Open Access Full Text Article

REVIEW

Neuropsychology of eating disorders: 1995–2012

Ignacio Jáuregui-Lobera

Nutrition and Bromatology, Pablo de Olavide University, Seville, Spain

Abstract: Eating disorders are considered psychiatric pathologies that are characterized by pathological worry related to body shape and weight. The lack of progress in treatment development, at least in part, reflects the fact that little is known about the pathophysiologic mechanisms that account for the development and persistence of eating disorders. The possibility that patients with eating disorders have a dysfunction of the central nervous system has been previously explored; several studies assessing the relationship between cognitive processing and certain eating behaviors have been conducted. These studies aim to achieve a better understanding of the pathophysiology of such diseases. The aim of this study was to review the current state of neuropsychological studies focused on eating disorders. This was done by means of a search process covering three relevant electronic databases, as well as an additional search on references included in the analyzed papers; we also mention other published reviews obtained by handsearching. Keywords: eating disorders, anorexia nervosa, bulimia nervosa, binge-eating disorder, neuropsychology, cognitive performance

Introduction

Neuropsychology studies the structure and function of the brain as far as they are related to specific psychological processes and behaviors. It is considered a clinical and experimental field of psychology, the aim of which is to study, assess, understand, and treat behaviors directly related to brain function.¹ Neuropsychology uses psychological, neurological, cognitive, behavioral, and physiological principles, techniques, and tests in order to evaluate patients' neurocognitive, behavioral, and emotional strengths and weaknesses without ignoring their relationship to normal and abnormal central nervous system functioning.²

Eating disorders (ED) are serious psychiatric pathologies. They are characterized by a pathological concern with body shape and weight above all. The lack of progress in treatment development, at least in part, reflects the fact that little is known about the pathophysiologic mechanisms that account for the development and persistence of ED. In contrast to the slow progress in understanding ED, basic knowledge of the neural basis of behavior has advanced rapidly in recent years, and this knowledge has begun to yield a better understanding of other serious mental illnesses.³ The possibility that there is a dysfunction of the central nervous system in patients with ED has been explored in several ways, including studies of neuropsychological test performance. Thus, the study assessing the relationship between cognitive processing and certain eating behaviors has been conducted, aiming to achieve a better understanding of the pathophysiology of ED.4

Correspondence: Ignacio Jáuregui-Lobera Nutrition and Bromatology, Pablo de Olivade University, Fernando IV 24-26 (bajo), Seville 41011, Spain Tel +34 954 280 789 Fax +34 954 278 167 Email ignacio-ja@telefonica.net

Dovencess http://dx.doi.org/10.2147/NDT.\$42714

submit your manuscript | www.dovepress.con

⁴¹⁵ © 2013 Jáuregui-Lobera, publisher and licensee Dove Medical Press Ltd. This is an Open Access article which permits unrestricted noncommercial use, provided the original work is properly cited.

The specific pathophysiology of ED is unknown, and it is likely that different factors are involved.⁴ To date, ED have been described on the basis of overt clinical phenotypes, a method that is perhaps not effective for exploring the specific etiology of these disorders.⁵ In order to identify causal factors, new ways of studying the diseases seem to be necessary.⁶ Some authors have suggested potential new focuses, including the study of endophenotypes and disease-associated traits, that are more useful in determining the relationship between underlying genes and neuropsychological functions.^{5,7} Some researchers (eg, Cavedini et al)⁸ state that neuropsychology has yet to produce an explanatory model of ED. Nevertheless, neuropsychological explorations are being used to improve the diagnosis, to obtain better ED data, and to develop more effective therapeutic strategies.⁹

The aim of this study was to review the current state of the neuropsychological studies focused on ED.

Materials and methods Search process

The search process covered three relevant electronic databases (MEDLINE, EMBASE, and PsycINFO). The general strategy included terms related to ED and neuropsychology. Next, the Medical Subjects Headings were used as well as the Boolean operators AND/OR. The shared MeSH terms were ((("Anorexia nervosa"[MeSH]) OR ("Bulimia nervosa"[MeSH]) OR ("Binge eating disorder"[MeSH])) AND (("Neuropsychology"[MeSH]) OR ("Memory"[MeSH]) OR ("Learning"[MeSH]) OR ("Attention"[MeSH]) OR ("Perception"[MeSH]) OR ("Cognition"[MeSH]) OR ("Concept formation"[MeSH]) OR ("Neuropsychological tests"[MeSH]) OR ("Neuropsychological tests"[MeSH]) OR ("Neuropsychological tasks"[MeSH]))).

Additional searches were carried out on the references included in the papers, published reviews, and via hand searching. Literature search was limited to articles published between 1995 and 2012.

Studies meeting the following criteria were included in the review: (1) studies focused on ED (anorexia nervosa [AN], bulimia nervosa [BN], and binge-eating disorder [BED]) and neuropsychology; and (2) controlled trials and randomized controlled trials. Applied exclusion criteria included: (1) descriptive studies or case reports and crosssectional studies; (2) interventions targeting populations with unspecified eating disorders (other than binge eating disorder); (3) participants with severe comorbidities; and (4) unavailable full text. Reviews and meta-analyses that fit the inclusion criteria were considered as other sources of articles.

The initial search yielded 129 references. These were combined in an EndNote 9 (Thomson Reuters, Carlsbad, CA, USA) library and screened on the basis of title and abstract; those clearly not meeting the review criteria were excluded as were duplicates. Thereafter, selected references were screened based on the full text. Reasons for exclusion were applied and 57 studies were finally included.

Selected studies are summarized in ascending order of publication year as well as with respect to the main diagnostic implied (Tables 1 and 2). Data extracted included journal reference, number of participants and age at enrollment, sex, tests, follow-up duration (when appropriate), and main outcome measurements related to the neuropsychology of ED.

Procedure

Taking into account previous recommendations,¹⁰ the content of the selected studies was analyzed considering the following functions: attention, memory and learning, visual perception/visuospatial ability, executive functions, and other functions. In addition, the analysis was based on each diagnostic as follows: AN and/or BN and/or BED.

Results AN

Attention

Considering sustained attention (attention maintained over time), Green et al did not find differences between AN patients and nonclinical participants with respect to attentional focus and the ability to maintain attention. They did so by means of a focused attention task as a measure of the Eriksen effect.¹¹ With regard to selective attention (intentional, focused attention), the Stroop Test, in its modified version (Emotional Stroop), and a wordrecognition test did not enable the authors to confirm specific cognitive deficits in AN patients.¹² Following this emotional Stroop paradigm, significant main effects of group (patients versus controls) and condition (xxxx [words made of xxxxs] neutral, fat, thin), and a significant interaction between group and condition have been reported. Patients with AN seem to have attentional bias to "fat" and "thin" words.13 With a modified color-naming Stroop task, AN patients, but not unrestrained or restrained eaters, have shown delayed color-naming latencies for both thin and fat word categories and, to a lesser extent, for high-caloric-density food words.¹⁴ With a similar attentional paradigm (eye tracking to examine attentive processes during free visual exploration of

Authors	Journal	Sample	Mean age and sex	Tests	Follow-up	Results
Kaye et al ⁵⁴	Int J Eat Disord	9 AN, 7 BN	22 AN, 19 BN	MMFT		AN patients took a greater amount of time to respond after
			Females			the sample stimulus item was presented. In contrast, the BN
						parteries responded more quickly area une sample ream was presented. AN were less impulsive than BN patients.
Gillberg et al ³³	Compr Psychiatry	51 AN, 51 C	21	WAIS-R	4.9 years	AN group scored lower than C on the object assembly
Green et al ^{II}	I Peuchint Res	12 AN 17 C	NR 26 AN 21 C	άντ ςρτ έδτ	10 weeks	subtest. AN groun recalled fewer words disclaved poorer
3)	Females	TFTT, IFRT		reaction times and motor speed than C group. Despite
						having gained weight there was no corresponding
201						improvement in cognitive performance.
Kingston et al"	Psychol Med	46 AN, 41 C	22.1 AN, 22 C	DS, LS, IMI, BD,	Until weight	AN group had a worse performance on attention,
			remales	stroop, PC, Key, TCF. SD. PM. AMT.	gaın of at least 10%	visuospatial capacity and immediate memory. Only attention improved with treatment.
				MP, SDMT		-
Sebastian et al ³⁰	Cogn Ther Res	10 ED	22.73	VS, FRMT		A memory bias for fatness words was found in ED
		30 WPG	21.23			patients.
		30 C	20.36			The three groups did not differ in the recall of non-fat or
			Females			neutral words.
Bradley et al ²⁸	J Clin Exp Neuropsychol	20 AN, 20 C	15.7	WISC-R	Mean	The two groups did not differ on any of the tests used.
		(12 AN retested	Females	LAT	8.5 months	AN patients who recovered performed significantly better
		after weight gain)		CFT, DSPALT,		than C on the Card Rotations Test and Coding F's.
				DVMS, CPT		
Cooper et al [%]	Br J Clin Psychol	12 AN, 12 BN,	Age NR	Stroop		Both patient groups showed attentional bias for eating-
		18 C	Females			and weight-related words. Only AN patients showed
						attentional bias for body shape-related words.
Hermans et al ²⁹	J Abnorm Psychol	12 AN, 12 C	21.08 AN	CR, WCT		Strong explicit memory bias for anorexia-related words in
			24.67 C			AN but not in C. There was no evidence for a similar bias
			Sex ratio (M/F),			in implicit memory.
Mathias et al ²⁰	J Clin Exp Neuropsychol	34 AN (26 inpatients,	22 AN, 20.08 C	WAIS-R, NART-R,		AN patients were deficient in their ability to recall
		8 outpatients)	Females	WMS-R, RAVLT, Rey,		meaningful prose and visuospatial information.
		31 C		АМТ, СОWA, ТМТ		
Rieger et al ¹⁶	Int J Eat Disord	16 AN, 17 BN, 32 C	20.9 AN. 23.9 BN,	VPDT		ED patients detected target probes more slowly than
			20.3 C			C when they appeared in the same location as stimulus
			Females			words connoting a thin physique. The contrary (faster
						detection) occurs with words connoting a large physique.
Sackville et al ¹⁴	Int J Eat Disord	20 AN, 53 C	19.07	Stroop		AN patients, but not unrestrained or restrained eaters,
		(ПЛК-НЛК)	Females			had delayed color-naming latencies for both thin and fat
						WOLD CALEGOLIES ALID, LO A LESSEL EXTENT, IOL NIGN CALOTIC density food words

