



# Concurrent and simultaneous polydrug use: latent class analysis of an Australian nationally representative sample of young adults

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**Background:** Alcohol use and illicit drug use peak during young adulthood (around 18–29 years of age), but comparatively little is known about polydrug use in nationally representative samples of young adults. Drawing on a nationally representative cross-sectional survey (Australian National Drug Strategy Household Survey), this study examines polydrug use patterns and associated psychosocial risk factors among young adults ( $n = 3,333$ ; age 19–29).

**Method:** The use of a broad range of licit and illicit drugs were examined, including alcohol, tobacco, cannabis, cocaine, hallucinogens, ecstasy, ketamine, GHB, inhalants, steroids, barbiturates, meth/amphetamines, heroin, methadone/buprenorphine, other opiates, painkillers, and tranquilizers/sleeping pills. Latent class analysis was employed to identify patterns of polydrug use.

**Results:** Polydrug use in this sample was best described using a 5-class solution. The majority of young adults predominantly used alcohol only (52.3%), alcohol and tobacco (34.18%). The other classes were cannabis, ecstasy, and licit drug use (9.4%), cannabis, amphetamine derivative, and licit drug use (2.8%), and sedative and alcohol use (1.3%). Young adult males with low education and/or high income were most at risk of polydrug use.

**Conclusion:** Almost half of young adults reported polydrug use, highlighting the importance of post-high school screening for key risk factors and polydrug use profiles, and the delivery of early intervention strategies targeting illicit drugs.

**Keywords:** young adults, polydrug use, latent class analysis, cluster, risk and protective factors, simultaneous

## INTRODUCTION

Young adulthood (18–29 years) is a high risk time for the use and misuse of licit and illicit substances, relative to other age groups (1). Alcohol is the most common drug used (86%) in this group, with 55% of young adult Australians drinking at levels that are at risk of alcohol-related injury arising from a single occasion in the past month (2). Though decreasing over recent decades, approximately 24% of young Australians smoke tobacco (2). The use of illicit drugs peaks in young adulthood, with rates two to three times higher than that of the general population (2). The use and misuse of individual drugs are associated with substantial costs to individuals, communities, and health systems (3–6).

In recent years there has been increased research attention on patterns of polydrug use [defined as the consumption of more than one drug during a specific time period; (7)] among young adults. Much of this research has been done on relatively small convenience samples (e.g., street-based, drug injecting, emergency room, and club patrons), where multiple drugs are frequently used

consecutively or simultaneously, often for their perceived counter-acting or complementary effects (8–15). These targeted studies are important because they highlight the near universal patterns of polydrug use that occur in specific populations (9, 10), and the substantially higher risks to mental and physical health faced by polydrug users compared to single drug users (16–18). However, polydrug use in specific populations tells us little about rates of polydrug use in wider populations.

Comparatively few studies have examined young adult polydrug use based on large scale population-based samples. Extant research has been conducted on younger Australians [less than 18 years; (19)], Latin Americans [mean age 35; (20)], adult American populations [mean age 46; (16)], adult populations from Great Britain [mean age 43 years; (21)], and large scale Australian twin registry studies [mean 30 years; (17)]. Among these studies, there is variation in the window of time used to define polydrug use. Some studies have defined polydrug use as the use of more than one drug in a lifetime (16, 17, 19), in the past year (21), and in the

past month (20). The narrower the window of time used to define polydrug use, the closer the potential proximity of use of different drug types. However, in large scale population studies, this results in smaller cell sizes for individuals reporting specific combinations of drug use, and this creates challenges for established procedures like latent cluster analysis, which rely on minimum cell sizes to reliably detect clusters. In part because of these statistical power challenges, and the inherent challenges in assessing temporally linked drug taking events in large scale surveys, most such studies utilize wide windows of time (most commonly lifetime prevalence) to detect polydrug use.

The first goal of the present study was to evaluate the nature and extent of polydrug use in a nationally representative population of Australian young adults (18–29 years of age) using two temporal windows to define polydrug use. The first was polydrug use in the previous year [hereafter termed *concurrent polydrug use*, consistent with earlier research; (22)], and the second was *simultaneous use*, a dimension rarely assessed in large scale population studies. One-year temporal windows for assessing polydrug use have the advantage of capturing developmentally sensitive periods, relative to lifetime prevalence, but are likely to retain statistical power to determine clusters, relative to shorter temporal windows. Simultaneous use potentially yields an understanding of how various combinations of drugs are used for their synergistic properties including the enhancement of specific effects and the suppression of unwanted effects ones (23, 24). A key concern about polydrug use is that the effect of individual drugs are often compounded, and harmful physiological effects accumulate in the body (23–26). We sought to describe the extent to which simultaneous use characterized usage patterns in young adults with recent experience in the use of different substances. The second goal of this study was to examine demographic and broad socio-emotional correlates of polydrug use clusters. Compared to adolescence, the lives of young adults often involve increased disposable income, heightened social independence, and fewer societal restrictions on the use of certain drugs (27, 28). In this study we explored the association of drug use clusters with socioeconomic status (29, 30), educational level, peer substance use (31), and mental health (16, 21).

