

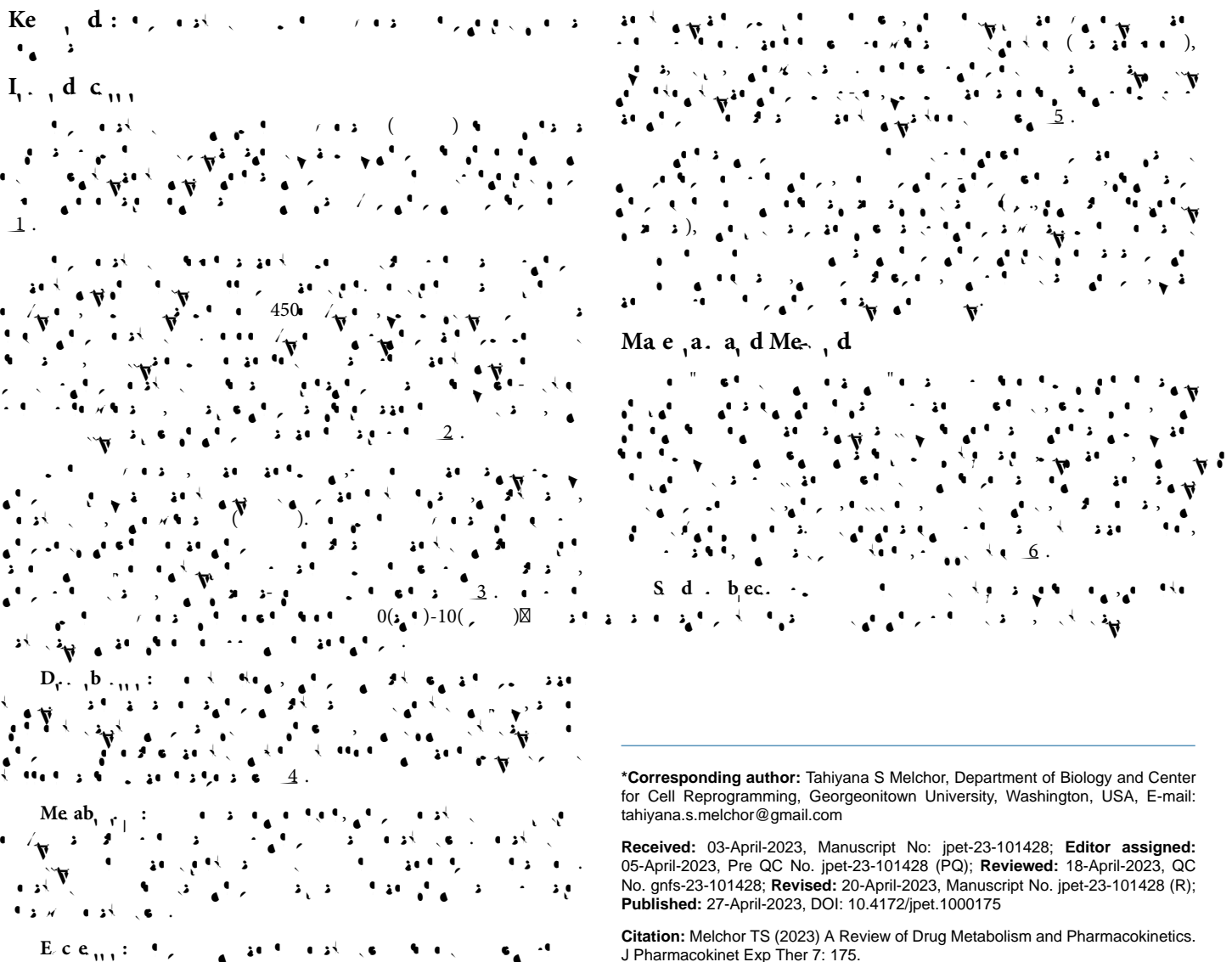
## A Review of Drug Metabolism and Pharmacokinetics

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### Abstract

Drug Metabolism and Pharmacokinetics (DMPK) is a field of study focused on understanding how drugs are processed and eliminated by the body. It encompasses drug metabolism, which involves the enzymatic conversion of drugs into metabolites, primarily in the liver. The metabolites can be inactive, active, or potentially toxic. Pharmacokinetics deals with the movement of drugs within the body, including their absorption, distribution, metabolism, and excretion (ADME). These processes influence the concentration of drugs in the bloodstream over time and are influenced by various factors such as drug formulation, route of administration, and patient-specific characteristics. Understanding DMPK is crucial for the development of safe and effective medications, dosage determination, assessment of drug-drug interactions, and evaluation of drug toxicity. Pharmaceutical companies utilize DMPK data during drug development to optimize drug design and dosing strategies, ensuring the safety and efficacy of medications.



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- (2019) In vitro glucuronidation of designer benzodiazepines by human UDP-glucuronyltransferases. *Drug Test Anal* 11:45-50.
7. Fountain NB, Krauss G, Isojarvi J, Dilley D, Doty P, et al. (2013) Safety and tolerability of adjunctive lacosamide intravenous loading dose in lacosamide-naive patients with partial-onset seizures. *Epilepsia* 54:58-65.
8. Cawello W, Boekens H, Bonn R (2012) Absorption, disposition, metabolic fate and elimination of the anti-epileptic drug lacosamide in humans: mass balance following intravenous and oral administration. *Eur J Drug Metab Pharmacokinet* 37:241-8.
9. Zhou X, Zhao Y, Wang J, Wang X, Chen C, et al. (2018) Resveratrol represses estrogen-induced mammary carcinogenesis through NRF2-UGT1A8-estrogen metabolic axis activation. *Biochem Pharmacol* 155:252-263.
10. Wu L, Chen Y, Liu H, Zhan Z, Liang Z, et al. (2018) Emodin-induced hepatotoxicity was exacerbated by probenecid through inhibiting UGTs and MRP2