Luster et al. f primute at al. 12 AN, 14 BN 12 AN mute at all and an input of all and all all and all all and all all and all all all and all all all and all all all all all all all all all al	Authors	Journal	Sample	Mean age and sex	Tests	Follow-up	Results
detail Int/Enr Bendes Females Females Females Per MMMT. RSPMT. detail Int JErr Deord IO N. IO C 159 AN. 16.14 C HT Men detail Encluid Adoles Pay 18 AN. 25 C 145 AN. 15.8 C CFT30. VOC. NS 10.83 days detail Encluid Adoles Pay 16 AN. 16 C 178 AN. 18.7 C Stroop 10.33 days detail Am Clin Psychology 16 AN. 16 C 178 AN. 18.7 C Stroop 10.33 days detail Am Clin Psychology 16 AN. 16 BN. 16 C 23.3 No. 77 Stroop NMT. W. 141. etail Int J Ear Deord 20 AN. 20 C 23.3 No. 77 Stroop NMT. AND. RAD. atail Int J Ear Deord 20 AN. 20 C 23.1 AN. 29.3 1 C Tailored version 14.5 Anor 887. atail Int J Ear Deord 28 AN 28.3 AN. 23.2 BN. MT. Th. C.AL. 3279 days atail Int J Ear Deord 20 AN, 23.3 1 C Tailored version 14.5 AN. 23.4 C Strong atail Int J Ear Deord 28 AN 28 AN. 23.3 L <td>auer et al¹⁸</td> <td>J Psychiatr Res</td> <td>12 AN, 14 BN</td> <td>19.7 AN, 21.9 BN</td> <td>LCT (d2), TMT, DTD,</td> <td>7 months</td> <td>Initial testing: AN and BN showed similar impaired</td>	auer et al ¹⁸	J Psychiatr Res	12 AN, 14 BN	19.7 AN, 21.9 BN	LCT (d2), TMT, DTD,	7 months	Initial testing: AN and BN showed similar impaired
details Int fait Disord DDST details Int Jait Disord IOAN. IOC 159 AN. IS. IS. C HET Men ere et al [®] En Child Adoles Pay IB AN. 25 C 153 AN. IS. IS. C CFT20, VOC. NS 1033 days ere et al [®] Int Jait Disord IB AN. 25 C 178 AN. IB. C Strop 145, months et al [®] Am Clin Psychiany IB AN. 25 C 178 AN. IB. C Strop 145, months et al [®] Am Clin Psychiany IB AN. IS C 178 AN. IB. C Stroop 145, months et al [®] Am Clin Psychiany IB AN. IS C 178 AN. IB 7 C Stroop 145, months et al [®] Am Clin Psychiany Int J Ear Disord 20 AN. 212 BN. IK. TCOMA, TMT, RAM. AN. IB 7 C Stroop et al [®] J Clin Exp Neuropsychoh Int J Ear Disord 20 AN. 212 BN. IK. AM. II. If. Ear Disord 20 AN. 200 C 213 AN. 23 BN. AMT, B. AMA. AMT, B. AMT, B. AMT, AMT, AMT, AMT, B. AMT, B. AMT, AMT, B. AMT, AMT, B. AMT, B.				Females	FPR, MVMT, RSPMT,		performance on attention and problem solving abilities.
details details Int Jara Disord IOAN: IOC IS3 ANI, IS3 C HET Man details Eur Child Adoles Pay IB ANI, 25 C IS3 ANI, IS3 C CFT20, VOC, INS IS3 Bays details Numoprotobiology I6 ANI, I6 C I78 ANI, I83 C Stroop IS3 Bays details Numoprotobiology I6 ANI, I6 C I78 ANI, I83 C Stroop IS3 Bays details Numoprotobiology I6 ANI, I6 C I78 ANI, I83 C Stroop IS3 Bays table Am Clin Psychiatry 39 ANI IR3 C Stroop Stroop IS3 Bays table Int J Ear Disord 20 ANI R 23.8 WASAT-3 WASAT-3 table J Cin Exp Neuropsychol I6 ANI, I6 BNI IG C 23.3 Stroop Stroop all J Int Neuropsychol I6 ANI, I6 C 23.4 23.2 BNI, WIT, IIT, CAL IIT, IT, CAL all Int J Int Neuropsychol I6 ANI, I6 C 23.4 Stroop IIT, CON, ANI, RST 23.79 days all Int Neuropsychol I6 ANI, I6 C <t< td=""><td></td><td></td><td></td><td></td><td>DLPST</td><td></td><td>Their mnemonic functions were preserved. Both</td></t<>					DLPST		Their mnemonic functions were preserved. Both
derails Int J Ent Disord IoAN. IOC I59 AN. IE.I4C HET Men cereral ¹⁰ Err Child Adoles Pay I8 AN. 25 C 145 AN. 158 C CFT20, VOC, NS 1338 days cereral ¹⁰ Neuropsycholology I6 AN. I6 C 778 AN. 187 C Stroop 145 Am. Tob cereral ¹⁰ Neuropsycholology I6 AN. I6 C 738 AN. 187 C Stroop 145 Amorths cereral ¹⁰ Am Clin Psycholology I6 AN. I6 C 738 AN. 187 C Stroop 145 months cereal ¹¹ Am Clin Psycholology I6 AN. I6 C 738 AN. 187 C Stroop 145 months cereal ¹¹ Int J Ent Disord 20 AN. E 23.8 WASAT. WASAT. int J Ent Disord 20 AN. E 23.8 WASAT. MM. A. Men int J Ent Disord 20 AN. 20 C 23.3 AN. 23.8N MMT-R. WIR. BSRT. 2379 days ceral ¹¹ Int J Ent Disord 20 AN. 20 C 23.1 AN. 29.31 C Tailoned version 2379 days cel ¹¹ Int J Ent Disord 20 AN. 20.2 SI. A. 23.1 AN. 23.4 C <							improved after treatment.
Genetal Ennels Familes Number of the Adoles Pay 145 anoths 145 anoths rec et al ¹¹ Neuropsycholology 16 AN, 16 C 778 AN, 18, 16 C 5170, VOC, NS 10383 days rec et al ¹¹ Neuropsycholology 16 AN, 16 C 778 AN, 18, 17 C Stroop 145 anoths rec et al ¹¹ Am Clin Psycholology 16 AN, 16 C 778 AN, 18, 17 C Stroop NaS, 11, WL 141, 14, 14, 14, 14, 14, 14, 14, 14, 14	Grunwald et al ⁵²	Int J Eat Disord	10 AN, 10 C	15.9 AN, 16.14 C	HET	Mean	AN patients showed poorer performance than control
nczetal ¹ Neuropsycholology I6 AN. I6 C 1/3 AN. I8 / C Stroop krali Am Clin Psychiatry 59 AN Familes WMS-III, W. I-II, Familes WMS-III, W. I-II, Familes krali Int J Ear Disord 20 AN-R 233 WMS-III, W. I-II, Familes WMS-III, W. I-II, Familes et al ¹¹ J Clin Exp Neuropsychol 16 AN. 16 BN. 16 C 23.1 WMS-T. S. WIT- S. S. MIT- C. COWA, AHO, Rey, Familes MMT- B. WIT- B. WIT- S. MIT- M. MI	Veumärker et al ⁵⁰	Eur Child Adoles Psv	18 AN. 25 C	Females 14.5 AN. 15.8 C	CFT20. VOC. NS	14.5 months 103.83 days	individuals. Initial resting: number-processing performance was lower
icz ect al ¹¹ Meuropsychobiology 16 AN, 16 C 17.8 AN, 18.7 C Stroop trail Am Clin Psychietry 59 AN 24.3 WAS-II, WL LI, Females WAS-II, WL LI, WAS-II, WL LI, WAS-II, WL LI, WAS-II, WL LI, 20 C 20 AN-R 23.3 WAS-RI, WL LI, WAS-RI, COWA, THT, WR AT-3 WAS-RI, WL LI, WR AT-3 trail J Clin Exp Neuropsychol 20 AN-R 23.3 WAS-RI, WL LI, WR AT-3 WAS-RI, WR AT-3 WAS-RI, WR AT-3 trail J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.3 AN, 23 BN, WR AT-3 MMT-8, WP BSRT, COWA, AHO, Rey, Females SDMT, TMT, CAL 11" Int J Ect Disord 20 AN, 20 C 29.12 AN, 29.31 C Tailored version 11" Int J Ect Disord 28 AN 28 AN, 20-31 C Tailored version 11" Int J Ect Disord 28 AN 29.12 AN, 29.31 C Tailored version ct al ¹¹ J Psychietr Res 32 AN, 29.31 C Tailored version ct al ¹² J Psychietr Res 32 AN, 29.31 C COMA, AHO, Rey 21 J Int Neuropsychol Soc 34 N, 177 C Drawing and C ct al ¹² Psychietr, Res 59 AN, 31 AN-P), Females Corry Soc 26 Psychietr, Res 28 AN, 31 AN-P), BC 21 AN, 23 A C C 26 Psychietr, Res 28 AN,				NR			in patients. With the treatment the prevalence of patients with a subnormal performance was similar to the normal
Image:	1endlewicz et al ¹²		16 AN, 16 C	17.8 AN, 18.7 C	Stroop		population. Authors did not find specific cognitive deficits in AN.
and mutual synony Construction Construction at all Int J Ear Disord 20 C 23.1 WRAT-3 at all J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.3 AN, 23 BN, WCAT WRAT-3 at all J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.3 AN, 23 BN, WCAT WCAT at all J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.3 AN, 23 BN, WCAT WCAT at all J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.3 AN, 23 BN, WCAT MCAT-3 at all J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.3 AN, 23 BN, WCAT MCAT-3 at all Int J Neuropsychol 16 AN, 17 C 23.3 AN, 23 BN, MCAL Tailored version at all Int J Ear Disord 28 AN 28 AN 28 AN 28 AN at all Int J Ear Disord 28 AN 28 AN 17 AN, 234 C 7 allored version at all Int J Ear Disord 28 AN 28 AN 17 AN, 234 C 7 allored version at all Int J Ear Disord 28 AN 28 AN 17 AN, 234 C 7 allored version at all Int J Ear Disord 28 AN 18 AN, 177 C Drawing and at all Plotner Res 32 AN 21 AN, 234 C 67 WST, 0AT, 82 C	laulase at al51		EQ AN	Females			Mild consistiv inscriments is sound tools
et al ¹³ Int J Ent Disord 20 AN-R 23.8 WRAT-3 C 2 23 AN 22 BN, WCAT 4000 W	ajicos er ai			Females	BVRT, COWA, TMT,		No conclusions about the reversibility with weight gain.
et al ¹⁵ Int J Ent Disord 20 AN-R 218 Stroop et al ¹⁶ J Clin Exp Neuropsychol I6 AN, 16 BN, 16 C 213 AN, 22 BN, MMT-B, WIP, BSRT, Emailes Mrth, CAL 253 C 2004A, AHQ, Rey, Females SPMT, THT, CAL 253 C 2004A, AHQ, Rey, Females Int J Neuropsychoph 20 AN, 20 C 29.12 AN, 29.31 C Tailored version of CDR Females Int J Ent Disord 28 AN 24.43 RBANS 24.43 RBANS 32.79 days Females 32 AN 17.7 C Drawing and 54.43 Females COPYING tasks 23 AN, 21.7 AN, 23.4 C GT, WCST 24.43 Females COPYING tasks 32 C Females COPYING tasks 32 C Females COPYING tasks 32 C Females 66.43 RBANS 70.7 OAT, 61.7 AN, 23.4 C GT, WCST 29 AN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 17.7 C Drawing and 67 CDR 28 C GT, MAN, 23.4 C GT, WCST 28 C GT, WCST					WRAT-3		
at al ¹² J Clin Exp Neuropsychol 16 AN, 16 BN, 16 C 23.1 WCST Females Famales MWT-B, WF, BSRT, 23.3 AN, 22 BN, 22 BN, AHO, Rey, Females SDMT, TMT, CAL 11° Int J Neuropsychoph 16 AN, 16 BN, 16 C 23.3 C COWA, AHO, Rey, Females 11° Int J Neuropsychoph 20 AN, 20 C 29.12 AN, 29.31 C Talored version 11° Int J Neuropsychoph 20 AN, 20 C 29.12 AN, 29.31 C Talored version 11° Int J Eat Disord 28 AN 28 AN 26.43 RANS 11° Eat Disord 28 AN 26.43 RANS 32.79 days 11° Eat Disord 29 AN 26.43 RANS 32.79 days 11 Eat Disord 28 AN 26.43 RANS 32.79 days 11 Eat Disord 28 AN 26.43 RANS 32.79 days 11 Fat Disord 28 AN 17.77 C Drawing and 26.43 Paychiatr Res 32.20 Females ContriContri27.43 20 Janeb Females Females ContriContri27.44 21.7 AN, 17.7 C Drawing and 26.43 Contri26.64 21.7 AN, 23.4 C GT, WST, OAT 27.7 <td>assino et al³⁵-</td> <td>Int J Eat Disord</td> <td>20 AN-R</td> <td>23.8</td> <td>Stroop</td> <td></td> <td>AN-R patients showed nonspecific attentional biases with</td>	assino et al ³⁵ -	Int J Eat Disord	20 AN-R	23.8	Stroop		AN-R patients showed nonspecific attentional biases with
et al ¹² J Clin Exp Neuropsychol I6 AN, I6 BN, I6 C 223 AN, 22 BN, MYT-B, WP, BSRT, 23 SC 50WA, AHQ, Rey, Females 50 YTT, THT, CAL al ¹⁹ Int J Neuropsychoph 20 AN, 20 C 29:12 AN, 29.31 C Tailored version Females MBANS 70 C 29:12 AN, 29.31 C Tailored version Females 64.3 RBANS 32.79 days females 64.3 RBANS 32.79 days et al ¹² J Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹² Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹³ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and et al ¹⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and t al J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, B 2 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, B 2 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, B 2 AN, 17 C BT, LITT C BT HIT			20 C	23.1	WCST		a decrease in abstraction capacity and cognitive flexibility
et al ⁻¹ J Luin Exp reuropsyction Io ANI, Io BNI, Io L.J. ANI, LZ INI, TIMT, CAL 31 ³ Int J Reuropsychoph 20 ANI, 20 C 29:12 ANI, 29:31 C Tailored version Females SDMT, TIMT, CAL Females SDMT, TIMT, CAL Females COWA, AHO, Rey, 25:3 C COWA, AHO, Rey, 25:4 3 ANI, 20:4 20:4 20:4 20:4 20:4 20:4 20:4 20:4	1			Females			similar to those patients with OCD.
III ¹⁰ Int J Neuropsychoph 20 AN, 20 C 29.12 AN, 29.31 C Tailored version Ial ¹² Int J Ear Disord 28 AN 20.43.12 AN, 29.31 C Tailored version Ial ¹² Int J Ear Disord 28 AN 26.43 RBANS 32.79 days Ial ¹² J Psychiatr Res 32 AN 18 AN, 177 C Drawing and Eanales 32 AN 18 AN, 177 C Drawing and et al ⁶ Psychiatry Res 32 AN 18 AN, 177 C Drawing and 26.43 RBANS 32 C Females copying tasks et al ⁶ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, 26 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, WCST WCT, MT, BT, a) J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NAT-R, TMT, BT,	vlurpny et al"	J UIIN EXP Neuropsychol		22.3 AIN, 22 BIN,	11101-B, VVIP, BSK1,		In the conditional-associative learning task, AIV patients
III Int / Neuropsychoph 20 AN, 20 C 29.12 AN, 29.31 C Tailored version ial ³ Int J Eat Disord 28 AN 26.43 RBANS 32.79 days tal ¹² J Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and Females 32.79 days tal ¹² J Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and Females 32.79 days et al ⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and Females 32.79 days et al ⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and Females 32.79 days et al ⁶ Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and Females 56.43 et al ⁶ Psychiatr Res 32 C Females Copying tasks a) J Int Neuropsychol Soc 34 N, 19 BN, 35 C 26.7 AN, 26.5 BN NART-R, TMT, BT a) J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN NART-R, TMT, BT				25.3 C	COWA, AHQ, Key,		displayed an impaired performance with neutral materia
II° Int J Neuropsychoph 20 AN, 20 C 29,12 AN, 29,31 C Tailored version cal ³² Int J Eat Disord 28 AN 26,43 RBANS 32.79 days tal ¹² J Psychiatr Res 32 AN 18 AN, 177 C Drawing and Females 32.79 days tal ¹² J Psychiatr Res 32 AN 18 AN, 177 C Drawing and Females 32.79 days et al ⁶ Psychiatr Res 32 AN 18 AN, 177 C Drawing and Copying tasks 32.79 days et al ⁶ Psychiatr Res 32 AN 18 AN, 177 C Drawing and Females Copying tasks et al ⁶ Psychiatry Res 59 AN 18 AN, 234 C GT, WST, OAT, WCST ia J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 267 AN, 265 BN, NCST NART-R, TMT, BT,				remales	SUMI, IMI, CAL		but not with individually threatening material. Such a
alis Int J Eart Disord 28 AN 26.43 RBANS 32.79 days t alia J Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and copying tasks 32.79 days t alia J Psychiatr Res 32 C Females copying tasks 32.79 days et ale Psychiatry Res 32 C Females Copying tasks 32.79 days et ale Psychiatry Res 32 C Females Copying tasks 32.79 days et ale Psychiatry Res 32 C Females Copying tasks 21.7 AN, 23.4 C GT, WST, OAT, WST, OAT, WST, OAT, BL, 33 AN-P), Females (96%-97%) ia Jint Neuropsychol Soc 34 N, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, MT, BT, BT, CBT HIT	end et al ¹⁹	Int I Neurobsvchoth	20 AN 20 C	29 12 AN 29 31 C	Tailored version		deficit was not evident in BN or in C. AN parients showed selective impairments on attention
tal ¹³ Int J Eat Disord 28 AN 26.4.3 RBANS 32.79 days Females 0.000 32.79 days at al ¹² J Psychiatr Res 32 AN 18 AN 17.7 C Drawing and 32 C Females copying tasks et al ⁶ Psychiatry Res 59 AN 21.7 AN 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 54.8 C 55.7 N, TH, TMT, BT, 24.8 C 55.7 N, TH, TMT, BT,		indona (ndo mon form		Eomolos			
t al ¹² J Psychiatr Res 32 AN 18 AN, 17.7 C Drawing and 32 C Females copying tasks et al ⁸ Psychiatry Res 59 AN 2.1.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C T Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 10 T Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT	100 Aoser et al ²⁵	Int J Eat Disord	28 AN	26.43	RBANS	32.79 days	Although standard scores were improved in the immediate
t al ¹² J Psychiatr Res 32 AN 13.7 C Drawing and 32 C Females copying tasks et al ⁸ Psychiatry Res 59 AN 21.7 AN 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C and 1011 Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, and				Females			memory, visuospatial/constructional, language, and
I alsolution J Psychiatr Res 32 AN 18 AN, 177 C Drawing and copying tasks 32 C Females copying tasks et al ⁶ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, WCST et al ⁶ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, WCST et al ⁶ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, WCST al J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NAT-R, TMT, BT, 24.8 C PST, VFT, CBT HIT							attention domain standard scores, only the immediate
et al ⁴² J Psychriotr Res 32 AN 18 AN, 177 C Drawing and 32 C Females copying tasks et al ⁸ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT	5						memory domain improvement was statistically significant.
et al ⁶ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%-97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT 24 C PST, VFT, CBT,	oieters et al ²²	J Psychiatr Res	32 AN	18 AN, 17.7 C -	Drawing and		AN patients were significantly faster in a drawing task
et al ⁶ <i>Psychiatry Res</i> 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT			32 C	Females	copying tasks		and showed shorter reaction times in copying tasks.
et al [®] Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 31 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT							Movement times did not differ significantly between the
et al ⁸ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 31 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C 54.8 C 24.7 C 74.7 C 7							two groups. In the most complex copying task, a signific
et al ⁸ Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT							group $ imes$ complexity interaction for reaction time (patien
et al ⁸ <i>Psychiatry Res</i> 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C 74 ST, VFT, CBT HIT							shorter) and re-inspection time (patients longer) was
et al [®] Psychiatry Res 59 AN 21.7 AN, 23.4 C GT, WST, OAT, (26 AN-R, 33 AN-P), Females (96%–97%) WCST 82 C ia J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT							found. Patients also made more errors than controls.
(26 AN-R, 33 AN-P), Females (96%-97%) WCST 82 C 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT	Cavedini et al ⁸	Psychiatry Res	59 AN	21.7 AN, 23.4 C	GT, WST, OAT,		AN patients showed deficits on decision-making. While
ia J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT			(26 AN-R, 33 AN-P),	Females (96%–97%)	WCST		AN-R patients showed an inability to produce an
ia J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT			82 C				advantageous long-term strategy in the GT, the AN-P
ia J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26.7 AN, 26.5 BN, NART-R, TMT, BT, 24.8 C PST, VFT, CBT HIT							patients did not follow either a clearly advantageous or
ria J Int Neuropsychol Soc 34 AN, 19 BN, 35 C 26./ AN, 26.5 BN, NAKI-K, IMI, BI, 24.8 C PST, VFT, CBT HIT							a clear disadvantageous behavioral strategy.
24.8 C PST, VFT, CBT HIT	l chanturia	J Int Neuropsychol Soc	34 AN, 19 BN, 35 C	26./ AN, 26.5 BN,	NARI-R, IMI, BI,		Anorectic patients show impairments on simple
	et al ^{36,78}			24.8 C	PST, VFT, CBT HIT		alternation and perceptual shift and bulimic patients sho

