EVPM ICSR(s)		Individual Case Safety Report Form				EudraVigilance	
General Information							
EudraVigilance Local Report Number		EU-E	C-10016892693				
Sender Type		Health professional					
Sender's Organisation		ASTRAZENECA AB					
Type of Report		Spontaneous					
Primary source country		Non-European Economic Area					
Reporter's qualification		Non-Healthcare Professional					
Case serious?		Yes					
Patient							
Age Group		Age Group (as per reporter)			Sex		
Reaction / Event							
MedDRA LLT		Duration		0	Outcome		Seriousness ¹
Vaccination adverse reaction			Fa		Fatal		death.
Drug Information							
Role ²	Drug		Duration	Dose	U	nits in Interval	Action taken
S	VAXZEVRIA - COVID-19 VACCINE ASTRAZENECA (CHADOX1 NCOV-19))					Unknown
Drug Information (cont.)							
Info ³	Drug	Indication		ı	Pharm. Form		Route of Admin.
	VAXZEVRIA - COVID-19 VACCINE ASTRAZENECA (CHADOX1 NCOV-19)		N/A		Solution for injection		Intramuscular use
Rechallenge matrix table							
Reaction/Event (MedDRA LLT)			Drug			Rechallenge? / Reaction recurred?	
Vaccination	on adverse reaction		/AXZEVRIA - COVID-19 VACCINE ASTRAZENECA CHADOX1 NCOV-19)			no - n/a	

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

² Drug role: S=suspect; C=concomitant; I=interacting; N=not administered

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