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## Effect of Common Excipients on Intestinal Drug Absorption in Wistar Rats

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Molecular Pharmaceutics, 2020, 17, 2310-2318.

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6	Physiologically Based Pharmacokinetic/Pharmacodynamic Modeling to Support Waivers of Clinical Studies: Current Status, Challenges, and Opportunities. <i>Molecular Pharmaceutics</i> , <b>2021</b> , 18, 1-17	5.6	3
5	Intestinal membrane transporter-mediated approaches to improve oral drug delivery. <i>Journal of Pharmaceutical Investigation</i> , <b>2021</b> , 51, 137-158	6.3	8
4	A differential equation based modelling approach to predict supersaturation and in vivo absorption from in vitro dissolution-absorption system (idas2) data. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , <b>2021</b> , 165, 1-12	5.7	1
3	An Assessment of Occasional Bio-Inequivalence for BCS1 and BCS3 Drugs: What are the Underlying Reasons?. <i>Journal of Pharmaceutical Sciences</i> , <b>2021</b> ,	3.9	
2	Interaction of Commonly Used Oral Molecular Excipients with P-glycoprotein. <i>AAPS Journal</i> , <b>2021</b> , 23, 106	3.7	1
1	High amount of lecithin facilitates oral delivery of a poorly soluble pyrazoloquinolinone ligand formulated in lipid nanoparticles: physicochemical, structural and pharmacokinetic performances. <b>2023</b> , 122613		1