ORIGINAL PAPER

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Regional differences in psychiatric disorders in Chile

Accepted: 7 September 2006 / Published online: 11 October 2006

Abstract Background Psychiatric epidemiological surveys in developing countries are rare and are frequently conducted in regions that are not necessarily representative of the entire country. In addition, in large countries with dispersed populations national rates may have low value for estimating the need for mental health services and programs. Methods The Chile Psychiatric Prevalence Study using the Composite International Diagnostic Interview was conducted in four distinct regions of the country on a stratified random sample of 2,978 people. Lifetime and 12-month prevalence and service utilization rates were estimated. Results Significant differences in the rates of major depressive disorder, substance abuse disorders, non-affective psychosis, and service utilization were found across the regions. The differential prevalence rates could not be accounted by sociodemographic differences between sites. Conclusions Regional differences across countries may exist that have both implications for prevalence rates and service utilization. Planning mental health services for population centers that span wide geographical areas based on studies conducted in a single region may be misleading, and may result in areas with high need being underserved.

Key words community prevalences – psychiatric disorders – regional differences – Chile

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Cross-national psychiatric epidemiological prevalence studies using similar diagnostic instruments have resulted in disparate rates for specific disorders [1]. The reasons for these differences in rates have been attributed to methodological issues between studies; socio-demographic factors such as socio-economic status differences between countries; and cultural differences. Cross-national comparative studies [2, 3] attempt to correct for socio-demographic variability; yet, differences persist. In some regions of the world, such as prevalence rates in Chinese based studies [4-6], the rates are markedly different than the rest of the world. This either suggests true differences in the rates of pathology or lack of cultural appropriateness of the diagnostic tools and the Western diagnostic systems utilized [7]. Understanding these cross-national differences may provide clues into the etiology of psychopathology.

Regional differences also exist within countries as evidenced by studies where the methodology is similar across geographic areas [8–11]. Regional differences in the presence of serious mental illness have been attributed to migration of the mentally ill [12, 13]; birth in urban areas [14]; and genetic pooling [15]. The most studied regional differences are those between urban and rural populations, where urban environmental adversity is argued to contribute to pathology [16, 17].

An understanding of regional differences in countries where the population is spread across large distances is relevant for health care planning. Potentially risk factors may differ across a country resulting in the need to address mental health needs on a regional basis rather than centrally. Furthermore, an understanding of geographic variability permits allocation of resources to be distributed in a proportional basis by need.

This issue is of particular importance to developing countries such as in Latin America, where frequently services are only provided centrally, or $\frac{1}{2}$ disproportionately to the wealthier regions of a country. In addition, epidemiological studies based on single regions of the country [18–20] are extrapolated to larger population bases for which they may or may not be representative.

Population studies about psychiatric disorders in Latin America, as well as other developing regions of the world, are rare. They are important, however, for understanding variations in patterns of disorders, underlying determinants, and service needs. Chile, given its rather extraordinary geography provides an important test of variations in disorder rates across a spatially dispersed population, and offers perhaps the best case example of a country where national rates would seemingly have low value for estimating the need for mental health services and programs.

The Chile Psychiatric Prevalence Study (CPPS) was developed to address issues regarding the prevalence and risk factors for mental illness based on a nationally representative sample, and service utilization. Chile has a population of approximately 16 millions people. The country is composed of 51 provinces grouped in 13 regions covering an area spanning 2 million km² (including Antarctica and Insular Territories) over a length of 8,000 km. The large distances between major population centers resulted in the CPPS being conducted in four regions of the country, Bio Bio, the south-central region containing the second largest city, Metropolitana, the north-central region which includes the capital Santiago, Tarapaca, the north, and Araucania, the south of the country, in order to obtain a representative sampling of the population of the nation. This report focuses on whether regional differences in the prevalence of psychiatric disorders and service utilization, if present, are due to factors other than socio-demographic differences between population centers.

Methods

Sample selection

The CPPS was based on a household stratified sample of people age 15 and older. A more detailed description of the methods used in the CPPS is available in earlier publications [21]. The sample frame was developed to be representative of the nation's population. Four regions and their most representative province and comunas were selected. These were subsequently subdivided into districts, and then randomly selected blocks. The number of households available on each block was enumerated. The 1992 census of each region was used to determine the number of households required on each block. A list of the inhabitants, age 15 and older, in each household was generated. Using pre-assigned Kish tables (Kish 1965) one person per household was selected from the list to be interviewed.

The survey was conducted by the University of Concepcion, Department of Psychiatry and Mental Health, between July 1992 and June 1999, with each site being completed in the following order based on funding: Bio Bio, Metropolitana, Tarapaca, and Araucania. A total of 2,987 individuals participated in the survey. Response rate did differ by site ($\chi^2 = 11.08$, df = 3, P < 0.02) with Metropolitana having the highest non-response rate 12.6% and Tarapaca the lowest 7.5%. A weight was used to account for the probability of the comuna, district, block, household, and respondent being selected. The data was adjusted to the 1992 census of each region based on age, gender, and marital status using a second weight.

