

RATES OF PERSONALITY PATHOLOGY AMONG CLINICAL AND NON-CLINICAL SAMPLES

SAIRA KHAN, ANILA KAMAL

National Institute of Psychology, Quaid-i-Azam University, Islamabad, Pakistan

CORRESPONDENCE: SAIRA KHAN, E-mail: sairakhan@nip.edu.pk

Submitted: May 08, 2018 Accepted: November 20, 2018

ABSTRACT

OBJECTIVE

To compare the rates of personality disorders among clinical and non-clinical samples.

STUDY DESIGN

Cross sectional study

PLACE AND DURATION OF STUDY

The study was carried out at National Institute of Psychology, Quaid-i-Azam University Islamabad from August 17, 2015 to June 10, 2017.

SUBJECTS AND METHODS

Employing the technique of convenience sampling, 408 individuals seeking treatment in psychiatric wards and OPDs of Rawalpindi, Islamabad and Lahore were approached. Formal permission was taken from hospital authorities. Informed consent was taken from participants. Similarly data was collected from 360 non-clinical individuals who were not on any psychiatric medicine. The age of the sample ranged from 18-59 years. For the sake of achieving equivalence, both groups were matched on age, gender, marital status, and socio-economic status. The current study utilizes the assessment of DSM-IV Personality Disorder Questionnaire (ADP-IV) to identify particular personality disorder.

RESULTS

Rate of Personality disorders in Clinical sample was 68.1%, whereas for non-Clinical sample it was 16.3 %. Over all rate of Personality disorders was 43.8 %.

CONCLUSION

The present research conclude that special consideration needs to be given to personality pathology in designing treatment plans for patients seeking treatment for mental illness.

KEY WORDS

Personality Co-Morbidity, Personality disorders

INTRODUCTION

Personality Disorders are referred to as relatively enduring patterns of behaviors that interferes markedly with individuals normal and adaptive functioning.1 There lies an underlying similarity in general criteria for diagnosing personality disorders in both International Classification of Disease (ICD) and Diagnostic and Statistical Manual (DSM). Broadly personality disorders have been classified into distinct clusters based on descriptive similarities.2 Cluster A (odd and eccentric disorders) include paranoid, schizoid, and schizotypal personality disorders. Cluster B (dramatic, emotional and erratic disorders) further include anti-social, borderline, histrionic and narcissistic personality disorder. Cluster C (anxious and fearful) include avoidant, dependent and obsessive compulsive personality disorders. Two other unspecified disorders include depressive and passive aggressive personality disorders.² Personality disorders have been strongly linked with other mental illness, poor treatment adherence, attempts of suicide and mortality.3

Studies conducted in different parts of the world indicated that prevalence of Personality disorders varies in different societies. Researches in Britain recorded an overall prevalence of 4.4% in community sample with obsessive compulsive, avoidant, schizoid and borderline being most prevalent and dependent and schizotypal as least prevalent.¹ Studies in United States indicated that overall prevalence of personality disorders varied between 5 to 10 % in community samples.⁴ The overall prevalence of Personality disorders for psychiatric out patients varied between 1.07 % for India to 60 % for Pakistan.⁵

Comorbidity within personality has been well documented by researchers. Individuals who meet the criteria for one Personality disorder are likely to be more vulnerable to other personality disorders as well. It might be from the similar cluster or from different as well. Comorbidity is referred to as conditions that include presence of more than one disorder at either single point of time or across life span. Researchers aimed at studying comorbidity indicated that 54% individuals had only one personality disorder, 22 % had two personality disorders, 11 % had three personality disorders and 14% had four to eight personality disorders.

Taking into account the comorbidity, researchers have devised a five point system of Classifying personality disorders (PD) based on severity and comorbidity ranging from 0 to 5. In this system 0 represent absence of Personality Disorder, 1 representing Personality Difficulty which includes not meeting the full criteria of any PD but having symptoms of Personality disorder, 2 represents either simply one personality disorder or comorbidity within similar cluster, 3 represents complex personality disorders with presence of two or more disorders from different clusters and 4 represents two or more personality disorders resulting in massive societal disturbance.

