publications have found positive associations, but negative results have also been reported [16,18,34]. Besides, three meta-analysis have been published to date, and all of them have suggested that there is some evidence that serum Hcy is associated with RVO. Nevertheless, authors indicate that these analyses should be interpreted cautiously because of marked heterogeneity between the study estimates and possible effect of publication bias. In fact, Li, et al. [15] have included in their systematic review and meta-analysis only 9 studies that reported an association between serum Hcy levels and RVO. They showed that
1 µmol/L increase in Hcy plasma levels was associated with a significant OR of 1.14 (95%CI, 1.07-1.21) in the random-effects model. However, the heterogeneity was higher, so no definite conclusions can be drawn.

In this scenario, the studies analyzing the association between Hcy and RVO have many methodological weaknesses, which may lead to misunderstanding this relationship. Thus, the difficulty to assess Hcy levels regarding to sample management issues (fasting status, extraction and conservation of the sample, etc.; substantial heterogeneity among studies (design, sample size, confounding variables); limitations in the selection of controls or no adequate matchinchg, and absence of population based-studies have been the main limitations found in these studies.

In a recent study from Spain, Martinez, et al. [35], analyzed the overall risk factors associated with RVO, and showed that hyperhomocysteinemia was higher in patients with RVO than in some cohorts from general population and similar to the results obtained in a cohort of venous thromboembolic disease. However, composition of control cohorts was rather heterogeneous, with some missing data for important variables, such as dyslipidemia, and above all, adjustment for confounding variables was lacking. Therefore, this study cannot be compared with ours.

We have designed a case-control study including a well-defined prospective cohort of patients with RVO and a subset of age- and sex-matched controls from the general population of our region. Besides, we were able to control for the main variables reported as confounders for the association between serum Hcy and RVO. We have found that patients with RVO have significant higher serum Hcy concentrations than controls, independently of sex, age, BMI, glomerular filtration rate, hypertension, dyslipidemia, diabetes mellitus, tobacco use and alcohol intake, and even of past history or cardiovascular or thromboembolic diseases. When stratified by sex, this finding was particularly relevant in the group of women. Furthermore, we also found that hyperhomocysteinemia (serum levels above 15 µmol/L) was associated with a 4.5-fold increase in the overall risk for developing RVO (2.6 in the case of men and 6.1 in the group of women).

We found some weaknesses in our study. Firstly, given its cross-sectional design it is not possible to determine the temporal nature of the observed relationships, for which prospective data are certainly needed. Secondly, those derived from the time interval after RVO and serum Hcy extraction. In fact, it has been pointed out that the vascular occlusive event itself could increase plasma Hcy concentration [20]. However, we have obtained the samples at least 4-5 days after the episode of RVO, and during the first 2-week, and this possible effect has therefore been minimized. Thirdly, methylenetetrahydrofolate reductase (MTHFR) C677T genotype has not been determined. Nevertheless, there was no evidence to suggest an association between homozygosity for the MTHFR C677T genotype and RVO [15].

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Conclusion

In summary, RVO may be considered as a manifestation of systemic atherosclerosis, and classical risk factors should be controlled, according to the current guidelines, to avoid relapses. We suggest that serum Hcy should be measured in RVO patients, in order to consider to treat hyperhomocysteinemia, given the few adverse effects of vitamin B12 and folate supplementation. Nevertheless, further prospective and controlled trials, with strong cardiovascular end-points, are needed to ascertain the usefulness of this therapeutic approach in these patients.

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