AUSTRALIAN CAPITAL TERRITORY

POISONS AND DRUGS ACT 1978

INSTRUMENT NO. 254 OF 2001

MODIFICATION

Under Section 46 of the *Poisons and Drugs Act 1978*, I, **Michael John Moore**, Minister for Health, Housing and Community Services hereby modify the drugs and poisons standard current on 1 September 2001 by withdrawing the entries listed in the attachment to this instrument.

Dated 29th day of August 2001.

Michael Moore
Minister for Health, Housing and Community Services

STANDARD FOR THE UNIFORM SCHEDULING OF DRUGS AND POISONS

No. 16

AMENDMENT No. 1

WITHDRAWN ENTRIES

Effective Date – 1 September 2001

PART 4 - THE SCHEDULES

Schedule 2 – New entries

*DATURA TATULA (stramonium) for oral use:

(a) in undivided preparations containing 0.025 per cent or less of the alkaloids of stramonium when labelled with a dose of 0.025 mg or less of the alkaloids of stramonium and a recommended daily dose of 0.5 mg or less of the alkaloids of stramonium; or

(b) in divided preparations containing 0.025 mg or less of the alkaloids of stramonium per dosage unit when labelled with a recommended daily dose of 0.5 mg or less of the alkaloids of stramonium;

except for smoking or burning.

Schedule 2 – Amendments

*ATROPINE – amend entry to read:

ATROPINE (excluding atropine methonitrate):

(a) for oral use:

(i) in undivided preparations containing 0.025 per cent or less of atropine when labelled with a dose of 0.025 mg or less of atropine and a recommended daily dose of 0.5 mg or less of atropine; or

(ii) in divided preparations containing 0.025 mg or less of atropine per dosage unit when labelled with a recommended daily dose of 0.5 mg or less of atropine; or

(b) in preparations containing atropine sulfate when packed and labelled for the treatment of organophosphorus poisoning:

(i) in tablets each containing 0.6 mg or less of atropine sulfate in packs of 20 tablets; or

(ii) in preparations for injection each containing 0.6 mg per ml or less of atropine sulfate in packs of 5.

*BELLADONNA – amend entry to read:

ATROPA BELLADONNA (belladonna):

(a) for external use in preparations containing 0.025 per cent or less of the alkaloids of belladonna; or

(b) for oral use:

(i) in undivided preparations containing 0.025 per cent or less of the alkaloids of belladonna when labelled with a dose of 0.025 mg or less of the alkaloids of belladonna and a recommended daily dose of 0.5 mg or less of the alkaloids of belladonna; or

(ii) in divided preparations containing 0.025 mg or less of the alkaloids of belladonna per dosage unit, when labelled with a recommended daily dose of 0.5 mg or less of the alkaloids of belladonna.

*DATURA – amend entry to read:

DATURA spp. for oral use **except** when separately specified in these Schedules:

(a) in undivided preparations containing 0.025 per cent or less of the alkaloids of datura when labelled with a dose of 0.3 mg or less of the alkaloids of datura and a recommended daily dose of 1 mg or less of the alkaloids of datura; or

(b) in divided preparations containing 0.3 mg or less of the alkaloids of datura per dosage unit when labelled with a recommended daily dose of 1 mg or less of the alkaloids of datura.

*DUBOISIA LEICHARDTII – amend entry to read:

DUBOISIA LEICHARDTII for oral use:

(a) in undivided preparations containing 0.025 per cent or less of the alkaloids of duboisia calculated as hyoscyamine when labelled with a dose of 0.025 mg or less of the alkaloids of duboisia calculated as hyoscyamine and a recommended daily dose of 0.5 mg or less of the alkaloids of duboisia calculated as hyoscyamine; or

(b) in divided preparations containing 0.025 mg or less of the alkaloids of duboisia calculated as hyoscyamine per dosage unit when labelled with a recommended daily dose of 0.5 mg or less of the alkaloids of duboisia calculated as hyoscyamine.

