

## Alcoholism – treating ‘the second sex’

Diana G Patterson, MD, MRCPsych, Consultant Psychiatrist, Area Mental Health Unit, Tyrone and Fermanagh Hospital, Omagh, Co. Tyrone, BT79 0NS, Northern Ireland.

Much of our theory surrounding the development and treatment of problem drinking women has derived from our knowledge of drinking in men. This has led to deficits in the mainstream treatment for problem drinking women.

Most treatments have been developed by male providers for the treatment of male patients.<sup>1</sup> The ‘male-as-norm’ bias defines male alcoholism as the standard by which female alcoholism is judged.<sup>2</sup> This has affected research, assessment and treatment standards for women. Weaknesses in our existing theoretical models spring not only from the fact that women have not been adequately studied, but also from the fact that we have stereotyped ideas about female alcoholics.<sup>3</sup> There has been an increase in the number of research and review reports addressing gender over the last number of years. Despite this, Gomberg<sup>3</sup> describes women as “the second sex” in theory development and diagnostic definition.

Throughout history women have been drinking for as long as men. In many cultures women drink as often as men and in some cultures women drink more than men, despite the fact that a given amount of alcohol has a greater physical impact on women. Heath<sup>5</sup> in a review of world-wide literature on women and alcohol suggests that norms, values, attitudes and expectations may be as important as physiological differences between the sexes with respect to treatment outcomes. Too narrow a focus on physical, psychological and social problems may have hampered our understanding of women’s diverse roles with respect to alcoholism.

A review of published material by Tonaetto *et al*<sup>6</sup> shows no sex differences in the outcome of treatment, but 72% of studies do not address gender. Treatment units and prevention strategies would benefit a greater number of female substance abusers if they were tailored to suit women’s specific needs.<sup>7</sup> Women usually display solitary drinking, neurotic and family problems, early health impairment and abuse of other drugs. Therapy may be enhanced by addressing these specific factors. Prevention programmes may also benefit by addressing them.

Changes have occurred in women’s drinking patterns over time but epidemiological studies have not shown the expected convergence of male and female drinking rates.

Antecedents to female drinking include difficulties in impulse control, depression and other psychiatric diagnoses such as eating disorders.<sup>4</sup> Societal attitudes may act as protective or destructive factors in women’s alcoholism.<sup>9</sup> Smith<sup>10</sup> considers factors which may affect a woman’s ability to enter, maintain and sustain treatment. The onset of heavy drinking tends to be at an older age in women, and they tend to have a shorter history at the time of presentation for treatment.<sup>11</sup>

Specific issues encountered by women include incest,<sup>12</sup> homelessness,<sup>13</sup> employment<sup>14</sup> and aggression, both as aggressor and as the object of aggression.<sup>15</sup> Pregnant or parenting female alcoholics have problems which may be best addressed by co-ordinated, comprehensive ‘family centred’ care.<sup>16</sup> We need to acknowledge the effects of alcohol on women’s relationships as daughters, partners and parents. Consideration must also be given to its role in violence against women. Primary care physicians have a unique role in the treatment of female alcoholics. They need accurate information about female alcoholism in order to overcome prejudice, and to enable them to offer the most appropriate treatment.<sup>17</sup>

Schmidt *et al*<sup>18</sup> have developed an extensive taxonomy on indicators of women’s alcohol problems. Important factors appear to relate to individual characteristics including psychological and behavioural issues. Social factors are important, in particular family or partner relationships. Physical appearance also seems to bear importance.<sup>19</sup> This review was drawn from the literature over a spell from 1970 to 1986 and the authors suggest that the resulting taxonomy may serve as a base line against which to assess material from other sources such as ethnographic studies or studies of the general population.

