# **RESEARCH ARTICLES**

# Pharmacists' and Pharmacy Students' Ability to Identify Drug-related Problems Using TIMER (Tool to Improve Medications in the Elderly via Review)

Sarah Snyder Lee, PharmD,<sup>a</sup> Ann K. Schwemm, PharmD,<sup>b</sup> Jeffrey Reist, PharmD,<sup>c,d</sup> Matthew Cantrell, PharmD,<sup>c,e</sup> Michael Andreski, MBA,<sup>c</sup> William R. Doucette, PhD,<sup>c</sup> Elizabeth A. Chrischilles, PhD,<sup>f</sup> and Karen B. Farris, PhD<sup>c</sup>

<sup>a</sup>Kansas University Medical Center <sup>b</sup>University of Michigan Health System <sup>c</sup>University of Iowa College of Pharmacy <sup>d</sup>University of Iowa Hospitals and Clinics <sup>e</sup>Veterans Affairs Medical Center, Iowa City <sup>f</sup>University of Iowa College of Public Health

Submitted February 2, 2008; accepted July 14, 2008; published May 27, 2009.

**Objective.** Determine the effectiveness of TIMER (Tool to Improve Medications in the Elderly via Review) in helping pharmacists and pharmacy students identify drug-related problems during patient medication reviews.

**Methods.** In a randomized, controlled study design, geriatric patient cases were sent to 136 pharmacists and 108 third-year pharmacy students who were asked to identify drug related-problems (DRPs) with and without using TIMER.

**Results.** Pharmacists identified more tool-related DRPs using TIMER (p = 0.027). Pharmacy students identified more tool-related DRPs using TIMER in the first case (p = 0.02), but not in the second. **Conclusion.** TIMER increased the number of DRPs identified by practicing pharmacists and pharmacy students during medication reviews of hypothetical patient cases.

Keywords: medication therapy management, drug-related problems, elderly, community pharmacy

# **INTRODUCTION**

Medicare Part D requires MTM for enrollees who use a high proportion of financial resources in the program.<sup>1</sup> Pharmacists are challenged to find ways to effectively and efficiently provide MTM services, given the high volume of prescriptions that they dispense. While there are MTM plans that assist pharmacists, these services are not always available.<sup>2-3</sup> Community pharmacists, who fill the vast majority of prescriptions, are in a unique position to provide important medication reviews.<sup>4-6</sup> An MTM tool may provide pharmacists with a systematic approach for conducting medication reviews and improve efficiency.

MTM services can be especially valuable to older adults.<sup>7-8</sup> The presence of polypharmacy and age-related physiological changes cause this population to experience more drug-related problems (DRPs).<sup>9</sup> Medication errors

**Corresponding Author:** Karen B, Farris, University of Iowa, S525 Pharmacy, 115 South Grand, Iowa City, IA, 52242. Tel: 319-384-4516. Fax: 319-353-5646. E-mail: karen-farris@uiowa.edu

consist of overuse, underuse, and misuse, and lead to more than 5% of hospital admissions in older adults.<sup>9-11</sup> Between 14% and 23% of older adults receive a medication that should not have been prescribed for them,<sup>12-14</sup> and 10% to 25% of patients have an adverse drug reaction or adverse drug event.<sup>15,16</sup> However, some medications, such as statins and angiotensin-converting enzyme (ACE) inhibitors, are under prescribed among older adults.<sup>17-21</sup>

In order to simplify medication reviews, screening tools have been developed. Available tools include Beer's list, Assessing Care of Vulnerable Elders guidelines, and the Medication Appropriateness Index.<sup>22-28</sup> These tools can rarely accommodate patients with multiple chronic diseases, multiple drug interactions, and/or organ-system insufficiency that may cause patient-specific medication problems. Medication reviews need to be done in a way that encompasses the whole patient, not just a particular disease, medication, or drug interaction. This complexity makes it nearly impossible to create a tool that is sufficiently specific and sensitive to identify drug therapy problems.<sup>28</sup>

In addition to practicing pharmacists, pharmacy students are trained to provide MTM,<sup>29,30</sup> but there is currently no standardized approach used to provide MTM services or to identify DRPs.<sup>31,32</sup> A tool may make the task of learning how to provide these services easier for pharmacy students and practicing pharmacists who are new to MTM. Having such a tool may also facilitate and improve the efficiency of identifying DRPs.

The objective of this study was to evaluate the Tool to Improve Medications in Elderly via Review (TIMER), a systematic approach to conducting medication reviews and improving pharmacists' ability to identify important drug-related problems among older adults. The specific aim of this 2-part study was to compare the number and types of drug-related problems identified by practicing pharmacists (part 1) and pharmacy students (part 2) with and without using the TIMER.

# **METHODS**

# **Development of the TIMER**

TIMER is a guide for pharmacists and pharmacy students to follow when conducting medication reviews. TIMER was developed by 2 of the authors (KF and EC), with input from a consensus panel of 4 regional experts who reviewed the tool and provided feedback. Using a scale ranging from strongly agree to strongly disagree, reviewers rated each section of the TIMER on whether the content was evidence-based, important, helpful/useful, and understandable. Feedback from reviewers resulted in several improvements to TIMER, for example, including drug-drug interactions based on both prevalence *and* severity rather than just on severity, and reducing the symptom timeframe to several months.

