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# The Relationship Between Type D Personality and the Complexity of Coronary Artery Disease

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**Purpose:** The relationship between personality traits and cardiovascular disease has gathered sustained interest over the last years, type –D personality (TDP) being significantly associated with coronary artery disease (CAD). However, data regarding the connection between the TDP and the severity of CAD disease is scarce. The aim of our study was to assess the relationship between TDP and the complexity of CAD, and to compare it with other sociodemographic and clinical features.

**Patients and Methods:** We conducted a cross-sectional case-control clinical-based study on 221 consecutive hospitalized patients with chest pain ( $60 \pm 10.2$  years; 131 men), referred for coronary angiography.

**Results:** TDP was identified in 42 (19%) patients, using the DS 14 scale. Symptomatology profile was evaluated using the SCL-90 scale. Syntax score was greater in the subgroup of patients with TDP in comparison to non-TDP subgroup ( $26.21 \pm 12.03$  vs  $15.49 \pm 8.89$ , respectively,  $p < 0.001$ ), and most of SCL-90 symptom dimensions have significantly higher levels in the subgroup of TDP with CAD patients (all  $p < 0.05$ ). Smoking ( $\beta = 0.132$ ,  $p = 0.037$ ), dyslipidemia ( $\beta = 0.149$ ,  $p = 0.013$ ), Diabetes Mellitus ( $\beta = 232$ ,  $p < 0.001$ ), NA dimension of TDP ( $\beta = 0.255$ ,  $p < 0.001$ ) and SI ( $\beta = 0.279$ ,  $p < 0.001$ ) dimension of TDP have a significant contribution to the complexity of CAD assessed by Syntax score.

**Conclusion:** TDP was associated with a more complex CAD assessed by Syntax score, and may represent a dynamic interface between the biological and psychological vulnerabilities and the symptoms of CAD.

**Keywords:** coronary artery disease, type D personality, Syntax score

## Introduction

Cardiovascular diseases continue to represent the leading cause of death worldwide, with being responsible for 45% of all deaths in Europe. Moreover, coronary heart disease by itself accounts for about 20% of all deaths in Europe annually, thereby surpassing even breast cancer in women and lung cancer in men.<sup>1,2</sup>

The relationship between cardiac diseases and different psychological features of the affected individuals should be regarded as more complex and bidirectional. Recently, the advances in research of cardiac diseases have led to the conclusion that depression should be recognized as a risk factor for poor prognosis in patients with acute coronary syndrome.<sup>3</sup> Thus, it was the first instance when a psychological factor was weighted to the same extent as other previously known traditional risk factors such as hypertension, family history, hypercholesterolemia, diabetes mellitus (DM) and smoking.<sup>4,5</sup>

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On the other hand, some of the personality traits that are more stable could play a more significant concurrent pathogenic role over time in coronary heart disease, rather than depression, which is considered as a temporary altered state of mood. Since the 1950s, there have been concerns related to identifying particular personality profiles and to correlate them with certain types of chronic medical conditions. Hence, Friedman and Rosenman's contribution is a turning point concerning research on the role of personality in CAD.<sup>6</sup> More recently, the type-D or "distressed" personality concept was conceived by Denollet to detect those traits that place individuals at increased risk of CAD. The type D personality (TDP) consists of two sub-dimensions, namely social inhibition (SI) and negative affectivity (NA), where NA refers to the tendency to experience feelings of dysphoria, anxiety, and irritability, while SI refers to the tendency to inhibit the expression of emotions or behaviors in social interactions to avoid disapproval by others. It has been noticed that the TDP is significantly associated with cardiovascular diseases, but also with diabetes mellitus (DM).<sup>7</sup> The TDP has elicited the researcher's interest, both as a vulnerable terrain for the heart disease occurrence and as a worsening factor that negatively impacts health outcomes of cardiovascular diseases, and especially outcomes following myocardial infarction.<sup>8,9</sup> However, a few studies have found no associations between Type D personality and adverse cardiovascular outcomes.<sup>10,11</sup>

The aim of our study was to assess the relationship between TDP and the complexity of CAD, and to compare it with other sociodemographic and clinical features, assessing the hypothesis that TDP could negatively impact CAD reflected by the Syntax score.

## Patients and Methods

### Procedure

A cross-sectional case-control clinical-based study was conducted in consecutive patients with chest pain and age of at least 18 referred to the Institute of Cardiovascular Diseases Timisoara for coronary angiography according to current guidelines: disabling stable angina pectoris (CCS III–IV) despite medical therapy, high-risk criteria on noninvasive testing, angina in conjunction with signs of heart failure, recurrence of angina after PCI/CABG, survivors of cardiac arrest, severe ventricular arrhythmias, inconclusive or conflicting results after noninvasive stress testing, unable to

undergo noninvasive testing (disability, illness, or morbid obesity), acute coronary syndrome.

