lisinopril ICI

Prescribing Notes

Consult data sheet before

prescribing.

Use: All graces of essential hypertension and renovascular hypertension. Congestive heart failure, adjunctive therapy

Presentation: Tablets containing 2.5mg, 5mg, 10mg or 20mg lisinopri ('Zestril'i.

Dosage and administration: Hypertension - initially 2 5mg dairy a 2 5mg dose seldom actileves a therapeutic response; adjust dose according to response. Maintenance usually 10-20mg once daily, Maximum is 40mg dazy. Duretic-treated patients - f possible stop gruretic 2-3 days before starting 'Zestru' Pesume diuretic later if desired

Congestive heart failure:
(adjunctive therapy) - initially
2.5mg daily in hospital under
close medical supervision,
increasing to 5-20mg once daily according to response

Impaired renal function: May require a lower maintenance dose. Zestni is dialysable.

Elderly patients: No change Contraindications: Pregnancy - stop therapy if sus-pected. Hypersensitivity to Zestni'. Patients reacting with angioneurotic dedema to previous ACE inhibitor treatment

vous ACE inhibitor treatment

Precautions: Assessment of
renal function is recommended.

Rena insufficiency renovascular hypertension: surgery,
anaesthesia: possibility of anaesthesia: possibility of hypotension especially in ischaemic heart disease and cereprovascular disease Compi nation with antihypertensives may increase hypotensive effect Sometimes increased blood urea and creatinine and/or cases of renai insufficiency if given with diuretics. Minimises thiazide-induced hypokalaema and hyperur caemia Potassium sup-piements potassium-sparing diuretics and potassium-containing sait substitutes not recommended. Indomethacin may reduce hypotensive effect. Afro-Carbbean patients. Use with authow tract obstruction or corpulmonale Monitor ithium serum leveis f lithium saits administered. Side effects: Dizziness, head ache, diarrhoea, fatigue, cough and hausea Less frequently, rash and asthenia. Rarely, angioneurotic dedema and other hypersensitivity reactions, renal falure symptomatic hypotension (especially if volume-depleted) severe hypotension (more likely if severe heart failure; calpitation; hyperkalaema; ncreases in liver enzymes and serum broubin (usually reversible on discontinuation of Zestril') and impotence.

Product licence numbers and basic NHS cost: 'Zestril' 2.5mg :29:0208) 28 tablets £7.84 5mg :29/0204; 28 tab ets. £9.83, 10mg :29/0205; 28 tablets. £12.13, 20mg :29/0206; 28 tablets, £20.96.

'Zestril' is a trademark

Further information is available from C: Pharmaceuticals (UK), Southoank. A deney Park, Vaccleskeid, Cheshire SK10 4TF.





Everyone has a certain zest for life. 'Zestril' lets your hypertensive patients enjoy theirs. 'Zestril' is an effective ACF innibitor from ICI. 'Zestril' is taken once daily, so encouraging good patient compliance. A valuable aspect of 'Zestril' therapy is that it is well

tolerated, and suitable for a wide range of hypertensive patients. Your hypertensive patients wat appreciate the convenient dosage and good tolerability of 'Zestril'. It's a therapy they'll find easy to ive with one which helps retain their zest for life.

Morning, noon and night:

Oh what a beautiful morning!



lisinopril ICI Once daily for 24 hour BP control

Antihypertensive therapy which helps retain that zest for life.

BMJ 1840-1990

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ABRIDGED PRODUCT INFORMATION Refer to Data Sheet before prescribing. INDICATIONS

All grades of essential hypertension and renovascular hypertension. Congestive heart failure (adjunctive therapy).

DOSAGE AND ADMINISTRATION

Hypertension: Initially, 5 mg once daily, reduce starting dose to 2.5 mg if over 65 years, on diuretics or renally impaired. Adjust dose according to response; maintenance usually 10-20 mg once daily. Maximum dose 40 mg daily.

Diuretic-treated patients – if possible stop diuretic two to three days before starting 'Innovace'. Resume diuretic later if required.

Congestive heart failure (adjunctive therapy): Initially, 2.5 mg daily in hospital under medical supervision, increasing to the usual maintenance dose of 10-20 mg once daily according to response.

Impaired renal function: May require a lower maintenance dose. 'Innovace' is dialysable.

CONTRA-INDICATIONS

Pregnancy – stop therapy if suspected. Hypersensitivity to 'Innovace'. Patients reacting with angioneurotic oedema to previous ACE-inhibitor treatment.

PRECAUTIONS

Assess renal function prior to therapy with 'Innovace' and during therapy where appropriate.

Renal insufficiency; possibility of hypotension especially in ischaemic heart disease or cerebrovascular disease or in volume-depleted patients; surgery/anaesthesia.

Combination with antihypertensives may increase hypotensive effect. In some patients with bilateral renal artery stenosis increased blood urea and creatinine has been seen, especially in patients treated with diuretics and/or those with renal insufficiency. Minimises thiazide-induced hypokalaemia and hyperuricaemia. Potassium supplements, potassium-sparing diuretics, and potassium-containing salt substitutes are not recommended.

Possible reduced response in Afro-Caribbean patients. Use with caution in breast-feeding mothers. Do not use in aortic stenosis, or outflow tract obstruction. Monitor serum levels of lithium, if lithium salts are given.

SIDE EFFECTS

Side effects include: dizziness, headache. Others include fatigue, asthenia, nausea, diarrhoea, muscle cramps, rash, cough.

Less commonly, angioneurotic oedema; other hypersensitivity reactions; renal failure; symptomatic hypotension (especially if volume-depleted); severe hypotension (more likely if severe heart failure); hyperkalaemia, hyponatraemia; increases in liver enzymes and serum bilirubin (usually reversible on discontinuation of 'Innovace'); paraesthesiae; impotence.

BASIC NHS COST

2.5 mg tablets, £10.00 for bottles of 50. 5 mg tablets, £7.86 for 28-day calendar pack; £14.03 for bottles of 50. 10 mg tablets, £11.03 for 28-day calendar pack; £19.69 for bottles of 50. 20 mg tablets, £13.10 for 28-day calendar pack; £23.40 for bottles of 50.

Product licence numbers:

 $2.5~\rm mg$ tablets, 0025/0120; 5 mg tablets, 0025/0194; 10 mg tablets, 0025/0195; 20 mg tablets, 0025/0196.

Issued February 1991.

8 denotes registered trademark of Merck & Co., Inc., Rahway, NJ, USA

References

- Kjekshus, J., and Swedberg, K., Amer. J. Cardiol., 1989, 63, 26D.
- Swedberg, K., and Kjekshus, J., Drugs, 1990, 39 (Suppl. 4), 49.
- CONSENSUS Trial Study Group, New Engl. J. Med., 1987, 316, 1429.



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