

Heroin intoxication in a dog

Fergal M McDermott^{1, 2}, E. E. Henriksson^{1, 3}, T. A. Wismer⁴

¹ Western College of Veterinary Medicine, University of Saskatchewan, Saskatoon, Canada

² Utrecht University, Utrecht, the Netherlands

³ Auburn University, Auburn, USA.

⁴ ASPCA Animal Poison Control Center, Urbana, USA.

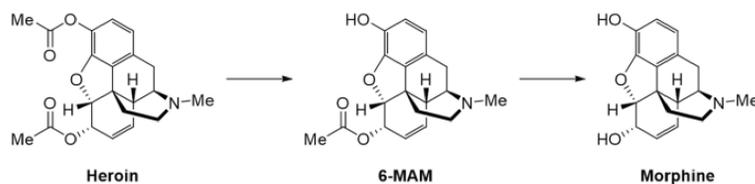



Figure 1: The metabolic pathway of heroin to morphine¹.

Background

The opioid crisis continues² and it is not an exclusively human problem. Reports of accidental opioid intoxication in dogs were determined to be higher in areas of higher opioid prescription³. The lethal heroin dose for dogs is reported to be 25 mg/kg⁴. This report describes the successful treatment of 40 mg/kg heroin intoxication in a dog and outlines insights in opioid intoxication.

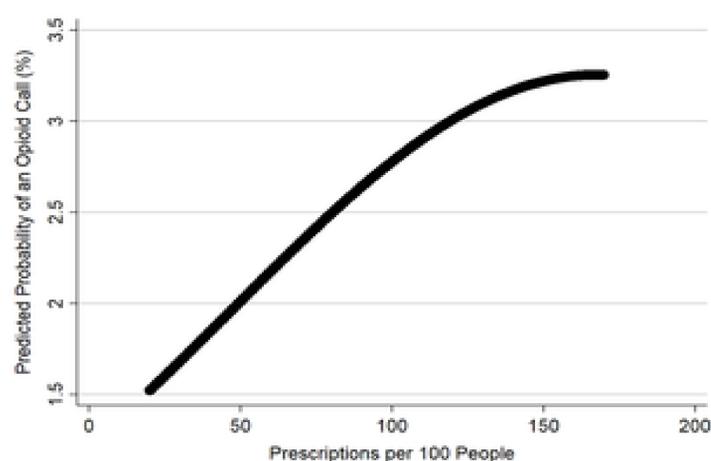


Fig 2. Predicted probability of opioid poisoning calls to the ASPCA poison control hotline plotted against county-level prescription rate (2006–2014)²

Case Presentation

An 11-month-old, intact male boxer cross was presented having become unconscious shortly after it consumed 1g of powdered heroin. On physical examination the dog was comatose and laterally recumbent. There was bilateral pupillary miosis and marked ptyalism. The heart-rate was 60 bpm and respiratory-rate was 20 bpm. The dog was normothermic (37.4 °C) and normotensive (163/103 (123) mmHg) with a 96% SpO₂.

Treatment

Intravenous naloxone (0.02 mg/kg) quickly (<1 min) reversed clinical signs. The owners were unable to afford hospitalization. They purchased naloxone which they were instructed to administer intranasally (IN) should signs of toxicity recur. The animal was discharged against medical advice (AMA) with no rebound opioid toxicity signs reported at 2-month follow-up.

Clinical Relevance

The classic toxidrome of opioid intoxication consists of respiratory and central nervous system depression and pupillary miosis⁵. Many substances (ethanol, antipsychotics, etc.) cause miosis and coma, without respiratory depression⁵.

Heroin is commonly mixed with other drugs, such as the increasingly consumed fentanyl². Delayed gastric emptying, enzyme saturation and the resulting zero-order elimination cause oral opioid intoxications to be unpredictable. The different pharmacokinetics of various opioid agents are rarely relevant in the event of oral overdose⁵. Therefore, the management of opioid intoxication, regardless of which opioid is involved, is founded on naloxone administration and respiratory support if necessary.

