

International Journal of Advanced Research in Science, Communication and Technology (IJARSCT)

International Open-Access, Double-Blind, Peer-Reviewed, Refereed, Multidisciplinary Online Journal

Volume 3, Issue 3, December 2023

Review On Prodrug : An Advance Approach for The Drug Design to Enhance the Therapeutic Efficacy

Abhishek V. Kadam¹, Prof. Nikita Bajad², Dr. Swati P. Deshmukh³, Govind S. Nirgunkar⁴, Vishal Singh R Thakur⁵.

Students, Shraddha Institute of Pharmacy, Washim, India^{1,4,5} Asst. Professor, Shraddha Institute of Pharmacy, Washim, India² Professor, Shraddha Institute of Pharmacy, Washim, India³

Abstract: Prodrugs are derivatives of drug molecules that are pharmacologically inactive but require either chemical or enzymatic transformation to release the active drug in vivo in order to exert a pharmacological effect. Prodrugs have better delivery properties that surpass the parent drug molecule. Prodrug concept is justified because it enables the active drug to overcome the barrier that would impede it from reaching the site of action to exert the required pharmacological activity. Some of the barriers that the prodrug approach helps to surmount are as follows, low bioavailability due to poor aqueous solubility (corticosteroids); poor permeability or absorption (ampicillin); high first pass metabolism (propranolol); metabolic instability leading to short half- life, (dopamine); poorsite specificity (anticancer agents); incomplete absorption (epinephrine); unfavorable organoleptic properties (chloramphenicol); difficulties during formulation and adverse effects and toxicity. The prodrug approach is rapidly becoming a crucial part in the stratagem of delivery of drugs. The prodrug strategy implementation in the last 20 y has led to a steady advancement in the biopharmaceutical, physicochemical and/or pharmacokinetic attributes of the pharmacologically active compounds.

Keywords: Prodrugs

REFERENCES

- [1]. Jarkko R, Krista L, Mikko G, Savolainen J. (Prodrugs: design and clinical applications). AAPS, 2008; 10(1): 92-102.
- [2]. Goodman DW. (Lisdexamfetamine dimesylate: the first prodrug stimulant). Psychiatry (Edgmont), 2007; 4(8): 39-45.
- [3]. Rautio J, Kumpulainen H, Heimbach T, et al. (Prodrugs: design and clinical applications). Nat Rev Drug Discov, 2008; 7(3): 255-70.
- [4]. Stella VJ, Charman WNA, Naringer VH. (Prodrugs Do they have advantages in clinical practice?). Drugs, 1985; 29: 455-73.
- [5]. Hsieh PW, Hung CF, Fang JY. (Current prodrug design for drug discovery). Curr Pharm Des, 2009; 15(19): 2236-50.
- [6]. Wagstaff AJ, Ibbotson T, Goa KL. (Capecitabine: a review of its pharmacology and therapeutic efficacy in the management of advanced breast cancer). Drugs, 2003; 63(2): 217-36.
- [7]. Bhosle D, Bharambe S, Gairola N, Dhaneshwar SS. (Mutual prodrug concept: fundamentals and applications). IJPSR, 2006; 68(3): 286-94.
- [8]. Ohlan S, Nanda S, Jagia M, Pathak DP. (Mutual prodrugs- a swot analysis). IJPSR, 2011; 2(4): 719-29.
- [9]. Manon B, Sharma PD. (Design, synthesis and evaluation of diclofenac-antioxidant mutual prodrugs as safer NSAIDs). Indian J Chem, 2009; 48B: 1279-87.
- [10]. Wright V. (A review of benorylate a new antirheumatic drug). Scand J Rheumatol Suppl 1975; 13: 5-8.

DOI: 10.48175/IJARSCT-14348



IJARSCT



International Journal of Advanced Research in Science, Communication and Technology (IJARSCT)

International Open-Access, Double-Blind, Peer-Reviewed, Refereed, Multidisciplinary Online Journal

Volume 3, Issue 3, December 2023

- [11]. Halen PK, Murumkar PR, Yadav MR, Giridhar R. (Prodrug designing of nsaids). Mini Rev. Med. Chem, 2009; 9(1): 124-39.
- [12]. Nazeruddin GM, Suryawanshi SB. (Synthesis of novel mutual pro-drugs by coupling of ibuprofen (nsaid) with sulfa drugs). J Chem Pharm Res, 2010; 2: 508-12.
- [13]. Zawilska JB, Wojcieszak J, Olejniczak AB. (Prodrugs: a challenge for the drug development). Pharmacol Rep, 2013; 65(1): 1–14.
- [14]. Gomes P, Vale N, Moreira R. (Cyclization-activated prodrugs). Molecules, 2007; 12: 2484- 506.
- [15]. Duggal S, Rathore P, Kanwar K. (Prodrug: novel approaches for antiinflammatory action of NSAID's). IJPT, 2012; 4(1): 1889-1908.
- [16]. Waller DG, George CF. (Prodrugs). Br J Clin Pharmacol, 1989; 28(5): 497-507.
- [17]. Verma A, Verma B, Prajapati SK, Tripathi K. (Prodrug as a chemical delivery system: a review). Asian J Research Chem, 2009; 2(2): 100-03.
- [18]. Huttunen KM, Rautio J. (Prodrugs an efficient way to breach delivery and targeting barriers). Curr Top Med Chem, 2011; 11(18): 2265-87.
- [19]. Malik P, Kadam RS, Cheruvu NP, Kompella UB. (Hydrophilic prodrug approach for reduced pigment binding and enhanced transscleral retinal delivery of celecoxib). Mol Pharm, 2012; 9(3): 605-14.
- [20]. Hu L. (Prodrugs: effective solutions for solubility, permeability and targeting challenges). IDrugs, 2004; 7(8): 736-42.
- [21]. Nicoll GDA, Falgueyret JP, Silva JM, et al. (Oxidative bioactivation of the lactol prodrug of a lactone cyclooxygenase-2 inhibitor). Drug Metab Dispos, 1999; 27(3): 403-9.
- [22]. Hu L. The prodrug approach to better targeting. Curr. Drug Discovery, 2004; 28-32.
- [23]. Yan Z, Sun J, Chang Y, et al. (Bifunctional peptidomimetic prodrugs of didanosine for improved intestinal permeability and enhanced acidic stability: synthesis, transpithelial transport, chemical stability and pharmacokinetics). Mol Pharm, 2011; 8(2): 319-29.
- [24]. Bai A, Meier GP, Wang Y, et al. (Prodrug modification increases potassium tricyclo[5.2.1.02,6]-decan-8-yl dithiocarbonate (D609) chemical stability and cytotoxicity against U937 leukemia cells). JPET, 2004; 309(3): 1051-59.