Personality disorders also show comorbidity with other mental illness which were previously reported on Axis I.⁸ Comparative studies indicate that both in patients and out-patients seeking treatment for other mental illnesses are likely to have an underlying comorbid personality disorder as well.^{8,9} Furthermore the prevalence of Personality disorders is much higher in clinical samples as compared with the non-clinical samples.⁸

Local literature is scarce on the topic so current research aimed at comparing the rate of Personality disorders in both clinical and non-clinical groups.

SUBJECTS AND METHODS

Participants

Employing the technique of Convenience sampling, data was collected from 408 clinical and 306 non-clinical individuals. The age of the sample ranged from 18-59 years. For the sake of equivalence groups were matched on age, education, marital status and Socioeconomic status. Clinical data was obtained from Psychiatric units of Pakistan Institute of Medical Sciences Islamabad, Military Hospital Islamabad, Capital Development Authority Hospital Islamabad and Punjab institute of Mental Health Lahore. It was ensured that only those individuals were included from the clinical sample who were identified as stable by psychiatrist and had an orientation of time and place. Employing the technique of non-probability convenience sampling non-clinical sample was also obtained from Rawalpindi, Islamabad and Lahore. It was ensured that participants from nonclinical sample are not on any psychiatric medicine from the past two year. An effort was made to have same age range, gender ratio and education level for both clinical and non-clinical groups to make the comparison valid.

Instruments

The present research uses the assessment of DSM-IV Personality Disorder Questionnaire (ADP-IV). ADP IV employs diagnostic algorithm, which identify the presence of particular symptom for a disorder. It assess each item on trait and distress associated with it. Trait is rated on a seven point Likert type scale and distress in rated on three point scale. In present study Trait score greater than 5 and distress score greater than 1 has been used to identify the diagnostic criteria according to DSM-IV-TR in light of threshold of symptoms for each PD. 9 It comprises of 94 items. The scale categorizes individuals with Cluster A disorders including Paranoid (7 items), Schizoid (7 items) and Schizotypal personality disorders (9 items), Cluster B including Borderline (10 items), Histrionic (8 items), narcissistic (9 items) and anti-social personality disorder (8 items) and Cluster C

including dependent (8 items), Obsessive compulsive (8 items) and avoidant personality disorder (7 items). It also has not otherwise specified Passive aggressive NOS-PA (7 items) and depressive personality disorders NOS DP (7 items). The scale provides scores of each sub-domain and over all cluster level as well. The scale has been translated and validated for Pakistani sample. ⁹

Procedure

After seeking formal permission from hospital administration data was collected from wards and OPDs of hospitals of Islamabad, Rawalpindi and Lahore. Participants were formally briefed about the purpose of the study and informed consent was sought from them. Individuals were assured about the confidentiality of their responses. Afterwards for the clinical sample items were read to the individuals and their responses were marked. For the sake of comparison demographically similar data was collected for non-clinical sample. Both of them were matched on age, education, marital status and Socio-economic status. In cases where individuals were educated questionnaires were given to the respondents. They were instructed to mark any item that they think is either difficult to comprehend or culturally inappropriate. But in cases where individuals were not educated items were read to them and response were marked as for the clinical sample. At the end participants were thanked for their cooperation.

RESULTS

57.59% of clinical sample comprised of males and 42.40% comprised of females. In contrast, for Non-Clinical sample 40.83% comprised of males and 59.16% comprised of females. 50% of the clinical sample was between the age of 18 to 35 where as 50% was between the age of 36 to 59. Whereas 48.33% from non-clinical sample were between the age of 18-35 and 51.66% were between the ages of 36 to 59. For clinical sample 23.52% of the sample comprised of individuals who did matric whereas for Non-Clinical 22.25% comprised of sample who had completed matric. 38% of individuals comprised of clinical sample who were single whereas for non-clinical 34% individuals were single.

Table I indicated that the rate of Personality disorders in clinical sample was 68.1%, whereas in non-clinical sample it was 16.3%. The combined rate of PDs for both clinical and non-clinical sample was 43.8%. Borderline personality disorder was most prevalent in clinical sample (f = 184) followed by paranoid (f = 123), Schizoid (f = 118) and Obsessive compulsive personality disorder (f = 111). Table 2 indicated significant differences on chi square for all PD's across gender where frequency of all PDs for males was higher as compared with females except for Borderline, Depressive and Passive aggressive PDs for clinical sample. The gender wise comparison was not possible in non clinical group because certain cells in cross table had count less than 5.

