Prognosis of adolescent partial syndromes of eating disorder

George C. Patton, Carolyn Coffey, John B. Carlin, Lena Sanci and Susan Sawyer

Background
Partial syndromes of eating disorder are common in adolescents but the health significance of these syndromes remains uncertain.

Aims
To document the health and social adjustment in young adulthood of females assessed as having a partial syndrome of eating disorder in adolescence.

Method
A community sample of 1943 participants was tracked over 10 years in an eight-wave cohort study. A partial syndrome was defined as the fulfilment of at least two DSM–IV criteria for either anorexia or bulimia nervosa at one assessment or more between the ages of 15 years and 17 years.

Results
Partial syndromes were found in 9.4% of 15- to 17-year-old female participants and 1.4% of males. There were few instances of progression of partial syndromes to fully fledged anorexia and bulimia nervosa. However, among those with partial syndromes depressive and anxiety symptoms were two to three times higher in young adulthood, substance misuse was common, and a majority of those with a partial syndrome of anorexia nervosa were still underweight in their mid-20s.

Conclusions
Given the level of subsequent psychopathology and social role impairment, there may be justification for initiating trials of preventive and early clinical intervention strategies for adolescent partial syndromes.

Declaration of interest
None.

Method
Between August 1992 and March 2003 we conducted an eight-wave cohort study of adolescent and young-adult health in the state of Victoria, Australia. Data collection protocols were approved by The Royal Children's Hospital's ethics in human research committee. The cohort was initially defined with a two-stage cluster sample in which we selected two classes at random from each of 44 schools drawn from a stratified frame of all schools (government, Catholic and independent) in the state (total number of students 60905). School retention rates to year
9 in the year of sampling were 98%. One class from each school entered the cohort in the latter part of the 9th school year (wave 1), with a second class entering 6 months later, early in the 10th school year (wave 2). Participants were subsequently reviewed at a further four 6-month intervals during their teenage years (waves 3 to 6) with two follow-up waves in young adulthood, at age 20–21 years (wave 7) and 24–25 years (wave 8). In waves 1 to 6, participants self-administered the questionnaire on laptop computers,17 with telephone follow-up of those absent from school. The 7th and 8th waves of data collection were undertaken with computer-assisted telephone interviews.

From a total sample of 2032 students, 1943 (96%) participated at least once during the first six (adolescent) waves. In the 7th wave, 1601 young adults (79% of the initial sample or 82% of teenage participants) were interviewed between April and December 1998. In the 8th wave, 1520 (75% of the initial sample, 78% of teenage participants) were interviewed between April 2002 and June 2005. Response rates are shown in Fig. 1. Reasons for non-completion at follow-up were refusal (wave 7, n=152; wave 8, n=269), loss of contact (wave 7, n=192; wave 8, n=152) and death (wave 7, n=2; wave 8, n=7). The mean ages at waves 1, 7 and 8 were 14.5 years (s.d.=0.5), 20.7 years (s.d.=0.5) and 24.1 years (s.d.=0.6) respectively.

Measures

Weight and height

Weight was measured to the nearest 0.1 kg, with participants wearing minimal school uniform and a further 1 kg deducted to account for this clothing. Height was taken with shoes removed using a stadiometer and measured to the nearest centimetre. Self-reported weights were used for those who had left school. At waves 7 and 8, weight and height were assessed using self-report, after prior notification that these two questions would be asked at interview.

Eating disorder

The Branched Eating Disorders Test (BET) was used to assess DSM–IV criteria for eating disorder.18 This test was designed for use in teenaged community samples and covers symptoms of eating disorder over the previous 4 weeks. Partial syndromes of eating disorder were based on the DSM–IV criteria for bulimia nervosa and anorexia nervosa, this being the approach most commonly adopted in the definition of these syndromes.19 A partial syndrome of anorexia nervosa was defined as meeting two of the following four criteria:

(a) low body weight, defined on the basis of a z score for body mass index (BMI) below the 5th percentile for age and gender, using the reference data from the international task-force for the standardisation of overweight and obesity in children and adolescents;20

(b) intense fear of gaining weight or becoming fat when under the 25th percentile z score for BMI for age and gender;

(c) disturbance in the experience of body weight, size and shape when under the 25th percentile z score for BMI for age and gender;

(d) amenorrhoea, defined as missing three consecutive menstrual periods.

