Irish addiction services - past, present and future

Eamon Keenan

Ir J Psych Med 2005; 22(4): 118-120

To discuss the development of addiction services into the future, one must first examine the historical response to the misuse of substances in this country. A brief overview reveals wide disparity between, the treatment of opioid dependence in the east of the country and the way statutory services deal with the significant problem of alcohol misuse. A comprehensive model is required to deal with addiction on a national basis that necessitates the amalgamation of the drug and alcohol services.

The National Advisory Committee on Drugs (NACD) in Ireland and the Drug and Alcohol Information and Research Unit (DAIRU) in Northern Ireland have documented the extent of the problem for the island of Ireland in 2002/2003. The lifetime prevalence for any illicit drug use is 19% in the age group 15-64 years and the lifetime prevalence for alcohol use is 91% for the same age group.¹ The main illicit drug is cannabis with 24.4% of young adults (15-34 years) reporting lifetime use. It is also interesting to note that 4% in this age group have used cocaine and 0.6% have used heroin.

The response of the health services in the early 1990s to the problem of heroin dependence was largely driven by public health. The growing numbers of injecting drug users infected with HIV led to the development of a harm reduction policy, encompassing needle exchange, education for drug users and methadone maintenance. Allied to this approach was the provision of significant resources by a government that recognised the drugs problem as a serious issue of public concern. There was the coming together of statutory, community and voluntary services to produce a communitybased model of addiction treatment involving primary care with a focus to provide services for the individual drug user at a local level. There are currently over 7500 individuals in receipt of methadone treatment in Ireland with approximately one third of these attending a general practitioner. The remainder are treated at a network of community based clinics many of which involve close partnership between the health services and the local community. There are approximately 60 such clinics in the greater Dublin area and six in the rest of the country located in Portlaoise, Athlone, Carlow, Waterford, Limerick and Galway.

This current approach to the treatment of substance misuse however has not been without its critics. Some have described it as an 'opioid only service', while others suggest

*Eamon Keenan, MB, MRCPsych, Clinical Director, Addiction Services, HSE, South Western Area, Bridge House, Cherry Orchard Hospital, Dublin 10.

*Correspondence SUBMITTED: JULY 27, 2005. that what has emerged is simply a highly developed and organised 'methadone delivery service'. In addition, more recently as resources for statutory services have remained static, community groups have been applying for funding for treatment initiatives via the National Drugs Strategy. While these groups do not have the concerns of the statutory services around staff ceilings, for example, it is vitally important that these treatment initiatives are provided by trained professionals and are subject to intensive scrutiny and audit so that they may conform to professional quality norms. Thus for example, a needle exchange programme provided by the community and funded by the National Drugs Strategy team should meet and maintain appropriate standards. Careful monitoring of the impact of such a development would be required by for example, documenting the levels of blood borne viruses within that community on an ongoing basis.

In contrast to the resources provided for the development of drug services, the provision of resources for alcohol services has been grossly inadequate. During this period of lack of resources, per capita drinking in Ireland increased dramatically. Between 1989 and 2000 per capita alcohol consumption in Ireland increased by 41% and in 2001 Ireland had the second highest per capita consumption of alcohol in the enlarged EU.² The SLAN survey in 2002 highlighted the fact that 30% of males and 22% of females consumed more than the recommended weekly limits of sensible alcohol consumption.³ The ESPAD report of 1999 showed that Irish 16 year olds are among the highest alcohol abusers in Europe in terms of binge drinking and drunkenness.⁴

The contribution of alcohol to morbidity and mortality is well documented. In 2002, the WHO identified alcohol related death and disability as accounting for 9.2% of all burden of disease in developed countries behind tobacco and hypertension.5 Irish healthcare costs for alcohol in 2003 were estimated at €433 million, however the vast majority of these costs were related to treating the consequences of alcohol abuse with only a small provision for alcohol treatment.6 In many cases, alcohol counsellors provide services for individuals with alcohol problems in isolation from other supports such as multidisciplinary mental health teams. For psychiatric services it should also be noted that in 2002 alcoholic disorder was the second highest cause of admission to psychiatric hospitals for males and the fourth highest cause for females.7 Developing an effective community based alcohol service would reduce considerably this admission rate.

