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'Portrait of Pearse Hutchinson' by Edward McGuire (oil on canvas, 124 x 120.5cm)
From the Collection at the Irish Museum of Modern Art, Royal Hospital Kilmainham, Dublin 8.
(on loan, private collection)

A brighter outlook for prescribers of fluoxetine



The original brand? - **25%** more expensive.¹
Is it time to change?

Gerozac®

fluoxetine

A more affordable therapy in the treatment of depression



GERARD
LABORATORIES

GEROZAC: (fluoxetine HCL) Abbreviated prescribing information: Presentation: Each capsule contains fluoxetine hydrochloride equivalent to 20 mg of fluoxetine. **Indications:** GEROZAC is indicated for the treatment of major depressive episodes. **Dose:** A dose of 20 mg/day is recommended and a maximum daily dose should not exceed 80 mg/day, which can be administered as single or divided dose, during or between meals. **Patients with renal or liver disease:** In cases of liver dysfunction or renal failure (GFR 10-50 ml/min), the dose should be reduced, e.g. to 20 mg every second day. **Children:** Fluoxetine capsules are not indicated for use in children and adolescents below the age of 18. **Elderly:** Caution is recommended when increasing the dose, which should rarely exceed 40 mg and should not exceed 60 mg. **Method of administration:** For oral administration. **Contraindications:** Concurrent treatment with MAOIs (monoamine oxidase inhibitors). Cautionary use with other antidepressants. Not to be used where there is severe renal failure (GFR < 10ml/min). Unstable or uncontrolled epilepsy. Not to be used by nursing mothers. Hypersensitivity to any of the ingredients. **Precautions:** As with all antidepressants risk of suicide particularly at the beginning of treatment due to the delay between treatment and clinical improvement. Concomitant use of tryptophan. Epilepsy, electroconvulsive therapy, cardiovascular disease, recent myocardial infarction, diabetes, alcohol, hepatic and renal insufficiency, and overdose. **Side-effects:** rash and allergic reaction, psychosis and mood shift towards manic phase, serotonin syndrome, inappropriate secretion of antidiuretic hormone, anorexia, weight loss, appetite loss, nausea, vomiting, diarrhoea, dry mouth, dyspepsia, constipation, headache, restlessness, insomnia, anxiety, dizziness, visual disturbance, drowsiness, confusion, tremor, sweating, sedation. Small increases in diastolic blood pressure and tachycardia as well as bradycardia. Hyperprolactinemia with galactorrhea, hyponatremias. Rare cases of increased ALTs and exceptional cytolytic or mixed hepatitis. **Product authorisation holder:** Generics (UK) Ltd, Station Close, Potters Bar, Herts, EN6 1TL, England. **Product authorisation number:** PA/405/36/1 Available only on prescription. **Date of preparation or last review:** January 2002. For full prescribing information please see the Summary of Product Characteristics. **Further information is available from:** Gerard Laboratories, 2004A Orchard Avenue, CityWest Business Campus, Naas Rd, Dublin 24. **FREEPHONE 1800 272 272.** Fax: 01 466 1912 **Reference:** 1. MIMS January 2002. **GMS REIMBURSABLE 1ST. FEBRUARY 2000.** Code No.: 26232

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EDITORIAL

3 Immigration: new challenges for psychiatry and mental health services in Ireland

Anthony W Clare

ORIGINAL PAPERS

4 Characteristics and treatment of asylum seekers reviewed by psychiatrists in an Irish inner city area

Noel Kennedy, Paula Jerrard-Dunne, Michael Gill, Marcus Webb

8 Infanticide and child murder – admissions to the Central Mental Hospital 1850-2000

Niamh Mulryan, Pat Gibbons, Art O'Connor

BRIEF REPORTS

13 Screening for excessive drinkers in a Hampshire general practice

Alistair James, Guy Edwards

REVIEWS

16 Psychiatric and medical co-morbidity in mitral valve prolapse

Efrain A Gonzalez, Claudia Pimental, Ruby A Natale, Benjamin A Toll, Ariel Soffer, Leonard Gralnik

