Individual Case Safety Rep	oort Form
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General Information									
EudraVig	ilance Local Report Number	EU-E0	EU-EC-10011259316						
Sender T	ype	Health	Health professional						
Sender's	Organisation	PFIZE	PFIZER S.R.L.						
Type of R	Report	Spont	Spontaneous						
Primary s	source country	Non-E	Non-European Economic Area						
Reporter'	s qualification	Health	Healthcare Professional						
Case seri	ious?	Yes	Yes						
Patient									
Age Group			Age Group (as per reporter)			Sex			
	12-17 Years						Female		
Reaction	on / Event								
MedDRA	MedDRA LLT		Duration	Outcome)	Seriousness ¹		
Chills				Fatal			death.		
Itching				Fatal			death.		
Rash generalised				Fatal			death.		
Drug In	formation								
Role ²	Drug		Duration	Dose	ι	Jnits in Interval	Action taken		
S	COMIRNATY - TOZINAMERAN		1.0 Days			Total	Not applicable		
Drug Information (cont.)									
Info ³	Drug		Indication F		Pl	narm. Form	Route of Admin.		
	COMIRNATY - TOZINAMERAN		COVID-19 immunisation				Intramuscular use		

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

 $^{2 \}quad \text{Drug role: } \textbf{S} \text{=-suspect; } \textbf{C} \text{=-concomitant; } \textbf{I} \text{=-interacting; } \textbf{N} \text{=-not administered}$

³ 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