The present study was based on the National Drug Strategy Household Survey (NDSHS), which is unique in that it is one of the few nationally representative studies that enables an examination of polydrug clusters, and an exploration of simultaneous drug use. The study employs latent class analysis (LCA), which is a probabilistic method of determining the size and structure of drug use profiles. This analytic method has rarely been used in large scale population research on young adults, but has been extensively used with other age groups (16, 17, 21). The study is also a rare examination of how a comprehensive range of drugs, including alcohol and tobacco, cluster together in young Australians.

## MATERIALS AND METHODS

### SAMPLE

The sample consisted of 3,333 participants in the 2007 NDSHS (32) who were aged between 19 and 29 years (42.2% males, mean age = 23.78, SD = 3.42). Of this initial sample, 322 (9.67%) participants were excluded due to non-response for any of the

drug-related items and four participants were excluded as they reported using a fictitious drug. The final sample size for the LCA was 3011. Among this sample, 292 participants had one or more missing data in the covariates. Full information maximum likelihood estimation was used to handle the missing data. A supplementary analysis with complete cases only was conducted to evaluate the robustness of the results.

### Measures

**Concurrent drug use.** This measure was based on NDSHS drug use frequencies for a broad range of drugs used in the previous 12 months. Tobacco use was assessed with the item “How often do you now smoke cigarettes, pipes, or other tobacco products?” with responses of “daily,” “at least weekly,” “less often than weekly,” “not at all but I have smoked in the last 12 months,” and “not at all and I have not smoked in the last 12 months.” These responses were recoded as 0 “No smoking in the last 12 months” and 1 “Smoking in last 12 months.” Alcohol consumption was measured using a single, yes/no response item: “Have you had an alcoholic drink of any kind in the last 12 months?” Yes/no response items were also used to assess for use of painkillers, tranquilizers/sleeping pills, steroids, barbiturates, and meth/amphetamine: “Have you used (drug type) for non-medical purpose in the last 12 months?” Consumption of all other drugs, including cannabis/marijuana, heroin, methadone/buprenorphine, cocaine, hallucinogens, ecstasy, ketamine, GHB, and inhalants, was assessed using yes/no questions: “Have you used (drug type) in the last 12 months?”

**Simultaneous drug use.** Simultaneous drug use was examined in a separate series of questions in which respondents reported using a drug in the past 12 months (except tobacco and alcohol), using the question “Which of the following did you use at the same time, on at least one occasion that you used (drug type)?” With a list of possible drugs (includes alcohol but not tobacco). This question identified pairs of drugs that are being used simultaneously at least once in the past 12 months [similar to previous studies, e.g., Ref. (26, 33, 34)]. To increase reliability of simultaneous drug use coding, it was coded as present only when (i) both drugs were used in the past 12 months, and (ii) simultaneous use were also reported in both drug types (e.g., amphetamines being used simultaneously with cannabis, and cannabis being used simultaneously with amphetamines).

**Demographic and psychosocial correlates.** The 10-item Kessler Psychological Distress Scale (35) was used to measure depressive and anxiety symptoms in the past 4 weeks [e.g., about how often did you feel tired out for no good reason? (5-point scale ranging from 1 “None” to 5 “All,” alpha = 0.86)]. Peer drug use was accessed with the item “About what proportion of your friends and acquaintances use any of the following? Alcohol/Tobacco/Cannabis” (same 5-point response scale), and was recoded into 0 “Less than half” or 1 “Half or more.” General health was measured with the item “In general, would you say your health is . . .” (1 “Excellent” to 5 “Poor”). Due to the low frequency of participants reporting poor health ( $n = 35$ , 1.16% of the analysis sample), the categories “fair” and “poor” were combined to form a single category. Demographic variables included in the analysis were:

sex (0 “male,” 1 “female”); marital status (1 “never married,” 2 “widowed/divorced/separated,” 3 “married/defacto”); employment status (1 “employed/self-employed,” 2 “unemployed,” 3 “home duties,” 4 “student,” 5 “retired/unable to work/other”); high school completion (0 “completed,” 1 “not completed”); income levels (0 “\$41,600 or above,” 1 “\$13,000–41,599,” 2 “\$12,999 or below,” 3 “prefer not to say/don’t know”); socioeconomic indices for areas (SEIFA, 1 “Least advantaged/1st quartile” to 2 “Most advantaged/fourth quartile” = 4), and regionality (1 “major cities,” 2 “inner regional,” 3 “outer regional,” 4 “remote/very remote”) (36).

