

Date of report 26 Nov 2019

# Reported case interaction between Ritonavir and Solifenacin

# Drugs suspected to be involved in the DDI

Perpetrator

**Ritonavir** 

Dose adjustment performed

No

Start date

Jan. 1, 2010

Daily Dose

200 (mg)

Administration Route

Oral

End date

Ongoing

Victim

**Solifenacin** 

Dose adjustment performed

No

Start date

Oct. 24, 2019

Daily Dose

5 (mg)

Administration Route

Oral

End date

Nov. 4, 2019

## Complete list of drugs taken by the patient

Antiretroviral treatment

Darunavir (with Ritonavir or Cobicistat)
Ritonavir
Etravirine
Raltegravir

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

Aciclovir, atorvastatin 10mg QD, Fenofibrate 145 mg QD, ramipril/Hydrochlorothiazide 5/25 mg QD, amlodipine 5mg QD, solifenacin 5 mg QD

# **Clinical case description**

Gender Age
Male 69

eGFR (mL/min) Liver function impairment

>60 No

#### Description

69 year-old male with HIV infection on treatment with darunavir/ritonavir + etravirine + raltegravir (multi-experienced, multi-resistant) since 2010. Prescribed with solifenacin 5 mg QD due to prostatic syndrome since 24/Oct/2019. The patient complains about dry mouth since he started taking solifenacin. On 4/Nov/2019 solifenacin was replaced by tamsulosin 0.4 mg QD, with resolution of symptoms.

### **Clinical Outcome**

## **Toxicity**

### **Drug Interaction Probability Scale (DIPS)**

Score

#### 5 - Probable

#### **Editorial Comment**

Solifenacin is metabolized by CYP3A4 and concentrations are likely to increase due to inhibition of CYP3A4 by ritonavir. It is recommended that solifenacin dosage should be limited to 5 mg once daily if coadministered with a strong CYP3A4 inhibitor such as ritonavir. Even so, close monitoring for adverse events (as in this patient) is recommended.

### **University of Liverpool Recommendation**

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

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