BMC Psychiatry



Research article Open Access

Weight management in a cohort of Irish inpatients with serious mental illness (SMI) using a modular behavioural programme. A preliminary service evaluation

Chris J Bushe*¹, Dermot McNamara¹, Cliff Haley², Mary Fleming McCrossan³ and Pat Devitt⁴

Address: ¹Eli Lilly and Company Ltd, Adelaide Road, Dublin, Ireland, ²Psychiatry Dept, Letterkenny General Hospital, Donegal, Ireland, ³Nursing Dept, Cleary House, Letterkenny, Donegal, Ireland and ⁴Psychiatry Dept, Adelaide and Meath Hospital, Tallaght, Dublin 24, Ireland

Email: Chris J Bushe* - bushe_chris@lilly.com; Dermot McNamara - mcnamara_dermot@lilly.com; Cliff Haley - Clifford.Haley@mailb.hse.ie; Mary Fleming McCrossan - Clifford.Haley@mailb.hse.ie; Pat Devitt - patrickdevitt@hotmail.com

Received: 29 February 2008 Accepted: 15 September 2008

* Corresponding author

Published: 15 September 2008

BMC Psychiatry 2008, 8:76 doi:10.1186/1471-244X-8-76

bine rsychiatry 2006, **6**.76 doi:10.1166/1471-244A-0-76

This article is available from: http://www.biomedcentral.com/1471-244X/8/76

© 2008 Bushe et al; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Background: Weight gain is commonly observed during psychotropic treatments for chronic forms of severe mental illness and is most rapid during the early treatment phases. All formats of behavioural weight intervention programmes have suggested that weight gain can be prevented or reversed in some patients. There is no data on these programmes in acutely unwell inpatients whom may be the major beneficiaries.

Methods: A modular behavioural intervention programme (Solutions for Wellness) used in SMI outpatients since 2002 in Ireland has been adapted for inpatient use. Preliminary data is reported from 5 centres in Ireland.

Results: In 47 inpatients the mean weight change was +0.26 kg (SD 2.02) with a median change of 0 kg. Mean follow-up was 23.7 (SD 21.6) days, and median 14 days (range 6–98 days). There was no difference in mean weight change in those patients involved for > 35 days compared with < 35 days (+0.26 kg; 0.25 kg; p = 0.5). Weight loss or maintenance was seen in 70% of patients.

Conclusion: These preliminary data are supportive of the concept that acutely unwell inpatients with SMI may engage with a behavioural weight programme. Weight change observed contrasts with the significant weight gain often seen in most subjects. Further clinical trials are warranted.

Background

There has been increasing recognition that cardiovascular mortality is significantly increased in patients with severe mental illness (SMI) [1-3]. There are complex reasons for this that include genetic predisposition, smoking, poor dietary choices, decreased physical activity and weight gain observed with most psychotropic treatments [4]. The

consensus view is that cardiovascular risk factors in SMI patients need to be reduced [2,5]. Weight gain prevention is one of the more easily defined and measured problem areas.

Patients with schizophrenia are generally of normal BMI at illness onset and males may even be underweight [6].

There is clear recognition that patients treated in their early phase of psychosis gain significant weight during the first year of treatment with the recent CAFÉ study reporting 80% olanzapine subjects, 50% quetiapine and 58% risperidone subjects gaining > 7% of their body weight in the first 12 months [7] in the absence of any preventative intervention. In addition although weight gain is maximal during the early months of initiating antipsychotic treatment, in a large 3 year naturalistic study an increasing number of subjects on all medications continued to gain weight during the 3 year study [8,9], and [10].

These efficacy based clinical trials both randomised and observational have to date not been able to encompass any preventative measures to reduce weight gain. Despite this a variety of weight and lifestyle management programmes have been developed worldwide in outpatients and broadly have produced good results [11-16]. The forms of intervention range from weekly group sessions using behavioural models through to one to one nursing sessions and telephone/internet based models [17].

In Ireland a lifestyle management programme Solutions for Wellness has been operating in outpatients with forms of severe mental illness (SMI) since 2002 and recently reported preliminary data in the outpatient subject cohort [15]. The programme has been slightly adapted to be used in an inpatient setting with the rationale of earlier intervention. We describe preliminary results from the initial cohort of patients evaluated. The premise being evaluated is that early weight gain may be prevented or reduced in some patients.