Diagnostic assessment

Composite International Diagnostic Interview (CIDI) versions 1.0 and 1.1 [22] were used to generate the diagnoses using well-trained lay interviewers. DSM-III-R [23] diagnostic criteria were employed. A section on health service utilization in the 6-months prior to the interview was also included. The Spanish translation was conducted using the protocol outlined by the World Health Organization (WHO) [24]. A validation study of the Chilean CIDI was found to have kappas that ranged from 0.52 for somatiform disorders up to 0.94 for affective disorders [25] using a sample of patients and volunteers for each CIDI section. After double entry of data and verification for logical inconsistencies diagnoses were generated using the CIDI computer programs for 1.0 and 1.1 [26]. The DSM-III-R diagnoses included in this report are all affective disorders; all anxiety disorders defined as panic disorder, agoraphobia, and generalized anxiety disorder; substance use disorder which does not include nicotine dependence, and any diagnosis. Lifetime and 12month prevalence rates were examined.

Interviewers and training

Social science university students in their senior year underwent training following the WHO protocol at the University of Concepcion, a WHO CIDI training and reference center. The 64 interviewers received over 80 h of instruction and practice sessions. Each interviewer had to conduct practice interviews with volunteer adult subjects with and without psychiatric disorders selected from local clinics, as well as a pilot interview on an individual in a non-selected household in the community. Approximately 80% of the interviews were audiotaped following the subject's consent, and 20% randomly reviewed for quality control.

Analysis procedures

The SUDAAN statistical package [27], Taylor series linearization method, was used to estimate the standard errors due to the sample design and the need for weighting. The analysis was conducted using procedures without replacement for non-respondents. The comuna and district selected were used as the defined strata. Chisquare analyses were used to examine the association of disorders and service utilization between regions. Logistic regression was used to adjust for socio-demographic differences across regions accounting for differential rates or service utilization. Additional analyses were conducted to examine urban-rural differences. The logistic regression analyses included gender, marital status, age group, education, and income as potential confounders. All results are presented as weighted data.

Results

The distribution of income and marital status were found to differ across the four regions of the country (see Table 1). The population of Bio Bio had significantly lower incomes than the other regions ($\chi^2 = 67.96$, df = 9, P < 0.0001). In addition, Bio Bio had the lowest rate of individuals who were separated or had annulled marriages ($\chi^2 = 25.31$, df = 12, P < 0.05). In two of the regions the rural population was under-represented relative to the census. In Bio

Table 1	Socio-demographic	characteristics	by	region
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	Bio Bio (<i>N</i> = 80	0)	Metropo $(N = 13)$	litana 63)	Tarapaca $(N = 30)$	і б)	Araucani (N = 50	ia 9)			
Disorders	%	SE	%	SE	%	SE	%	SE	χ^2	df	Р
Gender											
Male	48.2	1.0	46.4	1.4	48.9	1.0	47.4	1.5	3.23	3	0.38
Female	51.8	1.0	53.6	1.4	51.1	1.0	52.6	1.5			
Age											
15–24	26.7	1.6	24.9	2.1	27.3	2.3	27.0	7.4	7.3	15	0.94
25–34	25.4	1.6	26.1	1.4	27.1	0.6	23.2	4.6			
35–44	18.2	1.9	18.5	1.1	21.1	2.0	17.5	2.6			
45–54	12.1	1.4	12.8	1.2	11.8	0.4	12.3	1.2			
55–64	9.0	1.4	9.2	0.8	6.5	0.4	9.8	1.3			
65+	8.7	2.3	8.5	1.0	6.2	1.9	10.4	1.7			
Education											
No education	2.7	1.0	1.1	0.2	0.3	0.2	1.3	1.0	11.13	9	0.31
Basic	19.7	3.8	17.9	2.3	7.8	2.0	15.2	7.6			
Medium	52.3	3.3	47.9	1.6	52.7	1.5	38.2	7.3			
High	25.2	7.1	33.2	2.3	39.1	3.2	45.2	15.4			
Marital status											
Married	54.7	2.8	52.8	2.3	53.7	3.2	52.3	5.8	25.31	12	0.05
Widowed	5.2	1.1	4.7	0.7	3.3	1.3	5.8	0.2			
Separated/anulled	1.7	0.7	4.1	0.5	3.3	1.4	2.0	0.6			
Never Married	34.4	1.9	32.6	2.4	33.9	1.1	36.1	8.0			
Common Law	4.0	0.8	5.8	1.0	5.9	0.6	3.8	2.0			
Income											
U\$100–U\$400	70.5	7.1	54.5	3.9	12.2	4.8	47.7	15.4	67.96	9	0.0001
U\$401–U\$800	19.3	3.3	21.9	1.8	29.6	2.5	20.6	4.5			
U\$801–U\$1500	6.9	1.9	11.9	1.3	35.6	3.3	12.5	4.0			
U\$1501+	3.3	2.5	11.7	2.8	22.7	3.6	19.2	7.2			
Urban/rural											
Urban	97.3	3.1	99.2	0.9	98.9	1.4	92.3	5.0	3.1	3	0.39
Rural	2.7	3.1	0.8	0.9	1.1	1.4	7.7	5.0			

Bio 22% of population was rural the sample only included 2.7%, and in Araucania 38% of the population was rural and the sample only included 7.7%.