An important assumption made in developing TIMER was that its users have conducted a patient medication history so that a complete medication list is available. TIMER has 4 sections including cost-effective drug selection, adherence, medication safety, and attaining therapeutic goals, and covers the most common medication issues that affect older adults. Specific reference to the 8 DRPs commonly used in practice-based research studies was not included because TIMER was intended to encourage pharmacists to look beyond those DRPs and consider patients' symptomatology and complications among older adults.

Each of the 4 sections includes points to discuss with patients and suggested recommendations if a DRP is found. The section on cost-effective drug selection suggests generic and therapeutic substitution to ensure that patients are getting the most cost-effective medications. The section of adherence gives examples of how to question patients about adherence and provides specific recommendations. A section on medication safety addresses adverse drug effects, screening for symptomology, inappropriate medications, drug interactions, and Beer's criteria medications. When determining whether patients are attaining their therapeutic goals, TIMER contains guidance on cardiovascular risk reduction and complication management. The section on cardiovascular risk management outlines the major risk factors for coronary heart disease and evaluates treatment goals. The section on complication management identifies common complications seen in the geriatric population, how to screen for them, and recommendations for each complication (Appendix 1).

#### **Evaluation of TIMER Via Written Cases**

To evaluate TIMER, a 2-part randomized, controlled study was designed that involved practicing pharmacists and pharmacy students using the tool to assess hypothetical patient cases.

Patient Case 1 was developed by one of the authors (MA) and based on a case taken from IowaTeach, a University resource for faculty members to use in developing teaching activities. Both clinical and hypothetical patient cases are available in IowaTeach and many contain additional instructional materials such as test questions and teaching notes. Case 1 was an older adult presenting to a community pharmacist for MTM services. Two expert clinicians reviewed Case 1 and identified 13 DRPs, 6 of which were tool-related and 7 that were non-tool-related (Table 1). Tool-related DRPs were those covered in TIMER and non-tool-related DRPs were problems not included in TIMER. The non-tool-related DRPs identified by the experts were not eliminated from the case but were not expected to be affected because TIMER did not contain them.

Cases 2 and 3 were written by 2 of the authors (MC and JR). Both cases involved older adults presenting to pharmacists for MTM services. The authors identified 4 non-tool-related DRPs and 5 tool-related DRPs in their respective case, and this list later served as the key when coding DRPs (Table 1). Case 1 from the pharmacists study served as the basis for cases 2 and 3. The DRPs from case 1 were incorporated into cases 2 and 3, with substitution of medications. For example, warfarin interactions in cases 1 and 2 involved levothryoxin, while warfarin interactions in case 3 were caused by bismuth subsalicylate. Another example of modifications was the wrong drug used in the cases. In cases 1 and 2, the wrong drug was proposyphene and in case 3 it was diphenhydrame: both are Beer's list drugs.

Table 1. Drug-related Problems (DRPs) in Three Cases Evaluating TIMER

DRPs for Case 1 (Part 1):

- Dose too low: LDL not at goal with current dose of Lipitor (T)
- Wrong drug: proposyphene in elderly (T)
- Needs additional therapy: pain not controlled (T)
- Inappropriate adherence: Norvasc (T)
- Inappropriate adherence: KCl (T)
- Drug interaction: warfarin and levothyroxine (T)
- Possible drug interaction: calcium and levothyroxine based on administration
- Wrong drug: Prilosec for lactose intolerance
- Wrong drug: calcium carbonate vs calcium citrate when taken with PPI
- Needs additional therapy: obesity
- Wrong drug: Norvasc for diabetic hypertension
- Possible additional therapy needed: TSH levels need to be checked
- Dose too high: calcium 1000mg in one dose

DRPs for Case 2 (Part 2):

- Drug interaction: warfarin and levothyroxine (T)
- Inappropriate adherence: verapamil (T)
- Wrong drug: propoxyphene in elderly (T)
- Wrong drug: verapamil for diabetic hypertension (T)
- Dose too low: LDL not at goal with current dose of Lipitor (T)
- Wrong drug: Prilosec for lactose intolerance
- Dose too high: calcium 1000mg in one dose
- Needs additional therapy: obesity

• Possible wrong drug: metformin with reduced renal clearance

DRPs for Case 3 (Part 2):

- Dose too low: warfarin, IRN not at goal
- Adverse drug reaction: citalopram may be causing gastrointestinal side effects
- Adverse drug reaction: terazosin may be causing orthostatic hypotension
- Wrong drug: diphenhydramine in elderly (T)
- Inappropriate adherence: finesteride (T)
- Needs additional therapy: LDL not at goal with pravastatin (T)
- Drug interaction: warfarin and Pepto-Bismol (T)
- Theraputic conversion: generic less costly than Atacand (T)
- Unnecessary drug therapy: saw palmetto not effective

Abbreviations: TIMER = Tool to Improve Medications in the Elderly via Review; (T) = Tool-related DRP. A Tool-related DRP is one that is included in TIMER.