### Procedure

There were 23,356 respondents from households across all states and territories of Australia who participated in the NDSHS and 3,333 were in the targeted age range (between 19 and 29 years). Participants were randomly selected using a stratified design based on statistical local areas (36). The survey involved two modes of delivery – drop and collect (19,818 respondents) and computer-assisted telephone interview (CATI) (3,538 respondents). For each household, the respondent was the household member aged >12 years whose birthday was next to occur in the family. Further information on survey procedures is available in AIHW reports (37). The survey was approved by the Australian Institute of Health and Welfare Ethics Committee. Informed consent was obtained from all subjects. Access to these survey data was approved by the Australian Social Science Data Archive and by The University of Queensland Human Research Ethics Committee.

### Analysis

To identify patterns of polydrug use, LCA was performed on the last year use of nine drugs: alcohol, tobacco, cannabis, ecstasy, meth/amphetamine, pain-killer, tranquilizers/sleeping pills, cocaine, and hallucinogens. LCA is a technique that identifies sub-classes within a large population based on similarity of response to a set of measured variables. This technique is characterized by two sets of parameters: (1) The estimated proportion of each class in the population, and (2) the probability of an individual in a particular class using a certain drug.

Determination of number of classes was based on a number of fit criteria (38). First, the Bayesian Information Criterion (BIC) (39), Sample Size Adjusted Bayesian Information Criterion (SSABIC) (40), and the Akaike Information Criterion (AIC) (41) were compared across models with different number of classes. A lower value of these information criteria indicates a better balance model parsimony and model fit. Second, the Lo–Mendell–Rubin adjusted likelihood ratio test (42) was used to compare the fit of a model with  $k$  class with a model with  $k-1$  class. A significant  $p$ -value ( $p < 0.05$ ) from this test indicates a  $k$  class model fits the data better than a  $k-1$  class model. Third, the number of significant bivariate residuals was used to assess the validity of the local independent assumption of LCA. A large number of significant bivariate residuals indicates that this assumption is severely violated. Finally, the average posterior probabilities were used to evaluate the classification quality. Model fitting began with a one-class solution, and the number of classes was increased successively up to a 6-class solution. Once the optimal number of classes was determined, covariates were added to the LCA to examine their

associations with the latent class membership. In this study, data was prepared with STATA 11 (43) and analyses were performed with Mplus 6.01 (44).

## RESULTS

### LATENT CLASS ANALYSIS

Model fit statistics for 1–6-class solutions are presented in **Table 1**. A 3-class solution attained the lowest value of BIC, and a 5-class solution attained the lowest value of AIC and SSABIC. Results from the LMR-LRT suggested that the fit of the 4-class model was significantly better than a 3-class model but not worse than the 5-class model. Simulation suggested that the performance of SSABIC and LMR-LRT were similar and superior to other statistics criteria (45, 46). SSABIC suggested the 5-class solution was optimal but LMR-LRT suggested the 4-class solution was optimal. Since they pointed to different solutions and other fit statistics of these two solutions were very close, both were examined on the basis of interpretability (44). The 4-class solution identified a class with cannabis, amphetamine derivatives, and licit drug use but the 5-class model subdivided this class into two – a class with cannabis, amphetamine derivatives, and licit drug use and a class that primarily used sedatives and alcohol. This classification was consistent with previous findings (26). The 5-class solution was chosen as the optimal solution as it yielded classification that was clearly distinct and interpretable, and had adequate class sizes with high average posterior probabilities (>0.80).

Each of the five classes was described below using the probabilities of drugs use in the past 12 months (see **Figure 1**). For heuristic purposes, the nomenclature adopted for each class is based on the type and range of substances with posterior probabilities greater than 0.65.

**Class 1:** Participants in this class were predominantly alcohol users (0.80 probability of alcohol use), with a small probability (0.10) of tobacco use and nearly zero probabilities of other drug use. This class was labeled *Alcohol only*, and the prevalence estimate of this class was 52.3% ( $n = 1572$ ).

**Class 2:** These participants reported universal alcohol use (0.98), high probability of tobacco use (0.66), moderate probability of cannabis use (0.31), and negligible probabilities of other drug use (below 0.05). This class was labeled *Alcohol and tobacco*, and the prevalence estimate of this class was 34.18% ( $n = 1028$ ).