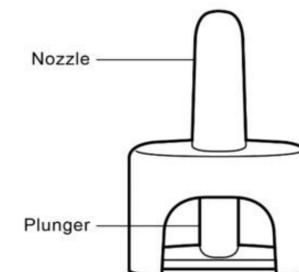


Figure 3: Pre-made IN naloxone devices. Injectable naloxone can be effective with unpredictable pharmacokinetics⁷.

Clinical Relevance (continued)

Naloxone is a safe drug and should be given if opioid intoxication is ever suspected. If clinical signs do not improve, opioid intoxication is unlikely. It is effective when given IN⁶. Atomized formulas are ideal, but the injectable formula is effective IN with less predictable results⁷. A suggested IN dose is 2–4 mg/25 kg⁸. If unavailable, butorphanol at 0.4 mg/kg may be useful⁹. The short duration of intravenous naloxone (90 minutes) in relation to the opioids involved in overdose means rebound opioid toxicities occur¹⁰. Large-scale human studies suggest a low risk in discharging a heroin intoxication patient AMA 1 hour post reversal, provided the patient has normal vital parameters and is neurologically appropriate¹¹.

References

- Zhang, T, Zheng, X., Kim, K. *et al.* Blocking drug activation as a therapeutic strategy to attenuate acute toxicity and physiological effects of heroin. *Sci Rep* 8, 16762 (2018). <https://doi.org/10.1038/s41598-018-35196-8>
- United Nations Office on Drugs and Crime. World drug report 2020. 25 June 2020. <https://wdr.unodc.org/wdr2020/index.html>
- Howard-Azzeh M, Pearl DL, O'Sullivan TL, Berke O. The identification of risk factors contributing to accidental opioid poisonings in companion dogs using data from a North American poison control center (2006–2014). *PLoS one*. 2020 Jan 29;15(1):e0227701.
- Volmer PA. "Recreational" drugs. In *Small animal toxicology* 2013 Jan 1 (pp. 309–334). WB Saunders.
- Boyer EW. Management of opioid analgesic overdose. *New England Journal of Medicine*. 2012 Jul 12;367(2):146–55.
- Wahler BM, Lerche P, Ricco Pereira CH, Bednarski RM, KuKanich B, Lakritz J, Aarnes TK. Pharmacokinetics and pharmacodynamics of intranasal and intravenous naloxone hydrochloride administration in healthy dogs. *American journal of veterinary research*. 2019 Jul;80(7):696–701.
- Krieter P, Chiang N, Gyaw S, Skolnick P, Crystal R, Keegan F, Aker J, Beck M, Harris J. Pharmacokinetic properties and human use characteristics of an FDA-approved intranasal naloxone product for the treatment of opioid overdose. *The Journal of Clinical Pharmacology*. 2016 Oct 1;56(10):1243–53.
- Tonozi C, McMichael M, Mitek A, Weir W, Smith M, Cornwell N. Non-Veterinary Emergency Care of Law Enforcement Canines at Mass Gathering Events A Pilot Training Course for Collegiate-Based EMS Providers. *J Coll Emerg Med Serv*. 2019;2(1):03.
- Hovda L, Brutlag A, Poppenga RH, Peterson K, editors. *Blackwell's Five-Minute Veterinary Consult Clinical Companion: Small Animal Toxicology*. John Wiley & Sons; 2016 May 2.
- Berkowitz BA. The relationship of pharmacokinetics to pharmacological activity: morphine, methadone and naloxone. *Clinical pharmacokinetics*. 1976 Jun 1;1(3):219–188.
- Willman MW, Liss DB, Schwarz ES, Mullins ME. Do heroin overdose patients require observation after receiving naloxone?. *Clinical toxicology*. 2017 Feb 7;55(2):81–7.