The recent Strategic Taskforce on Alcohol second report in 2004 gave specific recommendations to reduce consumption and limit harm within the population. These included, decreasing availability, increasing taxation and putting in place effective treatment services utilising specialist treat-

ment. The government has ignored these recommendations.

More recently dual diagnosis (co-existence of serious mental health disorder and an addictive disorder) has gained increasing recognition in this country as elsewhere. The National Household Study in the UK found psychological morbidity present in 12% of non-dependent individuals, 30% of alcohol dependent individuals and 45% of drug dependent individuals.9 In the Comorbidity of Substance Misuse and Mental Illness Collaborative study (COSMIC), 40% of patients managed in community mental health settings reported problem drug use and/or harmful alcohol use in the past year.10 Unless the individual receives appropriate interventions for both conditions then outcomes in this cohort will remain poor and relapses high.

Various models have been proposed to deal with dual diagnosis. These include the serial model (one condition treated after another), the parallel model (the two services work together to treat the individual with joint planning and liaison) and the integrated model (dedicated team to treat both conditions). Although the integrated model has found favour in the US, there is inconclusive evidence for this expensive approach." An integrated model may result in isolation of dual diagnosis from mainstream services and staff in mainstream services may become deskilled. A care programme approach as advocated by the Mental Health Policy Guide in the UK may have merit.12 This approach advocates co-ordination between mental health teams and substance misuse teams and closer links in terms of training and supervision and the augmentation of teams with a dual diagnosis worker.

In addition to the recognition of strong links between addiction services and mental health services, there are a number of links with other healthcare providers which would benefit patients. Primary care can play a pivotal role in the detection, evaluation and treatment of drug and alcohol abuse. While good links have been established between the drugs services and primary care, these could be strengthened in relation to alcohol services. Liaison between maternity and drug services improves antenatal care and decreases neonatal morbidity particularly in the case of opioid dependence.13

For HIV services, supervision of antiretroviral therapy at methadone treatment clinics can improve the adherence to and outcome for treatment.14 Closer links with hepatology services could improve hepatitis C treatment outcomes with supervision of interferon and ribavirin for infected drug users at addiction centers. For accident and emergency services it has been demonstrated that brief interventions for alcohol related problems on-site can reduce reattendance.¹⁵ Prison healthcare services and homelessness services will benefit from stronger links with addiction services given the high levels of substance misuse among these cohorts. Continuation of care in relation to individuals recently released from prison is a vital factor in the reduction of fatal overdose for this group. One further area where substance misuse services could contribute an intervention could be in services to the elderly. The problem of long-term benzodiazepine use and also alcohol abuse in individuals over the age of 65 has been recognised. Up to 20% of female general medical services patients in this age group have been shown to be in receipt of scripts for benzodiazepines over a three-month period.16 To date, no effective strategies have been implemented to address this concern.

The treatment of substance misuse problems in young people under the age of 18 is one area where Ireland can justifiably claim to be at the forefront of international practice. The appointment of consultant child and adolescent psychiatrists in substance misuse and the provision of some resources to develop a service is a welcome innovation and the outcomes for such a strategy are awaited with interest.

Clearly addiction problems impact significantly on a broad spectrum of other health domains. Currently the drug and alcohol services in Ireland are operating by and large independently of one another. The rational for combining these services, one of which has received significant funding for mainly political reasons while the other has received little funding, perhaps also for political reasons, is now becoming irrefutable. Co-dependence on drugs and alcohol occurs frequently among individuals particularly in younger age groups.17 Treatment approaches used in the management of addiction are similar. Professional competencies of staff working in the areas of drug and alcohol dependence are similar. A combined service would improve the support provided to primary care in the area of alcohol treatment. A combined service is more likely to meet the needs of both sets of patients. By integrating the services, resources for addiction could be lobbied for more effectively and allocated according to need particularly in the case of alcohol services nationally.