**Class 3:** Participants in this class universally reported using alcohol (1.00), very high probabilities of tobacco, cannabis, and ecstasy use (probabilities 0.70 or higher), moderate probability of meth/amphetamine and cocaine use (0.22–0.38) and low probabilities of hallucinogens and other drugs. This class was labeled *Cannabis, ecstasy, and licit drug use*, and the prevalence estimate of this class was 9.44% ( $n = 284$ ).

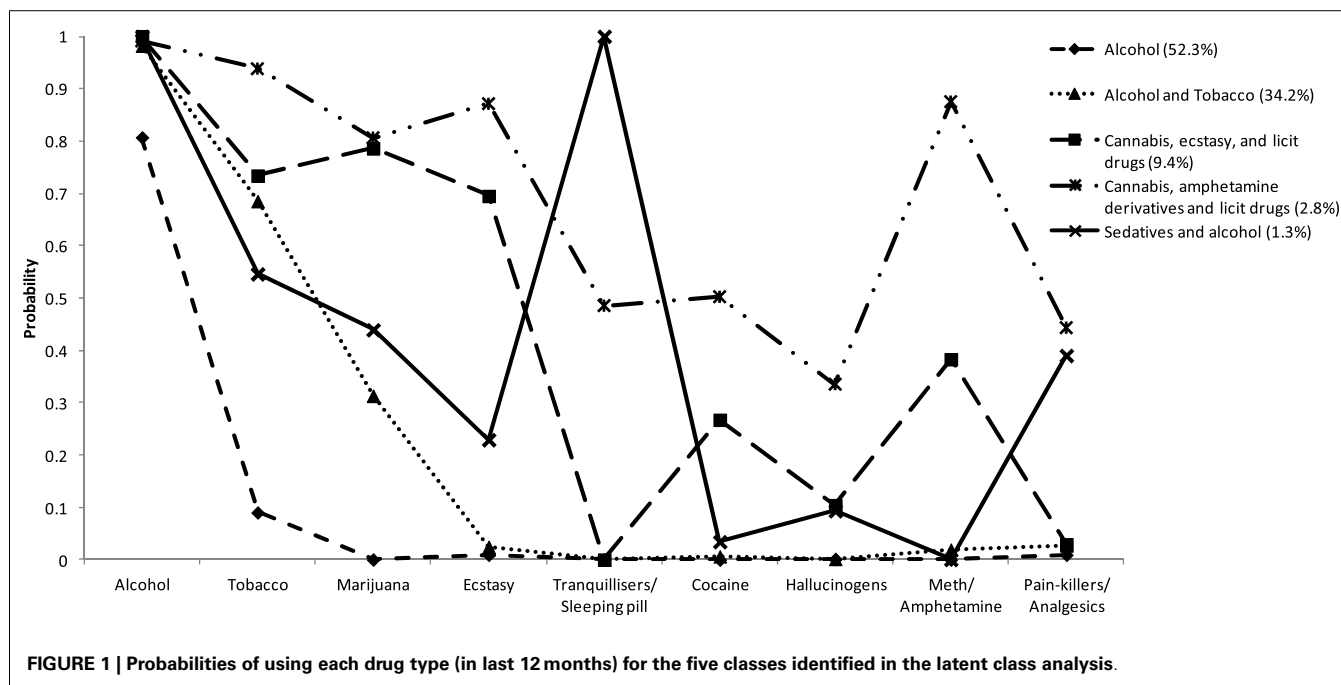
**Class 4:** This class was characterized by universal use of alcohol (0.99), very high probabilities of tobacco, cannabis, ecstasy, and meth/amphetamine use (0.70–0.89) and moderate to high probabilities of all other drug use. This class was labeled *Cannabis, amphetamine derivatives, and licit drug use*, and the prevalence estimate of this class was 2.79% ( $n = 83$ ).

**Class 5:** These participants reported universal alcohol and tranquilizer use (1.00), moderate to high probabilities of tobacco,

**Table 1 | Fit statistics from 1 to 6-class models.**

Class	Likelihood	df	AIC	BIC	SSABIC	p-Value (LMR-LRT)	No. of Sig. residuals
1	-7969.24	9	15956.47	16010.55	15981.95	N/A	98
2	-6859.55	19	13757.09	13871.26	13810.89	0.00	31
3	-6718.08	29	13494.15	13668.40	13576.26	0.00	4
4	-6683.10	39	13444.20	13678.54	13554.60	0.01	0
<b>5</b>	<b>-6658.02</b>	<b>49</b>	<b>13414.03</b>	<b>13708.46</b>	<b>13552.77</b>	<b>0.14</b>	<b>0</b>
6	-6644.71	59	14307.41	13761.93	13574.46	0.15	1

*Bolded text indicates the chosen model based on outlined criteria.*



cannabis, ecstasy, and pain-killer/analgesic use (0.23–0.54), and low probabilities of using other drugs (below 0.10). This class was labeled *Sedative and alcohol use*, and the prevalence estimate of this class was 1.31% ( $n = 39$ ).

