

- Spahn H, *et al.* Pharmacokinetics of amiloride in renal and hepatic disease. *Eur J Clin Pharmacol* 1987; **33**: 493–8.
- Sabanathan K, *et al.* A comparative study of the pharmacokinetics and pharmacodynamics of atenolol, hydrochlorothiazide and amiloride in normal young and elderly subjects and elderly hypertensive patients. *Eur J Clin Pharmacol* 1987; **32**: 53–60.
- Ismail Z, *et al.* The pharmacokinetics of amiloride-hydrochlorothiazide combination in the young and elderly. *Eur J Clin Pharmacol* 1989; **37**: 167–71.

## Uses and Administration

Amiloride is a weak diuretic that appears to act mainly on the distal renal tubules. It is described as potassium-sparing since, like spironolactone, it increases the excretion of sodium and reduces the excretion of potassium. Unlike spironolactone, however, it does not act by specifically antagonising aldosterone. Amiloride does not inhibit carbonic anhydrase. It takes effect about 2 hours after oral dosage and its diuretic action reaches a peak in 6 to 10 hours and has been reported to persist for about 24 hours.

Amiloride diminishes the kaliuretic effects of other diuretics, and may produce an additional natriuretic effect. It is mainly used as an adjunct to thiazide diuretics such as hydrochlorothiazide and loop diuretics such as furosemide, to conserve potassium in those at risk from hypokalaemia during the long-term treatment of oedema associated with hepatic cirrhosis (including ascites, p.1159) and heart failure (p.1165). It is also used with other diuretics in the treatment of hypertension (p.1171). Diuretic-induced hypokalaemia and its management, including the role of potassium-sparing diuretics such as amiloride, is discussed under Effects on Electrolyte Balance in the Adverse Effects of Hydrochlorothiazide, p.1308. Amiloride is sometimes used to manage hypokalaemia in primary hyperaldosteronism (p.1402).

Amiloride by inhalation has also been investigated in the management of cystic fibrosis patients with lung disease (see below).

In the treatment of oedema amiloride is given orally as the hydrochloride and doses are expressed in terms of the anhydrous substance. 1 mg of anhydrous hydrochloride is equivalent to about 1.14 mg of the hydrated substance. Treatment may be started with a dose of 5 to 10 mg daily, increased, if necessary, to a maximum of 20 mg daily. An initial dose of 2.5 mg once daily may be used in patients already taking other diuretics or antihypertensives. Similar doses to those given for oedema are used to reduce potassium loss in patients receiving thiazide or loop diuretics.

Potassium supplements should not be given.

**Cystic fibrosis.** Pulmonary disease is the major cause of mortality in cystic fibrosis (p.166). Experimental treatment aimed at modifying the pulmonary disease process has included giving amiloride by inhalation.<sup>1,2</sup> No evidence of pulmonary or systemic toxicity was seen in 14 patients treated for 25 weeks.<sup>1</sup> The mechanism of action is unclear but could be the sodium-channel blocking effect<sup>1</sup> or anti-inflammatory effects<sup>3</sup> of amiloride. Concern has been expressed<sup>4</sup> over possible consequences of the inhibition of endogenous urokinase by amiloride although others<sup>5</sup> considered this to be unlikely at the concentrations studied. However, a systematic review<sup>6</sup> found no evidence that amiloride was of clinical benefit.

- Knowles MR, *et al.* A pilot study of aerosolized amiloride for the treatment of lung disease in cystic fibrosis. *N Engl J Med* 1990; **322**: 1189–94.
- App EM, *et al.* Acute and long-term amiloride inhalation in cystic fibrosis lung disease: a rational approach to cystic fibrosis therapy. *Am Rev Respir Dis* 1990; **141**: 605–12.
- Gallo RL. Aerosolized amiloride for the treatment of lung disease in cystic fibrosis. *N Engl J Med* 1990; **323**: 996–7.
- Henkin J. Aerosolized amiloride for the treatment of lung disease in cystic fibrosis. *N Engl J Med* 1990; **323**: 997.
- Knowles MR, *et al.* Aerosolized amiloride for the treatment of lung disease in cystic fibrosis. *N Engl J Med* 1990; **323**: 997–8.
- Burrows E, *et al.* Sodium channel blockers for cystic fibrosis. Available in The Cochrane Database of Systematic Reviews; Issue 3. Chichester: John Wiley; 2006 (accessed 28/04/08).