All study participants were on appropriate medical therapy and underwent anamnesis, clinical examination, 12-lead electrocardiogram, blood samples, transthoracic echocardiogram, and coronary angiography, followed by psychological tests. The study was approved by Ethics Commission in Research - Development of the Institute of Cardiovascular Diseases Timisoara (registration number: 1658/28 March 2014). This study was conducted in accordance with the Declaration of Helsinki. Informed written consent was obtained from all patients.

### Sociodemographic, Case History and Major Risk Factors

Sociodemographic data were collected as follows: current age, gender, educational level, residency, professional and marital status. Concerning case history data and biological data, the heredity for cardiovascular diseases, hypertension, hypercholesterolemia, diabetes mellitus, smoking and unarmful current alcohol drinking (a drink is equivalent with 14 grams of pure alcohol; so, moderate drinking is defined as up to 1 drink per day for women and up to 2 drinks per day for men), were recorded according to current guidelines.<sup>12,13</sup>

### Echocardiography

Echocardiography was performed with a Vivid 9 ultrasonographic system (General Electric, Milwaukee, WI), in accordance with guidelines.<sup>14,15</sup> All images were digitally stored and analyzed off-line with EchoPac PC Dimension software (GE Medical). Left ventricular ejection fraction (LVEF) was calculated from apical two- and four-chamber views using LV volumes by the modified biplane Simpson rule.<sup>14</sup> Peak early (E) and E-velocity deceleration time were measured from the LV-inflow pattern at the tips of the mitral valve at end-expiratory time.<sup>15</sup>

Peak systolic ( $S'$ ) and peak early diastolic velocities ( $E'$ ) were obtained at the septal and lateral sites of the mitral annulus by pulse wave tissue Doppler imaging from apical four-chamber views. All velocities were recorded for three consecutive cardiac cycles during end-expiratory apnea, and the results were averaged. The  $E/E'$  ratio using the average of the septal and lateral  $E'$  was calculated.<sup>16,17</sup>

Speckle tracking echocardiography was also used for myocardial deformation measurements. Longitudinal peak systolic strain was determined from apical planes (four-,

three- and two-chamber view) in a 17 LV segment model. Value of global longitudinal strain (GLS) was obtained by averaging all 17 LV segments.<sup>18,19</sup> Off-line analysis was performed by two observers, who were blinded to clinical data.

## Coronary Angiography

Coronary angiography results were evaluated by an interventional cardiologist, who was blinded to the psychological tests. The complexity of CAD was determined by the SYNTAX score, a tool which takes into consideration the anatomical features of the coronary lesions.<sup>20,21</sup> The SYNTAX score is the sum of the points assigned to each individual lesion identified in the coronary tree with >50% narrowing in the diameter of the vessels that measure greater than 1.5 mm.

## Type D Personality

The DS 14 scale was used to assess TDP. This scale comprises 14 items, each item being scored between 0 = false and 4 = true, and consists of two subscales representing two distinct dimensions of TDP (Negative Affectivity and Social Inhibition). A score  $\geq 10$  on each subscale is required to meet the D personality.<sup>22,23</sup> Moreover, the DS14 was shown to have a good reliability in an extensive cross-cultural equivalence analysis including most European countries and English-speaking countries, except Russia, with a Cronbach's  $\alpha$  between 0.80 and 0.90 for NA and between 0.74 and 0.89 for SI.<sup>24</sup>

## SCL-90 (Symptom Check List – 90)

The SCL-90 is a multidimensional self-assessment inventory consisting of 90 items that cover nine symptomatology domains, such as somatization, obsessive-compulsive, interpersonal sensitivity, depression, anxiety, hostility, phobic anxiety, paranoid ideation, and psychoticism. Each item is scored on a Likert scale varying between 0 = not at all to 4 = extremely. The SCL-90 scale is not a diagnostic tool; thus, it only measures a symptomatologic profile over the previous seven days before the self-assessment.<sup>25</sup>

## Participants

Of 250 initially eligible patients for our study, 12 denied participating in the research and 17 patients were excluded due to missing data. Finally, after the inclusion and exclusion criteria were applied, a total number of 221 hospitalized subjects were enrolled to participate in the study. Patients with positive past or current personal history for

any psychiatric disorders, including alcohol or other psychoactive use disorders, the absence of a signed informed consent, confusional state or any mental ability to provide written informed consent, inadequate echocardiographic images, paced rhythm, mitral prosthesis, congenital heart disease, renal failure (defined as a serum creatinine >2 mg/dL), thyroid disorders, acute infections, stroke, or diabetic ketoacidosis, and age below 18 were not included in the study group.