Prior to adjusting for socio-demographic differences between the regions, a number of differences in prevalence rates were noted (see Tables 2, 3). Lifetime rates for major depressive disorder were markedly elevated in Tarapaca, 17.2%, and lowest in Bio Bio, 11.6% ($\chi^2 = 9.76$, df = 3, P < 0.04). Drug abuse, but not dependence, also had the highest prevalence rate in Tarapaca, 2.4% ($\chi^2 = 8.59$, df = 3, P < 0.05). Interestingly, in Araucania the rate of non-affective psychosis was the lowest ($\chi^2 = 11.45$, df = 3, P < 0.02). When females were examined the differential rates for major depressive disorder ($\chi^2 = 11.76$, df = 3, P < 0.02) and substance use disorders were noted ($\chi^2 = 10.88$, df = 3, P < 0.03). Among males the only lifetime difference in prevalence rates was for elevated alcohol abuse in Tarapaca ($\chi^2 = 9.17$, df = 3, P < 0.04). For 12-month prevalence the increased risk for major depression in Tarapaca persisted ($\chi^2 = 8.78$, df = 3, P < 0.05) for both genders combined and for females ($\chi^2 = 10.08$, df = 3, P < 0.03). The differences noted in the prevalence of substance use disorders were no longer evident at 12-months.

Using logistic regression controlling for sociodemographic variables the regional differences for major depression were maintained for both lifetime and 12-month prevalence, as well as among females in both prevalence periods. Males with lifetime prevalent affective disorders were also at increased risk in Tarapaca. In addition regional differences in lifetime prevalence for alcohol abuse, drug abuse, and nicotine dependence were found. Among females, regional differences in lifetime prevalence were noted for drug abuse, drug dependence, any alcohol or drug use disorder, and cognitive disorders, and among men alcohol abuse with increased risk among those residing in Tarapaca. The statistical differences in regional lifetime prevalence of non-affective psychosis for both genders combined, and females in both prevalence periods, persisted with the rates for Araucania remaining low. As the rates for not only major depression, but also alcohol and drug use disorders were elevated in Tarapaca, additional analyses were conducted controlling for comorbidity in the logistic regressions, the regional differences noted were not altered.

Differences in service utilization across the four regions were also found. Araucania had the lowest use of mental health services utilization ($\chi^2 = 0.03$, df = 3, P < 0.03), in particular in the non-specialized health care sector ($\chi^2 = 12.63$, df = 3, P < 0.02). The rates of service utilization by region are presented in

