

Deleted: **IMPORTANT – DELIVER IMMEDIATELY**

<b>ASEAN POST-MARKETING ALERT SYSTEM</b>		
<b>1. Level of confidentiality:</b> <input type="checkbox"/> Internal / restricted circulation <input type="checkbox"/> Public domain, specify website: <hr/>	<b>2. Issue:</b> <input type="checkbox"/> Adulteration <input type="checkbox"/> Quality defect <input type="checkbox"/> Safety aspects <input type="checkbox"/> Others, please specify: <hr/>	<b>3. Action:</b> <input type="checkbox"/> Labelling revision <input type="checkbox"/> Product recall <input type="checkbox"/> Product withdrawal <input type="checkbox"/> Registration cancelled <input type="checkbox"/> Registration suspended <input type="checkbox"/> Others, please specify: <hr/>
<b>4. Source / Type:</b>	<input type="checkbox"/> Signal from local ADR reports <input type="checkbox"/> Signal from local studies <input type="checkbox"/> Decision made by other regulatory authorities &/or industry <input type="checkbox"/> Scientific literature / studies <input type="checkbox"/> Others, please specify: <hr/>	
<b>Product information (whenever possible, please provide an image of the product)</b>		
<b>5. Product type:</b>	<input type="checkbox"/> Biologic <input type="checkbox"/> Cosmetic <input type="checkbox"/> Health supplement <input type="checkbox"/> Pharmaceutical <input type="checkbox"/> Traditional medicine <input type="checkbox"/> Others, please specify: <hr/>	
<b>6. Forensic classification in your country:</b>	<input type="checkbox"/> General sales list / Over-the-counter <input type="checkbox"/> Pharmacy only <input type="checkbox"/> Prescription only <input type="checkbox"/> Others, please specify: <hr/>	
<b>7. Brand / Product name (or drug class):</b>	<b>8. Alternative name (e.g. Chinese, Malay):</b>	
<b>9. Active ingredient / Generic name / Full formula (for cosmetics, traditional medicines etc.)*:</b>		
<b>10. Dosage form (if applicable):</b>	<b>11. Strength (if applicable):</b>	
<b>12. Pack size / Presentation:</b>	<b>13. Batch / Lot number:</b>	
<b>14. Expiry date (if applicable):</b>	<b>15. Date manufactured:</b>	
<b>16. Local licensed indication(s) / Intended use:</b>		
<b>17. Countries which the product is exported to:</b>		
<b>Company information</b>		
<b>18. Marketing authorisation holder / Product licence holder / Company responsible for putting product in the market:</b>		
<b>19. Name &amp; country of manufacturer or supplier:</b>		
<b>Details / Action(s) taken</b>		
<b>20. Background (e.g. reasons for recall, defect, ADR):</b>		

<b>21. Actions / proposed actions to be taken</b> (including the level of recall e.g. hospital, retail, consumers; type of recall e.g. temporary or permanent; date of withdrawal or recall):	
<b>Source of information</b>	
<b>22. Country reporting / Issuing authority:</b>	<b>23. Report reference no.:</b>
<b>24. Department / Designation of person issuing the alert:</b>	<b>25. Date of report:</b>
<b>26. Contact person</b>	
<b>Name:</b>	<b>Department / Designation:</b>
<b>Email address:</b>	<b>Telephone no.:</b> <b>Fax no.:</b>

\* Additional information e.g. press statement and photograph of product should be attached separately. Indicate NA whenever the field is not applicable.