Table 1
Rate of Personality Pathology among Clinical and Non-Clinical Sample

Sample	N	Diagnosed f (%)	Un-Diagnosed f(%)	
Clinical	408	278 (68.1%)	130 (31.8%)	
Non-Clinical	360	59 (16.3%)	301 (83.6%)	
Total	768	337 (43.8%)	431 (56.1%)	

Table 2 Chi Square Analysis of Personality Disorders as per Gender among Clinical Sampl

Clinical $(N = 408)$							
PD		Male Female		Female	χ2, df		
			f(%)	f(%)	7.2, uj		
1.	Par	Dia	84(20.59)	39 ((9.55)	8.24*, 1		
		Undiag	151(37.01)	134 (32.84)	6.241, 1		
2.	Sch	Dia	83(20.34)	35 (8.57)	11.03**,1		
	i	Undiag	152 (37.25)	138 (33.82)	11,05**,1		
3.	St	Dia	63 (15.44)	24 (5.88)	n one -		
1		Undiag	172 (42.15)	149 (36.51)	9.93*,1		
4.	AS	Dia	53 (12.99)	7 (1.72)			
		Undiag	182 (44.61)	166 (40.68)	29.20***,1		
5.	BPD	Dia	119 (29.16)	65 (15.93)	c nor :		
\neg		Undiag	116 (28.43)	108 (26,47)	6.87,1		
6.	His	Dia	30 (7.35)	5 (1.22)			
		Undiag	205 (50.24)	168 (41.17)	12.39***,1		
7.	Nar	Dia	48 (11.76)	7 (1.71)	22 02444		
		Undiag	187 (45.83)	168 (40.68)	22.92***,1		
8.	Avo	Dia	59 (14.46)	20 (4.90)	100000000000000000000000000000000000000		
		Undiag	176 (43.13)	153 (37.5)	11.17**,1		
9.	DE	Dia	51 (12.5)	18 (4.41)			
		Undiag	184 (45.09)	155 (37.99)	9.05*,1		
10.	OC	Dia	77 (18.87)	34 (8.33)	8.65*,1		
		Undiag	158 (38.72)	139 (34.06)			
11.	DEP	Dia	48 (11.76)	29 (7.10)	07.		
		Undiag	187 (45.83)	144 (35.29)	.87,1		
12.	PA	Dia	70 (17.15)	38 (9.31)	2121		
\neg		Undiag	165 (40.44)	135 (33.08)	3.13,1		

Note: PD = Personality Disorder, Par = Paranoid, Sch = Schizoid, St = Schizotypal, AS = Anti-social, BPD = Borderline, His = Histrionic, Nar = Narcissistic, Avo = Avoidant, DE = Dependent, OC = Obsessive Compulsive, DEP = Depressive, PA = Passive Aggressive, Dia = Diagnosed, Undiag = Undiagnosed

Results indicated that rate of personality pathology among clinical sample was much higher as compared with non-clinical sample. For clinical sample 67 individuals had considerable score on one personality disorder, 69 on two personality disorders, 49 on three personality disorders respectively (see table 3 for details). This cooccurrence of the personality characteristic can be from the same cluster as well as from different clusters. For non-clinical sample 35 individuals had considerable score on one personality disorder, 8 on two personality disorders and 7 on simultaneous four personality disorders (see table 3 for details).

DISCUSSION

Findings indicated that personality pathology was higher in clinical sample as compared with non-clinical sample. The most frequently occurring personality disorder in clinical sample was borderline

