



# Response Distortion in Forensic Inpatients with Antisocial Personality Disorder on the MMPI-2-RF Validity Scales

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## INTRODUCTION

According to the DSM-5, individuals diagnosed with antisocial personality disorder (ASPD) are at increased risk for malingering compared to those without the disorder.<sup>1</sup>

**Research has shown conflicting data on whether ASPD should be considered a risk factor for response distortion:**

- Kucharski et al. (2006) found criminal defendants diagnosed with ASPD scored significantly higher than those without ASPD on MMPI-2 overreporting Validity Scales and other validity indicators<sup>2</sup>
- Pierson et al. (2011) did not find evidence that individuals with ASPD were more likely to malingering than others<sup>3</sup>

This study examines whether ASPD can be supported as a **risk factor for overreporting** in an incompetent to stand trial (ICST) forensic inpatient sample, where individuals may have significant motivation to overreport.

We also examine whether there is empirical evidence for ASPD as a **risk factor for underreporting** in a not guilty by reason of insanity (NGRI) forensic inpatient sample, where individuals may have significant motivation to underreport.<sup>4</sup>

## HYPOTHESES

As compared to ICST patients without ASPD, **we expected patients adjudicated ICST and diagnosed with ASPD would score higher on:**

- MMPI-2-RF overreporting Validity Scales (F-r, Fp-r, Fs, FBS-r, RBS), especially those specific to overreported psychopathology (F-r, Fp-r)

As compared to NGRI patients without ASPD, **we expected patients adjudicated NGRI and diagnosed with ASPD would score higher on:**

- MMPI-2-RF underreporting Validity Scales (L-r, K-r), especially the scale designed to measure underreporting of psychopathology (K-r)

## METHOD

- **Participants.** Using archival records of forensic psychiatric inpatients admitted to a large maximum-security state psychiatric hospital, patients were separated into two groups: ICST and NGRI. A total of 146 patients were excluded due to non-content-based invalid responding (CNS  $\geq$  18 [raw], VRIN-r  $\geq$  80T, and/or TRIN-r  $\geq$  80T).<sup>5</sup>
- **Measure.** As part of forensic or clinical evaluations at the hospital, patients in both groups completed the MMPI-2 or MMPI-2-RF<sup>5</sup>. When necessary, MMPI-2 items were rescored into MMPI-2-RF scales.<sup>6</sup>
- **Procedure.** Uncontaminated diagnoses from the date of testing to were used to identify patients with and without antisocial personality disorder diagnoses that were rendered without access to MMPI-2/RF test results.
- Although a uniform procedure was not used across the hospital, diagnoses were rendered by a treatment team consisting of a psychiatrist, psychologist, social worker, and other clinical staff and based upon clinical records and observations made in this 24-hour facility.

**Table 1.**  
MMPI-2-RF Content-Based Validity Scale Scores for Patients Adjudicated Incompetent to Stand Trial (N = 196)

		Antisocial Personality Disorder				t	g
		No (n = 172)		Yes (n = 24)			
		M	SD	M	SD		
F-r	Infrequent Responses	<b>75.60</b>	28.03	<b>101.83</b>	26.36	4.33*	<b>0.94</b>
Fp-r	Infrequent Psychopathology Responses	<b>71.57</b>	25.59	<b>93.54</b>	26.64	3.92*	<b>0.85</b>
Fs	Infrequent Somatic Responses	<b>66.88</b>	24.09	<b>80.96</b>	23.29	2.69*	0.58
FBS-r	Symptom Validity	<b>60.67</b>	15.87	<b>70.79</b>	13.58	2.98*	0.65
RBS	Response Bias	<b>67.36</b>	21.99	<b>87.33</b>	24.69	4.11*	<b>0.89</b>
L-r	Uncommon Virtues	<b>64.11</b>	13.41	<b>56.83</b>	12.06	-2.52*	-0.55
K-r	Adjustment Validity	<b>49.93</b>	11.83	<b>40.54</b>	12.01	-3.64*	-0.79

Note. \*statistically significant t-test (two-tailed); p < .05. g = Hedges' g. All means as well as small ([0.20]-[0.49]), medium ([0.50]-[.79]), and large ([0.80+]) Hedges' g values are **bolded**. MMPI-2-RF (Minnesota Multiphasic Personality Inventory-2-Restructured Form).

**Table 2.**  
MMPI-2-RF Content-Based Validity Scale Scores for Patients Adjudicated Not Guilty By Reason of Insanity (N = 442)

		Antisocial Personality Disorder				t	g
		No (n = 382)		Yes (n = 60)			
		M	SD	M	SD		
F-r	Infrequent Responses	<b>61.73</b>	20.33	<b>64.33</b>	22.21	.91	0.13
Fp-r	Infrequent Psychopathology Responses	<b>58.07</b>	18.44	<b>60.38</b>	20.71	.89	0.12
Fs	Infrequent Somatic Responses	<b>55.38</b>	15.02	<b>56.88</b>	14.39	.72	0.10
FBS-r	Symptom Validity	<b>53.87</b>	11.61	<b>53.42</b>	11.29	-.28	-0.04
RBS	Response Bias	<b>57.55</b>	15.11	<b>57.73</b>	14.78	.09	0.01
L-r	Uncommon Virtues	<b>61.06</b>	13.43	<b>59.85</b>	14.32	-.65	-0.09
K-r	Adjustment Validity	<b>54.18</b>	10.93	<b>51.55</b>	12.02	-1.71	-0.24

Note. \*statistically significant t-test (two-tailed); p < .05. g = Hedges' g. All means as well as small ([0.20]-[0.49]), medium ([0.50]-[.79]), and large ([0.80+]) Hedges' g values are **bolded**. MMPI-2-RF (Minnesota Multiphasic Personality Inventory-2-Restructured Form).

## RESULTS

- As anticipated, results from the ICST sample showed statistically and practically significant differences, with **patients diagnosed with ASPD scoring higher** than those with out the disorder on overreporting scales (Table 1).
- Results from the NGRI sample did not show an anticipated pattern of greater underreporting in ASPD patients. There were **no statistical differences and very small practical differences** (Table 2).

## DISCUSSION

- This study supports the DSM-5 premise that ASPD is a risk factor for overreporting in the presence of an incentive to appear mentally ill.
- In the NGRI population, where there is little to gain from overreporting but potential gains from underreporting, scores were similar to non-ASPD patients, suggesting that ASPD is not a risk factor for underreporting in this context.
- **Limitations & Future Directions.** In addition to the small sample sizes of ASPD patients, diagnoses were not determined using a standardized method. Further studies would benefit from larger samples of ASPD patients as well as standardized assessments for diagnoses.
- This research should be extended by considering factors that distinguish ICST and NGRI patients, such as symptom severity and time in treatment, as well as classification accuracy metrics<sup>2</sup>.

## REFERENCES & ACKNOWLEDGEMENTS

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The statements and opinions reflected in this poster are those of the authors and do not constitute the official views or the official policy of DSH-Patton, the California Department of State Hospitals, or the State of California. This research was made possible by a grant from the University of Minnesota Press and was approved by the CA Department of Mental Health Committee for the Protection of Human Subjects.