**Table 2** shows the odd ratios and the associated 95% confidence intervals of covariates for class membership. Relative to the *Alcohol* class, being male, being older, not completing high school and having more income were significantly associated with membership in the *class 2 (alcohol and tobacco)*, *class 3 (cannabis, ecstasy, and licit drug use)*, and *class 4 (cannabis, amphetamine derivatives, and licit drug use)* ( $ps < 0.05$ ). Although being older was significantly associated with *class 5 (Sedatives and alcohol)*, sex, high school completion, and having more income were not. Young adults without partners and not living in the regional areas were also significant predictors of *class 3* and *class 4* membership. Individuals living in high socioeconomic status areas or those where were unemployed were significantly more likely to be in *class 3* ( $p < 0.05$ ).

Suboptimal general health was associated with *classes 2, 3, and 4* ( $ps < 0.05$ ), whilst psychological distress was significantly associated with the *classes 4 and 5* ( $ps < 0.05$ ). Both peer alcohol

and tobacco use were predictors of *classes 2, 3, and 4* ( $ps < 0.05$ ), while peer cannabis use was associated with *classes 2–5* ( $ps < 0.05$ ). Peer alcohol use was a stronger predictor of membership in *class 4* (95% CI: 4.37–53.33) than *class 2* (95% CI: 1.52–2.34) and peer cannabis use was a stronger predictor of membership of *class 4* (95% CI: 8.75–21.90) compared to all other classes.

Since there were missing data in the covariate variables and full information maximum likelihood estimation was used to handle the missing data, a supplementary analysis using participants with complete cases was done. The results were essentially the same – there were only negligible differences in the odds ratios estimates (less than 1% difference in most of the odds ratios). Therefore, the results obtained from full information maximum likelihood estimation were robust.

#### SIMULTANEOUS DRUG USE

Results for the analysis of simultaneous use (see **Table 3**) indicated that a large proportion of concurrent users engage in simultaneous drug use, especially combined use of alcohol and cannabis (18.45% of analysis sample). The most common other drug pairs included alcohol – ecstasy (9.54%), alcohol – meth/amphetamine (5.91%), cannabis – ecstasy (4.99%), and alcohol – cocaine (3.69%). The

**Table 2 | Relative risk ratio estimates and 95% confidence intervals from the multinomial logistic regression with the Alcohol class as the reference group.**

