MMPI-2-RF VRIN-r and TRIN-r Utility: Does Detected Invalid Responding Impact Substantive Scale Criterion Validity?

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INTRODUCTION

Few studies have examined the utility of the MMPI-2-RF non-content-based invalid (NCBI) responding validity indicators, Variable Response Inconsistency (VRIN-r; designed to detect variable responding) and True Response Inconsistency (TRIN-r; designed to detect fixed acquiescent or counteracquiescent responding).

- Handel, Ben-Porath, Tellegen, & Archer (2010) found VRIN-r and TRIN-r were sensitive to computergenerated NCBI responding.
- Burchett et al. (2015) found that utilizing VRIN-r and TRIN-r dramatically decreased interpretive errors in the presence of computer-generated NCBI responding.
- Mason et al. (2013) demonstrated the utility of VRIN-r and TRIN-r to detect random protocols in a simulation study.

This study aims to examine the validity of VRIN-r and TRIN-r in a real-world forensic inpatient sample, where actual invalid responding rates are unknown—but where it can be reasonably assumed that substantive scales *should* better distinguish between subgroups with and without relevant diagnoses in (a) a valid sample than in (b) a sample consisting of invalid fixed and/or variable responses.

HYPOTHESIS

We hypothesized Hedges' *g* values distinguishing those with and without relevant diagnoses would be larger for valid samples than for samples identified as non-content-based invalid by VRIN-r/TRIN-r elevations. Such results would support the utility of VRIN-r and TRIN-r to identify NCBI responding.

METHOD

- We used VRIN-r and TRIN-r to identify 184 forensic inpatients with NCBI protocols and compared them to 641 forensic inpatients with valid protocols.
- We used uncontaminated diagnoses from the date of testing to identify whether patients did or did not experience (1) internalizing dysfunction, (2) thought dysfunction, and (3) externalizing dysfunction disorders.
- Independent samples *t*-tests and Hedges' *g* values were examined to compare mean differences for valid and invalid groups on relevant MMPI-2-RF substantive scales.

Table 1. MMPI-2-RF Internalizing Dysfunction Scale Mean Scores for Subsamples with Valid and Non-Content-Based Invalid Protocols

	Valid Sample						Invalid Sample					
	No Internalizing Dysfunction Dx. $(n = 321)$		Internalizing Dysfunction Dx. (n = 320)			No Internalizing Dysfunction Dx $(n = 103)$						
	М	SD	M	SD	g_1	M	SD	M	SD	g_2		
EID Emotional/Internalizing Dysfunction	49	11	54	13	.39*	60	9	61	9	.12		
RCd Demoralization	52	11	56	12	.35*	63	11	64	11	.13		
RC2 Low Positive Emotions	51	12	54	14	.22*	57	15	54	13	19		
RC7 Dysfunctional Negative Emotions	47	11	52	11	.37*	61	14	64	14	.23		
SUI Suicidal/Death Ideation	50	11	54	15	.30*	65	20	68	20	.16		
HLP Helplessness/Hopelessness	49	11	53	14	.25*	60	13	60	14	.02		
SFD Self-Doubt	50	10	53	12	.33*	58	11	59	11	.09		
NFC Inefficacy	51	11	53	11	.25*	58	11	61	11	.26		
STW Stress/Worry	48	10	51	11	.33*	55	9	57	9	.23		
AXY Anxiety	52	13	56	14	.25*	67	17	72	18	.28		
ANP Anger Proneness	47	9	51	10	.35*	56	9	59	10	.36*		
BRF Behavior-Restricting Fears	52	12	56	14	.26*	71	17	72	18	.10		
MSF Multiple Specific Fears	49	9	52	9	.28*	53	8	54	9	.15		
NEGE-r Negative Emotionality/Neuroticism-Revised	48	10	53	11	.40*	58	11	62	11	.35*		
INTR-r Introversion/Low Positive Emotionality-Revised	51	12	52	12	.06	54	14	52	12	17		

Table 2. MMPI-2-RF Thought Dysfunction Scale Mean Scores for Subsamples with Valid and Non-Content-Based Invalid Protocols

	Valid Sample					Invalid Sample					
	No Thought Dysfunction Dx. $(n = 69)$		Thought Dysfunction Dx. $(n = 572)$			No Thought Dysfunction Dx. $(n = 17)$		Thought Dysfunction Dx. $(n = 167)$			
	M	SD	М	SD	$g_{\scriptscriptstyle 1}$	M	SD	M	SD	g_2	
THD Thought Dysfunction	54	14	58	15	.27*	75	17	79	18	.24	
RC6 Ideas of Persecution	61	16	62	16	.07	80	15	81	18	.05	
RC8 Aberrant Experiences	52	12	55	13	.23	69	15	72	17	.18	
PSYC-r Psychoticism-Revised	53	14	56	15	.22	73	16	78	20	.27	

Table 3. MMPI-2-RF Externalizing Dysfunction Scale Mean Scores for Subsamples with Valid and Non-Content-Based Invalid Protocols

		Valid Sample					Invalid Sample				
	Dysfund	No Externalizing Dysfunction Dx. $(n = 172)$		Externalizing Dysfunction Dx. (n = 469)		No Externalizing Dysfunction Dx. $(n = 45)$		Externalizing Dysfunction Dx. $(n = 139)$			
	M	SD	M	SD	g_1	M	SD	M	SD	g_2	
BXD Behavioral/Externalizing Dysfunction	54	10	59	11	.46*	58	13	66	12	.64*	
RC4 Antisocial Behavior	56	11	62	11	.48*	61	12	68	10	.62*	
RC9 Hypomanic Behavior	47	11	48	11	.10	50	15	58	13	.57*	
JCP Juvenile Conduct Problems	56	13	61	13	.44*	61	14	67	12	.51*	
SUB Substance Abuse	51	10	57	10	.65*	56	12	61	12	.39*	
AGG Aggression	48	10	50	11	.15	54	14	61	13	.49*	
ACT Activation	46	12	47	12	.11	51	13	56	13	.40*	
AGGR-r Aggressiveness-Revised	51	10	52	10	.03	51	11	55	11	.42*	
DISC-r Disconstraint-Revised	53	10	57	10	.43*	56	11	62	11	.50*	
							<u>-</u>	<i>(</i> 1.1		\	

Note for Tables 1-3. Valid sample descriptive results were originally reported in Romero et al. (2016). *p < .05. Hedges' g values (Hedges & Olkin, 1985) are in bold (small: |.20+|; medium: |.50+|; large: |.80+|).

RESULTS

- Most internalizing dysfunction results were in the anticipated direction, with larger differences in the valid than the invalid group.
- **Thought dysfunction** results were also in the anticipated direction, although *g* values were only somewhat smaller for the invalid group, as compared to the valid group.
- Few externalizing dysfunction scale results were in the anticipated direction, with most evidencing larger effects for the invalid group.

DISCUSSION

- The pattern of results from internalizing—and to a lesser degree, thought dysfunction—scales is consistent with the idea that VRIN-r and TRIN-r are able to separate valid and invalid groups.
- Notably, thought dysfunction mean scores were quite high in the invalid subsample, which may have been due to NCBI endorsement of these rarely endorsed items.
- The pattern within the externalizing scales was surprising, and suggests future studies should explore the possibility of mixed responding (e.g., partially random; partially exaggerated responding) on MMPI-2-RF scale scores.
- A post-hoc examination of results suggests our hypotheses may have been too simple. It may be more appropriate to expect NCBI responding to lead to a pattern of results that are *less consistent* with theory (i.e., a mix of larger, smaller, negative effects in a NCBI group) rather than *smaller* effects in particular.

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