

1. Reporte	r Details			nitial	□ Follow-up	
Reporter N	ame:		E-mail:			
Contact ad	ldress:		Telephone nur	mber:		
			Fax number:			
Type:	\Box Physician (Specialty):		□ Nurse			
	Pharmacist	□ Other Healthcare profe	essional (Please	specify)		
			2			
If reporter i	is a consumer, have they inf	ormed their physician of the	ne exposure?	\Box Yes	🗆 No	
Has the cor	nsumer provided permission	to contact their healthcare	professional?	□ Yes	🗆 No	
If yes, please provide healthcare professional contact details:						
Name: Type:				Telephone:		
Address:				Email:		

2. Patient Details			
Date of birth	Age	Height (cm)	Weight (kg)
(Day/Month/Year)			
	Yrs/mo.		

3.8	3. Suspect Product Details								
	Name	Strength	Dose	Route	Indication	Treatment Start date (day/month/year)	Treatment end date (day/month/year)	Lot	Exp. date
1.									
2.									
3.									

4. Pancreatic Disorder Details						
Symptoms:						
□ Abdominal pain	□ Malaise					
□ Nausea						
□ Jaundice	□ Fever					
□ Weight loss	□ Abdominal bleeding					
□ Diarrhoea	□ Change in colour or consistency of stools					
□ Itching	□ Bloating					
Other relevant symptoms:						
Please provide, the diagnosis of the patient:						
Does the patient have acute pancreatitis?						
\Box Yes \Box No						

Pancreatic effect



Targeted	
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If yes, please specify the severity.						
□ Mild □ Moderate □ Severe.						
Is the pancreatitis associated with any of the following?						
□ Interstitial edema □ Necrosis of pancrea	atic or peripancreatic tissue					
Is the pancreatitis associated with any of the following or	gan failures?					
□ respiratory failure □ cardiovascular failure	□ renal failure					
Is the pancreatitis associated with any of these local comp	olications?					
 Peripancreatic fluid collections Acute necrotic collections 	 Pancreatic pseudocysts Walled-off pancreatic necrosis 					
Other relevant details:						

5. Laboratory tests			
	Date (day/month/year)	Results	Normal Range
Amylase			
Lipase			
Ultrasonogram			
Endoscopic Ultrasound			
Abdominal X-Ray			
CT scan			
Magnetic Resonance			
Cholangiopancreatography			
Endoscopic Retrograde			
Cholangiopancreatography			
Liver function tests			
Other relevant test details:			

6. Medical History						
Patient's concomitant conditions, relevant medical history, known risk factors, relevant tests, laboratory data.						
🗆 Trauma	□ Diabetes					
□ Cholecystitis	□ Cholelithiasis					

Pancreatic effect

Targeted



□ Autoimmune disease	□ Cystic fibrosis
Pancreatitis	□ Hepatitis
□ Drug abuse	□ Colitis
	□ Toxic exposure
□ Drug abuse	□ Alcohol use: Glass/day
Other relevant medical history:	

7. Treatment for Pancreatic Disorders

8. Details of A	dverse Events				
Adverse Event	Start Date (day/month/year)	Stop Date (day/month/year)	Hospitalization	Outcome	Event Causality
			□ Yes	Recovered / Resolved	□ Related
			□ No	\Box Recovered / Resolved with	□ Not Related
			If yes, provide	Sequelae	🗆 Unknown
			dates of	□ Recovering /Resolving	
			hospitalization.	\Box Not Recovered /Not Resolved	
				□ Fatal	
				🗆 Unknown	
			□ Yes	□ Recovered / Resolved	□ Related
			□ No	\square Recovered / Resolved with	□ Not Related
			If yes, provide	Sequelae	□ Unknown
			dates of	□ Recovering /Resolving	
			hospitalization.	\Box Not Recovered /Not Resolved	
				□ Fatal	
				🗆 Unknown	
			\Box Yes	□ Recovered / Resolved	□ Related
			□ No	\square Recovered / Resolved with	□ Not Related
			If yes, provide	Sequelae	□ Unknown
			dates of	□ Recovering /Resolving	
			hospitalization.	□ Not Recovered /Not Resolved	

Pancreatic effect

Targeted



8. Details of Other Adverse Events							
Adverse Event	Start Date (day/month/year)	Stop Date (day/month/year)	Hospitalization	Outcome	Event Causality		
				□ Fatal			
				🗆 Unknown			

9. Concomitant Drugs & Therapies						
Name	Dose	Indication	Treatment start date (day/month/year)	Treatment end date (day/month/year)		

10. Completed By		
Name:	Signature:	Date (day/month/year):