

Adverse Drug Event Reporting Form		Send to: CHEPLAPHARM Arzneimittel GmbH Ziegelhof 24, 17489 Greifswald Phone: 03834 - 451 1329 Email: drugsafety@cheplapharm.com Fax: 03834 - 451 1349				
Privacy notice: As a pharmaceutical company we are legally bound to process and save information on possible side effects of our medicinal products and forward such information to competent authorities. Details can be found at: https://www.cheplapharm.com/datenschutz/ .						
I. PATIENT INFORMATION						
Initials* <div style="border: 1px solid black; width: 40px; height: 20px; margin: 2px;"></div>	Date of Birth* <div style="border: 1px solid black; width: 60px; height: 20px; margin: 2px;"></div>	Age* <div style="border: 1px solid black; width: 40px; height: 20px; margin: 2px;"></div>	Sex* male female	Height: cm Weight: kg	Study no yes:	
First/Middle/Last name Day Month Year						
* Please provide at least one of the above-mentioned identification details in order to be able to report the side effect(s) to competent authorities effectively.						
Relevant medical history, including pre-existing medical conditions:						
II. ADVERSE DRUG EVENT						
Diagnosis:						
				Start Date <div style="border: 1px solid black; width: 60px; height: 20px; margin: 2px;"></div>	End Date <div style="border: 1px solid black; width: 60px; height: 20px; margin: 2px;"></div>	
				Day Month Year	Day Month Year	
Further description of adverse event (symptoms, clinical signs, tests, related laboratory results including dates, affected body parts, therapy programme and treatment progress):						
Seriousness of adverse event (Check all that apply): Initial or prolonged hospitalisation Disability or permanent damage Congenital anomaly/Birth defects Life-threatening Death			Outcome of adverse event: Ongoing Recovering Fully recovered Permanent Unknown Date of death: <div style="border: 1px solid black; width: 60px; height: 20px; margin: 2px;"></div>			
			Day Month Year		Autopsy: yes no	
III. MEDICINAL PRODUCTS (Application prior to adverse event)						
Brand name/ Active substance/ Batch number	Manufacturer	Dosage	Route of application	Dates of use (From/To)	Indication	Causality suspected
I.						yes no
II.						yes no
III.						yes no
IV.						yes no
If you do not suspect a causal relationship between the reported adverse event and the CHEPLAPHARM product, please indicate the possible cause of effect (e.g. primary disease or comorbidity):						
IV. COMMENTS						
V. DATA OF THE REPORTING PERSON						
Name, Surname*:	Function*:	Physician Pharmacist Patient/Relative	Address*:			
Additionally reported to:						
Date, Signature: <div style="border: 1px solid black; width: 60px; height: 20px; margin: 2px;"></div>						
Day Month Year						
	Phone*: Email*:					
* Please provide the name of the reporter and (if the reporter is a health professional) the job title. For further enquiries it is necessary to provide at least one possibility to contact, i.e. phone, email and/or postal address.						