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Andreas Bernkop Schnürch

University of Innsbruck, Austria

Oral delivery of biologics-Back to the roots

W Their fast majority, however, is working through the parenteral route being less accepted and inconvenient, as the oral route for administration of biologics emerged to be problematic mostly due to the enzymatic barrier (I), the mucus gel barrier (II) and the absorption barrier (III) of the GI-tract. To overcome these barriers, a huge variety of strategies were established. Among these different strategies, lipophilic emulsifying delivery systems- having already been established more than 30 years ago for the oral administration of the peptide drug cyclosporine- are nowadays attracting more and more academic and industrial research groups, as the number of encouraging *in vivo* data and late stage clinical trials is strongly increasing. Among lipophilic emulsifying delivery systems in particular Self-Emulsifying Drug Delivery Systems (SEDDS) are in focus of research and development. Despite their hydrophilic character, biologics can be incorporated in the lipophilic phase of SEDDS via complexation with lipophilic excipients. Once emulsified in the GI-tract to lipid droplets in the size of 30-200 nm, SEDDS provide a protective effect towards a presystemic metabolism without taking the risk of any side effects. Furthermore, SEDDS can be produced very simply and cost effectively. Because of these properties, they seem to be a promising tool for oral administration of biologics.

Biography

Andreas Bernkop Schnürch completed his MSc in Pharmacy, Microbiology and Genetics at University of Vienna. He completed his Doctoral Degree in 1994. In 2003, he was appointed as a Chair in Pharmaceutical Technology at University of Innsbruck. He invented and pioneered thiolated polymers-thiomers-as a new generation of mucoadhesive polymers. Various medicines based on thiomers have already passed clinical trials and a first product will soon reach the global pharmaceutical market. He is the founder of several biotech companies and author of over 300 research articles and reviews.

Andreas.Bernkop@uibk.ac.at