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A new option in the management of Non-small Cell Lung Cancer



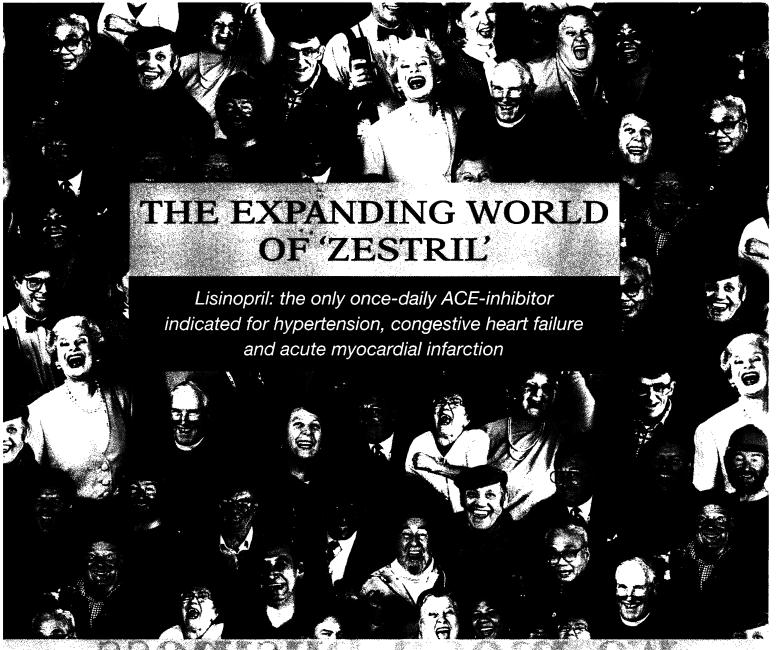
For details of a patient centred approach to the management of NSCLC contact Lilly Oncology on 01256 315461

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Abbreviated Prescribing Information: Gemzar - (Gemcitabine Inn) Presentations Glass vials containing gemcitabine hydrochloride, equivalent to 200mg or 1g of gemcitabine, as a sterile lyophilised powder. Uses Gemzar is indicated for the palliative treatment of adult patients with locally advanced or metastatic nonsmall cell lung cancer. Dosage and Administration (For full information, see data sheet or package inserts.) The drug is only administered intravenously and not more than once per week. The recommended dose is 1000mg/m² given by 30 minute intravenous infusion. This dose should be repeated once weekly for three weeks, followed by a one week rest period. This four week cycle is then repeated. Gemcitabine can be administered on an outpatient basis. Dosage reduction is based upon the amount of toxicity experienced by the patient. Patients receiving gemcitabine should be monitored every other week for platelet, leucocyte and granulocyte counts and, if necessary, the dose may be either reduced or withheld in the presence of haematological toxicity (see data sheet). The elderly: As for adults. Gemcitabine has been well tolerated by patients over the age of 65 years. Hepatic and renal impairment: Gemcitabine should be used with caution in patients with hepatic insufficiency or with impaired renal function. Periodic checks of liver and kidney functions, including transaminases and serum creatinine, should be performed in patients receiving gemcitabine. Contra-indications, Warnings, etc. Contra-indications: Known hypersenstivity to gemcitabine. Warnings: Prolongation of the infusion time and increased dosing frequency have been shown to increase toxicity. Myelosuppression is short-lived and usually does not result in dose reductions and rarely in discontinuation. There have been very rare cases of haemolytic uraemic syndrome (signs of microangiopathic haemolytic anaemia such as rapidly falling haemoglobin with concomitant thrombocytopenia, elevation of serum bilirubin or LDH). If this occurs, discontinue gemcitabine immediately. Precautions Usage in pregnancy or lactation. Avoid in pregnant or breast-feeding women, since safety has not been established. Experimental animal studies have shown reproductive toxicity. Patients should be cautioned against driving or operating machinery until they are sure they do not become somnolent. Adverse reactions. Haematological: Anaemia, leucopenia and thrombocytopenia. Thrombocythaemia. Gastro-intestinal: Abnormalities of liver transaminases, which were usually mild, non-progressive and rarely necessitated stopping treatment. Nausea or vomiting, which required therapy in about 20% of patients and was rarely dose-limiting. Renal: Mild proteinuria and haematuria, which was rarely clinically significant. A few cases of renal failure of uncertain aetiology have been reported. Allergic: Rash and pruritus were usually mild and not dose-limiting. Desquamation, vesiculation and ulceration have been reported rarely. Anaphylaxis has been reported rarely. Dyspnoea. Bronchospasm occurred in less than 1% of patients and was usually mild and transient but parenteral therapy may be required. Other: Influenza-like symptoms (fever, headache, back pain, chill, myalgia, asthenia and anorexia). Cough, rhinitis, malaise, sweating and insomnia. Oedema or peripheral oedema. A few cases of hypotension have been reported. Alopecia (usually minimal hair loss), somnolence, diarrhoea, oral toxicity (mainly soreness and erythema) and constipation. Rarely, injection site reaction. Legal Category POM Product Licence Numbers Vials 200mg: 0006/0301, Vials 1g: 0006/0302 Direct Hospital Price 200mg: £18.75 ex.VAT.; 1g: £93.75 ex.VAT. Full Prescribing Information is Available From Eli Lilly and Company Limited, Dextra Court, Chapel Hill, Basingstoke, Hampshire, RG21 5SY. Telephone: Basingstoke (01256) 315000 'GEMZAR' is a Lilly trade mark. Date of preparation October 1995.

