

FDACDER1719v2

609

515 of

Case ID: 8464514

**Case Information:** 

Case Type: EXPEDITED (15- eSub: Y HP: Country: AUS Outcomes: OT, (A)NDA/BLA: 014685/

DAY)

FDA Rcvd Date: 25-Aug-2014 Mfr Rcvd Date: 16-Mar-2012 Mfr Control #: AU-RANBAXY-2012R1-53740

**Patient Information:** 

Age: 35 YR Sex: Female Weight:

Suspect Products: Dose/

# Product Name Frequency Route Dosage Text Indications(s) Start Date End Date

Nortriptyline Hydrochloride
 Valerian
 Valerian
 Unknown
 UNK
 Product used for

unknown indication

# Product Name Dec ReC Lot# Exp Date NDC# MFR/Labeler

NDC# NDC# MFR/Labeler

NDC# NAME

NDC# RANBAXY

2 Valerian A

**Event Information:** 

Preferred Term ( MedDRA 🖄 Version: 17.0 ) ReC

Akathisia

Delirium

Drug interaction

Homicide

Suicidal ideation



FDACDER1720v2

Case ID: 8464514

#### **Event/Problem Narrative:**

This case, manufacturer control number 2012R1-53740 from AUSTRALIA refers to a 35 year old female with the following events (Preferred Terms): Akathisia and Homicide, Suicidal ideation and Delirium .

This case was deemed serious as the reported events were considered to be medically significant.

This case was reported in literature. The literature report consists of 10 case reports and this case refers to Patient of 10.

The patient received following medications which were considered as suspects: #1) Nortriptyline (Company suspect product) 12.5mg/day for psychological distress Therapy start date/stop date: Unknown Action taken: Withdrawn

#2) Valerian for unknown indication Dose and frequency: Unknown Therapy start date/stop date: Unknown

Action taken: Withdrawn

As per the report, eleven patients were identified in a study after developing severe akathisia during treatment with antidepressants for psychological distress and various other drugs [some drug details not stated]; they subsequently became suicidal and/or committed or attempted homicide. All were found to have variant CYP450 alleles. All patients recovered from akathisia on withdrawal of the suspect drugs.

A 35-year-old woman started receiving nortriptyline 12.5 mg/day; she had also been taking small doses of valerian for 1 month. Adding nortriptyline to valerian caused a toxic delirium. She developed severe akathisia and became suicidal, then committed homicide after 3 days.

The outcome of the event 'Akathisia' was reported as 'resolved' and the outcome of the event 'Homicide, Suicidal ideation and Delirium' were reported as 'unknown' at the time of this report.

Case outcome: Unknown

The case is deemed serious. Medical Reviewer considered the case to be possibly related to suspect drug due to its temporal association as per WHO UMC system for standardized causality assessment.

F/U#1: 16-Mar-2012 (Significant)

Print Time: 20-AUG-2015 11:18 AM



FDACDER1721v2

Case ID: 8464514

A follow up was received to the case with following significant informations as:

- # Authors details were updated.
- # Dechallenge for Nortriptyline (company suspect product) was updated to "Positive" from "Unknown"
- # Following event (PT) was added
- 1) Drug interaction with outcome as 'Resolved'
- # Outcome of the events (PT) "Homicide, Suicidal ideation and Delirium" were updated to "Resolved" from "Unknown"

As per follow up report,

Table 3. Information on subject's drug regimens, CYP 450 genotypes, criminal facts and outcomes for ten of those suffering violant akathisia

- # Subject number: (b)
- # Age; 35
- # Gender: Female
- # Drug, dose and treatment duration: Nortriptyline (12.5 mg/day) superimposed on valerian
- # Reason for treatment: Treated for distress due to husband's drinking
- # Genotypes # of null, diminished or ultrarapid metabolizer alleles: 2D6 \*4/\*41, 2C9 \*1/\*2, 2C19 \*1/\*17; One null allele, Two diminished alleles ,One ultrarapid allele
- # Circumstances of crime: Killed (b) (6) daughter in toxic delirium after 3 days
- # Outcome: Pleaded mental illness, misdiagnosed with schizophrenia and treated with more CYP450 substrates. Recovered on withdrawal but more drugs are still being prescribed on Tribunal orders with near fatal consequences