Methods

A group programme (Acute Solutions for Wellness) designed to address weight and other cardiovascular risk factors commenced in 2005 in a number of geographically diverse centres in Ireland in inpatients with acute forms of severe mental illness (SMI). Five centres of those taking part in the programme agreed to provide their data for a preliminary service evaluation. Data was not routinely recorded in other centres. There was no restriction placed on the precise diagnoses to which the intervention would be offered to but it was expected that most patients would have schizophrenia or severe affective disorders and was offered to suitable inpatients referred by the ward nursing staff. There was neither restriction on drug and non-drug treatments nor age restrictions. As this was not a clinical study total discretion for entry to the programme remained with the ward nursing staff. They chose subjects whom they considered were potentially suitable for the programme based on mental state. The groups commenced with a cohorts of patients at each centre whom progressed through the programme sequentially as a group.

The programme consisted of 8 modules to complete over 4 weeks, each module of 30 minutes covering dietary choices, appetite, physical activity and nutrition. The materials were adapted to be simplified from the outpatient version of the same service and in addition modules were designed to be 30 instead of 60 minutes each. The programme otherwise was unchanged. Groups were led by trained healthcare professionals usually nurse. The nurses received full training in the operation of the modular programme and worked from identical source material workbooks consisting of 20 page instructor guides describing each module individually. Patients also received written materials in the form of a 22 page workbook similarly describing in appropriate terminology each module separately. The 8 modules covered the topics:

- 1. Healthy living
- 2. Physical activity
- 3. The food pyramid
- 4. Recommended food servings
- 5. Fat and salt in your diet
- 6. Healthy and unhealthy eating habits
- 7. High fibre diet
- 8. Controlling your hunger

The patients received either twice weekly modules each 30 minutes or a single hourly session weekly combining two modules. Subject diagnoses were recorded from the notes.

Ethics

Ethical approval was not sought as this service provision of SFW was deemed to be part of good clinical practice and service evaluations may be considered exempt from ethical review. The only difference being that the service was provided in secondary care rather than in primary care. Informed consent was obtained from the patients to use their data in an anonymised manner for the purpose of a service evaluation.

Statistical Analysis

Our analysis has been predominantly descriptive, aiming to present the raw data as a series of weight or BMI results over a period of time. Our descriptive statistics are supported with standard deviations, median and significance is reported where appropriate.

Results

Since 2005 data is available from 47 patients (26 females, 21 males) from 5 centres. Diagnoses were Schizophrenia in 17/47, affective disorders 25/47, non-psychotic diagnoses (personality disorder, anxiety, alcohol misuse) 5/47. Baseline mean weight 78.4 kg (SD 15.2 kg) and median 77 kg (range 47–135 kg).

The mean weight change was +0.26 kg (SD 2.02) with a median change 0 kg. Paired Student's t-Test t = -0.624, df = 46, p = 0.950. Mean follow-up 23.7 (SD 21.6) days, median 14 days (range 6-98 days) with 16/47 engaging with the programme >=21 days (figure 1). The categorical weight changes at discontinuation with programme were weight gain in 14/47 (30%), weight maintenance in 21/47 (45%) and weight loss in 12/47 (25%). The categorical changes are shown in figure 2. The mean weight change in the 10/47 patients whom remained in the programme >=35 days was +0.25 kg, (SD 3.55) and in the 37/47 whom remained in the programme <=35 days +0.262 kg, (SD 1.43) (p = 0.50)

Discussion

In this preliminary report an adapted lifestyle intervention programme has found promising results. At discontinuation of engagement with the programme only 14/47 (30%) patients had gained weight during a mean follow up of 24 days (median 14 days) and the remainder either maintained their weight or lost weight. The median weight change was 0 kg. Furthermore there were no differences between those patients who remained in the programme less than 35 days compared to the group whom engaged for greater than 35 days (p = 0.5). These data may hold relevance in that a major determinant of length of engagement with SFW was the time to discharge of the

Categorical Weight Change

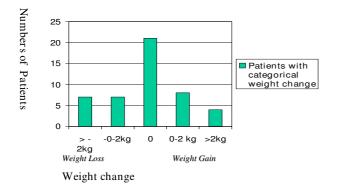
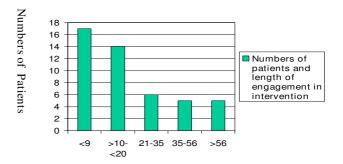


Figure I
Length of engagement with programme by patients.

