NAME OF THE MEDICINE

The active ingredient in Catapres is clonidine hydrochloride.

Clonidine hydrochloride has the following structural formula:

![Structural formula of clonidine hydrochloride](image)

DESCRIPTION

The chemical name for clonidine hydrochloride is 2,6-dichloro-N-2-imidazolidinylidene-benzenamine hydrochloride. The molecular formula is C₉H₉N₃Cl₂.HCl and the molecular weight is 266.56. The CAS number is 4205-91-8. The laboratory designation is ST 155.

Clonidine hydrochloride is a white or almost white, crystalline powder. It is soluble in ethanol, slightly soluble in chloroform and practically insoluble in ether. One gram is soluble in 13 mL of water (20°C).

Catapres 100 tablets contain the excipients maize starch, lactose, calcium hydrogen phosphate, colloidal anhydrous silica, povidone and stearic acid.

PHARMACOLOGY

Mode of Action

**Antihypertensive effect:** Catapres is predominantly an antihypertensive agent whose mechanism of action appears to be central alpha₂ adrenergic stimulation, as demonstrated in animal studies. This results in the inhibition of bulbar sympathetic cardioaccelerator and sympathetic vasoconstrictor centres, thereby causing a decrease in sympathetic outflow from the brain. There is an increase in vagal activity which produces a decrease in heart rate. There is also an increase in baroreceptor activity. Additionally Catapres stimulates peripheral alpha₁ adrenergic receptors. This is reflected by a small transient pressor effect (5-10 mmHg systolic blood pressure) following parenteral use. A transient rise in blood sugar may also occur following large doses of Catapres. The peripheral effects of Catapres generally require isolated organ type preparations for their demonstration, as in the intact animal or man, the central action predominates.

**Use in migraine prophylaxis and menopausal flushing:** Smaller doses of clonidine hydrochloride may be used for migraine prophylaxis and the alleviation of symptoms in menopausal flushing. The mechanism of action appears to be modification of the response of peripheral blood vessels to vasoconstrictor and vasodilator stimuli including noradrenaline, isoprenaline and angiotensin.
**Pharmacokinetic Studies**

Clonidine, the active ingredient of Catapres, is well absorbed from the gastrointestinal tract. Peak plasma concentrations are reached within 1-3 hours after oral administration. The duration of action varies from six to twelve hours, the duration of action being longer in the milder hypertensives.

The plasma half-life is 12-20 hours in patients with normal renal function. With impaired renal function it has been reported to increase to 18-48 hours.

The metabolic pathway of clonidine involves cleavage of the imidazolidine ring and the hydroxylation of the phenyl ring. Five metabolites have been identified in man and include para-hydroxy-clonidine and dichlorophenylguanidine.

Two-thirds of an administered dose is excreted in the urine (about half of which is unchanged Catapres) and the remainder is excreted in the faeces.

Given intravenously Catapres is effective within five minutes, has a maximum hypotensive action within 20 to 30 minutes, and the effect lasts for several hours. Following intramuscular administration, Catapres is effective within five to ten minutes. The maximum hypotensive effect is reached after 75 minutes and the duration of action is approximately five hours.

In a study designed to evaluate the pharmacokinetics of clonidine following administration of Catapres controlled release tablets (formulation not registered in Australia) in 30 patients (13 white patients, 6 black patients and 11 Hispanic patients), the pharmacokinetics was found to be similar between subjects from different racial groups.

The pharmacokinetics of clonidine are not influenced by food.

**INDICATIONS**

**Oral:**
- All grades of essential hypertension.
- Renal hypertension.

The prophylactic management of migraine or recurrent vascular headaches which occur in adult patients with a frequency of more than once a month and are not adequately relieved by appropriate therapy for the acute attack. Alleviation of symptoms due to localised vasodilatation in menopausal flushing.

**CONTRAINDICATIONS**

Catapres should not be used in patients with known hypersensitivity to the active ingredient, clonidine hydrochloride, and in patients with severe bradyarrhythmia resulting from either sick sinus syndrome or AV block of second or third degree.