AN patients showed shorter reaction times in copying tasks and shorter drawing time in the drawing task than normal controls, and this pattern persisted after weight restoration. Patients made more perseverative errors on the WCST, indicating a problem in set shifting. AN patients performed poorly in the IGT compared to the C and to the recovered AN participants.	Women with AN showed a significant deficit in abstract thinking performance, which could not be explained by a more general intellectual deficit or diminished information processing speed. The AN sample also showed a greater preoccupation with detail relative to the control group. AN subjects showed impairments in verbal abilities, cognitive efficiency, reading, mathematics and long-term	An patient of fat and the provident of the and the patients showed attentional bias to blocks of fat and thin words. AN patients scored significantly higher on the efficiency dimension score than the control group. The three clinical groups were significantly impaired on the IGT compared with the C group on both overall task performance and task learning; however, the three clinical	groups were not significantly different from each other. No significant differences in IGT performance were observed between patients and healthy controls, or between restrictive and purging types of anorexia nervosa. Cognitive impairments appear to normalize with refeeding and weight gain. AN patients showed mental rigidity in both verbal and nonverbal domains. AN patients allocated overall less attention to food pictures but showed no early attentional bias toward food pictures. Attentional engagement for food pictures was most pronounced in fasted healthy control subjects	Subtle deficits in cognitive flexibility were found in AN patients compared to C.After weight gain, the AN group improved relative to their baseline values in most of the variables but did not reach C values. They still showed slight impairments. (<i>Continued</i>)
2–5 months	Mean 6.4 years		89 days	2.8 months
Drawing and copying tasks WASI, CVLT, Stroop, TMT, COWA, WCST IGT, NART	Object assembly, GEFT, TMT, SWT SBWJ, HVLT-R, VRS (WMS-R)	Stroop task MFFT IGT	NART, IGT IntegNeuro cognitive battery TMT, WCST, HSCT, IGT Eye-tracking, food/ non-food pictures	P-ORT, TMT, DST, CFT-20-R
17.75 AN, 17.61 C Females 25.6 AN, 24 C Females 28.5 AN, 26.3 C Females Females	21.79 AN, 22.04 C Females 21.3 AN, 20.7 C Females	18–45 Females 26.80 AN, 25.71 BN, 27.27 C Females 29.09 AN, 29.94 BN, 52.11 Ob, 27.75 C Females	23.3 AN, 23 BN, 28 C Females 15.16 AN, NR for C 24.13 AN, 24.67 C NR 24.4 AN, 24.3 C Females	16.2 AN, 16.7 C Females
17 AN, 17 C 15 AN, 11 C 29 AN, 14 R-AN, 29 C	24 AN, 24 C 66 AN, 42 C	6 AN, 6 C 20 AN, 14 BN, 26 C 22 AN, 17 B, 18 Ob, 20 C	49 AN, 38 BN, 83 C 37 AN, 45 C 30 AN (restrictive), 30 C 19 AN, 38 C	30 AN, 28 C
J Clin Exp Neuropsychol J Int Neuropsychol Soc J Int Neuropsychol Soc	J Clin Exp Neuropsychol Pediatrics	Brain Imaging Psyquiatr Res J Int Neuropsychol Soc	Neuropsychology Int J Eat Disord BMC Psychiatry Biol Psychiatry	Psychoneuroendocrinology
Pieters et al ²³ Steinglass et al ³³⁷ Tchanturia et al ⁴⁷	Tokley et al ⁴⁰ Chui et al ⁵³	Redgrave et al ¹³ Southgate et al ³⁸ Brogan et al ⁴⁹	Guillaume et al ⁴⁵ Hatch et al ³² Abbate-Daga et al ¹³ Giel et al ¹⁵	Sarrar et al ²¹