References:

 Anderson H., Betticher D.C., et al Symptomatic Benefit from Gemcitabine and Other Chemotherapy In Advanced Non-Small Cell Lung Cancer; Changes in Performance Status and Tumour Related Symptoms. Anti-Cancer Drugs 1995; Vol 6(Suppl. 6):39-48 2. Anderson H., Thatcher N., et al Quality of Life and Symptomatic Benefit in patients with NSCLC. Lilly Oncology Global Medical Conference.22 - 35



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PRESENTATION: Tablets containing 2.5mg, 5mg, 10mg or 20mg lisinopril.

DOSAGE AND ADMINISTRATION: Hypertension Adults (inc elderly): initially 2.5mg daily, a 2.5mg dose seldom achieves a therapeutic response; adjust dose according to response. Maintenance usually 10-20mg once daily. Maximum is 40mg daily.

Diuretic-treated patients - if possible stop diuretic 2-3 days before starting 'Zestril'. Resume diuretic later if desired.

Congestive heart failure Adults: initially 2.5mg daily under close medical supervision (hospital initiation for severe or unstable heart failure and other patients at higher risk), increasing to 5-20mg once daily according to response. Monitor blood pressure and renal function.

Acute myocardial infarction Treatment may be started within 24 hours of symptoms. First dose is 5mg, followed by 5mg after 24 hours, 10mg after 48 hours and then 10mg once daily. Dosing should continue for six weeks. Lower dosage in patients with low systolic blood pressure (120mmHg or less) - see Data Sheet.

Renal impairment - may require lower maintenance dosage. 'Zestril' is dialysable.

Children - not recommended.

CONTRA-INDICATIONS: Pregnancy. Hypersensitivity to 'Zestril'. Patients with history of angioneurotic oedema to previous ACE-inhibitor therapy. Patients with aortic stenosis, cor pulmonale or outflow tract obstruction.

PRECAUTIONS: Assessment of renal function is recommended. Symptomatic hypotension may occur, particularly in volume depleted patients and congestive heart failure. Caution in patients with ischaemic heart or cerebrovascular disease; renal insufficiency; renovascular hypertension. Patients with a history of angioedema may be at increased risk of angioedema with an ACE inhibitor. Acute myocardial infarction patients with evidence of renal dysfunction or at risk of serious haemodynamic deterioration - see Data Sheet. Cough has been reported with ACE inhibitors. Renal impairment (usually reversible) may occur in some patients. Hypotension may occur during surgery or anaesthesia. Caution in nursing mothers. No paediatric experience. Afro-Caribbean patients may show reduced therapeutic response. Symptomatic hypotension can be minimised by discontinuing diuretic prior to 'Zestril'. Interaction with indomethacin and lithium. Potassium supplements, potassium sparing diuretics and potassium containing salt substitutes not recommended. Avoid concomitant use with high-flux dialysis membranes.

