

Date of report 20 Jun 2019

# Reported case interaction between Cobicistat and Tamsulosin

## **Drugs suspected to be involved in the DDI**

Perpetrator	Daily Dose
Cobicistat	150 (mg)
Dose adjustment performed No	Administration Route Oral
Start date	End date
July 23, 2018	Ongoing
Victim	Daily Dose
<b>Tamsulosin</b>	0.4 (mg)
Idilisulosili	0.4 (mg)
Dose adjustment performed <b>No</b>	Administration Route Oral
Start date	End date
Oct. 15, 2017	Ongoing

## **Complete list of drugs taken by the patient**

Antiretroviral treatment

Darunavir/Cobicistat/Emtricitabine/Tenofovir-AF

Complete list of all comedications taken by the patient, included that involved in the DDI

Tamsulosin, indapamide, enalapril, atorvastatin 20 mg qd, omeprazol, vitamin D

## **Clinical case description**

Gender	Age
Male	<b>72</b>
eGFR (mL/min) >60	Liver function impairment <b>No</b>

#### Description

72 year-old HIV patient on antiretroviral therapy with darunavir/cobicistat/FTC/TAF. Benign prostatic hypertrophy on treatment with tamsulosin 0.4 mg qd. Despite tamsulosin is metabolised mainly by CYP3A4, with potential increase in tansulosine exposure by cobicistat, no adverse events were observed in this patient using of low dose of tamsulosin.

### **Clinical Outcome**

No unwanted outcome

## **Editorial Comment**

The recommendation in the tamsulosin SmPC is that tamsulosin should be used with caution in combination with strong and moderate inhibitors of CYP3A4, and cobicistat is a strong inhibitor. There are data with ketoconazole showing a 2.8-fold increase in exposure. Although this particular case with no unwanted outcomes does not mean you can automatically extrapolate to other patients on tamsulosin, it suggests that the lower dose of 0.4 mg of tamsulosin may be adequate with both ritonavir and cobicistat.

## **University of Liverpool Recommendation**

Potential interaction - may require close monitoring, alteration of drug dosage or timing of administration

For more information <u>click here</u>