The used method of sampling to enroll patients was by convenience including consecutive patients in order to increase the sample size. Enrolled subjects were provided with the self-assessment scales, indicated in the measures section below. Patients were stratified based on the presence or absence of TDP.

## Statistical Analyses

Gender, family history of cardiovascular diseases, dyslipidemia, DM and smoking were recorded as dichotomous variables. The staged preexisting comorbid hypertension was quantified by using the ordinal variable starting with zero, signifying the lack of hypertension, and continuing with the different stages of hypertension. Current age and body mass index were treated as continuous variables. Numeric variables are presented as mean value  $\pm$  standard variation (SD) and compared by using Independent-Samples *t*-test. Descriptive statistics were done depending on two strata represented by the presence and absence of TDP. Dichotomized TDP was used in the majority of statistical analyses. According to the recommendations from other studies, the NA and SI subscales scores were treated as distinct ordinal variables only in the multiple linear regression model.<sup>26,27</sup> The Chi-squared test was used to compare the frequency of nominal variables. For continuous variables, Independent-Samples *t*-test was applied to compare means and for ordinal variables Mann-Whitney U-test was carried out to analyze mean rank differences. Partial correlations were calculated to estimate the relationship between DS 14 subscales scores and the Syntax score after controlling for anxiety and depression levels. In addition, multiple linear regression was performed to estimate the association of cardiovascular morbidity risk factors such as the family history of cardiovascular diseases, body mass index, smoking, DM, current age, gender, dyslipidemia, hypertension, NA and SI (as distinct ordinal variables) with CAD complexity reflected by Syntax score. For the statistical analyses, we used the software package SPSS version 11.5 (SPSS Inc,

**Table 1** The Frequency of TDP and the Differences of Mean Ranks for Negative Affectivity and Social Inhibition Based on the Presence or Absence of Type D Personality

Subjects Enrolled in Study, n (%)	Presence of TDP, n (%)	Absence of TDP, n (%)	Significance of Differences
221 (100%)	42 (19%)	179 (81%)	
Negative affectivity	169.17	97.35	$p < 0.001$
Social inhibition	197.37	90.73	$p < 0.001$

**Notes:** The level of significance for all analyses was set at  $\alpha = 0.05$ ; mean ranks are reported to either strata considered separately. Mann–Whitney U-test was performed.  
**Abbreviation:** TDP, type D personality.

Chicago, IL). A two-sided p-value of  $<0.05$  was accepted as statistically significant.

## Results

In our 221 consecutive hospitalized patients with chest pain ( $60 \pm 10.2$  years; 131 men), referred for coronary angiography, the final diagnosis was: acute ST-segment elevation myocardial infarction in 61 patients (27.6%), non-ST-elevation acute myocardial infarction in 25 patients (11.3%), unstable angina in 45 patients (20.3%) and stable angina associated with obstructive coronary artery disease in 51 patients (23%). Other 39 patients presented non-obstructive coronary artery diseases (21.7%) and 27 (12.2%) had a previous coronary revascularization. In our group, 148 patients' benefited of PCI (66.9%) and 34 of CABG (15.3%). The mean Syntax score in our study group was  $17.53 \pm 10.4$ .

By applying the DS 14 scale we identified 42 (19%) patients with TDP. There were significant differences between the mean ranks of NA (169.17 vs 97.35;  $p < 0.001$ ) and SI (197.37 vs 90.73;  $p < 0.001$ ) depending on the two analyzed strata (Table 1).

The sociodemographic characteristics were mostly similar between the two considered strata, making them comparable in this regard (Table 2). There was only one statistically significant difference between the means of the current age of males vs females ( $58.92 \pm 10.72$  vs  $62.12 \pm 9.49$ ,  $p = 0.023$ ). Moreover, CAD patients with TDP have tended to reside in urban regions that are probably more demanding for daily existence compared to rural areas, therefore shaping the personality profile during their psychological development.

The psychoactive substance use has not been significantly different between the non-TDP and TDP in our CAD patients. Neither current smokers (45.8% vs 52.4%, respectively,  $p = 0.443$ ) nor current alcohol users (27.4% vs 31.0%, respectively,  $p = 0.642$ ) made a difference between the compared strata.

