

THE BRITISH JOURNAI OF PSYCHIATRY

AUGUST 1999 VOL. 175

Helpline And Information Serv The Royal College Of Psychiati 17 Belgrave Square London SWIX 8PG Tel. 0171 235 2351 Eat. 138/152

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3 Based on 28 days treatment, prizes from NiMS, May 1999; 4, Partis M. et al. Int Clin. Psychophamacol. 1996; 11: 129-136. 5, Stahl SM. Ctafagram vs. sertuline vs. placolog. preliminary efficacy results. Poster presented at the APA metring. 1998; 6, Data on file, Lundock, Limited, to December 1998. 7. Taylor Nolson Scriptcount: precorption audit data. 6 months to March. 1999.

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The British Journal of Psychiatry is published monthly by the Royal College of Psychiatrists (a registered charity, registration number 228636). The BJP publishes original work in all fields of psychiatry. Manuscripts for publication should be sent to the Editor, British Journal of Psychiatry, 17 Belgrave Square, London SWIX 8PG. Queries, letters to the Editor and book reviews may also be sent electronically to sthakor@rcpsych.ac.uk.

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Full instructions to authors are given at the beginning of the January and July issues, and on the Web Site below. Copies are also available from the Journal Office.

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US Mailing Information

The British Journal of Psychiatry is published monthly by the Royal College of Psychiatrists. Subscription price is \$375. Second class postage paid at Rathway, NJ. Postmaster send address corrections to the British Journal of Psychiatry, c/o Mercury Airfreight International Ltd Inc., 2323 Randolph Avenue, Avenel, New Jersey 07001.

THThe paper used in this publication meets the minimum requirements of the American National Standard for Information Sciences – Permanence of Paper for Printed Library Materials, ANSI 23948-1984.

Typeset by Dobbie Typesetting Ltd, Tavistock.

Printed by Henry Ling Ltd. The Dorset Press, 23 High East Street, Dorchester, Dorset DTI IHD.

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Founded by J. C. Bucknill in 1853 as the Asylum Journal and known as the Journal of Mental Science from 1858 to 1963.

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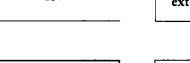
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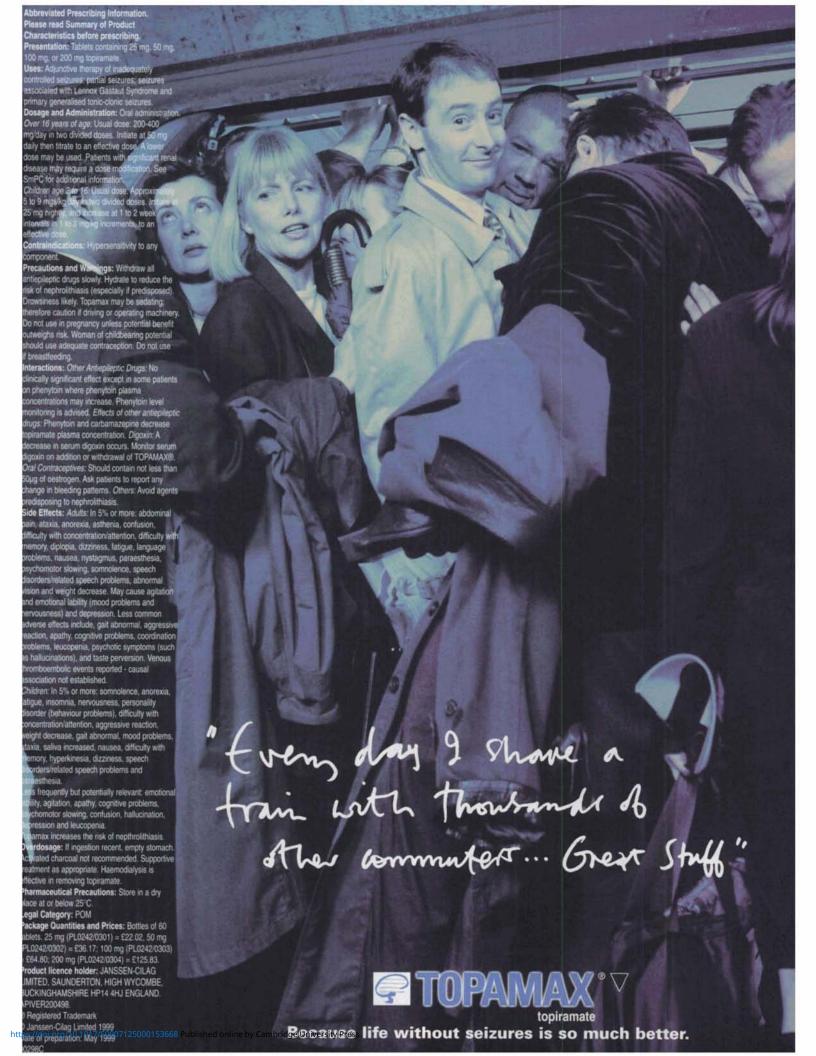


