

Psychosis in Indigenous populations of Cape York and the Torres Strait

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Despite mental illness being a major contributor to the health gap for Aboriginal and Torres Strait Islander adults,¹ reliable prevalence and severity data are limited. Although there has been work focusing on assessment² and service delivery,³ research has been limited to a small case-control study⁴ and studies of subgroups at elevated risk.^{5,6} As a rough indicator, in the 2005–06 financial year, male and female Indigenous Australians were hospitalised for psychiatric disorders at 2.4 and 1.5 times the rate for non-Indigenous Australians, respectively.⁷ The admission rate ratios for male and female Indigenous Australians for schizophrenia and delusional disorders were 2.7 and 2.5, and for mental disorders due to psychoactive substances, 4.5 and 3.3,⁷ respectively.

In this analysis, we aimed to describe and characterise treated psychotic disorders in the Indigenous population of Cape York and the Torres Strait Islands in Far North Queensland.

Methods

Study location and population

Cape York and the Torres Strait region cover 7.5% of Queensland, with a population estimate for 2009 of 24 049 — 0.5% of Queensland's population.⁸ Extrapolating from 2006 ethnic breakdown data,⁹ the region's Indigenous population in 2009 was estimated at 15 904 (39% Aboriginal, 45% Torres Strait Islander, and 16% both Aboriginal and Torres Strait Islander), of whom 10 217 were aged 15 years and older, and constitute the denominator for the calculation of prevalence rates in this study.

Data collection

In mid 2010, we collated data on all adult Indigenous patients in the region who were either symptomatic or in remission and receiving treatment for a psychotic disorder falling within the following International Classification of Diseases, 10th revision

Abstract

Objective: To describe and characterise treated psychotic disorders in the Indigenous populations of Cape York and the Torres Strait.

Design: Cross-sectional analysis of patients with a psychotic disorder identified by treating psychiatrists.

Setting and participants: Indigenous patients aged ≥ 15 years in Cape York and Torres Strait communities receiving treatment for a psychotic disorder over 3 months in 2010.

Main outcome measures: Prevalence of psychosis diagnoses, intellectual disability, and substance use comorbidities.

Results: 171 patients were included. The prevalence rate in this population was 1.68%, higher for males (2.60%) than females (0.89%), and twice as high in the Aboriginal (2.05%) than in the Torres Strait Islander (0.95%) population. High rates of alcohol and cannabis use were found. Comorbid intellectual disability was common and more frequent among Aboriginal patients.

Conclusions: The burden of psychosis in the Indigenous population of Cape York and the Torres Strait is high. Further research is needed to understand the social determinants of these disorders and to design effective social and clinical measures to alleviate this burden.

groups: organic-related (F06); substance use-related (F10–19); schizophrenia-related (F20–22); acute and transient psychoses (F23); and mood disorder-related (F25, F30–33). Participants were patients of the Rural and Remote Area Mental Health Service (Queensland Health) and were assessed by one of two of the authors (EMH and BDG) who have been psychiatrists in this region for 18 and 10 years, respectively. These clinicians review all cases of psychosis in the region, and patients were identified through the service's clinical database.

All information included in this study is routinely collected. Permission to collate for review was obtained from the Queensland Health Data Custodian and was approved by the Cairns Base Hospital Human Research Ethics Committee.

Although extensive data were collected, those relevant to this article are demographics, diagnoses, and clinicians' judgements regarding whether alcohol and cannabis were significant to the onset or current psychiatric condition, and the presence of intellectual disability.

Statistical methods

Prevalence rates were calculated by sex, age and ethnicity. Because a small

proportion of the sample identified as being both Aboriginal and Torres Strait Islander, prevalence rates by ethnicity were calculated in two ways. First, we removed data from participants who identified as being both Aboriginal and Torres Strait Islander from both the sample and population. Second, for a more conservative estimate, we collapsed the data from the "both" category into the Aboriginal or Torres Strait Islander categories at the sample and population levels using the proportions observed in the population.

This study is primarily descriptive, intending to highlight prevalence rates, between-group differences in prevalence rates, and a selection of within-sample associations that may provide clues as to why the rates are high. As such, univariate statistical analyses were used sparingly, and no attempt was made to explain the magnitude of the prevalence rates. The statistical significance of between-group differences was established using χ^2 tests with $P=0.05$ (two-sided).

Results

Of the 171 patients included in the study, 124 were reviewed or assessed during routine clinical community visits over 3 months in 2010 (72.5%).

