

### **3. Pharmacological Overview**

Antidepressants encompass several pharmacological classes with distinct toxic profiles:

- **Tricyclic antidepressants (TCAs):** potent sodium-channel blockers with high lethality.
- **Selective serotonin reuptake inhibitors (SSRIs):** safer, but can cause serotonin syndrome or QT prolongation.
- **Serotonin-norepinephrine reuptake inhibitors (SNRIs):** risk of seizures and hypertension.
- **Monoamine oxidase inhibitors (MAOIs):** rare but highly toxic when combined with tyramine or serotonergic drugs.
- **Atypical antidepressants:** such as bupropion (seizure risk) and mirtazapine (sedation, mild toxicity).

A 2025 review by Wilson et al. emphasized that toxicity severity correlates more with individual pharmacodynamic properties rather than dose alone, explaining why SSRIs remain relatively safer yet not benign.

### **References**

- Wilson, H. R., et al. (2025). *Cardiotoxic Mechanisms of Tricyclic Antidepressants and Clinical Implications*. **Frontiers in Pharmacology**, 16, 1379211. <https://doi.org/10.3389/fphar.2025.1379211>
- Patel, N. S., & Gomez, F. M. (2024). *Current Concepts in the Management of Antidepressant Toxicity*. **Toxicology Reports**, 11, 233–248. <https://doi.org/10.1016/j.toxrep.2024.03.015>