## Part 1: Pharmacists' Use of the TIMER

A randomized controlled study of practicing pharmacists who were members of a regional MTM network (Iowa, Minnesota, Nebraska, North Dakota, South Dakota, Wyoming and Montana) was conducted and half of the pharmacists were randomly selected to receive TIMER. All pharmacists received a printed copy of case 1 and a response form with instructions to identify DRPs in the patient case and write SOAP notes including recommendations. A document of consent to participate was also included in the packet. Participants were asked to return all materials to the investigators and by doing so indicated their informed consent. IRB approval for this study was obtained.

The packet of materials was mailed to 136 pharmacists in mid-April 2007 and a follow-up postcard was sent to non-responders 2 weeks later. A second packet with the same materials was mailed again in mid-June to nonresponders and a follow-up postcard was sent 1 month later. Responses obtained by mid-August 2007 were included in the analysis. Demographic information obtained when the MTM network was formed was linked to the responses in this study. These data included age, practice years, gender, average hours spent in pharmacy per week, and state in which they practiced.

An investigator (AS) coded the pharmacists' responses as either correctly identifying each of the 13 DRPs or not, using a pre-established set of coding rules. Only the DRPs were considered when coding the responses, and not the actions the pharmacist recommended. If a DRP was unclear, the study team reviewed it and consensus was reached. The data were entered into a spreadsheet and analyzed using SPSS. The 2 groups of pharmacists (those who received the TIMER and those who did not) were compared for age, years in practice, and hours worked per week using *t* tests, and for gender using the chi-square test. The numbers of tool-related and non-tool-related DRPs were summed for each respondent. The total number of tool-related and non-toolrelated DRPs identified per study group were compared, controlling for gender and practice years. A chi-square test also was used to compare each tool-related DRP identified with whether the pharmacist had received the TIMER.

## Part 2: Pharmacy Students' Use of the TIMER

In the second part of the study, third-year pharmacy students enrolled in the Pharmacy Practice Laboratory course at University of Iowa College of Pharmacy were asked to identify DRPs in 2 cases, 1 using and 1 not using TIMER. IRB approval for this study was obtained. As seating was randomly assigned at the start of the course, students were already assigned to 54 groups of 2. Students were informed that their answers would not affect their grade for the course. Students reviewed a study information sheet containing all elements of informed consent. Their submission of answers indicated informed consent. For the first 30 minutes of class time, each group of 2 students was assigned 1 of 2 patient cases and asked to identify drug-related problems. For the next 30 minutes, each group was given a second case along with the TIMER and again asked to identify DRPs. Groups of 2 were randomly assigned to receipt of TIMER for 1 of the 2 cases.

The students were asked to provide their age, pharmacy grade point average (GPA), gender, laboratory section, and a unique identifier, which allowed for statistical comparisons to be made later without compromising students' anonymity. Groups of 2 students were asked to list all DRPS that they could find in 30 minutes and state the action that would be taken for each DRP. As the objective was to determine whether TIMER improved students' ability to identify DRPs, only the DRPs were considered during coding and not students' proposed solutions.

Coding of responses was done by an investigator (SL) trained to examine the SOAP notes and identify the presence or absence of the DRPs. Each DRP was coded as either correctly identified or not correctly identified for each group. A set of coding rules for each case was developed by the investigator and reviewed by other investigators before coding was completed. The classification of a correct versus incorrect DRP was based on their written identification of a DRP, not necessarily how the student described it. For example, for the presence of nonadherence to verapamil in case 2, any mention of poor compliance with verapamil was coded as a "yes, the DRP was identified" whether the cause of noncompliance was attributed to side effect, inability to swallow, or patient thinking the drug did not work. Also, if poor compliance with another drug besides verapamil was identified, subjects were not given credit for identifying noncompliance with verapamil. Simply identifying the ADR of verapamil and constipation was not sufficient if compliance or patient education was not mentioned. Proposed actions or recommendations were not used to determine DRPs—the DRP had to be stated. If a DRP was unclear, the study team reviewed it and consensus was reached.

Results were entered into a spreadsheet and analyzed using SPSS. Subjects were divided into 2 study groups: those who received case 2 first and those who received case 3 first. Age, gender, and GPA of the student pharmacists in the 2 study groups were analyzed using chi-square and t tests. The number of tool and non-tool related DRPs identified by student groups were summed across both cases and used as dependent variables. Tool-related DRPs were those that were included in TIMER. The independent variables used in analyses were laboratory section of the student, whether TIMER was used, and patient case. A one-way ANOVA was used to calculate the difference in dependent variables between the 3 Pharmacy Practice Laboratory sections. Then the effect of TIMER and case on total tool-related and total non-tool-related DRPs identified was examined using 2-way ANOVA. We also analyzed the cases separately for significance of TIMER using 2-tailed t tests, as case was significant in the primary analysis. A p value <0.05 was considered significant.

# RESULTS

Eighty-seven of the 136 practicing pharmacists participated in the study. Of these, 41 had been given the TIMER and 46 had not. The average age of participants was  $37.0 \pm 10.6$  years; average time in practice was  $13.0 \pm 11.2$  years; hours worked per week were  $36.3 \pm 11.2$ hours, and 63.2% were female. There were no significant demographic differences between the 2 groups of pharmacists. Practice years and age were correlated (0.965), so age was not included in any further analyses.