Table 2 Lifetime prevalence rates of DSM-III-R disorders by region

	Bio Bio		Metropo	litana	Tarapaca	a	Araucan	ia		
Disorders	%	SE	%	SE	%	SE	%	SE	χ^2	Р
Affective disorders										
Major depressive episode	7.1	1.3	11.6	0.8	17.2	2.4	9.8	0.8	9.76	0.04
Manic episode	2.2	0.7	1.4	0.4	1.8	0.3	1.5	1.3	1.11	0.78
Dysthmia	7.5	1.1	7.3	1.2	12.2	1.7	6.0	3.3	3.65	0.32
Any affective disorder	13.6	2.2	15.4	1.2	23.2	1.9	14.0	3.2	3.88	0.30
Anxiety disorders										
Panic disorder	1.2	0.6	1.3	0.3	4.3	0.8	1.1	0.5	1.16	0.//
Agoraphobia without panic	14.2	2./	9.8	1.3	9.7	1.8	5.3	0.8	5.46	0.16
Any anyiety disorder	1.0	0./	3./ 14.0	0.5	2.0	0.4	5.U 0.6	0.0	5.15	0.19
Substance use disorder	19.2	5./	14.8	1.0	17.9	1.7	8.0	0.9	0.79	0.11
	28	0.8	2.0	0.5	16.9	26	7.2	14	8 20	0.06
Alcohol dependence	7.0	17	6.4	0.5	63	0.5	5.0	0.8	3 63	0.00
Drug abuse	0.6	0.3	1.5	0.4	2.4	0.5	0.1	0.0	8.59	0.05
Drug dependence	2.2	0.7	3.3	1.0	2.2	0.3	1.2	0.1	7.72	0.07
Nicotine dependence	2.9	0.6	2.1	0.6	6.5	0.2	5.4	1.0	7.86	0.07
Any alcohol or drug use disorder	11.0	1.8	11.2	1.0	24.3	2.9	12.3	1.8	1.94	0.59
Any substance use disorder	13.0	2.0	12.9	1.1	29.6	2.9	14.6	1.7	2.43	0.50
Other disorders										
Non-affective psychosis	2.1	0.6	2.3	0.5	0.8	0.2	0.1	0.0	11.45	0.02
Somatoform disorder	2.7	1.1	4.4	0.8	3.1	0.6	3.5	0.7	1.75	0.63
Cognitive disorder	4.5	1.8	3.7	0.8	0.7	0.8	1.0	0.4	5.70	0.15
Any CPPS disorder	32.2	4.1	30.8	1.6	44.4	1.9	28.9	3.1	2.51	0.49
Female										
Affective disorders	71	1 2	15.2	1 /	20.6	0.0	11 3	1 1	11 76	0.00
Major depressive episode	7.1	1.5	15.2	1.4	20.0	0.8	0.5	1.1	11./0	0.02
Dysthmia	2.0	2.0	1.0	0.5	5.0 17.2	0.0	0.5	0.5	5.02 2.43	0.52
Any affective disorder	16.4	2.0	21.0	1.9	78.3	0.0	16.0	4.9	3 60	0.50
Anxiety disorders	10.4	2.0	21.0	1.7	20.5	0.7	10.0	ч.)	5.00	0.52
Panic disorder	1.7	0.9	2.1	0.5	6.7	1.2	2.1	0.9	1.15	0.77
Agoraphobia without panic	20.3	3.2	13.6	2.0	12.2	0.2	7.0	2.4	4.88	0.20
Generalized anxiety disorder	2.5	1.0	6.4	1.0	3.8	0.8	4.4	0.4	5.98	0.13
Any anxiety disorder	26.4	3.9	20.9	2.5	24.8	2.1	12.3	2.0	5.19	0.18
Substance use disorders										
Alcohol abuse	0.5	0.4	1.1	0.6	2.6	0.4	1.0	0.4	2.06	0.57
Alcohol dependence	1.7	0.7	2.0	0.7	4.2	0.2	0.4	0.2	7.53	0.07
Drug abuse	0.0	0.0	0.8	0.3	2.3	0.5	0.0	0.0	7.38	0.08
Drug dependence	2.4	0.7	5.0	1.6	2./	0.4	0.2	0.2	6.86	0.10
Nicotine dependence	2.5	0.7	2.3	0.8	10.0	0.6	2.6	0.4	2.45	0.49
Any alconor of drug use disorder	5.0 5.0	0.9	7.1	1.5	0.9	0.0	1.0	0.0	11.00	0.02
Ally substance use disorder Other disorders	5.9	1.4	0.9	1.4	10.5	0.5	4.1	0.7	10.00	0.02
Non-affective psychosis	24	07	19	0.6	16	03	01	01	7 81	0.07
Somatoform disorder	2.4	0.7	5.3	1.1	3.6	0.7	4.4	1.7	4.10	0.27
Any CPPS disorder	35.6	3.8	34.7	2.5	37.8	1.1	23.4	4.0	3.02	0.40
Male										
Affective disorders										
Major depressive episode	7.1	1.5	7.4	1.3	13.6	4.6	8.3	1.5	6.84	0.10
Manic episode	1.8	1.0	1.0	0.4	0.5	0.6	2.6	2.2	1.18	0.76
Dysthmia	4.0	1.0	2.4	0.6	7.1	3.3	3.1	2.0	4.10	0.27
Any affective disorder	10.5	2.2	9.0	1.4	18.0	4.3	11.8	2.9	14.57	0.01
Anxiety disorders	0.7	0.5	0.4	0.2	1.0	0.2	0.0	0.0	4.20	0.24
Panic disorder	0.7	0.5	0.4	0.2	1.8	0.3	0.0	0.0	4.39	0.24
Agoraphobid without panic Generalized anxiety disorder	1.7	2.9	5.5 0.7	1.9	7.2	5.5 0.0	5.5 1 3	1.0	5.94 1.83	0.28
Any anyiety disorder	11.0	4.4	79	17	10.6	3.4	4.5	0.5	7 59	0.01
Substance use disorders	11.5	7.7	7.5	1.7	10.0	5.4	- 1 .J	0.5	1.59	0.07
Alcohol abuse	5.3	1.5	3.0	0.8	31.9	5.6	14.1	2.8	9.17	0.04
Alcohol dependence	12.5	3.7	11.6	1.9	8.4	1.2	10.2	1.8	1.11	0.77
Drug abuse	1.1	0.7	2.4	0.8	2.6	0.6	0.2	0.2	6.93	0.09
Drug dependence	1.9	1.1	1.5	0.6	1.6	0.4	2.3	0.5	1.40	0.71
Nicotine dependence	3.2	0.7	1.9	0.9	2.9	0.5	8.6	2.2	6.05	0.13
Any alcohol or drug use disorder	18.9	4.1	15.9	1.9	42.6	6.3	24.3	3.4	4.83	0.20
Any substance use disorder	20.5	4.3	17.6	2.1	43.3	5.8	26.2	2.9	5.55	0.16