	Class 2: alcohol and tobacco		Class 3: cannabis, ecstasy, and licit drug use		Class 4: cannabis, amphetamine derivatives, and licit drug use		Class 5: sedatives and alcohol	
	RRR <sup>a</sup>	95% CI	RRR <sup>a</sup>	95% CI	RRR <sup>a</sup>	95% CI	RRR <sup>a</sup>	95% CI
Age								
Continuous years	1.03*	(1.01–1.05)	1.07**	(1.03–1.12)	1.10*	(1.02–1.18)	1.14*	(1.01–1.28)
Sex (Ref: Male)								
Female	0.83**	(0.73–0.94)	0.69**	(0.54–0.89)	0.39***	(0.25–0.61)	0.96	(0.50–1.85)
High school (Ref: Completed)								
Not completed	1.44***	(1.24–1.68)	1.51**	(1.13–2.01)	2.03**	(1.32–3.11)	1.21	(0.51–2.87)
Marital status (Ref: Non-partnered)								
Partnered	0.95	(0.82–1.09)	0.56***	(0.43–0.75)	0.45**	(0.27–0.77)	0.46	(0.21–1.02)
Employment status (Ref: Self-employed/employed for wages)								
Students	0.94	(0.77–1.14)	1.09	(0.72–1.63)	1.78	(0.82–3.83)	1.15	(0.33–4.01)
Unemployed	1.25	(0.92–1.69)	2.03*	(1.16–3.57)	1.72	(0.75–3.94)	1.12	(0.23–5.46)
Other	1.02	(0.83–1.26)	0.90	(0.57–1.43)	1.24	(0.59–2.61)	0.36	(0.08–1.69)
Income level (Ref: \$41,600 or above)								
\$13,000–\$41,599	0.98	(0.84–1.15)	0.79	(0.59–1.05)	0.60*	(0.36–1.00)	1.14	(0.49–2.67)
\$12,999 or below	0.75*	(0.60–0.94)	0.49**	(0.31–0.76)	0.25**	(0.11–0.55)	0.29	(0.05–1.75)
Prefer not to say/don't know	0.64***	(0.50–0.80)	0.32***	(0.19–0.53)	0.21**	(0.09–0.51)	0.27	(0.05–1.55)
Socioeconomic status of area (Ref: Least advantaged)								
2nd quartile	0.94	(0.78–1.12)	1.06	(0.72–1.55)	1.04	(0.53–2.01)	1.01	(0.37–2.74)
3rd quartile	0.96	(0.81–1.14)	1.22	(0.85–1.75)	1.19	(0.65–2.20)	1.33	(0.52–3.38)
4th quartile	1.02	(0.86–1.22)	1.44*	(1.02–2.05)	1.55	(0.86–2.77)	1.33	(0.49–3.64)
Remoteness (Ref: Major cities)								
Inner regional	1.07	(0.91–1.26)	0.79	(0.56–1.11)	0.49*	(0.27–0.90)	0.75	(0.30–1.89)
Outer regional	1.04	(0.86–1.27)	0.54***	(0.36–0.81)	0.47*	(0.22–0.99)	1.20	(0.43–3.36)
Remote/very remote	0.85	(0.62–1.15)	0.46*	(0.25–0.86)	0.64	(0.24–1.70)	0.50	(0.07–3.78)
Depressive symptoms								
Continuous K10 scores	1.01	(1.00–1.02)	1.01	(0.99–1.04)	1.09***	(1.05–1.12)	1.10***	(1.05–1.16)
General health (Ref: Excellent)								
Very good	1.15	(0.98–1.34)	0.94	(0.67–1.34)	1.05	(0.54–2.07)	1.26	(0.45–3.57)
Good	1.54***	(1.30–1.81)	1.55*	(1.10–2.18)	2.29*	(1.21–4.36)	2.53	(0.90–7.12)
Fair/poor	1.60***	(1.23–2.09)	1.61	(0.97–2.66)	1.94	(0.89–4.22)	1.86	(0.46–7.47)
Peer alcohol use (Ref: Less than half)								
More than half	1.88***	(1.52–2.34)	3.81***	(2.02–7.18)	15.27***	(4.37–53.33)	1.93	(0.42–8.92)
Peer tobacco use (Ref: Less than half)								
More than half	1.88***	(1.66–2.13)	2.02***	(1.55–2.64)	3.65***	(2.14–6.22)	1.29	(0.67–2.50)
Peer cannabis use (Ref: Less than half)								
More than half	2.43***	(2.02–2.91)	8.61***	(6.56–11.31)	13.85***	(8.75–21.90)	3.45***	(1.76–6.77)

\*\*\**p* < 0.001; \*\**p* < 0.01; \**p* < 0.05.

<sup>a</sup>RRR, relative risk ratio.

results also showed that a very high proportion of concurrent users of two specific drugs also reported using those specific drugs simultaneously. This was especially notable for alcohol and cannabis, alcohol and ecstasy, and alcohol and meth/amphetamine (greater than 90% of concurrent users were simultaneous users for each of these drug pairs).

## DISCUSSION

The primary aim of this study was to examine patterns of poly-drug use (last 12 months) across a broad range of licit and illicit substances in a nationally representative sample of young adults. Over 25% of young adults in this study reported using drugs other than alcohol in the past 12 months. A 5-class solution resulted,

**Table 3 | Percentage of participants reporting concurrent use, simultaneous use, and proportion of concurrent users who were also simultaneous users.**

Top 10 simultaneous drug combinations	% Concurrent use (use of these two drugs at an time in the last 12 months)		% Simultaneous use (use of two specific drugs simultaneously)		Percentage of concurrent users also report simultaneous use (%)
	All (%)	95% CI	All (%)	95% CI	
Alcohol and cannabis	20.72	(19.27, 22.17)	18.45	(17.06, 19.84)	89
Alcohol and ecstasy	10.54	(9.44, 11.64)	9.54	(8.49, 10.59)	91
Alcohol and meth/amphetamine	6.52	(5.73, 7.51)	5.91	(5.07, 6.76)	91
Cannabis and ecstasy	7.42	(6.48, 8.35)	4.99	(4.21, 5.77)	67
Cannabis and meth/amphetamine	5.02	(4.24, 5.80)	4.09	(3.38, 4.80)	81
Alcohol and cocaine	4.16	(3.44, 4.87)	3.69	(3.02, 4.37)	89
Meth/amphetamines and ecstasy	4.69	(3.93, 5.45)	3.23	(2.59, 3.86)	69
Cannabis and cocaine	3.19	(2.56, 3.82)	1.76	(1.29, 2.23)	55
Cocaine and ecstasy	3.09	(2.47, 3.71)	1.63	(1.18, 2.08)	53
Alcohol and painkillers	3.22	(2.59, 3.86)	1.50	(1.06, 1.93)	47

