HSA Health Sciences Authority
-

CONFIDENTIAL

HSA Adverse Watch

SUSPECTED ADVERSE EVENTS Safer Health Products Through Reporting

I-PARTICUI ARS OF PATIENT

Name/init	ials:			NRIC/FIN/Passpor	t no.:					
		Weight:		Ethnic group:	_	iese	Indian	Malay	E E	urasiar
Sex:	Female:	Male	-	5 .	_					
II-DETAI	LS OF ADVERS	SE EVENT (AE)								
			7	Outcome: R					t yet rec	overed
Date of on		/ m m / y y)	L F	atal (Date of	death):		[_] Un	known	
		, ,		Sequelae (any pe		· –	-):
Descriptio	on of AE(s):				es)		known	
					_					
(Pleas	uspected pro	ame if known.	(e.g. PO 5 mg TDS, I	n, including route V 1g stat, for vaccines:	Date started	Date stopped		Indication	(s)	
For vac	cines, please indic	cate batch no.)	IM Dose 1/	/2/3/booster)						
2.										
3.										
		Concomi	tant products (ind	luding complement	ary medicin	es consumed				
1.	_	orvacc	ines administered	l at the same time ar	nd/or 3 mon	ths before)	1	_		
2.										
and photos	s of complementa	ary health products, v	/here applicable.							
III-MAN	AGEMENT OF	ADVERSE EVENT								
	ation (following onsider the reac) the AE): tion to be serious?	Yes Yes	No No	Alread	dy hospitalise	d before /	AE occurred		
<u> </u>				ous (please tick ✓ all						
	ient died due to	reaction	_	ed or prolonged inp			-1e			
_	threatening ngenital anoma	kz		ved persistent or sign cally significant, pleas			-			
		·		any significant, pleas	-					
	of product to ad									
	·	le Possible	Unlikely	Unconfirmed						
IV-PART	ICULARS OF	REPORTER				Name and a	address o	f place of practi	ice	
Name:			Signature:							
			-							
Contact n	0:		Email address:					Ref No. (fo	r official	use)

CONFIDENTIALITY

Any information related to the identities of the reporter and patient will be kept confidential.

WHAT TO REPORT

An Adverse Event (AE) is defined as a reaction which is noxious (harmful) and unintended, and which occurs at doses normally used in man for prophylaxis, diagnosis, or treatment of a disease, or for the modification of a physiological function.

HSA encourages the reporting of all suspected adverse events to health products (including herbal, traditional or alternative remedies). In particular, please report the following:

- 1. All serious adverse events which:
 - a. are life threatening or fatal,
 - b. require inpatient hospitalisation or prolong existing hospitalisation,
 - c. cause persistent incapacity or disability,
 - d. cause birth defect,
 - e. are medically significant.
- 2. All adverse events to recently marketed health products that have been introduced into Singapore in the recent 5 years, regardless of their nature and severity.

Please do not be deterred from reporting because some details are not known. You may send the completed AE Report Form (through your respective hospital pharmacies, if applicable) to the Vigilance and Compliance Branch, Health Products Regulation Group (see below for full address). Additional pages may be attached if required.

SUBMISSION OF FOLLOW-UP REPORTS

Any follow-up information for an AE that has already been reported can be sent to us on another form or via any other modes of reporting. Please indicate that it is a follow-up report. It is very important that follow-up reports are identified and linked to the original report.

HOW TO REPORT



Mail to: Adverse Event Management Unit Vigilance & Compliance Branch Health Products Regulation Group Health Sciences Authority 11 Biopolis Way #11-03 Helios Singapore 138667



Phone: (65) 6866 1111



Email: HSA_productsafety@hsa.gov.sg



Online Reporting: http://www.hsa.gov.sg/adverse-events



Adverse Event Management Unit Vigilance & Compliance Branch Health Products Regulation Group Health Sciences Authority 11 Biopolis Way, #11-03, Helios, Singapore 138667 http://www.hsa.gov.sg