(Continued)

	Bio Bio		Metropol	Metropolitana Tara		Tarapaca		Araucania		
Disorders	%	SE	%	SE	%	SE	%	SE	χ^2	Р
Other disorders Non-affective psychosis Somatoform disorder Any CPPS disorder	1.8 3.0 28.5	1.1 1.7 5.3	2.7 3.2 26.4	1.1 1.3 2.3	0.0 2.6 51.2	0.0 0.5 4.0	0.0 2.6 35.0	0.0 0.9 5.0	5.78 0.23 4.46	0.14 0.97 0.23

Non-affective psychosis includes schizophrenia, schizophreniform disorder, schizoaffective disorder, delusional disorder, and atypical psychosis Any CPSS disorder does not include nicotine dependence or cognitive disorder; χ^2 df = 3

Table 4. The lower use of services persisted after controlling for socio-demographic variables in a logistic regression.

When urban versus rural was examined across all sites no statistical differences in the rates of disorders were noted. In addition, there were no socio-demographic differences. The sample size of the rural population was small, 203.

Discussion

Regional differences that persisted after adjusting for potential confounders persisted in the CPPS. Major depression and substance use disorders were highly prevalent in Tarapaca. The high rates of substance use disorders, especially drugs, and were not surprising as the region bordering Bolivia and Peru is heavily involved in the drug trade. The increased rates of major depression, however, could not be accounted for by substance use disorder comorbidity. The differences in rates for non-affective psychosis, although may simply be due to a type 1 error, are nonetheless surprising as the Araucania region's population and our sample has a sizable proportion of Mapuche indigenous people. The Mapuche in earlier psychiatric literature were thought to be at increased risk for psychosis [28]. The small sample size of the rural population precluded finding statistically significant differences.

The utilization of health service was lowest in Araucania and Bio Bio. This may be consistent with the inequities in availability and access that do exist in health and mental health resources across different regions of Chile. The southern half of the country is the poorest and has the least resources; therefore, the lower rates may be due to a lack of access rather than demand. For example across the regions the number of available mental health beds 2001 in the public health service per 100,000 were Araucania 2.2; Bio Bio 4.8; Metropolitana 34.2; and Tarapaca 47.2. The number of primary care physicians per 100,000 populations also was lowest in Araucania, 57.0, compared to Bio Bio with 169.7, Tarapaca with 61.6 and Metropolitana with 185.8.

It could be argued that these regional differences are simply artifact due to sampling differences. Clearly the population investigated in Tarapaca is small for a prevalence study and may have resulted in rates that may prove unstable. Another potential limitation is that the four regions were investigated sequentially, with the potential for socio-cultural influences to impact on the rates during the intervals between data collection. The high proportion of lowincome individuals in the Bio Bio sample in comparison to the other sites and in particular Aracucania, the poorest region of the country, reflects the improved economic conditions in Chile during the course of data collection and supports a cohort effect. A cohort effect, however, is highly unlikely to explain the rates of psychopathology given that data collection was obtained from Araucania last, yet it has the highest rate of major depressive disorder.

Conclusion

Regional differences across countries may exist that have both implications for prevalence rates and service utilization. Planning mental health services for population centers that span wide geographical areas based on studies conducted in a single region may be misleading, and may result in areas with high need being underserved. Psychiatric epidemiological studies that are nationally representative of developing nations are needed that have a sufficient sampling frame to examine populations believed to be at high risk and regions where increased inequities may exist. Even the most recent epidemiological studies representing Latin America [6, 29] have ignored large segments of the population, such as those countries and regions of countries with large indigenous populations or segments of the population that are very poor. Fewer studies in the region have examined service needs and none have addressed regional differences in services. Data that is more representative of the Latin American population is needed in order to improve mental health services planning and addresses the large under-estimated treatment gap.

Acknowledgment The authors wish to thank the Pan American Health Organization/World Health Organization for their technical and financial support. We also wish to acknowledge the financial support of FONDECYT (No. 90-229, 92-233, 1971315, 1990325) and

