Letter to the Editor



Prognostic Factors in Patients with Acute Coronary Syndrome without ST-segment Elevation

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Dear Editor,

We read the article, "Prognostic factors in patients with acute coronary syndrome without ST-segment elevation" written by Santos JC¹. The authors concluded that the assessment of renal function and lymphocyte count provide potentially useful information for the prognostic stratification of patients with non-ST elevation ACS.

We know that patients with end-stage renal disease have a higher risk for and a worse outcome after myocardial infarction. However, lesser degrees of renal dysfunction also predict an adverse prognosis in patients with acute MI.

Several studies have shown the magnitude of this effect².

We also know that patients with a higher white blood cell (WBC) count – which is a marker of inflammation, have

an increased risk for adverse events, in-hospital mortality, and short- and long-term mortality after a non-ST elevation ACS, as well as for acute ST-elevation MI³. The value of the baseline WBC in patients with non-ST-elevation MI or unstable angina was evaluated in the TACTICS-TIMI 18 trial⁴, which concluded that higher WBC was associated with significantly lower TIMI flow grades, myocardial perfusion grades and more extensive coronary disease. This was found in relation to WBC but not to lymphocytes, because many factors in the ICU environment may lead to lymphocytopenia (trauma, hemorrhage, viral infections, etc.). Therefore, in order to establish a relationship between lymphocyte count and prognosis, further evaluation and objective evidence are needed.

Keywords

Acute Coronary Syndrome; Inflammation; Myocardial Infarction, Prognosis, Lymphopenia.

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Manuscript received 04/15/13; revised manuscript received 04/17/13; accepted 06/20/13.

DOI: 10.5935/abc.20130198

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Letter to the Editor

Reply

As regards Dr. İşcen's comments, we have some points to consider.

All patients diagnosed with unstable angina (UA) or non-ST segment elevation (STE) acute myocardial infarction (AMI) admitted to our hospital from Janurary to December 2010 were included in our study. In an attempt to eliminate possible confounders, patients presenting with malignant neoplasias, infectious diseases, autoimmune diseases, and those who had recently suffered a trauma or undergone surgery – conditions which could lead to leukocytosis, neutrophilia or lymphopenia, were excluded. The inclusion and exclusion criteria were described in the respective article¹.

We verified that complete white blood cell and neutrophil counts in our sample were not able to discriminate patients at a higher risk for adverse events (7781 \pm 3252 /mm³ vs. 8140 \pm 2835 /mm³, p = 0.5; and 5653 \pm 3058 /mm³ vs. 5220 \pm 2496 /mm³, p = 0.4, respectively). However, the logistic regression analysis showed an independent and significant relationship between lymphocyte count and

combined events (OR: 1.02; 95%CI: 1.01-1.04; p = 0.03). Some studies have suggested that the count of leukocytes and their subpopulations may predict fatal and non-fatal outcomes in patients with non-STE acute coronary syndrome^{2,3}. Cannon et al⁴ and Núñez et al⁵ demonstrated that the relationship between leukocytosis, neutrophilia and a worse prognosis is valid for patients with AMI, but not for those with UA. Lloyd-Jones et al⁶ and Zouridakis et al⁷ found that only lymphopenia, among the subpopulations of white blood cells, was associated with future cardiac events in these patients. Since, in our article¹, two thirds of our sample was comprised of patients with UA, we suggested that the lymphocyte count in this population probably provides a better identification of patients with a worse prognosis.

Sincerely, Jessica C M D'Almeida Santos Mário de Seixas Rocha Marcos da Silva Araújo

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