

MEDICATION PASS OBSERVATION REPORT

Facility  Staff Observed	Station/Wing Time Started	Page of Time Finished Date	* ERROR TYPE CODES 1. Omissions (Drugs ordered but not administered at least once) 2. Unauthorized Drug (Drugs administered without a physician's order) 3. Wrong Dose 4. Wrong Route of Administration 5. Wrong Dosage Form 6. Wrong Drug 7. Wrong Time 8. Mfg. Specs. and/or Professional Standard (below)																
(Rev. 09/06) Form # MP5513 These colors (Teal and Pink) are a trademark of Med-Pass, Inc.	RESIDENT DRUG/DOSE AS PASSED CURRENT DRUGS AS ORDERED		ERROR TYPE * <table border="1" style="width: 100%; border-collapse: collapse; text-align: center;"> <thead> <tr> <th>1</th><th>2</th><th>3</th><th>4</th><th>5</th><th>6</th><th>7</th><th>8</th><th>S/NS</th> </tr> </thead> </table>								1	2	3	4	5	6	7	8	S/NS
											1	2	3	4	5	6	7	8	S/NS
TOTAL NO OF MEDS PASSED	TOTAL NO. OF DRUGS ORDERED ➔				ERROR GRAND TOTALS ➔														
See explanation on reverse for error types and examples of significant and non-significant errors (S/NS)								PAGE ERROR TOTALS ➔											
CUMULATIVE ERROR TOTALS ➔ (This plus previous pages)								➤											
Type 8 - Medication Errors Due to Failure to Follow Manufacturers Specifications or Accepted Professional Standards (check each that was observed) (The pharmacist is responsible for supporting any cited errors of Accepted Professional Standards with contemporaneous literature and/or facility policies.)																			
<input type="checkbox"/> Failure to "shake well" products so labeled. <input type="checkbox"/> Failure to properly mix insulin suspensions (e.g., "rolling") without creating air bubbles. <input type="checkbox"/> Crushing tablets or capsules where manufacturer states "do not crush" [Exceptions - (a) if MD explains in the clinical record why crushing will not adversely affect patient, or; (b) if Facility can provide manufacturer or professional literature to justify why modification of dosage form will not compromise patient care] <input type="checkbox"/> Failure to administer adequate (i.e., 8 oz) fluids with medications when manufacturer so specifies (e.g., with administration of bulk laxatives, NSAIDs or solid/liquid potassium supplements). <input type="checkbox"/> Failure to administer medications with food or antacids when so specified by manufacturer (either before or with medication). This is especially important with NSAID medications. <input type="checkbox"/> Administration of medications immediately before, during or immediately after administration of enteral nutritional formulas (ENFs) without, as a minimum, (1) checking placement of nasogastric or gastrostomy tube (cited under F281); (2) pre and post-flushing of tube with at least 30 ml of preferably warm water (cited under F332/F333). <small>NOTE - if Dilantin (phenytoin) is being given immediately before, during or immediately after ENF, check clinical record to assure patient is not experiencing loss of seizure control or side effects such as sedation.</small> <input type="checkbox"/> Failure to properly administer ophthalmic products wherein either (a) contact of the product with the eye occurs, or; (b) insufficient time (3-5 minutes) is allowed between administration of multiple ophthalmic products. <input type="checkbox"/> Allowing resident to swallow sublingual tablets. <input type="checkbox"/> Failure to properly administer medication via metered dose inhalers (MDIs). Proper administration includes: (a) shaking MDI well; (b) positioning MDI 2 finger widths in front of resident's mouth (or using spacer); (c) having resident exhale first then take a slow, deep breath as MDI is activated; (d) holding breath for a count of 10 after inhalation before slowly exhaling, and; (e) waiting a minute between puffs if multiple puffs are ordered. <small>NOTE - if resident is unable to cooperate with the above procedures (e.g., due to dementia) this should not be considered an error.</small> <input type="checkbox"/> Other (please explain including potential for resident's discomfort or jeopardy to his or her health and safety). _____ <input type="checkbox"/> _____ <input type="checkbox"/> _____ <input type="checkbox"/> _____																			
<small>© 1993 MED-PASS, Inc. Reorder From: MED-PASS 800-438-8884</small>																			
<small>Conclusion/Recommendation(s)</small>																			
CALCULATED ERROR RATE		NUMBER OF ERRORS OBSERVED <small>(Significant + Non Significant)</small>		<input type="text"/>	X 100 =		<input type="text"/> %	Consultant Pharmacist/Observer											

The following numbers represent the error types coded on the front of this form.