In case of rare hereditary conditions that may be incompatible with an excipient of the product (please refer to Precautions) the use of the product is contraindicated.
PRECAUTIONS

Special care should be exercised in treating patients who have a history of depression or who have advanced cerebrovascular disease. Reduction of blood pressure in the latter circumstances may itself cause mental changes. Concurrent administration of tricyclic antidepressants may require adjustment of Catapres dosage.

Although a transient rise in blood sugar has been noted occasionally in humans treated with Catapres, which may be due to a pharmacologic alpha-adrenomimetic effect of the drug, no case of induced diabetes mellitus due to Catapres has been reported. Patients with clinical diabetes mellitus should be watched for a possible increase in their requirements of anti-diabetic therapy.

Catapres should be used with caution in patients with mild to moderate bradyarrhythmia such as low sinus rhythm, with disorders of cerebral or peripheral perfusion, polyneuropathy, and constipation.

No therapeutic effect of Catapres can be expected in the treatment of hypertension caused by phaeochromocytoma.

Since Catapres and its metabolites are extensively excreted in the urine, careful adjustment of dosage is required in patients with renal insufficiency (see Dosage).

As with other anti-hypertensives, treatment with Catapres should be monitored particularly carefully in patients with heart failure or severe coronary heart disease.

Termination of oral therapy should be gradual (e.g. over more than 7 days).

Sudden cessation of antihypertensive therapy is known to be associated in some instances with rebound hypertension which in some cases may be severe. This may occur with Catapres particularly in patients receiving more than the maximum recommended dose of 900 micrograms per day.

Following sudden discontinuation of Catapres after prolonged treatment with high doses, restlessness, palpitations, rapid rise in blood pressure, nervousness, tremor, headache or nausea have been reported.

If long-term treatment with a β-blocker needs to be interrupted, the β-blocker should be gradually phased out first, then clonidine.

Patients who wear contact lenses should be warned that treatment with Catapres may cause decreased lacrimation.

Catapres 100 tablets contain 324.9 mg of Lactose per maximum recommended daily dose. Patients with the rare hereditary conditions of galactose intolerance e.g. galactosaemia should not take this medicine.

Use in children and adolescents

The use and the safety of clonidin in children and adolescents has little supporting evidence in randomised controlled trials and therefore can not be recommended for use in this population.
In particular, when clonidine is used off-label concomitantly with methylphenidate in children with ADHD, serious adverse reactions, including death, have been observed. Therefore, clonidine in this combination is not recommended.

**Interactions with other medicines**

If the patient is on antihypertensive therapy, care should be taken as even a small dose of clonidine may further lower blood pressure and necessitate adjustment of the antihypertensive regime.

Where the antihypertensive agent is clonidine in the form of Catapres, it is illogical to add clonidine for the prophylaxis of migraine or the alleviation of symptoms in menopausal flushing. Catapres may potentiate the effects of alcohol, sedatives, hypnotics or other centrally active substances.

Although retinal, lens or corneal damage have not been detected with clonidine therapy, follow up procedures, such as ophthalmoscopy, are recommended.

Substances which raise blood pressure or induce a sodium and water retaining effect such as nonsteroidal anti-inflammatory drugs can reduce the therapeutic effect of clonidine.

Substances with $\alpha_2$-adrenergic receptor blocking properties, such as phentolamine or tolazoline, may abolish the $\alpha_2$-adrenergic receptor mediated effects of clonidine in a dose-dependent way.

Concomitant administration of drugs with a negative chronotropic or dromotropic effect such as $\beta$-blockers or digitalis glycosides can cause or potentiate bradycardiac rhythm disturbances.

It cannot be ruled out that concomitant administration of a $\beta$-blocker will cause or potentiate peripheral vascular disorders.

The antihypertensive effect of clonidine may be reduced or abolished and orthostatic regulation disturbances may be provoked or aggravated by concomitant administration of tricyclic antidepressants or neuroleptics with $\alpha$-receptor blocking effects.