SFW – Length of Involvement



Number of Days Engaged with Acute SFW

Figure 2
Categorical amounts of weight change in patients.

patients. At discharge there would be no certainty that continual engagement with any weight management of lifestyle intervention programme would be either offered or accepted.

There are few data which can be compared with our data as to our knowledge this is one of the first data sets that has examined a lifestyle intervention in an inpatient SMI cohort acutely unwell.

Some short term interventions have been successful in outpatients. In a 14-week open label study of patients switched to risperidone from olanzapine and randomised to receive a behavioural intervention, 27% of the active cohort lost > 5% body weight over 14 weeks [13] and in a 12-week intervention using a similar programme to Acute SFW in outpatients there was a mean weight loss of 2.7 kg [18]. The few studies that have examined interventions in early phases of illness are worth comparison. A study of 61 first-episode patients aged 16-45 years randomised to behavioural intervention or "usual care" over 12 weeks [19]. Only 40.7% of the intervention group gained > 7% of weight compared with 77% of the control group. Similar data was reported from a cohort of 50 patients randomised to intervention or usual care within 30 days of commencement on an antipsychotic and followed for 16 weeks [20]. 63% of the intervention cohort experienced no weight gain compared with 22% of the control patients (p = 0.009). In this study 70% of our cohort experienced no weight gain but only over mean 24 days follow up. A further encouraging finding was that the patients engaging with the programme for greater than 35 days had no greater weight gain than those engaging for less time. In a recent naturalistic study of an admixture of inpatients and outpatients (mean weight loss 2.64 +- 2.75 kg) BMI reduction over 12 weeks was greater in the outpatient cohort making these preliminary data even more encouraging and worthy of further research [21].

The literature on weight management programmes consists primarily of uncontrolled interventions, whose data seems promising and shorter term controlled trials but not conclusive of benefit. With a few reservations most data show encouraging results. A number of systematic reviews and reviews have consistently suggested some benefit from these programmes [14,22]. Recent controlled trials have also suggested more tangible gains from lifestyle programmes. Mauri et al, report a reduction in fasting insulin levels, insulin resistance (HOMA index) and improvement in insulin sensitivity but unchanged FPG in an olanzapine cohort randomised to a psycho educational programme for weight loss [23] for either 12 or 24 weeks. Menza found a reduction in glycosylated haemoglobin over 1 year using a multi-modal weight control programme (n = 31) from 5.35% to 5.11% (p = 0.001) [24].

The data from the UK programme Wellbeing support programme report 80% retention in the programme over 2 years and in addition to reduction of various cardiovascular risk factors; weight gain to any extent was measured in only 30.8 % [25]. Longer term compliance with these programmes may be crucial with recent data suggesting compliance is a significant factor [21,11], and [12]. An additional salient consideration is that Werneke et al in 2002 [26] and Faulkner in 2007 [27] concluded that there was not sufficient evidence to use any pharmacological agents for weight reduction or prevention.

In a Cochrane review in 2007 Faulkner found that only 5 trials out of 23 randomised interventions were behavioural in type and modest weight loss and prevention were reported [27]. The relevant caveats around these findings included small studies over short term periods with varying interventions, and longer term larger studies need to be undertaken before making definitive statements of benefit.

Limitations

There are however significant limitations in our data that will need to be resolved in future controlled clinical studies and in further service evaluations of this programme. Primarily this relates to the nature of this being a service evaluation and not a form of designed controlled trial or audit. These data may provide hypotheses for future controlled evaluations of such programmes in acutely unwell SMI patients. There are no data on the number of patients who were available for the intervention but were either not approached or declined. Thus it cannot be certain that our results can be extrapolated to other inpatient populations. There is also no data on the severity of the patient's

illness and there is no control group. All data obtained from this service evaluation has been included, however data on many demographics including pharmacological treatments, previous weight changes and BMI were not available for 38% of the patients and have thus not been reported. A final limitation is the short follow up period that fails to assess whether any benefits may persist.