A partial syndrome of bulimia nervosa was defined as meeting at least two of the following criteria:

(a) objective bingeing at least weekly for at least 4 weeks;

(b) use of any of the following for at least 4 weeks: self-induced vomiting at least twice weekly, laxatives at least twice weekly, diuretics at least twice weekly, daily fasting (12 h or longer) for at least 4 weeks or daily vigorous exercise to control weight;

(c) report of body weight and shape as extremely important to the participant’s sense of self.

Definitions of partial syndrome of bulimia nervosa using the BET have previously been shown high agreement with the Eating Disorders Examination in community sample of schoolgirls in Australia (sensitivity 1.0, specificity 0.99, positive predictive value 0.7).21 The BET was administered from wave 2 through to wave 6. At follow-up in wave 7 the BET was administered to female participants only. Therefore this report examines continuity of eating disorder in young females only.

Depression and anxiety

Symptoms of depression and anxiety were assessed from waves 1 to 7 using the Revised Clinical Interview Schedule (CIS–R).22 The total scores on this measure were dichotomised at a cut-off point of 11/12 to delineate a mixed depression–anxiety state at a lower threshold than syndromes of major depression and anxiety disorder but where clinical intervention would still be appropriate.23,24 Specific anxiety and depressive syndromes were also defined using the CIS–R.22 The 12-item General Health Questionnaire (GHQ)25 was used to assess these symptoms at wave 8: a cut-off score of 2/3 was used to define a group with high psychiatric morbidity.

Substance use (wave 8)

Nicotine dependence was assessed with the Fagerstrom Test for Nicotine Dependence and was defined as a score of 4 or more.26

Fig. 1 Sampling and ascertainment in the Victorian Adolescent Health Cohort, 1992–2003.
Alcohol misuse and dependence was assessed using the Composite International Diagnostic Interview 2.1, 12-month version (CIDI) to define DSM–IV categories.27 Cannabis use was assessed on the basis of self-reported frequency of use over the past year, with high use defined as reported use at least weekly. Amphetamine use was defined as self-reported use in the previous 12 months.

Analysis
Data collection was undertaken at a developmental point when young people are difficult to trace because of high mobility. Although the initial response was high and attrition low, 36% of respondents missed at least one wave of data collection in the adolescent phase (waves 1 to 6), leading to a potential bias in summary measures of exposure to cannabis and mental health problems calculated from these data. For example, when we examined characteristics of wave 7 non-completers using logistic regression models, male participants were overrepresented (OR=1.9, 95% CI 1.5–2.4), as were those with a background of parental divorce or separation (OR=1.8, 95% CI 1.4–2.5) and those who were daily smokers at study inception (OR=2.1, 95% CI 1.5–2.9). To address this selection bias, we used the method of multiple imputation,28 with five complete data-sets created by imputation under a multivariate normal model that incorporated all the outcome variables of interest measured at all waves of data collection, along with the fixed covariates of gender, age, rural/urban residence and parental education (available for all participants).29 Participants were classified according to whether they fell into categories of interest at least once during waves 1 to 6 (adolescence) and, separately, in waves 7 and 8 (young adulthood). Data analysis was undertaken using Stata version 8 for Windows. Univariate logistic regression analyses were used

### Table 1: Cumulative prevalence of partial-syndromal eating disorders in 1943 participants in the Victorian Adolescent Health Cohort Study for waves 3 to 6

<table>
<thead>
<tr>
<th>Eating disorder</th>
<th>Males (n=943)</th>
<th>Females (n=1000)</th>
<th>Total (n=1943)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>% (95% CI)</td>
<td>% (95% CI)</td>
<td>% (95% CI)</td>
</tr>
<tr>
<td>Bulimic disorder</td>
<td>72 (1.0–1.6)</td>
<td>6.4 (4.8–8.0)</td>
<td>3.7 (2.8–4.6)</td>
</tr>
<tr>
<td>Anorexic disorder</td>
<td>42 (0.0–0.8)</td>
<td>3.8 (2.5–5.1)</td>
<td>2.2 (1.4–3.0)</td>
</tr>
<tr>
<td>Any eating disorder</td>
<td>107 (1.4–2.1)</td>
<td>9.4 (7.3–11.4)</td>
<td>5.5 (4.3–6.7)</td>
</tr>
</tbody>
</table>

a. Frequencies obtained by averaging across five imputed data-sets.
b. Eight participants fulfilled criteria for both anorexia and bulimia partial syndromes at different waves.