The CAD appreciated by mean Syntax score was more complex in the subgroup of patients with TDP in comparison to non-TDP subgroup ( $26.21 \pm 12.03$  vs  $15.49 \pm 8.89$ , respectively,  $p < 0.001$ ) (Table 2).

We also analyzed the differences between the mean ranks of several distinct symptom dimensions of SCL-90 inventory depending upon the presence or absence of TDP, as shown in Table 3. Except for the obsessive-compulsive symptoms (126.73 vs 107.31, respectively;  $p = 0.076$ ), the rest of SCL-90 symptom dimensions (somatization, interpersonal sensitivity, depression, anxiety, hostility, phobic anxiety, paranoid ideation and psychoticism) have significantly higher levels in the subgroup of TDP with CAD patients (all  $p < 0.05$ ) (Table 3).

Given that, the higher level of psychopathological dimensions measured by the SCL-90 scale could have partially influenced the magnitude of the Syntax score, we calculated the partial correlations between the two dimensions of TDP with the Syntax scores after controlling for anxiety and depression scores as measured by the mentioned scale (Table 4).

Hence, both dimensions of TDP, NA ( $r = 0.178$ ,  $p = 0.008$  and  $r = 0.150$ ,  $p = 0.026$ , respectively) and SI ( $r = 0.253$ ,  $p < 0.008$  and  $r = 0.183$ ,  $p = 0.006$ , respectively) have remained statistically significant, correlated with the Syntax scores indicating a direct contribution of them.

The presence of acute myocardial infarction was significantly higher in the subgroup of patients with TDP (57.1%, 24 patients with TDP vs 34.6%, 62 patients without TDP,  $p = 0.007$ ). Table 2 also showed a higher incidence of occluded coronary vessels in patients with TDP ( $p < 0.001$ ).

The value of the echocardiographic parameters is also influenced by the presence of TDP in CAD patients (Table 5). These patients have more altered systolic dysfunction [LVEF ( $47.45 \pm 13.78\%$  vs  $51.7 \pm 9.48\%$ ,  $p = 0.06$ ),  $S'$  ( $7.05 \pm 2.57$  cm/s vs  $8.89 \pm 3.06$  cm/s,  $p < 0.001$ ) and global longitudinal strain

**Table 2** Sociodemographic, Case History and Clinical Data

Sociodemographic and Anamnesis Data	Non-TDP CAD Patients	TDP CAD Patients	Statistical Significance
n (%)	179 (81%)	42 (19%)	–
Average current age, mean (SD) <sup>a</sup>	60.08 (10.369)	60.81 (10.317)	p = 0.683
Gender – male, n (%)	107 (59.8%)	24 (57.1%)	p = 0.755
Educational level, mean ranks	111.56	108.60	p = 0.772
Residency - urban area, n (%)	110 (61.5%)	32 (76.2%)	p = 0.073
Professional status - Employed or Student, n (%)	57 (31.8%)	14 (33.3%)	p = 0.852
Marital status – with intimate partner, n (%)	148 (82.7%)	37 (88.1%)	p = 0.393
Positive family history of CVD, n (%)	24 (13.4%)	7 (16.7%)	p = 0.584
Current smokers <sup>b</sup> , n (%)	82 (45.8%)	22 (52.4%)	p = 0.443
Occasionally alcohol use (not abuse or dependence) <sup>b</sup> , n (%)	49 (27.4%)	13 (31.0%)	p = 0.642
Syntax score, mean (S.D.)	15.49 (8.892)	26.21 (12.038)	p < 0.001
The presence of myocardial infarction, n (%) <sup>c</sup>	62 (34.6%)	24 (57.1%)	p = 0.007
No. of occluded coronary vessels, mean ranks <sup>c</sup>	102.87	145.67	p < 0.001
Staged hypertension, mean ranks	108.90	119.96	p = 0.199
Dyslipidemia, n (%)	39 (21.8%)	6 (14.3%)	p = 0.277
Body mass index, mean (S.D.)	26.113 (3.1324)	26.595 (3.9177)	p = 0.394
Preexisting diabetes mellitus irrespective of type, n (%)	67 (37.4%)	14 (33.3%)	p = 0.620

**Notes:** The level of significance for all analyses was set at  $\alpha = 0.05$ ; the listed percentages and means are reported to either strata considered separately. <sup>a</sup>There was a statistically significant differences between the means of current age of males vs females [58.92 (S.D. = 10.724) vs 62.12 (S.D. = 9.494); p = 0.023]. <sup>b</sup>Based on self-reporting. <sup>c</sup>Cardiac status at the admission in Cardiac Intensive Care Unit.

