Potential and True Adverse Drug Event Reporting Logic

MEDICATION RELATED PROBLEM (MRP)	ADE / pADE CLASSIFICATION	ACTION / INTERVENTION
 A. No med error / event, but potentiated medical problem A. Drug dosing not adequate for treatment goals (dose, interval, or duration) Treatment not optimal based on current evidence / guidelines Monitoring standards not being followed A. No med error / event, but potentiated entified Med error/event DID NOT readed. Med error/event reached pating harm Med error/event (ADE) Event occurred, resulting in terns harm and requiring hospitaliza G. Event occurred, resulted in per harm / disability H. Event occurred, life-threatening 	 B. Med error/event DID NOT reach patient C. Med error/event reached patient, but no harm D. Med error/event reached patient, monitoring or intervention required to confirm no harm Adverse Drug Event (ADE) E. Event occurred, resulting in temporary harm and requiring intervention F. Event occurred, resulting in temporary harm and requiring hospitalization G. Event occurred, resulted in permanent 	101. DC drug(s)102. Substitute drug(s)103. Add drug(s)104. Change dose/dose interval105. Change duration of tx / qty106. Change PRN to schedule107. Change schedule to PRN108. Order lab / diagnostic test109. Educate patient110. Refer to other service111. Clarify Rx112. Substitute dosage form113. Make appt w/ provider114. Provide Rx compliance box115. Other
 15. Pharmacy / dispensing error 16. Medication overuse or misuse 17. Dose discrepancy between patient use & prescribed therapy 18. Using expired medication Monadherence and Patient Variables 19. Medication underuse / poor adherence	pADE SEVERITY RATING i. Potential for minimal (would require patient self management) or no harm ii. Potential for moderate harm (would require	resulting from the use of a drug medications; thrush due to inhaled corticosteroids; dizziness due to antihypertensives; INR out of range and patient has bleed or clot; patient has allergic reaction due to wrong drug prescribed; drug-drug interaction causes patient to have CNS side effects
 20. Dosage form is not reasonable for patient 21. Inadequate patient self-management of lifestyle and other non-drug variables 22. Patient dissatisfied or refuses treatment, No rational reason given 	 iii. Potential for moderate name (would require healthcare professional intervention or hospitalization to resolve) iii. Potential for severe harm (permanent disability or death) 	Potential Adverse Drug Event (pADE): Examples of pADEs include: duplication of therapy; an event that was identified and avoided omission of therapy; incorrect dose; wrong with appropriate interventions before drug/patient/route/time/ dosage form/technique; drug- affecting the patient interaction; inappropriate medication prescribed for indication indication

Adapted From: (1) From Patient-Centered Primary Care Collaborative (http://www.pcpcc.net/files/medmanagepub.pdf); (2) From NCC MERP

(http://www.nccmerp.org/medErrorCatIndex.html); (3) From Medicare Nursing Home Levels of Harm categories,

http://www.medicare.gov/NHCompare/static/related/incdrawlevelofharm.asp?language=English&version=default Steven Chen, PharmD, University of Southern California School of Pharmacy

28. Other

Miscellaneous

26. Illegible prescription

23. Drug not available in prescribed strength 24. Inadequate refills between scheduled visits

27. No follow-up appointment with PCP

25. Nonformulary / not cost effective drug choice