### Table 2: Prevalence of health outcomes in young adulthood of adolescent eating disorders among 999 living female cohort participants according to adolescent eating disorder status

<table>
<thead>
<tr>
<th>Outcome measured in young adult waves (waves 7 and 8)</th>
<th>No eating disorder (n=905)a</th>
<th>Partial syndromal eating disorder diagnosis across waves 3 to 6</th>
<th>Any eating disorder (n=94)b</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Prevalence</td>
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</tr>
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<td></td>
<td>% (95% CI)</td>
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</tr>
<tr>
<td>Partial syndromal eating disorders</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Anorexic</td>
<td>17 (0.4–2)</td>
<td>7 (0.0–1.3)</td>
<td>10 (0.0–2.2)</td>
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<tr>
<td>Bulimic</td>
<td>22 (0.2–3)</td>
<td>11 (2–19)</td>
<td>14 (0.8–26)</td>
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<tr>
<td>Any eating disorder</td>
<td>37 (1–4)</td>
<td>14 (4–25)</td>
<td>21 (2–40)</td>
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<tr>
<td>Psychiatric morbidity</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Depressive syndrome</td>
<td>129 (9–14)</td>
<td>29 (17–41)</td>
<td>32 (12–51)</td>
</tr>
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<td>219 (17–22)</td>
<td>47 (33–61)</td>
<td>47 (29–65)</td>
</tr>
<tr>
<td>Wave 7 (average age 20 years)</td>
<td></td>
<td></td>
<td></td>
</tr>
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to model associations, and Wald tests and related confidence intervals were used to assess statistical significance and precision.

### Results

Table 1 shows the estimated prevalence for partial and full syndromes of eating disorder between the ages of 15 years and 17 years (waves 3 to 6). Just under one in ten of the female participants fulfilled criteria for an eating disorder over this time. One participant fulfilled all criteria for anorexia nervosa during this time, and four fulfilled all criteria for bulimia nervosa. Eight participants fulfilled criteria for partial syndromes of anorexia and bulimia at different waves. The prevalence rate for partial syndromes in male participants was low, at just over 1%. The OR of an eating disorder was 7.9 (95% CI 4.1–15) times greater for female participants than for males.

Health outcomes for partial syndromes in young adulthood are shown in Table 2. Results are presented for outcomes at wave 7 (average age 20 years) and wave 8 (average age 24 years). Eating disorders at a partial syndrome level were found in 14% of those who had an adolescent bulimic partial syndrome and 21% of those who had an anorexic partial syndrome. The majority of young-adult cases from the adolescent bulimic group were also female participants according to adolescent eating disorder.

### Discussion

This study confirms that partial syndromes are common during adolescence. Around one in ten young females fulfilled criteria for these conditions at some time between the ages of 15 and 17 years. Around one in six of those with an adolescent eating disorder had a persisting partial syndrome as a young adult, consistent with an earlier report from the USA that a majority of such disorders do not persist into young adulthood. There was little to suggest that partial syndromes progress to clinical eating disorders in the medium term. Nevertheless, female participants with
A partial syndrome had a broad range of ongoing health and social problems as young adults. Depressive and anxiety syndromes were two to three times more common in this group. Weight disturbance persisted in those with adolescent anorexia partial syndromes, with over two-thirds remaining underweight in their mid-20s. Substance misuse was generally higher, with over twofold higher rates of nicotine dependence, alcohol use disorders and amphetamine use. The psychosexual profile of the partial syndrome group also differed, with high rates of early initiation of sexual intercourse and a tendency towards more early pregnancies. Those with adolescent eating disorder were less likely to have successfully completed their education and to have made a successful transition into employment or vocational training by their mid-20s.

