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Report № \_\_\_\_ / \_\_\_\_  
 initial  follow up  final  
(Should be filled by  
Pharmacovigilance Department)

## ADVERSE DRUG REACTION REPORT

ALL PROVIDED INFORMATION IS CONFIDENTIAL AND NON-DISCLOSURE WITH THE EXCEPTION  
OF THE CASES STIPULATED BY LAW

### INFORMATION ABOUT REPORTER (person, who reports about ADR)

Name:	Professional belonging:
Place of employment:	Address:
Telephone/Mobile:	E-mail:

### INFORMATION ABOUT PATIENT (CONSUMER)

Initials:	Sex: <input type="checkbox"/> male   <input type="checkbox"/> fem   <input type="checkbox"/> unknown	Weight (кг):	Age:	<input type="checkbox"/> unknown
Liver disease:	<input type="checkbox"/> Yes   <input type="checkbox"/> No   <input type="checkbox"/> unknown	Kidney disease:	<input type="checkbox"/> Yes   <input type="checkbox"/> No   <input type="checkbox"/> unknown	
Allergy:	<input type="checkbox"/> Yes (specify the allergen)   <input type="checkbox"/> No   <input type="checkbox"/> unknown	Pregnancy:	<input type="checkbox"/> Yes Term _____ weeks   <input type="checkbox"/> No   <input type="checkbox"/> unknown	
		Additional information:		

### SUSPECTED MEDICINAL PRODUCT (-S)

Product (trade name, dose, pharmaceutical form)	Batch	Frequency and method of administration	Indication	Date of start	Date of stop

### OTHER MEDICINES

Medicine (trade name, pharmaceutical form, dose, active substance)	Batch	Frequency and method of administration	Indication	Date of start	Date of stop

### INFORMATION ABOUT ADVERSE DRUG REACTION

Detailed description of ADR	<input type="checkbox"/> ADR are continuing	Start date: ( ____/____/____ )	Date of stop: ( ____/____/____ )
Did the ADR disappear after the drug was stopped? <input type="checkbox"/> Yes   <input type="checkbox"/> No   <input type="checkbox"/> Drug was not stopped			
Did the ADR reappear after the drug was reintroduced? <input type="checkbox"/> Yes   <input type="checkbox"/> No   <input type="checkbox"/> Drug was not reintroduced			
Actions taken to treat the ADR: <input type="checkbox"/> Drug withdrawal   <input type="checkbox"/> Dose reducing   <input type="checkbox"/> Co-treatment cessation   <input type="checkbox"/> None   <input type="checkbox"/> Medicinal therapy   <input type="checkbox"/> Non-medicinal therapy <input type="checkbox"/> Other (indicate):			
Treatment of ADR:			
Outcome: <input type="checkbox"/> Recovering without consequences   <input type="checkbox"/> Improvement of state   <input type="checkbox"/> State without changes   <input type="checkbox"/> Death   <input type="checkbox"/> Unknown <input type="checkbox"/> Recovering with the consequences (indicate):			
Criterion of seriousness: <input type="checkbox"/> Death (date ____/____/____ )   <input type="checkbox"/> Life threatening   <input type="checkbox"/> Hospitalization – initial or prolonged   <input type="checkbox"/> Disability <input type="checkbox"/> Congenital malformations   <input type="checkbox"/> Important medical event (indicate):			<input type="checkbox"/> None

Employee name: \_\_\_\_\_ Position: \_\_\_\_\_ Region: \_\_\_\_\_

Information receive date: \_\_\_\_\_ Date sent in company: \_\_\_\_\_ Signature: \_\_\_\_\_