The average tool-related DRPs identified by the respondents was  $3.4 \pm 1.0$  by those using the TIMER and  $3.0 \pm 1.0$  for those not using the TIMER (t = -2.26, p = 0.027). The overall model predicting tool-related DRPs using practice years, gender, and TIMER use was significant (F = 6.53, p = 0.001). Gender and TIMER use were significant (p = 0.007 for both), while number of practice years was not significant (p = 0.44). There was no significant difference in the number of DRPs identified

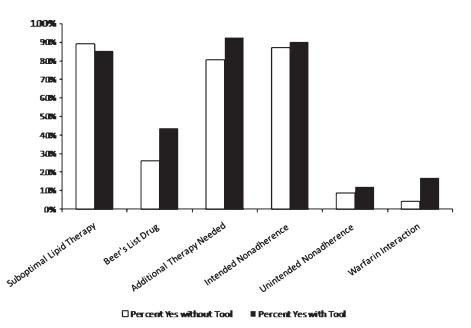


Figure 1. Pharmacists' Identification of Individual Tool-Related DRPs by Use of TIMER (Tool to Improve Medications in the Elderly via Review)

by pharmacists in Iowa compared to those in other states (p = 0.23). Both use of TIMER and practice years were associated with the number of non-tool-related DRPs identified (p = 0.027 and p = 0.030, respectively). When individual tool-related DRPs were analyzed according to use of TIMER, none showed a significant difference (Figure 1).

In part 2 of the study, the average age of the pharmacy students was  $25 \pm 4.2$  years; average GPA was  $3.3 \pm 0.4$ ; and 57.5% were female. There were no significant differences in demographics between the 2 groups of students.

The average number of tool-related DRPs identified by the pharmacy students was  $3.30 \pm 1.05$  using TIMER and  $2.96 \pm 1.13$  not using TIMER (p = 0.11). There was no significant difference in the number of DRPs identified by laboratory section (p = 0.31). The overall 2-way ANOVA model predicting tool-related DRPs using TIMER and case was significant (p < 0.001). Case was significant (p < 0.001), but use of TIMER (p = 0.07) and the interaction were not significant (p = 0.12). When the effect of TIMER for each case was examined separately, the number of tool-related DRPs identified was significantly different for case 2 (p < 0.001, Table 2). Finally, the effect of TIMER on each tool-related DRP was examined. Only the DRP involving Beers' list medications in case 2 showed a significant difference with use of TIMER (Figure 2).

# DISCUSSION

In this randomized study, practicing pharmacists who used TIMER were able to identify more DRPs than practicing pharmacists given the same patient case but not TIMER. TIMER may be helpful to pharmacists because it provides a structured way of reviewing patients' profile and medications. Importantly, it may identify

Table 2. Average Number of Tool-related and Non-tool-related Drug-Related Problems (DRPs) Identified by Pharmacy Students by Case

	TIMER Used?	No.	DRPs Identified, No. (SD)	Р
Case 2 Tool-related DRPs	No	26	3.7 (0.8)	0.85
	Yes	27	3.8 (0.9)	
Case 2 non-Tool-related DRPs	No	26	1.8 (0.9)	0.37
	Yes	27	2.1 (1.2)	
Case 3 Tool-related DRPs	No	27	2.2 (0.0)	0.02
	Yes	26	2.8 (0.9)	
Case 3 non-Tool-related DRPs	No	27	2.5 (0.8)	0.12
	Yes	26	2.9 (0.9)	

Abbreviations: TIMER = Tool to Improve Medication in Elderly via Review

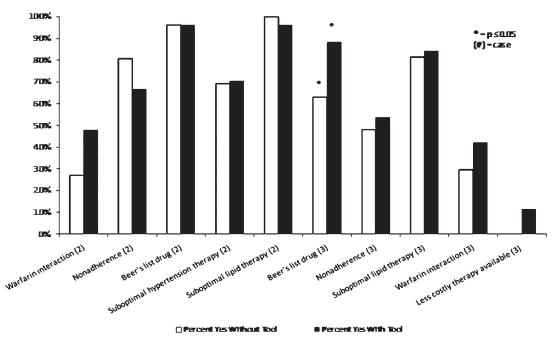


Figure 2. Student Pharmacists' Identification of Individual Tool-Related DRPs by Use of TIMER (Tool to Improve Medications in the Elderly via Review)

DRPs that pharmacists otherwise would not have considered, such as Beer's List medications or new symptoms attributable to adverse drug events. TIMER also increased the number of non-tool-related DRPs identified by pharmacists. TIMER appeared to increase DRPs where pharmacists were less likely to identify problems, namely unintended adherence, warfarin interactions, and Beer's list drugs – although these were identified by less than 50% of pharmacists.

In part 2 of the study, pharmacy students identified nearly half of the tool-related DRPs in the patient cases without using TIMER. This may have occurred because students were given the case to complete at their laboratory workstations where they had a vast number of reference texts and online resources available to them. Students also had uninterrupted time in class to complete the cases. However, pharmacy students appeared less likely to identify warfarin interactions, nonadherence, and less costly therapy when TIMER was not used.

