

MINISTERE DE LA SANTE PUBLIQUE

MINISTRY OF PUBLIC HEALTH

**Form for reporting adverse drug reaction(s) likely due to a drug or a health product used by humans**  
To be filled and returned to the provincial drug monitoring centre and to the Department of Pharmacy and Drugs

Fax: 00 237 22 23 39 33

E-mail : [pharmacovigilance\\_cam@yahoo.fr](mailto:pharmacovigilance_cam@yahoo.fr)

PATIENT TREATED :	Name (first 3 letters): _____	If new-born, Product taken: By the patient: <input type="checkbox"/> By breastfeeding: <input type="checkbox"/> By mother during pregnancy: <input type="checkbox"/> (specify the trimester): _____
	Given name (first letter): ____ Sex: _____ Age: _____ Weight: _____ Height: _____	
History / favourable factors: Pregnancy <input type="checkbox"/> Alcoholism <input type="checkbox"/> Hepathopathy <input type="checkbox"/> Allergy <input type="checkbox"/>		
Nephropathy <input type="checkbox"/> Tobacco addiction <input type="checkbox"/> Others (Specify) <input type="checkbox"/> _____		

**SUSPICIOUS MEDICATIONS (Including vaccines, solvents and herbal medicines):**

N°.	Name	Producer	Batch N°	Expiry date	Method of administration	Dosage	INDICATION/ aim of treatment	DURATION OF TREATMENT	
								Start	End
1									
2									

**ASSOCIATED PRODUCTS (Including vaccines, solvents and herbal medicines):**

N°.	Name	Producer	Batch N°	Expiry date	Method of administration	Dosage	INDICATION/ aim of treatment	DURATION OF TREATMENT	
								Start	End
1									
2									
3									

Has one or more products been stopped?: Yes <input type="checkbox"/> No <input type="checkbox"/> No information <input type="checkbox"/> If so, which: _____ Did reaction disappear after stopping? Yes <input type="checkbox"/> No <input type="checkbox"/> No information <input type="checkbox"/>	Was one or more products re-introduced? Yes <input type="checkbox"/> No <input type="checkbox"/> No information <input type="checkbox"/> If so, did reaction reappear, Yes <input type="checkbox"/> No <input type="checkbox"/> No information <input type="checkbox"/>
--	--

**ADVERSE REACTION:**

DATE OF OCCURRENCE	DURATION AND REACTION	SERIOUSNESS	EVOLUTION
		<input type="checkbox"/> Hospitalisation or prolongation of hospitalization	<input type="checkbox"/> Healing without after-effects
		<input type="checkbox"/> Incapacity or permanent invalidity	<input type="checkbox"/> Death induced by reaction
		<input type="checkbox"/> Involvement of vital prognosis	<input type="checkbox"/> Death unrelated to reaction
		<input type="checkbox"/> Death	<input type="checkbox"/> Subject yet to recover
			<input type="checkbox"/> Healing with after-effect
			<input type="checkbox"/> Death contributed to by reaction
			<input type="checkbox"/> Unknown

NATURE AND DESCRIPTION OF ADVERSE REACTION: (continue overleaf if necessary)

**REPORTER:**

Surname and given name: \_\_\_\_\_  
 Medical doctor  Pharmacist  Dentist  Midwife  Nurse  Others (specify) \_\_\_\_\_  
 Speciality (specify) \_\_\_\_\_  
 Address: \_\_\_\_\_  
 Tel: \_\_\_\_\_ Fax: \_\_\_\_\_ E-mail: \_\_\_\_\_  
 Date: \_\_\_\_\_ Signature: \_\_\_\_\_ Stamp: \_\_\_\_\_