Corrigan’s study, on page 48 of this issue, is timely and addresses some of these issues as they pertain to female drinkers in Ireland. This is a replication of Corrigan’s study of American alcoholic women in treatment.<sup>20</sup> The American study found that 84% of 150 women recruited into the study drank most often in their own homes. Professionals and those employed were more likely to drink in restaurants and bars. Two thirds of the women hid their drinking and most offered escapist rather than social reasons for their drinking. More

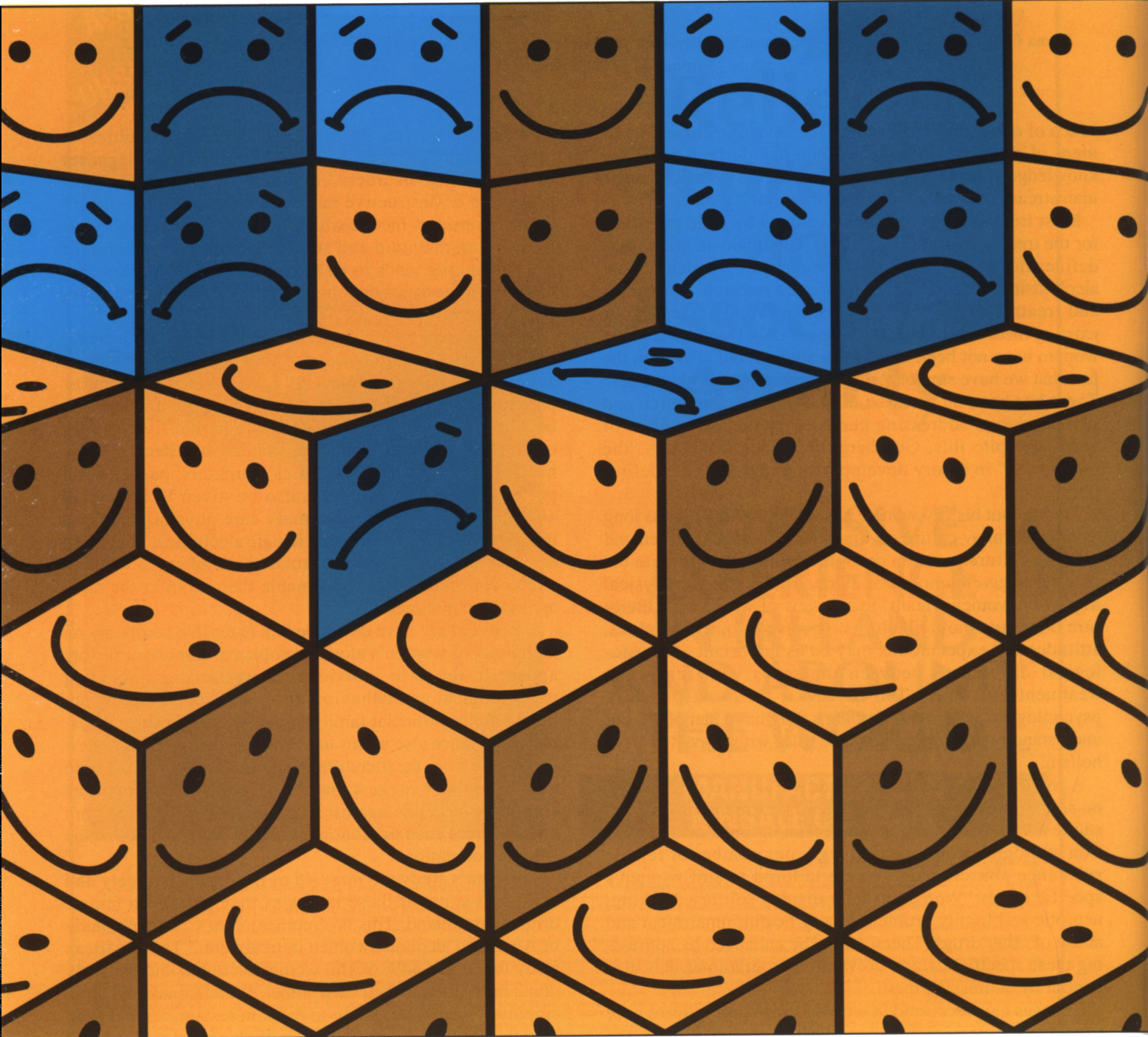
**Editor-in-Chief:** Brian Lawlor (Dublin). **Editors:** Timothy Dinan (London), David King (Belfast). **Deputy Editor:** Brian O’Shea (Dublin). **Associate Editors:** Ken Brown (Belfast), Patricia Casey (Dublin), Anthony Clare (Dublin), Stephen Cooper (Belfast), Thomas Fahy (Galway), Michael Fitzgerald (Dublin), Michael Kelleher (Cork), Brian Leonard (Galway), Roy McClelland (Belfast), Aidan McGennis (Dublin), Ciaran O’Boyle (Dublin), Eadbhard O’Callaghan (Dublin), Art O’Connor (Dublin), Ethna O’Gorman (Belfast), Ian Pullen (Edinburgh), David Sheehan (Tampa), Philip Snaith (Leeds), Hugh Staunton (Dublin), John Waddington (Dublin), Richard Williams (Calgary). **Statistical Editor:** Leslie Daly (Dublin). **Deputy Statistical Editor:** Ronan Conroy (Dublin).