SIDE EFFECTS: Hypotension, dizziness, headache, diarrhoea, cough, nausea, fatigue. Less frequently, rash, asthenia. Rarely, angioneurotic oedema and other hypersensitivity reactions, myocardial infarction or cerebrovascular accident possibly secondary to excessive hypotension in high risk patients, palpitations, tachycardia, abdominal pain, dry mouth, pancreatitis, hepatitis, jaundice, mood alterations, mental confusion, paraesthesia, bronchospasm, alopecia, urticaria, diaphoresis, pruritus, uraemia, oliguria/anuria, renal dysfunction, acute renal failure, impotence, haemolytic anaemia. A symptom complex which may include fever, vasculitis, myalgia, arthralgia/arthritis, positive ANA, elevated ESR, eosinophilia, leukocytosis; rash, photosensitivity or other dermatological manifestations may occur. Increases (usually reversible) in blood urea, serum creatinine, liver enzymes and serum bilirubin. Decreases in haemoglobin and haematocrit. Hyperkalaemia and hyponatraemia.

Anaphylactoid reactions during desensitisation treatment. Leucopenia and thrombocytopenia have occurred (causal relationship not established). LEGAL CATEGORY: POM

PRODUCT LICENCE NUMBERS AND BASIC NHS COSTS: 'Zestril' 2.5mg (12619/0084) 28 tablets £7.64; 5mg (12619/0085) 28 tablets, £9.58; 10mg (12619/0086) 28 tablets, £11.83; 20mg (12619/0087) 28 tablets, £13.38. 'Zestril' is a trademark, the property of ZENECA Limited.

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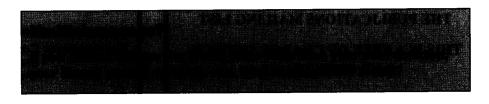
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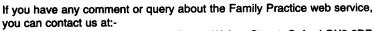
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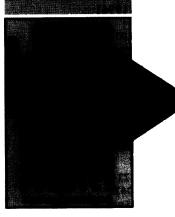
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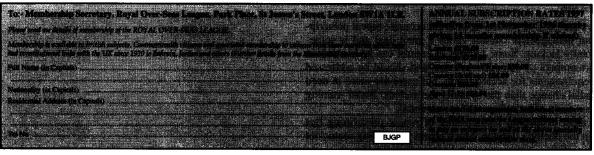
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Thursday, 28th November 1996

to be held at The Royal Pharmaceutical Society of Great Britain, London

'Primary/Secondary Care prescribing groups — experience so far and the way forward'

AGENDA

Commences: 09.30 am

The Public Health perspective of primary/ secondary care prescribing groups

Dr Martin Duerden, Public Health Registrar, Cambridge & Huntingdon Health Authority

Evidence-based medicine — values explored

Dr Andrew Moore, Editor - Bandolier Oxford Health Authority

Prescribing issues for hospitals which are driven by the interface

Mr Howard Rowe, Pharmacy Manager, East Glamorgan NHS Trust, East Glamorgan General Hospital

Management of Benign Prostatic Hyperplasia — a Public Health Perspective

Professor Peter Boyle, Director, Division of Epidemiology & Biostatistics, European Institute of Oncology, Milan

Future issues in prescribing

Mr Ian Carruthers, Chief Executive, Dorset Health Authority

The GP's perspective of the hospital/community prescribing debate

Dr David Phillips, General Practitioner and Medical Adviser, Gwent Health Authority

What opportunities exist for joint prescribing groups

Mr Norman Evans, Pharmaceutical Adviser, Merton, Sutton & Wandsworth Health Authority

What I expect from my joint prescribing group

Mr Graham Butland, Chief Executive, South Essex Health Authority

Concludes: 16.50 pm

Ticket price £30.00 To include VAT, coffee, lunch and post seminar refreshments

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A two day workshop to help participants improve their ability to facilitate the learning of communication skills. Open to any health care professional who is involved in teaching interviewing, assessment or counselling skills

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For further information contact:

Yvonne Hopkins, CRC Training Development Officer, CRC Psychological Medicine Group, Stanley House, Christie Hospital NHS Trust, Wilmslow Road, Manchester M20 4BX.

