Alternative syntheses of bromo-dragonFLY

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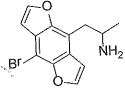
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Drug abuse in Australia has shifted from widespread opiate use towards the use of amphetamine-type stimulants (ATS), such as amphetamine ('speed'), methamphetamine (MA, 'ice'), and 3,4-methylenedioxymethamphetamine (MDMA, 'ecstasy').1 ATS are the second most commonly used illicit drugs in Australia, behind cannabis, and these drugs are strongly established on the global illicit drug market.1

Recent designer amphetamines have been in the spotlight, receiving significant attention on online drug-use forums, suggesting that these drugs may soon become a problem in Australia.2 The most potent hallucinogenic amphetamine analogue synthesised to date is the bromo-substituted benzodifuran; 2-amino-1-(8-bromobenzo[1,2-b;4,5-b']difuran-4yl]propane, known as 'bromo-dragonFLY' due to its molecular configuration.3

The reported syntheses for this drug surpass the capabilities of clandestine drug laboratories and clandestine chemists. This research project aims to pre-empt prospective clandestine synthesis, allowing the potential monitoring and/or restriction of the necessary precursor chemicals to prevent the illicit synthesis of bromo-dragonFLY.





Amphetamine ("speed")

Methamphetamine (MA, "jce")

Bromo dragonFLY

- 1. Australian Crime Commission, Illicit Drug Data Report 2009-10.
- 2. (a) Bluelight. http://www.bluelight.ru/vb/home.php (accessed 9th January 2012); (b) Chemistry Archive. http://www. erowid.org/archive/rhodium/chemistry/index.html (accessed 4th January 2012).
- 3. McLean, T. H.; Parrish, J. C.; Braden, M. R.; Marona-Lewicka, D.; Gallardo-Godoy, A.; Nichols, D. E., 1-Aminomethylbenzocycloalkanes: Conformationally Restricted Hallucinogenic Phenethylamine Analogues as Functionally Selective 5-HT2A Receptor Agonists. J. Med. Chem. 2006, 49 (19), 5794-5803.