The future development of addiction services requires a combined drug and alcohol service under the umbrella of mental health. Linkages with other health providers need to be strengthened through the appointment of liaison workers and improved communication and co-operation. Research into the area of addiction needs to be supported and developed further in order that effective and quality focused services can be implemented and maintained. This may require the setting up of a strong academic base and programme in the context of a National Addiction Centre similar to the UK. For services to be effective, sufficient resources are required to develop a comprehensive model. Under the new Health Services Executive structure, each local health area in the country will need access to an addiction service under the direction and leadership of a consultant psychiatrist in substance misuse. The success and experience of the drugs service in developing a community based treatment service involving statutory, community and voluntary agencies can provide a template for a similar approach in relation to alcohol services. This should complement and enhance the implementation of the national alcohol strategy.

Declaration of interest: None

References

- 1. NACD & DAIRU. Drug use in Ireland and Northern Ireland. First results from the 2002/2003 Drug prevalence study. Bulletin I and II, 2005.
- Health for all database. Geneva: World Health Organisation, 2001.
 Health Promotion Unit, Department of Health and Children and the Centre for Health Promotion Studies, National University of Ireland, Galway. Survey of lifestyle, attitudes and nutrition (SLAN). Galway: The National Health and Lifestyles Surveys, 2003. 4. Hibell B, Andersson B, Ahlstrom S, Balakireva O, Bjarnasson T, Kokkevi A, Morgan
- M. The 1999 ESPAD Report: Alcohol and other drug use among students in 30 European countries. The Swedish council for information on alcohol and other drugs. The Pompidou group at the Council of Europe, 2000.
- 5. The World Health Report 2002: Reducing risks, promoting healthy life. Geneva: World Health Organisation, 2002.
- 6. Byrne S. Update on estimates of the cost of alcohol problems in Ireland. Strategic taskforce on alcohol second report, 2004.
- 7. Daly A, Walsh D. Activities of the Irish Psychiatric Services 2002. Health Research

Board, 2003.

- 8. Strategic task force on alcohol, second report. Health Promotion Unit, Department of Health and Children, 2004.
- 9. Farrell M, Howes S, Bebbington P et al. Nicotine, alcohol and drug dependence and psychiatric comorbidity. Results of a national household survey. Br J Psychiatry 2001; 179: 432-37.
- 10. Weaver T, Madden P, Stimpson CG et al Comorbidity of substance misuse and mental illness in community mental health and substance misuse services. Br J Psychiatry 2003: 183: 304-13.
- 11. Tyrer P, Weaver T. Desperately seeking solutions: the search for appropriate treatment for comorbid substance misuse and psychosis. Psychiatric Bull 2004; 28: 1-
- 12. Mental health policy implementation guide: Dual diagnosis good practice guide.

London: Department of Health, 2002.

- 13. Scully M, Geoghegan N, Corcoran P, Tiernan M, Keenan E. Specialised drug liaison midwife services for pregnant opioid dependent women in Dublin, Ireland. J Subst Abuse Treat 2004; 26(1): 329-35.
- 14. Clarke S, Keenan E, Ryan M, Barry M. Mulcahy F. Directly Observed Therapy for Injecting Drug Users with HIV Infection. The AIDS Reader 2002; 12(7): 305-16.
- 15. Crawford MJ, Patton R, Touquet Ret al. Screening and referral for brief intervention of alcohol-misusing patients in an emergency department: a pragmatic randomised controlled trial. Lancet 2004; 364: 1334-39.
- 16. Keenan E. An Irish Perspective on Benzodiazepine Prescribing. Contribution to the Sensible Use of Benzodiazepines. Council of Europe Publishing, 2001; 53-63.