Table 3
Personality Disorders for Clinical and Non- Clinical Sample

			Clinica (N=408		Non-clinic (N=360)	al
	Personality D	isorder Clusters	f	%	f	%
1.	Cluster A	Undiagnosed	217	53.2	260	72.2
		Diagnosed on 1 PD	92	22.5	77	21.4
		Diagnosed on 2 PD	61	15.0	16	4.4
		Diagnosed on 3 PD	38	9.3	7	1.9
2.	Cluster B	Undiagnosed	204	50.0	336	93.3
		Diagnosed on 1 PD	132	32.4	19	5.3
		Diagnosed on 2 PD	32	7.8	260 77 16 7 336 19 5 0 0 4 314 3 32 3 8 6 5 301 35 8 8 3	1.4
		Diagnosed on 3 PD	22	5.4	0	0
		Diagnosed on 4 PD	18	4.4	0	0
3.	Cluster C	Undiagnosed	256	62.74	314	87.2
		Diagnosed on 1 PD	76	18.63	32	8.9
		Diagnosed on 2	45	11.03	8	2.2
		Diagnosed on 3	31	7.6	6	1.7
4.	Total Clusters	Undiagnosed	130 31.86 30	301	83.6	
		Diagnosed on 1 PD	67	16.4	35	9.7
		Diagnosed on 2 PD	69	16.9	8	2.2
		Diagnosed on 3 PD	49	12.0	3	0.8
		Diagnosed on 4 PD	30	7.35	7	1.9
		Diagnosed on 5 PD	17	4.17	3	0.8
		Diagnosed on 6 PD	11	2.7	2	0.6
		Diagnosed on 7 PD	7	1.7	1	0.3
		Diagnosed on 8 PD	9	2.2	0	0
		Diagnosed on 9 PD	13	3.2	0	0
		Diagnosed on 10 PD	6	1.5	0	0

personality disorder (f = 184) which can be attributed to the fact that among all personality disorders people with Borderline Personality tend to seek more treatment. Personality disorder in particular and individuals with Borderline line personality disorder in particular and cluster B overall tend to seek more treatment as these conditions are associated with marked degree of functional impairment as well as increased need to seek attention from significant others in society $^{\rm S,11,12}$. They also tend to have greater comorbidity with other disorders as well $^{\rm S,11}$.

Though low as compared with clinical sample but a substantial number of individuals showed scores on personality pathology in non-clinical sample as well. Past researchers conducted on community sample confirm this finding that prevalence of personality disorder is apparent in non-clinical samples'.

Personality Pathology was more frequent in males as compared with females across clinical sample except for Borderline Personality Disorder, Depressive and Passive Aggressive Personality Disorder.

Previous literature also supports these findings where cluster A specifically i.e. Paranoid, Schizoid and Schizotypal disorders are found to be more prevalent among males^{4, 5}. For Cluster B mixed findings are apparent. BPD has been found to be most prevalent of all PDs and recent literature is suggesting that no gender differences exist in prevalence of BPD though differences might be in symptom severity. Similar trend was apparent for Cluster C where more males had dependent, obsessive compulsive and Avoidant personality disorder.

The presence of co-occurrence in personality pathology has been well documented by previous researchers. SALTALIS. This comorbidity has been attributed to shared etiological factors which might include genetics, biological, environmental, psychological and temperamental factors. Researchers also argued that presence of one personality disorder makes an individual vulnerable for developing another personality disorder. Similarly researches also attributed co-morbidity to complication models which argued that though two disorders represent distinct entities but they are related to one another as one disorder remains in a remitted form and has a scar or complicated effect. Co-occurrence was apparent in both clinical and non-clinical groups with varying degree of complexity and severity. Researches indicated that presence of more than one personality disorder is likely to result in poor treatment outcomes.

The high rate of personality disorders clearly indicated that these need to be assessed properly at the time of diagnosis and designing of intervention plans. It has been evident that personality disorders are often overlooked and ignored by professionals as they are considered to be manifestation of other mental illness to be rated previously on Axis I¹⁷. Researchers further argue that as the symptoms of Axis I disorder improve, the level of associated personality pathology decreases¹⁸. It is critical to assess personality disorders at the initial stage because it has been strongly linked up with shortened life expectancy as well as poor adherence to treatment of the personality pathology can be helpful in modification, adaptation and designing of the treatment plans. ¹⁶

CONCLUSION

It can be concluded that the rate of personality pathology is much higher in clinical sample as compared with non-clinical sample. Further co-occurrence of two or more personality disorder characteristics was also evident. Special consideration needs to be given to the element of co-morbidity of personality pathology in planning treatment for better treatment outcomes.

LIMITATION AND SUGGESTIONS

For assessment of personality pathology, present study used a self-report measure. Further evidence can be gained by adding informant rated measures. It has been suggested that collateral information should be included when making diagnoses for personality disorders ADP IV is based upon categorical approach which ignores the presence of symptoms that do not fully meet the criteria for a particular disorder. Future studies could take in to account the empirical system of taxonomies for assessment of personality pathology.