**Abbreviated Prescribing Information: LUSTRAL® (sertraline)**

**Presentation:** Tablets containing 50mg or 100mg sertraline. **Indications:** Treatment of symptoms of depressive illness. Prevention of relapse or recurrence of depressive episodes. **Dosage:** LUSTRAL should be given as a single daily dose with food. The initial dose is 50mg and the usual therapeutic dose is 50mg or 100mg daily. Dosage can be further increased, if appropriate, to 150mg or a maximum of 200mg daily. Patients should be maintained on the lowest effective dose. **Use in children:** Not recommended. **Use in the elderly:** Usual adult dose. **Contra-indications:** Hypersensitivity to LUSTRAL. Hepatic insufficiency, unstable epilepsy and convulsant disorders, pregnancy and lactation. Do not use with, or within two weeks of ending treatment with, MAOIs. At least 7 days should elapse before starting any MAOI following discontinuation of LUSTRAL. **Precautions, Warnings:** Renal insufficiency, ECT, epilepsy, driving. LUSTRAL should not be administered with benzodiazepines or other tranquilizers in patients who drive or operate machinery. The patient should be monitored for signs of suicide or mania. LUSTRAL

has not been observed to produce dependence. **Drug Interactions:** Administer with caution in combination with other centrally active medication (e.g. lithium, tryptophan). Although LUSTRAL has been shown to have no adverse interaction with alcohol, concomitant use with alcohol is not recommended. The potential for LUSTRAL to interact with other highly protein bound drugs should be borne in mind. The potential of LUSTRAL to interact with e.g. propranolol and phenytoin has not been fully assessed. **Side-Effects:** Dry mouth, nausea, diarrhoea/loose stools, ejaculatory delay, tremor, increased sweating and dyspepsia. **Legal Category:** S1A. **Package Quantities:** 50mg tablet (PA 19/46/4) Calendar pack of 28; 100mg tablet (PA 19/46/5) Calendar pack of 28. Further information on request. **Invicta® Pharmaceuticals:** A Division of Pfizer Limited, Sandwich, Kent. **Invicta® Pharmaceuticals office in Dublin:** Pharmapark, Chapelizod, Dublin 20. Tel: Dublin 6268340. \*Trade Mark



 **LUSTRAL<sup>\*</sup> 50mg**  
sertraline

**Builds success right from the start**



than half reported a specific stressor in the period preceding their drinking, particularly emotional losses. Many were unhappy, lonely, anxious and depressed. Almost half of the subjects used other drugs while drinking. The outcome of treatment for these American women was much better than the outcome in comparable studies for men. Of 116 completed interviews 13 months after treatment, 41% had not had a drink since leaving treatment. A further 12% drank on only four occasions or less and 71% had had no alcohol in the previous month. Professional women and those in higher socio-economic groups were more likely to remain abstinent. The entire group showed improvement in symptoms of emotional ill health, but regression analysis suggested that this improvement was due to abstinence from alcohol.

Despite the fact that abstinence was the aim in treatment for the Irish women described in Corrigan's study in this issue, only 26% remained abstinent in the 13 months after treatment. A further 24% drank up to four times and 43% had had no drink in the previous six months. Self-esteem was found to be improved in those who were abstinent. Seasoned therapists will derive some reassurance from the 74% accuracy of staff predictions about clients outcome. The number of years working with alcoholics and the number of alcoholics treated were factors which influenced successful predictions.

The face of alcoholism treatment is changing as efforts are made to provide forms of treatment delivery which are cost-effective. It is imperative that such changes are based on relevant theory supported by adequate clinical research. In the course of conducting this research and in the course of designing appropriate prevention and treatment strategies, we must ensure that there is not further marginalisation of the second sex.