**Diabetes insipidus.** Thiazide diuretics are commonly used in nephrogenic diabetes insipidus (p.2179) and NSAIDs may also be employed; both result in an overall decrease in urine production. Hydrochlorothiazide with amiloride has been reported to be at least as effective as hydrochlorothiazide plus indometacin in 5 patients.<sup>1</sup> In addition, amiloride obviated the need for potassium supplements. Hydrochlorothiazide with amiloride was also effective and well tolerated in a group of 4 children with nephrogenic diabetes insipidus who were treated for up to 5 years.<sup>2</sup>

- Knoers N, Monnens LAH. Amiloride-hydrochlorothiazide versus indometacin-hydrochlorothiazide in the treatment of nephrogenic diabetes insipidus. *J Pediatr* 1990; **117**: 499–502.
- Kirchlechner V, *et al.* Treatment of nephrogenic diabetes insipidus with hydrochlorothiazide and amiloride. *Arch Dis Child* 1999; **80**: 548–52.

**Renal calculi.** Patients with idiopathic hypercalciuria and a history of renal calculi (p.2181) are usually given a thiazide diuretic such as hydrochlorothiazide to reduce calcium excretion. In patients with calcium oxalate calculi an inherited cellular defect in oxalate transport may also be involved and this might be corrected by amiloride.<sup>1</sup>

- Baggio B, *et al.* An inheritable anomaly of red-cell oxalate transport in "primary" calcium nephrolithiasis correctable with diuretics. *N Engl J Med* 1986; **314**: 599–604.

## Preparations

**BP 2008:** Amiloride Tablets; Co-amilofruse Tablets; Co-amilozide Oral Solution; Co-amilozide Tablets.

**USP 31:** Amiloride Hydrochloride and Hydrochlorothiazide Tablets; Amiloride Hydrochloride Tablets.

**Proprietary Preparations** (details are given in Part 3)

**Austral:** Kaluri; Midamor; **Austria:** Midamor; **Canada:** Midamor; **Cz:** Amiclaran; **Denm:** Amikal; **Fin:** Medamor; **Fr:** Modamide; **NZ:** Midamor; **Swed:** Midamor; **Switz:** Midamor; **UK:** Amilamont; **USA:** Midamor.