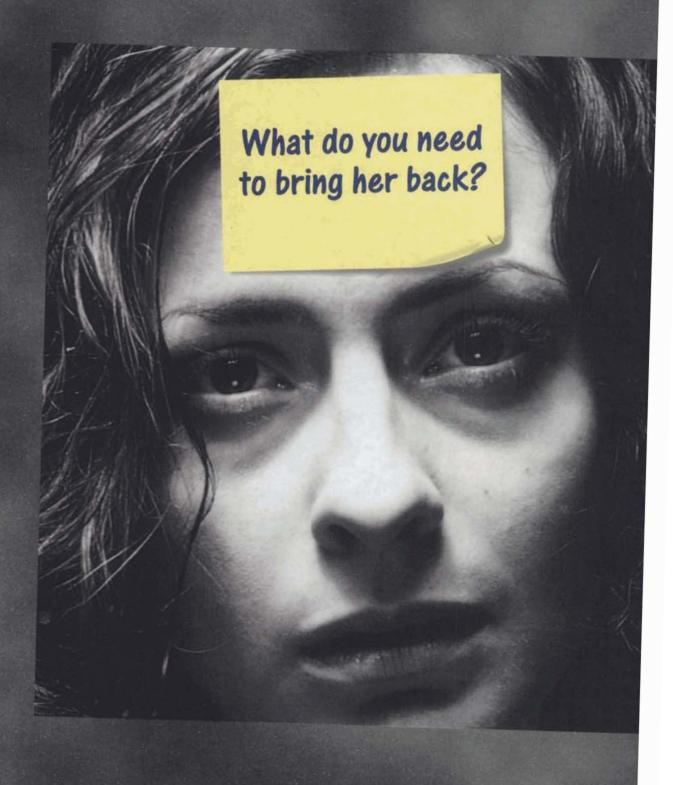
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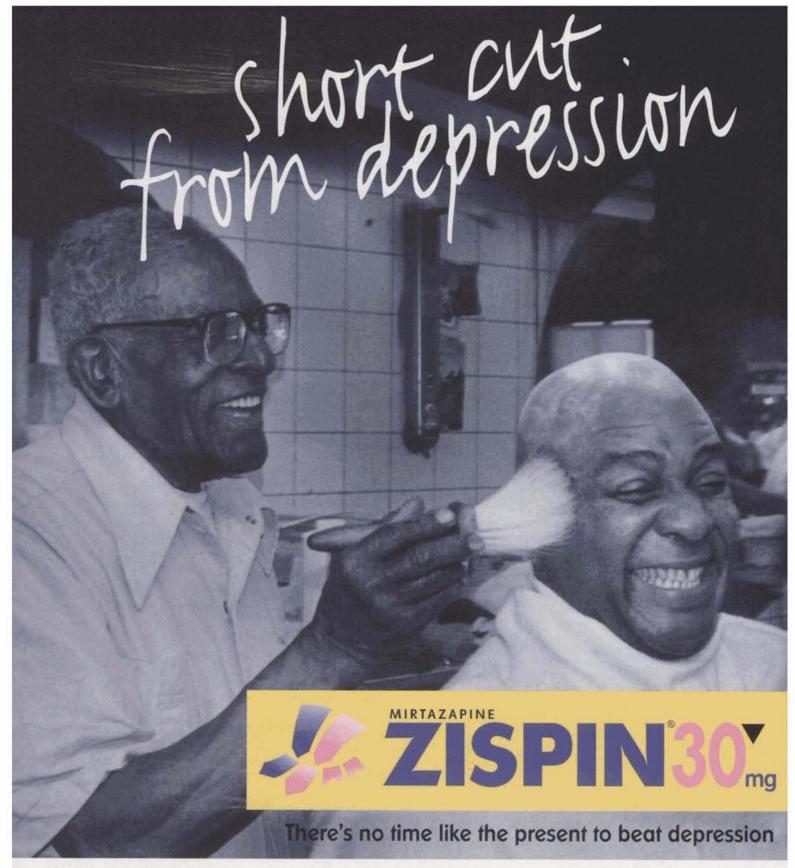


