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Drug interactions of dipeptidyl peptidase 4 inhibitors involving CYP enzymes and P-gp efflux pump

Naina Mohamed Pakkir Maideen

Abstract

Dipeptidyl peptidase 4 (DPP4) inhibitors are oral antidiabetic drugs approved to manage type 2 diabetes mellitus. Saxagliptin is a substrate of CYP3A4/5 enzymes while other DPP4 inhibitors such as Sitagliptin, Linagliptin, Gemigliptin and Teneligliptin are weak substrates of CYP3A4. DPP4 inhibitors have also been identified as the substrates of P-gp. Hence, the drugs inhibiting or inducing CYP3A4/5 enzymes and/or P-gp can alter the pharmacokinetics of DPP4 inhibitors. This review is aimed to identify the drugs

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With the exception of saxagliptin, a substrate of CYP3A4, **DPP-4 inhibitors** seem to be devoid of strong **DDIs involving CYP enzymes**. However, some evidence already exists that inhibition or induction of **drug transporters** (e.g. **P-gp**) can affect the pharmacokinetics of different **DPP-4 inhibitors**. More research in this area is needed.

Cited by: 84

Author: Aleksi Tornio, Mikko Niemi, Pertti J. Neuv...

Publish Year: 2012

Clinical pharmacology of dipeptidyl peptidase 4 inhibitors ...

onlinelibrary.wiley.com/doi/10.1111/1440-1681.12455/full

Dipeptidyl peptidase-4 inhibitors reduce plasma **DPP-4** activity by 70–90% in a sustained manner for 24 h with an increase of **GLP-1** levels (1.5- to 4-fold). They do not pass the blood–brain barrier, have no direct central effect on satiety, and in contrast with **GLP-1** ...

Published in: *Clinical and Experimental Pharmacology and Physiology* · 2015

Authors: Xiao Wu Chen · Zhi Xu He · Zhiwei Zhou · Tianxin Yang · Xueji Zhang · Yin Xue Yang

Affiliation: Southern Medical University · Guiyang Medical University · University of South Florida

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