1 Prevalence of psychotic disorder in the Indigenous populations (≥ 15 years old) of Cape York and the Torres Strait stratified by ethnicity, sex, and age group*

	Sample, no. (%)	Population, %	Prevalence, %
Ethnicity			
Aboriginal	108 (63.2%)	38.5%;† 46.1%‡	3.23%;† 2.05%‡
Torres Strait Islander	58 (33.9%)	45%;† 53.9%‡	1.50%;† 0.95%‡
Sex			
Male	124 (72.5%)	47.8%	2.60%
Female	47 (27.5%)	52.2%	0.89%
Age group			
15–29 years	54 (31.6%)	24.4%	1.39%
30–39 years	57 (33.3%)	14.3%	2.51%
≥ 40 years	60 (35.1%)	25.1%	1.51%

* All within-category comparisons are statistically significant at $P < 0.005$ level, youngest – oldest not significant. † "Both Aboriginal and Torres Strait Islander" category removed from calculations. ‡ "Both Aboriginal and Torres Strait Islander" category collapsed. ◆

2 Prevalence of psychotic disorders by sex and ethnicity*

Sex	Aboriginal (n = 110)	Torres Strait Islander (n = 61)	χ^2 ; P
Male (n = 124)	3.59%	1.57%	$\chi^2_1 = 20.2$; $P < 0.001$
Female (n = 47)	1.23%	0.67%	$\chi^2_1 = 2.53$; ns
χ^2 ; P	$\chi^2_1 = 26.5$; $P < 0.001$	$\chi^2_1 = 9.03$; $P = 0.003$	

ns = not significant. * "Both Aboriginal and Torres Strait Islander" category collapsed. ◆

Seventeen were discussed with staff but not seen (9.9%). Of the other 30 (17.5%), three were in forensic detention (1.8%), six in prison (3.5%), four in hospital (2.3%), and five in other institutional care (psychiatric rehabilitation, nursing home, acquired brain injury unit or supported accommodation) (2.9%). The remaining 12 included those who had recently moved or been lost to follow-up (7.0%). Previously diagnosed patients in remission and not receiving treatment were excluded.

The 171 patients included in the study were aged between 17 and 68 years. Primary diagnoses were: schizophrenia-related (106; 62.0%), substance use-related (39; 22.8%), mood disorder-related (14; 8.2%), organic-related (6; 4%) and acute and transient psychoses (6; 3.5%). The overall treated point prevalence for psychotic disorders for Indigenous residents of this region aged 15 years and older was 1.68%.

In our sample, some patients identified as being of both Aboriginal and Torres Strait Islander descent (3% v 16% for the region). Prevalence rates by ethnicity with the "both" category removed and proportionally collapsed are shown in Box 1.

Sex by ethnicity stratified prevalence rates using the collapsed approach are shown in Box 2. Box 3 provides information about the patients' histories of hospitalisation (previous 12 months) and incarceration (lifetime), and whether they received depot medication. Box 3 also includes clinical judgements about compliance, whether alcohol or cannabis was currently used at clinically significant levels, and whether their use contributed to the onset of the psychotic disorders.

The prevalence of intellectual disability in the sample and the magnitude of their ethnicity-related

differences presented in Box 3 were such that it was thought necessary to conduct further analyses on the 46 patients who were identified as having an intellectual disability. The results of these analyses show that the difference in proportions of Aboriginal and Torres Strait Islander patients identified with an intellectual disability was statistically significant (Box 4). A statistically significant difference was also revealed when this analysis was replicated with male patients only; but not with female patients only. There were no other statistically significant differences in intellectual disability between males and females.

Discussion

The treated prevalence rate for psychotic disorders in this Indigenous population is high, particularly among males and people of Aboriginal descent. Unfortunately, there is a dearth of articles that provide prevalence rates of psychotic disorders in the Australian context. Two of the more contemporary articles claim prevalence rates of about one-third of that calculated in this study. A study conducted at four urban centres, one each situated in Queensland (period prevalence rate, 0.42%), Victoria (0.59%), Western Australia (0.69%) and the Australian Capital Territory (0.39%), found a period prevalence rate (1 month) for psychotic disorders of 0.47%.¹⁰ A systematic review of 118 Australian and international studies found a period prevalence rate (1 to 12 months) for schizophrenia only of 0.55%, and a prevalence rate of 0.43% in rural centres.¹¹ Acknowledging the limitations of comparison with these