**Abbreviations:** CAD, coronary artery disease; CVD, cardiovascular disease; TDP, type D personality.

**Table 3** The Differences in Symptomatology Profiles in CAD Patients Depending Upon the Presence or Absence of Type D Personality

SCL-90 Domains of Symptomatology (Mean Ranks)	Non-TDP CAD Patients	TDP CAD Patients	Statistical Significance
n (%)	179 (81%)	42 (19%)	
Somatization	104.18	140.08	p = 0.001
Obsessive-compulsive	107.31	126.73	p = 0.076
Interpersonal sensitivity	101.63	150.94	p < 0.001
Depression	98.60	163.86	p < 0.001
Anxiety	102.74	146.19	p < 0.001
Hostility	104.61	138.21	p = 0.002
Phobic anxiety	102.69	146.43	p < 0.001
Paranoid ideation	102.41	147.60	p < 0.001
Psychoticism	100.91	154.01	p < 0.001

**Notes:** The level of significance for all analyses was set at  $\alpha = 0.05$ ; the listed percentages and means are reported to either strata considered separately. Mann–Whitney U-test was performed for all comparisons of SCL-90 domains of symptomatology.

**Abbreviations:** SCL-90, Symptom Check List – 90; CAD, coronary artery disease; TDP, type D personality.

**Table 4** The Partial Correlations Between the DS 14 Subscales Score and Syntax Score After Controlling for Anxiety and Depression Levels as Measured by SC-90 Scale

DS 14 Subscales Scores	Syntax Score	
	Controlling for Anxiety Levels as Measured on SCL-90 Scale	Controlling for Depression Levels as Measured on SCL-90 Scale
Negative affectivity	$r = 0.178$ ; $p = 0.008$	$r = 0.150$ ; $p = 0.026$
Social inhibition	$r = 0.253$ ; $p < 0.001$	$r = 0.183$ ; $p = 0.006$

**Abbreviations:** DS 14, the 14-item Type D Scale; SCL-90, Symptom Check List – 90.

**Table 5** Echocardiographic Parameters Depend on the Presence or Absence of TDP in CAD Patients

Echocardiographic Parameters in CAD Subjects		CAD Patients with TDP; Mean (S.D.)	CAD Patients without TDP; Mean (S.D.)	Significance of Differences
Systolic function	Left ventricular ejection fraction (%) <sup>a</sup>	47.45 (13.788)	51.77 (9.481)	$p = 0.060$
	Peak systolic mitral A=annular velocity (cm/s)	7.053 (2.575)	8.898 (3.069)	$p < 0.001$
	Global longitudinal strain (%)	- 14.067 (5.650)	- 15.938 (4.822)	$p = 0.030$
Diastolic function	The early diastolic transmitral velocity/early mitral annular diastolic velocity ratio (E/E') <sup>a</sup>	13.498 (4.155)	10.242 (3.254)	$p < 0.001$
	Left atrial volume (mL) <sup>a</sup>	85.79 (34.436)	71.03 (26.497)	$p = 0.012$

**Notes:** The level of significance for all analyses was set at  $\alpha = 0.05$ ; The listed means are reported to either strata considered separately. <sup>a</sup>Levene's Test for Equality of Variances statistically significant.

**Abbreviations:** CAD, coronary artery disease, S.D., standard deviation; TDP, type D personality.

( $-14.06 \pm 5.65\%$  vs  $-15.93 \pm 4.82\%$ ,  $p < 0.001$ ), more altered LV diastolic function assessed by E/E' index ( $13.49 \pm 4.15$  vs  $10.24 \pm 3.25$ ,  $p = 0.03$ ) and larger LAV ( $85.79 \pm 34.4$  mL vs  $71.03 \pm 26.49$  mL,  $p = 0.012$ ) (Table 5).

The number of occluded coronary vessels and the number of patients with myocardial infarction were significantly higher represented in CAD patients with TDP. They are quite intriguing results, even more so because the major cardiovascular risk factors (heredity, smoking, staged hypertension, dyslipidemia, DM) did not differ between the two compared strata (Table 2).

Table 6 outlines the contributing role of TDP and risk factors for CAD to the Syntax score.