Based on observations in patients in a state of delirium alcoholicum, it has been suggested that high intravenous doses of clonidine may increase the arrhythmogenic potential (QT-prolongation, ventricular fibrillation) of high intravenous doses of halopendol.

**Anaesthesia**

Abrupt withdrawal of Catapres is undesirable. Limited evidence suggests that it is unnecessary to withdraw Catapres before anaesthesia and that maintenance of therapy is preferable to abrupt withdrawal. In the peri-operative period Catapres can, where necessary, be administered parenterally until oral therapy is resumed.

Where therapy with Catapres is to be suspended before operation, withdrawal should be gradual (ie over more than 7 days) and monitored by regular observation of blood pressure.

**Carcinogenicity, Mutagenicity, Impairment of Fertility**

The carcinogenic potential of clonidine has not been assessed in an adequate range of studies. In rats, dietary administration of clonidine at doses up to 1.2 mg/kg/day (males) or
1.5 mg/kg/day (females) did not cause carcinogenic effects. These doses are 10-14 times the maximum recommended human daily dose of clonidine, based on body surface area.

Comprehensive investigations have not been performed to assess the potential genotoxic effects of clonidine. Clonidine showed no activity in the Ames test for mutagenicity.

Clonidine had no effect on fertility in male or female rats when administered orally at doses up to 0.15 mg/kg/day (35% higher than the maximum recommended total daily dose of clonidine in humans, based on body surface area).

**Use in Pregnancy:** Category B3

Clonidine hydrochloride has not shown teratogenic potential when tested in rats, but in some circumstances the incidence of embryonic and perinatal deaths was increased with doses comparable to those used clinically for antihypertensive therapy.

There are no well controlled studies in pregnant women, but the experience with clonidine hydrochloride since marketing does not include any positive evidence of adverse effect on the foetus. Since this experience cannot exclude such an effect, clonidine hydrochloride should be used during pregnancy only when the benefit clearly justifies the possible risk to the foetus.

Clonidine passes the placenta barrier, and may lower the heart rate of the foetus. Post partum, a transient rise in blood pressure in the newborn cannot be excluded.

Clonidine hydrochloride may also induce transitory elevation of blood glucose and impairment of glucose tolerance. Children born to mothers treated with clonidine hydrochloride during pregnancy should be specifically examined for changes in glucose metabolism.

During pregnancy the oral forms of Catapres are preferred, the intravenous injection of clonidine should be avoided.

**Use in Lactation**

Animal studies have shown that clonidine is secreted in the breast milk. As the effect on the newborn is not known, infants born to mothers being treated with Catapres should not be breast fed.

**Menopausal Flushing**

The efficacy of clonidine in the treatment of menopausal flushing has only been demonstrated in the first year after onset of symptoms.

**Instructions to be given to the Patient**

Patients should be advised not to drive cars or operate machinery until their competence to do so under treatment has been established. Prescribing instructions should be followed exactly for smooth control of blood pressure.

**Notice to Pharmacist**

Please do not remove or obliterate the patient advice labels on the carton.
ADVERSE EFFECTS

The following adverse events (regardless of causality) and incidences are based on a review of 22 clinical studies comprising 640 patients treated with clonidine hydrochloride.

Endocrine disorders:
≥0.01% and <0.1% gynaecomastia

Psychiatric disorders:
≥1% and <10% depression, sleep disorder
≥0.1% and <1% delusional perception, hallucination, nightmare
Not known confusional state, libido decreased

Nervous system disorders:
≥10% dizziness, sedation
≥1% and <10% headache
≥0.1% and <1% paraesthesia

Eye disorder:
≥0.01% and <0.1% lacrimation decreased
Not known accommodation disorder

Cardiac disorders:
≥0.1% and <1% sinus bradycardia
≥0.01% and <0.1% atrioventricular block
Not known bradyarrhythmia

Vascular disorders:
≥10% orthostatic hypotension
≥0.1% and <1% Raynaud's phenomenon