**Multingredient Arg:** Amilorid; Diflux; Diur Pot; Diurex A; Erolon A; Furdinex; Hidrenox A; Lasinid; Moduretic; Nuriban A; Plenacor D; Prenomod; Ren-Ur; Vericordin Compuesto; **Austral:** Amizide; Moduretic; **Austria:** Aldoretic; Amilorid/HCT; Amiloretic; Amiloid comp; Amilostad HCT; Lanuretic; Loradur; Moducrin; Moduretic; **Belg:** Belidral; Co-Amiloride; Frusamil; Kalten; Moduretic; **Braz:** Amiretic; Diupress; Diurezin-A; Diursa; Moduretic; **Canada:** Apo-Amilzide; Gen-Amilzide; Modure; Novamior; Nu-Amilzide; **Chile:** Furdinex; Hidrium; Hidropid; **Cz:** Amiclorid; Amilorid/HCT; Apo-Amilzide; Liorid; Loradur; Moduretic; Rhefluin; **Denm:** Amilco; Buram; Frusamil; Moduretic; Sparkal; **Fin:** Amifrid; Diuramin; Diurex; Milorid; Moduretic; Sparkal; **Fr:** Logirene; Moducrin; Moduretic; **Ger:** Amilocomp beta; Amiloretic; Amiloid comp; Amilorid/HCT; Amilozid; Aquaretic; Diaphal; Diursan; durarese; Esmalorid; Moducrin; Moduretic; Tensoflux; **Gr:** Frumil; Ividol; Moduretic; Tia-den; **Hong Kong:** Amilco; Amithiazide; Apo-Amilzide; Moducrin; Moduretic; Navipare; Sefaretic; **Hung:** Amiloid Comp; Amilozid-B; **India:** Biduret; Frumil; Hilpes-D; **Indon:** Lorid; **Ir:** Amilco; Buram; Fru-Co; Frumil; Lasorid; Moduretic; Moduret; **Israel:** Kaluri; **Ital:** Moduretic; **Malaysia:** Ami-Hydrotride; Amilzide; Apo-Amilzide; Moduretic; **Mex:** Moduretic; **Neth:** Moduretic; **Norw:** Moduretic; Normonix; **NZ:** Amizide; Frumil; **Pol:** Tialid; **Port:** Aldoretic; Amilone Comp; Chibreticof; Diurene; Moducrin; Moduretic; **S.Afr:** Adco-Retic; Amiloretic; Betaretic; Hexaretic; Moducrin; Moduretic; Servatrin; **Singapore:** Apo-Amilzide; **Spain:** Ameride; Diuzine; Kalten; **Swed:** Amioferm; Moduretic; Normonix; **Switz:** Agorex; Amilo-basan; Amiloride/HCT; Betadur; Co-Amilozid; Comiloid; Ecodurex; Escoretic; Frumil; Grodurex; Kalten; Moducrin; Moduretic; Rhefluin; **Thai:** Biduretic; Hydrozide Plus; Hyperetic; Moduretic; Milorex; Miretic; Modulan; Moduretic; Moure-M; Poli-Uretic; Renase; Sefaretic; **Turk:** Moduretic; **UK:** Amil-Co; Andil; Bunex A; Froop Co; Fru-Co; Frumil; Kalten; Komil; Lasorid; Moducrin; Moduret; Moduretic; Navipare; **USA:** Moduretic; **Venez:** Furdinex; Moduretic.

## Amiodarone (BAN, USAN, rINN)

Amiodaron; Amiodarona; Amiodaroni; Amiodaronum; L-3428; 51087-N; SKF-33134-A. 2-Butylbenzofuran-3-yl 4-(2-diethylaminoethoxy)-3,5-diiodophenyl ketone.

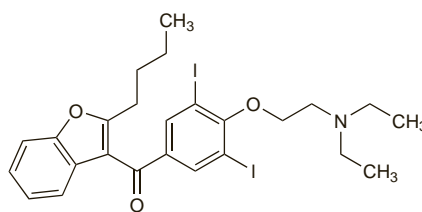
Амиодарон

$C_{25}H_{29}I_2NO_3$  = 645.3.

CAS — 1951-25-3.

ATC — C01BD01.

ATC Vet — QC01BD01.



## Amiodarone Hydrochloride (BANM, rINNM)

Amiodaron Hidroklorür; Amiodarone, chlorhydrate d'; Amiodaron-hidroklorid; Amiodaron-hydrochlorid; Amiodaronhidroklorid; Amiodaroni hidrochloridum; Amiodaronihydrochlorid; Amiodarona hidrochloridas; Hidrocloruro de amiodarona.

Амиодарона Гидрохлорид

$C_{25}H_{29}I_2NO_3 \cdot HCl$  = 681.8.

CAS — 19774-82-4.

ATC — C01BD01.

ATC Vet — QC01BD01.

**Pharmacopoies.** In Chin. and Eur. (see p.vii).

**Ph. Eur. 6.2** (Amiodarone Hydrochloride). A white or almost white, fine crystalline powder. Very slightly soluble in water; sparingly soluble in alcohol; freely soluble in dichloromethane;

soluble in methyl alcohol. Store at a temperature not exceeding 30°. Protect from light.

**Adsorption.** Amiodarone is known to be adsorbed by PVC, although the amount of adsorption has varied in different studies. A study<sup>1</sup> using amiodarone hydrochloride 600 micrograms/mL in glucose 5% found that the concentration fell by 10% in 3 hours followed by a steady decrease to 60% of the initial concentration after 5 days when stored in flexible PVC bags at ambient temperature.<sup>1</sup> However, another study<sup>2</sup> using amiodarone hydrochloride 1.8 to 2 mg/mL in glucose 5% found only that the concentration remained 97.3% of the initial value after 24 hours in PVC infusion bags. In the first study, perfusion of the solution through PVC giving sets resulted in the concentration falling to 82% after 15 minutes, whereas the second study found the concentration fell to 95.1% after 1 hour but then returned to the initial value. No loss was noted in either study when glass or rigid PVC containers were used, suggesting that the losses were caused by the plasticiser, di-2-ethylhexylphthalate (DEHP). Amiodarone may also leach out DEHP and other plasticisers, and it has been suggested that bags and tubing containing DEHP should not be used for giving amiodarone in order to minimise patient exposure.