REFERENCES

- Coid J, Yang M, Tyrer P, Roberts A, Ullrich S. Prevalence and correlates of personality disorder in Great Britain. Br J Psychiatry. 2006 May 1;188(5):423-31.
- American Psychiatric Association. Diagnostic criteria from dsmiv-tr. American Psychiatric Pub; 2000.
- Tyrer P, Reed GM, Crawford MJ. Classification, assessment, prevalence, and effect of personality disorder. The Lancet. 2015 Feb 27; 385(9969):717-26.
- Samuels J. Personality disorders: epidemiology and public health issues. Int. Rev. Psychiatry. 2011 Jun 1;23(3):223-33.
- Beckwith H, Moran PF, Reilly J. Personality disorder prevalence in psychiatric outpatients: a systematic literature review. Personality and mental health. 2014 May 1;8(2):91-101.
- Banerjee PJ, Gibbon S, Huband N. Assessment of personality disorder. Advances in psychiatric treatment. 2009 Sep 1;15(5):389-97.
- Tyrer P, Johnson T. Establishing the severity of personality disorder. American Journal of Psychiatry. 1996 Dec 1;153(12):1593-7.
- Hayward M, Moran P. Comorbidity of personality disorders and mental illnesses. Psychiatry. 2008 Mar 31;7(3):102-4.
- Schotte CK, Doncker DA, Dmitruk D, Mulders IV, D'Haenen H, Cosyns P. The ADP-IV questionnaire: differential validity and concordance with the semi-structured interview. Journal of Personality Disorders. 2004 Aug;18(4):405-19.
- Tyrer P. Personality diatheses: a superior explanation than disorder. Psychological medicine. 2007 Nov 1;37(11):1521-5.
- Skodol AE, Gunderson JG, McGlashan TH, Dyck IR, Stout RL, Bender DS, Grilo CM, Shea MT, Zanarini MC, Morey LC, Sanislow CA. Functional impairment in patients with schizotypal, borderline, avoidant, or obsessive-compulsive personality disorder. American Journal of Psychiatry. 2002 Feb 1;159(2):276-83.
- Keown P, Holloway F, Kuipers E. The prevalence of personality disorders, psychotic disorders and affective disorders amongst the patients seen by a community mental health team in London. Social psychiatry and psychiatric epidemiology. 2002 May 24;37(5):225-9.
- Newton-Howes G, Tyrer P, Anagnostakis K, Cooper S, Bowden-Jones O, Weaver T. The prevalence of personality disorder, its comorbidity with mental state disorders, and its clinical significance in community mental health teams. Social psychiatry and psychiatric epidemiology. 2010 Apr 1;45(4):453-60
- Dolan-Sewell RT, Krueger RF, Shea MT. Co-occurrence with syndrome disorders. Handbook of personality disorders: Theory, research, and treatment. 2001:84-104.
- Cailhol L, Damsa C, Bui E, Klein R, Adam E, Schmitt L, Andreoli A. Is assessing for borderline personality disorder useful in the referral after a suicide attempt?. L'Encephale. 2008 Jan;34(1):23-30
- McGilloway, Angela, Ruth E. Hall, Tennyson Lee, and Kamaldeep S. Bhui. "A systematic review of personality disorder, race and ethnicity: prevalence, aetiology and treatment." BMC psychiatry 10. no. 1 (2010): 33.
- Zimmerman M, Chelminski I, Young D. The frequency of personality disorders in psychiatric patients. Psychiatric Clinics of North America. 2008 Sep 30;31(3):405-20.

- Zimmerman M. Diagnosing personality disorders: A review of issues and research methods. Archives of General Psychiatry. 1994 Mar 1;51(3):225-45.
- Fok ML, Hayes RD, Chang CK, Stewart R, Callard FJ, Moran P. Life expectancy at birth and all-cause mortality among people with personality disorder. Journal of psychosomatic research. 2012 Aug 31;73(2):104-7.
- Trull TJ, Jahng S, Tomko RL, Wood PK, Sher KJ. Revised NESARC personality disorder diagnoses: gender, prevalence, and comorbidity with substance dependence disorders. Journal of personality disorders. 2010 Aug;24(4):412-26.

Sr. Author Name		Affiliation of Author Contribution		Signature
1	Saira Khan	National Institute of Psychology	Data Collection and Report writing	B.K.
2	Prof. Dr. Anila Kamal	National Institute of Psychology	Designing of Methodology and supervision	()