

Confidential

Number:

Report ID:

DESCRIPTION SUSPECTED ADVERSE REACTION			
START DATE ADVERSE REACTION	DAY	MONTH	YEAR
IF USED LESS THAN ONE DAY	HOURS	MINUTES	
Has the adverse event led to one of the following serious situation			
<input type="radio"/> yes <input type="radio"/> no		<input type="checkbox"/> decease <input type="checkbox"/> life threatening <input type="checkbox"/> hospitalization <input type="checkbox"/> congenital abnormality <input type="checkbox"/> permanent disability <input type="checkbox"/> other severe abnormalities	
FILL IN BELOW: COURSE AND SUPPLEMENTARY NOTES			
USED MEDICATION			
Medicine:			
MAH:		Country of origin:	
DATE OF ADMINISTRATION	DAY	MONTH	YEAR
DATA CLIENT			
DATE OF BIRTH	DAY	MONTH	YEAR
SEX	<input type="checkbox"/> MALE	<input type="checkbox"/> FEMALE	
WEIGHT	KG	PREGNANCY?	Yes No
LENGTH	CM		
MEDICAL HISORY			
DATA NOTIFIER			
NAME			
ADDRESS			
POSTAL CODE			
PLACE			
PHONE NO.			
E-MAIL			
Relationship to client			
ANNEXES			
Please add all annexes and possible additional information.			
Please mention the documents you sent along:			

SEND THIS FORM TO:

Orly Pharma BV, St. Jansweg 15, 5928 LW Venlo
Attn. Quality Management
By e-mail: quality@orlypharma.com

Signature