Male gender, age, BMI, heredity, hypertension did not reach a significant value in our study group (all  $p > 0.05$ ), while smoking ( $\beta = 0.132$ ,  $p = 0.037$ ), dyslipidemia ( $\beta = 0.149$ ,  $p = 0.013$ ), Diabetes Mellitus ( $\beta = 0.232$ ,  $p < 0.001$ ), NA dimension of TDP ( $\beta = 0.255$ ,  $p < 0.001$ ) and SI ( $\beta = 0.279$ ,  $p < 0.001$ ) dimension of TDP have a significant contribution to the complexity of CAD assessed by Syntax score.

## Discussion

In the present study, we analyzed the relationship between TDP and CAD complexity assessed by Syntax score in patients referred for coronary angiography. We demonstrate for the first time a more complex CAD in patients with TDP, irrespective of the anxiety and depression symptoms influences.

To date, numerous studies have revealed different psychological paradigms, including behavioral patterns, underpinning the connection between the TDP and the severity of CAD disease.<sup>8,9,28–30</sup> Our results are inconsistent with some previous studies, in which such a relationship was not traced. Probably, the reason is that in previous studies homogeneous groups of patients were examined (before CABG, PCI, during cardiac rehabilitation, etc.), and in the present study, the surveyed cohort is heterogeneous. We included all the patients with chest pain referred for coronary angiography after noninvasive evaluation. For diagnostic cath procedures, 12.5% of patients had a prior coronary revascularization, in comparison with 25% as reported in the CathPCI Registry.<sup>31</sup> In

**Table 6** The Contributing Role of TDP and Risk Factors for CAD to the Syntax Score

Risk Factors	Beta Standardized Coefficient	SE	p value	95% C.I. for B Unstandardized Coefficient
Male gender	-0.104	1.333	0.100	-4.829-0.425
Current age	0.114	0.062	0.065	-0.007-0.237
Body Mass Index	-0.027	0.183	0.640	-0.447-0.275
Smoking	0.132	1.317	0.037	0.166-5.357
Positive family history of CVD	0.027	1.758	0.641	-2.645-4.286
Hypertension (staged)	0.026	0.676	0.665	-1.039-1.625
Dyslipidemia	0.149	1.527	0.013	0.834-6.855
Preexisting Diabetes Mellitus (DM)	0.232	1.290	<0.001	2.467-7.552
NA dimension of type D personality	0.255	0.112	<0.001	0.213-0.655
SI dimension of type D personality	0.279	0.147	<0.001	0.346-0.924

**Notes:** Model summary (adjusted R square = 0.286; R square change = 0.318; F change = 9.809; significant F change <0.001).

**Abbreviations:** SE, standard error; CVD, cardiovascular disease; NA, negative affectivity; SI, social inhibition; DM, diabetes mellitus, irrespective of type; TDP, type D personality.

our series 27.6% patients presented acute ST-segment elevation myocardial infarction (versus 16% in CathPCI Registry), 11.3% non-ST-elevation acute myocardial infarction (versus 9% in CathPCI Registry), 20.3% unstable angina (33% in CathPCI Registry) and 23% stable angina (21% in CathPCI Registry). The percent of patients with non-obstructive coronary artery diseases was similar with this one from CathPCI Registry (21.7% vs 18%).

## Behavioral Patterns

It has been shown that the TDP measured by using the DS 14 scale was significantly associated with cardiovascular diseases.<sup>8,9</sup> However, the increasing comorbidity of major and minor depression with cardiovascular diseases has provided for sufficient research to investigate the mediating effect of depression on this category of physical diseases in individuals having a TDP.<sup>14,32-34</sup> In a previous study, TDP was correlated in patients with CAD with more unhealthy behaviors and less healthy behaviors, while controlling for anxiety and depression. These results are in line with our findings according to which TDP in CAD patients have had slightly more frequent addictive behaviors such as nicotine and alcohol use compared to non-TDP subset of CAD patients. Moreover, TDP may be closely related to the unfavorable evolution of CAD by engaging other unhealthy behaviors, such as low adherence to treatment, propensity for dysfunctional coping

styles and resistance to lifestyle changing or post-event cardiac rehabilitation.<sup>12,30,35,36</sup>

At the same time, the TDP has been studied concerning its capacity to exert a negative effect on the quality of life in cardiovascular patients. Indeed, the TDP may impair health-related quality of life in CAD patients by mediating the role of anxiety and depressive overlapping symptoms.<sup>30</sup> Thus, one study has found that TDP was significantly associated with poor self-perceived symptoms in patients with mild coronary abnormalities, which can be supposed to be deleterious over the long term.<sup>35-40</sup> A dyadic cross-sectional study has revealed that the TDP and perceived control were closely related to the health-related quality of life in both cardiac arrest survivors and their spouses.<sup>41</sup> The social support and coping style were shown to be hypothetically underlying mechanisms that can, in part, explain the relationships between TDP and physical symptoms, and TDP and perceived stress, respectively.<sup>42</sup>