Conclusion

Despite these limitations there is an increasing body of evidence that weight management interventions may potentially provide benefit [12,21] although caveats remain that most studies are small, use varying interventions and are mostly short term. Further clinical studies are required in acutely psychotic patients to determine if early weight gain can be prevented or ameliorated and benefits maintained longer term. Soon after diagnosis of psychosis and soon after drug initiation in more chronic subjects, may represent crucial times to intervene with regard to prevention of weight gain.

Competing interests

CB and DM are employees of Eli-Lilly UK.

Full details of the programme and its application can be obtained from via Email: - bushe_chris@lilly.com

This paper has been written by the named authors without editorial assistance.

Authors' contributions

CB, DM, PD, MF and CH conceived the analysis. CB analysed the data. PD, MF and CH have expertise in the SFW for outpatients. CB drafted the manuscript and all authors had input and approved the final version.

Acknowledgements and funding

Eli Lilly provided the funding for the SFW programme. CB and DM are employees of Eli Lilly.

References

- Osborn DP, Levy G, Nazareth I, Petersen I, Islam A, King MB: Relative risk of cardiovascular and cancer mortality in people with severe mental illness from the United Kingdom's General Practice Research Database. Arch Gen Psychiatry 2007, 64(2):242-9. Erratum in: Arch Gen Psychiatry. 2007 Jun; 64(6):736.
- Hennekens CH, Hennekens AR, Hollar D, Casey DÉ: Schizophrenia and increased risks of cardiovascular disease. Am Heart J 2005, 150(6):1115-21.
- Lawrence DM, Holman CD, Jablensky AV, Hobbs MS: Death rate from ischaemic heart disease in Western Australian psychiatric patients 1980–1998. Br / Psychiatry 2003, 182:31-6.
- Thakore JH, Mann JN, Vlahos I, Martin A, Reznek R: Increased visceral fat distribution in drug-naive and drug-free patients with schizophrenia. Int J Obes Relat Metab Disord 2002:137-41.
- McEvoy J, Meyer M, Goff D, Nasrallah H, Davis S, Sullivan L, et al.: Prevalence of the metabolic syndrome in patients with schizophrenia: baseline results from the Clinical Antipsychotic Trials of Intervention Effectiveness (CATIE) schizophrenia trial and comparison with national estimates from NHANES III. Schizophr Res 80(1):19-32. 2005 Dec 1;

