

1. Reporter	r Det	ails										Initial		□Foll	ow-up	
Reporter Na	ame:							Е	-mail:							
Contact address:							Т	Telephone number:								
								F	ax num	nber:						
Type:	☐ Physician (Specialty):							[☐ Consumer or other non healthcare professional							
	☐ Pharmacist								☐ Other (Specify)							
If reporter is a consumer, have they informed their physician of								of the	exposu	re?		Yes		□ No)	
Has the consumer provided permission to contact their healthcare								ncare pi	rofessio	onal?		☐ Yes		□ No)	
If yes, please provide healthcare professional contact details:																
Name:				Type:					Telephone:							
Address:										Email:						
2. Patient I	Detai	ls														
Date of birth			Age				Height			Weight						
(Day/Month/Year)				V					cm			kg				
Yrs/mo.																
3. Compan	3. Company Drug Section															
_		Strengt	h D	Dose Ro		oute Indica		tion Treat		tment Treatme		ent	Lot	Expiry		
											date	end date				
									(day/mo		onth/year) (day/mor		year)			
1.																
2.																
3.	3.															
4 5 4																
		verse Event Start Date	Ston	Data												
Adverse Ev	ent	(day/month/year)	Stop 3 (day/mon		HOSDIIAHZAHOI			ation		(Outcome		Event Causality			
									☐ Recovered / Resolved ☐ Rela				. 1			
					\Box N		vide	dates of	Recovered / Resolved with Sequelae			☐ Not Related☐ Unknown				
	hospitalization.			,	☐ Recovering /Resolving											
							□ Not Recovered /Not I			ot Resolved						
							☐ Fatal ☐ Unknown									
Site of Infe	ction		l		1				1 - 0.		-					
□ Bone				☐ Genitourinary					☐ Prostate							
□ Blood			☐ Hepatobiliary					☐ Respiratory								
☐ Cardiovascular			☐ HEENT					☐ Skin								
□ CNS			☐ Joint							☐ Othe	r, specify:					
☐ Gastrointestinal			□ Kidney													



5. Were there any complications caused by the infection?								
If yes, please provide details.								
6. Treatment								
Treatment provided for event:								
Action taken with Company Drug in resp	onse to event:							
7. Diagnostic Tests								
-	cation:	Findings:						
	ation:	Findings:						
-								
Radiographic Studies: Yes / No Date:	Location:	Findings:		202				
, ,	Cell count:	Culture:	Staining:	PCR:				
Cytology:								
Other diagnostic test results (e.g. complete b	lood cell count):							
8. Concomitant Drugs & Therapies								
9. 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).								
10. Completed By								
Name:	Signature:		Date (day/month/ye	ar)·				
	2151141410.		Date (day/intolitill ye					