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Reactions: Intermenstrual bleeding, increase in the size of the uterine fibromyomata, endometrial proliferation, excessive production of cervical mucus, aggravation of endometriosis, premenstrual-like syndrome. Breast tenderness, pain, enlargement, secretion. Nausea, vomiting, cholelithiasis, cholestatic jaundice. Thrombosis, rise of blood pressure. Chloasma, erythema nodosum, rash. Discomfort of the cornea if contact lenses are used. Headache, migraine, mood changes, sodium and water retention, reduced glucose tolerance, a change in body weight. Changes in liver function. Interactions: May diminish glucose tolerance.

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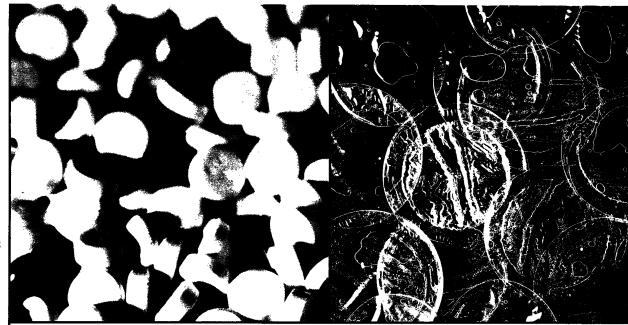
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Wednesday 11th December 1996
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Friday 22nd November 1996

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Thursday 23rd January 1997 Handling People Problems

Wednesday 29th January 1997

Report Writing

Thursday 12th

and Friday 13th December 1996

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Contact Dr. J. Scott-Jones, PO Box 360, Opotiki, New Zealand. Or E-mail jdpsj@wave.co.nz".

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Topics include Microbiological Health and Safety in the GP Surgery; Management of Infection; Communicable Disease Control in the Community

Wednesday 16 April 1997

Venue: National Motorcycle Museum, Solihull

For further details contact:
RCGP Conference & Courses Unit,
Royal College of General Practitioners, 14 Princes Gate,
Hyde Park, LondonSW7 1PU

Tel: 0171 823 9703 Fax: 0171 225 3047

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DEPRESSION AND PHYSICAL ILLNESS

A one-day conference on Friday 22 November 1996 organised by The Royal College of Psychiatrists

and

THE ROYAL SOCIETY OF MEDICINE

The first in a series of meetings on the theme of 'Psychiatry for non-Psychiatrists'. Particular aims are to highlight psychiatric problems arising in clinical practice with a special accent on those that occur as a result of organic illness. The first of this series addresses the general topic of depression and physical illness. The meeting has 5 CME credits.

Full programme details from:

Ms Alison Hamlett, Academic Conference Department The Royal Society of Medicine, 1 Wimpole Street, London W1M 8AE. Fax: 0171 290 2977.

Study Day on Sport & Health: Fitness for Children & Teenagers Friday 29 November 1996

Venue: RCGP, 14 Princes Gate, London SW7

The RCGP, in association with the Sports Council, is organising a Study Day which will look at the effects of exercise on the health of young people. Current research in this area will be discussed, and topics for discussion will include asthma management, weight control and mental health.

The Delegate Fee (inclusive of VAT) is £60.00. PGEA Applied for.

Study Day on HIV & AIDS

Friday 13 December 1996

The RCGP Working Party on HIV/AIDS is organising a Study Day which will address two important areas for health professionals - Home Care of Patients with HIV/AIDS and Issues around HIV and pregnancy.

The Delegate Fee (inclusive of VAT) is £30.00. PGEA Applied for.