Table I (Continued)	ued)				
Authors	Journal	Sample	Mean age and sex	Tests Follow-up	Results
Shott et al ³¹	Neuropsychology	21 AN, 19 C	25.2 AN, 27.3 C	Implicit category learning task (Gabor patches)	AN patients were less accurate on implicit category learning relative to C. Even when AN patients used the appropriate (ie, implicit) strategy they were still impaired
Cardi et al ⁵⁹	World J Biol Psychiatry	29 AN, I7 BN, I3 R-AN, 9 R-BN, 50 C	27.3 ED, 29.5 ED-R, 25.3 C Females	Dot probe task with faces	Patients with a lifetime diagnosis of ED showed an attentional bias to reject faces and a difficulty disengaging attention from these stimuli.
Abbreviations: Al test (revised); BVT, test-20 (revised); C(paired associates lea GEFT, group embec IFRT, immediate fre pathway in reverse; completion; PM, pro standard progressiw finger tapping task; WAIS-III, Wechsler of the WAIS, Germi WRAT-3, Wide ran	Abbreviations: AN, anorexia nervosa; AHQ, Annett handedness questionnaire; test (revised); BVT, Bakan vigilance task; C, control; CAL, conditional associative test-20 (revised); COWA, controlled oral word association test; CPT, continuous paired associates learning task; DST, digit symbol test; DTD, dual task design; DVY GET, group embedded figures test; GT, gambling task; HDR, high in dietary res IFRT, immediate free recall task; IGT, Iowa gambling test; LAT, lateral asymmetry pathway in reverse; MVMT, Munich, reverbal memory test; MWT-B, verbal intellige completion; PM, prose memory; p-ORT, cognitive flexibility computer based test; standard progressive matrices T; SBWJ, standard batteny of the Woodcock-Johns finger tapping task; TMT, trail making test; VT, verbal fluency task; VOC, vocabb WAIS-III, Wechsler adult intelligence scale III; WAIS-R, Wechsler adult intelligence of the WAIS. German translation; WISC-R, Wechsler intelligence scale for childr WRAT-3, Wide range achievement test 3; WST, Weigl's sorting test.	andedness questionnaire; AMT L conditional associative learr ion test; CPT, continuous per TD, dual task design; DVMS, E HDR, high in dietary restrain HDR, high in dietary restrain ti, LAT, lateral asymmetry test ity computer based test; P5T, ity computer based test; P5T, of the Woodcock-Johnson II uency task; VOC, vocabulary Vechsler adult intelligence scal telligence scale for children (t i sorting test.	T, Austin Maze test; BD, block ning; CBT, cat bat task; CDR, c formance test; CR, cued recall; Denman verbal memory scale; E c, HET, haptic exploration task it: HET, haptic exploration task tri: LCT (d2), letter cancellation NART, National adult reading picture set test; RAVLT, Rey a picture set test; NAVLT, Rey a listory; VPDT, visual probe dd lie (revised); WASI, Wechsler a revised); WL I-II, word lists I al	design; BN, bulimia nervosa; BSRT, Buschke's ognitive drug research; CFT, coding F's task; CVLT, California verbal learning test; DLPST, CVLT, California verbal learning test; DLPST, iD, eating disorders; F, females; FAT, focused a: HIT, haptic illusion task; HSCT, Hayling set nask (d2); LDR, low in dietary restraint; LS, test : NART-R, National adult reading test (fr uitory verbal learning test; RBANS, repeatabl bold digit modality test; SRT, simple reaction ta bbreviate scale of intelligence; WCT, word co dd II of the Wechsler memory scale III; WMS	Abbreviations: AN, anorexia nervosa; AHQ, Annett handedness questionnaire; AMT, Austin Maze test, BD, block design; BN, bulimia nervosa; BSRT, Buschke's selective reminding test, BVRT, Benton visual retention test (revised); BVT, Bakan vigilance task; C, control: CAL, conditional associative learning; CBT, cality invisor problem solving test; DFT, digit symbol presents; C, controlied oral word association test; CPT, continuous performance test; CR, cued recall; CVLT, California verbal learning test; DFT3, dialy living problem solving task; DS digit symbol; DSPAIT, digit symbol presents; DST, digit symbol test; DTD, dual task design; DVMS, Denman verbal menory scate; ED, eating disorders; F, females; FAT, focused attention task; HNLT-R, Hopkins verbal learning test; DST, digit symbol; test; CTT, and presents; DST, digit symbol; DSPAIT, free recall memory test; GET, group embedde figures test; GT, lowa gambling task; HDR, high in dietary restraint; HET, inpute exploration task; HNT, Munich verbal memory test; PMT, free recall memory test; GET, group embedde figures test; GT, lowa gambling test; LAT, and the verbal entry restraint to a divert version task; NMT, matching fiamiliar figure test; PS, repeatable battery of the assessment of neuropsychological status; RSPTT, Raven SMT, matching fiamiliar figure test; PS, sinch and and treading test; LAT, insultand test; DTT, valor completion; PM, prose memory; p-ORT, computer based etst; PST, picture settes; RAVIT, Rey auditory verbal learning test; RANS, repeatable battery of the assessment of neuropsychological status; RSPTT, Raven SMT, matching figure facting test; LAT, test assesting tist; SPMT, symbol approxes, PSMT, savenade and treading test; PTT, treading test; PST, siture settes; RAVIT, Rey auditory verbal learning test; RANS, repeatable battery of the assessment of neuropsychological status; RSPTT, haven standard brock prosters and and batters of the receall task; NTT, traiter assective transition; SPMT, symbol digit; SDMT, symbol digit; TSDMT, symbol digit;

food pictures versus non-food pictures), it has been shown that AN patients have more attentional disengagement to food pictures compared with control subjects. Attentional disengagement was positively related to the severity of the disorder (eg, lower body mass index [BMI]). Apart from the selective attention captured by "emotional pictures," this study reports that individuals with AN show no early vigilance (sustained attention) but do show later avoidance when confronted with food information.¹⁵ Words reflecting either a thin or a large physique and positively or negatively valenced emotion words have been used in a visual detection task with ED patients. Both AN and BN patients directed their attention away from stimulus words connoting a thin physique. In contrast, there was a trend to direct their attention towards stimulus words connoting a large physique. Comparing AN and BN, results reflected a tendency for AN patients to direct their attention toward positive emotion words while those with BN tended to direct their attention away from these words.¹⁶ The "divided attention" has been tested by means of the dual task design (Zimmermann and Fimm)¹⁷ in a 7-month follow-up study. Regarding the attentional demands, the level of performance increased, but it must be noted that only divided attention was impaired at the beginning of this study.¹⁸

In other tasks, predominantly measuring "different facets of attention" (eg, Trail Making Test and letter cancellation test), the level of performance improved as a function of time during treatment.¹⁸ Despite these functions not being impaired at the beginning of the study, other authors have reported that patients are significantly impaired on a number of performance measures related to attentional processes, simple reaction time, choice reaction time, derived "thinking" time, and digit vigilance.¹⁹ In line with Lauer et al,¹⁸ others did not find any significant differences in the attentional or mental tracking capacities (Trail Making Test, revised Wechsler Memory Scale [attention/concentration index], and digit symbol) between AN patients and control participants.²⁰ Moreover, other authors did not find any significant differences between AN patients and control participants using a digit symbol test either at baseline or follow-up.²¹

With respect to "psychomotor speed," Pieters et al reported that anorectic patients were significantly faster in a drawing task and showed shorter reaction times in copying tasks. In the most complex copying task, patients showed shorter reaction time and longer reinspection time with respect to control participants. In addition, patients committed more errors than control participants.²² In order to explore the effect of weight restoration, Pieters et al studied

420

Authors	Journal	Sample	Mean age and sex	Tests	Follow-up	Results
Kaye et al ⁵⁴	Int J Eat Disord	9 AN, 7 BN	22 AN, 19 BN Females	MFFT		AN patients took a greater amount of time to respond after the sample stimulus item was presented. In contrast, the BN patients responded more quickly after the sample item was
Black et al ⁶¹	Int J Eat Disord	16 BN, 29 C	23.8 BN (10	Stroop	2 weeks	presented. AN were less impulsive than BN patients. The data failed to show any specificity in the Stroop effect. Nor
š		(KE and non-KE)	Females			did the test provide a useful measure of treatment response.
Cooper et al [%]	Br J Clin Psychol	12 AN, 12 BN, 18 C	Age NR Females	Color naming Stroop		Both patient groups showed attentional bias for eating and weight related words. Only AN patients showed attentional hiss for body shone-relared words
Ferraro et al ⁵⁵	J Clin Psychol	23 BN, 28 C	18–41, NR	SDMT, FR, WCST, Eckman faces		Measures reflected marked impulsivity and problem-solving deficits in BN.
Lovell et al ⁶²	Br J Clin Psychol	31 AN, 24 BN, 23 R-AN, 11 R-BN, 33 C	25.48 AN, 26.92 BN, 29.30 R-AN, 34.36 R-BN, 24.72 C	Color naming Stroop test		Women currently suffering from bulimia, and those who had recovered from anorexia were found to be more distracted by shape concerns than women who had never suffered eating
	Int Eat Dicord	31 IVI C	Females	"[Dotoction	disorders and women who had recovered from bulimia.
joines-criescers et al ⁶³	ווור) בתר האסום		26.55 C Females	Stroop task	after 6–8 days	in a mixed computed (mixture of word types in each prock), patients took longer to color-name food/eating and weight' shape words than control words. With blocked presentation (with words from just one set in each block) this effect was
						magnined; and patients with builtma nervosa also showed increased naming-latency for "emotion" words.
Rieger et al ¹⁶	Int J Eat Disord	16 AN, 17 BN, 32 C	20.9 AN, 23.9 BN, 20.3 C Females	VPDT		ED patients detected target probes more slowly than C when they appeared in the same location as stimulus words connoting a thin physique. The contrary (faster detection)
Lauer et al ¹⁸	J Psychiatr Res	12 AN, 14 BN	19.7 AN, 21.9 BN Females	LCT (d2), TMT, DTD, FPR, MVMT, RSPM. DLPST	7 months	but in the second of the second secon
Steiger et al ⁶⁷	Int J Eat Disord	51 BN	27.35 Females	DAPP-BQ	Recording 8–22 days	Urge to binge was higher (on average) prior to eating binges than at comparable times on binge free days, and thus seemed to signal the potential for binge eating.
Carter et al ⁶⁴	Int J Eat Disord	98 BN	17—45 Females	Color naming Stroop test	6 weeks	Patients processed information more quickly following treatment and that, overall, patients processed food/body words more slowly than control words, but processed color words even more slowly.
Esplen et al ⁶⁶	Int J Eat Disord	50 BN	26.6 48 females	SRS, A/E MS, BPI-IRS		A lower level of soothing receptivity (indicating a decreased capacity for self-soothing) was correlated with a decreased capacity for evocative memory. A lower level of soothing receptivity and decreased capacity for evocative memory were associated with a greater experience of aloneness.

