Disturbed behaviour. Buspirone has been tried in various disorders for the control of symptoms such as agitation, aggression, and disruptive behaviour (see Disturbed Behaviour, p.954) but evidence of efficacy is limited. Nonetheless, in the management of dementia, some¹ consider that it might be worth trying in nonpsychotic patients with disturbed behaviour, especially those with mild symptoms or those intolerant or unresponsive to antipsychotics.

1. Rabins PV, et al. APA Work Group on Alzheimer's Disease and other Dementias. Steering Committee on Practice Guidelines. American Psychiatric Association practice guideline for the treatment of patients with Alzheimer's disease and other demen-tias. Second edition. Am J Psychiatry 2007; 164 (12 suppl): 5-56. Also available at: http://www.psychiatryonline.com/ pracGuide/loadGuidelinePdf.aspx?file=AlzPG101007 (ac-cessed 23/07/08) cessed 23/07/08)

Extrapyramidal disorders. Although there have been that buspirone may improve symptoms of drug-inreports duced dyskinesia (p.971), drugs with dopaminergic actions have mostly exacerbated symptoms and there are a few reports of extrapyramidal disorders with buspirone (see under Adverse Effects, above).

- Moss LE, et al. Buspirone in the treatment of tardive dyskinesia. J Clin Psychopharmacol 1993; 13: 204–9.
- 2. Bonifati V, et al. Buspirone in levodopa-induced dyskinesias. Clin Neuropharmacol 1994; 17: 73-82.

Substance dependence. ALCOHOL. Despite an early study¹ suggesting that buspirone could reduce alcohol craving in alcohol dependent patients, later studies2-4 have overall failed to confirm that buspirone improves abstinence or reduces alco-hol consumption. Although some studies^{4,5} have found that buspirone may improve certain psychopathological symptoms in these patients, others² have found no such benefit; a meta-analysis⁶ of 5 studies favoured the former interpretation.

The management of alcohol withdrawal and abstinence is discussed on p.1626.

- 1. Bruno F. Buspirone in the treatment of alcoholic patients. Psychopathology 1989; 22 (suppl 1): 49-59.
- 2. Malcolm R, et al. A placebo-controlled trial of buspirone in anxious inpatient alcoholics. Alcohol Clin Exp Res 1992; 16: 1007–13.
- George DT, et al. Buspirone does not promote long term absti-nence in alcoholics. Clin Pharmacol Ther 1995; 57: 161.
- Malec E, et al. Buspirone in the treatment of alcohol dependence: a placebo-controlled trial. Alcohol Clin Exp Res 1996; 20: ence: a p 307-12.
- 5. Kranzler HR. et al. Buspirone treatment of anxious alcoholics: a placebo-controlled trial. Arch Gen Psychiatry 1994; 51: 720-31.
- Malec TS, et al. Efficacy of buspirone in alcohol dependence: a review. Alcohol Clin Exp Res 1996; 20: 853–8.

NICOTINE. Buspirone has produced conflicting results1-5 in the management of smoking cessation (p.2354). Although some studies suggest that in the short-term buspirone can increase the numbers of patients who are able to cease smoking, it does not necessarily decrease withdrawal symptoms.

- 1. West R, et al. Effect of buspirone on cigarette withdrawal symptoms and short-term abstinence rates in a smokers clinic. Psy-chopharmacology (Berl) 1991; 104: 91-6.
- 2. Hilleman DE, et al. Effect of buspirone on withdrawal symptoms associated with smoking cessation. Arch Intern Med 1992; 152: 350-2.
- 3. Hilleman DE, et al. Comparison of fixed-dose transdermal nicotine, tapered-dose transdermal nicotine, and buspirone in smoking cessation. J Clin Pharmacol 1994; 34: 222-4.
- Schneider NG, et al. Efficacy of buspirone in smoking cessation: a placebo-controlled trial. Clin Pharmacol Ther 1996; 60: 568–75.
- Farid P, Abate MA. Buspirone use for smoking cessation. Ann Pharmacother 1998; 32: 1362–4.

OPIOIDS. Buspirone has been investigated in the management of opioid withdrawal (p.101) in dependent patients.

- 1. Rose JS, et al. Effects of buspirone in withdrawal from opiates. Am J Addict 2003; 12: 253-9.
- Buydens-Branchey L, et al. Efficacy of buspirone in the treat-ment of opioid withdrawal. J Clin Psychopharmacol 2005; 25: 230-6

Preparations

USP 31: Buspirone Hydrochloride Tablets.