Table 3	12-Month	prevalence	rates	of DSM-III-I	R disorders	bv	region
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	Bio Bio		Metropo	litana	Tarapaca	a	Araucan	ia		
Disorders	%	SE	%	SE	%	SE	%	SE	χ²	Р
Total										
Affective disorders										
Major depressive episode	4.1	0.9	7.8	0.8	10.3	1.9	5.1	1.1	8.78	0.05
Manic episode	1.9	0.7	1.1	0.3	1.8	0.3	1.0	0.9	2.05	0.57
Dysthmia	3.0	0./	4.1	1.2	7.9	2.2	3.1	2.2	6.10	0.13
Any affective disorder	7.8	1.0	10.8	1.5	15.0	1.0	7.5	2.9	5.19	0.18
Papic disorder	0.5	0.2	0.4	0.2	11	0.0	0.6	0.6	1 2 2	0.72
Agoraphobia without papic	0.5	0.5	0.4 6.1	0.2	4.1	0.9	0.0	0.0	1.52	0.75
Generalized anxiety disorder	1.0	0.5	23	0.6	11	0.3	17	0.5	2 41	0.24
Any anxiety disorder	11.1	1.7	9.2	1.6	11.9	1.6	3.8	1.3	5.69	0.15
Substance use disorders							510		5107	0115
Alcohol abuse	2.1	0.6	1.8	0.5	5.8	1.8	3.3	0.8	4.05	0.28
Alcohol dependence	5.2	1.5	4.8	0.7	3.7	0.4	2.8	0.5	4.26	0.26
Drug abuse	0.2	0.2	0.4	0.2	1.5	0.4	0.0	0.0	3.99	0.28
Drug dependence	1.5	0.8	1.9	0.5	1.8	0.3	0.4	0.4	2.91	0.42
Nicotine dependence	2.1	0.5	1.9	0.6	5.6	0.3	4.4	1.0	4.18	0.26
Any alcohol or drug use disorder	8.4	1.6	7.9	0.9	10.7	2.0	6.2	1.2	2.84	0.43
Any substance use disorder	10.0	1.6	9.4	1.0	15.7	2.1	8.2	1.6	2.38	0.51
Other disorders										
Non-affective psychosis	1.1	0.4	1.4	0.3	0.8	0.2	0.0	0.0	8.45	0.06
Somatoform disorder	1.8	0.8	3.9	0.6	3.1	0.6	3.0	0.6	2.95	0.42
Any CPPS disorder	23.9	3.5	23.0	1.5	25.3	1./	14./	3.9	2.84	0.43
Affective disorders										
Major depressive episode	18	0.0	10.5	13	1/1 8	0.8	53	24	10.08	0.03
Manic enisode	- 1 .0 2.6	1.0	10.5	0.5	3.0	0.0	0.2	0.2	6.25	0.03
Dysthmia	43	1.0	6.9	23	8.8	17	3.9	3.2	6.92	0.12
Any affective disorder	9.6	1.8	15.2	2.2	20.0	0.9	8.2	5.3	5.72	0.15
Anxiety disorders						•••				
Panic disorder	0.5	0.3	0.6	0.3	6.6	1.3	1.0	1.2	1.37	0.71
Agoraphobia without panic	11.9	3.0	8.4	2.4	10.4	0.3	3.4	1.0	5.78	0.14
Generalized anxiety disorder	1.4	0.8	3.9	1.0	2.2	0.6	2.7	0.7	3.24	0.37
Any anxiety disorder	17.3	2.3	13.0	2.5	17.5	1.2	6.3	1.7	5.90	0.14
Substance use disorders										
Alcohol abuse	0.5	0.4	1.0	0.5	0.9	0.2	0.6	0.5	0.93	0.82
Alcohol dependence	0.9	0.5	1.3	0.7	4.2	0.2	0.0	0.0	5.16	0.18
Drug abuse	0.0	0.0	0.2	0.2	0.5	0.1	0.0	0.0	2.48	0.49
Drug dependence	1.5	0.7	2.9	0.8	2.7	0.5	0.2	0.2	8.50	0.05
Nicotine dependence	2.0	0./	2.1 4.F	0.8	8.5	0.4	2.0	0.8	1./1	0.64
Any substance use disorder	2.9 1 Q	1.0	4.5 6.0	0.0	5.1 13.3	0.4	0.0	0.7	7.90 17.77	0.00
Ally substance use disorder Other disorders	4.7	1.4	0.0	0.0	12.2	0.9	2.0	0.7	14.74	0.01
Non-affective psychosis	15	07	12	0.5	16	03	0.0	0.0	7.05	0.09
Somatoform disorder	1.9	0.7	5.0	1.1	3.6	0.7	4.2	1.8	4.50	0.23
Any CPPS disorder	23.4	2.7	24.7	2.9	28.4	0.9	12.3	5.1	2.83	0.42
Male										
Affective disorders										
Major depressive episode	3.4	1.1	4.7	1.1	5.5	3.6	5.0	0.4	1.30	0.73
Manic episode	1.1	0.9	0.7	0.3	0.5	0.6	1.8	1.6	0.68	0.88
Dysthmia	1.6	0.8	0.8	0.3	7.0	3.3	2.1	2.0	4.78	0.21
Any affective disorder	5.9	1./	5./	1.1	9.8	2.8	6./	1.3	4.46	0.23
Anxiety disorders	0.0	0.4	0.2	0.2	1.0	0.4	0.0	0.0	2.07	0.20
Panic disorder	0.0	0.4	0.2	0.2	1.0	0.4	0.0	0.0	3.07	0.39
Agoraphobia without panic Constalized anxiety disorder	1./	0.9	5.4 0.4	1./	4.5	5.8 0.0	0.9	0.9	1.94	0.59
Any anyiety disorder	1.0	0.0	0.4 17	0.2	6.1	0.0	0.5	0.4	3.26	0.58
Substance use disorders	7.7	1.0	ч./	1.0	0.1	J.J	1.1	0.9	5.20	0.57
Alcohol abuse	3.7	1.2	2.7	0.8	10.9	3.7	6.3	1.2	6.20	0.12
Alcohol dependence	9.7	2.8	8.8	1.8	3.1	1.0	5.9	0.9	3.88	0.29
Drug abuse	0.4	0.4	0.7	0.5	2.6	0.6	0.0	0.0	3.47	0.34
Drug dependence	1.5	1.2	0.8	0.5	0.9	0.2	0.6	0.8	0.33	0.95
Nicotine dependence	2.2	0.7	1.8	0.8	2.5	0.5	7.0	2.0	3.08	0.39
Any alcohol or drug use disorder	14.4	3.2	11.8	2.0	16.6	4.4	12.1	1.6	1.25	0.74
Any substance use disorder	15.4	3.2	13.3	2.1	18.2	4.0	14.1	2.8	1.07	0.78