 17. O'Malley PM. Alcohol use among adolescents. Alcohol health and research world
- 1998; 22: 85-93.

Cover Image: 'Elevation' 2003

Stephen Vaughan's work examines the varied landscapes and environments both internal and external that mankind inhabits. He works with large plates using all the techniques of intaglio to create highly textured images. He likes to think of this as 'building' plates and in this sense there is a sculptural aspect to the work. It is an act of stimulation. At times his prints have also approximated the characteristics of paintings. Vaughan's work is multifaceted, culminating in the convergence of ideas, experiences and events, both current and historical. Stephen Vaughan graduated from the Crawford College of Art and Design with a BA in Fine Art and Printmaking, in 1994. 'Elevation' is one of two Vaughan prints in the St. James's Hospital art collection. Vaughan has exhibited his work extensively in Ireland and internationally. His artwork features in many public, private and corporate collections.

Guidelines for Authors

The Journal's aim is to publish original scientific contributions in psychiatry, psychological medicine (including surgery and obstetrics), and related basic sciences (neurosciences, biological, psychological, and social sciences). Its scope includes any subspecialties of the above, eg. behavioural pharmacology, biological psychiatry, child and adolescent psychiatry, mental handicap, forensic psychiatry, psychotherapies, psychiatry of old age, epidemiology, rehabilitation, psychometrics, substance misuse, sexual studies, linguistics, and the history, philosophy and economics of psychiatry.

The Journal will accept original papers, clinical case reports, brief research reports, review articles, perspective articles, historical papers, editorials, practice reviews (medical audits), letters to the editor and book reviews. Review articles are usually invited. Original data papers receive top priority for speedy publication. Manuscripts should be prepared in accordance with the guidelines of the International Committee of Medical Journal Editors.1 All case reports must have the patient's consent before an article can be published.

The page following the title page should carry an Abstract followed by a list of three to 10 Key Words or short phrases drawn, if possible, from the medical subject headings (MeSH) list of Index Medicus.

The Title, Key Words and Abstract should be chosen to help future literature searchers. The Abstract, up to 150 words for an unstructured or 250 words for the structured abstract,2 should state specifically the main purposes, procedures, findings and conclusions of the study, emphasising what is new or important. For original papers, brief research reports, medical audits and review articles, a structured abstract2 is required, using the headings Objectives, Methods, Results (Findings for review articles) and Conclusions.

Under the Abstract heading of Method, include wherever applicable the study design, setting, patients/participants (selection criteria, description), interventions, observational and analytical methods and main outcome measures. (For review articles specify the methods of literature search and selection). Under the Abstract heading of Results, give the most important specific data together with their statistical significance.

Timely references should highlight the study's relevance to current research or clinical practice. References to journal articles13 and to books46 illustrate the 'Vancouver' style,' ie. number references in the order they appear in the text, do not alphabetise. Journal titles should be abbreviated as in Index Medicus. The Uniform requirements for manuscripts submitted to biomedical journals* has two paragraphs on statistical guidelines. These have been explained by Bailar and Mosteller.3

Figures and graphs should be clear and of good quality, and should be accompanied by relevant data to facilitate redrawing where necessary. All materials sent for publication should be accompanied by a covering letter signed by all the authors, and such material will become the property of the httplournal until, and if, publication is refused. Material somefaged should not be sent elsewhere for publication. One copy of the manuscript should be retained by the author(s) for reference, and four copies of the manuscript and covering letter, one of these being the original, should be sent to:

The Editor, Irish Journal of Psychological Medicine, 25 Adelaide Street, Dun Laoghaire, Co Dublin, Ireland.

Electronic submissions

Manuscripts may be sent via email to the following address: psychological@medmedia.ie Full postal address, telephone and fax numbers should be included. Where possible tables, figures and text should be included in the same document. There is no need to also submit by post or fax.