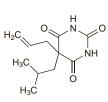
Proprietary Preparations (details are given in Part 3)

Proprietary Preparations (details are given in Part 3) Arg: Ansial'; Austral.: Buspar; Austria: Buspar; Belg:: Buspar; Braz.: An-sienon; Ansiate; Buspanith; Buspar; Canad.: Buspar; Buspirex; Chile: Paxon; Cz.: Anxiron†; Buspar; Denm.: Buspar; Stesiron†; Fin.: Buspar; Stesiron†; Fr.: Buspar; Ger.: Anxut: Bespar; Buspirex; Challe: Paxon; Lebilon; Ledion; Loxapin; Nadrifor; Nervostai; Nevrorestoi; Lontal; Pendium; Stressigai; Svitalark; Tendan†; Tensispes; Trafurith; Umolit; Hortba; Bergand; Buspar; Kalmiren; Hung:: Anxiron; Spitomin; India: Buscalm; Indon.: Tran-Q; Xiety, Irl.: Buspar; Israel: Buspirol†; Sorbon; Ital.: Ax-orent; Buspar; Buspinen†; Mex.: Buspar; Norw:: Buspar; Steisron†; NZ: Biron; Buspar; Pol.: Mabuson; Spamilan; Port:: Ansiten; Buspanit; Buscalma; Buspar; Buspinm; Istalbir(; Itagil; Paibeter; S.Afr.: Buspar; Thai:: Anxiolan; Turk:: Buspar; USA: Buspar; VSA:: Buspar; Switz:: Buspar; Thai:: Anxiolan; Turk:: Buspon; UK: Buspar; USA: Buspar; Yenez: Dalpas.

Butalbital (USAN, rINN)

Alisobumalum; Allylbarbital; Allylbarbituric Acid; Butalbitaali; Butalbitalum; Itobarbital; Tetrallobarbital. 5-Allyl-5-isobutylbarbituric acid.

Буталбитал $\dot{C}_{11}H_{16}N_2O_3 = 224.3.$ CAS — 77-26-9.



NOTE. The name Butalbital has also been applied to talbutal, the S-butyl analogue, which was formerly used as a hypnotic and sedative.

Compounded preparations of butalbital may be represented by the following names:

· Co-bucafAPAP (PEN)-butalbital, paracetamol, and caffeine Pharmacopoeias. In US.

USP 31 (Butalbital). A white odourless crystalline powder. Slightly soluble in cold water; soluble in boiling water; freely soluble in alcohol, in chloroform, and in ether; soluble in solutions of fixed alkalis and alkali carbonates. A saturated solution is acid to litmus.

Profile

Butalbital is a barbiturate with general properties similar to those of amobarbital (p.961). It has been used mainly in combination preparations with analgesics in the treatment of occasional ten-sion-type headaches, but other treatments are generally preferred.

Preparations

USP 31: Butalbital and Aspirin Tablets; Butalbital, Acetaminophen, and Caf-feine Capsules; Butalbital, Acetaminophen, and Caffeine Tablets; Butalbital, Aspirin, and Caffeine Capsules; Butalbital, Aspirin, and Caffeine Tablets; Butalbital, Aspirin, Caffeine, and Codeine Phosphate Capsules:

Proprietary Preparations (details are given in Part 3)

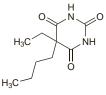
Multi-ingredient: Canad.: Fiorinal; Fiorinal C; ratio-Tecnal; ratio-Tecnal Multi-ingredient: Canad.: Honnal; Honnal; C; rato-lecnal; rato-lecnal C; Tirana; Tiranal C; Chile: Cafergot-PB; Penm:: Gynergen Comp; Ital:: Optalidom: S.Afr.: Cafergot-PB; Spain: Cafergot-PB; Switza:: Cafergot-PB; B; USA: Manphen with Codeine; Americet: Anolor: Ascomp with Co-deine: Bupap; Butex; Dolgic; Dolgic LQ; Dolgic Plus; Endolor; Esgic; Esgic-Plus; Fioricet; Fioricet with Codeine; Fiorinal; Fiorinal with Codeine: Marge-sic; Marten-Tab; Medgesic; Pacaps; Pinrenlin; Phrenlin; Margenlin w Caferine and Co-deine: Promacet; Prominol; Pyridium Plus; Repan; Repan; CF; Sedapap; Tenect: Tencon; Trellium Plus; Triad. Tencet; Tencon; Trellium Plus; Triad.