(Continued)

Table 3 continued

	Bio Bio		Metropoli	Metropolitana Tarapaca			Araucani	a		
Disorders	%	SE	%	SE	%	SE	%	SE	χ²	Р
Other disorders Non-affective psychosis Somatoform disorder Any CPPS disorder	0.8 1.6 21.8	0.4 1.2 5.3	1.6 2.6 20.5	0.7 1.0 2.5	0.0 2.6 22.0	0.0 0.5 3.0	0.0 1.8 17.3	0.0 0.8 3.4	5.75 2.05 1.12	0.14 0.57 0.77

Non-affective psychosis includes schizophrenia, schizophreniform disorder, schizoaffective disorder, delusional disorder, and atypical psychosis Any CPSS disorder does not include nicotine dependence or cognitive disorder; χ^2 df = 3

Table 4 Mental health service utilization in the	past 6-months by region among	g those with DSM-III-R 12-month p	prevalent disorder
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	Bio Bio		Metropol	Metropolitana T		Tarapaca		Araucania		
Disorders	%	SE	%	SE	%	SE	%	SE	χ^2	Р
Any MH service Non-specialized MH service Specialized MH service Substance service	17.6 15.2 5.2 0.1	1.1 1.0 0.9 0.1	21.1 17.1 6.7 0.3	2.5 1.8 1.3 0.2	25.9 24.3 5.6 0.4	0.3 0.4 1.1 0.1	12.0 10.7 3.4 0.1	1.9 1.6 1.1 0.1	10.09 12.63 4.26 2.87	0.03 0.02 0.26 0.43

 χ^2 df = 3, MH = Mental Health

Non-Specialized MH Services = primary care physicians; Specialized Mental Health Services = inpatient or outpatient services provided by a psychiatrist or a psychologist or formal substance abuse services; Substance abuse services included inpatient and outpatient detoxification or Alcoholics Anonymous

Dirección de Investigación de la Universidad de Concepción (No. 201.087.027-1.0).

References

- Bjil RV, de Graff R, Hiripi E, Kessler RC, Kohn R, Offord DR, Ustun TB, Vicente B, Vollebergh WAM, Walters EE, Wittchen HU (2003) The prevalence of treated and untreated mental disorders in five countries. Health Affairs 22:122–133
- WHO International Consortium in Psychiatric Epidemiology (2000) Cross-national comparisons of the prevalences and correlates of mental disorders. Bull World Health Organ 78:413–25
- Weissman MM, Bland RC, Canino GJ, Garavelli C, Greenwald S, Hwu HG, Joyce PR, Karam EG, Lee CK, Lellouch J, Lepine JP, Newman SC, Rubio-Stipec M, Wells JE, Wickramaratne PJ, Wittchen HU, Yeh EK (1996) Cross-national epidemiology of major depression and bipolar disorder. JAMA 276:293–299
- Hwu HG, Yeh EK, Chang LY (1989) Prevalence of psychiatric disorders in Taiwan defined by the Chinese Diagnostic Interview Schedule. Acta Psychiatr Scand 79:136–147
- 5. Wang CH, Liu WT, Zhang MY, Yu ESH, Xia ZY, Fernandez M, Lung CT, Xu CL, Qu GY (1992) Alcohol use, abuse, and dependency in Shanghai. In: Helzer JE, Canino GJ (eds) Alcoholism in North America, Europe, and Asia. Oxford University Press, New York
- WHO World Mental Health Survey Consortium (2004) Prevalence, severity, and unmet need for treatment of mental disorders in the World Health Organization World Mental Health Surveys. JAMA 291:2581–2590
- 7. Kleinman A (1997) Rethinking psychiatry from cultural category to personal experience. Simon and Schuster, New York
- Kessler RC, McGonagle KA, Zhao S, Nelson CB, Hughes M, Eshleman S, Wittchen HU, Kendler K (1994) Lifetime and 12month prevalence of DSM-III-R psychiatric disorders in the United States. Arch Gen Psychiatr 51:8–19
- 9. Robins LN, Regier DA (eds) (1989) Psychiatric disorders in America: the epidemiologic catchment area study. Free Press, New York