A participant can display more than one combination of polydrug use.

with all classes containing a very high probability of alcohol use, and classes named by the unique feature of each cluster. The classes were: *Alcohol only* (52.3%) *Alcohol and tobacco* (34.2%), *Cannabis, ecstasy and licit drug use* (9.4%), *Cannabis, amphetamine derivatives, and licit drug use* (2.8%), and *Sedatives and alcohol use* (1.3%). Additional analysis indicated that when young adults reported concurrent use of drug pairs (in the last 12 months), there was a very high probability (>90%) that the respective drugs were used simultaneously.

While alcohol use was evident in every cluster, the groups showed unique features in both the prominence of particular drug types and the breadth of drugs used. Class 2 was characterized by high probabilities of alcohol and tobacco use. Class 3 was characterized by a limited breadth of drugs used, and the use of two illicit drug types – cannabis and ecstasy. Class 4 was characterized by a similar breadth of drug type use, and was unique in terms of its high probability of meth/amphetamine use. Class 5 was unique in that the breadth of drug types was narrow, and this was the only class to have a high probability of sedative use.

In some earlier research (based on different age groups and from other countries), cluster analytic approaches have tended to identify clusters that primarily vary in the breadth of substance uses (i.e., clusters frequently characterized by minimal, limited, or extended use of different substances). The present findings suggest that, at least in the present sample, variability in high probability substances is limited (between two and five substances, including alcohol), and that several classes are characterized by “spikes” (high and unique probabilities for particular drugs) (see **Figure 1**). These two observations suggest that polydrug use clusters are not simply groupings based on increasing “deviance” in relation to drug use (i.e., subgroups representing those who have progressively expanded their field of preferred substances). Though beyond the scope of this research, the findings point to the possibility that certain subgroups prefer certain combinations of drugs or that there

are restraints or sanctions on the use of particular drugs outside the clusters established in this study. For example, subgroups may have positive and negative attitudes about certain drugs based on their legal status (accounting for the split of alcohol/alcohol and tobacco from other classes), the acceptability of common routes of administration [inhaling versus oral ingestion; (47)], or the perceived instrumental properties of particular drug types (e.g., to increase sociability, reduce anxiety). Further research is needed on mechanisms that may account for clustered use of particular drugs in nationally representative samples. The reasons why particular drugs cluster together may vary across subgroups or within subgroups.

Predictors of class membership (relative to the *alcohol* cluster) indicated consistent and relatively strong odd ratios for sex (being female less likely to be associated with polydrug use and this was significant for all clusters), non-completion of high school (for all except the *Sedative* and *alcohol* class), low income (associated with lower likelihood of polydrug use), and peer use of alcohol/tobacco/cannabis (consistently predicted membership of all clusters). There were weak or inconsistent effects for age, remoteness (though outer regional participants were less likely to be in clusters 3 and 4), marital status, and depressed mood. The odds of cluster membership relative to the alcohol group were generally non-significant for employment status and socioeconomic disadvantage (measured using a community-level government index). The effects for general health were in the expected directions (clusters 2–5 reporting lower health), although poor health ratings mostly did not reach statistical significance. In sum, polydrug use was generally associated with being male, having a very low education (high school not completed), having a high income relative to those on a low income, and having a peer group where the majority use alcohol, tobacco, or marijuana.

The secondary analyses exploring simultaneous use of drugs increased our confidence in the polydrug classes identified in the



LCA. The percentage of concurrent users who also use simultaneously was high, depending on drug pair endorsements. In line with previous studies [e.g., Ref. (26, 33, 34)], alcohol was simultaneously used with a wide variety of other drugs (notably cannabis, ecstasy, meth/amphetamine, cocaine; all  $\geq 89\%$ ). Less commonly used together were cannabis and cocaine, cocaine and ecstasy, and alcohol and painkillers, though prevalence within concurrent users was still high (47–55%). The high rates of simultaneous use of particular drug pairs points to the possibility that certain drugs are commonly used for their synergistic effects [e.g., stimulants used to reduce the depressant effects of alcohol; (23, 24, 48, 49)], or that craving for one drug type may increase cravings for another (50), or that supply chains increase the probability of expansion into other drug types.