Early studies of the outcomes of partial syndromes focused on clinical samples from which generalisation may be difficult, had limited follow-up or used uncertain measures of the symptoms of eating disorders.15,32 The term ‘partial syndrome’ defines syndromes that are likely to be similar to the DSM–IV category of ‘eating disorder not otherwise specified’. A close-to-representative community sample, high participation rates and frequent measures of symptoms during the teenage years are strengths of this study. The use of multiple imputation should have minimised possible biases arising from missing data. It nevertheless remains possible that patterns of association among those who were non-responders might not be fully captured by the imputation modelling, leading to some residual bias in the final estimates. One further limitation is the availability of only one point of assessment for eating disorders in young adulthood. Given that eating disorders may continue to fluctuate in intensity, it is possible that we have underestimated the degree of continuity from adolescence to young adulthood. The ongoing low weight of those who had had an anorexia syndrome is noteworthy in this respect. It would be consistent with continuing body image sensitivity and efforts to control weight even if the individual does not fulfil formal criteria for an eating disorder.

A majority of later health and adjustment problems in young females with an adolescent partial syndrome were not specifically related to ongoing eating disorder. Some of these problems might arise because of the presence of other psychiatric syndromes that commonly co-occur with eating disorders.52,53 Other problems such as educational failure might be the result of disruption to school attendance during the teenage years, a process to which the symptoms of eating disorder may contribute. It is also possible that adolescent partial syndromes are best viewed as a marker for an ongoing susceptibility to a range of psychiatric and behavioural disorders, rather than being linked specifically to anorexia and bulimia nervosa.

Later substance misuse was prominent in the partial syndrome group, suggesting that risk of substance misuse is not limited to those with fully fledged syndromes.53 Earlier reports have noted that a range of substances are commonly used by adolescents and young adults with weight concerns and dieting.52,53 In our study an association with tobacco and amphetamine use, but not with cannabis use, was prominent. This perhaps reflects an ongoing preoccupation with weight control, with tobacco and amphetamines being recruited for their effects on appetite and weight control. In contrast, cannabis intoxication often brings an increase in appetite that may be a disincentive for use.

A range of difficulties in social adjustment in young adulthood found in fully fledged eating disorders were also evident in those with partial syndromes.15,32 The pattern of psychosexual development differed somewhat from that found in fully fledged eating disorders, in which sexual milestones are typically delayed.38 It is, however, consistent with the finding of larger population-based surveys, in which bulimic symptoms have been linked to early menarche and in turn to earlier sexual activity.39 It is also possible that the impulsivity associated with bulimic syndromes has a role in early sexual activity and the trend towards higher rates of pregnancy. However, these factors cannot account for the apparently high rate of spontaneous first-trimester abortion. Earlier reports have found that active bulimia nervosa during pregnancy is associated with higher rates of miscarriage and that these differences were not accounted for by intercurrent substance misuse.40 Our findings raise the possibility that lower levels of eating disorder symptoms might also predict poor pregnancy outcomes.

Few partial syndromes appear to progress to clinical eating disorders in community settings. As a means of preventing full clinical syndromes of anorexia and bulimia nervosa, there appears little justification for early screening and intervention. However, partial syndromes are common and are linked to a range of later health and social problems. From this alternative perspective there is justification for giving clinical attention to this group. Yet clinical response is not likely to be easy. Although screening tools exist for use in community settings, those with probable eating disorder will often not accept referral for treatment.41 Eating disorders are not often routinely identified by general practitioners, and given their high prevalence these disorders present a major challenge for existing services.42 Treatment may well require the kind of multifaceted psychological approach advocated for the full syndromes of anorexia and bulimia nervosa.53 This may be one reason why those presenting for specific treatment of eating disorders in primary care are unlikely to persist with treatment if the clinical intervention is not sufficiently intense.44 However, several strategies of response seem worth considering. For particular subgroups such as young females who are pregnant or those with diabetes mellitus, screening and clinical interventions are more likely to meet with success.45,46 In primary care a greater attention to general psychosocial risk screening may be more effective in engaging this group than specific screening for eating disorders.47 For those identified as having a partial syndrome, there may be greater value in attending to symptoms of depression and anxiety, substance misuse and sexual health problems, as well as the social and educational difficulties so commonly associated with having a partial syndrome. Self-help interventions in non-clinical settings, tied to the promotion of health literacy around eating disorders, might also reduce the substantial disease burden associated with the most common eating disorders of adolescence.48,49 A recent finding that internet delivery of cognitive–behavioural therapy brought substantial reductions in symptoms and adverse consequences of common eating disorders for up to 2 years illustrates the potential of this approach.50

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