## Community Pharmacy Medication Safety Incident (Pharmacy Error) Report Form

This form is for use within the pharmacy to record details of medication safety incidents that relate to errors in the pharmacy (i.e. not external errors such as prescribing errors).

You may not have the necessary information to complete all parts of the form. The completed form is for internal use, but relevant parts of the report can be shared with the NHS via your normal reporting route, e.g. via your pharmacy superintendent or the <u>Learn from patient safety events (LFPSE) service</u>.

Pharmacy details							
Pharmacy/Branch name	Branch number (if applicable)						
Reference number from LFPSE report (obtained when completing the LFPSE report)							
Incident details							
Date of incident	Time of incident						
Describe what happened	Give as many details as necessary to enable others to understand the circumstances and be able to learn from the event. State facts only and <u>not</u> opinions.						
Degree of harm to the patient (severity)	Near miss No harm Low Moderate Severe Death						
Did any actions minimise the impact of the incident on the patient? (Please describe)							
If the patient took/used the medicine/medical device, what symptoms did they experience?							
Details of main patient affected	by incident						
Name							
Address							
Telephone number	Date of birth						
Sex	☐ Male   ☐ Female   ☐ Indeterminate   ☐ Unknown						
Ethnicity	☐ White   ☐ Mixed   ☐ Asian or Asian British						
	☐ Black or Black British ☐ Other ☐ Not stated/unknown						
Does the patient have any known/diagnosed impairments or disabilities?	□ Learning disabilities       □ Physical disabilities       □ None known         □ Sensory impairments       □ Other						
Contributing factors							
What were the apparent contributing factors?	<ul> <li>Communication factors (includes verbal, written and non-verbal between individuals, teams, and/or organisations)</li> <li>☐ Education and training factors (e.g. availability of training)</li> <li>☐ Equipment and resources factors (e.g. clear machine displays, poor working order, size, placement, ease of use)</li> <li>☐ Medication factors (where one or more drugs directly contributed to the incident)</li> <li>☐ Organisation and strategic factors (e.g. organisational structure, contractor / agency use, culture)</li> <li>☐ Patient factors (e.g. clinical condition, social / physical / psychological factors, relationships)</li> <li>☐ Task factors (includes work guidelines / procedures / policies, availability of decision making aids)</li> </ul>						



	Team and social factors (includes role definitions, leadership, support, and cultural factors)  Work and environment factors (e.g. poor/excess administration, physical environment, work
	load and hours of work, time pressures)
	Other
Describe any actions planned or	Unknown
Describe any actions planned or taken to prevent a reoccurrence	
In your view, what were the underlying causes or events which, if rectified, may prevent the incident from harming another patient?	
Incident details	
At what stage during the medication process did an actual or potential error occur?	<ul> <li>□ Prescribing</li> <li>□ Preparation of medicines in all locations / dispensing in a pharmacy</li> <li>□ Administration/supply of a medicine from a clinical area</li> <li>□ Monitoring/follow-up of medicine use</li> <li>□ Advice</li> <li>□ Supply or use of over-the-counter (OTC) medicine</li> </ul>
	Other (please specify)
Description of the medication incident Only choose one description.	Adverse drug reaction (when used as intended)  Contra-indication to the use of the medicine in relation to drugs or conditions  Mismatching between patient and medicine  Omitted medicine / ingredient  Patient allergic to treatment  Wrong / omitted / passed expiry date  Wrong / omitted patient information leaflet  Wrong / omitted verbal patient directions  Wrong / transposed / omitted medicine label  Wrong / unclear dose or strength  Wrong drug / medicine  Wrong formulation  Wrong frequency  Wrong method of preparation / supply  Wrong route  Wrong storage  Other  Unknown
Were there other important factors? Multiple choices allowed.	Poor transfer /transcription of information between paper and/or electronic forms Poor communication between care providers (verbal or written) Use of abbreviations(s) of drug name / strength / dose / directions (e.g. MTX, 1 mg, 1 po) Handwritten prescription / chart difficult to read Omitted signature of healthcare practitioner Patient / carer failure to follow instructions Failure of compliance aid / monitored dosage system (MDS) Failure of adequate medicines security (e.g. missing CD)
	Substance misuse (including alcohol)



Details of the correct medicine	Medicines with similar looking or sounding name   Poor labelling and packaging from a commercial manufacturer   Healthcare practitioner undertaking supplementary prescribing   Variance to guidelines for sound clinical reasons   Involving a medicine supplied under a Patient Group Direction (PGD)   Involving an OTC medicine   Failure in monitoring / assessing medicines therapy   Failure of clinical assessment equipment   Other   Unknown   Unknown								
Name of medicine / medical device	e (include brand name if applicabl	e)							
Form		Dose and	strength						
Route		Manufact	urer						
Batch number		Manufact	ured special?	Yes	☐ No				
Is this medicine a parallel import (PI)?		Yes	☐ No						
Details of the incorrect medicin	e / medical device associated v	with this incide	nt (if applicable	•)					
Name of medicine / medical device (include brand name if applicable)		e)							
Form		Dose and	Dose and strength						
Route		Manufact	Manufacturer						
Batch number		Manufact	ured special?	Yes	☐ No				
Is this medicine a parallel import (F	Yes	☐ No							
Staff involved in the incident									
Name of dispenser		Job title							
Staff status (e.g. locum, permanent)									
Name of accuracy-checker		Job title	Job title						
Staff status (e.g. locum, permanent)									
Responsible Pharmacist on duty									
Name of person responsible for completing this report		Job title							
Staff status (e.g. locum, permanen	t)								
Date report completed									
Action required									
Action requested by patient?		☐ No action	Teleph	none call	Letter	_			
Responsible Pharmacist notified?		Yes	☐ No						
Submit report to LFPSE?		Yes	☐ No						