'Subject 1 (35-year-old female; genotypes: CYP2D6 \*4/\*41, CYP2C9 \*1/\*2, and CYP2C19 \*1/\*17) was prescribed a low dose of the 2D6-metabolized tricyclic antidepressant nortriptyline (12.5 mg/day) 3 days prior to committing homicide. She was severely impaired in metabolizing this drug owing to the 2D6 combination of \*4 (null allele) and \*41 (reduced activity allele). This subject also had reduced metabolism through 2C9 (\*1/\*2) and increased (ultrarapid) metabolism through 2C19 (\*1/\*17), although the significance of these genotypes for nortriptyline metabolism is unclear.

Subject <sup>(b)</sup>, in her own words:

Print Time: 20-AUG-2015 11:18 AM



#### FDA - Adverse Event Reporting System (FAERS)

**FOIA Case Report Information** 

FDACDER1722v2

Case ID: 8464514



Nortriptyline is extensively metabolized in the liver, primarily by hydroxylation. There is a strong correlation between total plasma clearance of nortriptyline and 2D6 enzyme activity. Reduced clearance resulting from this subject's 2D6 genotype alone would lead to a build up of serum nortriptyline, which has been shown to be associated with loss of efficacy, toxicity, and increased adversity in several studies. Nortriptyline is also significantly metabolized through 3A4 as an additional pathway, according to in vitro and in vivo data. In the case of severely reduced metabolism through 2D6, as in this subject, 3A4 would have been the major pathway for metabolism. However, the nortriptyline was superimposed on valerian, which has been shown to be a 3A4 inhibitor in vitro and to significantly increase the maximum serum levels of a 3A4 substrate in vivo. The addition of a substrate to an inhibitor is one drug-drug interaction scenario (Pattern 2), as described in Armstrong et al. The addition of nortriptyline to valerian immediately provoked a toxic delirium in this subject; she became akathisic and suicidal and 3 days later she committed the homicide.

Conclusion: A detailed history and mental-state examination in these subjects presented here can exclude functional mental illness (which still has no biological markers) and confirm neurotoxicity. Pharmacogenetic evidence can assist with the rediagnosis of the population that causes the increased demand for mental health services. Restoring them to normality will reduce that demand and associated costs by taking pressure off ambulances, hospitals, prisons, and forensic wards. Akathisia homicides have been defended as instances of involuntary intoxication both with and without genetic evidence. Some perpetrators (and victims) succeed in receiving damages from the manufacturers for failure to warn. The Innocence Project examined evidence from crimes that took place before DNA testing was available. Personalized medicine in health care can bring about corresponding personalized justice in clinics, tribunals, courts and morgues."

Case outcome: Resolved.

Print Time: 20-AUG-2015 11:18 AM

Ranbaxy medical reviewers comment :~



FDACDER1723v2

Case ID: 8464514

The case is deemed serious. Medical Reviewer considered the case to be possibly related to suspect drug due to its temporal association as per WHO UMC system for standardized causality assessment.

Relevant Medical Histor	y:								
Disease/Surgical Procedure			Start Date	End Date	Continuing?				
Medical History Product(s)			Start Date	End Date	Indications	Indications		Events	
Relevant Laboratory Da Test Name	ta:	Result	Unit	Normal Low Ran	ge Norm	al High Range	Info A	vail	
Concomitant Products:	Dose/	Route	Dosage Text	Ir	ndications(s)	Start Date	End Date	Interval 1st	
	Frequency		g		(-,			Dose to Event	
Reporter Source:									
Study Report?: No	Sender Or	ganization: RA	NBAXY						
Print Time: 20-AUG-2015 11:18 AM			If a field is blan	If a field is blank, there is no data for that field				519 of 60	



FDACDER1724v2

Case ID: 8464514

Literature Text:

Lucire Y, Crotty. Antidepressant-induced Akathisia-related homicides associated with diminishing mutations in metabolizing genes of the CYP450 family. Pharmacogenomics and Personalized Medicine. 2011;4(1):65-81