

## **Benzodiazep**ines Benzodiazepine common names: barbs, benzos, downers, GHB, Georgia Home Boy, Grievous Bodily Harm, Liquid X, Nerve pills, phennies, R2, Reds, Roofies, Rophies, Tranks, Yellows<sup>2</sup> Most Commonly Abused (Valium, Xanax, Halcion, Ativan, Klonopin)<sup>2</sup> . **Benzodiazepines** Positive allosteric modulators of the GABAA-chloride receptor complex. Binding to the "benzodiazepine" GABA<sub>A</sub> receptor complex increases the frequency of opening of the chloride channels, facilitating inhibition of neuronal firing at the level of the limbic system, the brain stem reticular formation, and the cortex<sup>2</sup> **Characteristics** Onset of action is dependent on the kind of benzodiazepine. For diazepam, it is 15 (Depressant) minutes or less with an elimination half-life of 20-80 hours. For lorazepam, onset of action is 15-30 minutes with a half-life of 10-20 hours. For clonazepam, the onset of action is 15-30 minutes with a half-life of 18-50 hours.<sup>2</sup> Benzodiazepines may be detectable in the urine for approximately 2-4 days<sup>5</sup> Common signs and symptoms of intoxication can include: <sup>2</sup> Decreased motor coordination Sedation Decreased concentration Confusion and disorientation Presentation during Overdose intoxication Symptoms include hypotension, respiratory depression and comma<sup>2</sup> • Slurred speech, confusion, severe drowsiness, weakness and staggering, slow heartbeat, • breathing problems and unconsciousness Goal<sup>6</sup> Prevent severe respiratory depression • Monitor<sup>2,6</sup>: Assess level of disorientation and if possible time of last ingestion and amount consumed Monitoring and Monitor for falls risk support during Monitor vitals every 15 minutes initially and less frequently as acute symptoms subside • intoxication Supportive Interventions<sup>6</sup>: Ensure a quiet private space • Frequently orient client to reality and surroundings • Promote fluid and food intake as tolerated If Overdose<sup>2</sup>: Flumazenil injection (a benzodiazepine antagonist) reverses the hypnotic-sedative effects of benzodiazepines. Withdrawal Symptoms may include: presentation<sup>2</sup> Withdrawal occurs 1-2 Insomnia Headaches **Muscle Aches** Agitation days with a short acting Twitches Tremors Diaphoresis Anxiety agent (such as oxazepam, GI distress Perceptual Dysphoria Tachycardia alprazolam and lorazepam) and continues Changes 2-4 weeks or longer. \* Severe withdrawal symptoms may include paranoia and delirium Withdrawal occurs 2-7 days after the last dose \* Severe reactions such as grand mal or petit mal seizures , depersonalization, psychotic (of a long-acting agent) states, and coma may occur (especially with alprazolam) and continues for 2-8 weeks or longer.



Monitoring and support during withdrawal	<ul> <li>Goal<sup>6</sup></li> <li>Preserve respiratory and cardiovascular function and reduce withdrawal symptoms Monitor<sup>2,6</sup></li> <li>Monitor regularly for withdrawal symptoms</li> <li>Monitor mental status</li> <li>Monitor risk for falls</li> <li>Monitor hydration/nutrition and sleeping patterns</li> </ul> Supportive Interventions <sup>6</sup> <ul> <li>Provide reassurance and explanation of symptoms if necessary</li> <li>Provide a calm and quiet environment</li> <li>Withdrawals have also been managed by administering benzodiazepines regularly in gradually decreasing amounts (tapering)<sup>3</sup></li></ul>
Potential Complications	<ul> <li>May include:</li> <li>Benzodiazepines can cause extensions of the generalized sedative effect (e.g., fatigue, drowsiness)</li> <li>Impaired mental speed, central cognitive processing ability, memory and performance.</li> <li>Anterograde amnesia (more likely with higher doses).</li> <li>Chronic use can cause impaired visuospatial and visuomotor abilities.</li> <li>Confusion and disorientation</li> <li>Excessive doses can result in respiratory depression and apnea.<sup>2</sup></li> </ul>
Notable Drug interactions	<ul> <li>With Antidepressants<sup>2</sup></li> <li>Cyclic antidepressants (such as desipramine and imipramine) and benzodiazepines can contribute to increased plasma levels of the antidepressant. Hypothermia has also been reported.</li> <li>With SSRIs (fluoxetine, fluvoxamine and sertraline), there is decreased metabolism and increased plasma level of benzodiazepines.</li> <li>With Antigsychotics<sup>2</sup></li> <li>With Antigsychotics<sup>2</sup></li> <li>With Alcohol<sup>2</sup></li> <li>Potentiation of CNS effects</li> <li>Alprazolam reported to increase aggression in moderate alcohol drinkers</li> <li>Brain concentrations of various benzodiazepines altered by ethanol (triazolam, estazolam concentration dincreased)</li> <li>With Antigsychotics<sup>2</sup></li> <li>With Cozapine, there can be marked sedation, increased salivation, hypotension, delirium, and respiratory depression.</li> <li>With olanzapine, there may be a synergistic increase in somnolence. IM olanzapine and benzodiazepines can potentiate hypotension, bradycardia, and respiratory or CNS depression.</li> <li>With Lithium<sup>2</sup></li> <li>Increased incidence of sexual dysfunction (up to 49%) has been reported with the use of</li> </ul>



Psychiatric effects

- Benzodiazepines can contribute to depression, particularly with high doses and in people with a pre-existing mood disorder.
- Benzodiazepines sometimes have a disinhibiting effect, (especially among people with psychosis or certain personality disorders). <sup>1</sup>



## References

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