Dovepress

l able 2 (Continued)	ed)					
Authors	Journal	Sample	Mean age and sex	Tests	Follow-up	Results
Murphy et al ³²	J Clin Exp	16 AN, 16 BN, 16 C	22.3 AN, 22 BN,	MWT-B, WIP, BSRT,		In the conditional-associative learning task, AN patients
	Neuropsychol		25.3 C	COWA, AHQ, Rey,		displayed an impaired performance with neutral material but
			Females	SDMT, TMT, CAL		not with individually threatening material. Such a deficit was
						not evident in BN or in C.
Tchanturia	J Int	34 AN, I9 BN, 35 C	26.7 AN, 26.5 BN,	NART-R, TMT, BT,		Anorectic patients show impairments on simple alternation
et al ^{36,78}	Neuropsychol Soc		24.8 C	PST, VFT, CBT, HIT		and perceptual shift and bulimic patients show difficulties in
			NR			mental flexibility and perceptual shift.
Brand et al ⁶⁸	Neuropsychology	I5 BN, I5 C	21.86 BN,	GDT and NTB		On the GDT, patients with BN chose the disadvantageous
			21.64			alternatives more frequently than did C subjects. Performance
			Females			on the GDT was related to executive functioning but not to
						other neuropsychological functions, personality, or disease-
						specific variables in the BN group.
Mobbs et al ⁵⁷	Eat Behav	18 BN, 18 C	25.11 BN,	Go/no go affective		BN patients reacted faster than C in the task. BN patients
			24.28 C	shifting task		showed poorer discrimination ability than C and inhibition
			Females	1		problems (especially with targets related to food).
Southgate et al ³⁸	Psyciatr Res	20 AN, 14 BN, 26 C	26.80 AN, 25.71 BN,	MFFT		AN patients scored significantly higher on the efficiency
			27.27 C			dimension score than the control group.
			Females			
Liao et al ⁶⁹	J Clin Exp	29 AN, 26 BN,	28.5 AN, 27.8 BN,	IGT		BN patients performed poorly.
	Neuropsychol	51 C	29.4 C			
			Females			
Davis et al ⁷⁷	Appetite	65 obese patients	34.3 BED, 35.2	IGT, DDT		BED and obese patients performed worse than C, but did not
		with BED, 73 obese	non-BED, 31.8 C			differ from each other.
		patients without	Females			
		BED, 71 C				
Guillaume et al ⁴⁵	Neuropsychology	49 AN, 38 BN, 83 C	23.3 AN, 23 BN,	NART, IGT		No significant differences in IGT performance were observed
			28 C			between patients and healthy controls, or between restrictive
			Females			and purging types of anorexia nervosa.
Kemps et al ⁵⁶	J Clin Exp	13 BN, 13 C	22.17 BN,	Stroop, HSCT,		BN patients displayed significant impairments on all inhibition
	Neuropsychol		20.76 C	MFFT, BIS-11		measures and posited significantly higher impulsivity scores
			Females			than the controls.
Legenbauer et al ⁶⁵	J Clin Psychol	25 BN, 27 C	23.88 BN,	Recall and recognition		Poorer recognition and recall of body related stimuli was
			24.74 C	rates of body related,		found for BN in comparison to C, suggesting a memory bias.
			Females	food related and neutral		
				TV commercials		
Brogan et al ⁴⁹	J Int	22 AN, I7 B,	29.09 AN, 29.94 BN,	IGT		The three clinical groups were significantly impaired on
	Neuropsychol	18 Ob, 20 C	52.11 Ob, 27.75 C			the IGT compared with the C group on both overall task
	Soc		Females			performance and task learning; however, the three clinical
						groups were not significantly different from each other.
Mobbs et al ⁷⁶	Appetite	l6 obese patients	45.1, 39.3 and	Mental flexibility task		Obese patients made more errors and more omissions
		with BED, 16, obese	40.2 respectively			than controls in both food and body sections of the task.
		patients without	34 females			Obese patients with BED made significantly more errors and
		BED, 16 C				omissions than those without BED.

422

Patients with a lifetime diagnosis of ED showed an attentional

Dot probe task with

27.3 ED, 29.5 ED-R,

29 AN, 17 BN,

World J Bio

Cardi et al⁵⁹

	Psychiatry	13 R-AN, 9 R-BN,	25.3 C	faces (rejection or	bias to reject faces and a difficulty disengaging attention from
		50 C	Females	acceptance)	these stimuli.
Van den Eynde	J Clin Exp	40 BN, 30 EDNOS-	28.3 BN,	LCT (d2), Stroop	People with BN and EDNOS-BN performed as well as C
et al ⁵⁸	Neuropsychol	BN, 65 C	27.5 EDNOS-BN,	and go/no go task,	on all tasks. Attention task performance was poorer in the
			24 C	GDT	EDNOS-BN than in the BN group.
Abbreviations: A/E inventory: BSRT, Busc eating disorders reco paragraph recall; FR, f Munich verbal memor matrices test; SDMT,	MS, aloneness/evocative i hke's selective reminding <i>i</i> ered; EDNOS, eating diss ree recall; GDT, game of y test; MWT-B, verbal int symbol digits modalities tu	memory scale: AHQ, Annett I test; BT, Brixton test; C, contr orders not otherwise specifier dice task; HIT, haptic illusion t elligence; NART, National adu est; SRS, soothing receptivity :	handedness questionnaire: A rol; CAL, conditional associat d; DAPP-BQ, dimensional ass task; HSCT, Haylings sentenc ilt reading test; NART-R, Nat scale; TMT, trail making test;	N, anorexia nervosa; BED, ; BIS-I1, Barratt impulsivel ive learning; CBT, cat bat task; COWA, controlled ora essment of personality pathology basic questionnaire; e completion test; IGT, Iowa gambling task; LCT (d2) cional adult reading test (revised); NTB, neuropsycholo i VFT, verbal fluency task; VPDT, visual probe detectio	Abbreviations: A/E MS, aloneness/evocative memory scale: AHQ, Annett handedness questionnaire: AN, anorexia nervosa; BED, ; BIS-11, Barratt impulsiveness scale version 11; BN, bulimia nervosa; BPL/RS, BPI, basic personality inventory; BSRT, Buschke's selective reminding test; BT, Brixton test; C, control: CAL, conditional associative learning; CBT, cat bat task; COWA, controlled oral association test; DTD, delay discounting task; ED, eating disorders; ED-R, eating disorders recovered; EDNOS, eating disorders not otherwise specified; DAPP-BQ, dimensional assessment of personality pathology basic questionnaire; DLPST, daily living problem solving test; DTD, dual task design; FPR, free paragraph recall; FR, free recall; GDT, game of dice task; HIT, haptic illusion task; HSCT, Haylings sentence completion test; IGT, lowa gambling task; LCT (d2), letter cancellation task (d2); MFFT, matching familiar figures test; MWT, Munich verbal memory test; MWT-8, verbal intelligence; NART. National adult reading test; VART-R, National adult reading test; VFT, verbal fuency task; VPDT, visual probe detection task; WCST, Wisconsin cast; SS, soothing receptivity scale; TMT, trail making test; VFT, verbal fuency task; VPDT, visual probe detection task; WCST, Wisconsin cast conting test; WMT, trail making test; VFT, verbal fuency task; VPDT, visual probe detection task; WCST, Wisconsin cast conting test; ORT, verbal fuency task; VPDT, visual probe detection task; WCST, Wisconsin cast conting test; ORT, verbal fuency task; VPDT, visual probe detection task; WCST, Wisconsin cast conting test; ORT, wend to the

the performance of AN patients in drawing and copying tasks. Again, AN patients showed shorter reaction times in copying tasks and shorter drawing time in the drawing task compared to normal controls. This pattern persisted after weight gain.²³ This persistence has also been reported in motor tasks after weight recovery of AN patients.²⁴ Considering the effect of an inpatient treatment program for anorexia nervosa, the neuropsychological functioning improves during treatment with significant changes in psychomotor speed.²⁵

In summary, it seems that AN patients have attentional bias to "fat" and "thin" words as well as more attentional disengagement to food pictures. Patients with AN seem to be faster in drawing tasks and tend to show shorter reaction times in copying tasks. Comparing AN and BN, patients with AN tend to direct their attention towards positive emotion words while those with BN tend to direct their attention away from these words. We can conclude that AN patients show more relevant attention deficits in functions such as vigilance and selective attention.

Memory and learning

First of all, it must be noted that different authors study different types of memory with respect to ED. Thus, implicit and explicit memory, short- and long-term memory, and different aspects like learning, recall, recognition of different materials, etc, are usually mentioned.

In this regard, Mathias and Kent explored memory and learning by means of the revised Wechsler Memory Scale, Rey Auditory Verbal Learning Test, Austin Maze, and the Rey-Osterrieth Complex Figure Test. As a result, they found that patients with AN differed from control participants in their performance on the immediate and delayed trials of the logical memory subtests. Patients demonstrated a much poorer ability to recall verbal passages.²⁰ In a recent analysis of the neuropsychological profile of patients with AN, a relative weakness in visuospatial memory has been reported.²⁶ Green et al assessed the cognitive performance of AN patients including an immediate free recall task; patients recalled fewer words than nonclinical controls.¹¹ In addition, Kingston et al reported that anorectic patients had worse performance than controls in different functions including memory.²⁷ With respect to "long-term memory" or continuing storage of information (analyzing immediate word recall/delayed word recall, word recognition, and picture recognition), patients with AN produced a greater number of errors (words not present in the learnt list) and they showed lower sensitivity index in word and picture recognition. There were no differences in reaction times. In the same study,

Wechsler adult intelligence scale.

working memory (temporary storage and manipulation of the information necessary for different tasks) was explored by means of memory scanning and spatial working memory. In this case, patients had significantly longer reaction times.¹⁹ Working memory has been assessed by Lauer et al by means of material presented verbally (analyzing the backward memory span for digits) and material presented visually (analyzing the backward span). In this regard, AN patients showed normal performance.¹⁸ The work of Green et al showed no differences in the Bakan vigilance task when comparing AN patients and control subjects. The Bakan vigilance task has a high loading on the central executive component of the working memory model.¹¹

Considering "explicit memory," two different tasks (verbal and nonverbal) were used in a study by Bradley et al. In addition, different tests of memory were applied. While differences between AN patients and controls were observed with respect to verbal and nonverbal tasks (event related potentials (ERP) waveform amplitudes and latencies), there were no differences on neuropsychological measures, including memory tests.²⁸ Both "implicit" (word-completion test) and "explicit" (cued recall test) memory for shape-, weight-, and food-related words, have been analyzed in patients with AN. Results showed a strong explicit memory bias for anorexia-related words for patients with AN but not for nondieting controls. There was no evidence of a similar bias in implicit memory.²⁹ The explicit memory for fatness words has also been studied and a memory bias for these words was found among anorectic patients.³⁰ Short-term verbal memory (capacity to hold a small amount of information in mind in an available state for a short period of time) has been explored by carrying out a free paragraph recall task and the California Verbal Learning Test in a study by Lauer et al; patients with AN showed normal performance in these tasks.18

Another recent study analyzed implicit category learning. Patients with AN were less accurate when dealing with a task in which they and control participants were asked to categorize simple perceptual stimuli into one of two categories. Results showed that, even when patients used the appropriate (ie, implicit) strategy, they were impaired relative to controls when using the same strategy.³¹ Comparing BN patients and control subjects, AN patients have shown an impairment performance with neutral material but not with individually threatening material in a conditional associative learning task.³²

The main conclusions about this function may be summarized with the following results: AN patients show a

poorer ability to recall verbal passages and they tend to recall fewer words and commit a greater number of errors with longer reaction times. In addition, patients with AN show a strong explicit memory bias for anorexia related words. While AN patients maintain a normal learning memory capability, they show selective memory biases.

Visual perception and visuospatial ability

In the above mentioned study by Bradley et al, AN patients showed longer latencies for nonverbal (visual) tasks relative to verbal tasks, thus noting a theoretical difficulty in processing visual information.²⁸ In another study, AN patients showed a worse performance on tasks measuring visuospatial ability (block design and picture completion).²⁷ Gillberg et al have reported a worse performance of AN patients on the object assembly subtest of the Wechsler Abbreviated Scale Of Intelligence (revised) in contrast with a better result in the block design subtest.³³ Mathias et al assessed visuospatial ability using the Rey-Osterrieth complex figure test and the block design and object assembly subtests of the Wechsler Abbreviated Scale Of Intelligence (revised). There were no differences between AN patients and control participants.²⁰

In summary, compared to other functions, there is a shortage of studies on this area, some results suggesting visuospatial deficits in patients with ED.