In our study, the percent of CAD patients that fulfilled the criteria for TDP was similar to those found in other research, meaning that about one in five patients has the pre-existing maladaptive personality traits.<sup>27,28</sup> The socio-demographic characteristics were mostly similar between the two considered strata, making them comparable in this regard. However, CAD patients with TDP have tended to reside in urban regions that are probably more demanding for daily existence compared to rural areas, therefore

shaping the personality profile during their psychological development. Other large representative research has found that different types of personality disorders, classified according to DSM III – R, were more frequent in the urban areas than living on the outskirts.<sup>43</sup>

## Psychopathological Aspects

Concerning the higher levels of general psychopathological dimensions that we have found in CAD patients with TDP, including depression and anxiety, these findings have already been evidenced in other researches.<sup>30,32,33</sup> Therefore, the assumption that TDP negatively impacts the health outcomes of CAD patients, not singularly, but in a broader context of psychosocial factors clustered within the same individual is quite feasible.<sup>36</sup> Nonetheless, considering our results even after controlling for anxiety and depressive levels, the TDP has been significantly associated with the complexity of CAD assessed by Syntax score. On the upside, the propensity for maladaptive coping styles that have been documented by other researches, in conjunction with higher levels of psychopathologies, might support our result whereby the perceived stressful life events are significantly more self-reported by TDP in CAD patients in contrast to non-TDP controls.<sup>35,44</sup> It was the only anamnesis data revealed by our research that made the difference between the two compared subsets of CAD patients.

Regarding the negative influences of TDP on the cardiac status of CAD patients, several overlapped psychophysiological mechanisms may support this negative outcome. Given its innate component represented by temperament, which is mainly originating in cerebral neurotransmission profile, TDP may continuously alter the homeostasis of specific physiological parameters.<sup>45</sup> Intermittently, against the background of the continuous influence, the physiological imbalances can be amplified by the interposing of the anxiety and depressive symptoms, which in turn, are promoted by the presence of TDP.<sup>36</sup> Therefore, the mind–heart interactions may be influenced by several neuroendocrine and neurobiological pathways such as the hypothalamic–pituitary–adrenal and sympathoadrenal medullary axes dysfunction, chronic low-grade inflammation, endothelial cell dysfunction, neurotransmitters imbalances, altered level of neurotrophic factors and the disconnectivity of cortico-subcortical neural networks.<sup>46,47</sup> Higher comorbidity between distinct physical illnesses such as dyslipidemia, hypertension, DM, and obesity, and CAD and stress-related conditions,

including TDP, may also be impacted by the above-mentioned pathophysiological processes.

## Biological Substrate

From the biological standpoint, the TDP stems from the imbalance of the cerebral neurotransmitters reflected by altered levels of specific metabolites, such as tryptophan metabolite kynurenine.<sup>13</sup> It has been noticed that restricted sleep mediates the connection between TDP and a low or blunted cardiovascular response to the social stressors, increasing the risk for a range of adverse health outcomes even in healthy individuals.<sup>14,15</sup> Other research data have shown that TDP is more associated with endothelial dysfunction and inflammation plasma biomarkers.<sup>16–18</sup> Also, some studies have indirectly tied the TDP to coronary heart diseases through the intermediate action of dyslipidemia.<sup>19–21</sup>

TDP individuals may be subject to a biased interpretation of their health status due to dysfunctional cognitive schemas that underlay the negative affective experiences through the erroneous interpretation of objective reality, leading to a more amplified negative affectivity and autonomic over activity.<sup>48</sup> In a clinical study, the presence of TDP was independently associated with the occurrence of ventricular arrhythmias as a marker of sympathetic dysregulation.<sup>49</sup> In both healthy individuals and CAD patients, the presence of TDP has been associated with increased diurnal cortisol release.<sup>50,51</sup>

Intriguing results were attained regarding the current or past myocardial infarction and the number of occluded coronary vessels that were significantly higher represented in CAD patients with TDP. The more so because all the major risk factors like heredity, smoking, staged hypertension, dyslipidemia, body mass index, and DM did not differ between the two compared strata. Recently, research have repeatedly evidenced that atherosclerosis, especially CAD, are no longer considered as a simple lipid storage disorder. These might also be considered chronic inflammatory diseases, where increased levels of TNF- $\alpha$ , other cytokines and increased IFN- $\gamma$  transcription factors will lead to endothelial cell dysfunction and atherosclerotic plaques formation.<sup>52–55</sup>