- Almeida-Filho N, Mari JJ, Coutinho E, França JF, Fernandes J, Andreoli SB, Busnello ED (1977) Brazilian multicentric study of psychiatric morbidity. Methodological features and prevalence estimates. Br J Psychiatr 171:524–529
- Lewis G, Booth M (1992) Regional differences in mental health in Great Britain. J Epidemiol Community Health 46:608–611
- Dembling BP, Rovnyak V, Mackey S, Blank M (2002) Effect of geographic migration on SMI prevalence estimates. Ment Health Serv Res 4:7-12
- Harrison G, Glazebrook C, Brewin J, Cantwell R, Dalkin T, Fox R, Jones P, Medley I (1997) Increased incidence of psychotic disorders in migrants from the Caribbean to the United Kingdom. Psychol Med 27:799-806
- 14. Haukka J, Suvisaari J, Varilo T, Lönnqvist J (2001) Regional variation in the incidence of schizophrenia in Findland: a study of birth corhorts born from 1950 to 1969. Psychol Med 31:1045–1053
- Youssef HA, Kinsella A, Waddington JL (1991) Evidence for geographical variations in the prevalence of schizophrenia in rural Ireland. Arch Gen Psychiatr 48:254–255
- Marsella AJ (1998) Urbanization, mental health, and social deviancy. A review of issues and research. Am Psychol 53:624– 634
- Paykel E, Abbott R, Jenkins R, Brugha TT, Meltzer H (2003) Urban-rural mental health differences in Great Britain findings from the National Morbidity Survey. Int Rev Psychiatry 15:87– 107
- Caraveo-Anduaga JJ, Colmenares E, Saldivar GJ (1999) Morbilidad psiquiátrica en la ciudad de México: prevalencia y comorbilidad a lo largo de la vida. Salud Mental 34:62–67
- Andrade LH, Walters EE, Gentil V, Laurenti R (2002) Prevalence of ICD-10 mental disorders in a catchment area in the city of São Paulo, Brazil. Soc Psychiatr Psychiat Epidemiol 37:316– 325
- Hayashi S, Perales A, Sogi C, Warthon D, Llanos R, Novara J (1985) Prevalencia de vida de trastornos mentales en Independencia (Lima, Peru). Anales de Salud Mental 1:206-222
- Vicente B, Rioseco P, Saldivia S, Kohn R, Torres S (2002) Estudio Chileno de prevalencia de patología psiquiátrica (DSM-III-R/CIDI) (ECPP). Rev Med Chile 130:527–536

- 22. Robins LN, Wing J, Wittchen HU, Helzer JE, Babor TF, Burke J, Farmer A, Jablensky A, Pickens R, Regier DA, Sartorius N, Towle LH (1988) The composite international diagnostic interview: an epidemiologic instrument suitable for use in conjunction with different diagnostic systems and in different cultures. Arch Gen Psychiatr 45:1069–1077
- 23. American Psychiatric Association (1987) Diagnostic and statistical manual of mental disorders, DSM-III-R, 3rd edn. revised. American Psychiatric Association, Washington, DC
- 24. Sartorius N, Kuyken W (1994) Translation of health status instruments. In: Orley J, Kuyken W (eds) Quality of life assessment in health care settings, volume 1. Springer, Berlin
- 25. Vielma M, Vicente B, Rioseco P, Castro P, Castro N, Torres S (1992) Validación en Chile de la entrevista diagnóstica estandarizada para estudios epidemiológicos CIDI. Rev Psiquiatría 9:1039-1049
- 26. Pfister H, Wittchen HU (1990) CIDI core computer manual for data entry and diagnostic programmes for the composite international diagnostic interview (CIDI). World Health Organization, Geneva
- 27. Shah BV, Barnwell BG, Bieler GS (1997) SUDAAN user's manual. Release 7.5. Research Triangle Institute, Research Triangle Park, NC
- Muñoz L, Marconi J, Horwitz J, Naveillan P (1966) Crosscultural definitions applied to the study of functional psychoses in Chilean Mapuches. Br J Psychiatr 112:1205–1215
- 29. Kohn R, Levav I, de Almeida JM, Vicente B, Andrade L, Caraveo-Anduaga JJ, Saxena S, Saraceno B (2005) Mental disorders in Latin America and the Caribbean: a public health priority. Rev Panam Salud Publica 18:229-240