

Benzimidazole-Opioids Other Name: Nitazenes

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Introduction:

Recently, a number of synthetic substances of benzimidazole structural class are being trafficked and abused for their opioidlike effects. In the late 1950s, the pharmaceutical research laboratories of the Swiss chemical company CIBA Aktiengesellschaft synthesized numerous substances in this structural class. Since 2019, the abuse of benzimidazole-opioids as evidenced by their identification in toxicology cases, similar to other synthetic opioids, has resulted in adverse health effects including deaths. As the United States continues to experience an unprecedented epidemic of opioid misuse and abuse, the continued evolution and increased trafficking and popularity of new and deadly synthetic opioids from a variety of structural classes, including benzimidazoles, with no approved medical use are of public health concern.

Chemistry:

This class of substances contains a benzimidazole ring with an ethylamine at its 1-position and a benzyl group at its 2-position. Small structural modifications to this scaffold can produce a series of analogous substances, including the substances listed below (shown below):

` R₄ _ ´		R ₁	R ₂	R_3	R_4
	metonitazene etonitazene etodesnitazene protonitazene butonitazene clonitazene flunitazene etonitazpyne etonitazpyne	- NO2 H NO2 H NO2 NO	2 OCH ₃ OCH ₂ CH ₃ OCH ₂ CH ₃ OCH ₂ CH ₂ CH ₃ OCH ₂ CH ₂ CH ₃ OCH ₂ CH ₂ CH ₃ OCH(CH ₃) ₂ Cl F OCH ₂ CH ₃ OCH ₂ CH ₃	$\begin{array}{c} {\rm CH}_{2}{\rm CH}_{3} \\ {\rm CH}_{3} \\$	CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3 CH_2CH_3

Pharmacology:

Data obtained from pre-clinical studies demonstrate that benzimidazole-opioids exhibit pharmacological profile similar to that of etonitazene and other mu-opioid receptor agonists. In an antinociceptive study conducted in rodents, butonitazene, etodesnitazene, etonitazepipne, etonitazepyne, flunitazene, metodesnitazene, metonitazene, and protonitazene, similar to morphine, produced strong analgesic effects with varying potencies. As compared to morphine, flunitazene and metodesnitazene are equipotent, while butonitazene. etodesnitazene, etonitazepyne, metonitazene, and protonitazene are more potent as analgesics. Data from in vitro that butonitazene, etodesnitazene, studies showed etonitazepipne, etonitazepyne, flunitazene, metonitazene, metodesnitazene, and protonitazene, similar to fentanyl and morphine, bound to and activated the mu-opioid receptor, and thus acted as mu-opioid receptor agonists. Activation of the muopioid receptor butonitazene, metonitazene, by

metodesnitazene, and protonitazene involved interaction with β -arrestin-2. Mu-opioid receptor and β -arrestin-2 interaction has been implicated in adverse health effects of many opioid analgesics. It is well established that mu-opioid receptor agonists have a high potential for addiction and can produce dose-dependent respiratory depression and arrest. Abuse of these benzimidazole-opioids has led to their positive identification in several toxicological cases in the United States. Some of these benzimidazole-opioids have been positively identified in at least 94 toxicology cases.

User Population:

The population likely to abuse benzimidazole-opioids appears to be the same as those abusing prescription opioid analgesics, heroin, and other synthetic opioid substances. This is evidenced by the types of other drugs co-identified in some of the identified benzimidazole-opioids seizures and in fatal overdose cases. Toxicology analyses co-identified some of these benzimidazoleopioids with other opioids, stimulants, and benzodiazepines. Because abusers of these benzimidazole-opioids are likely to obtain them through unregulated sources, the identity, purity, and quantity are uncertain and inconsistent, thus posing significant adverse health risks to the users. Similar to other mu-opioid receptor agonists, the potential health and safety risks for users of these benzimidazole-opioids are high. Recent increase in positive of isotonitazene, metonitazene, identification and other benzimidazole-opioids in toxicology and post-mortem cases is a serious concern to the public safety.

Illicit Distribution:

On the illicit drug market, some of these benzimidazole-opioids have been identified in drug seizures. Reports of these substances to the National Forensic Laboratory Information System (NFLIS-Drug), began to re-emerge in 2019 after several reports of clonitazene and etonitazene in 1999-2004. Since 2019, there have been approximately 2,400 reports of benzimidazole-opioids to NFLIS-Drug. Furthermore, substances in this class have been co-identified with other psychoactive substances, including illicit opioids and benzodiazepines, in biological fluids. With no approved medical use, the positive identification of these substances in toxicology cases underscore the public health threat associated with their presence on the illicit drug market.

Control Status:

These benzimidazole-opioids are not approved for medical use in the United States. Ten of these benzimidazole-opioids are controlled in schedule I of the CSA; others may be treated under 21 U.S.C. 802(32)(A), if intended for human consumption.

Comments and additional information are welcomed by the Drug and Chemical Evaluation Section; Fax 571-362-4250, Telephone 571-362-3249, or E-mail DPE@dea.gov.