Respiratory, thoracic and mediastinal disorders:
≥0.01% and <0.1% nasal dryness

Gastrointestinal disorders:
≥10% dry mouth
≥1% and <10% constipation, nausea, salivary gland pain, vomiting
≥0.01% and <0.1% colonic pseudo-obstruction

Skin and subcutaneous tissue disorders:
≥0.1% and <1% pruritus, rash, urticaria
≥0.01% and <0.1% alopecia
Reproductive system and breast disorders:
≥1% and <10% erectile dysfunction

General disorders and administration site conditions:
≥1% and <10% fatigue
≥0.1% and <1% malaise

Investigations:
≥0.01% and <0.1% blood glucose increased

Most adverse effects are mild and tend to diminish with continued therapy.

Occasional reports of abnormal liver function tests and cases of hepatitis have also been reported.

DOSAGE AND ADMINISTRATION

The dosage recommendations are as follows:

Antihypertensive - initially 50-100 micrograms two to three times daily adjusted in small increments according to the patient's individual blood pressure response. If adequate control is not achieved with a daily dose of 600 micrograms of Catapres alone, additional therapy should be considered. Since the hypotensive effect of Catapres is dose dependent, it is usual to titrate the dose to satisfy the requirements for each patient. In impaired renal and hepatic function the half-life is prolonged and the dosage regime should be monitored carefully, as high variability in anti-hypertensive response is observed in patients with renal insufficiency. Since only a minimal amount of clonidine is removed during routine haemodialysis, there is no need to give supplemental clonidine following dialysis.

Migraine prophylaxis - initially 25 micrograms morning and evening. If necessary, after two weeks, this may be increased to 50 micrograms twice daily, then to a total daily dose of 150 micrograms. If the frequency of attacks is significantly reduced, consideration may be given to gradually ceasing therapy as remission may be sustained in a proportion of patients. Duration of treatment will depend upon the frequency and severity of attacks.

Menopausal flushing - initially 25 micrograms morning and evening. If after two weeks there has been no remission, increase to 50 micrograms twice daily. If necessary this may be increased to a total daily dose of 150 micrograms. Duration of treatment will depend upon the frequency and severity of attacks but long-term efficacy (longer than 8 weeks) in the treatment of menopausal flushing has not been established.

OVERDOSAGE

In case of poisoning or overdose, advice should be sought from the Poisons Information Centre (telephone 13 11 26).

The most important features of clonidine overdosage are likely to be bradycardia and sedation, respiratory depression including apnoea and somnolence including coma. Blood pressure response may be variable and may vary from severe hypotension (due to central sympathetic inhibition and vagal stimulation) to severe hypertension (due to direct alpha agonist activity). Treatment must therefore be appropriate to the clinical features (i.e. atropine followed by a pressor amine if necessary in patients with hypotension or an alpha
blocker such as phentolamine for patients with hypertension). Other features which may be seen include weakness, vomiting, diminished or absent reflexes, skin pallor, hypothermia, cardiac arrhythmias and constricted pupils with poor reaction to light.

**Management:** General supportive measures with regular checks of pulse, B.P., ECG, blood sugar and body temperature should be undertaken. The blood pressure should be monitored carefully for 48 hours following the overdosage, as a later hypertensive phase may be associated with declining blood levels of clonidine.

**PRESENTATION AND STORAGE CONDITIONS**

Catapres 100 tablets: scored white compressed tablets impressed with the symbol O1C/O1C on one side and the company symbol on the reverse side, each tablet containing 100 micrograms of clonidine hydrochloride. Blister packs of 100.

Catapres 100 tablets should be stored below 30°C.

**NAME AND ADDRESS OF THE SPONSOR**

Boehringer Ingelheim Pty Limited  
ABN 52 000 452 308  
78 Waterloo Road  
NORTH RYDE NSW 2113

**POISON SCHEDULE OF THE MEDICINE**

S4 – Prescription Only Medicine

**DATE OF APPROVAL**

TGA approval date: 28 January 2010