

1. R	Reporter	· Details								Initial		□Foll	ow-up
Reporter Name:					1	E-mail:							
Contact address:						-	Telephone number:						
					I	Fax number:							
Тур	pe: 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?							
Has the consumer provided permission to contact their healthcare professional?													
If y	es, pleas	se provide healtho	are profes	sional co	ntact deta	ails:							
Nar	ne:		7	Гуре:			Telephone:						
Ado	dress:		<u>'</u>						Email:				
2. P	Patient D			•			11 '	1.			XX 7	. 1.	
		Date of birth (Day/Month/Year)		Age		Height				Weight			
(Day/iviolitil/ i car)				Yrs/mo.			cm			kg			
2 (Jamman	y Drug Section											
<i>3.</i> (ompan	Name	Strength	Dose	Route	Indic	ation	Trea	tment	Treatme	ent	Lot	Expiry
	rame strengt		Suchgui	Dose Route		mare	start date			end date		Lot	Expiry
								(day/mo	onth/year)	(day/month/y	ear)		
1.													
2.													
3.													
4. D	Details of	f Adverse Event											
	verse Ev	ent Start Date	Stop D		Hospitaliz	ation		(Outcome	e	Ev	ent Ca	ausality
	(day/month/year		(day/month	(day/month/year)			☐ Recovered / Resolved			☐ Related			
				☐ No If yes, provide dates			of Recovered / Resolved With Sequelae			☐ Not Related			
					yes, proviae pitalization.	aates of			ng /Resolv	ving	∐ U	Inknown	l
						☐ Not Recovered /Not Resolved ☐ Fatal							
							☐ Unknown						
5. Interstitial Lung Disease (ILD)													
Signs and Symptoms of ILD (include onset date(s) for each sign and symptoms)													

Interstitial Lung Disease Targeted Questionnaire



	Result	Date (day/month/year)
Arterial Blood Gas		
Chest X-Ray		
Chest radiography and/or CT of the chest		
Pulmonary Function Tests		
Broncho-Alveolar Lavage		
Microbiology Culture		
Lung Biopsy		
Complete Blood Count		
Serum Biochemistry		
Blood Cultures		
Cytomegalovirus Titer		
Fungal Antigen		
C-Reactive Protein		
K1-6		
Echocardiographic evaluation		

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

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).

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