Besides cytokines, the serum high-sensitivity C-reactive protein has been found as a highly predictive factor of cardiovascular risk, irrespective of the low-density lipoprotein cholesterol level.<sup>56</sup> Low-grade chronic inflammation intercedes in all phases of the atherosclerosis process, starting with the accumulation of macrophage transformed into foam cells in the arterial wall continuing

with the fatty streak and fibrous plaque formation and ending with plaque fissuring, rupture, and thrombosis, all leading to an acute coronary event. It has been shown that atherosclerosis is accompanied by an excessive inflammatory response resulting in fibroproliferative reaction at the level of subintimal arterial space promoting subsequent thrombus formation as a direct consequence of various forms of injuries leading to an acute cardiac event.<sup>57,58</sup> Furthermore, traditional major cardiovascular risk factors (eg, smoking, hypertension, hyperglycemia, obesity or insulin resistance), together with stress-related psychopathological conditions, have the propensity to induce chronic inflammation resulting in endothelial activation through impaired nitric oxide production and loss of vasodilator and antithrombotic properties of the coronary endothelium.<sup>57,58</sup> Similarly, research has revealed that the heart is an active endocrine organ and a series of endocrine molecules (generically named cardiokines) are released by cardiomyocytes, fibroblasts, vascular endothelial cardiac cells and immune cells that reside in the heart.<sup>59</sup> Moreover, the immune functions at the heart level are also regulated by the autonomous nervous system through the neuro-cardiac axis by its two components. The sympathoadrenal axis stimulates immunological response mainly by the effect of NE on resident macrophages in addition to that on circulating monocytes. In opposite, the parasympathetic nerves mediate acetylcholine release both in the reticuloendothelial system and in the heart, where the neurotransmitter binds the nicotinic receptor expressed on resident cardiac macrophages, blocking the release of the main proinflammatory cytokines (ie, TNF- $\alpha$ , IL-1).<sup>60</sup> The TDP and CAD may partially share the same cascade of overactivated pro-inflammatory cytokines alongside the imbalanced functioning of sympathetic and parasympathetic, respectively, components of the autonomous nervous system. Altogether, these biological mechanisms may result in a worsened prognosis of the TDP CAD patients.<sup>9</sup>

It seems that specific and subtle immunological pathways can connect the stress-related disorders, such as depression, anxiety, or even negative affectivity component of TDP with cardiovascular diseases.

All converging biological and psychological mechanisms mentioned above have determined that TDP, alongside the major depression, has been included in the European Cardiovascular Prevention guideline as a risk factor to screen for atherosclerosis.<sup>61,62</sup> Based on a holistic perspective, the mixture of physical and psychological manifestations of CAD could be entirely integrated

into a systemic nosological concept where TDP is one of the stable parts of a whole syndrome entity. In these particular cases, amid the chronic manifestations of CAD patients, acute exacerbations periodically occur in both planes, psychologically mainly expressed as contextual symptoms of anxiety and depression, and physically marked by biological deteriorations of the cardiac parameters. Therefore, TDP should be considered as a critical psychological marker of a more unfavorable clinical course in CAD patients, characterized by a higher likelihood of negative outcomes and cardiac events and a poorer quality of life compared to their non-TDP counterparts.

Our results should be considered in the context of several limitations. The study population of patients undergoing coronary angiography was heterogeneous. Since this study is a cross-sectional design, the direction of the mediation between TDP and depressive symptoms cannot be confirmed. Despite the good response rate (95.2%), the non-consenting patients may have refused consent because of a higher level of depressive symptoms, leading to some bias in the results. This work was performed on a relatively small group of patients, and does not provide longer-term follow-up of the clinical implications of TDP. It is a single center study and its reproduction in other centers or by multicenter studies could support its validity.

## Conclusion

In patients referred for coronary angiography, TDP was associated with a more complex CAD assessed by Syntax score. Given its stability, TDP may be viewed as a dynamic interface between the underlying biological and psychological vulnerabilities and the bodily and psychological symptoms of CAD.

In other news, an interdisciplinary and personalized based therapeutic approach of patients suffering from CAD is more than necessary considering the psychological profile of each patient. Beyond the quality of life improvement, in well-defined cases, the psychological intervention, at times, combined with psychopharmacological treatment might result in extending the survival period in CAD patients with TDP by targeting the altered psychophysiological pathways. Further larger and prospective studies, including more biological, clinical and anamnestic data, are absolutely required.

## Disclosure

The authors report no conflicts of interest in this work.

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