Butobarbital (BAN)

Butethal; Butobarbitaali; Butobarbitalum; Butobarbitone. 5-Butyl-5-ethylbarbituric acid.

 $C_{10}H_{16}N_2O_3 = 212.2.$ CAS = 77-28-1. ATC = N05CA03.

ATC Vet - QN05CA03.



NOTE. Butobarbital should be distinguished from Butabarbital, which is Secbutabarbital (p.1027).

Dependence and Withdrawal

As for Amobarbital, p.962

Adverse Effects, Treatment, and Precautions As for Amobarbital, p.962.

Interactions

As for Amobarbital, p.962.

Antibacterials. The metabolism of butobarbital may be altered

by metronidazole.1

 Al Sharifi MA, et al. The effect of anti-amoebic drug therapy on the metabolism of butobarbitone. J Pharm Pharmacol 1982; 34: 126-7.

Pharmacokinetics

Butobarbital is metabolised in the liver mainly by hydroxylation; small amounts are excreted in the urine as unchanged drug. It has been reported to have a half-life of about 40 to 55 hours and to be about 26% bound to plasma proteins.

Uses and Administration

Butobarbital is a barbiturate with general properties similar to those of amobarbital (p.962). Its use can no longer be recommended because of the risk of its adverse effects and of dependence, although continued use may occasionally be considered necessary for severe intractable insomnia (p.957) in patients already taking it. It is given in usual oral doses of 100 to 200 mg at night.

Preparations

Proprietary Preparations (details are given in Part 3)

UK: Soneryl

Multi-ingredient: Cz.: Dinyl+; Fr.: Hypnasmine+.

Calcium Bromolactobionate

Bromolactobionato de calcio; Calcium Galactogluconate Bromide. Calcium bromide lactobionate hexahydrate.

 $Ca(C_{12}H_{21}O_{12})_{2},CaBr_{2},6H_{2}O = 1062.6.$ CAS — 33659-28-8 (anhydrous calcium bromolactobion-

ate).

Profile

Calcium bromolactobionate has sedative properties and has been given orally in the treatment of insomnia and anxiety disorders. The use of bromides is generally deprecated.

Overdosage. Bromide intoxication has been reported1 in a patient after overdosage with calcium bromolactobionate tablets. Danel VC, et al. Bromide intoxication and pseudohyperchlo-remia. Ann Pharmacother 2001; 35: 386–7.

Preparations

Proprietary Preparations (details are given in Part 3) Chile: Bromocalcio; Nervolta; Sedofantil; Cz.: Calabron†; Ital.: Calcibro-nat; Mex.: Calcibronat†; Mon.: Calcibronat; Venez.: Sedabron†.

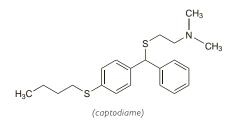
Captodiame Hydrochloride (BANM, pINNM)

Captodiame, Chlorhydrate de; Captodiami Hydrochloridum; Captodiamine Hydrochloride; Hidrocloruro de captodiamo. 2-(4-Butylthiobenzhydrylthio)ethyldimethylamine hydrochloride. Каптодиама Гидрохлорид

 $C_{21}H_{29}NS_2.HCl = 396.I.$ CAS = 486-17-9 (captodiame); 904-04-1 (captodiame hydrochloride)

ATC — N05BB02.

ATC Vet — QN05BB02.



Profile

Captodiame hydrochloride has been given in oral doses of 50 mg three times daily for the treatment of anxiety disorders (p.952).

Preparations

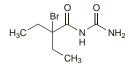
Proprietary Preparations (details are given in Part 3) Fr.: Covatine

Carbromal (BAN, HNN)

Bromodiethylacetylurea; Carbromalum; Karbromaali; Karbromal. N-(2-Bromo-2-ethylbutyryl)urea.

Карбромал $C_7H_{13}BrN_2O_2 = 237.1.$ CAS - 77-65-6. ATC - N05CM04.

ATC Vet - QN05CM04.



Profile

Carbromal is a bromureide with general properties similar to those of the barbiturates (see Amobarbital, p.961). It was formerly used for its hypnotic and sedative properties. Chronic use of carbromal could result in bromide accumulation and symptoms resembling bromism (see Bromides, p.2269). The use of bromides is generally deprecated.

Porphyria. Carbromal has been associated with acute attacks of porphyria and is considered unsafe in porphyric patients.