



Figure 1. The path to systemic oral bioavailability for peptides. The oral dosage form dissolves in the stomach or small intestine depending on whether it is enteric coated. The peptide is liberated and must survive stomach acid if released there, as well as pepsin. If it releases in the upper small intestine, it must survive exposure to pancreatic enzymes. Its permeability across the intestinal epithelium is likely to be of low capacity due to its large molecular weight and hydrophilicity. During the restricted journey across the epithelium it risks exposure to cytochrome P450 and cytosolic peptidases. If it permeates, it must then travel via the hepatic portal vein to the liver where it may undergo the liver first pass effect. If it arrives in the systemic circulation, the intact fraction that has made it from the original liberation from the dosage form is termed bioavailability and this value is expressed relative to an injectable administration route for the same peptide.

Oral Peptide Prodrugs Studies On Stability And Absorption

Katrin Pudersell

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Oral Peptide Prodrugs - Studies on Stability and Absorption Eve-Irene Lepist,2000 **Prodrugs** Valentino Stella, Ronald Borchardt, Michael Hageman, Reza Oliyai, Hans Maag, Jefferson Tilley, 2007-03-12 These volumes represent a comprehensive guide to prodrugs. They guide the reader through the current status of the prodrug concept and its many applications and highlight its many successes in overcoming formulation and delivery of problematic drugs. Replete with examples of approved and marketed prodrugs these volumes introduce the topic to the novice as well as professional in the design of prodrugs.

Prodrugs and Targeted Delivery Jarkko Rautio, 2011-01-31 This topical reference and handbook addresses the chemistry, pharmacology, toxicology and the patentability of prodrugs perfectly mirroring the integrated approach prevalent in today's drug design. It summarizes current experiences and strategies for the rational design of prodrugs beginning at the early stages of the development process as well as discussing organ and site selective prodrugs. Every company employing medicinal chemists will be interested in this practice oriented overview of a key strategy in modern drug discovery and development.

Acetamides—Advances in Research and Application: 2013 Edition, 2013-06-21 Acetamides Advances in Research and Application 2013 Edition is a ScholarlyBrief that delivers timely, authoritative, comprehensive and specialized information about Iodoacetamide in a concise format. The editors have built Acetamides Advances in Research and Application 2013 Edition on the vast information databases of ScholarlyNews. You can expect the information about Iodoacetamide in this book to be deeper than what you can access anywhere else as well as consistently reliable, authoritative, informed and relevant. The content of Acetamides Advances in Research and Application 2013 Edition has been produced by the world's leading scientists, engineers, analysts, research institutions and companies. All of the content is from peer-reviewed sources and all of it is written, assembled and edited by the editors at ScholarlyEditions and available exclusively from us. You now have a source you can cite with authority, confidence and credibility. More information is available at <http://www.ScholarlyEditions.com>

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