Executive functions

In a recent study, executive functions were explored using the Ravello Profile in a sample of patients with AN. Patients were within the average range on the assessment of executive functioning except for one measure of set shifting.26 This Ravello Profile has been suggested as a tool to define a common shared neuropsychological assessment battery.³⁴ Difficulties in abstraction and flexibility of thought have been reported in AN patients when compared with control participants.35 Lauer et al found that AN patients showed mild to moderate deficits, particularly on those tasks covering attentional demands and problem-solving abilities, which improved after several months of treatment.¹⁸ Considering cognitive flexibility, a different pattern has been reported for AN and BN patients: patients with AN show impairments on simple alternation and perceptual shift and BN patients show difficulties in mental flexibility and perceptual shift.³⁶ Difficulties of AN patients with set-shifting tasks have also been reported in other studies.³⁷ Other authors have suggested that AN patients perform better on local information processing tasks than on global processing tasks³⁸ and that they show a cognitive rigidity in both verbal and nonverbal domains.³⁹

In the same line of thinking, women with AN have shown a significant deficit in abstract thinking performance, which could not be explained by a more general intellectual deficit or diminished information processing speed. Patients with AN have also shown a greater preoccupation with detail relative to control participants.⁴⁰ This obsession for details has also been reported by other authors.⁴¹ These deficits in set shifting abilities have been considered independent of starvation in adults.⁴² With respect to the set shifting difficulties as traits linked to possible endophenotypes,43 recently, the set shifting impairment in AN has been reported to be probably unrelated to polymorphisms of SNAP-25 gene.44 In addition, the set shifting deficits have not been demonstrated in adolescent patients with AN.42 In contrast, Kingston et al did not find differences between AN patients and controls by means of cognitive flexibility tasks.27

Another explored function is decision making (the capacity to make decisions about a course of action). In this regard, Guillaume et al used the Iowa Gambling Task to analyze this function; they did not find significant differences between patients and controls.⁴⁵ By means of the same task, a lower decision-making capacity has been reported in both AN and BN patients.^{8,46} In addition, it must be noted that the scores on the Iowa Gambling Task seemed not to improve over time in AN patients.⁴⁷ Using the same task, it has been reported that, compared to control women, AN patients and recovered AN patients showed poor set shifting and decision-making skills.⁴⁸ Including obese patients, a similar impairment on the Iowa Gambling Task in AN and BN patients as well as in obese participants has been found.⁴⁹

Difficulties in abstraction and flexibility of thought along with an obsession for details are considered the main findings in AN patients. With respect to decision making, a lower decision-making capacity has been reported in both AN and BN patients.

Other functions

With respect to mathematic reasoning, Neumarker et al found that, initially, number processing performance was significantly lower in AN patients compared to controls.⁵⁰ However, when the patients restored their normal body weight, the prevalence of patients with a subnormal arithmetic performance was analogous to that in the normal population.

Different studies have failed to report significant differences between patients and controls considering verbal functions.^{20,28,33,51}

Bradley et al did not find learning deficits in digit–symbol paired associate learning.²⁸ Despite having observed a worse

performance on attention, visuospatial ability, and memory, Kingston et al did not find learning deficits.²⁷ In a study by Mathias et al, patients with AN were found to be deficient in the ability to recall meaningful prose and visuospatial information but not in other functions of learning.²⁰

Haptic explorations have been developed in AN patients with poorer performance than control individuals. In addition, reproduction quality was unchanged after weight gain and independent of BMI and intelligence. Mean exploration time was similar in AN patients and controls.⁵²

From a global perspective, impairments in verbal abilities, cognitive efficiency, reading, mathematics, and long term verbal memory have been reported among AN patients even years after diagnosis and with normal BMI.⁵³

ΒN

Neurocognition in BN is clearly under-researched compared to AN, and the most relevant focus has been the comparison between AN and BN patients with respect to impulsivity.^{54,55} In addition, the reported poor inhibitory control in BN patients has been at least partly attributed to an impulsive disposition.⁵⁶ Patients with BN tend to react faster than controls in tasks like go/no go affective shifting. They also have poorer discrimination ability than controls and show inhibition problems, particularly when the targets are related to food.⁵⁷

Attention

A recent study has focused on attention by means of a d2-letter cancellation task, among other functions. As a result, authors found out that patients with BN performed as well as healthy controls on the tasks. Attention task performance was poorer in eating disorders not otherwise specified, bulimic type, than in bulimic patients.⁵⁸

With a visual probe detection procedure, Rieger et al found a tendency for AN patients to direct their attention towards positive emotion words while those with BN tended to direct their attention away from these words.¹⁶

The major finding of Lauer et al was that patients with AN and BN did not differ with respect to their neuropsychological task profiles;¹⁸ both showed mild to moderate deficits, particularly in tasks covering attentional demands and problem-solving abilities.

Cardi et al⁵⁹ have reported that AN and BN patients show an attentional bias to rejecting faces and a difficulty disengaging attention from these stimuli. In addition, they have sustained attentional avoidance of accepting faces. In order to analyze the possible continuum of AN to BN to obesity, compared to

obese patients, AN patients (restrictive type) seem to be more attentive to angry faces and have difficulties in being attentive to positive expressions, while obese patients have shown problems in looking for or being attentive to negative expressions.⁶⁰

Patients with BN have shown worse performance in a symbol digit modalities test; despite being faster than controls, they made more errors.⁵⁵

Other authors have not found differences between BN patients and control participants by way of a modified Stroop test.⁶¹ Similarly, Lovell et al used an emotional Stroop task and determined that women currently suffering from BN and women who had recovered from AN were found to be more distracted by shape concerns than women who had never suffered ED and women who had recovered from BN.⁶² By means of food/eating, weight/shape, emotion, and neutral words in a Stroop task, Jones-Chesters et al reported that BN patients showed increased naming latency for emotion words.⁶³

In order to explore the effects of treatment, Carter et al studied a group of BN patients by means of a Stroop color naming task. Patients performed significantly faster on information processing tasks at posttreatment than at pretreatment and significantly slower on food/body words than on control words. In addition, patients performed significantly slower on color words than on food/body words.⁶⁴

In summary, patients with BN seem to show some attentional biases for weight- and shape-related words as well as an increased naming latency for emotion words.

Memory

Legenbauer et al⁶⁵ studied a group of BN patients who were exposed to body-related and neutral TV commercials then assessed recall and recognition rates. Poorer recognition and recall of body-related stimuli was found for BN patients compared to controls, suggesting a memory bias. Esplen et al⁶⁶ studied the evocative memory in BN by way of the Aloneness/Evocative Memory Scale. A lower level of soothing receptivity (indicating a decreased capacity for self-soothing) was correlated with a decreased capacity for evocative memory. A lower level of soothing receptivity and decreased capacity for evocative memory were associated with a greater experience of aloneness.

Short-term verbal memory has been assessed by way of a free paragraph recall task and the California Verbal Learning Test in a group of ED patients. After 16 weeks of therapy plus 8 weeks of outpatient status, the number of items recalled decreased in AN patients and increased in patients with BN.¹⁸ To summarize, poorer recognition and recall of bodyrelated stimuli have been found, suggesting possible memory biases in ED patients.

Executive functions

The study by Lauer et al reported that AN as well as BN patients showed mild to moderate deficits on tasks relating to problem-solving abilities.¹⁸

Taking into account impulsivity, Steiger et al found that binge eating is closely linked to dietary control in most BN individuals, but this may be less typical of individuals showing marked impulsivity.⁶⁷

Brand et al⁶⁸ explored the decision-making deficits in BN patients by means of the Game Of Dice Task. Patients chose the disadvantageous alternatives more frequently than did control subjects. Performance on the Game Of Dice Task was related to executive functioning but not to other neuropsychological functions, personality, or diseasespecific variables in the BN group. The authors stated that, in BN patients, decision-making abnormalities and executive reductions could be demonstrated and might be neuropsychological correlates of the patients' dysfunctional everyday life decision-making behavior.68 In the same line, Guillaume et al studied decision making by way of the Iowa Gambling Task including AN and BN patients as well as controls.45 These authors concluded that there was not reduced decision making in ED patients. Nevertheless, other authors have found that BN patients performed poorly in this task.⁶⁹

The main results in this area suggest possible decisionmaking abnormalities and executive reductions in BN patients.

BED

There are several studies based on samples comprising chocolate cravers,⁷⁰ fasting and non-fasting normal individuals,⁷¹ overweight/obese females,^{72,73} or subjects with different eating disorders.^{74,75} In these studies, different paradigms have been used, such as Stroop tasks, visual dot probe task, the visual search paradigm, or eye movement monitoring. Nevertheless, there is a shortage of studies specifically focused on BED.

Considering the keys to control unwanted behaviors and thoughts (attention, inhibitory control, mental flexibility), Mobbs et al⁷⁶ compared obese persons with and without BED by means of a food/body mental flexibility task. All patients made more errors and omissions than controls did. Obese patients with BED made more errors and omissions than those without BED. Another study, using the Iowa Gambling Task

and a delay discounting measure, reported that obese and BED patients had worse performance on both tasks compared to control participants, but did not differ from each other.⁷⁷

Discussion

Neuropsychological assessment of ED is being used in order to diagnose better and to conceptualize and design therapeutic plans. It is clear that the main efforts have been expended in AN. Another evident fact is that methodological limitations are more a rule than an exception in the literature regarding this field of study. Is there neuropsychological impairment in ED? Maybe or maybe not. Different types of ED, different populations, different tests, different follow-up periods, different severities, and so on, are hindrances to establishing an accurate answer to that question.

Perhaps the most important question is if the neuropsychological findings reported in ED are reversible with appropriate treatment (ie, are deficits an expression of traits or a mere consequence that emerged during the course of the disorder?). In a study by Green et al,¹¹ AN patients completed different neuropsychological tasks (on three occasions) over the course of 12 weeks of inpatient treatment. Following treatment, patients did not improve their cognitive performance. On the third occasion the mean BMI was 16.53, which represents undernutrition.¹¹ Sarrar et al²¹ studied the cognitive functions of AN patients before and after weight gain. The mean BMI at the final testing session was 17.4. Lauer et al¹⁸ included BN and AN patients in a study assessing their neuropsychological states before, during, and after a treatment. As a result, in the last testing session (7 months after the beginning of the treatment), the impaired cognitive functions improved similarly in AN and BN patients. The main finding of this study was the absence of association between cognitive and clinical rectifications, which led the authors to suggest the existence of mediating factors (eg, hormonal or metabolic). In this case, the weight status was expressed as a percentage of ideal body weight and changed from 70.1% to 86.8% and from 99.6% to 95.8% in AN and BN patients, respectively. Recently, Pieters et al²³ reported the persistence of some altered patterns after weight restoration (change of BMI from 14.56 to 18.90) in AN inpatients after an average stay of 131 days. Another study²⁵ showed that neuropsychological functioning improved over the course of treatment, but this improvement was not associated with a change in BMI (from 16.58 to 19.28 after a mean of 32.79 days). In other cases, the neuropsychological assessment was made a period of time after admission in the hospital (eg, 24.6 days in another study by Pieters et al).²² The study by Carter et al⁶⁴ with BN patients reported that patients performed significantly faster on information processing tasks posttreatment than pretreatment and significantly slower on food/body words than on control words. However, patients performed significantly slower on color words than on food/body words. Kingston et al²⁷ reported that, following treatment, AN patients improved relative to the control group only on tasks assessing attention. In addition, lower weight, but not duration of illness, was associated with poorer performance on tasks assessing flexibility/inhibition and memory. Tchanturia et al⁷⁸ analyzed set shifting tasks in AN patients, and difficulties in these tasks did not show any improvement following retesting after weight recovery.