All contributions are peer-reviewed by three anonymous assessors and, where relevant, by the Statistical Editor whom authors may contact for help. Assessments will be sent to the corresponding author usually within six weeks. Where revisions are sought prior to publication, authors are advised to return their revision (in quadruplicate if posting), incorporating any suggestions which they agree would improve their paper. The covering letter should respond to each comment, numbered, of each assessor, indicating where the revision deals with it, or why the authors disagree or cannot incorporate it.

Each assessor will then receive the authors' revision, covering letter and the previous comments of the other assessors. After the assessors' further comments have been received, the senior editors will make the final decision. including priority and time of publication, and the right to style and if necessary shorten material for publication.

Manuscripts are considered with the understanding that they have not been published previously, either in print or electronic format.

In the interest of accountability all financial and material support for the research and the work should be clearly stated.7 Authors of original data must take responsibility for the integrity of the data and accuracy of the data analysis. All authors must have full access to all the data in the study.

Authors should obtain permission from the individuals named in Acknowledgments, since readers may infer endorsement.

References

- International Committee of Medical Journal Editors. Uniform requirements for manuscripts submitted to biomedical journals. BMJ 1991; 302: 338-41
- 2. Haynes RB, Mulrow CD, Huth EJ, Altman DG, Gardner MJ. More information abstracts
- revisited. Ann Intern Med 1990; 113: 69-76.

 3. Bailar JC, Mosteller F. Guidelines for statistical reporting in articles for medical journals. Ann Intern Med 1988 Feb; 108(2): 266-73.

 4. Daly LE, Bourke GJ, McGilvray J. Interpretation and uses of medical statistics. 4th ed.
- Oxford: Blackwell Scientific Publications, 1991: 428-31
- 5. Gardner MJ, Altman DG, editors. Statistics with confidence confidence intervals and statistical guidelines. London: British Medical Journal, 1989: 103-5. [Note: British Medical Journal here is the publisher of a book, not the journal BMJ.
- 6. American Psychiatric Association. Diagnostic and statistical manual of mental disorders. Washington DC: American Psychiatric Association, 1987.
- 7. DeAngelis CD, Fontanarosa PB, Flanagin A. Reporting financial conflicts of interest and relationships between investigators and research sponsors. JAMA 2001; 286: 89-91.

Get patients with depression back in touch with life





VENLAFAXINE XI

First-line reconnection

Presentation: Efexor XL: capsules containing 75mg or 150mg ventafaxine (as hydrochloride) in an extended release formulation. Efexor: tablets containing 37.5mg or 75mg ventafaxine (as hydrochloride) Use: Treatment of depressive illiness, including depression accompanied by anxiety. Generalised Anxiety Disorder (GAD) primarily characterised by chronic and excessive worry and anxiety for at least 6 months; for the prevention of relapses of the initial episode of depression or for the prevention of the recurrence of new depressive episodes. Dosage: Adults (including the elderly): Depressive illness including depression accompanied by anxiety: Efexor XL: Usually 75mg, given once daily with food, increasing to 150mg once daily if necessary. The dose can be increased further to 225mg once a day. Dose increments should be made at intervals of approximately 2 weeks or more, but not less than 4 days. Efexor Usually 75mg (375mg bd) if mecessary. In more severely depressed patient, 150mg/day increasing every 2 to 3 days in up to 75mg/day increasing every 2 to 3 days in up to 75mg/day increasing every 2 to 3 days in up to 75mg/day increasing one of the prevention of recurrence of a new episode, is similar to that used during the reducing to usual dose consistent with patient response. Prevention of Relapse/recurrence: Usually, the dosage for prevention of relapse, or for prevention of recurrence of a new episode, is similar to that used during the index episode. Patients should be re-assessed regularly in order to evaluate the benefit of long-tarm therapy. Generalised Anxiety Disorder: Efexor XL: Usually 75mg, given once daily with food, increasing to 150mg once daily if necessary. The dose can be increased further to 225mg once a day. Dose increments should be made at intervals of approximately 2 weeks or more, but not less than 4 days. Discontinuation: Discontinua directed below 18 years of age. Moderate renal or moderate hepatic impairment: Doses should be reduced by 50%. Not recommended in severe renal or severe hepati