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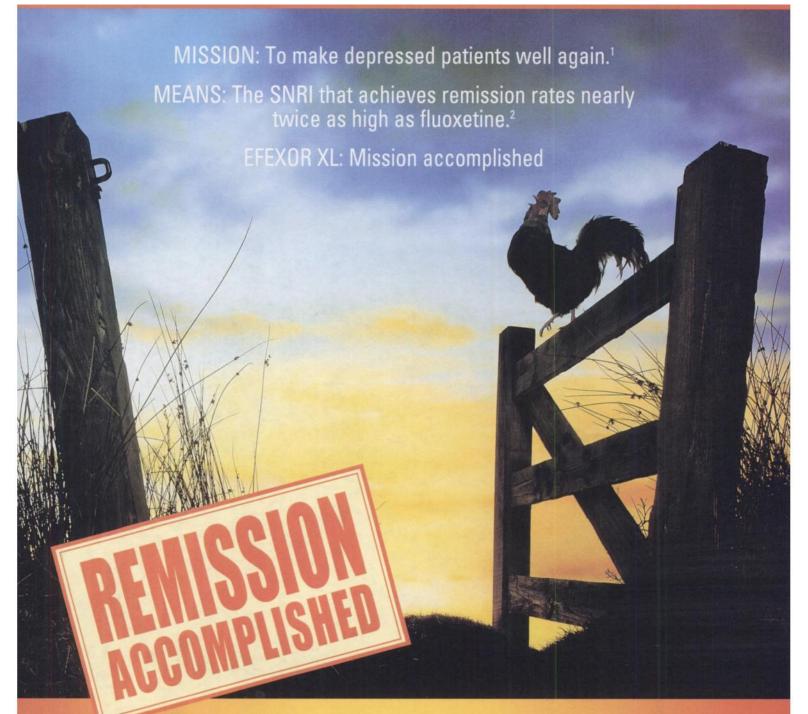
mellitus. Treatment should be discontinued if jaundice occurs. Moreover, as with other antidepressants, the following should be taken into account: worsening of psychotic symptoms can occur when antidepressants are administered to patients with schizophrenia or other psycholic disturbances; when the depressive phase of manicdepressive psychosis is being treated, it can transform into the manic phase. Zispin has sedative properties and may impair concentration and alertness. Interactions: Mirrazapine may potentiate the central nervous dampening action of alcohal; patients should therefore be advised to avoid alcohal during treatment with Zispin; Zispin should not be administered concomitantly with MAO inhibitors or within two weeks of cessation of therapy with these agents; Mirtazapine may potentiate the sedative effects of benzodiazepines; in vitro data suggest that clinically significant interactions are unlikely with mirrozap Pregnancy and lactation: The safety of Zispin in human pregnancy has not been established. Use during pregnancy is not recommended Warner of child bearing potential should employ an adequate method https://doi.org/10.1017/S0007125000153668 Published online by Cambridge University Press in nusing moments and recommended Adverse reactions: The following adverse effects have been reported: Common

