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Cognitive Symptom Trajectories of Forensic Inpatients with Psychotic Disorder Diagnoses with and without Comorbid Mood Symptoms

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Introduction

- Most forensic inpatients and roughly 1% of the U.S. population are diagnosed with psychotic disorders¹
- To better inform clinician assessment and treatment, cognitive symptom course should be determined as well as the impact of comorbid mood symptoms
- Two competing trajectory models exist for cognitive dysfunction: degenerative and developmental²⁻⁴
- Research on comorbid mood symptoms is limited and mixed⁵

Aims & Hypotheses

The current study sought to resolve the discrepancy between the trajectory models and explore the impact of comorbid mood symptoms within a large forensic inpatient sample diagnosed with

Hypotheses

- For our overall analyses, based on extant cross-sectional studies⁶, we hypothesized:
 - Young and Middle adults would show some cognitive impairment and Older adults would be the most impaired
- We conducted exploratory analyses to examine the impact of comorbid mood symptoms due to the mixed findings of current research

Method

Participants

- Sample consisted of 708 adult forensic inpatients (> 18 years old) living with schizophrenia spectrum disorder diagnoses Mean age = 40.20 years (SD = 10.72)
- Patients were divided into groups: Young Adult (18-34 years), Middle Adult (35-49 years), & Older Adult (≥ 50 years)
- For subsample analyses, the data set was split into two groups:
 - 1) Psychotic diagnoses only (n = 353)
 - 2) Psychotic + comorbid mood diagnoses (n = 355)

Measures

MMPI-2-RF

Variable Response Inconsistency Scale (VRIN-r) Validity scale; Indirect measure of cognitive problems^{7,8}

Cognitive Complaints Scale (COG)

Self-reported memory, attention, and concentration problems^{7,8}

Procedure

- Our cross-sectional study compared mean scores between age groups on VRIN-r and COG
- For subsample analyses, we conducted independent samples *t*-tests to compare those with and without comorbid mood diagnoses (within age bands)

Table 1: VRIN-r and COG Scores for Younger, Middle, and Older Patients with Psychotic Disorders

	Young (18-34 Years)			Middle (35-49 Years)			Older (≥ 50 Years)				
	n	M	SD	n	M	SD	n	M	SD	F	p
VRIN-r	236	59.94	17.10	338	59.36	14.12	134	57.12	15.03	1.52	.22
COG	148	51.18	11.62	216	50.37	9.95	95	51.84	10.54	0.51	.51

Note: Variable Response Inconsistency Scale (VRIN-r), Cognitive Complaints (COG). For Cognitive Complaints (COG) analyses, invalid protocols (CNS ≥ 18; VRIN-r ≥ 80; TRIN-r ≥ 80; F-r ≥ 120; Fp-r ≥ 100; RBS ≥ 80) were excluded.

Table 2: VRIN-r and COG Scores for Patients with Psychotic Disorders with and without Comorbid Mood Disorders

		Psychotic Only			Psychotic with Mood				
		n	M	SD	n	M	SD	t	p
Young	VRIN-r	116	59.03	17.18	120	60.82	17.05	-0.80	.42
(18-34)	COG	81	50.52	11.78	67	51.99	11.47	-0.76	.45
Middle	VRIN-r	172	59.44	13.64	166	59.29	14.64	0.10	.92
(34-49)	COG	116	50.59	10.03	100	50.12	9.89	0.34	.73
Older	VRIN-r	65	57.12	16.16	69	57.12	14.00	0.003	.998
(≥ 50)	COG	46	50.46	9.16	49	53.14	11.64	-1.25	.22

Note: Variable Response Inconsistency Scale (VRIN-r), Cognitive Complaints (COG). For Cognitive Complaints (COG) analyses, invalid protocols (CNS ≥ 18; VRIN-r ≥ 80; TRIN-r ≥ 80; F-r ≥ 120; Fp-r ≥ 100; RBS ≥ 80) were excluded.

Results & Discussion

Findings

- Found no significant differences for overall or subsample analyses
- Patients may not experience differences in cognitive dysfunction as they age and mood symptoms may not alter severity of cognitive dysfunction

Limitations

- Used indirect (VRIN-r) and self-report (COG) measures of cognitive dysfunction that may not be as sensitive to changes in cognitive symptom severity compared to neuropsychological tests
- Could not control for medication use or age of onset

Strengths

- Large sample of patients living with psychotic disorders
- Measured cognitive dysfunction in two distinct ways

Future Directions

- Use multimethod and direct measures of cognitive dysfunction as well as clinician- and family-ratings
- Study cognitive symptom trajectories in outpatient and community samples

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