#### References

1. Abbott AA. A feminist approach to substance abuse treatment and service delivery. *Social Work in Health Care* 1994; 19(3-4): 67-83.
2. Wilke D. Women and alcoholism: how a male-as-norm bias affects research, assessment and treatment. *Health and Social Work* 1994; 19(1): 29-35.
3. Nixon SJ. Recent developments in alcoholism: typologies in women. *Recent Developments in Alcoholism* 1993; 11: 305-23.
4. Gomberg ES. Recent developments in alcoholism: gender issues. *Recent Developments in Alcoholism* 1993; 11: 95-107.
5. Heath DB. Women and alcohol: cross-cultural perspectives. *J Sub Abuse* 1991; 3(2):175-85.
6. Toneatto A, Sobell LC, Sobell MB. Gender issues in the treatment of abusers of alcohol, nicotine and other drugs. *Journal of Substance Abuse* 1992; 4(2): 209-18.
7. Oppenheimer E. Alcohol and drug misuse among women – an overview. *Br J Psychiatry* 1991; (10) Supple: 36-44.
8. Nespor K. Treatment needs of alcohol-dependent women. *Int J Psychosom* 1990; 37(1-4): 50-2.
9. Blume SB. Chemical dependency in women: important issues. *Am J Drug Alcohol Abuse* 1990. 16(3-4): 297-307.
10. Smith L. Help seeking in alcohol-dependent females. *Alcohol and Alcoholism* 1992; 27(1): 3-9.
11. Ross HE. Alcohol and drug abuse in treated alcoholics: a comparison of men and women. *Alcohol Clin Exp Res* 1989; 13(6): 810-6.
12. Hurley DL. Women, alcohol and incest: an analytical review. *Journal of Studies on Alcohol* 1991; 52(3): 253-68.
13. Robertson MH. Homeless women with children. The role of alcohol and other drug abuse. *Am Psychol* 1991; 46(11): 1198-204.
14. Wilsnack RW, Wilsnack SC. Women, work and alcohol: failures of simple theories. *Alcoholism* 1992; 16(2): 172-9.
15. Gomberg ES. Alcohol, women and the expression of aggression. *J Stud Alcohol* 1993; 11 Supple: 89-95.
16. Finkelstein N. Treatment issues for alcohol – and drug-dependent pregnant and parenting women. *Health and Social Work* 1994; 19(1): 7-15.
17. Quinby PM, Graham AV. Substance abuse among women. *Primary Care* 1993; 20(1): 131-40.
18. Schmidt C, Klee L, Ames G. Review and analysis of literature on indicators of women's drinking problems. *Br J Addict* 1990; 85(2):179-92.
19. Klee L, Schmidt C, Ames G. Indicators of women's alcohol problems: what women themselves report. *Int J Addict* 1991; 26(8): 879-95.
20. Corrigan EM. *Alcoholic women in treatment*. New York: Oxford University Press, 1980.

Original manuscript received February 1, 1995  
 Manuscript accepted March 15, 1995

## The John Dunne Medal

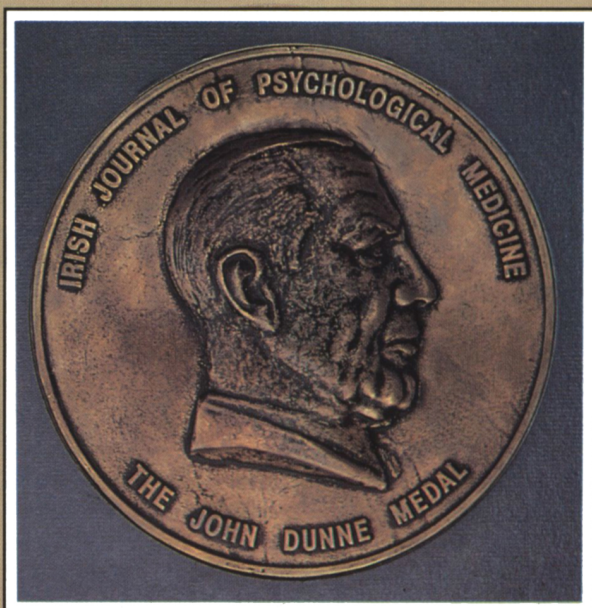
Sponsored by *Eli Lilly*

The **John Dunne Medal** is awarded annually by the *Irish Journal of Psychological Medicine*. Trainees from Ireland, Northern Ireland and Great Britain are eligible. The trainee shall have contributed substantially, though not necessarily as the first author, to an original paper (exceptionally an article in another category) published by the Journal in the previous year.

