## **Wisconsin Department of Justice**



**Tramadol Monograph** 

## **Tramadol - General Effects**

**Tramadol** (Ultram, Conzip, Nobligan) is a narcotic analgesic and central nervous system (CNS) depressant.

- 1. Tramadol is prescribed for the relief of moderate to severe pain.
  - a. Tramadol has a half-life of 4.3-8.3 hours.
  - b. Tramadol has a general therapeutic range, but experienced effects can vary based on an individual's prescription history.
    - i. Therapeutic range refers to the blood concentration expected to achieve the desired therapeutic effects. Due to many factors such as prescription history, dosage, tolerance, drug-drug interactions and use, an individual may exhibit signs of impairment even though blood concentrations fall within the therapeutic range.
  - c. Tramadol is equipotent to codeine but has less abuse potential as it causes less respiratory depression.
- 2. O-desmethyltramadol, an active metabolite of tramadol, has around 2-4 times the efficacy of tramadol.
  - a. O-desmethyltramadol has a half-life of 6-11 hours.
- 3. General effects of narcotic analgesics include but are not limited to: nausea, vomiting, respiratory depression, sedation, and mental clouding/mood swings.
  - a. General impairing effects of narcotic analgesics on driving include, but are not limited to: impaired divided attention, poor coordination, cognitive impairment, delayed reaction time, difficulty following direction, and falling asleep at the wheel.
- 4. The longer an individual uses a drug, the more they can build up a tolerance to its effects. Tolerance occurs when an individual no longer responds to the drug in the way that they initially responded. When an individual gains tolerance to a drug, a higher dose of the drug is necessary to achieve the same level or response initially achieved. As tolerance is gained, it may reduce some of the possible negative effects of a drug.
- 5. Drug metabolism (alcohol excluded) exhibits first order kinetics, or the elimination of a constant fraction of drug quantity per unit of time, which means that the amount eliminated is proportional to the drug concentration.
- 6. The use of more than one drug at a time may enhance the effects the drugs would otherwise have on their own, leading to greater impairment.

## References

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- 3. Moffat, A. C., Osselton, M. D., Widdop, B., & Watts, J. (2011). Clarke's analysis of drugs and poisons (3rd ed., Vol. 1). Pharmaceutical Press.
- 4. Levine, B., & Vina Spiehler. (2020). Pharmacokinetics and Pharmacodynamics. In B. Levine (Ed.), Principles of Forensic Toxicology (4th ed., pp. 77–93). essay, AACC Press.

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\*\* The interpretive information provided is not exhaustive nor meant to encompass all scenarios where toxicological results are reported. Interpretive information is meant to serve as a general guide for the reader and that for any given case, consultation with a forensic toxicologist is recommended. \*\*

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