

This form is for use within the pharmacy. It should **NOT** be sent to the NPSA.
To report the incident to the NPSA use the eForm which can be found at www.npsa.nhs.uk/eform
= *mandatory fields which must be completed*

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Describe any action which prevented the incident from reaching the patient or minimised the impact on the patient			
<i>If harm occurred, describe the injury (#)</i>			
Describe any apparent contributing factors (see guidance for examples)			
Describe any actions taken to prevent a reoccurrence			
In your view, what were the underlying causes or events which, if rectified, may prevent another incident?			
Has the GP been informed?	Yes <input type="checkbox"/> No <input type="checkbox"/>	Has the PCT/LHB been informed?	Yes <input type="checkbox"/> No <input type="checkbox"/>
Have you contacted your insurers?	Yes <input type="checkbox"/> No <input type="checkbox"/>	If yes, what advice have they given?	

If this is a medication incident, or a medicine contributed to the incident complete the box below

Details of medication involved in the incident		
At what stage of the medication process did the incident occur? (#)	Prescribing <input type="checkbox"/>	Dispensing <input type="checkbox"/> Monitoring therapy <input type="checkbox"/>
	Advice <input type="checkbox"/>	Other (please specify) <input type="checkbox"/>
Enter the medication incident code (#) (see guidance for codes)		
Medication details		
	correct/intended	incorrect (if applicable)
Medication name		
Form		
Dose/strength		
Route		
BNF Classification		
Is the medicine a manufactured special?	Yes <input type="checkbox"/>	No <input type="checkbox"/>
Is the medicine a parallel import?	Yes <input type="checkbox"/>	No <input type="checkbox"/>
Has a copy of the prescription (front & back) been retained?	Yes <input type="checkbox"/>	No <input type="checkbox"/>
Has a photocopy of the dispensing label in situ been made or has the patient returned the original container (if so, then retain)	Yes <input type="checkbox"/>	No <input type="checkbox"/>

If a medical device/medical product caused an incident or a device/product contributed to the incident, complete the box below

Details of medical device involved in the incident

Type of device (#) (e.g. dressing, blood testing kit etc.)

Product name

Model

Serial number

Manufacturer

Batch number

Expiry date

Date manufactured

Any defective equipment should be retained for inspection

Details of staff involved

(use a separate sheet if necessary)

Name

Job title (#)

Staff status (#) (e.g. locum, permanent etc.)

Role in incident (#)

Directly dispensing ☐

Assisting/checker ☐

Informed of incident ☐

Overall responsibility ☐

Name

Job title (#)

Staff status (#) (e.g. locum, permanent etc.)

Role in incident (#)

Directly dispensing ☐

Assisting/checker ☐

Informed of incident ☐

Overall responsibility ☐

Reporter details

Name

Job title (#)

Staff status (#) (e.g. locum, permanent etc.)

Role in incident (#)

Directly dispensing ☐

Assisting/checker ☐

Informed of incident ☐

Overall responsibility ☐

Date reported

Superintendent/Pharmacist in Charge follow up

(investigations, findings and planned actions)

Name		Job title (#)	
Staff status (#) (e.g. locum, permanent etc.)		Date	

Patient follow up	
Has feedback about the incident and the investigation been provided to the patient?	Yes <input type="checkbox"/> No <input type="checkbox"/>
Does the patient want to pursue the matter further? (if yes, explain the complaints procedure to the patient)	Yes <input type="checkbox"/> No <input type="checkbox"/>
What outcome does the patient want?	

Number of extra sheets attached	
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