3 Proportions of sample that are affected by selected attributes at the sample level and stratified by ethnicity and sex

	Sample	Ethnicity		Sex	
		Aboriginal	Torres Strait Islander	Male	Female
Alcohol: role in onset of disorder	48.8%	50.9%	45.1%	55.2%	32.6%
Alcohol: current clinical impact	18.7%	15.7%	22.4%	22.6%	8.5%
Cannabis: role in onset of disorder	54.0%	50.5%	58.8%	60.7%	37.0%
Cannabis: current clinical impact	32.2%	31.5%	32.8%	37.9%	17.0%
Hospitalisation (previous 12 months)	32.7%	34.3%	29.3%	29.8%	40.0%
Significant compliance problems	42.0%	42.6%	39.3%	45.8%	29.5%
Medication: depot administration	59.6%	69.4%	41.4%	62.8%	51.1%
Incarceration (lifetime)	31.8%	43.9%	10.3%	42.5%	4.4%
Comorbid intellectual disability	26.9%	38.9%	6.9%	29.0%	21.3%

4 Intellectual disability, stratified by ethnicity and sex

Sex	Ethnicity		$\chi^2_1; P$
	Aboriginal	Torres Strait Islander	
	<i>n</i> = 42	<i>n</i> = 4	$\chi^2_1 = 17.72; P < 0.001$
Male	<i>n</i> = 36	34	$\chi^2_1 = 16.6; P < 0.001$
Female	<i>n</i> = 10	8	$\chi^2_1 < 1.0; ns$
$\chi^2_1; P$	$\chi^2_1 < 1; ns$	$\chi^2_1 = 1.16; ns$	$\chi^2_1 < 1; ns$

ns = not significant. ◆

studies, the elevated prevalence rates in the Indigenous populations described here are stark.

Alcohol and cannabis use appear common in this clinical population, and have been shown elsewhere to have psychiatric implications for Indigenous populations. Research in a remote WA Aboriginal population demonstrated that heavy alcohol use was associated with psychotic symptoms.¹² However, comparison to this survey's findings may be complicated by differences between Cape York and the Torres Strait, including recent legislative restrictions on alcohol in Cape York. Although this would not have been relevant to the onset of psychosis (which preceded the restrictions for most patients), it has reduced quantities available since, and thus affected the impact on current clinical status. Widespread cannabis use is relatively recent, and the results are consistent with those in remote Northern Territory communities: cannabis use was more common among males (70%) than females (20%) (with 90% of users using heavily¹³), and was associated with psychotic⁵ and depressive symptoms.¹⁴

Nationally, 7.7% of Aboriginal and Torres Strait Islander people over 15 years of age have an intellectual disability,¹⁵ and roughly 57% of Australians with intellectual disability develop a psychiatric disorder.¹⁶ The prevalence of intellectual disability in our sample was 26.9%. This rate is substantially higher than that found in a WA study, which reported a prevalence of intellectual disability in a psychiatric sample of 1.8%.¹⁷ A review of admissions to the Cairns Base Hospital Acute Mental Health Ward found "cognitive difficulties" noted in the charts of 22% of Indigenous admissions in the 2004–05 financial year.¹⁸ Our study also showed a substantial difference between the rates of intellectual disability in the

Aboriginal (38.9%) and Torres Strait Islander (6.9%) patients. Evidence supporting acquired rather than genetic explanations for these differences have been explored elsewhere.¹⁹

Our study revealed several important findings, but is constrained by some methodological limitations. The most obvious of these is the reliance on clinical judgements, the potential for interrater reliability confounds, and potential effects of culturally informed biases (including potential for misdiagnosis among Indigenous patients with autism and intellectual disability²⁰). However, there is evidence of good agreement between skilled clinicians for the diagnosis of psychosis among patients with intellectual disability,²¹ and in our study, all judgements were made by psychiatrists with shared responsibility for patients for many years. Although this does not establish validity or reliability, it makes it more likely that case ascertainment and interrater reliability are high, and as accurate as can be collected using applied clinical methods.

This study has corroborated clinical impressions of a high burden of psychosis in the Indigenous populations of Cape York and the Torres Strait, particularly male Aboriginal Australians. Our finding is supported by hospitalisation data showing that psychotic disorders are common and increasing in the Indigenous population of Far North Queensland.⁸ These disorders are associated with substance misuse and a surprisingly high rate of intellectual disability. The findings indicate an urgent need for further research in this region, to extend this research to other Indigenous populations, to identify causal and perpetuating factors, and to develop effective social and clinical measures to alleviate this burden.

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