Besides a few studies that stated that cognitive deficits diminished after weight restoration,^{79–82} others^{11,27,78} have not observed such an improvement. What do neuropsychological deficits represent in ED? To date, this question remains unanswered. The only clear response is that there are severe methodological differences among studies. Are there state-related deficits?

The neuropsychological functions in ED have been accompanied by studies based on neuroimaging and neurophysiology in order to correlate structural and functional brain changes with neuropsychological findings.^{83,84} Due to the enormous amount of variables (weight, duration of illness, medications, etc), it is difficult to demonstrate the correlation between brain changes and functional changes. In order to establish a cause and effect relationship, it would be necessary to develop longitudinal neuroimaging studies. Is there a time limit of duration of weight loss, beyond which normalization of brain function would be more difficult? Would a longer period of normal eating and weight maintenance be required to improve cognitive functioning?⁴ In a recent study based on patients with early-onset AN, authors have suggested that neurobiological abnormalities at initial presentation predict neuropsychological status at follow-up, which might indicate a distinct neurodevelopmental subtype of early-onset AN.85

Different changes in AN patients are not specific. For example, Cooper and Todd have found no differences between AN and BN patients.⁸⁶ In addition, healthy individuals under a restrictive diet may suffer difficulties in sustained attention and short-term memory.⁸⁷ To some extent, it may be that some deficits observed in AN depend on food deprivation (with the corresponding biological consequences). The case of BN seems to be different, with respect to normal weight and overweight individuals. Binge episodes and purging behaviors would cause biological alterations, which, consequently, would alter performance on neuropsychological tasks.⁴

What is the clinical relevance of such a vast number of studies? Although the results give us some new practical knowledge, these types of studies remain substantially theoretical. Is it necessary to implement new forms of treatment to specifically focus on the neuropsychological impairment of these patients? The authors of one study have observed that patients with more cognitive deficits have a worse prognosis.⁷⁹

Former studies on the neuropsychology of ED highlighted the reversibility of neuropsychological impairments.^{18,27} Recent studies try to direct the attention to the neuropsychological impairments as predisposing factors and/or specific eating-disorder-related findings. An example of these efforts to search for ED endophenotypes are the several articles by Lopez et al regarding the concept of central coherence.^{88–90} Nevertheless, potential confounding factors, comorbid pathologies, use of different medications, etc, make it difficult to form definitive conclusions.⁴⁵ It seems that a jump is being made from the "consequences of malnutrition" to "predisposing factors to suffer ED". It must be noted that the unanimous consensus is that there are no gross neuropsychological deficits in AN.9 In addition, despite the persistence of impairments after weight recovery stated by some authors,^{11,27,78} another study has reported that the cognitive performance of AN patients can show improvement even after a period of 2 years following patient discharge.⁹¹ In a recent study focused on the first admission of adolescent patients with AN, cognitive impairments appear to normalize with refeeding and weight gain.92

In summary, the problem with the classification system of ED, the values of BMI considered in different studies, different sample sizes, the absence of ecological paradigms (eg, how neuropsychological deficits affect daily functioning), the possibility of previous neurological lesions (eg, perinatal), the subgroups of ED, duration of illness variability, and comorbid pathologies are some variables to consider before conclusions can be made. In addition, the classification systems of cognitive functions differ considerably among the different studies. As a result, the tests and tasks to assess the same function also differ among studies.

Conclusion

Different neuropsychological alterations have been described in ED, particularly in AN. Nevertheless, there are many inconsistencies among studies, mainly due to methodological biases. It remains unclear if some findings are related to traits or if they are a mere consequence of the core pathology (eg, malnutrition). To date, the clinical and therapeutic relevance of the neuropsychological findings in ED remains unclear. The main change in this field of study may be the view of neuropsychological impairments as predisposing factors of ED rather than a mere consequence of it. Some specific functions such as cognitive flexibility, problem solving, impulsiveness, etc, need to be related to the modern neuroimaging studies on ED in order to clarify the weight of the disposition and the consequences of each type of ED.

Acknowledgment

Thanks to the staff of the Eating Disorders Unit of the Behavioral Sciences Institute, Seville, Spain for its support.

Disclosure

The author reports no conflicts of interest in this work.

References

- Posner MI, DiGirolamo GJ. Cognitive neuroscience: origins and promise. *Psychol Bull*. 2000;126(6):873–889.
- NAN Definition Of A Clinical Neuropsychologist 2001. National Academy of Neuropsychology; 2001. Available from: http://www. nanonline.org/docs/PAIC/PDFs/NANPositionDefNeuro.pdf. Accessed March 16, 2013.
- Steinglass J, Walsh T. Habit learning and anorexia nervosa: a cognitive neuroscience hypothesis. *Int J Eat Disord*. 2006;39(4): 267–275.
- Duchesne M, Mattos P, Fontenelle LF, Veiga H, Rizo L, Appolinario JC. Neuropsychology of eating disorders: a systematic review of the literature. *Rev Bras Psiquiatr*. 2004;26(2):107–117. Portuguese.
- Steiger H, Bruce KR. Phenotypes, endophenotypes, and genotypes in bulimia spectrum eating disorders. *Can J Psychiatry*. 2007;52(4): 220–227.
- Södersten P, Bergh C, Zandian M. Understanding eating disorders. *Horm Behav.* 2006;50(4):572–578.
- Holliday J, Tchanturia K, Landau S, Collier D, Treasure J. Is impaired set-shifting an endophenotype of anorexia nervosa? *Am J Psychiatry*. 2005;162(12):2269–2275.
- Cavedini P, Bassi T, Ubbiali A, et al. Neuropsychological investigation of decision-making in anorexia nervosa. *Psychiatry Res.* 2004;127(3): 259–266.
- Tchanturia K, Campbell IC, Morris R, Treasure J. Neuropsychological studies in anorexia nervosa. *Int J Eat Disord*. 2005;37 Suppl:S72–S76; discussion S87–S89.
- Lezak MD, Howieson DB, Loring DW. Neuropsychological Assessment, 4th ed. New York, NY: Oxford University Press; 2004.
- Green MW, Elliman NA, Wakeling A, Rogers PJ. Cognitive functioning, weight change and therapy in anorexia nervosa. *J Psychiatr Res.* 1996;30(5):401–410.
- Mendlewicz L, Nef F, Simon Y. Selective handling of information in patients suffering from restrictive anorexia in an emotional Stroop test and a word recognition test. *Neuropsychobiology*. 2001;44(2):59–64.
- Redgrave GW, Bakker A, Bello NT, et al. Differential brain activation in anorexia nervosa to fat and thin words during a Stroop task. *Neuroreport*. 2008;19(12):1181–1185.
- Sackville T, Schotte DE, Touyz SW, Griffiths R, Beumont PJ. Conscious and preconscious processing of food, body weight and shape, and emotion-related words in women with anorexia nervosa. *Int J Eat Disord*. 1998;23(1):77–82.

- Giel KE, Friederich HC, Teufel M, Hautzinger M, Enck P, Zipfel S. Attentional processing of food pictures in individuals with anorexia nervosa – an eye-tracking study. *Biol Psychiatry*. 2011;69(7):661–667.
- Rieger E, Schotte DE, Touyz SW, Beumont PJ, Griffiths R, Russell J. Attentional biases in eating disorders, a visual probe detection procedure. *Int J Eat Disord*. 1998;23(2):199–205.
- Zimmermann P, Fimm B. Testbatterie zur Aufmerksamkeitsprüfung [Tests for Attentional Performance]. Würselen, Germany: Psytest; 1993. German.
- Lauer CJ, Gorzewski B, Gerlinghoff M, Backmund H, Zihl J. Neuropsychological assessments before and after treatment in patients with anorexia nervosa and bulimia nervosa. *J Psychiatr Res.* 1999;33(2): 129–138.
- Seed JA, McCue PM, Wesnes KA, Dahabra S, Young AH. Basal activity of the HPA axis and cognitive function in anorexia nervosa. *Int J Neuropsychopharmacol.* 2002;5(1):17–25.
- Mathias JL, Kent PS. Neuropsychological consequences of extreme weight loss and dietary restriction in patients with anorexia nervosa. *J Clin Exp Neuropsychol*. 1998;20(4):548–564.
- Sarrar L, Ehrlich S, Merle JV, Pfeiffer E, Lehmkuhl U, Schneider N. Cognitive flexibility and Agouti-related protein in adolescent patients with anorexia nervosa. *Psychoneuroendocrinology*. 2011;36(9):1396–1406.
- Pieters G, Sabbe B, Hulstijn W, Probst M, Vandereycken W, Peuskens J. Fast psychomotor functioning in underweight anorexia nervosa patients. *J Psychiatr Res.* 2003;37(6):501–508.
- Pieters G, Hulstijn W, Vandereycken W, et al. Fast psychomotor functioning in anorexia nervosa, effect of weight restoration. *J Clin Exp Neuropsychol.* 2005;27(8):931–942.
- Bosanac P, Kurlender S, Stojanovska L, et al. Neuropsychological study of underweight and "weight-recovered" anorexia nervosa compared with bulimia nervosa and normal controls. *Int J Eat Disord*. 2007;40(7): 613–621.
- Moser DJ, Benjamin ML, Bayless JD, et al. Neuropsychological functioning pretreatment and posttreatment in an inpatient eating disorders program. *Int J Eat Disord*. 2003;33(1):64–70.
- Stedal K, Rose M, Frampton I, Landrø NI, Lask B. The neuropsychological profile of children, adolescents, and young adults with anorexia nervosa. *Arch Clin Neuropsychol.* 2012;27(3):329–337.
- 27. Kingston K, Szmukler G, Andrewes D, Tress B, Desmond P. Neuropsychological and structural brain changes in anorexia nervosa before and after refeeding. *Psychol Med.* 1996;26(1):15–28.
- Bradley SJ, Taylor MJ, Rovet JF, et al. Assessment of brain function in adolescent anorexia nervosa before and after weight gain. J Clin Exp Neuropsychol. 1997;19(1):20–33.
- Hermans D, Pieters G, Eelen P. Implicit and explicit memory for shape, body weight and food-related words in patients with anorexia nervosa and non-dieting controls. *J Abnorm Psychol.* 1998;107(2):193–202.
- Sebastian SB, Williamson DA, Blouin DC. Memory bias for fatness stimuli in the eating disorders. *Cogn Ther Res.* 1996;20:275–286.
- Shott ME, Filoteo JV, Jappe LM, et al. Altered implicit category learning in anorexia nervosa. *Neuropsychology*. 2012;26(2):191–201.
- Murphy R, Nutzinger DO, Paul T, Leplow B. Dissociated conditionalassociative learning in anorexia nervosa. J Clin Exp Neuropsychol. 2002;24(2):176–186.
- Gillberg IC, Gillberg C, Råstam M, Johansson M. The cognitive profile of anorexia nervosa: a comparative study including a community-based sample. *Compr Psychiatry*. 1996;37(1):23–30.
- Rose M, Davis J, Frampton I, Lask B. The Ravello Profile: development of a global standard neuropsychological assessment for young people with anorexia nervosa. *Clin Child Psychol Psychiatry*. 2011;16(2): 195–202.
- Fassino S, Pieró A, Daga GA, Leombruni P, Mortara P, Rovera GG. Attentional biases and frontal functioning in anorexia nervosa. *Int J Eat Disord*. 2002;31(3):274–283.
- Tchanturia K, Anderluh MB, Morris RG, et al. Cognitive flexibility in anorexia nervosa and bulimia nervosa. J Int Neuropsychol Soc. 2004;10(4):513–520.