- Sorensen H, Mortensen EL, Reinisch JM, Mednick JA: Height, weight and body mass index in early adulthood and risk of schizophrenia. Acta Psychiatr Scand 2006, 114:49-54.
- McEvoy JP, Lieberman JA, Perkins DO, Hamer RM, Gu H, Lazarus A, Sweitzer D, Olexy C, Weiden P, Strakowski SD: Efficacy and tolerability of olanzapine, quetiapine, and risperidone in the treatment of early psychosis: a randomized, double-blind 52week comparison. Am J Psychiatry 2007, 164(7):1050-60.
- Lambert M, Haro JM, Novick D, Edgell ET, Kennedy L, Ratcliffe M, Naber D: Olanzapine vs. other antipsychotics in actual outpatient settings: six months tolerability results from the European Schizophrenia Out-patient Health Outcomes study. Acta Psychiatr Scand 2005, 111:232-43.
- Haddad P: Weight change with atypical antipsychotics in the treatment of schizophrenia. Psychopharmacol 2005, 19(6 Suppl):16-27.
- Haro JM, Suarez D, Novick D, Brown J, Usall J, Naber D, SOHO Study Group: Three-year antipsychotic effectiveness in the outpatient care of schizophrenia: observational versus randomized studies results. Eur Neuropsychopharmacol 2007, 17(4):235-44.
- Pendlebury J, Haddad P, Dursun S: Evaluation of a behavioural weight management programme for patients with severe mental illness: 3 year results. Hum Psychopharm Clin Exp 2005, 20:447-448.
- Pendlebury J, Bushe C, Wildgust H, Holt R: Long term maintenance of weight loss in patients with severe mental illness through a behavioural programme in UK. Acta Psychiatr Scand 2007, 115:286-294.
- Brar JS, Ganguli R, Pandina G, Turkoz I, Berry S, Mahmoud R: Effects of behavioral therapy on weight loss in overweight and obese patients with schizophrenia or schizoaffective disorder. J Clin Psychiatry 2005, 66:205-212.
- Bushe C, Haddad P, Peveler R, Pendlebury J: The role of lifestyle interventions and weight management in schizophrenia. Journal of Psychopharmacology 2005, 19(suppl):28-35.
- Bushe C, Haley C, McNamara D, Fleming-McCrossan M, Devitt P: Weight management by group interventions. The longer term outcomes at 20 months in an Irish cohort of patients with serious mental illness. Poster presented at 25th Biennial Congress of the Collegium Internationale Neuro-Psychopharmacologicum 9-13 July 2006; Chicago, IL.
- Evans S, Newton R, Higgins S: Nutritional intervention to prevent weight gain in patients commenced on olanzapine: a randomised controlled trial. Aus and N Zealand Journal of Psychiatry 2005, 39:479-486.
- Hoffmann V, Ahl J, Meyers A, Schuh L, Shults K, Collins D, Jensen L: Wellness intervention for patients with serious and persistent mental illness. J Clin Psych 2005, 66:1576-1579.
- Vreeland B, Minsky S, Menza M, Rigassio Radler D, Roemheld-Hamm B, Stern R: A program for managing weight gain associated with atypical antipsychotics. Psychiatr Serv 2003, 54(8):1155-7.
- Alvarez M, Gonzalez-Blanch C, Perez-Iglesias R, Perez-Pardal T, Martinez-Garcia O, et al.: Early intervention in antipsychotic-induced weight gain in first episode-psychosis. Schizophr Bull 2005, 21:S518.
- Ganguli R, Brar JS: Prevention of weight gain by behavioural interventions in patients starting novel antipsychotics. Schizophr Bull 2005, 21:S561.
- Lee SJ, Choi EJ, Kwon JS: A Naturalistic Multicenter Trial of a 12-Week Weight Management Program for Overweight and Obese Patients with Schizophrenia or Schizoaffective Disorder. J Clin Psychiatry in press. 2008 Feb 27
- Werneke U, Taylor D, Sanders TA, Wessely S: Behavioural management of antipsychotic-induced weight gain: a review. Acta Psychiatr Scand 2003, 108(4):252-9.
- Mauri M, Simoncini M, Castrogiovanni S, Iovieno N, Cecconi D, Dell'Agnello G, Quadrigli M, Rossi A, Donda P, Fagiolini A, Cassano GB: A psychoeducational program for weight loss in patients who have experienced weight gain during antipsychotic treatment with olanzapine. *Pharmacopsychiatry* 2008, 41(1):17-23.
- Menza M, Vreeland B, Minsky S, Gara M, Radler DR, Sakowitz M: Managing atypical antipsychotic-associated weight gain: 12-month data on a multimodal weight control program. J Clin Psychiatry 2004, 65(4):471-7.

- 25. Smith S, Yeomans D, Bushe CJ, Erikkson C, Holmes R, et al.: A Wellbeing Programme in Severe Mental Illness. Reducing Risk for Physical III-health: a post-programme service evaluation at 2 years. European Psychiatry 2007 in press.
- Werneke U, Taylor D, Sanders TA: Options for pharmacological management of obesity in patients treated with atypical antipsychotics. Int Clin Psychopharmacol 2002, 17(4):145-60.
- Faulkner G, Cohn T, Remington G: Interventions to reduce weight gain in schizophrenia. Cochrane Database of Systematic Reviews 2007:CD005148. DOI: 10.1002/14651858.CD005148.pub2

Pre-publication history

The pre-publication history for this paper can be accessed here:

http://www.biomedcentral.com/1471-244X/8/76/prepub

Publish with **Bio Med Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours you keep the copyright

Submit your manuscript here: http://www.biomedcentral.com/info/publishing_adv.asp