ERROR TYPE CODES (1 through 7)

S - SIGNIFICANT	NS - NON SIGNIFICANT										
1. OMISSIONS (Drugs ordered but not administered at least once) Examples: Haldol 1mg BID NS Motrin 400mg TID NS Quinidine 200mg TID S Multivitamin one daily NS Mylanta Suspension one oz. TID AC NS Nitrol Ointment one inch S Tearisol Drops 2 both eyes TID NS Metamucil one packet BID NS	4. WRONG ROUTE OF ADMINISTRATION Examples: Ordered Cortisporin Otic Administered Left eye S Drops 4 to 5 left ear QID										
2. UNAUTHORIZED DRUG (Drugs administered without a physician's order) Examples: Feosol NS Coumadin 4mg S Zyloprim 100mg NS Tylenol 5gr NS Motrin 400mg NS	5. WRONG DOSAGE FORM Examples: Ordered Colace Liquid 100mg BID Administered Capsule NS Mellaril 10mg tablet Concentrate NS* *(If correct dose was given) Dilantin Kapseals ** 100mg three Kapseals p.o. HS Prompt Phenytoin 100mg three capsules p.o. HS ** (Park Davis Kapseals have an extended rate of absorption. Prompt phenytoin capsules do not)										
3. WRONG DOSE Examples: <table border="1"><thead><tr><th>Ordered</th><th>Administered</th></tr></thead><tbody><tr><td>Timoptic 0.25% one drop in the left eye TID</td><td>Three drops in each eye NS</td></tr><tr><td>Digoxin 0.125mg everyday</td><td>0.25mg S</td></tr><tr><td>Amphojel 30cc QID</td><td>15cc NS</td></tr><tr><td>Dilantin 125 SUSP 12cc</td><td>2cc S</td></tr></tbody></table>	Ordered	Administered	Timoptic 0.25% one drop in the left eye TID	Three drops in each eye NS	Digoxin 0.125mg everyday	0.25mg S	Amphojel 30cc QID	15cc NS	Dilantin 125 SUSP 12cc	2cc S	6. WRONG DRUG Examples: Ordered Tums Administered Oscal NS Vibramycin Vancomycin S
Ordered	Administered										
Timoptic 0.25% one drop in the left eye TID	Three drops in each eye NS										
Digoxin 0.125mg everyday	0.25mg S										
Amphojel 30cc QID	15cc NS										
Dilantin 125 SUSP 12cc	2cc S										
	7. WRONG TIME Examples: Ordered Digoxin 0.25mg daily at 8 am Administered At 9:30 am NS Paracetamol 2 tablets 20 minutes before painful treatment 2 tablets given 3hrs after treatment S										

DEFINITION

Medication Error – the preparation or administration of drugs or biologicals which is not in accordance with:
(1) physician's orders;
(2) manufacturer's specifications (not recommendations) regarding the preparation and administration of the drug or biological;
(3) accepted professional standards and principles which apply to professionals providing services. Accepted professional standards and principles include various practice regulations in each State, and current commonly accepted health standards established by national organizations, boards and councils.

Significant Medication Error means one which causes the resident discomfort or jeopardizes his or her health. Criteria for judging significant medication errors as well as examples are provided above.

General Rules for Determining Significance. – The relative significance of medication errors is a matter of professional judgement. Observers who are responsible for assessing these requirements must be qualified to exercise such judgement (e.g., pharmacists, nurses). Follow three general rules in determining whether a medication error is significant or not:

- **RESIDENT CONDITION** – The resident's condition is an important factor to take into consideration. For example, a potent diuretic erroneously administered to a dehydrated resident may have serious consequences but if administered to a resident with a normal fluid balance may not. If the resident's condition requires rigid control, a single missed or wrong dose can be highly significant.
- **DRUG CATEGORY** – If the drug is from a category that usually requires the resident to be titrated to a specific blood level, a single medication error could alter that level and precipitate a reoccurrence of symptoms or toxicity. This is especially important with a drug that has a Narrow Therapeutic Index (NTI) (i.e., a drug in which the therapeutic dose is very close to the toxic dose). Examples of drugs with NTI are as follows: Anticonvulsants: phenytoin (Dilantin), carbamazepine (Tegretol), valproic acid (Depakote); Anticoagulants: warfarin (Coumadin), Antiarrhythmic: digoxin (Lanoxin); Antiasthmatics: theophylline (TheoDur); Antimanic Drugs: lithium salts (Eskalith, Lithobid). Examples of drug categories which require titration of resident blood levels include anticonvulsants, anticoagulants, antiarrhythmic, antianginal, and antiglaucoma agents.
- **FREQUENCY OF ERROR** – If an error is occurring with any frequency, there is more reason to classify the error as significant. For example, if a resident's drug was omitted several times, as verified by reconciling the number of tablets delivered with the number administered, classifying that error as significant would be more in order. This conclusion may be especially valid when taken in concert with the resident's condition and the drug category.