levels. Rare (<1/1000): Oedema and accompanying weight gain. Reversible agranulocytosis has been reported as a rare occurrence. (Orthostatic) hypotension. Exanthema. Mania, convulsions, tremot, myoclonus. **Overdosage:** Toxicity studies in animals suggest that clinically relevant cardiotoxic effects will not occur after overdosing with Zispin. Experience in clinical trials and from the market has shown that no serious adverse effects have been associated with Zispin in overdose. Symptoms of acute overdosage are confined to prolonged sedation. Cases of averdose should be freated by gastric lavage with appropriate symptomatic and supportive therapy for vital functions. Marketing authorization number: PL 0065/0145 Legal category: POM Basic NHS cost: £24 for 28 tablets of 30 mg

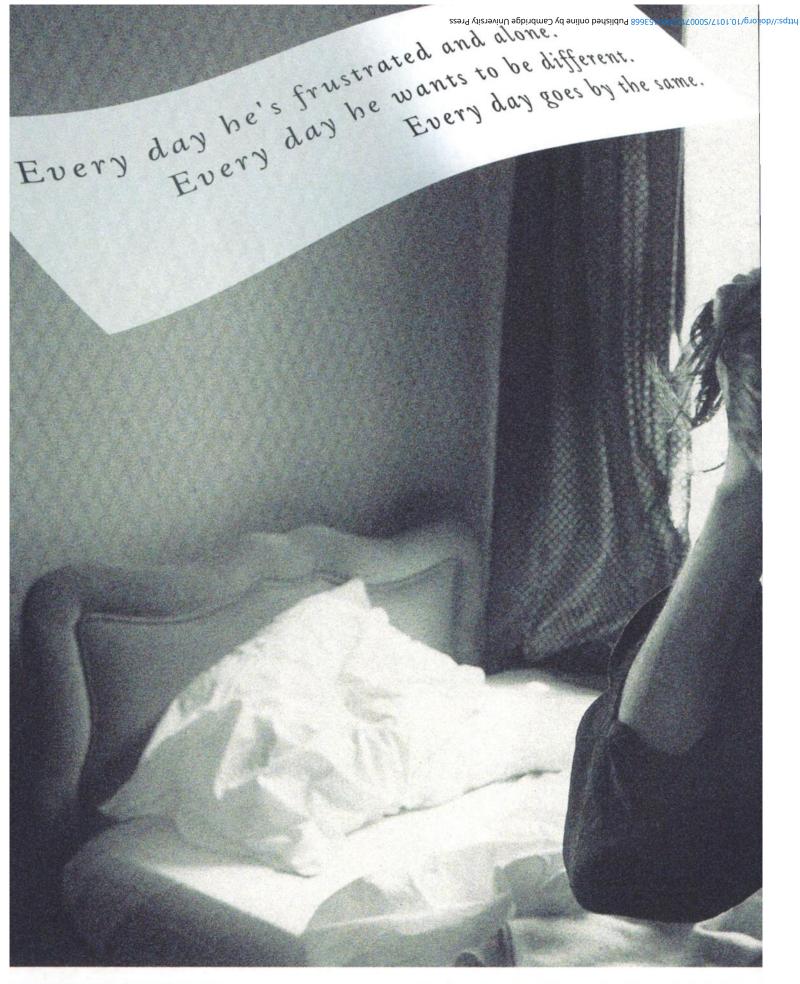
For further information, please contact



Nourypharma



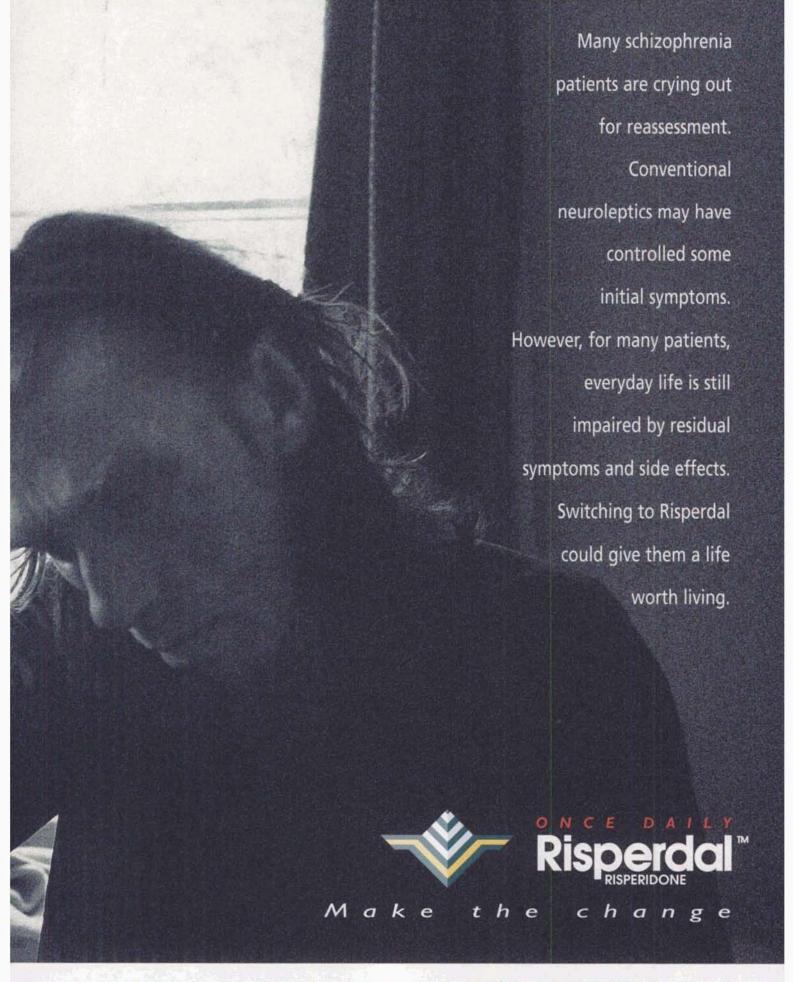
EFE OR XL ENLAFAXINE 75 mg o.d. Simply effective



SPERDAL™ ABBREVIATED PRESCRIBING INFORMATION

ease refer to Summary of Product Characteristics before prescribing Risperdal (risperidone). IES Schizophrenia. Other psychotic conditions, in which positive and/or negative symptoms are ominent. Alleviates affective symptoms of schizophrenia. DOSAGE. Adults: Once or twice daily. All tients, start with 2 mg/day. This may be increased to 4 mg/day on the second day. Some patients ay benefit from slower titration. Then can be maintained unchanged, or individualised, if needed.

by 0.5 mg bd to 1 to 2 mg bd. Well tolerated in elderly. Caution if renal and liver disease. Children: Not recommended. Contra-indications: Hypersensitivity. Precautions: Orthostatic hypotension. Cardiovascular disease. Drugs prolonging QT. Reduce dose if hypotension. If tardive dyskinesia, consider stopping all antipsychotic drugs. Parkinson's disease. Epilepsy. Advise of potential for weight gain. Advise not to drive or operate machinery if mental alertness affected. Pregnancy: Only if benefits outweigh risks. Lactation: Avoid. Interactions: Caution in combination with centrally