For further details of any of the above please contact: RCGP Courses, 14 Princes Gate, Hyde Park, London SW7 1PU. Tel: 0171 823 9703. Fax: 0171 225 3047. Email: Courses@RCGP.Org.UK



PANG

Pain and Nociception Group

Forthcoming Symposia for 1996 and 1997

Charing Cross & Westminster Medical School Margravine Road Fulham, London W6 8RP

Monday 2 December 1996

The economic consequences of pain

Monday 10 March 1997

Regional anaesthesia for surgery

Monday 12 May to Wednesday 14 May 1997

London chronic pain symposium

Friday 17 October 1997

Current controversies in the management

of cancer pain

Organisers: Dr

Dr P J D Evans & Dr D M Justins

Further information: Miss C O'Neil

Department of Anaesthesia Charing Cross Hospital

London W6 8RF

Tel: 0181 846 7017

Fax: 0181 846 7585

(Approved for CME) (Concessionary rates available)

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To ensure delivery of publications, gifts and record cards before Christmas all orders are to be received by the Sales Office at Princes Gate by Friday 13 December.

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Email: sales@rcgp.org.uk

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Royal College of General Practitioners 1996 Christmas Lecture for 15-18 Year Olds Thursday 12 December 1996

"Very General Practitioners: Primary and Secondary Care in Rural Africa"

PRESENTED BY:
Drs Richard and Carolyn Rigby MRCGP

Further details please contact:
The Royal College of General Practitioners, Courses Unit,
14 Princes Gate, Hyde Park, London SW7 1PU.

Tel: 0171 823 9703 or Fax: 0171 225 3047 Email address: courses@rcgp.org.uk

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morale and morality 4-6th APRIL 1997

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FOR FURTHER INFORMATION CONTACT:

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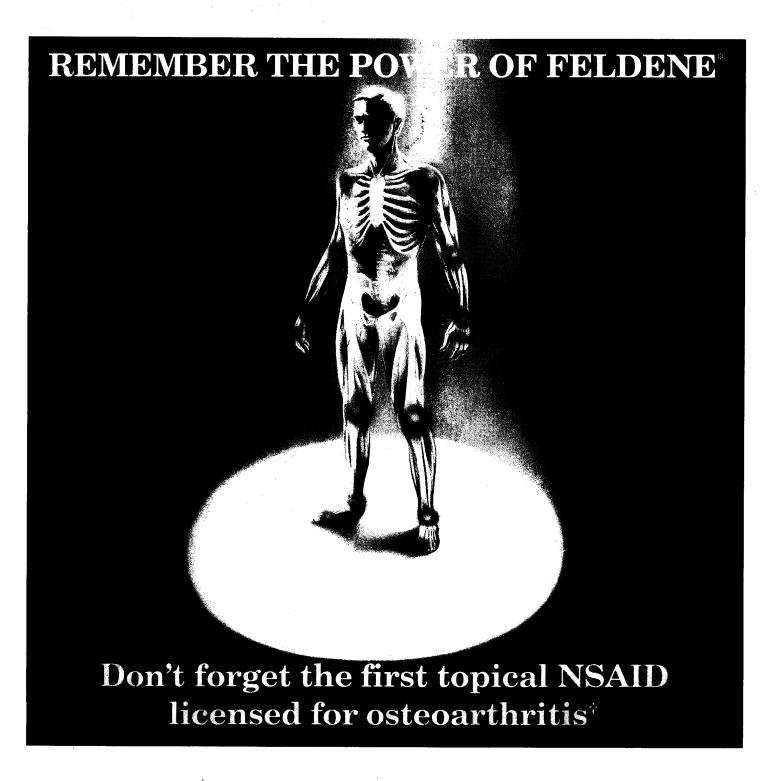
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dermatoses or infection. Use in children: Not recommended. Use in pregnancy and lactation: The safety of FELDENE Gel during pregnancy and lactation has not been established. Side- Effects: Mild or moderate local reactions at the application site. Systemic reactions occur infrequently including nausea and dyspepsia; rare reports of abdominal pain and gastritis and isolated reports of bronchospasm and dyspencea. Legal Category: POM. Package

Quantities and Basic NHS Cost: FELDENE Gel 112g tube £7.84 (PL 0057/0284). Further information on request. Pfizer Limited. Sandwich. Kent. FELDENE Gel is the first topical NSAID in the UK with a licensed indication for osteoarthritis.





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