In the analysis for pharmacy students, the use of TIMER was not significant, but analysis by case indicated that TIMER increased the number of DRPs identified in case 2. The differences in the effects of TIMER on the 2 cases were probably due to the differences in the authors of the cases. Although the authors of each case reviewed both cases to ensure they contained similar numbers and types of DRPs, the author of case 1 has been an instructor for 12 years and had written cases for examinations as well as laboratory activities, while the author of case 2 had been an instructor for 1 year. The author having less experience writing cases may have made the DRPs in the first case "easier" for students to find, and having 2 different authors write the cases may be a limitation of this study. Yet, given that each case had a randomized control group for comparison strengthens the findings.

In 2 of the 3 patient cases used in this study, the number of DRPs identified by subjects increased. The fact that 3 different cases were used in this study increases the generalizability of the findings because findings are not limited to 1 set of drug-related problems or 1 type of patient care. TIMER may be more useful in some patients' cases than others, as indicated by the results from pharmacy students. There was no single DRP that seemed more easily identifiable using TIMER.

The content validity of TIMER was established by experts who practice in geriatrics or family medicine. TIMER could be improved by including other DRPs typically identified by pharmacists such as high/low dose, ensuring an existing indication for all medications, and considering whether all indications are treated. However, we sought to include DRPs beyond those traditionally used by pharmacists. There are a number of companies that have created MTM plans or systems that prompt pharmacists when a DRP or problem is detected. These programs prompt pharmacists when an MTM service needs to be performed for a specific reason.<sup>1,2</sup> Typically, these prompts are generated from prescription claims analyses,

# American Journal of Pharmaceutical Education 2009; 73 (3) Article 52.

and we assert that pharmacists need to look at the entire patient when performing MTM. This reasoning was critical to the inclusion of a symptomology assessment in TIMER. It guides pharmacists through a broader review of patients and may serve as an added tool to any of these programs.

In order to use TIMER in practice, pharmacists may require training. We did not provide any training to participants in this study. Training could consist of how to read TIMER, how to apply it to individual patients, and how to examine each patient differently while using it. This might consist of a 30-minute session with an investigator, or a training program in print or online. TIMER also requires regular updates to reflect new guidelines, medications, and evidence.

#### Limitations

There are limitations to this study. Although the cases used were representative of typical cases seen in practice, TIMER has not been tested with actual patients. Recommendations made by pharmacists or pharmacy students were not included, and these are important in resolving DRPs. Also, how TIMER may affect the efficiency of providing an MTM service was not tested. Additionally, 3 different case writers were used, but cases 2 and 3 were based on case 1 and the latter 2 cases were each reviewed by 2 of the investigators. Finally, this study involved pharmacy students from one University and innovative community pharmacists from the upper Midwest and West, thereby limiting its generalizability.

# CONCLUSION

TIMER was effective in increasing the number of toolrelated DRPs found by pharmacists and by pharmacy students in hypothetical patient cases. TIMER may help pharmacists provide required MTM services for older adults.

# ACKNOWLEDGMENT

This study was conducted with a grant from the University of Iowa Older Adults CERT (Centers for Education & Research on Therapeutics) that is funded by the Agency for Healthcare Research and Quality (1 U18 HS016094-01) led by Dr. Elizabeth A. Chrischilles.

At the time of this study, Drs. Lee and Schwemm were fourth- and third-year pharmacy students, respectively, at the University of Iowa College of Pharmacy.

#### REFERENCES

1. Medicare Part D Medication Therapy Management (MTM) Programs 2008 FACT SHEET. Available at: http:// www.cms.hhs.gov/PrescriptionDrugCovContra/Downloads/ MTMFactSheet.pdf. Accessed May 7, 2009. 2. Outcomes Pharmaceutical Health Care, http:// www.getoutcomes.com/aspx/Pharmacists/

currentmtmopportunities.aspx, accessed 14 January 2009.

3. Helping community pharmacies thrive. Mirixa, Reston, Va. http:// www.mirixa.com/pharmacies.aspx Accessed April 30, 2009.

4. McGivney MS, Meyer SM, Duncan-Hewitt W, Hall DL, Goode JV, Smith RB. Medication therapy management: its relationship to patient counseling, disease management, and pharmaceutical care. *J Am Pharm Assoc.* 2007;47:620-8.

5. Hansen RA, Roth MT, Brouwer ES, Herndon S, Christensen DB. Medication therapy management services in North Carolina community pharmacies: current practice patterns and projected demand. *J Am Pharm Assoc.* 2006;46:700-6.

6. Doucette WR, McDonough RP, Klepser D, McCarthy R. Comprehensive medication therapy management: identifying and resolving drug-related issues in a community pharmacy. *Clin Ther.* 2005;27:1104-11.

7. Currie JD, Chrischilles EA, Kuehl AK, Buser RA. Effect of a training program on community pharmacists' detection of and intervention in drug-related problems. *J Am Pharm Assoc*. 1997;37:182-91.

8. Kassam R, Farris KB, Burback L, Volume CI, Cox CE, Cave A. Pharmaceutical care research and education project: pharmacists' interventions. *J Am Pharm Assoc.* 2001;41:401-10.