and liquid: Store below 30°C. Do not refrigerate. LEGAL CATEGORY POM. PRESENTATIONS, PACK SIZES, PRODUCT LICENCE NUMBERS & BASIC NHS COSTS 1 mg tablets (PL 0242/0186) 20: £13.45, 60: £40.35. 2 mg tablets (PL 0242/0187) 60: £79.56. 3 mg tablets (PL 0242/0188) 60: £117.00.4 mg tablets (PL 0242/0189) 60: £154.44. 6 mg tablets (PL 0242/0317) 28: £109.20.1 mg per ml solution: (PL 0242/0199) 100 ml: £65.00. FURTHER INFORMATION IS AVAILABLE FROM THE PRODUCT LICENCE HOLDER: Janssen-Cilag Ltd, Saunderton, High Wycombe, Buckinghamshire HP14 4HJ. APIVER200599

-- Life beyond Alzheimer's.



With Exelon, you can now help treat the symptoms of people with mild to moderately severe Alzheimer's disease.

While Exelon has not been shown to affect the disease process, six-month trials have established its effectiveness on key areas that Alzheimer's disease attacks - cognition, global functioning and activities of daily living.1

For carers and family, this could mean some relief from the demands for attention; for the sufferer, it could mean life beyond Alzheimer's.



Beyond cognition: improving functional ability.

EXELON Prescribing Information. Indications: Symptomatic freatment of mild to moderately severe Alzheimer's demential. Presentation: Capsules containing 1.5, 3, 4.5 or 6mg invastigmine. Dosage and Administration: Effective dose is 3 to 6mg hylice a day. Maintain patients on their highest well-tolerated dose. Maximum dose 6mg hylice adaly. Reassess patients regularly, Initial dose 1.5mg hylice daily, then build up dose, at a minimum of two week intervals, to 3mg hylice doily, 4.5mg hylice doily then 6mg hylice doily, if tolerated well. If adverse effects or weight decrease occur, these may respond to omitting one or more doses. If pensistent, doily dose should be temporarily reduced to previous well tolerated dose. Controllationations: Known hypersensitivity to invastigmine or excipients or any other carbamates derivatives; severe liver imporment. Special Warning & Precautions: Therapy should be initiated and supervised by a physician experienced in the diagnosis and treatment of Alzheimer's disease. A caregiver should be available to monitor compliance. There is no experience of use of EXELON in other types of demention/memory impairment. Nousea and vomiting may occur, particularly when initiating and/or increasing dose. Monitor any weight loss. Use with care in patients with Sick Shus Syndrome, conduction defects, active gostific or duodend ulcers, or those predisposed to ulcerative conditions, history of asthma or obstructive pulmonary disease, those predisposed to urinary obstruction and seizures. In renal and mild to moderate hepatic impairment, titrate dose individually. Safety in pregnancy not established; women should not breastfeed. Use in children not recommended. Interactions: May exaggerate effects of succinylcholine-type muscle relaxants during anaesthesia. Do not give with cholinomimetic drugs. May interfere with anticholinergic medications. No interactions were observed with digoxin, warfarin, diazepam, or fluoxetine (in healthy volunteers). Metabolic drug interactions unlikely, although tit may inhibit butrylcholinesterase mediated metabolism of other drugs. Undestrable Effects: Most commonous