impairment. Contra-indications: Concomitant use with MAOIs, hypersensitivity to venlataxine or other components, patients aged below 18 years.
Precautions: The risk of suicide should be considered in all patients. Use with caution in patients with myocardial infarction, unstable heart disease, renal or hepatic impairment, narrow angle glaucoma, mania, a history of epilepsy (discontinue in event of seizure), using neuroleptics or diuretics or predisposed to bleeding. Patients should not drive or operate machinery if their ability to do so is impaired. Possibility of postural hypotension (especially in the elderly). Prescribe smallest quantity of capsules or tablets according to good patient management. Blood pressure monitoring is recommended. Advise patients to notify their doctor should an allergy develop or if they become or intend to become pregnant. Patients with a history of drug abuse should be monitored carefully. Cholesterol measurement is recommended with long term use. Venlafaxine should not be used with veight loss agents. Usually not recommended during pregnancy or lactation. Interactions: MAOIs: do not use venlafaxine in combination with MAOIs or within 14 days of stopping MAOI treatment. Allow 7 days after stopping venlafaxine before starting am MAOI. Use with caution in elderly or hepatically impaired patients taking cimetion, in patients taking other CNS-active drugs in particular serotonergic drugs, clozapine or haloperidol; in patients taking warrafin and in patients taking direction, in patients taking overfacion, in advised with concurrent use of ECT. Side-effects: Most commonly occurring: constipation, neusea, asthenia, headeche, dizziness, dry mouth, insomnia, nervousness, somnolence, abnormal ejaculation/orgasm, sweating. Also reported: vasodilatation, hypotension/postural hypotension, hypertension, palepitation, syncope, ecchymosis, mucous membrane bleeding, Gl bleeding, anorexia, appetite decreased, diarrhoea, dyspepsia, vomiting, abdominal pain,

bruxism, abnormal dreams, chills, pyrexia, weight gain or loss, increased serum cholesterol hyponatraemia, increased liver enzymes, arthralgia, myalgia, muscle spasm, agitation, anxiety, confusion, hypertonia, paraesthesia, tremor, myoclonus, apathy, hallucinations, urinary frequency and retention, anorgasmia, eractile dysfunction, decreased libido, impotence, menstrual cycle disorders, menorrhagia, dyspnoea; prurits, rash, angioedema, maculopapular eruptions, urticaria, photosensitivity reactions, alopecia, mydriasis, tinnitus, abnormal vision/accommodation, alterat taste sensation. Hostility and suicidal ideation in paediatric patients. Rarely reported: thrombocytopenia, haemorthage, prolonged bleeding time, arrhythmias, hepatitis, SIADH, ataxia and disorders of balance and co-ordination, speech disorders including dysarthria, extrapyramidal disorders including dyskinesia, dystonia, mania or hypomania, neuroleptic malignant syndrome-like effects or serotonergic syndrome, galactorrhoea, erythema multiforme, Stevens-Johnson syndrome, very rarely anaphylaxis, blood dyscrasias, EGG changes, pancreatitis, increased prolactin, rhabdomyolysis, delirum, pulmonary esionphilas, Symptoms reported on discontinuation of venlafaxine were mostly non-serious and self-limiting and included dizziness, insomnia, nausea and nervousness. PA numbers: Efexor XI. 75mg capsule (PA 22/65/5) Efexor XI. 150mg capsule (PA 22/65/6) Efexor 37.5mg tablet (PA 22/65/4). Legal category: S1A. Further information is available upon request from: Wyeth Pharmaceuticals, M50 Business Park, Ballymount Road Ulper, Walkinstown, Dublin 12. Marketing Authorisation Holder. John Wyeth & Brother Limited, Taplow, Maidenhead, Berkshire, SL6 OPH. Date of preparation: 21 December 2004