A role of BDNF polymorphism in age-related recognition memory processes?

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INTRODUCTION

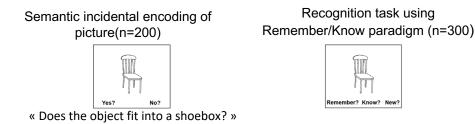
Episodic memory difficulties are frequently encountered in normal aging. However, the effect of brain-derived neurotrophic factor (BDNF) genetic polymorphism on decreased memory performance remains largely unclear. BDNF has a significant involvement in synaptic growth. Consequently, we have examined the concurrent effect of BDNF genetic polymorphism on recognition performance in young and old adults.

PARTICIPANTS & METHODS

Demographics			Genet	ic Variability		
	Young (N=51)	Old (N=60)		Val/Val	Met/Met Val/Met	Total
Sex (M/F)	29/22	29/31	Young (M+F)	28	22	50
Age	23.73 (3.07)	65.85 (4.47)	Old (M+F)	32	24	56
NSC	15,0 (1.93)	14.90 (2.67)	Total	60	46	106

Our participants were selected form a larger community dwelling cohort, which completed an episodic memory task in an fMRI setting.

Image acquisition: fMRI, 3T Siemens and Allegra scanners



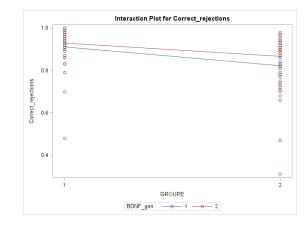


easy condition: item presented twice

hard condition: item presented once

RESULTS

Regression analyses on correct rejections



Note. 1: Young, 2: Old

	Estimate (SD)	F (df)	Р		Estimate (SD)	F (df)	Р
Group (young)	01 (.03)	0.15 (1,102)	.0003	Group (young)	03 (.01)	9.81 (1,102)	.002
BDNF (val/val)	04 (.02)	0.03 (1,102)	.13	BDNF (val/val)	.01 (.01)	0.95 (1,102)	.33
Interaction effect	.06 (.04)	.01 (1,102)	.50	Interaction effect	001 (.02)	.0 (1,102)	.95

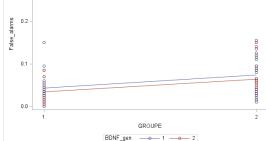
DISCUSSION

The absence of main or interaction genetic effects seems to indicate that the BDNF polymorphism does not influence recognition memory performance in aging. We will next investigate if specific brain activity patterns associated to compensatory processes are specifically observed in Val or Met carriers.





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Regression analyses on false alarms

Interaction Plot for False alarms

Note. 1: Young, 2: Old