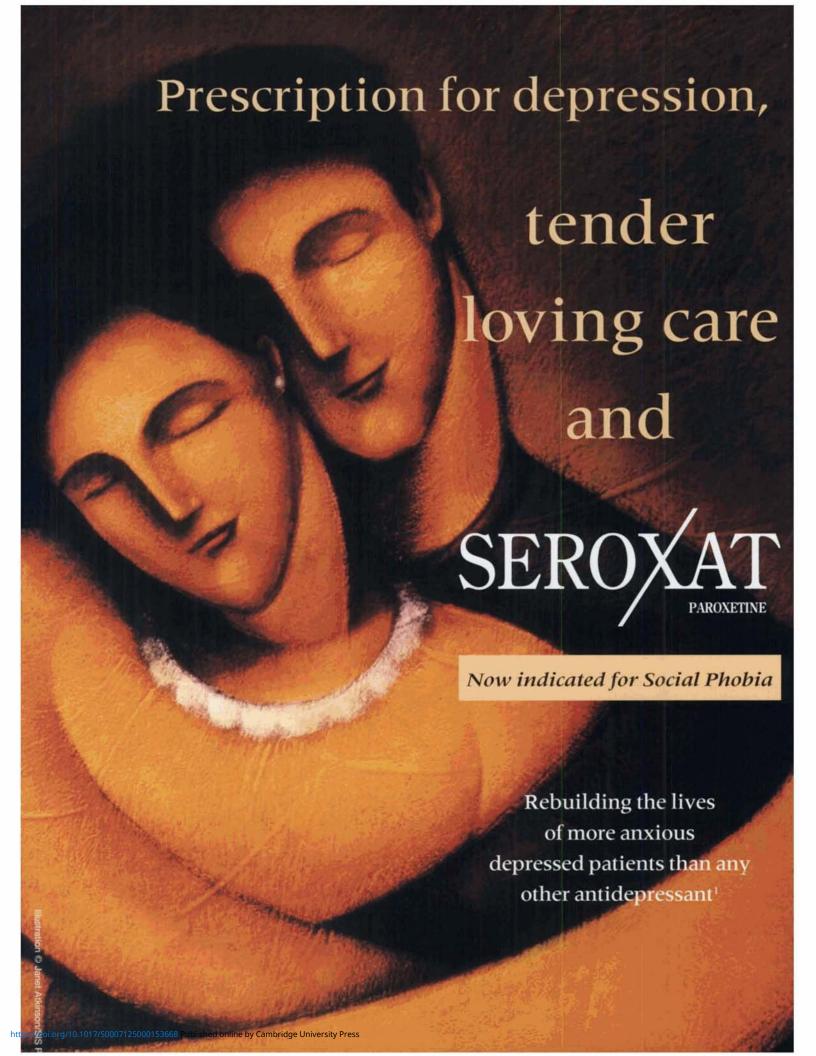
common effects (25% and ≥ placebo): abdominal pain, accidental trauma, agitation, confusion, depression, diarrhoea, dyspepsia, headache, insomnia, upper respiratory tract and urinary tract infections, increased sweating, malaise, weight loss, tremor, Rarely, angina pectoris. carriory inter interior is increased sweating, matures, weight loss items. Retailed to the gostrointestinal hoemorrhage and syncope. No notable abnormalities in laboratory values observed. Package Quantities and basic NHS Price: 1.5mg x 28, 531.50; 1.5mg x 56, 563.00; 3mg x 28, 531.50; 3mg x 56, 263.00; 4.5mg x 28, 531.50; 4.5mg x 56, 263.00; 6.7mg x 56, 263.00; 6 3mg, EU/1/98/066/004 - 5: 4.5mg, EU/1/98/066/007 - 8: 6mg, EU/1/98/066/010 - 11. Full prescribing information including Summary of Product Characteristics is available from: Novartis Pharmaceuticals UK Ltd., Frimley Business Park, Frimley, Camberley, Surrey, GU16 5SG.

Reference: 1. Corey-Bloom J, et al. International Journal of Geriatric Pyschopharmacology 1998;

Date of preparation: May 1999

Code No. EXE 99/20





PRESCRIBING INFORMATION

Prescribing information

Presentation: 'Seroxat' Tablets, PL 10592/0001-2, each containing either 20 or 30 mg paroxetine as the hydrochloride. 30 (OP) 20 mg tablets, £20.77; 30 (OP) 30 mg tablets, £31.16.

'Seroxat' Liquid, PL 10592/0092, containing 20 mg paroxetine as the hydrochloride per 10 ml. 150 ml (OP), £20.77.

