

General Information	
EudraVigilance Local Report Number	EU-EC-10012397734
Sender Type	Regulatory authority
Sender's Organisation	EEA Regulator
Type of Report	Spontaneous
Primary source country	European Economic Area
Reporter's qualification	Healthcare Professional
Case serious?	Yes

Patient		
Age Group	Age Group (as per reporter)	Sex
More than 85 Years		Male

Reaction / Event			
MedDRA LLT	Duration	Outcome	Seriousness ¹
Stroke		Fatal	death., hospital., other
Death	1.0 Days	Fatal	death.
Consciousness loss		Fatal	death., hospital., other

Drug Information					
Role ²	Drug	Duration	Dose	Units in Interval	Action taken
S	VAXZEVRIA - COVID-19 VACCINE ASTRAZENECA (CHADOX1 NCOV-19)				
C	APROVEL 300 MG - IRBESARTAN		1.0 {DF}	Days	
C	EBIXA 10 MG - MEMANTINE HYDROCHLORIDE		1.0 {DF}	Days	
C	- FOLIC ACID		1.0 {DF}	Days	
C	- FUROSEMIDE, FUROSEMIDE SODIUM		1.0 {DF}	Days	
C	- NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE		1.0 {DF}	Days	
C	- RISPERIDONE		1.0 {DF}	Days	
C	- ATORVASTATIN, ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM TRIHYDRATE		1.0 {DF}	Hours	

Drug Information (cont.)				
Info ³	Drug	Indication	Pharm. Form	Route of Admin.
	VAXZEVRIA - COVID-19 VACCINE ASTRAZENECA (CHADOX1 NCOV-19)	N/A		
	APROVEL 300 MG - IRBESARTAN	N/A		
	EBIXA 10 MG - MEMANTINE HYDROCHLORIDE	N/A		
	- FOLIC ACID	N/A		
	- FUROSEMIDE, FUROSEMIDE SODIUM	N/A		
	- NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE	N/A		
	- RISPERIDONE	N/A		
	- ATORVASTATIN, ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM TRIHYDRATE	N/A		

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Seriousness: **death**=results in death; **life threat**=life threatening; **hospital**=requires hospitalization/prolongation of hospitalization; **disability**=results in disability/incapacity; **congen**=congenital anomaly/birth defect; **other**=other medically important information; **(blank)**=non-serious

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Drug role: **S**=suspect; **C**=concomitant; **I**=interacting; **N**=not administered

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Additional Information on Drug: **1**=Counterfeit; **2**= Overdose; **3**=Drug taken by the father; **4**=Drug taken beyond expiry date; **5**=Batch and lot tested and found within specifications; **6**=Batch and lot tested and found not within specifications; **7**=Medication error; **8**=Misuse; **9**=Abuse; **10**=Occupational exposure; **11**=Off label use; **(blank)** =no additional information

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