- Steinglass JE, Walsh BT, Stern Y. Set shifting deficit in anorexia nervosa. J Int Neuropsychol Soc. 2006;12(3):431–435.
- Southgate L, Tchanturia K, Treasure T. Information processing bias in anorexia nervosa. *Psychiatry Res.* 2008;160(2):221–227.
- Abbate-Daga G, Buzzichelli S, Amianto F, et al. Cognitive flexibility in verbal and nonverbal domains and decision making in anorexia nervosa patients: a pilot study. *BMC Psychiatry*. 2011;11:162.
- Tokley M, Kemps E. Preoccupation with detail contributes to poor abstraction in women with anorexia nervosa. *J Clin Exp Neuropsychol*. 2007;29(7):734–741.
- Gillberg IC, Råstam M, Wentz E, Gillberg C. Cognitive and executive functions in anorexia nervosa ten years after onset of eating disorder, *J Clin Exp Neuropsychol*. 2007;29(2):170–178.
- Bühren K, Mainz V, Herpertz-Dahlmann B, et al. Cognitive flexibility in juvenile anorexia nervosa patients before and after weight recovery. *J Neural Trans*. 2012;119(9):1047–1057.
- Holliday J, Tchanturia K, Landau S, Collier D, Treasure J. Is impaired set shifting an endophenotype of anorexia nervosa? *Am J Psychiatry*. 2005;162(12):2269–2275.
- 44. Dmitrzak-Węglarz M, Słopień A, Tyszkiewicz M, Rybakowski F, Rajewski A, Hauser J. Polymorphisms of the SNAP-25 gene and performance on the Wisconsin Card Sorting Test in anorexia nervosa and in healthy adolescent participants. *Arch Psyc Psyc.* 2011;13(1):43–51.
- Guillaume S, Sang CN, Jaussent I, et al. Is decision making really impaired in eating disorders? *Neuropsychology*. 2010;24(6):808–812.
- Cavedini P, Zorzi C, Bassi T, et al. Decision-making functioning as a predictor of treatment outcome in anorexia nervosa. *Psychiatry Res.* 2006;145(2–3):179–187.
- 47. Tchanturia K, Liao PC, Uher R, Lawrence N, Treasure J, Campbell IC. An investigation of decision making in anorexia nervosa using the Iowa Gambling Task and skin conductance measurements. *J Int Neuropsychol Soc.* 2007;13(4):635–641.
- Danner UN, Sanders N, Smeets PA, et al. Neuropsychological weaknesses in anorexia nervosa: set-shifting, central coherence, and decision making in currently ill and recovered women. *Int J Eat Disord*. 2012; 45(5):685–694.
- Brogan A, Hevey D, Pignatti R. Anorexia, bulimia, and obesity, Shared decision making deficits on the Iowa Gambling Task (IGT). *J Int Neuropsychol Soc.* 2010;16(4):711–715.
- 50. Neumarker KJ, Bzufka WM, Dudeck U, Hein J, Neumarker U. Are there specific disabilities of number processing in adolescent patients with anorexia nervosa? Evidence from clinical and neuropsychological data when compared to morphometric measures from magnetic resonance imaging. *Eur Child Adolesc Psychiatry*. 2000;9(Suppl 2):111–121.
- Bayless JD, Kanz JE, Moser DJ, et al. Neuropsychological characteristics of patients in a hospital-based eating disorder program. *Ann Clin Psychiatry*. 2002;14(4):203–207.
- Grunwald M, Ettrich C, Krause W, et al. Haptic perception in anorexia nervosa before and after weight gain. J Clin Exp Neuropsychol. 2001;23(4):520–529.
- Chui HT, Christensen BK, Zipursky RB, et al. Cognitive function and brain structure in females with a history of adolescent-onset anorexia nervosa. *Pediatrics*. 2008;122(2):426–437.
- Kaye WH, Bastiani AM, Moss H. Cognitive style of patients with anorexia nervosa and bulimia nervosa. *Int J Eat Disord*. 1995;18(3):287–290.
- 55. Ferraro FR, Wonderlich S, Jocic Z. Performance variability as a new theoretical mechanism regarding eating disorders and cognitive processing. *J Clin Psychol.* 1997;53(2):117–121.
- Kemps E, Wilsdon A. Preliminary evidence for a role for impulsivity in cognitive disinhibition in bulimia nervosa. *J Clin Exp Neuropsychol*. 2010;32(5):515–521.
- Mobbs O, Van der Linden M, d'Acremont M, Perroud A. Cognitive deficits and biases for food and body in bulimia: investigation using an affective shifting task. *Eat Behav.* 2008;9(4):455–561.
- Van den Eynde F, Samarawickrema N, Kenyon M, et al. A study of neurocognition in bulimia nervosa and eating disorder not otherwise specified-bulimia type. *J Clin Exp Neuropsychol*. 2012;34(1):67–77.

- Cardi V, Matteo RD, Corfield F, Treasure J. Social reward and rejection sensitivity in eating disorders: An investigation of attentional bias and early experiences. *World J Biol Psychiatry*. Epub March 6, 2012.
- Cserjési R, Vermeulen N, Lénárd L, Luminet O. Reduced capacity in automatic processing of facial expression in restrictive anorexia nervosa and obesity. *Psychiatry Res.* 2011;188(2):253–537.
- Black CM, Wilson GT, Labouvie E, Heffernan K. Selective processing of eating disorder relevant stimuli: does the Stroop Test provide an objective measure of bulimia nervosa? *Int J Eat Disord*. 1997;22(3): 329–333.
- Lovell DM, Williams JM, Hill AB. Selective processing of shape-related words in women with eating disorders and those who have recovered. *Br J Clin Psychol.* 1997;36(Pt 3):421–432.
- Jones-Chesters MH, Monsell S, Cooper PJ. The disorder-salient stroop effect as a measure of psychopathology in eating disorders. *Int J Eat Disord*. 1998;24(1):65–82.
- Carter FA, Bulik CM, McIntosh VV, Joyce PR. Changes on the stroop test following treatment: relation to word type, treatment condition and treatment outcome among women with bulimia nervosa. *Int J Eat Disord*. 2000;28(4):349–355.
- Legenbauer T, Maul B, Rühl I, Kleinstäuber M, Hiller W. Memory bias for schema-related stimuli in individuals with bulimia nervosa. *J Clin Psychol.* 2010;66(3):302–316.
- Esplen MJ, Garfinkel P, Gallop R. Relationship between self-soothing, aloneness, and evocative memory in bulimia nervosa. *Int J Eat Disord*. 2000;27(1):96–100.
- Steiger H, Lehoux PM, Gauvin L. Impulsivity, dietary control and the urge to binge in bulimic syndromes. *Int J Eat Disord*. 1999;26(3): 261–274.
- Brand M, Franke-Sievert C, Jacoby GE, Markowitsch HJ, Tuschen-Caffier B. Neuropsychological correlates of decision making in patients with bulimia nervosa. *Neuropsychology*. 2007;21(6):742–750.
- Liao PC, Uher R, Lawrence N, et al. An examination of decision making in bulimia nervosa. J Clin Exp Neuropsychol. 2009;31(4):455–461.
- Kemps E, Tiggemann M. Attentional bias for craving-related (chocolate) food cues. *Exp Clin Psychopharmacol*. 2009;17(6):425–433.
- Leland DS, Pineda JA. Effects of food-related stimuli on visual spatial attention in fasting and nonfasting normal subjects: Behavior and electrophysiology. *Clin Neurophysiol*. 2006;117(1):67–84.
- Nijs IM, Muris P, Euser AS, Franken IHA. Differences in attention to food and food intake between overweight/obese and normal-weight females under conditions of hunger and satiety. *Appetite*. 2010;54(2): 243–254.
- Castellanos EH, Charboneau E, Dietrich MS, et al. Obese adults have visual attention bias for food cue images: evidence for altered reward system function. *Int J Obes (Lond)*. 2009;33(9):1063–1073.
- Shafran R, Lee M, Cooper Z, Palmer RL, Fairburn CG. Attentional bias in eating disorders. *Int J Eat Disord*. 2007;40(4):369–380.
- Smeets E, Roefs A, van Furth E, Jansen A. Attentional bias for body and food in eating disorders: increased distraction, speeded detection, or both? *Behav Res Ther.* 2008;46(2):229–238.

- Mobbs O, Iglesias K, Golay A, Van der Linden M. Cognitive deficits in obese persons with and without binge eating disorder. Investigation using a mental flexibility task. *Appetite*. 2011;57(1):263–271.
- Davis C, Patte K, Curtis C, Reid C. Immediate pleasures and future consequences. A neuropsychological study of binge eating and obesity. *Appetite*. 2010;54(1):208–213.
- Tchanturia K, Morris RG, Anderluh MB, Collier DA, Nikolaou V, Treasure J. Set shifting in anorexia nervosa, an examination before and after weight gain, in full recovery and relationship to childhood and adult OCPD traits. *J Psychiatr Res*. 2004;38(5):545–552.
- Hamsher Kde S, Halmi KA, Benton AL. Prediction of outcome in anorexia nervosa from neuropsychological status. *Psychiatry Res.* 1981;4(1):79–88.
- Kohlmeyer K, Lehmkuhl G, Poutska F. Computed tomography of anorexia nervosa. *AJNR Am J Neuroradiol*. 1983;4(3):437–438.
- Small A, Madero J, Teagno L. Intellect, perceptual characteristics, and weight gain in anorexia nervosa. J Clin Psychol. 1983;39(5): 780–782.
- Gillberg IC, Billstedt E, Wentz E, Anckarsäter H, Råstam M, Gillberg C. Attention, executive functions, and mentalizing in anorexia nervosa eighteen years after onset of eating disorder. *J Clin Exp Neuropsychol.* 2010;32(4):358–365.
- Jáuregui-Lobera I. Neuroimaging in eating disorders. *Neuropsychiatr Dis Treat*. 2011;7:577–784.
- Jáuregui-Lobera I. Electroencephalography in eating disorders. Neuropsychiatr Dis Treat. 2012;8:1–11.
- Frampton I, Hutchinson A, Watkins B, Lask B. Neurobiological status at initial presentation predicts neuropsychological functioning in early onset anorexia nervosa at four-year follow up. *Dev Neuropsychol*. 2012;37(1):76–83.
- Cooper M, Todd G. Selective processing of three types of stimuli in eating disorders. Br J Clin Psychol. 1997;36(Pt 2):279–281.
- 87. Green MW, McKenna FP. Developmental onset of eating related colornaming interference. *Int J Eat Disord*. 1993;13(4):391–397.
- Lopez CA, Tchanturia K, Stahl D, Treasure J. Central coherence in women with bulimia nervosa. *Int J Eat Disord*. 2008;41(4):340–347.
- Lopez CA, Tchanturia K, Stahl D, Booth R, Holliday J, Treasure J. An examination of the concept of central coherence in women with anorexia nervosa. *Int J Eat Disord*. 2008;41(2):143–152.
- Lopez CA, Tchanturia K, Stahl D, Treasure J. Weak central coherence in eating disorders: a step towards looking for an endophenotype of eating disorders. *J Clin Exp Neuropsychol*. 2009;31(1):117–125.
- Mikos AE, McDowell BD, Moser DJ, et al. Stability of neuropsychological performance in anorexia nervosa. *Ann Clin Psychiatry*. 2008;20(1):9–13.
- 92. Hatch A, Madden S, Kohn MR, et al. In first presentation adolescent anorexia nervosa, do cognitive markers of underweight status change with weight gain following a refeeding intervention? *Int J Eat Disord*. 2010;43(4):295–306.

Neuropsychiatric Disease and Treatment

Publish your work in this journal

Neuropsychiatric Disease and Treatment is an international, peerreviewed journal of clinical therapeutics and pharmacology focusing on concise rapid reporting of clinical or pre-clinical studies on a range of neuropsychiatric and neurological disorders. This journal is indexed on PubMed Central, the 'PsycINFO' database and CAS. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit http://www.dovepress.com/testimonials.php to read real quotes from published authors.

Submit your manuscript here: http://www.dovepress.com/neuropsychiatric-disease-and-treatment-journal

Dovepress