*DUBOISIA MYOPOROIDES – amend entry to read:

DUBOISIA MYOPOROIDES for oral use:

(a) in undivided preparations containing 0.025 per cent or less of the alkaloids of duboisia calculated as hyoscyamine when labelled with a dose of 0.025 mg or less of the alkaloids of duboisia calculated as hyoscyamine and a recommended daily dose of 0.5 mg or less of the alkaloids of duboisia calculated as hyoscyamine; or

(b) in divided preparations containing 0.025 mg or less of the alkaloids of duboisia calculated as hyoscyamine per dosage unit when labelled with a recommended daily dose of 0.5 mg or less of the alkaloids of duboisia calculated as hyoscyamine.

*HYOSCINE – amend entry to read:

HYOSCINE (excluding hyoscine butylbromide):

(a) for transdermal use in preparations containing 2 mg or less of hyoscine; or

(b) for oral use:

(i) in undivided preparations containing 0.025 per cent or less of hyoscine, when labelled with a dose of 0.3 mg or less of hyoscine and a recommended daily dose of 1 mg or less of hyoscine; or

(ii) in divided preparations containing 0.3 mg or less of hyoscine per dosage unit when labelled with a recommended daily dose of 1 mg or less of hyoscine.

HYOSCYAMINE – amend entry to read:

HYOSCYAMINE:

(a) for external use in preparations containing 0.025 per cent or less of hyoscyamine; or

(b) for oral use:

(i) in undivided preparations containing 0.025 per cent or less of hyoscyamine, when labelled with a dose of 0.025 mg or less of hyoscyamine and a recommended daily dose of 0.5 milligrams or less of hyoscyamine; or

(ii) in divided preparations containing 0.025 mg or less of hyoscyamine per dosage unit when labelled with a recommended daily dose of 0.5 mg or less hyoscyamine.

HYOSCYAMUS – amend entry to read:

HYOSCYAMUS NIGER for oral use:

(a) in undivided preparations containing 0.025 per cent or less of the alkaloids of hyoscyamus when labelled with a dose of 0.025 mg or less of the alkaloids of hyoscyamus and a recommended daily dose of 0.5 mg or less of the alkaloids of hyoscyamus; or

(b) in divided preparations containing 0.025 mg of the alkaloids of hyoscyamus or less per dosage unit when labelled with a recommended daily dose of 0.5 mg or less of the alkaloids of hyoscyamus.

*STRAMONIUM – amend entry to read:

DATURA STRAMONIUM (stramonium) for oral use when:

(a) in undivided preparations containing 0.025 per cent or less of the alkaloids of stramonium when labelled with a dose of 0.025 mg or less of the alkaloids of stramonium and a recommended daily dose of 0.5 mg or less of the alkaloids of stramonium; or

(b) in divided preparations containing 0.025 mg or less of the alkaloids of stramonium per dosage unit when labelled with a recommended daily dose of 0.5 mg or less of the alkaloids of stramonium;

except for smoking or burning.

Schedule 4 – New entries

*DATURA TATULA (stramonium) **except**:

(a) when included in Schedule 2; or

(b) for smoking or burning.

Schedule 4 – Amendments

*BELLADONNA – amend entry to read:

ATROPA BELLADONNA (belladonna) **except** when included in Schedule 2.

*DUBOISIA LEICHARDTII – amend entry to read:

DUBOISIA LEICHARDTII **except** when included in Schedule 2.

*DUBOISIA MYOPOROIDES – amend entry to read:

DUBOISIA MYOPOROIDES **except** when included in Schedule 2.

*HYOSYCAMUS – amend entry to read:

HYOSCYAMUS NIGER **except** when included in Schedule 2.

*STRAMONIUM – amend entry to read:

DATURA STRAMONIUM (stramonium) **except**:

(a) when included in Schedule 2; or

(b) for smoking or burning.