- Weir SJ, *et al.* Sorption of amiodarone to polyvinyl chloride infusion bags and administration sets. *Am J Hosp Pharm* 1985; **42**: 2679–83.
- Peters PG, Hayball PJ. A comparative analysis of the loss of amiodarone from small and large volume PVC and non-PVC infusion systems. *Anaesth Intensive Care* 1990; **18**: 241–5.

**Incompatibility.** Amiodarone injection has been reported to be incompatible with aminophylline,<sup>1</sup> flucloxacillin,<sup>2</sup> heparin,<sup>3</sup> and sodium bicarbonate.<sup>4</sup> A further study<sup>5</sup> reported incompatibility with ampicillin/sulbactam sodium, ceftazidime sodium, digoxin, furosemide, imipenem/cilastatin sodium, magnesium sulfate, piperacillin sodium, piperacillin/tazobactam sodium, potassium phosphate, and sodium phosphate. UK licensed product information states that it is incompatible with sodium chloride solutions.

- Hasegawa GR, Eder JF. Visual compatibility of amiodarone hydrochloride injection with other injectable drugs. *Am J Hosp Pharm* 1984; **41**: 1379–80.
- Taylor A, Lewis R. Amiodarone and injectable drug incompatibility. *Pharm J* 1992; **248**: 533.
- Cairns CJ. Incompatibility of amiodarone. *Pharm J* 1986; **236**: 68.
- Korth-Bradley JM. Incompatibility of amiodarone hydrochloride and sodium bicarbonate injections. *Am J Health-Syst Pharm* 1995; **52**: 2340.
- Chalmers JR, *et al.* Visual compatibility of amiodarone hydrochloride injection with various intravenous drugs. *Am J Health-Syst Pharm* 2001; **58**: 504–6.

**Stability.** An oral suspension prepared from tablets<sup>1</sup> and containing amiodarone hydrochloride 5 mg/mL was stable for 3 months at 4° and 6 weeks at 25°.

- Nahata MC. Stability of amiodarone in an oral suspension stored under refrigeration and at room temperature. *Ann Pharmacother* 1997; **31**: 851–2.

## Adverse Effects and Treatment

Adverse effects are common with amiodarone. Many are dose-related and reversible with reduction in dose; however, because of its long half-life this can take some time and adverse effects may develop after treatment is stopped.

Adverse cardiovascular effects associated with amiodarone include severe bradycardia, sinus arrest, and conduction disturbances. Severe hypotension may follow intravenous use, particularly (though not exclusively) at rapid infusion rates. Amiodarone may also produce ventricular tachyarrhythmias; torsade de pointes has been reported but appears to be less of a problem with amiodarone than other antiarrhythmics. Rarely, heart failure may be precipitated or aggravated.

Amiodarone reduces the peripheral transformation of thyroxine (T<sub>4</sub>) to tri-iodothyronine (T<sub>3</sub>) and increases the formation of reverse-T<sub>3</sub>. It can affect thyroid function and may induce hypo- or hyperthyroidism.

There have been reports of severe pulmonary toxicity including pulmonary fibrosis and interstitial pneumonitis. These effects are usually reversible on withdrawal of amiodarone but are potentially fatal.

Amiodarone can adversely affect the liver. There may be abnormal liver function tests and cirrhosis or hepatitis; fatalities have been reported.

Prolonged use of amiodarone causes the development of benign yellowish-brown corneal microdeposits in the majority of patients, sometimes associated with coloured haloes of light; these are reversible on stopping therapy. Photosensitivity reactions are also common and more rarely blue-grey discoloration of the skin may occur.