The bronze medal is named after Dr John Dunne, the first psychiatry professor in Ireland and the president in 1955 of the Royal Medico-Psychological Association. The bronze medal was sculpted by Robin Buick, ARHA. It has been exhibited by the Royal Hibernian Academy.

The international panel of adjudicators selects the medal winner on the basis of the paper's originality, method, and relevance for future research or clinical practice.

The first authors of articles published in December 1994, March 1995, June 1995 and September 1995, are invited to submit a trainee's name into the competition for the 1995 prize. The winner will be announced in the December 1995 issue.



Our thanks to *Eli Lilly* for their continued support





Treats the Present and Safeguards the Future

# Tegretol Retard

carbamazepine  
specially formulated to reduce peak plasma levels

For both generalised and partial seizures

**PRESCRIBING INFORMATION. Presentation:** TEGRETOL RETARD is a formulation which reduces the peak concentration of active substance in the plasma and also ensures that fluctuations in plasma concentration are reduced throughout the day. **Indications:** Epilepsy (generalised tonic-clonic and partial seizures). Trigeminal Neuralgia and other forms of deafferentation pain. Alcohol withdrawal symptoms. Treatment of mania and prophylaxis of manic-depressive illness. **Dosage:** Epilepsy: Adults: 100-200mg, once or twice daily, increasing slowly up to 800-1200mg daily, in divided doses. **Children:** See full prescribing information. Liquid is recommended for children under 5 years. It may be helpful to monitor drug levels: the optimum therapeutic plasma level ranges from 3-10mcg/ml (13-42 micromoles/L). TEGRETOL RETARD should not be chewed, but swallowed whole with a little water. For dosage of other indications: see full prescribing information. **Contra-indications:** Previous drug sensitivity to carbamazepine. Do not administer to patients with atrioventricular conduction abnormalities unless paced. **Precautions:** Blood counts should be performed before

commencing treatment, at weekly intervals during the first month and subsequently monthly for five months then two to four times a year. Liver function tests should be performed before treatment and periodically during therapy. Withdraw TEGRETOL if allergic skin reactions, deterioration of liver function, or severe, progressive or clinically manifest leucopenia occur. Caution in patients taking oral anticoagulants or concomitant anti-epileptic or lithium therapy, or requiring oral contraception. Macrolide antibiotics (e.g. erythromycin), isoniazid, some calcium antagonists (i.e. Verapamil, diltiazem), dextropropoxyphene, viloxazine and cimetidine may elevate carbamazepine levels. CNS side-effects may be exacerbated by alcohol. TEGRETOL may impair reactions of patients driving or operating machinery. Serum folic acid levels should be observed during anti-convulsant therapy. See full prescribing information. **SIDE-EFFECTS:** Occasionally dizziness, diplopia, headache, somnolence, ataxia, disorders of visual accommodation; confusion and agitation; dry mouth, nausea, diarrhoea or constipation; loss of appetite; generalised erythematous rash; leucopenia, thrombocytopenia; oedema, fever. Isolated cases exfoliative

dermatitis, Stevens-Johnson Syndrome, toxic epidermal necrolysis, hair loss; proteinuria, lymph-node enlargement and acute renal failure; agranulocytosis, aplastic anaemia and thromboembolism (blood count should be checked regularly in early stages of treatment). Rarely dose-dependent hyponatraemia, disturbance to cardiac conduction, hepatitis. **LEGAL CATEGORY:** S1B. Packs of TEGRETOL RETARD divisible tablets of 200mg (PA 11/1/5) in blister packs of 100; TEGRETOL RETARD divisible tablets of 400mg (PA 11/1/6) in blister packs of 100. ® denotes Registered Trademark. Full prescribing information is available from Geigy Pharmaceuticals, Beech House, Beech Hill Office Campus, Clonskeagh, Dublin 4. Telephone (01) 2601255. **DATE OF PREPARATION** February 1994. © Ciba Pharmaceuticals 1994.

**Geigy**

Your Partner in Health Care

G9403