The findings have important implications for prevention and early intervention programs and policies relating to young adult drug use. First, the clustering of alcohol and tobacco for a large minority of the sample points to the possibility that alcohol use is a high risk context for tobacco use (51). Though beyond the scope of this study, there is good evidence that alcohol consumption may erode smoking cessation attempts in young people (52), so smoking prevention programs may benefit from an increased focus on the risks of alcohol use. Second, the findings assist in the relative weighting of prevention strategies based on the prevalence of polydrug classes. It is clear that a substantial minority (approximately 12%) of young adults are concurrent users of several drug types, including cannabis, ecstasy, alcohol, and to a smaller extent meth/amphetamines. At least in Australia, the use of amphetamine derivatives (including ecstasy) is comparatively rare in the teenage years, so a focus on the risks associated with amphetamine derivatives (and simultaneous use of other drugs) upon reaching young adult may be useful in stemming the apparent expansion in the use of these drug types. Third, the proportion of young adults who use sedatives and alcohol is small but meaningful, and reinforces the importance of screening and surveillance strategies to detect vulnerable young adults, and strategies to reduce the inherent risks in combining sedatives and alcohol (Table 3). Finally, young adult males with low education were at elevated risk of poly drug use, pointing to the value of prevention programs instituted prior to the upper levels of high school and programs for young males made available in higher paid working environments.

The use of national surveys such as the NDSHS provides a large sample of cross-sectional overview of drug use in the general population. However, the sampling of individuals with a fixed home address or landline can fail to capture high risk young people, such as those in transient accommodation. This may lead to an underestimation of the actual rates of polydrug use in this group (53, 54). Because this was a cross-sectional survey, no conclusions can be made about mechanisms linking the use of different drugs, or the factors that may underlie cluster membership. It remains unclear whether people in particular polydrug classes have communalities in the development of their drug use profiles, or whether some classes merely represent a continuum of severity. This study did not involve measures of quantity or harm, and it is likely that only a very small proportion of this sample had any

form of drug dependence. Therefore, results may not generalize to groups with more significant drug problems. The study relies on self-report, although the confidential and anonymous nature of participation is likely to have reduced response bias. Although the overall sample size of this study was large, the sample sizes in the class “Cannabis, amphetamine, derivatives and illicit drugs” and “sedatives and alcohol” were relatively small and this limited the power to identify covariates that were associated with these two classes. Given the large number of possible permutation of simultaneous drug use, we only examined the prevalence of specific drug pairs. Lastly, due to the limitations of the survey design, we were not able to examine simultaneous use tobacco with other drugs.

## CONCLUSION

While the sole use of alcohol characterized most people in the sample, a substantial minority of young adults were concurrent users of several drug types. Young adults clustered on unique combinations of alcohol, tobacco, cannabis, ecstasy, meth/amphetamine, and sedatives. Simultaneous use of two substances was characteristic of concurrent users. Peer use of alcohol, tobacco, and cannabis were significant correlates of young adult polydrug use. Young males with low education and/or higher income are a high risk group that warrant a prevention and early intervention focus.

## AUTHOR CONTRIBUTIONS

Lake-Hui Quek drafted this manuscript under the supervision of Adrian B. Kelly and Angela White. Gary C. K. Chan conducted statistical analyses. Angela White participated in the design of the study and provided feedback at all stages. Jason P. Connor provided conceptual direction and feedback on manuscript drafts. Peter J. Baker supervised the statistical analyses. John B. Saunders participated in the design of the project and provided feedback on manuscript drafts. Adrian B. Kelly supervised the overall project, mentored Lake-Hui Quek in manuscript preparation, and assisted with writing the manuscript.

## ACKNOWLEDGMENTS

This study was supported through internal funds and via Australian Research Council Project DP130102015 to Adrian B. Kelly (first chief investigator). Jason P. Connor is supported by a National Health and Medical Research Council of Australia Career Development Fellowship (APP1031909). These funding bodies had no role in the design, analysis, or interpretation of the data. The data on which this study is based were collected as part of the National Drug Strategy Household Survey by the Australian Institute of Health and Welfare. The authors also wish to acknowledge the contribution of Dayna Smith for editing the manuscript and Charlotte Brackenridge who assisted with the literature search.

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

*Received: 19 August 2013; paper pending published: 24 September 2013; accepted: 14 November 2013; published online: 28 November 2013.*

*Citation: Quek L-H, Chan GCK, White A, Connor JP, Baker PJ, Saunders JB and Kelly AB (2013) Concurrent and simultaneous polydrug use: latent class analysis of an Australian nationally representative sample of young adults. Front. Public Health 1:61. doi: 10.3389/fpubh.2013.00061*

*This article was submitted to Epidemiology, a section of the journal Frontiers in Public Health.*

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