9. Budnitz DS, Pollock DA, Weidenback KN, Mendelsohn AB, Schroeder TJ, Annest JL. National surveillance of emergency department visits for outpatient adverse drug events. *JAMA*. 2006;296:2858-66.

10. Sullivan SD, Kreling DH, Hazlet TK. Noncompliance with medication regimens and subsequent hospitalizations: a literature analysis and cost of hospitalization estimate. *J Res Pharm Econ.* 1990;219-33.

11. Roughhead EE, Gilbert AL, Primrose JG, et al. Drug-related hospital admission: a review of the Australian studies published 1988-96. *Med J Australia*. 1998;168:405-8.

12. Caterino JM, Emond JA, Carmargo CA. Inappropriate medication administration to the acutely ill elderly: a nationwide emergency department study, 1992-2000. *J Am Geriatric Soc.* 2004;52: 1847-55.

13. Curtis LH, Ostbye T, Sendersky V, et al. Inappropriate prescribing for elderly Americans in a large outpatient population. *Arch Intern Med.* 2004;164:1621-5.

14. Zhan C, Correa-de-Araujo R, Bierman A, et al. Suboptimal prescribing in elderly outpatients: potentially harmful drug-drug and drug-disease combinations. *J Am Geriatric Soc.* 2005;53: 262-7.

15. Gandhi TK, Weingart SN, Borum J, et al. Adverse drug events in ambulatory care. *N Engl J Med.* 2003;348:1556-64.

16. Gurwitz JH, Field TS, Harrold LR, et al. Incidence and preventability of adverse drug events among older people in the ambulatory setting. *JAMA*. 2003;289:1107-16.

17. Higashi T, Shekelle PG, Solomon DH, et al. The quality of pharmacologic care for vulnerable older patients. *Ann Intern Med.* 2004;104:714-20.

18. Raffel OC, White HD. Drug insight: Statin use in the elderly. *Nat Clin Pract Cardiovasc Med.* 2006;3:318-28.

19. Kvan E, Pettersen KI, Landmark K, Reikvam A; INPHARM study investigators. Treatment with statins after acute myocardial infarction in patients >or=80 years: underuse despite general acceptance of drug therapy for secondary

prevention. Pharmacoepidemiol Drug Saf. 2006;15:261-7.

# American Journal of Pharmaceutical Education 2009; 73 (3) Article 52.

20. Tran CT, Laupacis A, Mamdani MM, Tu JV. Effect of age on the use of evidence-based therapies for acute myocardial infarction. *Am Heart J.* 2004;148:834-41.

21. Budnitz DS, Shehab N, Kegler SR, Richards CL. Medication use leading to emergency department visits for adverse drug events in older adults. *Ann Intern Med.* 2007;147:755-65.

22. Beers MH. Explicit criteria for determining potentially inappropriate medication use by the elderly. *Arch Intern Med.* 1997;157:1531-6.

23. Beers MH, Ouslander JG, Rollingher J, Reuben DB, Beck JC. Explicit criteria for determining inappropriate medication use in nursing home residents. *Arch Intern Med.* 1991;151:1825-32.

24. New Facts About Pharmacologic Management in Older Adults, Assessing Care of Vulnerable Elders, Pfizer Inc. Available at: http:// www.rand.org/health/projects/acove/acove3/ Accessed May 7, 2009 25. Hanlon JT, Schmader KE, Samsa GP, et al. A method for assessing drug therapy appropriateness. *J Clin Epidemiol.* 1992;45:1045-51.

26. Fitzgerald LS, Hanlon JT, Shelton PS, et al. Reliability of a modified medication appropriateness index in ambulatory older persons. *Ann Pharmacother*. 1997;31:543-8.

27. Kassam R, Martin LG, Farris KB. Reliability of a modified medication appropriateness index in community pharmacies. *Ann Pharmacother*. 2003;37:40-6.

28. De Smet PA, Denneboom W, Kramers C, Grol R. A composite screening tool for medication reviews of outpatients: general issues with specific examples. *Drugs Aging*. 2007; 24:733-60.

29. Dugan BD. Enhancing community pharmacy through advanced pharmacy practice experiences. *Am J Pharm Educ.* 2006;70:21.

30. Urmie JM, Farris KB, Herbert KE. Pharmacy students' knowledge of the Medicare drug benefit and intention to provide Medicare medication therapy management services. *Am J Pharm Educ.* 2007;71:41.

31. Cameron KA. Comment on medication management models and other pharmacist interventions: implications for policy and practice. *Home Health Care Serv Q.* 2005;24:73-85.

32. American Pharmacists Association. National Association of Chain Drug Stores Foundation. Medication Therapy Management in community pharmacy practice: core elements of an MTM service (version 1.0). *J Am Pharm Assoc.* 2005;45:573-9.



© 2009 University of Iowa

TIMER<sup>©</sup>: <u>Tool to Improve Medications in the Elderly via Review</u>

Confirm all current medications, including Prescription, Herbal remedies, Vitamins and OTCs, and how patients are taking them.

