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Letter to the editor

The lack of association between catechol-*O*-methyl-transferase Val108/158Met polymorphism and smoking in schizophrenia and alcohol dependence

Matea Nikolac^a, Marina Sagud^b, Gordana Nedic^a, Korona Nenadic Sviglin^c, Alma Mihaljevic Peles^b, Suzana Uzun^d, Bjanka Vuskan Cusa^e, Oliver Kozumplik^d, Maja Zivkovic^f, Maja Mustapic^a, Miro Jakovljevic^b, Mladen Pavlovic^g, Dorotea Muck-Seler^a, Fran Borovecki^h, Nela Pivac^{a*}

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*Corresponding author: Nela Pivac, Division of Molecular Medicine, Rudjer Boskovic Institute, Bijenicka 54, 10000 Zagreb, Croatia; Tel: ++385 1 4571207; Fax: ++385 1 4561010. E-mail: npivac@irb.hr Patients with schizophrenia and alcohol dependence are often smokers. There are inconsistencies across the literature data on the relationship between the functional catechol-*O*-methyl-transferase (COMT) Val108/158Met polymorphism and smoking and/or nicotine dependence (Redden et al., 2005; Tammimaki and Mannisto, 2010).

The study elucidated the association between the COMT Val108/158Met polymorphism and smoking in 828 ethnically homogenous non-related Caucasian patients with schizophrenia $(40.3\pm12.6 \text{ years old})$, 616 patients with alcohol dependence $(49.7\pm10.2 \text{ years old})$, diagnosed using a Structured Clinical Interview based on DSM-IV criteria, and in 1058 healthy control subjects $(40.5\pm15.3 \text{ years old})$. The COMT Val108/158Met polymorphism was genotyped using a Taqman-based allele-specific polymerase chain reaction assay (Applied Biosystems, Foster City, CA, USA).

COMT genotypes in the studied groups were in the Hardy-Weinberg equilibrium. The present smoking status differed significantly (χ^2 =327.71; *P*<0.001) between the studied groups, since male schizophrenic patients smoked more frequently, while healthy control women smoked less frequently than other subjects. The stepwise logistic regression (*OR*=1.56, 95% *CI*=1.10-2.23, *P*=0.014) and the χ^2 test (Table 1) revealed that COMT Val/Val genotype was significantly associated with smoking in healthy male subjects, since carriers of one (χ^2 =5.686; *P*=0.017) or two (χ^2 =9.783; *P*=0.008) Val alleles were more frequently current smokers.

The significant association between smoking, age and gender, with no interaction between smoking and COMT Val108/158Met variants in patients with schizophrenia and alcohol dependence might be explained by the different effects of nicotine on social and interpersonal factors in mental versus healthy subjects, or by the different interactions between dopamine activity and nicotine in the regions involved in the regulation of the reward pathways in healthy opposed to subjects with schizophrenia and alcohol dependence. Although the hypothesis of the study was that COMT Val108/158Met genotypes will be significantly associated with smoking in schizophrenia and alcohol dependence, our study, with sufficiently large sample size (N=2502) and power (higher than 0.800), matched for ethnicity and gender, showed for the first time the lack of significant association between smoking and COMT Val108/158Met polymorphism in schizophrenia, and confirmed no association between smoking and COMT Val108/158Met in alcoholism (Foroud et al., 2007). In line with previous results obtained in smaller groups (Nedic et al., 2010), COMT Val108/158Met polymorphism was significantly

associated with smoking in healthy male subjects, due to the over-representation of the Val/Val genotype in healthy male current smokers.

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COMT Val ^{108/158} Met genotype					
Male subjects			Female subjects		
Met/Met	Val/Met	Val/Val	Met/Met	Val/Met	Val/Val
N (%)	N (%)	N (%)	N (%)	N (%)	N (%)
56	93	69 ^a	15	39	15
(25.7)	(42.7)	(31.6)	(22.4)	(55.2)	(22.4)
108	267	111	78	134	73
(22.2)	(54.9)	(22.9)	(28.6)	(45.0)	(26.4)
104	185	93	29	58	36
(27.2)	(48.4)	(24.4)	(23.6)	(47.1)	(29.3)
47	87	42	31	68	48
(26.7)	(49.4)	(23.9)	(21.1)	(46.3)	(32.6)
75	154	72	23	30	15
(24.9)	(51.2)	(23.9)	(33.8)	(44.1)	(22.1)
52	103	37	14	27	14
(27.1)	(53.6)	(19.3)	(25.4)	(49.2)	(25.4)
	Met/Met N (%) 56 (25.7) 108 (22.2) 104 (27.2) 47 (26.7) 75 (24.9) 52	Male subject Met/Met Val/Met N (%) N (%) 56 93 (25.7) (42.7) 108 267 (22.2) (54.9) 104 185 (27.2) (48.4) 47 87 (26.7) (49.4) 75 154 (24.9) (51.2) 52 103	Male subjects Met/Met Val/Met Val/Val N (%) N (%) N (%) 56 93 69 ^a (25.7) (42.7) (31.6) 108 267 111 (22.2) (54.9) (22.9) 104 185 93 (27.2) (48.4) (24.4) 47 87 42 (26.7) (49.4) (23.9) 75 154 72 (24.9) (51.2) (23.9) 52 103 37	Male subjects Fe Met/Met Val/Met Val/Val Met/Met N (%) N (%) N (%) N (%) 56 93 69 ^a 15 (25.7) (42.7) (31.6) (22.4) 108 267 111 78 (22.2) (54.9) (22.9) (28.6) 104 185 93 29 (27.2) (48.4) (24.4) (23.6) 47 87 42 31 (26.7) (49.4) (23.9) (21.1) 75 154 72 23 (24.9) (51.2) (23.9) (33.8) 52 103 37 14	Male subjects Female subj Met/Met Val/Met Val/Val Met/Met Val/Met N (%) N (%) N (%) N (%) N (%) 56 93 69 ^a 15 39 (25.7) (42.7) (31.6) (22.4) (55.2) 108 267 111 78 134 (22.2) (54.9) (22.9) (28.6) (45.0) 104 185 93 29 58 (27.2) (48.4) (24.4) (23.6) (47.1) 47 87 42 31 68 (26.7) (49.4) (23.9) (21.1) (46.3) 75 154 72 23 30 (24.9) (51.2) (23.9) (33.8) (44.1) 52 103 37 14 27

Table 1. COMT Val108/158Met genotype counts and frequencies in male and female control

 subjects, patients with schizophrenia or alcohol dependence, further subdivided according to the

 smoking status into smokers and non-smokers.

^aP = 0.008 vs. healthy male non-smokers (χ^2 test).

COMT: catechol-O-methyl transferase; Met: methonine; Val: valine; N: genotype count.