21 Psychiatric services in Ireland: North and South

Marcus Webb, Roy McClelland, Glenda Mock

HISTORICAL

27 James Joyce and a North Dublin asylum

Aidan Collins

CASE REPORTS

29 Mental illness in migrants: diagnostic and therapeutic challenges

Larkin Feeney, Brendan D Kelly, Peter Whitty, Eadbhard O'Callaghan

32 A delusion of immortality – the dilemma of the Struldbruggs

Rachael Cullivan, Brian A Lawlor

22a John Dunne Medal

31 Guidelines for Authors

35 Letters to the Editor

37 Book Reviews

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Not just emerging from depression...



... but staying on track

Abbreviated Prescribing Information: LUSTRAL™ (sertraline) Presentation: Tablets containing 50mg or 100mg sertraline. **Indications:** Treatment of symptoms of depressive illness, including accompanying symptoms of anxiety. Prevention of relapse or recurrence of depressive episodes, including accompanying symptoms of anxiety. Obsessive compulsive disorder (OCD). Panic disorder, with or without agoraphobia. Post-traumatic stress disorder (PTSD). **Dosage:** Lustral should be given as a single daily dose. The initial dose in depression and OCD is 50mg and the usual antidepressant dose is 50mg. The initial dose in panic disorder and PTSD is 25mg, increasing to 50mg after one week. Dosage can be further increased, if appropriate, to a maximum of 200mg daily. Changes in dose should not be made more frequently than once per week given the 24 hour elimination half life of sertraline. 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 this group of drugs. 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 14 days should elapse before starting any MAOI following discontinuation of Lustral. **Precautions, warnings:** Renal insufficiency, ECT, epilepsy, driving. Lustral should be discontinued in a patient who develops seizures. Lustral should not be administered with benzodiazepines or other tranquilizers in patients who drive or operate machinery. Serotonergic drugs such as tryptophan or fenfluramine should be used with caution. The patient should be monitored for signs of suicide or mania. **Drug Interactions:** Caution with other centrally active medication. Lithium levels should be monitored. 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. Interactions with e.g. warfarin, diazepam, tolbutamide

and cimetidine have not been fully assessed. With warfarin prothrombin time should be monitored when Lustral is initiated or stopped. **Side-Effects:** Dry mouth, nausea, diarrhoea/loose stools, ejaculatory delay, tremor, increased sweating, dizziness, insomnia, somnolence, headache, anorexia and dyspepsia. Rarely, abnormal LFTs, hyponatraemia. The following have been reported with Lustral but may have no causal relationship: vomiting, abdominal pain, movement disorders, convulsions, menstrual irregularities, hyperprolactinaemia, galactorrhoea, rash and alopecia. Rarely, pancreatitis, serious liver events, altered platelet function, abnormal bleeding and purpura. As with other serotonin re-uptake inhibitors rare reports of agitation, confusion, depersonalisation, hallucinations, nervousness, postural hypotension, hypo/hypertension, tachycardia and arrhythmias. Withdrawal reactions have been reported with Lustral. Common symptoms include dizziness, paraesthesia, headache, anxiety and nausea. Abrupt discontinuation of treatment with Lustral should be avoided. The majority of symptoms experienced on withdrawal of Lustral are non-serious and self-limiting. **Legal Category:** S1A. **Package Quantities:** 50mg tablet (PA 822/1/4) Calendar pack of 28; 100mg tablet (PA 822/1/5) Calendar pack of 28. **Product Authorisation Holder:** Pfizer (Ireland) Limited, Parkway House, Ballymount Road Lower, Dublin 12, Republic of Ireland. **Further information on request:** Pfizer (Ireland) Limited. Date last revised: 17 March 1999.



62876

April 2001



LUSTRALTM 50^{MG}

sertraline

Antidepressant therapy that prevents relapse and recurrence

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