DIOVASCULAR RISK	MANAGEMENT					
	Increasing age (men > CHD in female first de Tobacco smoke/cigare	<b>T DISEASE (CHD)</b> (Exclusive of LDL 45 years; women >55 years), Male sex gree relative <65 years), tte smoking, high blood cholesterol, h us, Low HDL cholesterol (<40 mg/dI	x (gender), Family hi aypertension (BP >1	story of premature CHD		
Note: The Framing	* HDL cholesterol >6 MENT GOAL BY RISK CA ham risk equation, atten	0 mg/dL counts as a "negative" risk f: <b>MTEGORY.</b> Obtain the following medi upts to determine percent risk of a hea alculator.asp?usertype=prof	actor; its presence re cal history and test	results, reviewing accepted	l clinical values and cardiov	
CHD Risk Equi	ralent:: Clinical CHD, S	ymptomatic carotid artery disease, Peri	pheral arterial disea	se, Abdominal aortic aneu	rysm	
Diagnosis	Cardiova	scular Risk Category	Lipids: LDL	<u>Treatme</u> Lipids: Non-HDL (Total – HDL)	e <mark>nt Goal</mark> Blood Pressure	HbAlc
Diabetic	Diabetes is considered a	a CHD risk equivalent	<100 mg/dl		<130/80 mmHg	<7.0 9
Non- Diabetic		e (CHD) and CHD Risk Equivalent risk for CHD >20%)	<100 mg/dl	<130 mg/dl		
	Multiple (2+) Risk Fact (10	ors -year risk ≤20%)	<130 mg/dl	<160 mg/dl	<140/90 mmHg	
	0-1 Risk Factor	r Risk Factor <10%)	<160 mg/dl	<190 mg/dl	_	
MPLICATION MANAG	ement (review potential of	2 omplications)	. Consider need for	erence to current therapy additional or alternative th	Course of Action and/or under-treatment. herapy, and recommend the	erapy to phy
If the patient h MPLICATION MANAG Determine the pres Review common ge	ement (review potential a ence or absence of geriat riatric syndromes and ru	2 omplications) rric syndromes by asking, "Describe he ile out drug-induced causality.	. Consider need for	erence to current therapy additional or alternative tl eling lately?"	and/or under-treatment. herapy, and recommend the	erapy to phy
If the patient h MPLICATION MANAG . Determine the pres . Review common ge Common G Pain* 0-10 Numeric Pain	EMENT (review potential of ence or absence of geriar riatric syndromes and ru riatric Syndromes Mild 1-3/10	2 omplications) tric syndromes by asking, "Describe he	. Consider need for ww you have been fe 1.Clarify 2.Recon	erence to current therapy additional or alternative th eling lately?" Potentia the type of pain the patie umend acetaminophen.	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing.	erapy to phj
If the patient h MPLICATION MANAG Determine the pres Review common ge Common G Pain*	EMENT (review potential of ence or absence of geriar riatric syndromes and ru riatric Syndromes Mild 1-3/10	2 omplications) rric syndromes by asking, "Describe he ile out drug-induced causality.	. Consider need for wy ou have been fe 1.Clarify 2.Recon 3.Nono	erence to current therapy additional or alternative th eling lately?" Potentia the type of pain the patie	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing.	erapy to phy
If the patient h MPLICATION MANAG . Determine the pres . Review common ge Common G Pain* 0-10 Numeric Pain	EMENT (review potential ac ence or absence of geriat riatric syndromes and ru eriatric Syndromes Mild 1-3/10	2 omplications) rric syndromes by asking, "Describe he ile out drug-induced causality.	. Consider need for ww you have been fe 1.Clarify 2.Recon 3.Nonoj Opioi Refer	erence to current therapy additional or alternative th eling lately?" Potentia the type of pain the patie mend acetaminophen. sioid analgesic, fixed dose	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing. *** sic)	erapy to ph
If the patient h MPLICATION MANAG . Determine the pres . Review common ge Common G Pain* 0-10 Numeric Pain	EMENT (review potential a ence or absence of geriat riatric syndromes and ru eriatric Syndromes Mild 1-3/10 Moderate 4-6/10 Severe 7-10/10	2 omplications) rric syndromes by asking, "Describe he ile out drug-induced causality.	. Consider need for w you have been fe 1.Clarify 2.Recon 3.Nonoj Opioi Refer breakt 1.Reduc	erence to current therapy additional or alternative the eling lately?" Potentia the type of pain the patie mend acetaminophen. pioid analgesic, fixed dose d (consider adjunct analge to guideline for 24hr and	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing. *** sic) ****	erapy to ph
If the patient h MPLICATION MANAG . Determine the pres . Review common ge Common G Pain* 0-10 Numeric Pain Intensity Scale Constipation	EMENT (review potential a ence or absence of geriat riatric syndromes and ru eriatric Syndromes Mild 1-3/10 Moderate 4-6/10 Severe 7-10/10	2 amplications) tric syndromes by asking, "Describe he de out drug-induced causality. Drug -Induced Causes Opioids, Acetaminophen-NSAID	. Consider need for wyou have been fe 1.Clarify 2.Recon 3.Nonog Opioi Refer breakt 1.Reduc 2.Recon and/o	erence to current therapy additional or alternative the eling lately?" Potentia the type of pain the patie mend acetaminophen. bioid analgesic, fixed dose d (consider adjunct analge to guideline for 24hr and hrough pain management e drug-induced causes	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing. *** sic) *** lizing osmotic laxatives ose bulk-forming	erapy to phy
