

1. Reporter Details]			Initial		□Follow-up		
Reporter Name:								E-mail:							
Contact address:								Telephone number:							
								Fax number:							
Тур	ype: Physician (Specialty):							☐ Consumer or other non healthcare professional							
☐ Pharmacist								☐ Other (Specify)							
If reporter is a consumer, have they informed their physician of the							of the	ne exposure?			□ No				
Has	Has the consumer provided permission to contact their healthcare								e professional?			□ Yes		□ No	
If y	If yes, please provide healthcare professional contact details:														
Name:				Type:				Teleph			ione:				
Address:			_		En					Email:	nail:				
2. Patient Details															
2. F		te of birth			Age			Heig	ht			Wei	ight		
(Day/Month/Year)				1	-50		cm				kg				
				Yrs/mo.											
3. 0	3. Company Drug Section														
	Name		Strengtl	n Do	ose Route		Indi	dication Tre		atment Treatment		ent Lot		Expiry	
							start date			end date					
									(day/month/year		(day/month/year)				
1.															
2. 3.				 											
٦.															
4. D	4. Details of Adverse Event														
Adv	verse Event	Start Date (day/month/year)	Stop I			ospitaliz	talization		Outcome		e	Event Causality			
		⊠ Yes			☐ Recovered / Resol										
					☐ No If yes, provide dates			Recovered / Resovered / Resove					ot Related nknown		
				hospitalization.				☐ Recovering /Resol			ving		-		
								☐ Not Recovered /No☐ Fatal			ot Resolved				
					□ Unknown										
5 I.	ntaretitial I	una Disassa (l	II D)												
5. Interstitial Lung Disease (ILD) Signs and Symptoms of ILD (include onset date(s) for each sign and symptoms)															
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Chest radiography and/or CT of the chest (attach supporting chest radiography and/or CT of the chest)							
Pulmonary function testing (PFT) (attach suppor	ting PFT results)						
5 () (
Echocardiographic evaluation (attach supporting of	echocardiographic evaluation)						
Editoral and Grapine divariation (unach supporting t	cenocar anographic cramation)						
Other relevant test (include date examination was con	nduated and vaculty of ovamination)						
Other refevant test (include date examination was con	nducted and results of examination)						
6. Concomitant Drugs & Therapies							
7. Medical History							
Patient's concomitant conditions, relevant medical history	, known risk factors, relevant tests, laboratory data.	(Include information on familial disorders, known risk					
factors or conditions that may affect the outcome of the pregnancy e.g. alcohol, smoking, other substance consumption, hypertension, eclampsia, diabetes including							
gestational, infections during pregnancy, environmental or occupational exposure that may pose a risk factor).							
9. Completed By							
Name:	Signature:	Date (day/month/year):					
ivanic.	Signature.	Date (day/month).					