Indications: Treatment of symptoms of depressive illness of all types including depression accompanied by anxiety. Following satisfactory response, continuation is effective in preventing relapse. Treatment of symptoms and prevention of relapse of obsessive compulsive disorder (OCD). Treatment of symptoms and prevention of relapse of panic disorder with or without agoraphobia. Treatment of symptoms of social anxiety disorder/social phobia.

Dosage: Adults: Depression: 20 mg a day. Review response within two to three weeks and if necessary increase dose in 10 mg increments to a maximum of 50 mg according to response.

Obsessive compulsive disorder: 40 mg a day. Patients should be given 20 mg a day initially and the dose increased weekly in 10 mg increments. Some patients may benefit from a maximum dose of 60 mg a day.

Panic disorder: 40 mg a day. Patients should be given 10 mg a day initially and the dose increased weekly in 10 mg increments. Some patients may benefit from a maximum dose of 50 mg a day. Social anxiety disorder/social phobia: 20 mg a day. Patients should start on 20 mg and if no improvement after at least two weeks they may benefit from weekly 10 mg dose increases up to a maximum of 50 mg/day according to response. 'Seroxat' has been shown to be effective in 12 week placebo-controlled trials. There is only limited evidence of efficacy after 12 weeks' treatment.

Give orally once a day in the morning with food. The tablets should not be chewed. Continue treatment for a sufficient period, which should be at least four to six months after recovery for depression and may be longer for OCD and panic disorder. As with many psychoactive medications abrupt discontinuation should be avoided – see **Adverse reactions**.

Elderly: Dosing should commence at the adult starting dose and may be increased in weekly 10 mg increments up to a maximum of 40 mg a day according to response.

Children: Not recommended.

Severe renal impairment (creatinine clearance <30 ml/min) or severe hepatic impairment: 20 mg a day. Restrict incremental dosage if required to lower end of range.

Contra-indication: Hypersensitivity to paroxetine.

Precautions: History of mania. Cardiac conditions: caution. Caution in patients with epilepsy; stop treatment if seizures develop. Driving and operating machinery.

Drug interactions: Do not use with or within two weeks after MAO inhibitors; leave a two-week gap before starting MAO inhibitor treatment. Possibility of interaction with tryptophan. Great caution with warfarin and other oral anticoagulants. Use lower doses if given with drug metabolising enzyme inhibitors; adjust dosage if necessary with drug metabolising enzyme inducers. Alcohol is not advised. Use lithium with caution and monitor lithium levels. Increased adverse effects with phenytoin; similar possibility with other anticonvulsants.

Pregnancy and lactation: Use only if potential benefit outweighs possible risk.

Adverse reactions: In controlled trials most commonly nausea, somnolence, sweating, tremor, asthenia, dry mouth, insomnia, sexual dysfunction (including impotence and ejaculation disorders), dizziness, constipation and decreased appetite.

Also spontaneous reports of dizziness, vomiting, diarrhoea, restlessness, hallucinations, hypomania, rash including urticaria with pruritus or angioedema, and symptoms suggestive of postural hypotension. Extrapyramidal reactions reported infrequently; usually reversible abnormalities of liver function tests and hyponatraemia described rarely. Symptoms including dizziness, sensory disturbance, anxiety, sleep disturbances, agitation, tremor, nausea, sweating and confusion have been reported following abrupt discontinuation of 'Seroxat'. It is recommended that when antidepressant treatment is no longer required, gradual discontinuation by dose-tapering or alternate day dosing be considered.

Overdosage: Margin of safety from available data is wide. Symptoms include nausea, vomiting, tremor, dilated pupils, dry mouth, irritability, sweating and somnolence. No specific antidote. General treatment as for overdosage with any antidepressant. Early use of activated charcoal suggested.

Legal category: POM. 10.9.98



Welwyn Garden City, Hertfordshire AL7 1EY.

'Seroxat' is a trade mark.

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Reference: 1. Data on file.

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