If the patient h MPLICATION MANAG Determine the pres Review common ge Common G Pain* 0-10 Numeric Pain Intensity Scale Constipation Determine frequen combinations with	EMENT (review potential of ence or absence of gerial riatric syndromes and re eritatric Syndromes Mild 1-3/10 Moderate 4-6/10 Severe 7-10/10 cy of drugs linergic effects.	2 mplications) tric syndromes by asking, "Describe he le out drug-induced causality. Drug -Induced Causes Opioids, Acetaminophen-NSAID combinations*** Anticholinergics	. Consider need for wyou have been fe 1.Clarify 2.Recon 3.Nonoj Opioi Refer breakt 1.Reduc 2.Recon and/o agents 1.Recon	erence to current therapy additional or alternative th eling lately?" Potentia the type of pain the patie amend acetaminophen. oioid analgesic, fixed dose d (consider adjunct analges to guideline for 24hr and hrough pain management e drug-induced causes amend bowel regiment uti r stool softeners, lower do	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing. *** sic) **** lizing osmotic laxatives see bulk-forming ce. **	erapy to phy
If the patient h MPLICATION MANAG Determine the pres Review common ge Common G Pain* 0-10 Numeric Pain Intensity Scale Constipation Determine frequen combinations with producing anti-cho	EMENT (review potential of ence or absence of gerial riatric syndromes and re eritatric Syndromes Mild 1-3/10 Moderate 4-6/10 Severe 7-10/10 cy of drugs linergic effects.	2 complications) rric syndromes by asking, "Describe he ile out drug-induced causality. Drug -Induced Causes Opioids, Acetaminophen-NSAID combinations*** Anticholinergics Calcium supplementation**	. Consider need for w you have been fe 1. Clarify 2. Recon 3. Nonog Opioi Refer 1. Reduc 2. Recon and/o agents 1. Recon 2. Avoic 1. Recon 2. Avoic 1. Recon	erence to current therapy additional or alternative the eling lately?" Potentia the type of pain the patie mend acetaminophen. bioid analgesic, fixed dose' d (consider adjunct analge to guideline for 24hr and hrough pain management e drug-induced causes mend bowel regiment uti r stool softeners, lower dc with adequate liquid intal mend acetaminophen for I multiple NSAIDs.	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing. *** sic) **** lizing osmotic laxatives see bulk-forming ce. **	ıgly (use 50 need 120
If the patient h MPLICATION MANAG Determine the pres Review common ge Common G Pain* 0-10 Numeric Pain Intensity Scale Constipation Determine frequen combinations with producing anti-cho Muscular sorenes:	EMENT (review potential of ence or absence of gerial riatric syndromes and re eritatric Syndromes Mild 1-3/10 Moderate 4-6/10 Severe 7-10/10 cy of drugs linergic effects.	2 complications) rric syndromes by asking, "Describe he le out drug-induced causality.  Drug -Induced Causes  Opioids, Acetaminophen-NSAID combinations*** Anticholinergies Calcium supplementation**  Statins.	. Consider need for ww you have been fe 1. Clarify 2. Recon 3. Nono Opioi Refer breakt 1. Reduc 2. Recon and/o agents 1. Recon 2. Avoic 1. Recon 2. Recon and/o agents 1. Recon 2. Recon 1. Recon 2. Recon 1. Recon 2. Recon 1. Recon 2. Recon 1. Recon 3. Nono 0. Opioi 0.	erence to current therapy additional or alternative the eling lately?" Potentia the type of pain the patie amend acetaminophen, oioid analgesic, fixed dose d (consider adjunct analgesic) to guideline for 24hr and hrough pain management e drug-induced causes amend bowel regiment uti r stool softeners, lower do with adequate liquid intal amend acetaminophen for I multiple NSAIDS. nament measure bone den ments to prevent bone loor m daily and daily vitamin 1	and/or under-treatment. herapy, and recommend the al Course of Action nt is experiencing. **** sic) **** lizing osmotic laxatives see bulk-forming te, ** 'musculoskeletal pain. sity scan and treat accordin ss and rebuild bone). >age D at age 51-70 400 IU and , based upon review of mbination screening, jmen, vention	ıgly (use 50 need 120

Reference: Core competencies for the care of older patients: recommendations of the American Geriatrics Society. The Education Committee Writing Group of the American Geriatrics Society. Acad Med. 2000 Mar;75(3):252-5. \*Koda-Kimble, Mary Anne, Yee Young, Lloyd, Kradjan , Wayne A., Guglielmo, B. Joseph, Alldredge ,Brian K., Corelli, Robin L., 2005, *Applied Therapeutics: The Clinical Use of Drugs*, 8th Edition, Lippincott Williams & Wilkins, Philadelphia. \*\* Delafuente JC, Stewart RB, eds. Therapeutics in the elderly. 3rd ed. Cincinnati:Harvey Whitney Books Co., 2001. \*\*\* J Am Geriatr Soc. 1998 May;46(5):635-51. The management of chronic pain in older persons: AGS Panel on Chronic Pain in Older Persons. American Geriatrics Society. Page 2